

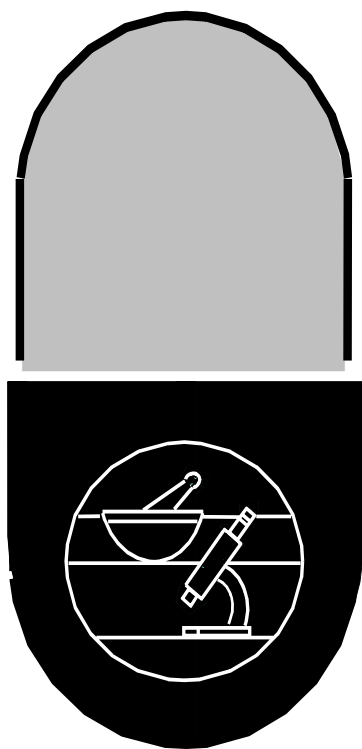
APPROVED DRUG PRODUCTS

With Therapeutic Equivalence Evaluations



The "Orange Book"

FDA data supplied by [DrugPatentWatch.com](https://www.drugpatentwatch.com)



APPROVED DRUG PRODUCTS

WITH

**THERAPEUTIC
EQUIVALENCE
EVALUATIONS**

26th EDITION

**THE PRODUCTS IN THIS LIST HAVE BEEN APPROVED UNDER
SECTION 505 OF THE FEDERAL FOOD, DRUG, AND COSMETIC ACT.**

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF PHARMACEUTICAL SCIENCE
OFFICE OF GENERIC DRUGS

2006

APPROVED DRUG PRODUCTS with THERAPEUTIC EQUIVALENCE EVALUATIONS

The products in this list have been approved under section 505 of the Federal Food, Drug, and Cosmetic Act. This volume is current through December 31, 2005.

26th EDITION



U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF PHARMACEUTICAL SCIENCE
OFFICE OF GENERIC DRUGS

2006

**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
APPROVED DRUG PRODUCTS
with
Therapeutic Equivalence Evaluations**

CONTENTS

	<i>PAGE</i>
PREFACE TO TWENTY SIXTH EDITION	iv
1 INTRODUCTION.....	v
1.1 Content and Exclusion.....	v
1.2 Therapeutic Equivalence-Related Terms	v
1.3 Statistical Criteria for Bioequivalence	vii
1.4 Reference Listed Drug.....	ix
1.5 General Policies and Legal Status	ix
1.6 Practitioner/User Responsibilities.....	x
1.7 Therapeutic Equivalence Evaluations Codes	xii
1.8 Description of Special Situations.....	xix
1.9 Therapeutic Equivalence Code Change for a Drug Entity.....	xxi
1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product	xxi
1.11 Discontinued Section.....	xxi
1.12 Changes to the Orange Book	xxii
1.13 Availability of the Edition.....	xxii
2 HOW TO USE THE DRUG PRODUCTS LISTS	2-1
2.1 Key Sections for Using the Drug Product Lists	2-1
2.2 Drug Product Illustration	2-3
2.3 Therapeutic Equivalence Evaluations Illustration	2-4
 DRUG PRODUCT LISTS	
Prescription Drug Product List	3-1
OTC Drug Product List	4-1
Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research List	5-1
Discontinued Drug Product List	6-1
Orphan Products Designations and Approvals List	7-1
Drug Products Which Must Demonstrate in vivo Bioavailability Only if Product Fails to Achieve Adequate Dissolution	8-1
 APPENDICES	
A. Product Name Index	A-1
B. Product Name Index Listed by Applicant	B-1
C. Uniform Terms	C-1
 PATENT AND EXCLUSIVITY INFORMATION ADDENDUM	 AD1
A. Patent and Exclusivity Lists	ADA1
B. Patent and Exclusivity Terms	ADB1

**FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
APPROVED DRUG PRODUCTS
with
Therapeutic Equivalence Evaluations**

PREFACE TO TWENTY SIXTH EDITION

The publication, *Approved Drug Products with Therapeutic Equivalence Evaluations* (the List, commonly known as the Orange Book), identifies drug products approved on the basis of safety and effectiveness by the Food and Drug Administration (FDA) under the Federal Food, Drug, and Cosmetic Act (the Act). Drugs on the market approved only on the basis of safety (covered by the ongoing Drug Efficacy Study Implementation [DESI] review [e.g., Donnatal® Tablets and Librax® Capsules] or pre-1938 drugs [e.g., Phenobarbital Tablets]) are not included in this publication. The main criterion for the inclusion of any product is that the product is the subject of an application with an effective approval that has not been withdrawn for safety or efficacy reasons. Inclusion of products on the List is independent of any current regulatory action through administrative or judicial means against a drug product. In addition, the List contains therapeutic equivalence evaluations for approved multisource prescription drug products. These evaluations have been prepared to serve as public information and advice to state health agencies, prescribers, and pharmacists to promote public education in the area of drug product selection and to foster containment of health care costs. Therapeutic equivalence evaluations in this publication are not official FDA actions affecting the legal status of products under the Act.

Background of the Publication. To contain drug costs, virtually every state has adopted laws and/or regulations that encourage the substitution of drug products. These state laws generally require either that substitution be limited to drugs on a specific list (the positive formulary approach) or that it be permitted for all drugs except those prohibited by a particular list (the negative formulary approach). Because of the number of requests in the late 1970s for FDA assistance in preparing both positive and negative formularies, it became apparent that FDA could not serve the needs of each state on an individual basis. The Agency also recognized that providing a single list based on common criteria would be preferable to evaluating drug products on the basis of differing definitions and criteria in various state laws. As a result, on May 31, 1978, the Commissioner of the Food and Drug Administration sent a letter to officials of each state stating FDA's intent to provide a list of all prescription drug products that are approved by FDA for safety and effectiveness, along with therapeutic equivalence determinations for multisource prescription products.

The List was distributed as a proposal in January 1979. It included only currently marketed prescription drug products approved by FDA through new drug applications (NDAs) and abbreviated new drug applications (ANDAs) under the provisions of Section 505 of the Act.

The therapeutic equivalence evaluations in the List reflect FDA's application of specific criteria to the multisource prescription drug products on the List approved under Section 505 of the Act. These evaluations are presented in the form of code letters that indicate the basis for the evaluation made. An explanation of the code appears in the *Introduction*.

A complete discussion of the background and basis of FDA's therapeutic equivalence evaluation policy was published in the *Federal Register* on January 12, 1979 (44 FR 2932). The final rule, which includes FDA's responses to the public comments on the proposal, was published in the *Federal Register* on October 31, 1980 (45 FR 72582). The first publication, October 1980, of

the final version of the List incorporated appropriate corrections and additions. Each subsequent edition has included the new approvals and made appropriate changes in data.

On September 24, 1984, the President signed into law the Drug Price Competition and Patent Term Restoration Act (1984 Amendments). The 1984 Amendments require that FDA, among other things, make publicly available a list of approved drug products with monthly supplements. The *Approved Drug Products with Therapeutic Equivalence Evaluations* publication and its monthly Cumulative Supplements satisfy this requirement. The *Addendum* to this publication identifies drugs that qualify under the 1984 Amendments for periods of exclusivity (during which ANDAs or applications described in Section 505(b)(2) of the Act for those drugs may not be submitted for a specified period of time and, if allowed to be submitted, would be tentatively approved) and provides patent information concerning the listed drugs which also may delay the approval of ANDAs or Section 505(b)(2) applications. The *Addendum* also provides additional information that may be helpful to those submitting a new drug application to the Agency.

The Agency intends to use this publication to further its objective of obtaining input and comment on the publication itself and related Agency procedures. Therefore, if you have comments on how the publication can be improved, please send them to the Director, Division of Labeling and Program Support, HFD-610, Office of Generic Drugs, Center for Drug and Evaluation and Research, 7500 Standish Place, Rockville, MD 20855. Comments received are publicly available to the extent allowable under the Freedom of Information regulations.

1. INTRODUCTION

1.1 Content and Exclusion

The List is composed of four parts: (1) approved prescription drug products with therapeutic equivalence evaluations; (2) approved over-the-counter (OTC) drug products for those drugs that may not be marketed without NDAs or ANDAs because they are not covered under existing OTC monographs; (3) drug products with approval under Section 505 of the Act administered by the Center for Biologics Evaluation and Research; and (4) a cumulative list of approved products that have been discontinued from marketing, have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued, have never been marketed, are for exportation, or are for military use.¹

This publication also includes indices of prescription and OTC drug products by trade or established name (if no trade name exists) and by applicant name (holder of the approved application). All established names for active ingredients generally conform to official compendial names or *United States Adopted Names* (USAN) as prescribed in (21 CFR 299.4(e)). The latter list includes applicants' names as abbreviated in this publication; in addition, a list of uniform terms is provided. An *Addendum* contains drug patent and exclusivity information for the Prescription and OTC Drug Product Lists, and for the Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research. The publication may include additional information that the Agency deems appropriate to disseminate.

Prior to the 6th Edition, the publication had excluded OTC drug products and drug products with approval under Section 505 of the Act administered by the Center for Biologics Evaluation and Research because the main purpose of the publication was to provide information to states regarding FDA's recommendation as to which generic prescription drug products were acceptable candidates for drug product selection. The 1984 Amendments required the Agency to begin publishing an up-to-date list of all marketed drug products, OTC as well as prescription, that have been approved for safety and efficacy and for which new drug applications are required.

Under the 1984 Amendments, some drug products were given tentative approvals. Prior to the effective date, the Agency will not include drug products with tentative approval in the List; however, they are available at <http://www.fda.gov/cder/ogd/approvals/default.htm>. When the tentative approval becomes a full approval through a subsequent action letter to the application holder, the Agency will list the drug product and the final, effective approval date in the appropriate approved drug product list.

Distributors or repackagers of products on the List are not identified. Because distributors or repackagers are not required to notify FDA when they shift their sources of supply from one approved manufacturer to another, it is not possible to maintain complete information linking product approval with the distributor or repackager handling the products.

1.2 Therapeutic Equivalence-Related Terms

Pharmaceutical Equivalents. Drug products are considered pharmaceutical equivalents if they contain the same active ingredient(s), are of the same dosage form, route of administration and are identical in strength or concentration (e.g., chlordiazepoxide hydrochloride, 5mg capsules).

¹ Newly approved products are added to parts 1, 2, or 3, of the List, depending on the prescription dispensing requirements (prescription or OTC) or approval authority, unless the Orange Book staff is otherwise notified before publication.

Pharmaceutically equivalent drug products are formulated to contain the same amount of active ingredient in the same dosage form and to meet the same or compendial or other applicable standards (i.e., strength, quality, purity, and identity), but they may differ in characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration time, and, within certain limits, labeling.

Pharmaceutical Alternatives. Drug products are considered pharmaceutical alternatives if they contain the same therapeutic moiety, but are different salts, esters, or complexes of that moiety, or are different dosage forms or strengths (e.g., tetracycline hydrochloride, 250mg capsules vs. tetracycline phosphate complex, 250mg capsules; quinidine sulfate, 200mg tablets vs. quinidine sulfate, 200mg capsules). Data are generally not available for FDA to make the determination of tablet to capsule bioequivalence. Different dosage forms and strengths within a product line by a single manufacturer are thus pharmaceutical alternatives, as are extended-release products when compared with immediate-release or standard-release formulations of the same active ingredient.

Therapeutic Equivalents. Drug products are considered to be therapeutic equivalents only if they are pharmaceutical equivalents and if they can be expected to have the same clinical effect and safety profile when administered to patients under the conditions specified in the labeling.

FDA classifies as therapeutically equivalent those products that meet the following general criteria: (1) they are approved as safe and effective; (2) they are pharmaceutical equivalents in that they (a) contain identical amounts of the same active drug ingredient in the same dosage form and route of administration, and (b) meet compendial or other applicable standards of strength, quality, purity, and identity; (3) they are bioequivalent in that (a) they do not present a known or potential bioequivalence problem, and they meet an acceptable *in vitro* standard, or (b) if they do present such a known or potential problem, they are shown to meet an appropriate bioequivalence standard; (4) they are adequately labeled; (5) they are manufactured in compliance with Current Good Manufacturing Practice regulations. *The concept of therapeutic equivalence, as used to develop the List, applies only to drug products containing the same active ingredient(s) and does not encompass a comparison of different therapeutic agents used for the same condition (e.g., ibuprofen vs. naproxen for the treatment of pain).* Any drug product in the List repackaged and/or distributed by other than the application holder is considered to be therapeutically equivalent to the application holder's drug product even if the application holder's drug product is single source or coded as non-equivalent (e.g., **BN**). Also, distributors or repackagers of an application holder's drug product are considered to have the same code as the application holder. Therapeutic equivalence determinations are not made for unapproved, off-label indications.

FDA considers drug products to be therapeutically equivalent if they meet the criteria outlined above, even though they may differ in certain other characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time and minor aspects of labeling (e.g., the presence of specific pharmacokinetic information) and storage conditions. When such differences are important in the care of a particular patient, it may be appropriate for the prescribing physician to require that a particular brand be dispensed as a medical necessity. With this limitation, however, FDA believes that products classified as therapeutically equivalent can be substituted with the full expectation that the substituted product will produce the same clinical effect and safety profile as the prescribed product.

Bioavailability. This term means the rate and extent to which the active ingredient or active moiety is absorbed from a drug product and becomes available at the site of action. For drug products that are not intended to be absorbed into the bloodstream, bioavailability may be assessed by

measurements intended to reflect the rate and extent to which the active ingredient or active moiety becomes available at the site of action.

Bioequivalent Drug Products. This term describes pharmaceutical equivalent or pharmaceutical alternative products that display comparable bioavailability when studied under similar experimental conditions. Section 505 (j)(7)(B) of the Act describes one set of conditions under which a test and reference listed drug (see Section 1.4) shall be considered bioequivalent:

the rate and extent of absorption of the test drug do not show a significant difference from the rate and extent of absorption of the reference drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses; or

the extent of absorption of the test drug does not show a significant difference from the extent of absorption of the reference drug when administered at the same molar dose of the therapeutic ingredient under similar experimental conditions in either a single dose or multiple doses and the difference from the reference drug in the rate of absorption of the drug is intentional, is reflected in its proposed labeling, is not essential to the attainment of effective body drug concentrations on chronic use, and is considered medically insignificant for the drug.

Where these above methods are not applicable (e.g., for drug products that are not intended to be absorbed into the bloodstream), other *in vivo* or *in vitro* test methods to demonstrate bioequivalence may be appropriate.

Bioequivalence may sometimes be demonstrated using an *in vitro* bioequivalence standard, especially when such an *in vitro* test has been correlated with human *in vivo* bioavailability data. In other situations, bioequivalence may sometimes be demonstrated through comparative clinical trials or pharmacodynamic studies.

1.3 Statistical Criteria for Bioequivalence

Under the Drug Price Competition and Patent Term Restoration Act of 1984, manufacturers seeking approval to market a generic drug product must submit data demonstrating that the drug product is bioequivalent to the pioneer (innovator) drug product. A major premise underlying the 1984 law is that bioequivalent drug products are therapeutically equivalent, and therefore, interchangeable.

Bioavailability refers to the rate and extent to which the active ingredient or therapeutic ingredient is absorbed from a drug product and becomes available at the site of drug action (Federal Food, Drug and Cosmetic Act, section 505(j)(8)). Bioequivalence refers to equivalent release of the same drug substance from two or more drug products or formulations. This leads to an equivalent rate and extent of absorption from these formulations. Underlying the concept of bioequivalence is the thesis that, if a drug product contains a drug substance that is chemically identical and is delivered to the site of action at the same rate and extent as another drug product, then it is equivalent and can be substituted for that drug product. Methods used to define bioequivalence can be found in 21 CFR 320.24, and include (1) pharmacokinetic (PK) studies, (2) pharmacodynamic (PD) studies, (3) comparative clinical trials, and (4) in-vitro studies. The choice of study used is based on the site of action of the drug and the ability of the study design to compare drug delivered to that site by the two products.

The standard bioequivalence (PK) study is conducted using a two-treatment crossover study design in a limited number of volunteers, usually 24 to 36 adults. Alternately, a four-period, replicate design crossover study may also be used. Single doses of the test and reference drug products are administered and blood or plasma levels of the drug are measured over time.

Pharmacokinetic parameters characterizing rate and extent of drug absorption are evaluated statistically. The PK parameters of interest are the resulting area under the plasma concentration-time curve (AUC), calculated to the last measured concentration ($AUC_{(0-t)}$) and extrapolated to infinity ($AUC_{(0-inf)}$), for extent of absorption; and the maximum or peak drug concentrations (C_{max}), for rate of absorption. Crossover studies may not be practical in drugs with a long half-life in the body, and a parallel study design may be used instead. Alternate study methods, such as in-vitro studies or equivalence studies with clinical or pharmacodynamic endpoints, are used for drug products where plasma concentrations are not useful to determine delivery of the drug substance to the site of activity (such as inhalers, nasal sprays and topical products applied to the skin).

The statistical methodology for analyzing these bioequivalence studies is called the two one-sided test procedure. Two situations are tested with this statistical methodology. The first of the two one-sided tests determines whether a generic product (test), when substituted for a brand-name product (reference) is significantly less bioavailable. The second of the two one-sided tests determines whether a brand-name product when substituted for a generic product is significantly less bioavailable. Based on the opinions of FDA medical experts, a difference of greater than 20% for each of the above tests was determined to be significant, and therefore, undesirable for all drug products. Numerically, this is expressed as a limit of test-product average/reference-product average of 80% for the first statistical test and a limit of reference-product average/test-product average of 80% for the second statistical test. By convention, all data is expressed as a ratio of the average response (AUC and C_{max}) for test/reference, so the limit expressed in the second statistical test is 125% (reciprocal of 80%).

For statistical reasons, all data is log-transformed prior to conducting statistical testing. In practice, these statistical tests are carried out using an analysis of variance procedure (ANOVA) and calculating a 90% confidence interval for each pharmacokinetic parameter (C_{max} and AUC). The confidence interval for both pharmacokinetic parameters, AUC and C_{max} , must be entirely within the 80% to 125% boundaries cited above. Because the mean of the study data lies in the center of the 90% confidence interval, the mean of the data is usually close to 100% (a test/reference ratio of 1). Different statistical criteria are sometimes used when bioequivalence is demonstrated through comparative clinical trials pharmacodynamic studies, or comparative in-vitro methodology.

The bioequivalence methodology and criteria described above simultaneously control for both, differences in the average response between test and reference, as well as the precision with which the average response in the population is estimated. This precision depends on the within-subject (normal volunteer or patient) variability in the pharmacokinetic parameters (AUC and C_{max}) of the two products and on the number of subjects in the study. The width of the 90% confidence interval is a reflection in part of the within-subject variability of the test and reference products in the bioequivalence study. A test product with no differences in the average response when compared to the reference might still fail to pass the bioequivalence criteria if the variability of one or both products is high and the bioequivalence study has insufficient statistical power (i.e., insufficient number of subjects). Likewise, a test product with low variability may pass the bioequivalence criteria, when there are somewhat larger differences in the average response.

This system of assessing bioequivalence of generic products assures that these substitutable products do not deviate substantially in in-vivo performance from the reference product. The Office of Generic Drugs has conducted two surveys to quantify the differences between generic and brand name products. The first survey included 224 bioequivalence studies submitted in approved applications during 1985 and 1986. The observed average differences between reference and generic products for AUC was 3.5% (JAMA, Sept. 4, 1987, Vol. 258, No. 9). The second survey included 127 bioequivalence studies submitted to the agency in 273 ANDAs approved in 1997.

The three measures reviewed include $AUC_{(0-t)}$, $AUC_{(0-inf)}$, and C_{max} . The observed average differences between the reference and generic products were $+ 3.47\%$ (SD 2.84) for $AUC_{(0-t)}$, $+ 3.25\%$ (SD 2.97) for $AUC_{(0-inf)}$, and $+ 4.29\%$ (SD 3.72) for C_{max} (JAMA, Dec. 1, 1999, Vol. 282, No. 21).

The primary concern from the regulatory point of view is the protection of the patient against approval of products that are not bioequivalent. The current practice of carrying out two one-sided tests at the 0.05 level of significance ensures that there is no more than a 5% chance that a generic product that is not truly equivalent to the reference will be approved.

1.4 Reference Listed Drug

A reference listed drug (21 CFR 314.94(a)(3)) means the listed drug identified by FDA as the drug product upon which an applicant relies in seeking approval of its ANDA.

FDA has identified in the Prescription Drug Product and OTC Drug Product Lists those reference listed drugs to which the *in vivo* bioequivalence (reference standard) and, in some instances, the *in vitro* bioequivalence of the applicant's product is compared. By designating a single reference listed drug as the standard to which all generic versions must be shown to be bioequivalent, FDA hopes to avoid possible significant variations among generic drugs and their brand name counterpart. Such variations could result if generic drugs were compared to different reference listed drugs. However, in some instances when listed drugs are approved for a single drug product, a product not designated as the reference listed drug and not shown to be bioequivalent to the reference listed drug may be shielded from generic competition. A firm wishing to market a generic version of a listed drug that is not designated as the reference listed drug may petition the Agency through the Citizen Petition procedure (see 21 CFR 10.25(a) and CFR 10.30). When the Citizen Petition is approved, the second listed drug will be designated as an additional reference listed drug and the petitioner may submit an Abbreviated New Drug Application citing the designated reference listed drug. Section 1.7, *Therapeutic Equivalence Evaluations Codes products meeting necessary bioequivalence requirements* explains the **AB**, **AB1**, **AB2**, **AB3**... coding system for multisource drug products listed under the same heading with two reference listed drugs.

In addition, there are two situations in which two listed drugs that have been shown to be bioequivalent to each other may both be designated as reference listed drugs. The first situation occurs when the *in vivo* determination of bioequivalence is self-evident and a waiver of the *in vivo* bioequivalence may be granted. The second situation occurs when the bioequivalence of two listed products may be determined through *in vitro* methodology. The reference listed drug is identified by the symbol "+" in the Prescription and Over-the-Counter (OTC) Drug Product Lists. These identified reference listed drugs represent the best judgment of the Division of Bioequivalence at this time. The Prescription and OTC Drug Product Lists identify reference drugs for oral dosage forms, injectables, ophthalmics, otics, and topical products. It is recommended that a firm planning to conduct an *in vivo* bioequivalence study, or planning to manufacture a batch of a drug product for which an *in vivo* waiver of bioequivalence will be requested, contact the Division of Bioequivalence, Office of Generic Drugs, to confirm the appropriate reference listed drug.

1.5 General Policies and Legal Status

The List contains public information and advice. It does not mandate the drug products which may be purchased, prescribed, dispensed, or substituted for one another, nor does it, conversely, mandate the products that should be avoided. To the extent that the List sets forth FDA's evaluations of the therapeutic equivalence of drug products that have been approved, it contains FDA's advice to the public, to practitioners and to the states regarding drug

product selection. These evaluations do not constitute determinations that any product is in violation of the Act or that any product is preferable to any other. Therapeutic equivalence evaluations are a scientific judgment based upon evidence, while generic substitution may involve social and economic policy administered by the states, intended to reduce the cost of drugs to consumers. To the extent that the List identifies drug products approved under Section 505 of the Act, it sets forth information that the Agency is required to publish and that the public is entitled to under the Freedom of Information Act. Exclusion of a drug product from the List does not necessarily mean that the drug product is either in violation of Section 505 of the Act, or that such a product is not safe or effective, or that such a product is not therapeutically equivalent to other drug products. Rather, the exclusion is based on the fact that FDA has not evaluated the safety, effectiveness, and quality of the drug product.

1.6 Practitioner/User Responsibilities

Professional care and judgment should be exercised in using the List.

Evaluations of therapeutic equivalence for prescription drugs are based on scientific and medical evaluations by FDA. Products evaluated as therapeutically equivalent can be expected, in the judgment of FDA, to have equivalent clinical effect and no difference in their potential for adverse effects when used under the conditions of their labeling. However, these products may differ in other characteristics such as shape, scoring configuration, release mechanisms, packaging, excipients (including colors, flavors, preservatives), expiration date/time, and, in some instances, labeling. If products with such differences are substituted for each other, there is a potential for patient confusion due to differences in color or shape of tablets, inability to provide a given dose using a partial tablet if the proper scoring configuration is not available, or decreased patient acceptance of certain products because of flavor. There may also be better stability of one product over another under adverse storage conditions, or allergic reactions in rare cases due to a coloring or a preservative ingredient, as well as differences in cost to the patient.

FDA evaluation of therapeutic equivalence in no way relieves practitioners of their professional responsibilities in prescribing and dispensing such products with due care and with appropriate information to individual patients. In those circumstances where the characteristics of a specific product, other than its active ingredient, are important in the therapy of a particular patient, the physician's specification of that product is appropriate. Pharmacists must also be familiar with the expiration dates/times and labeling directions for storage of the different products, particularly for reconstituted products, to assure that patients are properly advised when one product is substituted for another.

Multisource and single-source drug products. FDA has evaluated for therapeutic equivalence only multisource prescription drug products approved under Section 505 of the Act, which in most instances means those pharmaceutical equivalents available from more than one manufacturer. For such products, a therapeutic equivalence code is included and, in addition, product information is highlighted in bold face and underlined. Those products with approved applications that are single-source (i.e., there is only one approved product available for that active ingredient, dosage form, route of administration, and strength) are also included on the List, but no therapeutic equivalence code is included with such products. Any drug product in the List repackaged and/or distributed by other than the application holder is considered to be therapeutically equivalent to the application holder's drug product even if the application holder's drug product is single source or coded as non-equivalent (e.g., **BN**). Also, although not identified in the List, distributors or repackagers of an application holder's drug product are considered to have the same code as the application holder. The details of these codes and the policies underlying them are discussed in Section 1.7, *Therapeutic Equivalence Evaluations Codes*.

Products on the List are identified by the names of the holders of approved applications (applicants) who may not necessarily be the manufacturer of the product. The applicant may have had its product manufactured by a contract manufacturer and may simply be distributing the product for which it has obtained approval. In most instances, however, the manufacturer of the product is also the applicant. The name of the manufacturer is permitted by regulation to appear on the label, even when the manufacturer is not the marketer.

Although the products on the List are identified by the names of the applicants, circumstances, such as changing corporate ownership, have sometimes made identification of the applicant difficult. The Agency believes, based on continuing document review and communication with firms, that the applicant designations on the List are, in most cases, correct.

To relate firm name information on a product label to that on the List, the following should be noted: the applicant's name always appears on the List. This applies whether the applicant (firm name on the Form FDA 356h in the application) is the marketer (firm name in largest letters on the label) or not. However, the applicant's name may not always appear on the label of the product.

If the applicant is the marketer, its name appears on the List and on the label; if the applicant is not the marketer, and the Agency is aware of a corporate relationship (e.g., parent and subsidiary) between the applicant and the marketer, the name of the applicant appears on the List and both firm names may appear on the label. Firms with known corporate relationships are displayed in Appendix B. If there is no known corporate relationship between the applicant and the marketer, the applicant's name appears on the List; however, unless the applicant is the manufacturer, packager, or distributor, the applicant's name may not appear on the label. In this case, the practitioner, from labeling alone, will not be able to relate the marketed product to an applicant cited in the List, and hence to a specific approved drug product. In such cases, to assure that the product in question is the subject of an approved application, the firm named on the label should be contacted.

To relate trade name (proprietary name) information on a product label to that on the List, the following should be noted: if the applicant is the marketer, its name appears on the List and on the label; if the Agency is aware of a corporate relationship between the applicant and the marketer, the trade name (proprietary name) of the drug product (established drug name if no trade name exists) appears on the List. If a corporate relationship exists between an application holder and a marketer and both firms are distributing the drug product, the FDA reserves the right to select the trade name of either the marketer or the application holder to appear on the List. If there is no known corporate relationship between the applicant and the marketer, the established drug name appears on the List.

Every product on the List is subject at all times to regulatory action. From time to time, approved products may be found in violation of one or more provisions of the Act. In such circumstances, the Agency will commence appropriate enforcement action to correct the violation, if necessary, by securing removal of the product from the market by voluntary recall, seizure, or other enforcement actions. Such regulatory actions are, however, independent of the inclusion of a product on the List. The main criterion for inclusion of a product is that it has an application with an effective approval that has not been withdrawn for safety or efficacy reasons. FDA believes that retention of a violative product on the List will not have any significant adverse health consequences, because other legal mechanisms are available to the Agency to prevent the product's actual marketing. FDA may however, change a product's therapeutic equivalence rating if the circumstances giving rise to the violation change or otherwise call into question the data upon which the Agency's assessment of whether a product meets the criteria for therapeutic equivalence was made.

1.7 Therapeutic Equivalence Evaluations Codes

The coding system for therapeutic equivalence evaluations is constructed to allow users to determine quickly whether the Agency has evaluated a particular approved product as therapeutically equivalent to other pharmaceutically equivalent products (first letter) and to provide additional information on the basis of FDA's evaluations (second letter). With few exceptions, the therapeutic equivalence evaluation date is the same as the approval date.

The two basic categories into which multisource drugs have been placed are indicated by the first letter as follows:

A Drug products that FDA considers to be therapeutically equivalent to other pharmaceutically equivalent products, i.e., drug products for which:

- (1) there are no known or suspected bioequivalence problems. These are designated **AA, AN, AO, AP, or AT**, depending on the dosage form; or
- (2) actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bioequivalence. These are designated **AB**.

B Drug products that FDA at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products, i.e.,

drug products for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence. Often the problem is with specific dosage forms rather than with the active ingredients. These are designated **BC, BD, BE, BN, BP, BR, BS, BT, BX, or B***.

Individual drug products have been evaluated as therapeutically equivalent to the reference product in accordance with the definitions and policies outlined below:

"A" CODES

Drug products that are considered to be therapeutically equivalent to other pharmaceutically equivalent products.

"A" products are those for which actual or potential bioequivalence problems have been resolved with adequate *in vivo* and/or *in vitro* evidence supporting bio-equivalence. Drug products designated with an "A" code fall under one of two main policies:

- (1) for those active ingredients or dosage forms for which no *in vivo* bioequivalence issue is known or suspected, the information necessary to show bioequivalence between pharmaceutically equivalent products is presumed and considered self-evident based on other data in the application for some dosage forms (e.g., solutions) or satisfied for solid oral dosage forms by a showing that an acceptable *in vitro* dissolution standard is met. A therapeutically equivalent rating is assigned such products so long as they are manufactured in accordance with Current Good Manufacturing Practice regulations and meet the other requirements of their approved applications (these are designated **AA, AN, AO, AP, or AT**, depending on the dosage form, as described below); or

- (2) for those DESI drug products containing active ingredients or dosage forms that have been identified by FDA as having actual or potential bioequivalence problems, and for post-1962 drug products in a dosage form presenting a potential bioequivalence problem, an evaluation of therapeutic equivalence is assigned to pharmaceutical equivalents only if the approved application contains adequate scientific evidence establishing through *in vivo* and/or *in vitro* studies the bioequivalence of the product to a selected reference product (these products are designated as **AB**).

There are some general principles that may affect the substitution of pharmaceutically equivalent products in specific cases. Prescribers and dispensers of drugs should be alert to these principles so as to deal appropriately with situations that require professional judgment and discretion.

There may be labeling differences among pharmaceutically equivalent products that require attention on the part of the health professional. For example, pharmaceutically equivalent powders to be reconstituted for administration as oral or injectable liquids may vary with respect to their expiration time or storage conditions after reconstitution. An FDA evaluation that such products are therapeutically equivalent is applicable only when each product is reconstituted, stored, and used under the conditions specified in the labeling of that product.

The Agency will use notes in this publication to point out special situations such as potential differences between two drug products that have been evaluated as bioequivalent and otherwise therapeutically equivalent, when they should be brought to the attention of health professionals. These notes are contained in Section 1.8, *Description of Special Situations*.

For example, in rare instances, there may be variations among therapeutically equivalent products in their use or in conditions of administration. Such differences may be due to patent or exclusivity rights associated with such use. When such variations may, in the Agency's opinion, affect prescribing or substitution decisions by health professionals, a note will be added to Section 1.8.

Also, occasionally a situation may arise in which changes in a listed drug product after its approval (for example, a change in dosing interval) may have an impact on the substitutability of already approved generic versions of that product that were rated by the Agency as therapeutically equivalent to the listed product. When such changes in the listed drug product are considered by the Agency to have a significant impact on therapeutic equivalence, the Agency will change the therapeutic equivalence ratings for other versions of the drug product unless the manufacturers of those other versions of the product provide additional information to assure equivalence under the changed conditions. Pending receipt of the additional data, the Agency may add a note to Section 1.8, or, in rare cases, may even change the therapeutic equivalence rating.

In some cases (e.g., Isolyte® S w/ Dextrose 5% in Plastic Container and Plasma-Lyte® 148 and Dextrose 5% in Plastic Container), closely related products are listed as containing the same active ingredients, but in somewhat different amounts. In determining which of these products are pharmaceutically equivalent, the Agency has considered products to be pharmaceutically equivalent with labeled strengths of an ingredient that do not vary by more than 1%.

Different salts and esters of the same therapeutic moiety are regarded as pharmaceutical alternatives. For the purpose of this publication, such products are not considered to be therapeutically equivalent. There are no instances in this List where pharmaceutical alternatives are evaluated or coded with regard to therapeutic equivalence. Anhydrous and hydrated entities, as well as different polymorphs, are considered pharmaceutical equivalents and must meet the same standards and, where necessary, as in the case of

FDA data supplied by DrugPatentWatch.com
ampicillin/ampicillin trihydrate, their equivalence is supported by appropriate bioavailability/bioequivalence studies.

The codes in this book are not intended to preclude health care professionals from converting pharmaceutically different concentrations into pharmaceutical equivalents using accepted professional practice.

Where package size variations have therapeutic implications, products so packaged have not been considered pharmaceutically equivalent. For example, some oral contraceptives are supplied in 21-tablet and 28-tablet packets; the 28-tablet packets contain 7 placebo or iron tablets. These two packaging configurations are not regarded as pharmaceutically equivalent; thus, they are not designated as therapeutically equivalent.

Preservatives may differ among some therapeutically equivalent drug products. Differences in preservatives and other inactive ingredients do not affect FDA's evaluation of therapeutic equivalence except in cases where these components may influence bioequivalence or routes of administration.

The specific sub-codes for those drugs evaluated as therapeutically equivalent and the policies underlying these sub-codes follow:

AA Products in conventional dosage forms not presenting bioequivalence problems

Products coded as **AA** contain active ingredients and dosage forms that are not regarded as presenting either actual or potential bioequivalence problems or drug quality or standards issues. However, all oral dosage forms must, nonetheless, meet an appropriate *in vitro* bioequivalence standard that is acceptable to the Agency in order to be approved.

AB, AB1, AB2, AB3... Products meeting necessary bioequivalence requirements

Multisource drug products listed under the same heading (i.e., identical active ingredients(s), dosage form, and route(s) of administration) and having the same strength (see Section 1.2, *Therapeutic Equivalence-Related Terms, Pharmaceutical Equivalents*) generally will be coded **AB** if a study is submitted demonstrating bioequivalence.

In certain instances, a number is added to the end of the **AB** code to make a three character code (i.e., **AB1, AB2, AB3, etc.**). Three-character codes are assigned only in situations when more than one reference listed drug of the same strength has been designated under the same heading. Two or more reference listed drugs are generally selected only when there are at least two potential reference drug products which are not bioequivalent to each other. If a study is submitted that demonstrates bioequivalence to a specific listed drug product, the generic product will be given the same three-character code as the reference listed drug it was compared against.

For example, Adalat® CC (Miles) and Procardia XL® (Pfizer), extended-release tablets, are listed under the active ingredient nifedipine. These drug products, listed under the same heading, are not bioequivalent to each other. Generic drug products deemed by FDA to be bioequivalent to Adalat® CC and Procardia XL® have been approved, Adalat® CC and Procardia XL® have been assigned ratings of **AB1** and **AB2**, respectively. The generic drug products bioequivalent to Adalat® CC would be assigned a rating of **AB1** and those bioequivalent to Procardia XL® would be assigned a rating of **AB2**. (The assignment of an **AB1** or **AB2** rating to a specific product does not imply product preference.) Even though drug products of distributors and/or repackagers are not included in the List, they are considered therapeutically equivalent to the application holder's drug product if the application holder's drug product is rated either with an **AB** or three-character code or is single source in the List. Drugs coded as **AB** under a heading are considered therapeutically equivalent only to other drugs coded as **AB** under that heading. Drugs coded with a three-character code

under a heading are considered therapeutically equivalent only to other drugs coded with the same three-character code under that heading.

AN Solutions and powders for aerosolization

Uncertainty regarding the therapeutic equivalence of aerosolized products arises primarily because of differences in the drug delivery system. Solutions and powders intended for aerosolization that are marketed for use in any of several delivery systems are considered to be pharmaceutically and therapeutically equivalent and are coded **AN**. Those products that are compatible only with a specific delivery system or those products that are packaged in and with a specific delivery system are coded **BN**, unless they have met an appropriate bioequivalence standard. Solutions or suspensions in a specific delivery system will be coded **AN** if the bioequivalence standard is based upon *in vitro* methodology, if bioequivalence needs to be demonstrated by *in vivo* methodology then the drug products will be coded **AB**.

AO Injectable oil solutions

The absorption of drugs in injectable (parenteral) oil solutions may vary substantially with the type of oil employed as a vehicle and the concentration of the active ingredient. Injectable oil solutions are therefore considered to be pharmaceutically and therapeutically equivalent only when the active ingredient, its concentration, and the type of oil used as a vehicle are all identical.

AP Injectable aqueous solutions and, in certain instances, intravenous non-aqueous solutions

It should be noted that even though injectable (parenteral) products under a specific listing may be evaluated as therapeutically equivalent, there may be important differences among the products in the general category, *Injectable; Injection*. For example, some injectable products that are rated therapeutically equivalent are labeled for different routes of administration. In addition, some products evaluated as therapeutically equivalent may have different preservatives or no preservatives at all. Injectable products available as dry powders for reconstitution, concentrated sterile solutions for dilution, or sterile solutions ready for injection are all considered to be pharmaceutically and therapeutically equivalent provided they are designed to produce the same concentration prior to injection and are similarly labeled. Consistent with accepted professional practice, it is the responsibility of the prescriber, dispenser, or individual administering the product to be familiar with a product's labeling to assure that it is given only by the route(s) of administration stated in the labeling.

Certain commonly used large volume intravenous products in glass containers are not included on the List (e.g., dextrose injection 5%, dextrose injection 10%, sodium chloride injection 0.9%) since these products are on the market without FDA approval and the FDA has not published conditions for marketing such parenteral products under approved NDAs. When packaged in plastic containers, however, FDA regulations require approved applications prior to marketing. Approval then depends on, among other things, the extent of the available safety data involving the specific plastic component of the product. All large volume parenteral products are manufactured under similar standards, regardless of whether they are packaged in glass or plastic. Thus, FDA has no reason to believe that the packaging container of large volume parenteral drug products that are pharmaceutically equivalent would have any effect on their therapeutic equivalence.

The strength of parenteral drugs products is defined as the total drug content of the container. Until recently the strength of liquid parenteral drug products in the Orange Book have not been displayed. The concentration of the liquid parenteral drug product in the Orange Book has been shown as xmg/ml. The amount of dry powder or freeze dried powder in a container has always been identified as the strength.

With the finalization of the Waxman-Hatch amendments that characterized each strength of a drug product as a listed drug, it became evident that the format of the Orange Book should be changed to reflect each strength of a parenteral solution. To this end the OGD has started to display the strength of all new approvals of parenteral solutions. Previously we would have displayed only the concentration of an approved parenteral solution, e.g. 50mg/ml. If this drug product had a 20 ml and 60 ml container approved the two products would be shown as 1gm / 20ml (50mg/ml) and 3gm / 60ml (50mg/ml).

AT Topical products

There are a variety of topical dosage forms available for dermatologic, ophthalmic, otic, rectal, and vaginal administration, including creams, gels, lotions, oils, ointments, pastes, solutions, sprays and suppositories. Even though different topical dosage forms may contain the same active ingredient and potency, these dosage forms are not considered pharmaceutically equivalent. Therefore, they are not considered therapeutically equivalent. All solutions and DESI drug products containing the same active ingredient in the same topical dosage form for which a waiver of *in vivo* bioequivalence has been granted and for which chemistry and manufacturing processes are adequate to demonstrate bioequivalence, are considered therapeutically equivalent and coded **AT**. Pharmaceutically equivalent topical products that raise questions of bioequivalence, including all post-1962 non-solution topical drug products, are coded **AB** when supported by adequate bioequivalence data, and **BT** in the absence of such data.

"B" CODES

Drug products that FDA, at this time, considers not to be therapeutically equivalent to other pharmaceutically equivalent products.

"B" products, for which actual or potential bioequivalence problems have not been resolved by adequate evidence of bioequivalence, often have a problem with specific dosage forms rather than with the active ingredients. Drug products designated with a "B" code fall under one of three main policies:

- (1) the drug products contain active ingredients or are manufactured in dosage forms that have been identified by the Agency as having documented bio-equivalence problems or a significant potential for such problems and for which no adequate studies demonstrating bioequivalence have been submitted to FDA; or
- (2) the quality standards are inadequate or FDA has an insufficient basis to determine therapeutic equivalence; or
- (3) the drug products are under regulatory review.

The specific coding definitions and policies for the "B" sub-codes are as follows:

B* Drug products requiring further FDA investigation and review to determine therapeutic equivalence

The code **B*** is assigned to products previously assigned an **A** or **B** code when FDA receives new information that raises a significant question regarding therapeutic equivalence that can be resolved only through further Agency investigation and/or review of data and information submitted by the applicant. The **B*** code signifies that the Agency will take no position regarding the therapeutic equivalence of the product until the Agency completes its investigation and review.

BC Extended-release dosage forms (capsules, injectables and tablets)

Extended-release tablets are formulated in such a manner as to make the contained medicament available over an extended period of time following ingestion.

Although bioavailability studies have been conducted on these dosage forms, they may be subject to bioavailability differences, primarily because firms developing extended-release products for the same active ingredient rarely employ the same formulation approach. FDA, therefore, does not consider different extended-release dosage forms containing the same active ingredient in equal strength to be therapeutically equivalent unless equivalence between individual products in both rate and extent has been specifically demonstrated through appropriate bioequivalence studies. Extended-release products for which such bioequivalence data have not been submitted are coded **BC**, while those for which such data are available have been coded **AB**.

BD Active ingredients and dosage forms with documented bioequivalence problems

The **BD** code denotes products containing active ingredients with known bioequivalence problems and for which adequate studies have not been submitted to FDA demonstrating bioequivalence. Where studies showing bioequivalence have been submitted, the product has been coded **AB**.

BE Delayed-release oral dosage forms

Where the drug may be destroyed or inactivated by the gastric juice or where it may irritate the gastric mucosa, the use of "enteric" coatings is indicated. Such coatings are intended to delay the release of the medication until the tablet has passed through the stomach. Drug products in delayed-release dosage forms containing the same active ingredients are subject to significant differences in absorption. Unless otherwise specifically noted, the Agency considers different delayed-release products containing the same active ingredients as presenting a potential bioequivalence problem and codes these products **BE** in the absence of *in vivo* studies showing bioequivalence. If adequate *in vivo* studies have demonstrated the bioequivalence of specific delayed-release products, such products are coded **AB**.

BN Products in aerosol-nebulizer drug delivery systems

This code applies to drug solutions or powders that are marketed only as a component of, or as compatible with, a specific drug delivery system. There may, for example, be significant differences in the dose of drug and particle size delivered by different products of this type. Therefore, the Agency does not consider different metered aerosol dosage forms containing the same active ingredient(s) in equal strengths to be therapeutically equivalent unless the drug products meet an appropriate bioequivalence standard, such products are coded **AB**.

BP Active ingredients and dosage forms with potential bioequivalence problems

FDA's bioequivalence regulations (21 CFR 320.33) contain criteria and procedures for determining whether a specific active ingredient in a specific dosage form has a potential for causing a bioequivalence problem.

It is FDA's policy to consider an ingredient meeting these criteria as having a potential bioequivalence problem even in the absence of positive data demonstrating inequivalence. Pharmaceutically equivalent products containing these ingredients in oral dosage forms are coded **BP** until adequate *in vivo* bioequivalence data are submitted, such products are coded **AB**. Injectable suspensions containing an active ingredient suspended in an aqueous or oleaginous vehicle have also been coded **BP**. Injectable suspensions are subject to bioequivalence problems because differences in particle size, polymorphic structure of the suspended active ingredient, or the suspension formulation can significantly affect the rate of release and absorption. FDA does not consider pharmaceutical equivalents of these products bioequivalent without adequate evidence of bioequivalence, such products would be coded **AB**.

BR Suppositories or enemas that deliver drugs for systemic absorption

The absorption of active ingredients from suppositories or enemas that are intended to have a systemic effect (as distinct from suppositories administered for local effect) can vary significantly from product to product. Therefore, FDA considers pharmaceutically equivalent systemic suppositories or enemas bio-equivalent only if *in vivo* evidence of bioequivalence is available. In those cases where *in vivo* evidence is available, the product is coded **AB**. If such evidence is not available, the products are coded **BR**.

BS Products having drug standard deficiencies

If the drug standards for an active ingredient in a particular dosage form are found by FDA to be deficient so as to prevent an FDA evaluation of either pharmaceutical or therapeutic equivalence, all drug products containing that active ingredient in that dosage form are coded **BS**. For example, if the standards permit a wide variation in pharmacologically active components of the active ingredient such that pharmaceutical equivalence is in question, all products containing that active ingredient in that dosage form are coded **BS**.

BT Topical products with bioequivalence issues

This code applies mainly to post-1962 dermatologic, ophthalmic, otic, rectal, and vaginal products for topical administration, including creams, ointments, gels, lotions, pastes, and sprays, as well as suppositories not intended for systemic drug absorption. Topical products evaluated as having acceptable clinical performance, but that are not bioequivalent to other pharmaceutically equivalent products or that lack sufficient evidence of bioequivalence, will be coded **BT**.

BX Drug products for which the data are insufficient to determine therapeutic equivalence

The code **BX** is assigned to specific drug products for which the data that have been reviewed by the Agency are insufficient to determine therapeutic equivalence under the policies stated in this document. In these situations, the drug products are presumed to be therapeutically inequivalent until the Agency has determined that there is adequate information to make a full evaluation of therapeutic equivalence.

1.8 Description of Special Situations

Certain drugs listed in the Orange Book present special situations that merit further discussion. Following is a description of those special situations:

Amino Acid and Protein Hydrolysate Injections. These products differ in the amount and kinds of amino acids they contain and, therefore, are not considered pharmaceutical equivalents. For this reason, these products are not considered therapeutically equivalent. At the same time, the Agency believes that it is appropriate to point out that where nitrogen balance is the sole therapeutic objective and individual amino acid content is not a consideration, pharmaceutical alternatives with the same total amount of nitrogen content may be considered therapeutically equivalent.

Follitropin Alfa and Beta. Based on available data derived from physico-chemical tests and bioassay, follitropin alfa and follitropin beta are indistinguishable.

Gaviscon®. Gaviscon® is an OTC product which has been marketed since September 1970. The active ingredients in this product, aluminum hydroxide and magnesium trisilicate, were reviewed by the Agency's OTC Antacid Panel and were considered to be safe and effective ingredients (Category I) by that Panel. However, the tablet failed to pass the antacid test which is required of all antacid products. The Agency, therefore, placed the tablet in Category III for lack of effectiveness. A full NDA with clinical studies was submitted by Marion Laboratories, Inc., and approved by FDA on December 9, 1983. Gaviscon®'s activity in treating reflux acidity is made possible by the physical-chemical properties of the inactive ingredients, sodium bicarbonate and alginic acid. Therefore, *all ANDAs which cite Gaviscon® tablets as the listed drug must contain the inactive ingredients sodium bicarbonate and alginic acid.* A full NDA will be required to support the effectiveness of the drug product if different inactive ingredients are to be substituted for sodium bicarbonate or alginic acid or if different proportions of these ingredients are to be used.

Levothyroxine Sodium. Because there are multiple reference listed drugs of levothyroxine sodium tablets and some reference listed drugs' sponsors have conducted studies to establish their drugs' therapeutic equivalence to other reference listed drugs, FDA has determined that its usual practice of assigning two or three character TE codes may be potentially confusing and inadequate for these drug products. Accordingly, FDA provides the following explanation and chart of therapeutic equivalence evaluations for levothyroxine sodium drug products.

Levothyroxine Sodium (Mylan ANDA 76187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Unithroid (Jerome Stevens NDA 021210) tablets.

Levo-T (Alara NDA 021342), Levothyroxine Sodium (Mylan ANDA 76187) and Unithroid (Jerome Stevens NDA 021210) tablets have been determined to be therapeutically equivalent to corresponding strengths of Synthroid (Abbott NDA 021402) tablets.

Levo-T (Alara NDA 021342), Unithroid (Jerome Stevens NDA 021210) and Levothyroxine Sodium (Mylan ANDA 076187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levoxyl (King/Jones Pharma NDA 021301) tablets.

Novothyrox (Genpharm NDA 021292) requires further investigation and review to establish therapeutic equivalence to corresponding strengths of any other levothyroxine sodium drug products and is rated BX.

Levothroid (Lloyd NDA 021116) requires further investigation and review to establish therapeutic equivalence to corresponding strengths of any other levothyroxine sodium drug products and is rated BX.

Levolet (Vintage NDA 021137) requires further investigation and review to establish therapeutic equivalence to corresponding strengths of any other levothyroxine sodium drug products and is rated BX.

The chart outlines TE codes for all 0.025mg products with other products being similar. Therapeutic equivalence has been established between products that have the same AB+number TE code. More than one TE code may apply to some products. One common TE code indicates therapeutic equivalence between products.

Trade Name	Applicant	Potency	TE Code	Appl No	Product No
UNITHROID	STEVENS J	0.025MG	AB1	21210	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB1	76187	001
LEVOXYL	JONES PHARMA	0.025MG	AB1	21301	001
SYNTHROID	ABBOTT	0.025MG	AB1	21402	001
SYNTHROID	ABBOTT	0.025MG	AB2	21402	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB2	76187	001
LEVO-T	ALARA PHARM	0.025MG	AB2	21342	001
UNITHROID	STEVENS J	0.025MG	AB2	21210	001
LEVOXYL	JONES PHARMA	0.025MG	AB3	21301	001
LEVO-T	ALARA PHARM	0.025MG	AB3	21342	001
UNITHROID	STEVENS J	0.025MG	AB3	21210	001
LEVOTHYROXINE SODIUM	MYLAN	0.025MG	AB3	76187	001
NOVOTHYROX	GENPHARM	0.025MG	BX	21292	001
LEVOTHROID	LLOYD	0.025MG	BX	21116	001
LEVOLET	VINTAGE PHARMS	0.025MG	BX	21137	001

Patent Certification(s) Reference Listed Drug based upon a suitability petition. An abbreviated new drug application that refers to a Reference Listed Drug (RLD) approved pursuant to a suitability petition must demonstrate that the proposed product is bioequivalent to the RLD, and it must include appropriate patent certification(s) and an exclusivity statement with respect to the listed drug which served as the basis for the approved suitability petition.

Waived exclusivity. If a new drug application (NDA) submitted under section 505(b) of the Federal Food, Drug, and Cosmetic Act (Act) qualifies for exclusivity under sections 505(c)(3)(D) and 505(j)(5)(D), the exclusivity is listed in the Patent and Exclusivity Section of the Orange Book. If a drug product has received this exclusivity, the FDA will delay the approval of a 505(b)(2) application or an abbreviated new drug application (ANDA) under section 505(j) of the Act until the expiration of the exclusivity. If the listed drug is also protected by one or more patents, the approval date for the 505(b)(2) application or ANDA will be determined by the latest expiring patent or exclusivity listed in the Orange Book. However, the holder of the NDA may waive its exclusivity as to any or all 505(b)(2) and ANDA applications referencing the protected drug product. If an NDA sponsor waives its right to the exclusivity protection, qualified 505(b)(2) or ANDA applications may be approved without regard to the NDA holder's exclusivity. An NDA for which the holder has waived its exclusivity as to all 505(b)(2) and ANDA applications will be coded with a W in the Patent and Exclusivity Section of the Orange Book and be referred to this section. The applicant referencing this listed drug should indicate in the exclusivity statement that the holder of the listed drug has waived its exclusivity.

1.9 Therapeutic Equivalence Code Change for a Drug Entity

The Agency will use the following procedures when, in response to a petition or on its own initiative, it is considering a change in the therapeutic equivalence code for approved multi-source drug products. Such changes will generally occur when the Agency becomes aware of new scientific information affecting the therapeutic equivalence of an entire category of drug products in the List (e.g., information concerning the active ingredient or the dosage form), rather than information concerning a single drug product within the category. These procedures will be used when a change in therapeutic equivalence code is under consideration for all drug products found in the Prescription Drug Product List under a specific drug entity and dosage form. The change may be from the code signifying that the drug does not present a bioequivalence problem (e.g., **AA**) to a code signifying a bioequivalence problem (e.g., **BP**), or vice versa. This procedure does not apply to a change of a particular product code (e.g., a change from **BP** to **AB** or from **AB** to **BX**).

Before making a change in a therapeutic equivalence code for an entire category of drugs, the Agency will announce in the *Introduction* to the Cumulative Supplement that it is considering the change and will invite comment. Comments, along with scientific data, may be sent to the Director, Division of Bioequivalence, Office of Generic Drugs, Center for Drug Evaluation and Research, HFD-650, 7500 Standish Place, Rockville, MD 20855. The comment period will generally be 60 days in length, and the closing date for comments will be listed in the description of the proposed change for each drug entity.

The most useful type of scientific data submission is an *in vivo* bioavailability/bioequivalence study conducted on batches of the subject drug products. These submissions should present a full description of the analytical procedures and equipment used, a validation of the analytical methodology, including the standard curve, a description of the method of calculating results, and a description of the pharmacokinetic and statistical models used in analyzing the data. Anecdotal or testimonial information is the least useful to the Agency, and such submissions are discouraged. Copies of supporting reports published in the scientific literature or unpublished material, however, are welcome.

1.10 Change of the Therapeutic Equivalence Evaluation for a Single Product

The aforementioned procedure does not apply to a change in a single drug product code. For example, a change in a single drug product's code from **BP** to **AB** as a result of the submission of an acceptable bioequivalence study ordinarily will not be the subject of notice and comment. Likewise, a change in a single drug product's code from **AB** to **BX** (e.g., as a result of new information raising a significant question as to bioequivalence) does not require notice and comment. The Agency's responsibility to provide the public with the Agency's most current information related to therapeutic equivalence may require a change in a drug product's code prior to any formal notice and opportunity for the applicant to be heard. The publication in the *Federal Register* of a proposal to withdraw approval of a drug product will ordinarily result in a change in a product's code from **AB** to **BX** if this action has not already been taken.

1.11 Discontinued Section

Those drug products in the Discontinued Section of the Orange Book in which a determination has already been made that the products were not marketed or withdrawn for safety or efficacy reasons have "***Federal Register determination that product was not discontinued or withdrawn for safety or

efficacy reasons**" following the product strength. Those drug products are only reflective of citizen petitions approved since 1995. The identification of these drug products in the Discontinued Section of the Orange Book should avoid the submission of multiple citizen petitions for the same drug product

Generally, approved products are added to the Discontinued Section of the Orange Book when the applicant holder notifies the Orange Book staff of the products' not marketed status. Products may also be added if annual reports indicate the product is no longer marketed or other Agency administrative action (e.g., Withdrawal of an Application). Changes to the Orange Book are not affected by the drug registration and listing requirements of Section 510 of the Act.

1.12 Changes to the Orange Book

Every effort is made to ensure the Annual Edition is current and accurate. Applicant holders are requested to inform the FDA Orange Book Staff (OBS) of any changes or corrections. Please inform the OBS when products are no longer marketed. Notification of the Orange Book staff to include the newly approved product in the Discontinued Drug Product List rather than parts 1, 2 or 3 of the List (as discussed in Section 1.1) must occur by the end of the month in which the product is approved to ensure that the product is not included in the "active" portions of the next published Orange Book update

The OBS can be contacted by email at drugproducts@cder.fda.gov. Send Changes by FAX: 301-827-7337; mail to:

FDA/CDER Orange Book Staff
Office of Generic Drugs, HFD-610
7500 Standish Place
Rockville, MD 20855-2773

1.13 Availability of the Edition

Commencing with the 25th edition, the Annual Edition and current monthly Cumulative Supplements are available in a Portable Document Format (PDF) at the EOB home page, <http://www.fda.gov/cder/ob/default.htm>, by clicking on the [Annual Edition](#). The PDF annual format will duplicate previous paper versions except for the Orphan Products Designations and Approvals List. An annual subscription of the PDF format may be obtained from the U.S. Government Printing Office, 866-512-1800.

2. HOW TO USE THE DRUG PRODUCT LISTS

2.1 Key Sections for Using the Drug Product Lists

This publication contains illustrations, along with Drug Product Lists, indices, and lists of abbreviations and terms which facilitate their use.

Illustrations. The annotated *Drug Product Illustration*, see Section 2.2, and the *Therapeutic Equivalence Evaluations Illustration*, see Section 2.3, are offered to provide further clarification. These depict the format found in the Prescription Drug Product List (the only list in which therapeutic equivalence evaluation codes are displayed).

Drug Product Lists. The Prescription and OTC Drug Product Lists, arranged alphabetically by active ingredient(s), contain product identification information (active ingredients, dosage forms, routes of administration, product names, application holders, strengths) for single and multiple ingredient drug products. Also shown are the application number and drug product number (FDA internal computer data use only) and approval dates for those drug products approved on or after January 1, 1982.

The Discontinued Product List, arranged alphabetically by active ingredient(s), contain product identification information (dosage form, product name, strength, and application number).

If a prescription drug product is available from more than one source (multisource), a therapeutic equivalence code will appear in front of the applicant's name. If a product is therapeutically equivalent to one or more products or to an appropriate reference, it will be designated with a code beginning with "A" and the entry will be underlined and printed in bold font for emphasis.

Active ingredient headings for multiple ingredient (combination) drug products are arranged alphabetically. For purposes of this publication, this alphabetical sort takes precedence over United States Pharmacopeia official monograph order (i.e., Reserpine, Hydralazine Hydrochloride, Hydrochlorothiazide). For example, product information labeled as Reserpine, Hydrochlorothiazide and Hydralazine Hydrochloride appears under the active ingredient heading *Hydralazine Hydrochloride; Hydrochlorothiazide; Reserpine*. A cross-reference to the product information (for prescription and OTC products) appears for each additional active ingredient in the product. For combination drug products, the ingredient strengths are separated by semicolons and appear in the same relative sequence as the ingredients in the heading. Available strengths of the dosage form from an applicant appear on separate lines.

To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if necessary. Then, find the ingredient in the applicable Drug Product List. Proceed to the dosage form and route of administration and compare products within that ingredient heading only. Therapeutic equivalence or inequivalence for prescription products is determined on the basis of the therapeutic equivalence codes provided within that specific dosage form and route heading. The OTC Drug Product List, Discontinued Drug Product List, and Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research List have their data arranged

similarly.

The Discontinued Drug Product List contains approved products that have never been marketed, have been discontinued from marketing, are for military use, or have had their approvals withdrawn for other than safety or efficacy reasons subsequent to being discontinued from marketing. All products having a "@" in the 12th Cumulative Supplement of the 25th Edition List have been added to the Discontinued Drug Product List appearing in the 26th Edition. In addition, approved drug products that are not in the commercial distribution channel e.g., approved drug products in applications for export only are also listed in the Discontinued Section of the Orange Book.

Product Name Index (*Prescription and OTC Drug Product Lists*). This is an index of drug products by established or trade name. The second term of each entry indicates the active ingredient name under which product information can be found in the appropriate Drug Product List. For those drug products with multiple active ingredients, only the first active ingredient (in alphabetical order) will appear. OTC products are so designated.

Product Name Index Listed by Applicant (*Prescription and OTC Drug Product Lists*). This is an index that cross-references applicants to drug products. The bolded and underlined entry represents the applicant name abbreviation used in this publication. Each complete applicant name that is represented by the abbreviated name is marked with an asterisk (*). Listed under each complete applicant name is the first alphabetically arranged ingredient under which product information can be found in the appropriate Drug Product List. OTC products are so designated. To use the Drug Product Lists, determine by alphabetical order the ingredient under which the product information is listed, using the Product Name Index, if appropriate.

Uniform Terms. To improve readability, uniform terms are used to designate dosage forms, routes of administration, and abbreviations used to express strengths. These terms are listed in Appendix C. In some cases, the terms used may differ from those used in product labels and other labeling.

2.2 DRUG PRODUCT ILLUSTRATION

SINGLE INGREDIENT

ACTIVE INGREDIENT	MEPERIDINE HYDROCHLORIDE			
DOSAGE FORM; ROUTE OF ADMINISTRATION	INJECTABLE; INJECTION			
TRADE OR GENERIC NAMES	HEXANON			
REFERENCE LISTED DRUG	AP + METRO-PHYS	25MG/ML	N13111 001	
	AP +	50MG/ML	N13111 002	
	AP +	75MG/ML	N13111 003	
	AP +	100MG/ML	N13111 004	AUG 22, 1983
	MEPERIDINE HCL			JAN 04, 1985
	DONHARE PHARM			
THERAPEUTIC EQUIVALENCE (TE)	AP	25MG/ML	N42242 001	
CODE FOR MULTISOURCE PRODUCT	AP	50MG/ML	N42296 001	
	AP	75MG/ML	N42301 002	
	AP	100MG/ML	N42301 003	AUG 27, 1987
FINAL APPROVAL DATE				AUG 27, 1987
SINGLE SOURCE PRODUCT (NO TE CODE)	AP	JOHNSON MED	10MG/ML	N40000 001
		HOOVAC LLC	25MG/ML	N47222 001
		OBS PHARM	150MG/ML	N47100 001
APPLICANT				
AVAILABLE STRENGTH(S) OF A PRODUCT				
APPLICATION NUMBER AND PRODUCT NUMBER				
PRODUCT NUMBER IS FOR FDA INTERNAL COMPUTER DATA USE ONLY				

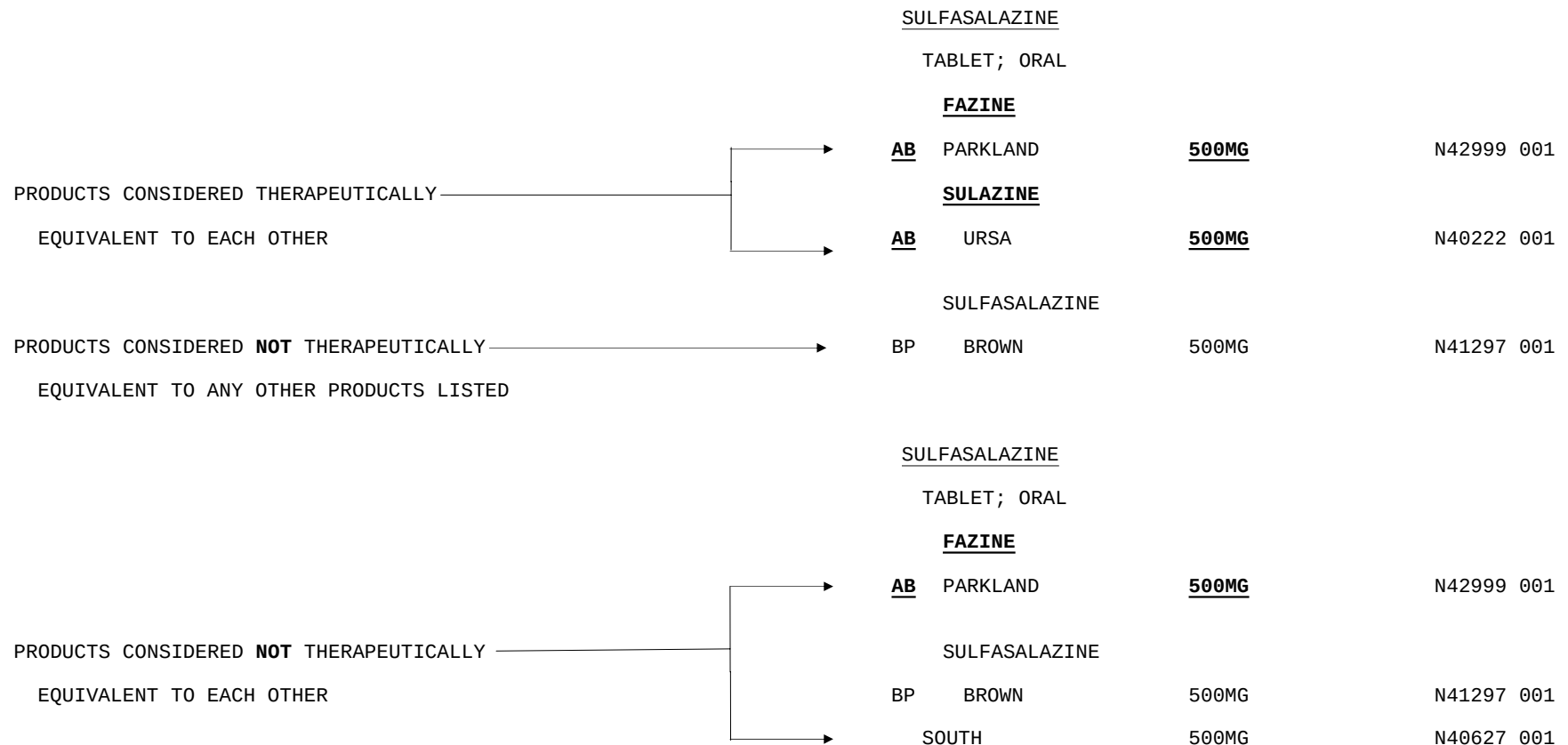
MULTIPLE INGREDIENTS WITH PRODUCT INFORMATION

ALPHABETICALLY SORTED BY				
ACTIVE INGREDIENT	HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE			
PRODUCT INFORMATION	TABLET; ORAL			
	HYDROCHLOROTHIAZIDE, RESERPINE AND HYDRALAZINE HCL			
	MADISON	25MG;15MG;0.1MG	N41290 001	
				JAN 18, 1982

THIS EXAMPLE IS FOR PURPOSES OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

2.3 THERAPEUTIC EQUIVALENCE EVALUATIONS ILLUSTRATION

DRUG PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "A") UNDER AN INGREDIENT AND DOSAGE FORM HEADING ARE CONSIDERED THERAPEUTICALLY EQUIVALENT ONLY TO OTHER PRODUCTS CODED **AB** (OR ANY CODE BEGINNING WITH AN "A") AND **NOT** TO THOSE CODED **BP** (OR ANY CODE BEGINNING WITH "B") AND ANY PRODUCTS NOT LISTED. DRUG PRODUCTS CODED **BP** (OR ANY CODE BEGINNING WITH A "B") ARE **NOT** CONSIDERED THERAPEUTICALLY EQUIVALENT TO ANY OTHER PRODUCT. FOR A COMPLETE EXPLANATION OF THE **TE** CODES REFER TO SECTION 1.7 OF THE INTRODUCTION.



NOTE: BOLD FONT AND UNDERLINING DENOTES MULTISOURCE PRODUCTS WHICH ARE CONSIDERED THERAPEUTICALLY EQUIVALENT.

THIS EXAMPLE IS FOR PURPOSES OF ILLUSTRATION ONLY. IT DOES NOT REPRESENT ACTUAL PRODUCTS FROM THE PRESCRIPTION DRUG PRODUCT LIST.

PRESCRIPTION DRUG PRODUCT LIST

3 - 1 (of 360)

ABACAVIR SULFATE

SOLUTION; ORAL					
ZIAGEN					
+ GLAXOSMITHKLINE	EQ 20MG BASE/ML	N20978	001	Dec 17, 1998	
TABLET; ORAL					
ZIAGEN					
+ GLAXOSMITHKLINE	EQ 300MG BASE	N20977	001	Dec 17, 1998	

ABACAVIR SULFATE; LAMIVUDINE

TABLET; ORAL					
EPZICOM					
+ SMITHKLINE BEECHAM	EQ 600MG BASE;300MG	N21652	001	Aug 02, 2004	

ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL					
TRIZIVIR					
+ GLAXOSMITHKLINE	EQ 300MG BASE;150MG;300MG	N21205	001	Nov 14, 2000	

ABARELIX

INJECTABLE; INTRAMUSCULAR					
PLENAXIS					
+ PRAECIS	100MG/VIAL	N21320	001	Nov 25, 2003	

ACAMPROSATE CALCIUM

TABLET, DELAYED RELEASE; ORAL					
CAMPRAL					
+ FOREST LABS	333MG	N21431	001	Jul 29, 2004	

ACARBOSE

TABLET; ORAL					
PRECOSE					
BAYER PHARMS	25MG	N20482	004	May 29, 1997	
	50MG	N20482	001	Sep 06, 1995	
+	100MG	N20482	002	Sep 06, 1995	

ACEBUTOLOL HYDROCHLORIDE

CAPSULE; ORAL

ACEBUTOLOL HYDROCHLORIDE

<u>AB</u>	ALPHAPHARM	<u>EQ 200MG BASE</u>	<u>N75047</u>	<u>001</u>	Dec 30, 1999
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>N75047</u>	<u>002</u>	Dec 30, 1999
<u>AB</u>	MYLAN	<u>EQ 200MG BASE</u>	<u>N74288</u>	<u>001</u>	Apr 24, 1995
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>N74288</u>	<u>002</u>	Apr 24, 1995
<u>AB</u>	WATSON LABS	<u>EQ 200MG BASE</u>	<u>N74007</u>	<u>001</u>	Oct 18, 1995
<u>AB</u>		<u>EQ 400MG BASE</u>	<u>N74007</u>	<u>002</u>	Oct 18, 1995

SECTRAL

<u>AB</u>	ESP PHARMA	<u>EQ 200MG BASE</u>	<u>N18917</u>	<u>001</u>	Dec 28, 1984
<u>AB</u>	+	<u>EQ 400MG BASE</u>	<u>N18917</u>	<u>003</u>	Dec 28, 1984

ACETAMINOPHEN; ASPIRIN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE

+ MIKART	150MG;180MG;30MG	N81096	001	Oct 26, 1990	
----------	------------------	--------	-----	--------------	--

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

BUCET

<u>AB</u>	MALLINCKRODT	<u>650MG;50MG</u>	<u>N88991</u>	<u>001</u>	Jun 28, 1985
	<u>PHRENILIN FORTE</u>				
<u>AB</u>	+	<u>650MG;50MG</u>	<u>N88831</u>	<u>001</u>	Jun 19, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 2 (of 360)

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

TENCON

<u>AB</u>	MALLINCKRODT	<u>650MG;50MG</u>	<u>N89405</u>	<u>001</u>	May 15, 1990
-----------	--------------	-------------------	---------------	------------	--------------

TABLET; ORAL

BUTAPAP

<u>AB</u>	MIKART	<u>325MG;50MG</u>	<u>N89987</u>	<u>001</u>	Oct 26, 1992
-----------	--------	-------------------	---------------	------------	--------------

<u>AB</u>		<u>650MG;50MG</u>	<u>N89988</u>	<u>001</u>	Oct 26, 1992
-----------	--	-------------------	---------------	------------	--------------

PHRENILIN

<u>AB</u>	+ VALEANT	<u>325MG;50MG</u>	<u>N87811</u>	<u>001</u>	Jun 19, 1985
-----------	-----------	-------------------	---------------	------------	--------------

SEDAPAP

<u>AB</u>	+ MAYRAND	<u>650MG;50MG</u>	<u>N88944</u>	<u>001</u>	Oct 17, 1985
-----------	-----------	-------------------	---------------	------------	--------------

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

ACETAMINOPHEN, BUTALBITAL, AND CAFFEINE

<u>AB</u>	+ MIKART	<u>325MG;50MG;40MG</u>	<u>N89007</u>	<u>001</u>	Mar 17, 1986
-----------	----------	------------------------	---------------	------------	--------------

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AB</u>	WEST WARD	<u>500MG;50MG;40MG</u>	<u>N40261</u>	<u>001</u>	Oct 28, 1998
-----------	-----------	------------------------	---------------	------------	--------------

ESGIC-PLUS

<u>AB</u>	+ MIKART	<u>500MG;50MG;40MG</u>	<u>N40085</u>	<u>001</u>	Mar 28, 1996
-----------	----------	------------------------	---------------	------------	--------------

SOLUTION; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

	+ MIKART	325MG/15ML;50MG/15ML;40MG/15ML	N40387	001	Jan 31, 2003
--	----------	--------------------------------	--------	-----	--------------

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

<u>AB</u>	MALLINCKRODT	<u>325MG;50MG;40MG</u>	<u>N87804</u>	<u>001</u>	Jan 24, 1985
-----------	--------------	------------------------	---------------	------------	--------------

<u>AB</u>	MIKART	<u>325MG;50MG;40MG</u>	<u>N89175</u>	<u>001</u>	Jan 21, 1987
-----------	--------	------------------------	---------------	------------	--------------

	+	750MG;50MG;40MG	N40496	001	Dec 23, 2003
--	---	-----------------	--------	-----	--------------

<u>AB</u>	MUTUAL PHARM	<u>325MG;50MG;40MG</u>	<u>N40601</u>	<u>001</u>	Jul 29, 2005
-----------	--------------	------------------------	---------------	------------	--------------

<u>AB</u>	VINTAGE PHARMS	<u>325MG;50MG;40MG</u>	<u>N40511</u>	<u>001</u>	Aug 27, 2003
-----------	----------------	------------------------	---------------	------------	--------------

<u>AB</u>		<u>500MG;50MG;40MG</u>	<u>N40513</u>	<u>001</u>	Aug 25, 2003
-----------	--	------------------------	---------------	------------	--------------

<u>AB</u>	WATSON LABS	<u>500MG;50MG;40MG</u>	<u>N40267</u>	<u>001</u>	Jul 30, 1998
-----------	-------------	------------------------	---------------	------------	--------------

<u>AB</u>	WEST WARD	<u>325MG;50MG;40MG</u>	<u>N89718</u>	<u>001</u>	Jun 12, 1995
-----------	-----------	------------------------	---------------	------------	--------------

<u>AB</u>		<u>500MG;50MG;40MG</u>	<u>N40336</u>	<u>001</u>	Aug 18, 1999
-----------	--	------------------------	---------------	------------	--------------

BUTALBITAL, APAP, AND CAFFEINE

<u>AB</u>	WATSON LABS	<u>325MG;50MG;40MG</u>	<u>N89536</u>	<u>001</u>	Feb 16, 1988
-----------	-------------	------------------------	---------------	------------	--------------

ESGIC-PLUS

<u>AB</u>	+ MIKART	<u>500MG;50MG;40MG</u>	<u>N89451</u>	<u>001</u>	May 23, 1988
-----------	----------	------------------------	---------------	------------	--------------

FIORICET

<u>AB</u>	+ WATSON PHARMS	<u>325MG;50MG;40MG</u>	<u>N88616</u>	<u>001</u>	Nov 09, 1984
-----------	-----------------	------------------------	---------------	------------	--------------

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN, BUTALBITAL, CAFFEINE, AND CODEINE PHOSPHATE

<u>AB</u>	VINTAGE PHARMS	<u>325MG;50MG;40MG;30MG</u>	<u>N75929</u>	<u>001</u>	Apr 22, 2002
-----------	----------------	-----------------------------	---------------	------------	--------------

BUTALBITAL, ACETAMINOPHEN, AND CAFFEINE WITH CODEINE PHOSPHATE

<u>AB</u>	WEST WARD	<u>325MG;50MG;40MG;30MG</u>	<u>N75618</u>	<u>001</u>	Mar 23, 2001
-----------	-----------	-----------------------------	---------------	------------	--------------

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE

<u>AB</u>	ANABOLIC LABS	<u>325MG;50MG;40MG;30MG</u>	<u>N76560</u>	<u>001</u>	Jun 10, 2004
-----------	---------------	-----------------------------	---------------	------------	--------------

FIORICET W/ CODEINE

<u>AB</u>	+ WATSON PHARMS	<u>325MG;50MG;40MG;30MG</u>	<u>N20232</u>	<u>001</u>	Jul 30, 1992
-----------	-----------------	-----------------------------	---------------	------------	--------------

PHRENILIN WITH CAFFEINE AND CODEINE

<u>AB</u>	VALEANT	<u>325MG;50MG;40MG;30MG</u>	<u>N74911</u>	<u>001</u>	Aug 22, 2001
-----------	---------	-----------------------------	---------------	------------	--------------

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

	+ MIKART	356.4MG;30MG;16MG	N40109	001	Aug 26, 1997
--	----------	-------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 3 (of 360)

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

TABLET; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

+ MIKART

712.8MG;60MG;32MG

N40316 001

Apr 28, 1999

ACETAMINOPHEN; CODEINE PHOSPHATE

SOLUTION; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA + ALPHARMA US PHARMS 120MG/5ML;12MG/5ML

N85861 001

AA CLONMEL 120MG/5ML;12MG/5ML

N40098 001

Sep 20, 1996

AA HI TECH PHARMA 120MG/5ML;12MG/5ML

N40119 001

Apr 26, 1996

AA MIKART 120MG/5ML;12MG/5ML

N89450 001

Oct 27, 1992

AA MORTON GROVE 120MG/5ML;12MG/5ML

N87006 001

AA PHARM ASSOC 120MG/5ML;12MG/5ML

N87508 001

ACETAMINOPHEN W/ CODEINE

AA ROXANE 120MG/5ML;12MG/5ML

N86366 001

SUSPENSION; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA + VALEANT 120MG/5ML;12MG/5ML

N86024 001

CAPITAL AND CODEINE

AA ALPHARMA US PHARMS 120MG/5ML;12MG/5ML

N85883 001

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

AA ANDRX PHARMS 300MG;15MG

N40443 001

Jan 22, 2003

AA 300MG;30MG

N40443 002

Jan 22, 2003

AA 300MG;60MG

N40443 003

Jan 22, 2003

AA + DURAMED PHARMS BARR 300MG;15MG

N40223 001

Nov 18, 1997

AA 300MG;30MG

N40223 002

Nov 18, 1997

AA 300MG;60MG

N40223 003

Nov 18, 1997

AA MALLINCKRODT 300MG;15MG

N40419 001

May 31, 2001

AA 300MG;30MG

N40419 002

May 31, 2001

AA 300MG;60MG

N40419 003

May 31, 2001

AA MIKART 300MG;30MG

N89238 001

Feb 25, 1986

+ 650MG;30MG

N89231 001

Mar 03, 1986

+ 650MG;60MG

N89363 001

Sep 09, 1991

AA MUTUAL PHARM 300MG;15MG

N89671 001

Feb 10, 1988

AA 300MG;30MG

N89672 001

Feb 10, 1988

AA PHARMERAL 300MG;30MG

N87762 001

Dec 10, 1982

AA RANBAXY 300MG;60MG

N87083 001

AA SANDOZ 300MG;30MG

N81250 001

Jul 16, 1992

AA 300MG;60MG

N81249 001

Jul 16, 1992

AA TEVA 300MG;15MG

N88627 001

Mar 06, 1985

AA 300MG;30MG

N88628 001

Mar 06, 1985

AA 300MG;60MG

N88629 001

Mar 06, 1985

AA VINTAGE 300MG;15MG

N89990 001

Sep 30, 1988

AA VINTAGE PHARMS 300MG;30MG

N89805 001

Sep 30, 1988

AA 300MG;60MG

N89828 001

Sep 30, 1988

AA WATSON LABS 300MG;15MG

N89997 001

Dec 28, 1994

AA 300MG;30MG

N89998 001

Dec 28, 1994

AA 300MG;60MG

N89999 001

Dec 28, 1994

ACETAMINOPHEN W/ CODEINE NO. 3

AA ROXANE 300MG;30MG

N84656 001

ACETAMINOPHEN W/ CODEINE PHOSPHATE #3

AA RANBAXY 300MG;30MG

N85868 001

CODRIX

+ ANDRX PHARMS 500MG;15MG

N40447 001

Feb 26, 2003

+ 500MG;30MG

N40441 001

Mar 27, 2003

+ 500MG;60MG

N40488 001

Mar 28, 2003

TYLENOL W/ CODEINE NO. 3

AA + ORTHO MCNEIL PHARM 300MG;30MG

N85055 003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 4 (of 360)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

ALLAY

<u>AA</u>	IVAX PHARMS	<u>500MG;5MG</u>	<u>N89907</u>	<u>001</u>	Jan 13, 1989
-----------	-------------	------------------	---------------	------------	--------------

HYDROCET

<u>AA</u>	MALLINCKRODT	<u>500MG;5MG</u>	<u>N89006</u>	<u>001</u>	Aug 09, 1985
-----------	--------------	------------------	---------------	------------	--------------

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	MALLINCKRODT	<u>500MG;5MG</u>	<u>N88956</u>	<u>001</u>	Jul 19, 1985
-----------	--------------	------------------	---------------	------------	--------------

<u>AA</u>	MIKART	<u>500MG;5MG</u>	<u>N81067</u>	<u>001</u>	Nov 30, 1989
-----------	--------	------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;5MG</u>	<u>N81068</u>	<u>001</u>	Nov 30, 1989
-----------	--	------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;5MG</u>	<u>N81069</u>	<u>001</u>	Nov 30, 1989
-----------	--	------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;5MG</u>	<u>N81070</u>	<u>001</u>	Nov 30, 1989
-----------	--	------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;5MG</u>	<u>N89008</u>	<u>001</u>	Feb 21, 1986
-----------	--	------------------	---------------	------------	--------------

LORCET-HD

<u>AA</u>	+ MALLINCKRODT	<u>500MG;5MG</u>	<u>N87336</u>	<u>001</u>	Jul 08, 1982
-----------	----------------	------------------	---------------	------------	--------------

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	KV PHARM	<u>500MG/15ML;7.5MG/15ML</u>	<u>N40366</u>	<u>001</u>	Jan 23, 2002
-----------	----------	------------------------------	---------------	------------	--------------

<u>AA</u>	MALLINCKRODT	<u>500MG/15ML;7.5MG/15ML</u>	<u>N40418</u>	<u>001</u>	Jun 27, 2001
-----------	--------------	------------------------------	---------------	------------	--------------

<u>AA</u>	+	<u>500MG/15ML;10MG/15ML</u>	<u>N40508</u>	<u>001</u>	Aug 29, 2003
-----------	---	-----------------------------	---------------	------------	--------------

<u>AA</u>	MIKART	<u>500MG/15ML;5MG/15ML</u>	<u>N81226</u>	<u>001</u>	Oct 27, 1992
-----------	--------	----------------------------	---------------	------------	--------------

<u>AA</u>	+	<u>500MG/15ML;5MG/15ML</u>	<u>N89557</u>	<u>001</u>	Apr 29, 1992
-----------	---	----------------------------	---------------	------------	--------------

<u>AA</u>	+	<u>500MG/15ML;7.5MG/15ML</u>	<u>N81051</u>	<u>001</u>	Aug 28, 1992
-----------	---	------------------------------	---------------	------------	--------------

	+	325MG/15ML;7.5MG/15ML	N40482	001	Sep 25, 2003
--	---	-----------------------	--------	-----	--------------

<u>AA</u>	PHARM ASSOC	<u>500MG/15ML;7.5MG/15ML</u>	<u>N40182</u>	<u>001</u>	Mar 13, 1998
-----------	-------------	------------------------------	---------------	------------	--------------

<u>AA</u>	VINTAGE PHARMS	<u>500MG/15ML;7.5MG/15ML</u>	<u>N40520</u>	<u>001</u>	Oct 30, 2003
-----------	----------------	------------------------------	---------------	------------	--------------

TABLET; ORAL

ANEXSIA

<u>AA</u>	MALLINCKRODT	<u>500MG;5MG</u>	<u>N89160</u>	<u>001</u>	Apr 23, 1987
-----------	--------------	------------------	---------------	------------	--------------

<u>AA</u>		<u>750MG;10MG</u>	<u>N40468</u>	<u>001</u>	Oct 31, 2002
-----------	--	-------------------	---------------	------------	--------------

ANEXSIA 5/325

<u>AA</u>	MALLINCKRODT	<u>325MG;5MG</u>	<u>N40409</u>	<u>001</u>	Oct 20, 2000
-----------	--------------	------------------	---------------	------------	--------------

ANEXSIA 7.5/325

<u>AA</u>	MALLINCKRODT	<u>325MG;7.5MG</u>	<u>N40405</u>	<u>001</u>	Sep 08, 2000
-----------	--------------	--------------------	---------------	------------	--------------

ANEXSIA 7.5/650

<u>AA</u>	MALLINCKRODT	<u>650MG;7.5MG</u>	<u>N89725</u>	<u>001</u>	Sep 30, 1987
-----------	--------------	--------------------	---------------	------------	--------------

CO-GESIC

<u>AA</u>	SCHWARZ PHARMA	<u>500MG;5MG</u>	<u>N87757</u>	<u>001</u>	May 03, 1982
-----------	----------------	------------------	---------------	------------	--------------

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

<u>AA</u>	ANDRX PHARMS	<u>500MG;5MG</u>	<u>N40493</u>	<u>001</u>	May 28, 2003
-----------	--------------	------------------	---------------	------------	--------------

<u>AA</u>		<u>660MG;10MG</u>	<u>N40495</u>	<u>001</u>	May 28, 2003
-----------	--	-------------------	---------------	------------	--------------

<u>AA</u>		<u>750MG;7.5MG</u>	<u>N40494</u>	<u>001</u>	May 28, 2003
-----------	--	--------------------	---------------	------------	--------------

<u>AA</u>	BARR	<u>500MG;2.5MG</u>	<u>N40307</u>	<u>001</u>	Jul 26, 2000
-----------	------	--------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;5MG</u>	<u>N40308</u>	<u>001</u>	Jul 26, 2000
-----------	--	------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;7.5MG</u>	<u>N40307</u>	<u>002</u>	Jul 26, 2000
-----------	--	--------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;10MG</u>	<u>N40309</u>	<u>001</u>	Jul 26, 2000
-----------	--	-------------------	---------------	------------	--------------

<u>AA</u>		<u>650MG;7.5MG</u>	<u>N40307</u>	<u>003</u>	Jul 26, 2000
-----------	--	--------------------	---------------	------------	--------------

<u>AA</u>		<u>650MG;10MG</u>	<u>N40307</u>	<u>004</u>	Jul 26, 2000
-----------	--	-------------------	---------------	------------	--------------

<u>AA</u>		<u>750MG;7.5MG</u>	<u>N40308</u>	<u>002</u>	Jul 26, 2000
-----------	--	--------------------	---------------	------------	--------------

<u>AA</u>	ENDO PHARMS	<u>500MG;5MG</u>	<u>N40281</u>	<u>001</u>	Sep 30, 1998
-----------	-------------	------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;7.5MG</u>	<u>N40280</u>	<u>001</u>	Sep 30, 1998
-----------	--	--------------------	---------------	------------	--------------

<u>AA</u>		<u>650MG;7.5MG</u>	<u>N40280</u>	<u>002</u>	Sep 30, 1998
-----------	--	--------------------	---------------	------------	--------------

<u>AA</u>		<u>650MG;10MG</u>	<u>N40280</u>	<u>003</u>	Sep 30, 1998
-----------	--	-------------------	---------------	------------	--------------

<u>AA</u>		<u>750MG;7.5MG</u>	<u>N40281</u>	<u>002</u>	Sep 30, 1998
-----------	--	--------------------	---------------	------------	--------------

	+	400MG;5MG	N40288	001	Nov 27, 1998
--	---	-----------	--------	-----	--------------

	+	400MG;7.5MG	N40288	002	Nov 27, 1998
--	---	-------------	--------	-----	--------------

	+	400MG;10MG	N40288	003	Nov 27, 1998
--	---	------------	--------	-----	--------------

<u>AA</u>	IVAX PHARMS	<u>500MG;5MG</u>	<u>N89696</u>	<u>001</u>	Apr 21, 1988
-----------	-------------	------------------	---------------	------------	--------------

<u>AA</u>	MALLINCKRODT	<u>325MG;10MG</u>	<u>N40400</u>	<u>001</u>	Jul 26, 2000
-----------	--------------	-------------------	---------------	------------	--------------

<u>AA</u>		<u>500MG;5MG</u>	<u>N40084</u>	<u>002</u>	Jun 01, 1995
-----------	--	------------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 5 (of 360)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL				
HYDROCODONE BITARTRATE AND ACETAMINOPHEN				
<u>AA</u>	MALLINCKRODT	<u>500MG; 7.5MG</u>	<u>N40201</u>	<u>001</u> Feb 27, 1998
<u>AA</u>		<u>500MG; 10MG</u>	<u>N40201</u>	<u>002</u> Feb 27, 1998
<u>AA</u>		<u>650MG; 10MG</u>	<u>N40084</u>	<u>004</u> Oct 16, 1996
<u>AA</u>	+	<u>660MG; 10MG</u>	<u>N40084</u>	<u>003</u> Jul 29, 1996
<u>AA</u>		<u>750MG; 7.5MG</u>	<u>N40084</u>	<u>001</u> Jun 01, 1995
<u>AA</u>	MIKART	<u>325MG; 7.5MG</u>	<u>N40432</u>	<u>001</u> Jan 22, 2003
<u>AA</u>	+	<u>500MG; 2.5MG</u>	<u>N89698</u>	<u>001</u> Aug 25, 1989
<u>AA</u>		<u>500MG; 5MG</u>	<u>N89271</u>	<u>001</u> Jul 16, 1986
<u>AA</u>		<u>500MG; 5MG</u>	<u>N89697</u>	<u>001</u> Jan 28, 1992
<u>AA</u>	+	<u>500MG; 7.5MG</u>	<u>N89699</u>	<u>001</u> Aug 25, 1989
<u>AA</u>	+	<u>650MG; 7.5MG</u>	<u>N89689</u>	<u>001</u> Jun 29, 1988
<u>AA</u>	+	<u>650MG; 10MG</u>	<u>N81223</u>	<u>001</u> May 29, 1992
<u>AA</u>	+	<u>300MG; 10MG</u>	<u>N40556</u>	<u>001</u> Jun 23, 2004
<u>AA</u>	MUTUAL PHARM	<u>500MG; 5MG</u>	<u>N40236</u>	<u>001</u> Sep 25, 1997
<u>AA</u>		<u>650MG; 7.5MG</u>	<u>N40240</u>	<u>002</u> Nov 26, 1997
<u>AA</u>		<u>650MG; 10MG</u>	<u>N40240</u>	<u>001</u> Nov 26, 1997
<u>AA</u>		<u>750MG; 7.5MG</u>	<u>N40236</u>	<u>002</u> Sep 25, 1997
<u>AA</u>	SANDOZ	<u>500MG; 5MG</u>	<u>N40149</u>	<u>001</u> Jan 27, 1997
<u>AA</u>		<u>750MG; 7.5MG</u>	<u>N40149</u>	<u>002</u> Jan 27, 1997
<u>AA</u>	UCB	<u>500MG; 10MG</u>	<u>N40210</u>	<u>001</u> Aug 13, 1997
<u>AA</u>		<u>650MG; 7.5MG</u>	<u>N40134</u>	<u>001</u> Nov 21, 1996
<u>AA</u>	VINTAGE PHARMS	<u>325MG; 10MG</u>	<u>N40355</u>	<u>001</u> May 31, 2000
<u>AA</u>		<u>500MG; 2.5MG</u>	<u>N40144</u>	<u>002</u> Apr 25, 1997
<u>AA</u>		<u>500MG; 5MG</u>	<u>N89831</u>	<u>001</u> Sep 07, 1988
<u>AA</u>		<u>500MG; 5MG</u>	<u>N89971</u>	<u>001</u> Dec 02, 1988
<u>AA</u>		<u>500MG; 7.5MG</u>	<u>N40144</u>	<u>001</u> Feb 22, 1996
<u>AA</u>		<u>500MG; 10MG</u>	<u>N40356</u>	<u>001</u> May 31, 2000
<u>AA</u>		<u>650MG; 7.5MG</u>	<u>N40155</u>	<u>001</u> Apr 14, 1997
<u>AA</u>		<u>650MG; 10MG</u>	<u>N40143</u>	<u>001</u> Feb 22, 1996
<u>AA</u>		<u>660MG; 10MG</u>	<u>N40358</u>	<u>001</u> May 31, 2000
<u>AA</u>		<u>750MG; 7.5MG</u>	<u>N40157</u>	<u>001</u> Apr 12, 1996
<u>AA</u>	WATSON LABS	<u>325MG; 10MG</u>	<u>N40248</u>	<u>002</u> Apr 28, 2000
<u>AA</u>		<u>500MG; 2.5MG</u>	<u>N40123</u>	<u>003</u> Mar 04, 1996
<u>AA</u>		<u>500MG; 2.5MG</u>	<u>N81079</u>	<u>001</u> Aug 30, 1991
<u>AA</u>		<u>500MG; 5MG</u>	<u>N40122</u>	<u>001</u> Mar 04, 1996
<u>AA</u>		<u>500MG; 5MG</u>	<u>N89883</u>	<u>001</u> Dec 01, 1988
<u>AA</u>		<u>500MG; 7.5MG</u>	<u>N40123</u>	<u>004</u> Mar 04, 1996
<u>AA</u>		<u>500MG; 7.5MG</u>	<u>N81080</u>	<u>001</u> Aug 30, 1991
<u>AA</u>		<u>500MG; 10MG</u>	<u>N40148</u>	<u>002</u> Feb 14, 1997
<u>AA</u>		<u>650MG; 7.5MG</u>	<u>N40094</u>	<u>001</u> Sep 29, 1995
<u>AA</u>		<u>650MG; 7.5MG</u>	<u>N40123</u>	<u>001</u> Mar 04, 1996
<u>AA</u>		<u>650MG; 10MG</u>	<u>N40094</u>	<u>002</u> Sep 29, 1995
<u>AA</u>		<u>650MG; 10MG</u>	<u>N40123</u>	<u>002</u> Mar 04, 1996
<u>AA</u>		<u>660MG; 10MG</u>	<u>N40094</u>	<u>003</u> Aug 08, 2000
<u>AA</u>		<u>750MG; 7.5MG</u>	<u>N40122</u>	<u>002</u> Mar 04, 1996
<u>AA</u>		<u>750MG; 7.5MG</u>	<u>N81083</u>	<u>001</u> Aug 30, 1991
<u>AA</u>	+	<u>750MG; 10MG</u>	<u>N40094</u>	<u>004</u> Mar 22, 1999
<u>HY - PHEN</u>				
<u>AA</u>	ASCHER	<u>500MG; 5MG</u>	<u>N87677</u>	<u>001</u> May 03, 1982
<u>LORTAB</u>				
<u>AA</u>	MALLINCKRODT	<u>500MG; 5MG</u>	<u>N87722</u>	<u>001</u> Jul 09, 1982
<u>AA</u>	+	<u>500MG; 10MG</u>	<u>N40100</u>	<u>001</u> Jan 26, 1996
<u>NORCO</u>				
<u>AA</u>	+	<u>325MG; 5MG</u>	<u>N40099</u>	<u>001</u> Jun 25, 1997
<u>AA</u>	+	<u>325MG; 7.5MG</u>	<u>N40148</u>	<u>003</u> Sep 12, 2000
<u>AA</u>	+	<u>325MG; 10MG</u>	<u>N40148</u>	<u>001</u> Feb 14, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 6 (of 360)

ACETAMINOPHEN; HYDROCODONE BITARTRATE

TABLET; ORAL

VICODIN

<u>AA</u>	+	ABBOTT	<u>500MG;5MG</u>	<u>N88058</u>	<u>001</u>	Jan 07, 1983
-----------	---	--------	------------------	---------------	------------	--------------

VICODIN ES

<u>AA</u>	+	ABBOTT	<u>750MG;7.5MG</u>	<u>N89736</u>	<u>001</u>	Dec 09, 1988
-----------	---	--------	--------------------	---------------	------------	--------------

VICODIN HP

<u>AA</u>		ABBOTT	<u>660MG;10MG</u>	<u>N40117</u>	<u>001</u>	Sep 23, 1996
-----------	--	--------	-------------------	---------------	------------	--------------

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>		AMIDE PHARM	<u>500MG;5MG</u>	<u>N40199</u>	<u>001</u>	Dec 30, 1998
-----------	--	-------------	------------------	---------------	------------	--------------

<u>AA</u>		BARR	<u>500MG;5MG</u>	<u>N40304</u>	<u>001</u>	Oct 02, 2000
-----------	--	------	------------------	---------------	------------	--------------

<u>AA</u>		DURAMED PHARMS BARR	<u>500MG;5MG</u>	<u>N40289</u>	<u>001</u>	Mar 16, 1999
-----------	--	---------------------	------------------	---------------	------------	--------------

<u>AA</u>		ENDO PHARMS	<u>500MG;5MG</u>	<u>N40303</u>	<u>001</u>	Dec 30, 1999
-----------	--	-------------	------------------	---------------	------------	--------------

<u>AA</u>		MALLINCKRODT	<u>500MG;5MG</u>	<u>N40257</u>	<u>001</u>	Aug 04, 1998
-----------	--	--------------	------------------	---------------	------------	--------------

<u>AA</u>		MUTUAL PHARM	<u>500MG;5MG</u>	<u>N40219</u>	<u>001</u>	Jan 22, 1998
-----------	--	--------------	------------------	---------------	------------	--------------

<u>AA</u>		VINTAGE PHARMS	<u>500MG;5MG</u>	<u>N40106</u>	<u>001</u>	Jul 30, 1996
-----------	--	----------------	------------------	---------------	------------	--------------

<u>AA</u>		WATSON LABS	<u>500MG;5MG</u>	<u>N40234</u>	<u>001</u>	Oct 30, 1997
-----------	--	-------------	------------------	---------------	------------	--------------

ROXILOX

<u>AA</u>		ROXANE	<u>500MG;5MG</u>	<u>N40061</u>	<u>001</u>	Jul 03, 1995
-----------	--	--------	------------------	---------------	------------	--------------

TYLOX

<u>AA</u>	+	ORTHO MCNEIL PHARM	<u>500MG;5MG</u>	<u>N88790</u>	<u>001</u>	Dec 12, 1984
-----------	---	--------------------	------------------	---------------	------------	--------------

SOLUTION; ORAL

ROXICET

	+	ROXANE	325MG/5ML;5MG/5ML	N89351	001	Dec 03, 1986
--	---	--------	-------------------	--------	-----	--------------

TABLET; ORAL

OXYCET

<u>AA</u>		MALLINCKRODT	<u>325MG;5MG</u>	<u>N87463</u>	<u>001</u>	Dec 07, 1983
-----------	--	--------------	------------------	---------------	------------	--------------

OXYCODONE AND ACETAMINOPHEN

<u>AA</u>		AMIDE PHARM	<u>325MG;5MG</u>	<u>N40203</u>	<u>001</u>	Mar 15, 1999
-----------	--	-------------	------------------	---------------	------------	--------------

<u>AA</u>		BARR	<u>325MG;5MG</u>	<u>N87406</u>	<u>001</u>	
-----------	--	------	------------------	---------------	------------	--

<u>AA</u>		DURAMED PHARMS BARR	<u>325MG;5MG</u>	<u>N40272</u>	<u>001</u>	Jun 30, 1998
-----------	--	---------------------	------------------	---------------	------------	--------------

<u>AA</u>		MALLINCKRODT	<u>325MG;7.5MG</u>	<u>N40545</u>	<u>001</u>	Jun 30, 2004
-----------	--	--------------	--------------------	---------------	------------	--------------

<u>AA</u>			<u>325MG;10MG</u>	<u>N40545</u>	<u>002</u>	Jun 30, 2004
-----------	--	--	-------------------	---------------	------------	--------------

<u>AA</u>			<u>500MG;7.5MG</u>	<u>N40550</u>	<u>001</u>	Jun 30, 2004
-----------	--	--	--------------------	---------------	------------	--------------

<u>AA</u>			<u>650MG;10MG</u>	<u>N40550</u>	<u>002</u>	Jun 30, 2004
-----------	--	--	-------------------	---------------	------------	--------------

	+	MIKART	300MG;2.5MG	N40608	001	Dec 30, 2005
--	---	--------	-------------	--------	-----	--------------

	+		300MG;5MG	N40608	002	Dec 30, 2005
--	---	--	-----------	--------	-----	--------------

	+		300MG;7.5MG	N40608	003	Dec 30, 2005
--	---	--	-------------	--------	-----	--------------

	+		300MG;10MG	N40608	004	Dec 30, 2005
--	---	--	------------	--------	-----	--------------

<u>AA</u>		VINTAGE PHARMS	<u>325MG;5MG</u>	<u>N40105</u>	<u>001</u>	Jul 30, 1996
-----------	--	----------------	------------------	---------------	------------	--------------

<u>AA</u>		WATSON LABS	<u>325MG;5MG</u>	<u>N40171</u>	<u>001</u>	Oct 30, 1997
-----------	--	-------------	------------------	---------------	------------	--------------

<u>AA</u>			<u>325MG;7.5MG</u>	<u>N40535</u>	<u>001</u>	Sep 05, 2003
-----------	--	--	--------------------	---------------	------------	--------------

<u>AA</u>			<u>325MG;10MG</u>	<u>N40535</u>	<u>002</u>	Sep 05, 2003
-----------	--	--	-------------------	---------------	------------	--------------

<u>AA</u>			<u>500MG;7.5MG</u>	<u>N40371</u>	<u>001</u>	Dec 29, 2000
-----------	--	--	--------------------	---------------	------------	--------------

<u>AA</u>			<u>650MG;10MG</u>	<u>N40371</u>	<u>002</u>	Dec 29, 2000
-----------	--	--	-------------------	---------------	------------	--------------

PERCOCET

<u>AA</u>		ENDO PHARMS	<u>325MG;5MG</u>	<u>N40330</u>	<u>002</u>	Jun 25, 1999
-----------	--	-------------	------------------	---------------	------------	--------------

<u>AA</u>	+		<u>325MG;5MG</u>	<u>N85106</u>	<u>002</u>	
-----------	---	--	------------------	---------------	------------	--

<u>AA</u>	+		<u>325MG;7.5MG</u>	<u>N40434</u>	<u>001</u>	Nov 23, 2001
-----------	---	--	--------------------	---------------	------------	--------------

<u>AA</u>	+		<u>325MG;10MG</u>	<u>N40434</u>	<u>002</u>	Nov 23, 2001
-----------	---	--	-------------------	---------------	------------	--------------

<u>AA</u>	+		<u>500MG;7.5MG</u>	<u>N40341</u>	<u>001</u>	Jul 26, 1999
-----------	---	--	--------------------	---------------	------------	--------------

<u>AA</u>	+		<u>650MG;10MG</u>	<u>N40341</u>	<u>002</u>	Jul 26, 1999
-----------	---	--	-------------------	---------------	------------	--------------

	+		325MG;2.5MG	N40330	001	Jun 25, 1999
--	---	--	-------------	--------	-----	--------------

ROXICET

<u>AA</u>		ROXANE	<u>325MG;5MG</u>	<u>N87003</u>	<u>001</u>	
-----------	--	--------	------------------	---------------	------------	--

ROXICET 5/500

	+	ROXANE	500MG;5MG	N89775	001	Jan 12, 1989
--	---	--------	-----------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 7 (of 360)

ACETAMINOPHEN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

<u>ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE</u>				
<u>AB</u>	AMIDE PHARM	<u>650MG;EQ 25MG BASE</u>	<u>N76202</u>	<u>001</u> Aug 02, 2002
<u>PENTAZOCINE HYDROCHLORIDE AND ACETAMINOPHEN</u>				
<u>AB</u>	WATSON LABS	<u>650MG;EQ 25MG BASE</u>	<u>N74699</u>	<u>001</u> Mar 24, 2000
<u>TALACEN</u>				
<u>AB</u>	+ SANOFI SYNTHELABO	<u>650MG;EQ 25MG BASE</u>	<u>N18458</u>	<u>001</u> Sep 23, 1982

ACETAMINOPHEN; PROPOXYPHENE HYDROCHLORIDE

TABLET; ORAL

<u>PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN</u>				
<u>AA</u>	MYLAN	<u>650MG;65MG</u>	<u>N83978</u>	<u>001</u>
<u>AA</u>	SANDOZ	<u>650MG;65MG</u>	<u>N89959</u>	<u>001</u> Jul 18, 1989
<u>AA</u>	VINTAGE PHARMS	<u>650MG;65MG</u>	<u>N40507</u>	<u>001</u> Jul 30, 2003
<u>AA</u>	WATSON LABS	<u>650MG;65MG</u>	<u>N40139</u>	<u>001</u> Dec 16, 1996
<u>WYGESIC</u>				
<u>AA</u>	+ LEITNER PHARMS	<u>650MG;65MG</u>	<u>N84999</u>	<u>001</u>

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

<u>DARVOCET A500</u>				
<u>AB</u>	XANODYNE PHARM	<u>500MG;100MG</u>	<u>N76429</u>	<u>001</u> Sep 10, 2003
<u>DARVOCET-N 100</u>				
<u>AB</u>	+ XANODYNE PHARM	<u>650MG;100MG</u>	<u>N17122</u>	<u>002</u>
<u>DARVOCET-N 50</u>				
<u>AB</u>	XANODYNE PHARM	<u>325MG;50MG</u>	<u>N17122</u>	<u>001</u>
<u>PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN</u>				
<u>AB</u>	ANDRX PHARMS	<u>500MG;100MG</u>	<u>N77196</u>	<u>001</u> Jun 28, 2005
<u>AB</u>		<u>650MG;100MG</u>	<u>N76609</u>	<u>001</u> Nov 16, 2004
<u>AB</u>	IVAX PHARMS	<u>650MG;100MG</u>	<u>N70146</u>	<u>001</u> Aug 02, 1985
<u>AB</u>	MALLINCKRODT	<u>650MG;100MG</u>	<u>N75738</u>	<u>001</u> Feb 02, 2001
<u>AB</u>	MUTUAL PHARM	<u>650MG;100MG</u>	<u>N70615</u>	<u>001</u> Mar 21, 1986
<u>AB</u>		<u>650MG;100MG</u>	<u>N70771</u>	<u>001</u> Mar 21, 1986
<u>AB</u>		<u>650MG;100MG</u>	<u>N70775</u>	<u>001</u> Mar 21, 1986
<u>AB</u>	MYLAN	<u>650MG;100MG</u>	<u>N70145</u>	<u>001</u> Jun 12, 1985
<u>AB</u>		<u>650MG;100MG</u>	<u>N72195</u>	<u>001</u> Feb 16, 1988
<u>AB</u>	PUREPAC PHARM	<u>650MG;100MG</u>	<u>N70910</u>	<u>001</u> Jan 02, 1987
<u>AB</u>	SANDOZ	<u>650MG;100MG</u>	<u>N70443</u>	<u>001</u> Jan 23, 1986
<u>AB</u>	TEVA	<u>650MG;100MG</u>	<u>N74119</u>	<u>001</u> Dec 19, 1994
<u>AB</u>	VINTAGE PHARMS	<u>325MG;50MG</u>	<u>N74843</u>	<u>002</u> Feb 15, 2001
<u>AB</u>		<u>500MG;100MG</u>	<u>N76750</u>	<u>001</u> Jun 28, 2004
<u>AB</u>		<u>650MG;100MG</u>	<u>N74843</u>	<u>001</u> Feb 12, 1997
		325MG;100MG	N76743	001 May 07, 2004

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

TABLET; ORAL

<u>TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN</u>				
<u>AB</u>	CARACO	<u>325MG;37.5MG</u>	<u>N77184</u>	<u>001</u> Dec 16, 2005
<u>AB</u>	KALI LABS	<u>325MG;37.5MG</u>	<u>N76475</u>	<u>001</u> Apr 21, 2005
<u>ULTRACET</u>				
<u>AB</u>	+ ORTHO MCNEIL PHARM	<u>325MG;37.5MG</u>	<u>N21123</u>	<u>001</u> Aug 15, 2001

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL

	DIAMOX			
	+ DURAMED PHARMS BARR	500MG	N12945	003 Apr 25, 1995

TABLET; ORAL

<u>ACETAZOLAMIDE</u>				
<u>AB</u>	LANNETT	<u>250MG</u>	<u>N84840</u>	<u>001</u>

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 8 (of 360)

ACETAZOLAMIDE

TABLET; ORAL

ACETAZOLAMIDE

<u>AB</u>	MUTUAL PHARM	<u>125MG</u>	<u>N89752</u>	<u>001</u>	Jun 22, 1988
<u>AB</u>		<u>250MG</u>	<u>N89753</u>	<u>001</u>	Jun 22, 1988
<u>AB</u>	TARO	<u>125MG</u>	<u>N40195</u>	<u>001</u>	May 28, 1997
<u>AB</u>		<u>250MG</u>	<u>N40195</u>	<u>002</u>	May 28, 1997
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>N88882</u>	<u>001</u>	Oct 22, 1985
<u>DIAMOX</u>					
<u>AB</u>	DURAMED PHARMS BARR	<u>125MG</u>	<u>N08943</u>	<u>001</u>	
<u>AB</u>	+	<u>250MG</u>	<u>N08943</u>	<u>002</u>	

ACETAZOLAMIDE SODIUM

INJECTABLE; INJECTION

ACETAZOLAMIDE SODIUM

<u>AP</u>	BEDFORD	<u>EQ 500MG BASE/VIAL</u>	<u>N40089</u>	<u>001</u>	Feb 28, 1995
<u>AP</u>	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>N40108</u>	<u>001</u>	Oct 30, 1995
<u>DIAMOX</u>					
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>N09388</u>	<u>001</u>	Dec 05, 1990

ACETIC ACID, GLACIAL

SOLUTION; IRRIGATION, URETHRAL

ACETIC ACID 0.25% IN PLASTIC CONTAINER

<u>AT</u>	B BRAUN	<u>250MG/100ML</u>	<u>N18161</u>	<u>001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>250MG/100ML</u>	<u>N18523</u>	<u>001</u>	Feb 19, 1982
<u>AT</u>	HOSPIRA	<u>250MG/100ML</u>	<u>N17656</u>	<u>001</u>	

SOLUTION/DROPS; OTIC

ACETASOL

<u>AT</u>	ALPHARMA US PHARMS	<u>2%</u>	<u>N87146</u>	<u>001</u>	
-----------	--------------------	-----------	---------------	------------	--

ACETIC ACID

<u>AT</u>	+	MORTON GROVE	<u>2%</u>	<u>N40166</u>	<u>001</u>	Jul 26, 1996
<u>AT</u>		TARO	<u>2%</u>	<u>N88638</u>	<u>001</u>	Sep 06, 1984
<u>AT</u>		VINTAGE	<u>2%</u>	<u>N40607</u>	<u>001</u>	Feb 24, 2005

ACETIC ACID, GLACIAL; ALUMINUM ACETATE

SOLUTION/DROPS; OTIC

ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE

+	BAUSCH AND LOMB	<u>2%;0.79%</u>	<u>N40063</u>	<u>001</u>	Feb 25, 1994
---	-----------------	-----------------	---------------	------------	--------------

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

ACETASOL HC

<u>AT</u>	ALPHARMA US PHARMS	<u>2%;1%</u>	<u>N87143</u>	<u>001</u>	Jan 13, 1982
-----------	--------------------	--------------	---------------	------------	--------------

HYDROCORTISONE AND ACETIC ACID

<u>AT</u>	MORTON GROVE	<u>2%;1%</u>	<u>N40168</u>	<u>001</u>	Aug 30, 1996
<u>AT</u>	TARO	<u>2%;1%</u>	<u>N88759</u>	<u>001</u>	Mar 04, 1985

VOSOL HC

<u>AT</u>	+	MEDPOINTE PHARM HLC	<u>2%;1%</u>	<u>N12770</u>	<u>001</u>	
-----------	---	---------------------	--------------	---------------	------------	--

ACETOHEXAMIDE

TABLET; ORAL

ACETOHEXAMIDE

<u>AB</u>	BARR	<u>250MG</u>	<u>N70869</u>	<u>001</u>	Feb 09, 1987
<u>AB</u>	+	<u>500MG</u>	<u>N70870</u>	<u>001</u>	Feb 09, 1987
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>N71893</u>	<u>001</u>	Nov 25, 1987
<u>AB</u>		<u>500MG</u>	<u>N71894</u>	<u>001</u>	Nov 25, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 9 (of 360)

ACETOHYDROXAMIC ACID

TABLET; ORAL

LITHOSTAT

+	MISSION PHARMA	250MG	N18749	001	May 31, 1983
---	----------------	-------	--------	-----	--------------

ACETYLCHOLINE CHLORIDE

FOR SOLUTION; OPHTHALMIC

MIOCHOL

+	NOVARTIS	20MG/VIAL	N16211	001	
---	----------	-----------	--------	-----	--

MIOCHOL-E

+	NOVARTIS	20MG/VIAL	N20213	001	Sep 22, 1993
---	----------	-----------	--------	-----	--------------

ACETYLCYSTEINE

INJECTABLE; INTRAVENOUS

ACETADOTE

+	CUMBERLAND PHARMS	6GM/30ML (200MG/ML)	N21539	001	Jan 23, 2004
---	-------------------	---------------------	--------	-----	--------------

SOLUTION; INHALATION, ORAL

ACETYLCYSTEINE

<u>AN</u>	BEDFORD	<u>10%</u>	<u>N72323</u>	<u>001</u>	Apr 30, 1992
<u>AN</u>		<u>20%</u>	<u>N72324</u>	<u>001</u>	Apr 30, 1992
<u>AN</u>	HOSPIRA	<u>10%</u>	<u>N73664</u>	<u>001</u>	Aug 30, 1994
<u>AN</u>		<u>20%</u>	<u>N74037</u>	<u>001</u>	Aug 30, 1994
<u>AN</u>	LUITPOLD	<u>10%</u>	<u>N72489</u>	<u>001</u>	Jul 28, 1995
<u>AN</u>		<u>20%</u>	<u>N72547</u>	<u>001</u>	Jul 28, 1995
<u>AN</u>	MAYNE PHARMA USA	<u>10%</u>	<u>N71364</u>	<u>001</u>	May 01, 1989
<u>AN</u>		<u>10%</u>	<u>N71365</u>	<u>002</u>	Jan 18, 2002
<u>AN</u>		<u>20%</u>	<u>N71364</u>	<u>002</u>	Jan 18, 2002
<u>AN</u>		<u>20%</u>	<u>N71365</u>	<u>001</u>	May 01, 1989

MUCOMYST

<u>AN</u>	+	APOTHECON	<u>10%</u>	<u>N13601</u>	<u>002</u>
<u>AN</u>	+		<u>20%</u>	<u>N13601</u>	<u>001</u>

ACITRETIN

CAPSULE; ORAL

SORIATANE

CONNETICS	10MG	N19821	001	Oct 28, 1996
-----------	------	--------	-----	--------------

+	25MG	N19821	002	Oct 28, 1996
---	------	--------	-----	--------------

ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

SEMPREX-D

+	UCB	8MG;60MG	N19806	001	Mar 25, 1994
---	-----	----------	--------	-----	--------------

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

<u>AB</u>	AAIPHARMA LLC	<u>200MG</u>	<u>N74833</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>	APOTEX	<u>200MG</u>	<u>N75677</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>	APOTHECON	<u>200MG</u>	<u>N74889</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>	CLONMEL HLTHCARE	<u>200MG</u>	<u>N74872</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>	GENPHARM	<u>200MG</u>	<u>N74977</u>	<u>001</u>	Apr 13, 1998
<u>AB</u>	IVAX PHARMS	<u>200MG</u>	<u>N74674</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>	MYLAN	<u>200MG</u>	<u>N74727</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>	PUREPAC PHARM	<u>200MG</u>	<u>N74906</u>	<u>001</u>	Aug 26, 1997
<u>AB</u>	RANBAXY	<u>200MG</u>	<u>N74975</u>	<u>001</u>	Sep 30, 1998
<u>AB</u>	STASON	<u>200MG</u>	<u>N75090</u>	<u>001</u>	Jan 26, 1999
<u>AB</u>	TEVA	<u>200MG</u>	<u>N74578</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>		<u>200MG</u>	<u>N74828</u>	<u>001</u>	Apr 22, 1997
<u>AB</u>	TEVA PHARMS	<u>200MG</u>	<u>N74914</u>	<u>001</u>	Nov 26, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 10 (of 360)

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

<u>AB</u>	WATSON LABS	<u>200MG</u>	<u>N75101</u>	<u>001</u>	Apr 15, 1998
-----------	-------------	--------------	---------------	------------	--------------

ZOVIRAX

<u>AB</u>	+ GLAXOSMITHKLINE	<u>200MG</u>	<u>N18828</u>	<u>001</u>	Jan 25, 1985
-----------	-------------------	--------------	---------------	------------	--------------

CREAM; TOPICAL

ZOVIRAX

	+ GLAXOSMITHKLINE	5%	N21478	001	Dec 30, 2002
--	-------------------	----	--------	-----	--------------

OINTMENT; TOPICAL

ZOVIRAX

	+ GLAXOSMITHKLINE	5%	N18604	001	Mar 29, 1982
--	-------------------	----	--------	-----	--------------

SUSPENSION; ORAL

ACYCLOVIR

<u>AB</u>	ALPHARMA US PHARMS	<u>200MG/5ML</u>	<u>N74738</u>	<u>001</u>	Apr 28, 1997
-----------	--------------------	------------------	---------------	------------	--------------

<u>AB</u>	HI TECH PHARMA	<u>200MG/5ML</u>	<u>N77026</u>	<u>001</u>	Jun 07, 2005
-----------	----------------	------------------	---------------	------------	--------------

ZOVIRAX

<u>AB</u>	+ GLAXOSMITHKLINE	<u>200MG/5ML</u>	<u>N19909</u>	<u>001</u>	Dec 22, 1989
-----------	-------------------	------------------	---------------	------------	--------------

TABLET; ORAL

ACYCLOVIR

<u>AB</u>	AAIPHARMA LLC	<u>400MG</u>	<u>N74946</u>	<u>001</u>	Nov 19, 1997
-----------	---------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N74946</u>	<u>002</u>	Nov 19, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	APOTEX	<u>400MG</u>	<u>N77309</u>	<u>001</u>	Sep 29, 2005
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N77309</u>	<u>002</u>	Sep 29, 2005
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	APOTHECON	<u>400MG</u>	<u>N74891</u>	<u>001</u>	Oct 31, 1997
-----------	-----------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N74891</u>	<u>002</u>	Oct 31, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	CARLSBAD	<u>400MG</u>	<u>N75382</u>	<u>001</u>	Apr 30, 1999
-----------	----------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N75382</u>	<u>002</u>	Apr 30, 1999
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	GENPHARM	<u>400MG</u>	<u>N74976</u>	<u>001</u>	Apr 13, 1998
-----------	----------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N74976</u>	<u>002</u>	Apr 13, 1998
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	IVAX PHARMS	<u>400MG</u>	<u>N74836</u>	<u>001</u>	Apr 22, 1997
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N74836</u>	<u>002</u>	Apr 22, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>400MG</u>	<u>N75211</u>	<u>001</u>	Sep 28, 1998
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N75211</u>	<u>002</u>	Sep 28, 1998
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	PUREPAC PHARM	<u>400MG</u>	<u>N74870</u>	<u>001</u>	Jun 05, 1997
-----------	---------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N74870</u>	<u>002</u>	Jun 05, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	RANBAXY	<u>400MG</u>	<u>N74980</u>	<u>001</u>	Sep 30, 1998
-----------	---------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N74980</u>	<u>002</u>	Sep 30, 1998
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>400MG</u>	<u>N74556</u>	<u>002</u>	Apr 22, 1997
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N74556</u>	<u>003</u>	Apr 22, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	TEVA PHARMS	<u>400MG</u>	<u>N75021</u>	<u>001</u>	Mar 18, 1998
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>800MG</u>	<u>N75021</u>	<u>002</u>	Mar 18, 1998
-----------	--	--------------	---------------	------------	--------------

ZOVIRAX

<u>AB</u>	GLAXOSMITHKLINE	<u>400MG</u>	<u>N20089</u>	<u>001</u>	Apr 30, 1991
-----------	-----------------	--------------	---------------	------------	--------------

<u>AB</u>	+	<u>800MG</u>	<u>N20089</u>	<u>002</u>	Apr 30, 1991
-----------	---	--------------	---------------	------------	--------------

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR

<u>AP</u>	SICOR PHARMS	<u>EQ 50MG BASE/ML</u>	<u>N75627</u>	<u>001</u>	Mar 28, 2001
-----------	--------------	------------------------	---------------	------------	--------------

ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE

+	BAXTER HLTHCARE	EQ 500MG BASE/VIAL	N74885	001	Dec 19, 1997
---	-----------------	--------------------	--------	-----	--------------

		EQ 1GM BASE/VIAL	N74885	002	Dec 19, 1997
--	--	------------------	--------	-----	--------------

ACYCLOVIR SODIUM

<u>AP</u>	+ AM PHARM PARTNERS	<u>EQ 50MG BASE/ML</u>	<u>N74930</u>	<u>001</u>	May 13, 1998
-----------	---------------------	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>N75015</u>	<u>001</u>	Apr 30, 1998
-----------	--	---------------------------	---------------	------------	--------------

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 500MG BASE/VIAL</u>	<u>N74913</u>	<u>001</u>	Oct 15, 1997
-----------	-----------------	---------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N74913</u>	<u>002</u>	Oct 15, 1997
-----------	--	-------------------------	---------------	------------	--------------

<u>AP</u>	BEDFORD	<u>EQ 500MG BASE/VIAL</u>	<u>N74596</u>	<u>002</u>	Apr 22, 1997
-----------	---------	---------------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 11 (of 360)

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

<u>AP</u>	BEDFORD	<u>EQ 1GM BASE/VIAL</u>	<u>N74596</u>	<u>001</u>	Apr 22, 1997
<u>AP</u>	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>N74663</u>	<u>001</u>	Apr 22, 1997
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>N74758</u>	<u>001</u>	Apr 22, 1997
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N74663</u>	<u>002</u>	Apr 22, 1997
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N74758</u>	<u>002</u>	Apr 22, 1997
<u>AP</u>	MAYNE PHARMA USA	<u>EQ 50MG BASE/ML</u>	<u>N75065</u>	<u>001</u>	Feb 25, 1999
	+	EQ 25MG BASE/ML	N74720	001	Apr 22, 1997
<u>AP</u>	SICOR PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>N74969</u>	<u>001</u>	Aug 26, 1997
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N74969</u>	<u>002</u>	Aug 26, 1997
<u>ZOVIRAX</u>					
<u>AP</u>	+ GLAXOSMITHKLINE	<u>EQ 500MG BASE/VIAL</u>	<u>N18603</u>	<u>001</u>	Oct 22, 1982
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N18603</u>	<u>002</u>	Jun 29, 1989

ADAPALENECREAM; TOPICAL
DIFFERIN

+ GALDERMA LABS LP 0.1% N20748 001 May 26, 2000

GEL; TOPICAL
DIFFERIN

+ GALDERMA LABS LP 0.1% N20380 001 May 31, 1996

ADEFOVIR DIPIVOXILTABLET; ORAL
HEPSERA

+ GILEAD 10MG N21449 001 Sep 20, 2002

ADENOSINE

INJECTABLE; INJECTION

ADENOCARD

<u>AP</u>	+ ASTELLAS	<u>3MG/ML</u>	<u>N19937</u>	<u>002</u>	Oct 30, 1989
	ADENOSCAN				
	+ ASTELLAS	3MG/ML	N20059	001	May 18, 1995
<u>ADENOSINE</u>					
<u>AP</u>	AM PHARM	<u>3MG/ML</u>	<u>N77133</u>	<u>001</u>	Apr 27, 2005
<u>AP</u>	BAXTER HLTHCARE	<u>3MG/ML</u>	<u>N76500</u>	<u>001</u>	Jun 16, 2004
<u>AP</u>		<u>3MG/ML</u>	<u>N76501</u>	<u>001</u>	Jun 16, 2004
<u>AP</u>	BEDFORD	<u>3MG/ML</u>	<u>N76404</u>	<u>001</u>	Jun 16, 2004
<u>AP</u>	SICOR PHARMS	<u>3MG/ML</u>	<u>N76564</u>	<u>001</u>	Jun 16, 2004

ALBENDAZOLETABLET; ORAL
ALBENZA

+ GLAXOSMITHKLINE 200MG N20666 001 Jun 11, 1996

ALBUMIN HUMANINJECTABLE; INJECTION
OPTISON

+ GE HEALTHCARE 10MG/ML N20899 001 Dec 31, 1997

ALBUMIN IODINATED I-125 SERUMINJECTABLE; INJECTION
JEANATOPE

ISO TEX	100UCI/10ML(10UCI/ML)	N17836	003	Jun 08, 2004
	500uCi/0.5ML	N17836	001	
+	1,000uCi/ML	N17836	002	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 12 (of 360)

ALBUMIN IODINATED I-131 SERUM

INJECTABLE; INJECTION

MEGATOPE

+	ISO TEX	0.5mCi/VIAL	N17837	001	
+		1mCi/VIAL	N17837	002	

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

<u>AB</u>	ARMSTRONG PHARMS	<u>0.09MG/INH</u>	<u>N72273</u>	<u>001</u>	Aug 14, 1996
<u>AB</u>	GENPHARM	<u>0.09MG/INH</u>	<u>N73045</u>	<u>001</u>	Aug 19, 1997
<u>AB</u>	IVAX PHARMS	<u>0.09MG/INH</u>	<u>N73272</u>	<u>001</u>	Dec 28, 1995
<u>AB</u>	PLIVA	<u>0.09MG/INH</u>	<u>N74072</u>	<u>001</u>	Aug 01, 1996
	PROVENTIL				
BN	+ SCHERING	0.09MG/INH	N17559	001	
	<u>VENTOLIN</u>				
<u>AB</u>	+ GLAXOSMITHKLINE	<u>0.09MG/INH</u>	<u>N18473</u>	<u>001</u>	

ALBUTEROL SULFATE

AEROSOL, METERED; INHALATION

PROAIR HFA

BX	+ IVAX RES	EQ 0.09MG BASE/INH	N21457	001	Oct 29, 2004
	PROVENTIL-HFA				
BX	+ 3M	EQ 0.09MG BASE/INH	N20503	001	Aug 15, 1996
	VENTOLIN HFA				
BX	+ GLAXOSMITHKLINE	EQ 0.09MG BASE/INH	N20983	001	Apr 19, 2001

SOLUTION; INHALATION

ACCUNEB

<u>AN</u>	+ DEY	<u>EQ 0.042% BASE</u>	<u>N20949</u>	<u>001</u>	Apr 30, 2001
	+	EQ 0.021% BASE	N20949	002	Apr 30, 2001

ALBUTEROL SULFATE

<u>AN</u>	ALPHARMA US PHARMS	<u>EQ 0.083% BASE</u>	<u>N73533</u>	<u>001</u>	Sep 26, 1995
<u>AN</u>	+ BAUSCH AND LOMB	<u>EQ 0.083% BASE</u>	<u>N75358</u>	<u>001</u>	Mar 29, 2000
<u>AN</u>	+	<u>EQ 0.5% BASE</u>	<u>N75050</u>	<u>001</u>	Jun 18, 1998
<u>AN</u>	+ DEY	<u>EQ 0.083% BASE</u>	<u>N72652</u>	<u>001</u>	Feb 21, 1992
<u>AN</u>	HI TECH PHARMA	<u>EQ 0.083% BASE</u>	<u>N75063</u>	<u>001</u>	Feb 09, 1999
<u>AN</u>		<u>EQ 0.5% BASE</u>	<u>N74543</u>	<u>001</u>	Jan 15, 1998
<u>AN</u>	IVAX PHARMS	<u>EQ 0.083% BASE</u>	<u>N75343</u>	<u>001</u>	Nov 09, 1999
<u>AN</u>	MORTON GROVE	<u>EQ 0.083% BASE</u>	<u>N75394</u>	<u>001</u>	Nov 22, 1999
<u>AN</u>	NEPHRON	<u>EQ 0.042% BASE</u>	<u>N76355</u>	<u>001</u>	Jun 28, 2004
<u>AN</u>		<u>EQ 0.083% BASE</u>	<u>N74880</u>	<u>001</u>	Sep 17, 1997
<u>AN</u>		<u>EQ 0.5% BASE</u>	<u>N75664</u>	<u>001</u>	Jun 26, 2001
<u>AN</u>	NOVEX	<u>EQ 0.5% BASE</u>	<u>N76391</u>	<u>001</u>	Apr 01, 2003
<u>AN</u>	RESPIRARE	<u>EQ 0.083% BASE</u>	<u>N76370</u>	<u>001</u>	Nov 24, 2003
<u>AN</u>	ROXANE	<u>EQ 0.083% BASE</u>	<u>N75129</u>	<u>001</u>	Feb 13, 2001

SYRUP; ORAL

ALBUTEROL SULFATE

<u>AA</u>	ALPHARMA US PHARMS	<u>EQ 2MG BASE/5ML</u>	<u>N74454</u>	<u>001</u>	Sep 25, 1995
<u>AA</u>	HI TECH PHARMA	<u>EQ 2MG BASE/5ML</u>	<u>N74749</u>	<u>001</u>	Jan 30, 1998
<u>AA</u>	+ TEVA	<u>EQ 2MG BASE/5ML</u>	<u>N73419</u>	<u>001</u>	Mar 30, 1992
<u>AA</u>	UDL	<u>EQ 2MG BASE/5ML</u>	<u>N75262</u>	<u>001</u>	Mar 30, 1999

TABLET; ORAL

ALBUTEROL SULFATE

<u>AB</u>	MUTUAL PHARM	<u>EQ 2MG BASE</u>	<u>N72636</u>	<u>001</u>	Dec 05, 1989
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N72637</u>	<u>001</u>	Dec 05, 1989
<u>AB</u>	+ MYLAN	<u>EQ 2MG BASE</u>	<u>N72894</u>	<u>002</u>	Jan 17, 1991
<u>AB</u>	+	<u>EQ 4MG BASE</u>	<u>N72894</u>	<u>001</u>	Jan 17, 1991
<u>AB</u>	PLIVA	<u>EQ 2MG BASE</u>	<u>N72316</u>	<u>001</u>	Dec 05, 1989
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N72317</u>	<u>001</u>	Dec 05, 1989

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 13 (of 360)

ALBUTEROL SULFATE

TABLET; ORAL

ALBUTEROL SULFATE

<u>AB</u>	SANDOZ	<u>EQ 2MG BASE</u>	<u>N72151</u>	<u>001</u>	Dec 05, 1989
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N72152</u>	<u>001</u>	Dec 05, 1989
<u>AB</u>	TEVA	<u>EQ 2MG BASE</u>	<u>N72619</u>	<u>001</u>	Dec 05, 1989
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N72779</u>	<u>001</u>	Jun 25, 1993
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N72938</u>	<u>001</u>	Mar 30, 1990
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N72620</u>	<u>001</u>	Dec 05, 1989
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N72780</u>	<u>001</u>	Jun 25, 1993
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N72939</u>	<u>001</u>	Mar 30, 1990
<u>AB</u>	WATSON LABS	<u>EQ 2MG BASE</u>	<u>N72629</u>	<u>001</u>	Jan 31, 1991
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N72764</u>	<u>001</u>	Aug 28, 1991
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N72630</u>	<u>001</u>	Jan 31, 1991
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N72765</u>	<u>001</u>	Aug 28, 1991

TABLET, EXTENDED RELEASE; ORAL

VOSPIRE ER

	ODYSSEY PHARMS	EQ 4MG BASE	N76130	002	Sep 26, 2002
+		EQ 8MG BASE	N76130	003	Sep 26, 2002

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

COMBIVENT

+	BOEHRINGER INGELHEIM	EQ 0.09MG BASE/INH;0.018MG/INH	N20291	001	Oct 24, 1996
---	----------------------	--------------------------------	--------	-----	--------------

SOLUTION; INHALATION

DUONEB

+	DEY	EQ 0.083% BASE;0.017%	N20950	001	Mar 21, 2001
---	-----	-----------------------	--------	-----	--------------

ALCLOMETASONE DIPROPIONATE

CREAM; TOPICAL

ACLOVATE

<u>AB</u>	+	GLAXOSMITHKLINE	<u>0.05%</u>	<u>N18707</u>	<u>001</u>	Dec 14, 1982
-----------	---	-----------------	--------------	---------------	------------	--------------

ALCLOMETASONE DIPROPIONATE

<u>AB</u>		ALTANA	<u>0.05%</u>	<u>N76973</u>	<u>001</u>	Jul 12, 2005
<u>AB</u>		TARO	<u>0.05%</u>	<u>N76587</u>	<u>001</u>	Sep 15, 2005

OINTMENT; TOPICAL

ACLOVATE

<u>AB</u>	+	GLAXOSMITHKLINE	<u>0.05%</u>	<u>N18702</u>	<u>001</u>	Dec 14, 1982
-----------	---	-----------------	--------------	---------------	------------	--------------

ALCLOMETASONE DIPROPIONATE

<u>AB</u>		ALTANA	<u>0.05%</u>	<u>N76884</u>	<u>001</u>	Jul 18, 2005
<u>AB</u>		TARO	<u>0.05%</u>	<u>N76730</u>	<u>001</u>	Jul 29, 2004

ALCOHOL; DEXTROSE

INJECTABLE; INJECTION

ALCOHOL 10% AND DEXTROSE 5%

+	B BRAUN	10ML/100ML;5GM/100ML	N04589	006	
---	---------	----------------------	--------	-----	--

ALCOHOL 5% AND DEXTROSE 5%

<u>AP</u>	+	B BRAUN	<u>5ML/100ML;5GM/100ML</u>	<u>N04589</u>	<u>004</u>
-----------	---	---------	----------------------------	---------------	------------

ALCOHOL 5% IN D5-W

<u>AP</u>		HOSPIRA	<u>5ML/100ML;5GM/100ML</u>	<u>N83263</u>	<u>001</u>
-----------	--	---------	----------------------------	---------------	------------

ALENDRONATE SODIUM

SOLUTION; ORAL

FOSAMAX

+	MERCK	EQ 70MG BASE/75ML	N21575	001	Sep 17, 2003
---	-------	-------------------	--------	-----	--------------

TABLET; ORAL

FOSAMAX

MERCK

	EQ 5MG BASE	N20560	003	Apr 25, 1997
	EQ 10MG BASE	N20560	001	Sep 29, 1995

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 14 (of 360)

ALENDRONATE SODIUM

TABLET; ORAL

FOSAMAX

MERCK

EQ 35MG BASE

N20560 004 Oct 20, 2000

EQ 40MG BASE

N20560 002 Sep 29, 1995

+

EQ 70MG BASE

N20560 005 Oct 20, 2000

ALENDRONATE SODIUM; CHOLECALCIFEROL

TABLET; ORAL

FOSAMAX PLUS D

+ MERCK

EQ 70MG BASE;2,800 IU

N21762 001 Apr 07, 2005

ALFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ALFENTAAP + AKORNEQ 0.5MG BASE/MLN19353 001 Dec 29, 1986ALFENTANILAP HOSPIRAEQ 0.5MG BASE/MLN75221 001 Oct 28, 1999ALFUZOSIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

UROXATRAL

+ SANOFI SYN RES

10MG

N21287 001 Jun 12, 2003

ALGLUCERASE

INJECTABLE; INJECTION

CEREDASE

+ GENZYME

80 UNITS/ML

N20057 003 Apr 05, 1991

ALITRETINOIN

GEL; TOPICAL

PANRETIN

+ LIGAND

EQ 0.1% BASE

N20886 001 Feb 02, 1999

ALLOPURINOL

TABLET; ORAL

ALLOPURINOLAB APOTEX100MGN77353 001 Sep 08, 2005AB300MGN77353 002 Sep 08, 2005AB MUTUAL PHARM100MGN71449 001 Jan 09, 1987AB300MGN71450 001 Jan 09, 1987AB MYLAN100MGN18659 001 Oct 24, 1986AB300MGN18659 002 Oct 24, 1986AB PAR PHARM100MGN70150 001 Dec 10, 1985AB300MGN70147 001 Dec 10, 1985AB SANDOZ100MGN70268 001 Dec 31, 1985AB300MGN70269 001 Dec 31, 1985AB VINTAGE PHARMS100MGN75798 001 Jun 27, 2003AB300MGN75798 002 Jun 27, 2003AB WATSON LABS100MGN18832 002 Sep 28, 1984AB300MGN18877 001 Sep 28, 1984LOPURINAB BASF100MGN71586 001 Apr 02, 1987AB300MGN71587 001 Apr 02, 1987ZYLOPRIMAB PROMETHEUS LABS100MGN16084 001AB +300MGN16084 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 15 (of 360)

ALLOPURINOL SODIUM

INJECTABLE; INJECTION

ALLOPURINOL SODIUM

<u>AP</u>	BEDFORD LABS	<u>EQ 500MG BASE/VIAL</u>	<u>N76870</u>	<u>001</u>	Aug 26, 2004
-----------	--------------	---------------------------	---------------	------------	--------------

ALOPRIM

<u>AP</u>	+ NABI	<u>EQ 500MG BASE/VIAL</u>	<u>N20298</u>	<u>001</u>	May 17, 1996
-----------	--------	---------------------------	---------------	------------	--------------

ALMOTRIPTAN MALATE

TABLET; ORAL

AXERT

ORTHO MCNEIL PHARM

EQ 6.25MG BASE

N21001 001

May 07, 2001

+

EQ 12.5MG BASE

N21001 002

May 07, 2001

ALOSETRON HYDROCHLORIDE

TABLET; ORAL

LOTRONEX

GLAXOSMITHKLINE

EQ 0.5MG BASE

N21107 002

Dec 23, 2003

+

EQ 1MG BASE

N21107 001

Feb 09, 2000

ALPHA-TOCOPHEROL ACETATE; ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN;
DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN
PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A PALMITATE; VITAMIN K

INJECTABLE; INJECTION

INFUVITE ADULT

+ SABEX 2002

2 IU/ML; 40MG/ML; 12UGM/ML; 40
 IU/ML; 1UGM/ML; 3MG/ML; 120UGM/ML; 8MG/ML; 1
 .2MG/ML; 0.72MG/ML; 1.2MG/ML; 660
 IU/ML; 0.03MG/ML

N21163 001

May 18, 2000

INJECTABLE; IV (INFUSION)

INFUVITE ADULT

+ SABEX 2002

2 IU/ML; 40MG/ML; 12UGM/ML; 40
 IU/ML; 1UGM/ML; 3MG/ML; 120UGM/ML; 8MG/ML; 1
 .2MG/ML; 0.72MG/ML; 1.2MG/ML; 660
 IU/ML; 30UGM/ML

N21559 001

Jun 16, 2003

ALPRAZOLAM

CONCENTRATE; ORAL

ALPRAZOLAM

+ ROXANE

1MG/ML

N74312 001

Oct 31, 1993

TABLET; ORAL

ALPRAZOLAM

<u>AB</u>	ALPHAPHARM	<u>0.25MG</u>	<u>N74046</u>	<u>001</u>	Oct 19, 1993
<u>AB</u>		<u>0.5MG</u>	<u>N74046</u>	<u>002</u>	Oct 19, 1993
<u>AB</u>		<u>1MG</u>	<u>N74046</u>	<u>003</u>	Oct 19, 1993
<u>AB</u>		<u>2MG</u>	<u>N74046</u>	<u>004</u>	May 07, 1997
<u>AB</u>	IVAX PHARMS	<u>0.25MG</u>	<u>N74294</u>	<u>001</u>	Jul 29, 1994
<u>AB</u>		<u>0.5MG</u>	<u>N74294</u>	<u>002</u>	Jul 29, 1994
<u>AB</u>		<u>1MG</u>	<u>N74294</u>	<u>003</u>	Jul 29, 1994
<u>AB</u>		<u>2MG</u>	<u>N74294</u>	<u>004</u>	Jul 29, 1994
<u>AB</u>	MYLAN	<u>0.25MG</u>	<u>N74215</u>	<u>001</u>	Jan 27, 1994
<u>AB</u>		<u>0.5MG</u>	<u>N74215</u>	<u>002</u>	Jan 27, 1994
<u>AB</u>		<u>1MG</u>	<u>N74215</u>	<u>003</u>	Jan 27, 1994
<u>AB</u>		<u>2MG</u>	<u>N74215</u>	<u>004</u>	Jan 27, 1994
<u>AB</u>	PUREPAC PHARM	<u>0.25MG</u>	<u>N74342</u>	<u>001</u>	Oct 31, 1993
<u>AB</u>		<u>0.5MG</u>	<u>N74342</u>	<u>002</u>	Oct 31, 1993
<u>AB</u>		<u>1MG</u>	<u>N74342</u>	<u>003</u>	Oct 31, 1993
<u>AB</u>		<u>2MG</u>	<u>N74342</u>	<u>004</u>	Oct 31, 1993
<u>AB</u>	SANDOZ	<u>0.25MG</u>	<u>N74112</u>	<u>001</u>	Dec 29, 1995
<u>AB</u>		<u>0.5MG</u>	<u>N74112</u>	<u>002</u>	Dec 29, 1995
<u>AB</u>		<u>1MG</u>	<u>N74112</u>	<u>003</u>	Dec 29, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 16 (of 360)

ALPRAZOLAM

TABLET; ORAL

ALPRAZOLAM

<u>AB</u>	SANDOZ	<u>2MG</u>	<u>N74909</u>	<u>001</u>	Mar 25, 1998
<u>AB</u>	TEVA	<u>0.25MG</u>	<u>N74085</u>	<u>001</u>	Feb 16, 1994
<u>AB</u>		<u>0.5MG</u>	<u>N74085</u>	<u>002</u>	Feb 16, 1994
<u>AB</u>		<u>1MG</u>	<u>N74085</u>	<u>003</u>	Feb 16, 1994
<u>AB</u>		<u>2MG</u>	<u>N74085</u>	<u>004</u>	Feb 26, 1996
<u>AB</u>	WATSON LABS	<u>0.25MG</u>	<u>N74456</u>	<u>001</u>	Aug 31, 1995
<u>AB</u>		<u>0.25MG</u>	<u>N74479</u>	<u>001</u>	Jan 21, 1997
<u>AB</u>		<u>0.5MG</u>	<u>N74456</u>	<u>002</u>	Aug 31, 1995
<u>AB</u>		<u>0.5MG</u>	<u>N74479</u>	<u>002</u>	Jan 21, 1997
<u>AB</u>		<u>1MG</u>	<u>N74456</u>	<u>003</u>	Aug 31, 1995
<u>AB</u>		<u>1MG</u>	<u>N74479</u>	<u>003</u>	Jan 21, 1997

XANAX

<u>AB</u>	PHARMACIA AND UPJOHN	<u>0.25MG</u>	<u>N18276</u>	<u>001</u>	
<u>AB</u>		<u>0.5MG</u>	<u>N18276</u>	<u>002</u>	
<u>AB</u>	+	<u>1MG</u>	<u>N18276</u>	<u>003</u>	
<u>AB</u>		<u>2MG</u>	<u>N18276</u>	<u>004</u>	Nov 27, 1985

TABLET, EXTENDED RELEASE; ORAL

XANAX XR

	PHARMACIA AND UPJOHN	0.5MG	N21434	001	Jan 17, 2003
		1MG	N21434	002	Jan 17, 2003
		2MG	N21434	003	Jan 17, 2003
+		3MG	N21434	004	Jan 17, 2003

TABLET, ORALLY DISINTEGRATING; ORAL

NIRAVAM

	SCHWARZ PHARMA	0.25MG	N21726	001	Jan 19, 2005
		0.5MG	N21726	002	Jan 19, 2005
		1MG	N21726	003	Jan 19, 2005
+		2MG	N21726	004	Jan 19, 2005

ALPROSTADIL

INJECTABLE; INJECTION

ALPROSTADIL

<u>AP</u>	BEDFORD	<u>0.5MG/ML</u>	<u>N74815</u>	<u>001</u>	Jan 20, 1998
<u>AP</u>	SICOR PHARMS	<u>0.5MG/ML</u>	<u>N75196</u>	<u>001</u>	Apr 30, 1999

CAVERJECT

<u>AP</u>	PHARMACIA AND UPJOHN	<u>0.01MG/VIAL</u>	<u>N20379</u>	<u>001</u>	Jul 06, 1995
<u>AP</u>	+	<u>0.02MG/VIAL</u>	<u>N20379</u>	<u>002</u>	Jul 06, 1995
<u>AP</u>	+	<u>0.04MG/VIAL</u>	<u>N20379</u>	<u>004</u>	May 19, 1997
		0.005MG/VIAL	N20379	003	Jun 27, 1996
		0.01MG/VIAL	N21212	001	Jun 11, 2002
		0.02MG/VIAL	N21212	002	Jun 11, 2002

EDEX

<u>AP</u>	SCHWARZ PHARMA	<u>0.01MG/VIAL</u>	<u>N20649</u>	<u>002</u>	Jun 12, 1997
<u>AP</u>		<u>0.02MG/VIAL</u>	<u>N20649</u>	<u>003</u>	Jun 12, 1997
<u>AP</u>	+	<u>0.04MG/VIAL</u>	<u>N20649</u>	<u>004</u>	Jun 12, 1997
+		0.01MG/VIAL	N20649	005	Jul 30, 1998
+		0.02MG/VIAL	N20649	006	Jul 30, 1998
+		0.04MG/VIAL	N20649	007	Jul 30, 1998

PROSTIN VR PEDIATRIC

<u>AP</u>	+	PHARMACIA AND UPJOHN	<u>0.5MG/ML</u>	<u>N18484</u>	<u>001</u>
-----------	---	----------------------	-----------------	---------------	------------

SUPPOSITORY; URETHRAL

MUSE

	VIVUS	0.125MG	N20700	001	Nov 19, 1996
		0.25MG	N20700	002	Nov 19, 1996
		0.5MG	N20700	003	Nov 19, 1996
+		1MG	N20700	004	Nov 19, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 17 (of 360)

ALTRETAMINE

CAPSULE; ORAL

HEXALEN

+ MGI PHARMA INC	50MG	N19926	001	Dec 26, 1990
------------------	------	--------	-----	--------------

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u> SANDOZ	100MG	<u>N71293</u>	<u>001</u>	Feb 18, 1987
------------------	-------	---------------	------------	--------------

<u>AB</u> + USL PHARMA	100MG	<u>N70589</u>	<u>001</u>	Aug 05, 1986
------------------------	-------	---------------	------------	--------------

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

<u>AA</u> + ALPHARMA US PHARMS	50MG/5ML	<u>N72655</u>	<u>001</u>	Oct 30, 1990
--------------------------------	----------	---------------	------------	--------------

<u>AA</u> + CAROLINA MEDCL	50MG/5ML	<u>N75819</u>	<u>001</u>	Sep 11, 2002
----------------------------	----------	---------------	------------	--------------

<u>AA</u> + HI TECH PHARMA	50MG/5ML	<u>N74170</u>	<u>001</u>	Oct 28, 1994
----------------------------	----------	---------------	------------	--------------

<u>AA</u> + MIKART	50MG/5ML	<u>N74028</u>	<u>001</u>	Jun 28, 1993
--------------------	----------	---------------	------------	--------------

<u>AA</u> + MORTON GROVE	50MG/5ML	<u>N75060</u>	<u>001</u>	Dec 24, 1998
--------------------------	----------	---------------	------------	--------------

<u>AA</u> + PHARM ASSOC	50MG/5ML	<u>N74509</u>	<u>001</u>	Jul 17, 1995
-------------------------	----------	---------------	------------	--------------

<u>AA</u> + SILARX	50MG/5ML	<u>N76352</u>	<u>001</u>	Sep 10, 2004
--------------------	----------	---------------	------------	--------------

<u>AA</u> + TEVA PHARMS	50MG/5ML	<u>N73115</u>	<u>001</u>	Aug 23, 1991
-------------------------	----------	---------------	------------	--------------

TABLET; ORAL

AMANTADINE HYDROCHLORIDE

<u>AB</u> USL PHARMA	100MG	<u>N76186</u>	<u>001</u>	Dec 16, 2002
----------------------	-------	---------------	------------	--------------

SYMMETREL

<u>AB</u> + ENDO PHARMS	100MG	<u>N18101</u>	<u>001</u>	
-------------------------	-------	---------------	------------	--

AMBENONIUM CHLORIDE

TABLET; ORAL

MYTELASE

+ SANOFI SYNTHELABO	10MG	N10155	002	
---------------------	------	--------	-----	--

AMCINONIDE

CREAM; TOPICAL

AMCINONIDE

<u>AB</u> ALTANA	0.1%	<u>N76065</u>	<u>001</u>	May 15, 2003
------------------	------	---------------	------------	--------------

<u>AB</u> TARO PHARM INDS	0.1%	<u>N76229</u>	<u>001</u>	May 31, 2002
---------------------------	------	---------------	------------	--------------

CYCLOCORT

<u>AB</u> + ASTELLAS	0.1%	<u>N18116</u>	<u>002</u>	
----------------------	------	---------------	------------	--

LOTION; TOPICAL

AMCINONIDE

<u>AB</u> ALTANA	0.1%	<u>N76329</u>	<u>001</u>	Nov 06, 2002
------------------	------	---------------	------------	--------------

CYCLOCORT

<u>AB</u> + ASTELLAS	0.1%	<u>N19729</u>	<u>001</u>	Jun 13, 1988
----------------------	------	---------------	------------	--------------

OINTMENT; TOPICAL

AMCINONIDE

<u>AB</u> ALTANA	0.1%	<u>N76096</u>	<u>001</u>	Nov 19, 2002
------------------	------	---------------	------------	--------------

<u>AB</u> TARO PHARM INDS	0.1%	<u>N76367</u>	<u>001</u>	Mar 19, 2003
---------------------------	------	---------------	------------	--------------

CYCLOCORT

<u>AB</u> + ASTELLAS	0.1%	<u>N18498</u>	<u>001</u>	
----------------------	------	---------------	------------	--

AMIFOSTINE

INJECTABLE; INJECTION

ETHYOL

+ MEDIMMUNE ONCOLOGY	500MG/VIAL	N20221	001	Dec 08, 1995
----------------------	------------	--------	-----	--------------

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

<u>AP</u> BEDFORD	EQ 50MG BASE/ML	<u>N63313</u>	<u>001</u>	Apr 11, 1994
-------------------	-----------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 18 (of 360)

AMIKACIN SULFATE

INJECTABLE; INJECTION

AMIKACIN SULFATE

<u>AP</u>	BEDFORD	<u>EQ 250MG BASE/ML</u>	<u>N63315</u>	<u>001</u>	Apr 11, 1994
<u>AP</u>	HOSPIRA	<u>EQ 50MG BASE/ML</u>	<u>N63263</u>	<u>001</u>	Nov 30, 1994
<u>AP</u>		<u>EQ 250MG BASE/ML</u>	<u>N63264</u>	<u>001</u>	Nov 30, 1994
<u>AP</u>		<u>EQ 250MG BASE/ML</u>	<u>N64098</u>	<u>001</u>	Jun 26, 1995
		EQ 62.5MG BASE/ML	N63283	001	Oct 31, 1994
<u>AP</u>	SICOR PHARMS	<u>EQ 50MG BASE/ML</u>	<u>N64045</u>	<u>001</u>	Sep 28, 1993
<u>AP</u>		<u>EQ 250MG BASE/ML</u>	<u>N64045</u>	<u>002</u>	Sep 28, 1993
	AMIKACIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
	HOSPIRA	EQ 500MG BASE/100ML	N64146	001	Apr 02, 1997
	<u>AMIKIN</u>				
<u>AP</u>	+ APOTHECON	<u>EQ 50MG BASE/ML</u>	<u>N62311</u>	<u>001</u>	
<u>AP</u>	+	<u>EQ 250MG BASE/ML</u>	<u>N62311</u>	<u>002</u>	

AMILORIDE HYDROCHLORIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE

<u>AB</u>	PAR PHARM	<u>5MG</u>	<u>N70346</u>	<u>001</u>	Jan 22, 1986
	<u>MIDAMOR</u>				
<u>AB</u>	+ MERCK	<u>5MG</u>	<u>N18200</u>	<u>001</u>	

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	BARR	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>N71111</u>	<u>001</u>	May 10, 1988
<u>AB</u>	MYLAN	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>N73209</u>	<u>001</u>	Oct 31, 1991
<u>AB</u>	SANDOZ	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>N73357</u>	<u>001</u>	Nov 27, 1991
<u>AB</u>	TEVA	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>N70795</u>	<u>001</u>	Apr 17, 1988
<u>AB</u>	WATSON LABS	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>N73334</u>	<u>001</u>	Jul 19, 1991
	<u>HYDRO-RIDE</u>				
<u>AB</u>	PAR PHARM	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>N70347</u>	<u>001</u>	Dec 25, 1990
	<u>MODURETIC 5-50</u>				
<u>AB</u>	+ MERCK	<u>EQ 5MG ANHYDROUS;50MG</u>	<u>N18201</u>	<u>001</u>	

AMINO ACIDS

INJECTABLE; INJECTION

AMINOSYN 10%

HOSPIRA

10% (10GM/100ML)

N17673 003

AMINOSYN 10% (PH6)

HOSPIRA

10% (10GM/100ML)

N17673 008

Nov 18, 1985

AMINOSYN 3.5%

HOSPIRA

3.5% (3.5GM/100ML)

N17789 004

AMINOSYN 5%

HOSPIRA

5% (5GM/100ML)

N17673 001

AMINOSYN 7%

HOSPIRA

7% (7GM/100ML)

N17673 002

AMINOSYN 7% (PH6)

HOSPIRA

7% (7GM/100ML)

N17673 006

Nov 18, 1985

AMINOSYN 8.5%

HOSPIRA

8.5% (8.5GM/100ML)

N17673 004

AMINOSYN 8.5% (PH6)

HOSPIRA

8.5% (8.5GM/100ML)

N17673 007

Nov 18, 1985

AMINOSYN II 10%

HOSPIRA

10% (10GM/100ML)

N19438 005

Apr 03, 1986

AMINOSYN II 10% IN PLASTIC CONTAINER

HOSPIRA

10% (10GM/100ML)

N20015 001

Dec 19, 1991

AMINOSYN II 15% IN PLASTIC CONTAINER

HOSPIRA

15% (15GM/100ML)

N20041 001

Dec 19, 1991

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 19 (of 360)

AMINO ACIDS

INJECTABLE; INJECTION

AMINOSYN II 7%				
HOSPIRA	7% (7GM/100ML)	N19438	003	Apr 03, 1986
AMINOSYN II 8.5%				
HOSPIRA	8.5% (8.5GM/100ML)	N19438	004	Apr 03, 1986
AMINOSYN-HBC 7%				
HOSPIRA	7% (7GM/100ML)	N19374	001	Jul 12, 1985
AMINOSYN-HF 8%				
HOSPIRA	8% (8GM/100ML)	N20345	001	Apr 04, 1996
AMINOSYN-PF 10%				
HOSPIRA	10% (10GM/100ML)	N19492	002	Oct 17, 1986
AMINOSYN-PF 7%				
HOSPIRA	7% (7GM/100ML)	N19398	001	Sep 06, 1985
AMINOSYN-RF 5.2%				
HOSPIRA	5.2% (5.2GM/100ML)	N18429	001	
BRANCHAMIN 4% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	4% (4GM/100ML)	N18684	001	Sep 28, 1984
CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER				
CLINTEC NUTR	15% (15GM/100ML)	N20512	001	Aug 30, 1996
FREAMINE HBC 6.9%				
B BRAUN	6.9% (6.9GM/100ML)	N16822	006	May 17, 1983
FREAMINE III 10%				
B BRAUN	10% (10GM/100ML)	N16822	005	
FREAMINE III 8.5%				
B BRAUN	8.5% (8.5GM/100ML)	N16822	004	
HEPATAMINE 8%				
B BRAUN	8% (8GM/100ML)	N18676	001	Aug 03, 1982
HEPATASOL 8%				
BAXTER HLTHCARE	8% (8GM/100ML)	N20360	001	Apr 04, 1996
NEPHRAMINE 5.4%				
B BRAUN	5.4% (5.4GM/100ML)	N17766	001	
NOVAMINE 11.4%				
HOSPIRA	11.4% (11.4GM/100ML)	N17957	003	Aug 09, 1982
NOVAMINE 15%				
HOSPIRA	15% (15GM/100ML)	N17957	004	Nov 28, 1986
PREMASOL 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	10% (10GM/100ML)	N75880	002	Jun 19, 2003
PREMASOL 6% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	6% (6GM/100ML)	N75880	001	Jun 19, 2003
PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER				
+ BAXTER HLTHCARE	20% (20GM/100ML)	N20849	001	Aug 26, 1998
RENAMIN W/O ELECTROLYTES				
BAXTER HLTHCARE	6.5% (6.5GM/100ML)	N17493	007	Oct 15, 1982
TRAVASOL 10% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	10% (10MG/100ML)	N18931	003	Aug 23, 1984
TRAVASOL 10% W/O ELECTROLYTES				
BAXTER HLTHCARE	10% (10GM/100ML)	N17493	006	
TRAVASOL 5.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	5.5% (5.5GM/100ML)	N18931	001	Aug 23, 1984
TRAVASOL 5.5% W/O ELECTROLYTES				
BAXTER HLTHCARE	5.5% (5.5GM/100ML)	N17493	004	
TRAVASOL 8.5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	8.5% (8.5GM/100ML)	N18931	002	Aug 23, 1984
TRAVASOL 8.5% W/O ELECTROLYTES				
BAXTER HLTHCARE	8.5% (8.5GM/100ML)	N17493	005	
TROPHAMINE				
+ B BRAUN	6% (6GM/100ML)	N19018	001	Jul 20, 1984
TROPHAMINE 10%				
+ B BRAUN	10% (10GM/100ML)	N19018	003	Sep 07, 1988

PRESCRIPTION DRUG PRODUCT LIST

3 - 20 (of 360)

AMINO ACIDS; CALCIUM ACETATE; GLYCERIN; MAGNESIUM ACETATE; PHOSPHORIC ACID;
POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PROCALAMINE

B BRAUN

3%;26MG/100ML;3GM/100ML;54MG/100ML;41MG N18582 001 May 08, 1982
/100ML;150MG/100ML;200MG/100ML;120MG/10
0ML

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE;
POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

HOSPIRA 3.5%;36.8MG/100ML;25GM/100ML;51MG/100ML N19683 001 Nov 07, 1988
;22.4MG/100ML;261MG/100ML;205MG/100ML

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER

HOSPIRA 4.25%;36.8MG/100ML;20GM/100ML;51MG/100M N19683 002 Nov 07, 1988
L;22.4MG/100ML;261MG/100ML;205MG/100ML

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

HOSPIRA 4.25%;36.8MG/100ML;25GM/100ML;51MG/100M N19683 003 Nov 07, 1988
L;22.4MG/100ML;261MG/100ML;205MG/100ML

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE,
DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

CLINIMIX E 2.75/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 2.75%;33MG/100ML;10GM/100ML;51MG/100ML; N20678 002 Mar 26, 1997
261MG/100ML;217MG/100ML;112MG/100ML

CLINIMIX E 2.75/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 2.75%;33MG/100ML;25GM/100ML;51MG/100ML; N20678 005 Mar 26, 1997
261MG/100ML;217MG/100ML;112MG/100ML

CLINIMIX E 2.75/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 2.75%;33MG/100ML;5GM/100ML;51MG/100ML;2 N20678 001 Mar 26, 1997
61MG/100ML;217MG/100ML;112MG/100ML

CLINIMIX E 4.25/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 4.25%;33MG/100ML;10GM/100ML;51MG/100ML; N20678 009 Mar 26, 1997
261MG/100ML;297MG/100ML;77MG/100ML

CLINIMIX E 4.25/20 SULFITE-FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 4.25%;33MG/100ML;20GM/100ML;51MG/100ML; N20678 011 Mar 26, 1997
261MG/100ML;297MG/100ML;77MG/100ML

CLINIMIX E 4.25/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 4.25%;33MG/100ML;25GM/100ML;51MG/100ML; N20678 012 Mar 26, 1997
261MG/100ML;297MG/100ML;77MG/100ML

CLINIMIX E 4.25/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 4.25%;33MG/100ML;5GM/100ML;51MG/100ML;2 N20678 008 Mar 26, 1997
61MG/100ML;297MG/100ML;77MG/100ML

CLINIMIX E 5/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5%;33MG/100ML;10GM/100ML;51MG/100ML;261 N20678 016 Mar 26, 1997
MG/100ML;340MG/100ML;59MG/100ML

CLINIMIX E 5/15 SULFITE-FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5%;33MG/100ML;15GM/100ML;51MG/100ML;261 N20678 017 Mar 26, 1997
MG/100ML;340MG/100ML;59MG/100ML

CLINIMIX E 5/20 SULFITE-FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5%;33MG/100ML;20GM/100ML;51MG/100ML;261 N20678 018 Mar 26, 1997
MG/100ML;340MG/100ML;59MG/100ML

CLINIMIX E 5/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5%;33MG/100ML;25GM/100ML;51MG/100ML;261 N20678 019 Mar 26, 1997
MG/100ML;340MG/100ML;59MG/100ML

CLINIMIX E 5/35 SULFITE-FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER

+ BAXTER HLTHCARE 5%;33MG/100ML;35GM/100ML;51MG/100ML;261 N20678 021 Mar 26, 1997
MG/100ML;340MG/100ML;59MG/100ML

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 21 (of 360)

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER			
HOSPIRA 3.5%;25GM/100ML	N19681 001	Nov 01, 1988	
AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER			
HOSPIRA 3.5%;5GM/100ML	N19681 002	Nov 01, 1988	
AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER			
HOSPIRA 4.25%;10GM/100ML	N19681 004	Nov 01, 1988	
AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER			
HOSPIRA 4.25%;20GM/100ML	N19681 005	Nov 01, 1988	
AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER			
HOSPIRA 4.25%;25GM/100ML	N19681 003	Nov 01, 1988	
AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER			
HOSPIRA 5%;25GM/100ML	N19681 006	Nov 01, 1988	
CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 2.75%;10GM/100ML	N20734 002	Sep 29, 1997	
CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 2.75%;25GM/100ML	N20734 005	Sep 29, 1997	
CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 2.75%;5GM/100ML	N20734 001	Sep 29, 1997	
CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 4.25%;10GM/100ML	N20734 008	Sep 29, 1997	
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 4.25%;20GM/100ML	N20734 010	Sep 29, 1997	
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 4.25%;25GM/100ML	N20734 011	Sep 29, 1997	
CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 4.25%;5GM/100ML	N20734 007	Sep 29, 1997	
CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 5%;10GM/100ML	N20734 014	Sep 29, 1997	
CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 5%;15GM/100ML	N20734 015	Sep 29, 1997	
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 5%;20GM/100ML	N20734 016	Sep 29, 1997	
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 5%;25GM/100ML	N20734 017	Sep 29, 1997	
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER			
BAXTER HLTHCARE 5%;35GM/100ML	N20734 018	Sep 29, 1997	

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN DEXTROSE 5% IN PLASTIC CONTAINER			
HOSPIRA 3.5%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML	N19682 001	Nov 01, 1988	
AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER			
HOSPIRA 4.25%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML	N19682 002	Nov 01, 1988	

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

FREAMINE III 8.5% W/ ELECTROLYTES			
B BRAUN 8.5%;110MG/100ML;230MG/100ML;10MG/100ML;440MG/100ML;690MG/100ML	N16822 007	Jul 01, 1988	

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M			
HOSPIRA 3.5%;21MG/100ML;40MG/100ML;128MG/100ML;234MG/100ML	N17789 003		

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 22 (of 360)

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

FREAMINE III 3% W/ ELECTROLYTES

B BRAUN	3%;54MG/100ML;40MG/100ML;150MG/100ML;20 0MG/100ML;120MG/100ML	N16822	003	
---------	--	--------	-----	--

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 10% W/ ELECTROLYTES

HOSPIRA	10%;102MG/100ML;45MG/100ML;522MG/100ML; 410MG/100ML	N19437	004	Apr 03, 1986
---------	--	--------	-----	--------------

AMINOSYN II 8.5% W/ ELECTROLYTES

HOSPIRA	8.5%;102MG/100ML;45MG/100ML;522MG/100ML ;410MG/100ML	N19437	005	Apr 03, 1986
---------	---	--------	-----	--------------

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

BAXTER HLTHCARE	3.5%;51MG/100ML;131MG/100ML;218MG/100ML ;35MG/100ML	N20177	001	Oct 23, 1995
-----------------	--	--------	-----	--------------

TRAVASOL 3.5% W/ ELECTROLYTES

BAXTER HLTHCARE	3.5%;51MG/100ML;131MG/100ML;218MG/100ML ;35MG/100ML	N17493	003	
-----------------	--	--------	-----	--

TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

BAXTER HLTHCARE	5.5%;102MG/100ML;522MG/100ML;431MG/100M L;224MG/100ML	N20173	001	Oct 27, 1995
-----------------	--	--------	-----	--------------

TRAVASOL 5.5% W/ ELECTROLYTES

BAXTER HLTHCARE	5.5%;102MG/100ML;522MG/100ML;431MG/100M L;224MG/100ML	N17493	001	
-----------------	--	--------	-----	--

TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER

BAXTER HLTHCARE	8.5%;102MG/100ML;522MG/100ML;594MG/100M L;154MG/100ML	N20173	002	Oct 27, 1995
-----------------	--	--------	-----	--------------

TRAVASOL 8.5% W/ ELECTROLYTES

BAXTER HLTHCARE	8.5%;102MG/100ML;522MG/100ML;594MG/100M L;154MG/100ML	N17493	002	
-----------------	--	--------	-----	--

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 7% W/ ELECTROLYTES

HOSPIRA	7%;102MG/100ML;522MG/100ML;410MG/100ML	N17789	002	
---------	--	--------	-----	--

AMINOSYN 8.5% W/ ELECTROLYTES

HOSPIRA	8.5%;102MG/100ML;522MG/100ML;410MG/100M L	N17673	005	
---------	--	--------	-----	--

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMICAR

<u>AP</u>	+ XANODYNE PHARM	<u>250MG/ML</u>	<u>N15229</u>	<u>002</u>	
-----------	------------------	-----------------	---------------	------------	--

AMINOCAPROIC ACID

<u>AP</u>	HOSPIRA	<u>250MG/ML</u>	<u>N70888</u>	<u>001</u>	Jun 16, 1988
-----------	---------	-----------------	---------------	------------	--------------

<u>AP</u>	LUITPOLD	<u>250MG/ML</u>	<u>N71192</u>	<u>001</u>	Dec 01, 1987
-----------	----------	-----------------	---------------	------------	--------------

AMINOCAPROIC ACID IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>250MG/ML</u>	<u>N70010</u>	<u>001</u>	Mar 09, 1987
-----------	---------	-----------------	---------------	------------	--------------

SYRUP; ORAL

AMICAR

<u>AA</u>	+ XANODYNE PHARM	<u>1.25GM/5ML</u>	<u>N15230</u>	<u>002</u>	
-----------	------------------	-------------------	---------------	------------	--

AMINOCAPROIC ACID

<u>AA</u>	MIKART	<u>1.25GM/5ML</u>	<u>N74759</u>	<u>001</u>	Sep 02, 1998
-----------	--------	-------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 23 (of 360)

AMINOCAPROIC ACID

TABLET; ORAL

AMICAR

<u>AB</u>	+	XANODYNE PHARM	<u>500MG</u>	<u>N15197</u>	<u>001</u>	
-----------	---	----------------	--------------	---------------	------------	--

AMINOCAPROIC

<u>AB</u>		MIKART	<u>500MG</u>	<u>N75602</u>	<u>001</u>	May 24, 2001
-----------	--	--------	--------------	---------------	------------	--------------

AMINOGLUTETHIMIDE

TABLET; ORAL

CYTADREN

+	NOVARTIS	250MG	N18202	001	
---	----------	-------	--------	-----	--

AMINOHIPPURATE SODIUM

INJECTABLE; INJECTION

AMINOHIPPURATE SODIUM

+	MERCK	20%	N05619	001	
---	-------	-----	--------	-----	--

AMINOLEVULINIC ACID HYDROCHLORIDE

SOLUTION; TOPICAL

LEVULAN

+	DUSA	20%	N20965	001	Dec 03, 1999
---	------	-----	--------	-----	--------------

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

<u>AP</u>		HOSPIRA	<u>25MG/ML</u>	<u>N87242</u>	<u>001</u>	Oct 26, 1983
-----------	--	---------	----------------	---------------	------------	--------------

<u>AP</u>	+		<u>25MG/ML</u>	<u>N87601</u>	<u>001</u>	Jul 23, 1982
-----------	---	--	----------------	---------------	------------	--------------

<u>AP</u>		INTL MEDICATION	<u>25MG/ML</u>	<u>N87209</u>	<u>001</u>	Feb 01, 1982
-----------	--	-----------------	----------------	---------------	------------	--------------

<u>AP</u>		LUITPOLD	<u>25MG/ML</u>	<u>N87240</u>	<u>001</u>	
-----------	--	----------	----------------	---------------	------------	--

<u>AP</u>			<u>25MG/ML</u>	<u>N87600</u>	<u>001</u>	
-----------	--	--	----------------	---------------	------------	--

<u>AP</u>		PHARMA SERVE NY	<u>25MG/ML</u>	<u>N87392</u>	<u>001</u>	Dec 15, 1983
-----------	--	-----------------	----------------	---------------	------------	--------------

<u>AP</u>		SICOR PHARMS	<u>25MG/ML</u>	<u>N81142</u>	<u>001</u>	Sep 25, 1991
-----------	--	--------------	----------------	---------------	------------	--------------

AMINOPHYLLINE IN SODIUM CHLORIDE 0.45%

+	HOSPIRA	100MG/100ML	N88147	002	May 03, 1983
---	---------	-------------	--------	-----	--------------

+		200MG/100ML	N88147	003	May 03, 1983
---	--	-------------	--------	-----	--------------

SOLUTION; ORAL

AMINOPHYLLINE

<u>AA</u>	+	ROXANE	<u>105MG/5ML</u>	<u>N88126</u>	<u>001</u>	Aug 19, 1983
-----------	---	--------	------------------	---------------	------------	--------------

AMINOPHYLLINE DYE FREE

<u>AA</u>	+	ALPHARMA US PHARMS	<u>105MG/5ML</u>	<u>N87727</u>	<u>001</u>	Apr 16, 1982
-----------	---	--------------------	------------------	---------------	------------	--------------

SUPPOSITORY; RECTAL

TRUPHYLLINE

	G AND W LABS	250MG	N85498	001	Mar 23, 1983
--	--------------	-------	--------	-----	--------------

+		500MG	N85498	002	Jan 03, 1983
---	--	-------	--------	-----	--------------

TABLET; ORAL

AMINOPHYLLINE

+	WEST WARD	100MG	N84540	001	
---	-----------	-------	--------	-----	--

+		200MG	N85003	001	
---	--	-------	--------	-----	--

AMINOSALICYLATE SODIUM

TABLET; ORAL

SODIUM P.A.S.

+	LANNETT	500MG	N80138	002	
---	---------	-------	--------	-----	--

AMINOSALICYLIC ACID

GRANULE, DELAYED RELEASE; ORAL

PASER

+	JACOBUS	4GM/PACKET	N74346	001	Jun 30, 1994
---	---------	------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 24 (of 360)

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

AMIODARONE HYDROCHLORIDE

<u>AP</u>	+	AM PHARM PARTNERS	<u>50MG/ML</u>	<u>N75761</u>	<u>001</u>	Oct 15, 2002
<u>AP</u>	+	APOTEX	<u>50MG/ML</u>	<u>N76394</u>	<u>001</u>	Apr 25, 2003
<u>AP</u>	+		<u>50MG/ML</u>	<u>N77161</u>	<u>001</u>	Apr 20, 2005
<u>AP</u>	+	BEDFORD	<u>50MG/ML</u>	<u>N76018</u>	<u>001</u>	Oct 15, 2002
<u>AP</u>	+	BEDFORD LABS	<u>50MG/ML</u>	<u>N76299</u>	<u>001</u>	Oct 24, 2002
<u>AP</u>	+	BEN VENUE	<u>50MG/ML</u>	<u>N76088</u>	<u>001</u>	Oct 15, 2002
<u>AP</u>	+	BIONICHE (CANADA)	<u>50MG/ML</u>	<u>N76217</u>	<u>001</u>	Oct 15, 2002
<u>AP</u>	+	HOSPIRA	<u>50MG/ML</u>	<u>N75955</u>	<u>001</u>	Oct 18, 2002
<u>AP</u>	+	INTL MEDICATION SYS	<u>50MG/ML</u>	<u>N21594</u>	<u>001</u>	Feb 04, 2004
<u>AP</u>	+	MAYNE PHARMA USA	<u>50MG/ML</u>	<u>N76108</u>	<u>001</u>	Oct 15, 2002
<u>AP</u>	+	SICOR PHARMS	<u>50MG/ML</u>	<u>N76163</u>	<u>001</u>	Sep 05, 2003
<u>CORDARONE</u>						
<u>AP</u>	+	WYETH PHARMS INC	<u>50MG/ML</u>	<u>N20377</u>	<u>001</u>	Aug 03, 1995

TABLET; ORAL

AMIODARONE HYDROCHLORIDE

<u>AB</u>		ALPHAPHARM	<u>200MG</u>	<u>N75188</u>	<u>001</u>	Feb 24, 1999
<u>AB</u>		AUROSAL PHARMS	<u>200MG</u>	<u>N77069</u>	<u>001</u>	Apr 08, 2005
<u>AB</u>			<u>400MG</u>	<u>N77069</u>	<u>002</u>	Apr 08, 2005
<u>AB</u>		BARR	<u>200MG</u>	<u>N75389</u>	<u>001</u>	Jan 25, 2001
<u>AB</u>		SANDOZ	<u>200MG</u>	<u>N75315</u>	<u>001</u>	Dec 23, 1998
<u>AB</u>			<u>400MG</u>	<u>N75315</u>	<u>002</u>	Jun 30, 2000
<u>AB</u>		TARO	<u>100MG</u>	<u>N75424</u>	<u>002</u>	Dec 18, 2002
<u>AB</u>			<u>200MG</u>	<u>N75424</u>	<u>001</u>	Mar 30, 2001
<u>AB</u>			<u>400MG</u>	<u>N76362</u>	<u>001</u>	Nov 29, 2002
			300MG	N76362	002	Dec 02, 2003
<u>AB</u>		TEVA	<u>200MG</u>	<u>N74895</u>	<u>001</u>	Apr 16, 1999
<u>AB</u>		TEVA PHARMS	<u>200MG</u>	<u>N74739</u>	<u>001</u>	Nov 30, 1998
<u>CORDARONE</u>						
<u>AB</u>	+	WYETH PHARMS INC	<u>200MG</u>	<u>N18972</u>	<u>001</u>	Dec 24, 1985
<u>PACERONE</u>						
<u>AB</u>		UPSHER SMITH	<u>100MG</u>	<u>N75135</u>	<u>002</u>	Apr 12, 2005
<u>AB</u>			<u>200MG</u>	<u>N75135</u>	<u>001</u>	Apr 30, 1998

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>		MUTUAL PHARM	<u>10MG</u>	<u>N89398</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>			<u>25MG</u>	<u>N89399</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>			<u>50MG</u>	<u>N89400</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>			<u>75MG</u>	<u>N89401</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>			<u>100MG</u>	<u>N89402</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>			<u>150MG</u>	<u>N89403</u>	<u>001</u>	Jul 14, 1987
<u>AB</u>		MYLAN	<u>10MG</u>	<u>N86157</u>	<u>001</u>	
<u>AB</u>			<u>25MG</u>	<u>N86010</u>	<u>001</u>	
<u>AB</u>			<u>50MG</u>	<u>N86009</u>	<u>001</u>	
<u>AB</u>			<u>75MG</u>	<u>N86011</u>	<u>001</u>	
<u>AB</u>			<u>100MG</u>	<u>N86158</u>	<u>001</u>	
<u>AB</u>			<u>150MG</u>	<u>N86153</u>	<u>001</u>	
<u>AB</u>		PLIVA	<u>10MG</u>	<u>N88883</u>	<u>001</u>	Sep 26, 1984
<u>AB</u>			<u>25MG</u>	<u>N88884</u>	<u>001</u>	Sep 26, 1984
<u>AB</u>			<u>50MG</u>	<u>N88885</u>	<u>001</u>	Sep 26, 1984
<u>AB</u>			<u>75MG</u>	<u>N88886</u>	<u>001</u>	Sep 26, 1984
<u>AB</u>			<u>100MG</u>	<u>N88887</u>	<u>001</u>	Sep 26, 1984
<u>AB</u>			<u>150MG</u>	<u>N88888</u>	<u>001</u>	Sep 26, 1984
<u>AB</u>		SANDOZ	<u>10MG</u>	<u>N85969</u>	<u>001</u>	
<u>AB</u>	+		<u>25MG</u>	<u>N85966</u>	<u>001</u>	
<u>AB</u>			<u>50MG</u>	<u>N85968</u>	<u>001</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 25 (of 360)

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>75MG</u>	<u>N85971</u>	<u>001</u>	
<u>AB</u>		<u>100MG</u>	<u>N85967</u>	<u>001</u>	
<u>AB</u>		<u>150MG</u>	<u>N85970</u>	<u>001</u>	
<u>AB</u>	TEVA	<u>10MG</u>	<u>N84910</u>	<u>003</u>	
<u>AB</u>		<u>25MG</u>	<u>N85031</u>	<u>001</u>	
<u>AB</u>		<u>100MG</u>	<u>N85836</u>	<u>001</u>	
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>N40218</u>	<u>001</u>	Sep 11, 1997
<u>AB</u>		<u>25MG</u>	<u>N40218</u>	<u>002</u>	Sep 11, 1997
<u>AB</u>		<u>50MG</u>	<u>N40218</u>	<u>003</u>	Sep 11, 1997
<u>AB</u>		<u>75MG</u>	<u>N40218</u>	<u>004</u>	Sep 11, 1997
<u>AB</u>		<u>100MG</u>	<u>N40218</u>	<u>005</u>	Sep 11, 1997
<u>AB</u>		<u>150MG</u>	<u>N40218</u>	<u>006</u>	Sep 11, 1997

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET; ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>EQ 12.5MG BASE;5MG</u>	<u>N71296</u>	<u>001</u>	Dec 10, 1986
<u>AB</u>		<u>EQ 25MG BASE;10MG</u>	<u>N71297</u>	<u>001</u>	Dec 10, 1986
<u>AB</u>	PAR PHARM	<u>EQ 12.5MG BASE;5MG</u>	<u>N72277</u>	<u>001</u>	May 09, 1988
<u>AB</u>		<u>EQ 25MG BASE;10MG</u>	<u>N72278</u>	<u>001</u>	May 09, 1988
<u>AB</u>	WATSON LABS	<u>EQ 12.5MG BASE;5MG</u>	<u>N72052</u>	<u>001</u>	Dec 16, 1988
<u>AB</u>		<u>EQ 25MG BASE;10MG</u>	<u>N72053</u>	<u>001</u>	Dec 16, 1988
<u>LIMBITROL</u>					
<u>AB</u>	VALEANT PHARM INTL	<u>EQ 12.5MG BASE;5MG</u>	<u>N16949</u>	<u>001</u>	
<u>LIMBITROL DS</u>					
<u>AB</u>	+ VALEANT PHARM INTL	<u>EQ 25MG BASE;10MG</u>	<u>N16949</u>	<u>002</u>	

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>10MG;2MG</u>	<u>N70336</u>	<u>001</u>	Nov 10, 1988
<u>AB</u>		<u>10MG;4MG</u>	<u>N71442</u>	<u>001</u>	Nov 10, 1988
<u>AB</u>		<u>25MG;2MG</u>	<u>N70337</u>	<u>001</u>	Nov 10, 1988
<u>AB</u>	+	<u>25MG;4MG</u>	<u>N70338</u>	<u>001</u>	Nov 10, 1988
<u>AB</u>	+	<u>50MG;4MG</u>	<u>N71443</u>	<u>001</u>	Nov 10, 1988
<u>AB</u>	SANDOZ	<u>10MG;2MG</u>	<u>N71062</u>	<u>001</u>	Nov 27, 1987
<u>AB</u>		<u>10MG;4MG</u>	<u>N71862</u>	<u>001</u>	Dec 21, 1987
<u>AB</u>		<u>25MG;2MG</u>	<u>N71063</u>	<u>001</u>	Nov 27, 1987
<u>AB</u>		<u>25MG;4MG</u>	<u>N71064</u>	<u>001</u>	Nov 27, 1987
<u>AB</u>		<u>50MG;4MG</u>	<u>N71863</u>	<u>001</u>	Dec 21, 1987
<u>AB</u>	WATSON LABS	<u>10MG;2MG</u>	<u>N73007</u>	<u>001</u>	Oct 17, 1991
<u>AB</u>		<u>10MG;4MG</u>	<u>N73009</u>	<u>001</u>	Oct 17, 1991
<u>AB</u>		<u>25MG;2MG</u>	<u>N73008</u>	<u>001</u>	Oct 17, 1991
<u>AB</u>		<u>25MG;4MG</u>	<u>N73010</u>	<u>001</u>	Oct 17, 1991

AMLEXANOX

PASTE; DENTAL

APHTHASOL

+	ULURU	5%	N20511	001	Dec 17, 1996
---	-------	----	--------	-----	--------------

PATCH; TOPICAL

AMLEXANOX

+	ULURU	2MG	N21727	001	Sep 29, 2004
---	-------	-----	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 26 (of 360)

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

<u>AB</u>	MYLAN	<u>EQ 2.5MG BASE</u>	<u>N76418</u>	<u>001</u>	Oct 03, 2005
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>N76418</u>	<u>002</u>	Oct 03, 2005
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N76418</u>	<u>003</u>	Oct 03, 2005
	<u>NORVASC</u>				
<u>AB</u>	PFIZER	<u>EQ 2.5MG BASE</u>	<u>N19787</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>N19787</u>	<u>002</u>	Jul 31, 1992
<u>AB</u>	+	<u>EQ 10MG BASE</u>	<u>N19787</u>	<u>003</u>	Jul 31, 1992

AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM

TABLET; ORAL

CADUET

PFIZER

EQ 2.5MG BASE;EQ 10MG BASE	N21540	009	Jul 29, 2004
EQ 2.5MG BASE;EQ 20MG BASE	N21540	010	Jul 29, 2004
EQ 2.5MG BASE;EQ 40MG BASE	N21540	011	Jul 29, 2004
EQ 5MG BASE;EQ 10MG BASE	N21540	001	Jan 30, 2004
EQ 5MG BASE;EQ 20MG BASE	N21540	002	Jan 30, 2004
EQ 5MG BASE;EQ 40MG BASE	N21540	003	Jan 30, 2004
EQ 5MG BASE;EQ 80MG BASE	N21540	004	Jan 30, 2004
EQ 10MG BASE;EQ 10MG BASE	N21540	005	Jan 30, 2004
EQ 10MG BASE;EQ 20MG BASE	N21540	006	Jan 30, 2004
EQ 10MG BASE;EQ 40MG BASE	N21540	007	Jan 30, 2004
EQ 10MG BASE;EQ 80MG BASE	N21540	008	Jan 30, 2004

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

LOTREL

NOVARTIS

EQ 2.5MG BASE;10MG	N20364	002	Mar 03, 1995
EQ 5MG BASE;10MG	N20364	003	Mar 03, 1995
EQ 5MG BASE;20MG	N20364	004	Mar 03, 1995
EQ 10MG BASE;20MG	N20364	005	Jun 20, 2002

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE IN PLASTIC CONTAINER

+ HOSPIRA

5MEQ/ML

N88366 001 Jun 13, 1984

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	CLAY PARK	<u>EQ 12% BASE</u>	<u>N75774</u>	<u>001</u>	May 01, 2002
<u>AB</u>	TARO	<u>EQ 12% BASE</u>	<u>N75883</u>	<u>001</u>	Apr 10, 2003
	<u>LAC-HYDRIN</u>				
<u>AB</u>	+ WESTWOOD SQUIBB	<u>EQ 12% BASE</u>	<u>N20508</u>	<u>001</u>	Aug 29, 1996

LOTION; TOPICAL

AMMONIUM LACTATE

<u>AB</u>	CLAY PARK	<u>EQ 12% BASE</u>	<u>N75570</u>	<u>001</u>	Jun 23, 2004
<u>AB</u>	PADDOCK	<u>EQ 12% BASE</u>	<u>N75575</u>	<u>001</u>	Jun 11, 2002
<u>AB</u>	TARO	<u>EQ 12% BASE</u>	<u>N76216</u>	<u>001</u>	May 28, 2004
	<u>LAC-HYDRIN</u>				
<u>AB</u>	+ WESTWOOD SQUIBB	<u>EQ 12% BASE</u>	<u>N19155</u>	<u>001</u>	Apr 24, 1985

AMOXAPINE

TABLET; ORAL

AMOXAPINE

<u>AB</u>	SANDOZ	<u>25MG</u>	<u>N72943</u>	<u>001</u>	Jun 28, 1991
<u>AB</u>		<u>50MG</u>	<u>N72944</u>	<u>001</u>	Jun 28, 1991

PRESCRIPTION DRUG PRODUCT LIST

3 - 27 (of 360)

AMOXAPINE

TABLET; ORAL

AMOXAPINE

<u>AB</u>	SANDOZ	<u>100MG</u>	<u>N72878</u>	<u>001</u>	Jun 28, 1991
<u>AB</u>		<u>150MG</u>	<u>N72879</u>	<u>001</u>	Jun 28, 1991
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>N72418</u>	<u>001</u>	May 11, 1989
<u>AB</u>		<u>25MG</u>	<u>N72688</u>	<u>001</u>	Aug 28, 1992
<u>AB</u>		<u>50MG</u>	<u>N72419</u>	<u>001</u>	May 11, 1989
<u>AB</u>		<u>50MG</u>	<u>N72689</u>	<u>001</u>	Aug 28, 1992
<u>AB</u>		<u>100MG</u>	<u>N72420</u>	<u>001</u>	May 11, 1989
<u>AB</u>		<u>100MG</u>	<u>N72690</u>	<u>001</u>	Aug 28, 1992
<u>AB</u>	+	<u>150MG</u>	<u>N72421</u>	<u>001</u>	May 11, 1989
<u>AB</u>		<u>150MG</u>	<u>N72691</u>	<u>001</u>	Aug 28, 1992

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

<u>AB</u>	AUROBINDO	<u>250MG</u>	<u>N65271</u>	<u>001</u>	Nov 09, 2005
<u>AB</u>		<u>500MG</u>	<u>N65271</u>	<u>002</u>	Nov 09, 2005
<u>AB</u>	BIOCHEMIE	<u>250MG</u>	<u>N64076</u>	<u>001</u>	Sep 30, 1994
<u>AB</u>		<u>500MG</u>	<u>N64076</u>	<u>002</u>	Sep 30, 1994
<u>AB</u>	CLONMEL	<u>250MG</u>	<u>N62884</u>	<u>001</u>	Feb 25, 1988
<u>AB</u>		<u>500MG</u>	<u>N62881</u>	<u>001</u>	Feb 25, 1988
<u>AB</u>	CONSOLIDATED PHARM	<u>250MG</u>	<u>N62058</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N62058</u>	<u>002</u>	
<u>AB</u>	RANBAXY	<u>250MG</u>	<u>N65016</u>	<u>001</u>	Apr 08, 1999
<u>AB</u>		<u>500MG</u>	<u>N65016</u>	<u>002</u>	Apr 08, 1999
<u>AB</u>	TEVA	<u>250MG</u>	<u>N61926</u>	<u>001</u>	
<u>AB</u>		<u>250MG</u>	<u>N62853</u>	<u>001</u>	Dec 22, 1987
<u>AB</u>		<u>500MG</u>	<u>N61926</u>	<u>003</u>	
<u>AB</u>		<u>500MG</u>	<u>N62854</u>	<u>001</u>	Dec 22, 1987

AMOXIL

<u>AB</u>	GLAXOSMITHKLINE	<u>250MG</u>	<u>N62216</u>	<u>001</u>	
<u>AB</u>	+	<u>500MG</u>	<u>N62216</u>	<u>004</u>	
<u>TRIMOX</u>					
<u>AB</u>	APOTHECON	<u>250MG</u>	<u>N61885</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N61885</u>	<u>002</u>	

FOR SUSPENSION; ORAL

AMOXICILLIN

<u>AB</u>	CLONMEL	<u>125MG/5ML</u>	<u>N62927</u>	<u>001</u>	Nov 25, 1988
<u>AB</u>		<u>250MG/5ML</u>	<u>N62927</u>	<u>002</u>	Nov 25, 1988
<u>AB</u>	CONSOLIDATED PHARM	<u>125MG/5ML</u>	<u>N62059</u>	<u>001</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>N62059</u>	<u>002</u>	
<u>AB</u>	RANBAXY	<u>200MG/5ML</u>	<u>N65113</u>	<u>001</u>	Nov 29, 2002
<u>AB</u>		<u>400MG/5ML</u>	<u>N65113</u>	<u>002</u>	Nov 29, 2002
<u>AB</u>	TEVA	<u>125MG/5ML</u>	<u>N61931</u>	<u>001</u>	
<u>AB</u>		<u>125MG/5ML</u>	<u>N62946</u>	<u>001</u>	Nov 01, 1988
<u>AB</u>		<u>200MG/5ML</u>	<u>N65119</u>	<u>001</u>	Dec 04, 2002
<u>AB</u>		<u>250MG/5ML</u>	<u>N61931</u>	<u>002</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>N63001</u>	<u>001</u>	Jan 06, 1989
<u>AB</u>		<u>400MG/5ML</u>	<u>N65119</u>	<u>002</u>	Dec 04, 2002

AMOXICILLIN PEDIATRIC

<u>AB</u>	TEVA	<u>50MG/ML</u>	<u>N61931</u>	<u>003</u>	Dec 01, 1982
<u>AMOXIL</u>					
<u>AB</u>	GLAXOSMITHKLINE	<u>50MG/ML</u>	<u>N62226</u>	<u>005</u>	
<u>AB</u>		<u>125MG/5ML</u>	<u>N62226</u>	<u>001</u>	
<u>AB</u>		<u>200MG/5ML</u>	<u>N50760</u>	<u>001</u>	Apr 15, 1999
<u>AB</u>	+	<u>250MG/5ML</u>	<u>N62226</u>	<u>002</u>	
<u>AB</u>	+	<u>400MG/5ML</u>	<u>N50760</u>	<u>002</u>	Apr 15, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 28 (of 360)

AMOXICILLIN

FOR SUSPENSION; ORAL

LAROTID

<u>AB</u>	GLAXOSMITHKLINE	<u>125MG/5ML</u>	<u>N62226</u>	<u>003</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>N62226</u>	<u>004</u>	
	<u>TRIMOX</u>				
<u>AB</u>	APOTHECON	<u>50MG/ML</u>	<u>N61886</u>	<u>001</u>	
<u>AB</u>		<u>125MG/5ML</u>	<u>N61886</u>	<u>002</u>	
<u>AB</u>		<u>125MG/5ML</u>	<u>N62885</u>	<u>001</u>	Mar 08, 1988
<u>AB</u>		<u>250MG/5ML</u>	<u>N61886</u>	<u>003</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>N62885</u>	<u>002</u>	Mar 08, 1988

TABLET; ORAL

AMOXICILLIN

<u>AB</u>	AUROBINDO	<u>500MG</u>	<u>N65256</u>	<u>001</u>	Nov 09, 2005
<u>AB</u>		<u>875MG</u>	<u>N65256</u>	<u>002</u>	Nov 09, 2005
<u>AB</u>	RANBAXY	<u>500MG</u>	<u>N65059</u>	<u>001</u>	Nov 24, 2000
<u>AB</u>		<u>875MG</u>	<u>N65059</u>	<u>002</u>	Nov 24, 2000
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>N65228</u>	<u>001</u>	Jul 13, 2005
<u>AB</u>		<u>875MG</u>	<u>N65228</u>	<u>002</u>	Jul 13, 2005
<u>AB</u>	TEVA	<u>500MG</u>	<u>N65056</u>	<u>001</u>	Sep 18, 2000
<u>AB</u>		<u>875MG</u>	<u>N65056</u>	<u>002</u>	Sep 18, 2000
	<u>AMOXIL</u>				
<u>AB</u>	GLAXOSMITHKLINE	<u>500MG</u>	<u>N50754</u>	<u>002</u>	Jul 10, 1998
<u>AB</u>	+	<u>875MG</u>	<u>N50754</u>	<u>001</u>	Jul 10, 1998

TABLET, CHEWABLE; ORAL

AMOXICILLIN

<u>AB</u>	CLONMEL	<u>125MG</u>	<u>N64139</u>	<u>001</u>	Jan 29, 1996
<u>AB</u>		<u>250MG</u>	<u>N64139</u>	<u>002</u>	Jan 29, 1996
<u>AB</u>	RANBAXY	<u>125MG</u>	<u>N65021</u>	<u>001</u>	Dec 23, 1999
<u>AB</u>		<u>200MG</u>	<u>N65060</u>	<u>001</u>	Nov 29, 2000
<u>AB</u>		<u>250MG</u>	<u>N65021</u>	<u>002</u>	Dec 23, 1999
<u>AB</u>		<u>400MG</u>	<u>N65060</u>	<u>002</u>	Nov 29, 2000
<u>AB</u>	TEVA	<u>125MG</u>	<u>N64013</u>	<u>002</u>	Sep 11, 1995
<u>AB</u>		<u>125MG</u>	<u>N64031</u>	<u>001</u>	Dec 19, 1996
<u>AB</u>		<u>250MG</u>	<u>N64013</u>	<u>001</u>	Dec 22, 1992
<u>AB</u>		<u>250MG</u>	<u>N64031</u>	<u>002</u>	Dec 19, 1996
	<u>AMOXIL</u>				
<u>AB</u>	GLAXOSMITHKLINE	<u>200MG</u>	<u>N50761</u>	<u>001</u>	Apr 15, 1999
<u>AB</u>	+	<u>400MG</u>	<u>N50761</u>	<u>002</u>	Apr 15, 1999

TABLET, FOR SUSPENSION; ORAL

DISPERMOX

	RANBAXY	200MG	N65080	002	Aug 11, 2003
+		400MG	N65080	001	Aug 11, 2003
+		600MG	N65159	001	Dec 04, 2003

AMOXICILLIN; CLARITHROMYCIN; LANSOPRAZOLE

CAPSULE, TABLET, CAPSULE, DELAYED REL PELLETS; ORAL

PREVPAC

+	TAP PHARM	500MG, N/A, N/A; N/A, 500MG, N/A; N/A, N/A, 30MG	N50757	001	Dec 02, 1997
---	-----------	--	--------	-----	--------------

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	HIKMA PHARMS	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>N65191</u>	<u>002</u>	Jan 25, 2005
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>N65191</u>	<u>001</u>	Jan 25, 2005
<u>AB</u>	LEK PHARMS	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>N65098</u>	<u>001</u>	Dec 16, 2002
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>N65098</u>	<u>002</u>	Dec 16, 2002
<u>AB</u>	RANBAXY	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>N65132</u>	<u>001</u>	Mar 19, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 29 (of 360)

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	RANBAXY	<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>N65132</u>	<u>002</u>	Mar 19, 2003
<u>AB</u>	SANDOZ	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>N65066</u>	<u>001</u>	Jun 05, 2002
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>N65066</u>	<u>002</u>	Jun 05, 2002
<u>AB</u>	TEVA	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>N65089</u>	<u>001</u>	May 25, 2004
<u>AB</u>		<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>N65089</u>	<u>002</u>	May 25, 2004
<u>AB</u>		<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>N65162</u>	<u>001</u>	Mar 12, 2004
	AUGMENTIN '125'				
	GLAXOSMITHKLINE	125MG/5ML;EQ 31.25MG BASE/5ML	N50575	001	Aug 06, 1984
	AUGMENTIN '200'				
<u>AB</u>	GLAXOSMITHKLINE	<u>200MG/5ML;EQ 28.5MG BASE/5ML</u>	<u>N50725</u>	<u>001</u>	May 31, 1996
	AUGMENTIN '250'				
	+ GLAXOSMITHKLINE	250MG/5ML;EQ 62.5MG BASE/5ML	N50575	002	Aug 06, 1984
	AUGMENTIN '400'				
<u>AB</u>	+ GLAXOSMITHKLINE	<u>400MG/5ML;EQ 57MG BASE/5ML</u>	<u>N50725</u>	<u>002</u>	May 31, 1996
	AUGMENTIN ES-600				
<u>AB</u>	+ GLAXOSMITHKLINE	<u>600MG/5ML;EQ 42.9MG BASE/5ML</u>	<u>N50755</u>	<u>001</u>	Jun 22, 2001

TABLET; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	LEK PHARMS	<u>500MG;EQ 125MG BASE</u>	<u>N65117</u>	<u>001</u>	Nov 27, 2002
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>N65093</u>	<u>001</u>	Nov 21, 2002
<u>AB</u>	RANBAXY	<u>500MG;EQ 125MG BASE</u>	<u>N65109</u>	<u>001</u>	Nov 04, 2002
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>N65102</u>	<u>001</u>	Sep 17, 2002
<u>AB</u>	SANDOZ	<u>250MG;EQ 125MG BASE</u>	<u>N65189</u>	<u>001</u>	Aug 23, 2005
<u>AB</u>		<u>500MG;EQ 125MG BASE</u>	<u>N65064</u>	<u>001</u>	Mar 15, 2002
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>N65063</u>	<u>001</u>	Mar 14, 2002
<u>AB</u>	TEVA	<u>500MG;EQ 125MG BASE</u>	<u>N65101</u>	<u>001</u>	Oct 30, 2002
<u>AB</u>		<u>875MG;EQ 125MG BASE</u>	<u>N65096</u>	<u>001</u>	Oct 29, 2002
	AUGMENTIN '250'				
<u>AB</u>	GLAXOSMITHKLINE	<u>250MG;EQ 125MG BASE</u>	<u>N50564</u>	<u>001</u>	Aug 06, 1984
	AUGMENTIN '500'				
<u>AB</u>	GLAXOSMITHKLINE	<u>500MG;EQ 125MG BASE</u>	<u>N50564</u>	<u>002</u>	Aug 06, 1984
	AUGMENTIN '875'				
<u>AB</u>	+ GLAXOSMITHKLINE	<u>875MG;EQ 125MG BASE</u>	<u>N50720</u>	<u>001</u>	Feb 13, 1996

TABLET, CHEWABLE; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

<u>AB</u>	RANBAXY	<u>200MG;EQ 28.5MG BASE</u>	<u>N65161</u>	<u>001</u>	Dec 03, 2003
<u>AB</u>		<u>400MG;EQ 57MG BASE</u>	<u>N65161</u>	<u>002</u>	Dec 03, 2003
<u>AB</u>	SANDOZ	<u>200MG;EQ 28.5MG BASE</u>	<u>N65065</u>	<u>001</u>	Apr 18, 2002
<u>AB</u>		<u>400MG;EQ 57MG BASE</u>	<u>N65065</u>	<u>002</u>	Apr 18, 2002
<u>AB</u>	TEVA	<u>200MG;EQ 28.5MG BASE</u>	<u>N65205</u>	<u>001</u>	Feb 09, 2005
<u>AB</u>		<u>400MG;EQ 57MG BASE</u>	<u>N65205</u>	<u>002</u>	Feb 09, 2005
	AUGMENTIN '125'				
	GLAXOSMITHKLINE	125MG;EQ 31.25MG BASE	N50597	001	Jul 22, 1985
	AUGMENTIN '200'				
<u>AB</u>	GLAXOSMITHKLINE	<u>200MG;EQ 28.5MG BASE</u>	<u>N50726</u>	<u>001</u>	May 31, 1996
	AUGMENTIN '250'				
	+ GLAXOSMITHKLINE	250MG;EQ 62.5MG BASE	N50597	002	Jul 22, 1985
	AUGMENTIN '400'				
<u>AB</u>	+ GLAXOSMITHKLINE	<u>400MG;EQ 57MG BASE</u>	<u>N50726</u>	<u>002</u>	May 31, 1996

TABLET, EXTENDED RELEASE; ORAL

AUGMENTIN XR

	+ GLAXOSMITHKLINE	1GM;EQ 62.5MG BASE	N50785	001	Sep 25, 2002
--	-------------------	--------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 30 (of 360)

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE;
DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

ADDERALL XR 10					
SHIRE	2.5MG;2.5MG;2.5MG;2.5MG	N21303	001	Oct 11, 2001	
ADDERALL XR 15					
SHIRE	3.75MG;3.75MG;3.75MG;3.75MG	N21303	006	May 22, 2002	
ADDERALL XR 20					
SHIRE	5MG;5MG;5MG;5MG	N21303	002	Oct 11, 2001	
ADDERALL XR 25					
SHIRE	6.25MG;6.25MG;6.25MG;6.25MG	N21303	004	May 22, 2002	
ADDERALL XR 30					
+ SHIRE	7.5MG;7.5MG;7.5MG;7.5MG	N21303	003	Oct 11, 2001	
ADDERALL XR 5					
SHIRE	1.25MG;1.25MG;1.25MG;1.25MG	N21303	005	May 22, 2002	

TABLET; ORAL

ADDERALL 10					
AB SHIRE	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N11522</u>	<u>007</u>	Feb 13, 1996	
ADDERALL 12.5					
AB SHIRE	<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>N11522</u>	<u>012</u>	Aug 31, 2000	
ADDERALL 15					
AB SHIRE	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>N11522</u>	<u>013</u>	Aug 31, 2000	
ADDERALL 20					
AB SHIRE	<u>5MG;5MG;5MG;5MG</u>	<u>N11522</u>	<u>008</u>	Feb 13, 1996	
ADDERALL 30					
AB + SHIRE	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N11522</u>	<u>010</u>	May 12, 1997	
ADDERALL 5					
AB SHIRE	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N11522</u>	<u>009</u>	May 12, 1997	
ADDERALL 7.5					
AB SHIRE	<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>N11522</u>	<u>011</u>	Aug 31, 2000	

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB ABRIKA PHARMS	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N40472</u>	<u>001</u>	Sep 30, 2003	
AB	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N40472</u>	<u>002</u>	Sep 30, 2003	
AB	<u>5MG;5MG;5MG;5MG</u>	<u>N40472</u>	<u>003</u>	Sep 30, 2003	
AB	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N40472</u>	<u>004</u>	Sep 30, 2003	
AB BARR	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N40422</u>	<u>001</u>	Feb 11, 2002	
AB	<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>N40422</u>	<u>005</u>	Mar 19, 2003	
AB	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N40422</u>	<u>002</u>	Feb 11, 2002	
AB	<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>N40422</u>	<u>006</u>	Mar 19, 2003	
AB	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>N40422</u>	<u>007</u>	Mar 19, 2003	
AB	<u>5MG;5MG;5MG;5MG</u>	<u>N40422</u>	<u>003</u>	Feb 11, 2002	
AB	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N40422</u>	<u>004</u>	Feb 11, 2002	
AB COREPHARMA	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N40444</u>	<u>001</u>	Jun 19, 2002	
AB	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N40444</u>	<u>002</u>	Jun 19, 2002	
AB	<u>5MG;5MG;5MG;5MG</u>	<u>N40444</u>	<u>003</u>	Jun 19, 2002	
AB	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N40444</u>	<u>004</u>	Jun 19, 2002	
AB MALLINCKRODT	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N40440</u>	<u>001</u>	Oct 07, 2003	
AB	<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>N40440</u>	<u>002</u>	Oct 07, 2003	
AB	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N40440</u>	<u>003</u>	Oct 07, 2003	
AB	<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>N40440</u>	<u>004</u>	Oct 07, 2003	
AB	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>N40440</u>	<u>005</u>	Oct 07, 2003	
AB	<u>5MG;5MG;5MG;5MG</u>	<u>N40440</u>	<u>006</u>	Oct 07, 2003	
AB	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N40440</u>	<u>007</u>	Oct 07, 2003	
AB MUTUAL PHARM	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N40480</u>	<u>001</u>	Sep 09, 2003	
AB	<u>1.875MG;1.875MG;1.875MG;1.875MG</u>	<u>N40480</u>	<u>002</u>	Sep 09, 2003	
AB	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N40480</u>	<u>003</u>	Sep 09, 2003	
AB	<u>3.125MG;3.125MG;3.125MG;3.125MG</u>	<u>N40480</u>	<u>004</u>	Sep 09, 2003	
AB	<u>3.75MG;3.75MG;3.75MG;3.75MG</u>	<u>N40480</u>	<u>005</u>	Sep 09, 2003	
AB	<u>5MG;5MG;5MG;5MG</u>	<u>N40480</u>	<u>006</u>	Sep 09, 2003	
AB	<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N40480</u>	<u>007</u>	Sep 09, 2003	
AB SANDOZ	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N40470</u>	<u>001</u>	Sep 27, 2002	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 31 (of 360)

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE;
DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

<u>DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE</u>					
<u>AB</u>	SANDOZ	<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N40439</u>	<u>001</u>	Jun 14, 2002
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>N40439</u>	<u>002</u>	Jun 14, 2002
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N40439</u>	<u>003</u>	Jun 14, 2002
<u>AB</u>	WATSON LABS	<u>1.25MG;1.25MG;1.25MG;1.25MG</u>	<u>N40456</u>	<u>001</u>	May 06, 2003
<u>AB</u>		<u>2.5MG;2.5MG;2.5MG;2.5MG</u>	<u>N40456</u>	<u>002</u>	May 06, 2003
<u>AB</u>		<u>5MG;5MG;5MG;5MG</u>	<u>N40456</u>	<u>003</u>	May 06, 2003
<u>AB</u>		<u>7.5MG;7.5MG;7.5MG;7.5MG</u>	<u>N40456</u>	<u>004</u>	May 06, 2003

AMPHOTERICIN B

CREAM; TOPICAL

FUNGIZONE

+ APOTHECON 3% N50314 001

INJECTABLE; INJECTION

AMPHOTERICIN B

<u>AP</u>	PHARMA TEK	<u>50MG/VIAL</u>	<u>N63206</u>	<u>001</u>	Apr 29, 1992
<u>AP</u>	SICOR PHARMS	<u>50MG/VIAL</u>	<u>N64062</u>	<u>001</u>	Mar 31, 1995

FUNGIZONE

<u>AP</u>	+ APOTHECON	<u>50MG/VIAL</u>	<u>N60517</u>	<u>001</u>	
-----------	-------------	------------------	---------------	------------	--

INJECTABLE, LIPID COMPLEX; INJECTION

ABELCET

+ ENZON 5MG/ML N50724 001 Nov 20, 1995

AMPHOTEC

+ THREE RIVERS PHARMS 50MG/VIAL N50729 001 Nov 22, 1996

+ 100MG/VIAL N50729 002 Nov 22, 1996

INJECTABLE, LIPOSOMAL; INJECTION

AMBISOME

+ ASTELLAS 50MG/VIAL N50740 001 Aug 11, 1997

LOTION; TOPICAL

FUNGIZONE

+ APOTHECON 3% N60570 001

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

<u>AP</u>	IBI	<u>EQ 250MG BASE/VIAL</u>	<u>N62719</u>	<u>001</u>	May 12, 1987
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>N62719</u>	<u>003</u>	May 12, 1987
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N62719</u>	<u>002</u>	May 12, 1987
<u>AP</u>	INSTITUTO BIOCHEMICO	<u>EQ 125MG BASE/VIAL</u>	<u>N62797</u>	<u>001</u>	Jul 12, 1993
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>N62797</u>	<u>002</u>	Jul 12, 1993
<u>AP</u>	+ SANDOZ	<u>EQ 125MG BASE/VIAL</u>	<u>N61395</u>	<u>001</u>	
<u>AP</u>	+	<u>EQ 250MG BASE/VIAL</u>	<u>N61395</u>	<u>002</u>	
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>N61395</u>	<u>003</u>	
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N61395</u>	<u>004</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N62738</u>	<u>001</u>	Feb 19, 1987
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>N61395</u>	<u>005</u>	
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>N62738</u>	<u>002</u>	Feb 19, 1987
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>N61395</u>	<u>006</u>	

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N65074</u>	<u>001</u>	Mar 19, 2002
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>N65074</u>	<u>002</u>	Mar 19, 2002
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>N65076</u>	<u>001</u>	Mar 19, 2002
<u>AP</u>	HANFORD GC	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N65176</u>	<u>001</u>	Nov 30, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 32 (of 360)

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

<u>AP</u>	HANFORD GC	<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>N65176</u>	<u>002</u>	Nov 30, 2005
<u>AP</u>		<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>N65188</u>	<u>001</u>	Nov 25, 2005
<u>AP</u>	INSTITUTO BIOCHIMICO	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N65222</u>	<u>001</u>	Nov 29, 2005
<u>AP</u>		<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>N65222</u>	<u>002</u>	Nov 29, 2005
<u>UNASYN</u>					
<u>AP</u>	+ PFIZER	<u>EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL</u>	<u>N50608</u>	<u>002</u>	Dec 31, 1986
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL</u>	<u>N50608</u>	<u>001</u>	Dec 31, 1986
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL</u>	<u>N50608</u>	<u>005</u>	Dec 10, 1993
		EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	N62901	001	Nov 23, 1988

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

<u>AB</u>	CLONMEL	<u>EQ 250MG BASE</u>	<u>N62883</u>	<u>001</u>	Feb 25, 1988
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N62882</u>	<u>001</u>	Feb 25, 1988
<u>PRINCIPEN</u>					
<u>AB</u>	APOTHECON	<u>EQ 250MG BASE</u>	<u>N61392</u>	<u>001</u>	
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N62888</u>	<u>001</u>	Mar 04, 1988
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N61392</u>	<u>002</u>	
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>N62888</u>	<u>002</u>	Mar 04, 1988

FOR SUSPENSION; ORAL

AMPICILLIN TRIHYDRATE

<u>AB</u>	CLONMEL	<u>EQ 125MG BASE/5ML</u>	<u>N62982</u>	<u>001</u>	Feb 10, 1989
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N62982</u>	<u>002</u>	Feb 10, 1989
<u>AB</u>	PUREPAC PHARM	<u>EQ 250MG BASE/5ML</u>	<u>N61980</u>	<u>002</u>	
<u>PRINCIPEN</u>					
<u>AB</u>	APOTHECON	<u>EQ 125MG BASE/5ML</u>	<u>N61394</u>	<u>002</u>	
<u>AB</u>	+	<u>EQ 250MG BASE/5ML</u>	<u>N61394</u>	<u>003</u>	
		EQ 100MG BASE/ML	N61394	001	

AMPRENAVIR

CAPSULE; ORAL

AGENERASE

+ GLAXOSMITHKLINE 50MG N21007 001 Apr 15, 1999

SOLUTION; ORAL

AGENERASE

+ GLAXOSMITHKLINE 15MG/ML N21039 001 Apr 15, 1999

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

AGRYLIN

<u>AB</u>	SHIRE	<u>EQ 0.5MG BASE</u>	<u>N20333</u>	<u>001</u>	Mar 14, 1997
<u>AB</u>	+	<u>EQ 1MG BASE</u>	<u>N20333</u>	<u>002</u>	Mar 14, 1997

ANAGRELIDE HYDROCHLORIDE

<u>AB</u>	BARR	<u>EQ 0.5MG BASE</u>	<u>N76530</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76530</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	IMPAX LABS	<u>EQ 0.5MG BASE</u>	<u>N76910</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76910</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	IVAX PHARMS	<u>EQ 0.5MG BASE</u>	<u>N76468</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76468</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	MYLAN	<u>EQ 0.5MG BASE</u>	<u>N76811</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76811</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	ROXANE	<u>EQ 0.5MG BASE</u>	<u>N76489</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76489</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>	SANDOZ	<u>EQ 0.5MG BASE</u>	<u>N76683</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76683</u>	<u>002</u>	Apr 18, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 33 (of 360)

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL

ANAGRELIDE HYDROCHLORIDE

<u>AB</u>	WATSON LABS	<u>EQ 0.5MG BASE</u>	<u>N76417</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76417</u>	<u>002</u>	Apr 18, 2005

ANASTROZOLE

TABLET; ORAL

ARIMIDEX

+	ASTRAZENECA	1MG	N20541	001	Dec 27, 1995
---	-------------	-----	--------	-----	--------------

ANISINDIONE

TABLET; ORAL

MIRADON

+	SCHERING	50MG	N10909	003	
---	----------	------	--------	-----	--

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS

APOKYN

+	VERNALIS	30MG/3ML (10MG/ML)	N21264	002	Apr 20, 2004
		20MG/2ML (10MG/ML)	N21264	001	Apr 20, 2004

APRACLONIDINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

IOPIDINE

+	ALCON	EQ 0.5% BASE	N20258	001	Jul 30, 1993
+		EQ 1% BASE	N19779	001	Dec 31, 1987

APREPITANT

CAPSULE; ORAL

EMEND

	MERCK	80MG	N21549	001	Mar 26, 2003
+		125MG	N21549	002	Mar 26, 2003

APROTININ BOVINE

INJECTABLE; INJECTION

TRASYLOL

+	BAYER PHARMS	10,000KIU/ML	N20304	001	Dec 29, 1993
---	--------------	--------------	--------	-----	--------------

ARGATROBAN

INJECTABLE; INJECTION

ARGATROBAN

+	ENCYSIVE	100MG/ML	N20883	001	Jun 30, 2000
---	----------	----------	--------	-----	--------------

ARGININE HYDROCHLORIDE

INJECTABLE; INJECTION

R-GENE 10

+	PHARMACIA AND UPJOHN	10GM/100ML	N16931	001	
---	----------------------	------------	--------	-----	--

ARIPIRAZOLE

SOLUTION; ORAL

ABILIFY

+	OTSUKA	1MG/ML	N21713	001	Dec 10, 2004
---	--------	--------	--------	-----	--------------

TABLET; ORAL

ABILIFY

+	OTSUKA	5MG	N21436	005	Nov 15, 2002
		10MG	N21436	001	Nov 15, 2002
+		15MG	N21436	002	Nov 15, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 34 (of 360)

ARIPIPIRAZOLE

TABLET; ORAL

ABILIFY

OTSUKA

20MG

N21436 003

Nov 15, 2002

+

30MG

N21436 004

Nov 15, 2002

ARSENIC TRIOXIDE

INJECTABLE; INJECTION

TRISENOX

+ CEPHALON

1MG/ML

N21248 001

Sep 25, 2000

ARTICAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

SEPTOCAINE

+ DEPROCO

4%;EQ 0.01MG BASE/ML

N20971 001

Apr 03, 2000

ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; DEXPANTHENOL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; TOCOPHEROL ACETATE; VITAMIN A; VITAMIN K

INJECTABLE; IV (INFUSION)

INFUVITE PEDIATRIC

+ SABEX 2002

80MG/VIAL;0.02MG/VIAL;400
IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.14MG/VIAL;
17MG/VIAL;1MG/VIAL;1.4MG/VIAL;1.2MG/
VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL

N21265 001

Feb 21, 2001

INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE)

+ SABEX 2002

80MG/VIAL;0.02MG/VIAL;400
IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.14MG/VIAL;
17MG/VIAL;1MG/VIAL;1.4MG/VIAL;1.2MG/
VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL

N21646 001

Jan 29, 2004

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PHYTONADIONE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

FOR SOLUTION; IV (INFUSION)

M.V.I. PEDIATRIC

+ MAYNE PHARMA USA

80MG/VIAL;0.02MG/VIAL;0.001MG/VIAL;5MG/
VIAL;0.01MG/VIAL;0.14MG/VIAL;17MG/VIAL;
0.2MG/VIAL;1MG/VIAL;1.4MG/VIAL;EQ
1.2MG BASE/VIAL;0.7MG/VIAL;7MG/VIAL

N18920 001

Sep 21, 2000

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I.-12 (WITHOUT VITAMIN K)

+ MAYNE PHARMA USA

20MG/ML;0.006MG/ML;0.05UGM/ML;1.5MG/ML;
0.0005MG/ML;0.06MG/ML;4MG/ML;0.6MG/ML;0
.36MG/ML;0.6MG/ML;0.1MG/ML;1MG/ML

N08809 006

Sep 09, 2004

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E; VITAMIN K

INJECTABLE; IV (INFUSION)

M.V.I. ADULT

+ MAYNE PHARMA USA

200MG/VIAL;0.06MG/VIAL;0.005MG/VIAL;15M
G/VIAL;0.005MG/VIAL;0.6MG/VIAL;40MG/VIA
L;6MG/VIAL;3.6MG/VIAL;6MG/VIAL;1MG/VIAL
;10MG/VIAL;0.15MG/VIAL

N21625 001

Jan 30, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 35 (of 360)

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E; VITAMIN K

INJECTABLE; IV (INFUSION)

M.V.I. ADULT (PHARMACY BULK PACKAGE)

+ MAYNE PHARMA USA	200MG/5ML; 0.06MG/5ML; 0.005MG/5ML; 15MG/5ML; 0.005MG/5ML; 0.6MG/5ML; 40MG/5ML; 6MG/5ML; 3.6MG/5ML; 6MG/5ML; 1MG/5ML; 10MG/5ML; 0.15MG/5ML	N21643	001	Feb 18, 2004
--------------------	--	--------	-----	--------------

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

FIORINAL

AB + WATSON PHARMS	325MG; 50MG; 40MG	N17534	005	Apr 16, 1986
--------------------	-------------------	--------	-----	--------------

LANORINAL

AB LANNETT	325MG; 50MG; 40MG	N86996	002	Oct 11, 1985
------------	-------------------	--------	-----	--------------

TABLET; ORAL

ASPIRIN AND CAFFEINE W/ BUTALBITAL

AB PUREPAC PHARM	325MG; 50MG; 40MG	N86710	002	Aug 23, 1983
------------------	-------------------	--------	-----	--------------

BUTAL COMPOUND

AB SANDOZ	325MG; 50MG; 40MG	N86398	002	Apr 06, 1984
-----------	-------------------	--------	-----	--------------

BUTALBITAL, ASPIRIN AND CAFFEINE

AB + WEST WARD	325MG; 50MG; 40MG	N86162	002	Feb 16, 1984
----------------	-------------------	--------	-----	--------------

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

AB ANABOLIC	325MG; 50MG; 40MG; 30MG	N75231	001	Nov 30, 2001
-------------	-------------------------	--------	-----	--------------

AB ENDO PHARMS	325MG; 50MG; 40MG; 30MG	N75351	001	Mar 05, 1999
----------------	-------------------------	--------	-----	--------------

AB STEVENS J	325MG; 50MG; 40MG; 30MG	N74951	001	Aug 31, 1998
--------------	-------------------------	--------	-----	--------------

AB WATSON LABS	325MG; 50MG; 40MG; 30MG	N74359	001	Aug 31, 1995
----------------	-------------------------	--------	-----	--------------

FIORINAL W/CODEINE NO 3

AB + WATSON PHARMS	325MG; 50MG; 40MG; 30MG	N19429	003	Oct 26, 1990
--------------------	-------------------------	--------	-----	--------------

ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

SYNALGOS-DC

+ LEITNER PHARMS	356.4MG; 30MG; 16MG	N11483	004	Sep 06, 1983
------------------	---------------------	--------	-----	--------------

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

INVAGESIC

AB SANDOZ	385MG; 30MG; 25MG	N74817	001	Nov 27, 1996
-----------	-------------------	--------	-----	--------------

INVAGESIC FORTE

AB SANDOZ	770MG; 60MG; 50MG	N74817	002	Nov 27, 1996
-----------	-------------------	--------	-----	--------------

NORGESIC

AB 3M	385MG; 30MG; 25MG	N13416	003	Oct 27, 1982
-------	-------------------	--------	-----	--------------

NORGESIC FORTE

AB + 3M	770MG; 60MG; 50MG	N13416	004	Oct 27, 1982
---------	-------------------	--------	-----	--------------

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

AB SANDOZ	385MG; 30MG; 25MG	N74654	001	Dec 31, 1996
-----------	-------------------	--------	-----	--------------

AB	770MG; 60MG; 50MG	N74654	002	Dec 31, 1996
----	-------------------	--------	-----	--------------

AB STEVENS J	385MG; 30MG; 25MG	N74988	001	Apr 30, 1999
--------------	-------------------	--------	-----	--------------

AB	770MG; 60MG; 50MG	N74988	002	Apr 30, 1999
----	-------------------	--------	-----	--------------

ORPHENGESIC

AB PAR PHARM	385MG; 30MG; 25MG	N75141	001	May 29, 1998
--------------	-------------------	--------	-----	--------------

ORPHENGESIC FORTE

AB PAR PHARM	770MG; 60MG; 50MG	N75141	002	May 29, 1998
--------------	-------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 36 (of 360)

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON COMPOUND-65

<u>AA</u>	+	XANODYNE PHARM	<u>389MG;32.4MG;65MG</u>	<u>N10996</u>	<u>007</u>	Mar 08, 1983
-----------	---	----------------	--------------------------	---------------	------------	--------------

PROPOXYPHENE COMPOUND 65

<u>AA</u>		TEVA	<u>389MG;32.4MG;65MG</u>	<u>N89025</u>	<u>001</u>	Mar 29, 1985
-----------	--	------	--------------------------	---------------	------------	--------------

ASPIRIN; CARISOPRODOL

TABLET; ORAL

CARISOPRODOL AND ASPIRIN

<u>AB</u>		AMIDE PHARM	<u>325MG;200MG</u>	<u>N40252</u>	<u>001</u>	Dec 10, 1997
-----------	--	-------------	--------------------	---------------	------------	--------------

<u>AB</u>		PAR PHARM	<u>325MG;200MG</u>	<u>N89594</u>	<u>001</u>	Mar 31, 1989
-----------	--	-----------	--------------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>325MG;200MG</u>	<u>N40116</u>	<u>001</u>	Apr 25, 1996
-----------	--	--------	--------------------	---------------	------------	--------------

SOMA COMPOUND

<u>AB</u>	+	MEDPOINTE PHARM HLC	<u>325MG;200MG</u>	<u>N12365</u>	<u>005</u>	Jul 11, 1983
-----------	---	---------------------	--------------------	---------------	------------	--------------

ASPIRIN; CARISOPRODOL; CODEINE PHOSPHATE

TABLET; ORAL

CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE

<u>AB</u>		AMIDE PHARM	<u>325MG;200MG;16MG</u>	<u>N40283</u>	<u>001</u>	Dec 29, 1998
-----------	--	-------------	-------------------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>325MG;200MG;16MG</u>	<u>N40118</u>	<u>001</u>	Apr 16, 1996
-----------	--	--------	-------------------------	---------------	------------	--------------

SOMA COMPOUND W/ CODEINE

<u>AB</u>	+	MEDPOINTE PHARM HLC	<u>325MG;200MG;16MG</u>	<u>N12366</u>	<u>002</u>	Jul 11, 1983
-----------	---	---------------------	-------------------------	---------------	------------	--------------

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

AGGRENOL

	+	BOEHRINGER INGELHEIM	25MG;200MG	N20884	001	Nov 22, 1999
--	---	----------------------	------------	--------	-----	--------------

ASPIRIN; HYDROCODONE BITARTRATE

TABLET; ORAL

AZDONE

	+	SCHWARZ PHARMA	500MG;5MG	N89420	001	Jan 25, 1988
--	---	----------------	-----------	--------	-----	--------------

ASPIRIN; MEPROBAMATE

TABLET; ORAL

EQUAGESIC

	+	LEITNER PHARMS	325MG;200MG	N11702	003	Dec 29, 1983
--	---	----------------	-------------	--------	-----	--------------

ASPIRIN; METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL AND ASPIRIN

<u>AB</u>	+	IVAX PHARMS	<u>325MG;400MG</u>	<u>N87211</u>	<u>001</u>	Dec 22, 1982
-----------	---	-------------	--------------------	---------------	------------	--------------

<u>AB</u>		PAR PHARM	<u>325MG;400MG</u>	<u>N89657</u>	<u>001</u>	Nov 04, 1988
-----------	--	-----------	--------------------	---------------	------------	--------------

<u>AB</u>		STEVENS J	<u>325MG;400MG</u>	<u>N81145</u>	<u>001</u>	Jan 31, 1995
-----------	--	-----------	--------------------	---------------	------------	--------------

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

OXYCODONE AND ASPIRIN

<u>AA</u>		MUTUAL PHARM	<u>325MG;4.5MG;0.38MG</u>	<u>N40260</u>	<u>001</u>	Jul 17, 1998
-----------	--	--------------	---------------------------	---------------	------------	--------------

<u>AA</u>			<u>325MG;4.5MG;0.38MG</u>	<u>N87794</u>	<u>001</u>	May 26, 1982
-----------	--	--	---------------------------	---------------	------------	--------------

<u>AA</u>		WATSON LABS	<u>325MG;4.5MG;0.38MG</u>	<u>N40255</u>	<u>001</u>	Feb 27, 1998
-----------	--	-------------	---------------------------	---------------	------------	--------------

PERCODAN

<u>AA</u>	+	ENDO PHARMS	<u>325MG;4.5MG;0.38MG</u>	<u>N07337</u>	<u>006</u>	
-----------	---	-------------	---------------------------	---------------	------------	--

PERCODAN-DEMI

<u>AA</u>	+	ENDO PHARMS	<u>325MG;2.25MG;0.19MG</u>	<u>N07337</u>	<u>005</u>	
-----------	---	-------------	----------------------------	---------------	------------	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 37 (of 360)

ASPIRIN; PRAVASTATIN SODIUM

TABLET, TABLET; ORAL

PRAVIGARD PAC (COPACKAGED)

BRISTOL MYERS SQUIBB	81MG, N/A; N/A, 40MG	N21387	002	Jun 24, 2003
	81MG, N/A; N/A, 80MG	N21387	003	Jun 24, 2003
	81MG, N/A; N/A, 20MG	N21387	001	Jun 24, 2003
	325MG, N/A; N/A, 80MG	N21387	006	Jun 24, 2003
	325MG, N/A; N/A, 20MG	N21387	004	Jun 24, 2003
+	325MG, N/A; N/A, 40MG	N21387	005	Jun 24, 2003

ATAZANAVIR SULFATE

CAPSULE; ORAL

REYATAZ

BRISTOL MYERS SQUIBB	EQ 100MG BASE	N21567	001	Jun 20, 2003
	EQ 150MG BASE	N21567	002	Jun 20, 2003
	EQ 200MG BASE	N21567	003	Jun 20, 2003
+				

ATENOLOL

INJECTABLE; INJECTION

TENORMIN

+	ASTRAZENECA	0.5MG/ML	N19058	001	Sep 13, 1989
---	-------------	----------	--------	-----	--------------

TABLET; ORAL

ATENOLOL

<u>AB</u>	CLONMEL HLTHCARE	<u>25MG</u>	<u>N74099</u>	<u>001</u>	Apr 28, 1992
<u>AB</u>		<u>50MG</u>	<u>N73542</u>	<u>001</u>	Dec 19, 1991
<u>AB</u>		<u>100MG</u>	<u>N73543</u>	<u>001</u>	Dec 19, 1991
<u>AB</u>	GENPHARM	<u>25MG</u>	<u>N74126</u>	<u>003</u>	Aug 26, 1998
<u>AB</u>		<u>50MG</u>	<u>N74126</u>	<u>001</u>	Mar 23, 1994
<u>AB</u>		<u>100MG</u>	<u>N74126</u>	<u>002</u>	Mar 23, 1994
<u>AB</u>	IPR	<u>25MG</u>	<u>N73646</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>		<u>50MG</u>	<u>N72303</u>	<u>001</u>	Jul 15, 1988
<u>AB</u>		<u>100MG</u>	<u>N72304</u>	<u>001</u>	Jul 15, 1988
<u>AB</u>	MARTEC	<u>50MG</u>	<u>N74127</u>	<u>001</u>	Feb 21, 1995
<u>AB</u>		<u>100MG</u>	<u>N74127</u>	<u>002</u>	Feb 21, 1995
<u>AB</u>	MUTUAL PHARM	<u>25MG</u>	<u>N74499</u>	<u>001</u>	Jul 30, 1997
<u>AB</u>		<u>50MG</u>	<u>N73475</u>	<u>001</u>	Mar 30, 1993
<u>AB</u>		<u>100MG</u>	<u>N73476</u>	<u>001</u>	Mar 30, 1993
<u>AB</u>	MYLAN	<u>25MG</u>	<u>N73457</u>	<u>002</u>	Apr 26, 1999
<u>AB</u>		<u>50MG</u>	<u>N73456</u>	<u>001</u>	Jan 24, 1992
<u>AB</u>		<u>100MG</u>	<u>N73457</u>	<u>001</u>	Jan 24, 1992
<u>AB</u>	PLIVA	<u>25MG</u>	<u>N74101</u>	<u>001</u>	Jul 17, 1997
<u>AB</u>		<u>50MG</u>	<u>N74101</u>	<u>002</u>	Jul 17, 1997
<u>AB</u>		<u>100MG</u>	<u>N74101</u>	<u>003</u>	Jul 17, 1997
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>N74052</u>	<u>001</u>	May 01, 1992
<u>AB</u>		<u>25MG</u>	<u>N74265</u>	<u>001</u>	Feb 28, 1994
<u>AB</u>		<u>50MG</u>	<u>N73025</u>	<u>001</u>	Sep 17, 1991
<u>AB</u>		<u>50MG</u>	<u>N74265</u>	<u>002</u>	Feb 28, 1994
<u>AB</u>		<u>100MG</u>	<u>N73026</u>	<u>001</u>	Sep 17, 1991
<u>AB</u>		<u>100MG</u>	<u>N74265</u>	<u>003</u>	Feb 28, 1994
<u>AB</u>	SCS	<u>50MG</u>	<u>N73676</u>	<u>001</u>	Oct 30, 1992
<u>AB</u>		<u>100MG</u>	<u>N73676</u>	<u>002</u>	Oct 30, 1992
<u>AB</u>	TEVA	<u>25MG</u>	<u>N74056</u>	<u>003</u>	Jul 19, 2004
<u>AB</u>		<u>50MG</u>	<u>N73315</u>	<u>001</u>	May 28, 1993
<u>AB</u>		<u>50MG</u>	<u>N74056</u>	<u>001</u>	Jan 18, 1995
<u>AB</u>		<u>100MG</u>	<u>N73316</u>	<u>001</u>	May 28, 1993
<u>AB</u>		<u>100MG</u>	<u>N74056</u>	<u>002</u>	Jan 18, 1995
<u>AB</u>		<u>50MG</u>	<u>N74120</u>	<u>001</u>	Feb 24, 1995
<u>AB</u>	TEVA PHARMS	<u>100MG</u>	<u>N74120</u>	<u>002</u>	Feb 24, 1995
<u>AB</u>		<u>50MG</u>	<u>N73352</u>	<u>001</u>	Dec 27, 1991
<u>AB</u>		<u>100MG</u>	<u>N73353</u>	<u>001</u>	Dec 27, 1991

PRESCRIPTION DRUG PRODUCT LIST

3 - 38 (of 360)

ATENOLOL

TABLET; ORAL

ATENOLOL

<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>N76900</u>	<u>001</u>	Jan 28, 2005
<u>AB</u>		<u>50MG</u>	<u>N76900</u>	<u>002</u>	Jan 28, 2005
<u>AB</u>		<u>100MG</u>	<u>N76900</u>	<u>003</u>	Jan 28, 2005
<u>TENORMIN</u>					
<u>AB</u>	ASTRAZENECA	<u>25MG</u>	<u>N18240</u>	<u>004</u>	Apr 09, 1990
<u>AB</u>		<u>50MG</u>	<u>N18240</u>	<u>001</u>	
<u>AB</u>	+	<u>100MG</u>	<u>N18240</u>	<u>002</u>	

ATENOLOL; CHLORTHALIDONE

TABLET; ORAL

ATENOLOL AND CHLORTHALIDONE

<u>AB</u>	IPR	<u>50MG; 25MG</u>	<u>N72301</u>	<u>001</u>	May 31, 1990
<u>AB</u>		<u>100MG; 25MG</u>	<u>N72302</u>	<u>001</u>	May 31, 1990
<u>AB</u>	MARTEC	<u>50MG; 25MG</u>	<u>N74404</u>	<u>001</u>	May 14, 1998
<u>AB</u>		<u>100MG; 25MG</u>	<u>N74404</u>	<u>002</u>	May 14, 1998
<u>AB</u>	MUTUAL PHARM	<u>50MG; 25MG</u>	<u>N73581</u>	<u>001</u>	Apr 29, 1993
<u>AB</u>		<u>100MG; 25MG</u>	<u>N73582</u>	<u>001</u>	Apr 29, 1993
<u>AB</u>	MYLAN	<u>50MG; 25MG</u>	<u>N74203</u>	<u>001</u>	Oct 31, 1993
<u>AB</u>		<u>100MG; 25MG</u>	<u>N74203</u>	<u>002</u>	Oct 31, 1993
<u>AB</u>	PLIVA	<u>50MG; 25MG</u>	<u>N74107</u>	<u>001</u>	Sep 24, 1997
<u>AB</u>		<u>100MG; 25MG</u>	<u>N74107</u>	<u>002</u>	Sep 24, 1997
<u>AB</u>	WATSON LABS	<u>50MG; 25MG</u>	<u>N73665</u>	<u>001</u>	Jul 02, 1992
<u>AB</u>		<u>100MG; 25MG</u>	<u>N73665</u>	<u>002</u>	Jul 02, 1992
<u>TENORETIC 100</u>					
<u>AB</u>	+ ASTRAZENECA	<u>100MG; 25MG</u>	<u>N18760</u>	<u>001</u>	Jun 08, 1984
<u>TENORETIC 50</u>					
<u>AB</u>	ASTRAZENECA	<u>50MG; 25MG</u>	<u>N18760</u>	<u>002</u>	Jun 08, 1984

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL

STRATTERA

LILLY

	10MG	N21411	002	Nov 26, 2002
	18MG	N21411	003	Nov 26, 2002
	25MG	N21411	004	Nov 26, 2002
	40MG	N21411	005	Nov 26, 2002
+	60MG	N21411	006	Nov 26, 2002
	80MG	N21411	007	Feb 14, 2005
	100MG	N21411	008	Feb 14, 2005

ATORVASTATIN CALCIUM

TABLET; ORAL

LIPITOR

PFIZER

	EQ 10MG BASE	N20702	001	Dec 17, 1996
	EQ 20MG BASE	N20702	002	Dec 17, 1996
	EQ 40MG BASE	N20702	003	Dec 17, 1996
+	EQ 80MG BASE	N20702	004	Apr 07, 2000

ATOVAQUONE

SUSPENSION; ORAL

MEPRON

+	GLAXOSMITHKLINE	750MG/5ML	N20500	001	Feb 08, 1995
---	-----------------	-----------	--------	-----	--------------

TABLET; ORAL

MEPRON

+	GLAXOSMITHKLINE	250MG	N20259	001	Nov 25, 1992
---	-----------------	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 39 (of 360)

ATOVAQUONE; PROGUANIL HYDROCHLORIDE

TABLET; ORAL

MALARONE

+ GLAXOSMITHKLINE 250MG;100MG N21078 001 Jul 14, 2000

MALARONE PEDIATRIC

+ GLAXOSMITHKLINE 62.5MG;25MG N21078 002 Jul 14, 2000

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATEAP BAXTER HLTHCARE 10MG/ML N74824 001 Sep 30, 1997AP BAXTER HLTHCARE CORP 10MG/ML N74753 001 Jan 23, 1997AP BEDFORD 10MG/ML N74901 001 Jul 18, 1997AP HOSPIRA 10MG/ML N74632 001 Dec 23, 1996AP MARSAM PHARMS LLC 10MG/ML N74945 001 Jul 28, 1998AP MAYNE PHARMA USA 10MG/ML N74740 001 Mar 28, 1997AP SICOR PHARMS 10MG/ML N74784 001 Jun 11, 1997ATRACURIUM BESYLATE PRESERVATIVE FREEAP BAXTER HLTHCARE CORP 10MG/ML N74768 001 Jan 23, 1997AP BEDFORD 10MG/ML N74900 001 Jul 18, 1997AP HOSPIRA 10MG/ML N74633 001 Dec 23, 1996AP 10MG/ML N74639 001 Mar 25, 1997AP MARSAM PHARMS LLC 10MG/ML N74944 001 Jul 28, 1998AP MAYNE PHARMA USA 10MG/ML N74741 001 Mar 28, 1997TRACRIUMAP + HOSPIRA 10MG/ML N18831 002 Jun 20, 1985TRACRIUM PRESERVATIVE FREEAP + HOSPIRA 10MG/ML N18831 001 Nov 23, 1983ATROPINE

INJECTABLE; INJECTION

ATROPEN

+ MERIDIAN MEDCL TECHN EQ 0.25MG SULFATE/0.3ML N17106 004 Sep 17, 2004

+ EQ 0.5MG SULFATE/0.7ML N17106 003 Jun 19, 2003

+ EQ 1MG SULFATE/0.7ML N17106 002 Jun 19, 2003

+ EQ 2MG SULFATE/0.7ML N17106 001

ATROPINE SULFATE

INJECTABLE; IM-IV-SC

ATROPINE SULFATE ANSYR PLASTIC SYRINGE

HOSPIRA 0.05MG/ML N21146 002 Jul 09, 2001

+ 0.1MG/ML N21146 001 Jul 09, 2001

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL

MOTOFEN

+ VALEANT 0.025MG;1MG N17744 002

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE

SCHERER RP 0.025MG;2.5MG N86440 001

SOLUTION; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATEAA ROXANE 0.025MG/5ML;2.5MG/5ML N87708 001 May 03, 1982LOMOTILAA + GD SEARLE LLC 0.025MG/5ML;2.5MG/5ML N12699 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 40 (of 360)

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE

<u>AA</u>	LANNETT	<u>0.025MG;2.5MG</u>	<u>N85372</u>	<u>001</u>	
-----------	---------	----------------------	---------------	------------	--

<u>AA</u>	MYLAN	<u>0.025MG;2.5MG</u>	<u>N85762</u>	<u>001</u>	
-----------	-------	----------------------	---------------	------------	--

<u>AA</u>	PAR PHARM	<u>0.025MG;2.5MG</u>	<u>N40357</u>	<u>001</u>	May 02, 2000
-----------	-----------	----------------------	---------------	------------	--------------

DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE

<u>AA</u>	IVAX PHARMS	<u>0.025MG;2.5MG</u>	<u>N86727</u>	<u>001</u>	
-----------	-------------	----------------------	---------------	------------	--

<u>AA</u>	SANDOZ	<u>0.025MG;2.5MG</u>	<u>N86173</u>	<u>001</u>	
-----------	--------	----------------------	---------------	------------	--

LOMOTIL

<u>AA</u>	+ GD SEARLE LLC	<u>0.025MG;2.5MG</u>	<u>N12462</u>	<u>001</u>	
-----------	-----------------	----------------------	---------------	------------	--

LONOX

<u>AA</u>	SANDOZ	<u>0.025MG;2.5MG</u>	<u>N85311</u>	<u>002</u>	
-----------	--------	----------------------	---------------	------------	--

ATROPINE SULFATE; EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

ENLON-PLUS

+	BAXTER HLTHCARE CORP	0.14MG/ML;10MG/ML	N19677	001	Nov 06, 1991
---	----------------------	-------------------	--------	-----	--------------

+		0.14MG/ML;10MG/ML	N19678	001	Nov 06, 1991
---	--	-------------------	--------	-----	--------------

AURANOFIN

CAPSULE; ORAL

RIDAURA

+	PROMETHEUS LABS	3MG	N18689	001	May 24, 1985
---	-----------------	-----	--------	-----	--------------

AZACITIDINE

INJECTABLE; SUBCUTANEOUS

VIDAZA

+	PHARMION	100MG/VIAL	N50794	001	May 19, 2004
---	----------	------------	--------	-----	--------------

AZATHIOPRINE

TABLET; ORAL

AZASAN

<u>AB</u>	AAIPHARMA LLC	<u>50MG</u>	<u>N75252</u>	<u>001</u>	Jun 07, 1999
-----------	---------------	-------------	---------------	------------	--------------

		25MG	N75252	002	Feb 03, 2003
--	--	------	--------	-----	--------------

		75MG	N75252	003	Feb 03, 2003
--	--	------	--------	-----	--------------

		100MG	N75252	004	Feb 03, 2003
--	--	-------	--------	-----	--------------

AZATHIOPRINE

<u>AB</u>	GENPHARM	<u>50MG</u>	<u>N75568</u>	<u>001</u>	Dec 13, 1999
-----------	----------	-------------	---------------	------------	--------------

<u>AB</u>	ROXANE	<u>50MG</u>	<u>N74069</u>	<u>001</u>	Feb 16, 1996
-----------	--------	-------------	---------------	------------	--------------

IMURAN

<u>AB</u>	+ PROMETHEUS LABS	<u>50MG</u>	<u>N16324</u>	<u>001</u>	
-----------	-------------------	-------------	---------------	------------	--

AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

AZATHIOPRINE SODIUM

+	BEDFORD	EQ 100MG BASE/VIAL	N74419	001	Mar 31, 1995
---	---------	--------------------	--------	-----	--------------

AZELAIC ACID

CREAM; TOPICAL

AZELEX

+	ALLERGAN	20%	N20428	001	Sep 13, 1995
---	----------	-----	--------	-----	--------------

GEL; TOPICAL

FINACEA

+	INTENDIS	15%	N21470	001	Dec 24, 2002
---	----------	-----	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 41 (of 360)

AZELASTINE HYDROCHLORIDESOLUTION/DROPS; OPHTHALMIC
OPTIVAR

+ MEDPOINTE PHARM HLC 0.05% N21127 001 May 22, 2000

SPRAY, METERED; NASAL
ASTELIN

+ MEDPOINTE PHARM HLC EQ 0.125MG BASE/SPRAY N20114 001 Nov 01, 1996

AZITHROMYCINFOR SUSPENSION; ORAL
ZITHROMAX

PFIZER EQ 100MG BASE/5ML N50710 001 Oct 19, 1995

+ EQ 200MG BASE/5ML N50710 002 Oct 19, 1995

+ EQ 1GM BASE/PACKET N50693 001 Sep 28, 1994

FOR SUSPENSION, EXTENDED RELEASE; ORAL
ZMAX

+ PFIZER GLOBAL EQ 2GM BASE/BOT N50797 001 Jun 10, 2005

INJECTABLE; INJECTION

AZITHROMYCINAP AM PHARM EQ 500MG BASE/VIAL N65179 001 Dec 13, 2005ZITHROMAXAP + PFIZER EQ 500MG BASE/VIAL N50733 001 Jan 30, 1997

TABLET; ORAL

AZITHROMYCINAB PLIVA EQ 250MG BASE N65225 001 Nov 14, 2005AB EQ 500MG BASE N65223 001 Nov 14, 2005AB EQ 600MG BASE N65218 001 Nov 14, 2005AB SANDOZ EQ 250MG BASE N65211 001 Nov 14, 2005AB EQ 500MG BASE N65212 001 Nov 14, 2005AB EQ 600MG BASE N65209 001 Nov 14, 2005AB TEVA EQ 250MG BASE N65153 001 Nov 14, 2005AB EQ 500MG BASE N65193 001 Nov 14, 2005AB EQ 600MG BASE N65150 001 Nov 14, 2005ZITHROMAXAB PFIZER EQ 250MG BASE N50711 001 Jul 18, 1996AB EQ 500MG BASE N50784 001 May 24, 2002AB + EQ 600MG BASE N50730 001 Jun 12, 1996AZTREONAM

INJECTABLE; INJECTION

AZACTAM

+ BRISTOL MYERS SQUIBB 500MG/VIAL N50580 001 Dec 31, 1986

+ 1GM/VIAL N50580 002 Dec 31, 1986

+ 2GM/VIAL N50580 003 Dec 31, 1986

AZACTAM IN PLASTIC CONTAINER

+ BRISTOL MYERS SQUIBB 20MG/ML N50632 002 May 24, 1989

+ 40MG/ML N50632 001 May 24, 1989

BACAMPICILLIN HYDROCHLORIDEFOR SUSPENSION; ORAL
SPECTROBID

+ PFIZER 125MG/5ML N50556 001 Mar 23, 1982

TABLET; ORAL

SPECTROBID

+ PFIZER 400MG N50520 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 42 (of 360)

BACITRACIN

INJECTABLE; INJECTION

BACIIM

<u>AP</u>	PHARMA TEK	<u>50,000 UNITS/VIAL</u>	<u>N64153</u>	<u>001</u>	May 09, 1997
-----------	------------	--------------------------	---------------	------------	--------------

BACITRACIN

<u>AP</u>	AM PHARM PARTNERS	<u>50,000 UNITS/VIAL</u>	<u>N65116</u>	<u>001</u>	Dec 03, 2002
-----------	-------------------	--------------------------	---------------	------------	--------------

<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>50,000 UNITS/VIAL</u>	<u>N60733</u>	<u>002</u>	
		10,000 UNITS/VIAL	N60733	001	

OINTMENT; OPHTHALMIC

BACITRACIN

	+ ALTANA	500 UNITS/GM	N61212	001	
--	----------	--------------	--------	-----	--

POWDER; FOR RX COMPOUNDING

BACI-RX

<u>AA</u>	PHARMA TEK	<u>5,000,000 UNITS/BOT</u>	<u>N61580</u>	<u>001</u>	
-----------	------------	----------------------------	---------------	------------	--

BACITRACIN

<u>AA</u>	APOTHEKERNES	<u>5,000,000 UNITS/BOT</u>	<u>N61699</u>	<u>001</u>	
-----------	--------------	----------------------------	---------------	------------	--

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

CORTISPORIN

<u>AT</u>	MONARCH PHARMS	<u>400 UNITS/GM;1%;EQ 3.5MG</u> <u>BASE/GM;10,000 UNITS/GM</u>	<u>N50416</u>	<u>002</u>	
-----------	----------------	---	---------------	------------	--

NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE

<u>AT</u>	+ BAUSCH AND LOMB	<u>400 UNITS/GM;1%;EQ 3.5MG</u> <u>BASE/GM;10,000 UNITS/GM</u>	<u>N64068</u>	<u>001</u>	Oct 30, 1995
-----------	-------------------	---	---------------	------------	--------------

OINTMENT; TOPICAL

CORTISPORIN

	+ MONARCH PHARMS	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;5,000 UNITS/GM	N50168	002	May 04, 1984
--	------------------	---	--------	-----	--------------

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN

<u>AT</u>	ALTANA	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N60764</u>	<u>002</u>	
-----------	--------	--	---------------	------------	--

NEOMYCIN AND POLYMYXIN B SULFATE AND BACITRACIN ZINC

<u>AT</u>	AKORN	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N65088</u>	<u>001</u>	Feb 06, 2004
-----------	-------	--	---------------	------------	--------------

NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC

<u>AT</u>	+ BAUSCH AND LOMB	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N64064</u>	<u>001</u>	Oct 30, 1995
-----------	-------------------	--	---------------	------------	--------------

NEOSPORIN

<u>AT</u>	MONARCH PHARMS	<u>400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N50417</u>	<u>001</u>	
-----------	----------------	--	---------------	------------	--

BACITRACIN ZINC; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN ZINC AND POLYMYXIN B SULFATE

<u>AT</u>	AKORN	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>N64028</u>	<u>001</u>	Jan 30, 1995
-----------	-------	-------------------------------------	---------------	------------	--------------

<u>AT</u>	ALTANA	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>N65022</u>	<u>001</u>	Feb 27, 2002
-----------	--------	-------------------------------------	---------------	------------	--------------

<u>AT</u>	+ BAUSCH AND LOMB	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>N64046</u>	<u>001</u>	Jan 26, 1995
-----------	-------------------	-------------------------------------	---------------	------------	--------------

POLYSPORIN

<u>AT</u>	MONARCH PHARMS	<u>500 UNITS/GM;10,000 UNITS/GM</u>	<u>N61229</u>	<u>001</u>	
-----------	----------------	-------------------------------------	---------------	------------	--

BACITRACIN; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE

	+ PHARMADERM	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	N62166	002	
--	--------------	--	--------	-----	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 43 (of 360)

BACLOFEN

INJECTABLE; INTRATHECAL
LIORESAL

+ MEDTRONIC	0.05MG/ML	N20075 003	Nov 07, 1996
+	0.5MG/ML	N20075 001	Jun 17, 1992
+	2MG/ML	N20075 002	Jun 17, 1992

TABLET; ORAL

BACLOFEN

<u>AB</u> ALPHAPHARM	<u>10MG</u>	<u>N77181</u> <u>001</u>	Jul 29, 2005
<u>AB</u>	<u>20MG</u>	<u>N77121</u> <u>002</u>	Jul 29, 2005
<u>AB</u> IVAX PHARMS	<u>10MG</u>	<u>N72234</u> <u>001</u>	Jul 21, 1988
<u>AB</u> +	<u>20MG</u>	<u>N72235</u> <u>001</u>	Jul 21, 1988
<u>AB</u> LANNETT	<u>20MG</u>	<u>N77241</u> <u>001</u>	Dec 20, 2005
<u>AB</u> USL PHARMA	<u>10MG</u>	<u>N74584</u> <u>001</u>	Aug 19, 1996
<u>AB</u>	<u>20MG</u>	<u>N74584</u> <u>002</u>	Aug 19, 1996
<u>AB</u> VINTAGE PHARMS	<u>10MG</u>	<u>N77156</u> <u>001</u>	Aug 30, 2005
<u>AB</u>	<u>20MG</u>	<u>N77068</u> <u>001</u>	Aug 30, 2005
<u>AB</u> WATSON LABS	<u>10MG</u>	<u>N72824</u> <u>001</u>	Sep 18, 1991
<u>AB</u>	<u>10MG</u>	<u>N73092</u> <u>001</u>	Jan 28, 1994
<u>AB</u>	<u>20MG</u>	<u>N72825</u> <u>001</u>	Sep 18, 1991

TABLET, ORALLY DISINTEGRATING; ORAL
KEMSTRO

SCHWARZ PHARMA	10MG	N21589 001	Oct 30, 2003
+	20MG	N21589 002	Oct 30, 2003

BALSALAZIDE DISODIUM

CAPSULE; ORAL
COLAZAL

+ SALIX PHARMS	750MG	N20610 001	Jul 18, 2000
----------------	-------	------------	--------------

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

QVAR 40			
+ 3M	0.04MG/INH	N20911 002	Sep 15, 2000
QVAR 80			
+ 3M	0.08MG/INH	N20911 001	Sep 15, 2000

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED; NASAL
BECONASE AQ

+ GLAXOSMITHKLINE	EQ 0.042MG DIPROP/SPRAY	N19389 001	Jul 27, 1987
-------------------	-------------------------	------------	--------------

BENZAEPRIIL HYDROCHLORIDE

TABLET; ORAL

BENZAEPRIIL HYDROCHLORIDE

<u>AB</u> ANDRX PHARMS	<u>5MG</u>	<u>N76267</u> <u>001</u>	Feb 11, 2004
<u>AB</u>	<u>10MG</u>	<u>N76267</u> <u>002</u>	Feb 11, 2004
<u>AB</u>	<u>20MG</u>	<u>N76267</u> <u>003</u>	Feb 11, 2004
<u>AB</u>	<u>40MG</u>	<u>N76267</u> <u>004</u>	Feb 11, 2004
<u>AB</u> GENPHARM	<u>5MG</u>	<u>N76476</u> <u>001</u>	Feb 11, 2004
<u>AB</u>	<u>10MG</u>	<u>N76476</u> <u>002</u>	Feb 11, 2004
<u>AB</u>	<u>20MG</u>	<u>N76476</u> <u>003</u>	Feb 11, 2004
<u>AB</u>	<u>40MG</u>	<u>N76476</u> <u>004</u>	Feb 11, 2004
<u>AB</u> IVAX PHARMS	<u>5MG</u>	<u>N76333</u> <u>001</u>	Feb 11, 2004
<u>AB</u>	<u>10MG</u>	<u>N76333</u> <u>002</u>	Feb 11, 2004
<u>AB</u>	<u>20MG</u>	<u>N76333</u> <u>003</u>	Feb 11, 2004
<u>AB</u>	<u>40MG</u>	<u>N76333</u> <u>004</u>	Feb 11, 2004
<u>AB</u> KV PHARM	<u>5MG</u>	<u>N76118</u> <u>001</u>	Feb 11, 2004
<u>AB</u>	<u>10MG</u>	<u>N76118</u> <u>002</u>	Feb 11, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 44 (of 360)

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

<u>AB</u>	KV PHARM	<u>20MG</u>	<u>N76118</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>N76118</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	MYLAN	<u>5MG</u>	<u>N76430</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>N76430</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>N76430</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>N76430</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	RANBAXY	<u>5MG</u>	<u>N76344</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>N76344</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>N76344</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>N76344</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>N76402</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>N76402</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>N76402</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>N76402</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	TEVA	<u>5MG</u>	<u>N76211</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG</u>	<u>N76211</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG</u>	<u>N76211</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>40MG</u>	<u>N76211</u>	<u>004</u>	Feb 11, 2004
<u>LOTENSIN</u>					
<u>AB</u>	NOVARTIS	<u>5MG</u>	<u>N19851</u>	<u>001</u>	Jun 25, 1991
<u>AB</u>		<u>10MG</u>	<u>N19851</u>	<u>002</u>	Jun 25, 1991
<u>AB</u>		<u>20MG</u>	<u>N19851</u>	<u>003</u>	Jun 25, 1991
<u>AB</u>	+	<u>40MG</u>	<u>N19851</u>	<u>004</u>	Jun 25, 1991

BENAZEPRIL HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ANDRX PHARMS	<u>5MG; 6.25MG</u>	<u>N76342</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>N76342</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>N76342</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>N76342</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	GENPHARM	<u>5MG; 6.25MG</u>	<u>N76612</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>N76612</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>N76612</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>N76612</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	IVAX PHARMS	<u>5MG; 6.25MG</u>	<u>N76348</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>N76348</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>N76348</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>N76348</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	MYLAN	<u>5MG; 6.25MG</u>	<u>N76688</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>N76688</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>N76688</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>N76688</u>	<u>004</u>	Feb 11, 2004
<u>AB</u>	RANBAXY	<u>5MG; 6.25MG</u>	<u>N77483</u>	<u>001</u>	Sep 08, 2005
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>N77483</u>	<u>002</u>	Sep 08, 2005
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>N77483</u>	<u>003</u>	Sep 08, 2005
<u>AB</u>		<u>20MG; 25MG</u>	<u>N77483</u>	<u>004</u>	Sep 08, 2005
<u>AB</u>	SANDOZ	<u>5MG; 6.25MG</u>	<u>N76631</u>	<u>001</u>	Feb 11, 2004
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>N76631</u>	<u>002</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>N76631</u>	<u>003</u>	Feb 11, 2004
<u>AB</u>		<u>20MG; 25MG</u>	<u>N76631</u>	<u>004</u>	Feb 11, 2004
<u>LOTENSIN HCT</u>					
<u>AB</u>	NOVARTIS	<u>5MG; 6.25MG</u>	<u>N20033</u>	<u>001</u>	May 19, 1992
<u>AB</u>		<u>10MG; 12.5MG</u>	<u>N20033</u>	<u>002</u>	May 19, 1992
<u>AB</u>		<u>20MG; 12.5MG</u>	<u>N20033</u>	<u>004</u>	May 19, 1992
<u>AB</u>	+	<u>20MG; 25MG</u>	<u>N20033</u>	<u>003</u>	May 19, 1992

PRESCRIPTION DRUG PRODUCT LIST

3 - 45 (of 360)

BENDROFLUMETHIAZIDE

TABLET; ORAL				
NATURETIN-5				
+	APOTHECON	5MG	N12164	002

BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL				
CORZIDE				
	KING PHARMS	5MG; 40MG	N18647	001
+		5MG; 80MG	N18647	002
				May 25, 1983
				May 25, 1983

BENZONATATE

CAPSULE; ORAL				
<u>BENZONATATE</u>				
<u>AA</u>	BANNER PHARMACAPS	<u>100MG</u>	<u>N81297</u>	<u>001</u>
				Jan 29, 1993
<u>TESSALON</u>				
<u>AA</u>	+ FOREST LABS	<u>100MG</u>	<u>N11210</u>	<u>001</u>
			<u>N11210</u>	<u>003</u>
				Jun 25, 1999

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL				
BENZACLIN				
BT	+ DERMIK LABS	5%; EQ 1% BASE	N50756	001
				Dec 21, 2000
	DUAC			
BT	+ STIEFEL	5%; EQ 1% BASE	N50741	001
				Aug 26, 2002

BENZOYL PEROXIDE; ERYTHROMYCIN

GEL; TOPICAL				
<u>BENZAMYCIN</u>				
<u>AB</u>	+ DERMIK LABS	<u>5%; 3%</u>	<u>N50557</u>	<u>001</u>
	BENZAMYCIN PAK			
	+ DERMIK LABS	5%; 3%	N50769	001
				Nov 27, 2000
<u>ERYTHROMYCIN AND BENZOYL PEROXIDE</u>				
<u>AB</u>	QLT USA	<u>5%; 3%</u>	<u>N65112</u>	<u>001</u>
				Mar 29, 2004

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL				
DIDREX				
+	PHARMACIA AND UPJOHN	50MG	N12427	002

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION				
EMETE-CON				
+	PFIZER	EQ 50MG BASE/VIAL	N16820	001

BENZTROPINE MESYLATE

INJECTABLE; INJECTION				
COGENTIN				
+	OVATION PHARMS	1MG/ML	N12015	001
TABLET; ORAL				
<u>BENZTROPINE MESYLATE</u>				
<u>AA</u>	COREPHARMA	<u>0.5MG</u>	<u>N72264</u>	<u>001</u>
				Feb 27, 1989
<u>AA</u>		<u>1MG</u>	<u>N72265</u>	<u>001</u>
				Feb 27, 1989
<u>AA</u>		<u>2MG</u>	<u>N72266</u>	<u>001</u>
				Feb 27, 1989
<u>AA</u>	MUTUAL PHARM	<u>1MG</u>	<u>N81264</u>	<u>001</u>
				Jan 23, 1992
<u>AA</u>		<u>2MG</u>	<u>N81265</u>	<u>001</u>
				Jan 23, 1992
<u>AA</u>	+ PAR PHARM	<u>0.5MG</u>	<u>N88877</u>	<u>001</u>
				Apr 11, 1985
<u>AA</u>	+	<u>1MG</u>	<u>N88894</u>	<u>001</u>
				Apr 11, 1985
<u>AA</u>	+	<u>2MG</u>	<u>N88895</u>	<u>001</u>
				Apr 11, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 46 (of 360)

BENZTROPINE MESYLATE

TABLET; ORAL

BENZTROPINE MESYLATE

<u>AA</u>	PLIVA	<u>0.5MG</u>	<u>N89058</u>	<u>001</u>	Aug 10, 1988
<u>AA</u>		<u>1MG</u>	<u>N89059</u>	<u>001</u>	Aug 10, 1988
<u>AA</u>		<u>2MG</u>	<u>N89060</u>	<u>001</u>	Aug 10, 1988
<u>AA</u>	USL PHARMA	<u>0.5MG</u>	<u>N40103</u>	<u>001</u>	Dec 12, 1996
<u>AA</u>		<u>1MG</u>	<u>N40103</u>	<u>002</u>	Dec 12, 1996
<u>AA</u>		<u>2MG</u>	<u>N40103</u>	<u>003</u>	Dec 12, 1996

BERACTANTSUSPENSION; INTRATRACHEAL
SURVANTA

+	ROSS LABS	25MG/ML	N20032	001	Jul 01, 1991
---	-----------	---------	--------	-----	--------------

BETAINE, ANHYDROUSFOR SOLUTION; ORAL
CYSTADANE

+	ORPHAN MEDCL	1GM/SC00PFUL	N20576	001	Oct 25, 1996
---	--------------	--------------	--------	-----	--------------

BETAMETHASONESYRUP; ORAL
CELESTONE

+	SCHERING	0.6MG/5ML	N14215	002	
---	----------	-----------	--------	-----	--

BETAMETHASONE ACETATE; BETAMETHASONE SODIUM PHOSPHATEINJECTABLE; INJECTION
CELESTONE SOLUSPAN

+	SCHERING	3MG/ML;EQ 3MG BASE/ML	N14602	001	
---	----------	-----------------------	--------	-----	--

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

ALPHATREX

<u>AB</u>	+	SAVAGE LABS	<u>EQ 0.05% BASE</u>	<u>N19138</u>	<u>001</u>	Jun 26, 1984
<u>BETAMETHASONE DIPROPIONATE</u>						
<u>AB</u>		ALPHARMA US PHARMS	<u>EQ 0.05% BASE</u>	<u>N70885</u>	<u>001</u>	Feb 03, 1987
<u>AB</u>		FOUGERA	<u>EQ 0.05% BASE</u>	<u>N19137</u>	<u>001</u>	Jun 26, 1984
<u>AB</u>		TARO	<u>EQ 0.05% BASE</u>	<u>N71143</u>	<u>001</u>	Jun 17, 1987
<u>AB</u>			<u>EQ 0.05% BASE</u>	<u>N73552</u>	<u>001</u>	Apr 30, 1992
<u>AB</u>		TEVA	<u>EQ 0.05% BASE</u>	<u>N71476</u>	<u>001</u>	Aug 10, 1987

CREAM, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>		ALTANA	<u>EQ 0.05% BASE</u>	<u>N76215</u>	<u>001</u>	Dec 09, 2003
<u>AB</u>		CLAY PARK	<u>EQ 0.05% BASE</u>	<u>N76592</u>	<u>001</u>	Dec 09, 2003
<u>AB</u>		QLT USA	<u>EQ 0.05% BASE</u>	<u>N76603</u>	<u>001</u>	Jan 23, 2004
<u>AB</u>		TARO	<u>EQ 0.05% BASE</u>	<u>N76543</u>	<u>001</u>	Dec 09, 2003
<u>DIPROLENE AF</u>						
<u>AB</u>	+	SCHERING	<u>EQ 0.05% BASE</u>	<u>N19555</u>	<u>001</u>	Apr 27, 1987

GEL, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>		ALTANA	<u>EQ 0.05% BASE</u>	<u>N75276</u>	<u>001</u>	May 13, 2003
<u>AB</u>		TARO	<u>EQ 0.05% BASE</u>	<u>N76508</u>	<u>001</u>	Dec 02, 2003
<u>DIPROLENE</u>						
<u>AB</u>	+	SCHERING	<u>EQ 0.05% BASE</u>	<u>N19408</u>	<u>002</u>	Nov 22, 1991

LOTION; TOPICAL

ALPHATREX

<u>AB</u>		SAVAGE LABS	<u>EQ 0.05% BASE</u>	<u>N70273</u>	<u>001</u>	Aug 12, 1985
<u>BETAMETHASONE DIPROPIONATE</u>						
<u>AB</u>		ALPHARMA US PHARMS	<u>EQ 0.05% BASE</u>	<u>N70281</u>	<u>001</u>	Jul 31, 1985

PRESCRIPTION DRUG PRODUCT LIST

3 - 47 (of 360)

BETAMETHASONE DIPROPIONATE

LOTION; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	CLAY PARK	<u>EQ 0.05% BASE</u>	<u>N72538</u>	<u>001</u>	Jan 31, 1990
<u>AB</u>	+ FOUGERA	<u>EQ 0.05% BASE</u>	<u>N70275</u>	<u>001</u>	Aug 12, 1985
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>N72276</u>	<u>001</u>	Aug 24, 1988
<u>AB</u>		<u>EQ 0.05% BASE</u>	<u>N74272</u>	<u>001</u>	Sep 30, 1994
<u>AB</u>	TEVA	<u>EQ 0.05% BASE</u>	<u>N71467</u>	<u>001</u>	Aug 10, 1987
<u>AB</u>	TEVA PHARMS	<u>EQ 0.05% BASE</u>	<u>N71882</u>	<u>001</u>	Jun 06, 1988

LOTION, AUGMENTED; TOPICAL
DIPROLENE

	+ SCHERING	EQ 0.05% BASE	N19716	001	Aug 01, 1988
--	------------	---------------	--------	-----	--------------

OINTMENT; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ALPHARMA US PHARMS	<u>EQ 0.05% BASE</u>	<u>N71012</u>	<u>001</u>	Feb 03, 1987
<u>AB</u>	+ FOUGERA	<u>EQ 0.05% BASE</u>	<u>N19141</u>	<u>001</u>	Sep 04, 1984
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>N74271</u>	<u>001</u>	Sep 15, 1994
<u>AB</u>	TEVA	<u>EQ 0.05% BASE</u>	<u>N71477</u>	<u>001</u>	Aug 10, 1987

OINTMENT, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

<u>AB</u>	ALPHARMA US PHARMS	<u>EQ 0.05% BASE</u>	<u>N74304</u>	<u>001</u>	Aug 31, 1995
<u>AB</u>	ALTANA	<u>EQ 0.05% BASE</u>	<u>N75373</u>	<u>001</u>	Jun 22, 1999
<u>AB</u>	TARO	<u>EQ 0.05% BASE</u>	<u>N76753</u>	<u>001</u>	Oct 12, 2004
<u>DIPROLENE</u>					
<u>AB</u>	+ SCHERING	<u>EQ 0.05% BASE</u>	<u>N18741</u>	<u>001</u>	Jul 27, 1983

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM; TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

<u>AB</u>	ALPHARMA US PHARMS	<u>EQ 0.05% BASE;1%</u>	<u>N76002</u>	<u>001</u>	Aug 02, 2002
<u>AB</u>	ALTANA	<u>EQ 0.05% BASE;1%</u>	<u>N75502</u>	<u>001</u>	Jun 05, 2001
<u>AB</u>	TARO	<u>EQ 0.05% BASE;1%</u>	<u>N75673</u>	<u>001</u>	May 29, 2001
<u>LOTTRISONE</u>					
<u>AB</u>	+ SCHERING	<u>EQ 0.05% BASE;1%</u>	<u>N18827</u>	<u>001</u>	Jul 10, 1984

LOTION; TOPICAL

CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE

<u>AB</u>	ALTANA PHARMA	<u>EQ 0.05% BASE;1%</u>	<u>N76516</u>	<u>001</u>	Jun 16, 2005
<u>AB</u>	TARO	<u>EQ 0.05% BASE;1%</u>	<u>N76493</u>	<u>001</u>	Jul 28, 2004
<u>LOTTRISONE</u>					
<u>AB</u>	+ SCHERING PLOUGH RES	<u>EQ 0.05% BASE;1%</u>	<u>N20010</u>	<u>001</u>	Dec 08, 2000

BETAMETHASONE VALERATEAEROSOL, FOAM; TOPICAL
LUXIQ

	+ CONNETICS	EQ 0.12% BASE	N20934	001	Feb 28, 1999
--	-------------	---------------	--------	-----	--------------

CREAM; TOPICAL

BETAMETHASONE VALERATE

<u>AB</u>	+ FOUGERA	<u>EQ 0.1% BASE</u>	<u>N18861</u>	<u>001</u>	Aug 31, 1983
<u>AB</u>	TARO	<u>EQ 0.1% BASE</u>	<u>N70062</u>	<u>001</u>	May 14, 1985
<u>BETA-VAL</u>					
<u>AB</u>	TEVA	<u>EQ 0.1% BASE</u>	<u>N18642</u>	<u>001</u>	Mar 24, 1983
<u>DERMABET</u>					
<u>AB</u>	TARO	<u>EQ 0.1% BASE</u>	<u>N72041</u>	<u>001</u>	Jan 06, 1988
<u>VALNAC</u>					
<u>AB</u>	ALPHARMA US PHARMS	<u>EQ 0.1% BASE</u>	<u>N70050</u>	<u>001</u>	Oct 10, 1984

LOTION; TOPICAL

BETAMETHASONE VALERATE

<u>AB</u>	ALPHARMA US PHARMS	<u>EQ 0.1% BASE</u>	<u>N70052</u>	<u>001</u>	Jul 31, 1985
<u>AB</u>	+ FOUGERA	<u>EQ 0.1% BASE</u>	<u>N18866</u>	<u>001</u>	Aug 31, 1983

PRESCRIPTION DRUG PRODUCT LIST

3 - 48 (of 360)

BETAMETHASONE VALERATE

LOTION; TOPICAL

BETAMETHASONE VALERATE

<u>AB</u>	TEVA PHARMS	<u>EQ 0.1% BASE</u>	<u>N71883</u>	<u>001</u>	Apr 22, 1988
-----------	-------------	---------------------	---------------	------------	--------------

BETA-VAL

<u>AB</u>	TEVA	<u>EQ 0.1% BASE</u>	<u>N70072</u>	<u>001</u>	Jun 27, 1985
-----------	------	---------------------	---------------	------------	--------------

OINTMENT; TOPICAL

BETAMETHASONE VALERATE

<u>AB</u>	ALPHARMA US PHARMS	<u>EQ 0.1% BASE</u>	<u>N70051</u>	<u>001</u>	Oct 10, 1984
-----------	--------------------	---------------------	---------------	------------	--------------

<u>AB</u>	+ FOUGERA	<u>EQ 0.1% BASE</u>	<u>N18865</u>	<u>001</u>	Aug 31, 1983
-----------	-----------	---------------------	---------------	------------	--------------

BETA-VAL

<u>AB</u>	TEVA	<u>EQ 0.1% BASE</u>	<u>N70069</u>	<u>001</u>	Dec 19, 1985
-----------	------	---------------------	---------------	------------	--------------

BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BETAXOLOL

<u>AT</u>	AKORN	<u>EQ 0.5% BASE</u>	<u>N75386</u>	<u>001</u>	Jun 30, 2000
-----------	-------	---------------------	---------------	------------	--------------

<u>AT</u>	NOVEX	<u>EQ 0.5% BASE</u>	<u>N75446</u>	<u>001</u>	Sep 28, 2000
-----------	-------	---------------------	---------------	------------	--------------

BETAXOLOL HYDROCHLORIDE

<u>AT</u>	BAUSCH AND LOMB	<u>EQ 0.5% BASE</u>	<u>N75630</u>	<u>001</u>	Apr 12, 2001
-----------	-----------------	---------------------	---------------	------------	--------------

BETOPTIC

<u>AT</u>	+ ALCON	<u>EQ 0.5% BASE</u>	<u>N19270</u>	<u>001</u>	Aug 30, 1985
-----------	---------	---------------------	---------------	------------	--------------

SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC S

	+ ALCON	EQ 0.25% BASE	N19845	001	Dec 29, 1989
--	---------	---------------	--------	-----	--------------

TABLET; ORAL

BETAXOLOL HYDROCHLORIDE

<u>AB</u>	AMIDE PHARM	<u>10MG</u>	<u>N75541</u>	<u>001</u>	Oct 22, 1999
-----------	-------------	-------------	---------------	------------	--------------

<u>AB</u>		<u>20MG</u>	<u>N75541</u>	<u>002</u>	Oct 22, 1999
-----------	--	-------------	---------------	------------	--------------

KERLONE

<u>AB</u>	LOREX	<u>10MG</u>	<u>N19507</u>	<u>001</u>	Oct 27, 1989
-----------	-------	-------------	---------------	------------	--------------

<u>AB</u>	+	<u>20MG</u>	<u>N19507</u>	<u>002</u>	Oct 27, 1989
-----------	---	-------------	---------------	------------	--------------

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

<u>AA</u>	AMIDE PHARM	<u>5MG</u>	<u>N40552</u>	<u>001</u>	Oct 28, 2004
-----------	-------------	------------	---------------	------------	--------------

<u>AA</u>		<u>10MG</u>	<u>N40553</u>	<u>001</u>	Oct 28, 2004
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>25MG</u>	<u>N40554</u>	<u>001</u>	Oct 28, 2004
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>50MG</u>	<u>N40551</u>	<u>001</u>	Oct 28, 2004
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>	UPSHER SMITH	<u>5MG</u>	<u>N40633</u>	<u>001</u>	Jun 01, 2005
-----------	--------------	------------	---------------	------------	--------------

<u>AA</u>		<u>10MG</u>	<u>N40634</u>	<u>001</u>	Jun 01, 2005
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>25MG</u>	<u>N40635</u>	<u>001</u>	Jun 01, 2005
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>50MG</u>	<u>N40636</u>	<u>001</u>	Jun 01, 2005
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>	WOCKHARDT	<u>5MG</u>	<u>N40532</u>	<u>001</u>	Sep 29, 2003
-----------	-----------	------------	---------------	------------	--------------

<u>AA</u>		<u>10MG</u>	<u>N40533</u>	<u>001</u>	Sep 29, 2003
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>25MG</u>	<u>N40534</u>	<u>001</u>	Sep 29, 2003
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>50MG</u>	<u>N40518</u>	<u>001</u>	Sep 29, 2003
-----------	--	-------------	---------------	------------	--------------

DUVOID

<u>AA</u>	WELLSPRING PHARM	<u>10MG</u>	<u>N86262</u>	<u>001</u>	
-----------	------------------	-------------	---------------	------------	--

<u>AA</u>		<u>25MG</u>	<u>N86263</u>	<u>001</u>	
-----------	--	-------------	---------------	------------	--

<u>AA</u>		<u>50MG</u>	<u>N85882</u>	<u>003</u>	
-----------	--	-------------	---------------	------------	--

URECHOLINE

<u>AA</u>	+ ODYSSEY PHARMS	<u>5MG</u>	<u>N89095</u>	<u>001</u>	Dec 19, 1985
-----------	------------------	------------	---------------	------------	--------------

<u>AA</u>	+	<u>10MG</u>	<u>N88440</u>	<u>001</u>	May 29, 1984
-----------	---	-------------	---------------	------------	--------------

<u>AA</u>	+	<u>25MG</u>	<u>N88441</u>	<u>001</u>	May 29, 1984
-----------	---	-------------	---------------	------------	--------------

<u>AA</u>	+	<u>50MG</u>	<u>N89096</u>	<u>001</u>	Dec 19, 1985
-----------	---	-------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 49 (of 360)

BEXAROTENE

CAPSULE; ORAL					
TARGRETIN					
+	LIGAND	75MG	N21055	001	Dec 29, 1999
GEL; TOPICAL					
TARGRETIN					
+	LIGAND	1%	N21056	001	Jun 28, 2000

BICALUTAMIDE

TABLET; ORAL					
CASODEX					
+	ASTRAZENECA	50MG	N20498	001	Oct 04, 1995

BIMATOPROST

SOLUTION/DROPS; OPHTHALMIC					
LUMIGAN					
+	ALLERGAN	0.03%	N21275	001	Mar 16, 2001

BIPERIDEN HYDROCHLORIDE

TABLET; ORAL					
AKINETON					
+	ABBOTT	2MG	N12003	001	

BISACODYL; POLYETHYLENE GLYCOL; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL					
HALFLYTELY					
+	BRAINTREE	5MG, N/A; N/A, 210GM; N/A, 0.74GM; N/A, 2.86GM; N/A, 5.6GM	N21551	001	May 10, 2004

BISMUTH SUBSALICYLATE; METRONIDAZOLE; TETRACYCLINE HYDROCHLORIDE

TABLET, CHEWABLE, TABLET, CAPSULE; ORAL					
HELIDAC					
+	PROMETHEUS LABS	262.4MG, N/A, N/A; N/A, 250MG, N/A; N/A, N/A, 500MG	N50719	001	Aug 15, 1996

BISOPROLOL FUMARATE

TABLET; ORAL					
<u>BISOPROLOL FUMARATE</u>					
<u>AB</u>	MUTUAL PHARM	<u>5MG</u>	<u>N75474</u>	<u>001</u>	Oct 25, 2002
<u>AB</u>		<u>10MG</u>	<u>N75474</u>	<u>002</u>	Oct 25, 2002
<u>AB</u>	MYLAN	<u>5MG</u>	<u>N75831</u>	<u>001</u>	Dec 14, 2005
<u>AB</u>		<u>10MG</u>	<u>N75831</u>	<u>002</u>	Dec 14, 2005
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>N75643</u>	<u>001</u>	Nov 16, 2000
<u>AB</u>		<u>10MG</u>	<u>N75643</u>	<u>002</u>	Nov 16, 2000
<u>AB</u>	TEVA PHARMS	<u>5MG</u>	<u>N75644</u>	<u>001</u>	Jun 26, 2001
<u>AB</u>		<u>10MG</u>	<u>N75644</u>	<u>002</u>	Jun 26, 2001
<u>ZEBETA</u>					
<u>AB</u>	DURAMED PHARMS BARR	<u>5MG</u>	<u>N19982</u>	<u>002</u>	Jul 31, 1992
<u>AB</u>	+	<u>10MG</u>	<u>N19982</u>	<u>001</u>	Jul 31, 1992

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL					
<u>BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE</u>					
<u>AB</u>	MYLAN	<u>2.5MG; 6.25MG</u>	<u>N75768</u>	<u>001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>N75768</u>	<u>002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG; 6.25MG</u>	<u>N75768</u>	<u>003</u>	Sep 25, 2000
<u>AB</u>	PUREPAC PHARM	<u>2.5MG; 6.25MG</u>	<u>N75672</u>	<u>001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG; 6.25MG</u>	<u>N75672</u>	<u>002</u>	Sep 25, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 50 (of 360)

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	PUREPAC PHARM	<u>10MG;6.25MG</u>	<u>N75672</u>	<u>003</u>	Sep 25, 2000
<u>AB</u>	SANDOZ	<u>2.5MG;6.25MG</u>	<u>N75527</u>	<u>001</u>	Sep 25, 2000
<u>AB</u>		<u>2.5MG;6.25MG</u>	<u>N75579</u>	<u>001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG;6.25MG</u>	<u>N75527</u>	<u>003</u>	Sep 25, 2000
<u>AB</u>		<u>5MG;6.25MG</u>	<u>N75579</u>	<u>002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG;6.25MG</u>	<u>N75527</u>	<u>002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG;6.25MG</u>	<u>N75579</u>	<u>003</u>	Sep 25, 2000
<u>AB</u>	TEVA	<u>2.5MG;6.25MG</u>	<u>N75686</u>	<u>001</u>	Jan 19, 2001
<u>AB</u>		<u>5MG;6.25MG</u>	<u>N75686</u>	<u>002</u>	Jan 19, 2001
<u>AB</u>		<u>10MG;6.25MG</u>	<u>N75686</u>	<u>003</u>	Jan 19, 2001
<u>AB</u>	WATSON LABS	<u>2.5MG;6.25MG</u>	<u>N75469</u>	<u>001</u>	Sep 25, 2000
<u>AB</u>		<u>5MG;6.25MG</u>	<u>N75469</u>	<u>002</u>	Sep 25, 2000
<u>AB</u>		<u>10MG;6.25MG</u>	<u>N75469</u>	<u>003</u>	Sep 25, 2000
	<u>ZIAC</u>				
<u>AB</u>	DURAMED PHARMS BARR	<u>2.5MG;6.25MG</u>	<u>N20186</u>	<u>003</u>	Mar 26, 1993
<u>AB</u>		<u>5MG;6.25MG</u>	<u>N20186</u>	<u>001</u>	Mar 26, 1993
<u>AB</u>	+	<u>10MG;6.25MG</u>	<u>N20186</u>	<u>002</u>	Mar 26, 1993

BIVALIRUDININJECTABLE; INTRAVENOUS
ANGIOMAX

+ MEDICINES CO 250MG/VIAL N20873 001 Dec 15, 2000

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLENOXANE

<u>AP</u>	+	BRISTOL MYERS SQUIBB	<u>EQ 15 UNITS BASE/VIAL</u>	<u>N50443</u>	<u>001</u>	
<u>AP</u>	+		<u>EQ 30 UNITS BASE/VIAL</u>	<u>N50443</u>	<u>002</u>	Sep 07, 1995
		<u>BLEOMYCIN</u>				
<u>AP</u>		BEDFORD	<u>EQ 15 UNITS BASE/VIAL</u>	<u>N65042</u>	<u>002</u>	Oct 17, 2001
<u>AP</u>			<u>EQ 30 UNITS BASE/VIAL</u>	<u>N65042</u>	<u>001</u>	Oct 17, 2001
<u>AP</u>		MAYNE PHARMA USA	<u>EQ 15 UNITS BASE/VIAL</u>	<u>N65031</u>	<u>001</u>	Mar 10, 2000
<u>AP</u>			<u>EQ 30 UNITS BASE/VIAL</u>	<u>N65031</u>	<u>002</u>	Mar 10, 2000
<u>AP</u>		SICOR PHARMS	<u>EQ 15 UNITS BASE/VIAL</u>	<u>N65033</u>	<u>001</u>	Jun 27, 2000
<u>AP</u>			<u>EQ 30 UNITS BASE/VIAL</u>	<u>N65033</u>	<u>002</u>	Jun 27, 2000
		<u>BLEOMYCIN SULFATE</u>				
<u>AP</u>		SICOR PHARMS	<u>EQ 15 UNITS BASE/VIAL</u>	<u>N64084</u>	<u>001</u>	Jun 01, 1996
<u>AP</u>			<u>EQ 30 UNITS BASE/VIAL</u>	<u>N64084</u>	<u>002</u>	Jun 01, 1996

BORTEZOMIBINJECTABLE; INTRAVENOUS
VELCADE

+ MILLENNIUM PHARMS 3.5MG/VIAL N21602 001 May 13, 2003

BOSENTANTABLET; ORAL
TRACLEERACTELION 62.5MG N21290 001 Nov 20, 2001
+ 125MG N21290 002 Nov 20, 2001BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE

<u>AP</u>	+	INTL MEDICATION	<u>50MG/ML</u>	<u>N70119</u>	<u>001</u>	Apr 29, 1986
<u>AP</u>		LUITPOLD	<u>50MG/ML</u>	<u>N70891</u>	<u>001</u>	Jul 26, 1988

PRESCRIPTION DRUG PRODUCT LIST

3 - 51 (of 360)

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>200MG/100ML</u>	<u>N19121</u>	<u>002</u>	Apr 29, 1986
<u>AP</u>	+		<u>400MG/100ML</u>	<u>N19121</u>	<u>003</u>	Apr 29, 1986
	+		100MG/100ML	N19121	001	Apr 29, 1986
<u>AP</u>	+	HOSPIRA	<u>200MG/100ML</u>	<u>N19008</u>	<u>002</u>	Apr 29, 1986
<u>AP</u>	+		<u>400MG/100ML</u>	<u>N19008</u>	<u>003</u>	Apr 29, 1986
<u>BRETYLIUM TOSYLATE IN PLASTIC CONTAINER</u>						
<u>AP</u>	+	HOSPIRA	<u>50MG/ML</u>	<u>N19030</u>	<u>001</u>	Apr 29, 1986

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN P

+	ALLERGAN	0.1%	N21770	001	Aug 19, 2005
+		0.15%	N21262	001	Mar 16, 2001

BRIMONIDINE TARTRATE

<u>AT</u>		ALCON	<u>0.2%</u>	<u>N76254</u>	<u>001</u>	Sep 16, 2003
<u>AT</u>	+	BAUSCH AND LOMB	<u>0.2%</u>	<u>N76260</u>	<u>001</u>	May 28, 2003
<u>AT</u>		IVAX PHARMS	<u>0.2%</u>	<u>N76372</u>	<u>001</u>	Sep 10, 2004

BRINZOLAMIDE

SUSPENSION/DROPS; OPHTHALMIC

AZOPT

+	ALCON	1%	N20816	001	Apr 01, 1998
---	-------	----	--------	-----	--------------

BROMFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC

XIBROM

+	ISTA PHARMS	0.09%	N21664	001	Mar 24, 2005
---	-------------	-------	--------	-----	--------------

BROMOCRIPTINE MESYLATE

CAPSULE; ORAL

BROMOCRIPTINE MESYLATE

<u>AB</u>		MYLAN	<u>EQ 5MG BASE</u>	<u>N77226</u>	<u>001</u>	Apr 04, 2005
		<u>PARLODEL</u>				
<u>AB</u>	+	NOVARTIS	<u>EQ 5MG BASE</u>	<u>N17962</u>	<u>002</u>	Mar 01, 1982

TABLET; ORAL

BROMOCRIPTINE MESYLATE

<u>AB</u>		LEK PHARMS	<u>EQ 2.5MG BASE</u>	<u>N74631</u>	<u>001</u>	Jan 13, 1998
<u>AB</u>		MYLAN	<u>EQ 2.5MG BASE</u>	<u>N76962</u>	<u>001</u>	Sep 24, 2004
		<u>PARLODEL</u>				
<u>AB</u>	+	NOVARTIS	<u>EQ 2.5MG BASE</u>	<u>N17962</u>	<u>001</u>	

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP; ORAL

AMBENYL

<u>AA</u>	+	FOREST LABS	<u>12.5MG/5ML;10MG/5ML</u>	<u>N09319</u>	<u>006</u>	Jan 10, 1984
		<u>MYBANIL</u>				
<u>AA</u>		MORTON GROVE	<u>12.5MG/5ML;10MG/5ML</u>	<u>N88626</u>	<u>001</u>	Oct 12, 1984

BROMPHENIRAMINE MALEATE

ELIXIR; ORAL

BROMPHENIRAMINE MALEATE

+	KV PHARM	2MG/5ML	N85466	001	
---	----------	---------	--------	-----	--

INJECTABLE; INJECTION

BROMPHENIRAMINE MALEATE

+	STERIS	10MG/ML	N83821	001	
---	--------	---------	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 52 (of 360)

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL

BROMFED-DM

<u>AA</u>	BRIGHTON PHARMS INC	<u>2MG/5ML;10MG/5ML;30MG/5ML</u>	<u>N89681</u>	<u>001</u>	Dec 22, 1988
-----------	---------------------	----------------------------------	---------------	------------	--------------

MYPHETANE DX

<u>AA</u>	+ MORTON GROVE	<u>2MG/5ML;10MG/5ML;30MG/5ML</u>	<u>N88811</u>	<u>001</u>	Jun 07, 1985
-----------	----------------	----------------------------------	---------------	------------	--------------

BUDESONIDE

CAPSULE; ORAL

ENTOCORT EC

+	ASTRAZENECA	3MG	N21324	001	Oct 02, 2001
---	-------------	-----	--------	-----	--------------

POWDER, METERED; INHALATION

PULMICORT

+	ASTRAZENECA	0.16MG/INH	N20441	002	Jun 24, 1997
---	-------------	------------	--------	-----	--------------

SPRAY, METERED; NASAL

RHINOCORT

	ASTRAZENECA	0.032MG/INH	N20746	001	Oct 01, 1999
--	-------------	-------------	--------	-----	--------------

+		0.064MG/INH	N20746	002	Oct 01, 1999
---	--	-------------	--------	-----	--------------

SUSPENSION; INHALATION

PULMICORT RESPULES

	ASTRAZENECA	0.25MG/2ML	N20929	001	Aug 08, 2000
--	-------------	------------	--------	-----	--------------

+		0.5MG/2ML	N20929	002	Aug 08, 2000
---	--	-----------	--------	-----	--------------

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

<u>AP</u>	BEDFORD	<u>0.25MG/ML</u>	<u>N74441</u>	<u>001</u>	Jan 27, 1995
-----------	---------	------------------	---------------	------------	--------------

<u>AP</u>	HOSPIRA	<u>0.25MG/ML</u>	<u>N74160</u>	<u>001</u>	Oct 30, 1997
-----------	---------	------------------	---------------	------------	--------------

<u>AP</u>		<u>0.25MG/ML</u>	<u>N74332</u>	<u>001</u>	Oct 31, 1994
-----------	--	------------------	---------------	------------	--------------

<u>AP</u>	SICOR PHARMS	<u>0.25MG/ML</u>	<u>N74613</u>	<u>001</u>	Nov 18, 1997
-----------	--------------	------------------	---------------	------------	--------------

BUMEX

<u>AP</u>	+ ROCHE	<u>0.25MG/ML</u>	<u>N18226</u>	<u>001</u>	Feb 28, 1983
-----------	---------	------------------	---------------	------------	--------------

TABLET; ORAL

BUMETANIDE

<u>AB</u>	IVAX PHARMS	<u>0.5MG</u>	<u>N74225</u>	<u>001</u>	Apr 24, 1995
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>1MG</u>	<u>N74225</u>	<u>002</u>	Apr 24, 1995
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>2MG</u>	<u>N74225</u>	<u>003</u>	Apr 24, 1995
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>N74700</u>	<u>001</u>	Nov 21, 1996
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>1MG</u>	<u>N74700</u>	<u>002</u>	Nov 21, 1996
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>2MG</u>	<u>N74700</u>	<u>003</u>	Nov 21, 1996
-----------	--	------------	---------------	------------	--------------

BUMEX

<u>AB</u>	ROCHE	<u>0.5MG</u>	<u>N18225</u>	<u>002</u>	Feb 28, 1983
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>		<u>1MG</u>	<u>N18225</u>	<u>001</u>	Feb 28, 1983
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	+	<u>2MG</u>	<u>N18225</u>	<u>003</u>	Jun 14, 1985
-----------	---	------------	---------------	------------	--------------

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>0.25%</u>	<u>N18053</u>	<u>002</u>	
-----------	---------	--------------	---------------	------------	--

<u>AP</u>		<u>0.25%</u>	<u>N70583</u>	<u>001</u>	Feb 17, 1987
-----------	--	--------------	---------------	------------	--------------

<u>AP</u>		<u>0.25%</u>	<u>N70586</u>	<u>001</u>	Mar 03, 1987
-----------	--	--------------	---------------	------------	--------------

<u>AP</u>		<u>0.25%</u>	<u>N70590</u>	<u>001</u>	Feb 17, 1987
-----------	--	--------------	---------------	------------	--------------

<u>AP</u>		<u>0.5%</u>	<u>N18053</u>	<u>001</u>	
-----------	--	-------------	---------------	------------	--

<u>AP</u>		<u>0.5%</u>	<u>N70584</u>	<u>001</u>	Feb 17, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AP</u>		<u>0.5%</u>	<u>N70597</u>	<u>001</u>	Mar 03, 1987
-----------	--	-------------	---------------	------------	--------------

<u>AP</u>		<u>0.5%</u>	<u>N70609</u>	<u>001</u>	Mar 03, 1987
-----------	--	-------------	---------------	------------	--------------

<u>AP</u>		<u>0.75%</u>	<u>N18053</u>	<u>003</u>	
-----------	--	--------------	---------------	------------	--

<u>AP</u>		<u>0.75%</u>	<u>N70585</u>	<u>001</u>	Mar 03, 1987
-----------	--	--------------	---------------	------------	--------------

<u>AP</u>		<u>0.75%</u>	<u>N70587</u>	<u>001</u>	Mar 03, 1987
-----------	--	--------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 53 (of 360)

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	INTL MEDICATED	<u>0.25%</u>	<u>N76012</u>	<u>001</u>	Jan 09, 2002
<u>AP</u>		<u>0.5%</u>	<u>N76012</u>	<u>002</u>	Jan 09, 2002
<u>AP</u>		<u>0.75%</u>	<u>N76012</u>	<u>003</u>	Jan 09, 2002

MARCAINE HYDROCHLORIDE

<u>AP</u>	+ HOSPIRA	<u>0.25%</u>	<u>N16964</u>	<u>001</u>	
<u>AP</u>	+	<u>0.5%</u>	<u>N16964</u>	<u>006</u>	

MARCAINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>	+ HOSPIRA	<u>0.25%</u>	<u>N16964</u>	<u>012</u>	
<u>AP</u>	+	<u>0.5%</u>	<u>N16964</u>	<u>005</u>	
<u>AP</u>	+	<u>0.75%</u>	<u>N16964</u>	<u>009</u>	

SENSORCAINE

<u>AP</u>	ASTRAZENECA	<u>0.25%</u>	<u>N18304</u>	<u>001</u>	
<u>AP</u>		<u>0.25%</u>	<u>N70552</u>	<u>001</u>	May 21, 1986
<u>AP</u>		<u>0.5%</u>	<u>N18304</u>	<u>002</u>	
<u>AP</u>		<u>0.5%</u>	<u>N70553</u>	<u>001</u>	May 21, 1986
<u>AP</u>		<u>0.75%</u>	<u>N18304</u>	<u>003</u>	
<u>AP</u>		<u>0.75%</u>	<u>N70554</u>	<u>001</u>	May 21, 1986

INJECTABLE; SPINAL

BUPIVACAINE

<u>AP</u>	HOSPIRA	<u>0.75%</u>	<u>N71810</u>	<u>001</u>	Dec 11, 1987
-----------	---------	--------------	---------------	------------	--------------

MARCAINE

<u>AP</u>	+ HOSPIRA	<u>0.75%</u>	<u>N18692</u>	<u>001</u>	May 04, 1984
-----------	-----------	--------------	---------------	------------	--------------

SENSORCAINE

<u>AP</u>	ASTRAZENECA	<u>0.75%</u>	<u>N71202</u>	<u>001</u>	Apr 15, 1987
-----------	-------------	--------------	---------------	------------	--------------

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

+	HOSPIRA	0.25%;0.005MG/ML	N71165	001	Jun 16, 1988
		0.25%;0.005MG/ML	N71166	001	Jun 16, 1988
		0.25%;0.005MG/ML	N71167	001	Jun 16, 1988
+		0.5%;0.005MG/ML	N71168	001	Jun 16, 1988
		0.5%;0.005MG/ML	N71169	001	Jun 16, 1988
		0.5%;0.005MG/ML	N71170	001	Jun 16, 1988
+		0.75%;0.005MG/ML	N71171	001	Jun 16, 1988

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE

<u>AP</u>	+ HOSPIRA	<u>0.25%;0.0091MG/ML</u>	<u>N16964</u>	<u>004</u>	
<u>AP</u>	+	<u>0.5%;0.0091MG/ML</u>	<u>N16964</u>	<u>008</u>	

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE

<u>AP</u>	+ HOSPIRA	<u>0.25%;0.0091MG/ML</u>	<u>N16964</u>	<u>013</u>	
<u>AP</u>	+	<u>0.5%;0.0091MG/ML</u>	<u>N16964</u>	<u>007</u>	
<u>AP</u>	+	<u>0.75%;0.0091MG/ML</u>	<u>N16964</u>	<u>010</u>	

SENSORCAINE

<u>AP</u>	ASTRAZENECA	<u>0.25%;0.0091MG/ML</u>	<u>N70966</u>	<u>001</u>	Oct 13, 1987
<u>AP</u>		<u>0.25%;0.0091MG/ML</u>	<u>N70967</u>	<u>001</u>	Oct 13, 1987
<u>AP</u>		<u>0.5%;0.0091MG/ML</u>	<u>N18304</u>	<u>004</u>	Sep 02, 1983
<u>AP</u>		<u>0.5%;0.0091MG/ML</u>	<u>N70968</u>	<u>001</u>	Oct 13, 1987
<u>AP</u>		<u>0.75%;0.0091MG/ML</u>	<u>N18304</u>	<u>005</u>	Sep 02, 1983

BUPIVACAINE HYDROCHLORIDE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DUOCAINE

+	AMPHASTAR	EQ 0.375% (37.5MG/10ML);EQ 1% (100MG/10ML)	N21496	001	May 23, 2003
---	-----------	--	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 54 (of 360)

BUPRENORPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPRENEX

<u>AP</u>	+	RECKITT BENCKISER	<u>EQ 0.3MG BASE/ML</u>	<u>N18401</u>	<u>001</u>	
-----------	---	-------------------	-------------------------	---------------	------------	--

BUPRENORPHINE HYDROCHLORIDE

<u>AP</u>		BEDFORD	<u>EQ 0.3MG BASE/ML</u>	<u>N76931</u>	<u>001</u>	Mar 02, 2005
<u>AP</u>		HOSPIRA	<u>EQ 0.3MG BASE/ML</u>	<u>N74137</u>	<u>001</u>	Jun 03, 1996

TABLET; SUBLINGUAL
SUBUTEX

		RECKITT BENCKISER	EQ 2MG BASE	N20732	002	Oct 08, 2002
	+		EQ 8MG BASE	N20732	003	Oct 08, 2002

BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDETABLET; SUBLINGUAL
SUBOXONE

		RECKITT BENCKISER	2MG;0.5MG	N20733	001	Oct 08, 2002
	+		8MG;2MG	N20733	002	Oct 08, 2002

BUPROPION HYDROCHLORIDE

TABLET; ORAL

BUPROPION HYDROCHLORIDE

<u>AB</u>		MYLAN	<u>75MG</u>	<u>N75491</u>	<u>001</u>	Apr 17, 2000
<u>AB</u>			<u>100MG</u>	<u>N75491</u>	<u>002</u>	Apr 17, 2000
<u>AB</u>		SANDOZ	<u>75MG</u>	<u>N75584</u>	<u>001</u>	Feb 07, 2000
<u>AB</u>			<u>75MG</u>	<u>N75613</u>	<u>002</u>	Oct 10, 2000
<u>AB</u>			<u>100MG</u>	<u>N75584</u>	<u>002</u>	Feb 07, 2000
<u>AB</u>			<u>100MG</u>	<u>N75613</u>	<u>001</u>	Oct 10, 2000
<u>AB</u>		TEVA	<u>75MG</u>	<u>N75310</u>	<u>001</u>	Nov 29, 1999
<u>AB</u>			<u>100MG</u>	<u>N75310</u>	<u>002</u>	Nov 29, 1999

WELLBUTRIN

<u>AB</u>		GLAXOSMITHKLINE	<u>75MG</u>	<u>N18644</u>	<u>002</u>	Dec 30, 1985
<u>AB</u>	+		<u>100MG</u>	<u>N18644</u>	<u>003</u>	Dec 30, 1985

TABLET, EXTENDED RELEASE; ORAL

BUPROPION HYDROCHLORIDE

<u>AB1</u>		IMPAX LABS	<u>100MG</u>	<u>N75913</u>	<u>001</u>	Jan 28, 2004
<u>AB1</u>			<u>150MG</u>	<u>N75913</u>	<u>002</u>	Mar 22, 2004
<u>AB1</u>			<u>200MG</u>	<u>N76711</u>	<u>001</u>	Dec 03, 2004
<u>AB2</u>			<u>150MG</u>	<u>N75914</u>	<u>001</u>	May 27, 2004
<u>AB1</u>		SANDOZ	<u>100MG</u>	<u>N75932</u>	<u>001</u>	Nov 25, 2003
<u>AB1</u>			<u>100MG</u>	<u>N76845</u>	<u>001</u>	Jul 14, 2005
<u>AB1</u>			<u>150MG</u>	<u>N75932</u>	<u>002</u>	Mar 22, 2004
<u>AB1</u>			<u>150MG</u>	<u>N76845</u>	<u>002</u>	Jul 14, 2005
<u>AB1</u>			<u>200MG</u>	<u>N75932</u>	<u>003</u>	Jun 22, 2005
<u>AB2</u>			<u>150MG</u>	<u>N76834</u>	<u>001</u>	Jul 14, 2005

WELLBUTRIN SR

<u>AB1</u>		GLAXOSMITHKLINE	<u>100MG</u>	<u>N20358</u>	<u>002</u>	Oct 04, 1996
<u>AB1</u>	+		<u>150MG</u>	<u>N20358</u>	<u>003</u>	Oct 04, 1996
<u>AB1</u>			<u>200MG</u>	<u>N20358</u>	<u>004</u>	Jun 14, 2002
			50MG	N20358	001	Oct 04, 1996

WELLBUTRIN XL

	+	SMITHKLINE BEECHAM	150MG	N21515	001	Aug 28, 2003
			300MG	N21515	002	Aug 28, 2003

ZYBAN

<u>AB2</u>	+	GLAXOSMITHKLINE	<u>150MG</u>	<u>N20711</u>	<u>003</u>	May 14, 1997
------------	---	-----------------	--------------	---------------	------------	--------------

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPAR

<u>AB</u>		BRISTOL MYERS SQUIBB	<u>5MG</u>	<u>N18731</u>	<u>001</u>	Sep 29, 1986
-----------	--	----------------------	------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 55 (of 360)

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPAR

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>10MG</u>	<u>N18731</u>	<u>002</u>	Sep 29, 1986
<u>AB</u>		<u>15MG</u>	<u>N18731</u>	<u>003</u>	Apr 22, 1996
<u>AB</u>	+	<u>30MG</u>	<u>N18731</u>	<u>004</u>	Apr 22, 1996
	<u>BUSPIRONE HYDROCHLORIDE</u>				
<u>AB</u>	EGIS	<u>5MG</u>	<u>N75119</u>	<u>001</u>	Mar 14, 2002
<u>AB</u>		<u>10MG</u>	<u>N75119</u>	<u>002</u>	Mar 14, 2002
<u>AB</u>		<u>15MG</u>	<u>N75119</u>	<u>003</u>	Jan 23, 2003
<u>AB</u>	KV PHARM	<u>5MG</u>	<u>N75572</u>	<u>001</u>	Feb 27, 2002
<u>AB</u>		<u>10MG</u>	<u>N75572</u>	<u>002</u>	Feb 27, 2002
<u>AB</u>		<u>15MG</u>	<u>N75572</u>	<u>003</u>	Feb 27, 2002
<u>AB</u>	MYLAN	<u>5MG</u>	<u>N75272</u>	<u>001</u>	Mar 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75272</u>	<u>002</u>	Mar 01, 2002
<u>AB</u>		<u>15MG</u>	<u>N75272</u>	<u>003</u>	Mar 28, 2001
<u>AB</u>		<u>30MG</u>	<u>N76008</u>	<u>001</u>	Jun 28, 2001
<u>AB</u>	PAR PHARM	<u>5MG</u>	<u>N75467</u>	<u>001</u>	Feb 28, 2002
<u>AB</u>		<u>7.5MG</u>	<u>N75467</u>	<u>002</u>	Mar 28, 2001
<u>AB</u>		<u>10MG</u>	<u>N75467</u>	<u>003</u>	Feb 28, 2002
<u>AB</u>		<u>15MG</u>	<u>N75467</u>	<u>004</u>	Feb 28, 2002
<u>AB</u>	PHARMEX PRODS	<u>5MG</u>	<u>N75388</u>	<u>001</u>	May 09, 2002
<u>AB</u>		<u>10MG</u>	<u>N75388</u>	<u>002</u>	May 09, 2002
<u>AB</u>		<u>15MG</u>	<u>N75388</u>	<u>003</u>	May 09, 2002
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>N75413</u>	<u>001</u>	Mar 19, 2002
<u>AB</u>		<u>10MG</u>	<u>N75413</u>	<u>002</u>	Mar 19, 2002
<u>AB</u>		<u>15MG</u>	<u>N75413</u>	<u>003</u>	Mar 19, 2002
<u>AB</u>	TEVA	<u>5MG</u>	<u>N75022</u>	<u>001</u>	Feb 28, 2002
<u>AB</u>		<u>10MG</u>	<u>N75022</u>	<u>002</u>	Feb 28, 2002
<u>AB</u>		<u>15MG</u>	<u>N75022</u>	<u>003</u>	Feb 28, 2002
<u>AB</u>		<u>30MG</u>	<u>N75022</u>	<u>004</u>	Mar 25, 2004
<u>AB</u>	TORPHARM	<u>5MG</u>	<u>N75521</u>	<u>001</u>	Apr 05, 2002
<u>AB</u>		<u>10MG</u>	<u>N75521</u>	<u>002</u>	Apr 05, 2002
<u>AB</u>		<u>15MG</u>	<u>N75521</u>	<u>003</u>	Apr 05, 2002
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>N74253</u>	<u>001</u>	Mar 28, 2001
<u>AB</u>		<u>10MG</u>	<u>N74253</u>	<u>002</u>	Mar 28, 2001
<u>AB</u>		<u>15MG</u>	<u>N74253</u>	<u>003</u>	Mar 13, 2002
<u>AB</u>	ZENITH GOLDLINE	<u>5MG</u>	<u>N75385</u>	<u>001</u>	Mar 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75385</u>	<u>002</u>	Mar 01, 2002
<u>AB</u>		<u>15MG</u>	<u>N75385</u>	<u>003</u>	Mar 01, 2002

BUSULFAN

INJECTABLE; INJECTION

BUSULFEX

+ ESP PHARMA 6MG/ML N20954 001 Feb 04, 1999

TABLET; ORAL

MYLERAN

+ GLAXOSMITHKLINE 2MG N09386 001

BUTABARBITAL SODIUM

ELIXIR; ORAL

BUTISOL SODIUM

+ MEDPOINTE PHARM HLC 30MG/5ML N85380 001

TABLET; ORAL

BUTABARBITAL

AA BUNDY 30MG N85550 001

BUTISOL SODIUM

AA + MEDPOINTE PHARM HLC 30MG N00793 004
+ 50MG N00793 003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 56 (of 360)

BUTABARBITAL SODIUM

TABLET; ORAL

SODIUM BUTABARBITAL

MARSHALL PHARMA 16.2MG
32.4MGN83524 001
N83858 001BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL

MENTAX

+ MYLAN BERTEK 1%

N20524 001 Oct 18, 1996

MENTAX-TC

+ MYLAN BERTEK 1%

N21408 001 Oct 17, 2002

BUTOCONAZOLE NITRATE

CREAM; VAGINAL

GYNAZOLE-1

+ KV PHARM 2%

N19881 001 Feb 07, 1997

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATEAP APOTEX 2MG/MLN75697 001 Oct 23, 2001AP BEDFORD 2MG/MLN75046 001 Aug 12, 1998AP HOSPIRA 1MG/MLN75559 001 Mar 20, 2000AP 2MG/MLN75559 002 Mar 20, 2000AP MAYNE PHARMA USA 1MG/MLN75342 001 Nov 04, 1999AP 2MG/MLN75342 002 Nov 04, 1999BUTORPHANOL TARTRATE PRESERVATIVE FREEAP APOTEX 1MG/MLN75695 001 Oct 23, 2001AP 2MG/MLN75695 002 Oct 23, 2001AP BEDFORD 1MG/MLN75045 001 Aug 12, 1998AP 2MG/MLN75045 002 Aug 12, 1998AP HOSPIRA 1MG/MLN74620 001 Jan 22, 1997AP 1MG/MLN74626 001 Jan 23, 1997AP 2MG/MLN74620 002 Jan 22, 1997AP 2MG/MLN74626 002 Jan 23, 1997AP MAYNE PHARMA USA 1MG/MLN75170 001 Sep 28, 1998AP 2MG/MLN75170 002 Sep 28, 1998STADOLAP + APOTHECON 2MG/MLN17857 004STADOL PRESERVATIVE FREEAP + APOTHECON 1MG/MLN17857 001AP + 2MG/MLN17857 002

SPRAY, METERED; NASAL

BUTORPHANOL TARTRATEAB + MYLAN 1MG/SPRAYN75759 001 Aug 08, 2001AB NOVEX 1MG/SPRAYN75499 001 Dec 04, 2002AB ROXANE 1MG/SPRAYN75824 001 Mar 12, 2002CABERGOLINE

TABLET; ORAL

CABERGOLINEAB PAR PHARM 0.5MGN76310 001 Dec 29, 2005DOSTINEXAB + PHARMACIA AND UPJOHN 0.5MGN20664 001 Dec 23, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 57 (of 360)

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS, ORAL
CAFCIT

+ MEAD JOHNSON	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N20793 001	Sep 21, 1999
----------------	------------------------------------	------------	--------------

SOLUTION; ORAL
CAFCIT

+ MEAD JOHNSON	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N20793 002	Apr 12, 2000
----------------	------------------------------------	------------	--------------

CAFFEINE; ERGOTAMINE TARTRATE

SUPPOSITORY; RECTAL
CAFERGOT

BR + NOVARTIS	100MG;2MG	N09000 002	
---------------	-----------	------------	--

MIGERGOT

BR G AND W LABS	100MG;2MG	N86557 001	Oct 04, 1983
-----------------	-----------	------------	--------------

TABLET; ORAL

CAFERGOT

<u>AA</u> + SANDOZ	<u>100MG;1MG</u>	<u>N84294</u> <u>001</u>	
--------------------	------------------	--------------------------	--

ERGOTAMINE TARTRATE AND CAFFEINE

<u>AA</u> MIKART	<u>100MG;1MG</u>	<u>N40590</u> <u>001</u>	Sep 16, 2005
------------------	------------------	--------------------------	--------------

<u>AA</u> WEST WARD	<u>100MG;1MG</u>	<u>N40510</u> <u>001</u>	Sep 17, 2004
---------------------	------------------	--------------------------	--------------

CALCIPOTRIENE

CREAM; TOPICAL
DOVONEX

+ BRISTOL MYERS SQUIBB	0.005%	N20554 001	Jul 22, 1996
------------------------	--------	------------	--------------

OINTMENT; TOPICAL
DOVONEX

+ BRISTOL MYERS SQUIBB	0.005%	N20273 001	Dec 29, 1993
------------------------	--------	------------	--------------

SOLUTION; TOPICAL
DOVONEX

+ BRISTOL MYERS SQUIBB	0.005%	N20611 001	Mar 03, 1997
------------------------	--------	------------	--------------

CALCITONIN SALMON RECOMBINANT

SPRAY, METERED; NASAL
FORTICAL

+ UNIGENE	200 IU/SPRAY	N21406 001	Aug 12, 2005
-----------	--------------	------------	--------------

CALCITONIN, SALMON

INJECTABLE; INJECTION
MIACALCIN

<u>AP</u> + NOVARTIS	<u>200 IU/ML</u>	<u>N17808</u> <u>002</u>	Mar 29, 1991
----------------------	------------------	--------------------------	--------------

SPRAY, METERED; NASAL
MIACALCIN

+ NOVARTIS	200 IU/SPRAY	N20313 002	Aug 17, 1995
------------	--------------	------------	--------------

CALCITRIOL

CAPSULE; ORAL
CALCITRIOL

<u>AB</u> TEVA	<u>0.25UGM</u>	<u>N75765</u> <u>001</u>	Oct 12, 2001
----------------	----------------	--------------------------	--------------

<u>AB</u>	<u>0.5UGM</u>	<u>N75765</u> <u>002</u>	Oct 12, 2001
-----------	---------------	--------------------------	--------------

ROCALTROL

<u>AB</u> ROCHE	<u>0.25UGM</u>	<u>N18044</u> <u>001</u>	
-----------------	----------------	--------------------------	--

<u>AB</u> +	<u>0.5UGM</u>	<u>N18044</u> <u>002</u>	
-------------	---------------	--------------------------	--

INJECTABLE; INJECTION
CALCIJEX

<u>AP</u> + ABBOTT	<u>0.001MG/ML</u>	<u>N18874</u> <u>001</u>	Sep 25, 1986
--------------------	-------------------	--------------------------	--------------

<u>AP</u> +	<u>0.002MG/ML</u>	<u>N18874</u> <u>002</u>	Sep 25, 1986
-------------	-------------------	--------------------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 58 (of 360)

CALCITRIOL

INJECTABLE; INJECTION

CALCITRIOL

<u>AP</u>	AM PHARM PARTNERS	<u>0.001MG/ML</u>	<u>N75836</u>	<u>001</u>	Dec 31, 2002
<u>AP</u>		<u>0.002MG/ML</u>	<u>N75836</u>	<u>002</u>	Dec 31, 2002
<u>AP</u>	LUITPOLD	<u>0.001MG/ML</u>	<u>N75746</u>	<u>001</u>	Sep 26, 2003
<u>AP</u>		<u>0.002MG/ML</u>	<u>N75746</u>	<u>002</u>	Sep 26, 2003
<u>AP</u>	LYNE	<u>0.001MG/ML</u>	<u>N76206</u>	<u>001</u>	Sep 17, 2003
<u>AP</u>	SICOR PHARMS	<u>0.001MG/ML</u>	<u>N75823</u>	<u>001</u>	Mar 31, 2003
<u>AP</u>		<u>0.002MG/ML</u>	<u>N75823</u>	<u>002</u>	Mar 31, 2003
<u>AP</u>	XANODYNE PHARM	<u>0.001MG/ML</u>	<u>N75766</u>	<u>001</u>	Feb 20, 2003
<u>AP</u>		<u>0.002MG/ML</u>	<u>N75766</u>	<u>002</u>	Feb 20, 2003

SOLUTION; ORAL

CALCITRIOL

<u>AA</u>	ROXANE	<u>1UGM/ML</u>	<u>N76242</u>	<u>001</u>	Jul 18, 2003
<u>ROCALTROL</u>					
<u>AA</u>	+ ROCHE	<u>1UGM/ML</u>	<u>N21068</u>	<u>001</u>	Nov 20, 1998

CALCIUM ACETATE

CAPSULE; ORAL

PHOSLO GELCAPS

	+ NABI	EQ 169MG CALCIUM	N21160	003	Apr 02, 2001
--	--------	------------------	--------	-----	--------------

TABLET; ORAL

PHOSLO

	+ NABI	EQ 169MG CALCIUM	N19976	001	Dec 10, 1990
--	--------	------------------	--------	-----	--------------

CALCIUM CARBONATE; RISEDRONATE SODIUM

TABLET, TABLET; ORAL

ACTONEL WITH CALCIUM (COPACKAGED)

	+ PROCTER AND GAMBLE	EQ 500MG BASE, N/A; N/A, 35MG	N21823	001	Aug 12, 2005
--	----------------------	-------------------------------	--------	-----	--------------

CALCIUM CHLORIDE

INJECTABLE; INJECTION

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER

	+ HOSPIRA	100MG/ML	N21117	001	Jan 28, 2000
--	-----------	----------	--------	-----	--------------

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

BSS PLUS

<u>AT</u>	+ ALCON	<u>0.154MG/ML;0.92MG/ML;0.184MG/ML;0.2MG/ML;0.38MG/ML;2.1MG/ML;7.14MG/ML;0.42MG/ML</u>	<u>N18469</u>	<u>001</u>	
-----------	---------	--	---------------	------------	--

ENDOSOL EXTRA

<u>AT</u>	+ AKORN	<u>0.154MG/ML;0.92MG/ML;0.184MG/ML;0.2MG/ML;0.38MG/ML;2.1MG/ML;7.14MG/ML;0.42MG/ML</u>	<u>N20079</u>	<u>001</u>	Nov 27, 1991
-----------	---------	--	---------------	------------	--------------

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER

	+ B BRAUN	37MG/100ML;5GM/100ML;31MG/100ML;120MG/100ML;330MG/100ML;88MG/100ML	N19864	001	Jun 10, 1993
--	-----------	--	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 59 (of 360)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER

+ B BRAUN	35MG/100ML; 5GM/100ML; 30MG/100ML; 74MG/100ML; 640MG/100ML; 500MG/100ML; 74MG/100ML	N19867 001	Dec 20, 1993
-----------	---	------------	--------------

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER

+ BAXTER HLTHCARE	37MG/100ML; 5GM/100ML; 30MG/100ML; 119MG/100ML; 161MG/100ML; 94MG/100ML; 138MG/100ML	N17390 001	
-------------------	--	------------	--

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION; INTRAPERITONEAL

DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER

+ B BRAUN	510MG/100ML; 30GM/100ML; 200MG/100ML; 9.2GM/100ML; 9.6GM/100ML	N18807 001	Aug 26, 1983
-----------	--	------------	--------------

+	510MG/100ML; 30GM/100ML; 200MG/100ML; 9.4GM/100ML; 11GM/100ML	N18807 003	Aug 26, 1983
---	---	------------	--------------

DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER

+ B BRAUN	510MG/100ML; 50GM/100ML; 200MG/100ML; 9.2GM/100ML; 9.6GM/100ML	N18807 002	Aug 26, 1983
-----------	--	------------	--------------

+	510MG/100ML; 50GM/100ML; 200MG/100ML; 9.4GM/100ML; 11GM/100ML	N18807 004	Aug 26, 1983
---	---	------------	--------------

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DELFLX W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5.67MG/100ML; 392MG/100ML	N18379 002	
----	-----------------	--	------------	--

AT		25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 5.67MG/100ML; 392MG/100ML	N18883 001	Nov 30, 1984
----	--	--	------------	--------------

DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5.38MG/100ML; 448MG/100ML	N18883 004	Nov 30, 1984
----	-----------------	--	------------	--------------

DELFLX W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	18.4MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 5.38MG/100ML; 448MG/100ML	N20171 001	Aug 19, 1992
----	-----------------	--	------------	--------------

DELFLX W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 5.67MG/100ML; 392MG/100ML	N18379 003	
----	-----------------	--	------------	--

AT		25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 5.67MG/100ML; 392MG/100ML	N18883 002	Nov 30, 1984
----	--	--	------------	--------------

DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5.38MG/100ML; 448MG/100ML	N18883 005	Nov 30, 1984
----	-----------------	--	------------	--------------

DELFLX W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	18.4MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 5.38MG/100ML; 448MG/100ML	N20171 002	Aug 19, 1992
----	-----------------	--	------------	--------------

DELFLX W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 3.5GM/100ML; 15.2MG/100ML; 5.67MG/100ML; 392MG/100ML	N18379 007	Jun 24, 1988
----	-----------------	--	------------	--------------

DELFLX W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 5.67MG/100ML; 392MG/100ML	N18379 001	
----	-----------------	---	------------	--

AT		25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 5.67MG/100ML; 392MG/100ML	N18883 003	Nov 30, 1984
----	--	---	------------	--------------

DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER

AT	FRESENIUS MEDCL	25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 5.38MG/100ML; 448MG/100ML	N18883 006	Nov 30, 1984
----	-----------------	---	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 60 (of 360)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL						
<u>DELFLX W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER</u>						
<u>AT</u>	FRESENIUS MEDCL	<u>18.4MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N20171</u>	<u>003</u>		Aug 19, 1992
<u>DELFLX-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	FRESENIUS MEDCL	<u>25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N18379</u>	<u>004</u>		Jul 07, 1982
<u>DELFLX-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	FRESENIUS MEDCL	<u>25.7MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N18379</u>	<u>005</u>		Jul 07, 1982
<u>DELFLX-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	FRESENIUS MEDCL	<u>25.7MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N18379</u>	<u>008</u>		Jun 24, 1988
<u>DELFLX-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>						
<u>AT</u>	FRESENIUS MEDCL	<u>25.7MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N18379</u>	<u>006</u>		Jul 07, 1982
DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER						
	B BRAUN	26MG/100ML; 1.5GM/100ML; 5MG/100ML; 530MG/100ML; 450MG/100ML	N18460	007		Jan 29, 1986
DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER						
	B BRAUN	26MG/100ML; 2.5GM/100ML; 5MG/100ML; 530MG/100ML; 450MG/100ML	N18460	005		Nov 02, 1983
DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER						
	B BRAUN	26MG/100ML; 4.25GM/100ML; 5MG/100ML; 530MG/100ML; 450MG/100ML	N18460	009		Jan 29, 1986
<u>DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML</u>	<u>N17512</u>	<u>001</u>		
<u>DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML</u>	<u>N17512</u>	<u>003</u>		
<u>DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML</u>	<u>N17512</u>	<u>002</u>		
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>18.3MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N20183</u>	<u>001</u>		Dec 04, 1992
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>18.3MG/100ML; 2.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N20183</u>	<u>002</u>		Dec 04, 1992
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>18.3MG/100ML; 3.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N20183</u>	<u>003</u>		Dec 04, 1992
<u>DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>18.3MG/100ML; 4.25GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N20183</u>	<u>004</u>		Dec 04, 1992
<u>DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML; 1.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML</u>	<u>N17512</u>	<u>007</u>		Jul 09, 1984
<u>DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML; 2.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML</u>	<u>N17512</u>	<u>008</u>		Jul 09, 1984
<u>DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML; 3.5GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML</u>	<u>N17512</u>	<u>010</u>		Nov 18, 1985
<u>DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML; 4.25GM/100ML; 15.2MG/100ML; 567MG/100ML; 392MG/100ML</u>	<u>N17512</u>	<u>009</u>		Jul 09, 1984
<u>DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>						
<u>AT</u>	BAXTER HLTHCARE	<u>18.3MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N17512</u>	<u>004</u>		
<u>AT</u>		<u>25.7MG/100ML; 1.5GM/100ML; 5.08MG/100ML; 538MG/100ML; 448MG/100ML</u>	<u>N20163</u>	<u>001</u>		Dec 04, 1992

PRESCRIPTION DRUG PRODUCT LIST

3 - 61 (of 360)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL					
<u>DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N17512</u>	<u>005</u>	
<u>AT</u>		<u>25.7MG/100ML;2.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N20163</u>	<u>002</u>	Dec 04, 1992
<u>DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML;3.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N17512</u>	<u>011</u>	Nov 18, 1985
<u>DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>					
<u>AT</u>	BAXTER HLTHCARE	<u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;</u> <u>538MG/100ML;448MG/100ML</u>	<u>N17512</u>	<u>006</u>	
<u>AT</u>		<u>25.7MG/100ML;4.25GM/100ML;5.08MG/100ML;</u> <u>538MG/100ML;448MG/100ML</u>	<u>N20163</u>	<u>003</u>	Dec 04, 1992
<u>INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	FRESENIUS	<u>18.4MG/100ML;1.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N20374</u>	<u>001</u>	Jun 13, 1994
<u>INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	FRESENIUS	<u>18.4MG/100ML;2.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N20374</u>	<u>002</u>	Jun 13, 1994
<u>INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER</u>					
<u>AT</u>	FRESENIUS	<u>18.4MG/100ML;3.5GM/100ML;5.08MG/100ML;5</u> <u>38MG/100ML;448MG/100ML</u>	<u>N20374</u>	<u>003</u>	Jun 13, 1994
<u>INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER</u>					
<u>AT</u>	FRESENIUS	<u>18.4MG/100ML;4.25GM/100ML;5.08MG/100ML;</u> <u>538MG/100ML;448MG/100ML</u>	<u>N20374</u>	<u>004</u>	Jun 13, 1994

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM SULFATE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INTRATHECAL					
ELLIOTTS B SOLUTION					
	+ QOL MEDCL	0.2MG/ML;0.8MG/ML;0.3MG/ML;0.3MG/ML;1.9 MG/ML;7.3MG/ML;0.2MG/ML	N20577	001	Sep 27, 1996

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION					
DEXTROSE 5% IN ACETATED RINGER'S IN PLASTIC CONTAINER					
	+ B BRAUN	20MG/100ML;5GM/100ML;30MG/100ML;380MG/1 00ML;600MG/100ML	N18258	001	

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION					
<u>DEXTROSE 5% AND RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>	HOSPIRA	<u>33MG/100ML;5GM/100ML;30MG/100ML;860MG/1</u> <u>00ML</u>	<u>N18254</u>	<u>001</u>	
<u>DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>	B BRAUN	<u>33MG/100ML;5GM/100ML;30MG/100ML;860MG/1</u> <u>00ML</u>	<u>N20000</u>	<u>001</u>	Apr 17, 1992
<u>AP</u>	BAXTER HLTHCARE	<u>33MG/100ML;5GM/100ML;30MG/100ML;860MG/1</u> <u>00ML</u>	<u>N16695</u>	<u>001</u>	

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION					
DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER					
	B BRAUN	10MG/100ML;2.5GM/100ML;15MG/100ML;300MG /100ML;160MG/100ML	N19634	001	Feb 24, 1988
<u>DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>	HOSPIRA	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/1</u> <u>00ML;310MG/100ML</u>	<u>N17608</u>	<u>001</u>	
<u>DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER</u>					
<u>AP</u>	B BRAUN	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/1</u> <u>00ML;310MG/100ML</u>	<u>N19634</u>	<u>003</u>	Feb 24, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 62 (of 360)

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;30MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N16679</u>	<u>001</u>	
	POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
	BAXTER HLTHCARE	20MG/100ML;5GM/100ML;105MG/100ML;600MG/100ML;310MG/100ML	N19367	002	Apr 05, 1985
		20MG/100ML;5GM/100ML;179MG/100ML;600MG/100ML;310MG/100ML	N19367	003	Apr 05, 1985
	<u>POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>				
<u>AP</u>	BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;254MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N19367</u>	<u>006</u>	Apr 05, 1985
	<u>POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>				
<u>AP</u>	BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N19367</u>	<u>004</u>	Apr 05, 1985
<u>AP</u>		<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N19367</u>	<u>005</u>	Apr 05, 1985
<u>AP</u>	HOSPIRA	<u>20MG/100ML;5GM/100ML;179MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N19685</u>	<u>002</u>	Oct 17, 1988
<u>AP</u>		<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N19685</u>	<u>008</u>	Oct 17, 1988
	<u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>				
<u>AP</u>	BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;254MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N19367</u>	<u>007</u>	Apr 05, 1985
	<u>POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER</u>				
<u>AP</u>	BAXTER HLTHCARE	<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N19367</u>	<u>008</u>	Apr 05, 1985
<u>AP</u>	HOSPIRA	<u>20MG/100ML;5GM/100ML;328MG/100ML;600MG/100ML;310MG/100ML</u>	<u>N19685</u>	<u>004</u>	Oct 17, 1988
	POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER				
	BAXTER HLTHCARE	20MG/100ML;5GM/100ML;105MG/100ML;600MG/100ML;310MG/100ML	N19367	001	Apr 05, 1985

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER

+	HOSPIRA	16.5MG/ML;25.4MG/ML;74.6MG/ML;121MG/ML;16.1MG/ML	N18895	001	Jul 20, 1984
---	---------	--	--------	-----	--------------

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN PLASTIC CONTAINER

+	B BRAUN	35MG/100ML;30MG/100ML;74MG/100ML;640MG/100ML;500MG/100ML;74MG/100ML	N19718	001	Sep 29, 1989
---	---------	---	--------	-----	--------------

SOLUTION; IRRIGATION

BSS

+	ALCON	0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9MG/ML;6.4MG/ML;1.7MG/ML	N20742	001	Dec 10, 1997
---	-------	---	--------	-----	--------------

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

PLASMA-LYTE R IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	36.8MG/100ML;30.5MG/100ML;74.6MG/100ML;640MG/100ML;496MG/100ML;89.6MG/100ML	N17438	001	
---	-----------------	---	--------	-----	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 63 (of 360)

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

SOLUTION; PERFUSION, CARDIAC

CARDIOPLEGIC IN PLASTIC CONTAINER

AT	BAXTER HLTHCARE	<u>17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML</u>	<u>N75323</u>	<u>001</u>	Apr 21, 2000
----	-----------------	--	---------------	------------	--------------

PLEGISOL IN PLASTIC CONTAINER

AT	+	HOSPIRA	<u>17.6MG/100ML; 325.3MG/100ML; 119.3MG/100ML; 643MG/100ML</u>	<u>N18608</u>	<u>001</u>	Feb 26, 1982
----	---	---------	--	---------------	------------	--------------

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

AP	B BRAUN	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N20002</u>	<u>001</u>	Apr 17, 1992
AP	BAXTER HLTHCARE	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N16693</u>	<u>001</u>	
AP	HOSPIRA	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N18251</u>	<u>001</u>	

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

AT	B BRAUN	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N18156</u>	<u>001</u>	
AT	BAXTER HLTHCARE	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N18495</u>	<u>001</u>	Feb 19, 1982
AT	HOSPIRA	<u>33MG/100ML; 30MG/100ML; 860MG/100ML</u>	<u>N17635</u>	<u>001</u>	

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

AP	B BRAUN	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N19632</u>	<u>001</u>	Feb 29, 1988
AP	BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N16682</u>	<u>001</u>	
AP	HOSPIRA	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N17641</u>	<u>001</u>	

SOLUTION; IRRIGATION

LACTATED RINGER'S IN PLASTIC CONTAINER

AT	B BRAUN	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N18681</u>	<u>001</u>	Dec 27, 1982
AT	BAXTER HLTHCARE	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N18494</u>	<u>001</u>	Feb 19, 1982
AT		<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N18921</u>	<u>001</u>	Apr 03, 1984
AT		<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N19933</u>	<u>001</u>	Aug 29, 1989
AT	HOSPIRA	<u>20MG/100ML; 30MG/100ML; 600MG/100ML; 310MG/100ML</u>	<u>N19416</u>	<u>001</u>	Jan 17, 1986

CALFACTANT

SUSPENSION; INTRATRACHEAL

INFASURF PRESERVATIVE FREE

+	ONY	35MG/ML	N20521	001	Jul 01, 1998
---	-----	---------	--------	-----	--------------

CANDESARTAN CILEXETIL

TABLET; ORAL

ATACAND

	ASTRAZENECA	4MG	N20838	001	Jun 04, 1998
+		8MG	N20838	002	Jun 04, 1998
		16MG	N20838	003	Jun 04, 1998
+		32MG	N20838	004	Jun 04, 1998

CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ATACAND HCT

	ASTRAZENECA	16MG; 12.5MG	N21093	001	Sep 05, 2000
+		32MG; 12.5MG	N21093	002	Sep 05, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 64 (of 360)

CAPECITABINE

TABLET; ORAL

XELODA

HLR

150MG

N20896 001

Apr 30, 1998

+

500MG

N20896 002

Apr 30, 1998

CAPREOMYCIN SULFATE

INJECTABLE; INJECTION

CAPASTAT SULFATE

+ LILLY

EQ 1GM BASE/VIAL

N50095 001

CAPTOPRIL

TABLET; ORAL

CAPOTENAB PAR PHARM12.5MGN18343 005

Jan 17, 1985

AB25MGN18343 002AB50MGN18343 001AB

+

100MGN18343 003CAPTOPRILAB

APOTHECON

12.5MGN74472 001

Mar 31, 1995

AB25MGN74472 002

Mar 31, 1995

AB50MGN74472 003

Mar 31, 1995

AB100MGN74472 004

Mar 31, 1995

AB

CLONMEL HLTHCARE

12.5MGN74423 001

Feb 13, 1996

AB25MGN74423 002

Feb 13, 1996

AB50MGN74423 003

Feb 13, 1996

AB100MGN74423 004

Feb 13, 1996

AB

EGIS PHARMS

12.5MGN74748 004

May 29, 1997

AB25MGN74748 002

May 29, 1997

AB50MGN74748 001

May 29, 1997

AB100MGN74748 003

May 29, 1997

AB

ENDO LABS

12.5MGN74418 001

Feb 13, 1996

AB25MGN74418 002

Feb 13, 1996

AB50MGN74418 003

Feb 13, 1996

AB100MGN74418 004

Feb 13, 1996

AB

IVAX PHARMS

12.5MGN74590 004

Aug 30, 1996

AB25MGN74590 002

Aug 30, 1996

AB50MGN74590 001

Aug 30, 1996

AB100MGN74590 003

Aug 30, 1996

AB

KALI LABS

12.5MGN74477 001

Feb 13, 1996

AB25MGN74477 002

Feb 13, 1996

AB50MGN74477 003

Feb 13, 1996

AB100MGN74477 004

Feb 13, 1996

AB

MYLAN

12.5MGN74434 001

Feb 13, 1996

AB25MGN74434 002

Feb 13, 1996

AB50MGN74434 003

Feb 13, 1996

AB100MGN74434 004

Feb 13, 1996

AB

SANDOZ

12.5MGN74363 001

Nov 09, 1995

AB12.5MGN74481 001

Feb 13, 1996

AB12.5MGN74519 001

Feb 13, 1996

AB25MGN74363 002

Nov 09, 1995

AB25MGN74481 002

Feb 13, 1996

AB25MGN74519 002

Feb 13, 1996

AB50MGN74363 003

Nov 09, 1995

AB50MGN74481 003

Feb 13, 1996

AB50MGN74519 003

Feb 13, 1996

AB100MGN74363 004

Nov 09, 1995

AB100MGN74481 004

Feb 13, 1996

AB100MGN74519 004

Feb 13, 1996

AB

STASON

12.5MGN74677 004

May 30, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 65 (of 360)

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

<u>AB</u>	STASON	<u>25MG</u>	<u>N74677</u>	<u>002</u>	May 30, 1997
<u>AB</u>		<u>50MG</u>	<u>N74677</u>	<u>001</u>	May 30, 1997
<u>AB</u>		<u>100MG</u>	<u>N74677</u>	<u>003</u>	May 30, 1997
<u>AB</u>	TEVA	<u>12.5MG</u>	<u>N74322</u>	<u>001</u>	Feb 13, 1996
<u>AB</u>		<u>12.5MG</u>	<u>N74433</u>	<u>001</u>	Feb 13, 1996
<u>AB</u>		<u>12.5MG</u>	<u>N74483</u>	<u>001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>N74322</u>	<u>002</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>N74433</u>	<u>002</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>N74483</u>	<u>002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>N74322</u>	<u>003</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>N74433</u>	<u>003</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>N74483</u>	<u>003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>N74322</u>	<u>004</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>N74433</u>	<u>004</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>N74483</u>	<u>004</u>	Feb 13, 1996
<u>AB</u>	TEVA PHARMS	<u>12.5MG</u>	<u>N74462</u>	<u>001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>N74462</u>	<u>002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>N74462</u>	<u>003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>N74462</u>	<u>004</u>	Feb 13, 1996
<u>AB</u>	TORPHARM	<u>12.5MG</u>	<u>N74737</u>	<u>001</u>	Oct 28, 1998
<u>AB</u>		<u>25MG</u>	<u>N74737</u>	<u>002</u>	Oct 28, 1998
<u>AB</u>		<u>50MG</u>	<u>N74737</u>	<u>003</u>	Oct 28, 1998
<u>AB</u>		<u>100MG</u>	<u>N74737</u>	<u>004</u>	Oct 28, 1998
<u>AB</u>	WATSON LABS	<u>12.5MG</u>	<u>N74386</u>	<u>001</u>	May 23, 1996
<u>AB</u>		<u>12.5MG</u>	<u>N74451</u>	<u>001</u>	Feb 13, 1996
<u>AB</u>		<u>12.5MG</u>	<u>N74576</u>	<u>001</u>	Apr 23, 1996
<u>AB</u>		<u>25MG</u>	<u>N74386</u>	<u>002</u>	May 23, 1996
<u>AB</u>		<u>25MG</u>	<u>N74451</u>	<u>002</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>N74576</u>	<u>002</u>	Apr 23, 1996
<u>AB</u>		<u>50MG</u>	<u>N74386</u>	<u>003</u>	May 23, 1996
<u>AB</u>		<u>50MG</u>	<u>N74451</u>	<u>003</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>N74576</u>	<u>003</u>	Apr 23, 1996
<u>AB</u>		<u>100MG</u>	<u>N74386</u>	<u>004</u>	May 23, 1996
<u>AB</u>		<u>100MG</u>	<u>N74451</u>	<u>004</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>N74576</u>	<u>004</u>	Apr 23, 1996
<u>AB</u>	WEST WARD	<u>12.5MG</u>	<u>N74505</u>	<u>001</u>	Feb 13, 1996
<u>AB</u>		<u>25MG</u>	<u>N74505</u>	<u>002</u>	Feb 13, 1996
<u>AB</u>		<u>50MG</u>	<u>N74505</u>	<u>003</u>	Feb 13, 1996
<u>AB</u>		<u>100MG</u>	<u>N74505</u>	<u>004</u>	Feb 13, 1996
<u>AB</u>	WOCKHARDT	<u>12.5MG</u>	<u>N74532</u>	<u>001</u>	Mar 28, 1997
<u>AB</u>		<u>25MG</u>	<u>N74532</u>	<u>002</u>	Mar 28, 1997
<u>AB</u>		<u>50MG</u>	<u>N74532</u>	<u>003</u>	Mar 28, 1997
<u>AB</u>		<u>100MG</u>	<u>N74532</u>	<u>004</u>	Mar 28, 1997

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPOZIDE 25/15

<u>AB</u>	APOTHECON	<u>25MG;15MG</u>	<u>N18709</u>	<u>001</u>	Oct 12, 1984
<u>AB</u>	<u>CAPOZIDE 25/25</u>				
<u>AB</u>	+ APOTHECON	<u>25MG;25MG</u>	<u>N18709</u>	<u>002</u>	Oct 12, 1984
<u>AB</u>	<u>CAPOZIDE 50/15</u>				
<u>AB</u>	+ APOTHECON	<u>50MG;15MG</u>	<u>N18709</u>	<u>004</u>	Oct 12, 1984
<u>AB</u>	<u>CAPOZIDE 50/25</u>				
<u>AB</u>	APOTHECON	<u>50MG;25MG</u>	<u>N18709</u>	<u>003</u>	Oct 12, 1984
<u>AB</u>	<u>CAPTOPRIL AND HYDROCHLOROTHIAZIDE</u>				
<u>AB</u>	ENDO LABS	<u>25MG;15MG</u>	<u>N74788</u>	<u>001</u>	Dec 29, 1997
<u>AB</u>		<u>25MG;25MG</u>	<u>N74788</u>	<u>002</u>	Dec 29, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 66 (of 360)

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL

CAPTOPRIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ENDO LABS	<u>50MG;15MG</u>	<u>N74788</u>	<u>004</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;25MG</u>	<u>N74788</u>	<u>003</u>	Dec 29, 1997
<u>AB</u>	IVAX PHARMS	<u>25MG;15MG</u>	<u>N75055</u>	<u>001</u>	Jun 18, 1998
<u>AB</u>		<u>25MG;25MG</u>	<u>N75055</u>	<u>002</u>	Jun 18, 1998
<u>AB</u>		<u>50MG;15MG</u>	<u>N75055</u>	<u>004</u>	Jun 18, 1998
<u>AB</u>		<u>50MG;25MG</u>	<u>N75055</u>	<u>003</u>	Jun 18, 1998
<u>AB</u>	MYLAN	<u>25MG;15MG</u>	<u>N74896</u>	<u>001</u>	Dec 29, 1997
<u>AB</u>		<u>25MG;25MG</u>	<u>N74896</u>	<u>002</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;15MG</u>	<u>N74896</u>	<u>004</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;25MG</u>	<u>N74896</u>	<u>003</u>	Dec 29, 1997
<u>AB</u>	TEVA	<u>25MG;15MG</u>	<u>N74827</u>	<u>001</u>	Dec 29, 1997
<u>AB</u>		<u>25MG;25MG</u>	<u>N74827</u>	<u>002</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;15MG</u>	<u>N74827</u>	<u>004</u>	Dec 29, 1997
<u>AB</u>		<u>50MG;25MG</u>	<u>N74827</u>	<u>003</u>	Dec 29, 1997

CARBACHOL

SOLUTION; INTRAOCULAR

CARBASTAT

<u>AT</u>	NOVARTIS	<u>0.01%</u>	<u>N73677</u>	<u>001</u>	Apr 28, 1995
	<u>MIOSTAT</u>				
<u>AT</u>	+ ALCON	<u>0.01%</u>	<u>N16968</u>	<u>001</u>	

CARBAMAZEPINE

CAPSULE, EXTENDED RELEASE; ORAL

CARBATROL

	SHIRE	100MG	N20712	003	Sep 30, 1997
		200MG	N20712	001	Sep 30, 1997
	+	300MG	N20712	002	Sep 30, 1997
	EQUETRO				
	SHIRE	100MG	N21710	001	Dec 10, 2004
		200MG	N21710	002	Dec 10, 2004
	+	300MG	N21710	003	Dec 10, 2004

SUSPENSION; ORAL

CARBAMAZEPINE

<u>AB</u>	MORTON GROVE	<u>100MG/5ML</u>	<u>N75714</u>	<u>001</u>	Jun 05, 2002
	<u>TEGRETOL</u>				
<u>AB</u>	+ NOVARTIS	<u>100MG/5ML</u>	<u>N18927</u>	<u>001</u>	Dec 18, 1987
	<u>TERIL</u>				
<u>AB</u>	TARO	<u>100MG/5ML</u>	<u>N76729</u>	<u>001</u>	Sep 20, 2004

TABLET; ORAL

CARBAMAZEPINE

<u>AB</u>	APOTEX	<u>200MG</u>	<u>N75948</u>	<u>001</u>	Feb 27, 2002
<u>AB</u>	CARACO	<u>200MG</u>	<u>N77272</u>	<u>002</u>	Dec 07, 2005
		100MG	N77272	001	Dec 07, 2005
		300MG	N77272	003	Dec 07, 2005
		400MG	N77272	004	Dec 07, 2005
<u>AB</u>	INWOOD LABS	<u>200MG</u>	<u>N70231</u>	<u>001</u>	Aug 14, 1986
<u>AB</u>	PLIVA	<u>200MG</u>	<u>N71479</u>	<u>001</u>	Jul 24, 1987
<u>AB</u>	PUREPAC PHARM	<u>200MG</u>	<u>N71696</u>	<u>001</u>	Nov 09, 1987
<u>AB</u>	TARO	<u>200MG</u>	<u>N74649</u>	<u>001</u>	Oct 03, 1996
	<u>EPITOL</u>				
<u>AB</u>	TEVA	<u>200MG</u>	<u>N70541</u>	<u>001</u>	Sep 17, 1986
	<u>TEGRETOL</u>				
<u>AB</u>	+ NOVARTIS	<u>200MG</u>	<u>N16608</u>	<u>001</u>	
	<u>TERIL</u>				
<u>AB</u>	TARO	<u>200MG</u>	<u>N76525</u>	<u>001</u>	Sep 26, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 67 (of 360)

CARBAMAZEPINE

TABLET, CHEWABLE; ORAL

CARBAMAZEPINE

<u>AB</u>	CARACO	<u>100MG</u>	<u>N75712</u>	<u>001</u>	Jul 05, 2001
<u>AB</u>	TARO PHARM INDS	<u>100MG</u>	<u>N75687</u>	<u>001</u>	Oct 24, 2000
	+	200MG	N75687	002	Jul 29, 2002
<u>AB</u>	WARNER CHILCOTT	<u>100MG</u>	<u>N71940</u>	<u>001</u>	Feb 01, 1988
	<u>EPITOL</u>				
<u>AB</u>	TEVA	<u>100MG</u>	<u>N73524</u>	<u>001</u>	Jul 29, 1992
	<u>TEGRETOL</u>				
<u>AB</u>	+ NOVARTIS	<u>100MG</u>	<u>N18281</u>	<u>001</u>	
	TABLET, EXTENDED RELEASE; ORAL				
	TEGRETOL-XR				
	NOVARTIS	100MG	N20234	001	Mar 25, 1996
		200MG	N20234	002	Mar 25, 1996
	+	400MG	N20234	003	Mar 25, 1996

CARBENICILLIN INDANYL SODIUM

TABLET; ORAL

GEOCILLIN

+	PFIZER	EQ 382MG BASE	N50435	001	
---	--------	---------------	--------	-----	--

CARBIDOPA

TABLET; ORAL

LODOSYN

+	BRISTOL MYERS SQUIBB	25MG	N17830	001	
---	----------------------	------	--------	-----	--

CARBIDOPA; ENTACAPONE; LEVODOPA

TABLET; ORAL

STALEVO 100

ORION PHARMA INC	25MG;200MG;100MG	N21485	002	Jun 11, 2003
------------------	------------------	--------	-----	--------------

STALEVO 150

+	ORION PHARMA INC	37.5MG;200MG;150MG	N21485	003	Jun 11, 2003
---	------------------	--------------------	--------	-----	--------------

STALEVO 50

ORION PHARMA INC	12.5MG;200MG;50MG	N21485	001	Jun 11, 2003
------------------	-------------------	--------	-----	--------------

CARBIDOPA; LEVODOPA

TABLET; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	PUREPAC PHARM	<u>10MG;100MG</u>	<u>N74260</u>	<u>001</u>	Sep 03, 1993
<u>AB</u>		<u>25MG;100MG</u>	<u>N74260</u>	<u>002</u>	Sep 03, 1993
<u>AB</u>		<u>25MG;250MG</u>	<u>N74260</u>	<u>003</u>	Sep 03, 1993
<u>AB</u>	SANDOZ	<u>10MG;100MG</u>	<u>N73586</u>	<u>001</u>	Jun 29, 1995
<u>AB</u>		<u>25MG;100MG</u>	<u>N73587</u>	<u>001</u>	Jun 29, 1995
<u>AB</u>		<u>25MG;250MG</u>	<u>N73620</u>	<u>001</u>	Jun 29, 1995
<u>AB</u>	TEVA	<u>10MG;100MG</u>	<u>N73618</u>	<u>001</u>	Aug 28, 1992
<u>AB</u>		<u>25MG;100MG</u>	<u>N73589</u>	<u>001</u>	Aug 28, 1992
<u>AB</u>		<u>25MG;250MG</u>	<u>N73607</u>	<u>001</u>	Aug 28, 1992
	<u>SINEMET</u>				
<u>AB</u>	BRISTOL MYERS SQUIBB	<u>10MG;100MG</u>	<u>N17555</u>	<u>001</u>	
<u>AB</u>		<u>25MG;100MG</u>	<u>N17555</u>	<u>003</u>	
<u>AB</u>	+	<u>25MG;250MG</u>	<u>N17555</u>	<u>002</u>	

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	IMPAX LABS	<u>25MG;100MG</u>	<u>N76521</u>	<u>001</u>	May 14, 2004
<u>AB</u>		<u>50MG;200MG</u>	<u>N76521</u>	<u>002</u>	May 14, 2004
<u>AB</u>	KV PHARM	<u>50MG;200MG</u>	<u>N76663</u>	<u>001</u>	Jun 24, 2004
<u>AB</u>	MYLAN	<u>25MG;100MG</u>	<u>N75091</u>	<u>002</u>	Apr 21, 2000
<u>AB</u>		<u>50MG;200MG</u>	<u>N75091</u>	<u>001</u>	Sep 30, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 68 (of 360)

CARBIDOPA; LEVODOPA

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

<u>AB</u>	TORPHARM	<u>25MG;100MG</u>	<u>N76212</u>	<u>001</u>	Jun 16, 2004
<u>AB</u>		<u>50MG;200MG</u>	<u>N76212</u>	<u>002</u>	Jun 16, 2004
	<u>SINEMET CR</u>				
<u>AB</u>	BRISTOL MYERS SQUIBB	<u>25MG;100MG</u>	<u>N19856</u>	<u>002</u>	Dec 24, 1992
<u>AB</u>	+	<u>50MG;200MG</u>	<u>N19856</u>	<u>001</u>	May 30, 1991

TABLET, FOR SUSPENSION; ORAL

CARBILEV

	RANBAXY	10MG;100MG	N76643	001	Jun 10, 2005
		25MG;100MG	N76643	002	Jun 10, 2005
+		25MG;250MG	N76643	003	Jun 10, 2005

TABLET, ORALLY DISINTEGRATING; ORAL

PARCOPA

	SCHWARZ PHARMA	10MG;100MG	N76699	001	Aug 27, 2004
		25MG;100MG	N76699	002	Aug 27, 2004
+		25MG;250MG	N76699	003	Aug 27, 2004

CARBINOXAMINE MALEATE

SOLUTION; ORAL

CARBINOXAMINE MALEATE

+	MIKART	4MG/5ML	N40458	001	Apr 25, 2003
---	--------	---------	--------	-----	--------------

TABLET; ORAL

CARBINOXAMINE MALEATE

+	MIKART	4MG	N40442	001	Mar 19, 2003
---	--------	-----	--------	-----	--------------

CARBOPLATIN

INJECTABLE; INJECTION

CARBOPLATIN

<u>AP</u>	<u>AM PHARM PARTNERS</u>	<u>50MG/VIAL</u>	<u>N76235</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/VIAL</u>	<u>N76235</u>	<u>002</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/VIAL</u>	<u>N76235</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>	BEDFORD	<u>50MG/VIAL</u>	<u>N76099</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/VIAL</u>	<u>N76099</u>	<u>002</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/VIAL</u>	<u>N76099</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>	PHARMACHEMIE	<u>50MG/VIAL</u>	<u>N76162</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>		<u>150MG/VIAL</u>	<u>N76162</u>	<u>002</u>	Oct 14, 2004
<u>AP</u>		<u>450MG/VIAL</u>	<u>N76162</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>	PLIVA	<u>50MG/VIAL</u>	<u>N76602</u>	<u>001</u>	Nov 16, 2004
<u>AP</u>		<u>150MG/VIAL</u>	<u>N76602</u>	<u>002</u>	Nov 16, 2004
<u>AP</u>		<u>450MG/VIAL</u>	<u>N76602</u>	<u>003</u>	Nov 16, 2004
<u>AP</u>	SANDOZ	<u>50MG/VIAL</u>	<u>N76959</u>	<u>001</u>	Mar 18, 2005
<u>AP</u>		<u>150MG/VIAL</u>	<u>N76959</u>	<u>002</u>	Mar 18, 2005
<u>AP</u>		<u>450MG/VIAL</u>	<u>N76959</u>	<u>003</u>	Mar 18, 2005
	<u>PARAPLATIN</u>				
<u>AP</u>	+ BRISTOL MYERS SQUIBB	<u>50MG/VIAL</u>	<u>N19880</u>	<u>001</u>	Mar 03, 1989
<u>AP</u>	+	<u>150MG/VIAL</u>	<u>N19880</u>	<u>002</u>	Mar 03, 1989
<u>AP</u>	+	<u>450MG/VIAL</u>	<u>N19880</u>	<u>003</u>	Mar 03, 1989

INJECTABLE; IV (INFUSION)

CARBOPLATIN

<u>AP</u>	AM PHARM	<u>EQ 50MG/5ML (10MG/ML)</u>	<u>N77247</u>	<u>001</u>	Oct 21, 2004
<u>AP</u>		<u>EQ 150MG/15ML (10MG/ML)</u>	<u>N77247</u>	<u>002</u>	Oct 21, 2004
<u>AP</u>		<u>EQ 450MG/45ML (10MG/ML)</u>	<u>N77247</u>	<u>003</u>	Oct 21, 2004
<u>AP</u>	BEDFORD LABS	<u>EQ 50MG/5ML (10MG/ML)</u>	<u>N77244</u>	<u>001</u>	Oct 15, 2004
<u>AP</u>		<u>EQ 150MG/15ML (10MG/ML)</u>	<u>N77244</u>	<u>002</u>	Oct 15, 2004
<u>AP</u>		<u>EQ 450MG/45ML (10MG/ML)</u>	<u>N77244</u>	<u>003</u>	Oct 15, 2004
<u>AP</u>	MAYNE PHARMA USA	<u>EQ 50MG/5ML (10MG/ML)</u>	<u>N76517</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>		<u>EQ 150MG/15ML (10MG/ML)</u>	<u>N76517</u>	<u>002</u>	Oct 14, 2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 69 (of 360)

CARBOPLATIN

INJECTABLE; IV (INFUSION)

CARBOPLATIN

<u>AP</u>	MAYNE PHARMA USA	<u>EQ 450MG/45ML (10MG/ML)</u>	<u>N76517</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>		<u>EQ 600MG/ML (10MG/ML)</u>	<u>N77059</u>	<u>001</u>	Nov 23, 2004
<u>AP</u>	PHARMACHEMIE	<u>EQ 50MG/5ML (10MG/ML)</u>	<u>N77269</u>	<u>001</u>	Oct 14, 2004
<u>AP</u>		<u>EQ 150MG/15ML (10MG/ML)</u>	<u>N77269</u>	<u>002</u>	Oct 14, 2004
<u>AP</u>		<u>EQ 450MG/45ML (10MG/ML)</u>	<u>N77269</u>	<u>003</u>	Oct 14, 2004
<u>AP</u>	SICOR PHARMS	<u>EQ 50MG/5ML (10MG/ML)</u>	<u>N77139</u>	<u>001</u>	Sep 21, 2005
<u>AP</u>		<u>EQ 150MG/15ML (10MG/ML)</u>	<u>N77139</u>	<u>002</u>	Sep 21, 2005
<u>AP</u>		<u>EQ 450MG/45ML (10MG/ML)</u>	<u>N77139</u>	<u>003</u>	Sep 21, 2005
<u>AP</u>		<u>EQ 600MG/60ML (10MG/ML)</u>	<u>N77139</u>	<u>004</u>	Sep 21, 2005
<u>AP</u>	SPECTRUM PHARMS	<u>EQ 50MG/5ML (10MG/ML)</u>	<u>N77096</u>	<u>001</u>	Jun 14, 2005
<u>AP</u>		<u>EQ 150MG/15ML (10MG/ML)</u>	<u>N77096</u>	<u>002</u>	Jun 14, 2005
<u>AP</u>		<u>EQ 450MG/45ML (10MG/ML)</u>	<u>N77096</u>	<u>003</u>	Jun 14, 2005
<u>PARAPLATIN</u>					
<u>AP</u>	+ BRISTOL MYERS SQUIBB	<u>EQ 50MG /5ML (10MG/ML)</u>	<u>N20452</u>	<u>001</u>	Jul 14, 2003
<u>AP</u>	+	<u>EQ 150MG /15ML (10MG/ML)</u>	<u>N20452</u>	<u>002</u>	Jul 14, 2003
<u>AP</u>	+	<u>EQ 450MG /45ML (10MG/ML)</u>	<u>N20452</u>	<u>003</u>	Jul 14, 2003
<u>AP</u>	+	<u>EQ 600MG /60ML (10MG/ML)</u>	<u>N20452</u>	<u>004</u>	Jan 15, 2004

CARBOPROST TROMETHAMINE

INJECTABLE; INJECTION

HEMABATE

+ PHARMACIA AND UPJOHN EQ 0.25MG BASE/ML N17989 001

CARISOPRODOL

TABLET; ORAL

CARISOPRODOL

<u>AA</u>	AMIDE PHARM	<u>350MG</u>	<u>N40188</u>	<u>001</u>	Mar 07, 1997
<u>AA</u>	COREPHARMA	<u>350MG</u>	<u>N40397</u>	<u>001</u>	Sep 21, 2000
<u>AA</u>	MUTUAL PHARM	<u>350MG</u>	<u>N89346</u>	<u>001</u>	Oct 17, 1991
<u>AA</u>	NEIL	<u>350MG</u>	<u>N40576</u>	<u>001</u>	Jun 07, 2005
<u>AA</u>	SANDOZ	<u>350MG</u>	<u>N81025</u>	<u>001</u>	Apr 13, 1989
<u>AA</u>	VINTAGE PHARMS	<u>350MG</u>	<u>N40245</u>	<u>001</u>	Sep 08, 1997
<u>AA</u>	WATSON LABS	<u>350MG</u>	<u>N40152</u>	<u>001</u>	Dec 03, 1996
<u>AA</u>		<u>350MG</u>	<u>N86179</u>	<u>001</u>	
<u>AA</u>		<u>350MG</u>	<u>N87499</u>	<u>001</u>	Apr 20, 1982
<u>AA</u>	WEST WARD	<u>350MG</u>	<u>N40124</u>	<u>001</u>	Jan 24, 1996
<u>SOMA</u>					
<u>AA</u>	+ MEDPOINTE PHARM HLC	<u>350MG</u>	<u>N11792</u>	<u>001</u>	

CARMUSTINE

IMPLANT; INTRACRANIAL

GLIADEL

+ MGI PHARMA INC 7.7MG N20637 001 Sep 23, 1996

INJECTABLE; INJECTION

BICNU

+ BRISTOL 100MG/VIAL N17422 001

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CARTEOLOL HYDROCHLORIDE

<u>AT</u>	ALCON	<u>1%</u>	<u>N75476</u>	<u>001</u>	Jan 03, 2000
<u>AT</u>	BAUSCH AND LOMB	<u>1%</u>	<u>N75546</u>	<u>001</u>	Jan 20, 2000
<u>AT</u>	NOVEX	<u>1%</u>	<u>N76097</u>	<u>001</u>	Feb 06, 2002
<u>OCUPRESS</u>					
<u>AT</u>	+ NOVARTIS	<u>1%</u>	<u>N19972</u>	<u>001</u>	May 23, 1990

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 70 (of 360)

CARVEDILOL

TABLET; ORAL

COREG

GLAXOSMITHKLINE	3.125MG	N20297	004	May 29, 1997
	6.25MG	N20297	003	Sep 14, 1995
+	12.5MG	N20297	002	Sep 14, 1995
	25MG	N20297	001	Sep 14, 1995

CASPOFUNGIN ACETATE

INJECTABLE; IV (INFUSION)

CANCIDAS

+ MERCK RES	50MG/VIAL	N21227	001	Jan 26, 2001
+	70MG/VIAL	N21227	002	Jan 26, 2001

CEFACLOR

CAPSULE; ORAL

CEFACLOR

<u>AB</u>	CARLSBAD	<u>EQ 250MG BASE</u>	<u>N65146</u>	<u>001</u>	Jan 22, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N65146</u>	<u>002</u>	Jan 22, 2004
<u>AB</u>	CEPH INTL	<u>EQ 250MG BASE</u>	<u>N62205</u>	<u>001</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N62205</u>	<u>002</u>	
<u>AB</u>	IVAX PHARMS	<u>EQ 250MG BASE</u>	<u>N64061</u>	<u>001</u>	Apr 27, 1995
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N64061</u>	<u>002</u>	Apr 27, 1995
<u>AB</u>	MARSAM PHARMS LLC	<u>EQ 250MG BASE</u>	<u>N64148</u>	<u>001</u>	May 23, 1996
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N64148</u>	<u>002</u>	May 23, 1996
<u>AB</u>	RANBAXY	<u>EQ 250MG BASE</u>	<u>N64156</u>	<u>001</u>	Aug 28, 1997
<u>AB</u> +		<u>EQ 500MG BASE</u>	<u>N64156</u>	<u>002</u>	Aug 28, 1997
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>N64145</u>	<u>001</u>	Jun 24, 1996
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N64145</u>	<u>002</u>	Jun 24, 1996

FOR SUSPENSION; ORAL

CEFACLOR

<u>AB</u>	CEPH INTL	<u>EQ 125MG BASE/5ML</u>	<u>N62206</u>	<u>001</u>	
<u>AB</u>		<u>EQ 187MG BASE/5ML</u>	<u>N62206</u>	<u>003</u>	Apr 20, 1988
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N62206</u>	<u>002</u>	
<u>AB</u>		<u>EQ 375MG BASE/5ML</u>	<u>N62206</u>	<u>004</u>	Apr 20, 1988
<u>AB</u>	IVAX PHARMS	<u>EQ 375MG BASE/5ML</u>	<u>N64070</u>	<u>001</u>	Apr 28, 1995
<u>AB</u>	MARSAM PHARMS LLC	<u>EQ 125MG BASE/5ML</u>	<u>N64204</u>	<u>001</u>	Feb 18, 1998
<u>AB</u>		<u>EQ 187MG BASE/5ML</u>	<u>N64205</u>	<u>001</u>	Feb 18, 1998
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N64206</u>	<u>001</u>	Feb 18, 1998
<u>AB</u>		<u>EQ 375MG BASE/5ML</u>	<u>N64207</u>	<u>001</u>	Feb 18, 1998
<u>AB</u>	RANBAXY	<u>EQ 125MG BASE/5ML</u>	<u>N64166</u>	<u>001</u>	Oct 02, 1997
<u>AB</u>		<u>EQ 187MG BASE/5ML</u>	<u>N64165</u>	<u>001</u>	Oct 02, 1997
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N64164</u>	<u>001</u>	Oct 02, 1997
<u>AB</u> +		<u>EQ 375MG BASE/5ML</u>	<u>N64155</u>	<u>001</u>	Oct 02, 1997

TABLET, CHEWABLE; ORAL

RANICLOR

RANBAXY	EQ 125MG BASE	N65092	001	Dec 22, 2003
	EQ 187MG BASE	N65092	002	Dec 22, 2003
	EQ 250MG BASE	N65092	003	Dec 22, 2003
+	EQ 375MG BASE	N65092	004	Dec 22, 2003

TABLET, EXTENDED RELEASE; ORAL

CEFACLOR

<u>AB</u> +	IVAX PHARMS	<u>EQ 500MG BASE</u>	<u>N65057</u>	<u>001</u>	Jan 05, 2001
<u>AB</u>	TEVA	<u>EQ 500MG BASE</u>	<u>N65058</u>	<u>002</u>	Sep 04, 2002
		EQ 375MG BASE	N65058	001	Sep 04, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 71 (of 360)

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

<u>AB</u>	APOTHECON	<u>EQ 500MG BASE</u>	<u>N62291</u>	<u>001</u>	
<u>AB</u>	IVAX PHARMS	<u>EQ 500MG BASE</u>	<u>N62766</u>	<u>001</u>	Mar 03, 1987
<u>AB</u>	RANBAXY	<u>EQ 500MG BASE</u>	<u>N65015</u>	<u>001</u>	Jun 22, 1999

DURICEF

<u>AB</u>	+ WARNER CHILCOTT	<u>EQ 500MG BASE</u>	<u>N50512</u>	<u>001</u>	
-----------	-------------------	----------------------	---------------	------------	--

FOR SUSPENSION; ORAL

CEFADROXIL

<u>AB</u>	RANBAXY	<u>EQ 125MG BASE/5ML</u>	<u>N65115</u>	<u>001</u>	Mar 26, 2003
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N65115</u>	<u>002</u>	Mar 26, 2003
<u>AB</u>		<u>EQ 500MG BASE/5ML</u>	<u>N65115</u>	<u>003</u>	Mar 26, 2003

DURICEF

<u>AB</u>	WARNER CHILCOTT	<u>EQ 125MG BASE/5ML</u>	<u>N50527</u>	<u>002</u>	
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N50527</u>	<u>003</u>	
<u>AB</u>	+	<u>EQ 500MG BASE/5ML</u>	<u>N50527</u>	<u>001</u>	

TABLET; ORAL

CEFADROXIL

<u>AB</u>	IVAX PHARMS	<u>EQ 1GM BASE</u>	<u>N62774</u>	<u>001</u>	Apr 08, 1987
<u>AB</u>	RANBAXY	<u>EQ 1GM BASE</u>	<u>N65018</u>	<u>001</u>	Apr 23, 1999

DURICEF

<u>AB</u>	+ WARNER CHILCOTT	<u>EQ 1GM BASE</u>	<u>N50528</u>	<u>001</u>	
-----------	-------------------	--------------------	---------------	------------	--

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

ANCEF

<u>AP</u>	+ GLAXOSMITHKLINE	<u>EQ 1GM BASE/VIAL</u>	<u>N50461</u>	<u>003</u>	
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>N50461</u>	<u>005</u>	

ANCEF IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	EQ 10MG BASE/ML	N63002	001	Mar 28, 1991
+		EQ 20MG BASE/ML	N63002	002	Mar 28, 1991

CEFAZOLIN

<u>AP</u>	ORCHID HLTHCARE	<u>EQ 1GM BASE</u>	<u>N65244</u>	<u>001</u>	Aug 12, 2005
<u>AP</u>		<u>EQ 10GM BASE</u>	<u>N65247</u>	<u>001</u>	Aug 12, 2005

CEFAZOLIN AND DEXTROSE

+	B BRAUN	EQ 500MG BASE/VIAL	N50779	001	Jul 27, 2000
+		EQ 1GM BASE/VIAL	N50779	002	Jul 27, 2000

CEFAZOLIN SODIUM

<u>AP</u>	+ AM PHARM PARTNERS	<u>EQ 500MG BASE/VIAL</u>	<u>N64169</u>	<u>001</u>	Aug 14, 1998
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N64169</u>	<u>002</u>	Aug 14, 1998
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>N64170</u>	<u>001</u>	Mar 18, 1998
<u>AP</u>	+	<u>EQ 20GM BASE/VIAL</u>	<u>N64170</u>	<u>002</u>	Mar 18, 1998
<u>AP</u>	APOTHECON	<u>EQ 500MG BASE/VIAL</u>	<u>N62831</u>	<u>001</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N62831</u>	<u>002</u>	Dec 09, 1988
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>N62831</u>	<u>003</u>	Sep 25, 1992
<u>AP</u>	GLAXOSMITHKLINE	<u>EQ 1GM BASE/VIAL</u>	<u>N64033</u>	<u>001</u>	Oct 31, 1993
<u>AP</u>	HANFORD GC	<u>EQ 500MG BASE/VIAL</u>	<u>N63214</u>	<u>001</u>	Dec 27, 1991
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>N63216</u>	<u>001</u>	Dec 27, 1991
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N63207</u>	<u>001</u>	Dec 27, 1991
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N63208</u>	<u>001</u>	Dec 27, 1991
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>N63209</u>	<u>001</u>	Dec 27, 1991
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 500MG BASE/VIAL</u>	<u>N65047</u>	<u>001</u>	Sep 18, 2001
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N65047</u>	<u>002</u>	Sep 18, 2001
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>N65143</u>	<u>001</u>	Oct 18, 2004
<u>AP</u>	ORCHID HLTHCARE	<u>EQ 500MG BASE/VIAL</u>	<u>N65226</u>	<u>001</u>	Apr 21, 2005
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N65226</u>	<u>002</u>	Apr 21, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 72 (of 360)

CEFDINIR

CAPSULE; ORAL					
OMNICEF					
+	ABBOTT	300MG	N50739	001	Dec 04, 1997
FOR SUSPENSION; ORAL					
OMNICEF					
	ABBOTT	125MG/5ML	N50749	001	Dec 04, 1997
+		250MG/5ML	N50749	002	Jul 29, 2004

CEFDITOREN PIVOXIL

TABLET; ORAL					
SPECTRACEF					
+	PURDUE PHARMA LP	200MG	N21222	001	Aug 29, 2001

CEFEPIME HYDROCHLORIDE (ARGININE FORMULATION)

INJECTABLE; INJECTION					
MAXIPIME					
+	BRISTOL MYERS SQUIBB	EQ 500MG BASE/VIAL	N50679	001	Jan 18, 1996
+		EQ 1GM BASE/VIAL	N50679	002	Jan 18, 1996
+		EQ 2GM BASE/VIAL	N50679	003	Jan 18, 1996

CEFIXIME

SUSPENSION; ORAL					
SUPRAX					
+	LUPIN	100MG/5ML	N65129	001	Feb 23, 2004

CEFOPERAZONE SODIUM

INJECTABLE; INJECTION					
CEFOBID					
+	PFIZER	EQ 1GM BASE/VIAL	N50551	001	Nov 18, 1982
+		EQ 2GM BASE/VIAL	N50551	002	Nov 18, 1982
+		EQ 10GM BASE/VIAL	N50551	003	Mar 05, 1990
CEFOBID IN PLASTIC CONTAINER					
+	PFIZER	EQ 20MG BASE/ML	N50613	002	Jul 31, 1987
+		EQ 40MG BASE/ML	N50613	001	Jul 23, 1986

CEFOTAXIME SODIUM

INJECTABLE; INJECTION					
<u>CEFOTAXIME</u>					
<u>AP</u>	AM PHARM PARTNERS	<u>EQ 500MG BASE/VIAL</u>	<u>N64200</u>	<u>001</u>	Mar 24, 2000
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N64200</u>	<u>002</u>	Mar 24, 2000
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>N64200</u>	<u>003</u>	Mar 24, 2000
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>N64201</u>	<u>001</u>	Mar 24, 2000
	+	EQ 20GM BASE/VIAL	N64201	002	Mar 24, 2000
<u>AP</u>	HIKMA	<u>EQ 500MG BASE/VIAL</u>	<u>N65072</u>	<u>001</u>	Nov 20, 2002
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N65072</u>	<u>002</u>	Nov 20, 2002
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>N65072</u>	<u>003</u>	Nov 20, 2002
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>N65071</u>	<u>001</u>	Nov 20, 2002
CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER					
+	B BRAUN	EQ 2GM BASE	N50792	001	Jul 29, 2004
CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER					
+	B BRAUN	EQ 1GM BASE	N50792	002	Jul 29, 2004
<u>CEFOTAXIME SODIUM</u>					
<u>AP</u>	LUPIN	<u>EQ 500MG BASE/VIAL</u>	<u>N65124</u>	<u>001</u>	Sep 24, 2003
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N65124</u>	<u>002</u>	Sep 24, 2003
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>N65124</u>	<u>003</u>	Sep 24, 2003
<u>CLAFORAN</u>					
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>N50547</u>	<u>001</u>	
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N50547</u>	<u>002</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 73 (of 360)

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CLAFORAN

<u>AP</u>	AVENTIS PHARMS	<u>EQ 1GM BASE/VIAL</u>	<u>N62659</u>	<u>001</u>	Jan 13, 1987
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>N50547</u>	<u>003</u>	
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>N62659</u>	<u>002</u>	Jan 13, 1987
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>N50547</u>	<u>004</u>	Dec 29, 1983
	CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER				
	+ AVENTIS PHARMS	EQ 20MG BASE/ML	N50596	002	May 20, 1985
	+	EQ 40MG BASE/ML	N50596	004	May 20, 1985

CEFOTETAN DISODIUM

INJECTABLE; INJECTION

CEFOTAN

+	ASTRAZENECA	EQ 1GM BASE/VIAL	N50588	001	Dec 27, 1985
		EQ 1GM BASE/VIAL	N63293	001	Apr 29, 1993
+		EQ 2GM BASE/VIAL	N50588	002	Dec 27, 1985
		EQ 2GM BASE/VIAL	N63293	002	Apr 29, 1993
+		EQ 10GM BASE/VIAL	N50588	003	Apr 25, 1988
	CEFOTAN IN PLASTIC CONTAINER				
+	ASTRAZENECA	EQ 20MG BASE/ML	N50694	002	Jul 30, 1993
+		EQ 40MG BASE/ML	N50694	001	Jul 30, 1993

CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

<u>AP</u>	+	AM PHARM PARTNERS	<u>EQ 1GM BASE/VIAL</u>	<u>N65012</u>	<u>001</u>	Jul 03, 2000
<u>AP</u>	+		<u>EQ 2GM BASE/VIAL</u>	<u>N65012</u>	<u>002</u>	Jul 03, 2000
<u>AP</u>	+		<u>EQ 10GM BASE/VIAL</u>	<u>N65011</u>	<u>001</u>	Jul 03, 2000
<u>AP</u>		BAXTER HLTHCARE	<u>EQ 1GM BASE/VIAL</u>	<u>N65051</u>	<u>001</u>	Sep 11, 2000
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>N65051</u>	<u>002</u>	Sep 11, 2000
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>N65050</u>	<u>001</u>	Sep 11, 2000

MEFOXIN

<u>AP</u>		MERCK	<u>EQ 1GM BASE/VIAL</u>	<u>N62757</u>	<u>001</u>	Jan 08, 1987
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>N62757</u>	<u>002</u>	Jan 08, 1987
	MEFOXIN IN PLASTIC CONTAINER					
	+ MERCK	EQ 20MG BASE/ML	N63182	001	Jan 25, 1993	
	+	EQ 40MG BASE/ML	N63182	002	Jan 25, 1993	

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

CEFPODOXIME PROXETIL

<u>AB</u>		RANBAXY	<u>EQ 50MG BASE/5ML</u>	<u>N65082</u>	<u>001</u>	May 31, 2002
<u>AB</u>			<u>EQ 100MG BASE/5ML</u>	<u>N65082</u>	<u>002</u>	May 31, 2002
	<u>VANTIN</u>					
<u>AB</u>		PHARMACIA AND UPJOHN	<u>EQ 50MG BASE/5ML</u>	<u>N50675</u>	<u>001</u>	Aug 07, 1992
<u>AB</u>	+		<u>EQ 100MG BASE/5ML</u>	<u>N50675</u>	<u>002</u>	Aug 07, 1992

TABLET; ORAL

CEFPODOXIME PROXETIL

<u>AB</u>		RANBAXY	<u>EQ 100MG BASE</u>	<u>N65083</u>	<u>001</u>	Aug 20, 2003
<u>AB</u>			<u>EQ 200MG BASE</u>	<u>N65083</u>	<u>002</u>	Aug 20, 2003
	<u>VANTIN</u>					
<u>AB</u>		PHARMACIA AND UPJOHN	<u>EQ 100MG BASE</u>	<u>N50674</u>	<u>001</u>	Aug 07, 1992
<u>AB</u>	+		<u>EQ 200MG BASE</u>	<u>N50674</u>	<u>002</u>	Aug 07, 1992

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

<u>AB</u>		LUPIN	<u>125MG/5ML</u>	<u>N65261</u>	<u>001</u>	Dec 19, 2005
<u>AB</u>			<u>250MG/5ML</u>	<u>N65261</u>	<u>002</u>	Dec 19, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 74 (of 360)

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

<u>AB</u>	ORCHID HLTHCARE	<u>125MG/5ML</u>	<u>N65284</u>	<u>002</u>	Dec 30, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>N65284</u>	<u>001</u>	Dec 30, 2005
<u>AB</u>	SANDOZ	<u>125MG/5ML</u>	<u>N65257</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>N65257</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>	TEVA PHARMS	<u>125MG/5ML</u>	<u>N65236</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>250MG/5ML</u>	<u>N65236</u>	<u>002</u>	Dec 08, 2005

CEFZIL

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>125MG/5ML</u>	<u>N50665</u>	<u>001</u>	Dec 23, 1991
<u>AB</u>	+	<u>250MG/5ML</u>	<u>N50665</u>	<u>002</u>	Dec 23, 1991

TABLET; ORAL

CEFPROZIL

<u>AB</u>	LUPIN	<u>250MG</u>	<u>N65276</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>500MG</u>	<u>N65276</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>	ORCHID HLTHCARE	<u>250MG</u>	<u>N65267</u>	<u>001</u>	Dec 19, 2005
<u>AB</u>		<u>500MG</u>	<u>N65267</u>	<u>002</u>	Dec 19, 2005
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>N65235</u>	<u>001</u>	Nov 14, 2005
<u>AB</u>		<u>500MG</u>	<u>N65235</u>	<u>002</u>	Nov 14, 2005
<u>AB</u>	TEVA	<u>250MG</u>	<u>N65208</u>	<u>001</u>	Dec 06, 2005
<u>AB</u>		<u>500MG</u>	<u>N65208</u>	<u>002</u>	Dec 06, 2005

CEFZIL

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>250MG</u>	<u>N50664</u>	<u>001</u>	Dec 23, 1991
<u>AB</u>	+	<u>500MG</u>	<u>N50664</u>	<u>002</u>	Dec 23, 1991

CEFTAZIDIME

INJECTABLE; INJECTION

CEFTAZIDIME

<u>AP</u>	ACS DOBFAR	<u>500MG/VIAL</u>	<u>N62640</u>	<u>001</u>	Nov 20, 1985
<u>AP</u>		<u>1GM/VIAL</u>	<u>N62640</u>	<u>002</u>	Nov 20, 1985
<u>AP</u>		<u>2GM/VIAL</u>	<u>N62640</u>	<u>003</u>	Nov 20, 1985
<u>AP</u>		<u>6GM/VIAL</u>	<u>N62640</u>	<u>004</u>	Feb 03, 1992

FORTAZ

<u>AP</u>	+	GLAXOSMITHKLINE	<u>500MG/VIAL</u>	<u>N50578</u>	<u>001</u>	Jul 19, 1985
<u>AP</u>	+		<u>1GM/VIAL</u>	<u>N50578</u>	<u>002</u>	Jul 19, 1985
<u>AP</u>	+		<u>2GM/VIAL</u>	<u>N50578</u>	<u>003</u>	Jul 19, 1985
<u>AP</u>	+		<u>6GM/VIAL</u>	<u>N50578</u>	<u>004</u>	Jul 19, 1985

TAZICEF

<u>AP</u>	HOSPIRA	<u>500MG/VIAL</u>	<u>N62662</u>	<u>001</u>	Mar 06, 1986
<u>AP</u>		<u>1GM/VIAL</u>	<u>N62662</u>	<u>002</u>	Mar 06, 1986
<u>AP</u>		<u>1GM/VIAL</u>	<u>N64032</u>	<u>001</u>	Oct 31, 1993
<u>AP</u>		<u>2GM/VIAL</u>	<u>N62662</u>	<u>003</u>	Mar 06, 1986
<u>AP</u>		<u>2GM/VIAL</u>	<u>N64032</u>	<u>002</u>	Oct 31, 1993
<u>AP</u>		<u>6GM/VIAL</u>	<u>N62662</u>	<u>004</u>	Mar 06, 1986

CEFTAZIDIME (ARGININE FORMULATION)

INJECTABLE; INJECTION

CEPTAZ

+	GLAXOSMITHKLINE	1GM/VIAL	N50646	002	Sep 27, 1990
+		2GM/VIAL	N50646	003	Sep 27, 1990
+		10GM/VIAL	N50646	004	Sep 27, 1990

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

CEFTAZIDIME SODIUM IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 20MG BASE/ML</u>	<u>N63221</u>	<u>002</u>	Apr 29, 1993
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N63221</u>	<u>003</u>	Apr 29, 1993
+		EQ 10MG BASE/ML	N63221	001	Apr 29, 1993

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 75 (of 360)

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

FORTAZ IN PLASTIC CONTAINER

<u>AP</u>	+	GLAXOSMITHKLINE	<u>EQ 20MG BASE/ML</u>	<u>N50634</u>	<u>002</u>	Apr 28, 1989
<u>AP</u>	+		<u>EQ 40MG BASE/ML</u>	<u>N50634</u>	<u>003</u>	Apr 28, 1989

CEFTIBUTEN DIHYDRATE

CAPSULE; ORAL

CEDAX

+	SHIONOGI	EQ 400MG BASE	N50685	002	Dec 20, 1995
---	----------	---------------	--------	-----	--------------

FOR SUSPENSION; ORAL

CEDAX

+	SHIONOGI	EQ 90MG BASE/5ML	N50686	001	Dec 20, 1995
---	----------	------------------	--------	-----	--------------

CEFTIZOXIME SODIUM

INJECTABLE; INJECTION

CEFIZOX

+	ASTELLAS	EQ 1GM BASE/VIAL	N50560	002	Sep 15, 1983
		EQ 1GM BASE/VIAL	N63294	002	Mar 31, 1994
+		EQ 2GM BASE/VIAL	N50560	003	Sep 15, 1983
		EQ 2GM BASE/VIAL	N63294	003	Mar 31, 1994
		EQ 10GM BASE/VIAL	N50560	005	Mar 19, 1993

CEFIZOX IN PLASTIC CONTAINER

+	ASTELLAS	EQ 20MG BASE/ML	N50589	003	Apr 13, 1995
+		EQ 40MG BASE/ML	N50589	004	Apr 13, 1995

CEFTRIAXONE SODIUM

INJECTABLE; IM-IV

CEFTRIAXONE

<u>AP</u>		<u>LUPIN</u>	<u>EQ 250MG BASE/VIAL</u>	<u>N65125</u>	<u>001</u>	Sep 30, 2003
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>N65125</u>	<u>002</u>	Sep 30, 2003
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>N65125</u>	<u>003</u>	Sep 30, 2003
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>N65125</u>	<u>004</u>	Sep 30, 2003
<u>AP</u>		<u>ORCHID HLTHCARE</u>	<u>EQ 250MG BASE/VIAL</u>	<u>N65230</u>	<u>001</u>	Aug 02, 2005
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>N65230</u>	<u>002</u>	Aug 02, 2005
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>N65230</u>	<u>003</u>	Aug 02, 2005
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>N65230</u>	<u>004</u>	Aug 02, 2005
<u>AP</u>		<u>SANDOZ</u>	<u>EQ 250MG BASE/VIAL</u>	<u>N65169</u>	<u>001</u>	May 09, 2005
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>N65169</u>	<u>002</u>	May 09, 2005
<u>AP</u>			<u>EQ 1GM BASE/VIAL</u>	<u>N65169</u>	<u>003</u>	May 09, 2005
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>N65169</u>	<u>004</u>	May 09, 2005
<u>AP</u>	+	<u>HLR</u>	<u>EQ 250MG BASE/VIAL</u>	<u>N50585</u>	<u>001</u>	Dec 21, 1984
<u>AP</u>	+		<u>EQ 500MG BASE/VIAL</u>	<u>N50585</u>	<u>002</u>	Dec 21, 1984
<u>AP</u>	+		<u>EQ 1GM BASE/VIAL</u>	<u>N50585</u>	<u>003</u>	Dec 21, 1984
<u>AP</u>	+		<u>EQ 2GM BASE/VIAL</u>	<u>N50585</u>	<u>004</u>	Dec 21, 1984

INJECTABLE; INJECTION

CEFTRIAXONE

<u>AP</u>		<u>ORCHID HLTHCARE</u>	<u>EQ 1GM BASE/VIAL</u>	<u>N65231</u>	<u>001</u>	Aug 02, 2005
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>N65231</u>	<u>002</u>	Aug 02, 2005
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>N65232</u>	<u>001</u>	Aug 02, 2005
<u>AP</u>		<u>SANDOZ</u>	<u>EQ 1GM BASE/VIAL</u>	<u>N65204</u>	<u>001</u>	May 03, 2005
<u>AP</u>			<u>EQ 2GM BASE/VIAL</u>	<u>N65204</u>	<u>002</u>	May 03, 2005
<u>AP</u>			<u>EQ 10GM BASE/VIAL</u>	<u>N65168</u>	<u>001</u>	May 17, 2005
<u>CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER</u>						
<u>AP</u>	+	<u>B BRAUN</u>	<u>EQ 1GM BASE/VIAL</u>	<u>N50796</u>	<u>001</u>	Apr 20, 2005
<u>AP</u>	+		<u>EQ 2GM BASE/VIAL</u>	<u>N50796</u>	<u>002</u>	Apr 20, 2005
<u>CEFTRIAXONE IN PLASTIC CONTAINER</u>						
<u>AP</u>		<u>BAXTER HLTHCARE</u>	<u>EQ 20MG BASE/ML</u>	<u>N65224</u>	<u>001</u>	Aug 23, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 76 (of 360)

CEFTRIAZONE SODIUM

INJECTABLE; INJECTION

CEFTRIAZONE IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 40MG BASE/ML</u>	<u>N65224</u>	<u>002</u>	Aug 23, 2005
	<u>ROCEPHIN</u>				
<u>AP</u>	+ HLR	<u>EQ 1GM BASE/VIAL</u>	<u>N62654</u>	<u>002</u>	Apr 30, 1987
<u>AP</u>	+	<u>EQ 2GM BASE/VIAL</u>	<u>N62654</u>	<u>003</u>	Apr 30, 1987
<u>AP</u>	+	<u>EQ 10GM BASE/VIAL</u>	<u>N50585</u>	<u>005</u>	Dec 21, 1984
	<u>ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER</u>				
<u>AP</u>	+ HLR	<u>EQ 20MG BASE/ML</u>	<u>N50624</u>	<u>002</u>	Feb 11, 1987
<u>AP</u>	+	<u>EQ 40MG BASE/ML</u>	<u>N50624</u>	<u>003</u>	Feb 11, 1987

CEFUROXIME AXETIL

FOR SUSPENSION; ORAL

CEFTIN

	GLAXOSMITHKLINE	EQ 125MG BASE/5ML	N50672	001	Jun 30, 1994
	+	EQ 250MG BASE/5ML	N50672	002	Apr 29, 1997

TABLET; ORAL

CEFTIN

<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 125MG BASE</u>	<u>N50605</u>	<u>001</u>	Dec 28, 1987
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N50605</u>	<u>002</u>	Dec 28, 1987
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>N50605</u>	<u>003</u>	Dec 28, 1987

CEFUROXIME AXETIL

<u>AB</u>	APOTEX	<u>EQ 250MG BASE</u>	<u>N65069</u>	<u>001</u>	Oct 02, 2002
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N65069</u>	<u>002</u>	Oct 02, 2002
<u>AB</u>	LUPIN	<u>EQ 250MG BASE</u>	<u>N65135</u>	<u>001</u>	Jul 25, 2003
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N65135</u>	<u>002</u>	Jul 25, 2003
<u>AB</u>	RANBAXY	<u>EQ 125MG BASE</u>	<u>N65043</u>	<u>003</u>	Feb 15, 2002
<u>AB</u>		<u>EQ 125MG BASE</u>	<u>N65118</u>	<u>001</u>	Apr 25, 2003
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N65043</u>	<u>002</u>	Feb 15, 2002
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N65118</u>	<u>002</u>	Apr 25, 2003
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N65043</u>	<u>001</u>	Feb 15, 2002
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N65118</u>	<u>003</u>	Apr 25, 2003
<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>	<u>N65126</u>	<u>001</u>	Oct 28, 2003
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N65126</u>	<u>002</u>	Oct 28, 2003
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>N65190</u>	<u>001</u>	Oct 18, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N65190</u>	<u>002</u>	Oct 18, 2004
<u>AB</u>	WOCKHARDT	<u>EQ 125MG BASE</u>	<u>N65166</u>	<u>001</u>	Jul 29, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N65166</u>	<u>002</u>	Jul 29, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N65166</u>	<u>003</u>	Jul 29, 2005

CEFUROXIME SODIUM

INJECTABLE; IM-IV

CEFUROXIME

<u>AB</u>	AM PHARM PARTNERS	<u>EQ 750MG BASE/VIAL</u>	<u>N65001</u>	<u>001</u>	May 30, 2001
<u>AB</u>	HIKMA FARMACEUTICA	<u>EQ 750MG BASE/VIAL</u>	<u>N65048</u>	<u>001</u>	Jan 09, 2004
<u>AB</u>	TEVA	<u>EQ 750MG BASE/VIAL</u>	<u>N64192</u>	<u>002</u>	Apr 16, 1998

CEFUROXIME SODIUM

<u>AB</u>	HANFORD GC	<u>EQ 750MG BASE/VIAL</u>	<u>N64125</u>	<u>001</u>	May 30, 1997
<u>AB</u>	MARSAM PHARMS LLC	<u>EQ 750MG BASE/VIAL</u>	<u>N64035</u>	<u>001</u>	Feb 26, 1993

ZINACEF

<u>AB</u>	+ GLAXOSMITHKLINE	<u>EQ 750MG BASE/VIAL</u>	<u>N50558</u>	<u>002</u>	Oct 19, 1983
-----------	-------------------	---------------------------	---------------	------------	--------------

INJECTABLE; INJECTION

CEFUROXIME

<u>AP</u>	AM PHARM PARTNERS	<u>EQ 1.5GM BASE/VIAL</u>	<u>N65001</u>	<u>002</u>	May 30, 2001
<u>AP</u>	HIKMA FARMACEUTICA	<u>EQ 1.5GM BASE/VIAL</u>	<u>N65048</u>	<u>002</u>	Jan 09, 2004
<u>AP</u>		<u>EQ 7.5GM BASE/VIAL</u>	<u>N65046</u>	<u>001</u>	Jan 09, 2004
<u>AP</u>	TEVA	<u>EQ 1.5GM BASE/VIAL</u>	<u>N64192</u>	<u>001</u>	Apr 16, 1998
<u>AP</u>		<u>EQ 7.5GM BASE/VIAL</u>	<u>N64191</u>	<u>001</u>	Apr 16, 1998

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 77 (of 360)

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER

<u>AP</u>	+	B BRAUN	<u>EQ 750MG BASE/VIAL</u>	<u>N50780</u>	<u>001</u>	Feb 21, 2001
<u>AP</u>	+		<u>EQ 1.5GM BASE/VIAL</u>	<u>N50780</u>	<u>002</u>	Feb 21, 2001

CEFUROXIME SODIUM

<u>AP</u>		AM PHARM PARTNERS	<u>EQ 7.5GM BASE/VIAL</u>	<u>N65002</u>	<u>001</u>	Sep 28, 1998
<u>AP</u>		HANFORD GC	<u>EQ 1.5GM BASE/VIAL</u>	<u>N64125</u>	<u>002</u>	May 30, 1997
<u>AP</u>			<u>EQ 7.5GM BASE/VIAL</u>	<u>N64124</u>	<u>001</u>	May 30, 1997
<u>AP</u>		MARSAM PHARMS LLC	<u>EQ 1.5GM BASE/VIAL</u>	<u>N64035</u>	<u>002</u>	Feb 26, 1993
<u>AP</u>			<u>EQ 7.5GM BASE/VIAL</u>	<u>N64036</u>	<u>001</u>	Feb 26, 1993

ZINACEF

<u>AP</u>	+	GLAXOSMITHKLINE	<u>EQ 1.5GM BASE/VIAL</u>	<u>N50558</u>	<u>003</u>	Oct 19, 1983
<u>AP</u>	+		<u>EQ 7.5GM BASE/VIAL</u>	<u>N50558</u>	<u>004</u>	Oct 23, 1986
		ZINACEF IN PLASTIC CONTAINER				
	+	GLAXOSMITHKLINE	EQ 15MG BASE/ML	N50643	001	Apr 28, 1989
	+		EQ 30MG BASE/ML	N50643	002	Apr 28, 1989

CELECOXIB

CAPSULE; ORAL

CELEBREX

		GD SEARLE LLC	100MG	N20998	001	Dec 31, 1998
			200MG	N20998	002	Dec 31, 1998
	+		400MG	N20998	003	Aug 29, 2002

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

<u>AB</u>		APOTHECON	<u>EQ 250MG BASE</u>	<u>N63063</u>	<u>001</u>	Sep 29, 1989
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N63063</u>	<u>002</u>	Sep 29, 1989
<u>AB</u>		AUROBINDO PHARMA LTD	<u>EQ 250MG BASE</u>	<u>N65253</u>	<u>001</u>	Nov 16, 2005
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N65253</u>	<u>002</u>	Nov 16, 2005
<u>AB</u>		BARR	<u>EQ 250MG BASE</u>	<u>N62773</u>	<u>001</u>	Jun 26, 1987
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N62775</u>	<u>001</u>	Apr 22, 1987
<u>AB</u>		BELCHER	<u>EQ 250MG BASE</u>	<u>N62713</u>	<u>001</u>	Jul 15, 1988
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N62713</u>	<u>002</u>	Jul 15, 1988
<u>AB</u>		CEPH INTL	<u>EQ 250MG BASE</u>	<u>N62118</u>	<u>001</u>	
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N62118</u>	<u>002</u>	
<u>AB</u>		IVAX PHARMS	<u>EQ 250MG BASE</u>	<u>N61969</u>	<u>001</u>	
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N61969</u>	<u>002</u>	
<u>AB</u>		LUPIN	<u>EQ 250MG BASE</u>	<u>N65229</u>	<u>001</u>	Nov 25, 2005
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N65229</u>	<u>002</u>	Nov 25, 2005
<u>AB</u>		ORCHID HLTHCARE	<u>EQ 250MG BASE</u>	<u>N65248</u>	<u>001</u>	Jun 28, 2005
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N65248</u>	<u>002</u>	Jun 28, 2005
<u>AB</u>		RANBAXY	<u>EQ 250MG BASE</u>	<u>N65007</u>	<u>001</u>	Sep 16, 1999
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N65007</u>	<u>002</u>	Sep 16, 1999
<u>AB</u>		STEVENS J	<u>EQ 250MG BASE</u>	<u>N62870</u>	<u>001</u>	Mar 17, 1988
<u>AB</u>		SUN PHARM INDS (IN)	<u>EQ 250MG BASE</u>	<u>N62791</u>	<u>001</u>	Jun 11, 1987
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N62791</u>	<u>002</u>	Jun 11, 1987
<u>AB</u>		TEVA	<u>EQ 250MG BASE</u>	<u>N62702</u>	<u>001</u>	Feb 13, 1987
<u>AB</u>			<u>EQ 250MG BASE</u>	<u>N62760</u>	<u>001</u>	Apr 24, 1987
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N62702</u>	<u>002</u>	Feb 13, 1987
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N62761</u>	<u>001</u>	Apr 24, 1987
<u>AB</u>		YUNG SHIN PHARM	<u>EQ 250MG BASE</u>	<u>N65152</u>	<u>001</u>	Feb 24, 2005
<u>AB</u>			<u>EQ 500MG BASE</u>	<u>N65152</u>	<u>002</u>	Feb 24, 2005
		<u>KEFLEX</u>				
<u>AB</u>		ADVANCIS PHARM	<u>EQ 250MG BASE</u>	<u>N50405</u>	<u>002</u>	
<u>AB</u>	+		<u>EQ 500MG BASE</u>	<u>N50405</u>	<u>003</u>	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 78 (of 360)

CEPHALEXIN

FOR SUSPENSION; ORAL

CEPHALEXIN

<u>AB</u>	BARR	<u>EQ 250MG BASE/5ML</u>	<u>N62777</u>	<u>001</u>	Aug 06, 1987
<u>AB</u>	CEPH INTL	<u>EQ 125MG BASE/5ML</u>	<u>N62117</u>	<u>002</u>	
<u>AB</u>	+	<u>EQ 250MG BASE/5ML</u>	<u>N62117</u>	<u>003</u>	
	+	EQ 100MG BASE/ML	N62117	001	
<u>AB</u>	LUPIN	<u>EQ 125MG BASE/5ML</u>	<u>N65234</u>	<u>001</u>	Aug 17, 2005
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N65234</u>	<u>002</u>	Aug 17, 2005
<u>AB</u>	RANBAXY	<u>EQ 125MG BASE/5ML</u>	<u>N65081</u>	<u>001</u>	Jul 27, 2001
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N65081</u>	<u>002</u>	Jul 27, 2001
<u>AB</u>	TEVA	<u>EQ 125MG BASE/5ML</u>	<u>N62703</u>	<u>001</u>	Feb 13, 1987
<u>AB</u>		<u>EQ 125MG BASE/5ML</u>	<u>N62767</u>	<u>001</u>	Jun 16, 1987
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N62703</u>	<u>002</u>	Feb 13, 1987
<u>AB</u>		<u>EQ 250MG BASE/5ML</u>	<u>N62768</u>	<u>001</u>	Jun 16, 1987

TABLET; ORAL

CEPHALEXIN

	TEVA	EQ 250MG BASE	N63023	001	Jan 12, 1989
	+	EQ 500MG BASE	N63024	001	Jan 12, 1989

TABLET, FOR SUSPENSION; ORAL

PANIXINE DISPERDOSE

	RANBAXY	EQ 125MG BASE	N65100	002	Sep 11, 2003
	+	EQ 250MG BASE	N65100	001	Sep 11, 2003

CEPHRADINE

CAPSULE; ORAL

ANSPOR

<u>AB</u>	GLAXOSMITHKLINE	<u>250MG</u>	<u>N61859</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N61859</u>	<u>002</u>	
	<u>CEPHRADINE</u>				
<u>AB</u>	TEVA	<u>250MG</u>	<u>N62683</u>	<u>001</u>	Jan 09, 1987
<u>AB</u>		<u>500MG</u>	<u>N62683</u>	<u>002</u>	Jan 09, 1987
	<u>VELOSEF</u>				
<u>AB</u>	APOTHECON	<u>250MG</u>	<u>N61764</u>	<u>001</u>	
<u>AB</u>	+	<u>500MG</u>	<u>N61764</u>	<u>002</u>	

FOR SUSPENSION; ORAL

ANSPOR

<u>AB</u>	GLAXOSMITHKLINE	<u>125MG/5ML</u>	<u>N61866</u>	<u>001</u>	
<u>AB</u>		<u>250MG/5ML</u>	<u>N61866</u>	<u>002</u>	
	<u>CEPHRADINE</u>				
<u>AB</u>	TEVA	<u>125MG/5ML</u>	<u>N62693</u>	<u>001</u>	Jan 09, 1987
<u>AB</u>		<u>250MG/5ML</u>	<u>N62693</u>	<u>002</u>	Jan 09, 1987
	<u>VELOSEF '125'</u>				
<u>AB</u>	+	<u>APOTHECON</u>	<u>125MG/5ML</u>	<u>N61763</u>	<u>001</u>
	<u>VELOSEF '250'</u>				
<u>AB</u>	+	<u>APOTHECON</u>	<u>250MG/5ML</u>	<u>N61763</u>	<u>002</u>

CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

ZYRTEC

	+	PFIZER	5MG/5ML	N20346	001	Sep 27, 1996
--	---	--------	---------	--------	-----	--------------

TABLET; ORAL

ZYRTEC

	PFIZER	5MG	N19835	001	Dec 08, 1995
	+	10MG	N19835	002	Dec 08, 1995

TABLET, CHEWABLE; ORAL

ZYRTEC

	PFIZER	5MG	N21621	001	Mar 16, 2004
	+	10MG	N21621	002	Mar 16, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 79 (of 360)

CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL		
Zyrtec-D 12 Hour		
+ PFIZER	5MG;120MG	N21150 001 Aug 10, 2001

CETRORELIX

INJECTABLE; INJECTION		
CETROTIDE		
+ SERONO INC	EQ 0.25MG BASE/ML	N21197 001 Aug 11, 2000
+	EQ 3MG BASE/ML	N21197 002 Aug 11, 2000

CETYL ALCOHOL; COLFOSCERIL PALMITATE; TYLOXAPOL

FOR SUSPENSION; INTRATRACHEAL		
EXOSURF NEONATAL		
+ GLAXOSMITHKLINE	12MG/VIAL;108MG/VIAL;8MG/VIAL	N20044 001 Aug 02, 1990

CEVIMELINE HYDROCHLORIDE

CAPSULE; ORAL		
EVOXAC		
+ DAIICHI	EQ 30MG BASE	N20989 002 Jan 11, 2000

CHLORAMBUCIL

TABLET; ORAL		
LEUKERAN		
+ SMITHKLINE BEECHAM	2MG	N10669 002

CHLORAMPHENICOL

OINTMENT; OPHTHALMIC		
CHLORAMPHENICOL		
+ ALTANA	1%	N60133 001

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION					
<u>CHLORAMPHENICOL SODIUM SUCCINATE</u>					
AP	AM PHARM PARTNERS	<u>EQ 1GM BASE/VIAL</u>	<u>N62365</u>	<u>001</u>	Aug 25, 1982
<u>CHLOROMYCETIN</u>					
AP	+ PARKEDALE	<u>EQ 1GM BASE/VIAL</u>	<u>N50155</u>	<u>001</u>	

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL					
<u>CHLORDIAZEPOXIDE HYDROCHLORIDE</u>					
<u>AB</u>	BARR	<u>5MG</u>	<u>N84768</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>N83116</u>	<u>001</u>	
<u>AB</u>		<u>25MG</u>	<u>N84769</u>	<u>001</u>	
<u>AB</u>	USL PHARMA	<u>10MG</u>	<u>N84623</u>	<u>001</u>	
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>N86383</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>N86294</u>	<u>001</u>	
<u>AB</u>		<u>25MG</u>	<u>N86382</u>	<u>001</u>	
<u>LIBRIUM</u>					
<u>AB</u>	VALEANT PHARM INTL	<u>5MG</u>	<u>N85461</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>N85472</u>	<u>001</u>	
<u>AB</u>	+	<u>25MG</u>	<u>N85475</u>	<u>001</u>	

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL					
<u>CHLORHEXIDINE GLUCONATE</u>					
<u>AT</u>	ALPHARMA US PHARMS	<u>0.12%</u>	<u>N74291</u>	<u>001</u>	Dec 28, 1995
<u>AT</u>	HI TECH PHARMA	<u>0.12%</u>	<u>N74356</u>	<u>001</u>	May 07, 1996
<u>AT</u>	JOHN O BUTLER CO	<u>0.12%</u>	<u>N76434</u>	<u>001</u>	Nov 29, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 80 (of 360)

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

<u>AT</u>	MORTON GROVE	<u>0.12%</u>	<u>N75006</u>	<u>001</u>	Mar 03, 2004
<u>AT</u>	NOVEX	<u>0.12%</u>	<u>N75561</u>	<u>001</u>	Nov 14, 2000
<u>AT</u>	TEVA	<u>0.12%</u>	<u>N74522</u>	<u>001</u>	Dec 15, 1995
<u>PERIDEX</u>					
<u>AT</u>	+ ZILA	<u>0.12%</u>	<u>N19028</u>	<u>001</u>	Aug 13, 1986
<u>PERIOGARD</u>					
<u>AT</u>	COLGATE	<u>0.12%</u>	<u>N73695</u>	<u>001</u>	Jan 14, 1994
TABLET; DENTAL					
PERIOCHIP					
	+ DEXCEL PHARMA	2.5MG	N20774	001	May 15, 1998

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHLOROPROCAINE HYDROCHLORIDE

<u>AP</u>	BEDFORD	<u>2%</u>	<u>N40273</u>	<u>001</u>	Sep 09, 1998
<u>AP</u>		<u>3%</u>	<u>N40273</u>	<u>002</u>	Sep 09, 1998
<u>AP</u>	HOSPIRA	<u>2%</u>	<u>N87447</u>	<u>001</u>	Apr 16, 1982
<u>AP</u>		<u>3%</u>	<u>N87446</u>	<u>001</u>	Apr 16, 1982
<u>NESACAINE</u>					
<u>AP</u>	ASTRAZENECA	<u>2%</u>	<u>N09435</u>	<u>002</u>	
	+	<u>1%</u>	N09435	001	
<u>NESACAINE-MPF</u>					
<u>AP</u>	+ ASTRAZENECA	<u>2%</u>	<u>N09435</u>	<u>006</u>	May 02, 1996
<u>AP</u>	+	<u>3%</u>	<u>N09435</u>	<u>007</u>	May 02, 1996

CHLOROQUINE PHOSPHATE

TABLET; ORAL

ARALEN

<u>AA</u>	+ SANOFI SYNTHELABO	<u>EQ 300MG BASE</u>	<u>N06002</u>	<u>001</u>	
<u>CHLOROQUINE PHOSPHATE</u>					
<u>AA</u>	IMPAX LABS	<u>EQ 150MG BASE</u>	<u>N80880</u>	<u>001</u>	
<u>AA</u>		<u>EQ 300MG BASE</u>	<u>N40516</u>	<u>001</u>	Aug 29, 2003
<u>AA</u>	+ WEST WARD	<u>EQ 150MG BASE</u>	<u>N83082</u>	<u>001</u>	
<u>AA</u>		<u>EQ 300MG BASE</u>	<u>N83082</u>	<u>002</u>	Sep 17, 1999

CHLOROTHIAZIDE

SUSPENSION; ORAL

DIURIL

+ MERCK 250MG/5ML N11870 001

TABLET; ORAL

CHLOROTHIAZIDE

<u>AB</u>	+ MYLAN	<u>250MG</u>	<u>N84388</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N84217</u>	<u>001</u>	
<u>AB</u>	WEST WARD	<u>250MG</u>	<u>N86028</u>	<u>001</u>	Jul 14, 1982
<u>AB</u>		<u>500MG</u>	<u>N87736</u>	<u>001</u>	Jul 14, 1982

CHLOROTHIAZIDE SODIUM

INJECTABLE; INJECTION

DIURIL

+ OVATION PHARMS EQ 500MG BASE/VIAL N11145 005

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

DIUPRES-250

BP MERCK 250MG;0.125MG N11635 003 Aug 26, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 81 (of 360)

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL				
DIUPRES-500				
BP	+ MERCK	500MG;0.125MG	N11635 006	Aug 26, 1987

CHLOROXINE

SHAMPOO; TOPICAL				
CAPITROL				
	+ WESTWOOD SQUIBB	2%	N17594 001	

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL				
CODEPREX				
	+ UCB	EQ 4MG MALEATE/5ML;EQ 20MG BASE/5ML	N21369 001	Jun 21, 2004

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL				
TUSSIONEX				
	+ UCB	EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML	N19111 001	Dec 31, 1987

CHLORPROMAZINE

SUPPOSITORY; RECTAL				
THORAZINE				
	+ GLAXOSMITHKLINE	25MG	N09149 024	
	+	100MG	N09149 033	

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL				
THORAZINE				
	+ GLAXOSMITHKLINE	200MG	N11120 019	
	+	300MG	N11120 020	

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

<u>AA</u>	ALPHARMA US PHARMS	<u>100MG/ML</u>	<u>N86863</u>	<u>001</u>	
<u>AA</u>	PHARM ASSOC	<u>30MG/ML</u>	<u>N40231</u>	<u>001</u>	Dec 30, 1999
<u>AA</u>		<u>100MG/ML</u>	<u>N40224</u>	<u>001</u>	Jan 26, 1999

CHLORPROMAZINE HYDROCHLORIDE INTENSOL

<u>AA</u>	ROXANE	<u>30MG/ML</u>	<u>N88157</u>	<u>001</u>	Apr 27, 1983
<u>AA</u>		<u>100MG/ML</u>	<u>N88158</u>	<u>001</u>	Apr 27, 1983

SONAZINE

<u>AA</u>	SANDOZ	<u>30MG/ML</u>	<u>N80983</u>	<u>004</u>	
<u>AA</u>		<u>100MG/ML</u>	<u>N80983</u>	<u>005</u>	

THORAZINE

<u>AA</u>	+ GLAXOSMITHKLINE	<u>30MG/ML</u>	<u>N09149</u>	<u>032</u>	
<u>AA</u>	+	<u>100MG/ML</u>	<u>N09149</u>	<u>043</u>	

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

<u>AP</u>	BAXTER HLTHCARE	<u>25MG/ML</u>	<u>N83329</u>	<u>001</u>	
-----------	-----------------	----------------	---------------	------------	--

THORAZINE

<u>AP</u>	+ GLAXOSMITHKLINE	<u>25MG/ML</u>	<u>N09149</u>	<u>011</u>	
-----------	-------------------	----------------	---------------	------------	--

SYRUP; ORAL

SONAZINE

<u>AA</u>	SANDOZ	<u>10MG/5ML</u>	<u>N83040</u>	<u>001</u>	
-----------	--------	-----------------	---------------	------------	--

THORAZINE

<u>AA</u>	+ GLAXOSMITHKLINE	<u>10MG/5ML</u>	<u>N09149</u>	<u>022</u>	
-----------	-------------------	-----------------	---------------	------------	--

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

BP	SANDOZ	10MG	N80439	001	
----	--------	------	--------	-----	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 82 (of 360)

CHLORPROMAZINE HYDROCHLORIDE

TABLET; ORAL			
CHLORPROMAZINE HYDROCHLORIDE			
BP	SANDOZ	25MG	N80439 002
BP		50MG	N80439 003
BP	+	100MG	N80439 004
BP		200MG	N80439 005
BP	USL PHARMA	10MG	N83386 001
BP		25MG	N84112 001
BP		50MG	N84113 001
BP		100MG	N84114 001
BP		200MG	N84115 001

CHLORPROPAMIDE

TABLET; ORAL			
CHLORPROPAMIDE			
<u>AB</u>	MYLAN	<u>100MG</u>	<u>N88548 001</u> Jun 01, 1984
<u>AB</u>		<u>250MG</u>	<u>N88549 001</u> Jun 01, 1984
<u>AB</u>	PAR PHARM	<u>100MG</u>	<u>N88175 001</u> Feb 27, 1984
<u>AB</u>		<u>250MG</u>	<u>N88176 001</u> Feb 27, 1984
<u>AB</u>	PLIVA	<u>100MG</u>	<u>N88921 001</u> Apr 12, 1985
<u>AB</u>		<u>250MG</u>	<u>N88922 001</u> Apr 12, 1985
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>N88725 001</u> Aug 31, 1984
<u>AB</u>		<u>250MG</u>	<u>N88726 001</u> Aug 31, 1984
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N88852 001</u> Sep 26, 1984
<u>AB</u>		<u>250MG</u>	<u>N88826 001</u> Sep 26, 1984
DIABINESE			
<u>AB</u>	PFIZER	<u>100MG</u>	<u>N11641 003</u>
<u>AB</u>	+	<u>250MG</u>	<u>N11641 006</u>

CHLORTHALIDONE

TABLET; ORAL			
CHLORTHALIDONE			
<u>AB</u>	MYLAN	<u>25MG</u>	<u>N87180 001</u>
<u>AB</u>	+	<u>50MG</u>	<u>N86831 001</u>
<u>AB</u>	PLIVA	<u>25MG</u>	<u>N88902 001</u> Sep 19, 1985
<u>AB</u>		<u>50MG</u>	<u>N88903 001</u> Sep 19, 1985
	THALITONE		
	+ MONARCH PHARMS	15MG	N19574 001 Dec 20, 1988

CHLORTHALIDONE; CLONIDINE HYDROCHLORIDE

TABLET; ORAL			
CLORPRES			
<u>AB</u>	MYLAN	<u>15MG;0.1MG</u>	<u>N71323 001</u> Feb 09, 1987
<u>AB</u>		<u>15MG;0.2MG</u>	<u>N71324 001</u> Feb 09, 1987
	+	15MG;0.3MG	N71325 001 Feb 09, 1987

CHLORZOXAZONE

TABLET; ORAL			
CHLORZOXAZONE			
<u>AA</u>	AMIDE PHARM	<u>250MG</u>	<u>N88928 001</u> May 08, 1987
<u>AA</u>		<u>500MG</u>	<u>N40113 001</u> Sep 29, 1995
<u>AA</u>	BARR	<u>500MG</u>	<u>N89895 001</u> May 04, 1988
<u>AA</u>	MUTUAL PHARM	<u>500MG</u>	<u>N89970 001</u> Sep 27, 1990
<u>AA</u>	OHM LABS	<u>250MG</u>	<u>N81298 001</u> Dec 29, 1993
<u>AA</u>		<u>500MG</u>	<u>N81299 001</u> Dec 29, 1993
<u>AA</u>	PAR PHARM	<u>250MG</u>	<u>N87981 001</u> Sep 20, 1983
<u>AA</u>	SANDOZ	<u>250MG</u>	<u>N89852 001</u> May 04, 1988
<u>AA</u>		<u>500MG</u>	<u>N89853 001</u> May 04, 1988
<u>AA</u>	TEVA	<u>500MG</u>	<u>N89859 001</u> May 04, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 83 (of 360)

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

<u>AA</u>	WATSON LABS	<u>500MG</u>	<u>N40137</u>	<u>001</u>	Aug 09, 1996
<u>AA</u>		<u>500MG</u>	<u>N81040</u>	<u>001</u>	Aug 22, 1989
	<u>PARAFON FORTE DSC</u>				
<u>AA</u>	+ ORTHO MCNEIL PHARM	<u>500MG</u>	<u>N11529</u>	<u>002</u>	Jun 15, 1987
	<u>STRIFON FORTE DSC</u>				
<u>AA</u>	FERNDAL LABS	<u>500MG</u>	<u>N81008</u>	<u>001</u>	Dec 23, 1988

CHOLESTYRAMINE

POWDER; ORAL

CHOLESTYRAMINE

<u>AB</u>	PAR PHARM	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N77204</u>	<u>002</u>	Aug 26, 2005
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>N77204</u>	<u>001</u>	Aug 26, 2005
<u>AB</u>	SANDOZ	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N74557</u>	<u>002</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>N74557</u>	<u>001</u>	Aug 15, 1996
<u>AB</u>	TEVA	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N74347</u>	<u>002</u>	May 28, 1998
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>N74347</u>	<u>001</u>	May 28, 1998
<u>AB</u>	TEVA PHARMS	<u>EQ 4GM RESIN/PACKET</u>	<u>N74554</u>	<u>001</u>	Oct 02, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N74554</u>	<u>002</u>	Oct 02, 1996
	<u>CHOLESTYRAMINE LIGHT</u>				
<u>AB</u>	PAR PHARM	<u>EQ 4GM RESIN/PACKET</u>	<u>N77203</u>	<u>001</u>	Aug 26, 2005
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N77203</u>	<u>002</u>	Aug 26, 2005
<u>AB</u>	SANDOZ	<u>EQ 4GM RESIN/PACKET</u>	<u>N74558</u>	<u>001</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N74558</u>	<u>002</u>	Aug 15, 1996
<u>AB</u>	TEVA	<u>EQ 4GM RESIN/PACKET</u>	<u>N74348</u>	<u>001</u>	May 28, 1998
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N74348</u>	<u>002</u>	May 28, 1998
<u>AB</u>	TEVA PHARMS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N74555</u>	<u>002</u>	Sep 30, 1998
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>N74555</u>	<u>001</u>	Sep 30, 1998
	<u>LOCHOLEST</u>				
<u>AB</u>	SANDOZ	<u>EQ 4GM RESIN/PACKET</u>	<u>N74561</u>	<u>001</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N74561</u>	<u>002</u>	Aug 15, 1996
	<u>LOCHOLEST LIGHT</u>				
<u>AB</u>	SANDOZ	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N74562</u>	<u>002</u>	Aug 15, 1996
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>N74562</u>	<u>001</u>	Aug 15, 1996
	<u>PREVALITE</u>				
<u>AB</u>	UPSHER SMITH	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N73263</u>	<u>002</u>	Oct 30, 1997
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>N73263</u>	<u>001</u>	Feb 22, 1996
	<u>QUESTRAN</u>				
<u>AB</u>	BRISTOL MYERS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N16640</u>	<u>003</u>	
<u>AB</u>	+	<u>EQ 4GM RESIN/PACKET</u>	<u>N16640</u>	<u>001</u>	
	<u>QUESTRAN LIGHT</u>				
<u>AB</u>	BRISTOL MYERS	<u>EQ 4GM RESIN/SCOOPFUL</u>	<u>N19669</u>	<u>003</u>	Dec 05, 1988
<u>AB</u>		<u>EQ 4GM RESIN/PACKET</u>	<u>N19669</u>	<u>001</u>	Dec 05, 1988

CHORIOGONADOTROPIN ALFA

INJECTABLE; SUBCUTANEOUS

OVIDREL

+	SERONO INC	EQ 0.25MG /0.5ML	N21149	002	Oct 06, 2003
---	------------	------------------	--------	-----	--------------

CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE IN PLASTIC CONTAINER

+	HOSPIRA	EQ 0.004MG CHROMIUM/ML	N18961	001	Jun 26, 1986
---	---------	------------------------	--------	-----	--------------

CHROMIC PHOSPHATE, P-32

INJECTABLE; INJECTION

PHOSPHOCOL P32

+	MALLINCKRODT	5mCi/ML	N17084	001	
---	--------------	---------	--------	-----	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 84 (of 360)

CICLOPIROX

CREAM; TOPICAL

CICLOPIROX

<u>AB</u>	ALTANA	<u>0.77%</u>	<u>N76435</u>	<u>001</u>	Dec 29, 2004
<u>AB</u>	TARO	<u>0.77%</u>	<u>N76790</u>	<u>001</u>	Apr 12, 2005

LOPROX

<u>AB</u>	+ MEDICIS	<u>0.77%</u>	<u>N18748</u>	<u>001</u>	Dec 30, 1982
-----------	-----------	--------------	---------------	------------	--------------

GEL; TOPICAL

LOPROX

	+ MEDICIS	0.77%	N20519	001	Jul 21, 1997
--	-----------	-------	--------	-----	--------------

SHAMPOO; TOPICAL

LOPROX

	+ MEDICIS	1%	N21159	001	Feb 28, 2003
--	-----------	----	--------	-----	--------------

SOLUTION; TOPICAL

PENLAC

	+ DERMIK LABS	8%	N21022	001	Dec 17, 1999
--	---------------	----	--------	-----	--------------

SUSPENSION; TOPICAL

CICLOPIROX

<u>AB</u>	ALTANA	<u>0.77%</u>	<u>N76422</u>	<u>001</u>	Aug 06, 2004
<u>AB</u>	TARO	<u>0.77%</u>	<u>N77092</u>	<u>001</u>	Aug 10, 2005

LOPROX

<u>AB</u>	+ MEDICIS	<u>0.77%</u>	<u>N19824</u>	<u>001</u>	Dec 30, 1988
-----------	-----------	--------------	---------------	------------	--------------

CIDOFVIR

INJECTABLE; INJECTION

VISTIDE

	+ GILEAD	EQ 75MG BASE/ML	N20638	001	Jun 26, 1996
--	----------	-----------------	--------	-----	--------------

CILASTATIN SODIUM; IMIPENEM

INJECTABLE; INJECTION

PRIMAXIN

	+ MERCK	EQ 250MG BASE/VIAL;250MG/VIAL	N50587	001	Nov 26, 1985
		EQ 250MG BASE/VIAL;250MG/VIAL	N62756	001	Jan 08, 1987
	+	EQ 500MG BASE/VIAL;500MG/VIAL	N50587	002	Nov 26, 1985
		EQ 500MG BASE/VIAL;500MG/VIAL	N50630	001	Dec 14, 1990
		EQ 500MG BASE/VIAL;500MG/VIAL	N62756	002	Jan 08, 1987
	+	EQ 750MG BASE/VIAL;750MG/VIAL	N50630	002	Dec 14, 1990

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

<u>AB</u>	ALPHAPHARM	<u>50MG</u>	<u>N77019</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>		<u>100MG</u>	<u>N77019</u>	<u>002</u>	Nov 23, 2004
<u>AB</u>	AMIDE PHARM	<u>100MG</u>	<u>N77028</u>	<u>002</u>	Nov 26, 2004
<u>AB</u>	APOTEX	<u>50MG</u>	<u>N77030</u>	<u>001</u>	Dec 10, 2004
<u>AB</u>		<u>100MG</u>	<u>N77030</u>	<u>002</u>	Dec 10, 2004
<u>AB</u>	COREPHARMA	<u>50MG</u>	<u>N77150</u>	<u>001</u>	Mar 11, 2005
<u>AB</u>		<u>100MG</u>	<u>N77022</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>	IVAX PHARMS	<u>100MG</u>	<u>N77020</u>	<u>002</u>	Mar 01, 2005
<u>AB</u>	ROXANE	<u>50MG</u>	<u>N77024</u>	<u>001</u>	May 17, 2005
<u>AB</u>		<u>100MG</u>	<u>N77024</u>	<u>002</u>	May 17, 2005
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N77310</u>	<u>001</u>	Nov 08, 2005
<u>AB</u>		<u>100MG</u>	<u>N77021</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>	TEVA	<u>50MG</u>	<u>N77027</u>	<u>001</u>	Nov 24, 2004
<u>AB</u>		<u>100MG</u>	<u>N77027</u>	<u>002</u>	Nov 24, 2004

PLETAL

<u>AB</u>	+ OTSUKA	<u>50MG</u>	<u>N20863</u>	<u>001</u>	Jan 15, 1999
<u>AB</u>	+	<u>100MG</u>	<u>N20863</u>	<u>002</u>	Jan 15, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 85 (of 360)

CIMETIDINE

TABLET; ORAL

CIMETIDINE

<u>AB</u>	<u>CLONMEL HLTHCARE</u>	<u>300MG</u>	<u>N74340</u>	<u>001</u>	Jun 23, 1995
<u>AB</u>		<u>400MG</u>	<u>N74340</u>	<u>002</u>	Jun 23, 1995
<u>AB</u>		<u>800MG</u>	<u>N74339</u>	<u>001</u>	Jun 23, 1995
<u>AB</u>	<u>ENDO PHARMS</u>	<u>200MG</u>	<u>N74281</u>	<u>001</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>N74281</u>	<u>002</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>N74281</u>	<u>003</u>	May 17, 1994
<u>AB</u>		<u>800MG</u>	<u>N74329</u>	<u>001</u>	May 17, 1994
<u>AB</u>	<u>IVAX PHARMS</u>	<u>200MG</u>	<u>N74424</u>	<u>001</u>	Jul 28, 1995
<u>AB</u>		<u>300MG</u>	<u>N74424</u>	<u>002</u>	Jul 28, 1995
<u>AB</u>		<u>400MG</u>	<u>N74424</u>	<u>003</u>	Jul 28, 1995
<u>AB</u>		<u>800MG</u>	<u>N74402</u>	<u>001</u>	May 30, 1995
<u>AB</u>		<u>800MG</u>	<u>N74424</u>	<u>004</u>	Jul 28, 1995
<u>AB</u>	<u>LEK PHARMS</u>	<u>300MG</u>	<u>N74250</u>	<u>002</u>	Jun 29, 1995
<u>AB</u>		<u>400MG</u>	<u>N74250</u>	<u>003</u>	Jun 29, 1995
<u>AB</u>		<u>800MG</u>	<u>N74250</u>	<u>004</u>	Jun 29, 1995
<u>AB</u>	<u>MYLAN</u>	<u>200MG</u>	<u>N74246</u>	<u>001</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>N74246</u>	<u>002</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>N74246</u>	<u>003</u>	May 17, 1994
<u>AB</u>		<u>800MG</u>	<u>N74246</u>	<u>004</u>	May 17, 1994
<u>AB</u>	<u>PLIVA</u>	<u>200MG</u>	<u>N74568</u>	<u>001</u>	Feb 27, 1997
<u>AB</u>		<u>300MG</u>	<u>N74568</u>	<u>002</u>	Feb 27, 1997
<u>AB</u>		<u>400MG</u>	<u>N74568</u>	<u>003</u>	Feb 27, 1997
<u>AB</u>		<u>800MG</u>	<u>N74566</u>	<u>001</u>	Feb 27, 1997
<u>AB</u>	<u>SANDOZ</u>	<u>200MG</u>	<u>N74100</u>	<u>001</u>	Jan 31, 1995
<u>AB</u>		<u>200MG</u>	<u>N74506</u>	<u>001</u>	Jan 24, 1996
<u>AB</u>		<u>300MG</u>	<u>N74100</u>	<u>002</u>	Jan 31, 1995
<u>AB</u>		<u>300MG</u>	<u>N74506</u>	<u>002</u>	Jan 24, 1996
<u>AB</u>		<u>400MG</u>	<u>N74100</u>	<u>003</u>	Jan 31, 1995
<u>AB</u>		<u>400MG</u>	<u>N74506</u>	<u>003</u>	Jan 24, 1996
<u>AB</u>		<u>800MG</u>	<u>N74100</u>	<u>004</u>	Jan 31, 1995
<u>AB</u>		<u>800MG</u>	<u>N74506</u>	<u>004</u>	Jan 24, 1996
<u>AB</u>	<u>TEVA</u>	<u>200MG</u>	<u>N74151</u>	<u>001</u>	May 17, 1994
<u>AB</u>		<u>200MG</u>	<u>N74365</u>	<u>001</u>	Feb 28, 1995
<u>AB</u>		<u>300MG</u>	<u>N74151</u>	<u>002</u>	May 17, 1994
<u>AB</u>		<u>300MG</u>	<u>N74365</u>	<u>002</u>	Feb 28, 1995
<u>AB</u>		<u>400MG</u>	<u>N74151</u>	<u>003</u>	May 17, 1994
<u>AB</u>		<u>400MG</u>	<u>N74365</u>	<u>003</u>	Feb 28, 1995
<u>AB</u>		<u>800MG</u>	<u>N74365</u>	<u>004</u>	Feb 28, 1995
<u>AB</u>		<u>800MG</u>	<u>N74463</u>	<u>001</u>	May 17, 1994
<u>AB</u>	<u>TORPHARM</u>	<u>200MG</u>	<u>N74890</u>	<u>001</u>	Dec 18, 1998
<u>AB</u>		<u>300MG</u>	<u>N74890</u>	<u>002</u>	Dec 18, 1998
<u>AB</u>		<u>400MG</u>	<u>N74890</u>	<u>003</u>	Dec 18, 1998
<u>AB</u>		<u>800MG</u>	<u>N74890</u>	<u>004</u>	Dec 18, 1998
<u>AB</u>	<u>WATSON LABS</u>	<u>200MG</u>	<u>N74349</u>	<u>001</u>	Aug 30, 1996
<u>AB</u>		<u>300MG</u>	<u>N74349</u>	<u>002</u>	Aug 30, 1996
<u>AB</u>		<u>400MG</u>	<u>N74349</u>	<u>003</u>	Aug 30, 1996
<u>AB</u>		<u>800MG</u>	<u>N74316</u>	<u>001</u>	Feb 28, 1996
<u>TAGAMET</u>					
<u>AB</u>	<u>GLAXOSMITHKLINE</u>	<u>200MG</u>	<u>N17920</u>	<u>002</u>	
<u>AB</u>		<u>300MG</u>	<u>N17920</u>	<u>003</u>	
<u>AB</u>		<u>400MG</u>	<u>N17920</u>	<u>004</u>	Dec 14, 1983
<u>AB</u>	<u>+</u>	<u>800MG</u>	<u>N17920</u>	<u>005</u>	Apr 30, 1986

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

<u>AP</u>	<u>CLONMEL HLTHCARE</u>	<u>EQ 300MG BASE/2ML</u>	<u>N74428</u>	<u>001</u>	Apr 25, 1996
-----------	-------------------------	--------------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 86 (of 360)

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

<u>AP</u>	ENDO PHARMS	<u>EQ 300MG BASE/2ML</u>	<u>N74005</u>	<u>001</u>	Aug 31, 1994
<u>AP</u>	HOSPIRA	<u>EQ 300MG BASE/2ML</u>	<u>N74344</u>	<u>001</u>	Jan 31, 1995
<u>AP</u>		<u>EQ 300MG BASE/2ML</u>	<u>N74345</u>	<u>001</u>	Jan 31, 1995
<u>AP</u>	SICOR PHARMS	<u>EQ 300MG BASE/2ML</u>	<u>N74252</u>	<u>001</u>	Nov 26, 1997

CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+	HOSPIRA	EQ 6MG BASE/ML	N74269	001	Dec 27, 1994
---	---------	----------------	--------	-----	--------------

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

<u>AA</u>	+	ALPHARMA US PHARMS	<u>EQ 300MG BASE/5ML</u>	<u>N74176</u>	<u>001</u>	Jun 01, 1994
<u>AA</u>		DURAMED PHARMS BARR	<u>EQ 300MG BASE/5ML</u>	<u>N75110</u>	<u>001</u>	Jun 18, 1998
<u>AA</u>		ENDO PHARMS	<u>EQ 300MG BASE/5ML</u>	<u>N74251</u>	<u>001</u>	Dec 22, 1994
<u>AA</u>		HI TECH PHARMA	<u>EQ 300MG BASE/5ML</u>	<u>N74664</u>	<u>001</u>	Oct 28, 1997
<u>AA</u>		MORTON GROVE	<u>EQ 300MG BASE/5ML</u>	<u>N74757</u>	<u>001</u>	Oct 17, 1997
<u>AA</u>		NOVEX	<u>EQ 300MG BASE/5ML</u>	<u>N75560</u>	<u>001</u>	Mar 15, 2000
<u>AA</u>		PHARM ASSOC	<u>EQ 300MG BASE/5ML</u>	<u>N74553</u>	<u>001</u>	Jan 27, 1997
<u>AA</u>		TEVA	<u>EQ 300MG BASE/5ML</u>	<u>N74610</u>	<u>001</u>	Sep 26, 1996
<u>AA</u>		TEVA PHARMS	<u>EQ 300MG BASE/5ML</u>	<u>N74859</u>	<u>001</u>	Jul 09, 1998

CINACALCET HYDROCHLORIDE

TABLET; ORAL

SENSIPAR

AMGEN

		EQ 30MG BASE	N21688	001	Mar 08, 2004
		EQ 60MG BASE	N21688	002	Mar 08, 2004
+		EQ 90MG BASE	N21688	003	Mar 08, 2004

CIPROFLOXACIN

FOR SUSPENSION; ORAL

CIPRO

BAYER PHARMS

		250MG/5ML	N20780	001	Sep 26, 1997
+		500MG/5ML	N20780	002	Sep 26, 1997

INJECTABLE; INJECTION

CIPRO

+	BAYER PHARMS	10MG/ML	N19847	001	Dec 26, 1990
	CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER				
+	BAYER PHARMS	200MG/100ML	N19857	001	Dec 26, 1990

CIPROFLOXACIN HYDROCHLORIDE

OINTMENT; OPHTHALMIC

CILOXAN

+	ALCON	EQ 0.3% BASE	N20369	001	Mar 30, 1998
---	-------	--------------	--------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

CILOXAN

<u>AT</u>	+	ALCON	<u>EQ 0.3% BASE</u>	<u>N19992</u>	<u>001</u>	Dec 31, 1990
-----------	---	-------	---------------------	---------------	------------	--------------

CIPROFLOXACIN

<u>AT</u>		BAUSCH AND LOMB	<u>EQ 0.3% BASE</u>	<u>N76754</u>	<u>001</u>	Jun 09, 2004
<u>AT</u>		HITECH PHARMA	<u>EQ 0.3% BASE</u>	<u>N76673</u>	<u>001</u>	Jan 21, 2005
<u>AT</u>		NOVEX	<u>EQ 0.3% BASE</u>	<u>N75928</u>	<u>001</u>	Jun 09, 2004

TABLET; ORAL

CIPRO

BAYER PHARMS

<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N19537</u>	<u>001</u>	Apr 08, 1996
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N19537</u>	<u>002</u>	Oct 22, 1987
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N19537</u>	<u>003</u>	Oct 22, 1987
<u>AB</u>	+	<u>EQ 750MG BASE</u>	<u>N19537</u>	<u>004</u>	Oct 22, 1987

CIPROFLOXACIN

BARR

<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N74124</u>	<u>001</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74124</u>	<u>002</u>	Jun 09, 2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 87 (of 360)

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPROFLOXACIN

<u>AB</u>	BARR	<u>EQ 750MG BASE</u>	<u>N74124</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>	CARLSBAD	<u>EQ 250MG BASE</u>	<u>N76126</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76126</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76126</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>	COBALT	<u>EQ 100MG BASE</u>	<u>N76794</u>	<u>001</u>	Feb 10, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N76794</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76794</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76794</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 100MG BASE</u>	<u>N75593</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N75593</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N75593</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N75593</u>	<u>001</u>	Jun 09, 2004
<u>AB</u>	GENPHARM	<u>EQ 250MG BASE</u>	<u>N75817</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N75817</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N75817</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>	HIKMA	<u>EQ 250MG BASE</u>	<u>N76558</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76558</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76558</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>	IVAX PHARMS	<u>EQ 250MG BASE</u>	<u>N76089</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76089</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76089</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>	MARTEC	<u>EQ 250MG BASE</u>	<u>N76138</u>	<u>001</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76138</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76138</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>	MYLAN	<u>EQ 250MG BASE</u>	<u>N75685</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N75685</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N75685</u>	<u>001</u>	Jun 09, 2004
<u>AB</u>	PLIVA	<u>EQ 100MG BASE</u>	<u>N76426</u>	<u>001</u>	Jun 15, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N76426</u>	<u>002</u>	Jun 15, 2005
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76426</u>	<u>003</u>	Jun 15, 2005
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76426</u>	<u>004</u>	Jun 15, 2005
<u>AB</u>	RANBAXY	<u>EQ 250MG BASE</u>	<u>N75747</u>	<u>001</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N75747</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N75747</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>	SANDOZ	<u>EQ 100MG BASE</u>	<u>N75939</u>	<u>001</u>	Mar 03, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N75939</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N76593</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N75939</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76593</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N75939</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76593</u>	<u>004</u>	Jun 09, 2004
<u>AB</u>	TARO	<u>EQ 100MG BASE</u>	<u>N76912</u>	<u>001</u>	Feb 18, 2005
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N76912</u>	<u>002</u>	Oct 06, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76912</u>	<u>003</u>	Oct 06, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76912</u>	<u>004</u>	Oct 06, 2004
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>N76136</u>	<u>001</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76136</u>	<u>002</u>	Jun 09, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76136</u>	<u>003</u>	Jun 09, 2004
<u>AB</u>	TORPHARM	<u>EQ 250MG BASE</u>	<u>N76896</u>	<u>001</u>	Nov 04, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76896</u>	<u>002</u>	Nov 04, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76896</u>	<u>003</u>	Nov 04, 2004
<u>AB</u>	UNIQUE PHARM LABS	<u>EQ 250MG BASE</u>	<u>N76639</u>	<u>001</u>	Sep 10, 2004
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N76639</u>	<u>002</u>	Sep 10, 2004
<u>AB</u>		<u>EQ 750MG BASE</u>	<u>N76639</u>	<u>003</u>	Sep 10, 2004

TABLET, EXTENDED RELEASE; ORAL

PROQUIN XR

+ ESPRIT PHARMA

EQ 500MG BASE

N21744 001

May 19, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 88 (of 360)

CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE

SUSPENSION/DROPS; OTIC

CIPRO HC

+ ALCON	EQ 0.2% BASE;1%	N20805	001	Feb 10, 1998
---------	-----------------	--------	-----	--------------

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPRO XR

+ BAYER PHARMS	212.6MG;EQ 287.5MG BASE	N21473	001	Dec 13, 2002
+	425.2MG;EQ 574.9MG BASE	N21473	002	Aug 28, 2003

CIPROFLOXACIN; DEXAMETHASONE

SUSPENSION/DROPS; OTIC

CIPRODEX

+ ALCON	0.3%;0.1%	N21537	001	Jul 18, 2003
---------	-----------	--------	-----	--------------

CISATRACURIUM BESYLATE

INJECTABLE; INJECTION

NIMBEX

+ ABBOTT	EQ 2MG BASE/ML	N20551	001	Dec 15, 1995
----------	----------------	--------	-----	--------------

NIMBEX PRESERVATIVE FREE

+ ABBOTT	EQ 2MG BASE/ML	N20551	003	Dec 15, 1995
----------	----------------	--------	-----	--------------

+	EQ 10MG BASE/ML	N20551	002	Dec 15, 1995
---	-----------------	--------	-----	--------------

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

<u>AP</u>	AM PHARM PARTNERS	<u>1MG/ML</u>	<u>N74735</u>	001	Jul 16, 1999
<u>AP</u>	BEDFORD	<u>1MG/ML</u>	<u>N75036</u>	001	Nov 07, 2000
<u>AP</u>		<u>10MG/VIAL</u>	<u>N74713</u>	001	Nov 14, 2000
<u>AP</u>		<u>50MG/VIAL</u>	<u>N74713</u>	002	Nov 14, 2000
<u>AP</u>	PHARMACHEMIE	<u>1MG/ML</u>	<u>N74656</u>	001	May 16, 2000
<u>AP</u>	SICOR PHARMS	<u>1MG/ML</u>	<u>N74814</u>	001	May 16, 2000

PLATINOL

<u>AP</u>	+ BRISTOL MYERS	<u>10MG/VIAL</u>	<u>N18057</u>	001	
<u>AP</u>	+	<u>50MG/VIAL</u>	<u>N18057</u>	002	

PLATINOL-AQ

<u>AP</u>	+ BRISTOL MYERS	<u>1MG/ML</u>	<u>N18057</u>	004	Nov 08, 1988
-----------	-----------------	---------------	---------------	-----	--------------

CITALOPRAM HYDROBROMIDE

SOLUTION; ORAL

CELEXA

<u>AA</u>	+ FOREST LABS	<u>EQ 10MG BASE/5ML</u>	<u>N21046</u>	001	Dec 22, 1999
-----------	---------------	-------------------------	---------------	-----	--------------

CITALOPRAM HYDROBROMIDE

<u>AA</u>	APOTEX	<u>EQ 10MG BASE/5ML</u>	<u>N77601</u>	001	Nov 15, 2005
<u>AA</u>	ROXANE	<u>EQ 10MG BASE/5ML</u>	<u>N77043</u>	001	Dec 13, 2004

TABLET; ORAL

CELEXA

<u>AB</u>	FOREST LABS	<u>EQ 10MG BASE</u>	<u>N20822</u>	001	Apr 27, 2000
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N20822</u>	002	Jul 17, 1998
<u>AB</u>	+	<u>EQ 40MG BASE</u>	<u>N20822</u>	003	Jul 17, 1998

CITALOPRAM HYDROBROMIDE

<u>AB</u>	AKYMA PHARMS	<u>EQ 10MG BASE</u>	<u>N77045</u>	003	Apr 29, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77045</u>	002	Apr 29, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77045</u>	001	Apr 29, 2005
<u>AB</u>	ALPHAPHARM	<u>EQ 10MG BASE</u>	<u>N77037</u>	001	Nov 05, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77037</u>	002	Nov 05, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77037</u>	003	Nov 05, 2004
<u>AB</u>	APOTEX	<u>EQ 10MG BASE</u>	<u>N77046</u>	001	Nov 24, 2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 89 (of 360)

CITALOPRAM HYDROBROMIDE

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

<u>AB</u>	APOTEX	<u>EQ 20MG BASE</u>	<u>N77046</u>	<u>002</u>	Nov 24, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77046</u>	<u>003</u>	Nov 24, 2004
<u>AB</u>	AUROBINDO	<u>EQ 10MG BASE</u>	<u>N77031</u>	<u>001</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77031</u>	<u>002</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77031</u>	<u>003</u>	Oct 28, 2004
<u>AB</u>	CARACO	<u>EQ 10MG BASE</u>	<u>N77032</u>	<u>001</u>	Nov 12, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77032</u>	<u>002</u>	Nov 12, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77032</u>	<u>003</u>	Nov 12, 2004
<u>AB</u>	COBALT	<u>EQ 10MG BASE</u>	<u>N77034</u>	<u>001</u>	Jun 30, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77034</u>	<u>002</u>	Jun 30, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77034</u>	<u>003</u>	Jun 30, 2005
<u>AB</u>	COREPHARMA	<u>EQ 10MG BASE</u>	<u>N77036</u>	<u>001</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77036</u>	<u>002</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77036</u>	<u>003</u>	Oct 28, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 10MG BASE</u>	<u>N77038</u>	<u>001</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77038</u>	<u>002</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77038</u>	<u>003</u>	Oct 28, 2004
<u>AB</u>	IVAX PHARMS	<u>EQ 10MG BASE</u>	<u>N77048</u>	<u>001</u>	Nov 16, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77048</u>	<u>002</u>	Nov 16, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77048</u>	<u>003</u>	Nov 16, 2004
<u>AB</u>	KALI LABS	<u>EQ 10MG BASE</u>	<u>N77042</u>	<u>001</u>	Nov 05, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77042</u>	<u>002</u>	Nov 05, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77042</u>	<u>003</u>	Nov 05, 2004
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>N77039</u>	<u>001</u>	Feb 03, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77039</u>	<u>002</u>	Feb 03, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77039</u>	<u>003</u>	Feb 03, 2005
<u>AB</u>	PLIVA	<u>EQ 10MG BASE</u>	<u>N77232</u>	<u>001</u>	Oct 31, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77232</u>	<u>002</u>	Oct 31, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77232</u>	<u>003</u>	Oct 31, 2005
<u>AB</u>	PUREPAC PHARM	<u>EQ 10MG BASE</u>	<u>N77033</u>	<u>001</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77033</u>	<u>002</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77033</u>	<u>003</u>	Oct 28, 2004
<u>AB</u>	ROXANE	<u>EQ 10MG BASE</u>	<u>N77041</u>	<u>001</u>	Nov 23, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77041</u>	<u>002</u>	Nov 23, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77041</u>	<u>003</u>	Nov 23, 2004
<u>AB</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>N77035</u>	<u>001</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N77040</u>	<u>001</u>	Aug 17, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77035</u>	<u>002</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77040</u>	<u>002</u>	Aug 17, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77035</u>	<u>003</u>	Oct 28, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77040</u>	<u>003</u>	Aug 17, 2005
<u>AB</u>	WATSON LABS	<u>EQ 10MG BASE</u>	<u>N77044</u>	<u>001</u>	Nov 05, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N77044</u>	<u>002</u>	Nov 05, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N77044</u>	<u>003</u>	Nov 05, 2004

TABLET, ORALLY DISINTEGRATING; ORAL

CITALOPRAM HYDROBROMIDE

BIOVAIL LABS INTL

EQ 10MG BASE

N21763 001

Dec 20, 2005

EQ 20MG BASE

N21763 002

Dec 20, 2005

+

EQ 40MG BASE

N21763 003

Dec 20, 2005

CITRIC ACID; GLUCONOLACTONE; MAGNESIUM CARBONATE

SOLUTION; IRRIGATION

RENACIDIN

+ UNITED GUARDIAN

6.602GM/100ML;198MG/100ML;3.177GM/100ML

N19481 001

Oct 02, 1990

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 90 (of 360)

CITRIC ACID; MAGNESIUM OXIDE; SODIUM CARBONATE

SOLUTION; IRRIGATION

IRRIGATING SOLUTION G IN PLASTIC CONTAINER

<u>AT</u>	+	BAXTER HLTHCARE	<u>3.24GM/100ML;380MG/100ML;430MG/100ML</u>	<u>N18519</u>	<u>001</u>	Jun 22, 1982
-----------	---	-----------------	---	---------------	------------	--------------

UROLOGIC G IN PLASTIC CONTAINER

<u>AT</u>	+	HOSPIRA	<u>3.24GM/100ML;380MG/100ML;430MG/100ML</u>	<u>N18904</u>	<u>001</u>	May 27, 1983
-----------	---	---------	---	---------------	------------	--------------

CLADRIBINE

INJECTABLE; INJECTION

CLADRIBINE

<u>AP</u>		AM PHARM	<u>1MG/ML</u>	<u>N76571</u>	<u>001</u>	Apr 22, 2004
-----------	--	----------	---------------	---------------	------------	--------------

<u>AP</u>		BEDFORD	<u>1MG/ML</u>	<u>N75405</u>	<u>001</u>	Feb 28, 2000
-----------	--	---------	---------------	---------------	------------	--------------

LEUSTATIN

<u>AP</u>	+	ORTHO BIOTECH	<u>1MG/ML</u>	<u>N20229</u>	<u>001</u>	Feb 26, 1993
-----------	---	---------------	---------------	---------------	------------	--------------

CLARITHROMYCIN

FOR SUSPENSION; ORAL

BIAXIN

		ABBOTT	125MG/5ML	N50698	001	Dec 23, 1993
--	--	--------	-----------	--------	-----	--------------

+			250MG/5ML	N50698	002	Dec 23, 1993
---	--	--	-----------	--------	-----	--------------

TABLET; ORAL

BIAXIN

<u>AB</u>	+	ABBOTT	<u>250MG</u>	<u>N50662</u>	<u>001</u>	Oct 31, 1991
-----------	---	--------	--------------	---------------	------------	--------------

<u>AB</u>	+		<u>500MG</u>	<u>N50662</u>	<u>002</u>	Oct 31, 1991
-----------	---	--	--------------	---------------	------------	--------------

CLARITHROMYCIN

<u>AB</u>		GENPHARM	<u>250MG</u>	<u>N65195</u>	<u>001</u>	Mar 11, 2005
-----------	--	----------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N65195</u>	<u>002</u>	Mar 11, 2005
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		IVAX PHARMS	<u>250MG</u>	<u>N65137</u>	<u>001</u>	May 31, 2005
-----------	--	-------------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N65137</u>	<u>002</u>	May 31, 2005
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		RANBAXY	<u>250MG</u>	<u>N65174</u>	<u>001</u>	Sep 24, 2004
-----------	--	---------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N65174</u>	<u>002</u>	Sep 24, 2004
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		ROXANE	<u>250MG</u>	<u>N65178</u>	<u>002</u>	May 25, 2004
-----------	--	--------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N65178</u>	<u>001</u>	May 25, 2004
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>250MG</u>	<u>N65144</u>	<u>001</u>	Oct 18, 2005
-----------	--	--------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N65136</u>	<u>001</u>	Aug 25, 2005
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		TEVA	<u>250MG</u>	<u>N65155</u>	<u>001</u>	May 31, 2005
-----------	--	------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N65155</u>	<u>002</u>	May 31, 2005
-----------	--	--	--------------	---------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

BIAXIN XL

<u>AB</u>	+	ABBOTT	<u>500MG</u>	<u>N50775</u>	<u>001</u>	Mar 03, 2000
-----------	---	--------	--------------	---------------	------------	--------------

CLARITHROMYCIN

<u>AB</u>		ANDRX PHARMS	<u>500MG</u>	<u>N65145</u>	<u>001</u>	Jun 24, 2004
-----------	--	--------------	--------------	---------------	------------	--------------

		RANBAXY	1GM	N65210	001	Jan 26, 2005
--	--	---------	-----	--------	-----	--------------

<u>AB</u>		TEVA	<u>500MG</u>	<u>N65154</u>	<u>001</u>	May 18, 2005
-----------	--	------	--------------	---------------	------------	--------------

CLARITHROMYCIN EXTENDED RELEASE

<u>AB</u>		SANDOZ	<u>500MG</u>	<u>N65250</u>	<u>001</u>	Aug 25, 2005
-----------	--	--------	--------------	---------------	------------	--------------

CLAVULANATE POTASSIUM; TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TIMENTIN

+		GLAXOSMITHKLINE	EQ 1GM BASE/VIAL;EQ 30GM BASE/VIAL	N50590	003	Aug 18, 1987
---	--	-----------------	------------------------------------	--------	-----	--------------

+			EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL	N50590	001	Apr 01, 1985
---	--	--	-------------------------------------	--------	-----	--------------

			EQ 100MG BASE/VIAL;EQ 3GM BASE/VIAL	N62691	001	Dec 19, 1986
--	--	--	-------------------------------------	--------	-----	--------------

+			EQ 200MG BASE/VIAL;EQ 3GM BASE/VIAL	N50590	002	Apr 01, 1985
---	--	--	-------------------------------------	--------	-----	--------------

TIMENTIN IN PLASTIC CONTAINER

+		GLAXOSMITHKLINE	EQ 100MG BASE/100ML;EQ 3GM BASE/100ML	N50658	001	Dec 15, 1989
---	--	-----------------	---------------------------------------	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 91 (of 360)

CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

<u>AA</u>	ALPHARMA US PHARMS	<u>EQ 0.5MG BASE/5ML</u>	<u>N74075</u>	<u>001</u>	Oct 31, 1993
<u>AA</u>	MORTON GROVE	<u>EQ 0.5MG BASE/5ML</u>	<u>N74863</u>	<u>001</u>	Mar 13, 1998
<u>AA</u>	NOVEX	<u>EQ 0.5MG BASE/5ML</u>	<u>N75703</u>	<u>001</u>	Nov 27, 2000
<u>AA</u>	SILARX	<u>EQ 0.5MG BASE/5ML</u>	<u>N74884</u>	<u>001</u>	Dec 17, 1997
<u>AA</u> +	TEVA	<u>EQ 0.5MG BASE/5ML</u>	<u>N73399</u>	<u>001</u>	Jun 30, 1994
<u>AA</u>	TEVA PHARMS	<u>EQ 0.5MG BASE/5ML</u>	<u>N73095</u>	<u>001</u>	Apr 21, 1992

TABLET; ORAL

CLEMASTINE FUMARATE

<u>AB</u>	SANDOZ	<u>2.68MG</u>	<u>N73459</u>	<u>001</u>	Oct 31, 1993
<u>AB</u> +	TEVA	<u>2.68MG</u>	<u>N73283</u>	<u>001</u>	Jan 31, 1992

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN HYDROCHLORIDE

<u>AB</u>	PHARMACIA AND UPJOHN	<u>EQ 75MG BASE</u>	<u>N50162</u>	<u>001</u>	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N50162</u>	<u>002</u>	
<u>AB</u> +		<u>EQ 300MG BASE</u>	<u>N50162</u>	<u>003</u>	Apr 14, 1988

CLINDAMYCIN HYDROCHLORIDE

<u>AB</u>	COREPHARMA	<u>EQ 150MG BASE</u>	<u>N65194</u>	<u>001</u>	Mar 22, 2004
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N65194</u>	<u>002</u>	Mar 22, 2004
<u>AB</u>	LANNETT	<u>EQ 75MG BASE</u>	<u>N65242</u>	<u>001</u>	Aug 12, 2005
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N65242</u>	<u>002</u>	Aug 12, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N65243</u>	<u>001</u>	Aug 12, 2005
<u>AB</u>	RANBAXY	<u>EQ 150MG BASE</u>	<u>N65061</u>	<u>001</u>	Feb 02, 2001
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N65061</u>	<u>002</u>	Feb 02, 2001
<u>AB</u>	TEVA	<u>EQ 75MG BASE</u>	<u>N63027</u>	<u>001</u>	Sep 20, 1989
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N63029</u>	<u>001</u>	Sep 20, 1989
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N63029</u>	<u>002</u>	Aug 05, 2005
<u>AB</u>	WATSON LABS	<u>EQ 150MG BASE</u>	<u>N63083</u>	<u>001</u>	Jul 31, 1991
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N63083</u>	<u>002</u>	Mar 18, 2003
<u>AB</u>	ZYDUS PHARMS USA	<u>EQ 75MG BASE</u>	<u>N65217</u>	<u>001</u>	Jan 31, 2005
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N65217</u>	<u>002</u>	Jan 31, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N65217</u>	<u>003</u>	Jan 31, 2005

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION; ORAL

CLEOCIN

+	PHARMACIA AND UPJOHN	EQ 75MG BASE/5ML	N62644	001	Apr 07, 1986
---	----------------------	------------------	--------	-----	--------------

CLINDAMYCIN PHOSPHATE

AEROSOL, FOAM; TOPICAL

EVOCLIN

+	CONNETICS	1%	N50801	001	Oct 22, 2004
---	-----------	----	--------	-----	--------------

CREAM; VAGINAL

CLEOCIN

<u>AB</u> +	PHARMACIA AND UPJOHN	<u>EQ 2% BASE</u>	<u>N50680</u>	<u>002</u>	Mar 02, 1998
-------------	----------------------	-------------------	---------------	------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AB</u>	ALTANA PHARMA	<u>EQ 2% BASE</u>	<u>N65139</u>	<u>001</u>	Dec 27, 2004
	CLINDESSE				
+	KV PHARM	EQ 2% BASE	N50793	001	Nov 30, 2004

GEL; TOPICAL

CLEOCIN T

<u>AB</u> +	PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N50615</u>	<u>001</u>	Jan 07, 1987
	CLINDAGEL				
BT	GALDERMA LABS LP	EQ 1% BASE	N50782	001	Nov 27, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 92 (of 360)

CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

CLINDAMYCIN PHOSPHATE

<u>AB</u>	ALTANA	<u>EQ 1% BASE</u>	<u>N64160</u>	<u>001</u>	Jan 28, 2000
-----------	--------	-------------------	---------------	------------	--------------

INJECTABLE; INJECTION

CLEOCIN PHOSPHATE

<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>EQ 150MG BASE/ML</u>	<u>N50441</u>	<u>001</u>	
-----------	------------------------	-------------------------	---------------	------------	--

<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>N62803</u>	<u>001</u>	Oct 16, 1987
-----------	--	-------------------------	---------------	------------	--------------

CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>EQ 6MG BASE/ML</u>	<u>N50639</u>	<u>001</u>	Aug 30, 1989
-----------	------------------------	-----------------------	---------------	------------	--------------

<u>AP</u>	+	<u>EQ 12MG BASE/ML</u>	<u>N50639</u>	<u>002</u>	Aug 30, 1989
-----------	---	------------------------	---------------	------------	--------------

<u>AP</u>	+	<u>EQ 18MG BASE/ML</u>	<u>N50639</u>	<u>003</u>	Apr 10, 1991
-----------	---	------------------------	---------------	------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 150MG BASE/ML</u>	<u>N62889</u>	<u>001</u>	Apr 25, 1988
-----------	-----------------	-------------------------	---------------	------------	--------------

<u>AP</u>	BEDFORD	<u>EQ 150MG BASE/ML</u>	<u>N65206</u>	<u>001</u>	Sep 24, 2004
-----------	---------	-------------------------	---------------	------------	--------------

<u>AP</u>	HOSPIRA	<u>EQ 150MG BASE/ML</u>	<u>N62800</u>	<u>001</u>	Jul 24, 1987
-----------	---------	-------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>N62801</u>	<u>001</u>	Jul 24, 1987
-----------	--	-------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 150MG BASE/ML</u>	<u>N62943</u>	<u>001</u>	Sep 29, 1988
-----------	--	-------------------------	---------------	------------	--------------

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

<u>AP</u>	+ AM PHARM PARTNERS	<u>EQ 900MG BASE/100ML</u>	<u>N50635</u>	<u>001</u>	Dec 22, 1989
-----------	---------------------	----------------------------	---------------	------------	--------------

LOTION; TOPICAL

CLEOCIN T

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N50600</u>	<u>001</u>	May 31, 1989
-----------	------------------------	-------------------	---------------	------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AB</u>	ALTANA	<u>EQ 1% BASE</u>	<u>N65067</u>	<u>001</u>	Jan 31, 2002
-----------	--------	-------------------	---------------	------------	--------------

SOLUTION; TOPICAL

CLEOCIN T

<u>AT</u>	+ PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N50537</u>	<u>001</u>	
-----------	------------------------	-------------------	---------------	------------	--

CLINDA-DERM

<u>AT</u>	PADDOCK	<u>EQ 1% BASE</u>	<u>N63329</u>	<u>001</u>	Sep 30, 1992
-----------	---------	-------------------	---------------	------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AT</u>	ALPHARMA US PHARMS	<u>EQ 1% BASE</u>	<u>N62811</u>	<u>001</u>	Sep 01, 1988
-----------	--------------------	-------------------	---------------	------------	--------------

<u>AT</u>	CLAY PARK	<u>EQ 1% BASE</u>	<u>N64050</u>	<u>001</u>	Nov 30, 1995
-----------	-----------	-------------------	---------------	------------	--------------

<u>AT</u>	FOUGERA	<u>EQ 1% BASE</u>	<u>N64159</u>	<u>001</u>	Jun 05, 1997
-----------	---------	-------------------	---------------	------------	--------------

<u>AT</u>	MORTON GROVE	<u>EQ 1% BASE</u>	<u>N63304</u>	<u>001</u>	Jul 15, 1997
-----------	--------------	-------------------	---------------	------------	--------------

<u>AT</u>	STIEFEL	<u>EQ 1% BASE</u>	<u>N64108</u>	<u>001</u>	Sep 27, 1996
-----------	---------	-------------------	---------------	------------	--------------

<u>AT</u>	TARO PHARM INDS	<u>EQ 1% BASE</u>	<u>N65184</u>	<u>001</u>	Mar 31, 2004
-----------	-----------------	-------------------	---------------	------------	--------------

SUPPOSITORY; VAGINAL

CLEOCIN

	+ PHARMACIA AND UPJOHN	100MG	N50767	001	Aug 13, 1999
--	------------------------	-------	--------	-----	--------------

SWAB; TOPICAL

CLEOCIN

<u>AT</u>	PHARMACIA AND UPJOHN	<u>EQ 1% BASE</u>	<u>N50537</u>	<u>002</u>	Feb 22, 1994
-----------	----------------------	-------------------	---------------	------------	--------------

CLINDAMYCIN PHOSPHATE

<u>AT</u>	CLAY PARK	<u>EQ 1% BASE</u>	<u>N65049</u>	<u>001</u>	May 25, 2000
-----------	-----------	-------------------	---------------	------------	--------------

CLINDETS

<u>AT</u>	STIEFEL	<u>EQ 1% BASE</u>	<u>N64136</u>	<u>001</u>	Sep 30, 1996
-----------	---------	-------------------	---------------	------------	--------------

CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

OLUX

	+ CONNETICS	0.05%	N21142	001	May 26, 2000
--	-------------	-------	--------	-----	--------------

CREAM; TOPICAL

CLOBETASOL PROPIONATE

<u>AB1</u>	ALPHARMA US PHARMS	<u>0.05%</u>	<u>N74139</u>	<u>001</u>	Aug 03, 1994
------------	--------------------	--------------	---------------	------------	--------------

<u>AB1</u>	FOUGERA	<u>0.05%</u>	<u>N74392</u>	<u>001</u>	Sep 30, 1996
------------	---------	--------------	---------------	------------	--------------

<u>AB1</u>	STIEFEL	<u>0.05%</u>	<u>N75338</u>	<u>001</u>	Feb 09, 2001
------------	---------	--------------	---------------	------------	--------------

<u>AB1</u>	TARO	<u>0.05%</u>	<u>N74249</u>	<u>001</u>	Jul 08, 1996
------------	------	--------------	---------------	------------	--------------

<u>AB1</u>	TEVA PHARMS	<u>0.05%</u>	<u>N74087</u>	<u>001</u>	Feb 16, 1994
------------	-------------	--------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 93 (of 360)

CLOBETASOL PROPIONATE

CREAM; TOPICAL

CLOBETASOL PROPIONATE (EMOLLIENT)

<u>AB2</u>	ALTANA	<u>0.05%</u>	<u>N75430</u>	<u>001</u>	May 26, 1999
<u>AB2</u>	STIEFEL	<u>0.05%</u>	<u>N75733</u>	<u>001</u>	Aug 22, 2001
<u>AB2</u>	TARO	<u>0.05%</u>	<u>N75633</u>	<u>001</u>	May 17, 2000
<u>CORMAX</u>					
<u>AB1</u>	HEALTHPOINT	<u>0.05%</u>	<u>N74220</u>	<u>001</u>	May 16, 1997
<u>EMBELINE E</u>					
<u>AB2</u>	HEALTHPOINT	<u>0.05%</u>	<u>N75325</u>	<u>001</u>	Dec 24, 1998
<u>TEMOVATE</u>					
<u>AB1</u> +	ALTANA	<u>0.05%</u>	<u>N19322</u>	<u>001</u>	Dec 27, 1985
<u>TEMOVATE E</u>					
<u>AB2</u> +	ALTANA	<u>0.05%</u>	<u>N20340</u>	<u>001</u>	Jun 17, 1994

GEL; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>N75368</u>	<u>001</u>	Feb 15, 2000
<u>AB</u>	STIEFEL	<u>0.05%</u>	<u>N75027</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>	TARO	<u>0.05%</u>	<u>N75279</u>	<u>001</u>	May 28, 1999
<u>EMBELINE</u>					
<u>AB</u>	HEALTHPOINT	<u>0.05%</u>	<u>N76141</u>	<u>001</u>	Apr 12, 2002
<u>TEMOVATE</u>					
<u>AB</u> +	ALTANA	<u>0.05%</u>	<u>N20337</u>	<u>001</u>	Apr 29, 1994

LOTION; TOPICAL

CLOBEX

+	GALDERMA LABS LP	0.05%	N21535	001	Jul 24, 2003
---	------------------	-------	--------	-----	--------------

OINTMENT; TOPICAL

CLOBETASOL PROPIONATE

<u>AB</u>	ALPHARMA US PHARMS	<u>0.05%</u>	<u>N74128</u>	<u>001</u>	Aug 03, 1994
<u>AB</u>	FOUGERA	<u>0.05%</u>	<u>N74407</u>	<u>001</u>	Feb 23, 1996
<u>AB</u>	STIEFEL	<u>0.05%</u>	<u>N75057</u>	<u>001</u>	Aug 12, 1998
<u>AB</u>	TARO	<u>0.05%</u>	<u>N74248</u>	<u>001</u>	Jul 12, 1996
<u>AB</u>	TEVA PHARMS	<u>0.05%</u>	<u>N74089</u>	<u>001</u>	Feb 16, 1994
<u>EMBELINE</u>					
<u>AB</u>	HEALTHPOINT	<u>0.05%</u>	<u>N74221</u>	<u>001</u>	Mar 31, 1995
<u>TEMOVATE</u>					
<u>AB</u> +	ALTANA	<u>0.05%</u>	<u>N19323</u>	<u>001</u>	Dec 27, 1985

SHAMPOO; TOPICAL

CLOBEX

+	GALDERMA LABS	0.05%	N21644	001	Feb 05, 2004
---	---------------	-------	--------	-----	--------------

SOLUTION; TOPICAL

CLOBETASOL PROPIONATE

<u>AT</u>	ALPHARMA US PHARMS	<u>0.05%</u>	<u>N74331</u>	<u>001</u>	Dec 15, 1995
<u>AT</u>	ALTANA	<u>0.05%</u>	<u>N75391</u>	<u>001</u>	Feb 08, 1999
<u>AT</u>	MORTON GROVE	<u>0.05%</u>	<u>N75205</u>	<u>001</u>	Nov 13, 1998
<u>AT</u>	QLT USA	<u>0.05%</u>	<u>N76977</u>	<u>001</u>	Aug 05, 2005
<u>AT</u>	TARO	<u>0.05%</u>	<u>N75224</u>	<u>001</u>	Nov 16, 1998
<u>AT</u>		<u>0.05%</u>	<u>N75363</u>	<u>001</u>	Dec 29, 2000
<u>EMBELINE</u>					
<u>AT</u>	DPT	<u>0.05%</u>	<u>N74222</u>	<u>001</u>	Dec 06, 1995
<u>TEMOVATE</u>					
<u>AT</u> +	ALTANA	<u>0.05%</u>	<u>N19966</u>	<u>001</u>	Feb 22, 1990

SPRAY; TOPICAL

CLOBEX

+	DOW PHARM SCI	0.05%	N21835	001	Oct 27, 2005
---	---------------	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 94 (of 360)

CLOCORTOLONE PIVALATE

CREAM; TOPICAL

CLODERM

+ HEALTHPOINT

0.1%

N17765 001

CLOFARABINE

INJECTABLE; IV (INFUSION)

CLOLAR

+ GENZYME

20MG/20ML (1MG/ML)

N21673 001

Dec 28, 2004

CLOFIBRATE

CAPSULE; ORAL

CLOFIBRATEAB BANNER PHARMACAPS500MGN73396 001

Mar 20, 1992

AB TEVA500MGN72600 001

Jul 25, 1991

AB + USL PHARMA500MGN70531 001

Jun 16, 1986

CLOMIPHENE CITRATE

TABLET; ORAL

CLOMIDAB + AVENTIS PHARMS50MGN16131 002CLOMIPHENE CITRATEAB PAR PHARM50MGN75528 001

Aug 30, 1999

MILOPHENEAB MILEX50MGN72196 001

Dec 20, 1988

SEROPHENEAB SERONO50MGN18361 001

Mar 22, 1982

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

ANAFRANILAB TYCO HLTHCARE25MGN19906 001

Dec 29, 1989

AB +50MGN19906 002

Dec 29, 1989

AB75MGN19906 003

Dec 29, 1989

CLOMIPRAMINE HYDROCHLORIDEAB MYLAN25MGN74947 001

Apr 30, 1998

AB50MGN74947 002

Apr 30, 1998

AB75MGN74947 003

Apr 30, 1998

AB SANDOZ25MGN74364 001

Mar 29, 1996

AB25MGN74953 001

Jun 25, 1997

AB50MGN74364 002

Mar 29, 1996

AB50MGN74953 002

Jun 25, 1997

AB75MGN74364 003

Mar 29, 1996

AB75MGN74953 003

Jun 25, 1997

AB TARO25MGN74694 001

Dec 31, 1996

AB50MGN74694 002

Dec 31, 1996

AB75MGN74694 003

Dec 31, 1996

AB TEVA25MGN74849 001

Apr 04, 1997

AB25MGN74958 001

Aug 26, 1997

AB50MGN74849 002

Apr 04, 1997

AB50MGN74958 002

Aug 26, 1997

AB75MGN74849 003

Apr 04, 1997

AB75MGN74958 003

Aug 26, 1997

AB WATSON LABS25MGN74600 001

Nov 27, 1996

AB25MGN74751 001

Sep 30, 1998

AB50MGN74600 002

Nov 27, 1996

AB50MGN74751 002

Sep 30, 1998

AB75MGN74600 003

Nov 27, 1996

AB75MGN74751 003

Sep 30, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 95 (of 360)

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

<u>AB</u>	ALPHAPHARM	<u>0.5MG</u>	<u>N74940</u>	<u>001</u>	Oct 30, 1997
<u>AB</u>		<u>1MG</u>	<u>N74940</u>	<u>002</u>	Oct 30, 1997
<u>AB</u>		<u>2MG</u>	<u>N74940</u>	<u>003</u>	Oct 30, 1997
<u>AB</u>	CARACO	<u>0.5MG</u>	<u>N75423</u>	<u>001</u>	Apr 27, 2001
<u>AB</u>		<u>1MG</u>	<u>N75423</u>	<u>002</u>	Apr 27, 2001
<u>AB</u>		<u>2MG</u>	<u>N75423</u>	<u>003</u>	Apr 27, 2001
<u>AB</u>	KALI LABS	<u>0.5MG</u>	<u>N77147</u>	<u>001</u>	May 02, 2005
<u>AB</u>		<u>1MG</u>	<u>N77147</u>	<u>002</u>	May 02, 2005
<u>AB</u>		<u>2MG</u>	<u>N77147</u>	<u>003</u>	May 02, 2005
<u>AB</u>	MYLAN	<u>0.5MG</u>	<u>N75150</u>	<u>001</u>	Oct 05, 1998
<u>AB</u>		<u>1MG</u>	<u>N75150</u>	<u>002</u>	Oct 05, 1998
<u>AB</u>		<u>2MG</u>	<u>N75150</u>	<u>003</u>	Oct 05, 1998
<u>AB</u>	PUREPAC PHARM	<u>0.5MG</u>	<u>N74869</u>	<u>001</u>	Oct 31, 1996
<u>AB</u>		<u>1MG</u>	<u>N74869</u>	<u>002</u>	Oct 31, 1996
<u>AB</u>		<u>2MG</u>	<u>N74869</u>	<u>003</u>	Oct 31, 1996
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>N74925</u>	<u>001</u>	Sep 30, 1997
<u>AB</u>		<u>0.5MG</u>	<u>N74979</u>	<u>001</u>	Aug 29, 1997
<u>AB</u>		<u>1MG</u>	<u>N74925</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>		<u>1MG</u>	<u>N74979</u>	<u>002</u>	Aug 29, 1997
<u>AB</u>		<u>2MG</u>	<u>N74925</u>	<u>003</u>	Sep 30, 1997
<u>AB</u>		<u>2MG</u>	<u>N74979</u>	<u>003</u>	Aug 29, 1997
<u>AB</u>	TEVA	<u>0.5MG</u>	<u>N74569</u>	<u>001</u>	Sep 10, 1996
<u>AB</u>		<u>0.5MG</u>	<u>N74920</u>	<u>001</u>	Aug 04, 1998
<u>AB</u>		<u>1MG</u>	<u>N74569</u>	<u>002</u>	Sep 10, 1996
<u>AB</u>		<u>1MG</u>	<u>N74920</u>	<u>002</u>	Aug 04, 1998
<u>AB</u>		<u>2MG</u>	<u>N74569</u>	<u>003</u>	Sep 10, 1996
<u>AB</u>		<u>2MG</u>	<u>N74920</u>	<u>003</u>	Aug 04, 1998
<u>AB</u>	TORPHARM	<u>0.5MG</u>	<u>N75468</u>	<u>001</u>	Oct 06, 2000
<u>AB</u>		<u>1MG</u>	<u>N75468</u>	<u>002</u>	Oct 06, 2000
<u>AB</u>		<u>2MG</u>	<u>N75468</u>	<u>003</u>	Oct 06, 2000
<u>AB</u>	WATSON LABS	<u>0.5MG</u>	<u>N74964</u>	<u>001</u>	Dec 30, 1997
<u>AB</u>		<u>1MG</u>	<u>N74964</u>	<u>002</u>	Dec 30, 1997
<u>AB</u>		<u>2MG</u>	<u>N74964</u>	<u>003</u>	Dec 30, 1997
<u>KLONOPIN</u>					
<u>AB</u>	ROCHE	<u>0.5MG</u>	<u>N17533</u>	<u>001</u>	
<u>AB</u>	+	<u>1MG</u>	<u>N17533</u>	<u>002</u>	
<u>AB</u>		<u>2MG</u>	<u>N17533</u>	<u>003</u>	

TABLET, ORALLY DISINTEGRATING; ORAL

CLONAZEPAM

<u>AB</u>	BARR	<u>0.125MG</u>	<u>N77194</u>	<u>001</u>	Aug 10, 2005
<u>AB</u>		<u>0.25MG</u>	<u>N77194</u>	<u>002</u>	Aug 10, 2005
<u>AB</u>		<u>0.5MG</u>	<u>N77194</u>	<u>003</u>	Aug 10, 2005
<u>AB</u>		<u>1MG</u>	<u>N77194</u>	<u>004</u>	Aug 10, 2005
<u>AB</u>		<u>2MG</u>	<u>N77194</u>	<u>005</u>	Aug 10, 2005
<u>AB</u>	KALI LABS	<u>0.125MG</u>	<u>N77171</u>	<u>001</u>	Aug 03, 2005
<u>AB</u>		<u>0.25MG</u>	<u>N77171</u>	<u>002</u>	Aug 03, 2005
<u>AB</u>		<u>0.5MG</u>	<u>N77171</u>	<u>003</u>	Aug 03, 2005
<u>AB</u>		<u>1MG</u>	<u>N77171</u>	<u>004</u>	Aug 03, 2005
<u>AB</u>		<u>2MG</u>	<u>N77171</u>	<u>005</u>	Aug 03, 2005
<u>KLONOPIN RAPIDLY DISINTEGRATING</u>					
<u>AB</u>	ROCHE	<u>0.125MG</u>	<u>N20813</u>	<u>001</u>	Dec 23, 1997
<u>AB</u>		<u>0.25MG</u>	<u>N20813</u>	<u>002</u>	Dec 23, 1997
<u>AB</u>		<u>0.5MG</u>	<u>N20813</u>	<u>003</u>	Dec 23, 1997
<u>AB</u>	+	<u>1MG</u>	<u>N20813</u>	<u>004</u>	Dec 23, 1997
<u>AB</u>		<u>2MG</u>	<u>N20813</u>	<u>005</u>	Dec 23, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 96 (of 360)

CLONIDINE

FILM, EXTENDED RELEASE; TRANSDERMAL

CATAPRES-TTS-1

BOEHRINGER INGELHEIM 0.1MG/24HR

N18891 001 Oct 10, 1984

CATAPRES-TTS-2

BOEHRINGER INGELHEIM 0.2MG/24HR

N18891 002 Oct 10, 1984

CATAPRES-TTS-3

+ BOEHRINGER INGELHEIM 0.3MG/24HR

N18891 003 Oct 10, 1984

CLONIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DURACLON

XANODYNE PHARM 0.1MG/ML

N20615 001 Oct 02, 1996

+ 0.5MG/ML

N20615 002 Apr 27, 1999

TABLET; ORAL

CATAPRES

AB BOEHRINGER INGELHEIM 0.1MG

N17407 001

AB 0.2MG

N17407 002

AB + 0.3MG

N17407 003

CLONIDINE HYDROCHLORIDE

AB CLONMEL HLTHCARE 0.1MG

N71783 001 Apr 05, 1988

AB 0.2MG

N71784 001 Apr 05, 1988

AB 0.3MG

N71785 001 Apr 05, 1988

AB MUTUAL PHARM 0.1MG

N70925 001 Sep 04, 1987

AB 0.2MG

N70924 001 Sep 04, 1987

AB 0.3MG

N70923 001 Sep 04, 1987

AB MYLAN 0.1MG

N70315 001 Jun 09, 1987

AB 0.2MG

N70316 001 Jun 09, 1987

AB 0.3MG

N70317 001 Jun 09, 1987

AB PUREPAC PHARM 0.1MG

N70974 001 Dec 16, 1986

AB 0.2MG

N70975 001 Dec 16, 1986

AB 0.3MG

N70976 001 Dec 16, 1986

AB SANDOZ 0.1MG

N70887 001 Aug 31, 1988

AB 0.2MG

N70886 001 Aug 31, 1988

AB 0.3MG

N71294 001 Aug 31, 1988

AB WATSON LABS 0.1MG

N70965 001 Jul 08, 1986

AB 0.2MG

N70964 001 Jul 08, 1986

AB 0.3MG

N70963 001 Jul 08, 1986

CLOPIDOGREL BISULFATE

TABLET; ORAL

PLAVIX

+ SANOFI SYNTHELABO EQ 75MG BASE

N20839 001 Nov 17, 1997

CLORAZEPATE DIPOTASSIUM

CAPSULE; ORAL

CLORAZEPATE DIPOTASSIUM

AB MYLAN 3.75MG

N71509 001 Oct 19, 1987

AB 7.5MG

N71510 001 Oct 19, 1987

AB + 15MG

N71511 001 Oct 19, 1987

AB SANDOZ 3.75MG

N72219 001 Aug 26, 1988

AB 15MG

N72112 001 Aug 26, 1988

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

AB MYLAN 3.75MG

N71856 001 Jul 17, 1987

AB 7.5MG

N71857 001 Jul 17, 1987

AB 15MG

N71858 001 Jul 17, 1987

AB RANBAXY 3.75MG

N76911 001 Sep 29, 2004

AB 7.5MG

N76911 002 Sep 29, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 97 (of 360)

CLORAZEPATE DIPOTASSIUM

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

<u>AB</u>	RANBAXY	<u>15MG</u>	<u>N76911</u>	<u>003</u>	Sep 29, 2004
<u>AB</u>	SANDOZ	<u>7.5MG</u>	<u>N72513</u>	<u>001</u>	May 11, 1990
<u>AB</u>		<u>15MG</u>	<u>N72514</u>	<u>001</u>	May 11, 1990
<u>AB</u>	TARO	<u>3.75MG</u>	<u>N75731</u>	<u>003</u>	Apr 27, 2000
<u>AB</u>		<u>7.5MG</u>	<u>N75731</u>	<u>002</u>	Apr 27, 2000
<u>AB</u>		<u>15MG</u>	<u>N75731</u>	<u>001</u>	Apr 27, 2000
<u>AB</u>	WATSON LABS	<u>3.75MG</u>	<u>N71852</u>	<u>001</u>	Feb 09, 1988
<u>AB</u>		<u>7.5MG</u>	<u>N71853</u>	<u>001</u>	Feb 09, 1988
<u>AB</u>		<u>15MG</u>	<u>N71854</u>	<u>001</u>	Feb 09, 1988
	<u>GEN-XENE</u>				
<u>AB</u>	ALRA	<u>3.75MG</u>	<u>N71787</u>	<u>001</u>	Apr 26, 1988
<u>AB</u>		<u>7.5MG</u>	<u>N71788</u>	<u>001</u>	Apr 26, 1988
<u>AB</u>		<u>15MG</u>	<u>N71789</u>	<u>001</u>	Apr 26, 1988
	<u>TRANXENE</u>				
<u>AB</u>	OVATION PHARMS	<u>3.75MG</u>	<u>N17105</u>	<u>006</u>	
<u>AB</u>		<u>7.5MG</u>	<u>N17105</u>	<u>007</u>	
<u>AB</u>	+	<u>15MG</u>	<u>N17105</u>	<u>008</u>	
	TRANXENE SD				
	OVATION PHARMS	11.25MG	N17105	005	
	+	22.5MG	N17105	004	

CLOTTRIMAZOLE

CREAM; TOPICAL

CLOTTRIMAZOLE

	+	TARO	1%	N72640	001	Aug 31, 1993
--	---	------	----	--------	-----	--------------

SOLUTION; TOPICAL

CLOTTRIMAZOLE

<u>AT</u>	TARO	<u>1%</u>	<u>N74580</u>	<u>001</u>	Jul 29, 1996
<u>AT</u>	TEVA	<u>1%</u>	<u>N73306</u>	<u>001</u>	Feb 28, 1995
	<u>MYCELEX</u>				
<u>AT</u>	+	BAYER PHARMS	<u>1%</u>	<u>N18181</u>	<u>001</u>

TROCHE/LOZENGE; ORAL

CLOTTRIMAZOLE

<u>AB</u>	PADDOCK	<u>10MG</u>	<u>N76763</u>	<u>001</u>	Oct 28, 2005
<u>AB</u>	ROXANE	<u>10MG</u>	<u>N76387</u>	<u>001</u>	Jul 29, 2004
	<u>MYCELEX</u>				
<u>AB</u>	+	BAYER PHARMS	<u>10MG</u>	<u>N18713</u>	<u>001</u>

CLOXACILLIN SODIUM

CAPSULE; ORAL

CLOXACILLIN SODIUM

<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>N62240</u>	<u>001</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N62240</u>	<u>002</u>	
	<u>CLOXAPEN</u>				
<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 250MG BASE</u>	<u>N61806</u>	<u>001</u>	
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>N61806</u>	<u>002</u>	

FOR SOLUTION; ORAL

CLOXACILLIN SODIUM

	+	TEVA	EQ 125MG BASE/5ML	N62268	001
--	---	------	-------------------	--------	-----

CLOZAPINE

TABLET; ORAL

CLOZAPINE

<u>AB</u>	CARACO	<u>25MG</u>	<u>N75713</u>	<u>001</u>	Nov 15, 2002
<u>AB</u>		<u>50MG</u>	<u>N75713</u>	<u>003</u>	Aug 19, 2005
<u>AB</u>		<u>100MG</u>	<u>N75713</u>	<u>002</u>	Nov 15, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 98 (of 360)

CLOZAPINE

TABLET; ORAL					
<u>CLOZAPINE</u>					
<u>AB</u>	GOLDLINE	<u>50MG</u>	<u>N76809</u>	<u>003</u>	Dec 16, 2005
<u>AB</u>		<u>100MG</u>	<u>N76809</u>	<u>002</u>	Dec 16, 2005
		200MG	N76809	001	Dec 16, 2005
<u>AB</u>	IVAX PHARMS	<u>25MG</u>	<u>N74949</u>	<u>001</u>	Nov 26, 1997
<u>AB</u>		<u>50MG</u>	<u>N74949</u>	<u>004</u>	Apr 25, 2005
<u>AB</u>		<u>100MG</u>	<u>N74949</u>	<u>002</u>	Nov 26, 1997
		12.5MG	N74949	003	Jul 31, 2003
<u>AB</u>	MYLAN	<u>25MG</u>	<u>N75417</u>	<u>001</u>	May 27, 1999
<u>AB</u>		<u>100MG</u>	<u>N75417</u>	<u>002</u>	May 27, 1999
<u>AB</u>	TEVA	<u>25MG</u>	<u>N75162</u>	<u>001</u>	Apr 26, 2005
<u>AB</u>		<u>100MG</u>	<u>N75162</u>	<u>002</u>	Apr 26, 2005
<u>CLOZARIL</u>					
<u>AB</u>	NOVARTIS	<u>25MG</u>	<u>N19758</u>	<u>001</u>	Sep 26, 1989
<u>AB</u>	+	<u>100MG</u>	<u>N19758</u>	<u>002</u>	Sep 26, 1989
TABLET, ORALLY DISINTEGRATING; ORAL					
FAZACLO ODT					
	ALAMO PHARMS	25MG	N21590	001	Feb 10, 2004
		50MG	N21590	003	Jun 03, 2005
	+	100MG	N21590	002	Feb 10, 2004

COBALT CHLORIDE, CO-57; CYANOCOBALAMIN; CYANOCOBALAMIN, CO-57; INTRINSIC FACTOR

N/A; N/A					
RUBRATOPE-57 KIT					
	BRACCO	N/A;N/A;N/A;N/A	N16089	001	

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL						
<u>PROMETH VC W/ CODEINE</u>						
<u>AA</u>	+	ALPHARMA US PHARMS	<u>10MG/5ML; 5MG/5ML; 6.25MG/5ML</u>	<u>N88764</u>	<u>001</u>	Oct 31, 1984
<u>PROMETHAZINE VC W/ CODEINE</u>						
<u>AA</u>		MORTON GROVE	<u>10MG/5ML; 5MG/5ML; 6.25MG/5ML</u>	<u>N88896</u>	<u>001</u>	Jan 04, 1985

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL						
<u>PROMETH W/ CODEINE</u>						
<u>AA</u>	+	ALPHARMA US PHARMS	<u>10MG/5ML; 6.25MG/5ML</u>	<u>N88763</u>	<u>001</u>	Oct 31, 1984
<u>PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE</u>						
<u>AA</u>		HI TECH PHARMA	<u>10MG/5ML; 6.25MG/5ML</u>	<u>N40151</u>	<u>001</u>	Aug 26, 1997
<u>AA</u>		PHARM ASSOC	<u>10MG/5ML; 6.25MG/5ML</u>	<u>N89647</u>	<u>001</u>	Dec 22, 1988
<u>PROMETHAZINE W/ CODEINE</u>						
<u>AA</u>		MORTON GROVE	<u>10MG/5ML; 6.25MG/5ML</u>	<u>N88875</u>	<u>001</u>	Dec 17, 1984

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL						
<u>TRIACIN-C</u>						
<u>AA</u>		ALPHARMA US PHARMS	<u>10MG/5ML;30MG/5ML;1.25MG/5ML</u>	<u>N88704</u>	<u>001</u>	Mar 22, 1985
<u>TRIPROLIDINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE AND CODEINE PHOSPHATE</u>						
<u>AA</u>	+	MORTON GROVE	<u>10MG/5ML;30MG/5ML;1.25MG/5ML</u>	<u>N88833</u>	<u>001</u>	Nov 16, 1984

COLCHICINE; PROBENECID

TABLET; ORAL						
COL-PROBENECID						
BP	+	WATSON LABS	0.5MG;500MG	N84279	001	
PROBENECID AND COLCHICINE						
BP		IVAX PHARMS	0.5MG;500MG	N83734	001	

PRESCRIPTION DRUG PRODUCT LIST

3 - 99 (of 360)

COLESEVELAM HYDROCHLORIDE

TABLET; ORAL				
WELCHOL				
+	SANKYO	625MG	N21176 001	May 26, 2000

COLESTIPOL HYDROCHLORIDE

GRANULE; ORAL				
COLESTID				
+	PHARMACIA AND UPJOHN	5GM/PACKET	N17563 004	Sep 22, 1995
		5GM/SC00PFUL	N17563 003	Sep 22, 1995
FLAVORED COLESTID				
	PHARMACIA AND UPJOHN	5GM/SC00PFUL	N17563 002	
+		5GM/PACKET	N17563 001	
TABLET; ORAL				
COLESTID				
+	PHARMACIA AND UPJOHN	1GM	N20222 001	Jul 19, 1994

COLISTIMETHATE SODIUM

INJECTABLE; INJECTION				
<u>COLISTIMETHATE</u>				
<u>AP</u>	PADDOCK	<u>EQ 150MG BASE/VIAL</u>	<u>N65177 001</u>	Mar 19, 2004
<u>AP</u>	PHARMA TEK	<u>EQ 150MG BASE/VIAL</u>	<u>N64216 001</u>	Feb 26, 1999
<u>COLY-MYCIN M</u>				
<u>AP</u>	+	PARKEDALE	<u>EQ 150MG BASE/VIAL</u>	<u>N50108 002</u>

COLISTIN SULFATE; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; THONZONIUM BROMIDE

SUSPENSION/DROPS; OTIC			
COLY-MYCIN S			
+	PARKEDALE	EQ 3MG BASE/ML;10MG/ML;EQ 3.3MG BASE/ML;0.5MG/ML	N50356 001

CONIVAPTAN HYDROCHLORIDE

INJECTABLE; IV (INFUSION)			
VAPRISOL			
+	ASTELLAS	20MG/4ML (5MG/ML)	N21697 001
			Dec 29, 2005

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE			
COPPER T MODEL TCU 380A			
+	DURAMED PHARMS BARR	309MG/COPPER	N18680 001
			Nov 15, 1984

CORTICORELIN OVINE TRIFLUTATE

INJECTABLE; INJECTION			
ACTHREL			
+	FERRING	EQ 0.1MG BASE/VIAL	N20162 001
			May 23, 1996

CORTICOTROPIN

INJECTABLE; INJECTION			
H.P. ACTHAR GEL			
+	QUESTCOR PHARMS	80 UNITS/ML	N08372 008

CORTISONE ACETATE

TABLET; ORAL			
CORTISONE ACETATE			
BP	+	PHARMACIA AND UPJOHN	25MG
			N08126 001
			5MG
			N08126 003
			10MG
			N08126 004
BP		WEST WARD	25MG
			N80776 002

PRESCRIPTION DRUG PRODUCT LIST

3 - 100 (of 360)

COSYNTROPIN

INJECTABLE; INJECTION
CORTROSYN

+ AMPHASTAR	0.25MG/VIAL	N16750	001	
-------------	-------------	--------	-----	--

CROMOLYN SODIUM

AEROSOL, METERED; INHALATION
INTAL

+ KING PHARMS	0.8MG/INH	N18887	001	Dec 05, 1985
---------------	-----------	--------	-----	--------------

SOLUTION; INHALATION

CROMOLYN SODIUM

<u>AN</u>	ALPHARMA US PHARMS	<u>10MG/ML</u>	<u>N75067</u>	001	Jul 19, 1999
<u>AN</u>	BAUSCH AND LOMB	<u>10MG/ML</u>	<u>N75585</u>	001	Dec 21, 2000
<u>AN</u>	DEY	<u>10MG/ML</u>	<u>N74209</u>	001	Apr 26, 1994
<u>AN</u>	IVAX PHARMS	<u>10MG/ML</u>	<u>N75271</u>	001	Jan 18, 2000
<u>AN</u>	MORTON GROVE	<u>10MG/ML</u>	<u>N75346</u>	001	Oct 25, 1999
<u>AN</u>	NOVEX	<u>10MG/ML</u>	<u>N75333</u>	001	Apr 30, 2002
<u>AN</u>	RESPIRARE	<u>10MG/ML</u>	<u>N76469</u>	001	Jun 17, 2005
<u>AN</u>	ROXANE	<u>10MG/ML</u>	<u>N75175</u>	001	Sep 30, 1999
<u>AN</u>	WARRICK PHARMS	<u>10MG/ML</u>	<u>N75437</u>	001	Apr 21, 2000

INTAL

<u>AN</u>	+ KING PHARMS	<u>10MG/ML</u>	<u>N18596</u>	001	May 28, 1982
-----------	---------------	----------------	---------------	-----	--------------

SOLUTION, CONCENTRATE; ORAL
GASTROCROM

+ UCB	100MG/5ML	N20479	001	Feb 29, 1996
-------	-----------	--------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

CROLOM

<u>AT</u>	BAUSCH AND LOMB	<u>4%</u>	<u>N74443</u>	001	Jan 30, 1995
-----------	-----------------	-----------	---------------	-----	--------------

CROMOLYN SODIUM

<u>AT</u>	AKORN	<u>4%</u>	<u>N74706</u>	001	Apr 29, 1998
<u>AT</u>	ALCON	<u>4%</u>	<u>N75282</u>	001	Jun 16, 1999
<u>AT</u>	NOVEX	<u>4%</u>	<u>N75615</u>	001	Jan 26, 2001

OPTICROM

<u>AT</u>	+ ALLERGAN	<u>4%</u>	<u>N18155</u>	001	Oct 03, 1984
-----------	------------	-----------	---------------	-----	--------------

CROTAMITON

CREAM; TOPICAL
EURAX

+ WESTWOOD SQUIBB	10%	N06927	001	
-------------------	-----	--------	-----	--

LOTION; TOPICAL

CROTAN

<u>AT</u>	SUMMERS	<u>10%</u>	<u>N87204</u>	001	
-----------	---------	------------	---------------	-----	--

EURAX

<u>AT</u>	+ WESTWOOD SQUIBB	<u>10%</u>	<u>N09112</u>	003	
-----------	-------------------	------------	---------------	-----	--

CUPRIC CHLORIDE

INJECTABLE; INJECTION
CUPRIC CHLORIDE IN PLASTIC CONTAINER

+ HOSPIRA	EQ 0.4MG COPPER/ML	N18960	001	Jun 26, 1986
-----------	--------------------	--------	-----	--------------

CYANOCOBALAMIN

GEL, METERED; NASAL
NASCOBAL

+ QOL MEDCL	0.5MG/INH	N19722	001	Nov 05, 1996
-------------	-----------	--------	-----	--------------

INJECTABLE; INJECTION

CYANOCOBALAMIN

<u>AP</u>	AM PHARM PARTNERS	<u>0.1MG/ML</u>	<u>N80557</u>	002	
<u>AP</u>	BIONICHE PHARMA	<u>1MG/ML</u>	<u>N40451</u>	001	Sep 23, 2003
<u>AP</u>	LUITPOLD	<u>1MG/ML</u>	<u>N80737</u>	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 101 (of 360)

CYANOCOBALAMIN

INJECTABLE; INJECTION

CYANOCOBALAMINAP STERIS 0.1MG/MLN80573 002AP 1MG/MLN80573 001RUBRAMIN PCAP + BRISTOL MYERS SQUIBB 1MG/MLN06799 010

Apr 28, 1988

VIBISONEAP AM PHARM PARTNERS 1MG/MLN80557 003SPRAY, METERED; NASAL
NASCOBAL

+ QOL MEDCL 0.5MG/SPRAY

N21642 001

Jan 31, 2005

CYANOCOBALAMIN, CO-57

CAPSULE; ORAL

RUBRATOPE-57

+ BRACCO 0.5-1uCi

N16089 002

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDEAB AMIDE PHARM 10MGN77209 001

Oct 04, 2005

AB MUTUAL PHARM 10MGN73541 001

May 23, 1995

AB MYLAN 10MGN73144 001

May 30, 1991

AB PLIVA 10MGN74421 001

Sep 29, 1995

AB SANDOZ 10MGN72854 001

Nov 19, 1991

AB 10MGN73683 001

Feb 26, 1993

AB WATSON LABS 10MGN71611 001

May 03, 1989

AB 10MGN73143 001

Nov 27, 1991

AB 10MGN74436 001

Nov 30, 1994

FLEXERILAB + MCNEIL CONS SPECLT 10MGN17821 002

5MG

N17821 001

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKPENTOLATEAT AKORN 1%N40164 001

Jan 13, 1997

AT 2%N40165 001

Jan 13, 1997

CYCLOGYLAT + ALCON 0.5%N84109 001AT + 1%N84110 001AT + 2%N84108 001CYCLOPENTOLATE HYDROCHLORIDEAT ALCON UNIVERSAL 1%N89162 001

Jan 24, 1991

PENTOLAIRAT BAUSCH AND LOMB 1%N40075 001

Apr 29, 1994

CYCLOPENTOLATE HYDROCHLORIDE; PHENYLEPHRINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYCLOMYDRIL

+ ALCON 0.2%;1%

N84300 001

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDEAP BAXTER HLTHCARE 100MG/VIALN88371 001

Jul 03, 1986

AP 200MG/VIALN88372 001

Jul 03, 1986

AP 500MG/VIALN88373 001

Jul 03, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 102 (of 360)

CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYCLOPHOSPHAMIDE

<u>AP</u>	BAXTER HLTHCARE	<u>1GM/VIAL</u>	<u>N88374</u>	<u>001</u>	Sep 24, 1986
	<u>LYOPHILIZED CYTOXAN</u>				
<u>AP</u>	+ BRISTOL MYERS SQUIBB	<u>100MG/VIAL</u>	<u>N12142</u>	<u>006</u>	Dec 05, 1985
<u>AP</u>	+	<u>200MG/VIAL</u>	<u>N12142</u>	<u>007</u>	Dec 10, 1985
<u>AP</u>	+	<u>500MG/VIAL</u>	<u>N12142</u>	<u>008</u>	Jan 04, 1984
<u>AP</u>	+	<u>1GM/VIAL</u>	<u>N12142</u>	<u>010</u>	Sep 24, 1985
<u>AP</u>	+	<u>2GM/VIAL</u>	<u>N12142</u>	<u>009</u>	Dec 10, 1984
	<u>NEOSAR</u>				
<u>AP</u>	SICOR PHARMS	<u>100MG/VIAL</u>	<u>N87442</u>	<u>001</u>	Feb 16, 1982
<u>AP</u>		<u>200MG/VIAL</u>	<u>N87442</u>	<u>002</u>	Feb 16, 1982
<u>AP</u>		<u>500MG/VIAL</u>	<u>N87442</u>	<u>003</u>	Feb 16, 1982
<u>AP</u>		<u>1GM/VIAL</u>	<u>N87442</u>	<u>004</u>	Jul 08, 1983
<u>AP</u>		<u>2GM/VIAL</u>	<u>N87442</u>	<u>005</u>	Mar 30, 1989

TABLET; ORAL

CYCLOPHOSPHAMIDE

<u>AB</u>	ROXANE	<u>25MG</u>	<u>N40032</u>	<u>001</u>	Aug 17, 1999
<u>AB</u>		<u>50MG</u>	<u>N40032</u>	<u>002</u>	Aug 17, 1999
	<u>CYTOXAN</u>				
<u>AB</u>	BRISTOL MYERS SQUIBB	<u>25MG</u>	<u>N12141</u>	<u>002</u>	
<u>AB</u>	+	<u>50MG</u>	<u>N12141</u>	<u>001</u>	

CYCLOSERINE

CAPSULE; ORAL

SEROMYCIN

+ LILLY 250MG N60593 001

CYCLOSPORINE

CAPSULE; ORAL

CYCLOSPORINE

<u>AB1</u>	IVAX PHARMS	<u>25MG</u>	<u>N65110</u>	<u>003</u>	Mar 29, 2005
<u>AB1</u>		<u>50MG</u>	<u>N65110</u>	<u>001</u>	Mar 29, 2005
<u>AB1</u>		<u>100MG</u>	<u>N65110</u>	<u>002</u>	Mar 29, 2005
<u>AB1</u>	PLIVA	<u>25MG</u>	<u>N65044</u>	<u>002</u>	Dec 20, 2000
<u>AB1</u>		<u>100MG</u>	<u>N65044</u>	<u>001</u>	Dec 20, 2000
<u>AB1</u>	SANDOZ	<u>25MG</u>	<u>N65017</u>	<u>002</u>	Jan 13, 2000
<u>AB1</u>		<u>100MG</u>	<u>N65017</u>	<u>001</u>	Jan 13, 2000
<u>AB2</u>	TORPHARM	<u>25MG</u>	<u>N65040</u>	<u>001</u>	May 09, 2002
<u>AB2</u>		<u>100MG</u>	<u>N65040</u>	<u>002</u>	May 09, 2002
	<u>GENGRAF</u>				
<u>AB1</u>	ABBOTT	<u>25MG</u>	<u>N65003</u>	<u>001</u>	May 12, 2000
<u>AB1</u>		<u>50MG</u>	<u>N65003</u>	<u>002</u>	May 12, 2000
<u>AB1</u>		<u>100MG</u>	<u>N65003</u>	<u>003</u>	May 12, 2000
	<u>NEORAL</u>				
<u>AB1</u>	NOVARTIS	<u>25MG</u>	<u>N50715</u>	<u>001</u>	Jul 14, 1995
<u>AB1</u>	+	<u>100MG</u>	<u>N50715</u>	<u>002</u>	Jul 14, 1995
	<u>SANDIMMUNE</u>				
<u>AB2</u>	NOVARTIS	<u>25MG</u>	<u>N50625</u>	<u>001</u>	Mar 02, 1990
<u>AB2</u>	+	<u>100MG</u>	<u>N50625</u>	<u>002</u>	Mar 02, 1990
<u>BX</u>		50MG	N50625	003	Nov 23, 1992

EMULSION; OPHTHALMIC

RESTASIS

+ ALLERGAN 0.05% N50790 001 Dec 23, 2002

INJECTABLE; INJECTION

CYCLOSPORINE

<u>AP</u>	BEDFORD	<u>50MG/ML</u>	<u>N65004</u>	<u>001</u>	Oct 29, 1999
<u>AP</u>	PHARMAFORCE	<u>50MG/ML</u>	<u>N65151</u>	<u>001</u>	Oct 07, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 103 (of 360)

CYCLOSPORINE

INJECTABLE; INJECTION

SANDIMMUNE

<u>AP</u>	+	NOVARTIS	<u>50MG/ML</u>	<u>N50573</u>	<u>001</u>	Nov 14, 1983
-----------	---	----------	----------------	---------------	------------	--------------

SOLUTION; ORAL

CYCLOSPORINE

<u>AB1</u>		ABBOTT	<u>100MG/ML</u>	<u>N65025</u>	<u>001</u>	Mar 03, 2000
------------	--	--------	-----------------	---------------	------------	--------------

<u>AB1</u>		IVAX PHARMS	<u>100MG/ML</u>	<u>N65078</u>	<u>001</u>	Mar 25, 2005
------------	--	-------------	-----------------	---------------	------------	--------------

<u>AB2</u>		MORTON GROVE	<u>100MG/ML</u>	<u>N65133</u>	<u>001</u>	Sep 17, 2004
------------	--	--------------	-----------------	---------------	------------	--------------

<u>AB1</u>		NOVEX	<u>100MG/ML</u>	<u>N65167</u>	<u>001</u>	Jan 05, 2005
------------	--	-------	-----------------	---------------	------------	--------------

<u>AB1</u>		PLIVA	<u>100MG/ML</u>	<u>N65054</u>	<u>001</u>	Dec 18, 2001
------------	--	-------	-----------------	---------------	------------	--------------

NEORAL

<u>AB1</u>	+	NOVARTIS	<u>100MG/ML</u>	<u>N50716</u>	<u>001</u>	Jul 14, 1995
------------	---	----------	-----------------	---------------	------------	--------------

SANDIMMUNE

<u>AB2</u>	+	NOVARTIS	<u>100MG/ML</u>	<u>N50574</u>	<u>001</u>	Nov 14, 1983
------------	---	----------	-----------------	---------------	------------	--------------

CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE

+	ALPHARMA US PHARMS	2MG/5ML	N86833	001	
---	--------------------	---------	--------	-----	--

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

<u>AA</u>		COREPHARMA	<u>4MG</u>	<u>N40537</u>	<u>001</u>	Sep 30, 2003
-----------	--	------------	------------	---------------	------------	--------------

<u>AA</u>	+	IVAX PHARMS	<u>4MG</u>	<u>N87056</u>	<u>001</u>	
-----------	---	-------------	------------	---------------	------------	--

<u>AA</u>		PAR PHARM	<u>4MG</u>	<u>N87129</u>	<u>001</u>	
-----------	--	-----------	------------	---------------	------------	--

<u>AA</u>		PLIVA	<u>4MG</u>	<u>N88205</u>	<u>001</u>	Jul 26, 1983
-----------	--	-------	------------	---------------	------------	--------------

CYSTEAMINE BITARTRATE

CAPSULE; ORAL

CYSTAGON

	MYLAN	EQ 50MG BASE	N20392	001	Aug 15, 1994
--	-------	--------------	--------	-----	--------------

+		EQ 150MG BASE	N20392	002	Aug 15, 1994
---	--	---------------	--------	-----	--------------

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

<u>AP</u>		AM PHARM	<u>100MG/ML</u>	<u>N76512</u>	<u>001</u>	Jan 15, 2004
-----------	--	----------	-----------------	---------------	------------	--------------

<u>AP</u>		BEDFORD	<u>100MG/VIAL</u>	<u>N71471</u>	<u>001</u>	Aug 02, 1989
-----------	--	---------	-------------------	---------------	------------	--------------

<u>AP</u>			<u>500MG/VIAL</u>	<u>N71472</u>	<u>001</u>	Aug 02, 1989
-----------	--	--	-------------------	---------------	------------	--------------

<u>AP</u>			<u>1GM/VIAL</u>	<u>N74245</u>	<u>001</u>	Aug 31, 1994
-----------	--	--	-----------------	---------------	------------	--------------

<u>AP</u>			<u>2GM/VIAL</u>	<u>N74245</u>	<u>002</u>	Aug 31, 1994
-----------	--	--	-----------------	---------------	------------	--------------

<u>AP</u>	+	MAYNE PHARMA USA	<u>100MG/ML</u>	<u>N75383</u>	<u>001</u>	Nov 22, 1999
-----------	---	------------------	-----------------	---------------	------------	--------------

+		20MG/ML	N71868	001	Jun 04, 1990
---	--	---------	--------	-----	--------------

+		20MG/ML	N72168	001	Aug 31, 1990
---	--	---------	--------	-----	--------------

+		20MG/ML	N72945	001	Feb 28, 1994
---	--	---------	--------	-----	--------------

CYTOSAR-U

<u>AP</u>		SICOR PHARMS	<u>100MG/VIAL</u>	<u>N75206</u>	<u>001</u>	Dec 30, 1998
-----------	--	--------------	-------------------	---------------	------------	--------------

<u>AP</u>	+		<u>500MG/VIAL</u>	<u>N75206</u>	<u>002</u>	Dec 30, 1998
-----------	---	--	-------------------	---------------	------------	--------------

<u>AP</u>	+		<u>1GM/VIAL</u>	<u>N75206</u>	<u>004</u>	Dec 30, 1998
-----------	---	--	-----------------	---------------	------------	--------------

<u>AP</u>	+		<u>2GM/VIAL</u>	<u>N75206</u>	<u>003</u>	Dec 30, 1998
-----------	---	--	-----------------	---------------	------------	--------------

INJECTABLE, LIPOSOMAL; INJECTION

DEPOCYT

+	SKYEPHARMA	10MG/ML	N21041	001	Apr 01, 1999
---	------------	---------	--------	-----	--------------

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

<u>AP</u>		AM PHARM PARTNERS	<u>100MG/VIAL</u>	<u>N75371</u>	<u>001</u>	Aug 27, 1999
-----------	--	-------------------	-------------------	---------------	------------	--------------

<u>AP</u>			<u>200MG/VIAL</u>	<u>N75371</u>	<u>002</u>	Aug 27, 1999
-----------	--	--	-------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 104 (of 360)

DACARBAZINE

INJECTABLE; INJECTION

DACARBAZINE

<u>AP</u>	BEDFORD	<u>200MG/VIAL</u>	<u>N75812</u>	<u>001</u>	Jun 15, 2001
<u>AP</u>		<u>500MG/VIAL</u>	<u>N75812</u>	<u>002</u>	Oct 31, 2002
<u>AP</u>	MAYNE PHARMA USA	<u>200MG/VIAL</u>	<u>N75940</u>	<u>001</u>	Oct 18, 2001
<u>AP</u>	SICOR PHARMS	<u>200MG/VIAL</u>	<u>N75259</u>	<u>002</u>	Aug 27, 1998
<u>AP</u>	+	<u>500MG/VIAL</u>	<u>N75259</u>	<u>001</u>	Sep 22, 2000
	<u>DTIC-DOME</u>				
<u>AP</u>	+	<u>100MG/VIAL</u>	<u>N17575</u>	<u>001</u>	
<u>AP</u>	+	<u>200MG/VIAL</u>	<u>N17575</u>	<u>002</u>	

DACTINOMYCIN

INJECTABLE; INJECTION

COSMEGEN

+	OVATION PHARMS	0.5MG/VIAL	N50682	001	
---	----------------	------------	--------	-----	--

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; IV (INFUSION)

SYNERCID

+	KING PHARMS	350MG/VIAL;150MG/VIAL	N50748	001	Sep 21, 1999
---	-------------	-----------------------	--------	-----	--------------

DALTEPARIN SODIUM

INJECTABLE; INJECTION

FRAGMIN

+	PHARMACIA AND UPJOHN	2,500 IU/0.2ML	N20287	001	Dec 22, 1994
+		5,000 IU/0.2ML	N20287	003	Mar 18, 1996
+		7,500 IU/0.3ML	N20287	005	Apr 04, 2002
+		10,000 IU/ML	N20287	004	Jan 30, 1998
+		95,000IU/3.8ML (25,000IU/ML)	N20287	006	Apr 04, 2002
+		95,000IU/9.5ML (10,000IU/ML)	N20287	007	Apr 04, 2002

DANAZOL

CAPSULE; ORAL

DANAZOL

<u>AB</u>	+	BARR	<u>200MG</u>	<u>N74582</u>	<u>001</u>	Aug 09, 1996
			50MG	N74582	003	May 29, 1998
			100MG	N74582	002	May 29, 1998
<u>AB</u>		LANNETT	<u>200MG</u>	<u>N77246</u>	<u>001</u>	Sep 28, 2005

DANTROLENE SODIUM

CAPSULE; ORAL

DANTRIUM

<u>AB</u>		PROCTER AND GAMBLE	<u>25MG</u>	<u>N17443</u>	<u>001</u>	
<u>AB</u>			<u>50MG</u>	<u>N17443</u>	<u>003</u>	
<u>AB</u>	+		<u>100MG</u>	<u>N17443</u>	<u>002</u>	
		<u>DANTROLENE SODIUM</u>				
<u>AB</u>		AMIDE PHARM	<u>25MG</u>	<u>N76686</u>	<u>001</u>	Oct 24, 2005
<u>AB</u>			<u>50MG</u>	<u>N76686</u>	<u>002</u>	Oct 24, 2005
<u>AB</u>			<u>100MG</u>	<u>N76686</u>	<u>003</u>	Oct 24, 2005
<u>AB</u>		IMPAX LABS	<u>25MG</u>	<u>N76856</u>	<u>001</u>	Mar 01, 2005
<u>AB</u>			<u>50MG</u>	<u>N76856</u>	<u>002</u>	Mar 01, 2005
<u>AB</u>			<u>100MG</u>	<u>N76856</u>	<u>003</u>	Mar 01, 2005

INJECTABLE; INJECTION

DANTRIUM

+	PROCTER AND GAMBLE	20MG/VIAL	N18264	001	
---	--------------------	-----------	--------	-----	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 105 (of 360)

DAPIPIRAZOLE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC
REV-EYES

+ ANGELINI	0.5%	N19849 001	Dec 31, 1990
------------	------	------------	--------------

DAPSONE

TABLET; ORAL

DAPSONE

JACOBUS

25MG

N86841 001

+

100MG

N86842 001

DAPTOMYCIN

INJECTABLE; IV (INFUSION)

CUBICIN

CUBIST

250MG/VIAL

N21572 001

Sep 12, 2003

+

500MG/VIAL

N21572 002

Sep 12, 2003

DARIFENACIN HYDROBROMIDE

TABLET, EXTENDED RELEASE; ORAL

ENABLEX

NOVARTIS

EQ 7.5MG BASE

N21513 001

Dec 22, 2004

+

EQ 15MG BASE

N21513 002

Dec 22, 2004

DAUNORUBICIN CITRATE

INJECTABLE, LIPOSOMAL; INJECTION

DAUNOXOME

+ GILEAD

EQ 2MG BASE/ML

N50704 002

Apr 08, 1996

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

CERUBIDINE

AP + BEDFORD

EQ 20MG BASE/VIAL

N64103 001

Feb 03, 1995

DAUNORUBICIN HYDROCHLORIDE

AP AM PHARM

EQ 5MG BASE/VIAL

N65034 001

Nov 20, 2001

AP AM PHARM PARTNERS

EQ 20MG BASE/VIAL

N65000 001

May 25, 1999

AP + BEDFORD

EQ 5MG BASE/ML

N50731 001

Jan 30, 1998

AP SICOR PHARMS

EQ 5MG BASE/ML

N65035 001

Jan 24, 2000

AP

EQ 20MG BASE/VIAL

N64212 001

Jun 23, 1998

+

EQ 50MG BASE/VIAL

N64212 002

May 03, 1999

DEFERASIROX

TABLET, FOR SUSPENSION; ORAL

EXJADE

NOVARTIS

125MG

N21882 001

Nov 02, 2005

250MG

N21882 002

Nov 02, 2005

+

500MG

N21882 003

Nov 02, 2005

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

AP HOSPIRA

500MG/VIAL

N76019 001

Mar 17, 2004

AP

2GM/VIAL

N76019 002

Mar 17, 2004

DESFERAL

AP + NOVARTIS

500MG/VIAL

N16267 001

AP +

2GM/VIAL

N16267 002

May 25, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 106 (of 360)

DELAVIDINE MESYLATETABLET; ORAL
RESCRIPTOR

AGOURON	100MG	N20705	001	Apr 04, 1997
+	200MG	N20705	002	Jul 14, 1999

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DECLOMYCINAB ESP PHARMA 150MGN50261 002AB + 300MGN50261 003DEMECLOCYCLINE HYDROCHLORIDEAB BARR 150MGN65171 001

Dec 13, 2004

AB 300MGN65171 002

Dec 13, 2004

AB IMPAX LABS 150MGN65094 001

Mar 22, 2004

AB 300MGN65094 002

Mar 22, 2004

DESFLURANE

LIQUID; INHALATION

SUPRANE

+	BAXTER HLTHCARE CORP	99.9%	N20118	001	Sep 18, 1992
---	----------------------	-------	--------	-----	--------------

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDEAB PLIVA 25MGN71800 001

Dec 08, 1987

AB 50MGN71801 001

Dec 08, 1987

AB 75MGN71802 001

Dec 08, 1987

AB 100MGN71803 001

May 29, 1997

AB 150MGN71804 001

May 29, 1997

AB SANDOZ 10MGN72099 001

May 24, 1988

AB 10MGN74430 001

Feb 09, 1996

AB 25MGN71601 001

Jun 05, 1987

AB 25MGN72100 001

May 24, 1988

AB 50MGN71588 001

Jun 05, 1987

AB 50MGN72101 001

May 24, 1988

AB 75MGN71602 001

Oct 05, 1987

AB 75MGN72102 001

Jun 20, 1988

AB 100MGN71766 001

Oct 05, 1987

AB 100MGN72103 001

Jun 20, 1988

AB 150MGN72104 001

Jun 20, 1988

AB 150MGN74430 002

Feb 09, 1996

NORPRAMINAB AVENTIS PHARMS 10MGN14399 007

Feb 11, 1982

AB 25MGN14399 001AB + 50MGN14399 003AB 75MGN14399 004AB + 100MGN14399 005AB 150MGN14399 006DESIRUDIN RECOMBINANT

INJECTABLE; SUBCUTANEOUS

IPRIVASK

+	CANYON	15MG/VIAL	N21271	001	Apr 04, 2003
---	--------	-----------	--------	-----	--------------

DESLORATADINE

SYRUP; ORAL

CLARINEX

+	SCHERING	0.5MG/ML	N21300	001	Sep 01, 2004
---	----------	----------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 107 (of 360)

DESLORATADINE

TABLET; ORAL

CLARINEX

+	SCHERING PLOUGH	5MG	N21165	001	Dec 21, 2001
---	-----------------	-----	--------	-----	--------------

TABLET, ORALLY DISINTEGRATING; ORAL

CLARINEX

	SCHERING	2.5MG	N21312	002	Jul 14, 2005
--	----------	-------	--------	-----	--------------

+		5MG	N21312	001	Jun 26, 2002
---	--	-----	--------	-----	--------------

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARINEX D 24 HOUR

+	SCHERING	5MG;240MG	N21605	001	Mar 03, 2005
---	----------	-----------	--------	-----	--------------

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

<u>AP</u>	+	AVENTIS	<u>0.004MG/ML</u>	<u>N18938</u>	<u>001</u>	Mar 30, 1984
-----------	---	---------	-------------------	---------------	------------	--------------

+		0.015MG/ML	N18938	002	Apr 25, 1995
---	--	------------	--------	-----	--------------

DESMOPRESSIN ACETATE

<u>AP</u>		BEDFORD	<u>0.004MG/ML</u>	<u>N74575</u>	<u>001</u>	Feb 18, 2000
-----------	--	---------	-------------------	---------------	------------	--------------

<u>AP</u>		HOSPIRA	<u>0.004MG/ML</u>	<u>N75220</u>	<u>001</u>	Aug 28, 2000
-----------	--	---------	-------------------	---------------	------------	--------------

<u>AP</u>		SICOR PHARMS	<u>0.004MG/ML</u>	<u>N74888</u>	<u>001</u>	Oct 15, 1997
-----------	--	--------------	-------------------	---------------	------------	--------------

DESMOPRESSIN ACETATE PRESERVATIVE FREE

<u>AP</u>		BEDFORD	<u>0.004MG/ML</u>	<u>N74574</u>	<u>001</u>	Feb 18, 2000
-----------	--	---------	-------------------	---------------	------------	--------------

SOLUTION; NASAL

DDAVP

+	AVENTIS	0.01%	N17922	001	
---	---------	-------	--------	-----	--

SPRAY, METERED; NASAL

DDAVP (NEEDS NO REFRIGERATION)

<u>AB</u>	+	AVENTIS	<u>0.01MG/SPRAY</u>	<u>N17922</u>	<u>003</u>	Aug 07, 1996
-----------	---	---------	---------------------	---------------	------------	--------------

DESMOPRESSIN ACETATE

<u>AB</u>	+	BAUSCH AND LOMB	<u>0.01MG/SPRAY</u>	<u>N74830</u>	<u>001</u>	Jan 25, 1999
-----------	---	-----------------	---------------------	---------------	------------	--------------

DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION)

<u>AB</u>		APOTEX	<u>0.01MG/SPRAY</u>	<u>N76703</u>	<u>001</u>	Jan 27, 2005
-----------	--	--------	---------------------	---------------	------------	--------------

MINIRIN

<u>AB</u>	+	FERRING	<u>0.01MG/SPRAY</u>	<u>N21333</u>	<u>001</u>	Sep 16, 2002
-----------	---	---------	---------------------	---------------	------------	--------------

STIMATE

+	ZLB BEHRING	0.15MG/SPRAY	N20355	001	Mar 07, 1994
---	-------------	--------------	--------	-----	--------------

TABLET; ORAL

DDAVP

<u>AB</u>		AVENTIS	<u>0.1MG</u>	<u>N19955</u>	<u>001</u>	Sep 06, 1995
-----------	--	---------	--------------	---------------	------------	--------------

<u>AB</u>	+		<u>0.2MG</u>	<u>N19955</u>	<u>002</u>	Sep 06, 1995
-----------	---	--	--------------	---------------	------------	--------------

DESMOPRESSIN ACETATE

<u>AB</u>		BARR	<u>0.1MG</u>	<u>N76470</u>	<u>001</u>	Jul 01, 2005
-----------	--	------	--------------	---------------	------------	--------------

<u>AB</u>			<u>0.2MG</u>	<u>N76470</u>	<u>002</u>	Jul 01, 2005
-----------	--	--	--------------	---------------	------------	--------------

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-21

DESOGESTREL AND ETHINYL ESTRADIOL

<u>AB</u>		DURAMED PHARMS BARR	<u>0.15MG;0.03MG</u>	<u>N75256</u>	<u>001</u>	Aug 12, 1999
-----------	--	---------------------	----------------------	---------------	------------	--------------

TABLET; ORAL-28

CYCLESSA

<u>AB</u>	+	ORGANON USA INC	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u>	<u>N21090</u>	<u>001</u>	Dec 20, 2000
-----------	---	-----------------	---	---------------	------------	--------------

DESOGEN

<u>AB</u>		ORGANON USA INC	<u>0.15MG;0.03MG</u>	<u>N20071</u>	<u>002</u>	Dec 10, 1992
-----------	--	-----------------	----------------------	---------------	------------	--------------

DESOGESTREL AND ETHINYL ESTRADIOL

<u>AB</u>		DURAMED PHARMS BARR	<u>0.15MG;0.03MG</u>	<u>N75256</u>	<u>002</u>	Aug 12, 1999
-----------	--	---------------------	----------------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 108 (of 360)

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-28

<u>DESOGESTREL AND ETHINYL ESTRADIOL</u>				
<u>AB</u>	<u>WATSON LABS</u>	<u>0.15MG;0.03MG</u>	<u>N76915</u> <u>001</u>	Jul 29, 2005
<u>AB</u>	<u>KARIVA</u>			
<u>AB</u>	<u>BARR</u>	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>N75863</u> <u>001</u>	Apr 05, 2002
<u>AB</u>	<u>MIRCETTE</u>			
<u>AB</u>	<u>+ ORGANON USA INC</u>	<u>0.15MG,N/A;0.02MG,0.01MG</u>	<u>N20713</u> <u>001</u>	Apr 22, 1998
<u>AB</u>	<u>ORTHO-CEPT</u>			
<u>AB</u>	<u>+ ORTHO MCNEIL PHARM</u>	<u>0.15MG;0.03MG</u>	<u>N20301</u> <u>002</u>	Dec 14, 1992
<u>AB</u>	<u>VELIVET</u>			
<u>AB</u>	<u>DURAMED PHARMS BARR</u>	<u>0.1MG,0.125MG,0.15MG;0.025MG,0.025MG,0.025MG</u>	<u>N76455</u> <u>001</u>	Feb 24, 2004

DESONIDE

CREAM; TOPICAL

<u>DESONIDE</u>				
<u>AB</u>	<u>TARO</u>	<u>0.05%</u>	<u>N73548</u> <u>001</u>	Jun 30, 1992
<u>AB</u>	<u>TEVA PHARMS</u>	<u>0.05%</u>	<u>N74027</u> <u>001</u>	Sep 28, 1992
<u>AB</u>	<u>DESOWEN</u>			
<u>AB</u>	<u>GALDERMA LABS LP</u>	<u>0.05%</u>	<u>N19048</u> <u>001</u>	Dec 14, 1984
<u>AB</u>	<u>TRIDESILON</u>			
<u>AB</u>	<u>+ CLAY PARK</u>	<u>0.05%</u>	<u>N17010</u> <u>001</u>	

LOTION; TOPICAL

<u>DESONIDE</u>				
<u>AB</u>	<u>ALTANA</u>	<u>0.05%</u>	<u>N75860</u> <u>001</u>	Mar 19, 2002
<u>AB</u>	<u>DESOWEN</u>			
<u>AB</u>	<u>+ GALDERMA LABS LP</u>	<u>0.05%</u>	<u>N72354</u> <u>001</u>	Jan 24, 1992

OINTMENT; TOPICAL

<u>DESONIDE</u>				
<u>AB</u>	<u>ALTANA</u>	<u>0.05%</u>	<u>N75751</u> <u>001</u>	Mar 12, 2001
<u>AB</u>	<u>TARO</u>	<u>0.05%</u>	<u>N74254</u> <u>001</u>	Aug 03, 1994
<u>AB</u>	<u>DESOWEN</u>			
<u>AB</u>	<u>GALDERMA LABS LP</u>	<u>0.05%</u>	<u>N71425</u> <u>001</u>	Jun 15, 1988
<u>AB</u>	<u>TRIDESILON</u>			
<u>AB</u>	<u>+ CLAY PARK</u>	<u>0.05%</u>	<u>N17426</u> <u>001</u>	

DESOXIMETASONE

CREAM; TOPICAL

<u>DESOXIMETASONE</u>				
<u>AB</u>	<u>CLAY PARK</u>	<u>0.25%</u>	<u>N76510</u> <u>001</u>	Jul 01, 2003
<u>AB</u>	<u>TARO</u>	<u>0.05%</u>	<u>N73210</u> <u>001</u>	Nov 30, 1990
<u>AB</u>		<u>0.25%</u>	<u>N73193</u> <u>001</u>	Nov 30, 1990
<u>AB</u>	<u>TOPICORT</u>			
<u>AB</u>	<u>+ TARO PHARMS NORTH</u>	<u>0.25%</u>	<u>N17856</u> <u>001</u>	
<u>AB</u>	<u>TOPICORT LP</u>			
<u>AB</u>	<u>+ TARO PHARMS NORTH</u>	<u>0.05%</u>	<u>N18309</u> <u>001</u>	

GEL; TOPICAL

<u>DESOXIMETASONE</u>				
<u>AB</u>	<u>CLAY PARK</u>	<u>0.05%</u>	<u>N77552</u> <u>001</u>	Jan 09, 2006
<u>AB</u>	<u>TARO</u>	<u>0.05%</u>	<u>N74904</u> <u>001</u>	Jul 14, 1998
<u>AB</u>	<u>TOPICORT</u>			
<u>AB</u>	<u>+ TARO PHARMS NORTH</u>	<u>0.05%</u>	<u>N18586</u> <u>001</u>	Mar 29, 1982

OINTMENT; TOPICAL

<u>DESOXIMETASONE</u>				
<u>AB</u>	<u>TARO</u>	<u>0.25%</u>	<u>N74286</u> <u>001</u>	Jun 07, 1996
<u>AB</u>	<u>TOPICORT</u>			
<u>AB</u>	<u>+ TARO PHARMS NORTH</u>	<u>0.25%</u>	<u>N18763</u> <u>001</u>	Sep 30, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 109 (of 360)

DEXAMETHASONE

CONCENTRATE; ORAL

DEXAMETHASONE INTENSOL

+	ROXANE	1MG/ML	N88252	001	Sep 01, 1983
---	--------	--------	--------	-----	--------------

ELIXIR; ORAL

DEXAMETHASONE

<u>AA</u>	ALPHARMA US PHARMS	<u>0.5MG/5ML</u>	<u>N84754</u>	<u>001</u>	
-----------	--------------------	------------------	---------------	------------	--

HEXADROL

<u>AA</u>	+	ORGANON USA INC	<u>0.5MG/5ML</u>	<u>N12674</u>	<u>001</u>
-----------	---	-----------------	------------------	---------------	------------

MYMETHASONE

<u>AA</u>	MORTON GROVE	<u>0.5MG/5ML</u>	<u>N88254</u>	<u>001</u>	Jul 27, 1983
-----------	--------------	------------------	---------------	------------	--------------

SOLUTION; ORAL

DEXAMETHASONE

+	ROXANE	0.5MG/5ML	N88248	001	Sep 01, 1983
---	--------	-----------	--------	-----	--------------

SUSPENSION/DROPS; OPHTHALMIC

MAXIDEX

+	ALCON	0.1%	N13422	001	
---	-------	------	--------	-----	--

TABLET; ORAL

DECADRON

<u>AB</u>	MERCK	<u>0.5MG</u>	<u>N11664</u>	<u>001</u>	
-----------	-------	--------------	---------------	------------	--

<u>AB</u>	+		<u>0.75MG</u>	<u>N11664</u>	<u>002</u>
-----------	---	--	---------------	---------------	------------

DEXAMETHASONE

BP	PAR PHARM	0.5MG	N88148	001	Apr 28, 1983
----	-----------	-------	--------	-----	--------------

BP		0.75MG	N88160	001	Apr 28, 1983
----	--	--------	--------	-----	--------------

BP		1.5MG	N88237	001	Apr 28, 1983
----	--	-------	--------	-----	--------------

BP		4MG	N88238	001	Apr 28, 1983
----	--	-----	--------	-----	--------------

BP		6MG	N88481	001	Nov 28, 1983
----	--	-----	--------	-----	--------------

		0.25MG	N88149	001	Apr 28, 1983
--	--	--------	--------	-----	--------------

<u>AB</u>	ROXANE	<u>0.5MG</u>	<u>N84611</u>	<u>001</u>	
-----------	--------	--------------	---------------	------------	--

<u>AB</u>		<u>0.75MG</u>	<u>N84613</u>	<u>001</u>	
-----------	--	---------------	---------------	------------	--

BP		1.5MG	N84610	001	
----	--	-------	--------	-----	--

BP		4MG	N84612	001	
----	--	-----	--------	-----	--

BP	+	6MG	N88316	001	Sep 15, 1983
----	---	-----	--------	-----	--------------

		1MG	N88306	001	Sep 15, 1983
--	--	-----	--------	-----	--------------

		2MG	N87916	001	Aug 26, 1982
--	--	-----	--------	-----	--------------

HEXADROL

BP	ORGANON USA INC	4MG	N12675	010	
----	-----------------	-----	--------	-----	--

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 4MG PHOSPHATE/ML</u>	<u>N84282</u>	<u>001</u>	
-----------	-----------------	----------------------------	---------------	------------	--

<u>AP</u>	ELKINS SINN	<u>EQ 10MG PHOSPHATE/ML</u>	<u>N87702</u>	<u>001</u>	Sep 07, 1982
-----------	-------------	-----------------------------	---------------	------------	--------------

DEXAMETHASONE SODIUM PHOSPHATE

<u>AP</u>	AM PHARM	<u>EQ 10MG PHOSPHATE/ML</u>	<u>N40491</u>	<u>001</u>	Apr 11, 2003
-----------	----------	-----------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 10MG PHOSPHATE/ML</u>	<u>N40572</u>	<u>001</u>	Apr 22, 2005
-----------	--	-----------------------------	---------------	------------	--------------

<u>AP</u>	AM PHARM PARTNERS	<u>EQ 4MG PHOSPHATE/ML</u>	<u>N84916</u>	<u>001</u>	
-----------	-------------------	----------------------------	---------------	------------	--

<u>AP</u>	+	LUITPOLD	<u>EQ 4MG PHOSPHATE/ML</u>	<u>N87440</u>	<u>001</u>
-----------	---	----------	----------------------------	---------------	------------

<u>AP</u>	+	SICOR PHARMS	<u>EQ 10MG PHOSPHATE/ML</u>	<u>N81126</u>	<u>001</u>
-----------	---	--------------	-----------------------------	---------------	------------

SOLUTION/DROPS; OPHTHALMIC, OTIC

DEXAMETHASONE SODIUM PHOSPHATE

<u>AT</u>	+	ALCON UNIVERSAL	<u>EQ 0.1% PHOSPHATE</u>	<u>N88771</u>	<u>001</u>
-----------	---	-----------------	--------------------------	---------------	------------

<u>AT</u>		BAUSCH AND LOMB	<u>EQ 0.1% PHOSPHATE</u>	<u>N40069</u>	<u>001</u>
-----------	--	-----------------	--------------------------	---------------	------------

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

MAXITROL

<u>AT</u>	+	FALCON PHARMS	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N50065</u>	<u>002</u>
-----------	---	---------------	--	---------------	------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 110 (of 360)

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE

<u>AT</u>	BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N64063</u>	<u>001</u>	Jul 25, 1994
<u>AT</u>	FOUGERA	<u>0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM</u>	<u>N62938</u>	<u>001</u>	Jul 31, 1989

SUSPENSION/DROPS; OPHTHALMIC

DEXASPORIN

<u>AT</u>	BAUSCH AND LOMB	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N64135</u>	<u>001</u>	Sep 13, 1995
-----------	-----------------	--	---------------	------------	--------------

MAXITROL

<u>AT</u>	ALCON	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N62341</u>	<u>001</u>	May 22, 1984
<u>AT</u>	+ FALCON PHARMS	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N50023</u>	<u>002</u>	

NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE

<u>AT</u>	ALCON UNIVERSAL	<u>0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N62721</u>	<u>001</u>	Nov 17, 1986
-----------	-----------------	--	---------------	------------	--------------

DEXAMETHASONE; TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBRADEX

	+ ALCON	0.1%;0.3%	N50616	001	Sep 28, 1988
--	---------	-----------	--------	-----	--------------

SUSPENSION/DROPS; OPHTHALMIC

TOBRADEX

<u>AB</u>	+ ALCON	<u>0.1%;0.3%</u>	<u>N50592</u>	<u>001</u>	Aug 18, 1988
-----------	---------	------------------	---------------	------------	--------------

TOBRAMYCIN AND DEXAMETHASONE

<u>AB</u>	BAUSCH AND LOMB	<u>0.1%;0.3%</u>	<u>N64134</u>	<u>001</u>	Oct 27, 1999
-----------	-----------------	------------------	---------------	------------	--------------

DEXCHLORPHENIRAMINE MALEATE

SYRUP; ORAL

MYLARAMINE

	+ MORTON GROVE	2MG/5ML	N88251	001	Mar 23, 1984
--	----------------	---------	--------	-----	--------------

TABLET; ORAL

DEXCHLORPHENIRAMINE MALEATE

	+ PLIVA	2MG	N88682	001	Jan 17, 1986
--	---------	-----	--------	-----	--------------

DEXMEDETOMIDINE

INJECTABLE; INJECTION

PRECEDEX

	+ HOSPIRA	EQ 100UGM BASE/ML	N21038	001	Dec 17, 1999
--	-----------	-------------------	--------	-----	--------------

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

FOCALIN XR

	NOVARTIS	5MG	N21802	001	May 26, 2005
		10MG	N21802	002	May 26, 2005
	+	20MG	N21802	003	May 26, 2005

TABLET; ORAL

FOCALIN

	NOVARTIS	2.5MG	N21278	001	Nov 13, 2001
		5MG	N21278	002	Nov 13, 2001
	+	10MG	N21278	003	Nov 13, 2001

DEXRAZOXANE HYDROCHLORIDE

INJECTABLE; INJECTION

DEXRAZOXANE

<u>AP</u>	BEDFORD	<u>EQ 250MG BASE/VIAL</u>	<u>N76068</u>	<u>001</u>	Sep 28, 2004
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>N76068</u>	<u>002</u>	Sep 28, 2004

ZINECARD

<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>EQ 250MG BASE/VIAL</u>	<u>N20212</u>	<u>001</u>	May 26, 1995
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>N20212</u>	<u>002</u>	May 26, 1995

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 111 (of 360)

DEXTROAMPHETAMINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

DEXEDRINE

<u>AB</u>	GLAXOSMITHKLINE	<u>5MG</u>	<u>N17078</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>N17078</u>	<u>002</u>	
<u>AB</u>	+	<u>15MG</u>	<u>N17078</u>	<u>003</u>	

DEXTROAMPHETAMINE SULFATE

<u>AB</u>	BARR	<u>5MG</u>	<u>N76137</u>	<u>001</u>	Jan 18, 2002
<u>AB</u>		<u>10MG</u>	<u>N76137</u>	<u>002</u>	Jan 18, 2002
<u>AB</u>		<u>15MG</u>	<u>N76137</u>	<u>003</u>	Jan 18, 2002
<u>AB</u>	MALLINCKRODT	<u>5MG</u>	<u>N76353</u>	<u>001</u>	May 06, 2003
<u>AB</u>		<u>10MG</u>	<u>N76353</u>	<u>002</u>	May 06, 2003
<u>AB</u>		<u>15MG</u>	<u>N76353</u>	<u>003</u>	May 06, 2003

TABLET; ORAL

DEXEDRINE

<u>AA</u>	GLAXOSMITHKLINE	<u>5MG</u>	<u>N84935</u>	<u>001</u>	
-----------	-----------------	------------	---------------	------------	--

DEXTROAMPHETAMINE SULFATE

<u>AA</u>	BARR	<u>5MG</u>	<u>N40361</u>	<u>001</u>	Jan 31, 2001
<u>AA</u>		<u>10MG</u>	<u>N40361</u>	<u>002</u>	Jan 31, 2001
<u>AA</u>	ENDO PHARMS	<u>5MG</u>	<u>N40299</u>	<u>001</u>	May 13, 1999
<u>AA</u>	KV PHARM	<u>5MG</u>	<u>N40365</u>	<u>001</u>	Oct 31, 2002
<u>AA</u>		<u>10MG</u>	<u>N40367</u>	<u>001</u>	Oct 31, 2002
<u>AA</u>	MALLINCKRODT	<u>5MG</u>	<u>N40436</u>	<u>001</u>	Jan 29, 2002
<u>AA</u>		<u>10MG</u>	<u>N40436</u>	<u>002</u>	Jan 29, 2002

DEXTROSTAT

<u>AA</u>	+	SHIRE RICHWOOD	<u>5MG</u>	<u>N84051</u>	<u>001</u>
<u>AA</u>	+		<u>10MG</u>	<u>N84051</u>	<u>002</u>

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH W/ DEXTROMETHORPHAN

<u>AA</u>	+	ALPHARMA US PHARMS	<u>15MG/5ML; 6.25MG/5ML</u>	<u>N88762</u>	<u>001</u>	Oct 31, 1984
-----------	---	--------------------	-----------------------------	---------------	------------	--------------

PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE

<u>AA</u>		HI TECH PHARMA	<u>15MG/5ML; 6.25MG/5ML</u>	<u>N40027</u>	<u>001</u>	Jul 31, 1996
-----------	--	----------------	-----------------------------	---------------	------------	--------------

PROMETHAZINE W/ DEXTROMETHORPHAN

<u>AA</u>		MORTON GROVE	<u>15MG/5ML; 6.25MG/5ML</u>	<u>N88864</u>	<u>001</u>	Jan 04, 1985
-----------	--	--------------	-----------------------------	---------------	------------	--------------

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 10% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>10GM/100ML</u>	<u>N19626</u>	<u>004</u>	Feb 02, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>10GM/100ML</u>	<u>N16694</u>	<u>001</u>	
<u>AP</u>	+	HOSPIRA	<u>10GM/100ML</u>	<u>N18080</u>	<u>001</u>	

DEXTROSE 20% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>20GM/100ML</u>	<u>N17521</u>	<u>004</u>	
<u>AP</u>	+	HOSPIRA	<u>20GM/100ML</u>	<u>N18564</u>	<u>001</u>	Mar 23, 1982

DEXTROSE 25%

+	HOSPIRA	250MG/ML	N19445	002	Nov 23, 1998
---	---------	----------	--------	-----	--------------

DEXTROSE 30% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>30GM/100ML</u>	<u>N17521</u>	<u>003</u>	
<u>AP</u>	+	HOSPIRA	<u>30GM/100ML</u>	<u>N19345</u>	<u>001</u>	Jan 26, 1985

DEXTROSE 40% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>40GM/100ML</u>	<u>N17521</u>	<u>002</u>	
<u>AP</u>	+	HOSPIRA	<u>40GM/100ML</u>	<u>N18562</u>	<u>001</u>	Mar 23, 1982

DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	B BRAUN	<u>50MG/ML</u>	<u>N16730</u>	<u>002</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N16730</u>	<u>001</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N19626</u>	<u>002</u>	Feb 02, 1988
<u>AP</u>	+	BAXTER HLTHCARE	<u>50MG/ML</u>	<u>N16673</u>	<u>003</u>	Oct 30, 1985
<u>AP</u>	+		<u>50MG/ML</u>	<u>N20179</u>	<u>002</u>	Dec 07, 1992

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 112 (of 360)

DEXTROSE

INJECTABLE; INJECTION

DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>5GM/100ML</u>	<u>N16673</u>	<u>001</u>	
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N20179</u>	<u>001</u>	Dec 07, 1992
<u>AP</u>	+	DHL	<u>5GM/100ML</u>	<u>N19971</u>	<u>001</u>	Sep 28, 1995
<u>AP</u>	+	HOSPIRA	<u>50MG/ML</u>	<u>N16367</u>	<u>002</u>	
<u>AP</u>	+		<u>50MG/ML</u>	<u>N19222</u>	<u>001</u>	Jul 13, 1984
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N19466</u>	<u>001</u>	Jul 15, 1985
<u>AP</u>	+		<u>5GM/100ML</u>	<u>N19479</u>	<u>001</u>	Sep 17, 1985

DEXTROSE 50% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>50GM/100ML</u>	<u>N17521</u>	<u>001</u>	
<u>AP</u>	+		<u>50GM/100ML</u>	<u>N20047</u>	<u>001</u>	Jul 02, 1991
<u>AP</u>		HOSPIRA	<u>500MG/ML</u>	<u>N19445</u>	<u>001</u>	Jun 03, 1986
<u>AP</u>	+		<u>50GM/100ML</u>	<u>N18563</u>	<u>001</u>	Mar 23, 1982
<u>AP</u>	+		<u>50GM/100ML</u>	<u>N19894</u>	<u>001</u>	Dec 26, 1989

DEXTROSE 60% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>60GM/100ML</u>	<u>N17521</u>	<u>005</u>	Mar 26, 1982
<u>AP</u>	+	HOSPIRA	<u>60GM/100ML</u>	<u>N19346</u>	<u>001</u>	Jan 25, 1985

DEXTROSE 70% IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>70GM/100ML</u>	<u>N17521</u>	<u>006</u>	Mar 26, 1982
<u>AP</u>	+		<u>70GM/100ML</u>	<u>N20047</u>	<u>003</u>	Jul 02, 1991
<u>AP</u>	+	HOSPIRA	<u>70GM/100ML</u>	<u>N18561</u>	<u>001</u>	Mar 23, 1982
<u>AP</u>	+		<u>70GM/100ML</u>	<u>N19893</u>	<u>001</u>	Dec 26, 1989

DEXTROSE; MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	5GM/100ML;32MG/100ML;128MG/100ML;234MG/100ML	N17385	001	
-----------------	--	--------	-----	--

DEXTROSE; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA	5GM/100ML;21MG/100ML;128MG/100ML;234MG/100ML	N17610	001	
---------	--	--------	-----	--

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION

ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;31MG/100ML;130MG/100ML;26MG/100ML;320MG/100ML	N19873	001	Jun 10, 1993
---------	---	--------	-----	--------------

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS

INJECTABLE; INJECTION

IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA	5GM/100ML;53MG/100ML;100MG/100ML;100MG/100ML;180MG/100ML;280MG/100ML;16MG/100ML	N19515	001	May 08, 1986
---------	---	--------	-----	--------------

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO.48 IN PLASTIC CONTAINER

BAXTER HLTHCARE	5GM/100ML;31MG/100ML;141MG/100ML;20MG/100ML;12MG/100ML;260MG/100ML	N17484	001	
-----------------	--	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 113 (of 360)

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM LACTATE; SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS

INJECTABLE; INJECTION

IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER			
HOSPIRA	5GM/100ML;30MG/100ML;141MG/100ML;15MG/100ML;260MG/100ML;25MG/100ML	N19513 001	May 08, 1986

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;30MG/100ML;97MG/100ML;220MG/100ML;140MG/100ML	N19844 001	Jun 10, 1993

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

ISOLYTE S IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;30MG/100ML;37MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML	N19843 001	Aug 09, 1993
NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER			
HOSPIRA	5GM/100ML;30MG/100ML;37MG/100ML;222MG/100ML;526MG/100ML;502MG/100ML	N17609 001	
PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	5GM/100ML;30MG/100ML;37MG/100ML;368MG/100ML;526MG/100ML;502MG/100ML	N17451 001	

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION

	<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER</u>		
AP	BRISTOL MYERS SQUIBB 5GM/100ML;75MG/100ML	N17634 004	
	<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER</u>		
AP	BRISTOL MYERS SQUIBB 5GM/100ML;150MG/100ML	N17634 001	
	<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER</u>		
AP	BRISTOL MYERS SQUIBB 5GM/100ML;224MG/100ML	N17634 003	
	<u>DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER</u>		
AP	BRISTOL MYERS SQUIBB 5GM/100ML;300MG/100ML	N17634 002	
	<u>POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
AP	B BRAUN 5GM/100ML;75MG/100ML	N18744 001	Nov 09, 1982
	<u>POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
AP	B BRAUN 5GM/100ML;150MG/100ML	N18744 002	Nov 09, 1982
AP	B BRAUN 5GM/100ML;150MG/100ML	N19699 004	Sep 29, 1989
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;220MG/100ML	N18744 003	Nov 09, 1982
	<u>POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
AP	B BRAUN 5GM/100ML;300MG/100ML	N18744 004	Nov 09, 1982
AP	B BRAUN 5GM/100ML;300MG/100ML	N19699 006	Sep 29, 1989
	POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER		
	HOSPIRA 5GM/100ML;149MG/100ML	N18371 001	
	<u>POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER</u>		
AP	HOSPIRA 5GM/100ML;224MG/100ML	N18371 003	
	POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER		
	HOSPIRA 5GM/100ML;298MG/100ML	N18371 002	

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM LACTATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS

INJECTABLE; INJECTION

IONOSOL T AND DEXTROSE 5% IN PLASTIC CONTAINER			
HOSPIRA	5GM/100ML;111MG/100ML;256MG/100ML;146MG/100ML;207MG/100ML	N19514 001	May 08, 1986

PRESCRIPTION DRUG PRODUCT LIST

3 - 114 (of 360)

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;150MG/100ML;130MG/100ML;280MG/100ML;91MG/100ML	N19870	001	Jun 10, 1993
---------	--	--------	-----	--------------

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

DEXTROSE 5% AND ELECTROLYTE NO 75 IN PLASTIC CONTAINER

BAXTER HLTHCARE	5GM/100ML;205MG/100ML;100MG/100ML;120MG/100ML;220MG/100ML	N18840	001	Jun 29, 1983
-----------------	---	--------	-----	--------------

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ

AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;200MG/100ML	N18037	006	Apr 13, 1982
AP		5GM/100ML;150MG/100ML;200MG/100ML	N18037	007	Apr 13, 1982

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K)

AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;200MG/100ML	N18037	004	
----	-----------------	-----------------------------------	--------	-----	--

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ

AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;200MG/100ML	N18037	008	Apr 13, 1982
----	-----------------	-----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K)

AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;200MG/100ML	N18037	001	
----	-----------------	-----------------------------------	--------	-----	--

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ

AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;200MG/100ML	N18037	005	Apr 13, 1982
----	-----------------	-----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ

AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;200MG/100ML	N18037	009	Apr 13, 1982
----	-----------------	-----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ

AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;200MG/100ML	N18037	002	
----	-----------------	----------------------------------	--------	-----	--

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K)

AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;200MG/100ML	N18037	003	
----	-----------------	-----------------------------------	--------	-----	--

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;330MG/100ML	N18629	005	Mar 23, 1982
----	-----------------	----------------------------------	--------	-----	--------------

AP		5GM/100ML;150MG/100ML;330MG/100ML	N18629	002	Mar 23, 1982
----	--	-----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;330MG/100ML	N18629	003	Mar 23, 1982
----	-----------------	-----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML;150MG/100ML;330MG/100ML	N18629	004	Mar 23, 1982
----	-----------------	-----------------------------------	--------	-----	--------------

AP		5GM/100ML;300MG/100ML;330MG/100ML	N18629	006	Mar 23, 1982
----	--	-----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML;224MG/100ML;330MG/100ML	N18629	007	Mar 23, 1982
----	-----------------	-----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;330MG/100ML	N18629	008	Mar 23, 1982
----	-----------------	-----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML;75MG/100ML;330MG/100ML	N18629	001	Mar 23, 1982
----	-----------------	----------------------------------	--------	-----	--------------

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER

AP	BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N18008	010	
----	-----------------	-----------------------------------	--------	-----	--

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;37MG/100ML;200MG/100ML	N19630	031	Feb 17, 1988
---------	-----------------------------------	--------	-----	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;37MG/100ML;450MG/100ML	N19630	037	Feb 17, 1988
---------	-----------------------------------	--------	-----	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;37MG/100ML;900MG/100ML	N19630	043	Feb 17, 1988
---------	-----------------------------------	--------	-----	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;37MG/100ML;110MG/100ML	N19630	001	Feb 17, 1988
---------	----------------------------------	--------	-----	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;37MG/100ML;200MG/100ML	N19630	007	Feb 17, 1988
---------	----------------------------------	--------	-----	--------------

POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;37MG/100ML;330MG/100ML	N19630	013	Feb 17, 1988
---------	----------------------------------	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 115 (of 360)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION			
	POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;37MG/100ML;450MG/100ML	N19630 019	Feb 17, 1988
	POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;37MG/100ML;900MG/100ML	N19630 025	Feb 17, 1988
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;75MG/100ML;200MG/100ML	N19630 032	Feb 17, 1988
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;75MG/100ML;450MG/100ML	N19630 038	Feb 17, 1988
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;75MG/100ML;900MG/100ML	N19630 044	Feb 17, 1988
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER		
	B BRAUN 3.3GM/100ML;75MG/100ML;300MG/100ML	N19630 049	May 07, 1992
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;75MG/100ML;110MG/100ML	N19630 002	Feb 17, 1988
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
AP	B BRAUN 5GM/100ML;75MG/100ML;200MG/100ML	N19630 008	Feb 17, 1988
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER		
AP	B BRAUN 5GM/100ML;75MG/100ML;330MG/100ML	N19630 014	Feb 17, 1988
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
AP	B BRAUN 5GM/100ML;75MG/100ML;450MG/100ML	N19630 020	Feb 17, 1988
	POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
AP	B BRAUN 5GM/100ML;75MG/100ML;900MG/100ML	N19630 026	Feb 17, 1988
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;110MG/100ML;200MG/100ML	N19630 033	Feb 17, 1988
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;110MG/100ML;450MG/100ML	N19630 039	Feb 17, 1988
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;110MG/100ML;900MG/100ML	N19630 045	Feb 17, 1988
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER		
	B BRAUN 3.3GM/100ML;110MG/100ML;300MG/100ML	N19630 050	May 07, 1992
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;110MG/100ML;110MG/100ML	N19630 003	Feb 17, 1988
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;110MG/100ML;200MG/100ML	N19630 009	Feb 17, 1988
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;110MG/100ML;330MG/100ML	N19630 015	Feb 17, 1988
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;110MG/100ML;450MG/100ML	N19630 021	Feb 17, 1988
	POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;110MG/100ML;900MG/100ML	N19630 027	Feb 17, 1988
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;150MG/100ML;200MG/100ML	N19630 034	Feb 17, 1988
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;150MG/100ML;450MG/100ML	N19630 040	Feb 17, 1988
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;150MG/100ML;900MG/100ML	N19630 046	Feb 17, 1988
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER		
	B BRAUN 3.3GM/100ML;150MG/100ML;300MG/100ML	N19630 051	May 07, 1992
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;150MG/100ML;110MG/100ML	N19630 004	Feb 17, 1988
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
AP	B BRAUN 5GM/100ML;150MG/100ML;200MG/100ML	N19630 010	Feb 17, 1988
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER		
AP	B BRAUN 5GM/100ML;150MG/100ML;330MG/100ML	N19630 016	Feb 17, 1988
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
AP	B BRAUN 5GM/100ML;150MG/100ML;450MG/100ML	N19630 022	Feb 17, 1988
	POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
AP	B BRAUN 5GM/100ML;150MG/100ML;900MG/100ML	N19630 028	Feb 17, 1988
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;220MG/100ML;200MG/100ML	N19630 035	Feb 17, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 116 (of 360)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION			
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;220MG/100ML;450MG/100ML	N19630 041	Feb 17, 1988
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;220MG/100ML;900MG/100ML	N19630 047	Feb 17, 1988
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER		
	B BRAUN 3.3GM/100ML;220MG/100ML;300MG/100ML	N19630 052	May 07, 1992
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;220MG/100ML;110MG/100ML	N19630 005	Feb 17, 1988
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;220MG/100ML;200MG/100ML	N19630 011	Feb 17, 1988
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;220MG/100ML;330MG/100ML	N19630 017	Feb 17, 1988
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;220MG/100ML;450MG/100ML	N19630 023	Feb 17, 1988
	POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;220MG/100ML;900MG/100ML	N19630 029	Feb 17, 1988
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;300MG/100ML;200MG/100ML	N19630 036	Feb 17, 1988
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;300MG/100ML;450MG/100ML	N19630 042	Feb 17, 1988
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
	B BRAUN 10GM/100ML;300MG/100ML;900MG/100ML	N19630 048	Feb 17, 1988
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER		
	B BRAUN 3.3GM/100ML;300MG/100ML;300MG/100ML	N19630 053	May 07, 1992
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER		
	B BRAUN 5GM/100ML;300MG/100ML;110MG/100ML	N19630 006	Feb 17, 1988
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER		
<u>AP</u>	B BRAUN 5GM/100ML;300MG/100ML;200MG/100ML	N19630 012	Feb 17, 1988
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER		
<u>AP</u>	B BRAUN 5GM/100ML;300MG/100ML;330MG/100ML	N19630 018	Feb 17, 1988
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
<u>AP</u>	B BRAUN 5GM/100ML;300MG/100ML;450MG/100ML	N19630 024	Feb 17, 1988
	POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
<u>AP</u>	B BRAUN 5GM/100ML;300MG/100ML;900MG/100ML	N19630 030	Feb 17, 1988
	POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER		
	HOSPIRA 5GM/100ML;74.5MG/100ML;225MG/100ML	N18365 002	Jul 05, 1983
	5GM/100ML;149MG/100ML;225MG/100ML	N18365 006	Mar 28, 1988
	POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER		
	HOSPIRA 5GM/100ML;74.5MG/100ML;300MG/100ML	N18876 001	Jan 17, 1986
	5GM/100ML;149MG/100ML;300MG/100ML	N18876 006	Mar 28, 1988
	POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
<u>AP</u>	BAXTER HLTHCARE 5GM/100ML;75MG/100ML;450MG/100ML	N18008 005	Apr 28, 1982
<u>AP</u>	5GM/100ML;150MG/100ML;450MG/100ML	N18008 006	Apr 28, 1982
<u>AP</u>	HOSPIRA 5GM/100ML;74.5MG/100ML;450MG/100ML	N18362 009	Jul 05, 1983
<u>AP</u>	5GM/100ML;74.5MG/100ML;450MG/100ML	N18362 005	Mar 28, 1988
	POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
<u>AP</u>	BAXTER HLTHCARE 5GM/100ML;75MG/100ML;900MG/100ML	N19308 004	Apr 05, 1985
<u>AP</u>	5GM/100ML;150MG/100ML;900MG/100ML	N19308 002	Apr 05, 1985
<u>AP</u>	HOSPIRA 5GM/100ML;74.5MG/100ML;900MG/100ML	N19691 002	Mar 24, 1988
<u>AP</u>	5GM/100ML;149MG/100ML;900MG/100ML	N19691 004	Mar 24, 1988
	POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER		
	HOSPIRA 5GM/100ML;224MG/100ML;225MG/100ML	N18365 008	Mar 28, 1988
	POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER		
	HOSPIRA 5GM/100ML;224MG/100ML;300MG/100ML	N18876 007	Mar 28, 1988
	POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER		
<u>AP</u>	HOSPIRA 5GM/100ML;224MG/100ML;450MG/100ML	N18362 006	Mar 28, 1988
	POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER		
<u>AP</u>	HOSPIRA 5GM/100ML;224MG/100ML;900MG/100ML	N19691 006	Mar 24, 1988
	POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER		
	HOSPIRA 5GM/100ML;149MG/100ML;225MG/100ML	N18365 001	

PRESCRIPTION DRUG PRODUCT LIST

3 - 117 (of 360)

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION						
	POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;298MG/100ML;225MG/100ML	N18365	009	Mar 28,	1988
	POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;298MG/100ML;300MG/100ML	N18876	008	Mar 28,	1988
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER						
<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML;150MG/100ML;450MG/100ML</u>	<u>N18008</u>	<u>007</u>	Apr 28,	1982
<u>AP</u>	HOSPIRA	<u>5GM/100ML;149MG/100ML;450MG/100ML</u>	<u>N18362</u>	<u>010</u>	Jul 05,	1983
<u>AP</u>		<u>5GM/100ML;298MG/100ML;450MG/100ML</u>	<u>N18362</u>	<u>007</u>	Mar 28,	1988
POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER						
<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML;150MG/100ML;900MG/100ML</u>	<u>N19308</u>	<u>005</u>	Apr 05,	1985
<u>AP</u>		<u>5GM/100ML;300MG/100ML;900MG/100ML</u>	<u>N19308</u>	<u>003</u>	Apr 05,	1985
<u>AP</u>	HOSPIRA	<u>5GM/100ML;149MG/100ML;900MG/100ML</u>	<u>N19691</u>	<u>005</u>	Mar 24,	1988
<u>AP</u>		<u>5GM/100ML;298MG/100ML;900MG/100ML</u>	<u>N19691</u>	<u>008</u>	Mar 24,	1988
	POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;149MG/100ML;300MG/100ML	N18876	002	Jan 17,	1986
	POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;224MG/100ML;225MG/100ML	N18365	003	Jul 05,	1983
	POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;224MG/100ML;300MG/100ML	N18876	003	Jan 17,	1986
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER						
<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML;224MG/100ML;450MG/100ML</u>	<u>N18008</u>	<u>008</u>	Apr 28,	1982
<u>AP</u>	HOSPIRA	<u>5GM/100ML;224MG/100ML;450MG/100ML</u>	<u>N18362</u>	<u>002</u>		
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER						
<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML;224MG/100ML;900MG/100ML</u>	<u>N19308</u>	<u>006</u>	Apr 05,	1985
<u>AP</u>	HOSPIRA	<u>5GM/100ML;224MG/100ML;900MG/100ML</u>	<u>N19691</u>	<u>007</u>	Mar 24,	1988
	POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;298MG/100ML;225MG/100ML	N18365	004	Jul 05,	1983
	POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;298MG/100ML;300MG/100ML	N18876	004	Mar 28,	1988
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER						
<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML;300MG/100ML;450MG/100ML</u>	<u>N18008</u>	<u>009</u>	Apr 28,	1982
<u>AP</u>	HOSPIRA	<u>5GM/100ML;298MG/100ML;450MG/100ML</u>	<u>N18362</u>	<u>003</u>		
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER						
<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML;300MG/100ML;900MG/100ML</u>	<u>N19308</u>	<u>007</u>	Apr 05,	1985
<u>AP</u>	HOSPIRA	<u>5GM/100ML;298MG/100ML;900MG/100ML</u>	<u>N19691</u>	<u>009</u>	Mar 24,	1988
	POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;74.5MG/100ML;225MG/100ML	N18365	005	Mar 28,	1988
		5GM/100ML;149MG/100ML;225MG/100ML	N18365	007	Mar 28,	1988
	POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER					
	HOSPIRA	5GM/100ML;74.5MG/100ML;300MG/100ML	N18876	005	Mar 28,	1988
		5GM/100ML;149MG/100ML;300MG/100ML	N18876	009	Mar 28,	1988
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER						
<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML;150MG/100ML;450MG/100ML</u>	<u>N18008</u>	<u>004</u>		
<u>AP</u>	HOSPIRA	<u>5GM/100ML;74.5MG/100ML;450MG/100ML</u>	<u>N18362</u>	<u>008</u>	Mar 28,	1988
<u>AP</u>		<u>5GM/100ML;149MG/100ML;450MG/100ML</u>	<u>N18362</u>	<u>004</u>	Mar 28,	1988
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER						
<u>AP</u>	BAXTER HLTHCARE	<u>5GM/100ML;150MG/100ML;900MG/100ML</u>	<u>N19308</u>	<u>001</u>	Apr 05,	1985
<u>AP</u>	HOSPIRA	<u>5GM/100ML;74.5MG/100ML;900MG/100ML</u>	<u>N19691</u>	<u>001</u>	Mar 24,	1988
<u>AP</u>		<u>5GM/100ML;149MG/100ML;900MG/100ML</u>	<u>N19691</u>	<u>003</u>	Mar 24,	1988

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION						
	DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER					
	B BRAUN	10GM/100ML;110MG/100ML	N19631	011	Feb 24,	1988
	DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER					
	B BRAUN	10GM/100ML;200MG/100ML	N19631	012	Feb 24,	1988
	DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER					
	B BRAUN	10GM/100ML;330MG/100ML	N19631	013	Feb 24,	1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 118 (of 360)

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION					
	DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
	B BRAUN	10GM/100ML;450MG/100ML	N19631	014	Feb 24, 1988
	<u>DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>10GM/100ML;900MG/100ML</u>	<u>N19631</u>	<u>015</u>	Feb 24, 1988
AP	BAXTER HLTHCARE	<u>10GM/100ML;900MG/100ML</u>	<u>N16696</u>	<u>001</u>	
	DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER				
	B BRAUN	2.5GM/100ML;110MG/100ML	N19631	001	Feb 24, 1988
	DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER				
	B BRAUN	2.5GM/100ML;200MG/100ML	N19631	002	Feb 24, 1988
	DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER				
	B BRAUN	2.5GM/100ML;330MG/100ML	N19631	003	Feb 24, 1988
	<u>DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>2.5GM/100ML;450MG/100ML</u>	<u>N19631</u>	<u>004</u>	Feb 24, 1988
AP	BAXTER HLTHCARE	<u>2.5GM/100ML;450MG/100ML</u>	<u>N16697</u>	<u>001</u>	
AP	HOSPIRA	<u>2.5GM/100ML;450MG/100ML</u>	<u>N18096</u>	<u>001</u>	
	DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
	B BRAUN	2.5GM/100ML;900MG/100ML	N19631	005	Feb 24, 1988
	DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
	B BRAUN	3.3GM/100ML;300MG/100ML	N19631	016	Jan 19, 1990
	DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER				
	B BRAUN	5GM/100ML;110MG/100ML	N19631	006	Feb 24, 1988
	<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;200MG/100ML</u>	<u>N19631</u>	<u>007</u>	Feb 24, 1988
	DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER				
	HOSPIRA	5GM/100ML;225MG/100ML	N17606	001	
	DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
	HOSPIRA	5GM/100ML;300MG/100ML	N17799	001	
	<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;330MG/100ML</u>	<u>N19631</u>	<u>008</u>	Feb 24, 1988
	<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;450MG/100ML</u>	<u>N19631</u>	<u>009</u>	Feb 24, 1988
AP	HOSPIRA	<u>5GM/100ML;450MG/100ML</u>	<u>N17607</u>	<u>001</u>	
	<u>DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	B BRAUN	<u>5GM/100ML;900MG/100ML</u>	<u>N19631</u>	<u>010</u>	Feb 24, 1988
AP	HOSPIRA	<u>5GM/100ML;900MG/100ML</u>	<u>N17585</u>	<u>001</u>	
	<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;200MG/100ML</u>	<u>N16689</u>	<u>001</u>	
	<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;330MG/100ML</u>	<u>N16687</u>	<u>001</u>	
	<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;450MG/100ML</u>	<u>N16683</u>	<u>001</u>	
	<u>DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>				
AP	BAXTER HLTHCARE	<u>5GM/100ML;900MG/100ML</u>	<u>N16678</u>	<u>001</u>	

DIATRIZOATE MEGLUMINE

INJECTABLE; INJECTION					
	<u>HYPaque</u>				
AP	+ GE HEALTHCARE	<u>60%</u>	<u>N16403</u>	<u>001</u>	
	<u>RENO-60</u>				
AP	+ BRACCO	<u>60%</u>	<u>N10040</u>	<u>016</u>	
	RENO-DIP				
	+ BRACCO	30%	N10040	012	
SOLUTION; URETERAL					
	RENO-30				
	BRACCO	30%	N10040	021	
SOLUTION; URETHRAL					
	<u>CYSTOGRAPIN</u>				
AT	BRACCO	<u>30%</u>	<u>N10040</u>	<u>018</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 119 (of 360)

DIATRIZOATE MEGLUMINE

SOLUTION; URETHRAL
CYSTOGRAPIN DILUTE
BRACCO

18%

N10040 022 Nov 09, 1982

HYPAQUE-CYSTO

AT GE HEALTHCARE

30%

N16403 003

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

HYPAQUE-76

AP GE HEALTHCARE

66%;10%

N86505 001

MD-76R

AP + MALLINCKRODT

66%;10%

N19292 001 Sep 29, 1989

RENOCAL-76

AP BRACCO

66%;10%

N89347 001 Jun 01, 1988

RENOGRAFIN-60

+ BRACCO

52%;8%

N10040 006

RENOGRAFIN-76

AP + BRACCO

66%;10%

N10040 001

SOLUTION; ORAL, RECTAL

GASTROGRAFIN

AA + BRACCO

66%;10%

N11245 003

MD-GASTROVIEW

AA MALLINCKRODT

66%;10%

N87388 001

DIATRIZOATE MEGLUMINE; IODIPAMIDE MEGLUMINE

SOLUTION; INTRAUTERINE
SINOGRAPHIN

+ BRACCO

52.7%;26.8%

N11324 002

DIATRIZOATE SODIUM

FOR SOLUTION; ORAL, RECTAL

HYPAQUE

GE HEALTHCARE

100%

N11386 001

INJECTABLE; INJECTION

HYPAQUE

+ GE HEALTHCARE

50%

N09561 001

DIAZEPAM

CONCENTRATE; ORAL
DIAZEPAM INTENSOL

+ ROXANE

5MG/ML

N71415 001 Apr 03, 1987

GEL; RECTAL

DIASTAT

VALEANT

2.5MG/0.5ML (5MG/ML)

N20648 001 Jul 29, 1997

DIASTAT ACUDIAL

VALEANT

10MG/2ML (5MG/ML)

N20648 007 Sep 15, 2005

+

20MG/4ML (5MG/ML)

N20648 006 Sep 15, 2005

INJECTABLE; INJECTION

DIAZEPAM

AP BAXTER HLTHCARE

5MG/ML

N71309 001 Jul 17, 1987

AP

5MG/ML

N71310 001 Jul 17, 1987

AP + HOSPIRA

5MG/ML

N71583 001 Oct 13, 1987

AP

5MG/ML

N71584 001 Oct 13, 1987

AP

5MG/ML

N72079 001 Dec 20, 1988

AP

MARSAM PHARMS LLC

5MG/ML

N72370 001 Jan 29, 1993

AP

5MG/ML

N72397 001 Jan 29, 1993

AP

PARENTA PHARMS

5MG/ML

N76815 001 Apr 15, 2004

AP

STERIS

5MG/ML

N70296 001 Feb 12, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 120 (of 360)

DIAZEPAM

SOLUTION; ORAL

DIAZEPAM

+	ROXANE	5MG/5ML	N70928	001	Apr 03, 1987
---	--------	---------	--------	-----	--------------

TABLET; ORAL

DIAZEPAM

<u>AB</u>	BARR	<u>2MG</u>	<u>N70152</u>	<u>001</u>	Nov 01, 1985
<u>AB</u>		<u>5MG</u>	<u>N70153</u>	<u>001</u>	Nov 01, 1985
<u>AB</u>		<u>10MG</u>	<u>N70154</u>	<u>001</u>	Nov 01, 1985
<u>AB</u>	CLONMEL HLTHCARE	<u>2MG</u>	<u>N70226</u>	<u>001</u>	Sep 26, 1985
<u>AB</u>	IVAX PHARMS	<u>2MG</u>	<u>N71307</u>	<u>001</u>	Dec 10, 1986
<u>AB</u>		<u>5MG</u>	<u>N71321</u>	<u>001</u>	Dec 10, 1986
<u>AB</u>		<u>10MG</u>	<u>N71322</u>	<u>001</u>	Dec 10, 1986
<u>AB</u>	MYLAN	<u>2MG</u>	<u>N70323</u>	<u>001</u>	Sep 04, 1985
<u>AB</u>		<u>5MG</u>	<u>N70324</u>	<u>001</u>	Sep 04, 1985
<u>AB</u>		<u>10MG</u>	<u>N70325</u>	<u>001</u>	Sep 04, 1985
<u>AB</u>	PAR PHARM	<u>2MG</u>	<u>N70462</u>	<u>001</u>	Feb 25, 1986
<u>AB</u>		<u>5MG</u>	<u>N70463</u>	<u>001</u>	Feb 25, 1986
<u>AB</u>	PUREPAC PHARM	<u>2MG</u>	<u>N70781</u>	<u>001</u>	Mar 19, 1986
<u>AB</u>		<u>5MG</u>	<u>N70706</u>	<u>001</u>	Mar 19, 1986
<u>AB</u>		<u>10MG</u>	<u>N70707</u>	<u>001</u>	Mar 19, 1986
<u>AB</u>	SANDOZ	<u>2MG</u>	<u>N70302</u>	<u>001</u>	Dec 20, 1985
<u>AB</u>		<u>5MG</u>	<u>N70303</u>	<u>001</u>	Dec 20, 1985
<u>AB</u>		<u>10MG</u>	<u>N70304</u>	<u>001</u>	Dec 20, 1985
<u>AB</u>	WATSON LABS	<u>2MG</u>	<u>N71134</u>	<u>001</u>	Feb 03, 1987
<u>AB</u>		<u>5MG</u>	<u>N71135</u>	<u>001</u>	Feb 03, 1987
<u>AB</u>		<u>10MG</u>	<u>N71136</u>	<u>001</u>	Feb 03, 1987

VALIUM

<u>AB</u>	ROCHE	<u>2MG</u>	<u>N13263</u>	<u>002</u>	
<u>AB</u>		<u>5MG</u>	<u>N13263</u>	<u>004</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N13263</u>	<u>006</u>	

DIAZOXIDE

INJECTABLE; INJECTION

HYPERSTAT

+	SCHERING	15MG/ML	N16996	001	
---	----------	---------	--------	-----	--

SUSPENSION; ORAL

PROGLYCEM

+	BAKER NORTON	50MG/ML	N17453	001	
---	--------------	---------	--------	-----	--

DICLOFENAC POTASSIUM

TABLET; ORAL

CATAFLAM

<u>AB</u>	+	NOVARTIS	<u>50MG</u>	<u>N20142</u>	<u>002</u>	Nov 24, 1993
-----------	---	----------	-------------	---------------	------------	--------------

DICLOFENAC POTASSIUM

<u>AB</u>	MUTUAL PHARM	<u>50MG</u>	<u>N75470</u>	<u>001</u>	Feb 21, 2002
<u>AB</u>	MYLAN	<u>50MG</u>	<u>N75463</u>	<u>001</u>	Jul 26, 1999
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N75229</u>	<u>001</u>	Nov 20, 1998
<u>AB</u>		<u>50MG</u>	<u>N75582</u>	<u>001</u>	Feb 23, 2001
<u>AB</u>	TEVA	<u>50MG</u>	<u>N75219</u>	<u>001</u>	Aug 06, 1998
<u>AB</u>	TORPHARM	<u>50MG</u>	<u>N76561</u>	<u>001</u>	Mar 18, 2004
<u>AB</u>	WATSON LABS	<u>50MG</u>	<u>N75152</u>	<u>001</u>	Nov 27, 1998

DICLOFENAC SODIUM

GEL; TOPICAL

SOLARAZE

+	BIOGLAN PHARMS CORP	3%	N21005	001	Oct 16, 2000
---	---------------------	----	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 121 (of 360)

DICLOFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC
VOLTAREN

+ NOVARTIS	0.1%	N20037	001	Mar 28, 1991
------------	------	--------	-----	--------------

TABLET, DELAYED RELEASE; ORAL

DICLOFENAC SODIUM

<u>AB</u>	ALPHAPHARM	50MG	N75281	002	Feb 12, 2002
<u>AB</u>		75MG	N75281	003	Feb 12, 2002
<u>AB</u>	CARLSBAD	25MG	N75185	002	Nov 13, 1998
<u>AB</u>		50MG	N75185	003	Nov 13, 1998
<u>AB</u>		75MG	N75185	001	Nov 13, 1998
<u>AB</u>	MARTEC	50MG	N74986	001	Feb 26, 1999
<u>AB</u>		75MG	N74986	002	Feb 26, 1999
<u>AB</u>	PLIVA	50MG	N74432	002	Jul 29, 1999
<u>AB</u>		75MG	N74432	003	Jul 29, 1999
<u>AB</u>	PUREPAC PHARM	50MG	N74514	001	Mar 26, 1996
<u>AB</u>		75MG	N74514	002	Mar 26, 1996
<u>AB</u>	ROXANE	25MG	N74391	001	Jun 29, 1995
<u>AB</u>		50MG	N74391	002	Jun 29, 1995
<u>AB</u>		75MG	N74391	003	Jun 29, 1995
<u>AB</u>	+ SANDOZ	25MG	N74376	001	Sep 28, 1995
<u>AB</u>	+	50MG	N74376	002	Sep 28, 1995
<u>AB</u>	+	75MG	N74394	001	Nov 30, 1995
<u>AB</u>	TEVA	50MG	N74723	001	Mar 30, 1999
<u>AB</u>		75MG	N74390	001	Aug 15, 1996
<u>AB</u>	TEVA PHARMS	25MG	N74459	001	Jun 25, 1997
<u>AB</u>		50MG	N74459	002	Jun 25, 1997
<u>AB</u>		75MG	N74459	003	Jun 25, 1997

TABLET, EXTENDED RELEASE; ORAL

DICLOFENAC SODIUM

<u>AB</u>	BIOVAIL	100MG	N75492	001	Feb 11, 2000
<u>AB</u>	DEXCEL LTD	100MG	N76201	001	Nov 06, 2002
<u>AB</u>	MYLAN	100MG	N76152	001	Dec 13, 2001
<u>AB</u>	PUREPAC PHARM	100MG	N75910	001	Jan 07, 2002

VOLTAREN-XR

<u>AB</u>	+ NOVARTIS	100MG	N20254	001	Mar 08, 1996
-----------	------------	-------	--------	-----	--------------

DICLOFENAC SODIUM; MISOPROSTOL

TABLET, DELAYED RELEASE; ORAL

ARTHROTEC

GD SEARLE LLC	50MG;0.2MG	N20607	001	Dec 24, 1997
+	75MG;0.2MG	N20607	002	Dec 24, 1997

DICLOXACILLIN SODIUM

CAPSULE; ORAL

DICLOXACILLIN SODIUM

<u>AB</u>	SANDOZ	EQ 250MG BASE	N61454	001	
<u>AB</u>	+	EQ 500MG BASE	N61454	003	
		EQ 125MG BASE	N61454	002	
<u>AB</u>	TEVA	EQ 250MG BASE	N62286	001	Jun 03, 1982
<u>AB</u>		EQ 500MG BASE	N62286	002	Jun 03, 1982

FOR SUSPENSION; ORAL

DICLOXACILLIN SODIUM

+ APOTHECON	EQ 62.5MG BASE/5ML	N61455	001	
-------------	--------------------	--------	-----	--

DICYCLOMINE HYDROCHLORIDE

CAPSULE; ORAL

BENTYL

<u>AB</u>	+ AXCAN SCANDIPHARM	10MG	N07409	003	Oct 15, 1984
-----------	---------------------	------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 122 (of 360)

DICYCLOMINE HYDROCHLORIDE

CAPSULE; ORAL

DICYCLOMINE HYDROCHLORIDE

<u>AB</u>	LANNETT	<u>10MG</u>	<u>N84285</u>	<u>001</u>	
<u>AB</u>	MYLAN	<u>10MG</u>	<u>N40319</u>	<u>001</u>	Sep 07, 1999
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N85082</u>	<u>001</u>	Jun 19, 1986
<u>AB</u>	WEST WARD	<u>10MG</u>	<u>N40204</u>	<u>001</u>	Feb 28, 1997

INJECTABLE; INJECTION

BENTYL

<u>AP</u>	+ AXCAN SCANDIPHARM	<u>10MG/ML</u>	<u>N08370</u>	<u>001</u>	Oct 15, 1984
-----------	---------------------	----------------	---------------	------------	--------------

BENTYL PRESERVATIVE FREE

<u>AP</u>	+ AXCAN SCANDIPHARM	<u>10MG/ML</u>	<u>N08370</u>	<u>002</u>	Oct 15, 1984
-----------	---------------------	----------------	---------------	------------	--------------

DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE-FREE)

<u>AP</u>	BEDFORD	<u>10MG/ML</u>	<u>N40465</u>	<u>001</u>	Jun 30, 2003
-----------	---------	----------------	---------------	------------	--------------

SYRUP; ORAL

BENTYL

<u>AA</u>	+ AXCAN SCANDIPHARM	<u>10MG/5ML</u>	<u>N07961</u>	<u>002</u>	Oct 15, 1984
-----------	---------------------	-----------------	---------------	------------	--------------

DICYCLOMINE HYDROCHLORIDE

<u>AA</u>	MIKART	<u>10MG/5ML</u>	<u>N40169</u>	<u>001</u>	Mar 24, 2005
-----------	--------	-----------------	---------------	------------	--------------

TABLET; ORAL

BENTYL

<u>AB</u>	+ AXCAN SCANDIPHARM	<u>20MG</u>	<u>N07409</u>	<u>001</u>	Oct 15, 1984
-----------	---------------------	-------------	---------------	------------	--------------

DICYCLOMINE HYDROCHLORIDE

<u>AB</u>	LANNETT	<u>20MG</u>	<u>N40230</u>	<u>001</u>	Feb 26, 1999
<u>AB</u>	MYLAN	<u>20MG</u>	<u>N40317</u>	<u>001</u>	Sep 07, 1999
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>N85223</u>	<u>001</u>	Jul 30, 1986
<u>AB</u>	WEST WARD	<u>20MG</u>	<u>N40161</u>	<u>001</u>	Oct 01, 1996

DIDANOSINE

CAPSULE, DELAYED REL PELLETS; ORAL

DIDANOSINE

<u>AB</u>	BARR	<u>200MG</u>	<u>N77167</u>	<u>001</u>	Dec 03, 2004
<u>AB</u>		<u>250MG</u>	<u>N77167</u>	<u>002</u>	Dec 03, 2004
<u>AB</u>		<u>400MG</u>	<u>N77167</u>	<u>003</u>	Dec 03, 2004

VIDEX EC

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>200MG</u>	<u>N21183</u>	<u>002</u>	Oct 31, 2000
<u>AB</u>		<u>250MG</u>	<u>N21183</u>	<u>003</u>	Oct 31, 2000
<u>AB</u>	+	<u>400MG</u>	<u>N21183</u>	<u>004</u>	Oct 31, 2000
		125MG	N21183	001	Oct 31, 2000

FOR SOLUTION; ORAL

VIDEX

+	BRISTOL MYERS SQUIBB	10MG/ML	N20156	001	Oct 09, 1991
---	----------------------	---------	--------	-----	--------------

TABLET, CHEWABLE; ORAL

VIDEX

	BRISTOL MYERS SQUIBB	25MG	N20154	002	Oct 09, 1991
		50MG	N20154	003	Oct 09, 1991
		100MG	N20154	004	Oct 09, 1991
		150MG	N20154	005	Oct 09, 1991
+		200MG	N20154	006	Oct 28, 1999

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL

TENUATE

+	AVENTIS PHARMS	25MG	N11722	002	
---	----------------	------	--------	-----	--

TABLET, EXTENDED RELEASE; ORAL

TENUATE DOSPAN

+	AVENTIS PHARMS	75MG	N12546	001	
---	----------------	------	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 123 (of 360)

DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

<u>AB1</u>	ALTANA	<u>0.05%</u>	<u>N75187</u>	<u>001</u>	Mar 30, 1998
<u>AB2</u>		<u>0.05%</u>	<u>N76263</u>	<u>001</u>	Dec 20, 2002
<u>AB1</u>	TARO	<u>0.05%</u>	<u>N75508</u>	<u>001</u>	Apr 24, 2000

FLORONE

BX + PHARMACIA AND UPJOHN 0.05%

N17741 001

FLORONE E

<u>AB2</u>	+ PHARMACIA AND UPJOHN	<u>0.05%</u>	<u>N19259</u>	<u>001</u>	Aug 28, 1985
------------	------------------------	--------------	---------------	------------	--------------

PSORCON

<u>AB1</u>	+ DERMIK LABS	<u>0.05%</u>	<u>N20205</u>	<u>001</u>	Nov 20, 1992
------------	---------------	--------------	---------------	------------	--------------

OINTMENT; TOPICAL

DIFLORASONE DIACETATE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>N75374</u>	<u>001</u>	Apr 27, 1999
<u>AB</u>	TARO	<u>0.05%</u>	<u>N75331</u>	<u>001</u>	May 14, 1999

FLORONE

+ PHARMACIA AND UPJOHN 0.05%

N17994 001

PSORCON

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>0.05%</u>	<u>N19260</u>	<u>001</u>	Aug 28, 1985
-----------	------------------------	--------------	---------------	------------	--------------

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

<u>AB</u>	SANDOZ	<u>500MG</u>	<u>N74604</u>	<u>001</u>	Jun 10, 1996
<u>AB</u>	TEVA	<u>250MG</u>	<u>N73679</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>		<u>500MG</u>	<u>N73673</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>N74400</u>	<u>001</u>	Jul 17, 1997
<u>AB</u>		<u>500MG</u>	<u>N74400</u>	<u>002</u>	Jul 17, 1997

DOLOBID

<u>AB</u>	MERCK	<u>250MG</u>	<u>N18445</u>	<u>001</u>	Apr 19, 1982
<u>AB</u>	+	<u>500MG</u>	<u>N18445</u>	<u>002</u>	Apr 19, 1982

DIGOXIN

CAPSULE; ORAL

LANOXICAPS

SMITHKLINE BEECHAM 0.05MG

N18118 002 Jul 26, 1982

0.1MG

N18118 003 Jul 26, 1982

+

0.2MG

N18118 001 Jul 26, 1982

ELIXIR; ORAL

DIGOXIN

+ ROXANE 0.05MG/ML

N21648 001 Aug 26, 2004

INJECTABLE; INJECTION

DIGOXIN

<u>AP</u>	BAXTER HLTHCARE	<u>0.25MG/ML</u>	<u>N83391</u>	<u>001</u>	
<u>AP</u>	HOSPIRA	<u>0.25MG/ML</u>	<u>N40093</u>	<u>001</u>	May 16, 1996
<u>AP</u>		<u>0.25MG/ML</u>	<u>N40206</u>	<u>001</u>	Aug 28, 1998
<u>AP</u>	SABEX 2002	<u>0.25MG/ML</u>	<u>N40481</u>	<u>001</u>	Aug 21, 2003

DIGOXIN PEDIATRIC

<u>AP</u>	HOSPIRA	<u>0.1MG/ML</u>	<u>N40092</u>	<u>001</u>	Apr 25, 1996
-----------	---------	-----------------	---------------	------------	--------------

LANOXIN

<u>AP</u>	+ GLAXOSMITHKLINE	<u>0.25MG/ML</u>	<u>N09330</u>	<u>002</u>	
-----------	-------------------	------------------	---------------	------------	--

LANOXIN PEDIATRIC

<u>AP</u>	+ GLAXOSMITHKLINE	<u>0.1MG/ML</u>	<u>N09330</u>	<u>004</u>	
-----------	-------------------	-----------------	---------------	------------	--

TABLET; ORAL

DIGOXIN

<u>AB</u>	AMIDE PHARM	<u>0.125MG</u>	<u>N40282</u>	<u>001</u>	Dec 23, 1999
<u>AB</u>		<u>0.25MG</u>	<u>N40282</u>	<u>002</u>	Dec 23, 1999
<u>AB</u>	CARACO	<u>0.125MG</u>	<u>N76363</u>	<u>001</u>	Jan 31, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 124 (of 360)

DIGOXIN

TABLET; ORAL

DIGOXIN

<u>AB</u>	CARACO	<u>0.25MG</u>	<u>N76363</u>	<u>002</u>	Jan 31, 2003
<u>AB</u>	STEVENS J	<u>0.125MG</u>	<u>N76268</u>	<u>001</u>	Jul 26, 2002
<u>AB</u>		<u>0.25MG</u>	<u>N76268</u>	<u>002</u>	Jul 26, 2002

LANOXIN

<u>AB</u>	SMITHKLINE BEECHAM	<u>0.125MG</u>	<u>N20405</u>	<u>002</u>	Sep 30, 1997
<u>AB</u>	+	<u>0.25MG</u>	<u>N20405</u>	<u>004</u>	Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE

INJECTABLE; INJECTION

D.H.E. 45

<u>AP</u>	+	VALEANT	<u>1MG/ML</u>	<u>N05929</u>	<u>001</u>	
<u>AP</u>		BEDFORD LABS	<u>1MG/ML</u>	<u>N40453</u>	<u>001</u>	Jun 09, 2003
<u>AP</u>		PADDOCK	<u>1MG/ML</u>	<u>N40475</u>	<u>001</u>	Apr 28, 2003

SPRAY, METERED; NASAL
MIGRANAL

+	VALEANT	0.5MG/INH	N20148	001	Dec 08, 1997
---	---------	-----------	--------	-----	--------------

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CARDIZEM CD

<u>AB3</u>	BIOVAIL	<u>120MG</u>	<u>N20062</u>	<u>001</u>	Aug 10, 1992
<u>AB3</u>		<u>180MG</u>	<u>N20062</u>	<u>002</u>	Dec 27, 1991
<u>AB3</u>		<u>240MG</u>	<u>N20062</u>	<u>003</u>	Dec 27, 1991
<u>AB3</u>		<u>300MG</u>	<u>N20062</u>	<u>004</u>	Dec 27, 1991
BC	+	360MG	N20062	005	Aug 24, 1999

CARTIA XT

<u>AB3</u>	ANDRX PHARMS	<u>120MG</u>	<u>N74752</u>	<u>002</u>	Jul 09, 1998
<u>AB3</u>		<u>180MG</u>	<u>N74752</u>	<u>001</u>	Jul 09, 1998
<u>AB3</u>		<u>240MG</u>	<u>N74752</u>	<u>003</u>	Jul 09, 1998
<u>AB3</u>		<u>300MG</u>	<u>N74752</u>	<u>004</u>	Jul 09, 1998

DILACOR XR

<u>AB2</u>	WATSON LABS	<u>120MG</u>	<u>N20092</u>	<u>001</u>	May 29, 1992
<u>AB2</u>		<u>180MG</u>	<u>N20092</u>	<u>002</u>	May 29, 1992
<u>AB2</u>	+	<u>240MG</u>	<u>N20092</u>	<u>003</u>	May 29, 1992

DILT-CD

<u>AB3</u>	TORPHARM	<u>120MG</u>	<u>N76151</u>	<u>001</u>	May 20, 2004
<u>AB3</u>		<u>180MG</u>	<u>N76151</u>	<u>002</u>	May 20, 2004
<u>AB3</u>		<u>240MG</u>	<u>N76151</u>	<u>003</u>	May 20, 2004
<u>AB3</u>		<u>300MG</u>	<u>N76151</u>	<u>004</u>	May 20, 2004

DILTIAZEM HYDROCHLORIDE

<u>AB2</u>	ANDRX	<u>120MG</u>	<u>N74852</u>	<u>001</u>	Oct 10, 1997
<u>AB2</u>		<u>180MG</u>	<u>N74852</u>	<u>002</u>	Oct 10, 1997
<u>AB2</u>		<u>240MG</u>	<u>N74852</u>	<u>003</u>	Oct 10, 1997
<u>AB3</u>	BIOVAIL	<u>120MG</u>	<u>N20939</u>	<u>001</u>	Jan 28, 2000
<u>AB3</u>		<u>120MG</u>	<u>N75116</u>	<u>001</u>	Dec 23, 1999
<u>AB3</u>		<u>180MG</u>	<u>N20939</u>	<u>002</u>	Jan 28, 2000
<u>AB3</u>		<u>180MG</u>	<u>N75116</u>	<u>002</u>	Dec 23, 1999
<u>AB3</u>		<u>240MG</u>	<u>N20939</u>	<u>003</u>	Jan 28, 2000
<u>AB3</u>		<u>240MG</u>	<u>N75116</u>	<u>003</u>	Dec 23, 1999
<u>AB3</u>		<u>300MG</u>	<u>N20939</u>	<u>004</u>	Jan 28, 2000
<u>AB3</u>		<u>300MG</u>	<u>N75116</u>	<u>004</u>	Dec 23, 1999
<u>AB2</u>	MYLAN	<u>120MG</u>	<u>N75124</u>	<u>002</u>	Mar 18, 1998
<u>AB2</u>		<u>180MG</u>	<u>N75124</u>	<u>003</u>	Mar 18, 1998
<u>AB2</u>		<u>240MG</u>	<u>N75124</u>	<u>001</u>	Mar 18, 1998
		60MG	N74910	001	May 02, 1997
		90MG	N74910	002	May 02, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 125 (of 360)

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

DILTIAZEM HYDROCHLORIDE

	+	MYLAN	120MG	N74910	003	May 02, 1997
<u>AB3</u>		PUREPAC PHARM	<u>120MG</u>	<u>N74984</u>	<u>001</u>	Dec 20, 1999
<u>AB3</u>			<u>180MG</u>	<u>N74984</u>	<u>002</u>	Dec 20, 1999
<u>AB3</u>			<u>240MG</u>	<u>N74984</u>	<u>003</u>	Dec 20, 1999
<u>AB3</u>			<u>300MG</u>	<u>N74984</u>	<u>004</u>	Dec 20, 1999
<u>AB2</u>		TORPHARM	<u>120MG</u>	<u>N74943</u>	<u>003</u>	Dec 19, 2000
<u>AB2</u>			<u>180MG</u>	<u>N74943</u>	<u>002</u>	Dec 19, 2000
<u>AB2</u>			<u>240MG</u>	<u>N74943</u>	<u>001</u>	Aug 06, 1998
		<u>TAZTIA XT</u>				
<u>AB4</u>		ANDRX PHARMS	<u>120MG</u>	<u>N75401</u>	<u>001</u>	Apr 10, 2003
<u>AB4</u>			<u>180MG</u>	<u>N75401</u>	<u>002</u>	Apr 10, 2003
<u>AB4</u>			<u>240MG</u>	<u>N75401</u>	<u>003</u>	Apr 10, 2003
<u>AB4</u>			<u>300MG</u>	<u>N75401</u>	<u>004</u>	Apr 10, 2003
<u>AB4</u>			<u>360MG</u>	<u>N75401</u>	<u>005</u>	Apr 10, 2003
		<u>TIAZAC</u>				
<u>AB4</u>		BIOVAIL	<u>120MG</u>	<u>N20401</u>	<u>001</u>	Sep 11, 1995
<u>AB4</u>			<u>180MG</u>	<u>N20401</u>	<u>002</u>	Sep 11, 1995
<u>AB4</u>			<u>240MG</u>	<u>N20401</u>	<u>003</u>	Sep 11, 1995
<u>AB4</u>			<u>300MG</u>	<u>N20401</u>	<u>004</u>	Sep 11, 1995
<u>AB4</u>			<u>360MG</u>	<u>N20401</u>	<u>005</u>	Sep 11, 1995
	+		420MG	N20401	006	Oct 16, 1998

INJECTABLE; INJECTION

CARDIZEM

<u>AP</u>	+	BIOVAIL	<u>100MG/VIAL</u>	<u>N20792</u>	<u>001</u>	Sep 05, 1997
<u>AP</u>	+	BIOVAIL LABS INTL	<u>5MG/ML</u>	<u>N20027</u>	<u>001</u>	Oct 24, 1991
	+		25MG/VIAL	N20027	003	Aug 18, 1995

DILTIAZEM HYDROCHLORIDE

<u>AP</u>		APOTEX	<u>5MG/ML</u>	<u>N75375</u>	<u>001</u>	Sep 30, 1999
<u>AP</u>	+	BEDFORD	<u>5MG/ML</u>	<u>N74617</u>	<u>001</u>	Feb 28, 1996
<u>AP</u>		HOSPIRA	<u>5MG/ML</u>	<u>N74941</u>	<u>001</u>	Apr 15, 1998
<u>AP</u>			<u>5MG/ML</u>	<u>N75004</u>	<u>001</u>	Feb 16, 2000
<u>AP</u>			<u>100MG/VIAL</u>	<u>N75853</u>	<u>001</u>	Dec 17, 2002
<u>AP</u>		INTL MEDICATION	<u>5MG/ML</u>	<u>N75749</u>	<u>001</u>	Nov 21, 2001
<u>AP</u>		MAYNE PHARMA USA	<u>5MG/ML</u>	<u>N75106</u>	<u>001</u>	Apr 29, 1999
<u>AP</u>		SICOR PHARMS	<u>5MG/ML</u>	<u>N74894</u>	<u>001</u>	Aug 26, 1997
	+		10MG/ML	N74894	002	Apr 19, 2002
<u>AP</u>		TAYLOR PHARMA	<u>5MG/ML</u>	<u>N75086</u>	<u>001</u>	Apr 09, 1998

TABLET; ORAL

CARDIZEM

<u>AB</u>		BIOVAIL LABS INTL	<u>30MG</u>	<u>N18602</u>	<u>001</u>	Nov 05, 1982
<u>AB</u>			<u>60MG</u>	<u>N18602</u>	<u>002</u>	Nov 05, 1982
<u>AB</u>			<u>90MG</u>	<u>N18602</u>	<u>003</u>	Dec 08, 1986
<u>AB</u>	+		<u>120MG</u>	<u>N18602</u>	<u>004</u>	Dec 08, 1986

DILTIAZEM HYDROCHLORIDE

<u>AB</u>		CLONMEL HLTHCARE	<u>30MG</u>	<u>N74093</u>	<u>001</u>	Nov 05, 1992
<u>AB</u>			<u>60MG</u>	<u>N74093</u>	<u>002</u>	Nov 05, 1992
<u>AB</u>			<u>90MG</u>	<u>N74093</u>	<u>003</u>	Nov 05, 1992
<u>AB</u>			<u>120MG</u>	<u>N74093</u>	<u>004</u>	Nov 05, 1992
<u>AB</u>		IVAX PHARMS	<u>30MG</u>	<u>N74168</u>	<u>001</u>	Mar 03, 1995
<u>AB</u>			<u>60MG</u>	<u>N74168</u>	<u>002</u>	Mar 03, 1995
<u>AB</u>			<u>90MG</u>	<u>N74168</u>	<u>003</u>	Mar 03, 1995
<u>AB</u>			<u>120MG</u>	<u>N74168</u>	<u>004</u>	Mar 03, 1995
<u>AB</u>		MYLAN	<u>30MG</u>	<u>N73185</u>	<u>001</u>	Nov 05, 1992
<u>AB</u>			<u>60MG</u>	<u>N73186</u>	<u>001</u>	Nov 05, 1992
<u>AB</u>			<u>90MG</u>	<u>N72837</u>	<u>001</u>	Nov 05, 1992
<u>AB</u>			<u>120MG</u>	<u>N72838</u>	<u>001</u>	Nov 05, 1992
<u>AB</u>		TEVA	<u>30MG</u>	<u>N74185</u>	<u>001</u>	May 31, 1995

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 126 (of 360)

DILTIAZEM HYDROCHLORIDE

TABLET; ORAL

DILTIAZEM HYDROCHLORIDE

<u>AB</u>	TEVA	<u>60MG</u>	<u>N74185</u>	<u>002</u>	May 31, 1995
<u>AB</u>		<u>90MG</u>	<u>N74185</u>	<u>003</u>	May 31, 1995
<u>AB</u>		<u>120MG</u>	<u>N74185</u>	<u>004</u>	May 31, 1995
TABLET, EXTENDED RELEASE; ORAL					
CARDIZEM LA					
	BIOVAIL LABS INTL	120MG	N21392	001	Feb 06, 2003
		180MG	N21392	002	Feb 06, 2003
		240MG	N21392	003	Feb 06, 2003
		300MG	N21392	004	Feb 06, 2003
		360MG	N21392	005	Feb 06, 2003
	+	420MG	N21392	006	Feb 06, 2003

DILTIAZEM MALATE; ENALAPRIL MALEATE

TABLET, EXTENDED RELEASE; ORAL

TECZEM

+	BIOVAIL	EQ 180MG HCL;5MG	N20507	001	Oct 04, 1996
---	---------	------------------	--------	-----	--------------

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

<u>AP</u>	AM PHARM	<u>50MG/ML</u>	<u>N40519</u>	<u>001</u>	Jun 23, 2004
<u>AP</u>	+	STERIS	<u>N80615</u>	<u>001</u>	

DIMERCAPROL

INJECTABLE; INJECTION

BAL

+	AKORN	10%	N05939	001	
---	-------	-----	--------	-----	--

DIMETHYL SULFOXIDE

SOLUTION; INTRAVESICAL

DIMETHYL SULFOXIDE

<u>AT</u>	BIONICHE (CANADA)	<u>50%</u>	<u>N76185</u>	<u>001</u>	Nov 29, 2002
<u>RIMSO-50</u>					
<u>AT</u>	+	EDWARDS LIFESCIENCES	<u>N17788</u>	<u>001</u>	

DINOPROSTONE

GEL; ENDOCERVICAL

PREPIDIL

+	PHARMACIA AND UPJOHN	0.5MG/3GM	N19617	001	Dec 09, 1992
---	----------------------	-----------	--------	-----	--------------

INSERT, EXTENDED RELEASE; VAGINAL

CERVIDIL

+	CONTROLLED THERAP	10MG	N20411	001	Mar 30, 1995
---	-------------------	------	--------	-----	--------------

SUPPOSITORY; VAGINAL

PROSTIN E2

+	PHARMACIA AND UPJOHN	20MG	N17810	001	
---	----------------------	------	--------	-----	--

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

<u>AA</u>	BARR	<u>50MG</u>	<u>N80738</u>	<u>001</u>	
<u>AA</u>	LNK	<u>25MG</u>	<u>N87977</u>	<u>001</u>	Jan 27, 1983
<u>AA</u>		<u>50MG</u>	<u>N87978</u>	<u>001</u>	Jan 27, 1983
<u>AA</u>	MUTUAL PHARM	<u>25MG</u>	<u>N84506</u>	<u>001</u>	
<u>AA</u>		<u>25MG</u>	<u>N89488</u>	<u>001</u>	Jan 02, 1987
<u>AA</u>		<u>50MG</u>	<u>N89489</u>	<u>001</u>	Jan 02, 1987
<u>AA</u>	+	SANDOZ	<u>N80832</u>	<u>001</u>	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 127 (of 360)

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

<u>AA</u>	+	SANDOZ	<u>50MG</u>	<u>N80832</u>	<u>002</u>	
<u>AA</u>		VALEANT PHARM INTL	<u>50MG</u>	<u>N80592</u>	<u>001</u>	
<u>AA</u>		WATSON LABS	<u>25MG</u>	<u>N80728</u>	<u>001</u>	
<u>AA</u>			<u>50MG</u>	<u>N80727</u>	<u>001</u>	

ELIXIR; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

<u>AA</u>	+	PHARM ASSOC	<u>12.5MG/5ML</u>	<u>N87513</u>	<u>001</u>	Feb 10, 1982
-----------	---	-------------	-------------------	---------------	------------	--------------

INJECTABLE; INJECTION

BENADRYL

<u>AP</u>	+	PARKE DAVIS	<u>50MG/ML</u>	<u>N06146</u>	<u>002</u>	
-----------	---	-------------	----------------	---------------	------------	--

BENADRYL PRESERVATIVE FREE

<u>AP</u>	+	PARKE DAVIS	<u>50MG/ML</u>	<u>N09486</u>	<u>001</u>	
-----------	---	-------------	----------------	---------------	------------	--

DIPHENHYDRAMINE HYDROCHLORIDE

<u>AP</u>		AM PHARM PARTNERS	<u>50MG/ML</u>	<u>N40466</u>	<u>001</u>	May 28, 2002
<u>AP</u>		BAXTER HLTHCARE	<u>50MG/ML</u>	<u>N80817</u>	<u>002</u>	
<u>AP</u>		BIONICHE PHARMA	<u>50MG/ML</u>	<u>N40498</u>	<u>001</u>	Jul 12, 2005
<u>AP</u>		HOSPIRA	<u>50MG/ML</u>	<u>N40140</u>	<u>001</u>	Nov 20, 1998
<u>AP</u>		STERIS	<u>50MG/ML</u>	<u>N80873</u>	<u>002</u>	
	+		10MG/ML	N80873	001	

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>		INTL MEDICATION	<u>50MG/ML</u>	<u>N84094</u>	<u>001</u>	
<u>AP</u>		STERIS	<u>50MG/ML</u>	<u>N80873</u>	<u>003</u>	

DIPIVEFRIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKPRO

<u>AT</u>		AKORN	<u>0.1%</u>	<u>N74382</u>	<u>001</u>	Sep 29, 1995
-----------	--	-------	-------------	---------------	------------	--------------

DIPIVEFRIN HYDROCHLORIDE

<u>AT</u>		BAUSCH AND LOMB	<u>0.1%</u>	<u>N74188</u>	<u>001</u>	May 19, 1995
<u>AT</u>		FALCON PHARMS	<u>0.1%</u>	<u>N73636</u>	<u>001</u>	Jun 30, 1994

PROPINE

<u>AT</u>	+	ALLERGAN	<u>0.1%</u>	<u>N18239</u>	<u>001</u>	
-----------	---	----------	-------------	---------------	------------	--

DIPYRIDAMOLE

INJECTABLE; INJECTION

DIPYRIDAMOLE

<u>AP</u>		AM PHARM PARTNERS	<u>5MG/ML</u>	<u>N74956</u>	<u>001</u>	Sep 30, 1998
<u>AP</u>		APOTEX	<u>5MG/ML</u>	<u>N75769</u>	<u>001</u>	Nov 27, 2002
<u>AP</u>		BAXTER HLTHCARE	<u>5MG/ML</u>	<u>N74521</u>	<u>001</u>	Oct 18, 1996
<u>AP</u>	+	BEDFORD	<u>5MG/ML</u>	<u>N74939</u>	<u>001</u>	Apr 13, 1998
<u>AP</u>		HOSPIRA	<u>5MG/ML</u>	<u>N74601</u>	<u>001</u>	Dec 19, 1997
<u>AP</u>		SICOR PHARMS	<u>5MG/ML</u>	<u>N74952</u>	<u>001</u>	Nov 26, 1997

TABLET; ORAL

DIPYRIDAMOLE

<u>AB</u>		BARR	<u>25MG</u>	<u>N87184</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>			<u>50MG</u>	<u>N87716</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>			<u>75MG</u>	<u>N87717</u>	<u>001</u>	Oct 03, 1990
<u>AB</u>		CLONMEL HLTHCARE	<u>50MG</u>	<u>N89000</u>	<u>001</u>	Feb 05, 1991
<u>AB</u>		PUREPAC PHARM	<u>25MG</u>	<u>N89425</u>	<u>001</u>	Jul 12, 1990
<u>AB</u>		SANDOZ	<u>25MG</u>	<u>N86944</u>	<u>002</u>	Apr 16, 1991
<u>AB</u>			<u>50MG</u>	<u>N87562</u>	<u>001</u>	Feb 25, 1992
<u>AB</u>			<u>75MG</u>	<u>N87561</u>	<u>001</u>	Feb 25, 1992
<u>AB</u>		WATSON LABS	<u>50MG</u>	<u>N87160</u>	<u>001</u>	Jun 07, 1996

PERSANTINE

<u>AB</u>		BOEHRINGER INGELHEIM	<u>25MG</u>	<u>N12836</u>	<u>003</u>	Dec 22, 1986
<u>AB</u>			<u>50MG</u>	<u>N12836</u>	<u>004</u>	Feb 06, 1987

PRESCRIPTION DRUG PRODUCT LIST

3 - 128 (of 360)

DIPYRIDAMOLE

TABLET; ORAL

PERSANTINE

<u>AB</u>	+ BOEHRINGER INGELHEIM	<u>75MG</u>	<u>N12836</u>	<u>005</u>	Feb 06, 1987
-----------	------------------------	-------------	---------------	------------	--------------

DISOPYRAMIDE PHOSPHATE

CAPSULE; ORAL

DISOPYRAMIDE PHOSPHATE

<u>AB</u>	IVAX PHARMS	<u>EQ 100MG BASE</u>	<u>N70186</u>	<u>001</u>	Nov 18, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N70187</u>	<u>001</u>	Nov 18, 1985
<u>AB</u>	SANDOZ	<u>EQ 100MG BASE</u>	<u>N70470</u>	<u>001</u>	Dec 10, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N70471</u>	<u>001</u>	Dec 10, 1985
<u>AB</u>	TEVA	<u>EQ 100MG BASE</u>	<u>N70101</u>	<u>001</u>	Feb 22, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N70102</u>	<u>001</u>	Feb 22, 1985
<u>AB</u>	WATSON LABS	<u>EQ 100MG BASE</u>	<u>N70173</u>	<u>001</u>	May 31, 1985
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N70174</u>	<u>001</u>	May 31, 1985

<u>NORPACE</u>					
<u>AB</u>	GD SEARLE LLC	<u>EQ 100MG BASE</u>	<u>N17447</u>	<u>001</u>	
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N17447</u>	<u>002</u>	

CAPSULE, EXTENDED RELEASE; ORAL

DISOPYRAMIDE PHOSPHATE

<u>AB</u>	KV PHARM	<u>EQ 150MG BASE</u>	<u>N71200</u>	<u>001</u>	Dec 15, 1987
-----------	----------	----------------------	---------------	------------	--------------

<u>NORPACE CR</u>					
<u>AB</u>	+	<u>EQ 150MG BASE</u>	<u>N18655</u>	<u>002</u>	Jul 20, 1982
		EQ 100MG BASE	N18655	001	Jul 20, 1982

DISULFIRAM

TABLET; ORAL

ANTABUSE

	+ ODYSSEY PHARMS	250MG	N88482	001	Dec 08, 1983
--	------------------	-------	--------	-----	--------------

DIVALPROEX SODIUM

CAPSULE, DELAYED REL PELLETS; ORAL

DEPAKOTE

	+ ABBOTT	EQ 125MG VALPROIC ACID	N19680	001	Sep 12, 1989
--	----------	------------------------	--------	-----	--------------

TABLET, DELAYED RELEASE; ORAL

DEPAKOTE

	ABBOTT	EQ 125MG VALPROIC ACID	N18723	003	Oct 26, 1984
		EQ 250MG VALPROIC ACID	N18723	001	Mar 10, 1983
+		EQ 500MG VALPROIC ACID	N18723	002	Mar 10, 1983

TABLET, EXTENDED RELEASE; ORAL

DEPAKOTE ER

	ABBOTT	EQ 250MG VALPROIC ACID	N21168	002	May 31, 2002
+		EQ 500MG VALPROIC ACID	N21168	001	Aug 04, 2000

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

<u>AP</u>	BEDFORD	<u>EQ 12.5MG BASE/ML</u>	<u>N74277</u>	<u>001</u>	Oct 31, 1994
<u>AP</u>	HOSPIRA	<u>EQ 12.5MG BASE/ML</u>	<u>N74086</u>	<u>001</u>	Nov 29, 1993
<u>AP</u>		<u>EQ 12.5MG BASE/ML</u>	<u>N74292</u>	<u>001</u>	Feb 16, 1995
<u>AP</u>		<u>EQ 1.25GM BASE/100ML</u>	<u>N74634</u>	<u>001</u>	Sep 27, 1996
<u>AP</u>	LUITPOLD	<u>EQ 12.5MG BASE/ML</u>	<u>N74545</u>	<u>001</u>	Jun 25, 1998
<u>AP</u>	MARSAM PHARMS LLC	<u>EQ 12.5MG BASE/ML</u>	<u>N74279</u>	<u>001</u>	Feb 18, 1998
<u>AP</u>		<u>EQ 12.5MG BASE/ML</u>	<u>N74995</u>	<u>001</u>	Mar 31, 1998
<u>AP</u>	SICOR PHARMS	<u>EQ 12.5MG BASE/ML</u>	<u>N74206</u>	<u>001</u>	Oct 19, 1993
<u>AP</u>	STERIS	<u>EQ 12.5MG BASE/ML</u>	<u>N74114</u>	<u>001</u>	Nov 30, 1993

<u>DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%</u>					
<u>AP</u>	+	<u>HOSPIRA</u>	<u>EQ 50MG BASE/100ML</u>	<u>N20269</u>	<u>001</u> Oct 19, 1993

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 129 (of 360)

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%

AP	+	HOSPIRA	<u>EQ 100MG BASE/100ML</u>	<u>N20269</u>	<u>002</u>	Oct 19, 1993
AP	+		<u>EQ 200MG BASE/100ML</u>	<u>N20269</u>	<u>003</u>	Oct 19, 1993

DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

AP	+	BAXTER HLHCARE	<u>EQ 50MG BASE/100ML</u>	<u>N20255</u>	<u>001</u>	Oct 19, 1993
AP	+		<u>EQ 100MG BASE/100ML</u>	<u>N20255</u>	<u>003</u>	Oct 19, 1993
AP	+		<u>EQ 200MG BASE/100ML</u>	<u>N20255</u>	<u>004</u>	Oct 19, 1993
AP	+		<u>EQ 400MG BASE/100ML</u>	<u>N20255</u>	<u>005</u>	Oct 19, 1993
AP	+	HOSPIRA	<u>EQ 50MG BASE/100ML</u>	<u>N20201</u>	<u>003</u>	Oct 19, 1993
AP	+		<u>EQ 100MG BASE/100ML</u>	<u>N20201</u>	<u>002</u>	Oct 19, 1993
AP	+		<u>EQ 200MG BASE/100ML</u>	<u>N20201</u>	<u>001</u>	Oct 19, 1993
AP	+		<u>EQ 400MG BASE/100ML</u>	<u>N20201</u>	<u>006</u>	Jul 07, 1994

DOCETAXEL

INJECTABLE; INJECTION

TAXOTERE

+	AVENTIS PHARMS	EQ 40MG BASE/ML	N20449	001	May 14, 1996
---	----------------	-----------------	--------	-----	--------------

DOFETILIDE

CAPSULE; ORAL

TIKOSYN

PFIZER

		0.125MG	N20931	001	Oct 01, 1999
		0.25MG	N20931	002	Oct 01, 1999
+		0.5MG	N20931	003	Oct 01, 1999

DOLASETRON MESYLATE MONOHYDRATE

INJECTABLE; INJECTION

ANZEMET

AVENTIS PHARMS

		EQ 12.5MG BASE/0.625ML	N20624	002	Sep 11, 1997
+		EQ 100MG BASE/5ML	N20624	001	Sep 11, 1997
+		EQ 500MG BASE/25ML	N20624	003	Dec 11, 2001

TABLET; ORAL

ANZEMET

AVENTIS PHARMS

		EQ 50MG BASE	N20623	001	Sep 11, 1997
+		EQ 100MG BASE	N20623	002	Sep 11, 1997

DONEPEZIL HYDROCHLORIDE

TABLET; ORAL

ARICEPT

EISAI MEDCL RES

		5MG	N20690	002	Nov 25, 1996
+		10MG	N20690	001	Nov 25, 1996

TABLET, ORALLY DISINTEGRATING; ORAL

ARICEPT ODT

EISAI MEDCL RES

		5MG	N21720	001	Oct 18, 2004
+		10MG	N21720	002	Oct 18, 2004

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

AP	+	HOSPIRA	<u>40MG/ML</u>	<u>N18132</u>	<u>001</u>	
AP	+		<u>40MG/ML</u>	<u>N74403</u>	<u>001</u>	May 23, 1996
AP	+		<u>80MG/ML</u>	<u>N18132</u>	<u>004</u>	Jul 09, 1982
AP	+		<u>80MG/100ML</u>	<u>N18132</u>	<u>002</u>	Feb 04, 1982
AP	+		<u>160MG/100ML</u>	<u>N18132</u>	<u>003</u>	Feb 04, 1982
AP	+	INTL MEDICATION	<u>40MG/ML</u>	<u>N18014</u>	<u>001</u>	
AP	+	LUITPOLD	<u>40MG/ML</u>	<u>N70799</u>	<u>001</u>	Feb 11, 1987
AP	+		<u>80MG/ML</u>	<u>N70820</u>	<u>001</u>	Feb 11, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 130 (of 360)

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOPAMINE HYDROCHLORIDE

<u>AP</u>	+	LUITPOLD	<u>160MG/ML</u>	<u>N70826</u>	<u>001</u>	Feb 11, 1987
<u>AP</u>	+	SICOR PHARMS	<u>40MG/ML</u>	<u>N72999</u>	<u>001</u>	Oct 23, 1991
<u>AP</u>	+		<u>80MG/ML</u>	<u>N73000</u>	<u>001</u>	Oct 23, 1991
<u>DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%</u>						
<u>AP</u>	+	B BRAUN	<u>80MG/100ML</u>	<u>N19099</u>	<u>002</u>	Oct 15, 1986
<u>AP</u>	+		<u>320MG/100ML</u>	<u>N19099</u>	<u>004</u>	Oct 15, 1986
<u>DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER</u>						
<u>AP</u>	+	B BRAUN	<u>160MG/100ML</u>	<u>N19099</u>	<u>003</u>	Oct 15, 1986
	+		40MG/100ML	N19099	001	Oct 15, 1986
<u>DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>						
<u>AP</u>	+	BAXTER HLTHCARE	<u>80MG/100ML</u>	<u>N19615</u>	<u>001</u>	Mar 27, 1987
<u>AP</u>	+		<u>160MG/100ML</u>	<u>N19615</u>	<u>002</u>	Mar 27, 1987
<u>AP</u>	+		<u>320MG/100ML</u>	<u>N19615</u>	<u>003</u>	Mar 27, 1987
	+		640MG/100ML	N19615	004	Mar 27, 1987
<u>AP</u>	+	HOSPIRA	<u>80MG/100ML</u>	<u>N18826</u>	<u>001</u>	Sep 30, 1983
<u>AP</u>	+		<u>160MG/100ML</u>	<u>N18826</u>	<u>002</u>	Sep 30, 1983
<u>AP</u>	+		<u>320MG/100ML</u>	<u>N18826</u>	<u>003</u>	Sep 30, 1983

DORZOLAMIDE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

TRUSOPT

+	MERCK	EQ 2% BASE	N20408	001	Dec 09, 1994
---	-------	------------	--------	-----	--------------

DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

COSOPT

+	MERCK	EQ 2% BASE;EQ 0.5% BASE	N20869	001	Apr 07, 1998
---	-------	-------------------------	--------	-----	--------------

DOXAPRAM HYDROCHLORIDE

INJECTABLE; INJECTION

DOPRAM

<u>AP</u>	+	BAXTER HLTHCARE CORP	<u>20MG/ML</u>	<u>N14879</u>	<u>001</u>	
<u>DOXAPRAM HYDROCHLORIDE</u>						
<u>AP</u>		BEDFORD	<u>20MG/ML</u>	<u>N76266</u>	<u>001</u>	Jan 10, 2003
<u>AP</u>		STERIS	<u>20MG/ML</u>	<u>N73529</u>	<u>001</u>	Jan 30, 1992

DOXAZOSIN MESYLATE

TABLET; ORAL

CARDURA

<u>AB</u>	+	PFIZER	<u>EQ 1MG BASE</u>	<u>N19668</u>	<u>001</u>	Nov 02, 1990
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>N19668</u>	<u>002</u>	Nov 02, 1990
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>N19668</u>	<u>003</u>	Nov 02, 1990
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>N19668</u>	<u>004</u>	Nov 02, 1990

DOXAZOSIN MESYLATE

<u>AB</u>		CLONMEL HLTHCARE	<u>EQ 1MG BASE</u>	<u>N76161</u>	<u>001</u>	Jun 10, 2004
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>N76161</u>	<u>002</u>	Jun 10, 2004
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>N76161</u>	<u>003</u>	Jun 10, 2004
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>N76161</u>	<u>004</u>	Jun 10, 2004
<u>AB</u>		GENPHARM	<u>EQ 1MG BASE</u>	<u>N75466</u>	<u>001</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>N75466</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>N75466</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>N75466</u>	<u>004</u>	Oct 18, 2000
<u>AB</u>		IVAX PHARMS	<u>EQ 1MG BASE</u>	<u>N75453</u>	<u>001</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 2MG BASE</u>	<u>N75453</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 4MG BASE</u>	<u>N75453</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>			<u>EQ 8MG BASE</u>	<u>N75453</u>	<u>004</u>	Oct 18, 2000
<u>AB</u>		KV PHARM	<u>EQ 1MG BASE</u>	<u>N75609</u>	<u>001</u>	Oct 18, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 131 (of 360)

DOXAZOSIN MESYLATE

TABLET; ORAL

DOXAZOSIN MESYLATE

<u>AB</u>	KV PHARM	<u>EQ 2MG BASE</u>	<u>N75609</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75609</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75609</u>	<u>004</u>	Oct 18, 2000
<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>N75509</u>	<u>001</u>	Oct 19, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75509</u>	<u>002</u>	Oct 19, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75509</u>	<u>003</u>	Oct 19, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75509</u>	<u>004</u>	Oct 19, 2000
<u>AB</u>	PLIVA	<u>EQ 1MG BASE</u>	<u>N75750</u>	<u>001</u>	Jun 08, 2001
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75750</u>	<u>002</u>	Jun 08, 2001
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75750</u>	<u>003</u>	Jun 08, 2001
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75750</u>	<u>004</u>	Jun 08, 2001
<u>AB</u>	PUREPAC PHARM	<u>EQ 1MG BASE</u>	<u>N75574</u>	<u>001</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75574</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75574</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75574</u>	<u>004</u>	Oct 18, 2000
<u>AB</u>	SANDOZ	<u>EQ 1MG BASE</u>	<u>N75432</u>	<u>001</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N75646</u>	<u>001</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75432</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75646</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75432</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75646</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75432</u>	<u>004</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75646</u>	<u>004</u>	Oct 18, 2000
<u>AB</u>	TEVA	<u>EQ 1MG BASE</u>	<u>N75353</u>	<u>001</u>	Jan 12, 2001
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N75536</u>	<u>001</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75353</u>	<u>002</u>	Jan 12, 2001
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75536</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75353</u>	<u>003</u>	Jan 12, 2001
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75536</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75353</u>	<u>004</u>	Jan 12, 2001
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75536</u>	<u>004</u>	Oct 18, 2000
<u>AB</u>	TORPHARM	<u>EQ 1MG BASE</u>	<u>N75580</u>	<u>001</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75580</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75580</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75580</u>	<u>004</u>	Oct 18, 2000
<u>AB</u>	WATSON LABS	<u>EQ 1MG BASE</u>	<u>N75426</u>	<u>001</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75426</u>	<u>002</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N75426</u>	<u>003</u>	Oct 18, 2000
<u>AB</u>		<u>EQ 8MG BASE</u>	<u>N75426</u>	<u>004</u>	Oct 18, 2000
TABLET, EXTENDED RELEASE; ORAL					
CARDURA XL					
	PFIZER	EQ 4MG BASE	N21269	001	Feb 22, 2005
	+	EQ 8MG BASE	N21269	002	Feb 22, 2005

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>N70789</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N70790</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N70791</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N70792</u>	<u>001</u>	May 13, 1986
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N70793</u>	<u>001</u>	May 13, 1986
<u>AB</u>	PAR PHARM	<u>EQ 10MG BASE</u>	<u>N71697</u>	<u>001</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N71437</u>	<u>001</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N71595</u>	<u>001</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N71608</u>	<u>001</u>	Nov 09, 1987
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N71422</u>	<u>001</u>	Nov 09, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 132 (of 360)

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

<u>AB</u>	PAR PHARM	<u>EQ 150MG BASE</u>	<u>N71669</u>	<u>001</u>	Nov 09, 1987
<u>AB</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>N71487</u>	<u>001</u>	Mar 02, 1987
<u>AB</u>	WATSON LABS	<u>EQ 10MG BASE</u>	<u>N71485</u>	<u>001</u>	Apr 30, 1987
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N72985</u>	<u>001</u>	Mar 29, 1991
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N71486</u>	<u>001</u>	Apr 30, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N72986</u>	<u>001</u>	Mar 29, 1991
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N71238</u>	<u>001</u>	Apr 30, 1987
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N72987</u>	<u>001</u>	Mar 29, 1991
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N71326</u>	<u>001</u>	Apr 30, 1987
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N71239</u>	<u>001</u>	Apr 30, 1987
<u>SINEQUAN</u>					
<u>AB</u>	PFIZER	<u>EQ 10MG BASE</u>	<u>N16798</u>	<u>003</u>	
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>N16798</u>	<u>001</u>	
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N16798</u>	<u>002</u>	
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N16798</u>	<u>006</u>	
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N16798</u>	<u>005</u>	
<u>AB</u>		<u>EQ 150MG BASE</u>	<u>N16798</u>	<u>007</u>	

CONCENTRATE; ORAL

DOXEPIN HYDROCHLORIDE

<u>AA</u>	MORTON GROVE	<u>EQ 10MG BASE/ML</u>	<u>N71918</u>	<u>001</u>	Jul 20, 1988
<u>AA</u>	PHARM ASSOC	<u>EQ 10MG BASE/ML</u>	<u>N75924</u>	<u>001</u>	Jan 15, 2004
<u>AA</u>	SILARX	<u>EQ 10MG BASE/ML</u>	<u>N74721</u>	<u>001</u>	Dec 29, 1998
<u>AA</u>	TEVA PHARMS	<u>EQ 10MG BASE/ML</u>	<u>N71609</u>	<u>001</u>	Nov 09, 1987
<u>SINEQUAN</u>					
<u>AA</u>	+	<u>EQ 10MG BASE/ML</u>	<u>N17516</u>	<u>001</u>	

CREAM; TOPICAL

ZONALON

+	BRADLEY PHARMS	5%	N20126	001	Apr 01, 1994
---	----------------	----	--------	-----	--------------

DOXERCALCIFEROL

CAPSULE; ORAL

HECTOROL

	GENZYME	0.5UGM	N20862	002	Apr 23, 2004
+		2.5UGM	N20862	001	Jun 09, 1999

INJECTABLE; INJECTION

HECTOROL

+	GENZYME	2UGM/ML	N21027	001	Apr 06, 2000
---	---------	---------	--------	-----	--------------

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN PFS

<u>AP</u>	+	PHARMACIA AND UPJOHN	<u>2MG/ML</u>	<u>N50629</u>	<u>001</u>	Dec 23, 1987
<u>AP</u>			<u>2MG/ML</u>	<u>N63165</u>	<u>001</u>	Jan 30, 1991
<u>AP</u>	+		<u>200MG/100ML</u>	<u>N50629</u>	<u>002</u>	May 03, 1988
<u>AP</u>			<u>200MG/100ML</u>	<u>N63165</u>	<u>002</u>	Jan 30, 1991

DOXORUBICIN HYDROCHLORIDE

<u>AP</u>	AM PHARM PARTNERS	<u>2MG/ML</u>	<u>N63277</u>	<u>001</u>	Oct 26, 1995
<u>AP</u>	BEDFORD	<u>2MG/ML</u>	<u>N62975</u>	<u>001</u>	Mar 17, 1989
<u>AP</u>		<u>10MG/VIAL</u>	<u>N62921</u>	<u>001</u>	Mar 17, 1989
<u>AP</u>		<u>20MG/VIAL</u>	<u>N62921</u>	<u>002</u>	Mar 17, 1989
<u>AP</u>		<u>50MG/VIAL</u>	<u>N62921</u>	<u>003</u>	Mar 17, 1989
<u>AP</u>		<u>200MG/100ML</u>	<u>N64097</u>	<u>001</u>	Sep 13, 1994
<u>AP</u>	PHARMACHEMIE	<u>2MG/ML</u>	<u>N63336</u>	<u>001</u>	Feb 28, 1995
<u>AP</u>		<u>10MG/VIAL</u>	<u>N63097</u>	<u>001</u>	May 21, 1990
<u>AP</u>		<u>20MG/VIAL</u>	<u>N63097</u>	<u>002</u>	May 21, 1990
<u>AP</u>		<u>50MG/VIAL</u>	<u>N63097</u>	<u>003</u>	May 21, 1990

PRESCRIPTION DRUG PRODUCT LIST

3 - 133 (of 360)

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

<u>AP</u>	PHARMACHEMIE	<u>200MG/100ML</u>	<u>N63336</u>	<u>004</u>	Feb 28, 1995
<u>AP</u>	SICOR PHARMS	<u>2MG/ML</u>	<u>N64140</u>	<u>001</u>	Jul 28, 1995
<u>AP</u>		<u>200MG/100ML</u>	<u>N64140</u>	<u>002</u>	Jul 28, 1995
<u>RUBEX</u>					
<u>AP</u>	BRISTOL MYERS SQUIBB	<u>10MG/VIAL</u>	<u>N62926</u>	<u>001</u>	Apr 13, 1989
<u>AP</u>		<u>50MG/VIAL</u>	<u>N62926</u>	<u>002</u>	Apr 13, 1989
		100MG/VIAL	N62926	003	Apr 13, 1989

INJECTABLE, LIPOSOMAL; INJECTION

DOXIL

+	ALZA	2MG/ML	N50718	001	Nov 17, 1995
---	------	--------	--------	-----	--------------

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>N65055</u>	<u>001</u>	Dec 01, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N65055</u>	<u>004</u>	Apr 18, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65055</u>	<u>002</u>	Dec 01, 2000
		EQ 150MG BASE	N65055	003	Jul 15, 2005
<u>AB</u>	RANBAXY	<u>EQ 50MG BASE</u>	<u>N65053</u>	<u>001</u>	Nov 22, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N65053</u>	<u>003</u>	Sep 10, 2003
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65053</u>	<u>002</u>	Nov 22, 2000
<u>AB</u>	SANDOZ	<u>EQ 50MG BASE</u>	<u>N65032</u>	<u>001</u>	Jun 30, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65032</u>	<u>002</u>	Jun 30, 2000
<u>AB</u>	WATSON LABS	<u>EQ 50MG BASE</u>	<u>N65041</u>	<u>001</u>	Apr 28, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65041</u>	<u>002</u>	Apr 28, 2000
<u>MONODOX</u>					
<u>AB</u>	OCLASSEN	<u>EQ 50MG BASE</u>	<u>N50641</u>	<u>002</u>	Feb 10, 1992
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N50641</u>	<u>001</u>	Dec 29, 1989

FOR SUSPENSION; ORAL

VIBRAMYCIN

+	PFIZER	EQ 25MG BASE/5ML	N50006	001	
---	--------	------------------	--------	-----	--

TABLET; ORAL

DOXYCYCLINE

<u>AB</u>	LANNETT	<u>EQ 50MG BASE</u>	<u>N65285</u>	<u>001</u>	Dec 08, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65285</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>	PAR PHARM	<u>EQ 50MG BASE</u>	<u>N65070</u>	<u>001</u>	Dec 15, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65070</u>	<u>002</u>	Dec 15, 2000
	+	EQ 150MG BASE	N65070	004	Jul 14, 2005

DOXYCYCLINE CALCIUM

SUSPENSION; ORAL

VIBRAMYCIN

+	PFIZER	EQ 50MG BASE/5ML	N50480	001	
---	--------	------------------	--------	-----	--

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	IVAX PHARMS	<u>EQ 50MG BASE</u>	<u>N62500</u>	<u>001</u>	Sep 11, 1984
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N62500</u>	<u>002</u>	Sep 11, 1984
<u>AB</u>	MUTUAL PHARM	<u>EQ 50MG BASE</u>	<u>N62675</u>	<u>001</u>	Jul 10, 1986
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N62676</u>	<u>001</u>	Jul 10, 1986
<u>AB</u>	MYLAN	<u>EQ 50MG BASE</u>	<u>N62337</u>	<u>001</u>	Mar 29, 1982
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N62337</u>	<u>002</u>	Mar 29, 1982
<u>AB</u>	WATSON LABS	<u>EQ 50MG BASE</u>	<u>N61717</u>	<u>001</u>	
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N62031</u>	<u>002</u>	Oct 13, 1982
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N61717</u>	<u>002</u>	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 134 (of 360)

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	WATSON LABS	<u>EQ 100MG BASE</u>	<u>N62031</u>	<u>001</u>	
<u>AB</u>	WEST WARD	<u>EQ 50MG BASE</u>	<u>N62396</u>	<u>002</u>	Nov 07, 1984
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N62396</u>	<u>001</u>	May 07, 1984
	+	EQ 20MG BASE	N65103	001	May 13, 2005
	<u>VIBRAMYCIN</u>				
<u>AB</u>	PFIZER	<u>EQ 50MG BASE</u>	<u>N50007</u>	<u>001</u>	
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N50007</u>	<u>002</u>	

CAPSULE, DELAYED RELEASE; ORAL

DORYX

<u>AB</u>	FH FAULding CO LTD	<u>EQ 75MG BASE</u>	<u>N50582</u>	<u>002</u>	Aug 13, 2001
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N50582</u>	<u>001</u>	Jul 22, 1985
<u>AB</u>	WARNER CHILCOTT	<u>EQ 100MG BASE</u>	<u>N62653</u>	<u>001</u>	Oct 30, 1985
	<u>DOXYCYCLINE HYCLATE</u>				
<u>AB</u>	SANDOZ	<u>EQ 75MG BASE</u>	<u>N65281</u>	<u>001</u>	Dec 21, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65281</u>	<u>002</u>	Dec 21, 2005

INJECTABLE; INJECTION

DOXY 100

	+	AM PHARM PARTNERS	EQ 100MG BASE/VIAL	N62475	001	Dec 09, 1983
		DOXY 200				
	+	AM PHARM PARTNERS	EQ 200MG BASE/VIAL	N62475	002	Dec 09, 1983

LIQUID, EXTENDED RELEASE; PERIODONTAL

ATRIDOX

	+	QLT USA	EQ 10% W/W	N50751	001	Sep 03, 1998
--	---	---------	------------	--------	-----	--------------

TABLET; ORAL

DOXYCYCLINE HYCLATE

<u>AB</u>	COREPHARMA	<u>EQ 20MG BASE</u>	<u>N65182</u>	<u>001</u>	May 13, 2005
<u>AB</u>	IVAX PHARMS	<u>EQ 20MG BASE</u>	<u>N65163</u>	<u>001</u>	May 13, 2005
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N62505</u>	<u>001</u>	Sep 11, 1984
<u>AB</u>	LANNETT	<u>EQ 20MG BASE</u>	<u>N65277</u>	<u>001</u>	Nov 10, 2005
<u>AB</u>	MUTUAL PHARM	<u>EQ 100MG BASE</u>	<u>N62677</u>	<u>001</u>	Jul 10, 1986
<u>AB</u>	MUTUAL PHARMA	<u>EQ 20MG BASE</u>	<u>N65134</u>	<u>001</u>	May 13, 2005
<u>AB</u>	MYLAN	<u>EQ 100MG BASE</u>	<u>N62432</u>	<u>001</u>	Feb 15, 1983
<u>AB</u>	TRUXTON INC	<u>EQ 100MG BASE</u>	<u>N62269</u>	<u>002</u>	Nov 08, 1982
<u>AB</u>	VINTAGE PHARMS	<u>EQ 100MG BASE</u>	<u>N62538</u>	<u>001</u>	Apr 07, 1986
<u>AB</u>	WATSON LABS	<u>EQ 100MG BASE</u>	<u>N62421</u>	<u>001</u>	Feb 02, 1983
<u>AB</u>	WEST WARD	<u>EQ 100MG BASE</u>	<u>N65095</u>	<u>001</u>	Jul 02, 2003

PERIOSTAT

<u>AB</u>	+	COLLAGENEX PHARMS	<u>EQ 20MG BASE</u>	<u>N50783</u>	<u>001</u>	Feb 02, 2001
-----------	---	-------------------	---------------------	---------------	------------	--------------

VIBRA-TABS

<u>AB</u>	+	PFIZER	<u>EQ 100MG BASE</u>	<u>N50533</u>	<u>001</u>	
-----------	---	--------	----------------------	---------------	------------	--

TABLET, DELAYED RELEASE; ORAL

DORYX

		FAULding	EQ 75MG BASE	N50795	001	May 06, 2005
	+		EQ 100MG BASE	N50795	002	May 06, 2005

DRONABINOL

CAPSULE; ORAL

MARINOL

		UNIMED	2.5MG	N18651	001	May 31, 1985
			5MG	N18651	002	May 31, 1985
	+		10MG	N18651	003	May 31, 1985

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

<u>AP</u>	HOSPIRA	<u>2.5MG/ML</u>	<u>N71981</u>	<u>001</u>	Feb 29, 1988
-----------	---------	-----------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 135 (of 360)

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

<u>AP</u>	HOSPIRA	<u>2.5MG/ML</u>	<u>N72272</u>	<u>001</u>	Aug 31, 1995
<u>AP</u>	LUITPOLD	<u>2.5MG/ML</u>	<u>N72123</u>	<u>001</u>	Oct 24, 1988
<u>AP</u>		<u>2.5MG/ML</u>	<u>N72335</u>	<u>001</u>	Oct 24, 1988

INAPSINE

<u>AP</u>	+ AKORN	<u>2.5MG/ML</u>	<u>N16796</u>	<u>001</u>	
-----------	---------	-----------------	---------------	------------	--

DROSPIRENONE; ESTRADIOL

TABLET; ORAL

ANGELIQ

+	BERLEX	0.5MG;1MG	N21355	002	Sep 28, 2005
---	--------	-----------	--------	-----	--------------

DROSPIRENONE; ETHINYL ESTRADIOL

TABLET; ORAL-28

YASMIN

+	BERLEX	3MG;0.03MG	N21098	001	May 11, 2001
---	--------	------------	--------	-----	--------------

DULOXETINE HYDROCHLORIDE

CAPSULE, DELAYED REL PELLETS; ORAL

CYMBALTA

LILLY

EQ 20MG BASE

N21427	001	Aug 03, 2004
--------	-----	--------------

EQ 30MG BASE

N21427	002	Aug 03, 2004
--------	-----	--------------

+

EQ 60MG BASE

N21427	004	Aug 03, 2004
--------	-----	--------------

DUTASTERIDE

CAPSULE; ORAL

AVODART

+	GLAXOSMITHKLINE	0.5MG	N21319	001	Nov 20, 2001
---	-----------------	-------	--------	-----	--------------

DYPHYLLINE

TABLET; ORAL

DILOR

BP	SAVAGE LABS	200MG	N84514	001	
----	-------------	-------	--------	-----	--

DILOR-400

BP	+ SAVAGE LABS	400MG	N84751	001	
----	---------------	-------	--------	-----	--

LUFYLLIN

BP	MEDPOINTE PHARM HLC	200MG	N84566	001	
----	---------------------	-------	--------	-----	--

BP		400MG	N84566	002	
----	--	-------	--------	-----	--

ECHOTHIOPHATE IODIDE

FOR SOLUTION; OPHTHALMIC

PHOSPHOLINE IODIDE

+	AYERST	0.125%	N11963	001	
---	--------	--------	--------	-----	--

ECONAZOLE NITRATE

CREAM; TOPICAL

ECONAZOLE NITRATE

<u>AB</u>	ALTANA	<u>1%</u>	<u>N76075</u>	<u>001</u>	Nov 26, 2002
-----------	--------	-----------	---------------	------------	--------------

<u>AB</u>	CLAY PARK	<u>1%</u>	<u>N76479</u>	<u>001</u>	Jun 23, 2004
-----------	-----------	-----------	---------------	------------	--------------

<u>AB</u>	HEALTHPOINT	<u>1%</u>	<u>N76574</u>	<u>001</u>	Dec 17, 2004
-----------	-------------	-----------	---------------	------------	--------------

<u>AB</u>	TARO	<u>1%</u>	<u>N76005</u>	<u>001</u>	Nov 26, 2002
-----------	------	-----------	---------------	------------	--------------

SPECTAZOLE

<u>AB</u>	+ JOHNSON AND JOHNSON	<u>1%</u>	<u>N18751</u>	<u>001</u>	Dec 23, 1982
-----------	-----------------------	-----------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 136 (of 360)

EDETATE CALCIUM DISODIUMINJECTABLE; INJECTION
CALCIUM DISODIUM VERSENATE

+ 3M 200MG/ML

N08922 001

EDETATE DISODIUM

INJECTABLE; INJECTION

EDETATE DISODIUM

AP APOTEX 150MG/ML

N40376 001

Nov 04, 2002

AP BIONICHE (CANADA) 150MG/ML

N40437 001

Jul 09, 2002

ENDRATE

AP + HOSPIRA 150MG/ML

N11355 001

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION

EDROPHONIUM CHLORIDE

AP HOSPIRA 10MG/ML

N40131 001

Feb 24, 1998

ENLON

AP BAXTER HLTHCARE CORP 10MG/ML

N88873 001

Aug 06, 1985

TENSILON

AP + VALEANT PHARM INTL 10MG/ML

N07959 001

TENSILON PRESERVATIVE FREE

AP + VALEANT PHARM INTL 10MG/ML

N07959 002

EFAVIRENZ

CAPSULE; ORAL

SUSTIVA

BRISTOL MYERS SQUIBB 50MG

N20972 001

Sep 17, 1998

100MG

N20972 002

Sep 17, 1998

+

200MG

N20972 003

Sep 17, 1998

TABLET; ORAL

SUSTIVA

+ BRISTOL MYERS SQUIBB 600MG

N21360 002

Feb 01, 2002

EFLORNITHINE HYDROCHLORIDE

CREAM; TOPICAL

VANIQA

+ SKINMEDICA 13.9%

N21145 001

Jul 27, 2000

ELETRIPTAN HYDROBROMIDE

TABLET; ORAL

RELPAK

PFIZER IRELAND EQ 20MG BASE

N21016 001

Dec 26, 2002

+

EQ 40MG BASE

N21016 002

Dec 26, 2002

EMEDASTINE DIFUMARATE

SOLUTION/DROPS; OPHTHALMIC

EMADINE

+ ALCON 0.05%

N20706 001

Dec 29, 1997

EMTRICITABINE

CAPSULE; ORAL

EMTRIVA

+ GILEAD 200MG

N21500 001

Jul 02, 2003

SOLUTION; ORAL

EMTRIVA

+ GILEAD 10MG/ML

N21896 001

Sep 28, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 137 (of 360)

EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

TRUVADA

+ GILEAD

200MG;300MG

N21752 001

Aug 02, 2004

ENALAPRIL MALEATE

TABLET; ORAL

ENALAPRIL MALEATE

<u>AB</u>	GENPHARM	<u>2.5MG</u>	<u>N75472</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75472</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75472</u>	<u>003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75472</u>	<u>004</u>	Aug 22, 2000
<u>AB</u>	KRKA DD NOVO MESTO	<u>2.5MG</u>	<u>N75370</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75370</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75369</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75369</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>	LEK PHARMS	<u>2.5MG</u>	<u>N75496</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75496</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75459</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75459</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>N75480</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75480</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75480</u>	<u>003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75480</u>	<u>004</u>	Aug 22, 2000
<u>AB</u>	RANBAXY	<u>2.5MG</u>	<u>N75556</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75556</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75556</u>	<u>003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75556</u>	<u>004</u>	Aug 22, 2000
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>N75048</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>2.5MG</u>	<u>N75621</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75048</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75621</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75048</u>	<u>003</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75621</u>	<u>003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75048</u>	<u>004</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75621</u>	<u>004</u>	Aug 22, 2000
<u>AB</u>	TARO	<u>2.5MG</u>	<u>N75657</u>	<u>001</u>	Jan 23, 2001
<u>AB</u>		<u>5MG</u>	<u>N75657</u>	<u>002</u>	Jan 23, 2001
<u>AB</u>		<u>10MG</u>	<u>N75657</u>	<u>003</u>	Jan 23, 2001
<u>AB</u>		<u>20MG</u>	<u>N75657</u>	<u>004</u>	Jan 23, 2001
<u>AB</u>	TEVA	<u>2.5MG</u>	<u>N75479</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75479</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75479</u>	<u>003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75479</u>	<u>004</u>	Aug 22, 2000
<u>AB</u>	TORPHARM	<u>2.5MG</u>	<u>N75178</u>	<u>002</u>	Mar 23, 2001
<u>AB</u>		<u>5MG</u>	<u>N75178</u>	<u>001</u>	Mar 23, 2001
<u>AB</u>		<u>10MG</u>	<u>N75178</u>	<u>003</u>	Mar 23, 2001
<u>AB</u>		<u>20MG</u>	<u>N75178</u>	<u>004</u>	Mar 23, 2001
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>N75501</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75501</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75501</u>	<u>003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75501</u>	<u>004</u>	Aug 22, 2000
<u>AB</u>	WOCKHARDT	<u>2.5MG</u>	<u>N75483</u>	<u>001</u>	Aug 22, 2000
<u>AB</u>		<u>5MG</u>	<u>N75483</u>	<u>002</u>	Aug 22, 2000
<u>AB</u>		<u>10MG</u>	<u>N75483</u>	<u>003</u>	Aug 22, 2000
<u>AB</u>		<u>20MG</u>	<u>N75483</u>	<u>004</u>	Aug 22, 2000
<u>VASOTEC</u>					
<u>AB</u>	BIOVAIL LABS INTL	<u>2.5MG</u>	<u>N18998</u>	<u>005</u>	Jul 26, 1988
<u>AB</u>		<u>5MG</u>	<u>N18998</u>	<u>001</u>	Dec 24, 1985
<u>AB</u>		<u>10MG</u>	<u>N18998</u>	<u>002</u>	Dec 24, 1985

PRESCRIPTION DRUG PRODUCT LIST

3 - 138 (of 360)

ENALAPRIL MALEATE

TABLET; ORAL

VASOTEC

<u>AB</u>	+ BIOVAIL LABS INTL	<u>20MG</u>	<u>N18998</u>	<u>003</u>	Dec 24, 1985
-----------	---------------------	-------------	---------------	------------	--------------

ENALAPRIL MALEATE; FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

LEXXEL

	+ ASTRAZENECA	5MG; 2.5MG	N20668	002	Oct 28, 1998
	+	5MG; 5MG	N20668	001	Dec 27, 1996

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	APOTEX	<u>5MG; 12.5MG</u>	<u>N76486</u>	<u>001</u>	Oct 27, 2004
<u>AB</u>		<u>10MG; 25MG</u>	<u>N76486</u>	<u>002</u>	Oct 27, 2004
<u>AB</u>	DR REDDYS LABS LTD	<u>5MG; 12.5MG</u>	<u>N75909</u>	<u>001</u>	Oct 15, 2001
<u>AB</u>		<u>10MG; 25MG</u>	<u>N75909</u>	<u>002</u>	Oct 15, 2001
<u>AB</u>	MYLAN	<u>5MG; 12.5MG</u>	<u>N75624</u>	<u>001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG; 25MG</u>	<u>N75624</u>	<u>002</u>	Sep 18, 2001
<u>AB</u>	SANDOZ	<u>5MG; 12.5MG</u>	<u>N76116</u>	<u>001</u>	Sep 19, 2001
<u>AB</u>		<u>10MG; 25MG</u>	<u>N76116</u>	<u>002</u>	Sep 19, 2001
<u>AB</u>	TARO PHARM INDS	<u>5MG; 12.5MG</u>	<u>N75788</u>	<u>001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG; 25MG</u>	<u>N75788</u>	<u>002</u>	Sep 18, 2001
<u>AB</u>	TEVA	<u>5MG; 12.5MG</u>	<u>N75727</u>	<u>001</u>	Sep 18, 2001
<u>AB</u>		<u>10MG; 25MG</u>	<u>N75727</u>	<u>002</u>	Sep 18, 2001
	<u>VASERETIC</u>				
<u>AB</u>	BIOVAIL LABS INTL	<u>5MG; 12.5MG</u>	<u>N19221</u>	<u>003</u>	Jul 12, 1995
<u>AB</u>	+	<u>10MG; 25MG</u>	<u>N19221</u>	<u>001</u>	Oct 31, 1986

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

<u>AP</u>	BEDFORD	<u>1.25MG/ML</u>	<u>N75634</u>	<u>001</u>	Aug 22, 2000
<u>AP</u>	HOSPIRA	<u>1.25MG/ML</u>	<u>N75456</u>	<u>001</u>	Aug 22, 2000
<u>AP</u>		<u>1.25MG/ML</u>	<u>N75458</u>	<u>001</u>	Aug 22, 2000
<u>AP</u>	MAYNE PHARMA USA	<u>1.25MG/ML</u>	<u>N75571</u>	<u>001</u>	Aug 22, 2000
<u>AP</u>	SICOR PHARMS	<u>1.25MG/ML</u>	<u>N75578</u>	<u>001</u>	Aug 22, 2000
	<u>VASOTEC</u>				
<u>AP</u>	+ BIOVAIL LABS INTL	<u>1.25MG/ML</u>	<u>N19309</u>	<u>001</u>	Feb 09, 1988

ENFLURANE

LIQUID; INHALATION

ENFLURANE

<u>AN</u>	MINRAD	<u>99.9%</u>	<u>N74396</u>	<u>001</u>	Jul 29, 1994
	<u>ETHRANE</u>				
<u>AN</u>	+ BAXTER HLTHCARE CORP	<u>99.9%</u>	<u>N17087</u>	<u>001</u>	

ENFUVRTIDE

INJECTABLE; SUBCUTANEOUS

FUZEON

	+ ROCHE	90MG/VIAL	N21481	001	Mar 13, 2003
--	---------	-----------	--------	-----	--------------

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

LOVENOX

	AVENTIS	300MG/3ML (100MG/ML)	N20164	009	Jan 23, 2003
--	---------	----------------------	--------	-----	--------------

LOVENOX (PRESERVATIVE FREE)

	AVENTIS	30MG/0.3ML (100MG/ML)	N20164	001	Mar 29, 1993
--	---------	-----------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 139 (of 360)

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

LOVENOX (PRESERVATIVE FREE)

AVENTIS	40MG/0.4ML (100MG/ML)	N20164	002	Jan 30, 1998
	60MG/0.6ML (100MG/ML)	N20164	003	Mar 27, 1998
	80MG/0.8ML (100MG/ML)	N20164	004	Mar 27, 1998
+	100MG/ML (100MG/ML)	N20164	005	Mar 27, 1998
	120MG/0.8ML (150MG/ML)	N20164	007	Jun 02, 2000
	150MG/ML (150MG/ML)	N20164	008	Jun 02, 2000

ENTACAPONE

TABLET; ORAL

COMTAN

+ ORION	200MG	N20796	001	Oct 19, 1999
---------	-------	--------	-----	--------------

ENTECAVIR

SOLUTION; ORAL

BARACLUDE

+ BRISTOL MYERS SQUIBB	0.05MG/ML	N21798	001	Mar 29, 2005
------------------------	-----------	--------	-----	--------------

TABLET; ORAL

BARACLUDE

BRISTOL MYERS SQUIBB	0.5MG	N21797	001	Mar 29, 2005
+	1MG	N21797	002	Mar 29, 2005

EPINASTINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ELESTAT

+ ALLERGAN	0.05%	N21565	001	Oct 16, 2003
------------	-------	--------	-----	--------------

EPINEPHRINE

INJECTABLE; IM-SC

TWINJECT

+ VERUS PHARMS	EQ 0.15MG /DELIVERY	N20800	002	May 28, 2004
----------------	---------------------	--------	-----	--------------

TWINJECT 0.3

+ VERUS PHARMS	EQ 0.3MG /DELIVERY	N20800	001	May 30, 2003
----------------	--------------------	--------	-----	--------------

INJECTABLE; INTRAMUSCULAR

EPIPEN

+ MERIDIAN MEDCL TECHN	0.3MG/DELIVERY	N19430	001	Dec 22, 1987
------------------------	----------------	--------	-----	--------------

EPIPEN JR.

+ MERIDIAN MEDCL TECHN	0.15MG/DELIVERY	N19430	002	Dec 22, 1987
------------------------	-----------------	--------	-----	--------------

EPINEPHRINE BITARTRATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIGNOSPAN FORTE

+ DEPROCO	EQ 0.02MG BASE/ML;2%	N88389	001	Jan 22, 1985
-----------	----------------------	--------	-----	--------------

LIGNOSPAN STANDARD

+ DEPROCO	EQ 0.01MG BASE/ML;2%	N88390	001	Jan 22, 1985
-----------	----------------------	--------	-----	--------------

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CITANEST FORTE

+ DENTSPLY PHARM	0.005MG/ML;4%	N21383	001	
------------------	---------------	--------	-----	--

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

AP	EASTMAN KODAK	0.01MG/ML;2%	N40057	002	Feb 26, 1993
AP		0.02MG/ML;2%	N40057	001	Feb 26, 1993
AP	GRAHAM CHEM	0.01MG/ML;2%	N80504	004	Oct 19, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 140 (of 360)

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

AP	GRAHAM CHEM	0.02MG/ML;2%	N80504	005	Oct 19, 1983
AP	HOSPIRA	0.005MG/ML;0.5%	N89635	001	Jun 21, 1988
AP		0.005MG/ML;1%	N89649	001	Jun 21, 1988
AP		0.005MG/ML;1.5%	N88571	001	Sep 13, 1985
AP		0.005MG/ML;1.5%	N89645	001	Jun 21, 1988
AP		0.005MG/ML;1.5%	N89650	001	Jun 21, 1988
AP		0.005MG/ML;2%	N89651	001	Jun 21, 1988
AP		0.01MG/ML;1%	N89644	001	Jun 21, 1988
AP		0.01MG/ML;2%	N89646	001	Jun 21, 1988

LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE

AP	DELL LABS	0.01MG/ML;1%	N83389	001	
AP		0.01MG/ML;2%	N83390	001	

LIDOCATON

AP	PHARMATON	0.01MG/ML;2%	N84729	001	Aug 17, 1983
----	-----------	--------------	--------	-----	--------------

OCTOCAINE

AP	SEPTODONT	0.01MG/ML;2%	N84048	001	
AP	+	0.02MG/ML;2%	N84048	002	

XYLOCAINE W/ EPINEPHRINE

AP	+	ASTRAZENECA	0.005MG/ML;0.5%	N06488	012	
AP	+		0.005MG/ML;1%	N06488	018	Nov 13, 1986
AP	+		0.005MG/ML;1.5%	N06488	017	
AP	+		0.005MG/ML;2%	N06488	019	Nov 13, 1986
AP	+		0.01MG/ML;1%	N06488	004	
	+	DENTSPLY PHARM	0.01MG/ML;2%	N21381	001	
		0.02MG/ML;2%	N21381	002		

PATCH; IONTOPHORESIS

LIDOSITE TOPICAL SYSTEM KIT

+	VYTERIS	1.05MG/PATCH;100MG/PATCH	N21504	001	May 06, 2004
---	---------	--------------------------	--------	-----	--------------

SOLUTION; IONTOPHORESIS

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

+	EMPI	0.01MG/ML;2%	N21486	001	Oct 26, 2004
---	------	--------------	--------	-----	--------------

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ELLECE

+	PFIZER INC	2MG/ML	N50778	001	Sep 15, 1999
---	------------	--------	--------	-----	--------------

EPLERENONE

TABLET; ORAL

INSPRA

	GD SEARLE LLC	25MG	N21437	001	Sep 27, 2002
+		50MG	N21437	002	Sep 27, 2002

EPOPROSTENOL SODIUM

INJECTABLE; INJECTION

FLOLAN

+	GLAXOSMITHKLINE	EQ 0.5MG BASE/VIAL	N20444	001	Sep 20, 1995
+		EQ 1.5MG BASE/VIAL	N20444	002	Sep 20, 1995

EPROSARTAN MESYLATE

TABLET; ORAL

TEVETEN

	KOS LIFE	EQ 400MG BASE	N20738	005	Dec 22, 1997
+		EQ 600MG BASE	N20738	006	May 27, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 141 (of 360)

EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

TEVETEN HCT

KOS LIFE

600MG;12.5MG

N21268 001

Nov 01, 2001

+

600MG;25MG

N21268 002

Nov 01, 2001

EPTIFIBATIDE

INJECTABLE; INJECTION

INTEGRILIN

+ SCHERING

2MG/ML

N20718 001

May 18, 1998

+

75MG/100ML

N20718 002

May 18, 1998

ERGOCALCIFEROL

CAPSULE; ORAL

DRISDOLAA + SANOFI SYNTHELABO50,000 IUN03444 001VITAMIN DAA BANNER PHARMACAPS50,000 IUN80704 001ERGOLOID MESYLATES

TABLET; ORAL

ERGOLOID MESYLATESAB MUTUAL PHARM1MGN81113 001

Oct 31, 1991

HYDERGINEAB + NOVARTIS1MGN17993 001

TABLET; SUBLINGUAL

ERGOLOID MESYLATESAA WATSON LABS0.5MGN87233 001GERIMALAA WATSON LABS0.5MGN86189 001AA1MGN86188 001HYDERGINEAA + NOVARTIS0.5MGN09087 002AA

+

1MGN09087 001HYDROGENATED ERGOT ALKALOIDSAA IVAX PHARMS1MGN87185 001ERGOTAMINE TARTRATE

TABLET; SUBLINGUAL

ERGOMAR

+ HARVEST PHARMS

2MG

N87693 001

Feb 24, 1983

ERLOTINIB HYDROCHLORIDE

TABLET; ORAL

TARCEVA

OSI PHARMS

EQ 25MG BASE

N21743 001

Nov 18, 2004

EQ 100MG BASE

N21743 002

Nov 18, 2004

+

EQ 150MG BASE

N21743 003

Nov 18, 2004

ERTAPENEM SODIUM

INJECTABLE; INTRAMUSCULAR, IV (INFUSION)

INVANZ

+ MERCK

EQ 1GM BASE/VIAL

N21337 001

Nov 21, 2001

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL

ERYCAB + MAYNE PHARMA USA250MGN50536 001AB WARNER CHILCOTT250MGN62338 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 142 (of 360)

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL

ERYTHROMYCIN

<u>AB</u>	ABBOTT	<u>250MG</u>	<u>N62746</u>	<u>001</u>	Dec 22, 1986
<u>AB</u>	BARR	<u>250MG</u>	<u>N63098</u>	<u>001</u>	May 04, 1989

GEL; TOPICAL

E-GLADES

<u>AT</u>	GLADES PHARMS	<u>2%</u>	<u>N65009</u>	<u>001</u>	Mar 18, 2002
-----------	---------------	-----------	---------------	------------	--------------

ERYGEL

<u>AT</u>	+ MERZ PHARMS	<u>2%</u>	<u>N50617</u>	<u>001</u>	Oct 21, 1987
-----------	---------------	-----------	---------------	------------	--------------

ERYTHROMYCIN

<u>AT</u>	ALTANA	<u>2%</u>	<u>N64184</u>	<u>001</u>	Sep 30, 1997
-----------	--------	-----------	---------------	------------	--------------

<u>AT</u>	STIEFEL	<u>2%</u>	<u>N63211</u>	<u>001</u>	Jan 29, 1993
-----------	---------	-----------	---------------	------------	--------------

LOTION; TOPICAL

E-SOLVE 2

	+ SYOSSET	2%	N62467	001	Jul 03, 1985
--	-----------	----	--------	-----	--------------

OINTMENT; OPHTHALMIC

ERYTHROMYCIN

<u>AT</u>	AKORN	<u>0.5%</u>	<u>N64030</u>	<u>001</u>	Jul 18, 1996
-----------	-------	-------------	---------------	------------	--------------

<u>AT</u>	BAUSCH AND LOMB	<u>0.5%</u>	<u>N64067</u>	<u>001</u>	Jul 29, 1994
-----------	-----------------	-------------	---------------	------------	--------------

<u>AT</u>	+ FOUGERA	<u>0.5%</u>	<u>N62447</u>	<u>001</u>	Sep 26, 1983
-----------	-----------	-------------	---------------	------------	--------------

OINTMENT; TOPICAL

AKNE-MYCIN

	+ HEALTHPOINT	2%	N50584	001	Jan 10, 1985
--	---------------	----	--------	-----	--------------

SOLUTION; TOPICAL

A/T/S

<u>AT</u>	AVENTIS PHARMS	<u>2%</u>	<u>N62405</u>	<u>001</u>	Nov 18, 1982
-----------	----------------	-----------	---------------	------------	--------------

ERYDERM

<u>AT</u>	ABBOTT	<u>2%</u>	<u>N62290</u>	<u>001</u>	
-----------	--------	-----------	---------------	------------	--

ERYTHRA-DERM

<u>AT</u>	PADDOCK	<u>2%</u>	<u>N62687</u>	<u>001</u>	Feb 05, 1988
-----------	---------	-----------	---------------	------------	--------------

ERYTHROMYCIN

<u>AT</u>	+ ALTANA	<u>2%</u>	<u>N64187</u>	<u>001</u>	Sep 30, 1997
-----------	----------	-----------	---------------	------------	--------------

<u>AT</u>	CLAY PARK	<u>2%</u>	<u>N63038</u>	<u>001</u>	Jan 11, 1991
-----------	-----------	-----------	---------------	------------	--------------

<u>AT</u>	MORTON GROVE	<u>2%</u>	<u>N62825</u>	<u>001</u>	Oct 23, 1987
-----------	--------------	-----------	---------------	------------	--------------

SWAB; TOPICAL

C-SOLVE-2

<u>AT</u>	IVAX PHARMS	<u>2%</u>	<u>N62751</u>	<u>001</u>	Jul 30, 1993
-----------	-------------	-----------	---------------	------------	--------------

ERYCETTE

<u>AT</u>	+ J AND J	<u>2%</u>	<u>N50594</u>	<u>001</u>	Feb 15, 1985
-----------	-----------	-----------	---------------	------------	--------------

ERYTHROMYCIN

<u>AT</u>	STIEFEL	<u>2%</u>	<u>N64126</u>	<u>001</u>	Jul 03, 1996
-----------	---------	-----------	---------------	------------	--------------

<u>AT</u>		<u>2%</u>	<u>N64128</u>	<u>001</u>	Jul 03, 1996
-----------	--	-----------	---------------	------------	--------------

TABLET; ORAL

ERYTHROMYCIN

	ABBOTT	250MG	N61621	001	
	+	500MG	N61621	002	

TABLET, COATED PARTICLES; ORAL

PCE

	ABBOTT	333MG	N50611	001	Sep 09, 1986
	+	500MG	N50611	002	Aug 22, 1990

TABLET, DELAYED RELEASE; ORAL

E-BASE

<u>AB</u>	BARR	<u>333MG</u>	<u>N63086</u>	<u>001</u>	May 15, 1990
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N62999</u>	<u>001</u>	Nov 25, 1988
-----------	--	--------------	---------------	------------	--------------

E-MYCIN

<u>AB</u>	ABBOTT	<u>250MG</u>	<u>N60272</u>	<u>001</u>	
-----------	--------	--------------	---------------	------------	--

<u>AB</u>		<u>333MG</u>	<u>N60272</u>	<u>002</u>	
-----------	--	--------------	---------------	------------	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 143 (of 360)

ERYTHROMYCIN

TABLET, DELAYED RELEASE; ORAL

ERY-TAB

<u>AB</u>	+	ABBOTT	<u>250MG</u>	<u>N62298</u>	<u>001</u>	
<u>AB</u>	+		<u>333MG</u>	<u>N62298</u>	<u>003</u>	Mar 29, 1982
<u>AB</u>	+		<u>500MG</u>	<u>N62298</u>	<u>002</u>	

ERYTHROMYCIN ESTOLATE

SUSPENSION; ORAL

ERYTHROMYCIN ESTOLATE

	ALPHARMA US PHARMS	EQ 125MG BASE/5ML	N62353	001	Nov 18, 1982
+		EQ 250MG BASE/5ML	N62409	001	Dec 16, 1982

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL

E.E.S.

<u>AB</u>		ABBOTT	<u>EQ 200MG BASE/5ML</u>	<u>N50207</u>	<u>001</u>	
<u>AB</u>		ABBOTT	<u>EQ 200MG BASE/5ML</u>	<u>N50207</u>	<u>003</u>	Mar 30, 1987
	+		EQ 400MG BASE/5ML	N50207	002	

ERYTHROMYCIN ETHYLSUCCINATE

<u>AB</u>		BARR	<u>EQ 200MG BASE/5ML</u>	<u>N62055</u>	<u>001</u>	
SUSPENSION; ORAL						
		<u>E.E.S. 200</u>				
<u>AB</u>		ABBOTT	<u>EQ 200MG BASE/5ML</u>	<u>N61639</u>	<u>001</u>	
		<u>E.E.S. 400</u>				
<u>AB</u>	+	ABBOTT	<u>EQ 400MG BASE/5ML</u>	<u>N61639</u>	<u>002</u>	
<u>ERYTHROMYCIN ETHYLSUCCINATE</u>						
<u>AB</u>		ALPHARMA US PHARMS	<u>EQ 200MG BASE/5ML</u>	<u>N62200</u>	<u>001</u>	
<u>AB</u>			<u>EQ 400MG BASE/5ML</u>	<u>N62200</u>	<u>002</u>	
<u>PEDIAMYCIN</u>						
<u>AB</u>		ROSS LABS	<u>EQ 200MG BASE/5ML</u>	<u>N62304</u>	<u>001</u>	
		<u>PEDIAMYCIN 400</u>				
<u>AB</u>		ROSS LABS	<u>EQ 400MG BASE/5ML</u>	<u>N62304</u>	<u>002</u>	

TABLET; ORAL

E.E.S. 400

<u>AB</u>	+	ABBOTT	<u>EQ 400MG BASE</u>	<u>N61905</u>	<u>002</u>	Aug 12, 1982
<u>ERYTHROMYCIN ETHYLSUCCINATE</u>						
<u>AB</u>		MYLAN	<u>EQ 400MG BASE</u>	<u>N62847</u>	<u>001</u>	Sep 14, 1988

TABLET, CHEWABLE; ORAL

E.E.S.

<u>AB</u>	+	ABBOTT	<u>EQ 200MG BASE</u>	<u>N50297</u>	<u>002</u>	
<u>ERYPED</u>						
<u>AB</u>		ABBOTT	<u>EQ 200MG BASE</u>	<u>N50297</u>	<u>003</u>	Jul 05, 1988

ERYTHROMYCIN ETHYLSUCCINATE; SULFISOXAZOLE ACETYL

GRANULE; ORAL

ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL

<u>AB</u>		BARR	<u>EQ 200MG BASE/5ML;EQ 600MG BASE/5ML</u>	<u>N62759</u>	<u>001</u>	May 20, 1988
<u>ERYZOLE</u>						
<u>AB</u>		ALRA	<u>EQ 200MG BASE/5ML;EQ 600MG BASE/5ML</u>	<u>N62758</u>	<u>001</u>	Jun 15, 1988
<u>PEDIAZOLE</u>						
<u>AB</u>	+	ROSS LABS	<u>EQ 200MG BASE/5ML;EQ 600MG BASE/5ML</u>	<u>N50529</u>	<u>001</u>	

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

<u>AP</u>	+	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>N50182</u>	<u>002</u>	
<u>AP</u>			<u>EQ 500MG BASE/VIAL</u>	<u>N50609</u>	<u>001</u>	Sep 24, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 144 (of 360)

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

<u>AP</u>	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>N62638</u>	<u>001</u>	Oct 31, 1986
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N50182</u>	<u>003</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N50609</u>	<u>002</u>	Sep 24, 1986
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N62638</u>	<u>002</u>	Oct 31, 1986

ERYTHROMYCIN LACTOBIONATE

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 500MG BASE/VIAL</u>	<u>N62993</u>	<u>001</u>	May 09, 1989
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N62993</u>	<u>002</u>	May 09, 1989
<u>AP</u>	SICOR PHARMS	<u>EQ 500MG BASE/VIAL</u>	<u>N63253</u>	<u>001</u>	Jul 30, 1993
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N63253</u>	<u>002</u>	Jul 30, 1993

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROCIN STEARATE

<u>AB</u>	ABBOTT	<u>EQ 250MG BASE</u>	<u>N60359</u>	<u>001</u>	
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>N60359</u>	<u>003</u>	

ERYTHROMYCIN STEARATE

<u>AB</u>	BARR	<u>EQ 250MG BASE</u>	<u>N61591</u>	<u>001</u>	
<u>AB</u>	MYLAN	<u>EQ 250MG BASE</u>	<u>N61505</u>	<u>001</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N61505</u>	<u>002</u>	

ESCITALOPRAM OXALATE

SOLUTION; ORAL

LEXAPRO

+	FOREST LABS	EQ 5MG BASE/5ML	N21365	001	Nov 27, 2002
---	-------------	-----------------	--------	-----	--------------

TABLET; ORAL

LEXAPRO

	FOREST LABS	5MG	N21323	001	Aug 14, 2002
		10MG	N21323	002	Aug 14, 2002
+		20MG	N21323	003	Aug 14, 2002

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

<u>AP</u>	+	BAXTER HLTHCARE CORP	<u>10MG/ML</u>	<u>N19386</u>	<u>006</u>	Feb 25, 2003
	+		20MG/ML	N19386	007	May 28, 2003
	+		250MG/ML	N19386	002	Dec 31, 1986
		BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER				
	+	BAXTER HLTHCARE CORP	2GM/100ML	N19386	005	Jan 27, 2003
		BREVIBLOC IN PLASTIC CONTAINER				
	+	BAXTER HLTHCARE CORP	1GM/100ML	N19386	004	Feb 16, 2001

ESMOLOL HYDROCHLORIDE

<u>AP</u>	AM PHARM	<u>10MG/ML</u>	<u>N76573</u>	<u>001</u>	May 02, 2005
<u>AP</u>	BEDFORD LABS	<u>10MG/ML</u>	<u>N76323</u>	<u>001</u>	Aug 10, 2004
<u>AP</u>	PHARMAFORCE	<u>10MG/ML</u>	<u>N76474</u>	<u>001</u>	May 02, 2005

ESOMEPRAZOLE MAGNESIUM

CAPSULE, DELAYED REL PELLETS; ORAL

NEXIUM

	ASTRAZENECA	EQ 20MG BASE	N21153	001	Feb 20, 2001
+		EQ 40MG BASE	N21153	002	Feb 20, 2001

ESOMEPRAZOLE SODIUM

INJECTABLE; INTRAVENOUS

NEXIUM IV

+	ASTRAZENECA	EQ 20MG BASE/VIAL	N21689	001	Mar 31, 2005
+		EQ 40MG BASE/VIAL	N21689	002	Mar 31, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 145 (of 360)

ESTAZOLAM

TABLET; ORAL

ESTAZOLAM

<u>AB</u>	IVAX PHARMS	<u>1MG</u>	<u>N74826</u>	<u>001</u>	Jul 03, 1997
<u>AB</u>		<u>2MG</u>	<u>N74826</u>	<u>002</u>	Jul 03, 1997
<u>AB</u>	TEVA	<u>1MG</u>	<u>N74921</u>	<u>001</u>	Jul 10, 1997
<u>AB</u>		<u>2MG</u>	<u>N74921</u>	<u>002</u>	Jul 10, 1997
<u>AB</u>	WATSON LABS	<u>1MG</u>	<u>N74818</u>	<u>001</u>	Aug 19, 1997
<u>AB</u>		<u>2MG</u>	<u>N74818</u>	<u>002</u>	Aug 19, 1997
<u>PROSOM</u>					
<u>AB</u>	ABBOTT	<u>1MG</u>	<u>N19080</u>	<u>001</u>	Dec 26, 1990
<u>AB</u>	+	<u>2MG</u>	<u>N19080</u>	<u>002</u>	Dec 26, 1990

ESTRADIOL

CREAM; VAGINAL

ESTRACE

+	WARNER CHILCOTT	0.01%	N86069	001	Jan 31, 1984
---	-----------------	-------	--------	-----	--------------

FILM, EXTENDED RELEASE; TRANSDERMAL

ALORA

BX	WATSON LABS	0.025MG/24HR	N20655	004	Apr 05, 2002
BX		0.05MG/24HR	N20655	001	Dec 20, 1996
BX		0.075MG/24HR	N20655	002	Dec 20, 1996
BX		0.1MG/24HR	N20655	003	Dec 20, 1996

CLIMARA

<u>AB2</u>	+	BERLEX	<u>0.025MG/24HR</u>	<u>N20375</u>	<u>004</u>	Mar 05, 1999
<u>AB2</u>	+		<u>0.05MG/24HR</u>	<u>N20375</u>	<u>001</u>	Dec 22, 1994
<u>AB2</u>	+		<u>0.075MG/24HR</u>	<u>N20375</u>	<u>003</u>	Mar 23, 1998
<u>AB2</u>	+		<u>0.1MG/24HR</u>	<u>N20375</u>	<u>002</u>	Dec 22, 1994
BX	+		0.0375MG/24HR	N20375	005	May 27, 2003
	+		0.06MG/24HR	N20375	006	May 27, 2003

ESTRADERM

BX	+	NOVARTIS	0.05MG/24HR	N19081	002	Sep 10, 1986
BX	+		0.1MG/24HR	N19081	003	Sep 10, 1986

ESTRADIOL

<u>AB2</u>	MYLAN TECHNOLOGIES	<u>0.025MG/24HR</u>	<u>N75182</u>	<u>003</u>	Jan 26, 2005
<u>AB2</u>		<u>0.05MG/24HR</u>	<u>N75233</u>	<u>001</u>	Feb 24, 2000
<u>AB2</u>		<u>0.075MG/24HR</u>	<u>N75182</u>	<u>002</u>	Jan 26, 2005
<u>AB2</u>		<u>0.1MG/24HR</u>	<u>N75182</u>	<u>001</u>	Feb 24, 2000

MENOSTAR

+	BERLEX LABS	0.014MG/24HR	N21674	001	Jun 08, 2004
---	-------------	--------------	--------	-----	--------------

VIVELLE

<u>AB1</u>	NOVARTIS	<u>0.05MG/24HR</u>	<u>N20323</u>	<u>002</u>	Oct 28, 1994
<u>AB1</u>		<u>0.1MG/24HR</u>	<u>N20323</u>	<u>004</u>	Oct 28, 1994

VIVELLE-DOT

<u>AB1</u>	+	NOVARTIS	<u>0.05MG/24HR</u>	<u>N20538</u>	<u>006</u>	Jan 08, 1999
<u>AB1</u>	+		<u>0.1MG/24HR</u>	<u>N20538</u>	<u>008</u>	Jan 08, 1999
BX	+		0.025MG/24HR	N20538	009	May 03, 2002
BX	+		0.0375MG/24HR	N20538	005	Jan 08, 1999
BX	+		0.075MG/24HR	N20538	007	Jan 08, 1999

GEL, METERED; TOPICAL

ESTROGEL

+	SOLVAY	0.06%	N21166	002	Feb 09, 2004
---	--------	-------	--------	-----	--------------

INSERT, EXTENDED RELEASE; VAGINAL

ESTRING

+	PHARMACIA AND UPJOHN	0.0075MG/24HR	N20472	001	Apr 26, 1996
---	----------------------	---------------	--------	-----	--------------

TABLET; ORAL

ESTRACE

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>0.5MG</u>	<u>N81295</u>	<u>001</u>	Jun 30, 1993
<u>AB</u>		<u>1MG</u>	<u>N84499</u>	<u>001</u>	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 146 (of 360)

ESTRADIOL

TABLET; ORAL

ESTRACE

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>2MG</u>	<u>N84500</u>	<u>001</u>	
-----------	---	----------------------	------------	---------------	------------	--

ESTRADIOL

<u>AB</u>		APPLIED ANAL	<u>0.5MG</u>	<u>N40138</u>	<u>001</u>	Jan 30, 1998
<u>AB</u>			<u>1MG</u>	<u>N40138</u>	<u>002</u>	Jan 30, 1998
<u>AB</u>			<u>2MG</u>	<u>N40138</u>	<u>003</u>	Jan 30, 1998
<u>AB</u>		BARR	<u>0.5MG</u>	<u>N40197</u>	<u>001</u>	Oct 22, 1997
<u>AB</u>			<u>1MG</u>	<u>N40197</u>	<u>002</u>	Oct 22, 1997
<u>AB</u>			<u>2MG</u>	<u>N40197</u>	<u>003</u>	Oct 22, 1997
<u>AB</u>		MYLAN	<u>0.5MG</u>	<u>N40326</u>	<u>001</u>	Apr 21, 1999
<u>AB</u>			<u>1MG</u>	<u>N40326</u>	<u>002</u>	Apr 21, 1999
<u>AB</u>			<u>2MG</u>	<u>N40326</u>	<u>003</u>	Apr 21, 1999
<u>AB</u>		USL PHARMA	<u>0.5MG</u>	<u>N40297</u>	<u>001</u>	Apr 17, 2002
<u>AB</u>			<u>1MG</u>	<u>N40297</u>	<u>002</u>	Apr 17, 2002
<u>AB</u>			<u>2MG</u>	<u>N40297</u>	<u>003</u>	Apr 17, 2002
<u>AB</u>		WATSON LABS	<u>0.5MG</u>	<u>N40114</u>	<u>003</u>	Mar 14, 1996
<u>AB</u>			<u>1MG</u>	<u>N40114</u>	<u>001</u>	Mar 14, 1996
<u>AB</u>			<u>2MG</u>	<u>N40114</u>	<u>002</u>	Mar 14, 1996

GYNODIOL

<u>AB</u>		DURAMED PHARMS BARR	<u>0.5MG</u>	<u>N40212</u>	<u>001</u>	Dec 29, 1997
<u>AB</u>			<u>1MG</u>	<u>N40212</u>	<u>002</u>	Dec 29, 1997
<u>AB</u>			<u>2MG</u>	<u>N40212</u>	<u>004</u>	Dec 29, 1997
			1.5MG	N40212	003	Dec 29, 1997

INNOFEM

<u>AB</u>		NOVO NORDISK INC	<u>0.5MG</u>	<u>N40312</u>	<u>001</u>	Nov 19, 1999
<u>AB</u>			<u>1MG</u>	<u>N40312</u>	<u>002</u>	Nov 19, 1999
<u>AB</u>			<u>2MG</u>	<u>N40312</u>	<u>003</u>	Nov 19, 1999

TABLET; VAGINAL

VAGIFEM

+	NOVO NORDISK INC	25UGM	N20908	001	Mar 26, 1999
---	------------------	-------	--------	-----	--------------

ESTRADIOL ACETATE

INSERT, EXTENDED RELEASE; VAGINAL

FEMRING

	GALEN LTD	EQ 0.05MG BASE/24HR	N21367	001	Mar 20, 2003
+		EQ 0.1MG BASE/24HR	N21367	002	Mar 20, 2003

TABLET; ORAL

FEMTRACE

	WARNER CHILCOTT	0.45MG	N21633	001	Aug 20, 2004
		0.9MG	N21633	002	Aug 20, 2004
+		1.8MG	N21633	003	Aug 20, 2004

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION

DEPO-ESTRADIOL

<u>AO</u>	+	PHARMACIA AND UPJOHN	<u>5MG/ML</u>	<u>N85470</u>	<u>003</u>
-----------	---	----------------------	---------------	---------------	------------

ESTRADIOL CYPIONATE

<u>AO</u>		STERIS	<u>5MG/ML</u>	<u>N85620</u>	<u>001</u>
-----------	--	--------	---------------	---------------	------------

ESTRADIOL HEMIHYDRATE

EMULSION; TOPICAL

ESTRASORB

+	NOVAVAX	0.25%	N21371	001	Oct 09, 2003
---	---------	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 147 (of 360)

ESTRADIOL VALERATE

INJECTABLE; INJECTION

DELESTROGEN

<u>AO</u>	+	KING PHARMS	<u>20MG/ML</u>	<u>N09402</u>	<u>004</u>	
<u>AO</u>	+		<u>40MG/ML</u>	<u>N09402</u>	<u>003</u>	
	+		10MG/ML	N09402	002	

ESTRADIOL VALERATE

<u>AO</u>		STERIS	<u>20MG/ML</u>	<u>N83547</u>	<u>001</u>	
<u>AO</u>			<u>40MG/ML</u>	<u>N83714</u>	<u>001</u>	

ESTRADIOL; LEVONORGESTRELFILM, EXTENDED RELEASE; TRANSDERMAL
CLIMARA PRO

+	BERLEX	0.045MG/24HR;0.015MG/24HR	N21258	001	Nov 21, 2003
---	--------	---------------------------	--------	-----	--------------

ESTRADIOL; NORETHINDRONE ACETATEFILM, EXTENDED RELEASE; TRANSDERMAL
COMBIPATCH

	NOVARTIS	0.05MG/24HR;0.14MG/24HR	N20870	001	Aug 07, 1998
+		0.05MG/24HR;0.25MG/24HR	N20870	002	Aug 07, 1998

TABLET; ORAL
ACTIVELLA

+	NOVO NORDISK INC	1MG;0.5MG	N20907	001	Nov 18, 1998
---	------------------	-----------	--------	-----	--------------

ESTRADIOL; NORGESTIMATE

TABLET; ORAL

ESTRADIOL AND NORGESTIMATE

<u>AB</u>		BARR	<u>1MG, 1MG; N/A, 0.09MG</u>	<u>N76812</u>	<u>001</u>	Apr 29, 2005
-----------	--	------	------------------------------	---------------	------------	--------------

PREFEST

<u>AB</u>	+	DURAMED	<u>1MG, 1MG; N/A, 0.09MG</u>	<u>N21040</u>	<u>001</u>	Oct 22, 1999
-----------	---	---------	------------------------------	---------------	------------	--------------

ESTRAMUSTINE PHOSPHATE SODIUMCAPSULE; ORAL
EMCYT

+	PHARMACIA AND UPJOHN	EQ 140MG PHOSPHATE	N18045	001	
---	----------------------	--------------------	--------	-----	--

ESTROGENS, CONJUGATEDCREAM; TOPICAL, VAGINAL
PREMARIN

+	WYETH PHARMS INC	0.625MG/GM	N20216	001	
---	------------------	------------	--------	-----	--

INJECTABLE; INJECTION
PREMARIN

+	WYETH PHARMS INC	25MG/VIAL	N10402	001	
---	------------------	-----------	--------	-----	--

TABLET; ORAL
PREMARIN

	WYETH PHARMS INC	0.3MG	N04782	003	
		0.45MG	N04782	006	Jul 16, 2003
+		0.625MG	N04782	004	
		0.9MG	N04782	005	Jan 26, 1984
+		1.25MG	N04782	001	

ESTROGENS, CONJUGATED SYNTHETIC ATABLET; ORAL
CENESTIN

	DURAMED	0.3MG	N20992	001	Jun 21, 2002
		0.45MG	N20992	005	Feb 05, 2004
		0.625MG	N20992	002	Mar 24, 1999
		0.9MG	N20992	003	Mar 24, 1999
+		1.25MG	N20992	004	Mar 13, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 148 (of 360)

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28

PREMPHASE 14/14

+ WYETH PHARMS INC	0.625MG; 0.625MG; N/A, 5MG	N20527	002	Nov 17, 1995
PREMPRO				
+ WYETH PHARMS INC	0.3MG; 1.5MG	N20527	005	Jun 04, 2003
+	0.45MG; 1.5MG	N20527	004	Mar 12, 2003
+	0.625MG; 2.5MG	N20527	001	Nov 17, 1995
+	0.625MG; 5MG	N20527	003	Jan 09, 1998

ESTROGENS, ESTERIFIED

TABLET; ORAL

MENEST

MONARCH PHARMS	0.3MG	N84951	001	
	0.625MG	N84948	001	
	1.25MG	N84950	001	
+	2.5MG	N84949	001	

ESTRONE

INJECTABLE; INJECTION

ESTRONE

+ STERIS	5MG/ML	N85239	001	
----------	--------	--------	-----	--

ESTROPIPATE

CREAM; VAGINAL

OGEN

+ PHARMACIA AND UPJOHN	1.5MG/GM	N84710	001	
------------------------	----------	--------	-----	--

TABLET; ORAL

ESTROPIPATE

<u>AB</u>	BARR	<u>0.75MG</u>	<u>N40135</u>	<u>001</u>	Nov 27, 1996
<u>AB</u>		<u>1.5MG</u>	<u>N40135</u>	<u>002</u>	Nov 27, 1996
<u>AB</u>		<u>3MG</u>	<u>N40135</u>	<u>003</u>	Nov 27, 1996
<u>AB</u>	DURAMED PHARMS BARR	<u>0.75MG</u>	<u>N40296</u>	<u>001</u>	Nov 01, 1999
<u>AB</u>		<u>1.5MG</u>	<u>N40296</u>	<u>002</u>	Nov 01, 1999
<u>AB</u>		<u>3MG</u>	<u>N40296</u>	<u>003</u>	Nov 01, 1999
<u>AB</u>	MYLAN	<u>0.75MG</u>	<u>N40359</u>	<u>001</u>	Aug 26, 1999
<u>AB</u>		<u>1.5MG</u>	<u>N40359</u>	<u>002</u>	Aug 26, 1999
<u>AB</u>		<u>3MG</u>	<u>N40359</u>	<u>003</u>	Aug 26, 1999
<u>AB</u>	WATSON LABS	<u>0.75MG</u>	<u>N81213</u>	<u>001</u>	Sep 23, 1993
<u>AB</u>		<u>1.5MG</u>	<u>N81214</u>	<u>001</u>	Sep 23, 1993
<u>AB</u>		<u>3MG</u>	<u>N81215</u>	<u>001</u>	Sep 23, 1993
<u>AB</u>		<u>6MG</u>	<u>N81216</u>	<u>001</u>	Sep 23, 1993
	<u>OGEN .625</u>				
<u>AB</u>	PHARMACIA AND UPJOHN	<u>0.75MG</u>	<u>N83220</u>	<u>001</u>	
	<u>OGEN 1.25</u>				
<u>AB</u>	PHARMACIA AND UPJOHN	<u>1.5MG</u>	<u>N83220</u>	<u>002</u>	
	<u>OGEN 2.5</u>				
<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>3MG</u>	<u>N83220</u>	<u>003</u>	
	<u>OGEN 5</u>				
<u>AB</u>	PHARMACIA AND UPJOHN	<u>6MG</u>	<u>N83220</u>	<u>004</u>	
	<u>ORTHO-EST</u>				
<u>AB</u>	SUN PHARM INDS (IN)	<u>0.75MG</u>	<u>N89567</u>	<u>001</u>	Feb 27, 1991
<u>AB</u>		<u>1.5MG</u>	<u>N89582</u>	<u>001</u>	Jul 17, 1991

ESZOPICLONE

TABLET; ORAL

LUNESTA

SEPRACOR	1MG	N21476	001	Dec 15, 2004
	2MG	N21476	002	Dec 15, 2004
+	3MG	N21476	003	Dec 15, 2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 149 (of 360)

ETHACRYNATE SODIUM

INJECTABLE; INJECTION

EDECIN

+ MERCK

EQ 50MG BASE/VIAL

N16093 001

ETHACRYNIC ACID

TABLET; ORAL

EDECIN

+ MERCK

25MG

N16092 001

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

ETHAMBUTOL HYDROCHLORIDEAB BARR 400MGN76057 001

Nov 26, 2001

AB WEST WARD 100MGN75095 001

Nov 30, 1999

AB + 400MGN75095 002

Nov 30, 1999

MYAMBUTOLAB STAT TRADE 100MGN16320 001AB 400MGN16320 003ETHANOLAMINE OLEATE

INJECTABLE; INJECTION

ETHAMOLIN

+ QOL MEDCL

50MG/ML

N19357 001

Dec 22, 1988

ETHINYL ESTRADIOL; ETHYNODIOL DIACETATE

TABLET; ORAL-21

DEMULEN 1/35-21AB + GD SEARLE LLC 0.035MG;1MGN18168 001DEMULEN 1/50-21AB + GD SEARLE LLC 0.05MG;1MGN16927 001ZOVIA 1/35E-21AB WATSON LABS 0.035MG;1MGN72720 001

Dec 30, 1991

ZOVIA 1/50E-21AB WATSON LABS 0.05MG;1MGN72722 001

Dec 30, 1991

TABLET; ORAL-28

DEMULEN 1/35-28AB + GD SEARLE LLC 0.035MG;1MGN18160 001DEMULEN 1/50-28AB GD SEARLE LLC 0.05MG;1MGN16936 001KELNORAB BARR 0.035MG;1MGN76785 001

May 23, 2005

ZOVIA 1/35E-28AB WATSON LABS 0.035MG;1MGN72721 001

Dec 30, 1991

ZOVIA 1/50E-28AB WATSON LABS 0.05MG;1MGN72723 001

Dec 30, 1991

ETHINYL ESTRADIOL; ETONOGESTREL

RING; VAGINAL

NUVARING

+ ORGANON USA INC

0.015MG;0.12MG

N21187 001

Oct 03, 2001

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

SEASONALE

+ DURAMED

0.03MG;0.15MG

N21544 001

Sep 05, 2003

TABLET; ORAL-28

ALESSEAB1 + WYETH PHARMS INC 0.02MG;0.1MGN20683 002

Mar 27, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 150 (of 360)

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-28

<u>AVIANE-28</u>					
<u>AB1</u>	DURAMED PHARMS BARR	<u>0.02MG;0.1MG</u>	<u>N75796</u>	<u>001</u>	Apr 30, 2001
<u>ENPRESSE-28</u>					
<u>AB</u>	DURAMED PHARMS BARR	<u>0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG</u>	<u>N75809</u>	<u>002</u>	Jul 16, 2001
<u>LESSINA-28</u>					
<u>AB2</u>	BARR	<u>0.02MG;0.1MG</u>	<u>N75803</u>	<u>002</u>	Mar 20, 2002
<u>LEVLITE</u>					
<u>AB2</u> +	BERLEX	<u>0.02MG;0.1MG</u>	<u>N20860</u>	<u>002</u>	Jul 13, 1998
<u>LEVONORGESTREL AND ETHINYL ESTRADIOL</u>					
<u>AB1</u>	BARR	<u>0.02MG;0.1MG</u>	<u>N75862</u>	<u>002</u>	Apr 29, 2003
<u>AB1</u>	WATSON LABS	<u>0.02MG;0.1MG</u>	<u>N76625</u>	<u>001</u>	Nov 18, 2004
<u>LEVORA 0.15/30-28</u>					
<u>AB</u>	WATSON LABS	<u>0.03MG;0.15MG</u>	<u>N73594</u>	<u>001</u>	Dec 13, 1993
<u>NORDETTE-28</u>					
<u>AB</u> +	DURAMED	<u>0.03MG;0.15MG</u>	<u>N18782</u>	<u>001</u>	Jul 21, 1982
<u>PORTIA-28</u>					
<u>AB</u>	BARR	<u>0.03MG;0.15MG</u>	<u>N75866</u>	<u>002</u>	May 23, 2002
<u>TRIPHASIL-28</u>					
<u>AB</u> +	WYETH PHARMS INC	<u>0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG</u>	<u>N19190</u>	<u>001</u>	Nov 01, 1984
<u>TRIVORA-28</u>					
<u>AB</u>	WATSON LABS	<u>0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG</u>	<u>N74538</u>	<u>002</u>	Dec 18, 1997

ETHINYL ESTRADIOL; NORELGESTROMIN

FILM, EXTENDED RELEASE; TRANSDERMAL

ORTHO EVRA

+ ORTHO MCNEIL PHARM 0.02MG/24HR;0.15MG/24HR N21180 001 Nov 20, 2001

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

BALZIVA-21					
+ BARR		0.035MG;0.4MG	N76198	001	Apr 22, 2004
<u>BREVICON 21-DAY</u>					
<u>AB</u> +	WATSON LABS	<u>0.035MG;0.5MG</u>	<u>N17566</u>	<u>001</u>	
<u>GENCEPT 10/11-21</u>					
<u>AB</u>	BARR	<u>0.035MG,0.035MG;0.5MG,1MG</u>	<u>N72694</u>	<u>001</u>	Feb 28, 1992
<u>NORETHIN 1/35E-21</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG;1MG</u>	<u>N71480</u>	<u>001</u>	Apr 12, 1988
<u>NORETHINDRONE AND ETHINYL ESTRADIOL</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG;0.5MG</u>	<u>N70684</u>	<u>001</u>	Jan 29, 1987
<u>AB</u>		<u>0.035MG;1MG</u>	<u>N70685</u>	<u>001</u>	Jan 29, 1987
<u>NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG,0.035MG;0.5MG,1MG</u>	<u>N71043</u>	<u>001</u>	Apr 01, 1988
<u>NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)</u>					
+ WATSON LABS		0.035MG,0.035MG;0.5MG,1MG	N71041	001	Sep 24, 1991
<u>NORINYL 1+35 21-DAY</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG;1MG</u>	<u>N17565</u>	<u>001</u>	
<u>NORTREL 0.5/35-21</u>					
<u>AB</u>	BARR	<u>0.035MG;0.5MG</u>	<u>N72692</u>	<u>001</u>	Feb 28, 1992
<u>NORTREL 1/35-21</u>					
<u>AB</u>	BARR	<u>0.035MG;1MG</u>	<u>N72693</u>	<u>001</u>	Feb 28, 1992
<u>NORTREL 7/7/7</u>					
	BARR	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	N75478	001	Aug 30, 2002
<u>ARANELLE</u>					
<u>AB</u>	BARR	<u>0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG</u>	<u>N76783</u>	<u>001</u>	Sep 29, 2004

TABLET; ORAL-28

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 151 (of 360)

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

<u>BALZIVA-28</u>					
<u>AB</u>	BARR	<u>0.035MG;0.4MG</u>	<u>N76238</u>	<u>001</u>	Apr 22, 2004
<u>BREVICON 28-DAY</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG;0.5MG</u>	<u>N17743</u>	<u>001</u>	
<u>GENCEPT 10/11-28</u>					
<u>AB</u>	BARR	<u>0.035MG,0.035MG;0.5MG,1MG</u>	<u>N72697</u>	<u>001</u>	Feb 28, 1992
<u>MODICON 28</u>					
<u>AB</u>	ORTHO MCNEIL PHARM	<u>0.035MG;0.5MG</u>	<u>N17735</u>	<u>001</u>	
<u>NORETHIN 1/35E-28</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG;1MG</u>	<u>N71481</u>	<u>001</u>	Apr 12, 1988
<u>NORETHINDRONE AND ETHINYL ESTRADIOL</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG;0.5MG</u>	<u>N70686</u>	<u>001</u>	Jan 29, 1987
<u>AB</u>		<u>0.035MG;1MG</u>	<u>N70687</u>	<u>001</u>	Jan 29, 1987
<u>NORETHINDRONE AND ETHINYL ESTRADIOL (10/11)</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG,0.035MG;0.5MG,1MG</u>	<u>N71044</u>	<u>001</u>	Apr 01, 1988
<u>NORETHINDRONE AND ETHINYL ESTRADIOL (7/14)</u>					
	WATSON LABS	0.035MG,0.035MG;0.5MG,1MG	N71042	001	Sep 24, 1991
<u>NORINYL 1+35 28-DAY</u>					
<u>AB</u>	WATSON LABS	<u>0.035MG;1MG</u>	<u>N17565</u>	<u>002</u>	
<u>NORTREL 0.5/35-28</u>					
<u>AB</u>	BARR	<u>0.035MG;0.5MG</u>	<u>N72695</u>	<u>001</u>	Feb 28, 1992
<u>NORTREL 1/35-28</u>					
<u>AB</u>	BARR	<u>0.035MG;1MG</u>	<u>N72696</u>	<u>001</u>	Feb 28, 1992
<u>NORTREL 7/7/7</u>					
<u>AB</u>	BARR	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>N75478</u>	<u>002</u>	Aug 30, 2002
<u>ORTHO-NOVUM 1/35-28</u>					
<u>AB</u>	+ ORTHO MCNEIL PHARM	<u>0.035MG;1MG</u>	<u>N17919</u>	<u>002</u>	
<u>ORTHO-NOVUM 10/11-28</u>					
<u>AB</u>	+ ORTHO MCNEIL PHARM	<u>0.035MG,0.035MG;0.5MG,1MG</u>	<u>N18354</u>	<u>002</u>	Jan 11, 1982
<u>ORTHO-NOVUM 7/7/7-28</u>					
<u>AB</u>	+ ORTHO MCNEIL PHARM	<u>0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG</u>	<u>N18985</u>	<u>002</u>	Apr 04, 1984
<u>OVCON-35</u>					
<u>AB</u>	WARNER CHILCOTT	<u>0.035MG;0.4MG</u>	<u>N17716</u>	<u>001</u>	
<u>OVCON-50</u>					
	+ WARNER CHILCOTT	0.05MG;1MG	N17576	001	
<u>TRI-NORINYL 28-DAY</u>					
<u>AB</u>	+ WATSON LABS	<u>0.035MG,0.035MG,0.035MG;0.5MG,1MG,0.5MG</u>	<u>N18977</u>	<u>002</u>	Apr 13, 1984

TABLET, CHEWABLE; ORAL-28

OVCON-35

	+ WARNER CHILCOTT	0.035MG;0.4MG	N21490	001	Nov 14, 2003
--	-------------------	---------------	--------	-----	--------------

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

FEMHRT

	WARNER CHILCOTT	0.0025MG;0.5MG	N21065	001	Jan 14, 2005
	+	0.005MG;1MG	N21065	002	Oct 15, 1999

TABLET; ORAL-21

JUNEL 1.5/30

<u>AB</u>	BARR	<u>0.03MG;1.5MG</u>	<u>N76381</u>	<u>001</u>	May 30, 2003
------------------	------	---------------------	---------------	------------	--------------

JUNEL 1/20

<u>AB</u>	BARR	<u>0.02MG;1MG</u>	<u>N76380</u>	<u>001</u>	May 30, 2003
------------------	------	-------------------	---------------	------------	--------------

LOESTRIN 21 1.5/30

<u>AB</u>	+ WARNER CHILCOTT	<u>0.03MG;1.5MG</u>	<u>N17875</u>	<u>001</u>	
------------------	-------------------	---------------------	---------------	------------	--

LOESTRIN 21 1/20

<u>AB</u>	+ WARNER CHILCOTT	<u>0.02MG;1MG</u>	<u>N17876</u>	<u>001</u>	
------------------	-------------------	-------------------	---------------	------------	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 152 (of 360)

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-28						
ESTROSTEP FE						
	+ WARNER CHILCOTT	0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG	N20130	002		Oct 09, 1996
	<u>JUNEL FE 1.5/30</u>					
<u>AB</u>	BARR	<u>0.03MG; 1.5MG</u>	<u>N76064</u>	<u>001</u>		Sep 18, 2003
	<u>JUNEL FE 1/20</u>					
<u>AB</u>	BARR	<u>0.02MG; 1MG</u>	<u>N76081</u>	<u>001</u>		Sep 18, 2003
	<u>LOESTRIN FE 1.5/30</u>					
<u>AB</u>	+ WARNER CHILCOTT	<u>0.03MG; 1.5MG</u>	<u>N17355</u>	<u>001</u>		
	<u>LOESTRIN FE 1/20</u>					
<u>AB</u>	+ WARNER CHILCOTT	<u>0.02MG; 1MG</u>	<u>N17354</u>	<u>001</u>		
	<u>MICROGESTIN FE 1.5/30</u>					
<u>AB</u>	WATSON LABS	<u>0.03MG; 1.5MG</u>	<u>N75548</u>	<u>001</u>		Feb 05, 2001
	<u>MICROGESTIN FE 1/20</u>					
<u>AB</u>	WATSON LABS	<u>0.02MG; 1MG</u>	<u>N75647</u>	<u>001</u>		Feb 05, 2001
<u>NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE</u>						
<u>AB</u>	ANDRX PHARMS	<u>0.02MG; 1MG</u>	<u>N77077</u>	<u>001</u>		May 20, 2005
<u>AB</u>		<u>0.03MG; 1.5MG</u>	<u>N77075</u>	<u>001</u>		Apr 28, 2005

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28						
<u>ORTHO CYCLEN-28</u>						
<u>AB</u>	+ JOHNSON RW	<u>0.035MG; 0.25MG</u>	<u>N19653</u>	<u>002</u>		Dec 29, 1989
	<u>ORTHO TRI-CYCLEN</u>					
<u>AB</u>	+ JOHNSON RW	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>N19697</u>	<u>001</u>		Jul 03, 1992
	ORTHO TRI-CYCLEN LO					
	+ ORTHO MCNEIL PHARM	0.025MG, 0.025MG, 0.25MG; 0.18MG, 0.215MG, 0.25MG	N21241	001		Aug 22, 2002
	<u>PREVIFEM</u>					
<u>AB</u>	ANDRX PHARMS	<u>0.035MG; 0.25MG</u>	<u>N76334</u>	<u>001</u>		Jan 09, 2004
	<u>SPRINTEC</u>					
<u>AB</u>	BARR	<u>0.035MG; 0.25MG</u>	<u>N75804</u>	<u>001</u>		Sep 25, 2002
	<u>TRI-PREVIFEM</u>					
<u>AB</u>	ANDRX PHARMS	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>N76335</u>	<u>001</u>		Mar 26, 2004
	<u>TRI-SPRINTEC</u>					
<u>AB</u>	BARR	<u>0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG</u>	<u>N75808</u>	<u>001</u>		Dec 29, 2003

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21						
<u>CRYSELLE</u>						
<u>AB</u>	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>N75840</u>	<u>001</u>		Nov 30, 2001
	<u>LOW-OGESTREL-21</u>					
<u>AB</u>	+ WATSON LABS	<u>0.03MG; 0.3MG</u>	<u>N75288</u>	<u>001</u>		Jul 28, 1999
	OGESTREL 0.5/50-21					
	+ WATSON LABS	0.05MG; 0.5MG	N75406	001		Dec 15, 1999
TABLET; ORAL-28						
<u>CRYSELLE</u>						
<u>AB</u>	DURAMED PHARMS BARR	<u>0.03MG; 0.3MG</u>	<u>N75840</u>	<u>002</u>		Nov 30, 2001
	<u>LO/OVRAL-28</u>					
<u>AB</u>	WYETH PHARMS INC	<u>0.03MG; 0.3MG</u>	<u>N17802</u>	<u>001</u>		
	<u>LOW-OGESTREL-28</u>					
<u>AB</u>	WATSON LABS	<u>0.03MG; 0.3MG</u>	<u>N75288</u>	<u>002</u>		Jul 28, 1999
	OGESTREL 0.5/50-28					
	+ WATSON LABS	0.05MG; 0.5MG	N75406	002		Dec 15, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 153 (of 360)

ETHIODIZED OIL

OIL; INTRALYMPHATIC

ETHIODOL

SAVAGE LABS

99%

N09190 001

ETHIONAMIDE

TABLET; ORAL

TRECATOR-SC

+ WYETH PHARMS INC

250MG

N13026 002

ETHOSUXIMIDE

CAPSULE; ORAL

ETHOSUXIMIDEAB BANNER PHARMACAPS250MGN40430 001

Oct 28, 2002

ZARONTINAB + PARKE DAVIS250MGN12380 001

SYRUP; ORAL

ETHOSUXIMIDEAA MIKART250MG/5MLN40506 001

Dec 22, 2003

AA PHARM ASSOC250MG/5MLN40253 001

Nov 22, 2000

AA TEVA PHARMS250MG/5MLN81306 001

Jul 30, 1993

ZARONTINAA + PARKE DAVIS250MG/5MLN80258 001ETHOTOIN

TABLET; ORAL

PEGANONE

+ OVATION PHARMS

250MG

N10841 001

ETIDRONATE DISODIUM

TABLET; ORAL

DIDRONELAB PROCTER AND GAMBLE200MGN17831 001AB +400MGN17831 002ETIDRONATE DISODIUMAB GENPHARM200MGN75800 001

Jan 24, 2003

AB400MGN75800 002

Jan 24, 2003

ETODOLAC

CAPSULE; ORAL

ETODOLACAB AAIPHARMA LLC300MGN74929 001

Jan 30, 1998

AB ENDO PHARMS200MGN74842 001

Jul 17, 1997

AB300MGN74842 002

Jul 17, 1997

AB GENPHARM200MGN75071 001

Sep 30, 1998

AB300MGN75071 002

Sep 30, 1998

AB MYLAN200MGN74932 001

May 16, 1997

AB300MGN74932 002

May 16, 1997

AB SANDOZ200MGN74840 001

Aug 29, 1997

AB200MGN74942 001

Sep 30, 1997

AB300MGN74840 002

Aug 29, 1997

AB300MGN74942 002

Sep 30, 1997

AB TARO200MGN75078 001

Apr 30, 1998

AB300MGN75078 002

Apr 30, 1998

AB TEVA200MGN75126 001

Sep 16, 1999

AB300MGN75126 002

Sep 16, 1999

AB TORPHARM200MGN75419 001

Jul 28, 2000

AB300MGN75419 002

Jul 28, 2000

AB WATSON LABS200MGN74844 001

Dec 23, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 154 (of 360)

ETODOLAC

CAPSULE; ORAL

ETODOLAC

<u>AB</u>	WATSON LABS	<u>300MG</u>	<u>N74844</u>	<u>002</u>	Dec 23, 1997
-----------	-------------	--------------	---------------	------------	--------------

LODINE

<u>AB</u>	+ WYETH PHARMS INC	<u>300MG</u>	<u>N18922</u>	<u>003</u>	Jan 31, 1991
-----------	--------------------	--------------	---------------	------------	--------------

TABLET; ORAL

ETODOLAC

<u>AB</u>	AAIPHARMA LLC	<u>400MG</u>	<u>N74927</u>	<u>001</u>	Oct 30, 1997
-----------	---------------	--------------	---------------	------------	--------------

<u>AB</u>	APOTEX	<u>400MG</u>	<u>N76004</u>	<u>001</u>	Dec 03, 2002
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N76004</u>	<u>002</u>	Dec 03, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	ENDO PHARMS	<u>400MG</u>	<u>N74841</u>	<u>001</u>	Jun 27, 1997
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>	GENPHARM	<u>400MG</u>	<u>N75012</u>	<u>001</u>	Sep 30, 1998
-----------	----------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N75012</u>	<u>002</u>	Sep 30, 1998
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	IVAX PHARMS	<u>400MG</u>	<u>N74883</u>	<u>001</u>	Feb 28, 1997
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N74883</u>	<u>002</u>	Nov 20, 1998
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>400MG</u>	<u>N75104</u>	<u>001</u>	Feb 06, 1998
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N75104</u>	<u>002</u>	Nov 20, 1998
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	PUREPAC PHARM	<u>400MG</u>	<u>N74819</u>	<u>001</u>	Feb 28, 1997
-----------	---------------	--------------	---------------	------------	--------------

<u>AB</u>	RANBAXY	<u>400MG</u>	<u>N75226</u>	<u>001</u>	Nov 24, 1998
-----------	---------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N75226</u>	<u>002</u>	Nov 24, 1998
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>400MG</u>	<u>N74839</u>	<u>001</u>	Jul 11, 1997
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>400MG</u>	<u>N74846</u>	<u>001</u>	Feb 28, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>400MG</u>	<u>N74903</u>	<u>001</u>	Apr 11, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N74903</u>	<u>002</u>	Apr 19, 1999
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	TARO PHARM INDS	<u>400MG</u>	<u>N75074</u>	<u>001</u>	Mar 11, 1998
-----------	-----------------	--------------	---------------	------------	--------------

<u>AB</u>	+	<u>500MG</u>	<u>N75074</u>	<u>002</u>	Apr 25, 2000
-----------	---	--------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>400MG</u>	<u>N74847</u>	<u>001</u>	Apr 23, 1999
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>400MG</u>	<u>N75009</u>	<u>001</u>	Nov 26, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N74847</u>	<u>002</u>	Apr 23, 1999
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N75009</u>	<u>002</u>	Dec 28, 1999
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	WATSON LABS	<u>400MG</u>	<u>N74892</u>	<u>001</u>	Apr 16, 1997
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>400MG</u>	<u>N75069</u>	<u>001</u>	Apr 16, 1998
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N74892</u>	<u>002</u>	Oct 29, 1998
-----------	--	--------------	---------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC

<u>AB</u>	ANDRX PHARMS	<u>400MG</u>	<u>N75829</u>	<u>001</u>	Nov 30, 2001
-----------	--------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N75829</u>	<u>002</u>	Nov 30, 2001
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>400MG</u>	<u>N75943</u>	<u>001</u>	Jul 26, 2002
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N75943</u>	<u>002</u>	Jul 26, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>600MG</u>	<u>N75943</u>	<u>003</u>	Jul 26, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	TARO	<u>400MG</u>	<u>N76174</u>	<u>001</u>	Mar 13, 2003
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N76174</u>	<u>002</u>	Mar 13, 2003
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>600MG</u>	<u>N76174</u>	<u>003</u>	Mar 13, 2003
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>400MG</u>	<u>N75665</u>	<u>003</u>	Feb 05, 2001
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N75665</u>	<u>002</u>	Jul 31, 2000
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	+	<u>600MG</u>	<u>N75665</u>	<u>001</u>	Jul 31, 2000
-----------	---	--------------	---------------	------------	--------------

ETOMIDATE

INJECTABLE; INJECTION

AMIDATE

<u>AP</u>	+	HOSPIRA	<u>2MG/ML</u>	<u>N18227</u>	<u>001</u>	Sep 07, 1982
-----------	---	---------	---------------	---------------	------------	--------------

ETOMIDATE

<u>AP</u>		BEDFORD	<u>2MG/ML</u>	<u>N74593</u>	<u>001</u>	Nov 04, 1996
-----------	--	---------	---------------	---------------	------------	--------------

ETOPOSIDE

CAPSULE; ORAL

ETOPOSIDE

<u>AB</u>	GENPHARM	<u>50MG</u>	<u>N75635</u>	<u>001</u>	Sep 19, 2001
-----------	----------	-------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 155 (of 360)

ETOPOSIDE

CAPSULE; ORAL

VEPESID

<u>AB</u>	+	BRISTOL MYERS SQUIBB	<u>50MG</u>	<u>N19557</u>	<u>001</u>	Dec 30, 1986
-----------	---	----------------------	-------------	---------------	------------	--------------

INJECTABLE; INJECTION

ETOPOSIDE

<u>AP</u>		AM PHARM PARTNERS	<u>20MG/ML</u>	<u>N74983</u>	<u>001</u>	Sep 30, 1998
<u>AP</u>		BEDFORD	<u>20MG/ML</u>	<u>N74290</u>	<u>001</u>	Jul 17, 1995
<u>AP</u>		HOSPIRA	<u>20MG/ML</u>	<u>N74320</u>	<u>001</u>	Aug 30, 1995
<u>AP</u>			<u>20MG/ML</u>	<u>N74351</u>	<u>001</u>	Aug 30, 1995
<u>AP</u>		MARSAM PHARMS LLC	<u>20MG/ML</u>	<u>N74968</u>	<u>001</u>	Jan 09, 1998
<u>AP</u>		PHARMACHEMIE	<u>20MG/ML</u>	<u>N74227</u>	<u>001</u>	Feb 22, 1996
<u>AP</u>		SICOR PHARMS	<u>20MG/ML</u>	<u>N74284</u>	<u>001</u>	Feb 10, 1994
<u>AP</u>			<u>20MG/ML</u>	<u>N74529</u>	<u>001</u>	Jul 24, 1996
<u>AP</u>		SUPERGEN	<u>20MG/ML</u>	<u>N74513</u>	<u>001</u>	Mar 14, 1996

TOPOSAR

<u>AP</u>		SICOR PHARMS	<u>20MG/ML</u>	<u>N74166</u>	<u>001</u>	Feb 27, 1995
-----------	--	--------------	----------------	---------------	------------	--------------

VEPESID

<u>AP</u>	+	BRISTOL MYERS SQUIBB	<u>20MG/ML</u>	<u>N18768</u>	<u>001</u>	Nov 10, 1983
-----------	---	----------------------	----------------	---------------	------------	--------------

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION

ETOPOPHOS PRESERVATIVE FREE

	+	BRISTOL MYERS SQUIBB	EQ 100MG BASE/VIAL	N20457	001	May 17, 1996
--	---	----------------------	--------------------	--------	-----	--------------

EXEMESTANE

TABLET; ORAL

AROMASIN

	+	PHARMACIA AND UPJOHN	25MG	N20753	001	Oct 21, 1999
--	---	----------------------	------	--------	-----	--------------

EXENATIDE SYNTHETIC

INJECTABLE; SUBCUTANEOUS

BYETTA

	+	AMYLIN	300UGM/1.2ML (250UGM/ML)	N21773	001	Apr 28, 2005
	+		600UGM/2.4ML (250UGM/ML)	N21773	002	Apr 28, 2005

EZETIMIBE

TABLET; ORAL

ZETIA

	+	MSP SINGAPORE	10MG	N21445	001	Oct 25, 2002
--	---	---------------	------	--------	-----	--------------

EZETIMIBE; SIMVASTATIN

TABLET; ORAL

VYTORIN

		MSP SINGAPORE	10MG;10MG	N21687	001	Jul 23, 2004
			10MG;20MG	N21687	002	Jul 23, 2004
			10MG;40MG	N21687	003	Jul 23, 2004
	+		10MG;80MG	N21687	004	Jul 23, 2004

FAMCICLOVIR

TABLET; ORAL

FAMVIR

		NOVARTIS	125MG	N20363	003	Dec 11, 1995
			250MG	N20363	001	Apr 26, 1996
	+		500MG	N20363	002	Jun 29, 1994

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 156 (of 360)

FAMOTIDINE

FOR SUSPENSION; ORAL

PEPCID

+ MERCK

40MG/5ML

N19527 001

Feb 02, 1987

INJECTABLE; INJECTION

FAMOTIDINEAP AM PHARM PARTNERS10MG/MLN75709 001

Apr 16, 2001

AP APOTEX10MG/MLN75942 001

Aug 02, 2002

AP BAXTER HLTHCARE10MG/MLN75488 001

Apr 16, 2001

AP BEDFORD10MG/MLN75799 001

Apr 30, 2002

AP BEDFORD10MG/MLN75651 001

Apr 16, 2001

AP BEDFORD10MG/MLN75684 001

Apr 16, 2001

AP HOSPIRA10MG/MLN75870 001

Nov 23, 2001

AP HOSPIRA10MG/MLN75905 001

Nov 23, 2001

FAMOTIDINE PRESERVATIVE FREEAP AM PHARM PARTNERS10MG/MLN75813 001

Apr 16, 2001

AP APOTEX10MG/MLN76324 001

Nov 27, 2002

AP BAXTER HLTHCARE10MG/MLN75486 001

Apr 16, 2001

AP BEDFORD10MG/MLN75789 001

Apr 30, 2002

AP BEDFORD10MG/MLN75622 001

Apr 16, 2001

AP BEN VENUE10MG/MLN75825 001

Apr 17, 2001

FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK)AP APOTEX10MG/MLN76322 001

Nov 27, 2002

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINERAP BAXTER HLTHCARE0.4MG/MLN75591 001

May 10, 2001

PEPCIDAP + MERCK10MG/MLN19510 001

Nov 04, 1986

PEPCID PRESERVATIVE FREEAP + MERCK10MG/MLN19510 004

Nov 04, 1986

PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINERAP + MERCK0.4MG/MLN20249 001

Feb 18, 1994

TABLET; ORAL

FAMOTIDINEAB CARLSBAD20MGN75805 001

Apr 16, 2001

AB CARLSBAD40MGN75805 002

Apr 16, 2001

AB DR REDDYS LABS LTD20MGN75718 001

Apr 16, 2001

AB DR REDDYS LABS LTD40MGN75718 002

Apr 16, 2001

AB GENPHARM20MGN75457 001

Apr 18, 2001

AB GENPHARM40MGN75457 002

Apr 18, 2001

AB IVAX PHARMS20MGN75511 001

Apr 16, 2001

AB IVAX PHARMS40MGN75511 002

Apr 16, 2001

AB MUTUAL PHARM20MGN75639 002

Dec 12, 2001

AB MUTUAL PHARM40MGN75639 001

Dec 12, 2001

AB MYLAN20MGN75704 001

Apr 16, 2001

AB MYLAN40MGN75704 002

Apr 16, 2001

AB PERRIGO20MGN77352 002

Jul 27, 2005

AB PERRIGO40MGN77352 001

Jul 27, 2005

AB PUREPAC PHARM20MGN75650 001

Sep 14, 2001

AB PUREPAC PHARM40MGN75650 002

Sep 14, 2001

AB SANDOZ20MGN75302 001

Apr 16, 2001

AB SANDOZ20MGN75607 001

May 10, 2001

AB SANDOZ20MGN75793 001

Apr 16, 2001

AB SANDOZ40MGN75302 002

Apr 16, 2001

AB SANDOZ40MGN75607 002

May 10, 2001

AB SANDOZ40MGN75793 002

Apr 16, 2001

AB TEVA20MGN75311 001

Apr 16, 2001

AB TEVA40MGN75311 002

Apr 16, 2001

AB TORPHARM20MGN75611 001

Jul 23, 2001

AB TORPHARM40MGN75611 002

Jul 23, 2001

AB WATSON LABS20MGN75062 002

Apr 16, 2001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 157 (of 360)

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

<u>AB</u>	WATSON LABS	<u>40MG</u>	<u>N75062</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>	WOCKHARDT	<u>20MG</u>	<u>N75786</u>	<u>001</u>	Apr 16, 2001
<u>AB</u>		<u>40MG</u>	<u>N75786</u>	<u>002</u>	Apr 16, 2001

PEPCID

<u>AB</u>	MERCK	<u>20MG</u>	<u>N19462</u>	<u>001</u>	Oct 15, 1986
<u>AB</u>	+	<u>40MG</u>	<u>N19462</u>	<u>002</u>	Oct 15, 1986

TABLET, ORALLY DISINTEGRATING; ORAL

FLUXID

	SCHWARZ PHARMA	20MG	N21712	001	Sep 24, 2004
+		40MG	N21712	002	Sep 24, 2004

FELBAMATE

SUSPENSION; ORAL

FELBATOL

+	MEDPOINTE	600MG/5ML	N20189	003	Jul 29, 1993
---	-----------	-----------	--------	-----	--------------

TABLET; ORAL

FELBATOL

	MEDPOINTE	400MG	N20189	001	Jul 29, 1993
+		600MG	N20189	002	Jul 29, 1993

FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

FELODIPINE

<u>AB</u>	MUTUAL PHARM	<u>2.5MG</u>	<u>N75896</u>	<u>001</u>	Nov 02, 2004
<u>AB</u>		<u>5MG</u>	<u>N75896</u>	<u>002</u>	Nov 02, 2004
<u>AB</u>		<u>10MG</u>	<u>N75896</u>	<u>003</u>	Nov 02, 2004

PLENDIL

<u>AB</u>	ASTRAZENECA	<u>2.5MG</u>	<u>N19834</u>	<u>004</u>	Sep 22, 1994
<u>AB</u>		<u>5MG</u>	<u>N19834</u>	<u>001</u>	Jul 25, 1991
<u>AB</u>	+	<u>10MG</u>	<u>N19834</u>	<u>002</u>	Jul 25, 1991

FENOFIBRATE

CAPSULE; ORAL

ANTARA (MICRONIZED)

	RELIANT PHARMS INC	43MG	N21695	001	Nov 30, 2004
+		130MG	N21695	003	Nov 30, 2004

FENOFIBRATE (MICRONIZED)

<u>AB</u>	IMPAX LABS	<u>67MG</u>	<u>N75868</u>	<u>001</u>	Oct 27, 2003
<u>AB</u>		<u>134MG</u>	<u>N75868</u>	<u>002</u>	Oct 27, 2003
<u>AB</u>		<u>200MG</u>	<u>N75868</u>	<u>003</u>	Oct 27, 2003
<u>AB</u>	TEVA	<u>67MG</u>	<u>N75753</u>	<u>001</u>	Sep 03, 2002
<u>AB</u>		<u>134MG</u>	<u>N75753</u>	<u>002</u>	Apr 09, 2002
<u>AB</u>	+	<u>200MG</u>	<u>N75753</u>	<u>003</u>	Apr 09, 2002

TABLET; ORAL

FENOFIBRATE

<u>AB</u>	PAR PHARM	<u>107MG</u>	<u>N76520</u>	<u>002</u>	Dec 29, 2005
<u>AB</u>	RANBAXY	<u>54MG</u>	<u>N76635</u>	<u>001</u>	Oct 31, 2005
<u>AB</u>		<u>107MG</u>	<u>N76635</u>	<u>002</u>	Oct 31, 2005
<u>AB</u>		<u>160MG</u>	<u>N76635</u>	<u>003</u>	Oct 31, 2005
<u>AB</u>	TEVA	<u>54MG</u>	<u>N76433</u>	<u>001</u>	May 13, 2005
<u>AB</u>		<u>160MG</u>	<u>N76433</u>	<u>002</u>	May 13, 2005

TRICOR

<u>AB</u>	ABBOTT	<u>54MG</u>	<u>N21203</u>	<u>001</u>	Sep 04, 2001
<u>AB</u>	+	<u>160MG</u>	<u>N21203</u>	<u>003</u>	Sep 04, 2001
		48MG	N21656	001	Nov 05, 2004
+		145MG	N21656	002	Nov 05, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 158 (of 360)

FENOFIBRATE

TABLET; ORAL					
TRIGLIDE					
BX	SKYEPHARMA	160MG	N21350 002	May 07, 2005	
		50MG	N21350 001	May 07, 2005	

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION					
<u>CORLOPAM</u>					
AP	+ HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>N19922 001</u>	Sep 23, 1997	
<u>FENOLDOPAM MESYLATE</u>					
AP	BEDFORD LABS	<u>EQ 10MG BASE/ML</u>	<u>N76582 001</u>	Oct 12, 2004	
AP	PHARMAFORCE	<u>EQ 10MG BASE/ML</u>	<u>N76656 001</u>	Dec 01, 2003	
AP	SABEX 2002	<u>EQ 10MG BASE/ML</u>	<u>N77155 001</u>	Feb 15, 2005	

FENOPROFEN CALCIUM

CAPSULE; ORAL					
<u>FENOPROFEN CALCIUM</u>					
AB	PAR PHARM	<u>EQ 200MG BASE</u>	<u>N72437 001</u>	Aug 22, 1988	
AB		<u>EQ 300MG BASE</u>	<u>N72438 001</u>	Aug 22, 1988	
AB	SANDOZ	<u>EQ 200MG BASE</u>	<u>N72394 001</u>	Oct 17, 1988	
AB		<u>EQ 300MG BASE</u>	<u>N72395 001</u>	Oct 17, 1988	
<u>NALFON</u>					
AB	+ PEDINOL	<u>EQ 300MG BASE</u>	<u>N17604 002</u>		
<u>NALFON 200</u>					
AB	PEDINOL	<u>EQ 200MG BASE</u>	<u>N17604 003</u>		
TABLET; ORAL					
<u>FENOPROFEN CALCIUM</u>					
AB	CLONMEL HLTHCARE	<u>EQ 600MG BASE</u>	<u>N72326 001</u>	Aug 17, 1988	
AB	IVAX PHARMS	<u>EQ 600MG BASE</u>	<u>N72557 001</u>	Aug 29, 1988	
AB	MUTUAL PHARM	<u>EQ 600MG BASE</u>	<u>N72902 001</u>	Dec 21, 1990	
AB	+ MYLAN	<u>EQ 600MG BASE</u>	<u>N72267 001</u>	Aug 17, 1988	
AB	PAR PHARM	<u>EQ 600MG BASE</u>	<u>N72429 001</u>	Aug 17, 1988	
AB	PUREPAC PHARM	<u>EQ 600MG BASE</u>	<u>N72274 001</u>	May 02, 1988	
AB	SANDOZ	<u>EQ 600MG BASE</u>	<u>N72396 001</u>	Oct 17, 1988	
AB	WATSON LABS	<u>EQ 600MG BASE</u>	<u>N72407 001</u>	Aug 17, 1988	
AB		<u>EQ 600MG BASE</u>	<u>N72602 001</u>	Oct 11, 1988	

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL					
<u>DURAGESIC-100</u>					
AB	ALZA	<u>100UGM/HR</u>	<u>N19813 001</u>	Aug 07, 1990	
	DURAGESIC-12				
	ALZA	12.5UGM/HR	N19813 005	Feb 04, 2005	
<u>DURAGESIC-25</u>					
AB	+ ALZA	<u>25UGM/HR</u>	<u>N19813 004</u>	Aug 07, 1990	
<u>DURAGESIC-50</u>					
AB	ALZA	<u>50UGM/HR</u>	<u>N19813 003</u>	Aug 07, 1990	
<u>DURAGESIC-75</u>					
AB	ALZA	<u>75UGM/HR</u>	<u>N19813 002</u>	Aug 07, 1990	
<u>FENTANYL</u>					
AB	MYLAN TECHNOLOGIES	<u>25UGM/HR</u>	<u>N76258 001</u>	Jan 28, 2005	
AB		<u>50UGM/HR</u>	<u>N76258 002</u>	Jan 28, 2005	
AB		<u>75UGM/HR</u>	<u>N76258 003</u>	Jan 28, 2005	
AB		<u>100UGM/HR</u>	<u>N76258 004</u>	Jan 28, 2005	

FENTANYL CITRATE

INJECTABLE; INJECTION					
<u>FENTANYL CITRATE</u>					
AP	HOSPIRA	<u>EQ 0.05MG BASE/ML</u>	<u>N19115 001</u>	Jan 12, 1985	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 159 (of 360)

FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE PRESERVATIVE FREE

<u>AP</u>	+	BAXTER HLTHCARE	<u>EQ 0.05MG BASE/ML</u>	<u>N19101</u>	<u>001</u>	Jul 11, 1984
-----------	---	-----------------	--------------------------	---------------	------------	--------------

<u>AP</u>		HOSPIRA	<u>EQ 0.05MG BASE/ML</u>	<u>N72786</u>	<u>001</u>	Sep 24, 1991
-----------	--	---------	--------------------------	---------------	------------	--------------

SUBLIMAZE PRESERVATIVE FREE

<u>AP</u>	+	AKORN MFG	<u>EQ 0.05MG BASE/ML</u>	<u>N16619</u>	<u>001</u>	
-----------	---	-----------	--------------------------	---------------	------------	--

TROCHE/LOZENGE; TRANSMUCOSAL

ACTIQ

CEPHALON

EQ 0.2MG BASE

N20747 001

Nov 04, 1998

+

EQ 0.4MG BASE

N20747 002

Nov 04, 1998

EQ 0.6MG BASE

N20747 003

Nov 04, 1998

EQ 0.8MG BASE

N20747 004

Nov 04, 1998

EQ 1.2MG BASE

N20747 005

Nov 04, 1998

EQ 1.6MG BASE

N20747 006

Nov 04, 1998

FERRIC HEXACYANOFERRATE(II)

CAPSULE; ORAL

RADIOGARDASE (PRUSSIAN BLUE)

+ HEYL CHEMISCH 500MG

N21626 001

Oct 02, 2003

FERUMOXIDES

INJECTABLE; INJECTION

FERIDEX I.V.

+ ADV MAGNETICS EQ 11.2MG IRON/ML

N20416 001

Aug 30, 1996

FERUMOXSiL

SUSPENSION; ORAL

GASTROMARK

+ ADV MAGNETICS EQ 0.175MG IRON/ML

N20410 001

Dec 06, 1996

FEXOFENADINE HYDROCHLORIDE

CAPSULE; ORAL

ALLEGRA

<u>AB</u>	+	AVENTIS PHARMS	<u>60MG</u>	<u>N20625</u>	<u>001</u>	Jul 25, 1996
-----------	---	----------------	-------------	---------------	------------	--------------

FEXOFENADINE HYDROCHLORIDE

<u>AB</u>		BARR	<u>60MG</u>	<u>N76169</u>	<u>001</u>	Jul 13, 2005
-----------	--	------	-------------	---------------	------------	--------------

TABLET; ORAL

ALLEGRA

<u>AB</u>		AVENTIS PHARMS	<u>30MG</u>	<u>N20872</u>	<u>001</u>	Feb 25, 2000
-----------	--	----------------	-------------	---------------	------------	--------------

<u>AB</u>			<u>60MG</u>	<u>N20872</u>	<u>002</u>	Feb 25, 2000
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>	+		<u>180MG</u>	<u>N20872</u>	<u>004</u>	Feb 25, 2000
-----------	---	--	--------------	---------------	------------	--------------

FEXOFENADINE HYDROCHLORIDE

<u>AB</u>		BARR	<u>30MG</u>	<u>N76191</u>	<u>001</u>	Aug 31, 2005
-----------	--	------	-------------	---------------	------------	--------------

<u>AB</u>			<u>60MG</u>	<u>N76191</u>	<u>002</u>	Aug 31, 2005
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>180MG</u>	<u>N76191</u>	<u>003</u>	Aug 31, 2005
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		TEVA	<u>30MG</u>	<u>N76447</u>	<u>001</u>	Sep 01, 2005
-----------	--	------	-------------	---------------	------------	--------------

<u>AB</u>			<u>60MG</u>	<u>N76447</u>	<u>002</u>	Sep 01, 2005
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>180MG</u>	<u>N76447</u>	<u>003</u>	Sep 01, 2005
-----------	--	--	--------------	---------------	------------	--------------

FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALLEGRA D 24 HOUR

+ AVENTIS PHARMS 180MG;240MG

N21704 001

Oct 19, 2004

ALLEGRA-D 12 HOUR

<u>AB</u>	+	AVENTIS PHARMS	<u>60MG;120MG</u>	<u>N20786</u>	<u>001</u>	Dec 24, 1997
-----------	---	----------------	-------------------	---------------	------------	--------------

FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE

<u>AB</u>		BARR	<u>60MG;120MG</u>	<u>N76236</u>	<u>001</u>	Apr 14, 2005
-----------	--	------	-------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 160 (of 360)

FINASTERIDE

TABLET; ORAL

PROPECIA

+ MERCK 1MG N20788 001 Dec 19, 1997

PROSCAR

+ MERCK 5MG N20180 001 Jun 19, 1992

FLAVOXATE HYDROCHLORIDE

TABLET; ORAL

FLAVOXATE HYDROCHLORIDEAB HAMPTON LAINE LLC 100MG N76835 001 Nov 30, 2005AB IMPAX PHARMS 100MG N76234 001 Aug 28, 2003AB PADDOCK 100MG N76831 001 Dec 16, 2004URISPASAB + ORTHO MCNEIL PHARM 100MG N16769 001FLECAINIDE ACETATE

TABLET; ORAL

FLECAINIDE ACETATEAB ALPHAPHARM 50MG N75442 001 Jul 31, 2001AB 100MG N75442 002 Jul 31, 2001AB 150MG N75442 003 Jul 31, 2001AB BARR 50MG N75882 001 Oct 28, 2002AB 100MG N75882 002 Oct 28, 2002AB 150MG N75882 003 Oct 28, 2002AB RANBAXY 50MG N76421 001 Mar 28, 2003AB 100MG N76421 002 Mar 28, 2003AB 150MG N76421 003 Mar 28, 2003AB ROXANE 50MG N76278 001 Jan 14, 2003AB 100MG N76278 002 Jan 14, 2003AB 150MG N76278 003 Jan 14, 2003AB SANDOZ 50MG N76030 001 Oct 28, 2002AB 100MG N76030 002 Oct 28, 2002AB 150MG N76030 003 Oct 28, 2002TAMBOCORAB 3M 50MG N18830 004 Aug 23, 1988AB 100MG N18830 001 Oct 31, 1985AB + 150MG N18830 003 Jun 03, 1988FLOXURIDINE

INJECTABLE; INJECTION

FLOXURIDINEAP AM PHARM PARTNERS 500MG/VIAL N75837 001 Feb 22, 2001AP BEDFORD 500MG/VIAL N75387 001 Apr 16, 2000FUDRAP + MAYNE PHARMA USA 500MG/VIAL N16929 001FLUCONAZOLE

FOR SUSPENSION; ORAL

DIFLUCANAB PFIZER 50MG/5ML N20090 001 Dec 23, 1993AB + 200MG/5ML N20090 002 Dec 23, 1993FLUCONAZOLEAB RANBAXY 50MG/5ML N76332 001 Jul 29, 2004AB 200MG/5ML N76332 002 Jul 29, 2004AB ROXANE 50MG/5ML N76246 001 Jul 29, 2004AB 200MG/5ML N76246 002 Jul 29, 2004

INJECTABLE; INJECTION

DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINERAP + PFIZER 200MG/100ML N19950 003 Sep 29, 1992

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 161 (of 360)

FLUCONAZOLE

INJECTABLE; INJECTION

<u>DIFLUCAN IN SODIUM CHLORIDE 0.9%</u>			
<u>AP</u>	+ PFIZER	<u>200MG/100ML</u>	<u>N19950</u> <u>001</u> Jan 29, 1990
<u>DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>	+ PFIZER	<u>200MG/100ML</u>	<u>N19950</u> <u>002</u> Jan 29, 1990
<u>FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>			
<u>AP</u>	APOTEX	<u>200MG/100ML</u>	<u>N76888</u> <u>001</u> Mar 25, 2005
<u>AP</u>	HOSPIRA	<u>200MG/100ML</u>	<u>N76304</u> <u>001</u> Jul 29, 2004
<u>FLUCONAZOLE IN SODIUM CHLORIDE 0.9%</u>			
<u>AP</u>	AM PHARM PARTNERS	<u>200MG/100ML</u>	<u>N76145</u> <u>001</u> Jul 29, 2004
<u>AP</u>	BEDFORD	<u>200MG/100ML</u>	<u>N76087</u> <u>001</u> Jul 29, 2004
<u>AP</u>	HIKMA FARMACEUTICA	<u>200MG/100ML</u>	<u>N76736</u> <u>001</u> Aug 23, 2005
<u>AP</u>	SICOR PHARMS	<u>200MG/100ML</u>	<u>N76653</u> <u>001</u> Jul 29, 2004
<u>FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
<u>AP</u>	APOTEX	<u>200MG/100ML</u>	<u>N76889</u> <u>001</u> Mar 25, 2005
<u>AP</u>	BAXTER HLTHCARE	<u>200MG/100ML</u>	<u>N76766</u> <u>001</u> Jul 29, 2004
<u>AP</u>	HOSPIRA	<u>200MG/100ML</u>	<u>N76303</u> <u>001</u> Jul 29, 2004
<u>AP</u>	MAYNE PHARMA USA	<u>200MG/100ML</u>	<u>N76617</u> <u>001</u> Jul 29, 2004
<u>AP</u>	SICOR PHARMS	<u>200MG/100ML</u>	<u>N76837</u> <u>001</u> Jan 13, 2005

TABLET; ORAL

<u>DIFLUCAN</u>			
<u>AB</u>	PFIZER	<u>50MG</u>	<u>N19949</u> <u>001</u> Jan 29, 1990
<u>AB</u>		<u>100MG</u>	<u>N19949</u> <u>002</u> Jan 29, 1990
<u>AB</u>		<u>150MG</u>	<u>N19949</u> <u>004</u> Jun 30, 1994
<u>AB</u>	+	<u>200MG</u>	<u>N19949</u> <u>003</u> Jan 29, 1990
<u>FLUCONAZOLE</u>			
<u>AB</u>	DR REDDYS LABS INC	<u>50MG</u>	<u>N76658</u> <u>001</u> Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76658</u> <u>002</u> Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76658</u> <u>003</u> Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76658</u> <u>004</u> Jul 29, 2004
<u>AB</u>	GEDEON RICHTER USA	<u>50MG</u>	<u>N76432</u> <u>001</u> Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76432</u> <u>002</u> Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76432</u> <u>003</u> Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76432</u> <u>004</u> Jul 29, 2004
<u>AB</u>	GENPHARM	<u>50MG</u>	<u>N76042</u> <u>001</u> Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76042</u> <u>002</u> Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76042</u> <u>003</u> Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76042</u> <u>004</u> Jul 29, 2004
<u>AB</u>	IVAX PHARMS	<u>50MG</u>	<u>N76077</u> <u>001</u> Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76077</u> <u>002</u> Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76077</u> <u>003</u> Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76077</u> <u>004</u> Jul 29, 2004
<u>AB</u>	MYLAN	<u>50MG</u>	<u>N76351</u> <u>001</u> Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76351</u> <u>002</u> Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76351</u> <u>003</u> Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76351</u> <u>004</u> Jul 29, 2004
<u>AB</u>	PLIVA	<u>50MG</u>	<u>N76424</u> <u>001</u> Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76424</u> <u>002</u> Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76424</u> <u>003</u> Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76424</u> <u>004</u> Jul 29, 2004
<u>AB</u>	RANBAXY	<u>50MG</u>	<u>N76386</u> <u>001</u> Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76386</u> <u>002</u> Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76386</u> <u>003</u> Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76386</u> <u>004</u> Jul 29, 2004
<u>AB</u>	ROXANE	<u>50MG</u>	<u>N76213</u> <u>001</u> Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76213</u> <u>002</u> Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76213</u> <u>003</u> Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76213</u> <u>004</u> Jul 29, 2004
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N76086</u> <u>001</u> Jul 29, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 162 (of 360)

FLUCONAZOLE

TABLET; ORAL

FLUCONAZOLE

<u>AB</u>	SANDOZ	<u>100MG</u>	<u>N76086</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76086</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76086</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>	TARO	<u>50MG</u>	<u>N76507</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76507</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76507</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76507</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>	TEVA	<u>50MG</u>	<u>N74681</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N74681</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N74681</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N74681</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>	TORPHARM	<u>50MG</u>	<u>N76665</u>	<u>001</u>	Jul 29, 2004
<u>AB</u>		<u>100MG</u>	<u>N76665</u>	<u>002</u>	Jul 29, 2004
<u>AB</u>		<u>150MG</u>	<u>N76665</u>	<u>003</u>	Jul 29, 2004
<u>AB</u>		<u>200MG</u>	<u>N76665</u>	<u>004</u>	Jul 29, 2004
<u>AB</u>	UNIQUE PHARM LABS	<u>50MG</u>	<u>N76957</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>		<u>100MG</u>	<u>N76957</u>	<u>002</u>	Sep 28, 2005
<u>AB</u>		<u>200MG</u>	<u>N76957</u>	<u>003</u>	Sep 28, 2005

FLUCYTOSINE

CAPSULE; ORAL

ANCOBON

VALEANT

+

250MG

N17001 001

500MG

N17001 002

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARA

<u>AP</u>	+ BERLEX	<u>50MG/VIAL</u>	<u>N20038</u>	<u>001</u>	Apr 18, 1991
<u>AP</u>	<u>FLUDARABINE PHOSPHATE</u>				
<u>AP</u>	SICOR PHARMS	<u>50MG/VIAL</u>	<u>N76349</u>	<u>001</u>	Aug 28, 2003
	+	50MG/2ML (25MG/ML)	N76661	001	Apr 28, 2004

FLUDEOXYGLUCOSE F-18

INJECTABLE; INTRAVENOUS

FLUDEOXYGLUCOSE F 18

+ WEILL MEDCL COLL

10-100mCi/ML

N21768 001

Aug 05, 2004

FLUDEOXYGLUCOSE F18

+ NORTH SHORE LIJ

20-200mCi/ML

N21870 001

Aug 19, 2005

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLORINEF

<u>AB</u>	+ KING PHARMS	<u>0.1MG</u>	<u>N10060</u>	<u>001</u>	
<u>AB</u>	<u>FLUDROCORTISONE ACETATE</u>				
<u>AB</u>	BARR	<u>0.1MG</u>	<u>N40425</u>	<u>001</u>	Jan 21, 2003
<u>AB</u>	IMPAX LABS	<u>0.1MG</u>	<u>N40431</u>	<u>001</u>	Mar 18, 2002

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

<u>AP</u>	AM PHARM	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N76955</u>	<u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>N76955</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>	APOTEX	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N76755</u>	<u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>N76755</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>	BAXTER HLTHCARE	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N76787</u>	<u>002</u>	Oct 12, 2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 163 (of 360)

FLUMAZENIL

INJECTABLE; INJECTION

FLUMAZENIL

<u>AP</u>	BAXTER HLTHCARE	<u>1MG/10ML (0.1MG/ML)</u>	<u>N76787</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>	BEDFORD LABS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N76256</u>	<u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>N76256</u>	<u>001</u>	Oct 12, 2004
<u>AP</u>	SABEX 2002	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N77071</u>	<u>001</u>	May 03, 2005
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>N77071</u>	<u>002</u>	May 03, 2005
<u>AP</u>	SICOR PHARMS	<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N76589</u>	<u>002</u>	Oct 12, 2004
<u>AP</u>		<u>1MG/10ML (0.1MG/ML)</u>	<u>N76589</u>	<u>001</u>	Oct 12, 2004
<u>ROMAZICON</u>					
<u>AP</u>	+ HLR	<u>1MG/10ML (0.1MG/ML)</u>	<u>N20073</u>	<u>001</u>	Dec 20, 1991
<u>AP</u>		<u>0.5MG/5ML (0.1MG/ML)</u>	<u>N20073</u>	<u>002</u>	Dec 20, 1991

FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROBID

+ ROCHE PALO	0.25MG/INH	N18340	001	Aug 17, 1984
--------------	------------	--------	-----	--------------

SPRAY, METERED; NASAL

FLUNISOLIDE

+ BAUSCH AND LOMB	0.025MG/SPRAY	N74805	001	Feb 20, 2002
-------------------	---------------	--------	-----	--------------

NASAREL

+ IVAX RES	0.029MG/SPRAY	N20409	001	Mar 08, 1995
------------	---------------	--------	-----	--------------

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	FOUGERA	<u>0.01%</u>	<u>N88170</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>		<u>0.025%</u>	<u>N88169</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>	G AND W LABS	<u>0.01%</u>	<u>N89526</u>	<u>001</u>	Jul 26, 1988
<u>AT</u>		<u>0.025%</u>	<u>N89525</u>	<u>001</u>	Jul 26, 1988
<u>AT</u>	TARO	<u>0.01%</u>	<u>N87102</u>	<u>001</u>	Apr 27, 1982
<u>AT</u>		<u>0.025%</u>	<u>N87104</u>	<u>001</u>	Apr 27, 1982
<u>SYNALAR</u>					
<u>AT</u>	+ MEDICIS	<u>0.01%</u>	<u>N12787</u>	<u>004</u>	
<u>AT</u>	+	<u>0.025%</u>	<u>N12787</u>	<u>005</u>	
<u>AT</u>	+	<u>0.025%</u>	<u>N12787</u>	<u>002</u>	

IMPLANT; INTRAVITREAL

RETISERT

+ BAUSCH AND LOMB	0.59MG	N21737	001	Apr 08, 2005
-------------------	--------	--------	-----	--------------

OIL; TOPICAL

DERMA-SMOOTH/FS

+ HILL DERMAC	0.01%	N19452	001	Feb 03, 1988
---------------	-------	--------	-----	--------------

OIL/DROPS; OTIC

FLUOCINOLONE ACETONIDE

+ HILL DERMAC	0.01%	N21930	001	Nov 09, 2005
---------------	-------	--------	-----	--------------

OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	FOUGERA	<u>0.025%</u>	<u>N88168</u>	<u>001</u>	Dec 16, 1982
<u>AT</u>	G AND W LABS	<u>0.025%</u>	<u>N89524</u>	<u>001</u>	Jul 26, 1988
<u>AT</u>	TARO	<u>0.025%</u>	<u>N40041</u>	<u>001</u>	Sep 15, 1994
<u>SYNALAR</u>					
<u>AT</u>	+ MEDICIS	<u>0.025%</u>	<u>N13960</u>	<u>001</u>	

SHAMPOO; TOPICAL

FS SHAMPOO

+ GALDERMA LABS LP	0.01%	N20001	001	Aug 27, 1990
--------------------	-------	--------	-----	--------------

SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	FOUGERA	<u>0.01%</u>	<u>N88167</u>	<u>001</u>	Dec 16, 1982
-----------	---------	--------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 164 (of 360)

FLUOCINOLONE ACETONIDE

SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

<u>AT</u>	TARO	<u>0.01%</u>	<u>N89124</u>	<u>001</u>	Sep 11, 1985
	<u>SYNALAR</u>				
<u>AT</u>	+ MEDICIS	<u>0.01%</u>	<u>N15296</u>	<u>001</u>	

FLUOCINOLONE ACETONIDE; HYDROQUINONE; TRETINOIN

CREAM; TOPICAL

TRI-LUMA

	+ GALDERMA LABS LP	0.01%;4%;0.05%	N21112	001	Jan 18, 2002
--	--------------------	----------------	--------	-----	--------------

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

<u>AB1</u>	ALPHARMA US PHARMS	<u>0.05%</u>	<u>N73085</u>	<u>001</u>	Feb 14, 1992
<u>AB1</u>	FOUGERA	<u>0.05%</u>	<u>N73030</u>	<u>001</u>	Oct 17, 1994
<u>AB1</u>	TARO	<u>0.05%</u>	<u>N19117</u>	<u>001</u>	Jun 26, 1984
<u>AB1</u>		<u>0.05%</u>	<u>N71500</u>	<u>001</u>	Jun 10, 1987
<u>AB1</u>	TEVA	<u>0.05%</u>	<u>N72488</u>	<u>001</u>	Feb 06, 1989

FLUOCINONIDE EMULSIFIED BASE

<u>AB2</u>	ALPHARMA US PHARMS	<u>0.05%</u>	<u>N74204</u>	<u>001</u>	Jun 13, 1995
<u>AB2</u>	ALTANA	<u>0.05%</u>	<u>N76586</u>	<u>001</u>	Jun 23, 2004
<u>AB2</u>	TARO	<u>0.05%</u>	<u>N72494</u>	<u>001</u>	Jan 19, 1989
<u>AB2</u>	TEVA	<u>0.05%</u>	<u>N72490</u>	<u>001</u>	Feb 07, 1989

LIDEX

<u>AB1</u>	+ MEDICIS	<u>0.05%</u>	<u>N16908</u>	<u>002</u>	
------------	-----------	--------------	---------------	------------	--

LIDEX-E

<u>AB2</u>	+ MEDICIS	<u>0.05%</u>	<u>N16908</u>	<u>003</u>	
------------	-----------	--------------	---------------	------------	--

VANOS

	+ MEDICIS	0.1%	N21758	001	Feb 11, 2005
--	-----------	------	--------	-----	--------------

GEL; TOPICAL

FLUOCINONIDE

<u>AB</u>	FOUGERA	<u>0.05%</u>	<u>N72933</u>	<u>001</u>	Dec 30, 1994
<u>AB</u>	TARO	<u>0.05%</u>	<u>N74935</u>	<u>001</u>	Jul 29, 1997
<u>AB</u>	TEVA	<u>0.05%</u>	<u>N72537</u>	<u>001</u>	Feb 07, 1989

LIDEX

<u>AB</u>	+ MEDICIS	<u>0.05%</u>	<u>N17373</u>	<u>001</u>	
-----------	-----------	--------------	---------------	------------	--

OINTMENT; TOPICAL

FLUOCINONIDE

<u>AB</u>	ALTANA	<u>0.05%</u>	<u>N74905</u>	<u>001</u>	Aug 26, 1997
<u>AB</u>	TARO	<u>0.05%</u>	<u>N75008</u>	<u>001</u>	Jun 30, 1999
<u>AB</u>	TEVA	<u>0.05%</u>	<u>N73481</u>	<u>001</u>	Dec 27, 1991

LIDEX

<u>AB</u>	+ MEDICIS	<u>0.05%</u>	<u>N16909</u>	<u>002</u>	
-----------	-----------	--------------	---------------	------------	--

SOLUTION; TOPICAL

FLUOCINONIDE

<u>AT</u>	ALPHARMA US PHARMS	<u>0.05%</u>	<u>N71535</u>	<u>001</u>	Dec 02, 1988
<u>AT</u>	FOUGERA	<u>0.05%</u>	<u>N72934</u>	<u>001</u>	Feb 27, 1995
<u>AT</u>	TARO	<u>0.05%</u>	<u>N72857</u>	<u>001</u>	Aug 02, 1989
<u>AT</u>		<u>0.05%</u>	<u>N74799</u>	<u>001</u>	Dec 31, 1996
<u>AT</u>	TEVA	<u>0.05%</u>	<u>N72511</u>	<u>001</u>	Feb 07, 1989
<u>AT</u>	TEVA PHARMS	<u>0.05%</u>	<u>N72522</u>	<u>001</u>	Sep 28, 1990

LIDEX

<u>AT</u>	+ MEDICIS	<u>0.05%</u>	<u>N18849</u>	<u>001</u>	Apr 06, 1984
-----------	-----------	--------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 165 (of 360)

FLUOROMETHOLONE

OINTMENT; OPHTHALMIC				
FML				
	+ ALLERGAN	0.1%	N17760 001	Sep 04, 1985
SUSPENSION/DROPS; OPHTHALMIC				
<u>FLUOR-OP</u>				
<u>AB</u>	NOVARTIS	<u>0.1%</u>	<u>N70185</u> <u>001</u>	Feb 27, 1986
<u>FML</u>				
<u>AB</u>	+ ALLERGAN	<u>0.1%</u>	<u>N16851</u> <u>002</u>	Jul 28, 1982
FML FORTE				
	ALLERGAN	0.25%	N19216 001	Apr 23, 1986

FLUOROMETHOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC				
FLAREX				
	+ ALCON	0.1%	N19079 001	Feb 11, 1986

FLUOROMETHOLONE ACETATE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC				
TOBRASONE				
	+ ALCON	0.1%;0.3%	N50628 001	Jul 21, 1989

FLUOROMETHOLONE; SULFACETAMIDE SODIUM

SUSPENSION/DROPS; OPHTHALMIC				
FML-S				
	+ ALLERGAN	0.1%;10%	N19525 001	Sep 29, 1989

FLUOROURACIL

CREAM; TOPICAL				
CARAC				
	+ DERMIK LABS	0.5%	N20985 001	Oct 27, 2000
EFUDEX				
	+ VALEANT PHARM INTL	5%	N16831 003	
FLUOROPLEX				
	+ ALLERGAN HERBERT	1%	N16988 001	
INJECTABLE; INJECTION				
<u>ADRUCIL</u>				
<u>AP</u>	SICOR PHARMS	<u>50MG/ML</u>	<u>N40023</u> <u>001</u>	Oct 18, 1991
<u>AP</u>	+	<u>50MG/ML</u>	<u>N81225</u> <u>001</u>	Aug 28, 1991
<u>FLUOROURACIL</u>				
<u>AP</u>	AM PHARM	<u>50MG/ML</u>	<u>N40291</u> <u>001</u>	Mar 24, 1999
<u>AP</u>	AM PHARM PARTNERS	<u>50MG/ML</u>	<u>N40278</u> <u>001</u>	Sep 30, 1998
<u>AP</u>		<u>50MG/ML</u>	<u>N40279</u> <u>001</u>	Sep 30, 1998
<u>AP</u>		<u>50MG/ML</u>	<u>N40379</u> <u>001</u>	Nov 15, 2000
<u>AP</u>	BEDFORD	<u>50MG/ML</u>	<u>N89508</u> <u>001</u>	Jan 26, 1988
<u>AP</u>	SICOR PHARMS	<u>50MG/ML</u>	<u>N40333</u> <u>001</u>	Jan 27, 2000
<u>AP</u>		<u>50MG/ML</u>	<u>N40334</u> <u>001</u>	Feb 25, 2000
<u>AP</u>	STERIS	<u>50MG/ML</u>	<u>N87792</u> <u>001</u>	Oct 13, 1982
<u>AP</u>	+ VALEANT	<u>50MG/ML</u>	<u>N12209</u> <u>001</u>	
SOLUTION; TOPICAL				
<u>EFUDEX</u>				
<u>AT</u>	+ VALEANT PHARM INTL	<u>2%</u>	<u>N16831</u> <u>001</u>	
<u>AT</u>	+	<u>5%</u>	<u>N16831</u> <u>002</u>	
<u>FLUOROURACIL</u>				
<u>AT</u>	TARO	<u>2%</u>	<u>N76526</u> <u>001</u>	Nov 05, 2003
<u>AT</u>		<u>5%</u>	<u>N76526</u> <u>002</u>	Nov 05, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 166 (of 360)

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE

<u>AB</u>	ALPHAPHARM	<u>EQ 10MG BASE</u>	<u>N75577</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75577</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>	BARR	<u>EQ 10MG BASE</u>	<u>N74803</u>	<u>002</u>	Jan 30, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N74803</u>	<u>001</u>	Aug 02, 2001
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N76251</u>	<u>001</u>	May 18, 2005
<u>AB</u>	CARLSBAD	<u>EQ 10MG BASE</u>	<u>N76022</u>	<u>001</u>	Jan 30, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N76022</u>	<u>002</u>	Jan 30, 2002
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 10MG BASE</u>	<u>N75465</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75465</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N75465</u>	<u>003</u>	Aug 02, 2001
<u>AB</u>	IVAX PHARMS	<u>EQ 10MG BASE</u>	<u>N75245</u>	<u>002</u>	Jan 31, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75245</u>	<u>001</u>	Jan 31, 2002
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N75245</u>	<u>003</u>	Sep 28, 2004
<u>AB</u>	MALLINCKRODT	<u>EQ 10MG BASE</u>	<u>N75658</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75658</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>	MUTUAL PHARMA	<u>EQ 10MG BASE</u>	<u>N75787</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75787</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>N75207</u>	<u>001</u>	Jan 30, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75207</u>	<u>002</u>	Jan 30, 2002
<u>AB</u>	PAR PHARM	<u>EQ 10MG BASE</u>	<u>N76922</u>	<u>001</u>	Dec 16, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N76922</u>	<u>002</u>	Dec 16, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N76922</u>	<u>003</u>	Dec 16, 2004
<u>AB</u>	PLIVA	<u>EQ 10MG BASE</u>	<u>N76001</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N76001</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>	RANBAXY	<u>EQ 10MG BASE</u>	<u>N76165</u>	<u>001</u>	Feb 01, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N76165</u>	<u>002</u>	Feb 01, 2002
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N76990</u>	<u>001</u>	Dec 13, 2004
<u>AB</u>	RXELITE	<u>EQ 10MG BASE</u>	<u>N75464</u>	<u>001</u>	Jan 30, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75464</u>	<u>002</u>	Jan 30, 2002
<u>AB</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>N75049</u>	<u>001</u>	Aug 02, 2001
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N75807</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75807</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>N75452</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75452</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N75452</u>	<u>003</u>	Jan 29, 2002
<u>AB</u>	WATSON LABS	<u>EQ 10MG BASE</u>	<u>N75662</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75662</u>	<u>002</u>	Jan 29, 2002

FLUOXETINE HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>EQ 20MG BASE</u>	<u>N75049</u>	<u>002</u>	Jan 29, 2002
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N75049</u>	<u>003</u>	Jan 29, 2002

PROZAC

<u>AB</u>	LILLY	<u>EQ 10MG BASE</u>	<u>N18936</u>	<u>006</u>	Dec 23, 1992
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N18936</u>	<u>001</u>	Dec 29, 1987
<u>AB</u>	+	<u>EQ 40MG BASE</u>	<u>N18936</u>	<u>003</u>	Jun 15, 1999
	SARAFEM				
	LILLY	EQ 10MG BASE	N18936	007	Jul 06, 2000
	+	EQ 20MG BASE	N18936	008	Jul 06, 2000

CAPSULE, DELAYED REL PELLETS; ORAL

PROZAC WEEKLY

	+	LILLY	EQ 90MG BASE	N21235	001	Feb 26, 2001
--	---	-------	--------------	--------	-----	--------------

SOLUTION; ORAL

FLUOXETINE

<u>AA</u>	ALPHARMA US PHARMS	<u>EQ 20MG BASE/5ML</u>	<u>N75690</u>	<u>001</u>	Jan 31, 2002
<u>AA</u>	MALLINCKRODT	<u>EQ 20MG BASE/5ML</u>	<u>N75920</u>	<u>001</u>	Jan 29, 2002
<u>AA</u>	NOVEX	<u>EQ 20MG BASE/5ML</u>	<u>N75292</u>	<u>001</u>	Feb 07, 2002
<u>AA</u>	PHARM ASSOC	<u>EQ 20MG BASE/5ML</u>	<u>N76015</u>	<u>001</u>	Jan 30, 2002
<u>AA</u>	TEVA	<u>EQ 20MG BASE/5ML</u>	<u>N75506</u>	<u>001</u>	Aug 02, 2001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 167 (of 360)

FLUOXETINE HYDROCHLORIDE

SOLUTION; ORAL

FLUOXETINE HYDROCHLORIDE

<u>AA</u>	HI TECH PHARMA	<u>EQ 20MG BASE/5ML</u>	<u>N75525</u>	<u>001</u>	Jun 27, 2002
<u>AA</u>	MORTON GROVE	<u>EQ 20MG BASE/5ML</u>	<u>N75514</u>	<u>001</u>	Aug 29, 2002
<u>AA</u>	PAR PHARM	<u>EQ 20MG BASE/5ML</u>	<u>N76458</u>	<u>001</u>	May 14, 2004
<u>PROZAC</u>					
<u>AA</u>	+ LILLY	<u>EQ 20MG BASE/5ML</u>	<u>N20101</u>	<u>001</u>	Apr 24, 1991

TABLET; ORAL

FLUOXETINE

<u>AB</u>	IVAX PHARMS	<u>EQ 40MG BASE</u>	<u>N75865</u>	<u>003</u>	Aug 30, 2004
<u>FLUOXETINE HYDROCHLORIDE</u>					
<u>AB</u>	ALPHAPHARM	<u>EQ 10MG BASE</u>	<u>N75755</u>	<u>001</u>	Aug 02, 2001
	+	EQ 20MG BASE	N75755	002	Aug 02, 2001
<u>AB</u>	BARR	<u>EQ 10MG BASE</u>	<u>N75810</u>	<u>001</u>	Feb 01, 2002
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 10MG BASE</u>	<u>N76006</u>	<u>001</u>	Jan 30, 2002
<u>AB</u>	IVAX PHARMS	<u>EQ 10MG BASE</u>	<u>N75865</u>	<u>001</u>	Feb 28, 2002
<u>AB</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>N76024</u>	<u>001</u>	Jan 29, 2002
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>N75872</u>	<u>001</u>	Jan 29, 2002
<u>PROZAC</u>					
<u>AB</u>	+ LILLY	<u>EQ 10MG BASE</u>	<u>N20974</u>	<u>001</u>	Mar 09, 1999

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

SYMBYAX

LILLY

		EQ 25MG BASE;EQ 6MG BASE	N21520	002	Dec 24, 2003
		EQ 25MG BASE;EQ 12MG BASE	N21520	004	Dec 24, 2003
	+	EQ 50MG BASE;EQ 6MG BASE	N21520	003	Dec 24, 2003
		EQ 50MG BASE;EQ 12MG BASE	N21520	005	Dec 24, 2003

FLUOXYMESTERONE

TABLET; ORAL

FLUOXYMESTERONE

BP	USL PHARMA	10MG	N88342	001	Oct 21, 1983
HALOTESTIN					
BP	+ PHARMACIA AND UPJOHN	10MG	N10611	010	
		2MG	N10611	002	
		5MG	N10611	006	

FLUPHENAZINE DECANOATE

INJECTABLE; IM-SC

FLUPHENAZINE DECANOATE

<u>AO</u>	APOTEX	<u>25MG/ML</u>	<u>N75918</u>	<u>001</u>	Aug 17, 2001
INJECTABLE; INJECTION					
<u>FLUPHENAZINE DECANOATE</u>					
<u>AO</u>	AM PHARM PARTNERS	<u>25MG/ML</u>	<u>N71413</u>	<u>001</u>	Jul 14, 1987
<u>AO</u>	BEDFORD	<u>25MG/ML</u>	<u>N74531</u>	<u>001</u>	Aug 30, 1996
<u>AO</u>	SICOR PHARMS	<u>25MG/ML</u>	<u>N74795</u>	<u>001</u>	Sep 10, 1996
<u>PROLIXIN DECANOATE</u>					
<u>AO</u>	+ BRISTOL MYERS SQUIBB	<u>25MG/ML</u>	<u>N16727</u>	<u>001</u>	

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AA</u>	PHARM ASSOC	<u>5MG/ML</u>	<u>N74725</u>	<u>001</u>	Sep 16, 1996
<u>AA</u>	TEVA PHARMS	<u>5MG/ML</u>	<u>N73058</u>	<u>001</u>	Aug 30, 1991
<u>PROLIXIN</u>					
<u>AA</u>	APOTHECON	<u>5MG/ML</u>	<u>N70533</u>	<u>001</u>	Nov 07, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 168 (of 360)

FLUPHENAZINE HYDROCHLORIDE

ELIXIR; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AA</u>	PHARM ASSOC	<u>2.5MG/5ML</u>	<u>N40146</u>	<u>001</u>	Aug 21, 1996
<u>AA</u>	TEVA PHARMS	<u>2.5MG/5ML</u>	<u>N81310</u>	<u>001</u>	Apr 29, 1993

PROLIXIN

<u>AA</u>	+ APOTHECON	<u>2.5MG/5ML</u>	<u>N12145</u>	<u>003</u>	
-----------	-------------	------------------	---------------	------------	--

INJECTABLE; INJECTION

FLUPHENAZINE HYDROCHLORIDE

<u>AP</u>	AM PHARM PARTNERS	<u>2.5MG/ML</u>	<u>N89556</u>	<u>001</u>	Apr 16, 1987
-----------	-------------------	-----------------	---------------	------------	--------------

PROLIXIN

<u>AP</u>	+ APOTHECON	<u>2.5MG/ML</u>	<u>N11751</u>	<u>005</u>	
-----------	-------------	-----------------	---------------	------------	--

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>1MG</u>	<u>N89801</u>	<u>001</u>	Aug 12, 1988
<u>AB</u>		<u>2.5MG</u>	<u>N89802</u>	<u>001</u>	Aug 12, 1988
<u>AB</u>		<u>5MG</u>	<u>N89803</u>	<u>001</u>	Aug 12, 1988
<u>AB</u>		<u>10MG</u>	<u>N89804</u>	<u>001</u>	Aug 12, 1988
<u>AB</u>	PAR PHARM	<u>1MG</u>	<u>N89740</u>	<u>001</u>	Aug 25, 1988
<u>AB</u>		<u>2.5MG</u>	<u>N89741</u>	<u>001</u>	Aug 25, 1988
<u>AB</u>		<u>5MG</u>	<u>N89742</u>	<u>001</u>	Aug 25, 1988
<u>AB</u>		<u>10MG</u>	<u>N89743</u>	<u>001</u>	Aug 25, 1988
<u>AB</u>	SANDOZ	<u>1MG</u>	<u>N89583</u>	<u>001</u>	Oct 16, 1987
<u>AB</u>		<u>2.5MG</u>	<u>N89584</u>	<u>001</u>	Oct 16, 1987
<u>AB</u>		<u>5MG</u>	<u>N89585</u>	<u>001</u>	Oct 16, 1987
<u>AB</u>		<u>10MG</u>	<u>N89586</u>	<u>001</u>	Oct 16, 1987

PROLIXIN

<u>AB</u>	APOTHECON	<u>1MG</u>	<u>N11751</u>	<u>004</u>	
<u>AB</u>		<u>2.5MG</u>	<u>N11751</u>	<u>001</u>	
<u>AB</u>		<u>5MG</u>	<u>N11751</u>	<u>003</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N11751</u>	<u>002</u>	

FLURANDRENOLIDE

CREAM; TOPICAL

CORDRAN SP

+	OCLASSEN	0.025%	N12806	003	
+		0.05%	N12806	002	

LOTION; TOPICAL

CORDRAN

+	WATSON LABS	0.05%	N13790	001	
---	-------------	-------	--------	-----	--

OINTMENT; TOPICAL

CORDRAN

+	OCLASSEN	0.025%	N12806	004	
+		0.05%	N12806	001	

TAPE; TOPICAL

CORDRAN

+	OCLASSEN	0.004MG/SQ CM	N16455	001	
---	----------	---------------	--------	-----	--

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

DALMANE

<u>AB</u>	VALEANT PHARM INTL	<u>15MG</u>	<u>N16721</u>	<u>001</u>	
<u>AB</u>	+	<u>30MG</u>	<u>N16721</u>	<u>002</u>	

FLURAZEPAM HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>15MG</u>	<u>N70344</u>	<u>001</u>	Nov 27, 1985
<u>AB</u>		<u>30MG</u>	<u>N70345</u>	<u>001</u>	Nov 27, 1985
<u>AB</u>	PAR PHARM	<u>15MG</u>	<u>N70444</u>	<u>001</u>	Mar 20, 1986
<u>AB</u>		<u>30MG</u>	<u>N70445</u>	<u>001</u>	Mar 20, 1986
<u>AB</u>	SANDOZ	<u>15MG</u>	<u>N71716</u>	<u>001</u>	Jul 31, 1991

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 169 (of 360)

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>30MG</u>	<u>N71717</u>	<u>001</u>	Jul 31, 1991
<u>AB</u>	WATSON LABS	<u>15MG</u>	<u>N71205</u>	<u>001</u>	Nov 25, 1986
<u>AB</u>		<u>30MG</u>	<u>N71068</u>	<u>001</u>	Nov 25, 1986
<u>AB</u>		<u>30MG</u>	<u>N72369</u>	<u>001</u>	Mar 30, 1989
<u>AB</u>	WEST WARD	<u>15MG</u>	<u>N71107</u>	<u>001</u>	Dec 08, 1986
<u>AB</u>		<u>30MG</u>	<u>N71108</u>	<u>001</u>	Dec 08, 1986

FLURBIPROFEN

TABLET; ORAL

ANSAID

<u>AB</u>	PHARMACIA AND UPJOHN	<u>50MG</u>	<u>N18766</u>	<u>002</u>	Oct 31, 1988
<u>AB</u>	+	<u>100MG</u>	<u>N18766</u>	<u>003</u>	Oct 31, 1988

FLURBIPROFEN

<u>AB</u>	CARACO	<u>50MG</u>	<u>N75058</u>	<u>001</u>	Apr 27, 2001
<u>AB</u>		<u>100MG</u>	<u>N75058</u>	<u>002</u>	Apr 27, 2001
<u>AB</u>	MYLAN	<u>50MG</u>	<u>N74358</u>	<u>001</u>	Jun 20, 1994
<u>AB</u>		<u>100MG</u>	<u>N74358</u>	<u>002</u>	Jun 20, 1994
<u>AB</u>	PLIVA	<u>50MG</u>	<u>N74647</u>	<u>001</u>	Apr 01, 1997
<u>AB</u>		<u>100MG</u>	<u>N74647</u>	<u>002</u>	Apr 01, 1997
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N74448</u>	<u>001</u>	Jul 28, 1995
<u>AB</u>		<u>100MG</u>	<u>N74448</u>	<u>002</u>	Jul 28, 1995
<u>AB</u>	TEVA	<u>50MG</u>	<u>N74405</u>	<u>002</u>	May 24, 1995
<u>AB</u>		<u>100MG</u>	<u>N74405</u>	<u>001</u>	May 24, 1995
<u>AB</u>		<u>100MG</u>	<u>N74431</u>	<u>001</u>	May 31, 1995
<u>AB</u>	WARNER CHILCOTT	<u>100MG</u>	<u>N74560</u>	<u>002</u>	May 16, 1997

FLURBIPROFEN SODIUM

SOLUTION/DROPS; OPHTHALMIC

FLURBIPROFEN SODIUM

<u>AT</u>	AVENTIS BEHRING	<u>0.03%</u>	<u>N74447</u>	<u>001</u>	Jan 04, 1995
	<u>OCUFEN</u>				
<u>AT</u>	+	<u>ALLERGAN</u>	<u>N19404</u>	<u>001</u>	Dec 31, 1986

FLUTAMIDE

CAPSULE; ORAL

EULEXIN

<u>AB</u>	+	SCHERING	<u>125MG</u>	<u>N18554</u>	<u>001</u>	Jan 27, 1989
-----------	---	----------	--------------	---------------	------------	--------------

FLUTAMIDE

<u>AB</u>	BARR	<u>125MG</u>	<u>N75820</u>	<u>001</u>	Sep 18, 2001
<u>AB</u>	GENPHARM	<u>125MG</u>	<u>N76224</u>	<u>001</u>	May 09, 2003
<u>AB</u>	IVAX PHARMS	<u>125MG</u>	<u>N75780</u>	<u>001</u>	Sep 19, 2001
<u>AB</u>	SANDOZ	<u>125MG</u>	<u>N75818</u>	<u>001</u>	Sep 18, 2001
<u>AB</u>	TEVA	<u>125MG</u>	<u>N75298</u>	<u>001</u>	Sep 18, 2001

FLUTICASONE PROPIONATE

AEROSOL, METERED; INHALATION

FLOVENT

+	GLAXOSMITHKLINE	0.044MG/INH	N20548	001	Mar 27, 1996
+		0.11MG/INH	N20548	002	Mar 27, 1996
+		0.22MG/INH	N20548	003	Mar 27, 1996
	FLOVENT HFA				
+	GLAXOSMITHKLINE	0.044MG/INH	N21433	003	May 14, 2004
+		0.11MG/INH	N21433	002	May 14, 2004
+		0.22MG/INH	N21433	001	May 14, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 170 (of 360)

FLUTICASONE PROPIONATE

CREAM; TOPICAL					
<u>CUTIVATE</u>					
<u>AB</u>	+	ALTANA	<u>0.05%</u>	<u>N19958</u>	<u>001</u> Dec 18, 1990
<u>FLUTICASONE PROPIONATE</u>					
<u>AB</u>		ALTANA	<u>0.05%</u>	<u>N76451</u>	<u>001</u> May 14, 2004
<u>AB</u>		CLAY PARK	<u>0.05%</u>	<u>N76793</u>	<u>001</u> May 14, 2004
<u>AB</u>		KV PHARM	<u>0.05%</u>	<u>N76865</u>	<u>001</u> Sep 10, 2004
<u>AB</u>		QLT USA	<u>0.05%</u>	<u>N76633</u>	<u>001</u> May 14, 2004
LOTION; TOPICAL					
CUTIVATE					
	+	ALTANA	0.05%	N21152	001 Mar 31, 2005
OINTMENT; TOPICAL					
<u>CUTIVATE</u>					
<u>AB</u>	+	ALTANA	<u>0.005%</u>	<u>N19957</u>	<u>001</u> Dec 14, 1990
<u>FLUTICASONE PROPIONATE</u>					
<u>AB</u>		ALTANA	<u>0.005%</u>	<u>N76300</u>	<u>001</u> May 14, 2004
<u>AB</u>		CLAY PARK	<u>0.005%</u>	<u>N76668</u>	<u>001</u> May 14, 2004
<u>AB</u>		TARO PHARM INDS	<u>0.005%</u>	<u>N77145</u>	<u>001</u> Jun 14, 2005
POWDER; INHALATION					
FLOVENT					
	+	GLAXOSMITHKLINE	0.044MG/INH	N20549	001 Nov 07, 1997
	+		0.088MG/INH	N20549	002 Nov 07, 1997
	+		0.22MG/INH	N20549	003 Nov 07, 1997
FLOVENT DISKUS 100					
	+	GLAXOSMITHKLINE	0.1MG/INH	N20833	002 Sep 29, 2000
FLOVENT DISKUS 250					
	+	GLAXOSMITHKLINE	0.25MG/INH	N20833	003 Sep 29, 2000
FLOVENT DISKUS 50					
	+	GLAXOSMITHKLINE	0.05MG/INH	N20833	001 Sep 29, 2000
SPRAY, METERED; NASAL					
FLONASE					
	+	GLAXOSMITHKLINE	0.05MG/SPRAY	N20121	001 Oct 19, 1994

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

POWDER; INHALATION					
ADVAIR DISKUS 100/50					
	+	GLAXOSMITHKLINE	0.1MG/INH;EQ 0.05MG BASE/INH	N21077	001 Aug 24, 2000
ADVAIR DISKUS 250/50					
	+	GLAXOSMITHKLINE	0.25MG/INH;EQ 0.05MG BASE/INH	N21077	002 Aug 24, 2000
ADVAIR DISKUS 500/50					
	+	GLAXOSMITHKLINE	0.5MG/INH;EQ 0.05MG BASE/INH	N21077	003 Aug 24, 2000

FLUVASTATIN SODIUM

CAPSULE; ORAL					
LESCOL					
		NOVARTIS	EQ 20MG BASE	N20261	001 Dec 31, 1993
	+		EQ 40MG BASE	N20261	002 Dec 31, 1993
TABLET, EXTENDED RELEASE; ORAL					
LESCOL XL					
	+	NOVARTIS	80MG	N21192	001 Oct 06, 2000

FLUVOXAMINE MALEATE

TABLET; ORAL					
<u>FLUVOXAMINE MALEATE</u>					
<u>AB</u>		BARR	<u>25MG</u>	<u>N75897</u>	<u>001</u> Jan 25, 2001
<u>AB</u>			<u>50MG</u>	<u>N75897</u>	<u>002</u> Jan 25, 2001
<u>AB</u>			<u>100MG</u>	<u>N75897</u>	<u>003</u> Jan 25, 2001
<u>AB</u>		GENPHARM	<u>50MG</u>	<u>N75950</u>	<u>001</u> Oct 15, 2001

PRESCRIPTION DRUG PRODUCT LIST

3 - 171 (of 360)

FLUVOXAMINE MALEATE

TABLET; ORAL

FLUVOXAMINE MALEATE

<u>AB</u>	GENPHARM	<u>100MG</u>	<u>N75950</u>	<u>002</u>	Oct 15, 2001
<u>AB</u>	IVAX PHARMS	<u>25MG</u>	<u>N75898</u>	<u>001</u>	Mar 12, 2001
<u>AB</u>		<u>50MG</u>	<u>N75898</u>	<u>002</u>	Mar 12, 2001
<u>AB</u>		<u>100MG</u>	<u>N75898</u>	<u>003</u>	Mar 12, 2001
<u>AB</u>	MUTUAL PHARM	<u>25MG</u>	<u>N76125</u>	<u>001</u>	Apr 29, 2002
<u>AB</u>		<u>50MG</u>	<u>N76125</u>	<u>002</u>	Apr 29, 2002
<u>AB</u>		<u>100MG</u>	<u>N76125</u>	<u>003</u>	Apr 29, 2002
<u>AB</u>	MYLAN	<u>25MG</u>	<u>N75889</u>	<u>001</u>	Nov 29, 2000
<u>AB</u>		<u>50MG</u>	<u>N75889</u>	<u>002</u>	Nov 29, 2000
<u>AB</u>		<u>100MG</u>	<u>N75889</u>	<u>003</u>	Nov 29, 2000
<u>AB</u>	PUREPAC PHARM	<u>25MG</u>	<u>N75901</u>	<u>001</u>	Dec 28, 2000
<u>AB</u>		<u>50MG</u>	<u>N75901</u>	<u>002</u>	Dec 28, 2000
<u>AB</u>		<u>100MG</u>	<u>N75901</u>	<u>003</u>	Dec 28, 2000
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>N75887</u>	<u>001</u>	Jan 05, 2001
<u>AB</u>		<u>25MG</u>	<u>N75888</u>	<u>001</u>	Nov 29, 2000
<u>AB</u>		<u>50MG</u>	<u>N75887</u>	<u>002</u>	Jan 05, 2001
<u>AB</u>		<u>50MG</u>	<u>N75888</u>	<u>002</u>	Nov 29, 2000
<u>AB</u>		<u>100MG</u>	<u>N75887</u>	<u>003</u>	Jan 05, 2001
<u>AB</u>	+	<u>100MG</u>	<u>N75888</u>	<u>003</u>	Nov 29, 2000
<u>AB</u>	SYNTHON PHARMS	<u>25MG</u>	<u>N75899</u>	<u>001</u>	Jan 17, 2001
<u>AB</u>		<u>50MG</u>	<u>N75899</u>	<u>002</u>	Jan 17, 2001
<u>AB</u>		<u>100MG</u>	<u>N75899</u>	<u>003</u>	Jan 17, 2001
<u>AB</u>	TEVA	<u>25MG</u>	<u>N75893</u>	<u>001</u>	Sep 10, 2002
<u>AB</u>		<u>50MG</u>	<u>N75893</u>	<u>002</u>	Sep 10, 2002
<u>AB</u>		<u>100MG</u>	<u>N75893</u>	<u>003</u>	Sep 10, 2002
<u>AB</u>	TORPHARM	<u>25MG</u>	<u>N75902</u>	<u>001</u>	May 07, 2001
<u>AB</u>		<u>50MG</u>	<u>N75902</u>	<u>002</u>	May 07, 2001
<u>AB</u>		<u>100MG</u>	<u>N75902</u>	<u>003</u>	May 07, 2001
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>N75894</u>	<u>001</u>	Apr 18, 2001
<u>AB</u>		<u>50MG</u>	<u>N75894</u>	<u>002</u>	Apr 18, 2001
<u>AB</u>		<u>100MG</u>	<u>N75894</u>	<u>003</u>	Apr 18, 2001

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

<u>AP</u>	+	AM PHARM PARTNERS	<u>5MG/ML</u>	<u>N89202</u>	<u>001</u>	Feb 18, 1986
<u>AP</u>		LOCH	<u>5MG/ML</u>	<u>N81066</u>	<u>001</u>	Dec 29, 1993

TABLET; ORAL

FOLIC ACID

<u>AA</u>		MUTUAL PHARM	<u>1MG</u>	<u>N40582</u>	<u>001</u>	Jul 18, 2005
<u>AA</u>		PHARM FORM	<u>1MG</u>	<u>N85061</u>	<u>001</u>	
<u>AA</u>		PHARMAX	<u>1MG</u>	<u>N40625</u>	<u>001</u>	Jul 21, 2005
<u>AA</u>		TABLICAPS	<u>1MG</u>	<u>N83133</u>	<u>002</u>	
<u>AA</u>		TRIGEN	<u>1MG</u>	<u>N40514</u>	<u>001</u>	Jun 14, 2005
<u>AA</u>	+	WATSON LABS	<u>1MG</u>	<u>N80680</u>	<u>001</u>	
<u>AA</u>		WEST WARD	<u>1MG</u>	<u>N80600</u>	<u>001</u>	
<u>FOLICET</u>						
<u>AA</u>		MISSION PHARMA	<u>1MG</u>	<u>N87438</u>	<u>001</u>	

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

+	ORGANON USA INC	75 IU/0.5ML	N21273	001	Aug 26, 2005
+		150 IU/0.18ML	N21211	003	Feb 11, 2004
+		150 IU/0.5ML	N21273	002	Aug 26, 2005
+		300 IU/0.36ML	N21211	001	Mar 23, 2004
+		600 IU/0.72ML	N21211	002	Mar 23, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 172 (of 360)

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

+	ORGANON USA INC	900 IU/1.08ML	N21211	004	Feb 11, 2005
	GONAL-F				
+	SERONO INC	450 IU/VIAL	N20378	005	Mar 26, 2004
	GONAL-F RFF				
+	SERONO INC	75 IU/VIAL	N21765	002	Mar 25, 2004
	GONAL-F RFF PEN				
+	SERONO INC	300 IU/0.5ML	N21684	001	May 25, 2004
+		450 IU/0.75ML	N21684	002	May 25, 2004
+		900 IU/1.5ML	N21684	003	May 25, 2004

FOMEPIZOLE

INJECTABLE; INJECTION

ANTIZOL

+	ORPHAN MEDCL	1GM/ML	N20696	001	Dec 04, 1997
---	--------------	--------	--------	-----	--------------

FONDAPARINUX SODIUM

INJECTABLE; SUBCUTANEOUS

ARIXTRA

+	GLAXOSMITHKLINE	2.5MG/0.5ML	N21345	001	Dec 07, 2001
+		5MG/0.4ML	N21345	002	May 28, 2004
+		7.5MG/0.6ML	N21345	003	May 28, 2004
+		10MG/0.8ML	N21345	004	May 28, 2004

FORMOTEROL FUMARATE

POWDER; INHALATION

FORADIL

+	NOVARTIS	0.012MG/INH	N20831	001	Feb 16, 2001
---	----------	-------------	--------	-----	--------------

FOSAMPRENAVIR CALCIUM

TABLET; ORAL

LEXIVA

+	GLAXOSMITHKLINE	EQ 700MG BASE	N21548	001	Oct 20, 2003
---	-----------------	---------------	--------	-----	--------------

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

<u>AP</u>	PHARMAFORCE	<u>2.4GM/100ML</u>	<u>N77174</u>	<u>001</u>	May 31, 2005
------------------	-------------	---------------------------	----------------------	-------------------	--------------

FOSCAVIR

<u>AP</u>	+	ASTRAZENECA	<u>2.4GM/100ML</u>	<u>N20068</u>	<u>001</u>	Sep 27, 1991
------------------	---	-------------	---------------------------	----------------------	-------------------	--------------

FOSFOMYCIN TROMETHAMINE

FOR SUSPENSION; ORAL

MONUROL

+	ZAMBON	EQ 3GM BASE/PACKET	N50717	001	Dec 19, 1996
---	--------	--------------------	--------	-----	--------------

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

<u>AB</u>	ANDRX PHARMS	<u>10MG</u>	<u>N76620</u>	<u>001</u>	Oct 15, 2004
<u>AB</u>		<u>20MG</u>	<u>N76620</u>	<u>002</u>	Oct 15, 2004
<u>AB</u>		<u>40MG</u>	<u>N76620</u>	<u>003</u>	Oct 15, 2004
<u>AB</u>	APOTEX	<u>10MG</u>	<u>N76906</u>	<u>001</u>	May 17, 2005
<u>AB</u>		<u>20MG</u>	<u>N76906</u>	<u>002</u>	May 17, 2005
<u>AB</u>		<u>40MG</u>	<u>N76906</u>	<u>003</u>	May 17, 2005
<u>AB</u>	INVAGEN PHARMS	<u>10MG</u>	<u>N77222</u>	<u>001</u>	Apr 20, 2005
<u>AB</u>		<u>20MG</u>	<u>N77222</u>	<u>002</u>	Apr 20, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 173 (of 360)

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

<u>AB</u>	INVAGEN PHARMS	<u>40MG</u>	<u>N77222</u>	<u>003</u>	Apr 20, 2005
<u>AB</u>	RANBAXY	<u>10MG</u>	<u>N76580</u>	<u>001</u>	Apr 23, 2004
<u>AB</u>		<u>20MG</u>	<u>N76580</u>	<u>002</u>	Apr 23, 2004
<u>AB</u>		<u>40MG</u>	<u>N76580</u>	<u>003</u>	Apr 23, 2004
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>N76188</u>	<u>001</u>	Oct 08, 2004
<u>AB</u>		<u>10MG</u>	<u>N76483</u>	<u>001</u>	Apr 23, 2004
<u>AB</u>		<u>20MG</u>	<u>N76188</u>	<u>002</u>	Oct 08, 2004
<u>AB</u>		<u>20MG</u>	<u>N76483</u>	<u>002</u>	Apr 23, 2004
<u>AB</u>		<u>40MG</u>	<u>N76188</u>	<u>003</u>	Oct 08, 2004
<u>AB</u>		<u>40MG</u>	<u>N76483</u>	<u>003</u>	Apr 23, 2004
<u>AB</u>	TEVA	<u>10MG</u>	<u>N76139</u>	<u>001</u>	Nov 25, 2003
<u>AB</u>		<u>20MG</u>	<u>N76139</u>	<u>002</u>	Nov 25, 2003
<u>AB</u>		<u>40MG</u>	<u>N76139</u>	<u>003</u>	Nov 25, 2003
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N76987</u>	<u>001</u>	Dec 23, 2004
<u>AB</u>		<u>20MG</u>	<u>N76987</u>	<u>002</u>	Dec 23, 2004
<u>AB</u>		<u>40MG</u>	<u>N76987</u>	<u>003</u>	Dec 23, 2004
<u>MONOPRIL</u>					
<u>AB</u>	BRISTOL MYERS SQUIBB	<u>10MG</u>	<u>N19915</u>	<u>002</u>	May 16, 1991
<u>AB</u>		<u>20MG</u>	<u>N19915</u>	<u>003</u>	May 16, 1991
<u>AB</u>	+	<u>40MG</u>	<u>N19915</u>	<u>004</u>	Mar 28, 1995

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

<u>AB</u>	ANDRX PHARMS	<u>10MG;12.5MG</u>	<u>N76608</u>	<u>001</u>	Dec 03, 2004
<u>AB</u>		<u>20MG;12.5MG</u>	<u>N76608</u>	<u>002</u>	Dec 03, 2004
<u>AB</u>	RANBAXY	<u>10MG;12.5MG</u>	<u>N76739</u>	<u>001</u>	Dec 17, 2004
<u>AB</u>		<u>20MG;12.5MG</u>	<u>N76739</u>	<u>002</u>	Dec 17, 2004
<u>AB</u>	SANDOZ	<u>10MG;12.5MG</u>	<u>N76961</u>	<u>001</u>	Sep 28, 2005
<u>AB</u>		<u>20MG;12.5MG</u>	<u>N76961</u>	<u>002</u>	Sep 28, 2005
<u>AB</u>	WATSON LABS	<u>10MG;12.5MG</u>	<u>N77144</u>	<u>001</u>	Aug 16, 2005
<u>AB</u>		<u>20MG;12.5MG</u>	<u>N77144</u>	<u>002</u>	Aug 16, 2005
<u>MONOPRIL - HCT</u>					
<u>AB</u>	BRISTOL MYERS SQUIBB	<u>10MG;12.5MG</u>	<u>N20286</u>	<u>002</u>	Nov 30, 1994
<u>AB</u>	+	<u>20MG;12.5MG</u>	<u>N20286</u>	<u>001</u>	Nov 30, 1994

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

CEREBYX

+	PARKE DAVIS	EQ 50MG PHENYTOIN NA/ML	N20450	001	Aug 05, 1996
---	-------------	-------------------------	--------	-----	--------------

FROVATRIPTAN SUCCINATE

TABLET; ORAL

FROVA

+	ENDO PHARMS	EQ 2.5MG BASE	N21006	001	Nov 08, 2001
---	-------------	---------------	--------	-----	--------------

FULVESTRANT

INJECTABLE; INTRAMUSCULAR

FASLODEX

+	ASTRAZENECA	50MG/ML	N21344	001	Apr 25, 2002
---	-------------	---------	--------	-----	--------------

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

<u>AP</u>	AM PHARM PARTNERS	<u>10MG/ML</u>	<u>N18902</u>	<u>001</u>	May 22, 1984
<u>AP</u>	BAXTER HLTHCARE	<u>10MG/ML</u>	<u>N71439</u>	<u>001</u>	Sep 14, 1990

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 174 (of 360)

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

<u>AP</u>	HOSPIRA	<u>10MG/ML</u>	<u>N18667</u>	<u>001</u>	May 28, 1982
<u>AP</u>		<u>10MG/ML</u>	<u>N70578</u>	<u>001</u>	Jul 08, 1987
<u>AP</u>		<u>10MG/ML</u>	<u>N72080</u>	<u>001</u>	Aug 13, 1991
<u>AP</u>		<u>10MG/ML</u>	<u>N74337</u>	<u>001</u>	Oct 31, 1994
<u>AP</u>		<u>10MG/ML</u>	<u>N75241</u>	<u>001</u>	May 28, 1999
<u>AP</u>	INTL MEDICATION	<u>10MG/ML</u>	<u>N18025</u>	<u>001</u>	
<u>AP</u>	+ LUITPOLD	<u>10MG/ML</u>	<u>N18579</u>	<u>001</u>	Nov 30, 1983

SOLUTION; ORAL

FUROSEMIDE

<u>AA</u>	MORTON GROVE	<u>10MG/ML</u>	<u>N70655</u>	<u>001</u>	Oct 02, 1987
<u>AA</u>	+ ROXANE	<u>10MG/ML</u>	<u>N70434</u>	<u>001</u>	Apr 22, 1987
		40MG/5ML	N70433	001	Apr 22, 1987

TABLET; ORAL

FUROSEMIDE

<u>AB</u>	EXCELLIUM	<u>20MG</u>	<u>N77293</u>	<u>001</u>	Nov 09, 2005
<u>AB</u>		<u>40MG</u>	<u>N77293</u>	<u>002</u>	Nov 09, 2005
<u>AB</u>		<u>80MG</u>	<u>N77293</u>	<u>003</u>	Nov 09, 2005
<u>AB</u>	INTL MEDICATION	<u>20MG</u>	<u>N18753</u>	<u>001</u>	Feb 28, 1984
<u>AB</u>		<u>40MG</u>	<u>N18753</u>	<u>002</u>	Feb 28, 1984
<u>AB</u>	IVAX PHARMS	<u>20MG</u>	<u>N18413</u>	<u>001</u>	Nov 30, 1983
<u>AB</u>		<u>40MG</u>	<u>N18413</u>	<u>002</u>	Nov 30, 1983
<u>AB</u>	MYLAN	<u>20MG</u>	<u>N18487</u>	<u>001</u>	
<u>AB</u>		<u>40MG</u>	<u>N18487</u>	<u>002</u>	
<u>AB</u>		<u>80MG</u>	<u>N70082</u>	<u>001</u>	Oct 29, 1986
<u>AB</u>	ROXANE	<u>20MG</u>	<u>N18823</u>	<u>001</u>	Nov 10, 1983
<u>AB</u>		<u>40MG</u>	<u>N18823</u>	<u>002</u>	Nov 10, 1983
<u>AB</u>		<u>80MG</u>	<u>N70086</u>	<u>001</u>	Jan 24, 1986
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>N18569</u>	<u>002</u>	
<u>AB</u>		<u>40MG</u>	<u>N18569</u>	<u>001</u>	
<u>AB</u>		<u>80MG</u>	<u>N18569</u>	<u>005</u>	Aug 14, 1984
<u>AB</u>	VINTAGE PHARMS	<u>20MG</u>	<u>N76796</u>	<u>001</u>	Mar 26, 2004
<u>AB</u>		<u>40MG</u>	<u>N76796</u>	<u>002</u>	Mar 26, 2004
<u>AB</u>		<u>80MG</u>	<u>N76796</u>	<u>003</u>	Mar 26, 2004
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>N70412</u>	<u>001</u>	Feb 26, 1986
<u>AB</u>		<u>20MG</u>	<u>N70449</u>	<u>001</u>	Nov 22, 1985
<u>AB</u>		<u>20MG</u>	<u>N71379</u>	<u>001</u>	Jan 02, 1987
<u>AB</u>		<u>40MG</u>	<u>N70450</u>	<u>001</u>	Nov 22, 1985
<u>AB</u>		<u>80MG</u>	<u>N70528</u>	<u>001</u>	Jan 07, 1986
<u>AB</u>		<u>80MG</u>	<u>N71594</u>	<u>001</u>	Feb 09, 1988
<u>LASIX</u>					
<u>AB</u>	AVENTIS PHARMS	<u>20MG</u>	<u>N16273</u>	<u>002</u>	
<u>AB</u>		<u>40MG</u>	<u>N16273</u>	<u>001</u>	
<u>AB</u>	+	<u>80MG</u>	<u>N16273</u>	<u>003</u>	

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

<u>AB</u>	APOTEX	<u>100MG</u>	<u>N75360</u>	<u>001</u>	Apr 06, 2005
<u>AB</u>		<u>300MG</u>	<u>N75360</u>	<u>002</u>	Apr 06, 2005
<u>AB</u>		<u>400MG</u>	<u>N75360</u>	<u>003</u>	Apr 06, 2005
<u>AB</u>	IVAX PHARMS	<u>100MG</u>	<u>N75477</u>	<u>001</u>	Mar 23, 2005
<u>AB</u>		<u>300MG</u>	<u>N75477</u>	<u>002</u>	Mar 23, 2005
<u>AB</u>		<u>400MG</u>	<u>N75477</u>	<u>003</u>	Mar 23, 2005
<u>AB</u>	MUTUAL PHARM	<u>100MG</u>	<u>N76537</u>	<u>001</u>	Jun 30, 2005
<u>AB</u>		<u>300MG</u>	<u>N76537</u>	<u>002</u>	Jun 30, 2005
<u>AB</u>		<u>400MG</u>	<u>N76537</u>	<u>003</u>	Jun 30, 2005
<u>AB</u>	PUREPAC PHARM	<u>100MG</u>	<u>N75350</u>	<u>001</u>	Sep 12, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 175 (of 360)

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

<u>AB</u>	PUREPAC PHARM	<u>300MG</u>	<u>N75350</u>	<u>002</u>	Sep 12, 2003
<u>AB</u>		<u>400MG</u>	<u>N75350</u>	<u>003</u>	Sep 12, 2003
<u>AB</u>	RANBAXY	<u>100MG</u>	<u>N76606</u>	<u>001</u>	Oct 07, 2005
<u>AB</u>		<u>300MG</u>	<u>N76606</u>	<u>002</u>	Oct 07, 2005
<u>AB</u>		<u>400MG</u>	<u>N76606</u>	<u>003</u>	Oct 07, 2005
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>N75539</u>	<u>001</u>	Apr 06, 2005
<u>AB</u>		<u>300MG</u>	<u>N75539</u>	<u>002</u>	Apr 06, 2005
<u>AB</u>		<u>400MG</u>	<u>N75539</u>	<u>003</u>	Apr 06, 2005
<u>AB</u>	TEVA PHARMS	<u>100MG</u>	<u>N75435</u>	<u>001</u>	Oct 08, 2004
<u>AB</u>		<u>300MG</u>	<u>N75435</u>	<u>002</u>	Oct 08, 2004
<u>AB</u>		<u>400MG</u>	<u>N75435</u>	<u>003</u>	Oct 08, 2004
<u>NEURONTIN</u>					
<u>AB</u>	PFIZER PHARMS	<u>100MG</u>	<u>N20235</u>	<u>001</u>	Dec 30, 1993
<u>AB</u>		<u>300MG</u>	<u>N20235</u>	<u>002</u>	Dec 30, 1993
<u>AB</u>	+	<u>400MG</u>	<u>N20235</u>	<u>003</u>	Dec 30, 1993

SOLUTION; ORAL

NEURONTIN

+	PARKE DAVIS	250MG/5ML	N21129	001	Mar 02, 2000
---	-------------	-----------	--------	-----	--------------

TABLET; ORAL

GABAPENTIN

<u>AB</u>	IVAX PHARMS	<u>600MG</u>	<u>N76017</u>	<u>004</u>	Apr 29, 2005
<u>AB</u>		<u>800MG</u>	<u>N76017</u>	<u>005</u>	Apr 29, 2005
		100MG	N76017	001	Apr 28, 2004
		300MG	N76017	002	Apr 28, 2004
		400MG	N76017	003	Apr 28, 2004
<u>AB</u>	PUREPAC PHARM	<u>600MG</u>	<u>N75694</u>	<u>001</u>	Oct 21, 2004
<u>AB</u>		<u>800MG</u>	<u>N75694</u>	<u>002</u>	Oct 21, 2004
<u>AB</u>	RANBAXY	<u>600MG</u>	<u>N76605</u>	<u>001</u>	Sep 14, 2005
<u>AB</u>		<u>800MG</u>	<u>N76605</u>	<u>002</u>	Sep 14, 2005
<u>AB</u>	TEVA	<u>600MG</u>	<u>N75827</u>	<u>001</u>	Dec 15, 2004
<u>AB</u>		<u>800MG</u>	<u>N75827</u>	<u>002</u>	Dec 15, 2004
<u>NEURONTIN</u>					
<u>AB</u>	PFIZER PHARMS	<u>600MG</u>	<u>N20882</u>	<u>001</u>	Oct 09, 1998
<u>AB</u>	+	<u>800MG</u>	<u>N20882</u>	<u>002</u>	Oct 09, 1998

GADOBENATE DIMEGLUMINE

INJECTABLE; INTRAVENOUS
MULTIHANCE

+	BRACCO	2.645GM/5ML (529MG/ML)	N21357	001	Nov 23, 2004
+		5.29GM/10ML (529MG/ML)	N21357	002	Nov 23, 2004
+		7.935GM/15ML (529MG/ML)	N21357	003	Nov 23, 2004
+		10.58GM/20ML (529MG/ML)	N21357	004	Nov 23, 2004
MULTIHANCE MULTIPACK					
+	BRACCO	26.45GM/50ML (529MG/ML)	N21358	001	Nov 23, 2004
+		52.9GM/100ML (529MG/ML)	N21358	002	Nov 23, 2004

GADODIAMIDE

INJECTABLE; INJECTION
OMNISCAN

+	GE HEALTHCARE	287MG/ML	N20123	001	Jan 08, 1993
---	---------------	----------	--------	-----	--------------

GADOPENTETATE DIMEGLUMINE

INJECTABLE; INJECTION
MAGNEVIST

+	BERLEX	469.01MG/ML	N19596	001	Jun 02, 1988
+		469.01MG/ML	N21037	001	Mar 10, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 176 (of 360)

GADOTERIDOL

INJECTABLE; INJECTION

PROHANCE

+	BRACCO	279.3MG/ML	N20131	001	Nov 16, 1992
---	--------	------------	--------	-----	--------------

PROHANCE MULTIPACK

+	BRACCO	279.3MG/ML	N21489	001	Oct 09, 2003
---	--------	------------	--------	-----	--------------

GADOVERSETAMIDE

INJECTABLE; INJECTION

OPTIMARK

+	MALLINCKRODT	330.9MG/ML	N20937	001	Dec 08, 1999
---	--------------	------------	--------	-----	--------------

+		330.9MG/ML	N20975	001	Dec 08, 1999
---	--	------------	--------	-----	--------------

OPTIMARK IN PLASTIC CONTAINER

+	MALLINCKRODT	330.9MG/ML	N20976	001	Dec 08, 1999
---	--------------	------------	--------	-----	--------------

GALANTAMINE HYDROBROMIDE

CAPSULE, EXTENDED RELEASE; ORAL

RAZADYNE ER

+	JANSSEN	EQ 8MG BASE	N21615	001	Dec 22, 2004
---	---------	-------------	--------	-----	--------------

		EQ 16MG BASE	N21615	002	Dec 22, 2004
--	--	--------------	--------	-----	--------------

		EQ 24MG BASE	N21615	003	Dec 22, 2004
--	--	--------------	--------	-----	--------------

SOLUTION; ORAL

RAZADYNE

+	JANSSEN PHARMA	4MG/ML	N21224	001	Jun 22, 2001
---	----------------	--------	--------	-----	--------------

TABLET; ORAL

RAZADYNE

+	JANSSEN PHARMA	EQ 4MG BASE	N21169	001	Feb 28, 2001
---	----------------	-------------	--------	-----	--------------

		EQ 8MG BASE	N21169	002	Feb 28, 2001
--	--	-------------	--------	-----	--------------

		EQ 12MG BASE	N21169	003	Feb 28, 2001
--	--	--------------	--------	-----	--------------

GALLIUM CITRATE, GA-67

INJECTABLE; INJECTION

GALLIUM CITRATE GA 67

BS	BRISTOL MYERS SQUIBB	2mCi/ML	N17478	001	
----	----------------------	---------	--------	-----	--

BS	MALLINCKRODT	2mCi/ML	N18058	001	
----	--------------	---------	--------	-----	--

GALLIUM NITRATE

INJECTABLE; INJECTION

GANITE

+	GENTA	25MG/ML	N19961	002	Jan 17, 1991
---	-------	---------	--------	-----	--------------

GANCICLOVIR

CAPSULE; ORAL

CYTOVENE

<u>AB</u>	ROCHE PALO	<u>250MG</u>	<u>N20460</u>	<u>001</u>	Dec 22, 1994
-----------	------------	--------------	---------------	------------	--------------

<u>AB</u>	+	<u>500MG</u>	<u>N20460</u>	<u>002</u>	Dec 12, 1997
-----------	---	--------------	---------------	------------	--------------

GANCICLOVIR

<u>AB</u>	RANBAXY	<u>250MG</u>	<u>N76457</u>	<u>001</u>	Jun 27, 2003
-----------	---------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N76457</u>	<u>002</u>	Jun 27, 2003
-----------	--	--------------	---------------	------------	--------------

IMPLANT; IMPLANTATION

VITRASERT

+	BAUSCH AND LOMB	4.5MG	N20569	001	Mar 04, 1996
---	-----------------	-------	--------	-----	--------------

GANCICLOVIR SODIUM

INJECTABLE; INJECTION

CYTOVENE IV

<u>AP</u>	+	ROCHE PALO	<u>EQ 500MG BASE/VIAL</u>	<u>N19661</u>	<u>001</u>	Jun 23, 1989
-----------	---	------------	---------------------------	---------------	------------	--------------

GANCICLOVIR SODIUM

<u>AP</u>	BEDFORD	<u>EQ 500MG BASE/VIAL</u>	<u>N76222</u>	<u>001</u>	Jul 16, 2003
-----------	---------	---------------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 177 (of 360)

GANIRELIX ACETATE

INJECTABLE; INJECTION
GANIRELIX ACETATE INJECTION

+ ORGANON USA INC	EQ 250UGM BASE/0.5ML	N21057 001	Jul 29, 1999
-------------------	----------------------	------------	--------------

GATIFLOXACIN

INJECTABLE; INJECTION
TEQUIN

+ BRISTOL MYERS SQUIBB	400MG/40ML(10MG/ML)	N21062 004	Dec 17, 1999
------------------------	---------------------	------------	--------------

TEQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER

+ BRISTOL MYERS SQUIBB	200MG/100ML(2MG/ML)	N21062 001	Dec 17, 1999
------------------------	---------------------	------------	--------------

+	400MG/200ML(2MG/ML)	N21062 002	Dec 17, 1999
---	---------------------	------------	--------------

SOLUTION/DROPS; OPHTHALMIC
ZYMAR

+ ALLERGAN	0.3%	N21493 001	Mar 28, 2003
------------	------	------------	--------------

SUSPENSION; ORAL
TEQUIN

+ BRISTOL MYERS SQUIBB	200MG/5ML	N21678 001	Aug 27, 2004
------------------------	-----------	------------	--------------

TABLET; ORAL
TEQUIN

BRISTOL MYERS SQUIBB	200MG	N21061 001	Dec 17, 1999
----------------------	-------	------------	--------------

+	400MG	N21061 002	Dec 17, 1999
---	-------	------------	--------------

GEFITINIB

TABLET; ORAL
IRESSA

+ ASTRAZENECA	250MG	N21399 001	May 05, 2003
---------------	-------	------------	--------------

GEMCITABINE HYDROCHLORIDE

INJECTABLE; INJECTION
GEMZAR

+ LILLY	EQ 200MG BASE/VIAL	N20509 001	May 15, 1996
---------	--------------------	------------	--------------

+	EQ 1GM BASE/VIAL	N20509 002	May 15, 1996
---	------------------	------------	--------------

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

<u>AB</u>	MYLAN	<u>600MG</u>	<u>N74452</u>	<u>001</u>	Feb 16, 1995
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>600MG</u>	<u>N74615</u>	<u>001</u>	Sep 29, 1995
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>600MG</u>	<u>N74256</u>	<u>001</u>	Oct 31, 1993
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>	TORPHARM	<u>600MG</u>	<u>N75034</u>	<u>001</u>	Jul 20, 1998
-----------	----------	--------------	---------------	------------	--------------

<u>AB</u>	WATSON LABS	<u>600MG</u>	<u>N74442</u>	<u>001</u>	Apr 28, 1995
-----------	-------------	--------------	---------------	------------	--------------

LOPID

<u>AB</u>	+ PFIZER PHARMS	<u>600MG</u>	<u>N18422</u>	<u>003</u>	Nov 20, 1986
-----------	-----------------	--------------	---------------	------------	--------------

GEMIFLOXACIN MESYLATE

TABLET; ORAL
FACTIVE

+ OSCIENT	EQ 320MG BASE	N21158 001	Apr 04, 2003
-----------	---------------	------------	--------------

GEMTUZUMAB OZOGAMICIN

INJECTABLE; INJECTION
MYLOTARG

+ WYETH PHARMS INC	5MG/VIAL	N21174 001	May 17, 2000
--------------------	----------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 178 (of 360)

GENTAMICIN SULFATE

CREAM; TOPICAL

GENTAMICIN

<u>AT</u>	CLAY PARK	<u>EQ 0.1% BASE</u>	<u>N62307</u>	<u>001</u>	
-----------	-----------	---------------------	---------------	------------	--

GENTAMICIN SULFATE

<u>AT</u>	+ FOUGERA	<u>EQ 0.1% BASE</u>	<u>N62531</u>	<u>001</u>	Jul 05, 1984
<u>AT</u>	TARO	<u>EQ 0.1% BASE</u>	<u>N62427</u>	<u>001</u>	May 26, 1983

INJECTABLE; INJECTION

GENTAMICIN SULFATE

<u>AP</u>	+ AM PHARM PARTNERS	<u>EQ 10MG BASE/ML</u>	<u>N62356</u>	<u>001</u>	Mar 04, 1982
<u>AP</u>	+	<u>EQ 40MG BASE/ML</u>	<u>N62356</u>	<u>002</u>	Mar 04, 1982
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N62366</u>	<u>001</u>	Aug 04, 1983
<u>AP</u>	HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>N62420</u>	<u>001</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>N62612</u>	<u>004</u>	Feb 20, 1986
<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N62420</u>	<u>002</u>	Aug 15, 1983
<u>AP</u>	PHARM SPEC	<u>EQ 40MG BASE/ML</u>	<u>N62340</u>	<u>001</u>	Mar 28, 1983

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>EQ 0.8MG BASE/ML</u>	<u>N62814</u>	<u>001</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 1.2MG BASE/ML</u>	<u>N62814</u>	<u>002</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 1.4MG BASE/ML</u>	<u>N62814</u>	<u>003</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 1.6MG BASE/ML</u>	<u>N62814</u>	<u>004</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 1.8MG BASE/ML</u>	<u>N62814</u>	<u>005</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>N62814</u>	<u>006</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 2.4MG BASE/ML</u>	<u>N62814</u>	<u>007</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 40MG BASE/100ML</u>	<u>N62814</u>	<u>008</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 60MG BASE/100ML</u>	<u>N62814</u>	<u>009</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 70MG BASE/100ML</u>	<u>N62814</u>	<u>010</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 80MG BASE/100ML</u>	<u>N62814</u>	<u>011</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 90MG BASE/100ML</u>	<u>N62814</u>	<u>012</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 100MG BASE/100ML</u>	<u>N62814</u>	<u>013</u>	Aug 28, 1987
<u>AP</u>		<u>EQ 120MG BASE/100ML</u>	<u>N62814</u>	<u>014</u>	Aug 28, 1987
<u>AP</u>	HOSPIRA	<u>EQ 1.2MG BASE/ML</u>	<u>N62414</u>	<u>001</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 1.4MG BASE/ML</u>	<u>N62414</u>	<u>002</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 1.6MG BASE/ML</u>	<u>N62414</u>	<u>003</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 1.8MG BASE/ML</u>	<u>N62414</u>	<u>004</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>N62414</u>	<u>005</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 60MG BASE/100ML</u>	<u>N62414</u>	<u>006</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 70MG BASE/100ML</u>	<u>N62414</u>	<u>007</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 80MG BASE/100ML</u>	<u>N62414</u>	<u>008</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 90MG BASE/100ML</u>	<u>N62414</u>	<u>009</u>	Aug 15, 1983
<u>AP</u>		<u>EQ 100MG BASE/100ML</u>	<u>N62414</u>	<u>010</u>	Aug 15, 1983

ISOTONIC GENTAMICIN SULFATE IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 0.8MG BASE/ML</u>	<u>N62373</u>	<u>001</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 1.2MG BASE/ML</u>	<u>N62373</u>	<u>007</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 1.6MG BASE/ML</u>	<u>N62373</u>	<u>008</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 2MG BASE/ML</u>	<u>N62373</u>	<u>009</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 2.4MG BASE/ML</u>	<u>N62373</u>	<u>010</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 40MG BASE/100ML</u>	<u>N62373</u>	<u>003</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 60MG BASE/100ML</u>	<u>N62373</u>	<u>004</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 80MG BASE/100ML</u>	<u>N62373</u>	<u>002</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 100MG BASE/100ML</u>	<u>N62373</u>	<u>005</u>	Sep 07, 1982
<u>AP</u>		<u>EQ 120MG BASE/100ML</u>	<u>N62373</u>	<u>006</u>	Sep 07, 1982

OINTMENT; OPHTHALMIC

GENTAMICIN SULFATE

<u>AT</u>	+ AKORN	<u>EQ 0.3% BASE</u>	<u>N64093</u>	<u>001</u>	Aug 31, 1995
<u>AT</u>	ALTANA	<u>EQ 0.3% BASE</u>	<u>N65024</u>	<u>001</u>	Jul 30, 2004

OINTMENT; TOPICAL

GENTAMICIN

<u>AT</u>	+ CLAY PARK	<u>EQ 0.1% BASE</u>	<u>N62351</u>	<u>001</u>	Feb 18, 1982
-----------	-------------	---------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 179 (of 360)

GENTAMICIN SULFATE

OINTMENT; TOPICAL

GENTAMICIN SULFATE

<u>AT</u>	FOUGERA	<u>EQ 0.1% BASE</u>	<u>N62533</u>	<u>001</u>	Oct 05, 1984
<u>AT</u>	TARO	<u>EQ 0.1% BASE</u>	<u>N62477</u>	<u>001</u>	Dec 23, 1983

SOLUTION/DROPS; OPHTHALMIC

GENOPTIC

<u>AT</u>	ALLERGAN	<u>EQ 0.3% BASE</u>	<u>N62452</u>	<u>001</u>	Oct 10, 1984
-----------	----------	---------------------	---------------	------------	--------------

GENTACIDIN

<u>AT</u>	NOVARTIS	<u>EQ 0.3% BASE</u>	<u>N62480</u>	<u>001</u>	Mar 30, 1984
-----------	----------	---------------------	---------------	------------	--------------

GENTAK

<u>AT</u>	AKORN	<u>EQ 0.3% BASE</u>	<u>N64163</u>	<u>001</u>	Oct 12, 2001
-----------	-------	---------------------	---------------	------------	--------------

GENTAMICIN SULFATE

<u>AT</u>	AKORN	<u>EQ 0.3% BASE</u>	<u>N62635</u>	<u>001</u>	Jan 08, 1987
<u>AT</u>	ALTANA	<u>EQ 0.3% BASE</u>	<u>N65121</u>	<u>001</u>	Jan 30, 2004
<u>AT</u>	+ BAUSCH AND LOMB	<u>EQ 0.3% BASE</u>	<u>N64048</u>	<u>001</u>	May 11, 1994
<u>AT</u>	FALCON PHARMS	<u>EQ 0.3% BASE</u>	<u>N62196</u>	<u>001</u>	

GENTAMICIN SULFATE; PREDNISOLONE ACETATE

OINTMENT; OPHTHALMIC

PRED-G

	+ ALLERGAN	EQ 0.3% BASE;0.6%	N50612	001	Dec 01, 1989
--	------------	-------------------	--------	-----	--------------

SUSPENSION/DROPS; OPHTHALMIC

PRED-G

	+ ALLERGAN	EQ 0.3% BASE;1%	N50586	001	Jun 10, 1988
--	------------	-----------------	--------	-----	--------------

GLATIRAMER ACETATE

FOR SOLUTION; SUBCUTANEOUS

COPAXONE

	+ TEVA	20MG/VIAL	N20622	001	Dec 20, 1996
--	--------	-----------	--------	-----	--------------

INJECTABLE; SUBCUTANEOUS

COPAXONE

	+ TEVA	20MG/ML	N20622	002	Feb 12, 2002
--	--------	---------	--------	-----	--------------

GLIMEPIRIDE

TABLET; ORAL

AMARYL

<u>AB</u>	+ AVENTIS PHARMS	<u>1MG</u>	<u>N20496</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>		<u>2MG</u>	<u>N20496</u>	<u>002</u>	Nov 30, 1995
<u>AB</u>		<u>4MG</u>	<u>N20496</u>	<u>003</u>	Nov 30, 1995

GLIMEPIRIDE

<u>AB</u>	COREPHARMA	<u>1MG</u>	<u>N77274</u>	<u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>N77274</u>	<u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>N77274</u>	<u>003</u>	Oct 06, 2005
<u>AB</u>	DR REDDYS LABS LTD	<u>1MG</u>	<u>N77091</u>	<u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>N77091</u>	<u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>N77091</u>	<u>003</u>	Oct 06, 2005
<u>AB</u>	INVAGEN PHARMS	<u>1MG</u>	<u>N77295</u>	<u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>N77295</u>	<u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>N77295</u>	<u>003</u>	Oct 06, 2005
<u>AB</u>	MYLAN	<u>1MG</u>	<u>N77624</u>	<u>001</u>	Nov 28, 2005
<u>AB</u>		<u>2MG</u>	<u>N77624</u>	<u>002</u>	Nov 28, 2005
<u>AB</u>		<u>4MG</u>	<u>N77624</u>	<u>003</u>	Nov 28, 2005
<u>AB</u>	PAR PHARM	<u>1MG</u>	<u>N77370</u>	<u>001</u>	Dec 23, 2005
<u>AB</u>		<u>2MG</u>	<u>N77370</u>	<u>002</u>	Dec 23, 2005
<u>AB</u>		<u>4MG</u>	<u>N77370</u>	<u>003</u>	Dec 23, 2005
<u>AB</u>		<u>8MG</u>	<u>N77370</u>	<u>004</u>	Dec 23, 2005
<u>AB</u>	RANBAXY	<u>1MG</u>	<u>N76875</u>	<u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>N76875</u>	<u>002</u>	Oct 06, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 180 (of 360)

GLIMEPIRIDE

TABLET; ORAL

GLIMEPIRIDE

<u>AB</u>	RANBAXY	<u>4MG</u>	<u>N76875</u>	<u>003</u>	Oct 06, 2005
<u>AB</u>		<u>8MG</u>	<u>N76875</u>	<u>004</u>	Oct 06, 2005
		3MG	N77366	001	Oct 06, 2005
		6MG	N77366	002	Oct 06, 2005
<u>AB</u>	TEVA	<u>1MG</u>	<u>N76802</u>	<u>001</u>	Oct 06, 2005
<u>AB</u>		<u>2MG</u>	<u>N76802</u>	<u>002</u>	Oct 06, 2005
<u>AB</u>		<u>4MG</u>	<u>N76802</u>	<u>003</u>	Oct 06, 2005

GLIMEPIRIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDARYL

SB PHARMC

	1MG; 4MG	N21700	001	Nov 23, 2005
	2MG; 4MG	N21700	002	Nov 23, 2005
+	4MG; 4MG	N21700	003	Nov 23, 2005

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

<u>AB</u>	ALPHAPHARM	<u>5MG</u>	<u>N74438</u>	<u>001</u>	Jun 20, 1995
<u>AB</u>		<u>10MG</u>	<u>N74438</u>	<u>002</u>	Jun 20, 1995
<u>AB</u>	ENDO PHARMS	<u>5MG</u>	<u>N74378</u>	<u>001</u>	Nov 28, 1994
<u>AB</u>		<u>10MG</u>	<u>N74378</u>	<u>002</u>	Nov 28, 1994
<u>AB</u>	IVAX PHARMS	<u>5MG</u>	<u>N74497</u>	<u>001</u>	Aug 31, 1995
<u>AB</u>		<u>10MG</u>	<u>N74497</u>	<u>002</u>	Aug 31, 1995
<u>AB</u>	KALI LABS	<u>5MG</u>	<u>N74550</u>	<u>001</u>	Sep 11, 1997
<u>AB</u>		<u>10MG</u>	<u>N74550</u>	<u>002</u>	Sep 11, 1997
<u>AB</u>	MYLAN	<u>5MG</u>	<u>N74226</u>	<u>001</u>	May 10, 1994
<u>AB</u>		<u>10MG</u>	<u>N74226</u>	<u>002</u>	May 10, 1994
<u>AB</u>	PLIVA	<u>5MG</u>	<u>N74619</u>	<u>001</u>	Apr 04, 1997
<u>AB</u>		<u>10MG</u>	<u>N74619</u>	<u>002</u>	Apr 04, 1997
<u>AB</u>	SANDOZ	<u>5MG</u>	<u>N74305</u>	<u>001</u>	Apr 07, 1995
<u>AB</u>		<u>5MG</u>	<u>N74542</u>	<u>001</u>	Jun 20, 1995
<u>AB</u>		<u>10MG</u>	<u>N74305</u>	<u>002</u>	Apr 07, 1995
<u>AB</u>		<u>10MG</u>	<u>N74542</u>	<u>002</u>	Jun 20, 1995
<u>AB</u>	TEVA	<u>5MG</u>	<u>N74387</u>	<u>001</u>	Mar 04, 1996
<u>AB</u>		<u>10MG</u>	<u>N74387</u>	<u>002</u>	Mar 04, 1996
<u>AB</u>	TORPHARM	<u>5MG</u>	<u>N75795</u>	<u>001</u>	Jun 13, 2001
<u>AB</u>		<u>10MG</u>	<u>N75795</u>	<u>002</u>	Jun 13, 2001
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>N74223</u>	<u>001</u>	Feb 27, 1995
<u>AB</u>		<u>5MG</u>	<u>N74370</u>	<u>001</u>	Nov 22, 1994
<u>AB</u>		<u>10MG</u>	<u>N74223</u>	<u>002</u>	Feb 27, 1995
<u>AB</u>		<u>10MG</u>	<u>N74370</u>	<u>002</u>	Nov 22, 1994
<u>AB</u>	<u>GLUCOTROL</u>				
<u>AB</u>	PFIZER	<u>5MG</u>	<u>N17783</u>	<u>001</u>	May 08, 1984
<u>AB</u>	+	<u>10MG</u>	<u>N17783</u>	<u>002</u>	May 08, 1984

TABLET, EXTENDED RELEASE; ORAL

GLIPIZIDE

<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>N76467</u>	<u>001</u>	Sep 08, 2003
<u>AB</u>		<u>10MG</u>	<u>N76467</u>	<u>002</u>	Nov 07, 2003
<u>AB</u>	<u>GLUCOTROL XL</u>				
<u>AB</u>	PFIZER	<u>5MG</u>	<u>N20329</u>	<u>001</u>	Apr 26, 1994
<u>AB</u>	+	<u>10MG</u>	<u>N20329</u>	<u>002</u>	Apr 26, 1994
		2.5MG	N20329	003	Aug 10, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 181 (of 360)

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>	COREPHARMA	<u>2.5MG;250MG</u>	<u>N77507</u>	<u>001</u>	Oct 27, 2005
<u>AB</u>		<u>2.5MG;500MG</u>	<u>N77507</u>	<u>002</u>	Oct 27, 2005
<u>AB</u>		<u>5MG;500MG</u>	<u>N77507</u>	<u>003</u>	Oct 27, 2005
<u>AB</u>	TEVA PHARMS	<u>2.5MG;250MG</u>	<u>N77270</u>	<u>001</u>	Oct 28, 2005
<u>AB</u>		<u>2.5MG;500MG</u>	<u>N77270</u>	<u>002</u>	Oct 28, 2005
<u>AB</u>		<u>5MG;500MG</u>	<u>N77270</u>	<u>003</u>	Oct 28, 2005
<u>METAGLIP</u>					
<u>AB</u>	BRISTOL MYERS SQUIBB	<u>2.5MG;250MG</u>	<u>N21460</u>	<u>001</u>	Oct 21, 2002
<u>AB</u>		<u>2.5MG;500MG</u>	<u>N21460</u>	<u>002</u>	Oct 21, 2002
<u>AB</u>	+	<u>5MG;500MG</u>	<u>N21460</u>	<u>003</u>	Oct 21, 2002

GLUCAGON HYDROCHLORIDE RECOMBINANT

INJECTABLE; INJECTION

GLUCAGON

+	NOVO NORDISK	EQ 1MG BASE/VIAL	N20918	001	Jun 22, 1998
---	--------------	------------------	--------	-----	--------------

GLUCAGON RECOMBINANT

INJECTABLE; INJECTION

GLUCAGON

+	LILLY	1MG/VIAL	N20928	001	Sep 11, 1998
---	-------	----------	--------	-----	--------------

GLYBURIDE

TABLET; ORAL

DIABETA

BX	AVENTIS PHARMS	1.25MG	N17532	001	May 01, 1984
BX		2.5MG	N17532	002	May 01, 1984
BX		5MG	N17532	003	May 01, 1984

GLYBURIDE

<u>AB</u>	AMIDE PHARM	<u>1.5MG</u>	<u>N75947</u>	<u>001</u>	Nov 14, 2002
<u>AB</u>		<u>3MG</u>	<u>N75947</u>	<u>002</u>	Nov 14, 2002
<u>AB</u>		<u>6MG</u>	<u>N75947</u>	<u>003</u>	Nov 14, 2002
<u>AB</u>	COREPHARMA	<u>1.25MG</u>	<u>N76257</u>	<u>001</u>	Jun 27, 2002
<u>AB</u>		<u>2.5MG</u>	<u>N76257</u>	<u>002</u>	Jun 27, 2002
<u>AB</u>		<u>5MG</u>	<u>N76257</u>	<u>003</u>	Jun 27, 2002
<u>AB</u>	TEVA	<u>1.25MG</u>	<u>N74388</u>	<u>001</u>	Aug 29, 1995
<u>AB</u>		<u>2.5MG</u>	<u>N74388</u>	<u>002</u>	Aug 29, 1995
<u>AB</u>		<u>5MG</u>	<u>N74388</u>	<u>003</u>	Aug 29, 1995

GLYBURIDE (MICRONIZED)

<u>AB</u>	CLONMEL HLTHCARE	<u>1.5MG</u>	<u>N74591</u>	<u>001</u>	Dec 22, 1997
<u>AB</u>		<u>3MG</u>	<u>N74591</u>	<u>002</u>	Dec 22, 1997
<u>AB</u>		<u>4.5MG</u>	<u>N74591</u>	<u>003</u>	Dec 22, 1997
<u>AB</u>		<u>6MG</u>	<u>N74591</u>	<u>004</u>	Dec 22, 1997
<u>AB</u>	HIKMA	<u>1.5MG</u>	<u>N75890</u>	<u>001</u>	Jul 31, 2003
<u>AB</u>		<u>3MG</u>	<u>N75890</u>	<u>002</u>	Jul 31, 2003
<u>AB</u>		<u>6MG</u>	<u>N75890</u>	<u>003</u>	Jul 31, 2003
<u>AB</u>	MYLAN	<u>1.5MG</u>	<u>N74792</u>	<u>001</u>	Jun 26, 1998
<u>AB</u>		<u>3MG</u>	<u>N74792</u>	<u>002</u>	Jun 26, 1998
<u>AB</u>		<u>6MG</u>	<u>N74792</u>	<u>003</u>	Aug 17, 1999
<u>AB</u>	SANDOZ	<u>1.5MG</u>	<u>N75174</u>	<u>001</u>	Jun 22, 1998
<u>AB</u>		<u>3MG</u>	<u>N75174</u>	<u>002</u>	Jun 22, 1998
<u>AB</u>	TEVA	<u>1.5MG</u>	<u>N74686</u>	<u>001</u>	Apr 20, 1999
<u>AB</u>		<u>3MG</u>	<u>N74686</u>	<u>002</u>	Apr 20, 1999
<u>AB</u>		<u>4.5MG</u>	<u>N74686</u>	<u>003</u>	Apr 20, 1999
<u>AB</u>		<u>6MG</u>	<u>N74686</u>	<u>004</u>	Apr 20, 1999

GLYNASE

<u>AB</u>	PHARMACIA AND UPJOHN	<u>1.5MG</u>	<u>N20051</u>	<u>001</u>	Mar 04, 1992
<u>AB</u>		<u>3MG</u>	<u>N20051</u>	<u>002</u>	Mar 04, 1992

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 182 (of 360)

GLYBURIDE

TABLET; ORAL

GLYNASE

<u>AB</u>	+	PHARMACIA AND UPJOHN	<u>6MG</u>	<u>N20051</u>	<u>004</u>	Sep 24, 1993
-----------	---	----------------------	------------	---------------	------------	--------------

MICRONASE

<u>AB</u>		PHARMACIA AND UPJOHN	<u>1.25MG</u>	<u>N17498</u>	<u>001</u>	May 01, 1984
-----------	--	----------------------	---------------	---------------	------------	--------------

<u>AB</u>			<u>2.5MG</u>	<u>N17498</u>	<u>002</u>	May 01, 1984
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>	+		<u>5MG</u>	<u>N17498</u>	<u>003</u>	May 01, 1984
-----------	---	--	------------	---------------	------------	--------------

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLUCOVANCE

<u>AB</u>		BRISTOL MYERS SQUIBB	<u>1.25MG; 250MG</u>	<u>N21178</u>	<u>001</u>	Jul 31, 2000
-----------	--	----------------------	----------------------	---------------	------------	--------------

<u>AB</u>			<u>2.5MG; 500MG</u>	<u>N21178</u>	<u>002</u>	Jul 31, 2000
-----------	--	--	---------------------	---------------	------------	--------------

<u>AB</u>	+		<u>5MG; 500MG</u>	<u>N21178</u>	<u>003</u>	Jul 31, 2000
-----------	---	--	-------------------	---------------	------------	--------------

GLYBURIDE AND METFORMIN HYDROCHLORIDE

<u>AB</u>		COREPHARMA	<u>1.25MG; 250MG</u>	<u>N76731</u>	<u>001</u>	Nov 19, 2004
-----------	--	------------	----------------------	---------------	------------	--------------

<u>AB</u>			<u>2.5MG; 500MG</u>	<u>N76731</u>	<u>002</u>	Nov 19, 2004
-----------	--	--	---------------------	---------------	------------	--------------

<u>AB</u>			<u>5MG; 500MG</u>	<u>N76731</u>	<u>003</u>	Nov 19, 2004
-----------	--	--	-------------------	---------------	------------	--------------

<u>AB</u>		IVAX PHARMS	<u>1.25MG; 250MG</u>	<u>N76345</u>	<u>001</u>	Feb 18, 2004
-----------	--	-------------	----------------------	---------------	------------	--------------

<u>AB</u>			<u>2.5MG; 500MG</u>	<u>N76345</u>	<u>002</u>	Feb 18, 2004
-----------	--	--	---------------------	---------------	------------	--------------

<u>AB</u>			<u>5MG; 500MG</u>	<u>N76345</u>	<u>003</u>	Feb 18, 2004
-----------	--	--	-------------------	---------------	------------	--------------

<u>AB</u>		PUREPAC PHARM	<u>1.25MG; 250MG</u>	<u>N76716</u>	<u>001</u>	Jun 28, 2005
-----------	--	---------------	----------------------	---------------	------------	--------------

<u>AB</u>			<u>2.5MG; 500MG</u>	<u>N76716</u>	<u>002</u>	Jun 28, 2005
-----------	--	--	---------------------	---------------	------------	--------------

<u>AB</u>			<u>5MG; 500MG</u>	<u>N76716</u>	<u>003</u>	Jun 28, 2005
-----------	--	--	-------------------	---------------	------------	--------------

<u>AB</u>		TEVA	<u>1.25MG; 250MG</u>	<u>N76821</u>	<u>001</u>	Jan 27, 2005
-----------	--	------	----------------------	---------------	------------	--------------

<u>AB</u>			<u>2.5MG; 500MG</u>	<u>N76821</u>	<u>002</u>	Jan 27, 2005
-----------	--	--	---------------------	---------------	------------	--------------

<u>AB</u>			<u>5MG; 500MG</u>	<u>N76821</u>	<u>003</u>	Jan 27, 2005
-----------	--	--	-------------------	---------------	------------	--------------

GLYCINE

SOLUTION; IRRIGATION

AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER

<u>AT</u>		BAXTER HLTHCARE	<u>1.5GM/100ML</u>	<u>N17865</u>	<u>001</u>	
-----------	--	-----------------	--------------------	---------------	------------	--

GLYCINE 1.5% IN PLASTIC CONTAINER

<u>AT</u>		B BRAUN	<u>1.5GM/100ML</u>	<u>N16784</u>	<u>001</u>	
-----------	--	---------	--------------------	---------------	------------	--

<u>AT</u>		HOSPIRA	<u>1.5GM/100ML</u>	<u>N17633</u>	<u>001</u>	
-----------	--	---------	--------------------	---------------	------------	--

<u>AT</u>			<u>1.5GM/100ML</u>	<u>N18315</u>	<u>001</u>	
-----------	--	--	--------------------	---------------	------------	--

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

<u>AP</u>		LUITPOLD	<u>0.2MG/ML</u>	<u>N89335</u>	<u>001</u>	Jul 23, 1986
-----------	--	----------	-----------------	---------------	------------	--------------

ROBINUL

<u>AP</u>	+	BAXTER HLTHCARE CORP	<u>0.2MG/ML</u>	<u>N17558</u>	<u>001</u>	
-----------	---	----------------------	-----------------	---------------	------------	--

TABLET; ORAL

GLYCOPYRROLATE

<u>AA</u>		COREPHARMA	<u>1MG</u>	<u>N40568</u>	<u>001</u>	Dec 22, 2004
-----------	--	------------	------------	---------------	------------	--------------

<u>AA</u>			<u>2MG</u>	<u>N40568</u>	<u>002</u>	Dec 22, 2004
-----------	--	--	------------	---------------	------------	--------------

ROBINUL

<u>AA</u>	+	FIRST HORIZON	<u>1MG</u>	<u>N12827</u>	<u>001</u>	
-----------	---	---------------	------------	---------------	------------	--

ROBINUL FORTE

<u>AA</u>	+	FIRST HORIZON	<u>2MG</u>	<u>N12827</u>	<u>002</u>	
-----------	---	---------------	------------	---------------	------------	--

GONADOTROPIN, CHORIONIC

INJECTABLE; INJECTION

CHORIONIC GONADOTROPIN

<u>AP</u>	+	AM PHARM PARTNERS	<u>10,000 UNITS/VIAL</u>	<u>N17067</u>	<u>002</u>	
-----------	---	-------------------	--------------------------	---------------	------------	--

<u>AP</u>	+	WATSON LABS (UTAH)	<u>10,000 UNITS/VIAL</u>	<u>N17016</u>	<u>007</u>	
-----------	---	--------------------	--------------------------	---------------	------------	--

PREGNYL

<u>AP</u>	+	ORGANON USA INC	<u>10,000 UNITS/VIAL</u>	<u>N17692</u>	<u>001</u>	
-----------	---	-----------------	--------------------------	---------------	------------	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 183 (of 360)

GOSERELIN ACETATE

IMPLANT; IMPLANTATION

ZOLADEX

+	ASTRAZENECA	EQ 3.6MG BASE	N19726	001	Dec 29, 1989
+		EQ 10.8MG BASE	N20578	001	Jan 11, 1996

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

<u>AT</u>	+	BAUSCH AND LOMB	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>N64047</u>	<u>001</u>	Jan 31, 1996
-----------	---	-----------------	---	---------------	------------	--------------

<u>AT</u>		PHARMAFORCE	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>N65187</u>	<u>001</u>	Oct 28, 2005
-----------	--	-------------	---	---------------	------------	--------------

NEOSPORIN

<u>AT</u>	+	MONARCH PHARMS	<u>0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML</u>	<u>N60582</u>	<u>001</u>	
-----------	---	----------------	---	---------------	------------	--

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

KYTRIL

+	ROCHE	EQ 0.1MG BASE/ML (EQ 0.1MG BASE/ML)	N20239	003	Sep 17, 2004
+		EQ 1MG BASE/ML (EQ 1MG BASE/ML)	N20239	004	Mar 11, 1994
+		EQ 4MG BASE/4ML (EQ 1MG BASE /ML)	N20239	002	Mar 11, 1994

SOLUTION; ORAL

KYTRIL

+	ROCHE	EQ 2MG BASE/10ML	N21238	001	Jun 27, 2001
---	-------	------------------	--------	-----	--------------

TABLET; ORAL

KYTRIL

+	ROCHE	EQ 1MG BASE	N20305	001	Mar 16, 1995
---	-------	-------------	--------	-----	--------------

GRISEOFULVIN, MICROCRYSTALLINE

SUSPENSION; ORAL

GRIFULVIN V

<u>AB</u>	+	J AND J	<u>125MG/5ML</u>	<u>N62483</u>	<u>001</u>	Jan 26, 1984
-----------	---	---------	------------------	---------------	------------	--------------

GRISEOFULVIN

<u>AB</u>		STIEFEL	<u>125MG/5ML</u>	<u>N65200</u>	<u>001</u>	Mar 02, 2005
-----------	--	---------	------------------	---------------	------------	--------------

TABLET; ORAL

FULVICIN-U/F

<u>AB</u>	+	SCHERING	<u>250MG</u>	<u>N60569</u>	<u>002</u>	
-----------	---	----------	--------------	---------------	------------	--

<u>AB</u>	+		<u>500MG</u>	<u>N60569</u>	<u>001</u>	
-----------	---	--	--------------	---------------	------------	--

GRIFULVIN V

<u>AB</u>		J AND J	<u>250MG</u>	<u>N62279</u>	<u>002</u>	
-----------	--	---------	--------------	---------------	------------	--

<u>AB</u>			<u>500MG</u>	<u>N62279</u>	<u>003</u>	
-----------	--	--	--------------	---------------	------------	--

		125MG	N62279	001	
--	--	-------	--------	-----	--

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL

GRIS-PEG

PEDINOL

125MG	N50475	001
-------	--------	-----

+	250MG	N50475	002
---	-------	--------	-----

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

<u>AB</u>		IVAX PHARMS	<u>EQ 4MG BASE</u>	<u>N74149</u>	<u>001</u>	Apr 07, 1995
-----------	--	-------------	--------------------	---------------	------------	--------------

<u>AB</u>			<u>EQ 8MG BASE</u>	<u>N74149</u>	<u>002</u>	Apr 07, 1995
-----------	--	--	--------------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>EQ 4MG BASE</u>	<u>N74517</u>	<u>001</u>	Sep 30, 1998
-----------	--	--------	--------------------	---------------	------------	--------------

<u>AB</u>			<u>EQ 8MG BASE</u>	<u>N74517</u>	<u>002</u>	Sep 30, 1998
-----------	--	--	--------------------	---------------	------------	--------------

<u>AB</u>		TEVA PHARMS	<u>EQ 4MG BASE</u>	<u>N74267</u>	<u>001</u>	Jun 01, 1994
-----------	--	-------------	--------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 184 (of 360)

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

<u>AB</u>	TEVA PHARMS	<u>EQ 8MG BASE</u>	<u>N74267</u>	<u>002</u>	Jun 01, 1994
<u>AB</u>	WATSON LABS	<u>EQ 4MG BASE</u>	<u>N74025</u>	<u>001</u>	Feb 28, 1994
<u>AB</u>	+	<u>EQ 8MG BASE</u>	<u>N74025</u>	<u>002</u>	Feb 28, 1994

GUANFACINE HYDROCHLORIDE

TABLET; ORAL

GUANFACINE HYDROCHLORIDE

<u>AB</u>	AMIDE PHARM	<u>EQ 1MG BASE</u>	<u>N74673</u>	<u>001</u>	Feb 28, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N74673</u>	<u>002</u>	Feb 28, 1997
<u>AB</u>	GENPHARM	<u>EQ 1MG BASE</u>	<u>N75109</u>	<u>001</u>	Nov 25, 1998
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N75109</u>	<u>002</u>	Nov 25, 1998
<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>N74796</u>	<u>001</u>	Jan 27, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N74796</u>	<u>002</u>	Jan 27, 1997
<u>AB</u>	WATSON LABS	<u>EQ 1MG BASE</u>	<u>N74145</u>	<u>001</u>	Oct 17, 1995
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N74762</u>	<u>001</u>	Jun 25, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N74145</u>	<u>002</u>	Oct 17, 1995
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N74762</u>	<u>002</u>	Jun 25, 1997
<u>TENEX</u>					
<u>AB</u>	ESP PHARMA	<u>EQ 1MG BASE</u>	<u>N19032</u>	<u>001</u>	Oct 27, 1986
<u>AB</u>	+	<u>EQ 2MG BASE</u>	<u>N19032</u>	<u>002</u>	Nov 07, 1988

GUANIDINE HYDROCHLORIDE

TABLET; ORAL

GUANIDINE HYDROCHLORIDE

SCHERING	125MG	N01546	001
----------	-------	--------	-----

HALCINONIDE

CREAM; TOPICAL

HALOG

+	WESTWOOD SQUIBB	0.1%	N17556	001
---	-----------------	------	--------	-----

OINTMENT; TOPICAL

HALOG

+	WESTWOOD SQUIBB	0.1%	N17824	001
---	-----------------	------	--------	-----

SOLUTION; TOPICAL

HALOG

WESTWOOD SQUIBB	0.1%	N17823	001
-----------------	------	--------	-----

HALOBETASOL PROPIONATE

CREAM; TOPICAL

HALOBETASOL PROPIONATE

<u>AB</u>	AGIS INDS	<u>0.05%</u>	<u>N77123</u>	<u>001</u>	Dec 16, 2004
<u>AB</u>	ALTANA	<u>0.05%</u>	<u>N77001</u>	<u>001</u>	Dec 16, 2004
<u>AB</u>	TARO	<u>0.05%</u>	<u>N77227</u>	<u>001</u>	Aug 04, 2005
<u>ULTRAVATE</u>					
<u>AB</u>	+ WESTWOOD SQUIBB	<u>0.05%</u>	<u>N19967</u>	<u>001</u>	Dec 27, 1990

OINTMENT; TOPICAL

HALOBETASOL PROPIONATE

<u>AB</u>	AGIS INDS	<u>0.05%</u>	<u>N76872</u>	<u>001</u>	Dec 16, 2004
<u>AB</u>	ALPHARMA US PHARMS	<u>0.05%</u>	<u>N77109</u>	<u>001</u>	Jun 14, 2005
<u>AB</u>	ALTANA	<u>0.05%</u>	<u>N76903</u>	<u>001</u>	Dec 16, 2004
<u>AB</u>	TARO	<u>0.05%</u>	<u>N76994</u>	<u>001</u>	Dec 16, 2004
<u>ULTRAVATE</u>					
<u>AB</u>	+ WESTWOOD SQUIBB	<u>0.05%</u>	<u>N19968</u>	<u>001</u>	Dec 17, 1990

PRESCRIPTION DRUG PRODUCT LIST

3 - 185 (of 360)

HALOPERIDOL

TABLET; ORAL

HALOPERIDOL

<u>AB</u>	MYLAN	<u>0.5MG</u>	<u>N70276</u>	<u>001</u>	Jun 10, 1986
<u>AB</u>		<u>1MG</u>	<u>N70277</u>	<u>001</u>	Jun 10, 1986
<u>AB</u>		<u>2MG</u>	<u>N70278</u>	<u>001</u>	Jun 10, 1986
<u>AB</u>		<u>5MG</u>	<u>N70279</u>	<u>001</u>	Jun 10, 1986
<u>AB</u>	PAR PHARM	<u>0.5MG</u>	<u>N71233</u>	<u>001</u>	Nov 03, 1986
<u>AB</u>		<u>1MG</u>	<u>N71234</u>	<u>001</u>	Nov 03, 1986
<u>AB</u>		<u>2MG</u>	<u>N71235</u>	<u>001</u>	Nov 03, 1986
<u>AB</u>		<u>5MG</u>	<u>N71236</u>	<u>001</u>	Nov 03, 1986
<u>AB</u>		<u>10MG</u>	<u>N71237</u>	<u>001</u>	Jul 20, 1987
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>N71206</u>	<u>001</u>	Nov 17, 1986
<u>AB</u>		<u>1MG</u>	<u>N71207</u>	<u>001</u>	Nov 17, 1986
<u>AB</u>	+	<u>2MG</u>	<u>N71208</u>	<u>001</u>	Nov 17, 1986
<u>AB</u>		<u>5MG</u>	<u>N71209</u>	<u>001</u>	Nov 17, 1986
<u>AB</u>		<u>10MG</u>	<u>N71210</u>	<u>001</u>	Mar 11, 1988
		20MG	N71211	001	Mar 11, 1988
<u>AB</u>	WATSON LABS	<u>0.5MG</u>	<u>N70981</u>	<u>001</u>	Mar 06, 1987
<u>AB</u>		<u>2MG</u>	<u>N70983</u>	<u>001</u>	Mar 06, 1987
<u>AB</u>		<u>5MG</u>	<u>N70984</u>	<u>001</u>	Mar 06, 1987

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION

HALDOL

<u>AO</u>	+ ORTHO MCNEIL	<u>EQ 50MG BASE/ML</u>	<u>N18701</u>	<u>001</u>	Jan 14, 1986
<u>AO</u>	+	<u>EQ 100MG BASE/ML</u>	<u>N18701</u>	<u>002</u>	Jan 31, 1997
	<u>HALOPERIDOL DECANOATE</u>				
<u>AO</u>	AM PHARM PARTNERS	<u>EQ 50MG BASE/ML</u>	<u>N74893</u>	<u>001</u>	Dec 19, 1997
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>N74893</u>	<u>002</u>	Dec 19, 1997
<u>AO</u>	APOTEX	<u>EQ 50MG BASE/ML</u>	<u>N75440</u>	<u>001</u>	Feb 28, 2000
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>N75440</u>	<u>002</u>	Feb 28, 2000
<u>AO</u>	BEDFORD	<u>EQ 50MG BASE/ML</u>	<u>N74811</u>	<u>001</u>	Jan 30, 1998
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>N75305</u>	<u>001</u>	Sep 28, 1998
<u>AO</u>	SABEX 2002	<u>EQ 50MG BASE/ML</u>	<u>N76463</u>	<u>001</u>	Jun 24, 2005
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>N76463</u>	<u>002</u>	Jun 24, 2005
<u>AO</u>	SICOR PHARMS	<u>EQ 50MG BASE/ML</u>	<u>N75393</u>	<u>001</u>	May 11, 1999
<u>AO</u>		<u>EQ 100MG BASE/ML</u>	<u>N75393</u>	<u>002</u>	May 11, 1999

HALOPERIDOL LACTATE

CONCENTRATE; ORAL

HALOPERIDOL

<u>AA</u>	PHARM ASSOC	<u>EQ 2MG BASE/ML</u>	<u>N73037</u>	<u>001</u>	Feb 26, 1993
<u>AA</u>	SILARX	<u>EQ 2MG BASE/ML</u>	<u>N73364</u>	<u>001</u>	Sep 28, 1993
<u>AA</u>	+ TEVA PHARMS	<u>EQ 2MG BASE/ML</u>	<u>N71617</u>	<u>001</u>	Dec 01, 1988

INJECTABLE; INJECTION

HALDOL

<u>AP</u>	+ ORTHO MCNEIL	<u>EQ 5MG BASE/ML</u>	<u>N15923</u>	<u>001</u>	
	<u>HALOPERIDOL</u>				
<u>AP</u>	AM PHARM PARTNERS	<u>EQ 5MG BASE/ML</u>	<u>N75689</u>	<u>001</u>	Mar 09, 2001
<u>AP</u>	APOTEX	<u>EQ 5MG BASE/ML</u>	<u>N76774</u>	<u>001</u>	Aug 25, 2004
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N76791</u>	<u>001</u>	Aug 25, 2004
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N76828</u>	<u>001</u>	Aug 25, 2004
<u>AP</u>	BEDFORD	<u>EQ 5MG BASE/ML</u>	<u>N75858</u>	<u>001</u>	Jun 18, 2001
<u>AP</u>	SABEX 2002	<u>EQ 5MG BASE/ML</u>	<u>N76464</u>	<u>001</u>	Sep 29, 2004
<u>AP</u>	SICOR PHARMS	<u>EQ 5MG BASE/ML</u>	<u>N76035</u>	<u>001</u>	Aug 29, 2001
<u>AP</u>	STERIS	<u>EQ 5MG BASE/ML</u>	<u>N70714</u>	<u>001</u>	May 17, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 186 (of 360)

HALOPERIDOL LACTATE

SOLUTION; ORAL

HALOPERIDOL LACTATE

UDL

EQ 1MG BASE/ML

N74536 001

Nov 02, 1995

HALOTHANE

LIQUID; INHALATION

HALOTHANEAN HALOCARBON99.99%N80810 001AN + HOSPIRA99.99%N83254 001HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN LOCK FLUSHAP AM PHARM10 UNITS/MLN17029 007

May 06, 1982

AP100 UNITS/MLN17029 006AP HOSPIRA10 UNITS/MLN05264 001AP10 UNITS/MLN40082 001

Feb 28, 1995

AP10 UNITS/MLN88097 001

Apr 28, 1983

AP10 UNITS/MLN88346 001

May 18, 1983

AP100 UNITS/MLN40082 002

Feb 28, 1995

AP100 UNITS/MLN88098 001

Apr 28, 1983

AP100 UNITS/MLN88347 001

May 18, 1983

HEPARIN LOCK FLUSH IN PLASTIC CONTAINERAP HOSPIRA10 UNITS/MLN05264 015

May 21, 1985

AP100 UNITS/MLN05264 016

May 21, 1985

HEPARIN SODIUMAP AM PHARM1,000 UNITS/MLN17029 001AP1,000 UNITS/MLN17979 001AP5,000 UNITS/MLN17651 006AP10,000 UNITS/MLN17029 003AP10,000 UNITS/MLN17979 002AP +20,000 UNITS/MLN17029 004AP

BAXTER HLTHCARE

1,000 UNITS/MLN17037 001AP5,000 UNITS/0.5MLN17037 013

Apr 07, 1986

AP5,000 UNITS/MLN17037 002AP10,000 UNITS/MLN17037 003AP HOSPIRA5,000 UNITS/MLN88100 001

Apr 28, 1983

AP MARSAM PHARMS LLC1,000 UNITS/MLN40008 001

Oct 10, 1995

AP + PHARMACIA AND UPJOHN1,000 UNITS/MLN04570 001AP +5,000 UNITS/MLN04570 002AP +10,000 UNITS/MLN04570 003HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINERAP BAXTER HLTHCARE200 UNITS/100MLN18609 001

Apr 28, 1982

HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINERAP B BRAUN200 UNITS/100MLN19953 001

Jul 20, 1992

AP HOSPIRA200 UNITS/100MLN18916 010

Jun 23, 1989

HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINERAP HOSPIRA10,000 UNITS/100MLN19339 003

Mar 27, 1985

HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

HOSPIRA

5,000 UNITS/100ML

N19339 001

Mar 27, 1985

HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

HOSPIRA

5,000 UNITS/100ML

N18916 006

Jan 31, 1984

HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINERAP BAXTER HLTHCARE200 UNITS/100MLN18609 002

Apr 28, 1982

HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINERAP HOSPIRA200 UNITS/100MLN18916 011

Jun 23, 1989

HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINERAP BAXTER HLTHCARE4,000 UNITS/100MLN18814 001

Oct 31, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 187 (of 360)

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>4,000 UNITS/100ML</u>	<u>N19952</u>	<u>001</u>	Jul 20, 1992
<u>AP</u>	HOSPIRA	<u>4,000 UNITS/100ML</u>	<u>N19805</u>	<u>001</u>	Jan 25, 1989

HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	<u>5,000 UNITS/100ML</u>	<u>N18814</u>	<u>003</u>	Jul 09, 1985
<u>AP</u>		<u>10,000 UNITS/100ML</u>	<u>N18814</u>	<u>004</u>	Jul 02, 1987

HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	<u>5,000 UNITS/100ML</u>	<u>N19952</u>	<u>004</u>	Jul 20, 1992
<u>AP</u>		<u>10,000 UNITS/100ML</u>	<u>N19952</u>	<u>005</u>	Jul 20, 1992
<u>AP</u>	HOSPIRA	<u>5,000 UNITS/100ML</u>	<u>N19339</u>	<u>004</u>	Mar 27, 1985
<u>AP</u>		<u>5,000 UNITS/100ML</u>	<u>N19805</u>	<u>002</u>	Jan 25, 1989
<u>AP</u>		<u>10,000 UNITS/100ML</u>	<u>N19339</u>	<u>002</u>	Mar 27, 1985

HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

	HOSPIRA	5,000 UNITS/100ML	N18916	007	Jan 31, 1984
		10,000 UNITS/100ML	N18916	008	Jan 31, 1984

HEPARIN SODIUM IN PLASTIC CONTAINER

<u>AP</u>	AM PHARM	<u>1,000 UNITS/ML</u>	<u>N17029</u>	<u>013</u>	Dec 05, 1985
<u>AP</u>		<u>5,000 UNITS/ML</u>	<u>N17029</u>	<u>014</u>	Dec 05, 1985
<u>AP</u>		<u>10,000 UNITS/ML</u>	<u>N17029</u>	<u>015</u>	Dec 05, 1985
<u>AP</u>		<u>20,000 UNITS/ML</u>	<u>N17029</u>	<u>016</u>	Dec 05, 1985

HEPARIN SODIUM PRESERVATIVE FREE

<u>AP</u>	+ AM PHARM	<u>1,000 UNITS/ML</u>	<u>N17029</u>	<u>010</u>	Apr 28, 1986
<u>AP</u>	+ HOSPIRA	<u>2,500 UNITS/ML</u>	<u>N05264</u>	<u>014</u>	Apr 07, 1986
<u>AP</u>	+	<u>10,000 UNITS/ML</u>	<u>N89522</u>	<u>001</u>	May 04, 1987
	+	2,000 UNITS/ML	N05264	013	Apr 07, 1986

HEPFLUSH-10

<u>AP</u>	AM PHARM	<u>10 UNITS/ML</u>	<u>N17651</u>	<u>009</u>	Jun 26, 1984
-----------	----------	--------------------	---------------	------------	--------------

HEP-LOCK

<u>AP</u>	+ BAXTER HLTHCARE	<u>10 UNITS/ML</u>	<u>N17037</u>	<u>007</u>	
<u>AP</u>	+	<u>100 UNITS/ML</u>	<u>N17037</u>	<u>006</u>	

HEP-LOCK U/P

<u>AP</u>	BAXTER HLTHCARE	<u>10 UNITS/ML</u>	<u>N17037</u>	<u>010</u>	Jun 10, 1983
<u>AP</u>		<u>100 UNITS/ML</u>	<u>N17037</u>	<u>011</u>	Jun 10, 1983

HEXACHLOROPHENE

EMULSION; TOPICAL

PHISOHEX

	+ SANOFI SYNTHELABO	3%	N06882	001	
--	---------------------	----	--------	-----	--

SPONGE; TOPICAL

PRE-OP

<u>AT</u>	+ DAVIS AND GECK	<u>480MG</u>	<u>N17433</u>	<u>001</u>	
-----------	------------------	--------------	---------------	------------	--

PRE-OP II

<u>AT</u>	DAVIS AND GECK	<u>480MG</u>	<u>N17433</u>	<u>002</u>	
-----------	----------------	--------------	---------------	------------	--

HISTRELIN ACETATE

IMPLANT; SUBCUTANEOUS

VANTAS

	+ VALERA	50MG	N21732	001	Oct 12, 2004
--	----------	------	--------	-----	--------------

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

HYCODAN

<u>AA</u>	+ ENDO PHARMS	<u>1.5MG/5ML;5MG/5ML</u>	<u>N05213</u>	<u>002</u>	Jul 26, 1988
-----------	---------------	--------------------------	---------------	------------	--------------

HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

<u>AA</u>	IVAX PHARMS	<u>1.5MG/5ML;5MG/5ML</u>	<u>N40285</u>	<u>001</u>	Jul 19, 1999
-----------	-------------	--------------------------	---------------	------------	--------------

HYDROCODONE COMPOUND

<u>AA</u>	ALPHARMA US PHARMS	<u>1.5MG/5ML;5MG/5ML</u>	<u>N88017</u>	<u>001</u>	Jul 05, 1983
-----------	--------------------	--------------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 188 (of 360)

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

MYCODONE

<u>AA</u>	MORTON GROVE	<u>1.5MG/5ML; 5MG/5ML</u>	<u>N88008</u>	<u>001</u>	Mar 03, 1983
-----------	--------------	---------------------------	---------------	------------	--------------

TABLET; ORAL

HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE

<u>AA</u>	AMIDE PHARM	<u>1.5MG; 5MG</u>	<u>N40295</u>	<u>001</u>	Dec 01, 2000
-----------	-------------	-------------------	---------------	------------	--------------

HYCODAN

<u>AA</u>	+ ENDO PHARMS	<u>1.5MG; 5MG</u>	<u>N05213</u>	<u>001</u>	Jul 26, 1988
-----------	---------------	-------------------	---------------	------------	--------------

TUSSIGON

<u>AA</u>	JONES PHARMA	<u>1.5MG; 5MG</u>	<u>N88508</u>	<u>001</u>	Jul 30, 1985
-----------	--------------	-------------------	---------------	------------	--------------

HYALURONIDASE

INJECTABLE; INJECTION

AMPHADASE

+	AMPHASTAR PHARM	150 UNITS/ML	N21665	001	Oct 26, 2004
---	-----------------	--------------	--------	-----	--------------

HYDASE

+	PRIMAPHARM	150 UNITS/ML	N21716	001	Oct 25, 2005
---	------------	--------------	--------	-----	--------------

VITRASE

	ISTA PHARMS	200 UNITS/VIAL	N21640	002	Dec 02, 2004
--	-------------	----------------	--------	-----	--------------

+		6,200 UNITS/VIAL	N21640	001	May 05, 2004
---	--	------------------	--------	-----	--------------

HYALURONIDASE RECOMBINANT HUMAN

INJECTABLE; INJECTION

HYLENEX RECOMBINANT

+	HALOZYME THERAP	150 UNITS/ML	N21859	001	Dec 02, 2005
---	-----------------	--------------	--------	-----	--------------

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE

<u>AP</u>	AM PHARM PARTNERS	<u>20MG/ML</u>	<u>N40388</u>	<u>001</u>	Mar 13, 2001
-----------	-------------------	----------------	---------------	------------	--------------

<u>AP</u>	+ LUITPOLD	<u>20MG/ML</u>	<u>N40136</u>	<u>001</u>	Jun 30, 1997
-----------	------------	----------------	---------------	------------	--------------

<u>AP</u>	SICOR PHARMS	<u>20MG/ML</u>	<u>N40373</u>	<u>001</u>	Feb 23, 2000
-----------	--------------	----------------	---------------	------------	--------------

TABLET; ORAL

APRESOLINE

<u>AA</u>	+ NOVARTIS	<u>10MG</u>	<u>N08303</u>	<u>004</u>	
-----------	------------	-------------	---------------	------------	--

<u>AA</u>	+	<u>25MG</u>	<u>N08303</u>	<u>001</u>	
-----------	---	-------------	---------------	------------	--

<u>AA</u>	+	<u>50MG</u>	<u>N08303</u>	<u>002</u>	
-----------	---	-------------	---------------	------------	--

<u>AA</u>	+	<u>100MG</u>	<u>N08303</u>	<u>005</u>	
-----------	---	--------------	---------------	------------	--

HYDRALAZINE HYDROCHLORIDE

<u>AA</u>	MUTUAL PHARM	<u>10MG</u>	<u>N89359</u>	<u>001</u>	Jul 25, 1986
-----------	--------------	-------------	---------------	------------	--------------

<u>AA</u>		<u>25MG</u>	<u>N89258</u>	<u>001</u>	May 05, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>50MG</u>	<u>N89259</u>	<u>001</u>	May 05, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>	PAR PHARM	<u>10MG</u>	<u>N87836</u>	<u>001</u>	Oct 05, 1982
-----------	-----------	-------------	---------------	------------	--------------

<u>AA</u>		<u>25MG</u>	<u>N86961</u>	<u>002</u>	
-----------	--	-------------	---------------	------------	--

<u>AA</u>		<u>50MG</u>	<u>N86962</u>	<u>001</u>	
-----------	--	-------------	---------------	------------	--

<u>AA</u>		<u>100MG</u>	<u>N88391</u>	<u>001</u>	Sep 27, 1983
-----------	--	--------------	---------------	------------	--------------

<u>AA</u>	PLIVA	<u>10MG</u>	<u>N89097</u>	<u>001</u>	Dec 18, 1985
-----------	-------	-------------	---------------	------------	--------------

<u>AA</u>		<u>25MG</u>	<u>N88467</u>	<u>001</u>	May 01, 1984
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>50MG</u>	<u>N88468</u>	<u>001</u>	May 01, 1984
-----------	--	-------------	---------------	------------	--------------

<u>AA</u>		<u>100MG</u>	<u>N89098</u>	<u>001</u>	Dec 18, 1985
-----------	--	--------------	---------------	------------	--------------

<u>AA</u>	SANDOZ	<u>10MG</u>	<u>N83241</u>	<u>001</u>	
-----------	--------	-------------	---------------	------------	--

<u>AA</u>		<u>25MG</u>	<u>N83560</u>	<u>001</u>	
-----------	--	-------------	---------------	------------	--

<u>AA</u>		<u>50MG</u>	<u>N83561</u>	<u>001</u>	
-----------	--	-------------	---------------	------------	--

<u>AA</u>	WATSON LABS	<u>25MG</u>	<u>N84504</u>	<u>001</u>	
-----------	-------------	-------------	---------------	------------	--

<u>AA</u>		<u>50MG</u>	<u>N84503</u>	<u>001</u>	
-----------	--	-------------	---------------	------------	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 189 (of 360)

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

APRESAZIDE

<u>AB</u>	NOVARTIS	<u>25MG;25MG</u>	<u>N84735</u>	<u>001</u>	
<u>AB</u>		<u>50MG;50MG</u>	<u>N84810</u>	<u>001</u>	

HYDRA-ZIDE

<u>AB</u>	PAR PHARM	<u>25MG;25MG</u>	<u>N88957</u>	<u>001</u>	Oct 21, 1985
<u>AB</u>		<u>50MG;50MG</u>	<u>N88946</u>	<u>001</u>	Oct 21, 1985
	+	100MG;50MG	N88961	001	Oct 21, 1985

HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE

TABLET; ORAL

BIDIL

	+	NITROMED	37.5MG;20MG	N20727	001	Jun 23, 2005
--	---	----------	-------------	--------	-----	--------------

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

<u>AB</u>	IVAX PHARMS	<u>12.5MG</u>	<u>N77005</u>	<u>001</u>	Jul 13, 2005
<u>AB</u>	MYLAN	<u>12.5MG</u>	<u>N75640</u>	<u>001</u>	Jan 28, 2000
<u>AB</u>	VINTAGE PHARMS	<u>12.5MG</u>	<u>N75907</u>	<u>001</u>	Sep 17, 2002

MICROZIDE

<u>AB</u>	+	WATSON LABS	<u>12.5MG</u>	<u>N20504</u>	<u>001</u>	Dec 27, 1996
-----------	---	-------------	---------------	---------------	------------	--------------

SOLUTION; ORAL

HYDROCHLOROTHIAZIDE

	+	ROXANE	50MG/5ML	N88587	001	Jul 02, 1984
--	---	--------	----------	--------	-----	--------------

TABLET; ORAL

ESIDRIX

<u>AB</u>	NOVARTIS	<u>25MG</u>	<u>N11793</u>	<u>005</u>	
<u>AB</u>		<u>50MG</u>	<u>N11793</u>	<u>008</u>	

HYDROCHLOROTHIAZIDE

<u>AB</u>	CLONMEL HLTHCARE	<u>25MG</u>	<u>N87059</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>N87068</u>	<u>001</u>	
<u>AB</u>	IVAX PHARMS	<u>25MG</u>	<u>N83177</u>	<u>001</u>	
<u>AB</u>	+	<u>50MG</u>	<u>N83177</u>	<u>002</u>	
<u>AB</u>	PHARM FORM	<u>25MG</u>	<u>N85181</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>N85182</u>	<u>001</u>	
<u>AB</u>	PUREPAC PHARM	<u>25MG</u>	<u>N85054</u>	<u>002</u>	
<u>AB</u>		<u>50MG</u>	<u>N85208</u>	<u>001</u>	
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>N87565</u>	<u>001</u>	Mar 09, 1982
<u>AB</u>		<u>50MG</u>	<u>N84912</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>N85219</u>	<u>001</u>	
<u>AB</u>	VINTAGE PHARMS	<u>25MG</u>	<u>N40412</u>	<u>001</u>	Mar 29, 2002
<u>AB</u>		<u>50MG</u>	<u>N40412</u>	<u>002</u>	Mar 29, 2002
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>N81189</u>	<u>001</u>	Jan 24, 1992

ORETIC

<u>AB</u>	ABBOTT	<u>50MG</u>	<u>N11971</u>	<u>002</u>	
-----------	--------	-------------	---------------	------------	--

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

	SANOFI SYNTHELABO	12.5MG;150MG	N20758	002	Sep 30, 1997
		12.5MG;300MG	N20758	003	Aug 31, 1998
	+	25MG;300MG	N20758	004	Mar 15, 2005

HYDROCHLOROTHIAZIDE; LISINOPRIL

TABLET; ORAL

LISINOPRIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	APOTEX	<u>12.5MG;10MG</u>	<u>N76674</u>	<u>001</u>	Oct 05, 2004
-----------	--------	--------------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 190 (of 360)

HYDROCHLOROTHIAZIDE; LISINOPRIL

TABLET; ORAL

LISINOPRIL AND HYDROCHLOROTHIAZIDE

<u>AB</u>	APOTEX	<u>12.5MG;20MG</u>	<u>N76674</u>	<u>002</u>	Oct 05, 2004
<u>AB</u>		<u>25MG;20MG</u>	<u>N76674</u>	<u>003</u>	Oct 05, 2004
<u>AB</u>	IVAX PHARMS	<u>12.5MG;10MG</u>	<u>N75776</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N75776</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N75776</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>	MYLAN	<u>12.5MG;10MG</u>	<u>N76113</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N76113</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N76113</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>	PUREPAC PHARM	<u>12.5MG;10MG</u>	<u>N76230</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N76230</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N76230</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>	RANBAXY	<u>12.5MG;10MG</u>	<u>N76007</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N76007</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N76007</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>	SANDOZ	<u>12.5MG;10MG</u>	<u>N75926</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;10MG</u>	<u>N76262</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N75926</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N76262</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N75926</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N76262</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>	TEVA	<u>12.5MG;10MG</u>	<u>N75869</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N75869</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N75869</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>12.5MG;10MG</u>	<u>N76194</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N76194</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N76194</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>	WEST WARD	<u>12.5MG;10MG</u>	<u>N76265</u>	<u>001</u>	Jul 08, 2002
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N76265</u>	<u>002</u>	Jul 08, 2002
<u>AB</u>		<u>25MG;20MG</u>	<u>N76265</u>	<u>003</u>	Jul 08, 2002
	<u>PRINZIDE</u>				
<u>AB</u>	MERCK	<u>12.5MG;10MG</u>	<u>N19778</u>	<u>003</u>	Nov 18, 1993
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N19778</u>	<u>001</u>	Feb 16, 1989
<u>AB</u>	+	<u>25MG;20MG</u>	<u>N19778</u>	<u>002</u>	Feb 16, 1989
	<u>ZESTORETIC</u>				
<u>AB</u>	ASTRAZENECA	<u>12.5MG;10MG</u>	<u>N19888</u>	<u>003</u>	Nov 18, 1993
<u>AB</u>		<u>12.5MG;20MG</u>	<u>N19888</u>	<u>001</u>	Sep 20, 1990
<u>AB</u>		<u>25MG;20MG</u>	<u>N19888</u>	<u>002</u>	Jul 20, 1989

HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM

TABLET; ORAL

HYZAAR

MERCK

12.5MG;50MG

N20387 001

Apr 28, 1995

12.5MG;100MG

N20387 003

Oct 20, 2005

+

25MG;100MG

N20387 002

Nov 10, 1998

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDORIL 15

<u>AB</u>	MERCK	<u>15MG;250MG</u>	<u>N13402</u>	<u>001</u>
-----------	-------	-------------------	---------------	------------

ALDORIL 25

<u>AB</u>	MERCK	<u>25MG;250MG</u>	<u>N13402</u>	<u>002</u>
-----------	-------	-------------------	---------------	------------

ALDORIL D30

<u>AB</u>	MERCK	<u>30MG;500MG</u>	<u>N13402</u>	<u>003</u>
-----------	-------	-------------------	---------------	------------

ALDORIL D50

<u>AB</u>	+	MERCK	<u>50MG;500MG</u>	<u>N13402</u>	<u>004</u>
-----------	---	-------	-------------------	---------------	------------

METHYLDOPA AND HYDROCHLOROTHIAZIDE

<u>AB</u>	MYLAN	<u>15MG;250MG</u>	<u>N70264</u>	<u>001</u>	Jan 23, 1986
-----------	-------	-------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 191 (of 360)

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

<u>AB</u>	MYLAN	<u>25MG;250MG</u>	<u>N70265</u>	<u>001</u>	Jan 23, 1986
<u>AB</u>	PAR PHARM	<u>15MG;250MG</u>	<u>N70616</u>	<u>001</u>	Feb 02, 1987
<u>AB</u>		<u>25MG;250MG</u>	<u>N70612</u>	<u>001</u>	Feb 02, 1987
<u>AB</u>		<u>30MG;500MG</u>	<u>N70613</u>	<u>001</u>	Feb 02, 1987
<u>AB</u>		<u>50MG;500MG</u>	<u>N70614</u>	<u>001</u>	Feb 02, 1987
<u>AB</u>	SANDOZ	<u>15MG;250MG</u>	<u>N70182</u>	<u>001</u>	Jan 15, 1986
<u>AB</u>		<u>25MG;250MG</u>	<u>N70183</u>	<u>001</u>	Jan 15, 1986
<u>AB</u>		<u>30MG;500MG</u>	<u>N70543</u>	<u>001</u>	Jan 15, 1986
<u>AB</u>		<u>50MG;500MG</u>	<u>N70544</u>	<u>001</u>	Jan 15, 1986
<u>AB</u>	WATSON LABS	<u>15MG;250MG</u>	<u>N70958</u>	<u>001</u>	Feb 06, 1989
<u>AB</u>		<u>25MG;250MG</u>	<u>N70959</u>	<u>001</u>	Jan 19, 1989
<u>AB</u>		<u>30MG;500MG</u>	<u>N71069</u>	<u>001</u>	Jan 19, 1989
<u>AB</u>		<u>50MG;500MG</u>	<u>N70960</u>	<u>001</u>	Feb 06, 1989

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

LOPRESSOR HCT

<u>AB</u>	NOVARTIS	<u>25MG;50MG</u>	<u>N18303</u>	<u>001</u>	Dec 31, 1984
<u>AB</u>		<u>25MG;100MG</u>	<u>N18303</u>	<u>002</u>	Dec 31, 1984
<u>AB</u>	+	<u>50MG;100MG</u>	<u>N18303</u>	<u>003</u>	Dec 31, 1984
	<u>METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE</u>				
<u>AB</u>	MYLAN	<u>25MG;50MG</u>	<u>N76792</u>	<u>001</u>	Aug 20, 2004
<u>AB</u>		<u>25MG;100MG</u>	<u>N76792</u>	<u>002</u>	Aug 20, 2004
<u>AB</u>		<u>50MG;100MG</u>	<u>N76792</u>	<u>003</u>	Aug 20, 2004

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

UNIRETIC

	SCHWARZ PHARMA	12.5MG;7.5MG	N20729	001	Jun 27, 1997
		12.5MG;15MG	N20729	003	Feb 14, 2002
	+	25MG;15MG	N20729	002	Jun 27, 1997

HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR HCT

	SANKYO	12.5MG;20MG	N21532	002	Jun 05, 2003
		12.5MG;40MG	N21532	003	Jun 05, 2003
	+	25MG;40MG	N21532	005	Jun 05, 2003

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

INDERIDE-40/25

<u>AB</u>	+	WYETH PHARMS INC	<u>25MG;40MG</u>	<u>N18031</u>	<u>001</u>	
	<u>PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE</u>					
<u>AB</u>		BARR	<u>25MG;40MG</u>	<u>N70704</u>	<u>001</u>	Oct 01, 1986
<u>AB</u>			<u>25MG;80MG</u>	<u>N70705</u>	<u>001</u>	Oct 01, 1986
<u>AB</u>		MYLAN	<u>25MG;40MG</u>	<u>N70946</u>	<u>001</u>	Mar 04, 1987
<u>AB</u>			<u>25MG;80MG</u>	<u>N70947</u>	<u>001</u>	Apr 01, 1987
<u>AB</u>		PLIVA	<u>25MG;40MG</u>	<u>N72042</u>	<u>001</u>	Mar 14, 1988
<u>AB</u>			<u>25MG;80MG</u>	<u>N72043</u>	<u>001</u>	Mar 14, 1988
<u>AB</u>		PUREPAC PHARM	<u>25MG;40MG</u>	<u>N70851</u>	<u>001</u>	May 15, 1986
<u>AB</u>			<u>25MG;80MG</u>	<u>N70852</u>	<u>001</u>	May 15, 1986
<u>AB</u>		SANDOZ	<u>25MG;40MG</u>	<u>N71060</u>	<u>001</u>	Aug 26, 1987
<u>AB</u>			<u>25MG;80MG</u>	<u>N71061</u>	<u>001</u>	Aug 26, 1987
<u>AB</u>		WATSON LABS	<u>25MG;40MG</u>	<u>N70301</u>	<u>001</u>	Apr 18, 1986
<u>AB</u>			<u>25MG;80MG</u>	<u>N70305</u>	<u>001</u>	Apr 18, 1986

PRESCRIPTION DRUG PRODUCT LIST

3 - 192 (of 360)

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL				
<u>ACCURETIC</u>				
<u>AB</u>	PFIZER PHARMS	<u>12.5MG;EQ 10MG BASE</u>	<u>N20125</u>	<u>001</u> Dec 28, 1999
<u>AB</u>		<u>12.5MG;EQ 20MG BASE</u>	<u>N20125</u>	<u>002</u> Dec 28, 1999
<u>AB</u>	+	<u>25MG;EQ 20MG BASE</u>	<u>N20125</u>	<u>003</u> Dec 28, 1999
<u>QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE</u>				
<u>AB</u>	MYLAN	<u>12.5MG;EQ 10MG BASE</u>	<u>N77093</u>	<u>001</u> Mar 28, 2005
<u>AB</u>		<u>12.5MG;EQ 20MG BASE</u>	<u>N77093</u>	<u>002</u> Mar 28, 2005
<u>AB</u>		<u>25MG;EQ 20MG BASE</u>	<u>N77093</u>	<u>003</u> Mar 28, 2005
<u>QUINARETIC</u>				
<u>AB</u>	AMIDE PHARM	<u>12.5MG;EQ 10MG BASE</u>	<u>N76374</u>	<u>001</u> Mar 31, 2004
<u>AB</u>		<u>12.5MG;EQ 20MG BASE</u>	<u>N76374</u>	<u>002</u> Mar 31, 2004
<u>AB</u>		<u>25MG;EQ 20MG BASE</u>	<u>N76374</u>	<u>003</u> Mar 31, 2004

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL				
<u>ALDACTAZIDE</u>				
<u>AB</u>	GD SEARLE LLC	<u>25MG;25MG</u>	<u>N12616</u>	<u>004</u> Dec 30, 1982
	+	<u>50MG;50MG</u>	<u>N12616</u>	<u>005</u> Dec 30, 1982
<u>SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE</u>				
<u>AB</u>	MUTUAL PHARM	<u>25MG;25MG</u>	<u>N87267</u>	<u>001</u>
<u>AB</u>		<u>25MG;25MG</u>	<u>N89534</u>	<u>001</u> Jul 02, 1987
<u>AB</u>	MYLAN	<u>25MG;25MG</u>	<u>N86513</u>	<u>001</u>
<u>AB</u>	SANDOZ	<u>25MG;25MG</u>	<u>N86881</u>	<u>001</u>
<u>AB</u>	WATSON LABS	<u>25MG;25MG</u>	<u>N87398</u>	<u>001</u>

HYDROCHLOROTHIAZIDE; TELMISARTAN

TABLET; ORAL				
MICARDIS HCT				
	BOEHRINGER INGELHEIM	12.5MG;40MG	N21162	001 Nov 17, 2000
		12.5MG;80MG	N21162	002 Nov 17, 2000
	+	25MG;80MG	N21162	003 Apr 19, 2004

HYDROCHLOROTHIAZIDE; TIMOLOL MALEATE

TABLET; ORAL				
TIMOLIDE 10-25				
	+	MERCK 25MG;10MG	N18061	001

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL				
<u>DYAZIDE</u>				
<u>AB</u>	+	GLAXOSMITHKLINE <u>25MG;37.5MG</u>	<u>N16042</u>	<u>003</u> Mar 03, 1994
<u>TRIAMTERENE AND HYDROCHLOROTHIAZIDE</u>				
<u>AB</u>	BARR	<u>25MG;37.5MG</u>	<u>N74970</u>	<u>001</u> Jan 06, 1998
<u>AB</u>	DURAMED PHARMS BARR	<u>25MG;37.5MG</u>	<u>N75052</u>	<u>001</u> Jun 18, 1999
<u>AB</u>	IVAX PHARMS	<u>25MG;50MG</u>	<u>N74259</u>	<u>001</u> Mar 30, 1995
<u>AB</u>	MYLAN	<u>25MG;37.5MG</u>	<u>N74701</u>	<u>001</u> Jun 07, 1996
<u>AB</u>	SANDOZ	<u>25MG;37.5MG</u>	<u>N74821</u>	<u>001</u> Jun 05, 1997
<u>AB</u>	+	<u>25MG;50MG</u>	<u>N73191</u>	<u>001</u> Jul 31, 1991
TABLET; ORAL				
<u>MAXZIDE</u>				
<u>AB</u>	+	MYLAN BERTEK <u>50MG;75MG</u>	<u>N19129</u>	<u>001</u> Oct 22, 1984
<u>MAXZIDE-25</u>				
<u>AB</u>	MYLAN BERTEK	<u>25MG;37.5MG</u>	<u>N19129</u>	<u>003</u> May 13, 1988
<u>TRIAMTERENE AND HYDROCHLOROTHIAZIDE</u>				
<u>AB</u>	BARR	<u>25MG;37.5MG</u>	<u>N71251</u>	<u>002</u> May 05, 1998
<u>AB</u>		<u>50MG;75MG</u>	<u>N71251</u>	<u>001</u> Apr 17, 1988
<u>AB</u>	PLIVA	<u>25MG;37.5MG</u>	<u>N74026</u>	<u>001</u> Apr 26, 1996
<u>AB</u>		<u>50MG;75MG</u>	<u>N73467</u>	<u>001</u> Jan 31, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 193 (of 360)

HYDROCHLOROTHIAZIDE; TRIAMTERENE

TABLET; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

<u>AB</u>	SANDOZ	<u>25MG;37.5MG</u>	<u>N73281</u>	<u>001</u>	Apr 30, 1992
<u>AB</u>		<u>50MG;75MG</u>	<u>N72011</u>	<u>001</u>	Jun 17, 1988
<u>AB</u>	WATSON LABS	<u>25MG;37.5MG</u>	<u>N73449</u>	<u>001</u>	Sep 23, 1993
<u>AB</u>		<u>50MG;75MG</u>	<u>N71851</u>	<u>001</u>	Nov 30, 1988
<u>AB</u>		<u>50MG;75MG</u>	<u>N71969</u>	<u>001</u>	Apr 17, 1988

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

DIOVAN HCT

NOVARTIS

	12.5MG;80MG	N20818	001	Mar 06, 1998
	12.5MG;160MG	N20818	002	Mar 06, 1998
+	25MG;160MG	N20818	003	Jan 17, 2002

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN

<u>AB</u>	ANDRX PHARMS	<u>7.5MG;200MG</u>	<u>N76604</u>	<u>001</u>	Dec 31, 2003
<u>AB</u>	INTERPHARM	<u>7.5MG;200MG</u>	<u>N76642</u>	<u>001</u>	Oct 12, 2004
	+	5MG;200MG	N76642	002	Mar 18, 2004
<u>AB</u>	TEVA	<u>7.5MG;200MG</u>	<u>N76023</u>	<u>001</u>	Apr 11, 2003
	<u>VICOPROFEN</u>				
<u>AB</u>	+ ABBOTT	<u>7.5MG;200MG</u>	<u>N20716</u>	<u>001</u>	Sep 23, 1997

HYDROCORTISONE

CREAM; TOPICAL

ALA-CORT

<u>AT</u>	DEL RAY LABS	<u>1%</u>	<u>N80706</u>	<u>006</u>	
-----------	--------------	-----------	---------------	------------	--

ANUSOL HC

<u>AT</u>	SALIX PHARMS	<u>2.5%</u>	<u>N88250</u>	<u>001</u>	Jun 06, 1984
-----------	--------------	-------------	---------------	------------	--------------

HI-COR

<u>AT</u>	C AND M PHARMA	<u>2.5%</u>	<u>N80483</u>	<u>001</u>	
-----------	----------------	-------------	---------------	------------	--

HYDROCORTISONE

<u>AT</u>	ALPHARMA US PHARMS	<u>1%</u>	<u>N87795</u>	<u>001</u>	May 03, 1983
<u>AT</u>		<u>2.5%</u>	<u>N89682</u>	<u>001</u>	Mar 10, 1988
<u>AT</u>	CLAY PARK	<u>2.5%</u>	<u>N85025</u>	<u>001</u>	
<u>AT</u>	EVERYLIFE	<u>1%</u>	<u>N80452</u>	<u>002</u>	
<u>AT</u>	FOUGERA	<u>1%</u>	<u>N80693</u>	<u>003</u>	
<u>AT</u>		<u>2.5%</u>	<u>N89414</u>	<u>001</u>	Dec 16, 1986
<u>AT</u>	INGRAM PHARM	<u>0.5%</u>	<u>N80456</u>	<u>002</u>	
<u>AT</u>		<u>1%</u>	<u>N80456</u>	<u>003</u>	
<u>AT</u>	TARO	<u>1%</u>	<u>N86155</u>	<u>001</u>	
<u>AT</u>		<u>2.5%</u>	<u>N88799</u>	<u>001</u>	Nov 09, 1984
<u>AT</u>	VINTAGE PHARMS	<u>2.5%</u>	<u>N40503</u>	<u>001</u>	Mar 12, 2004

HYTONE

<u>AT</u>	+ DERMIK LABS	<u>1%</u>	<u>N80472</u>	<u>003</u>	
-----------	---------------	-----------	---------------	------------	--

<u>AT</u>	+	<u>2.5%</u>	<u>N80472</u>	<u>004</u>	
-----------	---	-------------	---------------	------------	--

PENECORT

<u>AT</u>	ALLERGAN HERBERT	<u>1%</u>	<u>N88216</u>	<u>001</u>	Jun 06, 1984
-----------	------------------	-----------	---------------	------------	--------------

SYNACORT

<u>AT</u>	MEDICIS	<u>1%</u>	<u>N87458</u>	<u>001</u>	
-----------	---------	-----------	---------------	------------	--

<u>AT</u>		<u>2.5%</u>	<u>N87457</u>	<u>001</u>	
-----------	--	-------------	---------------	------------	--

ENEMA; RECTAL

COLOCORT

<u>AB</u>	PADDOCK	<u>100MG/60ML</u>	<u>N75172</u>	<u>001</u>	Dec 03, 1999
-----------	---------	-------------------	---------------	------------	--------------

CORTENEMA

<u>AB</u>	+ SOLVAY	<u>100MG/60ML</u>	<u>N16199</u>	<u>001</u>	
-----------	----------	-------------------	---------------	------------	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 194 (of 360)

HYDROCORTISONE

ENEMA; RECTAL

HYDROCORTISONE

<u>AB</u>	TEVA PHARMS	<u>100MG/60ML</u>	<u>N74171</u>	<u>001</u>	May 27, 1994
-----------	-------------	-------------------	---------------	------------	--------------

LOTION; TOPICAL

ALA-CORT

<u>AT</u>	DEL RAY LABS	<u>1%</u>	<u>N83201</u>	<u>001</u>	
-----------	--------------	-----------	---------------	------------	--

ALA-SCALP

	DEL RAY LABS	2%	N83231	001	
--	--------------	----	--------	-----	--

CETACORT

<u>AT</u>	HEALTHPOINT	<u>0.5%</u>	<u>N80426</u>	<u>002</u>	
-----------	-------------	-------------	---------------	------------	--

<u>AT</u>		<u>1%</u>	<u>N80426</u>	<u>001</u>	
-----------	--	-----------	---------------	------------	--

HYDROCORTISONE

<u>AT</u>	ALTANA	<u>2.5%</u>	<u>N40351</u>	<u>001</u>	Jul 25, 2000
-----------	--------	-------------	---------------	------------	--------------

<u>AT</u>	TARO	<u>2.5%</u>	<u>N40247</u>	<u>001</u>	Jul 23, 1999
-----------	------	-------------	---------------	------------	--------------

<u>AT</u>	VINTAGE PHARMS	<u>2.5%</u>	<u>N40417</u>	<u>001</u>	Jul 30, 2003
-----------	----------------	-------------	---------------	------------	--------------

HYTONE

<u>AT</u>	+ DERMIK LABS	<u>1%</u>	<u>N80473</u>	<u>003</u>	
-----------	---------------	-----------	---------------	------------	--

<u>AT</u>	+	<u>2.5%</u>	<u>N80473</u>	<u>004</u>	Nov 30, 1982
-----------	---	-------------	---------------	------------	--------------

NUTRACORT

<u>AT</u>	HEALTHPOINT	<u>1%</u>	<u>N80443</u>	<u>003</u>	
-----------	-------------	-----------	---------------	------------	--

<u>AT</u>		<u>2.5%</u>	<u>N87644</u>	<u>001</u>	Aug 24, 1982
-----------	--	-------------	---------------	------------	--------------

STIE-CORT

<u>AT</u>	STIEFEL	<u>1%</u>	<u>N89066</u>	<u>001</u>	Nov 25, 1985
-----------	---------	-----------	---------------	------------	--------------

<u>AT</u>		<u>2.5%</u>	<u>N89074</u>	<u>001</u>	Nov 26, 1985
-----------	--	-------------	---------------	------------	--------------

OINTMENT; TOPICAL

HYDROCORTISONE

<u>AT</u>	ALPHARMA US PHARMS	<u>1%</u>	<u>N87796</u>	<u>001</u>	Oct 13, 1982
-----------	--------------------	-----------	---------------	------------	--------------

<u>AT</u>	ALTANA	<u>1%</u>	<u>N80692</u>	<u>001</u>	
-----------	--------	-----------	---------------	------------	--

<u>AT</u>	CLAY PARK	<u>2.5%</u>	<u>N85027</u>	<u>001</u>	
-----------	-----------	-------------	---------------	------------	--

<u>AT</u>	FOUGERA	<u>2.5%</u>	<u>N81203</u>	<u>001</u>	May 28, 1993
-----------	---------	-------------	---------------	------------	--------------

<u>AT</u>	TARO	<u>1%</u>	<u>N86257</u>	<u>001</u>	
-----------	------	-----------	---------------	------------	--

<u>AT</u>		<u>2.5%</u>	<u>N40310</u>	<u>001</u>	Dec 29, 2000
-----------	--	-------------	---------------	------------	--------------

HYDROCORTISONE IN ABSORBASE

<u>AT</u>	CAROLINA MEDCL	<u>1%</u>	<u>N88138</u>	<u>001</u>	Sep 06, 1985
-----------	----------------	-----------	---------------	------------	--------------

POWDER; FOR RX COMPOUNDING

HYDROCORTISONE

	+ PHARMA TEK	100%	N85982	001	
--	--------------	------	--------	-----	--

SOLUTION; TOPICAL

PENECORT

<u>AT</u>	ALLERGAN HERBERT	<u>1%</u>	<u>N88214</u>	<u>001</u>	Jun 06, 1984
-----------	------------------	-----------	---------------	------------	--------------

TEXACORT

<u>AT</u>	+ SIRIUS LABS	<u>1%</u>	<u>N80425</u>	<u>001</u>	
-----------	---------------	-----------	---------------	------------	--

	+	2.5%	N81271	001	Apr 17, 1992
--	---	------	--------	-----	--------------

TABLET; ORAL

CORTEF

BP	PHARMACIA AND UPJOHN	10MG	N08697	001	
----	----------------------	------	--------	-----	--

BP	+	20MG	N08697	002	
----	---	------	--------	-----	--

		5MG	N08697	003	
--	--	-----	--------	-----	--

HYDROCORTISONE

BP	WEST WARD	20MG	N83365	001	
----	-----------	------	--------	-----	--

HYDROCORTONE

BP	MERCK	10MG	N08506	007	
----	-------	------	--------	-----	--

BP	+	20MG	N08506	011	
----	---	------	--------	-----	--

HYDROCORTISONE ACETATE

AEROSOL, METERED; RECTAL

CORTIFOAM

	+ SCHWARZ PHARMA	10%	N17351	001	Feb 10, 1982
--	------------------	-----	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 195 (of 360)

HYDROCORTISONE ACETATE

CREAM; TOPICAL

HYDROCORTISONE ACETATE

+ FERNDAL LABS 2.5% N40259 001 Jul 29, 1999

MICORT-HC

+ FERNDAL LABS 2% N40398 001 Mar 29, 2002

2.5% N40396 001 Feb 27, 2001

LOTION; TOPICAL

DRICORT

+ INGRAM PHARM 0.5% N86207 001

PASTE; TOPICAL

ORABASE HCA

COLGATE 0.5% N83205 001

POWDER; FOR RX COMPOUNDING

HYDROCORTISONE ACETATE

PHARMA TEK 100% N85981 001

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL

CORTISPORIN

+ MONARCH PHARMS 0.5%;EQ 3.5MG BASE/GM;10,000 UNITS/GM N50218 001 Aug 09, 1985

HYDROCORTISONE ACETATE; OXYTETRACYCLINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

TERRA-CORTIL

+ PFIZER 1.5%;EQ 5MG BASE/ML N61016 001

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

AEROSOL, METERED; TOPICAL

EPIFOAM

BX SCHWARZ PHARMA 1%;1% N86457 001

HYDROCORTISONE ACETATE 1% AND PRAMOXINE HYDROCHLORIDE 1%

BX BOCA PHARMA 1%;1% N89440 001 May 17, 1988

PROCTOFOAM HC

BX SCHWARZ PHARMA 1%;1% N86195 001

CREAM; TOPICAL

PRAMOSONE

FERNDAL LABS 0.5%;1% N83778 001

1%;1% N85368 001

LOTION; TOPICAL

PRAMOSONE

FERNDAL LABS 1%;1% N85980 001

2.5%;1% N85979 001

HYDROCORTISONE ACETATE; UREA

CREAM; TOPICAL

CARMOL HCAT + KENWOOD LABS 1%;10% N80505 001U-CORTAT TARO 1%;10% N89472 001 Jun 13, 1988HYDROCORTISONE BUTYRATE

CREAM; TOPICAL

HYDROCORTISONE BUTYRATEAB TARO PHARM INDS 0.1% N76654 001 Aug 03, 2005LOCOIDAB + FERNDAL LABS 0.1% N18514 001 Mar 31, 1982

LOCOID LIPOCREAM

+ FERNDAL LABS 0.1% N20769 001 Sep 08, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 196 (of 360)

HYDROCORTISONE BUTYRATE

OINTMENT; TOPICAL

HYDROCORTISONE BUTYRATE

<u>AB</u>	TARO	<u>0.1%</u>	<u>N76842</u>	<u>001</u>	Dec 27, 2004
	<u>LOCOID</u>				
<u>AB</u>	+ FERNDAL LABS	<u>0.1%</u>	<u>N18652</u>	<u>001</u>	Oct 29, 1982

SOLUTION; TOPICAL

HYDROCORTISONE BUTYRATE

<u>AT</u>	TARO PHARM INDS	<u>0.1%</u>	<u>N76364</u>	<u>001</u>	Jan 14, 2004
	<u>LOCOID</u>				
<u>AT</u>	+ FERNDAL LABS	<u>0.1%</u>	<u>N19116</u>	<u>001</u>	Feb 25, 1987

HYDROCORTISONE PROBUTATE

CREAM; TOPICAL

PANDEL

	+ SAVAGE LABS	0.1%	N20453	001	Feb 28, 1997
--	---------------	------	--------	-----	--------------

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

A-HYDROCORT

<u>AP</u>	HOSPIRA	<u>EQ 100MG BASE/VIAL</u>	<u>N85929</u>	<u>001</u>	
<u>AP</u>		<u>EQ 250MG BASE/VIAL</u>	<u>N85930</u>	<u>001</u>	
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>N85931</u>	<u>001</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N85932</u>	<u>001</u>	

SOLU-CORTEF

<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>EQ 100MG BASE/VIAL</u>	<u>N09866</u>	<u>001</u>	
<u>AP</u>	+	<u>EQ 250MG BASE/VIAL</u>	<u>N09866</u>	<u>002</u>	
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>N09866</u>	<u>003</u>	
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N09866</u>	<u>004</u>	

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE

<u>AB</u>	CLAY PARK	<u>0.2%</u>	<u>N75666</u>	<u>001</u>	May 24, 2000
<u>AB</u>	TARO	<u>0.2%</u>	<u>N75042</u>	<u>001</u>	Aug 25, 1998
<u>AB</u>	TEVA PHARMS	<u>0.2%</u>	<u>N74489</u>	<u>001</u>	Aug 12, 1998
	<u>WESTCORT</u>				
<u>AB</u>	+ WESTWOOD SQUIBB	<u>0.2%</u>	<u>N17950</u>	<u>001</u>	

OINTMENT; TOPICAL

HYDROCORTISONE VALERATE

<u>AB</u>	ALTANA	<u>0.2%</u>	<u>N75085</u>	<u>001</u>	Jul 31, 2001
<u>AB</u>	TARO	<u>0.2%</u>	<u>N75043</u>	<u>001</u>	Aug 25, 1998
	<u>WESTCORT</u>				
<u>AB</u>	+ WESTWOOD SQUIBB	<u>0.2%</u>	<u>N18726</u>	<u>001</u>	Aug 08, 1983

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

CORTISPORIN

<u>AT</u>	+ MONARCH PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N50479</u>	<u>001</u>	
	<u>NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE</u>				
<u>AT</u>	ALCON	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N62423</u>	<u>001</u>	Aug 25, 1983
<u>AT</u>	BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N64053</u>	<u>001</u>	Dec 29, 1995
<u>AT</u>	PHARMAFORCE	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N65216</u>	<u>001</u>	Oct 31, 2005

SUSPENSION/DROPS; OPHTHALMIC

CORTISPORIN

	+ MONARCH PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	N50169	001	
	<u>NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE</u>				
<u>AT</u>	ALCON UNIVERSAL	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N62874</u>	<u>001</u>	May 11, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 197 (of 360)

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SUSPENSION/DROPS; OTIC

CORTISPORIN

<u>AT</u>	+	MONARCH PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N60613</u>	<u>001</u>	
<u>NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE</u>						
<u>AT</u>		ALCON UNIVERSAL	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N62488</u>	<u>001</u>	Nov 06, 1985
<u>OTICAIR</u>						
<u>AT</u>		BAUSCH AND LOMB	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N64065</u>	<u>001</u>	Aug 28, 1996
<u>PEDIOTIC</u>						
<u>AT</u>		MONARCH PHARMS	<u>1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML</u>	<u>N62822</u>	<u>001</u>	Sep 29, 1987

HYDROFLUMETHIAZIDE

TABLET; ORAL

HYDROFLUMETHIAZIDE

<u>AB</u>		PAR PHARM	<u>50MG</u>	<u>N88850</u>	<u>001</u>	May 31, 1985
<u>SALURON</u>						
<u>AB</u>	+	SHIRE	<u>50MG</u>	<u>N11949</u>	<u>001</u>	

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

DILAUDID-HP

<u>AP</u>	+	ABBOTT	<u>10MG/ML</u>	<u>N19034</u>	<u>001</u>	Jan 11, 1984
	+		250MG/VIAL	N19034	002	Aug 04, 1994

HYDROMORPHONE HYDROCHLORIDE

<u>AP</u>		HOSPIRA	<u>10MG/ML</u>	<u>N74598</u>	<u>001</u>	Jun 19, 1997
<u>AP</u>		MAYNE PHARMA USA	<u>10MG/ML</u>	<u>N76444</u>	<u>001</u>	Apr 25, 2003
<u>AP</u>		STERIS	<u>10MG/ML</u>	<u>N74317</u>	<u>001</u>	Aug 23, 1995

SOLUTION; ORAL

DILAUDID

<u>AA</u>	+	ABBOTT	<u>5MG/5ML</u>	<u>N19891</u>	<u>001</u>	Dec 07, 1992
<u>HYDROMORPHONE HYDROCHLORIDE</u>						
<u>AA</u>		ROXANE	<u>5MG/5ML</u>	<u>N74653</u>	<u>001</u>	Jul 29, 1998

TABLET; ORAL

DILAUDID

<u>AB</u>	+	ABBOTT	<u>8MG</u>	<u>N19892</u>	<u>001</u>	Dec 07, 1992
<u>HYDROMORPHONE HYDROCHLORIDE</u>						
<u>AB</u>		AMIDE PHARM	<u>8MG</u>	<u>N76723</u>	<u>001</u>	Oct 18, 2005
<u>AB</u>		KV PHARM	<u>8MG</u>	<u>N77311</u>	<u>003</u>	Nov 09, 2005
			2MG	N77311	001	Nov 09, 2005
			4MG	N77311	002	Nov 09, 2005
<u>AB</u>		MALLINCKRODT	<u>8MG</u>	<u>N76855</u>	<u>001</u>	Dec 23, 2004
<u>AB</u>		ROXANE	<u>8MG</u>	<u>N74597</u>	<u>001</u>	Jul 29, 1998

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

HYDROXOCOBALAMIN

	+	STERIS	1MG/ML	N85998	001	
--	---	--------	--------	--------	-----	--

HYDROXYAMPHETAMINE HYDROBROMIDE; TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC

PAREMYD

	+	AKORN	1%;0.25%	N19261	001	Jan 30, 1992
--	---	-------	----------	--------	-----	--------------

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

<u>AB</u>		MYLAN	<u>200MG</u>	<u>N40274</u>	<u>001</u>	May 29, 1998
<u>AB</u>		SANDOZ	<u>200MG</u>	<u>N40104</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>			<u>200MG</u>	<u>N40150</u>	<u>001</u>	Jan 27, 1996

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 198 (of 360)

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

<u>AB</u>	TEVA PHARMS	<u>200MG</u>	<u>N40081</u>	<u>001</u>	Sep 30, 1994
<u>AB</u>	WATSON LABS	<u>200MG</u>	<u>N40133</u>	<u>001</u>	Nov 30, 1995

PLAQUENIL

<u>AB</u>	+ SANOFI SYNTHELABO	<u>200MG</u>	<u>N09768</u>	<u>001</u>	
-----------	---------------------	--------------	---------------	------------	--

HYDROXYPROPYL CELLULOSE

INSERT; OPHTHALMIC

LACRISERT

+	MERCK	5MG	N18771	001	
---	-------	-----	--------	-----	--

HYDROXYUREA

CAPSULE; ORAL

DROXIA

	BRISTOL MYERS SQUIBB	200MG	N16295	002	Feb 25, 1998
		300MG	N16295	003	Feb 25, 1998
+		400MG	N16295	004	Feb 25, 1998

HYDREA

<u>AB</u>	+ BRISTOL MYERS SQUIBB	<u>500MG</u>	<u>N16295</u>	<u>001</u>	
-----------	------------------------	--------------	---------------	------------	--

HYDROXYUREA

<u>AB</u>	BARR	<u>250MG</u>	<u>N75143</u>	<u>002</u>	Sep 21, 2000
<u>AB</u>		<u>500MG</u>	<u>N75143</u>	<u>001</u>	Oct 16, 1998
<u>AB</u>	+ DURAMED PHARMS BARR	<u>250MG</u>	<u>N75020</u>	<u>002</u>	Jun 26, 2000
<u>AB</u>		<u>500MG</u>	<u>N75020</u>	<u>001</u>	Jul 30, 1998
<u>AB</u>	PAR PHARM	<u>500MG</u>	<u>N75340</u>	<u>001</u>	Feb 24, 1999
<u>AB</u>	ROXANE	<u>500MG</u>	<u>N74476</u>	<u>001</u>	Aug 18, 1995

TABLET; ORAL

HYDROXYUREA

+	BARR	1GM	N75734	001	Aug 29, 2000
---	------	-----	--------	-----	--------------

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

<u>AP</u>	AM PHARM PARTNERS	<u>25MG/ML</u>	<u>N87329</u>	<u>001</u>	
<u>AP</u>		<u>50MG/ML</u>	<u>N87329</u>	<u>002</u>	
<u>AP</u>	HOSPIRA	<u>25MG/ML</u>	<u>N87416</u>	<u>001</u>	
<u>AP</u>		<u>50MG/ML</u>	<u>N87546</u>	<u>001</u>	
<u>AP</u>	LUITPOLD	<u>25MG/ML</u>	<u>N87408</u>	<u>001</u>	
<u>AP</u>		<u>50MG/ML</u>	<u>N87408</u>	<u>002</u>	
<u>AP</u>	STERIS	<u>50MG/ML</u>	<u>N85779</u>	<u>001</u>	
	<u>VISTARIL</u>				
<u>AP</u>	+ PFIZER	<u>25MG/ML</u>	<u>N11111</u>	<u>001</u>	
<u>AP</u>	+	<u>50MG/ML</u>	<u>N11111</u>	<u>002</u>	

SYRUP; ORAL

ATARAX

<u>AA</u>	+ ROERIG	<u>10MG/5ML</u>	<u>N10485</u>	<u>001</u>	
-----------	----------	-----------------	---------------	------------	--

HYDROXYZINE HYDROCHLORIDE

<u>AA</u>	ALPHARMA US PHARMS	<u>10MG/5ML</u>	<u>N86880</u>	<u>001</u>	
<u>AA</u>	HI TECH PHARMA	<u>10MG/5ML</u>	<u>N40010</u>	<u>001</u>	Oct 28, 1994
<u>AA</u>	MORTON GROVE	<u>10MG/5ML</u>	<u>N87294</u>	<u>001</u>	Apr 12, 1982
<u>AA</u>	VINTAGE PHARMS	<u>10MG/5ML</u>	<u>N40391</u>	<u>001</u>	Apr 10, 2002

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AB</u>	AMIDE PHARM	<u>10MG</u>	<u>N40600</u>	<u>001</u>	Dec 28, 2004
<u>AB</u>		<u>25MG</u>	<u>N40602</u>	<u>001</u>	Dec 28, 2004
<u>AB</u>		<u>50MG</u>	<u>N40604</u>	<u>001</u>	Dec 28, 2004
<u>AB</u>	MUTUAL PHARM	<u>10MG</u>	<u>N89381</u>	<u>001</u>	May 19, 1986

PRESCRIPTION DRUG PRODUCT LIST

3 - 199 (of 360)

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

<u>AB</u>	MUTUAL PHARM	<u>25MG</u>	<u>N89382</u>	<u>001</u>	May 19, 1986
<u>AB</u>		<u>50MG</u>	<u>N89383</u>	<u>001</u>	May 19, 1986
<u>AB</u>	+ PLIVA	<u>10MG</u>	<u>N88617</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>	+	<u>25MG</u>	<u>N88618</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>	+	<u>50MG</u>	<u>N88619</u>	<u>001</u>	Jan 10, 1986
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>N87869</u>	<u>001</u>	Dec 20, 1982
<u>AB</u>		<u>25MG</u>	<u>N87870</u>	<u>001</u>	Dec 20, 1982
<u>AB</u>		<u>50MG</u>	<u>N87871</u>	<u>001</u>	Dec 20, 1982
<u>AB</u>	VINTAGE PHARMS	<u>10MG</u>	<u>N40579</u>	<u>001</u>	May 27, 2005
<u>AB</u>		<u>25MG</u>	<u>N40574</u>	<u>001</u>	May 27, 2005
<u>AB</u>		<u>50MG</u>	<u>N40580</u>	<u>001</u>	May 27, 2005
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N81149</u>	<u>001</u>	Mar 18, 1994
<u>AB</u>		<u>10MG</u>	<u>N88348</u>	<u>001</u>	Sep 15, 1983
<u>AB</u>		<u>25MG</u>	<u>N81150</u>	<u>001</u>	Mar 18, 1994
<u>AB</u>		<u>25MG</u>	<u>N88349</u>	<u>001</u>	Sep 15, 1983
<u>AB</u>		<u>50MG</u>	<u>N81151</u>	<u>001</u>	Mar 18, 1994
<u>AB</u>		<u>50MG</u>	<u>N88350</u>	<u>001</u>	Sep 15, 1983

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

<u>AB</u>	BARR	<u>EQ 25MG HCL</u>	<u>N88496</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>		<u>EQ 50MG HCL</u>	<u>N88487</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>		<u>EQ 100MG HCL</u>	<u>N88488</u>	<u>001</u>	Jun 15, 1984
<u>AB</u>	IVAX PHARMS	<u>EQ 25MG HCL</u>	<u>N87761</u>	<u>001</u>	Mar 05, 1982
<u>AB</u>		<u>EQ 50MG HCL</u>	<u>N87760</u>	<u>001</u>	Mar 05, 1982
<u>AB</u>	SANDOZ	<u>EQ 25MG HCL</u>	<u>N81127</u>	<u>001</u>	Jun 28, 1991
<u>AB</u>		<u>EQ 25MG HCL</u>	<u>N87479</u>	<u>001</u>	
<u>AB</u>		<u>EQ 50MG HCL</u>	<u>N86183</u>	<u>001</u>	
<u>AB</u>	WATSON LABS	<u>EQ 25MG HCL</u>	<u>N40156</u>	<u>001</u>	Jul 15, 1996
<u>AB</u>		<u>EQ 25MG HCL</u>	<u>N81165</u>	<u>001</u>	Jul 31, 1991
<u>AB</u>		<u>EQ 50MG HCL</u>	<u>N40156</u>	<u>002</u>	Jul 15, 1996
<u>VISTARIL</u>					
<u>AB</u>	PFIZER	<u>EQ 25MG HCL</u>	<u>N11459</u>	<u>002</u>	
<u>AB</u>	+	<u>EQ 50MG HCL</u>	<u>N11459</u>	<u>004</u>	
<u>AB</u>		<u>EQ 100MG HCL</u>	<u>N11459</u>	<u>006</u>	

SUSPENSION; ORAL

VISTARIL

+	PFIZER	EQ 25MG HCL/5ML	N11795	001	
---	--------	-----------------	--------	-----	--

IBANDRONATE SODIUM

TABLET; ORAL

BONIVA

+	ROCHE	EQ 2.5MG BASE	N21455	001	May 16, 2003
+		EQ 150MG BASE	N21455	002	Mar 24, 2005

IBUPROFEN

SUSPENSION; ORAL

CHILDREN'S ADVIL

BX	WYETH CONS	100MG/5ML	N19833	002	Sep 19, 1989
<u>IBUPROFEN</u>					
<u>AB</u>	ALPHARMA US PHARMS	<u>100MG/5ML</u>	<u>N74978</u>	<u>001</u>	Mar 25, 1998
<u>AB</u>	PERRIGO R AND D	<u>100MG/5ML</u>	<u>N76925</u>	<u>001</u>	Sep 23, 2004
<u>MOTRIN</u>					
<u>AB</u>	+ MCNEIL CONS SPECLT	<u>100MG/5ML</u>	<u>N19842</u>	<u>001</u>	Sep 19, 1989

PRESCRIPTION DRUG PRODUCT LIST

3 - 200 (of 360)

IBUPROFEN

TABLET; ORAL

<u>IBU</u>				
<u>AB</u>	BASF	<u>400MG</u>	<u>N18197</u>	<u>001</u>
<u>AB</u>		<u>400MG</u>	<u>N70083</u>	<u>001</u> Feb 22, 1985
<u>AB</u>		<u>600MG</u>	<u>N70099</u>	<u>001</u> Mar 29, 1985
<u>AB</u>		<u>800MG</u>	<u>N70745</u>	<u>001</u> Jul 23, 1986
<u>IBUPROFEN</u>				
<u>AB</u>	BASF	<u>400MG</u>	<u>N75682</u>	<u>001</u> Nov 14, 2001
<u>AB</u>		<u>600MG</u>	<u>N75682</u>	<u>002</u> Nov 14, 2001
<u>AB</u>		<u>800MG</u>	<u>N75682</u>	<u>003</u> Nov 14, 2001
<u>AB</u>	DR REDDYS LABS INC	<u>400MG</u>	<u>N76112</u>	<u>001</u> Oct 31, 2001
<u>AB</u>		<u>600MG</u>	<u>N76112</u>	<u>002</u> Oct 31, 2001
<u>AB</u>		<u>800MG</u>	<u>N76112</u>	<u>003</u> Oct 31, 2001
<u>AB</u>	INTERPHARM	<u>400MG</u>	<u>N71334</u>	<u>001</u> Nov 25, 1986
<u>AB</u>		<u>600MG</u>	<u>N71335</u>	<u>001</u> Nov 25, 1986
<u>AB</u>		<u>800MG</u>	<u>N71935</u>	<u>001</u> Oct 13, 1987
<u>AB</u>	MUTUAL PHARM	<u>300MG</u>	<u>N71230</u>	<u>001</u> Oct 22, 1986
<u>AB</u>		<u>400MG</u>	<u>N71231</u>	<u>001</u> Oct 22, 1986
<u>AB</u>		<u>600MG</u>	<u>N71232</u>	<u>001</u> Oct 22, 1986
<u>AB</u>		<u>800MG</u>	<u>N72004</u>	<u>001</u> Nov 18, 1987
<u>AB</u>	MYLAN	<u>400MG</u>	<u>N70045</u>	<u>001</u> Sep 24, 1985
<u>AB</u>		<u>600MG</u>	<u>N70057</u>	<u>001</u> Sep 24, 1985
<u>AB</u>		<u>800MG</u>	<u>N71999</u>	<u>001</u> Dec 03, 1987
<u>AB</u>	OHM LABS	<u>400MG</u>	<u>N70818</u>	<u>001</u> Dec 26, 1985
<u>AB</u>	PAR PHARM	<u>400MG</u>	<u>N70329</u>	<u>001</u> Aug 06, 1985
<u>AB</u>		<u>600MG</u>	<u>N70330</u>	<u>001</u> Aug 06, 1985
<u>AB</u>		<u>800MG</u>	<u>N70986</u>	<u>001</u> Jul 25, 1986
<u>AB</u>	PERRIGO R AND D	<u>400MG</u>	<u>N77114</u>	<u>001</u> Jul 18, 2005
<u>AB</u>		<u>600MG</u>	<u>N77114</u>	<u>002</u> Jul 18, 2005
<u>AB</u>		<u>800MG</u>	<u>N77114</u>	<u>003</u> Jul 18, 2005
<u>AB</u>	PHARM FORM	<u>300MG</u>	<u>N71266</u>	<u>001</u> Oct 15, 1986
<u>AB</u>		<u>400MG</u>	<u>N71267</u>	<u>001</u> Oct 15, 1986
<u>AB</u>		<u>600MG</u>	<u>N71268</u>	<u>001</u> Oct 15, 1986
<u>AB</u>		<u>800MG</u>	<u>N72300</u>	<u>001</u> Jul 01, 1988
<u>AB</u>	PLIVA	<u>400MG</u>	<u>N71666</u>	<u>001</u> Jun 18, 1987
<u>AB</u>		<u>600MG</u>	<u>N71667</u>	<u>001</u> Jun 18, 1987
<u>AB</u>		<u>800MG</u>	<u>N71668</u>	<u>001</u> Jun 18, 1987
<u>AB</u>	SANDOZ	<u>300MG</u>	<u>N70734</u>	<u>001</u> Jun 12, 1986
<u>AB</u>		<u>400MG</u>	<u>N70735</u>	<u>001</u> Jun 12, 1986
<u>AB</u>		<u>600MG</u>	<u>N70736</u>	<u>001</u> Jun 12, 1986
<u>AB</u>		<u>800MG</u>	<u>N72169</u>	<u>001</u> Dec 11, 1987
<u>AB</u>	VINTAGE PHARMS	<u>400MG</u>	<u>N71644</u>	<u>001</u> Feb 01, 1988
<u>AB</u>	WATSON LABS	<u>400MG</u>	<u>N70436</u>	<u>001</u> Aug 21, 1985
<u>AB</u>		<u>600MG</u>	<u>N70437</u>	<u>001</u> Aug 21, 1985
<u>AB</u>		<u>800MG</u>	<u>N71547</u>	<u>001</u> Jul 02, 1987
<u>IBUPROHM</u>				
<u>AB</u>	OHM LABS	<u>400MG</u>	<u>N70469</u>	<u>001</u> Aug 29, 1985
<u>IBU - TAB</u>				
<u>AB</u>	ALRA	<u>400MG</u>	<u>N71058</u>	<u>001</u> Aug 11, 1988
<u>AB</u>		<u>600MG</u>	<u>N71059</u>	<u>001</u> Aug 11, 1988
<u>MOTRIN</u>				
<u>AB</u>	MCNEIL CONS SPECLT	<u>300MG</u>	<u>N17463</u>	<u>003</u>
<u>AB</u>		<u>400MG</u>	<u>N17463</u>	<u>002</u>
<u>AB</u>		<u>600MG</u>	<u>N17463</u>	<u>004</u>
<u>AB</u>	+	<u>800MG</u>	<u>N17463</u>	<u>005</u> May 22, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 201 (of 360)

IBUPROFEN; OXYCODONE HYDROCHLORIDETABLET; ORAL
COMBUNOX

+ FOREST LABS 400MG;5MG N21378 001 Nov 26, 2004

IBUTILIDE FUMARATEINJECTABLE; INJECTION
CORVERT

+ PHARMACIA AND UPJOHN 0.1MG/ML N20491 001 Dec 28, 1995

ICODEXTRINSOLUTION; INTRAPERITONEAL
EXTRANEAL

+ BAXTER HLTHCARE 7.5GM/100ML N21321 001 Dec 20, 2002

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN PFSAP + PHARMACIA AND UPJOHN 1MG/ML N50734 001 Feb 17, 1997

IDARUBICIN HYDROCHLORIDE

+ SICOR PHARMS 5MG/VIAL N65037 003 May 01, 2002

+ 10MG/VIAL N65037 002 May 01, 2002

+ 20MG/VIAL N65037 001 May 01, 2002

IDARUBICIN HYDROCHLORIDE PFSAP SICOR PHARMS 1MG/ML N65036 001 May 01, 2002IDOXURIDINE

SOLUTION/DROPS; OPHTHALMIC

DENDRIDAT + ALCON 0.1% N14169 001HERPLEXAT + ALLERGAN 0.1% N13935 002IFOSFAMIDEINJECTABLE; INJECTION
IFOSFAMIDE

+ AM PHARM PARTNERS 1GM/VIAL N76078 001 May 28, 2002

+ 3GM/VIAL N76078 002 May 28, 2002

IFOSFAMIDE; MESNAINJECTABLE; INJECTION
IFEX/MESNEX KIT

+ BRISTOL MYERS SQUIBB 1GM/VIAL;100MG/ML N19763 003 Oct 10, 1992

+ 3GM/VIAL;100MG/ML N19763 004 Oct 10, 1992

INJECTABLE; INTRAVENOUS
IFOSFAMIDE/MESNA KIT

+ SICOR PHARMS EQ 1GM /20ML(50MG/ML);EQ 1GM /10ML(100MG/ML) N75874 001 Feb 26, 2002

+ EQ 1GM/10ML (100MG/ML) N75874 002 Feb 26, 2002

ILOPROSTSOLUTION; INHALATION
VENTAVIS

+ COTHERIX 10UGM/ML (10UGM/ML) N21779 002 Dec 08, 2005

+ 20UGM/2ML (10UGM/ML) N21779 001 Dec 29, 2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 202 (of 360)

IMATINIB MESYLATE

TABLET; ORAL

GLEEVEC

NOVARTIS

100MG

N21588 001

Apr 18, 2003

+

400MG

N21588 002

Apr 18, 2003

IMIGLUCERASE

INJECTABLE; INJECTION

CEREZYME

GENZYME

200 UNITS/VIAL

N20367 001

May 23, 1994

+

400 UNITS/VIAL

N20367 002

Sep 22, 1999

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDEAB MUTUAL PHARM10MGN81048 001

Jun 05, 1990

AB25MGN81049 001

Jun 05, 1990

AB50MGN81050 001

Jun 05, 1990

AB PAR PHARM10MGN88292 001

Oct 21, 1983

AB10MGN89422 001

Jul 14, 1987

AB25MGN88262 001

Oct 21, 1983

AB25MGN89497 001

Jul 14, 1987

AB50MGN88276 001

Oct 21, 1983

AB SANDOZ10MGN84936 002AB25MGN83745 001AB50MGN84937 001TOFRANILAB TYCO HLTHCARE10MGN87844 001

May 22, 1984

AB25MGN87845 001

May 22, 1984

AB +50MGN87846 001

May 22, 1984

IMIPRAMINE PAMOATE

CAPSULE; ORAL

TOFRANIL-PM

TYCO HLTHCARE

EQ 75MG HCL

N17090 001

EQ 100MG HCL

N17090 004

EQ 125MG HCL

N17090 003

+

EQ 150MG HCL

N17090 002

IMIQUIMOD

CREAM; TOPICAL

ALDARA

+ 3M

5%

N20723 001

Feb 27, 1997

INAMRINONE LACTATE

INJECTABLE; INJECTION

AMRINONEAP BEDFORDEQ 5MG BASE/MLN75513 001

May 09, 2000

AP + HOSPIRAEQ 5MG BASE/MLN74616 001

Aug 03, 1998

AMRINONE LACTATEAP BAXTER HLTHCARE CORPEQ 5MG BASE/MLN75542 001

May 10, 2000

INDAPAMIDE

TABLET; ORAL

INDAPAMIDEAB ALPHAPHARM1.25MGN75105 001

Jul 23, 1998

AB2.5MGN75105 002

Jul 23, 1998

AB IVAX PHARMS1.25MGN74299 002

Apr 29, 1996

AB2.5MGN74299 001

Jul 27, 1995

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 203 (of 360)

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

<u>AB</u>	MYLAN	<u>1.25MG</u>	<u>N74461</u>	<u>002</u>	Mar 26, 1997
<u>AB</u>		<u>2.5MG</u>	<u>N74461</u>	<u>001</u>	Mar 27, 1996
<u>AB</u>	PUREPAC PHARM	<u>1.25MG</u>	<u>N74722</u>	<u>001</u>	Jun 17, 1996
<u>AB</u>		<u>2.5MG</u>	<u>N74722</u>	<u>002</u>	Jun 17, 1996
<u>AB</u>	SANDOZ	<u>1.25MG</u>	<u>N74594</u>	<u>001</u>	May 23, 1996
<u>AB</u>		<u>2.5MG</u>	<u>N74594</u>	<u>002</u>	May 23, 1996
<u>AB</u>	TEVA	<u>1.25MG</u>	<u>N74498</u>	<u>002</u>	Feb 12, 1998
<u>AB</u>		<u>2.5MG</u>	<u>N74498</u>	<u>001</u>	Oct 31, 1996
<u>AB</u>	TRIGEN	<u>1.25MG</u>	<u>N75201</u>	<u>001</u>	Dec 04, 1998
<u>AB</u>		<u>2.5MG</u>	<u>N75201</u>	<u>002</u>	Dec 04, 1998
<u>AB</u>	WATSON LABS	<u>1.25MG</u>	<u>N74585</u>	<u>001</u>	Sep 26, 1996
<u>AB</u>		<u>2.5MG</u>	<u>N74585</u>	<u>002</u>	Sep 26, 1996
<u>LOZOL</u>					
<u>AB</u>	AVENTIS	<u>1.25MG</u>	<u>N18538</u>	<u>002</u>	Apr 29, 1993
<u>AB</u>	+	<u>2.5MG</u>	<u>N18538</u>	<u>001</u>	Jul 06, 1983

INDINAVIR SULFATE

CAPSULE; ORAL

CRIXIVAN

MERCK

EQ 100MG BASE

N20685 006 Apr 19, 2000

EQ 200MG BASE

N20685 003 Mar 13, 1996

EQ 333MG BASE

N20685 005 Dec 17, 1998

+

EQ 400MG BASE

N20685 001 Mar 13, 1996

INDIUM IN 111 CHLORIDE

INJECTABLE; INJECTION

INDICLOR

+ AMERSHAM

2mCi/0.2ML

N19862 001 Dec 29, 1992

INDIUM IN 111 CHLORIDE

+ MALLINCKRODT

5mCi/0.5ML

N19841 001 Sep 27, 1994

INDIUM IN-111 OXYQUINOLINE

INJECTABLE; INJECTION

INDIUM IN-111 OXYQUINOLINE

GE HEALTHCARE

1mCi/ML

N19044 001 Dec 24, 1985

INDIUM IN-111 PENTETATE DISODIUM

INJECTABLE; INTRATHECAL

MPI INDIUM DTPA IN 111

GE HEALTHCARE

1mCi/ML

N17707 001 Feb 18, 1982

INDIUM IN-111 PENTETREOTIDE KIT

INJECTABLE; INJECTION

OCTREOSCAN

MALLINCKRODT

3mCi/ML

N20314 001 Jun 02, 1994

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN

+ AKORN

25MG/VIAL

N11525 001

INDOMETHACIN

CAPSULE; ORAL

INDOCIN

<u>AB</u>	MERCK	<u>25MG</u>	<u>N16059</u>	<u>001</u>	
<u>AB</u>	+	<u>50MG</u>	<u>N16059</u>	<u>002</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 204 (of 360)

INDOMETHACIN

CAPSULE; ORAL					
<u>INDO-LEMMON</u>					
<u>AB</u>	TEVA	<u>25MG</u>	<u>N70266</u>	<u>001</u>	Nov 07, 1985
<u>AB</u>		<u>50MG</u>	<u>N70267</u>	<u>001</u>	Nov 07, 1985
<u>INDOMETHACIN</u>					
<u>AB</u>	CLONMEL HLTHCARE	<u>25MG</u>	<u>N18851</u>	<u>001</u>	May 18, 1984
<u>AB</u>		<u>50MG</u>	<u>N18851</u>	<u>002</u>	May 18, 1984
<u>AB</u>	IVAX PHARMS	<u>25MG</u>	<u>N70719</u>	<u>001</u>	Feb 12, 1986
<u>AB</u>		<u>50MG</u>	<u>N70756</u>	<u>001</u>	Feb 12, 1986
<u>AB</u>	MUTUAL PHARM	<u>25MG</u>	<u>N70899</u>	<u>001</u>	Feb 09, 1987
<u>AB</u>		<u>50MG</u>	<u>N70900</u>	<u>001</u>	Feb 09, 1987
<u>AB</u>	MYLAN	<u>25MG</u>	<u>N18858</u>	<u>001</u>	Apr 20, 1984
<u>AB</u>		<u>50MG</u>	<u>N18858</u>	<u>002</u>	Apr 20, 1984
<u>AB</u>		<u>50MG</u>	<u>N70624</u>	<u>001</u>	Sep 04, 1985
<u>AB</u>	PAR PHARM	<u>25MG</u>	<u>N18829</u>	<u>002</u>	Aug 06, 1984
<u>AB</u>		<u>50MG</u>	<u>N18829</u>	<u>001</u>	Aug 06, 1984
<u>AB</u>		<u>50MG</u>	<u>N70651</u>	<u>001</u>	Mar 05, 1986
<u>AB</u>	PLIVA	<u>25MG</u>	<u>N71148</u>	<u>001</u>	Mar 18, 1987
<u>AB</u>		<u>50MG</u>	<u>N71149</u>	<u>001</u>	Mar 18, 1987
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>N70673</u>	<u>001</u>	Apr 29, 1987
<u>AB</u>		<u>50MG</u>	<u>N70674</u>	<u>001</u>	Apr 29, 1987
<u>AB</u>	TEVA	<u>25MG</u>	<u>N71342</u>	<u>001</u>	Apr 18, 1988
<u>AB</u>		<u>50MG</u>	<u>N71343</u>	<u>001</u>	Apr 18, 1988

CAPSULE, EXTENDED RELEASE; ORAL					
<u>INDOCIN SR</u>					
<u>AB</u>	+ SANDOZ	<u>75MG</u>	<u>N74464</u>	<u>001</u>	May 28, 1998
<u>INDOMETHACIN</u>					
<u>AB</u>	INWOOD LABS	<u>75MG</u>	<u>N72410</u>	<u>001</u>	Mar 15, 1989
SUPPOSITORY; RECTAL					
INDOMETHEGAN					
	+ G AND W LABS	50MG	N73314	001	Aug 31, 1992
SUSPENSION; ORAL					
INDOCIN					
	+ MERCK	25MG/5ML	N18332	001	Oct 10, 1985

INDOMETHACIN SODIUM

INJECTABLE; INJECTION					
INDOCIN I.V.					
	+ OVATION PHARMS	EQ 1MG BASE/VIAL	N18878	001	Jan 30, 1985

INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS					
NOVOLOG MIX 70/30					
	+ NOVO NORDISK INC	70 UNITS/ML;30 UNITS/ML	N21172	001	Nov 01, 2001

INSULIN ASPART RECOMBINANT

INJECTABLE; SUBCUTANEOUS					
NOVOLOG					
	+ NOVO NORDISK INC	100 UNITS/ML	N20986	001	Jun 07, 2000

INSULIN DETEMIR RECOMBINANT

INJECTABLE; SUBCUTANEOUS					
LEVEMIR					
	+ NOVO NORDISK INC	100 UNITS/ML	N21536	001	Jun 16, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 205 (of 360)

INSULIN GLARGINE RECOMBINANT

INJECTABLE; INJECTION				
LANTUS				
+ AVENTIS PHARMS	100 UNITS/ML	N21081	001	Apr 20, 2000

INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT

INJECTABLE; INJECTION				
HUMALOG MIX 50/50				
+ LILLY	50 UNITS/ML;50 UNITS/ML	N21018	001	Dec 22, 1999
HUMALOG MIX 75/25				
+ LILLY	75 UNITS/ML;25 UNITS/ML	N21017	001	Dec 22, 1999

INSULIN LISPRO RECOMBINANT

INJECTABLE; INJECTION				
HUMALOG				
+ LILLY	100 UNITS/ML	N20563	001	Jun 14, 1996
HUMALOG PEN				
+ LILLY	100 UNITS/ML	N20563	002	Aug 06, 1998

INSULIN PURIFIED PORK

INJECTABLE; INJECTION				
ILETIN II				
+ LILLY	500 UNITS/ML	N18344	002	

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION				
HUMULIN R				
+ LILLY	500 UNITS/ML	N18780	004	Mar 31, 1994

IOBENGUANE SULFATE I 131

INJECTABLE; INJECTION				
IOBENGUANE SULFATE I 131				
CIS	2.3mCi/ML	N20084	001	Mar 25, 1994

IODIPAMIDE MEGLUMINE

INJECTABLE; INJECTION				
CHOLOGRAFIN MEGLUMINE				
+ BRACCO	52%	N09321	003	

IODIXANOL

INJECTABLE; INJECTION				
VISIPAQUE 270				
+ GE HEALTHCARE	55%	N20351	001	Mar 22, 1996
	55%	N20808	001	Aug 29, 1997
VISIPAQUE 320				
+ GE HEALTHCARE	65.2%	N20351	002	Mar 22, 1996
	65.2%	N20808	002	Aug 29, 1997

IOHEXOL

INJECTABLE; INJECTION				
OMNIPAQUE 140				
+ GE HEALTHCARE	30.2%	N18956	005	Nov 30, 1988
SOLUTION; INJECTION, ORAL				
OMNIPAQUE 350				
+ GE HEALTHCARE	75.5%	N18956	004	Dec 26, 1985
	75.5%	N20608	003	Oct 24, 1995
SOLUTION; INJECTION, ORAL, RECTAL				
OMNIPAQUE 180				
+ GE HEALTHCARE	38.8%	N18956	001	Dec 26, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 206 (of 360)

IOHEXOL

SOLUTION; INJECTION, ORAL, RECTAL

OMNIPAQUE 240

+	GE HEALTHCARE	51.8%	N18956	002	Dec 26, 1985
		51.8%	N20608	001	Oct 24, 1995

OMNIPAQUE 300

+	GE HEALTHCARE	64.7%	N18956	003	Dec 26, 1985
		64.7%	N20608	002	Oct 24, 1995

IOPAMIDOL

INJECTABLE; INJECTION

IOPAMIDOL-200

<u>AP</u>	COOK IMAGING	<u>41%</u>	<u>N74881</u>	<u>001</u>	Jul 28, 2000
<u>AP</u>	HOSPIRA	<u>41%</u>	<u>N74898</u>	<u>001</u>	Dec 30, 1997

IOPAMIDOL-200 IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>41%</u>	<u>N74636</u>	<u>001</u>	Dec 30, 1997
-----------	---------	------------	---------------	------------	--------------

IOPAMIDOL-250

<u>AP</u>	AM PHARM PARTNERS	<u>51%</u>	<u>N74679</u>	<u>001</u>	Apr 02, 1997
<u>AP</u>	COOK IMAGING	<u>51%</u>	<u>N74881</u>	<u>002</u>	Jul 28, 2000
<u>AP</u>	HOSPIRA	<u>51%</u>	<u>N74898</u>	<u>002</u>	Dec 30, 1997
<u>AP</u>		<u>51%</u>	<u>N75005</u>	<u>001</u>	Feb 24, 1998

IOPAMIDOL-250 IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>51%</u>	<u>N74636</u>	<u>002</u>	Dec 30, 1997
-----------	---------	------------	---------------	------------	--------------

IOPAMIDOL-300

<u>AP</u>	AM PHARM PARTNERS	<u>61%</u>	<u>N74679</u>	<u>002</u>	Apr 02, 1997
<u>AP</u>	COOK IMAGING	<u>61%</u>	<u>N74881</u>	<u>003</u>	Jul 28, 2000
<u>AP</u>	HOSPIRA	<u>61%</u>	<u>N74898</u>	<u>003</u>	Dec 30, 1997
<u>AP</u>		<u>61%</u>	<u>N75005</u>	<u>002</u>	Feb 24, 1998

IOPAMIDOL-300 IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>61%</u>	<u>N74636</u>	<u>003</u>	Dec 30, 1997
<u>AP</u>		<u>61%</u>	<u>N74637</u>	<u>001</u>	Apr 03, 1997

IOPAMIDOL-370

<u>AP</u>	AM PHARM PARTNERS	<u>76%</u>	<u>N74679</u>	<u>003</u>	Apr 02, 1997
<u>AP</u>	COOK IMAGING	<u>76%</u>	<u>N74881</u>	<u>004</u>	Jul 28, 2000
<u>AP</u>	HOSPIRA	<u>76%</u>	<u>N74898</u>	<u>004</u>	Dec 30, 1997
<u>AP</u>		<u>76%</u>	<u>N75005</u>	<u>003</u>	Feb 24, 1998

IOPAMIDOL-370 IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>76%</u>	<u>N74636</u>	<u>004</u>	Dec 30, 1997
-----------	---------	------------	---------------	------------	--------------

ISOVUE-200

<u>AP</u>	+	BRACCO	<u>41%</u>	<u>N18735</u>	<u>006</u>	Jul 07, 1987
-----------	---	--------	------------	---------------	------------	--------------

ISOVUE-250

<u>AP</u>	+	BRACCO	<u>51%</u>	<u>N18735</u>	<u>007</u>	Jul 06, 1992
<u>AP</u>	+		<u>51%</u>	<u>N20327</u>	<u>002</u>	Oct 12, 1994

ISOVUE-300

<u>AP</u>	+	BRACCO	<u>61%</u>	<u>N18735</u>	<u>002</u>	Dec 31, 1985
<u>AP</u>	+		<u>61%</u>	<u>N20327</u>	<u>003</u>	Oct 12, 1994

ISOVUE-370

<u>AP</u>	+	BRACCO	<u>76%</u>	<u>N18735</u>	<u>003</u>	Dec 31, 1985
<u>AP</u>	+		<u>76%</u>	<u>N20327</u>	<u>004</u>	Oct 12, 1994

ISOVUE-M 200

+	BRACCO	41%	N18735	001	Dec 31, 1985
---	--------	-----	--------	-----	--------------

ISOVUE-M 300

+	BRACCO	61%	N18735	004	Dec 31, 1985
---	--------	-----	--------	-----	--------------

IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST (PHARMACY BULK)

+	BERLEX	49.9%	N21425	003	Mar 12, 2004
+		62.3%	N21425	001	Sep 20, 2002
+		76.9%	N21425	002	Sep 20, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 207 (of 360)

IOPROMIDE

INJECTABLE; INJECTION

ULTRAVIST 150

+	BERLEX	31.2%	N20220	004	May 10, 1995
---	--------	-------	--------	-----	--------------

ULTRAVIST 240

+	BERLEX	49.9%	N20220	003	May 10, 1995
---	--------	-------	--------	-----	--------------

ULTRAVIST 300

+	BERLEX	62.3%	N20220	002	May 10, 1995
---	--------	-------	--------	-----	--------------

ULTRAVIST 370

+	BERLEX	76.9%	N20220	001	May 10, 1995
---	--------	-------	--------	-----	--------------

IOTHALAMATE MEGLUMINE

INJECTABLE; INJECTION

CONRAY

+	MALLINCKRODT	60%	N13295	001	
---	--------------	-----	--------	-----	--

CONRAY 30

+	MALLINCKRODT	30%	N16983	001	
---	--------------	-----	--------	-----	--

CONRAY 43

+	MALLINCKRODT	43%	N13295	002	
---	--------------	-----	--------	-----	--

SOLUTION; INTRAVESICAL

CYSTO-CONRAY II

MALLINCKRODT	17.2%	N17057	002		
--------------	-------	--------	-----	--	--

IOTHALAMATE SODIUM

INJECTABLE; INJECTION

CONRAY 400

+	MALLINCKRODT	66.8%	N14295	001	
---	--------------	-------	--------	-----	--

IOTHALAMATE SODIUM, I-125

INJECTABLE; INJECTION

GLOFIL-125

QOL MEDCL	250-300uCi/ML	N17279	001		
-----------	---------------	--------	-----	--	--

IOVERSOL

INJECTABLE; INJECTION

OPTIRAY 160

+	MALLINCKRODT	34%	N19710	003	Dec 30, 1988
---	--------------	-----	--------	-----	--------------

OPTIRAY 240

+	MALLINCKRODT	51%	N19710	002	Dec 30, 1988
---	--------------	-----	--------	-----	--------------

OPTIRAY 300

+	MALLINCKRODT	64%	N19710	004	Jan 22, 1992
---	--------------	-----	--------	-----	--------------

+		64%	N20923	004	May 13, 1999
---	--	-----	--------	-----	--------------

OPTIRAY 320

+	MALLINCKRODT	68%	N19710	001	Dec 30, 1988
---	--------------	-----	--------	-----	--------------

OPTIRAY 350

+	MALLINCKRODT	74%	N19710	005	Jan 22, 1992
---	--------------	-----	--------	-----	--------------

+		74%	N20923	003	May 28, 1998
---	--	-----	--------	-----	--------------

IOXAGLATE MEGLUMINE; IOXAGLATE SODIUM

INJECTABLE; INJECTION

HEXABRIX

+	MALLINCKRODT	39.3%;19.6%	N18905	002	Jul 26, 1985
---	--------------	-------------	--------	-----	--------------

IOXILAN

INJECTABLE; INJECTION

OXILAN-300

GUERBET	62%	N20316	001	Dec 21, 1995	
---------	-----	--------	-----	--------------	--

OXILAN-350

GUERBET	73%	N20316	002	Dec 21, 1995	
---------	-----	--------	-----	--------------	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 208 (of 360)

IPRATROPIUM BROMIDE

AEROSOL, METERED; INHALATION

ATROVENT

+ BOEHRINGER INGELHEIM 0.018MG/INH

N19085 001 Dec 29, 1986

ATROVENT HFA

+ BOEHRINGER INGELHEIM 0.021MG/INH

N21527 001 Nov 27, 2004

SOLUTION; INHALATION

ATROVENTAN + BOEHRINGER INGELHEIM 0.02%N20228 001 Sep 29, 1993IPRATROPIUM BROMIDEAN ALPHARMA US PHARMS 0.02%N75111 001 Apr 22, 1999AN BAUSCH AND LOMB 0.02%N75835 001 Oct 15, 2001AN DEY 0.02%N74755 001 Jan 10, 1997AN HOLOPACK INTL 0.02%N75693 001 Jan 26, 2001AN IVAX PHARMS 0.02%N75313 001 Feb 07, 2000AN NEPHRON 0.02%N75562 001 Sep 27, 2001AN NOVEX 0.02%N75441 001 Mar 28, 2001AN RESPIRARE 0.02%N76291 001 May 09, 2005AN ROXANE 0.02%N75867 001 Jul 22, 2002AN RXELITE 0.02%N77072 001 Jul 19, 2005AN WARRICK PHARMS 0.02%N75507 001 Jan 19, 2001

SPRAY, METERED; NASAL

ATROVENTAB + BOEHRINGER INGELHEIM 0.021MG/SPRAYN20393 001 Oct 20, 1995AB + 0.042MG/SPRAYN20394 001 Oct 20, 1995IPRATROPIUM BROMIDEAB BAUSCH AND LOMB 0.021MG/SPRAYN76025 001 Mar 31, 2003AB 0.042MG/SPRAYN76103 001 Mar 31, 2003AB DEY 0.021MG/SPRAYN75552 001 Mar 31, 2003AB 0.042MG/SPRAYN75553 001 Mar 31, 2003AB NOVEX 0.021MG/SPRAYN76156 001 Apr 18, 2003AB 0.042MG/SPRAYN76155 001 Apr 18, 2003AB ROXANE 0.021MG/SPRAYN76664 001 Nov 05, 2003AB 0.042MG/SPRAYN76598 001 Nov 05, 2003IRBESARTAN

TABLET; ORAL

AVAPRO

SANOFI SYNTHELABO 75MG

N20757 001 Sep 30, 1997

150MG

N20757 002 Sep 30, 1997

+ 300MG

N20757 003 Sep 30, 1997

IRINOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

CAMPTOSAR

+ PHARMACIA AND UPJOHN 20MG/ML

N20571 001 Jun 14, 1996

IRON DEXTRAN

INJECTABLE; INJECTION

DEXFERRUM

BP LUITPOLD EQ 50MG IRON/ML

N40024 001 Feb 23, 1996

INFED

BP + WATSON LABS (UTAH) EQ 50MG IRON/ML

N17441 001

PROFERDEX

BP NEW RIVER EQ 50MG IRON/ML

N17807 001

PRESCRIPTION DRUG PRODUCT LIST

3 - 209 (of 360)

IRON SUCROSE

INJECTABLE; INTRAVENOUS
VENOFER

+ LUITPOLD	EQ 100MG BASE/5ML(EQ 20MG BASE/ML)	N21135	001	Nov 06, 2000
	EQ 50MG BASE/2.5ML(EQ 20MG BASE/ML)	N21135	002	Mar 20, 2005
	EQ 75MG BASE/3.75ML(EQ 20MG BASE/ML)	N21135	003	Mar 29, 2005

ISOCARBOXAZID

TABLET; ORAL
MARPLAN

+ OXFORD PHARM	10MG	N11961	001	
----------------	------	--------	-----	--

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION
BETA-2

+ NEPHRON	1%	N86711	001	
-----------	----	--------	-----	--

ISOFLURANE

LIQUID; INHALATION

FORANE

<u>AN</u> + BAXTER HLTHCARE CORP	<u>99.9%</u>	<u>N17624</u>	<u>001</u>	
----------------------------------	--------------	---------------	------------	--

ISOFLURANE

<u>AN</u> HALOCARBON PRODS	<u>99.9%</u>	<u>N75225</u>	<u>001</u>	Oct 20, 1999
<u>AN</u> HOSPIRA	<u>99.9%</u>	<u>N74097</u>	<u>001</u>	Jan 25, 1993
<u>AN</u> MARSAM PHARMS LLC	<u>99.9%</u>	<u>N74393</u>	<u>001</u>	May 12, 1995
<u>AN</u> MINRAD	<u>99.9%</u>	<u>N74416</u>	<u>001</u>	Sep 30, 1994
<u>AN</u> RHODIA	<u>99.9%</u>	<u>N74502</u>	<u>001</u>	Jun 27, 1995

ISONIAZID

INJECTABLE; INJECTION

ISONIAZID

<u>AP</u> SABEX 2002	<u>100MG/ML</u>	<u>N40648</u>	<u>001</u>	Jul 05, 2005
----------------------	-----------------	---------------	------------	--------------

NYDRAZID

<u>AP</u> + SANDOZ	<u>100MG/ML</u>	<u>N08662</u>	<u>001</u>	
--------------------	-----------------	---------------	------------	--

SYRUP; ORAL

ISONIAZID

+ CAROLINA MEDCL	50MG/5ML	N88235	001	Nov 10, 1983
------------------	----------	--------	-----	--------------

TABLET; ORAL

ISONIAZID

<u>AA</u> BARR	<u>100MG</u>	<u>N80936</u>	<u>001</u>	
<u>AA</u>	<u>300MG</u>	<u>N80937</u>	<u>002</u>	
<u>AA</u> DURAMED PHARMS BARR	<u>300MG</u>	<u>N88119</u>	<u>001</u>	Mar 17, 1983
<u>AA</u> MIKART	<u>100MG</u>	<u>N40090</u>	<u>001</u>	Jun 26, 1997
<u>AA</u>	<u>300MG</u>	<u>N40090</u>	<u>002</u>	Jun 26, 1997
<u>AA</u> MUTUAL PHARM	<u>300MG</u>	<u>N83633</u>	<u>001</u>	
<u>AA</u> + SANDOZ	<u>100MG</u>	<u>N08678</u>	<u>002</u>	
<u>AA</u> +	<u>300MG</u>	<u>N08678</u>	<u>003</u>	
<u>AA</u> WATSON LABS	<u>300MG</u>	<u>N80521</u>	<u>001</u>	
<u>AA</u> WEST WARD	<u>100MG</u>	<u>N80212</u>	<u>001</u>	
<u>AA</u>	<u>300MG</u>	<u>N87425</u>	<u>001</u>	

LANIAZID

<u>AA</u> LANNETT	<u>100MG</u>	<u>N80140</u>	<u>002</u>	
<u>AA</u>	<u>300MG</u>	<u>N89776</u>	<u>001</u>	Jun 13, 1988
	50MG	N80140	001	

ISONIAZID; PYRAZINAMIDE; RIFAMPIN

TABLET; ORAL

RIFATER

+ AVENTIS PHARMS	50MG;300MG;120MG	N50705	001	May 31, 1994
------------------	------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 210 (of 360)

ISONIAZID; RIFAMPIN

CAPSULE; ORAL

RIFAMATE

<u>AB</u>	+	AVENTIS PHARMS	<u>150MG;300MG</u>	<u>N61884</u>	<u>001</u>	
-----------	---	----------------	--------------------	---------------	------------	--

RIFAMPIN AND ISONIAZID

<u>AB</u>		WESTWARD	<u>150MG;300MG</u>	<u>N65221</u>	<u>001</u>	Jul 29, 2005
-----------	--	----------	--------------------	---------------	------------	--------------

ISOPROTERENOL HYDROCHLORIDE

INJECTABLE; INJECTION

ISOPROTERENOL HYDROCHLORIDE

<u>AP</u>		HOSPIRA	<u>0.2MG/ML</u>	<u>N83346</u>	<u>001</u>	
			0.02MG/ML	N83283	001	

<u>AP</u>		INTL MEDICATION	<u>0.2MG/ML</u>	<u>N83724</u>	<u>001</u>	
-----------	--	-----------------	-----------------	---------------	------------	--

ISUPREL

<u>AP</u>	+	HOSPIRA	<u>0.2MG/ML</u>	<u>N10515</u>	<u>001</u>	
-----------	---	---------	-----------------	---------------	------------	--

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL

DILATRATE-SR

	+	SCHWARZ PHARMA	40MG	N19790	001	Sep 02, 1988
--	---	----------------	------	--------	-----	--------------

TABLET; ORAL

ISORDIL

<u>AB</u>		BIOVAIL	<u>5MG</u>	<u>N12093</u>	<u>007</u>	Jul 29, 1988
-----------	--	---------	------------	---------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>N12093</u>	<u>002</u>	Jul 29, 1988
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>20MG</u>	<u>N12093</u>	<u>006</u>	Jul 29, 1988
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>	+		<u>30MG</u>	<u>N12093</u>	<u>005</u>	Jul 29, 1988
-----------	---	--	-------------	---------------	------------	--------------

	+		40MG	N12093	001	Jul 29, 1988
--	---	--	------	--------	-----	--------------

ISOSORBIDE DINITRATE

<u>AB</u>		PAR PHARM	<u>5MG</u>	<u>N86923</u>	<u>001</u>	Mar 12, 1987
-----------	--	-----------	------------	---------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>N86925</u>	<u>001</u>	Mar 12, 1987
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>20MG</u>	<u>N87537</u>	<u>001</u>	Oct 02, 1987
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>30MG</u>	<u>N87946</u>	<u>001</u>	Jan 12, 1988
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>5MG</u>	<u>N86221</u>	<u>001</u>	Jan 07, 1988
-----------	--	--------	------------	---------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>N86223</u>	<u>001</u>	Jan 07, 1988
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>20MG</u>	<u>N89367</u>	<u>001</u>	Apr 07, 1988
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>		WATSON LABS	<u>5MG</u>	<u>N86034</u>	<u>001</u>	Jan 06, 1988
-----------	--	-------------	------------	---------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>N86032</u>	<u>001</u>	Jan 07, 1988
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>		WEST WARD	<u>5MG</u>	<u>N86067</u>	<u>001</u>	Oct 29, 1987
-----------	--	-----------	------------	---------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>N86066</u>	<u>001</u>	Oct 29, 1987
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>20MG</u>	<u>N88088</u>	<u>001</u>	Nov 02, 1987
-----------	--	--	-------------	---------------	------------	--------------

TABLET; SUBLINGUAL

ISORDIL

<u>AB</u>		BIOVAIL	<u>2.5MG</u>	<u>N12940</u>	<u>004</u>	Jul 29, 1988
-----------	--	---------	--------------	---------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>N12940</u>	<u>003</u>	Jul 29, 1988
-----------	--	--	------------	---------------	------------	--------------

	+		10MG	N12940	005	Jul 29, 1988
--	---	--	------	--------	-----	--------------

ISOSORBIDE DINITRATE

<u>AB</u>		SANDOZ	<u>2.5MG</u>	<u>N86225</u>	<u>001</u>	Feb 19, 1988
-----------	--	--------	--------------	---------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>N86222</u>	<u>001</u>	Feb 19, 1988
-----------	--	--	------------	---------------	------------	--------------

<u>AB</u>		WATSON LABS	<u>2.5MG</u>	<u>N86033</u>	<u>001</u>	Feb 26, 1988
-----------	--	-------------	--------------	---------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>N86031</u>	<u>001</u>	Sep 29, 1987
-----------	--	--	------------	---------------	------------	--------------

<u>AB</u>		WEST WARD	<u>2.5MG</u>	<u>N86054</u>	<u>001</u>	Oct 29, 1987
-----------	--	-----------	--------------	---------------	------------	--------------

<u>AB</u>			<u>5MG</u>	<u>N86055</u>	<u>001</u>	Nov 02, 1987
-----------	--	--	------------	---------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

ISOSORBIDE DINITRATE

	+	INWOOD LABS	40MG	N40009	001	Dec 30, 1998
--	---	-------------	------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 211 (of 360)

ISOSORBIDE MONONITRATE

TABLET; ORAL

ISMO

<u>AB</u>	ESP PHARMA	<u>20MG</u>	<u>N19091</u>	<u>001</u>	Dec 30, 1991
<u>ISOSORBIDE MONONITRATE</u>					
<u>AB</u>	PUREPAC PHARM	<u>10MG</u>	<u>N75037</u>	<u>002</u>	Oct 30, 1998
<u>AB</u>		<u>20MG</u>	<u>N75037</u>	<u>001</u>	Oct 30, 1998
<u>AB</u>	TEVA	<u>20MG</u>	<u>N75147</u>	<u>001</u>	Nov 27, 1998
<u>AB</u>	WEST WARD	<u>20MG</u>	<u>N75361</u>	<u>001</u>	Oct 05, 2000
<u>MONOKET</u>					
<u>AB</u>	SCHWARZ	<u>10MG</u>	<u>N20215</u>	<u>002</u>	Jun 30, 1993
<u>AB</u>	+	<u>20MG</u>	<u>N20215</u>	<u>001</u>	Jun 30, 1993

TABLET, EXTENDED RELEASE; ORAL

IMDUR

<u>AB</u>	SCHERING PLOUGH	<u>30MG</u>	<u>N20225</u>	<u>001</u>	Aug 12, 1993
<u>AB</u>		<u>60MG</u>	<u>N20225</u>	<u>002</u>	Aug 12, 1993
<u>AB</u>	+	<u>120MG</u>	<u>N20225</u>	<u>003</u>	Mar 30, 1995
<u>ISOSORBIDE MONONITRATE</u>					
<u>AB</u>	BRIGHTSTONE	<u>60MG</u>	<u>N75166</u>	<u>001</u>	Oct 07, 1999
<u>AB</u>	DEXCEL LTD	<u>60MG</u>	<u>N75522</u>	<u>001</u>	Apr 17, 2000
<u>AB</u>	ELAN PHARM	<u>60MG</u>	<u>N75041</u>	<u>001</u>	Sep 22, 1998
<u>AB</u>	KREMERS URBAN	<u>30MG</u>	<u>N75155</u>	<u>002</u>	Jan 13, 2000
<u>AB</u>		<u>60MG</u>	<u>N75155</u>	<u>001</u>	Oct 30, 1998
<u>AB</u>		<u>120MG</u>	<u>N75155</u>	<u>003</u>	Aug 04, 2000
<u>AB</u>	KV PHARM	<u>30MG</u>	<u>N75395</u>	<u>001</u>	Mar 16, 2000
<u>AB</u>		<u>60MG</u>	<u>N75395</u>	<u>002</u>	Mar 16, 2000
<u>AB</u>		<u>120MG</u>	<u>N75395</u>	<u>003</u>	Mar 16, 2000
<u>AB</u>	PUREPAC PHARM	<u>30MG</u>	<u>N75306</u>	<u>001</u>	Dec 31, 1998
<u>AB</u>		<u>60MG</u>	<u>N75306</u>	<u>002</u>	Dec 31, 1998
<u>AB</u>	WEST WARD	<u>60MG</u>	<u>N76813</u>	<u>001</u>	Jan 07, 2005

ISOSULFAN BLUE

INJECTABLE; INJECTION

LYMPHAZURIN

+ US SURGCL 1% N18310 001

ISOTRETINOIN

CAPSULE; ORAL

ACCUTANE

<u>AB</u>	HLR	<u>10MG</u>	<u>N18662</u>	<u>002</u>	May 07, 1982
<u>AB</u>	+	<u>20MG</u>	<u>N18662</u>	<u>004</u>	Mar 28, 1983
<u>AB</u>	+	<u>40MG</u>	<u>N18662</u>	<u>003</u>	May 07, 1982
<u>AMNESTEEM</u>					
<u>AB</u>	GENPHARM	<u>10MG</u>	<u>N75945</u>	<u>001</u>	Nov 08, 2002
<u>AB</u>		<u>20MG</u>	<u>N75945</u>	<u>002</u>	Nov 08, 2002
<u>AB</u>		<u>40MG</u>	<u>N75945</u>	<u>003</u>	Nov 08, 2002
<u>CLARAVIS</u>					
<u>AB</u>	BARR	<u>10MG</u>	<u>N76356</u>	<u>001</u>	Apr 11, 2003
<u>AB</u>		<u>20MG</u>	<u>N76135</u>	<u>002</u>	Apr 11, 2003
<u>AB</u>		<u>40MG</u>	<u>N76135</u>	<u>001</u>	Apr 11, 2003
<u>SOTRET</u>					
<u>AB</u>	RANBAXY	<u>10MG</u>	<u>N76041</u>	<u>001</u>	Dec 24, 2002
<u>AB</u>		<u>20MG</u>	<u>N76041</u>	<u>002</u>	Dec 24, 2002
<u>AB</u>		<u>40MG</u>	<u>N76041</u>	<u>003</u>	Dec 24, 2002
		30MG	N76503	001	Jun 20, 2003

ISRADIPINE

CAPSULE; ORAL

DYNACIRC

<u>AB</u>	+	RELIAANT PHARMS	<u>2.5MG</u>	<u>N19546</u>	<u>001</u>	Dec 20, 1990
-----------	---	-----------------	--------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 212 (of 360)

ISRADIPINE

CAPSULE; ORAL

ISRADIPINE

<u>AB</u>	ABRIKA PHARMS	<u>2.5MG</u>	<u>N77317</u>	<u>001</u>	Jan 05, 2006
		5MG	N77317	002	Jan 05, 2006

TABLET, EXTENDED RELEASE; ORAL

DYNACIRC CR

	RELIANT PHARMS	5MG	N20336	001	Jun 01, 1994
+		10MG	N20336	002	Jun 01, 1994

ITRACONAZOLE

CAPSULE; ORAL

ITRACONAZOLE

<u>AB</u>	SANDOZ	<u>100MG</u>	<u>N76104</u>	<u>001</u>	May 28, 2004
-----------	--------	--------------	---------------	------------	--------------

SPORANOX

<u>AB</u>	+ JANSSEN PHARMA	<u>100MG</u>	<u>N20083</u>	<u>001</u>	Sep 11, 1992
-----------	------------------	--------------	---------------	------------	--------------

INJECTABLE; INJECTION

SPORANOX

+	JANSSEN PHARMA	10MG/ML	N20966	001	Mar 30, 1999
---	----------------	---------	--------	-----	--------------

SOLUTION; ORAL

SPORANOX

+	JANSSEN PHARMA	10MG/ML	N20657	001	Feb 21, 1997
---	----------------	---------	--------	-----	--------------

IVERMECTIN

TABLET; ORAL

STROMEKTOL

	MERCK	3MG	N50742	002	Oct 08, 1998
+		6MG	N50742	001	Nov 22, 1996

KANAMYCIN SULFATE

INJECTABLE; INJECTION

KANAMYCIN

<u>AP</u>	AM PHARM PARTNERS	<u>EQ 500MG BASE/2ML</u>	<u>N65111</u>	<u>001</u>	Dec 17, 2002
-----------	-------------------	--------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 1GM BASE/3ML</u>	<u>N65111</u>	<u>002</u>	Dec 17, 2002
-----------	--	------------------------	---------------	------------	--------------

KANTREX

<u>AP</u>	+ APOTHECON	<u>EQ 500MG BASE/2ML</u>	<u>N61901</u>	<u>001</u>	
-----------	-------------	--------------------------	---------------	------------	--

<u>AP</u>	+	<u>EQ 1GM BASE/3ML</u>	<u>N61901</u>	<u>002</u>	
-----------	---	------------------------	---------------	------------	--

	+	EQ 75MG BASE/2ML	N61901	003	
--	---	------------------	--------	-----	--

KETAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

KETALAR

<u>AP</u>	+ PARKEDALE	<u>EQ 50MG BASE/ML</u>	<u>N16812</u>	<u>002</u>	
-----------	-------------	------------------------	---------------	------------	--

<u>AP</u>	+	<u>EQ 100MG BASE/ML</u>	<u>N16812</u>	<u>003</u>	
-----------	---	-------------------------	---------------	------------	--

	+	EQ 10MG BASE/ML	N16812	001	
--	---	-----------------	--------	-----	--

KETAMINE HYDROCHLORIDE

<u>AP</u>	BEDFORD	<u>EQ 50MG BASE/ML</u>	<u>N74524</u>	<u>001</u>	Mar 22, 1996
-----------	---------	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 100MG BASE/ML</u>	<u>N74524</u>	<u>002</u>	Mar 22, 1996
-----------	--	-------------------------	---------------	------------	--------------

<u>AP</u>	BIONICHE (CANADA)	<u>EQ 50MG BASE/ML</u>	<u>N76092</u>	<u>002</u>	Dec 28, 2001
-----------	-------------------	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 100MG BASE/ML</u>	<u>N76092</u>	<u>003</u>	Oct 25, 2002
-----------	--	-------------------------	---------------	------------	--------------

<u>AP</u>	HOSPIRA	<u>EQ 50MG BASE/ML</u>	<u>N74549</u>	<u>001</u>	Jun 27, 1996
-----------	---------	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 100MG BASE/ML</u>	<u>N74549</u>	<u>002</u>	Jun 27, 1996
-----------	--	-------------------------	---------------	------------	--------------

KETOCONAZOLE

CREAM; TOPICAL

KETOCONAZOLE

<u>AB</u>	ALTANA	<u>2%</u>	<u>N76294</u>	<u>001</u>	Apr 28, 2004
-----------	--------	-----------	---------------	------------	--------------

<u>AB</u>	+ TEVA	<u>2%</u>	<u>N75581</u>	<u>001</u>	Apr 25, 2000
-----------	--------	-----------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 213 (of 360)

KETOCONAZOLE

CREAM; TOPICAL

KETOZOLE

<u>AB</u>	TARO	<u>2%</u>	<u>N75638</u>	<u>001</u>	Dec 18, 2002
-----------	------	-----------	---------------	------------	--------------

SHAMPOO; TOPICAL

KETOCONAZOLE

<u>AB</u>	CLAY PARK	<u>2%</u>	<u>N76419</u>	<u>001</u>	Jan 07, 2004
-----------	-----------	-----------	---------------	------------	--------------

<u>AB</u>	QLT USA	<u>2%</u>	<u>N76942</u>	<u>001</u>	Apr 11, 2005
-----------	---------	-----------	---------------	------------	--------------

NIZORAL

<u>AB</u>	+ MCNEIL CONS SPECLT	<u>2%</u>	<u>N19927</u>	<u>001</u>	Aug 31, 1990
-----------	----------------------	-----------	---------------	------------	--------------

TABLET; ORAL

KETOCONAZOLE

<u>AB</u>	AAIPHARMA LLC	<u>200MG</u>	<u>N75341</u>	<u>001</u>	Jul 27, 1999
-----------	---------------	--------------	---------------	------------	--------------

<u>AB</u>	MUTUAL PHARMA	<u>200MG</u>	<u>N75314</u>	<u>001</u>	Jun 15, 1999
-----------	---------------	--------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>200MG</u>	<u>N75597</u>	<u>001</u>	Dec 23, 1999
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>	PLIVA	<u>200MG</u>	<u>N75362</u>	<u>001</u>	Jun 15, 1999
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>	TARO	<u>200MG</u>	<u>N75319</u>	<u>001</u>	Jun 15, 1999
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>200MG</u>	<u>N74971</u>	<u>001</u>	Jun 15, 1999
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>200MG</u>	<u>N75273</u>	<u>001</u>	Jun 15, 1999
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	TORPHARM	<u>200MG</u>	<u>N75912</u>	<u>001</u>	Jan 10, 2002
-----------	----------	--------------	---------------	------------	--------------

NIZORAL

<u>AB</u>	+ JANSSEN PHARMA	<u>200MG</u>	<u>N18533</u>	<u>001</u>	
-----------	------------------	--------------	---------------	------------	--

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

<u>AB</u>	CLONMEL HLTHCARE	<u>25MG</u>	<u>N74014</u>	<u>001</u>	Jan 29, 1993
-----------	------------------	-------------	---------------	------------	--------------

<u>AB</u>		<u>50MG</u>	<u>N74014</u>	<u>002</u>	Jan 29, 1993
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>75MG</u>	<u>N74014</u>	<u>003</u>	Jan 29, 1993
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>50MG</u>	<u>N74035</u>	<u>002</u>	Dec 31, 1996
-----------	-------	-------------	---------------	------------	--------------

<u>AB</u>		<u>75MG</u>	<u>N74035</u>	<u>003</u>	Dec 31, 1996
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N74024</u>	<u>001</u>	Dec 29, 1995
-----------	--------	-------------	---------------	------------	--------------

<u>AB</u>		<u>75MG</u>	<u>N74024</u>	<u>002</u>	Dec 29, 1995
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>25MG</u>	<u>N73515</u>	<u>001</u>	Dec 22, 1992
-----------	------	-------------	---------------	------------	--------------

<u>AB</u>		<u>50MG</u>	<u>N73516</u>	<u>001</u>	Dec 22, 1992
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	+	<u>75MG</u>	<u>N73517</u>	<u>001</u>	Dec 22, 1992
-----------	---	-------------	---------------	------------	--------------

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

<u>AB</u>	ANDRX PHARMS	<u>100MG</u>	<u>N75270</u>	<u>002</u>	Mar 24, 1999
-----------	--------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>N75270</u>	<u>003</u>	Mar 24, 1999
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>200MG</u>	<u>N75270</u>	<u>001</u>	Mar 24, 1999
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	ELAN PHARM	<u>200MG</u>	<u>N74879</u>	<u>001</u>	Dec 10, 1997
-----------	------------	--------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>100MG</u>	<u>N75679</u>	<u>003</u>	Feb 20, 2002
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>		<u>150MG</u>	<u>N75679</u>	<u>002</u>	Feb 20, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>200MG</u>	<u>N75679</u>	<u>001</u>	Feb 20, 2002
-----------	--	--------------	---------------	------------	--------------

ORUVAIL

<u>AB</u>	+ WYETH PHARMS INC	<u>200MG</u>	<u>N19816</u>	<u>001</u>	Sep 24, 1993
-----------	--------------------	--------------	---------------	------------	--------------

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

<u>AP</u>	AM PHARM PARTNERS	<u>15MG/ML</u>	<u>N75784</u>	<u>001</u>	Jan 11, 2002
-----------	-------------------	----------------	---------------	------------	--------------

<u>AP</u>		<u>30MG/ML</u>	<u>N75784</u>	<u>002</u>	Jan 11, 2002
-----------	--	----------------	---------------	------------	--------------

<u>AP</u>	AMPHASTAR PHARM	<u>15MG/ML</u>	<u>N76209</u>	<u>001</u>	Jul 21, 2004
-----------	-----------------	----------------	---------------	------------	--------------

<u>AP</u>		<u>30MG/ML</u>	<u>N76209</u>	<u>002</u>	Jul 21, 2004
-----------	--	----------------	---------------	------------	--------------

<u>AP</u>	APOTEX	<u>15MG/ML</u>	<u>N75631</u>	<u>002</u>	Jun 29, 2001
-----------	--------	----------------	---------------	------------	--------------

<u>AP</u>		<u>15MG/ML</u>	<u>N76722</u>	<u>001</u>	Jul 27, 2004
-----------	--	----------------	---------------	------------	--------------

<u>AP</u>		<u>30MG/ML</u>	<u>N75626</u>	<u>001</u>	Jul 24, 2001
-----------	--	----------------	---------------	------------	--------------

<u>AP</u>		<u>30MG/ML</u>	<u>N75631</u>	<u>001</u>	Jun 29, 2001
-----------	--	----------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 214 (of 360)

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

<u>AP</u>	APOTEX	<u>30MG/ML</u>	<u>N76722</u>	<u>002</u>	Jul 27, 2004
<u>AP</u>		<u>30MG/ML</u>	<u>N77201</u>	<u>001</u>	Oct 14, 2005
<u>AP</u>	BAXTER HLTHCARE	<u>15MG/ML</u>	<u>N75772</u>	<u>001</u>	Jul 21, 2004
<u>AP</u>		<u>30MG/ML</u>	<u>N75772</u>	<u>002</u>	Jul 21, 2004
<u>AP</u>	BAXTER HLTHCARE CORP	<u>15MG/ML</u>	<u>N75299</u>	<u>001</u>	Nov 03, 1999
<u>AP</u>		<u>30MG/ML</u>	<u>N75299</u>	<u>002</u>	Nov 03, 1999
<u>AP</u>	+ BEDFORD	<u>15MG/ML</u>	<u>N75222</u>	<u>001</u>	Apr 26, 1999
<u>AP</u>	+	<u>30MG/ML</u>	<u>N75222</u>	<u>002</u>	Apr 26, 1999
<u>AP</u>	+	<u>30MG/ML</u>	<u>N75228</u>	<u>001</u>	Apr 26, 1999
<u>AP</u>	HOSPIRA	<u>15MG/ML</u>	<u>N74802</u>	<u>001</u>	Jun 05, 1997
<u>AP</u>		<u>15MG/ML</u>	<u>N74993</u>	<u>001</u>	Jan 27, 1999
<u>AP</u>		<u>30MG/ML</u>	<u>N74802</u>	<u>002</u>	Jun 05, 1997
<u>AP</u>		<u>30MG/ML</u>	<u>N74993</u>	<u>002</u>	Jan 27, 1999
<u>AP</u>	SABEX 2002	<u>15MG/ML</u>	<u>N76271</u>	<u>001</u>	Oct 06, 2004
<u>AP</u>		<u>30MG/ML</u>	<u>N76271</u>	<u>002</u>	Oct 06, 2004

SOLUTION/DROPS; OPHTHALMIC

ACULAR

+ ALLERGAN 0.5% N19700 001 Nov 09, 1992

ACULAR LS

+ ALLERGAN 0.4% N21528 001 May 30, 2003

ACULAR PRESERVATIVE FREE

+ ALLERGAN 0.5% N20811 001 Nov 03, 1997

TABLET; ORAL

KETOROLAC TROMETHAMINE

<u>AB</u>	MYLAN	<u>10MG</u>	<u>N74761</u>	<u>001</u>	May 16, 1997
<u>AB</u>	PLIVA	<u>10MG</u>	<u>N75284</u>	<u>001</u>	Jun 23, 1999
<u>AB</u>	TEVA	<u>10MG</u>	<u>N74754</u>	<u>001</u>	May 16, 1997
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N74955</u>	<u>001</u>	Sep 19, 1997
	<u>TORADOL</u>				
<u>AB</u>	+ ROCHE PALO	<u>10MG</u>	<u>N19645</u>	<u>001</u>	Dec 20, 1991

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

ZADITOR

+ NOVARTIS EQ 0.025% BASE N21066 001 Jul 02, 1999

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

<u>AP</u>	APOTEX	<u>5MG/ML</u>	<u>N76051</u>	<u>001</u>	Jul 05, 2002
<u>AP</u>	BEDFORD	<u>5MG/ML</u>	<u>N75303</u>	<u>001</u>	May 28, 1999
<u>AP</u>	HOSPIRA	<u>5MG/ML</u>	<u>N75239</u>	<u>001</u>	Nov 29, 1999
<u>AP</u>		<u>5MG/ML</u>	<u>N75240</u>	<u>001</u>	Nov 29, 1999
<u>AP</u>	MAYNE PHARMA USA	<u>5MG/ML</u>	<u>N75242</u>	<u>001</u>	Sep 30, 1999
<u>AP</u>	TAYLOR	<u>5MG/ML</u>	<u>N75431</u>	<u>001</u>	Nov 29, 1999
<u>AP</u>		<u>5MG/ML</u>	<u>N75524</u>	<u>001</u>	Nov 29, 1999
	<u>TRANDATE</u>				
<u>AP</u>	+ PROMETHEUS LABS	<u>5MG/ML</u>	<u>N19425</u>	<u>001</u>	Dec 31, 1985

TABLET; ORAL

LABETALOL HYDROCHLORIDE

<u>AB</u>	IVAX PHARMS	<u>100MG</u>	<u>N74787</u>	<u>001</u>	Aug 03, 1998
<u>AB</u>		<u>200MG</u>	<u>N74787</u>	<u>002</u>	Aug 03, 1998
<u>AB</u>		<u>300MG</u>	<u>N74787</u>	<u>003</u>	Aug 03, 1998
<u>AB</u>	MUTUAL PHARM	<u>100MG</u>	<u>N75215</u>	<u>001</u>	Jul 29, 1999
<u>AB</u>		<u>200MG</u>	<u>N75215</u>	<u>002</u>	Jul 29, 1999
<u>AB</u>		<u>300MG</u>	<u>N75215</u>	<u>003</u>	Jul 29, 1999

PRESCRIPTION DRUG PRODUCT LIST

3 - 215 (of 360)

LABETALOL HYDROCHLORIDE

TABLET; ORAL

LABETALOL HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>100MG</u>	<u>N75113</u>	<u>001</u>	Aug 04, 1998
<u>AB</u>		<u>200MG</u>	<u>N75113</u>	<u>002</u>	Aug 04, 1998
<u>AB</u>		<u>300MG</u>	<u>N75113</u>	<u>003</u>	Aug 04, 1998
<u>AB</u>	TEVA	<u>100MG</u>	<u>N74989</u>	<u>001</u>	Sep 30, 1998
<u>AB</u>		<u>200MG</u>	<u>N74989</u>	<u>002</u>	Sep 30, 1998
<u>AB</u>		<u>300MG</u>	<u>N74989</u>	<u>003</u>	Sep 30, 1998
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N75133</u>	<u>001</u>	Aug 03, 1998
<u>AB</u>		<u>200MG</u>	<u>N75133</u>	<u>002</u>	Aug 03, 1998
<u>AB</u>		<u>300MG</u>	<u>N75133</u>	<u>003</u>	Aug 03, 1998
	<u>TRANDATE</u>				
<u>AB</u>	PROMETHEUS LABS	<u>100MG</u>	<u>N18716</u>	<u>001</u>	May 24, 1985
<u>AB</u>		<u>200MG</u>	<u>N18716</u>	<u>002</u>	Aug 01, 1984
<u>AB</u>	+	<u>300MG</u>	<u>N18716</u>	<u>003</u>	Aug 01, 1984

LACTULOSE

FOR SOLUTION; ORAL
LACTULOSE

+	INALCO	10GM/PACKET	N74712	001	Dec 10, 1997
+		20GM/PACKET	N74712	002	Dec 10, 1997

SOLUTION; ORAL

CONSTILAC

<u>AA</u>	ALRA	<u>10GM/15ML</u>	<u>N71054</u>	<u>001</u>	Jul 26, 1988
-----------	------	------------------	---------------	------------	--------------

CONSTULOSE

<u>AA</u>	+	ALPHARMA US PHARMS	<u>10GM/15ML</u>	<u>N70288</u>	<u>001</u>	Aug 15, 1988
-----------	---	--------------------	------------------	---------------	------------	--------------

EVALOSE

<u>AA</u>	TEVA PHARMS	<u>10GM/15ML</u>	<u>N73497</u>	<u>001</u>	May 28, 1993
-----------	-------------	------------------	---------------	------------	--------------

LACTULOSE

<u>AA</u>	HI TECH PHARMA	<u>10GM/15ML</u>	<u>N74076</u>	<u>001</u>	Jul 03, 1995
<u>AA</u>	MORTON GROVE	<u>10GM/15ML</u>	<u>N74602</u>	<u>001</u>	Nov 14, 1996
<u>AA</u>	NOVEX	<u>10GM/15ML</u>	<u>N75911</u>	<u>001</u>	Feb 21, 2002
<u>AA</u>	PHARM ASSOC	<u>10GM/15ML</u>	<u>N74623</u>	<u>001</u>	Jul 30, 1996
<u>AA</u>	ROXANE	<u>10GM/15ML</u>	<u>N73591</u>	<u>001</u>	May 29, 1992
<u>AA</u>	VINTAGE PHARMS	<u>10GM/15ML</u>	<u>N75993</u>	<u>001</u>	Jul 26, 2001
<u>AA</u>	VISTAPHARM	<u>10GM/15ML</u>	<u>N74138</u>	<u>001</u>	Sep 30, 1992

LAXILOSE

<u>AA</u>	TECHNILAB	<u>10GM/15ML</u>	<u>N73686</u>	<u>001</u>	May 28, 1993
-----------	-----------	------------------	---------------	------------	--------------

SOLUTION; ORAL, RECTAL

ACILAC

<u>AA</u>	TECHNILAB	<u>10GM/15ML</u>	<u>N73685</u>	<u>001</u>	May 28, 1993
-----------	-----------	------------------	---------------	------------	--------------

CHOLAC

<u>AA</u>	ALRA	<u>10GM/15ML</u>	<u>N71331</u>	<u>001</u>	Jul 26, 1988
-----------	------	------------------	---------------	------------	--------------

ENULOSE

<u>AA</u>	+	ALPHARMA US PHARMS	<u>10GM/15ML</u>	<u>N71548</u>	<u>001</u>	Aug 15, 1988
-----------	---	--------------------	------------------	---------------	------------	--------------

GENERLAC

<u>AA</u>	MORTON GROVE	<u>10GM/15ML</u>	<u>N74603</u>	<u>001</u>	Oct 31, 1996
-----------	--------------	------------------	---------------	------------	--------------

HEPTALAC

<u>AA</u>	TEVA PHARMS	<u>10GM/15ML</u>	<u>N73504</u>	<u>001</u>	May 28, 1993
-----------	-------------	------------------	---------------	------------	--------------

LACTULOSE

<u>AA</u>	HI TECH PHARMA	<u>10GM/15ML</u>	<u>N74077</u>	<u>001</u>	Jul 03, 1995
<u>AA</u>	NOVEX	<u>10GM/15ML</u>	<u>N76645</u>	<u>001</u>	Jul 28, 2003

LAMIVUDINE

SOLUTION; ORAL

EPIVIR

+	GLAXOSMITHKLINE	10MG/ML	N20596	001	Nov 17, 1995
---	-----------------	---------	--------	-----	--------------

EPIVIR-HBV

+	GLAXOSMITHKLINE	5MG/ML	N21004	001	Dec 08, 1998
---	-----------------	--------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 216 (of 360)

LAMIVUDINE

TABLET; ORAL

EPIVIR

GLAXOSMITHKLINE	150MG	N20564	001	Nov 17, 1995	
+	300MG	N20564	003	Jun 24, 2002	
EPIVIR-HBV					
+	GLAXOSMITHKLINE	100MG	N21003	001	Dec 08, 1998

LAMIVUDINE; ZIDOVUDINE

TABLET; ORAL

COMBIVIR

+	GLAXOSMITHKLINE	150MG;300MG	N20857	001	Sep 26, 1997
---	-----------------	-------------	--------	-----	--------------

LAMOTRIGINE

TABLET; ORAL

LAMICTAL

+	GLAXOSMITHKLINE	25MG	N20241	005	Dec 27, 1994
		100MG	N20241	001	Dec 27, 1994
		150MG	N20241	002	Dec 27, 1994
		200MG	N20241	003	Dec 27, 1994

TABLET, CHEWABLE; ORAL

LAMICTAL CD

	GLAXOSMITHKLINE	2MG	N20764	004	Sep 08, 2000
		5MG	N20764	001	Aug 24, 1998
+		25MG	N20764	002	Aug 24, 1998

LANSOPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

PREVACID

TAP PHARM	15MG	N20406	001	May 10, 1995
+	30MG	N20406	002	May 10, 1995

FOR SUSPENSION, DELAYED RELEASE; ORAL

PREVACID

TAP PHARM	15MG/PACKET	N21281	001	May 03, 2001
+	30MG/PACKET	N21281	002	May 03, 2001

INJECTABLE; INTRAVENOUS

PREVACID IV

+	TAP PHARM	30MG/VIAL	N21566	001	May 27, 2004
---	-----------	-----------	--------	-----	--------------

TABLET, DELAYED RELEASE, ORALLY DISINTEGRATING; ORAL

PREVACID

TAP PHARM	15MG	N21428	001	Aug 30, 2002
+	30MG	N21428	002	Aug 30, 2002

LANSOPRAZOLE; NAPROXEN

CAPSULE, DELAYED REL PELLETS, TABLET; ORAL

NAPRAPAC 250 (COPACKAGED)

TAP PHARM	15MG,N/A;N/A,250MG	N21507	002	Nov 14, 2003
-----------	--------------------	--------	-----	--------------

NAPRAPAC 375 (COPACKAGED)

TAP PHARM	15MG,N/A;N/A,375MG	N21507	003	Nov 14, 2003
-----------	--------------------	--------	-----	--------------

NAPRAPAC 500 (COPACKAGED)

+	TAP PHARM	15MG,N/A;N/A,500MG	N21507	004	Nov 14, 2003
---	-----------	--------------------	--------	-----	--------------

LANTHANUM CARBONATE

TABLET, CHEWABLE; ORAL

FOSRENOL

SHIRE	EQ 250MG BASE	N21468	001	Oct 26, 2004
	EQ 500MG BASE	N21468	002	Oct 26, 2004
	EQ 750MG BASE	N21468	003	Nov 23, 2005
+	EQ 1GM BASE	N21468	004	Nov 23, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 217 (of 360)

LATANOPROST

SOLUTION/DROPS; OPHTHALMIC
XALATAN

+ PHARMACIA AND UPJOHN	0.005%	N20597	001	Jun	05, 1996
------------------------	--------	--------	-----	-----	----------

LEFLUNOMIDE

TABLET; ORAL

ARAVA

<u>AB</u> AVENTIS PHARMS	<u>10MG</u>	<u>N20905</u>	<u>001</u>	Sep	10, 1998
<u>AB</u> +	<u>20MG</u>	<u>N20905</u>	<u>002</u>	Sep	10, 1998
+	100MG	N20905	003	Sep	10, 1998

LEFLUNOMIDE

<u>AB</u> APOTEX	<u>10MG</u>	<u>N77090</u>	<u>001</u>	Sep	13, 2005
<u>AB</u>	<u>20MG</u>	<u>N77090</u>	<u>002</u>	Sep	13, 2005
<u>AB</u> BARR	<u>10MG</u>	<u>N77083</u>	<u>001</u>	Sep	13, 2005
<u>AB</u>	<u>20MG</u>	<u>N77083</u>	<u>002</u>	Sep	13, 2005
<u>AB</u> KALI LABS	<u>10MG</u>	<u>N77086</u>	<u>001</u>	Sep	13, 2005
<u>AB</u>	<u>20MG</u>	<u>N77086</u>	<u>002</u>	Sep	13, 2005
<u>AB</u> SANDOZ	<u>10MG</u>	<u>N77087</u>	<u>001</u>	Sep	13, 2005
<u>AB</u>	<u>20MG</u>	<u>N77087</u>	<u>002</u>	Sep	13, 2005
<u>AB</u> TEVA PHARMS	<u>10MG</u>	<u>N77084</u>	<u>001</u>	Sep	13, 2005
<u>AB</u>	<u>20MG</u>	<u>N77084</u>	<u>002</u>	Sep	13, 2005

LENALIDOMIDE

CAPSULE; ORAL
REVLIMID

CELGENE	5MG	N21880	001	Dec	27, 2005
+	10MG	N21880	002	Dec	27, 2005

LEPIRUDIN RECOMBINANT

INJECTABLE; INJECTION
REFLUDAN

+ BERLEX	50MG/VIAL	N20807	001	Mar	06, 1998
----------	-----------	--------	-----	-----	----------

LETROZOLE

TABLET; ORAL
FEMARA

+ NOVARTIS	2.5MG	N20726	001	Jul	25, 1997
------------	-------	--------	-----	-----	----------

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

<u>AP</u> BEDFORD	<u>EQ 50MG BASE/VIAL</u>	<u>N89384</u>	<u>001</u>	Sep	14, 1987
<u>AP</u>	<u>EQ 100MG BASE/VIAL</u>	<u>N89717</u>	<u>001</u>	Mar	28, 1988
<u>AP</u> + MAYNE PHARMA USA	<u>EQ 350MG BASE/VIAL</u>	<u>N08107</u>	<u>005</u>	Apr	05, 1989
<u>AP</u> PHARMACHEMIE	<u>EQ 350MG BASE/VIAL</u>	<u>N40262</u>	<u>001</u>	Dec	15, 1999
<u>AP</u> PHARMACHEMIE USA	<u>EQ 50MG BASE/VIAL</u>	<u>N89628</u>	<u>001</u>	Apr	17, 1997
<u>AP</u> SICOR PHARMS	<u>EQ 50MG BASE/VIAL</u>	<u>N81278</u>	<u>001</u>	Sep	28, 1993
<u>AP</u>	<u>EQ 100MG BASE/VIAL</u>	<u>N81277</u>	<u>001</u>	Sep	28, 1993
<u>AP</u>	<u>EQ 350MG BASE/VIAL</u>	<u>N40174</u>	<u>001</u>	Jun	12, 1997

LEUCOVORIN CALCIUM PRESERVATIVE FREE

<u>AP</u> AM PHARM PARTNERS	<u>EQ 200MG BASE/VIAL</u>	<u>N40258</u>	<u>001</u>	Feb	26, 1999
+	EQ 500MG BASE/VIAL	N40286	001	Feb	26, 1999
<u>AP</u> + BEDFORD	<u>EQ 10MG BASE/ML</u>	<u>N40347</u>	<u>001</u>	Apr	25, 2000
<u>AP</u> +	<u>EQ 200MG BASE/VIAL</u>	<u>N40056</u>	<u>001</u>	May	23, 1995
<u>AP</u>	<u>EQ 350MG BASE/VIAL</u>	<u>N40335</u>	<u>001</u>	Apr	20, 2000
<u>AP</u> LUITPOLD	<u>EQ 50MG BASE/VIAL</u>	<u>N40338</u>	<u>001</u>	Jan	31, 2001
<u>AP</u> SICOR PHARMS	<u>EQ 10MG BASE/ML</u>	<u>N40332</u>	<u>001</u>	Jun	28, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 218 (of 360)

LEUCOVORIN CALCIUM

TABLET; ORAL

LEUCOVORIN CALCIUM

<u>AB</u>	BARR	<u>EQ 5MG BASE</u>	<u>N71198</u>	<u>001</u>	Sep 24, 1987
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N71199</u>	<u>001</u>	Sep 24, 1987
<u>AB</u>	PAR PHARM	<u>EQ 5MG BASE</u>	<u>N74544</u>	<u>001</u>	Aug 28, 1997
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N74544</u>	<u>002</u>	Aug 28, 1997
<u>AB</u>	PHARMACHEMIE	<u>EQ 5MG BASE</u>	<u>N73099</u>	<u>001</u>	Mar 28, 1997
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N73101</u>	<u>001</u>	Mar 28, 1997
<u>AB</u>	ROXANE	<u>EQ 5MG BASE</u>	<u>N72733</u>	<u>001</u>	Feb 22, 1993
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N72734</u>	<u>001</u>	Feb 22, 1993
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>N72735</u>	<u>001</u>	Feb 22, 1993
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>N72736</u>	<u>001</u>	Feb 22, 1993
<u>AB</u>	SANDOZ	<u>EQ 15MG BASE</u>	<u>N75327</u>	<u>001</u>	Mar 24, 1999
<u>AB</u>	XANODYNE PHARM	<u>EQ 10MG BASE</u>	<u>N71962</u>	<u>001</u>	Nov 19, 1987
<u>AB</u>		<u>EQ 15MG BASE</u>	<u>N71104</u>	<u>001</u>	Mar 04, 1987
<u>BX</u>		EQ 5MG BASE	N18459	001	Jan 30, 1986

LEUPROLIDE ACETATE

IMPLANT; IMPLANTATION

VIADUR

+	ALZA	EQ 65MG BASE	N21088	001	Mar 03, 2000
---	------	--------------	--------	-----	--------------

INJECTABLE; INJECTION

LEUPROLIDE ACETATE

<u>AP</u>	BEDFORD LABS	<u>1MG/0.2ML</u>	<u>N74728</u>	<u>001</u>	Aug 04, 1998
<u>AP</u>	GENZYME	<u>1MG/0.2ML</u>	<u>N75721</u>	<u>001</u>	Nov 29, 2001
<u>AP</u>	SICOR PHARMS	<u>1MG/0.2ML</u>	<u>N75471</u>	<u>001</u>	Oct 25, 2000

LUPRON

<u>AP</u>	+	TAP PHARM	<u>1MG/0.2ML</u>	<u>N19010</u>	<u>001</u>	Apr 09, 1985
		LUPRON DEPOT				
	+	TAP PHARM	3.75MG/VIAL	N20011	001	Oct 22, 1990
	+		7.5MG/VIAL	N19732	001	Jan 26, 1989
	+		22.5MG/VIAL	N20517	001	Dec 22, 1995
		LUPRON DEPOT-3				
	+	TAP PHARM	11.25MG/VIAL	N20708	001	Mar 07, 1997
		LUPRON DEPOT-4				
	+	TAP PHARM	30MG/VIAL	N20517	002	May 30, 1997
		LUPRON DEPOT-PED				
	+	TAP PHARM	7.5MG/VIAL	N20263	002	Apr 16, 1993
	+		11.25MG/VIAL	N20263	005	Jan 21, 1994
	+		15MG/VIAL	N20263	006	Jan 21, 1994

INJECTABLE; SUBCUTANEOUS

ELIGARD

+	QLT USA	7.5MG/VIAL	N21343	001	Jan 23, 2002
+		22.5MG/VIAL	N21379	001	Jul 24, 2002
+		30MG/VIAL	N21488	001	Feb 13, 2003
+		45MG/VIAL	N21731	001	Dec 14, 2004

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

XOPENEX

+	SEPRACOR	EQ 0.0103% BASE	N20837	003	Jan 30, 2002
+		EQ 0.021% BASE	N20837	001	Mar 25, 1999
+		EQ 0.042% BASE	N20837	002	Mar 25, 1999
+		EQ 0.25% BASE	N20837	004	Jul 18, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 219 (of 360)

LEVALBUTEROL TARTRATE

AEROSOL, METERED; INHALATION

XOPENEX HFA

+ SEPRACOR	EQ 0.045MG BASE/INH	N21730	001	Mar 11, 2005
------------	---------------------	--------	-----	--------------

LEVETIRACETAM

SOLUTION; ORAL

KEPPRA

+ UCB	100MG/ML	N21505	001	Jul 15, 2003
-------	----------	--------	-----	--------------

TABLET; ORAL

KEPPRA

UCB	250MG	N21035	001	Nov 30, 1999
	500MG	N21035	002	Nov 30, 1999
+	750MG	N21035	003	Nov 30, 1999

LEVOBUNOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AKBETA

<u>AT</u> AKORN	<u>0.25%</u>	<u>N74779</u>	<u>001</u>	Oct 29, 1996
-----------------	--------------	---------------	------------	--------------

<u>AT</u>	<u>0.5%</u>	<u>N74780</u>	<u>001</u>	Oct 29, 1996
-----------	-------------	---------------	------------	--------------

BETAGAN

<u>AT</u> + ALLERGAN	<u>0.25%</u>	<u>N19814</u>	<u>001</u>	Jun 28, 1989
----------------------	--------------	---------------	------------	--------------

<u>AT</u> +	<u>0.5%</u>	<u>N19219</u>	<u>002</u>	Dec 19, 1985
-------------	-------------	---------------	------------	--------------

LEVOBUNOLOL HYDROCHLORIDE

<u>AT</u> BAUSCH AND LOMB	<u>0.25%</u>	<u>N74307</u>	<u>001</u>	Mar 04, 1994
---------------------------	--------------	---------------	------------	--------------

<u>AT</u>	<u>0.5%</u>	<u>N74326</u>	<u>001</u>	Mar 04, 1994
-----------	-------------	---------------	------------	--------------

<u>AT</u> FALCON PHARMS	<u>0.25%</u>	<u>N74851</u>	<u>001</u>	Oct 28, 1996
-------------------------	--------------	---------------	------------	--------------

<u>AT</u>	<u>0.5%</u>	<u>N74850</u>	<u>001</u>	Oct 28, 1996
-----------	-------------	---------------	------------	--------------

<u>AT</u> NOVEX	<u>0.25%</u>	<u>N75473</u>	<u>001</u>	Aug 03, 2000
-----------------	--------------	---------------	------------	--------------

<u>AT</u>	<u>0.5%</u>	<u>N75475</u>	<u>001</u>	Aug 03, 2000
-----------	-------------	---------------	------------	--------------

LEVOCABASTINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

LIVOSTIN

+ NOVARTIS	EQ 0.05% BASE	N20219	001	Nov 10, 1993
------------	---------------	--------	-----	--------------

LEVOCARNITINE

INJECTABLE; INJECTION

CARNITOR

<u>AP</u> + SIGMA TAU	<u>200MG/ML</u>	<u>N20182</u>	<u>001</u>	Dec 16, 1992
-----------------------	-----------------	---------------	------------	--------------

LEVOCARNITINE

<u>AP</u> BEDFORD	<u>200MG/ML</u>	<u>N75567</u>	<u>001</u>	Mar 29, 2001
-------------------	-----------------	---------------	------------	--------------

<u>AP</u> LUITPOLD	<u>200MG/ML</u>	<u>N75861</u>	<u>001</u>	Jun 22, 2001
--------------------	-----------------	---------------	------------	--------------

<u>AP</u> SICOR PHARMS	<u>200MG/ML</u>	<u>N75881</u>	<u>001</u>	Mar 29, 2001
------------------------	-----------------	---------------	------------	--------------

SOLUTION; ORAL

CARNITOR

<u>AA</u> + SIGMA TAU	<u>1GM/10ML</u>	<u>N19257</u>	<u>001</u>	Apr 10, 1986
-----------------------	-----------------	---------------	------------	--------------

LEVOCARNITINE

<u>AA</u> LYNE	<u>1GM/10ML</u>	<u>N76851</u>	<u>001</u>	Aug 10, 2004
----------------	-----------------	---------------	------------	--------------

TABLET; ORAL

CARNITOR

<u>AB</u> + SIGMA TAU	<u>330MG</u>	<u>N18948</u>	<u>001</u>	Dec 27, 1985
-----------------------	--------------	---------------	------------	--------------

LEVOCARNITINE

<u>AB</u> COREPHARMA	<u>330MG</u>	<u>N76858</u>	<u>001</u>	Sep 20, 2004
----------------------	--------------	---------------	------------	--------------

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVAQUIN

+ ORTHO MCNEIL PHARM	EQ 500MG /20ML(25MG/ML)	N20635	001	Dec 20, 1996
----------------------	-------------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 220 (of 360)

LEVOFLOXACIN

INJECTABLE; INJECTION

LEVAQUIN

+	ORTHO MCNEIL PHARM	EQ 750MG /30ML(25MG/ML)	N20635	004	Dec 20, 1996
	LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER				
+	ORTHO MCNEIL PHARM	EQ 250MG /50ML(5MG/ML)	N20635	002	Dec 20, 1996
+		EQ 500MG /100ML(5MG/ML)	N20635	003	Dec 20, 1996
+		EQ 750MG /150ML(5MG/ML)	N20635	005	Dec 20, 1996

SOLUTION; ORAL

LEVAQUIN

+	ORTHO MCNEIL PHARM	250MG/10ML	N21721	001	Oct 21, 2004
---	--------------------	------------	--------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

IQUIX

+	SANTEN	1.5%	N21571	001	Mar 01, 2004
---	--------	------	--------	-----	--------------

QUIXIN

+	SANTEN	0.5%	N21199	001	Aug 18, 2000
---	--------	------	--------	-----	--------------

TABLET; ORAL

LEVAQUIN

<u>AB</u>	+	ORTHO MCNEIL PHARM	<u>750MG</u>	<u>N20634</u>	<u>003</u>	Sep 08, 2000
			250MG	N20634	001	Dec 20, 1996
			500MG	N20634	002	Dec 20, 1996

LEVOFLOXACIN

<u>AB</u>		TEVA	<u>750MG</u>	<u>N76361</u>	<u>003</u>	Jan 26, 2005
-----------	--	------	--------------	---------------	------------	--------------

LEVONORDEFIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINE W/ NEO-COBEFRIN

<u>AP</u>	+	EASTMAN KODAK	<u>0.05MG/ML;2%</u>	<u>N12125</u>	<u>002</u>	
-----------	---	---------------	---------------------	---------------	------------	--

ISOCAINE HYDROCHLORIDE W/ LEVONORDEFIN

<u>AP</u>		NOVOCOL	<u>0.05MG/ML;2%</u>	<u>N84697</u>	<u>001</u>	
-----------	--	---------	---------------------	---------------	------------	--

MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFIN

<u>AP</u>		GRAHAM CHEM	<u>0.05MG/ML;2%</u>	<u>N84850</u>	<u>002</u>	Oct 21, 1983
-----------	--	-------------	---------------------	---------------	------------	--------------

POLOCAINE W/ LEVONORDEFIN

<u>AP</u>		DENTSPLY PHARM	<u>0.05MG/ML;2%</u>	<u>N89517</u>	<u>001</u>	Apr 14, 1988
-----------	--	----------------	---------------------	---------------	------------	--------------

SCANDONEST L

<u>AP</u>		DEPROCO	<u>0.05MG/ML;2%</u>	<u>N88388</u>	<u>001</u>	Oct 10, 1984
-----------	--	---------	---------------------	---------------	------------	--------------

LEVONORGESTREL

IMPLANT; IMPLANTATION

NORPLANT II

BX	+	POPULATION COUNCIL	75MG/IMPLANT	N20544	001	Nov 01, 1996
----	---	--------------------	--------------	--------	-----	--------------

INTRAUTERINE DEVICE; INTRAUTERINE

MIRENA

+	BERLEX	52MG	N21225	001	Dec 06, 2000
---	--------	------	--------	-----	--------------

TABLET; ORAL

PLAN B

+	DURAMED	0.75MG	N21045	001	Jul 28, 1999
---	---------	--------	--------	-----	--------------

LEVORPHANOL TARTRATE

TABLET; ORAL

LEVO-DROMORAN

<u>AB</u>	+	VALEANT PHARM INTL	<u>2MG</u>	<u>N08720</u>	<u>001</u>	Dec 19, 1991
-----------	---	--------------------	------------	---------------	------------	--------------

LEVORPHANOL TARTRATE

<u>AB</u>		ROXANE	<u>2MG</u>	<u>N74278</u>	<u>001</u>	Mar 31, 2000
-----------	--	--------	------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 221 (of 360)

LEVOTHYROXINE SODIUM**

Refer to Preface Section 1.8 Levothyroxine Sodium for amplifying information

TABLET; ORAL

LEVOLET

BX	VINTAGE	0.025MG	N21137	001	Jun 06, 2003
BX		0.05MG	N21137	002	Jun 06, 2003
BX		0.075MG	N21137	003	Jun 06, 2003
BX		0.088MG	N21137	004	Jun 06, 2003
BX		0.1MG	N21137	005	Jun 06, 2003
BX		0.112MG	N21137	006	Jun 06, 2003
BX		0.125MG	N21137	007	Jun 06, 2003
BX		0.137MG	N21137	008	Jun 06, 2003
BX		0.15MG	N21137	009	Jun 06, 2003
BX		0.175MG	N21137	010	Jun 06, 2003
BX		0.2MG	N21137	011	Jun 06, 2003
BX		0.3MG	N21137	012	Jun 06, 2003

LEVO-T

-->	ALARA PHARM	<u>AB2, AB3 0.025MG</u>	<u>N21342</u>	<u>001</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.05MG</u>	<u>N21342</u>	<u>002</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.075MG</u>	<u>N21342</u>	<u>003</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.088MG</u>	<u>N21342</u>	<u>004</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.1MG</u>	<u>N21342</u>	<u>005</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.112MG</u>	<u>N21342</u>	<u>006</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.125MG</u>	<u>N21342</u>	<u>007</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.137MG</u>	<u>N21342</u>	<u>012</u>	Dec 08, 2003
-->		<u>AB2, AB3 0.15MG</u>	<u>N21342</u>	<u>008</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.175MG</u>	<u>N21342</u>	<u>009</u>	Mar 01, 2002
-->		<u>AB2, AB3 0.2MG</u>	<u>N21342</u>	<u>010</u>	Mar 01, 2002
--> +		<u>AB2, AB3 0.3MG</u>	<u>N21342</u>	<u>011</u>	Mar 01, 2002

LEVOTHROID

BX	LLOYD	0.025MG	N21116	001	Oct 24, 2002
BX		0.05MG	N21116	002	Oct 24, 2002
BX		0.075MG	N21116	003	Oct 24, 2002
BX		0.088MG	N21116	010	Oct 24, 2002
BX		0.1MG	N21116	004	Oct 24, 2002
BX		0.112MG	N21116	011	Oct 24, 2002
BX		0.125MG	N21116	005	Oct 24, 2002
BX		0.137MG	N21116	012	Dec 07, 2004
BX		0.15MG	N21116	006	Oct 24, 2002
BX		0.175MG	N21116	007	Oct 24, 2002
BX		0.2MG	N21116	008	Oct 24, 2002
BX +		0.3MG	N21116	009	Oct 24, 2002

LEVOTHYROXINE SODIUM

<u>AB2</u>	GENPHARM	<u>0.025MG</u>	<u>N76752</u>	<u>001</u>	Jun 16, 2005
<u>AB2</u>		<u>0.05MG</u>	<u>N76752</u>	<u>002</u>	Jun 16, 2005
<u>AB2</u>		<u>0.075MG</u>	<u>N76752</u>	<u>003</u>	Jun 16, 2005
<u>AB2</u>		<u>0.088MG</u>	<u>N76752</u>	<u>004</u>	Jun 16, 2005
<u>AB2</u>		<u>0.1MG</u>	<u>N76752</u>	<u>005</u>	Jun 16, 2005
<u>AB2</u>		<u>0.112MG</u>	<u>N76752</u>	<u>006</u>	Jun 16, 2005
<u>AB2</u>		<u>0.125MG</u>	<u>N76752</u>	<u>007</u>	Jun 16, 2005
<u>AB2</u>		<u>0.15MG</u>	<u>N76752</u>	<u>008</u>	Jun 16, 2005
<u>AB2</u>		<u>0.175MG</u>	<u>N76752</u>	<u>009</u>	Jun 16, 2005
<u>AB2</u>		<u>0.2MG</u>	<u>N76752</u>	<u>010</u>	Jun 16, 2005
<u>AB2</u>		<u>0.3MG</u>	<u>N76752</u>	<u>011</u>	Jun 16, 2005
-->	MYLAN	<u>AB1, AB2, AB3 0.025MG</u>	<u>N76187</u>	<u>001</u>	Jun 05, 2002
-->		<u>AB1, AB2, AB3 0.05MG</u>	<u>N76187</u>	<u>002</u>	Jun 05, 2002
-->		<u>AB1, AB2, AB3 0.075MG</u>	<u>N76187</u>	<u>003</u>	Jun 05, 2002
-->		<u>AB1, AB2, AB3 0.088MG</u>	<u>N76187</u>	<u>004</u>	Jun 05, 2002
-->		<u>AB1, AB2, AB3 0.1MG</u>	<u>N76187</u>	<u>005</u>	Jun 05, 2002
-->		<u>AB1, AB2, AB3 0.112MG</u>	<u>N76187</u>	<u>006</u>	Jun 05, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 222 (of 360)

LEVOTHYROXINE SODIUM**

Refer to Preface Section 1.8 Levothyroxine Sodium for amplifying information

TABLET; ORAL

<u>LEVOTHYROXINE SODIUM</u>					
-->	MYLAN	--> AB1,AB2,AB3 0.125MG	<u>N76187</u>	<u>007</u>	Jun 05, 2002
-->		--> AB1,AB2,AB3 0.15MG	<u>N76187</u>	<u>008</u>	Jun 05, 2002
-->		--> AB1,AB2,AB3 0.175MG	<u>N76187</u>	<u>009</u>	Jun 05, 2002
-->		--> AB1,AB2,AB3 0.2MG	<u>N76187</u>	<u>010</u>	Jun 05, 2002
-->		--> AB1,AB2,AB3 0.3MG	<u>N76187</u>	<u>011</u>	Jun 05, 2002
<u>LEVOXYL</u>					
-->	JONES PHARMA	--> AB1,AB3 0.025MG	<u>N21301</u>	<u>001</u>	May 25, 2001
-->		--> AB1,AB3 0.05MG	<u>N21301</u>	<u>002</u>	May 25, 2001
-->		--> AB1,AB3 0.075MG	<u>N21301</u>	<u>003</u>	May 25, 2001
-->		--> AB1,AB3 0.088MG	<u>N21301</u>	<u>004</u>	May 25, 2001
-->		--> AB1,AB3 0.1MG	<u>N21301</u>	<u>005</u>	May 25, 2001
-->		--> AB1,AB3 0.112MG	<u>N21301</u>	<u>006</u>	May 25, 2001
-->		--> AB1,AB3 0.125MG	<u>N21301</u>	<u>007</u>	May 25, 2001
-->		--> AB1,AB3 0.15MG	<u>N21301</u>	<u>009</u>	May 25, 2001
-->		--> AB1,AB3 0.175MG	<u>N21301</u>	<u>010</u>	May 25, 2001
-->		--> AB1,AB3 0.2MG	<u>N21301</u>	<u>011</u>	May 25, 2001
--> +		--> AB1,AB3 0.3MG	<u>N21301</u>	<u>012</u>	May 25, 2001
<u>AB3</u>		<u>0.137MG</u>	<u>N21301</u>	<u>008</u>	May 25, 2001
NOVOTHYROX					
BX	GENPHARM	0.025MG	N21292	001	May 31, 2002
BX		0.05MG	N21292	002	May 31, 2002
BX		0.075MG	N21292	003	May 31, 2002
BX		0.088MG	N21292	004	May 31, 2002
BX		0.1MG	N21292	005	May 31, 2002
BX		0.112MG	N21292	006	May 31, 2002
BX		0.125MG	N21292	007	May 31, 2002
BX		0.137MG	N21292	008	May 31, 2002
BX		0.15MG	N21292	009	May 31, 2002
BX		0.175MG	N21292	010	May 31, 2002
BX		0.2MG	N21292	011	May 31, 2002
BX +		0.3MG	N21292	012	May 31, 2002
<u>SYNTHROID</u>					
-->	ABBOTT	--> AB1,AB2 0.025MG	<u>N21402</u>	<u>001</u>	Jul 24, 2002
-->		--> AB1,AB2 0.05MG	<u>N21402</u>	<u>002</u>	Jul 24, 2002
-->		--> AB1,AB2 0.075MG	<u>N21402</u>	<u>003</u>	Jul 24, 2002
-->		--> AB1,AB2 0.088MG	<u>N21402</u>	<u>004</u>	Jul 24, 2002
-->		--> AB1,AB2 0.1MG	<u>N21402</u>	<u>005</u>	Jul 24, 2002
-->		--> AB1,AB2 0.112MG	<u>N21402</u>	<u>006</u>	Jul 24, 2002
-->		--> AB1,AB2 0.125MG	<u>N21402</u>	<u>007</u>	Jul 24, 2002
-->		--> AB1,AB2 0.15MG	<u>N21402</u>	<u>009</u>	Jul 24, 2002
-->		--> AB1,AB2 0.175MG	<u>N21402</u>	<u>010</u>	Jul 24, 2002
-->		--> AB1,AB2 0.2MG	<u>N21402</u>	<u>012</u>	Jul 24, 2002
--> +		--> AB1,AB2 0.3MG	<u>N21402</u>	<u>011</u>	Jul 24, 2002
<u>AB2</u>		<u>0.137MG</u>	<u>N21402</u>	<u>008</u>	Jul 24, 2002
<u>UNITHROID</u>					
-->	STEVENS J	--> AB1,AB2,AB3 0.025MG	<u>N21210</u>	<u>001</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.05MG	<u>N21210</u>	<u>002</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.075MG	<u>N21210</u>	<u>003</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.088MG	<u>N21210</u>	<u>004</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.1MG	<u>N21210</u>	<u>005</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.112MG	<u>N21210</u>	<u>006</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.125MG	<u>N21210</u>	<u>007</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.15MG	<u>N21210</u>	<u>008</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.175MG	<u>N21210</u>	<u>009</u>	Aug 21, 2000
-->		--> AB1,AB2,AB3 0.2MG	<u>N21210</u>	<u>010</u>	Aug 21, 2000
--> +		--> AB1,AB2,AB3 0.3MG	<u>N21210</u>	<u>011</u>	Aug 21, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 223 (of 360)

LIDOCAINE

FILM, EXTENDED RELEASE; BUCCAL					
LIDOCAINE					
AT	+	NOVEN	46.1MG/PATCH	N20575	002 May 21, 1996
OINTMENT; TOPICAL					
<u>LIDOCAINE</u>					
AT	+	FOUGERA	5%	N80198	001
AT		GRAHAM CHEM	5%	N80210	001
AT		TARO	5%	N86724	001
PATCH; TOPICAL					
LIDODERM					
	+	TEIKOKU PHARMA USA	5%	N20612	001 Mar 19, 1999

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION					
<u>LIDOCAINE HYDROCHLORIDE</u>					
AP		GRAHAM CHEM	2%	N80504	001
AP		HOSPIRA	0.5%	N88328	001 May 17, 1984
AP			1%	N40013	001 Jun 23, 1995
AP			1%	N83158	001
AP			1%	N88329	001 May 17, 1984
AP			2%	N40078	001 Jun 23, 1995
AP			2%	N83158	002
AP			2%	N88294	001 May 17, 1984
AP			2%	N88331	001 May 17, 1984
AP			20%	N83158	003
AP		INTL MEDICATION	1%	N83173	001
AP			2%	N83173	002
AP		LUITPOLD	1%	N80850	001
AP			2%	N83198	001
<u>LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
AP		B BRAUN	200MG/100ML	N19830	002 Apr 08, 1992
AP		BAXTER HLTHCARE	200MG/100ML	N18461	002
<u>LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5%</u>					
AP		HOSPIRA	200MG/100ML	N83158	005
<u>LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
AP		HOSPIRA	200MG/100ML	N18388	001
<u>LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
AP		B BRAUN	400MG/100ML	N19830	003 Apr 08, 1992
AP		BAXTER HLTHCARE	400MG/100ML	N18461	003
<u>LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5%</u>					
AP		HOSPIRA	400MG/100ML	N83158	006
<u>LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
AP		HOSPIRA	400MG/100ML	N18388	002
<u>LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
AP		B BRAUN	800MG/100ML	N19830	004 Apr 08, 1992
AP		BAXTER HLTHCARE	800MG/100ML	N18461	004 Feb 22, 1982
<u>LIDOCAINE HYDROCHLORIDE 0.8% IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
AP		HOSPIRA	800MG/100ML	N18388	003 Nov 05, 1982
<u>LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER</u>					
AP		AM PHARM PARTNERS	1%	N88586	001 Jul 24, 1985
AP		HOSPIRA	0.5%	N88325	001 Jul 31, 1984
AP			1%	N88299	001 Jul 31, 1984
AP			1.5%	N88326	001 Jul 31, 1984
AP			2%	N88327	001 Jul 31, 1984
AP			10%	N88367	001 Jul 31, 1984
AP			20%	N88368	001 Jul 31, 1984
<u>LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE</u>					
AP		AM PHARM	2%	N17584	001
AP			4%	N17584	002

PRESCRIPTION DRUG PRODUCT LIST

3 - 224 (of 360)

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION			
<u>LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE</u>			
<u>AP</u>	AM PHARM PARTNERS	<u>1%</u>	<u>N80404</u> <u>002</u>
<u>AP</u>		<u>2%</u>	<u>N80404</u> <u>003</u>
<u>AP</u>	HOSPIRA	<u>1%</u>	<u>N80408</u> <u>001</u>
<u>AP</u>		<u>1.5%</u>	<u>N80408</u> <u>002</u>
<u>AP</u>		<u>4%</u>	<u>N88295</u> <u>001</u>
<u>AP</u>	INTL MEDICATION	<u>20%</u>	<u>N17702</u> <u>001</u>
<u>LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER</u>			
<u>AP</u>	HOSPIRA	<u>1%</u>	<u>N40302</u> <u>001</u>
<u>AP</u>		<u>2%</u>	<u>N40302</u> <u>002</u>
<u>LIDOPEN</u>			
<u>AP</u>	MERIDIAN MEDCL TECHN	<u>10%</u>	<u>N17549</u> <u>001</u>
<u>XYLOCAINE</u>			
<u>AP</u>	+ ASTRAZENECA	<u>0.5%</u>	<u>N06488</u> <u>008</u>
<u>AP</u>	+	<u>1%</u>	<u>N06488</u> <u>007</u>
<u>AP</u>	+	<u>1.5%</u>	<u>N06488</u> <u>010</u>
<u>AP</u>	+ DENTSPLY PHARM	<u>2%</u>	<u>N21380</u> <u>001</u>
<u>XYLOCAINE 4% PRESERVATIVE FREE</u>			
<u>AP</u>	+ ASTRAZENECA	<u>4%</u>	<u>N10417</u> <u>001</u>
<u>XYLOCAINE PRESERVATIVE FREE</u>			
<u>AP</u>	ASTRAZENECA	<u>1%</u>	<u>N16801</u> <u>005</u>
<u>AP</u>	+	<u>2%</u>	<u>N16801</u> <u>001</u>
<u>AP</u>		<u>4%</u>	<u>N16801</u> <u>002</u>
<u>AP</u>	+	<u>10%</u>	<u>N16801</u> <u>003</u>
<u>AP</u>	+	<u>20%</u>	<u>N16801</u> <u>004</u>
INJECTABLE; SPINAL			
	LIDOCAINE HYDROCHLORIDE	5% AND DEXTROSE 7.5%	
	+ HOSPIRA	5%	N83914 001
JELLY; TOPICAL			
<u>LIDOCAINE HYDROCHLORIDE</u>			
<u>AT</u>	AKORN	<u>2%</u>	<u>N40433</u> <u>001</u>
<u>AT</u>	INTL MEDICATION	<u>2%</u>	<u>N86283</u> <u>001</u>
<u>AT</u>	TEVA PHARMS	<u>2%</u>	<u>N81318</u> <u>001</u>
<u>XYLOCAINE</u>			
<u>AT</u>	+ ASTRAZENECA	<u>2%</u>	<u>N08816</u> <u>001</u>
SOLUTION; ORAL			
<u>LIDOCAINE HYDROCHLORIDE</u>			
<u>AT</u>	HI TECH PHARMA	<u>2%</u>	<u>N40014</u> <u>001</u>
<u>AT</u>	MORTON GROVE	<u>2%</u>	<u>N87872</u> <u>001</u>
<u>LIDOCAINE HYDROCHLORIDE VISCOUS</u>			
<u>AT</u>	ALPHARMA US PHARMS	<u>2%</u>	<u>N86578</u> <u>001</u>
<u>LIDOCAINE VISCOUS</u>			
<u>AT</u>	ROXANE	<u>2%</u>	<u>N88802</u> <u>001</u>
<u>XYLOCAINE VISCOUS</u>			
<u>AT</u>	+ ASTRAZENECA	<u>2%</u>	<u>N09470</u> <u>001</u>
SOLUTION; TOPICAL			
<u>ANESTACON</u>			
<u>AT</u>	+ POLYMEDICA	<u>2%</u>	<u>N80429</u> <u>001</u>
<u>LARYNG-O-JET KIT</u>			
<u>AT</u>	INTL MEDICATION	<u>4%</u>	<u>N86364</u> <u>001</u>
<u>LIDOCAINE HYDROCHLORIDE</u>			
<u>AT</u>	MORTON GROVE	<u>4%</u>	<u>N87881</u> <u>001</u>
<u>AT</u>	ROXANE	<u>4%</u>	<u>N88803</u> <u>001</u>
<u>LTA II KIT</u>			
<u>AT</u>	HOSPIRA	<u>4%</u>	<u>N80409</u> <u>001</u>
<u>AT</u>		<u>4%</u>	<u>N88542</u> <u>001</u>
<u>PEDIATRIC LTA KIT</u>			
<u>AT</u>	HOSPIRA	<u>2%</u>	<u>N85995</u> <u>001</u>

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 225 (of 360)

LIDOCAINE HYDROCHLORIDE

SOLUTION; TOPICAL

XYLOCAINE 4% PRESERVATIVE FREE

<u>AT</u>	+	ASTRAZENECA	<u>4%</u>	<u>N10417</u>	<u>002</u>	
-----------	---	-------------	-----------	---------------	------------	--

LIDOCAINE HYDROCHLORIDE; OXYTETRACYCLINE

INJECTABLE; INJECTION

TERRAMYCIN

+	PFIZER	2%;50MG/ML	N60567	001	
---	--------	------------	--------	-----	--

+		2%;125MG/ML	N60567	002	
---	--	-------------	--------	-----	--

LIDOCAINE; PRILOCAINE

CREAM; TOPICAL

EMLA

<u>AB</u>	+	ASTRAZENECA	<u>2.5%;2.5%</u>	<u>N19941</u>	<u>001</u>	Dec 30, 1992
-----------	---	-------------	------------------	---------------	------------	--------------

LIDOCAINE AND PRILOCAINE

<u>AB</u>		ALTANA	<u>2.5%;2.5%</u>	<u>N76453</u>	<u>001</u>	Aug 18, 2003
-----------	--	--------	------------------	---------------	------------	--------------

<u>AB</u>		HI TECH PHARMA	<u>2.5%;2.5%</u>	<u>N76290</u>	<u>001</u>	Sep 25, 2003
-----------	--	----------------	------------------	---------------	------------	--------------

<u>AB</u>		QLT USA	<u>2.5%;2.5%</u>	<u>N76320</u>	<u>001</u>	Aug 27, 2003
-----------	--	---------	------------------	---------------	------------	--------------

GEL; PERIODONTAL

ORAQIX

+	DENTSPLY PHARM	2.5%;2.5%	N21451	001	Dec 19, 2003
---	----------------	-----------	--------	-----	--------------

LIDOCAINE; TETRACAINE

PATCH; TOPICAL

SYNERA

+	ZARS	70MG;70MG	N21623	001	Jun 23, 2005
---	------	-----------	--------	-----	--------------

LINCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

LINCOCIN

+	PHARMACIA AND UPJOHN	EQ 300MG BASE/ML	N50317	001	
---	----------------------	------------------	--------	-----	--

LINDANE

LOTION; TOPICAL

LINDANE

<u>AT</u>	+	ALPHARMA US PHARMS	<u>1%</u>	<u>N87313</u>	<u>001</u>	
-----------	---	--------------------	-----------	---------------	------------	--

<u>AT</u>		MORTON GROVE	<u>1%</u>	<u>N88190</u>	<u>001</u>	Aug 16, 1984
-----------	--	--------------	-----------	---------------	------------	--------------

SHAMPOO; TOPICAL

LINDANE

<u>AT</u>	+	ALPHARMA US PHARMS	<u>1%</u>	<u>N87266</u>	<u>001</u>	
-----------	---	--------------------	-----------	---------------	------------	--

<u>AT</u>		MORTON GROVE	<u>1%</u>	<u>N88191</u>	<u>001</u>	Sep 18, 1984
-----------	--	--------------	-----------	---------------	------------	--------------

LINEZOLID

FOR SUSPENSION; ORAL

ZYVOX

+	PHARMACIA AND UPJOHN	100MG/5ML	N21132	001	Apr 18, 2000
---	----------------------	-----------	--------	-----	--------------

INJECTABLE; INJECTION

ZYVOX

+	PHARMACIA AND UPJOHN	200MG/100ML	N21131	001	Apr 18, 2000
---	----------------------	-------------	--------	-----	--------------

TABLET; ORAL

ZYVOX

+	PHARMACIA AND UPJOHN	600MG	N21130	002	Apr 18, 2000
---	----------------------	-------	--------	-----	--------------

LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

LIOTHYRONINE SODIUM

<u>AP</u>		PHARMAFORCE	<u>EQ 0.01MG BASE/ML</u>	<u>N76923</u>	<u>001</u>	Aug 17, 2005
-----------	--	-------------	--------------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 226 (of 360)

LIOTHYRONINE SODIUM

INJECTABLE; INJECTION

TRIOSTAT

<u>AP</u> + JONES PHARMA	<u>EQ 0.01MG BASE/ML</u>	<u>N20105</u>	<u>001</u>	Dec 31, 1991
--------------------------	--------------------------	---------------	------------	--------------

TABLET; ORAL

CYTOMEL

KING PHARMS

EQ 0.005MG BASE

N10379 001

EQ 0.025MG BASE

N10379 002

+

EQ 0.05MG BASE

N10379 003

LIOTRIX (T4;T3)

TABLET; ORAL

THYROLAR-0.25

FOREST LABS

0.0125MG;0.0031MG

N16807 001

THYROLAR-0.5

FOREST LABS

0.025MG;0.0063MG

N16807 005

THYROLAR-1

FOREST LABS

0.05MG;0.0125MG

N16807 004

THYROLAR-2

FOREST LABS

0.1MG;0.025MG

N16807 002

THYROLAR-3

+ FOREST LABS

0.15MG;0.0375MG

N16807 003

LISINOPRIL

TABLET; ORAL

LISINOPRIL

<u>AB</u>	APOTEX	<u>2.5MG</u>	<u>N76102</u>	<u>001</u>	Sep 30, 2002
<u>AB</u>		<u>5MG</u>	<u>N76102</u>	<u>002</u>	Sep 30, 2002
<u>AB</u>		<u>10MG</u>	<u>N76102</u>	<u>003</u>	Sep 30, 2002
<u>AB</u>		<u>20MG</u>	<u>N76102</u>	<u>004</u>	Sep 30, 2002
<u>AB</u>		<u>30MG</u>	<u>N76102</u>	<u>005</u>	Sep 30, 2002
<u>AB</u>		<u>40MG</u>	<u>N76102</u>	<u>006</u>	Sep 30, 2002
<u>AB</u>	IVAX PHARMS	<u>2.5MG</u>	<u>N75752</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N75752</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75752</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N75752</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N75752</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N75752</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	LEK PHARMS	<u>2.5MG</u>	<u>N75999</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N75999</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75999</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N75999</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N75999</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N75999</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	LUPIN	<u>2.5MG</u>	<u>N77321</u>	<u>001</u>	Sep 09, 2005
<u>AB</u>		<u>5MG</u>	<u>N77321</u>	<u>002</u>	Sep 09, 2005
<u>AB</u>		<u>10MG</u>	<u>N77321</u>	<u>003</u>	Sep 09, 2005
<u>AB</u>		<u>20MG</u>	<u>N77321</u>	<u>004</u>	Sep 09, 2005
<u>AB</u>		<u>30MG</u>	<u>N77321</u>	<u>005</u>	Sep 09, 2005
<u>AB</u>		<u>40MG</u>	<u>N77321</u>	<u>006</u>	Sep 09, 2005
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>N76071</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N76071</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N76071</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N76071</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N76071</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N76071</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	PAR PHARM	<u>2.5MG</u>	<u>N75743</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N75743</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75743</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N75743</u>	<u>004</u>	Jul 01, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 227 (of 360)

LISINOPRIL

TABLET; ORAL

LISINOPRIL

<u>AB</u>	PAR PHARM	<u>30MG</u>	<u>N75743</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N75743</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	PUREPAC PHARM	<u>2.5MG</u>	<u>N76180</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N76180</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N76180</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N76164</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N76164</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N76164</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>	RANBAXY	<u>2.5MG</u>	<u>N75944</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N75944</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75944</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N75944</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N75944</u>	<u>006</u>	Feb 11, 2003
<u>AB</u>		<u>40MG</u>	<u>N75944</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>N75903</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>2.5MG</u>	<u>N75994</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N75903</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N75994</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75903</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75994</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N75903</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N75994</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N75903</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N75994</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N75903</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N75994</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	TEVA	<u>2.5MG</u>	<u>N75783</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N75783</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N75783</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N75783</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N75783</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N75783</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>N76059</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N76059</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N76059</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N76059</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N76059</u>	<u>005</u>	Jul 01, 2002
<u>AB</u>		<u>40MG</u>	<u>N76059</u>	<u>006</u>	Jul 01, 2002
<u>AB</u>	WEST WARD	<u>2.5MG</u>	<u>N76063</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>		<u>5MG</u>	<u>N76063</u>	<u>002</u>	Jul 01, 2002
<u>AB</u>		<u>10MG</u>	<u>N76063</u>	<u>003</u>	Jul 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N76063</u>	<u>004</u>	Jul 01, 2002
<u>AB</u>		<u>30MG</u>	<u>N76063</u>	<u>006</u>	Jun 27, 2003
<u>AB</u>		<u>40MG</u>	<u>N76063</u>	<u>005</u>	Jul 01, 2002
<u>PRINIVIL</u>					
<u>AB</u>	MERCK	<u>2.5MG</u>	<u>N19558</u>	<u>006</u>	Jan 28, 1994
<u>AB</u>		<u>5MG</u>	<u>N19558</u>	<u>001</u>	Dec 29, 1987
<u>AB</u>		<u>10MG</u>	<u>N19558</u>	<u>002</u>	Dec 29, 1987
<u>AB</u>		<u>20MG</u>	<u>N19558</u>	<u>003</u>	Dec 29, 1987
<u>AB</u>		<u>40MG</u>	<u>N19558</u>	<u>004</u>	Oct 25, 1988
<u>ZESTRIL</u>					
<u>AB</u>	ASTRAZENECA	<u>2.5MG</u>	<u>N19777</u>	<u>005</u>	Apr 29, 1993
<u>AB</u>		<u>5MG</u>	<u>N19777</u>	<u>001</u>	May 19, 1988
<u>AB</u>		<u>10MG</u>	<u>N19777</u>	<u>002</u>	May 19, 1988
<u>AB</u>		<u>20MG</u>	<u>N19777</u>	<u>003</u>	May 19, 1988
<u>AB</u>		<u>30MG</u>	<u>N19777</u>	<u>006</u>	Jan 20, 1999
<u>AB</u>	+	<u>40MG</u>	<u>N19777</u>	<u>004</u>	May 19, 1988

PRESCRIPTION DRUG PRODUCT LIST

3 - 228 (of 360)

LITHIUM CARBONATE

CAPSULE; ORAL

ESKALITH

<u>AB</u>	GLAXOSMITHKLINE	<u>300MG</u>	<u>N16860</u>	<u>001</u>	
	<u>LITHIUM CARBONATE</u>				
<u>AB</u>	APOTEX	<u>300MG</u>	<u>N76795</u>	<u>001</u>	Nov 22, 2004
<u>AB</u>	ROXANE	<u>150MG</u>	<u>N17812</u>	<u>002</u>	Jan 28, 1987
<u>AB</u>		<u>300MG</u>	<u>N17812</u>	<u>001</u>	
	+	600MG	N17812	003	Jan 28, 1987
<u>AB</u>	WEST WARD	<u>150MG</u>	<u>N76243</u>	<u>002</u>	Feb 24, 2003
<u>AB</u>		<u>300MG</u>	<u>N76243</u>	<u>001</u>	Jun 27, 2002

TABLET; ORAL

LITHIUM CARBONATE

<u>AB</u>	+	PFIZER	<u>300MG</u>	<u>N16834</u>	<u>001</u>	
<u>AB</u>		ROXANE	<u>300MG</u>	<u>N18558</u>	<u>001</u>	Jan 29, 1982

TABLET, EXTENDED RELEASE; ORAL

ESKALITH CR

<u>AB</u>	+	GLAXOSMITHKLINE	<u>450MG</u>	<u>N18152</u>	<u>001</u>	Mar 29, 1982
		<u>LITHIUM CARBONATE</u>				
<u>AB</u>		BARR	<u>300MG</u>	<u>N76170</u>	<u>001</u>	Jun 10, 2002
<u>AB</u>			<u>450MG</u>	<u>N76366</u>	<u>001</u>	Aug 21, 2003
<u>AB</u>		ROXANE	<u>300MG</u>	<u>N76832</u>	<u>001</u>	Oct 28, 2004
<u>AB</u>			<u>450MG</u>	<u>N76691</u>	<u>001</u>	Jan 05, 2004
<u>AB</u>		WEST WARD	<u>450MG</u>	<u>N76490</u>	<u>001</u>	Jun 17, 2003
		<u>LITHOBID</u>				
<u>AB</u>	+	JDS PHARMS	<u>300MG</u>	<u>N18027</u>	<u>001</u>	

LITHIUM CITRATE

SYRUP; ORAL

LITHIUM CITRATE

<u>AA</u>		MORTON GROVE	<u>EQ 300MG CARBONATE/5ML</u>	<u>N70755</u>	<u>001</u>	May 21, 1986
<u>AA</u>	+	ROXANE	<u>EQ 300MG CARBONATE/5ML</u>	<u>N18421</u>	<u>001</u>	

LODOXAMIDE TROMETHAMINE

SOLUTION/DROPS; OPHTHALMIC

ALOMIDE

	+	ALCON	EQ 0.1% BASE	N20191	001	Sep 23, 1993
--	---	-------	--------------	--------	-----	--------------

LOMEFLOXACIN HYDROCHLORIDE

TABLET; ORAL

MAXAQUIN

	+	PHARMACIA	EQ 400MG BASE	N20013	001	Feb 21, 1992
--	---	-----------	---------------	--------	-----	--------------

LOMUSTINE

CAPSULE; ORAL

CEENU

		BRISTOL MYERS SQUIBB	10MG	N17588	001	
			40MG	N17588	002	
	+		100MG	N17588	003	

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

IMODIUM

<u>AB</u>	+	MCNEIL CONS SPECLT	<u>2MG</u>	<u>N17694</u>	<u>001</u>	
		<u>LOPERAMIDE HYDROCHLORIDE</u>				
<u>AB</u>		MYLAN	<u>2MG</u>	<u>N72741</u>	<u>001</u>	Sep 18, 1991
<u>AB</u>		SANDOZ	<u>2MG</u>	<u>N72993</u>	<u>001</u>	Aug 28, 1992
<u>AB</u>		TEVA	<u>2MG</u>	<u>N73122</u>	<u>001</u>	Aug 30, 1991
<u>AB</u>			<u>2MG</u>	<u>N73192</u>	<u>001</u>	Apr 30, 1992

PRESCRIPTION DRUG PRODUCT LIST

3 - 229 (of 360)

LOPINAVIR; RITONAVIR

CAPSULE; ORAL					
KALETRA					
+	ABBOTT	133.3MG;33.3MG	N21226	001	Sep 15, 2000
SOLUTION; ORAL					
KALETRA					
+	ABBOTT	80MG/ML;20MG/ML	N21251	001	Sep 15, 2000
TABLET; ORAL					
KALETRA					
+	ABBOTT	200MG;50MG	N21906	001	Oct 28, 2005

LORACARBEF

CAPSULE; ORAL					
LORABID					
	KING PHARMS	200MG	N50668	001	Dec 31, 1991
+		400MG	N50668	002	Apr 05, 1996
FOR SUSPENSION; ORAL					
LORABID					
	KING PHARMS	100MG/5ML	N50667	001	Dec 31, 1991
+		200MG/5ML	N50667	002	Dec 31, 1991

LORAZEPAM

CONCENTRATE; ORAL					
LORAZEPAM INTENSOL					
+	ROXANE	2MG/ML	N72755	001	Jun 28, 1991
INJECTABLE; INJECTION					
<u>ATIVAN</u>					
<u>AP</u>	+	BAXTER HLTHCARE CORP	<u>2MG/ML</u>	<u>N18140</u>	<u>001</u>
<u>AP</u>	+		<u>4MG/ML</u>	<u>N18140</u>	<u>002</u>
<u>LORAZEPAM</u>					
<u>AP</u>		BEDFORD	<u>2MG/ML</u>	<u>N77076</u>	<u>001</u>
<u>AP</u>			<u>4MG/ML</u>	<u>N77076</u>	<u>002</u>
<u>AP</u>		HOSPIRA	<u>2MG/ML</u>	<u>N74243</u>	<u>001</u>
<u>AP</u>			<u>2MG/ML</u>	<u>N74280</u>	<u>001</u>
<u>AP</u>			<u>2MG/ML</u>	<u>N74282</u>	<u>001</u>
<u>AP</u>			<u>2MG/ML</u>	<u>N74300</u>	<u>001</u>
<u>AP</u>			<u>4MG/ML</u>	<u>N74243</u>	<u>002</u>
<u>AP</u>			<u>4MG/ML</u>	<u>N74280</u>	<u>002</u>
<u>AP</u>			<u>4MG/ML</u>	<u>N74282</u>	<u>002</u>
<u>AP</u>			<u>4MG/ML</u>	<u>N74300</u>	<u>003</u>
<u>AP</u>		INTL MEDICATION SYS	<u>2MG/ML</u>	<u>N76150</u>	<u>001</u>
<u>AP</u>		STERIS	<u>2MG/ML</u>	<u>N74276</u>	<u>001</u>
<u>AP</u>			<u>4MG/ML</u>	<u>N74276</u>	<u>002</u>
<u>LORAZEPAM PRESERVATIVE FREE</u>					
<u>AP</u>		BEDFORD LABS	<u>2MG/ML</u>	<u>N77074</u>	<u>001</u>
<u>AP</u>			<u>4MG/ML</u>	<u>N77074</u>	<u>002</u>
SOLUTION; ORAL					
LORAZEPAM					
	ROXANE	0.5MG/5ML	N74648	001	Mar 18, 1997
TABLET; ORAL					
<u>ATIVAN</u>					
<u>AB</u>		BIOVAIL	<u>0.5MG</u>	<u>N17794</u>	<u>001</u>
<u>AB</u>			<u>1MG</u>	<u>N17794</u>	<u>002</u>
<u>AB</u>	+		<u>2MG</u>	<u>N17794</u>	<u>003</u>
<u>LORAZEPAM</u>					
<u>AB</u>		MUTUAL PHARM	<u>0.5MG</u>	<u>N72553</u>	<u>001</u>
<u>AB</u>			<u>1MG</u>	<u>N72554</u>	<u>001</u>
<u>AB</u>			<u>2MG</u>	<u>N72555</u>	<u>001</u>
<u>AB</u>		MYLAN	<u>0.5MG</u>	<u>N71589</u>	<u>001</u>

PRESCRIPTION DRUG PRODUCT LIST

3 - 230 (of 360)

LORAZEPAM

TABLET; ORAL				
<u>LORAZEPAM</u>				
<u>AB</u>	MYLAN	<u>1MG</u>	<u>N71590</u>	<u>001</u> Oct 13, 1987
<u>AB</u>		<u>2MG</u>	<u>N71591</u>	<u>001</u> Oct 13, 1987
<u>AB</u>	PUREPAC PHARM	<u>0.5MG</u>	<u>N71403</u>	<u>001</u> Apr 21, 1987
<u>AB</u>		<u>1MG</u>	<u>N71404</u>	<u>001</u> Apr 21, 1987
<u>AB</u>		<u>2MG</u>	<u>N71141</u>	<u>001</u> Apr 21, 1987
<u>AB</u>	RANBAXY	<u>0.5MG</u>	<u>N76045</u>	<u>001</u> Aug 29, 2001
<u>AB</u>		<u>1MG</u>	<u>N76045</u>	<u>002</u> Aug 29, 2001
<u>AB</u>		<u>2MG</u>	<u>N76045</u>	<u>003</u> Aug 29, 2001
<u>AB</u>	SANDOZ	<u>0.5MG</u>	<u>N71193</u>	<u>001</u> Apr 15, 1988
<u>AB</u>		<u>1MG</u>	<u>N71194</u>	<u>001</u> Apr 15, 1988
<u>AB</u>		<u>2MG</u>	<u>N71195</u>	<u>001</u> Apr 15, 1988
<u>AB</u>	WATSON LABS	<u>0.5MG</u>	<u>N71117</u>	<u>001</u> Jul 24, 1986
<u>AB</u>		<u>0.5MG</u>	<u>N72926</u>	<u>001</u> Oct 31, 1991
<u>AB</u>		<u>1MG</u>	<u>N71118</u>	<u>001</u> Jul 24, 1986
<u>AB</u>		<u>1MG</u>	<u>N72927</u>	<u>001</u> Oct 31, 1991
<u>AB</u>		<u>2MG</u>	<u>N71110</u>	<u>001</u> Jul 24, 1986
<u>AB</u>		<u>2MG</u>	<u>N72928</u>	<u>001</u> Oct 31, 1991

LOSARTAN POTASSIUM

TABLET; ORAL				
COZAAR				
	MERCK	25MG	N20386	001 Apr 14, 1995
		50MG	N20386	002 Apr 14, 1995
	+	100MG	N20386	003 Oct 13, 1998

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS; OPHTHALMIC				
ALREX				
	+ BAUSCH AND LOMB	0.2%	N20803	001 Mar 09, 1998
LOTEMAX				
	+ BAUSCH AND LOMB	0.5%	N20583	001 Mar 09, 1998

LOTEPREDNOL ETABONATE; TOBRAMYCIN

SUSPENSION/DROPS; OPHTHALMIC				
ZYLET				
	+ BAUSCH AND LOMB	0.5%;0.3%	N50804	001 Dec 14, 2004

LOVASTATIN

TABLET; ORAL				
<u>LOVASTATIN</u>				
<u>AB</u>	CARLSBAD	<u>10MG</u>	<u>N75991</u>	<u>001</u> Jun 05, 2002
<u>AB</u>		<u>20MG</u>	<u>N75991</u>	<u>002</u> Jun 05, 2002
<u>AB</u>		<u>40MG</u>	<u>N75991</u>	<u>003</u> Jun 05, 2002
<u>AB</u>	GENPHARM	<u>10MG</u>	<u>N75935</u>	<u>001</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>N75935</u>	<u>002</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>N75935</u>	<u>003</u> Dec 17, 2001
<u>AB</u>	MYLAN	<u>10MG</u>	<u>N75451</u>	<u>001</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>N75451</u>	<u>002</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>N75451</u>	<u>003</u> Dec 17, 2001
<u>AB</u>	PUREPAC PHARM	<u>10MG</u>	<u>N75828</u>	<u>001</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>N75828</u>	<u>002</u> Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>N75828</u>	<u>003</u> Dec 17, 2001
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>N75300</u>	<u>001</u> Dec 17, 2001
<u>AB</u>		<u>10MG</u>	<u>N75636</u>	<u>001</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>N75300</u>	<u>002</u> Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>N75636</u>	<u>002</u> Dec 17, 2001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 231 (of 360)

LOVASTATIN

TABLET; ORAL

LOVASTATIN

<u>AB</u>	SANDOZ	<u>40MG</u>	<u>N75300</u>	<u>003</u>	Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>N75636</u>	<u>003</u>	Dec 17, 2001
<u>AB</u>	TEVA	<u>10MG</u>	<u>N75551</u>	<u>003</u>	Dec 17, 2001
<u>AB</u>		<u>20MG</u>	<u>N75551</u>	<u>002</u>	Dec 17, 2001
<u>AB</u>		<u>40MG</u>	<u>N75551</u>	<u>001</u>	Dec 17, 2001
	<u>MEVACOR</u>				
<u>AB</u>	MERCK	<u>10MG</u>	<u>N19643</u>	<u>002</u>	Mar 28, 1991
<u>AB</u>		<u>20MG</u>	<u>N19643</u>	<u>003</u>	Aug 31, 1987
<u>AB</u>	+	<u>40MG</u>	<u>N19643</u>	<u>004</u>	Dec 14, 1988

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV

	ANDRX LABS LLC	10MG	N21316	001	Jun 26, 2002
		20MG	N21316	002	Jun 26, 2002
		40MG	N21316	003	Jun 26, 2002
	+	60MG	N21316	004	Jun 26, 2002

LOVASTATIN; NIACIN

TABLET, EXTENDED RELEASE; ORAL

ADVICOR

+	KOS LIFE	20MG;500MG	N21249	001	Dec 17, 2001
+		20MG;1GM	N21249	003	Dec 17, 2001

LOXAPINE SUCCINATE

CAPSULE; ORAL

LOXAPINE

<u>AB</u>	MYLAN	<u>EQ 5MG BASE</u>	<u>N76762</u>	<u>001</u>	Nov 01, 2004
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N76762</u>	<u>002</u>	Nov 01, 2004
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N76762</u>	<u>003</u>	Nov 01, 2004
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N76762</u>	<u>004</u>	Nov 01, 2004

LOXAPINE SUCCINATE

<u>AB</u>	AMIDE PHARM	<u>EQ 5MG BASE</u>	<u>N76868</u>	<u>001</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N76868</u>	<u>002</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N76868</u>	<u>003</u>	Aug 04, 2005
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N76868</u>	<u>004</u>	Aug 04, 2005
<u>AB</u>	WATSON LABS	<u>EQ 5MG BASE</u>	<u>N72204</u>	<u>001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N72205</u>	<u>001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N72206</u>	<u>001</u>	Jun 15, 1988
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N72062</u>	<u>001</u>	Jun 15, 1988

LOXITANE

<u>AB</u>	WATSON LABS	<u>EQ 5MG BASE</u>	<u>N17525</u>	<u>001</u>	
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N17525</u>	<u>002</u>	
<u>AB</u>	+	<u>EQ 25MG BASE</u>	<u>N17525</u>	<u>003</u>	
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N17525</u>	<u>004</u>	

LUTROPIN ALFA

INJECTABLE; SUBCUTANEOUS

LUVERIS

+	SERONO INC	75 IU/VIAL	N21322	001	Oct 08, 2004
---	------------	------------	--------	-----	--------------

MAFENIDE ACETATE

CREAM; TOPICAL

SULFAMYLON

+	MYLAN BERTEK	EQ 85MG BASE/GM	N16763	001	
---	--------------	-----------------	--------	-----	--

FOR SOLUTION; TOPICAL

SULFAMYLON

+	MYLAN BERTEK	5%	N19832	003	Jun 05, 1998
---	--------------	----	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 232 (of 360)

MAGNESIUM ACETATE TETRAHYDRATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

PLASMA-LYTE 56 IN PLASTIC CONTAINER

BAXTER HLTHCARE 32MG/100ML;128MG/100ML;234MG/100ML N19047 001 Jun 15, 1984

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

B BRAUN 30MG/100ML;37MG/100ML;0.82MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML;12MG/100ML N19696 001 Sep 29, 1989

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

ISOLYTE S IN PLASTIC CONTAINERAP B BRAUN 30MG/100ML;37MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML N19711 001 Sep 29, 1989

NORMOSOL-R IN PLASTIC CONTAINER

HOSPIRA 30MG/100ML;37MG/100ML;222MG/100ML;526MG/100ML;502MG/100ML N17586 001

PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINERAP BAXTER HLTHCARE 30MG/100ML;37MG/100ML;368MG/100ML;526MG/100ML;502MG/100ML N17378 001PLASMA-LYTE A IN PLASTIC CONTAINERAP BAXTER HLTHCARE 30MG/100ML;37MG/100ML;368MG/100ML;526MG/100ML;502MG/100ML N17378 002 Nov 22, 1982

SOLUTION; IRRIGATION

PHYSIOLYTE IN PLASTIC CONTAINER

B BRAUN 30MG/100ML;37MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML N19024 001 Jun 08, 1984

PHYSIOSOL IN PLASTIC CONTAINER

HOSPIRA 30MG/100ML;37MG/100ML;222MG/100ML;526MG/100ML;502MG/100ML N17637 002 Jul 08, 1982

PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER

HOSPIRA 30MG/100ML;37MG/100ML;222MG/100ML;526MG/100ML;502MG/100ML N18406 002 Jul 08, 1982

MAGNESIUM SULFATE

INJECTABLE; INJECTION

MAGNESIUM SULFATEAP + AM PHARM PARTNERS 500MG/ML N19316 001 Sep 08, 1986AP HOSPIRA 500MG/ML N75151 001 Apr 25, 2000

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER

+ HOSPIRA 1GM/100ML N20488 001 Jul 11, 1995

+ 2GM/100ML N20488 002 Jul 11, 1995

MAGNESIUM SULFATE IN PLASTIC CONTAINER

+ HOSPIRA 80MG/ML N20309 002 Jun 24, 1994

+ 4GM/100ML N20309 001 Jun 24, 1994

MAGNESIUM SULFATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

TIS-U-SOLAT BAXTER HLTHCARE 20MG/100ML;40MG/100ML;6.25MG/100ML;800MG/100ML;8.75MG/100ML N18508 001 Feb 19, 1982TIS-U-SOL IN PLASTIC CONTAINERAT BAXTER HLTHCARE 20MG/100ML;40MG/100ML;6.25MG/100ML;800MG/100ML;8.75MG/100ML N18336 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 233 (of 360)

MALATHIONLOTION; TOPICAL
OVIDE

+ TARO PHARMS NORTH 0.5% N18613 001 Aug 02, 1982

MANGANESE CHLORIDEINJECTABLE; INJECTION
MANGANESE CHLORIDE IN PLASTIC CONTAINER
HOSPIRA EQ 0.1MG MANGANESE/ML

N18962 001 Jun 26, 1986

MANNITOL

INJECTABLE; INJECTION

MANNITOL 10%AP B BRAUN 10GM/100MLN16080 002MANNITOL 10% IN PLASTIC CONTAINERAP B BRAUN 10GM/100MLN20006 002 Jul 26, 1993AP HOSPIRA 10GM/100MLN19603 002 Jan 08, 1987MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATERAP B BRAUN 10GM/100MLN16080 006MANNITOL 15%AP B BRAUN 15GM/100MLN16080 003MANNITOL 15% IN PLASTIC CONTAINERAP B BRAUN 15GM/100MLN20006 003 Jul 26, 1993AP HOSPIRA 15GM/100MLN19603 003 Jan 08, 1990MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%AP B BRAUN 15GM/100MLN16080 005MANNITOL 20%AP B BRAUN 20GM/100MLN14738 001AP 20GM/100MLN16080 004MANNITOL 20% IN PLASTIC CONTAINERAP B BRAUN 20GM/100MLN20006 004 Jul 26, 1993AP HOSPIRA 20GM/100MLN19603 004 Jan 08, 1990MANNITOL 25%AP AM PHARM PARTNERS 12.5GM/50MLN80677 001AP HOSPIRA 12.5GM/50MLN16269 006 Aug 25, 1994AP INTL MEDICATION 12.5GM/50MLN83051 001AP LUITPOLD 12.5GM/50MLN87409 001 Jan 21, 1982MANNITOL 5%AP B BRAUN 5GM/100MLN16080 001MANNITOL 5% IN PLASTIC CONTAINERAP B BRAUN 5GM/100MLN20006 001 Jul 26, 1993AP HOSPIRA 5GM/100MLN19603 001 Jan 08, 1987MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%AP B BRAUN 5GM/100MLN16080 007OSMITROL 10% IN WATERAP BAXTER HLTHCARE 10GM/100MLN13684 002OSMITROL 10% IN WATER IN PLASTIC CONTAINERAP BAXTER HLTHCARE 10GM/100MLN13684 006OSMITROL 15% IN WATERAP BAXTER HLTHCARE 15GM/100MLN13684 004OSMITROL 15% IN WATER IN PLASTIC CONTAINERAP BAXTER HLTHCARE 15GM/100MLN13684 008OSMITROL 20% IN WATERAP BAXTER HLTHCARE 20GM/100MLN13684 003OSMITROL 20% IN WATER IN PLASTIC CONTAINERAP BAXTER HLTHCARE 20GM/100MLN13684 007OSMITROL 5% IN WATERAP BAXTER HLTHCARE 5GM/100MLN13684 001OSMITROL 5% IN WATER IN PLASTIC CONTAINERAP BAXTER HLTHCARE 5GM/100MLN13684 005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 234 (of 360)

MANNITOL

SOLUTION; IRRIGATION
 RESECTISOL IN PLASTIC CONTAINER
 B BRAUN 5GM/100ML

N16772 002

MANNITOL; SORBITOL

SOLUTION; IRRIGATION
SORBITOL-MANNITOL IN PLASTIC CONTAINER

AT HOSPIRA 540MG/100ML; 2.7GM/100ML
AT 540MG/100ML; 2.7GM/100ML

N17636 001N18316 001MAPROTILINE HYDROCHLORIDE

TABLET; ORAL
MAPROTILINE HYDROCHLORIDE

AB MYLAN 25MG
AB + 50MG
AB 75MG
AB WATSON LABS 25MG
AB 50MG
AB 75MG

N72284 001N72285 001N72286 001N72162 001N72163 001N72164 001

Oct 03, 1988

Oct 03, 1988

Oct 03, 1988

Jun 01, 1988

Jun 01, 1988

Jun 01, 1988

MEBENDAZOLE

TABLET, CHEWABLE; ORAL
 MEBENDAZOLE
 + TEVA PHARMS 100MG

N73580 001

Jan 04, 1995

MECAMYLAMINE HYDROCHLORIDE

TABLET; ORAL
 INVERSINE
 + TARGACEPT 2.5MG

N10251 001

MECASERMIN RECOMBINANT

INJECTABLE; SUBCUTANEOUS
 INCRELEX
 + TERCICA 40MG/4ML (10MG/ML)

N21839 001

Aug 30, 2005

MECASERMIN RINFABATE RECOMBINANT

INJECTABLE; SUBCUTANEOUS
 IPLEX
 + INSMED 36MG/0.6ML

N21884 001

Dec 12, 2005

MECHLORETHAMINE HYDROCHLORIDE

INJECTABLE; INJECTION
 MUSTARGEN
 + OVATION PHARMS 10MG/VIAL

N06695 001

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

ANTIVERT

AA + PFIZER 12.5MG
AA + 25MG
AA + 50MG

N10721 006N10721 004N10721 001

Jan 20, 1982

MECLIZINE HYDROCHLORIDE

AA BUNDY 12.5MG
AA 25MG
AA IVAX PHARMS 12.5MG
AA 25MG
AA PAR PHARM 12.5MG
AA 25MG

N84382 001N84872 001N84975 001N84657 001N87127 001N87128 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 235 (of 360)

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

MECLIZINE HYDROCHLORIDE

<u>AA</u>	PAR PHARM	<u>50MG</u>	<u>N89674</u>	<u>001</u>	Mar 31, 1988
<u>AA</u>	PLIVA	<u>12.5MG</u>	<u>N88732</u>	<u>001</u>	Dec 11, 1985
<u>AA</u>		<u>25MG</u>	<u>N88734</u>	<u>001</u>	Dec 11, 1985
<u>AA</u>	SANDOZ	<u>12.5MG</u>	<u>N84843</u>	<u>002</u>	May 22, 1989
<u>AA</u>		<u>25MG</u>	<u>N84092</u>	<u>003</u>	May 22, 1989
<u>AA</u>	VINTAGE PHARMS	<u>12.5MG</u>	<u>N40179</u>	<u>001</u>	Jan 30, 1997
<u>AA</u>		<u>25MG</u>	<u>N40179</u>	<u>002</u>	Jan 30, 1997
<u>AA</u>	WATSON LABS	<u>25MG</u>	<u>N85740</u>	<u>001</u>	

TABLET, CHEWABLE; ORAL

ANTIVERT

<u>AA</u>	+ PFIZER	<u>25MG</u>	<u>N10721</u>	<u>005</u>	
<u>MECLIZINE HYDROCHLORIDE</u>					
<u>AA</u>	PLIVA	<u>25MG</u>	<u>N88733</u>	<u>001</u>	Dec 11, 1985

MECLOFENAMATE SODIUM

CAPSULE; ORAL

MECLOFENAMATE SODIUM

<u>AB</u>	MYLAN	<u>EQ 50MG BASE</u>	<u>N71080</u>	<u>001</u>	Sep 03, 1986
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N71081</u>	<u>001</u>	Sep 03, 1986
<u>AB</u>	SANDOZ	<u>EQ 50MG BASE</u>	<u>N72262</u>	<u>001</u>	Nov 29, 1988
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N72263</u>	<u>001</u>	Nov 29, 1988
<u>AB</u>	WATSON LABS	<u>EQ 50MG BASE</u>	<u>N71468</u>	<u>001</u>	Apr 15, 1987
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N71469</u>	<u>001</u>	Apr 15, 1987

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

<u>AB</u>	+ PHARMACIA AND UPJOHN	<u>150MG/ML</u>	<u>N20246</u>	<u>001</u>	Oct 29, 1992
	+	400MG/ML	N12541	003	

MEDROXYPROGESTERONE ACETATE

<u>AB</u>	SICOR PHARMS	<u>150MG/ML</u>	<u>N76552</u>	<u>001</u>	Oct 27, 2004
<u>AB</u>		<u>150MG/ML</u>	<u>N76553</u>	<u>001</u>	Jul 28, 2004

INJECTABLE; SUBCUTANEOUS

DEPO-SUBQ PROVERA 104

	+ PFIZER	104MG/0.65ML	N21583	001	Dec 17, 2004
--	----------	--------------	--------	-----	--------------

TABLET; ORAL

MEDROXYPROGESTERONE ACETATE

<u>AB</u>	BARR	<u>2.5MG</u>	<u>N40159</u>	<u>001</u>	Aug 09, 1996
<u>AB</u>		<u>5MG</u>	<u>N40159</u>	<u>002</u>	Aug 09, 1996
<u>AB</u>		<u>10MG</u>	<u>N40159</u>	<u>003</u>	Aug 09, 1996
<u>AB</u>	DURAMED PHARMS BARR	<u>2.5MG</u>	<u>N40311</u>	<u>001</u>	Dec 01, 1999
<u>AB</u>		<u>5MG</u>	<u>N40311</u>	<u>002</u>	Dec 01, 1999
<u>AB</u>		<u>10MG</u>	<u>N40311</u>	<u>003</u>	Dec 01, 1999
BP	USL PHARMA	10MG	N88484	001	Jul 26, 1984

PROVERA

<u>AB</u>	PHARMACIA AND UPJOHN	<u>2.5MG</u>	<u>N11839</u>	<u>001</u>	
<u>AB</u>		<u>5MG</u>	<u>N11839</u>	<u>003</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N11839</u>	<u>004</u>	

MEDRYSONE

SUSPENSION; OPHTHALMIC

HMS

	+ ALLERGAN	1%	N16624	003	
--	------------	----	--------	-----	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 236 (of 360)

MEFENAMIC ACID

CAPSULE; ORAL

PONSTEL

+ FIRST HORIZON 250MG

N15034 003

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

LARIAM

AB + ROCHE 250MG

N19591 001 May 02, 1989

MEFLOQUINE

AB BARR 250MG

N76392 001 Dec 29, 2003

MEFLOQUINE HYDROCHLORIDE

AB ROXANE 250MG

N76523 001 Oct 01, 2004

AB SANDOZ 250MG

N76175 001 Feb 20, 2002

MEGESTROL ACETATE

SUSPENSION; ORAL

MEGACE

AB + BRISTOL MYERS SQUIBB 40MG/ML

N20264 001 Sep 10, 1993

MEGACE ES

+ PAR PHARM 125MG/ML

N21778 001 Jul 05, 2005

MEGESTROL ACETATE

AB MORTON GROVE 40MG/ML

N76721 001 Nov 01, 2004

AB PAR PHARM 40MG/ML

N75671 001 Jul 25, 2001

AB ROXANE 40MG/ML

N75997 001 Feb 15, 2002

AB TEVA PHARMS 40MG/ML

N75681 001 May 05, 2003

TABLET; ORAL

MEGACE

AB BRISTOL MYERS SQUIBB 20MG

N16979 001

AB + 40MG

N16979 002

MEGESTROL ACETATE

AB BARR 20MG

N74621 002 Aug 16, 1996

AB 40MG

N74621 001 Nov 30, 1995

AB PAR PHARM 20MG

N72422 001 Aug 08, 1988

AB 40MG

N72423 001 Aug 08, 1988

AB ROXANE 20MG

N74458 001 Sep 29, 1995

AB 40MG

N74458 002 Sep 29, 1995

AB TEVA 40MG

N74745 001 Feb 27, 1998

MELOXICAM

SUSPENSION; ORAL

MOBIC

+ BOEHRINGER INGELHEIM 7.5MG/5ML

N21530 001 Jun 01, 2004

TABLET; ORAL

MOBIC

BOEHRINGER INGELHEIM 7.5MG

N20938 001 Apr 13, 2000

+ 15MG

N20938 002 Aug 23, 2000

MELPHALAN

TABLET; ORAL

ALKERAN

+ GLAXOSMITHKLINE 2MG

N14691 002

MELPHALAN HYDROCHLORIDE

INJECTABLE; INJECTION

ALKERAN

+ GLAXOSMITHKLINE EQ 50MG BASE/VIAL

N20207 001 Nov 18, 1992

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 237 (of 360)

MEMANTINE HYDROCHLORIDE

SOLUTION; ORAL

NAMENDA

+	FOREST LABS	2MG/ML	N21627	001	Apr 18, 2005
---	-------------	--------	--------	-----	--------------

TABLET; ORAL

NAMENDA

	FOREST LABS	5MG	N21487	001	Oct 16, 2003
--	-------------	-----	--------	-----	--------------

+		10MG	N21487	002	Oct 16, 2003
---	--	------	--------	-----	--------------

MENOTROPINS (FSH; LH)

INJECTABLE; IM-SC

REPRONEX

+	FERRING	75 IU/VIAL;75 IU/VIAL	N21047	001	Aug 27, 1999
---	---------	-----------------------	--------	-----	--------------

INJECTABLE; SUBCUTANEOUS

MENOPUR

+	FERRING	75 IU/VIAL;75 IU/VIAL	N21663	001	Oct 29, 2004
---	---------	-----------------------	--------	-----	--------------

MEPENZOLATE BROMIDE

TABLET; ORAL

CANTIL

+	AVENTIS PHARMS	25MG	N10679	003	
---	----------------	------	--------	-----	--

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

<u>AP</u>	+	HOSPIRA	<u>25MG/ML</u>	<u>N21171</u>	<u>001</u>	
-----------	---	---------	----------------	---------------	------------	--

<u>AP</u>	+		<u>50MG/ML</u>	<u>N21171</u>	<u>002</u>	
-----------	---	--	----------------	---------------	------------	--

<u>AP</u>	+		<u>75MG/ML</u>	<u>N21171</u>	<u>003</u>	
-----------	---	--	----------------	---------------	------------	--

<u>AP</u>	+		<u>100MG/ML</u>	<u>N21171</u>	<u>004</u>	
-----------	---	--	-----------------	---------------	------------	--

MEPERIDINE HYDROCHLORIDE

<u>AP</u>		BAXTER HLTHCARE CORP	<u>25MG/ML</u>	<u>N80455</u>	<u>007</u>	
-----------	--	----------------------	----------------	---------------	------------	--

<u>AP</u>			<u>50MG/ML</u>	<u>N80455</u>	<u>008</u>	
-----------	--	--	----------------	---------------	------------	--

<u>AP</u>			<u>75MG/ML</u>	<u>N80455</u>	<u>009</u>	
-----------	--	--	----------------	---------------	------------	--

<u>AP</u>			<u>100MG/ML</u>	<u>N80455</u>	<u>010</u>	
-----------	--	--	-----------------	---------------	------------	--

<u>AP</u>		ELKINS SINN	<u>25MG/ML</u>	<u>N80445</u>	<u>001</u>	
-----------	--	-------------	----------------	---------------	------------	--

<u>AP</u>			<u>50MG/ML</u>	<u>N80445</u>	<u>002</u>	
-----------	--	--	----------------	---------------	------------	--

<u>AP</u>			<u>75MG/ML</u>	<u>N80445</u>	<u>003</u>	
-----------	--	--	----------------	---------------	------------	--

<u>AP</u>			<u>100MG/ML</u>	<u>N80445</u>	<u>004</u>	
-----------	--	--	-----------------	---------------	------------	--

<u>AP</u>		STERIS	<u>50MG/ML</u>	<u>N73444</u>	<u>001</u>	Mar 17, 1992
-----------	--	--------	----------------	---------------	------------	--------------

<u>AP</u>			<u>100MG/ML</u>	<u>N73445</u>	<u>001</u>	Mar 17, 1992
-----------	--	--	-----------------	---------------	------------	--------------

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

<u>AP</u>		ASTRAZENECA	<u>10MG/ML</u>	<u>N81002</u>	<u>001</u>	Jul 30, 1993
-----------	--	-------------	----------------	---------------	------------	--------------

<u>AP</u>	+	HOSPIRA	<u>10MG/ML</u>	<u>N88432</u>	<u>001</u>	Aug 16, 1984
-----------	---	---------	----------------	---------------	------------	--------------

<u>AP</u>		INTL MEDICATION	<u>10MG/ML</u>	<u>N81309</u>	<u>001</u>	Aug 30, 1993
-----------	--	-----------------	----------------	---------------	------------	--------------

<u>AP</u>		MALLINCKRODT	<u>10MG/ML</u>	<u>N40163</u>	<u>001</u>	May 12, 1997
-----------	--	--------------	----------------	---------------	------------	--------------

<u>AP</u>		STERIS	<u>10MG/ML</u>	<u>N73443</u>	<u>001</u>	Mar 17, 1992
-----------	--	--------	----------------	---------------	------------	--------------

SYRUP; ORAL

DEMEROL

<u>AA</u>	+	SANOFI SYNTHELABO	<u>50MG/5ML</u>	<u>N05010</u>	<u>005</u>	
-----------	---	-------------------	-----------------	---------------	------------	--

MEPERIDINE HYDROCHLORIDE

<u>AA</u>		ROXANE	<u>50MG/5ML</u>	<u>N88744</u>	<u>001</u>	Jan 30, 1985
-----------	--	--------	-----------------	---------------	------------	--------------

TABLET; ORAL

DEMEROL

<u>AA</u>	+	SANOFI SYNTHELABO	<u>50MG</u>	<u>N05010</u>	<u>001</u>	
-----------	---	-------------------	-------------	---------------	------------	--

<u>AA</u>	+		<u>100MG</u>	<u>N05010</u>	<u>004</u>	
-----------	---	--	--------------	---------------	------------	--

MEPERIDINE HYDROCHLORIDE

<u>AA</u>		AMIDE PHARM	<u>50MG</u>	<u>N40331</u>	<u>001</u>	May 28, 1999
-----------	--	-------------	-------------	---------------	------------	--------------

<u>AA</u>			<u>100MG</u>	<u>N40331</u>	<u>002</u>	May 28, 1999
-----------	--	--	--------------	---------------	------------	--------------

<u>AA</u>		BARR	<u>50MG</u>	<u>N88639</u>	<u>001</u>	Jul 02, 1984
-----------	--	------	-------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 238 (of 360)

MEPERIDINE HYDROCHLORIDE

TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

<u>AA</u>	BARR	<u>100MG</u>	<u>N88640</u>	<u>001</u>	Sep 19, 1984
<u>AA</u>	CARACO	<u>50MG</u>	<u>N40446</u>	<u>001</u>	Aug 08, 2002
<u>AA</u>		<u>100MG</u>	<u>N40446</u>	<u>002</u>	Aug 08, 2002
<u>AA</u>	DURAMED PHARMS BARR	<u>50MG</u>	<u>N40318</u>	<u>001</u>	Oct 05, 1999
<u>AA</u>		<u>100MG</u>	<u>N40318</u>	<u>002</u>	Oct 05, 1999
<u>AA</u>	MALLINCKRODT	<u>50MG</u>	<u>N40352</u>	<u>001</u>	Jun 13, 2000
<u>AA</u>		<u>100MG</u>	<u>N40352</u>	<u>002</u>	Jun 13, 2000
<u>AA</u>	MUTUAL PHARM	<u>50MG</u>	<u>N80448</u>	<u>001</u>	
<u>AA</u>		<u>100MG</u>	<u>N80448</u>	<u>002</u>	
<u>AA</u>	ROXANE	<u>50MG</u>	<u>N40110</u>	<u>001</u>	Mar 12, 1997
<u>AA</u>		<u>100MG</u>	<u>N40110</u>	<u>002</u>	Mar 12, 1997
<u>AA</u>	VINTAGE PHARMS	<u>50MG</u>	<u>N40191</u>	<u>001</u>	Dec 17, 1998
<u>AA</u>		<u>100MG</u>	<u>N40191</u>	<u>002</u>	Dec 17, 1998
<u>AA</u>	WATSON LABS	<u>50MG</u>	<u>N40186</u>	<u>001</u>	Jun 30, 1997
<u>AA</u>		<u>100MG</u>	<u>N40186</u>	<u>002</u>	Jun 30, 1997

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CARBOCAINE

<u>AP</u>	+	EASTMAN KODAK	<u>3%</u>	<u>N12125</u>	<u>003</u>	
<u>AP</u>	+	HOSPIRA	<u>1%</u>	<u>N12250</u>	<u>001</u>	
<u>AP</u>	+		<u>1.5%</u>	<u>N12250</u>	<u>005</u>	
<u>AP</u>	+		<u>2%</u>	<u>N12250</u>	<u>002</u>	

ISOCAINE HYDROCHLORIDE

<u>AP</u>		NOVOCOL	<u>3%</u>	<u>N80925</u>	<u>001</u>	
-----------	--	---------	-----------	---------------	------------	--

MEPIVACAINE HYDROCHLORIDE

<u>AP</u>		GRAHAM CHEM	<u>3%</u>	<u>N83559</u>	<u>001</u>	
<u>AP</u>		STERIS	<u>1%</u>	<u>N88769</u>	<u>001</u>	Nov 20, 1984
<u>AP</u>			<u>2%</u>	<u>N88770</u>	<u>001</u>	Nov 20, 1984

POLOCAINE

<u>AP</u>		ASTRAZENECA	<u>1%</u>	<u>N89407</u>	<u>001</u>	Dec 01, 1986
<u>AP</u>			<u>2%</u>	<u>N89410</u>	<u>001</u>	Dec 01, 1986
<u>AP</u>		DENTSPLY PHARM	<u>3%</u>	<u>N88653</u>	<u>001</u>	Aug 21, 1984

POLOCAINE-MPF

<u>AP</u>		ASTRAZENECA	<u>1%</u>	<u>N89406</u>	<u>001</u>	Dec 01, 1986
<u>AP</u>			<u>1.5%</u>	<u>N89408</u>	<u>001</u>	Dec 01, 1986
<u>AP</u>			<u>2%</u>	<u>N89409</u>	<u>001</u>	Dec 01, 1986

SCANDONEST PLAIN

<u>AP</u>		DEPROCO	<u>3%</u>	<u>N88387</u>	<u>001</u>	Oct 10, 1984
-----------	--	---------	-----------	---------------	------------	--------------

MEPROBAMATE

TABLET; ORAL

MEPROBAMATE

	+	ROXANE	600MG	N84332	001	
<u>AA</u>		SANDOZ	<u>200MG</u>	<u>N14547</u>	<u>002</u>	
<u>AA</u>			<u>400MG</u>	<u>N14547</u>	<u>001</u>	
<u>AA</u>			<u>400MG</u>	<u>N80655</u>	<u>001</u>	
<u>AA</u>		SCHERER LABS	<u>400MG</u>	<u>N83343</u>	<u>001</u>	
<u>AA</u>		TABLICAPS	<u>400MG</u>	<u>N83494</u>	<u>001</u>	
<u>AA</u>		WATSON LABS	<u>200MG</u>	<u>N83304</u>	<u>001</u>	
<u>AA</u>			<u>200MG</u>	<u>N85720</u>	<u>001</u>	
<u>AA</u>			<u>400MG</u>	<u>N83308</u>	<u>001</u>	
<u>AA</u>			<u>400MG</u>	<u>N85721</u>	<u>001</u>	
<u>MILTOWN</u>						
<u>AA</u>	+	MEDPOINTE PHARM HLC	<u>200MG</u>	<u>N09698</u>	<u>004</u>	
<u>AA</u>	+		<u>400MG</u>	<u>N09698</u>	<u>002</u>	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 239 (of 360)

MEPROBAMATE

TABLET; ORAL

TRANMEP

<u>AA</u>	SOLVAY	<u>400MG</u>	<u>N16249</u>	<u>001</u>	
-----------	--------	--------------	---------------	------------	--

MEQUINOL; TRETINOIN

SOLUTION; TOPICAL

SOLAGE

+	BARRIER	2%;0.01%	N20922	001	Dec 10, 1999
---	---------	----------	--------	-----	--------------

MERCAPTOPURINE

TABLET; ORAL

MERCAPTOPURINE

<u>AB</u>	PROMETHEUS LABS	<u>50MG</u>	<u>N40461</u>	<u>001</u>	Feb 11, 2004
-----------	-----------------	-------------	---------------	------------	--------------

<u>AB</u>	ROXANE	<u>50MG</u>	<u>N40528</u>	<u>001</u>	Feb 13, 2004
-----------	--------	-------------	---------------	------------	--------------

PURINETHOL

<u>AB</u>	+ TEVA	<u>50MG</u>	<u>N09053</u>	<u>002</u>	
-----------	--------	-------------	---------------	------------	--

MERCAPTOPURINE ANHYDROUS

TABLET; ORAL

MERCAPTOPURINE

<u>AB</u>	MYLAN	<u>50MG</u>	<u>N40594</u>	<u>001</u>	Jul 01, 2005
-----------	-------	-------------	---------------	------------	--------------

MEROPENEM

INJECTABLE; INJECTION

MERREM I.V.

+	ASTRAZENECA	500MG/VIAL	N50706	003	Jun 21, 1996
---	-------------	------------	--------	-----	--------------

+		1GM/VIAL	N50706	001	Jun 21, 1996
---	--	----------	--------	-----	--------------

MESALAMINE

CAPSULE, EXTENDED RELEASE; ORAL

PENTASA

SHIRE

250MG

N20049	001	May 10, 1993
--------	-----	--------------

+		500MG	N20049	002	Jul 08, 2004
---	--	-------	--------	-----	--------------

ENEMA; RECTAL

MESALAMINE

<u>AB</u>	CLAY PARK	<u>4GM/60ML</u>	<u>N76751</u>	<u>001</u>	Sep 17, 2004
-----------	-----------	-----------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>4GM/60ML</u>	<u>N76841</u>	<u>001</u>	Sep 30, 2004
-----------	------	-----------------	---------------	------------	--------------

ROWASA

<u>AB</u>	+ SOLVAY	<u>4GM/60ML</u>	<u>N19618</u>	<u>001</u>	Dec 24, 1987
-----------	----------	-----------------	---------------	------------	--------------

SUPPOSITORY; RECTAL

CANASA

AXCAN SCANDIPHARM

500MG

N21252	001	Jan 05, 2001
--------	-----	--------------

+		1GM	N21252	002	Nov 05, 2004
---	--	-----	--------	-----	--------------

TABLET, DELAYED RELEASE; ORAL

ASACOL

+	PROCTER AND GAMBLE	400MG	N19651	001	Jan 31, 1992
---	--------------------	-------	--------	-----	--------------

MESNA

INJECTABLE; INTRAVENOUS

MESNA

<u>AP</u>	AM PHARM PARTNERS	<u>100MG/ML</u>	<u>N75811</u>	<u>001</u>	Apr 26, 2001
-----------	-------------------	-----------------	---------------	------------	--------------

<u>AP</u>	BEDFORD	<u>100MG/ML</u>	<u>N75739</u>	<u>001</u>	Jan 09, 2004
-----------	---------	-----------------	---------------	------------	--------------

<u>AP</u>	SICOR PHARMS	<u>100MG/ML</u>	<u>N75764</u>	<u>001</u>	Apr 27, 2001
-----------	--------------	-----------------	---------------	------------	--------------

MESNEX

<u>AP</u>	+ BAXTER HLTHCARE	<u>100MG/ML</u>	<u>N19884</u>	<u>001</u>	Dec 30, 1988
-----------	-------------------	-----------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 240 (of 360)

MESNA

TABLET; ORAL

MESNEX

+ BAXTER HLTHCARE	400MG	N20855	001	Mar 21, 2002
-------------------	-------	--------	-----	--------------

MESTRANOL; NORETHINDRONE

TABLET; ORAL-21

NORETHIN 1/50M-21

AB WATSON LABS	0.05MG;1MG	N71539	001	Apr 12, 1988
----------------	------------	--------	-----	--------------

NORETHINDRONE AND MESTRANOL

AB WATSON LABS	0.05MG;1MG	N70758	001	Jul 01, 1988
----------------	------------	--------	-----	--------------

NORINYL 1+50 21-DAY

AB + WATSON LABS	0.05MG;1MG	N13625	002	
------------------	------------	--------	-----	--

TABLET; ORAL-28

NORETHIN 1/50M-28

AB WATSON LABS	0.05MG;1MG	N71540	001	Apr 12, 1988
----------------	------------	--------	-----	--------------

NORETHINDRONE AND MESTRANOL

AB WATSON LABS	0.05MG;1MG	N70759	001	Jul 01, 1988
----------------	------------	--------	-----	--------------

NORINYL 1+50 28-DAY

AB WATSON LABS	0.05MG;1MG	N16659	001	
----------------	------------	--------	-----	--

ORTHO-NOVUM 1/50 28

AB ORTHO MCNEIL PHARM	0.05MG;1MG	N16709	001	
-----------------------	------------	--------	-----	--

METAPROTERENOL SULFATE

AEROSOL, METERED; INHALATION

ALUPENT

+ BOEHRINGER INGELHEIM	0.65MG/INH	N16402	001	
------------------------	------------	--------	-----	--

SOLUTION; INHALATION

ALUPENT

AN + BOEHRINGER INGELHEIM	0.4%	N18761	002	Oct 10, 1986
---------------------------	------	--------	-----	--------------

AN +	0.6%	N18761	001	Jun 30, 1983
------	------	--------	-----	--------------

METAPROTERENOL SULFATE

AN DEY	0.4%	N71786	001	Aug 05, 1988
--------	------	--------	-----	--------------

AN	0.6%	N70804	001	Aug 17, 1987
----	------	--------	-----	--------------

AN MORTON GROVE	0.4%	N75586	001	May 30, 2002
-----------------	------	--------	-----	--------------

AN	0.6%	N75586	002	May 30, 2002
----	------	--------	-----	--------------

AN NEPHRON	0.4%	N71855	001	Jul 14, 1988
------------	------	--------	-----	--------------

AN	0.6%	N71726	001	Jul 14, 1988
----	------	--------	-----	--------------

AN NOVEX	0.4%	N75402	001	Feb 28, 2001
----------	------	--------	-----	--------------

AN	0.6%	N75403	001	Feb 28, 2001
----	------	--------	-----	--------------

SYRUP; ORAL

METAPROTERENOL SULFATE

AA + MORTON GROVE	10MG/5ML	N74702	001	Mar 24, 1997
-------------------	----------	--------	-----	--------------

AA NOVEX	10MG/5ML	N75235	001	Jan 27, 2000
----------	----------	--------	-----	--------------

AA SILARX	10MG/5ML	N73632	001	Jul 22, 1992
-----------	----------	--------	-----	--------------

TABLET; ORAL

METAPROTERENOL SULFATE

AB PAR PHARM	10MG	N72024	001	Jun 28, 1988
--------------	------	--------	-----	--------------

AB +	20MG	N72025	001	Jun 28, 1988
------	------	--------	-----	--------------

AB TEVA	10MG	N72519	001	Mar 30, 1990
---------	------	--------	-----	--------------

AB	20MG	N72520	001	Mar 30, 1990
----	------	--------	-----	--------------

AB WATSON LABS	10MG	N73013	001	Jan 31, 1991
----------------	------	--------	-----	--------------

AB	20MG	N72795	001	Jan 31, 1991
----	------	--------	-----	--------------

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

ARAMINE

AP + MERCK	EQ 10MG BASE/ML	N09509	002	Dec 22, 1987
------------	-----------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 241 (of 360)

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

METARAMINOL BITARTRATE

<u>AP</u>	AM PHARM PARTNERS	<u>EQ 10MG BASE/ML</u>	<u>N80722</u>	<u>001</u>	
-----------	-------------------	------------------------	---------------	------------	--

METAXALONE

TABLET; ORAL

SKELAXIN

+	JONES PHARMA INC	800MG	N13217	003	Aug 30, 2002
---	------------------	-------	--------	-----	--------------

METFORMIN HYDROCHLORIDE

SOLUTION; ORAL

RIOMET

+	RANBAXY	500MG/5ML	N21591	001	Sep 11, 2003
---	---------	-----------	--------	-----	--------------

TABLET; ORAL

GLUCOPHAGE

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>500MG</u>	<u>N20357</u>	<u>001</u>	Mar 03, 1995
-----------	----------------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N20357</u>	<u>002</u>	Mar 03, 1995
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	+	<u>1GM</u>	<u>N20357</u>	<u>005</u>	Nov 05, 1998
-----------	---	------------	---------------	------------	--------------

METFORMIN HYDROCHLORIDE

<u>AB</u>	ALPHAPHARM	<u>500MG</u>	<u>N75969</u>	<u>001</u>	Jan 29, 2002
-----------	------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75969</u>	<u>002</u>	Jan 29, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75969</u>	<u>003</u>	Jan 29, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	ANDRX PHARMS	<u>500MG</u>	<u>N75961</u>	<u>001</u>	Jan 25, 2002
-----------	--------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75961</u>	<u>002</u>	Jan 25, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75961</u>	<u>003</u>	Jan 25, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	AUROBINDO	<u>500MG</u>	<u>N77095</u>	<u>001</u>	Jan 14, 2005
-----------	-----------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N77095</u>	<u>002</u>	Jan 14, 2005
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N77095</u>	<u>003</u>	Jan 14, 2005
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	BARR	<u>500MG</u>	<u>N75971</u>	<u>001</u>	Jan 25, 2002
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75971</u>	<u>002</u>	Jan 25, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75971</u>	<u>003</u>	Jan 25, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	CARACO	<u>500MG</u>	<u>N75967</u>	<u>001</u>	Jan 29, 2002
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75967</u>	<u>002</u>	Jan 29, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75967</u>	<u>003</u>	Jan 29, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	GENPHARM	<u>500MG</u>	<u>N75973</u>	<u>001</u>	Jan 25, 2002
-----------	----------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75973</u>	<u>002</u>	Jan 25, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75973</u>	<u>003</u>	Jan 25, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	GOLDLINE	<u>500MG</u>	<u>N75972</u>	<u>001</u>	Jan 24, 2002
-----------	----------	--------------	---------------	------------	--------------

<u>AB</u>		<u>625MG</u>	<u>N75972</u>	<u>005</u>	Jan 24, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>750MG</u>	<u>N75972</u>	<u>004</u>	Jan 24, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75972</u>	<u>002</u>	Jan 24, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75972</u>	<u>003</u>	Jan 24, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	MUTUAL PHARMA	<u>500MG</u>	<u>N76038</u>	<u>001</u>	Feb 21, 2002
-----------	---------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N76038</u>	<u>002</u>	Feb 21, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N76038</u>	<u>003</u>	Feb 21, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>500MG</u>	<u>N75976</u>	<u>001</u>	Jan 24, 2002
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75976</u>	<u>002</u>	Jan 24, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75976</u>	<u>003</u>	Jan 24, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	PUREPAC PHARM	<u>500MG</u>	<u>N76033</u>	<u>001</u>	Jan 24, 2002
-----------	---------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N76033</u>	<u>002</u>	Jan 24, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N76033</u>	<u>003</u>	Jan 24, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>500MG</u>	<u>N75965</u>	<u>001</u>	Jan 25, 2002
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N75985</u>	<u>001</u>	Jan 25, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75965</u>	<u>002</u>	Jan 25, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>850MG</u>	<u>N75985</u>	<u>002</u>	Jan 25, 2002
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75965</u>	<u>003</u>	Jan 25, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N75985</u>	<u>003</u>	Jan 25, 2002
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>500MG</u>	<u>N75978</u>	<u>001</u>	Jan 25, 2002
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N76328</u>	<u>001</u>	Dec 16, 2002
-----------	--	--------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 242 (of 360)

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

<u>AB</u>	TEVA	<u>850MG</u>	<u>N75978</u>	<u>002</u>	Jan 25, 2002
<u>AB</u>		<u>850MG</u>	<u>N76328</u>	<u>002</u>	Dec 16, 2002
<u>AB</u>		<u>1GM</u>	<u>N75978</u>	<u>003</u>	Nov 05, 2002
<u>AB</u>		<u>1GM</u>	<u>N76328</u>	<u>003</u>	Dec 16, 2002
<u>AB</u>	TORPHARM	<u>500MG</u>	<u>N75984</u>	<u>001</u>	Apr 23, 2002
<u>AB</u>		<u>850MG</u>	<u>N75984</u>	<u>002</u>	Apr 23, 2002
<u>AB</u>		<u>1GM</u>	<u>N75984</u>	<u>003</u>	Apr 23, 2002
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>N75979</u>	<u>001</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>N75979</u>	<u>002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>N75979</u>	<u>003</u>	Jan 24, 2002
<u>AB</u>	ZENITH GOLDLINE	<u>500MG</u>	<u>N75975</u>	<u>001</u>	Jan 24, 2002
<u>AB</u>		<u>625MG</u>	<u>N75975</u>	<u>004</u>	Jan 24, 2002
<u>AB</u>		<u>750MG</u>	<u>N75975</u>	<u>005</u>	Jan 24, 2002
<u>AB</u>		<u>850MG</u>	<u>N75975</u>	<u>002</u>	Jan 24, 2002
<u>AB</u>		<u>1GM</u>	<u>N75975</u>	<u>003</u>	Jan 24, 2002
<u>AB</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>N77064</u>	<u>001</u>	Apr 18, 2005
<u>AB</u>		<u>850MG</u>	<u>N77064</u>	<u>002</u>	Apr 18, 2005
<u>AB</u>		<u>1GM</u>	<u>N77064</u>	<u>003</u>	Apr 18, 2005

TABLET, EXTENDED RELEASE; ORAL

FORTAMET

BX	ANDRX LABS LLC	500MG	N21574	001	Apr 27, 2004
BX	+	1GM	N21574	002	Apr 27, 2004

GLUCOPHAGE XR

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>500MG</u>	<u>N21202</u>	<u>001</u>	Oct 13, 2000
<u>AB</u>	+	<u>750MG</u>	<u>N21202</u>	<u>004</u>	Apr 11, 2003
	GLUMETZA				
BX	BIOVAIL	500MG	N21748	001	Jun 03, 2005
BX		1GM	N21748	002	Jun 03, 2005

METFORMIN HYDROCHLORIDE

<u>AB</u>	ANDRX PHARMS	<u>500MG</u>	<u>N76172</u>	<u>001</u>	Jun 16, 2004
<u>AB</u>		<u>750MG</u>	<u>N76869</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	APOTEX	<u>500MG</u>	<u>N76706</u>	<u>001</u>	Dec 14, 2004
<u>AB</u>		<u>750MG</u>	<u>N76706</u>	<u>002</u>	Dec 29, 2005
<u>AB</u>	BARR	<u>500MG</u>	<u>N76496</u>	<u>001</u>	Nov 25, 2005
<u>AB</u>		<u>750MG</u>	<u>N76863</u>	<u>001</u>	Oct 14, 2004
<u>AB</u>	COBALT	<u>500MG</u>	<u>N76818</u>	<u>001</u>	Dec 14, 2004
<u>AB</u>	IMPAX LABS	<u>500MG</u>	<u>N76249</u>	<u>001</u>	Jul 30, 2004
<u>AB</u>		<u>750MG</u>	<u>N76985</u>	<u>001</u>	Sep 13, 2005
<u>AB</u>	IVAX PHARMS	<u>500MG</u>	<u>N76545</u>	<u>001</u>	Dec 01, 2003
<u>AB</u>	MUTUAL PHARM	<u>500MG</u>	<u>N77124</u>	<u>001</u>	Dec 21, 2005
<u>AB</u>	MYLAN	<u>500MG</u>	<u>N76650</u>	<u>001</u>	Sep 13, 2005
<u>AB</u>		<u>750MG</u>	<u>N77113</u>	<u>001</u>	Sep 08, 2005
<u>AB</u>	PUREPAC PHARM	<u>500MG</u>	<u>N76450</u>	<u>001</u>	Oct 01, 2004
<u>AB</u>		<u>750MG</u>	<u>N76878</u>	<u>001</u>	Apr 13, 2005
<u>AB</u>	RANBAXY	<u>500MG</u>	<u>N76413</u>	<u>001</u>	Jun 18, 2004
<u>AB</u>		<u>750MG</u>	<u>N77211</u>	<u>001</u>	Jun 29, 2005
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>N76223</u>	<u>001</u>	Dec 14, 2004
<u>AB</u>		<u>500MG</u>	<u>N76873</u>	<u>001</u>	Dec 14, 2004
<u>AB</u>	TEVA	<u>500MG</u>	<u>N76269</u>	<u>001</u>	Jun 18, 2004
<u>AB</u>		<u>750MG</u>	<u>N76864</u>	<u>001</u>	Apr 12, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>500MG</u>	<u>N77060</u>	<u>001</u>	Apr 20, 2005
<u>AB</u>		<u>750MG</u>	<u>N77078</u>	<u>001</u>	Apr 21, 2005

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDAMET

	SB PHARMC0	500MG;EQ 1MG BASE	N21410	001	Oct 10, 2002
--	------------	-------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 243 (of 360)

METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDAMET

SB PHARMC0

500MG;EQ 2MG BASE

N21410 002

Oct 10, 2002

500MG;EQ 4MG BASE

N21410 003

Oct 10, 2002

1GM;EQ 2MG BASE

N21410 004

Aug 25, 2003

+

1GM;EQ 4MG BASE

N21410 005

Aug 25, 2003

METFORMIN; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

ACTOPLUS MET

TAKEDA GLOBAL

500MG;EQ 15MG BASE

N21842 001

Aug 29, 2005

+

850MG;EQ 15MG BASE

N21842 002

Aug 29, 2005

METHACHOLINE CHLORIDE

FOR SOLUTION; INHALATION

PROVOCHOLINE

+ METHAPHARM

100MG/VIAL

N19193 001

Oct 31, 1986

METHADONE HYDROCHLORIDE

CONCENTRATE; ORAL

METHADONE HYDROCHLORIDEAA ROXANE10MG/MLN40180 001

Apr 30, 1998

AA UDL10MG/MLN40088 001

Nov 30, 1994

METHADONE HYDROCHLORIDE INTENSOLAA ROXANE10MG/MLN89897 001

Sep 06, 1988

METHADOSEAA + MALLINCKRODT10MG/MLN17116 002

INJECTABLE; INJECTION

DOLOPHINE HYDROCHLORIDE

+ XANODYNE PHARM

10MG/ML

N21624 001

POWDER; FOR RX COMPOUNDING

METHADONE HYDROCHLORIDE

MALLINCKRODT

50GM/BOT

N06383 002

100GM/BOT

N06383 003

500GM/BOT

N06383 004

SOLUTION; ORAL

METHADONE HYDROCHLORIDE

+ ROXANE

5MG/5ML

N87393 001

+

10MG/5ML

N87997 001

Aug 30, 1982

TABLET; ORAL

DOLOPHINE HYDROCHLORIDEAA + ROXANE5MGN06134 002AA +10MGN06134 010METHADONE HYDROCHLORIDEAA MALLINCKRODT5MGN40517 001

Apr 27, 2004

AA10MGN40517 002

Apr 27, 2004

AA40MGN77142 001

Jul 12, 2005

AA

ROXANE

5MGN88108 001

Mar 08, 1983

AA10MGN88109 001

Mar 08, 1983

AA

+

40MGN17058 001AA40MGN74081 001

Apr 28, 1995

AA

SANDOZ

10MGN40241 002

May 29, 1998

AA40MGN75082 001

Mar 25, 1998

METHADOSEAA MALLINCKRODT5MGN40050 001

Apr 15, 1993

AA10MGN40050 002

Apr 15, 1993

AA40MGN74184 001

Apr 29, 1993

PRESCRIPTION DRUG PRODUCT LIST

3 - 244 (of 360)

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL
DESOXYN

+ OVATION PHARMS	5MG		N05378	002	
------------------	-----	--	--------	-----	--

METHAZOLAMIDE

TABLET; ORAL

METHAZOLAMIDE

<u>AB</u>	MIKART	25MG	<u>N40062</u>	<u>001</u>	Jan 27, 1994
<u>AB</u>	+	50MG	<u>N40062</u>	<u>002</u>	Jan 27, 1994
<u>AB</u>	SANDOZ	25MG	<u>N40036</u>	<u>001</u>	Jun 30, 1993
<u>AB</u>		25MG	<u>N40102</u>	<u>001</u>	Aug 28, 1996
<u>AB</u>		50MG	<u>N40036</u>	<u>002</u>	Jun 30, 1993
<u>AB</u>		50MG	<u>N40102</u>	<u>002</u>	Aug 28, 1996
<u>AB</u>	TEVA PHARMS	25MG	<u>N40001</u>	<u>001</u>	Jun 30, 1993
<u>AB</u>		50MG	<u>N40001</u>	<u>002</u>	Jun 30, 1993

METHENAMINE HIPPURATE

TABLET; ORAL

HIPREX

<u>AB</u>	+ AVENTIS PHARMS	1GM	<u>N17681</u>	<u>001</u>	
-----------	------------------	-----	---------------	------------	--

METHENAMINE HIPPURATE

<u>AB</u>	COREPHARMA	1GM	<u>N76411</u>	<u>001</u>	Jun 20, 2003
-----------	------------	-----	---------------	------------	--------------

UREX

<u>AB</u>	VATRING PHARMS	1GM	<u>N16151</u>	<u>001</u>	
-----------	----------------	-----	---------------	------------	--

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

<u>AB</u>	CEDAR PHARMS	5MG	<u>N40547</u>	<u>001</u>	Feb 18, 2005
<u>AB</u>		10MG	<u>N40547</u>	<u>002</u>	Feb 18, 2005
		15MG	N40619	003	Jul 12, 2005
		20MG	N40547	004	Feb 18, 2005
<u>AB</u>	GENPHARM	5MG	<u>N40350</u>	<u>001</u>	Mar 29, 2000
<u>AB</u>	+	10MG	<u>N40350</u>	<u>002</u>	Mar 29, 2000
<u>AB</u>	JONES PHARMA	5MG	<u>N40320</u>	<u>001</u>	Mar 31, 2000
<u>AB</u>		10MG	<u>N40320</u>	<u>002</u>	Mar 31, 2000
<u>AB</u>	SANDOZ	5MG	<u>N40411</u>	<u>001</u>	Mar 27, 2001
<u>AB</u>		10MG	<u>N40411</u>	<u>002</u>	Mar 27, 2001

METHOCARBAMOL

INJECTABLE; INJECTION

METHOCARBAMOL

<u>AP</u>	STERIS	100MG/ML	<u>N86459</u>	<u>001</u>	
-----------	--------	----------	---------------	------------	--

ROBAXIN

<u>AP</u>	+ BAXTER HLTHCARE CORP	100MG/ML	<u>N11790</u>	<u>001</u>	
-----------	------------------------	----------	---------------	------------	--

TABLET; ORAL

METHOCARBAMOL

<u>AA</u>	IMPAX LABS	500MG	<u>N84927</u>	<u>001</u>	
<u>AA</u>		750MG	<u>N84928</u>	<u>001</u>	
<u>AA</u>	LANNETT	500MG	<u>N84756</u>	<u>002</u>	Mar 31, 2003
<u>AA</u>		750MG	<u>N84756</u>	<u>001</u>	
<u>AA</u>	PAR PHARM	500MG	<u>N86989</u>	<u>001</u>	
<u>AA</u>		750MG	<u>N86988</u>	<u>001</u>	
<u>AA</u>	SANDOZ	500MG	<u>N84616</u>	<u>001</u>	
<u>AA</u>		500MG	<u>N87283</u>	<u>001</u>	
<u>AA</u>		750MG	<u>N84615</u>	<u>001</u>	
<u>AA</u>		750MG	<u>N87282</u>	<u>001</u>	
<u>AA</u>	TABLICAPS	500MG	<u>N84846</u>	<u>001</u>	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 245 (of 360)

METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL

<u>AA</u>	VINTAGE PHARMS	<u>500MG</u>	<u>N40489</u>	<u>001</u>	Jan 29, 2003
<u>AA</u>		<u>750MG</u>	<u>N40489</u>	<u>002</u>	Jan 29, 2003
<u>AA</u>	WATSON LABS	<u>500MG</u>	<u>N84277</u>	<u>001</u>	
<u>AA</u>		<u>500MG</u>	<u>N85180</u>	<u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>N84276</u>	<u>002</u>	
<u>AA</u>		<u>750MG</u>	<u>N85192</u>	<u>001</u>	
<u>AA</u>	WEST WARD	<u>500MG</u>	<u>N85159</u>	<u>001</u>	
<u>AA</u>		<u>750MG</u>	<u>N85123</u>	<u>001</u>	
<u>ROBAXIN</u>					
<u>AA</u>	+ SCHWARZ PHARMA	<u>500MG</u>	<u>N11011</u>	<u>004</u>	
<u>ROBAXIN-750</u>					
<u>AA</u>	+ SCHWARZ PHARMA	<u>750MG</u>	<u>N11011</u>	<u>006</u>	

METHOHEXITAL SODIUMINJECTABLE; INJECTION
BREVITAL SODIUM

+	KING PHARMS	500MG/VIAL	N11559	001	
+		2.5GM/VIAL	N11559	002	
+		5GM/VIAL	N11559	003	

METHOTREXATEINJECTABLE; INJECTION
METHOTREXATE PRESERVATIVE FREE

+	BEDFORD	EQ 1GM BASE/VIAL	N40632	001	Aug 12, 2005
---	---------	------------------	--------	-----	--------------

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE

<u>AP</u>	+ MAYNE PHARMA USA	<u>EQ 50MG BASE/2ML (25 MG/ML)</u>	<u>N11719</u>	<u>010</u>	Dec 15, 2004
<u>METHOTREXATE PRESERVATIVE FREE</u>					
<u>AP</u>	+ MAYNE PHARMA USA	<u>EQ 1GM BASE/40ML (25 MG/ML)</u>	<u>N11719</u>	<u>012</u>	Apr 13, 2005
<u>METHOTREXATE SODIUM</u>					
<u>AP</u>	BEDFORD	<u>EQ 50 MG BASE/2ML (25 ML/ML)</u>	<u>N89340</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>		<u>EQ 100MG BASE/4ML (25 MG/ML)</u>	<u>N89341</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>		<u>EQ 200MG BASE/8ML (25 MG/ML)</u>	<u>N89342</u>	<u>001</u>	Sep 16, 1986
<u>AP</u>		<u>EQ 250MG BASE/10ML (25 MG/ML)</u>	<u>N89343</u>	<u>001</u>	Sep 16, 1986

TABLET; ORAL

METHOTREXATE SODIUM

<u>AB</u>	BARR	<u>EQ 2.5MG BASE</u>	<u>N81099</u>	<u>001</u>	Oct 15, 1990
<u>AB</u>	+ CLONMEL HLTHCARE	<u>EQ 2.5MG BASE</u>	<u>N08085</u>	<u>002</u>	
<u>AB</u>	DURAMED PHARMS BARR	<u>EQ 2.5MG BASE</u>	<u>N40233</u>	<u>001</u>	Jun 17, 1999
<u>AB</u>	MYLAN	<u>EQ 2.5MG BASE</u>	<u>N81235</u>	<u>001</u>	May 15, 1992
<u>AB</u>	ROXANE	<u>EQ 2.5MG BASE</u>	<u>N40054</u>	<u>001</u>	Aug 01, 1994
TREXALL					
	BARR	EQ 5MG BASE	N40385	001	Mar 21, 2001
		EQ 7.5MG BASE	N40385	002	Mar 21, 2001
		EQ 10MG BASE	N40385	003	Mar 21, 2001
+		EQ 15MG BASE	N40385	004	Mar 21, 2001

METHOXSALLENCAPSULE; ORAL
8-MOP

+	VALEANT PHARM INTL	10MG	N09048	001	
OXSORALEN-ULTRA					
+	VALEANT PHARM INTL	10MG	N19600	001	Oct 30, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 246 (of 360)

METHOXSALEN

INJECTABLE; INJECTION

UVADEX

+	THERAKOS	0.02MG/ML	N20969	001	Feb 25, 1999
---	----------	-----------	--------	-----	--------------

LOTION; TOPICAL

OXSORALEN

+	VALEANT PHARM INTL	1%	N09048	002	
---	--------------------	----	--------	-----	--

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

PAMINE

+	BRADLEY PHARMS	2.5MG	N08848	001	
---	----------------	-------	--------	-----	--

PAMINE FORTE

+	BRADLEY PHARMS	5MG	N08848	002	Mar 25, 2003
---	----------------	-----	--------	-----	--------------

METHSUXIMIDE

CAPSULE; ORAL

CELONTIN

	PARKE DAVIS	150MG	N10596	007	
--	-------------	-------	--------	-----	--

+		300MG	N10596	008	
---	--	-------	--------	-----	--

METHYLCLOTHIAZIDE

TABLET; ORAL

ENDURON

<u>AB</u>	ABBOTT	<u>2.5MG</u>	<u>N12524</u>	<u>001</u>	
-----------	--------	--------------	---------------	------------	--

<u>AB</u>	+	<u>5MG</u>	<u>N12524</u>	<u>004</u>	
-----------	---	------------	---------------	------------	--

METHYLCLOTHIAZIDE

<u>AB</u>	IVAX PHARMS	<u>2.5MG</u>	<u>N87913</u>	<u>001</u>	Jun 03, 1982
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>5MG</u>	<u>N87672</u>	<u>001</u>	Aug 17, 1982
-----------	-------	------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>N89835</u>	<u>001</u>	Aug 18, 1988
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>5MG</u>	<u>N89837</u>	<u>001</u>	Aug 18, 1988
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>N88724</u>	<u>001</u>	Sep 06, 1984
-----------	-------------	------------	---------------	------------	--------------

METHYL AMINOLEVULINATE HYDROCHLORIDE

CREAM; TOPICAL

METVIXIA

+	PHOTOCURE ASA	EQ 16.8% BASE	N21415	001	Jul 27, 2004
---	---------------	---------------	--------	-----	--------------

METHYLDOPA

TABLET; ORAL

METHYLDOPA

<u>AB</u>	ACCORD HLTH	<u>250MG</u>	<u>N70084</u>	<u>001</u>	Oct 15, 1985
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N70085</u>	<u>001</u>	Oct 15, 1985
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	IVAX PHARMS	<u>250MG</u>	<u>N70098</u>	<u>001</u>	Feb 20, 1986
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N70343</u>	<u>001</u>	Feb 20, 1986
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>250MG</u>	<u>N70075</u>	<u>001</u>	Apr 18, 1985
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>	+	<u>500MG</u>	<u>N70076</u>	<u>001</u>	Apr 18, 1985
-----------	---	--------------	---------------	------------	--------------

<u>AB</u>	PAR PHARM	<u>125MG</u>	<u>N70535</u>	<u>001</u>	Jan 02, 1987
-----------	-----------	--------------	---------------	------------	--------------

<u>AB</u>		<u>250MG</u>	<u>N70536</u>	<u>001</u>	Jan 02, 1987
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N70537</u>	<u>001</u>	Jan 02, 1987
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	PLIVA	<u>125MG</u>	<u>N72126</u>	<u>001</u>	Jul 07, 1988
-----------	-------	--------------	---------------	------------	--------------

<u>AB</u>		<u>250MG</u>	<u>N72127</u>	<u>001</u>	Jul 07, 1988
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N72128</u>	<u>001</u>	Jul 07, 1988
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>125MG</u>	<u>N71700</u>	<u>001</u>	Mar 02, 1988
-----------	--------	--------------	---------------	------------	--------------

<u>AB</u>		<u>250MG</u>	<u>N18934</u>	<u>001</u>	Jun 29, 1984
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N18934</u>	<u>002</u>	Jun 29, 1984
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>N70703</u>	<u>001</u>	Jun 06, 1986
-----------	-------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>500MG</u>	<u>N70625</u>	<u>001</u>	Jun 06, 1986
-----------	--	--------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 247 (of 360)

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

METHYLDOPATE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>50MG/ML</u>	<u>N70698</u>	<u>001</u>	Jun 15, 1987
<u>AP</u>		<u>50MG/ML</u>	<u>N70699</u>	<u>001</u>	Jun 15, 1987
<u>AP</u>	+ LUITPOLD	<u>50MG/ML</u>	<u>N71279</u>	<u>001</u>	Oct 02, 1987
<u>AP</u>	SICOR PHARMS	<u>50MG/ML</u>	<u>N72974</u>	<u>001</u>	Nov 22, 1991

METHYLERGONOVINE MALEATE

INJECTABLE; INJECTION

METHERGINE

+ NOVARTIS 0.2MG/ML

N06035 004

TABLET; ORAL

METHERGINE

+ NOVARTIS 0.2MG

N06035 003

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

METADATE CD

BX	UCB	10MG	N21259	003	May 27, 2003
BX		20MG	N21259	001	Apr 03, 2001
BX	+	30MG	N21259	002	Jun 19, 2003
	RITALIN LA				
BX	NOVARTIS	10MG	N21284	004	Apr 10, 2004
BX		20MG	N21284	001	Jun 05, 2002
BX		30MG	N21284	002	Jun 05, 2002
	+	40MG	N21284	003	Jun 05, 2002

SOLUTION; ORAL

METHYLIN

+ MALLINCKRODT BAKER 5MG/5ML

N21419 001 Dec 19, 2002

+ 10MG/5ML

N21419 002 Dec 19, 2002

TABLET; ORAL

METHYLPHENIDATE HCL

<u>AB</u>	UCB	<u>5MG</u>	<u>N86429</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>N85799</u>	<u>001</u>	
<u>AB</u>		<u>20MG</u>	<u>N86428</u>	<u>001</u>	

METHYLPHENIDATE HYDROCHLORIDE

<u>AB</u>	MALLINCKRODT	<u>5MG</u>	<u>N40300</u>	<u>001</u>	Nov 27, 1998
<u>AB</u>		<u>10MG</u>	<u>N40300</u>	<u>002</u>	Nov 27, 1998
<u>AB</u>		<u>20MG</u>	<u>N40300</u>	<u>003</u>	Nov 27, 1998
<u>AB</u>	PUREPAC PHARM	<u>5MG</u>	<u>N40321</u>	<u>001</u>	Feb 05, 2002
<u>AB</u>		<u>10MG</u>	<u>N40321</u>	<u>002</u>	Feb 05, 2002
<u>AB</u>		<u>20MG</u>	<u>N40321</u>	<u>003</u>	Feb 05, 2002
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>N40220</u>	<u>001</u>	Aug 29, 1997
<u>AB</u>		<u>10MG</u>	<u>N40220</u>	<u>002</u>	Aug 29, 1997
<u>AB</u>		<u>20MG</u>	<u>N40220</u>	<u>003</u>	Aug 29, 1997
	RITALIN				
<u>AB</u>	NOVARTIS	<u>5MG</u>	<u>N10187</u>	<u>003</u>	
<u>AB</u>		<u>10MG</u>	<u>N10187</u>	<u>006</u>	
<u>AB</u>	+	<u>20MG</u>	<u>N10187</u>	<u>010</u>	

TABLET, CHEWABLE; ORAL

METHYLIN

MALLINCKRODT 2.5MG

N21475 001 Apr 15, 2003

5MG

N21475 002 Apr 15, 2003

+ 10MG

N21475 003 Apr 15, 2003

TABLET, EXTENDED RELEASE; ORAL

CONCERTA

ALZA 18MG

N21121 001 Aug 01, 2000

27MG

N21121 004 Apr 01, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 248 (of 360)

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL					
CONCERTA					
	ALZA	36MG	N21121	002	Aug 01, 2000
	+	54MG	N21121	003	Dec 08, 2000
<u>METADATE ER</u>					
<u>AB</u>	UCB	<u>10MG</u>	<u>N40306</u>	<u>001</u>	Oct 20, 1999
<u>AB</u>	+	<u>20MG</u>	<u>N89601</u>	<u>001</u>	Jun 01, 1988
<u>METHYLIN ER</u>					
<u>AB</u>	MALLINCKRODT	<u>10MG</u>	<u>N75629</u>	<u>001</u>	May 09, 2000
<u>AB</u>		<u>20MG</u>	<u>N75629</u>	<u>002</u>	May 09, 2000
<u>METHYLPHENIDATE HYDROCHLORIDE</u>					
<u>AB</u>	PUREPAC PHARM	<u>20MG</u>	<u>N75450</u>	<u>001</u>	Dec 21, 2001
<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>N40410</u>	<u>001</u>	Feb 09, 2001
<u>RITALIN-SR</u>					
<u>AB</u>	NOVARTIS	<u>20MG</u>	<u>N18029</u>	<u>001</u>	Mar 30, 1982

METHYLPREDNISOLONE

TABLET; ORAL					
<u>MEDROL</u>					
<u>AB</u>	PHARMACIA AND UPJOHN	<u>4MG</u>	<u>N11153</u>	<u>001</u>	
<u>AB</u>		<u>8MG</u>	<u>N11153</u>	<u>004</u>	
<u>AB</u>		<u>16MG</u>	<u>N11153</u>	<u>003</u>	
<u>AB</u>		<u>24MG</u>	<u>N11153</u>	<u>005</u>	
<u>AB</u>	+	<u>32MG</u>	<u>N11153</u>	<u>006</u>	
<u>METHYLPREDNISOLONE</u>					
<u>AB</u>	DURAMED PHARMS BARR	<u>4MG</u>	<u>N88497</u>	<u>001</u>	Feb 21, 1984
<u>AB</u>	PAR PHARM	<u>16MG</u>	<u>N89207</u>	<u>001</u>	Apr 25, 1988
<u>AB</u>		<u>24MG</u>	<u>N89208</u>	<u>001</u>	Apr 25, 1988
<u>AB</u>		<u>32MG</u>	<u>N89209</u>	<u>001</u>	Apr 25, 1988
<u>AB</u>	SANDOZ	<u>4MG</u>	<u>N40194</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>	TRIGEN	<u>4MG</u>	<u>N40189</u>	<u>001</u>	Oct 31, 1997
<u>AB</u>		<u>8MG</u>	<u>N40189</u>	<u>002</u>	Oct 31, 1997
<u>AB</u>	VINTAGE PHARMS	<u>4MG</u>	<u>N40183</u>	<u>001</u>	Dec 22, 1998
<u>AB</u>	WATSON LABS	<u>4MG</u>	<u>N40232</u>	<u>001</u>	Oct 16, 1997

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION					
<u>DEPO-MEDROL</u>					
<u>AB</u>	+	<u>PHARMACIA AND UPJOHN 40MG/ML</u>	<u>N11757</u>	<u>001</u>	
<u>AB</u>	+	<u>80MG/ML</u>	<u>N11757</u>	<u>004</u>	
	+	<u>20MG/ML</u>	N11757	002	
<u>METHYLPREDNISOLONE ACETATE</u>					
<u>AB</u>	SICOR PHARMS	<u>40MG/ML</u>	<u>N40557</u>	<u>001</u>	Feb 23, 2005
<u>AB</u>		<u>80MG/ML</u>	<u>N40557</u>	<u>002</u>	Feb 23, 2005

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION					
<u>A-METHAPRED</u>					
<u>AP</u>	HOSPIRA	<u>EQ 40MG BASE/VIAL</u>	<u>N40664</u>	<u>001</u>	Dec 20, 2005
<u>AP</u>		<u>EQ 40MG BASE/VIAL</u>	<u>N85853</u>	<u>001</u>	
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>N40665</u>	<u>001</u>	Dec 20, 2005
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>N85855</u>	<u>001</u>	
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>N85854</u>	<u>001</u>	
<u>AP</u>		<u>EQ 500MG BASE/VIAL</u>	<u>N89173</u>	<u>001</u>	Aug 18, 1987
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N85852</u>	<u>001</u>	
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N89174</u>	<u>001</u>	Aug 18, 1987
<u>METHYLPREDNISOLONE SODIUM SUCCINATE</u>					
<u>AP</u>	AM PHARM	<u>EQ 40MG BASE/VIAL</u>	<u>N40583</u>	<u>001</u>	Jul 30, 2004
<u>AP</u>		<u>EQ 125MG BASE/VIAL</u>	<u>N40583</u>	<u>002</u>	Jul 30, 2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 249 (of 360)

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

<u>AP</u>	AM PHARM	<u>EQ 1GM BASE/VIAL</u>	<u>N40612</u>	<u>001</u>	Aug 12, 2004
<u>AP</u>	SICOR PHARMS	<u>EQ 125MG BASE/VIAL</u>	<u>N81266</u>	<u>001</u>	Nov 30, 1992
<u>SOLU-MEDROL</u>					
<u>AP</u>	+ PHARMACIA AND UPJOHN	<u>EQ 40MG BASE/VIAL</u>	<u>N11856</u>	<u>003</u>	
<u>AP</u>	+	<u>EQ 125MG BASE/VIAL</u>	<u>N11856</u>	<u>004</u>	
<u>AP</u>	+	<u>EQ 500MG BASE/VIAL</u>	<u>N11856</u>	<u>005</u>	
<u>AP</u>	+	<u>EQ 1GM BASE/VIAL</u>	<u>N11856</u>	<u>006</u>	
	+	EQ 2GM BASE/VIAL	N11856	007	Feb 27, 1985

METHYLTESTOSTERONE

CAPSULE; ORAL

TESTRED

BP	+ VALEANT PHARM INTL	10MG	N83976	001	
	VIRILON				
BP	STAR PHARMS FL	10MG	N87750	001	Nov 24, 1982

TABLET; ORAL

ANDROID 10

BP	VALEANT PHARM INTL	10MG	N86450	001	
----	--------------------	------	--------	-----	--

ANDROID 25

BP	+ VALEANT PHARM INTL	25MG	N87147	001	
----	----------------------	------	--------	-----	--

METHYLTESTOSTERONE

BP	IMPAX LABS	10MG	N80767	002	
----	------------	------	--------	-----	--

BP		25MG	N84310	001	
----	--	------	--------	-----	--

METIPRANOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

METIPRANOLOL

<u>AT</u>	FALCON PHARMS	<u>0.3%</u>	<u>N75720</u>	<u>001</u>	Aug 06, 2001
-----------	---------------	-------------	---------------	------------	--------------

OPTIPRANOLOL

<u>AT</u>	+ BAUSCH AND LOMB	<u>0.3%</u>	<u>N19907</u>	<u>001</u>	Dec 29, 1989
-----------	-------------------	-------------	---------------	------------	--------------

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>EQ 5MG BASE/ML</u>	<u>N70505</u>	<u>001</u>	Jun 23, 1989
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N73117</u>	<u>001</u>	Jan 17, 1991
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N73118</u>	<u>001</u>	Jan 17, 1991
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N74147</u>	<u>001</u>	Aug 02, 1996
<u>AP</u>	MAYNE PHARMA USA	<u>EQ 5MG BASE/ML</u>	<u>N70847</u>	<u>001</u>	Nov 07, 1988
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N71291</u>	<u>001</u>	Mar 03, 1989
<u>AP</u>	SICOR PHARMS	<u>EQ 5MG BASE/ML</u>	<u>N73135</u>	<u>001</u>	Nov 27, 1991
<u>REGLAN</u>					
<u>AP</u>	+ BAXTER HLTHCARE CORP	<u>EQ 5MG BASE/ML</u>	<u>N17862</u>	<u>001</u>	

SOLUTION; ORAL

METOCLOPRAMIDE

<u>AA</u>	JVL	<u>EQ 5MG BASE/5ML</u>	<u>N74703</u>	<u>001</u>	Oct 31, 1997
<u>AA</u>	VISTAPHARM	<u>EQ 5MG BASE/5ML</u>	<u>N75051</u>	<u>001</u>	Jan 26, 2001

METOCLOPRAMIDE HYDROCHLORIDE

<u>AA</u>	ALPHARMA US PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>N71340</u>	<u>001</u>	Aug 18, 1988
<u>AA</u>	PHARM ASSOC	<u>EQ 5MG BASE/5ML</u>	<u>N72744</u>	<u>001</u>	May 28, 1991
<u>AA</u>	PHARM VENTURES	<u>EQ 5MG BASE/5ML</u>	<u>N71402</u>	<u>001</u>	Jun 25, 1993
<u>AA</u>	ROXANE	<u>EQ 5MG BASE/5ML</u>	<u>N72038</u>	<u>001</u>	Dec 05, 1988
<u>AA</u>	SILARX	<u>EQ 5MG BASE/5ML</u>	<u>N73680</u>	<u>001</u>	Oct 27, 1992
<u>AA</u>	+ TEVA	<u>EQ 5MG BASE/5ML</u>	<u>N70819</u>	<u>001</u>	Jul 10, 1987
<u>AA</u>		<u>EQ 5MG BASE/5ML</u>	<u>N71315</u>	<u>001</u>	Jun 30, 1993

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 250 (of 360)

METOCLOPRAMIDE HYDROCHLORIDE

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

<u>AB</u>	PLIVA	<u>EQ 5MG BASE</u>	<u>N72750</u>	<u>001</u>	Dec 28, 1995
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N71250</u>	<u>001</u>	Feb 03, 1988
<u>AB</u>	PUREPAC PHARM	<u>EQ 10MG BASE</u>	<u>N70581</u>	<u>001</u>	Oct 17, 1985
<u>AB</u>	SANDOZ	<u>EQ 5MG BASE</u>	<u>N74478</u>	<u>001</u>	Oct 05, 1995
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N72215</u>	<u>001</u>	Jan 30, 1990
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N74478</u>	<u>002</u>	Oct 05, 1995
<u>AB</u>	TEVA	<u>EQ 5MG BASE</u>	<u>N72801</u>	<u>001</u>	Jun 15, 1993
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N70184</u>	<u>001</u>	Jul 29, 1985
<u>AB</u>	WATSON LABS	<u>EQ 10MG BASE</u>	<u>N70511</u>	<u>001</u>	Jan 22, 1986
<u>REGLAN</u>					
<u>AB</u>	SCHWARZ PHARMA	<u>EQ 5MG BASE</u>	<u>N17854</u>	<u>002</u>	May 05, 1987
<u>AB</u>	+	<u>EQ 10MG BASE</u>	<u>N17854</u>	<u>001</u>	

TABLET, ORALLY DISINTEGRATING; ORAL

REGLAN ODT

	SCHWARZ PHARMA	EQ 5MG BASE	N21793	001	Jun 10, 2005
+		EQ 10MG BASE	N21793	002	Jun 10, 2005

METOLAZONE

TABLET; ORAL

METOLAZONE

<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>N76698</u>	<u>001</u>	Dec 23, 2003
<u>AB</u>		<u>5MG</u>	<u>N76698</u>	<u>002</u>	Oct 19, 2004
<u>AB</u>		<u>10MG</u>	<u>N76698</u>	<u>003</u>	Oct 19, 2004
<u>AB</u>	ROXANE	<u>10MG</u>	<u>N76482</u>	<u>002</u>	Apr 29, 2004
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>N76732</u>	<u>001</u>	Dec 19, 2003
<u>AB</u>		<u>5MG</u>	<u>N76466</u>	<u>001</u>	Dec 19, 2003
<u>AB</u>		<u>10MG</u>	<u>N76466</u>	<u>002</u>	Dec 19, 2003
<u>AB</u>	TEVA	<u>2.5MG</u>	<u>N76600</u>	<u>001</u>	Jan 06, 2004
<u>AB</u>		<u>5MG</u>	<u>N76833</u>	<u>001</u>	Mar 01, 2004
<u>AB</u>		<u>10MG</u>	<u>N75543</u>	<u>003</u>	Dec 24, 2003
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N76891</u>	<u>001</u>	Jul 21, 2004
<u>ZAROXOLYN</u>					
<u>AB</u>	UCB	<u>2.5MG</u>	<u>N17386</u>	<u>001</u>	
<u>AB</u>	+	<u>5MG</u>	<u>N17386</u>	<u>002</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N17386</u>	<u>003</u>	

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

TOPROL-XL

	ASTRAZENECA	EQ 25MG TARTRATE	N19962	004	Feb 05, 2001
+		EQ 50MG TARTRATE	N19962	001	Jan 10, 1992
		EQ 100MG TARTRATE	N19962	002	Jan 10, 1992
+		EQ 200MG TARTRATE	N19962	003	Jan 10, 1992

METOPROLOL TARTRATE

INJECTABLE; INJECTION

LOPRESSOR

<u>AP</u>	+	NOVARTIS	<u>1MG/ML</u>	<u>N18704</u>	<u>001</u>	Mar 30, 1984
<u>METOPROLOL TARTRATE</u>						
<u>AP</u>		BEDFORD LABS	<u>1MG/ML</u>	<u>N76495</u>	<u>001</u>	Jul 07, 2003
<u>AP</u>		HOSPIRA	<u>1MG/ML</u>	<u>N74133</u>	<u>001</u>	Dec 21, 1993
<u>AP</u>			<u>1MG/ML</u>	<u>N75160</u>	<u>001</u>	Jul 06, 1998
<u>AP</u>		STERIS	<u>1MG/ML</u>	<u>N74032</u>	<u>001</u>	Dec 21, 1993

TABLET; ORAL

LOPRESSOR

<u>AB</u>	NOVARTIS	<u>50MG</u>	<u>N17963</u>	<u>001</u>	
-----------	----------	-------------	---------------	------------	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 251 (of 360)

METOPROLOL TARTRATE

TABLET; ORAL				
<u>LOPRESSOR</u>				
<u>AB</u>	NOVARTIS	<u>100MG</u>	<u>N17963</u>	<u>002</u>
<u>METOPROLOL TARTRATE</u>				
<u>AB</u>	APOTHECON	<u>50MG</u>	<u>N74258</u>	<u>001</u> Jan 27, 1994
<u>AB</u>		<u>100MG</u>	<u>N74258</u>	<u>002</u> Jan 27, 1994
<u>AB</u>	CARACO	<u>25MG</u>	<u>N76670</u>	<u>001</u> Jan 15, 2004
<u>AB</u>		<u>50MG</u>	<u>N74644</u>	<u>001</u> Dec 10, 1996
<u>AB</u>		<u>100MG</u>	<u>N74644</u>	<u>002</u> Dec 10, 1996
<u>AB</u>	MUTUAL PHARM	<u>50MG</u>	<u>N73653</u>	<u>001</u> Dec 21, 1993
<u>AB</u>		<u>100MG</u>	<u>N73654</u>	<u>001</u> Dec 21, 1993
<u>AB</u>	MYLAN	<u>25MG</u>	<u>N76704</u>	<u>001</u> Jan 16, 2004
<u>AB</u>		<u>50MG</u>	<u>N73666</u>	<u>001</u> Dec 21, 1993
<u>AB</u>		<u>50MG</u>	<u>N76704</u>	<u>002</u> Jan 16, 2004
<u>AB</u>		<u>100MG</u>	<u>N73666</u>	<u>002</u> Dec 21, 1993
<u>AB</u>	+	<u>100MG</u>	<u>N76704</u>	<u>003</u> Jan 16, 2004
<u>AB</u>	PAR PHARM	<u>50MG</u>	<u>N74453</u>	<u>001</u> Apr 27, 1995
<u>AB</u>		<u>100MG</u>	<u>N74453</u>	<u>002</u> Apr 27, 1995
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N73288</u>	<u>001</u> Mar 25, 1994
<u>AB</u>		<u>100MG</u>	<u>N73289</u>	<u>001</u> Mar 25, 1994
<u>AB</u>	TEVA	<u>50MG</u>	<u>N74141</u>	<u>001</u> Jan 31, 1995
<u>AB</u>		<u>50MG</u>	<u>N74143</u>	<u>001</u> Sep 30, 1994
<u>AB</u>		<u>100MG</u>	<u>N74141</u>	<u>002</u> Jan 31, 1995
<u>AB</u>		<u>100MG</u>	<u>N74143</u>	<u>002</u> Sep 30, 1994
<u>AB</u>	TEVA PHARMS	<u>50MG</u>	<u>N74333</u>	<u>001</u> Jan 27, 1994
<u>AB</u>		<u>100MG</u>	<u>N74333</u>	<u>002</u> Jan 27, 1994
<u>AB</u>	WATSON LABS	<u>50MG</u>	<u>N74217</u>	<u>001</u> May 27, 1994
<u>AB</u>		<u>100MG</u>	<u>N74217</u>	<u>002</u> May 27, 1994

METRONIDAZOLE

CAPSULE; ORAL				
<u>FLAGYL</u>				
<u>AB</u>	+	GD SEARLE LLC	<u>375MG</u>	<u>N20334</u> <u>001</u> May 03, 1995
<u>METRONIDAZOLE</u>				
<u>AB</u>		KALI LABS	<u>375MG</u>	<u>N76522</u> <u>001</u> Jan 29, 2004
CREAM; TOPICAL				
<u>METROCREAM</u>				
<u>AB</u>	+	GALDERMA LABS LP	<u>0.75%</u>	<u>N20531</u> <u>001</u> Sep 20, 1995
<u>METRONIDAZOLE</u>				
<u>AB</u>		ALTANA	<u>0.75%</u>	<u>N76408</u> <u>001</u> May 28, 2004
		NORITATE		
	+	DERMIK LABS	1%	N20743 001 Sep 26, 1997
GEL; TOPICAL				
METROGEL				
	+	GALDERMA LABS LP	0.75%	N19737 001 Nov 22, 1988
	+		1%	N21789 001 Jun 30, 2005
GEL; VAGINAL				
METROGEL-VAGINAL				
BX	+	3M	0.75%	N20208 001 Aug 17, 1992
VANDAZOLE				
BX		TEVA PHARMS	0.75%	N21806 001 May 20, 2005
INJECTABLE; INJECTION				
<u>FLAGYL I.V. RTU IN PLASTIC CONTAINER</u>				
<u>AP</u>	+	BAXTER HLTHCARE	<u>500MG/100ML</u>	<u>N18657</u> <u>001</u>
<u>AP</u>	+	GD SEARLE LLC	<u>500MG/100ML</u>	<u>N18353</u> <u>002</u>
<u>METRO I.V. IN PLASTIC CONTAINER</u>				
<u>AP</u>	+	B BRAUN	<u>500MG/100ML</u>	<u>N18900</u> <u>001</u> Sep 29, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 252 (of 360)

METRONIDAZOLE

INJECTABLE; INJECTION

METRONIDAZOLE IN PLASTIC CONTAINER

<u>AP</u>	+	HOSPIRA	<u>500MG/100ML</u>	<u>N18890</u>	<u>002</u>	Nov 18, 1983
-----------	---	---------	--------------------	---------------	------------	--------------

LOTION; TOPICAL

METROLOTION

	+	GALDERMA LABS LP	0.75%	N20901	001	Nov 24, 1998
--	---	------------------	-------	--------	-----	--------------

TABLET; ORAL

FLAGYL

<u>AB</u>		GD SEARLE LLC	<u>250MG</u>	<u>N12623</u>	<u>001</u>	
-----------	--	---------------	--------------	---------------	------------	--

<u>AB</u>	+		<u>500MG</u>	<u>N12623</u>	<u>003</u>	
-----------	---	--	--------------	---------------	------------	--

METRONIDAZOLE

<u>AB</u>		IVAX PHARMS	<u>250MG</u>	<u>N18517</u>	<u>001</u>	
-----------	--	-------------	--------------	---------------	------------	--

<u>AB</u>			<u>500MG</u>	<u>N18517</u>	<u>002</u>	May 05, 1982
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		LNK	<u>250MG</u>	<u>N19029</u>	<u>001</u>	Apr 10, 1984
-----------	--	-----	--------------	---------------	------------	--------------

<u>AB</u>		MUTUAL PHARM	<u>250MG</u>	<u>N70772</u>	<u>001</u>	Jul 16, 1986
-----------	--	--------------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N70773</u>	<u>001</u>	Jul 16, 1986
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		PAR PHARM	<u>250MG</u>	<u>N18845</u>	<u>001</u>	Aug 18, 1983
-----------	--	-----------	--------------	---------------	------------	--------------

<u>AB</u>			<u>250MG</u>	<u>N70040</u>	<u>001</u>	Jan 29, 1985
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N18930</u>	<u>001</u>	Aug 18, 1983
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N70039</u>	<u>001</u>	Jan 29, 1985
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		PLIVA	<u>250MG</u>	<u>N70027</u>	<u>001</u>	Nov 06, 1984
-----------	--	-------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N70033</u>	<u>001</u>	Dec 06, 1984
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>250MG</u>	<u>N18620</u>	<u>001</u>	Mar 04, 1982
-----------	--	--------	--------------	---------------	------------	--------------

<u>AB</u>			<u>250MG</u>	<u>N18740</u>	<u>001</u>	Oct 22, 1982
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N18620</u>	<u>002</u>	Jun 02, 1983
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N18740</u>	<u>002</u>	Oct 22, 1982
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		TEVA	<u>250MG</u>	<u>N70035</u>	<u>001</u>	Dec 20, 1984
-----------	--	------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N70044</u>	<u>001</u>	Feb 08, 1985
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		WATSON LABS	<u>250MG</u>	<u>N18764</u>	<u>001</u>	Sep 17, 1982
-----------	--	-------------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N18764</u>	<u>002</u>	Dec 20, 1982
-----------	--	--	--------------	---------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

FLAGYL ER

	+	GD SEARLE LLC	750MG	N20868	001	Nov 26, 1997
--	---	---------------	-------	--------	-----	--------------

METRONIDAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

FLAGYL I.V.

	+	GD SEARLE LLC	EQ 500MG BASE/VIAL	N18353	001	
--	---	---------------	--------------------	--------	-----	--

METYRAPONE

CAPSULE; ORAL

METOPIRONE

	+	NOVARTIS	250MG	N12911	002	Aug 09, 1996
--	---	----------	-------	--------	-----	--------------

METYROSINE

CAPSULE; ORAL

DEMSEER

	+	MERCK	250MG	N17871	001	
--	---	-------	-------	--------	-----	--

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

<u>AB</u>		SANDOZ	<u>150MG</u>	<u>N74450</u>	<u>001</u>	May 16, 1996
-----------	--	--------	--------------	---------------	------------	--------------

<u>AB</u>			<u>200MG</u>	<u>N74450</u>	<u>002</u>	May 16, 1996
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>			<u>250MG</u>	<u>N74450</u>	<u>003</u>	May 16, 1996
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		TEVA	<u>150MG</u>	<u>N74377</u>	<u>001</u>	May 16, 1995
-----------	--	------	--------------	---------------	------------	--------------

<u>AB</u>			<u>200MG</u>	<u>N74377</u>	<u>002</u>	May 16, 1995
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>			<u>250MG</u>	<u>N74377</u>	<u>003</u>	May 16, 1995
-----------	--	--	--------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 253 (of 360)

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>N74711</u>	<u>001</u>	Feb 26, 1997
<u>AB</u>		<u>150MG</u>	<u>N74865</u>	<u>001</u>	Apr 13, 1998
<u>AB</u>		<u>200MG</u>	<u>N74711</u>	<u>002</u>	Feb 26, 1997
<u>AB</u>		<u>200MG</u>	<u>N74865</u>	<u>002</u>	Apr 13, 1998
<u>AB</u>		<u>250MG</u>	<u>N74711</u>	<u>003</u>	Feb 26, 1997
<u>AB</u>		<u>250MG</u>	<u>N74865</u>	<u>003</u>	Apr 13, 1998
<u>MEXITIL</u>					
<u>AB</u>	BOEHRINGER INGELHEIM	<u>150MG</u>	<u>N18873</u>	<u>002</u>	Dec 30, 1985
<u>AB</u>		<u>200MG</u>	<u>N18873</u>	<u>003</u>	Dec 30, 1985
<u>AB</u>	+	<u>250MG</u>	<u>N18873</u>	<u>004</u>	Dec 30, 1985

MICAFUNGIN SODIUM

INJECTABLE; IV (INFUSION)

MYCAMINE

+	ASTELLAS	50MG/VIAL	N21506	002	Mar 16, 2005
---	----------	-----------	--------	-----	--------------

MICONAZOLE NITRATE

CREAM; TOPICAL

MONISTAT-DERM

+	JOHNSON AND JOHNSON	2%	N17494	001	
---	---------------------	----	--------	-----	--

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

<u>AB</u>	<u>ALPHARMA US PHARMS</u>	<u>200MG</u>	<u>N73508</u>	<u>001</u>	Nov 19, 1993
	<u>MONISTAT 3</u>				
<u>AB</u>	<u>+ ORTHO MCNEIL PHARM</u>	<u>200MG</u>	<u>N18888</u>	<u>001</u>	Aug 15, 1984

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

<u>AP</u>	AM PHARM PARTNERS	<u>EQ 1MG BASE/ML</u>	<u>N75154</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75154</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>	APOTEX	<u>EQ 1MG BASE/ML</u>	<u>N75637</u>	<u>001</u>	Oct 31, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75637</u>	<u>002</u>	Oct 31, 2000
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 1MG BASE/ML</u>	<u>N75243</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75243</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 1MG BASE/ML</u>	<u>N75324</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75324</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>	BEDFORD	<u>EQ 1MG BASE/ML</u>	<u>N75247</u>	<u>002</u>	Jun 23, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75247</u>	<u>001</u>	Jun 23, 2000
<u>AP</u>	BEN VENUE	<u>EQ 1MG BASE/ML</u>	<u>N75421</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75421</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>N75293</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>N75409</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>N75856</u>	<u>001</u>	Jun 13, 2002
<u>AP</u>	+	<u>EQ 1MG BASE/ML</u>	<u>N75857</u>	<u>001</u>	Jul 22, 2002
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75293</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75409</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75856</u>	<u>002</u>	Jun 13, 2002
<u>AP</u>	+	<u>EQ 5MG BASE/ML</u>	<u>N75857</u>	<u>002</u>	Jul 22, 2002
<u>AP</u>	INTL MEDICATED	<u>EQ 1MG BASE/ML</u>	<u>N76144</u>	<u>001</u>	Jan 26, 2005
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N76144</u>	<u>002</u>	Jan 26, 2005
<u>AP</u>	INTL MEDICATION	<u>EQ 1MG BASE/ML</u>	<u>N76020</u>	<u>001</u>	Jul 16, 2004
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N76020</u>	<u>002</u>	Jul 16, 2004
<u>AP</u>	MAYNE PHARMA USA	<u>EQ 1MG BASE/ML</u>	<u>N75396</u>	<u>001</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75396</u>	<u>002</u>	Jun 20, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75484</u>	<u>001</u>	Jun 20, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 254 (of 360)

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

<u>AP</u>	TAYLOR	<u>EQ 5MG BASE/ML</u>	<u>N75481</u>	<u>001</u>	Jun 30, 2000
<u>AP</u>	TAYLOR PHARMA	<u>EQ 1MG BASE/ML</u>	<u>N75494</u>	<u>001</u>	Jun 30, 2000
<u>AP</u>		<u>EQ 5MG BASE/ML</u>	<u>N75494</u>	<u>002</u>	Jun 30, 2000

SYRUP; ORAL

MIDAZOLAM HYDROCHLORIDE

<u>AA</u>	APOTEX	<u>EQ 2MG BASE/ML</u>	<u>N77115</u>	<u>001</u>	Sep 09, 2005
<u>AA</u>	HI TECH PHARMA	<u>EQ 2MG BASE/ML</u>	<u>N75958</u>	<u>001</u>	Sep 04, 2003
<u>AA</u>	PADDOCK	<u>EQ 2MG BASE/ML</u>	<u>N76379</u>	<u>001</u>	May 02, 2005
<u>AA</u>	RANBAXY	<u>EQ 2MG BASE/ML</u>	<u>N76058</u>	<u>001</u>	Mar 15, 2002
<u>AA</u> +	ROXANE	<u>EQ 2MG BASE/ML</u>	<u>N75873</u>	<u>001</u>	Apr 30, 2002

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

<u>AB</u>	IMPAX PHARMS	<u>2.5MG</u>	<u>N76449</u>	<u>001</u>	May 27, 2004
<u>AB</u>		<u>5MG</u>	<u>N76449</u>	<u>002</u>	May 27, 2004
<u>AB</u>		<u>10MG</u>	<u>N76449</u>	<u>003</u>	Dec 16, 2005
<u>AB</u>	MYLAN	<u>2.5MG</u>	<u>N76577</u>	<u>001</u>	Sep 10, 2003
<u>AB</u>		<u>5MG</u>	<u>N76577</u>	<u>002</u>	Sep 10, 2003
<u>AB</u>		<u>10MG</u>	<u>N76577</u>	<u>003</u>	Sep 10, 2003
<u>AB</u>	SANDOZ	<u>2.5MG</u>	<u>N76514</u>	<u>001</u>	Sep 11, 2003
<u>AB</u>		<u>5MG</u>	<u>N76514</u>	<u>002</u>	Sep 11, 2003
<u>AB</u>		<u>10MG</u>	<u>N76514</u>	<u>003</u>	Jul 02, 2004
	<u>ORVATEN</u>				
<u>AB</u>	UPSHER SMITH	<u>2.5MG</u>	<u>N76725</u>	<u>001</u>	Nov 03, 2004
<u>AB</u>		<u>5MG</u>	<u>N76725</u>	<u>002</u>	Nov 03, 2004
<u>AB</u>		<u>10MG</u>	<u>N76725</u>	<u>003</u>	Nov 03, 2004
	<u>PROAMATINE</u>				
<u>AB</u>	SHIRE	<u>2.5MG</u>	<u>N19815</u>	<u>001</u>	Sep 06, 1996
<u>AB</u> +		<u>5MG</u>	<u>N19815</u>	<u>002</u>	Sep 06, 1996
<u>AB</u>		<u>10MG</u>	<u>N19815</u>	<u>003</u>	Mar 20, 2002

MIFEPRISTONE

TABLET; ORAL

MIFEPREX

+	DANCO LABS LLC	200MG	N20687	001	Sep 28, 2000
---	----------------	-------	--------	-----	--------------

MIGLITOL

TABLET; ORAL

GLYSET

	PHARMACIA AND UPJOHN	25MG	N20682	001	Dec 18, 1996
		50MG	N20682	002	Dec 18, 1996
+		100MG	N20682	003	Dec 18, 1996

MIGLUSTAT

CAPSULE; ORAL

ZAVESCA

+	ACTELION PHARMS	100MG	N21348	001	Jul 31, 2003
---	-----------------	-------	--------	-----	--------------

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

<u>AP</u>	AM PHARM PARTNERS	<u>EQ 1MG BASE/ML</u>	<u>N75936</u>	<u>001</u>	May 28, 2002
<u>AP</u>	APOTEX	<u>EQ 1MG BASE/ML</u>	<u>N76427</u>	<u>001</u>	Sep 21, 2004
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 1MG BASE/ML</u>	<u>N75530</u>	<u>001</u>	May 28, 2002
<u>AP</u>	BAXTER HLTHCARE CORP	<u>EQ 1MG BASE/ML</u>	<u>N75852</u>	<u>001</u>	May 28, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 255 (of 360)

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE

<u>AP</u>	BEDFORD	<u>EQ 1MG /ML</u>	<u>N75660</u>	<u>001</u>	May 28, 2002
<u>AP</u>	BIONICHE (CANADA)	<u>EQ 1MG BASE/ML</u>	<u>N76428</u>	<u>001</u>	Jun 16, 2003
<u>AP</u>	HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>N75884</u>	<u>001</u>	May 28, 2002
<u>AP</u>	INTL MEDICATED	<u>EQ 1MG BASE/ML</u>	<u>N76013</u>	<u>001</u>	Aug 02, 2002
<u>AP</u>	MAYNE PHARMA USA	<u>EQ 1MG BASE/ML</u>	<u>N75830</u>	<u>001</u>	May 28, 2002
<u>MILRINONE LACTATE IN 5% DEXTROSE IN PLASTIC CONTAINER</u>					
<u>AP</u>	APOTEX	<u>EQ 20MG BASE/100ML</u>	<u>N77151</u>	<u>001</u>	Jul 20, 2005
<u>MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>	B BRAUN	<u>EQ 20MG BASE/100ML</u>	<u>N76414</u>	<u>001</u>	Aug 18, 2004
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 20MG BASE/100ML</u>	<u>N75510</u>	<u>001</u>	May 28, 2002
<u>AP</u>		<u>EQ 20MG BASE/100ML</u>	<u>N75834</u>	<u>001</u>	May 28, 2002
<u>AP</u>	HOSPIRA	<u>EQ 20MG BASE/100ML</u>	<u>N75885</u>	<u>001</u>	May 28, 2002
<u>MILRINONE LACTOSE IN 5% DEXTROSE</u>					
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 20MG BASE/100ML</u>	<u>N76259</u>	<u>001</u>	Aug 08, 2002
<u>PRIMACOR</u>					
<u>AP</u>	+ SANOFI SYNTHELABO	<u>EQ 1MG BASE/ML</u>	<u>N19436</u>	<u>001</u>	Dec 31, 1987
<u>PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+ SANOFI SYNTHELABO	<u>EQ 20MG BASE/100ML</u>	<u>N20343</u>	<u>003</u>	Aug 09, 1994

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DYNACIN

<u>AB</u>	MEDICIS	<u>EQ 75MG BASE</u>	<u>N63067</u>	<u>002</u>	Sep 15, 1999
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N63067</u>	<u>001</u>	Jul 31, 1990

MINOCIN

<u>AB</u>	WYETH PHARMS INC	<u>EQ 50MG BASE</u>	<u>N50649</u>	<u>001</u>	May 31, 1990
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N50649</u>	<u>002</u>	May 31, 1990

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	IMPAX LABS	<u>EQ 50MG BASE</u>	<u>N65005</u>	<u>001</u>	Mar 23, 1999
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N65005</u>	<u>003</u>	Apr 18, 2001
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65005</u>	<u>002</u>	Mar 23, 1999
<u>AB</u>	RANBAXY	<u>EQ 50MG BASE</u>	<u>N65062</u>	<u>001</u>	Nov 30, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N65062</u>	<u>002</u>	Nov 30, 2000
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65062</u>	<u>003</u>	Nov 30, 2000
<u>AB</u>	TEVA	<u>EQ 50MG BASE</u>	<u>N63011</u>	<u>001</u>	Mar 02, 1992
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N63009</u>	<u>002</u>	Aug 12, 2003
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N63009</u>	<u>001</u>	Mar 02, 1992
<u>AB</u>	WATSON LABS	<u>EQ 50MG BASE</u>	<u>N63181</u>	<u>001</u>	Dec 30, 1991
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N63065</u>	<u>002</u>	Jun 10, 1999
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N63065</u>	<u>001</u>	Dec 30, 1991

POWDER, EXTENDED RELEASE; DENTAL

ARESTIN

	+ ORAPHARMA	<u>EQ 1MG BASE</u>	<u>N50781</u>	<u>001</u>	Feb 16, 2001
--	-------------	--------------------	---------------	------------	--------------

TABLET; ORAL

MINOCYCLINE HYDROCHLORIDE

<u>AB</u>	MEDICIS	<u>EQ 50MG BASE</u>	<u>N65131</u>	<u>001</u>	Apr 16, 2003
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N65131</u>	<u>002</u>	Apr 16, 2003
<u>AB</u>	+	<u>EQ 100MG BASE</u>	<u>N65131</u>	<u>003</u>	Apr 16, 2003
<u>AB</u>	RANBAXY	<u>EQ 50MG BASE</u>	<u>N65156</u>	<u>001</u>	Jan 06, 2004
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N65156</u>	<u>002</u>	Jan 06, 2004
<u>AB</u>		<u>EQ 100MG BASE</u>	<u>N65156</u>	<u>003</u>	Jan 06, 2004

MINOXIDIL

TABLET; ORAL

LONITEN

<u>AB</u>	PHARMACIA AND UPJOHN	<u>2.5MG</u>	<u>N18154</u>	<u>001</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N18154</u>	<u>003</u>	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 256 (of 360)

MINOXIDIL

TABLET; ORAL

MINOXIDIL

<u>AB</u>	MUTUAL PHARM	<u>2.5MG</u>	<u>N72708</u>	<u>001</u>	Dec 14, 1995
<u>AB</u>		<u>10MG</u>	<u>N72709</u>	<u>001</u>	Dec 14, 1995
<u>AB</u>	PAR PHARM	<u>2.5MG</u>	<u>N71826</u>	<u>001</u>	Nov 14, 1988
<u>AB</u>		<u>10MG</u>	<u>N71839</u>	<u>001</u>	Nov 14, 1988
<u>AB</u>	WATSON LABS	<u>2.5MG</u>	<u>N71344</u>	<u>001</u>	Mar 03, 1987
<u>AB</u>		<u>10MG</u>	<u>N71345</u>	<u>001</u>	Mar 03, 1987

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

<u>AB</u>	ALPHAPHARM	<u>15MG</u>	<u>N76176</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>N76176</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>N76176</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	AMIDE PHARM	<u>15MG</u>	<u>N76241</u>	<u>001</u>	Jun 25, 2003
<u>AB</u>		<u>30MG</u>	<u>N76241</u>	<u>002</u>	Jun 25, 2003
<u>AB</u>		<u>45MG</u>	<u>N76241</u>	<u>003</u>	Jun 25, 2003
<u>AB</u>	ANDRX PHARMS	<u>15MG</u>	<u>N76336</u>	<u>001</u>	Jun 20, 2003
<u>AB</u>		<u>30MG</u>	<u>N76336</u>	<u>002</u>	Jun 20, 2003
<u>AB</u>		<u>45MG</u>	<u>N76336</u>	<u>003</u>	Jun 20, 2003
<u>AB</u>	AUROBINDO	<u>7.5MG</u>	<u>N76921</u>	<u>001</u>	Oct 22, 2004
<u>AB</u>		<u>15MG</u>	<u>N76921</u>	<u>002</u>	Oct 22, 2004
<u>AB</u>		<u>30MG</u>	<u>N76921</u>	<u>003</u>	Oct 22, 2004
<u>AB</u>		<u>45MG</u>	<u>N76921</u>	<u>004</u>	Oct 22, 2004
<u>AB</u>	CARACO	<u>7.5MG</u>	<u>N76541</u>	<u>004</u>	Apr 22, 2004
<u>AB</u>		<u>15MG</u>	<u>N76541</u>	<u>001</u>	Apr 22, 2004
<u>AB</u>		<u>30MG</u>	<u>N76541</u>	<u>002</u>	Apr 22, 2004
<u>AB</u>		<u>45MG</u>	<u>N76541</u>	<u>003</u>	Apr 22, 2004
<u>AB</u>	IVAX PHARMS	<u>15MG</u>	<u>N76244</u>	<u>001</u>	Dec 22, 2003
<u>AB</u>		<u>30MG</u>	<u>N76244</u>	<u>002</u>	Dec 22, 2003
<u>AB</u>		<u>45MG</u>	<u>N76244</u>	<u>003</u>	Dec 22, 2003
<u>AB</u>	MYLAN	<u>15MG</u>	<u>N76122</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>N76122</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>N76122</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	PUREPAC PHARM	<u>15MG</u>	<u>N76308</u>	<u>001</u>	Jun 20, 2003
<u>AB</u>		<u>30MG</u>	<u>N76308</u>	<u>002</u>	Jun 20, 2003
<u>AB</u>		<u>45MG</u>	<u>N76308</u>	<u>003</u>	Jun 20, 2003
<u>AB</u>	ROXANE	<u>15MG</u>	<u>N76270</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>N76270</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>N76270</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	SANDOZ	<u>15MG</u>	<u>N76189</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>15MG</u>	<u>N76219</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>N76189</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>N76219</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>N76189</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>N76219</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	TEVA	<u>15MG</u>	<u>N76119</u>	<u>001</u>	Jan 24, 2003
<u>AB</u>		<u>30MG</u>	<u>N76119</u>	<u>002</u>	Jan 24, 2003
<u>AB</u>		<u>45MG</u>	<u>N76119</u>	<u>003</u>	Jun 19, 2003
<u>AB</u>	WATSON LABS	<u>15MG</u>	<u>N76312</u>	<u>001</u>	Jun 19, 2003
<u>AB</u>		<u>30MG</u>	<u>N76312</u>	<u>002</u>	Jun 19, 2003
<u>AB</u>		<u>45MG</u>	<u>N76312</u>	<u>003</u>	Jun 19, 2003
<u>REMERON</u>					
<u>AB</u>	+ ORGANON USA INC	<u>15MG</u>	<u>N20415</u>	<u>001</u>	Jun 14, 1996
<u>AB</u>		<u>30MG</u>	<u>N20415</u>	<u>002</u>	Jun 14, 1996
<u>AB</u>		<u>45MG</u>	<u>N20415</u>	<u>003</u>	Mar 17, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 257 (of 360)

MIRTAZAPINE

TABLET, ORALLY DISINTEGRATING; ORAL

MIRTAZAPINE

<u>AB</u>	AMIDE PHARM	<u>15MG</u>	<u>N76689</u>	<u>001</u>	Aug 31, 2005
<u>AB</u>		<u>30MG</u>	<u>N76689</u>	<u>002</u>	Aug 31, 2005
<u>AB</u>		<u>45MG</u>	<u>N76689</u>	<u>003</u>	Aug 31, 2005
<u>AB</u>	AUROBINDO PHARMA LTD	<u>15MG</u>	<u>N77376</u>	<u>002</u>	Dec 08, 2005
<u>AB</u>		<u>30MG</u>	<u>N77376</u>	<u>003</u>	Dec 08, 2005
<u>AB</u>	BARR	<u>15MG</u>	<u>N76307</u>	<u>001</u>	Dec 17, 2003
<u>AB</u>		<u>30MG</u>	<u>N76307</u>	<u>002</u>	Dec 17, 2003
<u>AB</u>	TEVA	<u>15MG</u>	<u>N76901</u>	<u>001</u>	Jun 28, 2005
<u>AB</u>		<u>30MG</u>	<u>N76901</u>	<u>002</u>	Jun 28, 2005
<u>AB</u>		<u>45MG</u>	<u>N76901</u>	<u>003</u>	Jun 28, 2005
<u>REMERON SOLTAB</u>					
<u>AB</u>	+ ORGANON USA INC	<u>15MG</u>	<u>N21208</u>	<u>001</u>	Jan 12, 2001
<u>AB</u>		<u>30MG</u>	<u>N21208</u>	<u>002</u>	Jan 12, 2001
<u>AB</u>		<u>45MG</u>	<u>N21208</u>	<u>003</u>	Jan 12, 2001

MISOPROSTOL

TABLET; ORAL

CYTOTEC

<u>AB</u>	GD SEARLE LLC	<u>0.1MG</u>	<u>N19268</u>	<u>003</u>	Sep 21, 1990
<u>AB</u>	+	<u>0.2MG</u>	<u>N19268</u>	<u>001</u>	Dec 27, 1988
<u>MISOPROSTOL</u>					
<u>AB</u>	IVAX PHARMS	<u>0.1MG</u>	<u>N76095</u>	<u>001</u>	Jul 10, 2002
<u>AB</u>		<u>0.2MG</u>	<u>N76095</u>	<u>002</u>	Jul 10, 2002

MITOMYCIN

INJECTABLE; INJECTION

MITOMYCIN

<u>AP</u>	BAXTER HLTHCARE	<u>5MG/VIAL</u>	<u>N64180</u>	<u>001</u>	Dec 23, 1999
<u>AP</u>		<u>20MG/VIAL</u>	<u>N64180</u>	<u>002</u>	Dec 23, 1999
<u>AP</u>	BEDFORD	<u>5MG/VIAL</u>	<u>N64117</u>	<u>001</u>	Apr 19, 1995
<u>AP</u>		<u>20MG/VIAL</u>	<u>N64117</u>	<u>002</u>	Apr 19, 1995
<u>AP</u>	SUPERGEN	<u>5MG/VIAL</u>	<u>N64144</u>	<u>001</u>	Apr 30, 1998
<u>AP</u>		<u>20MG/VIAL</u>	<u>N64144</u>	<u>002</u>	Apr 30, 1998
<u>MUTAMYCIN</u>					
<u>AP</u>	+ BRISTOL MYERS	<u>5MG/VIAL</u>	<u>N62336</u>	<u>001</u>	
<u>AP</u>	+	<u>20MG/VIAL</u>	<u>N62336</u>	<u>002</u>	
	+	40MG/VIAL	N62336	003	Mar 10, 1988
	MYTOZYTREX				
	+ SUPERGEN	5MG/VIAL	N50763	001	Nov 14, 2002

MITOTANE

TABLET; ORAL

LYSODREN

+	BRISTOL	500MG	N16885	001	
---	---------	-------	--------	-----	--

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

NOVANTRONE

+	SERONO INC	EQ 20MG BASE/10 MG (2MG/ML)	N19297	001	Dec 23, 1987
+		EQ 25MG BASE/12.5ML (2MG/ML)	N19297	002	Dec 23, 1987
+		EQ 30MG BASE/15ML (2MG/ML)	N19297	003	Dec 23, 1987

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACRON

+	ABBOTT	EQ 2MG BASE/ML	N20098	001	Jan 22, 1992
---	--------	----------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 258 (of 360)

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER

+	ABBOTT	EQ 50MG BASE/100ML	N20098	003	Jan 22, 1992
---	--------	--------------------	--------	-----	--------------

MODAFINIL

TABLET; ORAL

PROVIGIL

CEPHALON 100MG

N20717 001 Dec 24, 1998

+		200MG	N20717	002	Dec 24, 1998
---	--	-------	--------	-----	--------------

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDEAB TEVA 7.5MGN76204 001 May 08, 2003AB 15MGN76204 002 May 08, 2003UNIVASCAB SCHWARZ PHARMA 7.5MGN20312 001 Apr 19, 1995AB + 15MGN20312 002 Apr 19, 1995MOLINDONE HYDROCHLORIDE

TABLET; ORAL

MOBAN

ENDO PHARMS 5MG

N17111 004

10MG

N17111 005

+		25MG	N17111	006	
---	--	------	--------	-----	--

50MG

N17111 007

MOMETASONE FUROATE

CREAM; TOPICAL

ELOCONAB + SCHERING 0.1%N19625 001 May 06, 1987MOMETASONE FUROATEAB ALTANA 0.1%N76171 001 Apr 08, 2005AB TARO 0.1%N76679 001 Dec 21, 2004

LOTION; TOPICAL

ELOCONAB + SCHERING 0.1%N19796 001 Mar 30, 1989MOMETASONE FUROATEAB AGIS INDS 0.1%N77180 001 Apr 06, 2005

OINTMENT; TOPICAL

ELOCONAB + SCHERING 0.1%N19543 001 Apr 30, 1987MOMETASONE FUROATEAB ALTANA 0.1%N77061 001 Mar 28, 2005AB CLAY PARK 0.1%N76067 001 Mar 18, 2002AB QLT USA 0.1%N76481 001 Nov 14, 2003AB TARO 0.1%N76624 001 Dec 03, 2004

POWDER; INHALATION

ASMANEX TWISTHALER

+	SCHERING	0.22MG/INH	N21067	001	Mar 30, 2005
---	----------	------------	--------	-----	--------------

MOMETASONE FUROATE MONOHYDRATE

SPRAY, METERED; NASAL

NASONEX

+	SCHERING PLOUGH	EQ 0.05MG BASE/SPRAY	N20762	001	Oct 01, 1997
---	-----------------	----------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 259 (of 360)

MONOBENZONECREAM; TOPICAL
BENOQUIN

+ VALEANT PHARM INTL 20%

N08173 003

MONTELUKAST SODIUMGRANULE; ORAL
SINGLAIR

+ MERCK EQ 4MG BASE/PACKET

N21409 001 Jul 26, 2002

TABLET; ORAL
SINGLAIR

+ MERCK EQ 10MG BASE

N20829 002 Feb 20, 1998

TABLET, CHEWABLE; ORAL
SINGLAIR

MERCK EQ 4MG BASE

N20830 002 Mar 03, 2000

+ EQ 5MG BASE

N20830 001 Feb 20, 1998

MORICIZINE HYDROCHLORIDETABLET; ORAL
ETHMOZINE
SHIRE

200MG

N19753 001 Jun 19, 1990

250MG

N19753 002 Jun 19, 1990

+ 300MG

N19753 003 Jun 19, 1990

MORPHINE SULFATECAPSULE, EXTENDED RELEASE; ORAL
AVINZA

BX LIGAND 30MG

N21260 001 Mar 20, 2002

BX 60MG

N21260 002 Mar 20, 2002

90MG

N21260 003 Mar 20, 2002

+ 120MG

N21260 004 Mar 20, 2002

KADIAN

BX ALPHARMA US PHARMS 30MG

N20616 004 Mar 09, 2001

BX 60MG

N20616 005 Mar 09, 2001

20MG

N20616 001 Jul 03, 1996

50MG

N20616 002 Jul 03, 1996

+ 100MG

N20616 003 Jul 03, 1996

INJECTABLE; INJECTION

ASTRAMORPH PFAP ASTRAZENECA 0.5MG/MLN71050 001 Oct 07, 1986AP 0.5MG/MLN71051 001 Oct 07, 1986AP 1MG/MLN71052 001 Oct 07, 1986AP 1MG/MLN71053 001 Oct 07, 1986DURAMORPH PFAP + BAXTER HLTHCARE 0.5MG/MLN18565 001 Sep 18, 1984AP + 1MG/MLN18565 002 Sep 18, 1984

INFUMORPH

+ BAXTER HLTHCARE 10MG/ML

N18565 003 Jul 19, 1991

+ 25MG/ML

N18565 004 Jul 19, 1991

MORPHINE SULFATEAP + HOSPIRA 0.5MG/MLN19917 001 Oct 30, 1992AP 0.5MG/MLN71849 001 May 11, 1988AP 0.5MG/MLN73509 001 Sep 30, 1992AP + 1MG/MLN19916 001 Oct 30, 1992AP 1MG/MLN71850 001 May 11, 1988AP 1MG/MLN73510 001 Sep 30, 1992AP MALLINCKRODT 1MG/MLN20631 001 Jul 03, 1996

+ 2MG/ML

N20631 002 Jul 03, 1996

+ MERIDIAN MEDCL TECHN 15MG/ML

N19999 001 Jul 12, 1990

PRESCRIPTION DRUG PRODUCT LIST

3 - 260 (of 360)

MORPHINE SULFATE

INJECTABLE; INJECTION

MORPHINE SULFATE

<u>AP</u>	STERIS	<u>0.5MG/ML</u>	<u>N73373</u>	<u>001</u>	Sep 30, 1991
<u>AP</u>		<u>0.5MG/ML</u>	<u>N73375</u>	<u>001</u>	Sep 30, 1991
<u>AP</u>		<u>1MG/ML</u>	<u>N73374</u>	<u>001</u>	Sep 30, 1991
<u>AP</u>		<u>1MG/ML</u>	<u>N73376</u>	<u>001</u>	Sep 30, 1991

INJECTABLE, LIPOSOMAL; EPIDURAL

DEPODUR

	SKYEPHARMA	10MG/ML	N21671	001	May 18, 2004
		15MG/1.5ML (10MG/ML)	N21671	002	May 18, 2004
+		20MG/2ML (10MG/ML)	N21671	003	May 18, 2004

TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

<u>AB</u>	AB GENERICS	<u>15MG</u>	<u>N74862</u>	<u>001</u>	Jul 07, 1998
<u>AB</u>		<u>30MG</u>	<u>N74862</u>	<u>002</u>	Jul 07, 1998
<u>AB</u>		<u>60MG</u>	<u>N74862</u>	<u>003</u>	Jul 07, 1998
<u>AB</u>		<u>100MG</u>	<u>N74769</u>	<u>001</u>	Jul 02, 1998
<u>AB</u>		<u>200MG</u>	<u>N74769</u>	<u>002</u>	Jul 02, 1998
<u>AB</u>	ENDO PHARMS	<u>15MG</u>	<u>N75295</u>	<u>001</u>	Oct 28, 1998
<u>AB</u>		<u>30MG</u>	<u>N75295</u>	<u>002</u>	Oct 28, 1998
<u>AB</u>		<u>60MG</u>	<u>N75295</u>	<u>003</u>	Oct 28, 1998
<u>AB</u>		<u>100MG</u>	<u>N75295</u>	<u>004</u>	Sep 15, 2000
<u>AB</u>		<u>200MG</u>	<u>N75295</u>	<u>005</u>	Sep 15, 2000
<u>AB</u>	KV PHARM	<u>15MG</u>	<u>N76733</u>	<u>001</u>	May 19, 2004
<u>AB</u>		<u>30MG</u>	<u>N76720</u>	<u>002</u>	Dec 23, 2005
<u>AB</u>		<u>60MG</u>	<u>N76720</u>	<u>001</u>	May 19, 2004
<u>AB</u>	MALLINCKRODT	<u>15MG</u>	<u>N76412</u>	<u>001</u>	Jul 31, 2003
<u>AB</u>		<u>30MG</u>	<u>N76412</u>	<u>002</u>	Jul 31, 2003
<u>AB</u>		<u>60MG</u>	<u>N76412</u>	<u>003</u>	Jul 31, 2003
<u>AB</u>		<u>100MG</u>	<u>N76438</u>	<u>001</u>	Jul 03, 2003
<u>AB</u>		<u>200MG</u>	<u>N76438</u>	<u>002</u>	Jul 03, 2003
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N75656</u>	<u>001</u>	Jan 30, 2001
	<u>MS CONTIN</u>				
<u>AB</u>	PURDUE FREDERICK	<u>15MG</u>	<u>N19516</u>	<u>003</u>	Sep 12, 1989
<u>AB</u>		<u>30MG</u>	<u>N19516</u>	<u>001</u>	May 29, 1987
<u>AB</u>	+	<u>60MG</u>	<u>N19516</u>	<u>002</u>	Apr 08, 1988
<u>AB</u>		<u>100MG</u>	<u>N19516</u>	<u>004</u>	Jan 16, 1990
<u>AB</u>		<u>200MG</u>	<u>N19516</u>	<u>005</u>	Nov 08, 1993
	ORAMORPH SR				
BC	XANODYNE PHARM	15MG	N19977	004	Nov 23, 1994
BC		30MG	N19977	001	Aug 15, 1991
BC	+	60MG	N19977	002	Aug 15, 1991
BC		100MG	N19977	003	Aug 15, 1991

MOXIFLOXACIN HYDROCHLORIDE

INJECTABLE; IV (INFUSION)

AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER

+	BAYER PHARMS	160MG/100ML	N21277	001	Nov 30, 2001
---	--------------	-------------	--------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

VIGAMOX

+	ALCON	0.5%	N21598	001	Apr 15, 2003
---	-------	------	--------	-----	--------------

TABLET; ORAL

AVELOX

+	BAYER PHARMS	EQ 400MG BASE	N21085	001	Dec 10, 1999
---	--------------	---------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 261 (of 360)

MUPIROCIN

OINTMENT; TOPICAL

BACTROBAN

<u>AB</u>	+	GLAXOSMITHKLINE	<u>2%</u>	<u>N50591</u>	<u>001</u>	Dec 31, 1987
		CENTANY				

BX		JOHNSON AND JOHNSON	2%	N50788	001	Dec 04, 2002
----	--	---------------------	----	--------	-----	--------------

MUPIROCIN

<u>AB</u>		ALTANA	<u>2%</u>	<u>N65192</u>	<u>001</u>	Nov 30, 2005
-----------	--	--------	-----------	---------------	------------	--------------

<u>AB</u>		CLAY PARK	<u>2%</u>	<u>N65123</u>	<u>001</u>	Nov 07, 2003
-----------	--	-----------	-----------	---------------	------------	--------------

<u>AB</u>		TARO	<u>2%</u>	<u>N65170</u>	<u>001</u>	Sep 23, 2005
-----------	--	------	-----------	---------------	------------	--------------

<u>AB</u>		TEVA	<u>2%</u>	<u>N65085</u>	<u>001</u>	Nov 07, 2003
-----------	--	------	-----------	---------------	------------	--------------

MUPIROCIN CALCIUM

CREAM, AUGMENTED; TOPICAL

BACTROBAN

	+	GLAXOSMITHKLINE	2%	N50746	001	Dec 11, 1997
--	---	-----------------	----	--------	-----	--------------

OINTMENT; NASAL

BACTROBAN

	+	GLAXOSMITHKLINE	EQ 2% BASE	N50703	001	Sep 18, 1995
--	---	-----------------	------------	--------	-----	--------------

MYCOPHENOLATE MOFETIL

CAPSULE; ORAL

CELLCEPT

	+	ROCHE PALO	250MG	N50722	001	May 03, 1995
--	---	------------	-------	--------	-----	--------------

SUSPENSION; ORAL

CELLCEPT

	+	ROCHE PALO	200MG/ML	N50759	001	Oct 01, 1998
--	---	------------	----------	--------	-----	--------------

TABLET; ORAL

CELLCEPT

	+	ROCHE PALO	500MG	N50723	001	Jun 19, 1997
--	---	------------	-------	--------	-----	--------------

MYCOPHENOLATE MOFETIL HYDROCHLORIDE

INJECTABLE; INJECTION

CELLCEPT

	+	ROCHE PALO	500MG/VIAL	N50758	001	Aug 12, 1998
--	---	------------	------------	--------	-----	--------------

MYCOPHENOLIC ACID

TABLET, DELAYED RELEASE; ORAL

MYFORTIC

		NOVARTIS	180MG	N50791	001	Feb 27, 2004
--	--	----------	-------	--------	-----	--------------

	+		360MG	N50791	002	Feb 27, 2004
--	---	--	-------	--------	-----	--------------

NABUMETONE

TABLET; ORAL

NABUMETONE

<u>AB</u>		COPLEY PHARM	<u>750MG</u>	<u>N75179</u>	<u>001</u>	Jun 06, 2000
-----------	--	--------------	--------------	---------------	------------	--------------

<u>AB</u>		IVAX PHARMS	<u>500MG</u>	<u>N76009</u>	<u>001</u>	Jan 24, 2003
-----------	--	-------------	--------------	---------------	------------	--------------

<u>AB</u>			<u>750MG</u>	<u>N76009</u>	<u>002</u>	Jan 24, 2003
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>500MG</u>	<u>N75280</u>	<u>001</u>	Feb 25, 2002
-----------	--	--------	--------------	---------------	------------	--------------

<u>AB</u>			<u>500MG</u>	<u>N75590</u>	<u>001</u>	Feb 25, 2002
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>			<u>750MG</u>	<u>N75280</u>	<u>002</u>	Feb 25, 2002
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>			<u>750MG</u>	<u>N75590</u>	<u>002</u>	Feb 25, 2002
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		TEVA	<u>500MG</u>	<u>N75189</u>	<u>001</u>	May 26, 2000
-----------	--	------	--------------	---------------	------------	--------------

<u>AB</u>			<u>750MG</u>	<u>N75189</u>	<u>002</u>	Sep 24, 2001
-----------	--	--	--------------	---------------	------------	--------------

RELAFEN

<u>AB</u>		GLAXOSMITHKLINE	<u>500MG</u>	<u>N19583</u>	<u>001</u>	Dec 24, 1991
-----------	--	-----------------	--------------	---------------	------------	--------------

<u>AB</u>	+		<u>750MG</u>	<u>N19583</u>	<u>002</u>	Dec 24, 1991
-----------	---	--	--------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 262 (of 360)

NADOLOL

TABLET; ORAL

CORGARD

<u>AB</u>	APOTHECON	<u>20MG</u>	<u>N18063</u>	<u>005</u>	Oct 28, 1986
<u>AB</u>		<u>40MG</u>	<u>N18063</u>	<u>001</u>	
<u>AB</u>		<u>80MG</u>	<u>N18063</u>	<u>002</u>	
<u>AB</u>		<u>120MG</u>	<u>N18063</u>	<u>003</u>	
<u>AB</u>	+	<u>160MG</u>	<u>N18063</u>	<u>004</u>	

NADOLOL

<u>AB</u>	IVAX PHARMS	<u>20MG</u>	<u>N74229</u>	<u>001</u>	Aug 30, 1996
<u>AB</u>		<u>40MG</u>	<u>N74229</u>	<u>002</u>	Aug 30, 1996
<u>AB</u>		<u>80MG</u>	<u>N74255</u>	<u>001</u>	Jan 24, 1996
<u>AB</u>		<u>120MG</u>	<u>N74255</u>	<u>002</u>	Jan 24, 1996
<u>AB</u>		<u>160MG</u>	<u>N74255</u>	<u>003</u>	Jan 24, 1996
<u>AB</u>	MYLAN	<u>20MG</u>	<u>N74172</u>	<u>001</u>	Oct 31, 1993
<u>AB</u>		<u>40MG</u>	<u>N74172</u>	<u>002</u>	Oct 31, 1993
<u>AB</u>		<u>80MG</u>	<u>N74172</u>	<u>003</u>	Oct 31, 1993
<u>AB</u>	SANDOZ	<u>20MG</u>	<u>N74501</u>	<u>001</u>	Nov 09, 1995
<u>AB</u>		<u>40MG</u>	<u>N74501</u>	<u>002</u>	Nov 09, 1995
<u>AB</u>		<u>80MG</u>	<u>N74501</u>	<u>003</u>	Nov 09, 1995
<u>AB</u>	TEVA PHARMS	<u>80MG</u>	<u>N74368</u>	<u>001</u>	Aug 31, 1994
<u>AB</u>		<u>120MG</u>	<u>N74368</u>	<u>002</u>	Aug 31, 1994
<u>AB</u>		<u>160MG</u>	<u>N74368</u>	<u>003</u>	Aug 31, 1994

NAFARELIN ACETATE

SPRAY, METERED; NASAL

SYNAREL

+	GD SEARLE LLC	EQ 0.2MG BASE/SPRAY	N19886	001	Feb 13, 1990
---	---------------	---------------------	--------	-----	--------------

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

<u>AP</u>	SANDOZ	<u>EQ 1GM BASE/VIAL</u>	<u>N62527</u>	<u>002</u>	Aug 02, 1984
<u>AP</u>		<u>EQ 1GM BASE/VIAL</u>	<u>N62732</u>	<u>001</u>	Dec 23, 1986
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>N62527</u>	<u>003</u>	Aug 02, 1984
<u>AP</u>		<u>EQ 2GM BASE/VIAL</u>	<u>N62732</u>	<u>002</u>	Dec 23, 1986
<u>AP</u>		<u>EQ 10GM BASE/VIAL</u>	<u>N62527</u>	<u>004</u>	Aug 02, 1984
+		EQ 500MG BASE/VIAL	N62527	001	Aug 02, 1984
	NALLPEN IN PLASTIC CONTAINER				
+	BAXTER HLTHCARE	EQ 20MG BASE/ML	N50655	001	Oct 31, 1989
+		EQ 2GM BASE/100ML	N50655	002	Oct 31, 1989

NAFTIFINE HYDROCHLORIDE

CREAM; TOPICAL

NAFTIN

	MERZ PHARMS	1%	N19599	001	Feb 29, 1988
--	-------------	----	--------	-----	--------------

GEL; TOPICAL

NAFTIN

+	MERZ PHARMS	1%	N19356	001	Jun 18, 1990
---	-------------	----	--------	-----	--------------

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>10MG/ML</u>	<u>N70914</u>	<u>001</u>	Feb 03, 1989
<u>AP</u>		<u>10MG/ML</u>	<u>N70915</u>	<u>001</u>	Feb 03, 1989
<u>AP</u>		<u>20MG/ML</u>	<u>N70916</u>	<u>001</u>	Feb 03, 1989
<u>AP</u>		<u>20MG/ML</u>	<u>N70918</u>	<u>001</u>	Feb 03, 1989
<u>AP</u>	MAYNE PHARMA USA	<u>10MG/ML</u>	<u>N74471</u>	<u>001</u>	Mar 19, 1998
<u>AP</u>		<u>20MG/ML</u>	<u>N74471</u>	<u>002</u>	Mar 19, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 263 (of 360)

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NUBAIN

<u>AP</u>	+ ENDO PHARMS	<u>10MG/ML</u>	<u>N18024</u>	<u>001</u>	
<u>AP</u>	+	<u>20MG/ML</u>	<u>N18024</u>	<u>002</u>	May 27, 1982

NALIDIXIC ACID

TABLET; ORAL

NALIDIXIC ACID

<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>N71936</u>	<u>001</u>	Jun 29, 1988
<u>AB</u>		<u>500MG</u>	<u>N72061</u>	<u>001</u>	Jun 29, 1988
<u>AB</u>		<u>1GM</u>	<u>N71919</u>	<u>001</u>	Jun 29, 1988

NEGGRAM

<u>AB</u>	SANOFI SYNTHELABO	<u>250MG</u>	<u>N14214</u>	<u>002</u>	
<u>AB</u>		<u>500MG</u>	<u>N14214</u>	<u>004</u>	
<u>AB</u>	+	<u>1GM</u>	<u>N14214</u>	<u>005</u>	

NALMEFENE HYDROCHLORIDE

INJECTABLE; INJECTION

REVEX

	+ BAXTER HLTHCARE CORP	EQ 0.1MG BASE/ML	N20459	001	Apr 17, 1995
	+	EQ 1MG BASE/ML	N20459	002	Apr 17, 1995

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>0.02MG/ML</u>	<u>N70171</u>	<u>001</u>	Sep 24, 1986
<u>AP</u>		<u>0.02MG/ML</u>	<u>N70252</u>	<u>001</u>	Jan 16, 1987
<u>AP</u>		<u>0.02MG/ML</u>	<u>N70253</u>	<u>001</u>	Jan 16, 1987
<u>AP</u>		<u>0.4MG/ML</u>	<u>N70172</u>	<u>001</u>	Sep 24, 1986
<u>AP</u>		<u>0.4MG/ML</u>	<u>N70254</u>	<u>001</u>	Jan 07, 1987
<u>AP</u>		<u>0.4MG/ML</u>	<u>N70255</u>	<u>001</u>	Jan 07, 1987
<u>AP</u>		<u>0.4MG/ML</u>	<u>N70256</u>	<u>001</u>	Jan 07, 1987
<u>AP</u>		<u>0.4MG/ML</u>	<u>N70257</u>	<u>001</u>	Jan 07, 1987
<u>AP</u>	INTL MEDICATION	<u>0.4MG/ML</u>	<u>N70639</u>	<u>001</u>	Sep 24, 1986
<u>AP</u>		<u>1MG/ML</u>	<u>N72076</u>	<u>001</u>	Mar 24, 1988

NARCAN

<u>AP</u>	+ ENDO PHARMS	<u>0.02MG/ML</u>	<u>N16636</u>	<u>002</u>	
<u>AP</u>	+	<u>0.4MG/ML</u>	<u>N16636</u>	<u>001</u>	
<u>AP</u>	+	<u>1MG/ML</u>	<u>N16636</u>	<u>003</u>	Jun 14, 1982

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

NALOXONE HYDROCHLORIDE AND PENTAZOCAINE

<u>AB</u>	AMIDE PHARM	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>N75735</u>	<u>001</u>	Jul 11, 2001
-----------	-------------	-----------------------------------	---------------	------------	--------------

PENTAZOCINE AND NALOXONE HYDROCHLORIDES

<u>AB</u>	RANBAXY	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>N75523</u>	<u>001</u>	Mar 17, 2000
<u>AB</u>	WATSON LABS	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>N74736</u>	<u>001</u>	Jan 21, 1997

TALWIN NX

<u>AB</u>	+ SANOFI SYNTHELABO	<u>EQ 0.5MG BASE;EQ 50MG BASE</u>	<u>N18733</u>	<u>001</u>	Dec 16, 1982
-----------	---------------------	-----------------------------------	---------------	------------	--------------

NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

<u>AB</u>	AMIDE PHARM	<u>50MG</u>	<u>N75274</u>	<u>001</u>	May 26, 1999
<u>AB</u>	BARR	<u>50MG</u>	<u>N74918</u>	<u>001</u>	May 08, 1998
<u>AB</u>	MALLINCKRODT	<u>50MG</u>	<u>N76264</u>	<u>002</u>	Mar 22, 2002
		25MG	N76264	001	Mar 22, 2002
	+	100MG	N76264	003	Mar 22, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 264 (of 360)

NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N75434</u>	<u>001</u>	Mar 08, 2000
<u>AB</u>	<u>REVIA</u>				
<u>AB</u>	+ DURAMED	<u>50MG</u>	<u>N18932</u>	<u>001</u>	Nov 20, 1984

NANDROLONE DECANOATE

INJECTABLE; INJECTION

NANDROLONE DECANOATE

+	STERIS	50MG/ML	N86385	001	Jan 13, 1984
+		100MG/ML	N86598	001	Jan 13, 1984
+		200MG/ML	N88128	001	Dec 05, 1983

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

ALBALON

<u>AT</u>	+ ALLERGAN	<u>0.1%</u>	<u>N80248</u>	<u>001</u>	
<u>AT</u>	<u>NAFAZAIR</u>				
<u>AT</u>	BAUSCH AND LOMB	<u>0.1%</u>	<u>N40073</u>	<u>001</u>	May 25, 1994
<u>AT</u>	<u>NAPHAZOLINE HYDROCHLORIDE</u>				
<u>AT</u>	TAYLOR	<u>0.1%</u>	<u>N83590</u>	<u>001</u>	
<u>AT</u>	<u>VASOCON</u>				
<u>AT</u>	NOVARTIS	<u>0.1%</u>	<u>N80235</u>	<u>002</u>	Mar 24, 1983

NAPROXEN

SUSPENSION; ORAL

NAPROSYN

<u>AB</u>	+ ROCHE PALO	<u>25MG/ML</u>	<u>N18965</u>	<u>001</u>	Mar 23, 1987
<u>AB</u>	<u>NAPROXEN</u>				
<u>AB</u>	ROXANE	<u>25MG/ML</u>	<u>N74190</u>	<u>001</u>	Mar 30, 1994

TABLET; ORAL

NAPROSYN

<u>AB</u>	ROCHE PALO	<u>250MG</u>	<u>N17581</u>	<u>002</u>	
<u>AB</u>		<u>375MG</u>	<u>N17581</u>	<u>003</u>	
<u>AB</u>	+	<u>500MG</u>	<u>N17581</u>	<u>004</u>	Apr 15, 1982
<u>AB</u>	<u>NAPROXEN</u>				
<u>AB</u>	CLONMEL HLTHCARE	<u>250MG</u>	<u>N74410</u>	<u>001</u>	Apr 28, 1995
<u>AB</u>		<u>375MG</u>	<u>N74410</u>	<u>002</u>	Apr 28, 1995
<u>AB</u>		<u>500MG</u>	<u>N74410</u>	<u>003</u>	Apr 28, 1995
<u>AB</u>	INTERPHARM	<u>250MG</u>	<u>N75927</u>	<u>001</u>	Dec 18, 2001
<u>AB</u>		<u>375MG</u>	<u>N75927</u>	<u>002</u>	Dec 18, 2001
<u>AB</u>		<u>500MG</u>	<u>N75927</u>	<u>003</u>	Dec 18, 2001
<u>AB</u>	MYLAN	<u>250MG</u>	<u>N74121</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>N74121</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>N74121</u>	<u>003</u>	Dec 21, 1993
<u>AB</u>	PERRIGO R AND D	<u>250MG</u>	<u>N77339</u>	<u>001</u>	Apr 27, 2005
<u>AB</u>		<u>375MG</u>	<u>N77339</u>	<u>002</u>	Apr 27, 2005
<u>AB</u>		<u>500MG</u>	<u>N77339</u>	<u>003</u>	Apr 27, 2005
<u>AB</u>	PLIVA	<u>250MG</u>	<u>N74182</u>	<u>001</u>	Jun 27, 1996
<u>AB</u>		<u>375MG</u>	<u>N74182</u>	<u>002</u>	Jun 27, 1996
<u>AB</u>		<u>500MG</u>	<u>N74182</u>	<u>003</u>	Jun 27, 1996
<u>AB</u>	SANDOZ	<u>250MG</u>	<u>N74140</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>N74140</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>N74140</u>	<u>003</u>	Dec 21, 1993
<u>AB</u>	TEVA	<u>250MG</u>	<u>N74129</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>250MG</u>	<u>N74201</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>250MG</u>	<u>N74216</u>	<u>001</u>	Apr 11, 1996
<u>AB</u>		<u>375MG</u>	<u>N74129</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>N74201</u>	<u>002</u>	Dec 21, 1993

PRESCRIPTION DRUG PRODUCT LIST

3 - 265 (of 360)

NAPROXEN

TABLET; ORAL

NAPROXEN

<u>AB</u>	TEVA	<u>375MG</u>	<u>N74216</u>	<u>002</u>	Apr 11, 1996
<u>AB</u>		<u>500MG</u>	<u>N74129</u>	<u>003</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>N74201</u>	<u>003</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>N74216</u>	<u>003</u>	Apr 11, 1996
<u>AB</u>	TEVA PHARMS	<u>250MG</u>	<u>N74207</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>375MG</u>	<u>N74207</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>500MG</u>	<u>N74207</u>	<u>003</u>	Dec 21, 1993
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>N74163</u>	<u>001</u>	Feb 10, 1995
<u>AB</u>		<u>250MG</u>	<u>N74457</u>	<u>001</u>	May 31, 1995
<u>AB</u>		<u>375MG</u>	<u>N74163</u>	<u>002</u>	Feb 10, 1995
<u>AB</u>		<u>375MG</u>	<u>N74457</u>	<u>002</u>	May 31, 1995
<u>AB</u>		<u>500MG</u>	<u>N74163</u>	<u>003</u>	Feb 10, 1995
<u>AB</u>		<u>500MG</u>	<u>N74457</u>	<u>003</u>	May 31, 1995
<u>AB</u>	WESTWARD	<u>250MG</u>	<u>N76494</u>	<u>001</u>	Jan 14, 2004
<u>AB</u>		<u>375MG</u>	<u>N76494</u>	<u>002</u>	Jan 14, 2004
<u>AB</u>		<u>500MG</u>	<u>N76494</u>	<u>003</u>	Jan 14, 2004

TABLET, DELAYED RELEASE; ORAL

EC-NAPROSYN

<u>AB</u>	+ ROCHE PALO	<u>375MG</u>	<u>N20067</u>	<u>002</u>	Oct 14, 1994
<u>AB</u>	+	<u>500MG</u>	<u>N20067</u>	<u>003</u>	Oct 14, 1994

NAPROXEN

<u>AB</u>	ALPHAPHARM	<u>375MG</u>	<u>N75390</u>	<u>001</u>	Apr 19, 2001
<u>AB</u>		<u>500MG</u>	<u>N75390</u>	<u>002</u>	Apr 19, 2001
<u>AB</u>	PLIVA	<u>375MG</u>	<u>N75337</u>	<u>001</u>	May 26, 1999
<u>AB</u>		<u>500MG</u>	<u>N75337</u>	<u>002</u>	May 26, 1999
<u>AB</u>	PUREPAC PHARM	<u>375MG</u>	<u>N74936</u>	<u>001</u>	Feb 24, 1998
<u>AB</u>		<u>500MG</u>	<u>N74936</u>	<u>002</u>	Feb 24, 1998
<u>AB</u>	SANDOZ	<u>375MG</u>	<u>N75061</u>	<u>001</u>	Feb 18, 1998
<u>AB</u>		<u>500MG</u>	<u>N75061</u>	<u>002</u>	Feb 18, 1998
<u>AB</u>	TEVA	<u>375MG</u>	<u>N75227</u>	<u>001</u>	Jun 30, 1998
<u>AB</u>		<u>500MG</u>	<u>N75227</u>	<u>002</u>	Jun 30, 1998

NAPROXEN SODIUM

TABLET; ORAL

ANAPROX

<u>AB</u>	ROCHE PALO	<u>EQ 250MG BASE</u>	<u>N18164</u>	<u>001</u>	
-----------	------------	----------------------	---------------	------------	--

ANAPROX DS

<u>AB</u>	+ ROCHE PALO	<u>EQ 500MG BASE</u>	<u>N18164</u>	<u>003</u>	Sep 30, 1987
-----------	--------------	----------------------	---------------	------------	--------------

NAPROXEN SODIUM

<u>AB</u>	HIKMA	<u>EQ 250MG BASE</u>	<u>N74480</u>	<u>002</u>	Feb 18, 1998
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74480</u>	<u>001</u>	May 14, 1996
<u>AB</u>	MYLAN	<u>EQ 250MG BASE</u>	<u>N74367</u>	<u>001</u>	Aug 31, 1994
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74367</u>	<u>002</u>	Aug 31, 1994
<u>AB</u>	PLIVA	<u>EQ 250MG BASE</u>	<u>N74242</u>	<u>001</u>	Jun 20, 1996
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74242</u>	<u>002</u>	Jun 20, 1996
<u>AB</u>	SANDOZ	<u>EQ 250MG BASE</u>	<u>N74162</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N74495</u>	<u>001</u>	Dec 05, 1994
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74162</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74495</u>	<u>002</u>	Dec 05, 1994
<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>N74142</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N74198</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74142</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74198</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>	TEVA PHARMS	<u>EQ 250MG BASE</u>	<u>N74289</u>	<u>001</u>	Jan 27, 1994
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74289</u>	<u>002</u>	Jan 27, 1994
<u>AB</u>	WATSON LABS	<u>EQ 250MG BASE</u>	<u>N74195</u>	<u>001</u>	Dec 21, 1993
<u>AB</u>		<u>EQ 250MG BASE</u>	<u>N74455</u>	<u>001</u>	May 31, 1995

PRESCRIPTION DRUG PRODUCT LIST

3 - 266 (of 360)

NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

<u>AB</u>	WATSON LABS	<u>EQ 500MG BASE</u>	<u>N74195</u>	<u>002</u>	Dec 21, 1993
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N74455</u>	<u>002</u>	May 31, 1995

TABLET, EXTENDED RELEASE; ORAL

NAPRELAN

<u>AB</u>	+ STAT TRADE	<u>EQ 375MG BASE</u>	<u>N20353</u>	<u>001</u>	Jan 05, 1996
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>N20353</u>	<u>002</u>	Jan 05, 1996

NAPROXEN SODIUM

<u>AB</u>	ANDRX PHARMS	<u>EQ 375MG BASE</u>	<u>N75416</u>	<u>002</u>	Apr 23, 2003
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N75416</u>	<u>001</u>	Aug 27, 2002

NARATRIPTAN HYDROCHLORIDE

TABLET; ORAL

AMERGE

	GLAXOSMITHKLINE	EQ 1MG BASE	N20763	002	Feb 10, 1998
	+	EQ 2.5MG BASE	N20763	001	Feb 10, 1998

NATAMYCIN

SUSPENSION; OPHTHALMIC

NATACYN

	+ ALCON	5%	N50514	001	
--	---------	----	--------	-----	--

NATEGLINIDE

TABLET; ORAL

STARLIX

	NOVARTIS	60MG	N21204	001	Dec 22, 2000
	+	120MG	N21204	002	Dec 22, 2000

NEDOCROMIL SODIUM

AEROSOL, METERED; INHALATION

TILADE

	+ KING PHARMS	1.75MG/INH	N19660	001	Dec 30, 1992
--	---------------	------------	--------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

ALOCRI

	+ ALLERGAN	2%	N21009	001	Dec 08, 1999
--	------------	----	--------	-----	--------------

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS INC	<u>50MG</u>	<u>N76309</u>	<u>001</u>	Sep 16, 2003
<u>AB</u>		<u>100MG</u>	<u>N76309</u>	<u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>N76309</u>	<u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>N76309</u>	<u>004</u>	Sep 16, 2003
<u>AB</u>		<u>250MG</u>	<u>N76309</u>	<u>005</u>	Sep 16, 2003
<u>AB</u>	IVAX PHARMS	<u>50MG</u>	<u>N75763</u>	<u>001</u>	Sep 16, 2003
<u>AB</u>		<u>100MG</u>	<u>N75763</u>	<u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>N75763</u>	<u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>N75763</u>	<u>004</u>	Sep 16, 2003
<u>AB</u>		<u>250MG</u>	<u>N75763</u>	<u>005</u>	Sep 16, 2003
<u>AB</u>	MYLAN	<u>100MG</u>	<u>N76129</u>	<u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>N76129</u>	<u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>N76129</u>	<u>004</u>	Sep 16, 2003
<u>AB</u>		<u>250MG</u>	<u>N76129</u>	<u>005</u>	Sep 16, 2003
<u>AB</u>	RANBAXY	<u>50MG</u>	<u>N76409</u>	<u>001</u>	Sep 16, 2003
<u>AB</u>		<u>100MG</u>	<u>N76409</u>	<u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>N76409</u>	<u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>N76409</u>	<u>004</u>	Sep 16, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 267 (of 360)

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

<u>AB</u>	RANBAXY	<u>250MG</u>	<u>N76409</u>	<u>005</u>	Sep 16, 2003
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N76072</u>	<u>001</u>	Sep 16, 2003
<u>AB</u>		<u>50MG</u>	<u>N76302</u>	<u>001</u>	Sep 16, 2003
<u>AB</u>		<u>100MG</u>	<u>N76072</u>	<u>002</u>	Sep 16, 2003
<u>AB</u>		<u>100MG</u>	<u>N76302</u>	<u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>N76072</u>	<u>003</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>N76302</u>	<u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>N76072</u>	<u>004</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>N76302</u>	<u>004</u>	Sep 16, 2003
<u>AB</u>		<u>250MG</u>	<u>N76072</u>	<u>005</u>	Sep 16, 2003
<u>AB</u>		<u>250MG</u>	<u>N76302</u>	<u>005</u>	Sep 16, 2003
<u>AB</u>	TEVA	<u>50MG</u>	<u>N76037</u>	<u>001</u>	Sep 16, 2003
<u>AB</u>		<u>100MG</u>	<u>N76037</u>	<u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>N76037</u>	<u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>N76037</u>	<u>004</u>	Sep 16, 2003
<u>AB</u>	+	<u>250MG</u>	<u>N76037</u>	<u>005</u>	Sep 16, 2003
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N76073</u>	<u>002</u>	Sep 16, 2003
<u>AB</u>		<u>150MG</u>	<u>N76073</u>	<u>003</u>	Sep 16, 2003
<u>AB</u>		<u>200MG</u>	<u>N76073</u>	<u>004</u>	Sep 16, 2003
<u>AB</u>		<u>250MG</u>	<u>N76073</u>	<u>005</u>	Sep 16, 2003

NELARABINE

INJECTABLE; IV (INFUSION)

ARRANON

+	GLAXOSMITHKLINE	250MG/50ML (5MG/ML)	N21877	001	Oct 28, 2005
---	-----------------	---------------------	--------	-----	--------------

NELFINAVIR MESYLATE

FOR SUSPENSION; ORAL

VIRACEPT

+	AGOURON	EQ 50MG BASE/SCOOPFUL	N20778	001	Mar 14, 1997
---	---------	-----------------------	--------	-----	--------------

TABLET; ORAL

VIRACEPT

+	AGOURON	EQ 250MG BASE	N20779	001	Mar 14, 1997
---	---------	---------------	--------	-----	--------------

+		EQ 625MG BASE	N21503	001	Apr 30, 2003
---	--	---------------	--------	-----	--------------

NEOMYCIN SULFATE

POWDER; FOR RX COMPOUNDING

NEO-RX

	PHARMA TEK	100%	N61579	001	
--	------------	------	--------	-----	--

SOLUTION; ORAL

NEO-FRADIN

+	PHARMATEK	EQ 87.5MG BASE/5ML	N65010	001	May 23, 2002
---	-----------	--------------------	--------	-----	--------------

TABLET; ORAL

NEOMYCIN SULFATE

+	TEVA	EQ 350MG BASE	N60304	001	
---	------	---------------	--------	-----	--

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOSPORIN G.U. IRRIGANT

+	MONARCH PHARMS	EQ 40MG BASE/ML;200,000 UNITS/ML	N60707	001	
---	----------------	----------------------------------	--------	-----	--

NEOMYCIN SULFATE; POLYMYXIN B SULFATE; PREDNISOLONE ACETATE

SUSPENSION/DROPS; OPHTHALMIC

POLY-PRED

+	ALLERGAN	EQ 0.35% BASE;10,000 UNITS/ML;0.5%	N50081	002	
---	----------	------------------------------------	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 268 (of 360)

NEPAFENAC

SUSPENSION/DROPS; OPHTHALMIC

NEVANAC

+	ALCON	0.1%	N21862	001	Aug 19, 2005
---	-------	------	--------	-----	--------------

NESIRITIDE RECOMBINANT

FOR SOLUTION; INTRAVENOUS

NATRECOR

+	SCIOS	1.5MG/VIAL	N20920	001	Aug 10, 2001
---	-------	------------	--------	-----	--------------

NEVIRAPINE

SUSPENSION; ORAL

VIRAMUNE

+	BOEHRINGER INGELHEIM	50MG/5ML	N20933	001	Sep 11, 1998
---	----------------------	----------	--------	-----	--------------

TABLET; ORAL

VIRAMUNE

+	BOEHRINGER INGELHEIM	200MG	N20636	001	Jun 21, 1996
---	----------------------	-------	--------	-----	--------------

NIACIN

TABLET; ORAL

NIACIN

<u>AA</u>	WOCKHARDT	<u>500MG</u>	<u>N81134</u>	<u>001</u>	Apr 28, 1992
-----------	-----------	--------------	---------------	------------	--------------

NIACOR

<u>AA</u>	+	UPSHER SMITH	<u>500MG</u>	<u>N40378</u>	<u>001</u>	May 03, 2000
-----------	---	--------------	--------------	---------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

NIACIN

<u>AB</u>	BARR	<u>500MG</u>	<u>N76378</u>	<u>001</u>	Apr 26, 2005
-----------	------	--------------	---------------	------------	--------------

<u>AB</u>		<u>750MG</u>	<u>N76378</u>	<u>002</u>	Apr 26, 2005
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>1GM</u>	<u>N76250</u>	<u>001</u>	Apr 14, 2005
-----------	--	------------	---------------	------------	--------------

NIASPAN

<u>AB</u>	+	KOS LIFE	<u>500MG</u>	<u>N20381</u>	<u>002</u>	Jul 28, 1997
-----------	---	----------	--------------	---------------	------------	--------------

<u>AB</u>	+		<u>750MG</u>	<u>N20381</u>	<u>003</u>	Jul 28, 1997
-----------	---	--	--------------	---------------	------------	--------------

<u>AB</u>	+		<u>1GM</u>	<u>N20381</u>	<u>004</u>	Jul 28, 1997
-----------	---	--	------------	---------------	------------	--------------

NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL

CARDENE

<u>AB</u>	ROCHE PALO	<u>20MG</u>	<u>N19488</u>	<u>001</u>	Dec 21, 1988
-----------	------------	-------------	---------------	------------	--------------

<u>AB</u>	+		<u>N19488</u>	<u>002</u>	Dec 21, 1988
-----------	---	--	---------------	------------	--------------

NICARDIPINE HYDROCHLORIDE

<u>AB</u>	GENPHARM	<u>20MG</u>	<u>N74928</u>	<u>001</u>	Mar 19, 1998
-----------	----------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N74928</u>	<u>002</u>	Mar 19, 1998
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	IVAX PHARMS	<u>20MG</u>	<u>N74439</u>	<u>001</u>	Dec 10, 1996
-----------	-------------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N74439</u>	<u>002</u>	Dec 10, 1996
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	MYLAN	<u>20MG</u>	<u>N74642</u>	<u>001</u>	Jul 18, 1996
-----------	-------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N74642</u>	<u>002</u>	Jul 18, 1996
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>20MG</u>	<u>N74540</u>	<u>001</u>	Oct 28, 1996
-----------	------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N74540</u>	<u>002</u>	Oct 28, 1996
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	WATSON LABS	<u>20MG</u>	<u>N74670</u>	<u>001</u>	Oct 28, 1996
-----------	-------------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N74670</u>	<u>002</u>	Oct 28, 1996
-----------	--	-------------	---------------	------------	--------------

CAPSULE, EXTENDED RELEASE; ORAL

CARDENE SR

+	ROCHE PALO	30MG	N20005	001	Feb 21, 1992
---	------------	------	--------	-----	--------------

+		45MG	N20005	002	Feb 21, 1992
---	--	------	--------	-----	--------------

+		60MG	N20005	003	Feb 21, 1992
---	--	------	--------	-----	--------------

INJECTABLE; INJECTION

CARDENE

+	ESP PHARMA	2.5MG/ML	N19734	001	Jan 30, 1992
---	------------	----------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 269 (of 360)

NICOTINEFILM, EXTENDED RELEASE; TRANSDERMAL
NICOTINE

SANO	7MG/24HR	N74645	001	Oct 20, 1997
	14MG/24HR	N74611	001	Oct 20, 1997
	21MG/24HR	N74612	001	Oct 20, 1997

INHALANT; ORAL
NICOTROL

+ PHARMACIA AND UPJOHN	4MG/CARTRIDGE	N20714	001	May 02, 1997
------------------------	---------------	--------	-----	--------------

SPRAY, METERED; NASAL
NICOTROL

+ PHARMACIA AND UPJOHN	0.5MG/SPRAY	N20385	001	Mar 22, 1996
------------------------	-------------	--------	-----	--------------

NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

<u>AB</u>	FLEMINGTON PHARM	<u>10MG</u>	<u>N72781</u>	<u>001</u>	Jul 30, 1993
<u>AB</u>	PUREPAC PHARM	<u>10MG</u>	<u>N72579</u>	<u>001</u>	Jan 08, 1991
<u>AB</u>		<u>20MG</u>	<u>N72556</u>	<u>001</u>	Sep 20, 1990
<u>AB</u>	SCHERER RP	<u>10MG</u>	<u>N73250</u>	<u>001</u>	Oct 08, 1991
<u>AB</u>		<u>20MG</u>	<u>N74045</u>	<u>001</u>	Apr 30, 1992
<u>AB</u>	TEVA	<u>10MG</u>	<u>N72651</u>	<u>001</u>	Feb 19, 1992

PROCARDIA

<u>AB</u>	PFIZER	<u>10MG</u>	<u>N18482</u>	<u>001</u>	
<u>AB</u>	+	<u>20MG</u>	<u>N18482</u>	<u>002</u>	Jul 24, 1986

TABLET, EXTENDED RELEASE; ORAL

ADALAT CC

<u>AB1</u>	BAYER PHARMS	<u>30MG</u>	<u>N20198</u>	<u>001</u>	Apr 21, 1993
<u>AB1</u>	+	<u>60MG</u>	<u>N20198</u>	<u>002</u>	Apr 21, 1993
<u>AB1</u>	+	<u>90MG</u>	<u>N20198</u>	<u>003</u>	Apr 21, 1993

NIFEDIPINE

<u>AB1</u>	BIOVAIL	<u>30MG</u>	<u>N75269</u>	<u>001</u>	Dec 04, 2000
<u>AB1</u>		<u>60MG</u>	<u>N75269</u>	<u>002</u>	Dec 04, 2000
<u>AB1</u>		<u>90MG</u>	<u>N76070</u>	<u>001</u>	Aug 16, 2002
<u>AB2</u>		<u>30MG</u>	<u>N75289</u>	<u>002</u>	Feb 06, 2001
<u>AB2</u>		<u>60MG</u>	<u>N75289</u>	<u>001</u>	Sep 27, 2000
<u>AB1</u>	ELAN PHARM	<u>30MG</u>	<u>N75128</u>	<u>001</u>	Mar 10, 2000
<u>AB1</u>		<u>60MG</u>	<u>N75659</u>	<u>001</u>	Oct 26, 2001
<u>AB2</u>	MARTEC	<u>90MG</u>	<u>N75414</u>	<u>003</u>	Mar 23, 2004
<u>AB2</u>	OSMOTICA PHARM	<u>30MG</u>	<u>N77127</u>	<u>001</u>	Nov 21, 2005
<u>AB2</u>		<u>60MG</u>	<u>N77127</u>	<u>002</u>	Nov 21, 2005

PROCARDIA XL

<u>AB2</u>	PFIZER	<u>30MG</u>	<u>N19684</u>	<u>001</u>	Sep 06, 1989
<u>AB2</u>		<u>60MG</u>	<u>N19684</u>	<u>002</u>	Sep 06, 1989
<u>AB2</u>	+	<u>90MG</u>	<u>N19684</u>	<u>003</u>	Sep 06, 1989

NILUTAMIDE

TABLET; ORAL

NILANDRON

+ AVENTIS PHARMS	150MG	N20169	002	Apr 30, 1999
------------------	-------	--------	-----	--------------

NIMODIPINE

CAPSULE; ORAL

NIMOTOP

+ BAYER PHARMS	30MG	N18869	001	Dec 28, 1988
----------------	------	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 270 (of 360)

NISOLDIPINE

TABLET, EXTENDED RELEASE; ORAL
SULAR

+ FIRST HORIZON	10MG	N20356	001	Feb 02, 1995
	20MG	N20356	002	Feb 02, 1995
+	30MG	N20356	003	Feb 02, 1995
+	40MG	N20356	004	Feb 02, 1995

NITAZOXANIDE

FOR SUSPENSION; ORAL
ALINIA

+ ROMARK	100MG/5ML	N21498	001	Nov 22, 2002
----------	-----------	--------	-----	--------------

TABLET; ORAL
ALINIA

+ ROMARK	500MG	N21497	001	Jul 21, 2004
----------	-------	--------	-----	--------------

NITISINONE

CAPSULE; ORAL
ORFADIN

SWEDISH ORPHAN	2MG	N21232	001	Jan 18, 2002
	5MG	N21232	002	Jan 18, 2002
+	10MG	N21232	003	Jan 18, 2002

NITRIC OXIDE

GAS; INHALATION
INOMAX

INO	100PPM	N20845	002	Dec 23, 1999
+	800PPM	N20845	003	Dec 23, 1999

NITROFURANTOIN

SUSPENSION; ORAL
FURADANTIN

+ FIRST HORIZON	25MG/5ML	N09175	001	
-----------------	----------	--------	-----	--

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL
MACRODANTIN

<u>AB</u>	PROCTER AND GAMBLE	<u>25MG</u>	<u>N16620</u>	<u>003</u>	
<u>AB</u>		<u>50MG</u>	<u>N16620</u>	<u>001</u>	
<u>AB</u>	+	<u>100MG</u>	<u>N16620</u>	<u>002</u>	
	<u>NITROFURANTOIN</u>				
<u>AB</u>	IVAX PHARMS	<u>50MG</u>	<u>N73671</u>	<u>001</u>	Jan 28, 1993
<u>AB</u>		<u>100MG</u>	<u>N73652</u>	<u>001</u>	Jan 28, 1993
<u>AB</u>	MYLAN	<u>50MG</u>	<u>N74967</u>	<u>001</u>	Jul 09, 1997
<u>AB</u>		<u>100MG</u>	<u>N77025</u>	<u>001</u>	Aug 18, 2004
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>N74336</u>	<u>001</u>	Jan 25, 1995
<u>AB</u>		<u>50MG</u>	<u>N74336</u>	<u>002</u>	Jan 25, 1995
<u>AB</u>		<u>100MG</u>	<u>N74336</u>	<u>003</u>	Jan 25, 1995
<u>AB</u>	WATSON LABS	<u>25MG</u>	<u>N73696</u>	<u>001</u>	Dec 31, 1992
<u>AB</u>		<u>50MG</u>	<u>N73696</u>	<u>002</u>	Dec 31, 1992
<u>AB</u>		<u>100MG</u>	<u>N73696</u>	<u>003</u>	Dec 31, 1992

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL
MACROBID

<u>AB</u>	+ PROCTER AND GAMBLE	<u>75MG;25MG</u>	<u>N20064</u>	<u>001</u>	Dec 24, 1991
	<u>NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)</u>				
<u>AB</u>	MYLAN	<u>75MG;25MG</u>	<u>N76648</u>	<u>001</u>	Mar 22, 2004
<u>AB</u>	RANBAXY	<u>75MG;25MG</u>	<u>N76951</u>	<u>001</u>	Mar 30, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 271 (of 360)

NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS)

<u>AB</u>	SANDOZ	<u>75MG;25MG</u>	<u>N77066</u>	<u>001</u>	Apr 05, 2005
-----------	--------	------------------	---------------	------------	--------------

NITROFURAZONE

OINTMENT; TOPICAL

NITROFURAZONE

+ TARO

0.2%

N86156 001

NITROGLYCERIN

FILM, EXTENDED RELEASE; TRANSDERMAL

MINITRAN

<u>AB1</u>	3M	<u>0.1MG/HR</u>	<u>N89771</u>	<u>001</u>	Aug 30, 1996
<u>AB1</u>		<u>0.2MG/HR</u>	<u>N89772</u>	<u>001</u>	Aug 30, 1996
<u>AB1</u>		<u>0.4MG/HR</u>	<u>N89773</u>	<u>001</u>	Aug 30, 1996
<u>AB1</u>		<u>0.6MG/HR</u>	<u>N89774</u>	<u>001</u>	Aug 30, 1996

NITRO-DUR

<u>AB1</u> +	KEY PHARMS	<u>0.1MG/HR</u>	<u>N20145</u>	<u>001</u>	Apr 04, 1995
<u>AB1</u> +		<u>0.2MG/HR</u>	<u>N20145</u>	<u>002</u>	Apr 04, 1995
<u>AB1</u> +		<u>0.4MG/HR</u>	<u>N20145</u>	<u>004</u>	Apr 04, 1995
<u>AB1</u> +		<u>0.6MG/HR</u>	<u>N20145</u>	<u>005</u>	Apr 04, 1995
BX +		0.8MG/HR	N20145	006	Apr 04, 1995
+		0.3MG/HR	N20145	003	Apr 04, 1995

NITROGLYCERIN

<u>AB2</u>	HERCON LABS	<u>0.2MG/HR</u>	<u>N89884</u>	<u>001</u>	Oct 30, 1998
<u>AB2</u>		<u>0.4MG/HR</u>	<u>N89885</u>	<u>001</u>	Oct 30, 1998
<u>AB2</u>		<u>0.6MG/HR</u>	<u>N89886</u>	<u>001</u>	Oct 30, 1998
<u>AB1</u>	KREMERS URBAN	<u>0.2MG/HR</u>	<u>N75115</u>	<u>001</u>	Aug 10, 2004
<u>AB1</u>		<u>0.4MG/HR</u>	<u>N75115</u>	<u>002</u>	Aug 10, 2004
<u>AB1</u>	MYLAN TECHNOLOGIES	<u>0.1MG/HR</u>	<u>N75076</u>	<u>001</u>	Nov 12, 1999
<u>AB1</u>		<u>0.2MG/HR</u>	<u>N75073</u>	<u>001</u>	Nov 12, 1999
<u>AB1</u>		<u>0.4MG/HR</u>	<u>N75075</u>	<u>001</u>	Nov 12, 1999
<u>AB1</u>		<u>0.6MG/HR</u>	<u>N74992</u>	<u>001</u>	Nov 12, 1999
<u>AB2</u> +		<u>0.1MG/HR</u>	<u>N75033</u>	<u>001</u>	Feb 06, 1998
<u>AB2</u> +		<u>0.2MG/HR</u>	<u>N74609</u>	<u>001</u>	Aug 30, 1996
<u>AB2</u> +		<u>0.4MG/HR</u>	<u>N74607</u>	<u>001</u>	Aug 30, 1996
<u>AB2</u> +		<u>0.6MG/HR</u>	<u>N74559</u>	<u>001</u>	Aug 30, 1996

INJECTABLE; INJECTION

NITROGLYCERIN

<u>AP</u> +	HOSPIRA	<u>5MG/ML</u>	<u>N18531</u>	<u>001</u>	
+	LUITPOLD	5MG/ML	N72034	001	May 24, 1988

NITROGLYCERIN IN DEXTROSE 5%

<u>AP</u> +	BAXTER HLTHCARE	<u>10MG/100ML</u>	<u>N19970</u>	<u>001</u>	Dec 29, 1989
<u>AP</u> +		<u>20MG/100ML</u>	<u>N19970</u>	<u>002</u>	Dec 29, 1989
<u>AP</u> +		<u>40MG/100ML</u>	<u>N19970</u>	<u>003</u>	Dec 29, 1989
<u>AP</u>	HOSPIRA	<u>10MG/100ML</u>	<u>N71846</u>	<u>001</u>	Aug 31, 1990
<u>AP</u>		<u>20MG/100ML</u>	<u>N71847</u>	<u>001</u>	Aug 31, 1990
<u>AP</u>		<u>40MG/100ML</u>	<u>N71848</u>	<u>001</u>	Aug 31, 1990

OINTMENT; TRANSDERMAL

NITROGLYCERIN

+ FOUGERA

2%

N87355 001 Jul 08, 1988

SPRAY, METERED; SUBLINGUAL

NITROLINGUAL PUMPSPRAY

+ POHL BOSKAMP

0.4MG/SPRAY

N18705 002 Jan 10, 1997

TABLET; SUBLINGUAL

NITROSTAT

PFIZER PHARMS

0.3MG

N21134 001 May 01, 2000

0.4MG

N21134 002 May 01, 2000

PRESCRIPTION DRUG PRODUCT LIST

3 - 272 (of 360)

NITROGLYCERIN

TABLET; SUBLINGUAL

NITROSTAT

+ PFIZER PHARMS	0.6MG	N21134	003	May 01, 2000
-----------------	-------	--------	-----	--------------

NIZATIDINE

CAPSULE; ORAL

AXID

<u>AB</u> RELIANT PHARMS	<u>150MG</u>	<u>N19508</u>	<u>001</u>	Apr 12, 1988
--------------------------	--------------	---------------	------------	--------------

<u>AB</u> +	<u>300MG</u>	<u>N19508</u>	<u>002</u>	Apr 12, 1988
-------------	--------------	---------------	------------	--------------

NIZATIDINE

<u>AB</u> DR REDDYS LABS LTD	<u>150MG</u>	<u>N77314</u>	<u>001</u>	Sep 15, 2005
------------------------------	--------------	---------------	------------	--------------

<u>AB</u>	<u>300MG</u>	<u>N77314</u>	<u>002</u>	Sep 15, 2005
-----------	--------------	---------------	------------	--------------

<u>AB</u> GENPHARM	<u>150MG</u>	<u>N75934</u>	<u>001</u>	Jul 09, 2002
--------------------	--------------	---------------	------------	--------------

<u>AB</u>	<u>300MG</u>	<u>N75934</u>	<u>002</u>	Jul 09, 2002
-----------	--------------	---------------	------------	--------------

<u>AB</u> MYLAN	<u>150MG</u>	<u>N75806</u>	<u>001</u>	Jul 05, 2002
-----------------	--------------	---------------	------------	--------------

<u>AB</u>	<u>300MG</u>	<u>N75806</u>	<u>002</u>	Jul 05, 2002
-----------	--------------	---------------	------------	--------------

<u>AB</u> SANDOZ	<u>150MG</u>	<u>N76178</u>	<u>001</u>	Jul 05, 2002
------------------	--------------	---------------	------------	--------------

<u>AB</u>	<u>300MG</u>	<u>N76178</u>	<u>002</u>	Jul 05, 2002
-----------	--------------	---------------	------------	--------------

<u>AB</u> TEVA	<u>150MG</u>	<u>N75668</u>	<u>001</u>	Sep 12, 2002
----------------	--------------	---------------	------------	--------------

<u>AB</u>	<u>300MG</u>	<u>N75668</u>	<u>002</u>	Sep 12, 2002
-----------	--------------	---------------	------------	--------------

<u>AB</u> TORPHARM	<u>150MG</u>	<u>N76383</u>	<u>001</u>	Jan 23, 2003
--------------------	--------------	---------------	------------	--------------

<u>AB</u>	<u>300MG</u>	<u>N76383</u>	<u>002</u>	Jan 23, 2003
-----------	--------------	---------------	------------	--------------

<u>AB</u> WATSON LABS	<u>150MG</u>	<u>N75616</u>	<u>001</u>	Jul 09, 2002
-----------------------	--------------	---------------	------------	--------------

<u>AB</u>	<u>300MG</u>	<u>N75616</u>	<u>002</u>	Jul 09, 2002
-----------	--------------	---------------	------------	--------------

<u>AB</u> ZENITH GOLDLINE	<u>150MG</u>	<u>N75461</u>	<u>001</u>	Jul 08, 2002
---------------------------	--------------	---------------	------------	--------------

<u>AB</u>	<u>300MG</u>	<u>N75461</u>	<u>002</u>	Jul 08, 2002
-----------	--------------	---------------	------------	--------------

SOLUTION; ORAL

AXID

+ BRAINTREE	15MG/ML	N21494	001	May 25, 2004
-------------	---------	--------	-----	--------------

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

LEVOPHED

<u>AP</u> + HOSPIRA	<u>EQ 1MG BASE/ML</u>	<u>N07513</u>	<u>001</u>	
---------------------	-----------------------	---------------	------------	--

NOREPINEPHRINE BITARTRATE

<u>AP</u> BEDFORD	<u>EQ 1MG BASE/ML</u>	<u>N40462</u>	<u>001</u>	Oct 31, 2003
-------------------	-----------------------	---------------	------------	--------------

<u>AP</u> PHARMAFORCE	<u>EQ 1MG BASE/ML</u>	<u>N40522</u>	<u>001</u>	Sep 30, 2004
-----------------------	-----------------------	---------------	------------	--------------

<u>AP</u> SICOR PHARMS	<u>EQ 1MG BASE/ML</u>	<u>N40455</u>	<u>001</u>	Mar 03, 2003
------------------------	-----------------------	---------------	------------	--------------

NORETHINDRONE

TABLET; ORAL-28

CAMILA

<u>AB1</u> BARR	<u>0.35MG</u>	<u>N76177</u>	<u>001</u>	Oct 21, 2002
-----------------	---------------	---------------	------------	--------------

ERRIN

<u>AB2</u> BARR	<u>0.35MG</u>	<u>N76225</u>	<u>001</u>	Oct 21, 2002
-----------------	---------------	---------------	------------	--------------

MICRONOR

<u>AB2</u> + ORTHO MCNEIL PHARM	<u>0.35MG</u>	<u>N16954</u>	<u>001</u>	
---------------------------------	---------------	---------------	------------	--

NOR-QD

<u>AB1</u> + WATSON LABS (UTAH)	<u>0.35MG</u>	<u>N17060</u>	<u>001</u>	
---------------------------------	---------------	---------------	------------	--

NORETHINDRONE ACETATE

TABLET; ORAL

AYGESTIN

<u>AB</u> + DURAMED PHARMS BARR	<u>5MG</u>	<u>N18405</u>	<u>001</u>	Apr 21, 1982
---------------------------------	------------	---------------	------------	--------------

NORETHINDRONE ACETATE

<u>AB</u> BARR	<u>5MG</u>	<u>N75951</u>	<u>001</u>	May 25, 2001
----------------	------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 273 (of 360)

NORFLOXACIN

TABLET; ORAL

NOROXIN

+ MERCK

400MG

N19384 002

Oct 31, 1986

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

NORTRIPTYLINE HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>EQ 10MG BASE</u>	<u>N74234</u>	<u>001</u>	Jul 26, 1993
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N74234</u>	<u>002</u>	Jul 26, 1993
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N74234</u>	<u>003</u>	Jul 26, 1993
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N74234</u>	<u>004</u>	Jul 26, 1993
<u>AB</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>N74054</u>	<u>001</u>	Dec 31, 1992
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N74835</u>	<u>001</u>	Jun 30, 1997
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N74054</u>	<u>002</u>	Dec 31, 1992
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N74835</u>	<u>002</u>	Jun 30, 1997
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N74054</u>	<u>003</u>	Dec 31, 1992
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N74835</u>	<u>003</u>	Jun 30, 1997
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N74054</u>	<u>004</u>	Dec 31, 1992
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N74835</u>	<u>004</u>	Jun 30, 1997
<u>AB</u>	TARO	<u>EQ 10MG BASE</u>	<u>N75520</u>	<u>004</u>	May 08, 2000
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N75520</u>	<u>003</u>	May 08, 2000
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N75520</u>	<u>001</u>	May 08, 2000
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N75520</u>	<u>002</u>	May 08, 2000
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>N73667</u>	<u>001</u>	Apr 11, 1996
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N74132</u>	<u>001</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N73667</u>	<u>002</u>	Apr 11, 1996
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N74132</u>	<u>002</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N73667</u>	<u>003</u>	Apr 11, 1996
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N74132</u>	<u>003</u>	Mar 27, 1995
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N73667</u>	<u>004</u>	Apr 11, 1996
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N74132</u>	<u>004</u>	Mar 27, 1995
<u>AB</u>	WATSON LABS	<u>EQ 10MG BASE</u>	<u>N73553</u>	<u>001</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N73554</u>	<u>001</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N73555</u>	<u>001</u>	Mar 30, 1992
<u>AB</u>		<u>EQ 75MG BASE</u>	<u>N73556</u>	<u>001</u>	Mar 30, 1992
<u>PAMELOR</u>					
<u>AB</u>	TYCO HLTHCARE	<u>EQ 10MG BASE</u>	<u>N18013</u>	<u>001</u>	
<u>AB</u>		<u>EQ 25MG BASE</u>	<u>N18013</u>	<u>002</u>	
<u>AB</u>		<u>EQ 50MG BASE</u>	<u>N18013</u>	<u>004</u>	
<u>AB</u>	+	<u>EQ 75MG BASE</u>	<u>N18013</u>	<u>003</u>	

SOLUTION; ORAL

AVENTYL HYDROCHLORIDE

<u>AA</u>	+ RANBAXY	<u>EQ 10MG BASE/5ML</u>	<u>N14685</u>	<u>001</u>	
<u>NORTRIPTYLINE HYDROCHLORIDE</u>					
<u>AA</u>	PHARM ASSOC	<u>EQ 10MG BASE/5ML</u>	<u>N75606</u>	<u>001</u>	Aug 28, 2000
<u>PAMELOR</u>					
<u>AA</u>	TYCO HLTHCARE	<u>EQ 10MG BASE/5ML</u>	<u>N18012</u>	<u>001</u>	

NYSTATIN

CREAM; TOPICAL

MYCOSTATIN

<u>AT</u>	+ WESTWOOD SQUIBB	<u>100,000 UNITS/GM</u>	<u>N60575</u>	<u>001</u>	
<u>NYSTATIN</u>					
<u>AT</u>	ALPHARMA US PHARMS	<u>100,000 UNITS/GM</u>	<u>N62949</u>	<u>001</u>	Jun 13, 1988
<u>AT</u>	ALTANA	<u>100,000 UNITS/GM</u>	<u>N62129</u>	<u>001</u>	
<u>AT</u>	CLAY PARK	<u>100,000 UNITS/GM</u>	<u>N62225</u>	<u>001</u>	
<u>AT</u>	TARO	<u>100,000 UNITS/GM</u>	<u>N62457</u>	<u>001</u>	Jul 28, 1983
<u>AT</u>		<u>100,000 UNITS/GM</u>	<u>N64022</u>	<u>001</u>	Jan 29, 1993

PRESCRIPTION DRUG PRODUCT LIST

3 - 274 (of 360)

NYSTATIN

OINTMENT; TOPICAL

NYSTATIN

<u>AT</u>	ALPHARMA US PHARMS	<u>100,000 UNITS/GM</u>	<u>N62840</u>	<u>001</u>	Nov 13, 1987
<u>AT</u>	+ ALTANA	<u>100,000 UNITS/GM</u>	<u>N62124</u>	<u>002</u>	Sep 23, 1982
<u>AT</u>	CLAY PARK	<u>100,000 UNITS/GM</u>	<u>N62472</u>	<u>001</u>	Feb 13, 1984

POWDER; ORAL

NYSTATIN

	+ PADDOCK	100%	N62613	001	Nov 26, 1985
--	-----------	------	--------	-----	--------------

POWDER; TOPICAL

MYCOSTATIN

<u>AT</u>	+ WESTWOOD SQUIBB	<u>100,000 UNITS/GM</u>	<u>N60578</u>	<u>001</u>	
<u>AT</u>	<u>NYSTATIN</u>				
<u>AT</u>	KALI LABS	<u>100,000 UNITS/GM</u>	<u>N65138</u>	<u>001</u>	Jul 23, 2004
<u>AT</u>	TANON	<u>100,000 UNITS/GM</u>	<u>N65203</u>	<u>001</u>	Jul 15, 2004
<u>AT</u>	UPSHER SMITH	<u>100,000 UNITS/GM</u>	<u>N65183</u>	<u>001</u>	May 03, 2005
<u>AT</u>	X GEN PHARMS	<u>100,000 UNITS/GM</u>	<u>N65175</u>	<u>001</u>	Dec 17, 2004

NYSTOP

<u>AT</u>	PADDOCK	<u>100,000 UNITS/GM</u>	<u>N64118</u>	<u>001</u>	Aug 16, 1996
-----------	---------	-------------------------	---------------	------------	--------------

SUSPENSION; ORAL

NILSTAT

<u>AA</u>	+ GLENMARK PHARMA	<u>100,000 UNITS/ML</u>	<u>N50299</u>	<u>001</u>	
<u>AA</u>	<u>NYSTATIN</u>				
<u>AA</u>	ALPHARMA US PHARMS	<u>100,000 UNITS/ML</u>	<u>N62349</u>	<u>001</u>	Jul 14, 1982
<u>AA</u>	BAUSCH AND LOMB	<u>100,000 UNITS/ML</u>	<u>N64042</u>	<u>001</u>	Feb 28, 1994
<u>AA</u>	FOUGERA	<u>100,000 UNITS/ML</u>	<u>N62517</u>	<u>001</u>	Jun 07, 1984
<u>AA</u>	+ MORTON GROVE	<u>100,000 UNITS/ML</u>	<u>N62512</u>	<u>001</u>	Oct 29, 1984
<u>AA</u>	VINTAGE PHARMS	<u>100,000 UNITS/ML</u>	<u>N65148</u>	<u>001</u>	Jun 28, 2005
<u>AA</u>	VISTAPHARM	<u>100,000 UNITS/ML</u>	<u>N64142</u>	<u>001</u>	Jun 25, 1998

TABLET; ORAL

MYCOSTATIN

<u>AA</u>	+ APOTHECON	<u>500,000 UNITS</u>	<u>N60574</u>	<u>001</u>	
<u>AA</u>	<u>NYSTATIN</u>				
<u>AA</u>	MUTUAL PHARM	<u>500,000 UNITS</u>	<u>N62838</u>	<u>001</u>	Dec 22, 1988
<u>AA</u>	PAR PHARM	<u>500,000 UNITS</u>	<u>N62474</u>	<u>001</u>	Dec 22, 1983
<u>AA</u>	TEVA	<u>500,000 UNITS</u>	<u>N62506</u>	<u>001</u>	Jan 16, 1984

TABLET; VAGINAL

NYSTATIN

	+ ODYSSEY PHARMS	100,000 UNITS	N62615	001	Oct 17, 1985
--	------------------	---------------	--------	-----	--------------

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYKACET

<u>AT</u>	ALPHARMA US PHARMS	<u>100,000 UNITS/GM;0.1%</u>	<u>N62367</u>	<u>001</u>	May 28, 1985
<u>AT</u>	<u>NYSTATIN AND TRIAMCINOLONE ACETONIDE</u>				
<u>AT</u>	+ TARO	<u>100,000 UNITS/GM;0.1%</u>	<u>N62364</u>	<u>001</u>	Dec 22, 1987
<u>AT</u>	<u>NYSTATIN-TRIAMCINOLONE ACETONIDE</u>				
<u>AT</u>	FOUGERA	<u>100,000 UNITS/GM;0.1%</u>	<u>N62599</u>	<u>001</u>	Oct 08, 1985

OINTMENT; TOPICAL

MYKACET

<u>AT</u>	ALPHARMA US PHARMS	<u>100,000 UNITS/GM;0.1%</u>	<u>N62733</u>	<u>001</u>	Mar 09, 1987
<u>AT</u>	<u>NYSTATIN AND TRIAMCINOLONE ACETONIDE</u>				
<u>AT</u>	+ TARO	<u>100,000 UNITS/GM;0.1%</u>	<u>N63305</u>	<u>001</u>	Mar 29, 1993
<u>AT</u>	<u>NYSTATIN-TRIAMCINOLONE ACETONIDE</u>				
<u>AT</u>	FOUGERA	<u>100,000 UNITS/GM;0.1%</u>	<u>N62602</u>	<u>001</u>	Oct 09, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 275 (of 360)

OCTREOTIDE ACETATE

INJECTABLE; INJECTION

OCTREOTIDE ACETATE

<u>AP</u>	BEDFORD	<u>EQ 0.2MG BASE/ML</u>	<u>N76330</u>	<u>001</u>	Apr 08, 2005
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>N76330</u>	<u>002</u>	Apr 08, 2005
<u>AP</u>	SICOR PHARMS	<u>EQ 0.05MG BASE/ML</u>	<u>N75957</u>	<u>001</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>N75957</u>	<u>002</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 0.2MG BASE/ML</u>	<u>N75959</u>	<u>001</u>	Nov 21, 2005
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>N75957</u>	<u>003</u>	Oct 03, 2005
<u>AP</u>		<u>EQ 1MG BASE/ML</u>	<u>N75959</u>	<u>002</u>	Nov 21, 2005
<u>OCTREOTIDE ACETATE (PRESERVATIVE FREE)</u>					
<u>AP</u>	BEDFORD	<u>EQ 0.05MG BASE/ML</u>	<u>N76313</u>	<u>001</u>	Mar 28, 2005
<u>AP</u>		<u>EQ 0.1MG BASE/ML</u>	<u>N76313</u>	<u>003</u>	Mar 28, 2005
<u>AP</u>		<u>EQ 0.5MG BASE/ML</u>	<u>N76313</u>	<u>002</u>	Mar 28, 2005
<u>SANDOSTATIN</u>					
<u>AP</u>	+ NOVARTIS	<u>EQ 0.05MG BASE/ML</u>	<u>N19667</u>	<u>001</u>	Oct 21, 1988
<u>AP</u>	+	<u>EQ 0.1MG BASE/ML</u>	<u>N19667</u>	<u>002</u>	Oct 21, 1988
<u>AP</u>	+	<u>EQ 0.2MG BASE/ML</u>	<u>N19667</u>	<u>004</u>	Jun 12, 1991
<u>AP</u>	+	<u>EQ 0.5MG BASE/ML</u>	<u>N19667</u>	<u>003</u>	Oct 21, 1988
<u>AP</u>	+	<u>EQ 1MG BASE/ML</u>	<u>N19667</u>	<u>005</u>	Jun 12, 1991
	SANDOSTATIN LAR				
	NOVARTIS	EQ 10MG BASE/VIAL	N21008	001	Nov 25, 1998
		EQ 20MG BASE/VIAL	N21008	002	Nov 25, 1998
	+	EQ 30MG BASE/VIAL	N21008	003	Nov 25, 1998

OFLOXACIN

INJECTABLE; INJECTION

OFLOXACIN

+	BEDFORD	40MG/ML	N75762	001	Jan 16, 2002
---	---------	---------	--------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

OCUFLOX

<u>AT</u>	+ ALLERGAN	<u>0.3%</u>	<u>N19921</u>	<u>001</u>	Jul 30, 1993
<u>OFLOXACIN</u>					
<u>AT</u>	ALCON	<u>0.3%</u>	<u>N76231</u>	<u>001</u>	May 14, 2004
<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>N76622</u>	<u>001</u>	May 14, 2004
<u>AT</u>	HI TECH PHARMA	<u>0.3%</u>	<u>N76615</u>	<u>001</u>	May 14, 2004
<u>AT</u>	NOVEX	<u>0.3%</u>	<u>N76513</u>	<u>001</u>	May 14, 2004
<u>AT</u>	PHARMAFORCE	<u>0.3%</u>	<u>N76830</u>	<u>001</u>	Aug 31, 2004

SOLUTION/DROPS; OTIC

FLOXIN OTIC

<u>AT</u>	+ DAIICHI	<u>0.3%</u>	<u>N20799</u>	<u>001</u>	Dec 16, 1997
<u>OFLOXACIN</u>					
<u>AT</u>	NOVEX	<u>0.3%</u>	<u>N76527</u>	<u>001</u>	Nov 18, 2005

TABLET; ORAL

FLOXIN

<u>AB</u>	ORTHO MCNEIL PHARM	<u>200MG</u>	<u>N19735</u>	<u>001</u>	Dec 28, 1990
<u>AB</u>		<u>300MG</u>	<u>N19735</u>	<u>002</u>	Dec 28, 1990
<u>AB</u>	+	<u>400MG</u>	<u>N19735</u>	<u>003</u>	Dec 28, 1990
<u>OFLOXACIN</u>					
<u>AB</u>	PAR PHARM	<u>200MG</u>	<u>N76093</u>	<u>001</u>	Sep 02, 2003
<u>AB</u>		<u>300MG</u>	<u>N76093</u>	<u>002</u>	Sep 02, 2003
<u>AB</u>		<u>400MG</u>	<u>N76093</u>	<u>003</u>	Sep 02, 2003
<u>AB</u>	RANBAXY	<u>200MG</u>	<u>N76220</u>	<u>001</u>	Sep 02, 2003
<u>AB</u>		<u>300MG</u>	<u>N76220</u>	<u>002</u>	Sep 02, 2003
<u>AB</u>		<u>400MG</u>	<u>N76220</u>	<u>003</u>	Sep 02, 2003
<u>AB</u>	TEVA	<u>200MG</u>	<u>N76182</u>	<u>001</u>	Sep 02, 2003
<u>AB</u>		<u>300MG</u>	<u>N76182</u>	<u>002</u>	Sep 02, 2003
<u>AB</u>		<u>400MG</u>	<u>N76182</u>	<u>003</u>	Sep 02, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 276 (of 360)

OLANZAPINE

INJECTABLE; INTRAMUSCULAR

ZYPREXA

+ LILLY	10MG/VIAL	N21253	001	Mar 29, 2004
---------	-----------	--------	-----	--------------

TABLET; ORAL

ZYPREXA

LILLY	2.5MG	N20592	001	Sep 30, 1996
+	5MG	N20592	002	Sep 30, 1996
	7.5MG	N20592	003	Sep 30, 1996
	10MG	N20592	004	Sep 30, 1996
	15MG	N20592	005	Sep 09, 1997
	20MG	N20592	006	Sep 09, 1997

TABLET, ORALLY DISINTEGRATING; ORAL
ZYPREXA ZYDIS

+ LILLY	5MG	N21086	001	Apr 06, 2000
	10MG	N21086	002	Apr 06, 2000
	15MG	N21086	003	Apr 06, 2000
	20MG	N21086	004	Apr 06, 2000

OLMESARTAN MEDOXOMIL

TABLET; ORAL

BENICAR

SANKYO	5MG	N21286	001	Apr 25, 2002
	20MG	N21286	003	Apr 25, 2002
+	40MG	N21286	004	Apr 25, 2002

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC
OLOPATADINE HYDROCHLORIDE

+ ALCON	0.2%	N21545	001	Dec 22, 2004
	PATANOL			
+	ALCON	N20688	001	Dec 18, 1996

OLSALAZINE SODIUM

CAPSULE; ORAL

DIPENTUM

+ UCB	250MG	N19715	001	Jul 31, 1990
-------	-------	--------	-----	--------------

OMEGA-3-ACID ETHYL ESTERS

CAPSULE; ORAL

OMACOR

+ RELIANT PHARMS INC	1GM	N21654	001	Nov 10, 2004
----------------------	-----	--------	-----	--------------

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

OMEPRAZOLE

<u>AB</u>	IMPAX LABS	<u>10MG</u>	<u>N75785</u>	<u>001</u>	Nov 08, 2002
<u>AB</u>		<u>20MG</u>	<u>N75785</u>	<u>002</u>	Nov 08, 2002
<u>AB</u>	KREMERS URBAN DEV	<u>10MG</u>	<u>N75410</u>	<u>001</u>	Nov 01, 2002
<u>AB</u>		<u>20MG</u>	<u>N75410</u>	<u>002</u>	Nov 01, 2002
<u>AB</u>	LEK PHARMS	<u>10MG</u>	<u>N75757</u>	<u>001</u>	Jan 28, 2003
<u>AB</u>		<u>20MG</u>	<u>N75757</u>	<u>002</u>	Jan 28, 2003
<u>AB</u>	MYLAN	<u>10MG</u>	<u>N75876</u>	<u>001</u>	May 29, 2003
<u>AB</u>		<u>20MG</u>	<u>N75876</u>	<u>002</u>	May 29, 2003
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>N75791</u>	<u>001</u>	Dec 23, 2002
<u>AB</u>		<u>20MG</u>	<u>N75791</u>	<u>002</u>	Dec 23, 2002
<u>AB</u>	TORPHARM	<u>10MG</u>	<u>N76048</u>	<u>001</u>	Oct 06, 2003
<u>AB</u>		<u>20MG</u>	<u>N76048</u>	<u>002</u>	Oct 06, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 277 (of 360)

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

PRILOSEC

<u>AB</u>	ASTRAZENECA	<u>10MG</u>	<u>N19810</u>	<u>003</u>	Oct 05, 1995
<u>AB</u>	+	<u>20MG</u>	<u>N19810</u>	<u>001</u>	Sep 14, 1989
	+	40MG	N19810	002	Jan 15, 1998

FOR SUSPENSION; ORAL

ZEGERID

+	SANTARUS	20MG/PACKET	N21636	001	Jun 15, 2004
+		40MG/PACKET	N21706	001	Dec 21, 2004

ONDANSETRON

TABLET, ORALLY DISINTEGRATING; ORAL

ZOFRAN ODT

	GLAXOSMITHKLINE	EQ 4MG BASE	N20781	001	Jan 27, 1999
+		EQ 8MG BASE	N20781	002	Jan 27, 1999

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ZOFRAN

+	GLAXOSMITHKLINE	EQ 2MG BASE/ML	N20007	001	Jan 04, 1991
	ZOFRAN IN PLASTIC CONTAINER				
+	GLAXOSMITHKLINE	EQ 0.64MG BASE/ML	N20403	001	Jan 31, 1995
	ZOFRAN PRESERVATIVE FREE				
+	GLAXOSMITHKLINE	EQ 2MG BASE/ML	N20007	003	Dec 10, 1993

SOLUTION; ORAL

ZOFRAN

+	GLAXOSMITHKLINE	EQ 4MG BASE/5ML	N20605	001	Jan 24, 1997
---	-----------------	-----------------	--------	-----	--------------

TABLET; ORAL

ZOFRAN

	GLAXOSMITHKLINE	EQ 4MG BASE	N20103	001	Dec 31, 1992
		EQ 8MG BASE	N20103	002	Dec 31, 1992
+		EQ 24MG BASE	N20103	003	Aug 27, 1999

ORLISTAT

CAPSULE; ORAL

XENICAL

+	HLR	120MG	N20766	001	Apr 23, 1999
---	-----	-------	--------	-----	--------------

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

NORFLEX

<u>AP</u>	+	3M	<u>30MG/ML</u>	<u>N13055</u>	<u>001</u>	
		<u>ORPHENADRINE CITRATE</u>				
<u>AP</u>		BEDFORD LABS	<u>30MG/ML</u>	<u>N40463</u>	<u>001</u>	Mar 04, 2003
<u>AP</u>		STERIS	<u>30MG/ML</u>	<u>N84779</u>	<u>001</u>	Mar 15, 1982
<u>AP</u>			<u>30MG/ML</u>	<u>N87062</u>	<u>001</u>	

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATE

<u>AB</u>		IMPAX PHARMS	<u>100MG</u>	<u>N40368</u>	<u>001</u>	Jun 23, 2000
<u>AB</u>		KIEL	<u>100MG</u>	<u>N40249</u>	<u>001</u>	Jan 29, 1999
<u>AB</u>		SANDOZ	<u>100MG</u>	<u>N40284</u>	<u>001</u>	Jun 19, 1998
<u>AB</u>	+		<u>100MG</u>	<u>N40327</u>	<u>001</u>	Feb 15, 2000

OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

TAMIFLU

+	ROCHE	EQ 75MG BASE	N21087	001	Oct 27, 1999
---	-------	--------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 278 (of 360)

OSELTAMIVIR PHOSPHATE

FOR SUSPENSION; ORAL

TAMIFLU

+ ROCHE

EQ 12MG BASE/ML

N21246 001

Dec 14, 2000

OXACILLIN SODIUM

CAPSULE; ORAL

BACTOCILLAB GLAXOSMITHKLINEEQ 250MG BASEN61336 001AB +EQ 500MG BASEN61336 002OXACILLIN SODIUMAB TEVAEQ 250MG BASEN62222 001ABEQ 500MG BASEN62222 002

FOR SOLUTION; ORAL

BACTOCILLAA GLAXOSMITHKLINEEQ 250MG BASE/5MLN62321 001OXACILLIN SODIUMAA TEVAEQ 250MG BASE/5MLN62252 001

INJECTABLE; INJECTION

BACTOCILL IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

EQ 20MG BASE/ML

N50640 001

Oct 26, 1989

+

EQ 40MG BASE/ML

N50640 002

Oct 26, 1989

OXACILLIN SODIUMAP MARSAM PHARMS LLCEQ 250MG BASE/VIALN62856 001

Oct 26, 1988

APEQ 500MG BASE/VIALN62856 002

Oct 26, 1988

APEQ 1GM BASE/VIALN62856 003

Oct 26, 1988

APEQ 2GM BASE/VIALN62856 004

Oct 26, 1988

APEQ 10GM BASE/VIALN62984 001

Sep 29, 1988

EQ 4GM BASE/VIAL

N62856 005

Oct 26, 1988

AP + SANDOZEQ 250MG BASE/VIALN61490 001AP +EQ 500MG BASE/VIALN61490 002AP +EQ 1GM BASE/VIALN61490 003APEQ 1GM BASE/VIALN62737 001

Dec 23, 1986

AP +EQ 2GM BASE/VIALN61490 004APEQ 2GM BASE/VIALN62737 002

Dec 23, 1986

AP +EQ 10GM BASE/VIALN61490 006

May 09, 1991

OXALIPLATIN

INJECTABLE; IV (INFUSION)

ELOXATIN

+ SANOFI SYNTHELABO

50MG/VIAL

N21492 001

Aug 09, 2002

+

50MG/10ML (5MG/ML)

N21759 001

Jan 31, 2005

+

100MG/VIAL

N21492 002

Aug 09, 2002

+

100MG/20ML (5MG/ML)

N21759 002

Jan 31, 2005

OXANDROLONE

TABLET; ORAL

OXANDRIN

SAVIENT PHARMS

2.5MG

N13718 001

+

10MG

N13718 002

Nov 05, 2001

OXAPROZIN

TABLET; ORAL

DAYPROAB + GD SEARLE LLC600MGN18841 004

Oct 29, 1992

OXAPROZINAB APOTEX600MGN75987 001

Sep 02, 2004

AB

CARACO

600MGN75844 001

Jan 03, 2002

AB

DR REDDYS LABS LTD

600MGN75855 001

Jan 31, 2001

PRESCRIPTION DRUG PRODUCT LIST

3 - 279 (of 360)

OXAPROZIN

TABLET; ORAL

OXAPROZIN

<u>AB</u>	GENPHARM	<u>600MG</u>	<u>N75847</u>	<u>001</u>	Feb 28, 2001
<u>AB</u>	IVAX PHARMS	<u>600MG</u>	<u>N75846</u>	<u>001</u>	May 13, 2002
<u>AB</u>	MYLAN	<u>600MG</u>	<u>N75851</u>	<u>001</u>	Aug 17, 2001
<u>AB</u>	PUREPAC PHARM	<u>600MG</u>	<u>N75843</u>	<u>001</u>	Oct 03, 2001
<u>AB</u>	SANDOZ	<u>600MG</u>	<u>N75842</u>	<u>001</u>	Apr 12, 2001
<u>AB</u>		<u>600MG</u>	<u>N75845</u>	<u>001</u>	Jan 31, 2001
<u>AB</u>		<u>600MG</u>	<u>N75850</u>	<u>001</u>	Apr 27, 2001
<u>AB</u>	TEVA	<u>600MG</u>	<u>N75849</u>	<u>001</u>	Jul 03, 2002
<u>AB</u>	WATSON LABS	<u>600MG</u>	<u>N75848</u>	<u>001</u>	Feb 09, 2001

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

<u>AB</u>	IVAX PHARMS	<u>10MG</u>	<u>N70943</u>	<u>001</u>	Aug 03, 1987
<u>AB</u>		<u>15MG</u>	<u>N70944</u>	<u>001</u>	Aug 03, 1987
<u>AB</u>	+	<u>30MG</u>	<u>N70945</u>	<u>001</u>	Aug 03, 1987
<u>AB</u>	PUREPAC PHARM	<u>10MG</u>	<u>N72251</u>	<u>001</u>	Apr 14, 1988
<u>AB</u>		<u>15MG</u>	<u>N72252</u>	<u>001</u>	Apr 14, 1988
<u>AB</u>		<u>30MG</u>	<u>N72253</u>	<u>001</u>	Apr 14, 1988
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>N71813</u>	<u>001</u>	Apr 19, 1988
<u>AB</u>		<u>15MG</u>	<u>N71756</u>	<u>001</u>	Apr 19, 1988
<u>AB</u>		<u>30MG</u>	<u>N71814</u>	<u>001</u>	Apr 19, 1988
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N72952</u>	<u>001</u>	Sep 28, 1990
<u>AB</u>		<u>15MG</u>	<u>N72953</u>	<u>001</u>	Sep 28, 1990
<u>AB</u>		<u>30MG</u>	<u>N72954</u>	<u>001</u>	Sep 28, 1990

OXCARBAZEPINE

SUSPENSION; ORAL

TRILEPTAL

+	NOVARTIS	300MG/5ML	N21285	001	May 25, 2001
---	----------	-----------	--------	-----	--------------

TABLET; ORAL

TRILEPTAL

	NOVARTIS	150MG	N21014	001	Jan 14, 2000
		300MG	N21014	002	Jan 14, 2000
+		600MG	N21014	003	Jan 14, 2000

OXICONAZOLE NITRATE

CREAM; TOPICAL

OXISTAT

+	ALTANA	EQ 1% BASE	N19828	001	Dec 30, 1988
---	--------	------------	--------	-----	--------------

LOTION; TOPICAL

OXISTAT

+	ALTANA	EQ 1% BASE	N20209	001	Sep 30, 1992
---	--------	------------	--------	-----	--------------

OXTRIPHYLLINE

TABLET, EXTENDED RELEASE; ORAL

CHOLEDYL SA

+	PARKE DAVIS	400MG	N87863	001	May 24, 1983
+	WARNER CHILCOTT	600MG	N86742	001	

OXYBUTYNIN

FILM, EXTENDED RELEASE; TRANSDERMAL

OXYTROL

+	WATSON LABS (UTAH)	3.9MG/24HR	N21351	002	Feb 26, 2003
---	--------------------	------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 280 (of 360)

OXYBUTYNIN CHLORIDE

SYRUP; ORAL

DITROPAN

<u>AA</u>	+ ALZA	<u>5MG/5ML</u>	<u>N18211</u>	<u>001</u>	
<u>OXYBUTYNIN CHLORIDE</u>					
<u>AA</u>	MIKART	<u>5MG/5ML</u>	<u>N75039</u>	<u>001</u>	Jan 29, 1999
<u>AA</u>	MORTON GROVE	<u>5MG/5ML</u>	<u>N74868</u>	<u>001</u>	Feb 12, 1997
<u>AA</u>	NOVEX	<u>5MG/5ML</u>	<u>N74997</u>	<u>001</u>	Oct 15, 1997
<u>AA</u>	PHARM ASSOC	<u>5MG/5ML</u>	<u>N75137</u>	<u>001</u>	Dec 18, 1998
<u>AA</u>	SILARX	<u>5MG/5ML</u>	<u>N74520</u>	<u>001</u>	Mar 29, 1996
<u>AA</u>	VINTAGE PHARMS	<u>5MG/5ML</u>	<u>N76682</u>	<u>001</u>	Dec 28, 2004

TABLET; ORAL

DITROPAN

<u>AB</u>	+ ALZA	<u>5MG</u>	<u>N17577</u>	<u>001</u>	
<u>OXYBUTYNIN CHLORIDE</u>					
<u>AB</u>	PLIVA	<u>5MG</u>	<u>N71655</u>	<u>001</u>	Nov 14, 1988
<u>AB</u>	USL PHARMA	<u>5MG</u>	<u>N74625</u>	<u>001</u>	Jul 31, 1996
<u>AB</u>	VINTAGE PHARMS	<u>5MG</u>	<u>N75079</u>	<u>001</u>	Oct 31, 1997

TABLET, EXTENDED RELEASE; ORAL

DITROPAN XL

	ALZA	5MG	N20897	001	Dec 16, 1998
		10MG	N20897	002	Dec 16, 1998
+		15MG	N20897	003	Jun 22, 1999

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	AMIDE PHARM	<u>15MG</u>	<u>N76636</u>	<u>001</u>	Feb 06, 2004
<u>AB</u>		<u>30MG</u>	<u>N76636</u>	<u>002</u>	Feb 06, 2004
<u>AB</u>	KV PHARM	<u>15MG</u>	<u>N77290</u>	<u>003</u>	Dec 08, 2005
<u>AB</u>		<u>30MG</u>	<u>N77290</u>	<u>005</u>	Dec 08, 2005
		5MG	N77290	001	Dec 08, 2005
		10MG	N77290	002	Dec 08, 2005
		20MG	N77290	004	Dec 08, 2005
<u>AB</u>	MALLINCKRODT	<u>15MG</u>	<u>N76758</u>	<u>001</u>	Jun 30, 2004
<u>AB</u>		<u>30MG</u>	<u>N76758</u>	<u>002</u>	Jun 30, 2004
<u>ROXICODONE</u>					
<u>AB</u>	+ XANODYNE PHARM	<u>15MG</u>	<u>N21011</u>	<u>001</u>	Aug 31, 2000
<u>AB</u>		<u>30MG</u>	<u>N21011</u>	<u>002</u>	Aug 31, 2000

TABLET, EXTENDED RELEASE; ORAL

OXYCODONE HYDROCHLORIDE

<u>AB</u>	ENDO PHARMS	<u>10MG</u>	<u>N75923</u>	<u>001</u>	Mar 23, 2004
<u>AB</u>		<u>20MG</u>	<u>N75923</u>	<u>002</u>	Mar 23, 2004
<u>AB</u>		<u>40MG</u>	<u>N75923</u>	<u>003</u>	Mar 23, 2004
<u>AB</u>		<u>80MG</u>	<u>N75923</u>	<u>004</u>	Sep 29, 2004
<u>AB</u>	IMPAX LABS	<u>80MG</u>	<u>N76318</u>	<u>001</u>	Sep 27, 2004
<u>AB</u>	IMPAX PHARMS	<u>10MG</u>	<u>N76446</u>	<u>001</u>	Dec 06, 2005
<u>AB</u>		<u>20MG</u>	<u>N76446</u>	<u>002</u>	Dec 06, 2005
<u>AB</u>		<u>40MG</u>	<u>N76446</u>	<u>003</u>	Dec 06, 2005
<u>AB</u>	TEVA	<u>10MG</u>	<u>N76610</u>	<u>001</u>	Dec 06, 2005
<u>AB</u>		<u>20MG</u>	<u>N76610</u>	<u>002</u>	Dec 06, 2005
<u>AB</u>		<u>40MG</u>	<u>N76610</u>	<u>003</u>	Dec 06, 2005
<u>AB</u>		<u>80MG</u>	<u>N76168</u>	<u>001</u>	Mar 23, 2004
<u>OXYCONTIN</u>					
<u>AB</u>	PURDUE PHARMA LP	<u>10MG</u>	<u>N20553</u>	<u>001</u>	Dec 12, 1995
<u>AB</u>		<u>20MG</u>	<u>N20553</u>	<u>002</u>	Dec 12, 1995
<u>AB</u>	+	<u>40MG</u>	<u>N20553</u>	<u>003</u>	Dec 12, 1995
<u>AB</u>		<u>80MG</u>	<u>N20553</u>	<u>004</u>	Jan 06, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 281 (of 360)

OXYMETHOLONE

TABLET; ORAL		
ANADROL-50		
+ UNIMED PHARMS	50MG	N16848 001

OXYMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION		
NUMORPHAN		
+ ENDO PHARMS	1MG/ML	N11707 002

OXYTETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL		
TERRAMYCIN		
+ PFIZER	EQ 250MG BASE	N50286 002

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC		
TERRAMYCIN W/ POLYMYXIN B SULFATE		
+ PFIZER	EQ 5MG BASE/GM;10,000 UNITS/GM	N61015 001

OXYTOCIN

INJECTABLE; INJECTION		
<u>OXYTOCIN</u>		
<u>AP</u> + AM PHARM PARTNERS	<u>10USP UNITS/ML</u>	<u>N18248 001</u>
<u>PITOCIN</u>		
<u>AP</u> + PARKEDALE	<u>10USP UNITS/ML</u>	<u>N18261 001</u>

PACLITAXEL

FOR SUSPENSION; IV (INFUSION)		
ABRAXANE		
+ AM BIOSCIENCE	100MG/VIAL	N21660 001 Jan 07, 2005
INJECTABLE; INJECTION		
<u>PACLITAXEL</u>		
<u>AP</u>	BAKER NORTON	<u>6MG/ML</u> <u>N75184 001</u> Jan 25, 2002
<u>AP</u>	BEDFORD	<u>6MG/ML</u> <u>N75190 001</u> Jan 28, 2002
<u>AP</u>	IVAX PHARMS	<u>6MG/ML</u> <u>N75297 001</u> Jan 25, 2002
<u>AP</u>	MAYNE PHARMA USA	<u>6MG/ML</u> <u>N76131 001</u> May 08, 2002
<u>AP</u>		<u>6MG/ML</u> <u>N76233 001</u> Aug 01, 2002
<u>AP</u>	MYLAN	<u>6MG/ML</u> <u>N75278 001</u> Jan 25, 2002
<u>AP</u>	SUPERGEN	<u>6MG/ML</u> <u>N75436 001</u> Nov 12, 2004
<u>TAXOL</u>		
<u>AP</u> +	BRISTOL MYERS SQUIBB	<u>6MG/ML</u> <u>N20262 001</u> Dec 29, 1992

PALONOSETRON HYDROCHLORIDE

INJECTABLE; INTRAVENOUS		
ALOXI		
+ HELSINN HLTHCARE	EQ 0.25MG BASE/5ML	N21372 001 Jul 25, 2003

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION		
<u>AREDIA</u>		
<u>AP</u> +	NOVARTIS	<u>30MG/VIAL</u> <u>N20036 001</u> Oct 31, 1991
<u>AP</u> +		<u>90MG/VIAL</u> <u>N20036 004</u> May 06, 1993
<u>PAMIDRONATE DISODIUM</u>		
<u>AP</u>	AESGEN	<u>30MG/VIAL</u> <u>N75594 001</u> May 06, 2002
<u>AP</u>		<u>90MG/VIAL</u> <u>N75594 002</u> May 06, 2002
<u>AP</u>	AM PHARM PARTNERS	<u>30MG/VIAL</u> <u>N75773 001</u> May 06, 2002
<u>AP</u>		<u>30MG /10ML (3MG/ML)</u> <u>N76207 001</u> May 17, 2002
<u>AP</u>		<u>90MG/VIAL</u> <u>N75773 002</u> May 06, 2002

PRESCRIPTION DRUG PRODUCT LIST

3 - 282 (of 360)

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

<u>AP</u>	AM PHARM PARTNERS	<u>90MG /10ML (9MG/ML)</u>	<u>N76207</u>	<u>002</u>	May 17, 2002
<u>AP</u>	+ BEDFORD	<u>30MG /10ML (3MG/ML)</u>	<u>N21113</u>	<u>001</u>	Mar 04, 2002
<u>AP</u>		<u>30MG/VIAL</u>	<u>N75290</u>	<u>001</u>	Apr 30, 2001
<u>AP</u>	+	<u>90MG /10ML (9MG/ML)</u>	<u>N21113</u>	<u>002</u>	Mar 04, 2002
<u>AP</u>		<u>90MG/VIAL</u>	<u>N75290</u>	<u>003</u>	Apr 30, 2001
<u>AP</u>	+ MAYNE PHARMA USA	<u>30MG/10ML (3MG/ML)</u>	<u>N75841</u>	<u>001</u>	Jun 27, 2002
<u>AP</u>	+	<u>60MG/10ML (6MG/ML)</u>	<u>N75841</u>	<u>002</u>	Jun 27, 2002
<u>AP</u>	+	<u>90MG/10ML (9MG/ML)</u>	<u>N75841</u>	<u>003</u>	Jun 27, 2002
<u>AP</u>	SICOR PHARMS	<u>30MG /10ML (3MG/ML)</u>	<u>N76153</u>	<u>001</u>	Mar 27, 2002
<u>AP</u>		<u>90MG /10ML (9MG/ML)</u>	<u>N76153</u>	<u>002</u>	Mar 27, 2002

PANCURONIUM BROMIDE

INJECTABLE; INJECTION

PANCURONIUM

<u>AP</u>	ELKINS SINN	<u>1MG/ML</u>	<u>N72058</u>	<u>001</u>	Mar 23, 1988
<u>AP</u>		<u>2MG/ML</u>	<u>N72059</u>	<u>001</u>	Mar 23, 1988
<u>AP</u>		<u>2MG/ML</u>	<u>N72060</u>	<u>001</u>	Mar 23, 1988

PANCURONIUM BROMIDE

<u>AP</u>	HOSPIRA	<u>1MG/ML</u>	<u>N72320</u>	<u>001</u>	Jan 19, 1989
<u>AP</u>		<u>2MG/ML</u>	<u>N72321</u>	<u>001</u>	Jan 19, 1989
<u>AP</u>	+ SICOR PHARMS	<u>1MG/ML</u>	<u>N72759</u>	<u>001</u>	Jul 31, 1990
<u>AP</u>	+	<u>2MG/ML</u>	<u>N72760</u>	<u>001</u>	Jul 31, 1990

PANTOPRAZOLE SODIUM

INJECTABLE; IV (INFUSION)

PROTONIX IV

+	WYETH PHARMS INC	EQ 40MG BASE/VIAL	N20988	001	Mar 22, 2001
---	------------------	-------------------	--------	-----	--------------

TABLET, DELAYED RELEASE; ORAL

PROTONIX

	WYETH PHARMS INC	EQ 20MG BASE	N20987	002	Jun 12, 2001
+		EQ 40MG BASE	N20987	001	Feb 02, 2000

PARICALCITOL

CAPSULE; ORAL

ZEMPLAR

	ABBOTT	1UGM	N21606	001	May 26, 2005
		2UGM	N21606	002	May 26, 2005
+		4UGM	N21606	003	May 26, 2005

INJECTABLE; INJECTION

ZEMPLAR

	ABBOTT	0.002MG/ML	N20819	002	Feb 01, 2000
+		0.005MG/ML	N20819	001	Apr 17, 1998

PAROMOMYCIN SULFATE

CAPSULE; ORAL

HUMATIN

<u>AA</u>	+ KING PHARMS	<u>EQ 250MG BASE</u>	<u>N62310</u>	<u>001</u>	
<u>AA</u>	CARACO	<u>EQ 250MG BASE</u>	<u>N64171</u>	<u>001</u>	Jun 30, 1997

PAROXETINE HYDROCHLORIDE

SUSPENSION; ORAL

PAXIL

+	GLAXOSMITHKLINE	EQ 10MG BASE/5ML	N20710	001	Jun 25, 1997
---	-----------------	------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 283 (of 360)

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

<u>AB</u>	ALPHAPHARM	<u>EQ 10MG BASE</u>	<u>N75716</u>	<u>001</u>	Mar 08, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75716</u>	<u>002</u>	Mar 08, 2004
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>N75716</u>	<u>003</u>	Mar 08, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N75716</u>	<u>004</u>	Mar 08, 2004
<u>AB</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>N75566</u>	<u>001</u>	Mar 08, 2004
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75566</u>	<u>002</u>	Mar 08, 2004
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>N75566</u>	<u>003</u>	Mar 08, 2004
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N75566</u>	<u>004</u>	Mar 08, 2004
<u>AB</u>	TEVA	<u>EQ 10MG BASE</u>	<u>N76618</u>	<u>001</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N76618</u>	<u>002</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>N76618</u>	<u>003</u>	Aug 15, 2005
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N76618</u>	<u>004</u>	Aug 15, 2005
<u>AB</u>	TORPHARM	<u>EQ 10MG BASE</u>	<u>N75356</u>	<u>001</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N75356</u>	<u>002</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>N75356</u>	<u>003</u>	Jul 30, 2003
<u>AB</u>		<u>EQ 40MG BASE</u>	<u>N75356</u>	<u>004</u>	Jul 30, 2003
	<u>PAXIL</u>				
<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 10MG BASE</u>	<u>N20031</u>	<u>001</u>	Dec 29, 1992
<u>AB</u>		<u>EQ 20MG BASE</u>	<u>N20031</u>	<u>002</u>	Dec 29, 1992
<u>AB</u>		<u>EQ 30MG BASE</u>	<u>N20031</u>	<u>003</u>	Dec 29, 1992
<u>AB</u>	+	<u>EQ 40MG BASE</u>	<u>N20031</u>	<u>005</u>	Dec 29, 1992

TABLET, EXTENDED RELEASE; ORAL

PAXIL CR

	GLAXOSMITHKLINE	EQ 12.5MG BASE	N20936	001	Feb 16, 1999
		EQ 25MG BASE	N20936	002	Feb 16, 1999
	+	EQ 37.5MG BASE	N20936	003	Dec 06, 2000

PAROXETINE MESYLATE

TABLET; ORAL

PEXEVA

	SYNTHON PHARMS	EQ 10MG BASE	N21299	001	Jul 03, 2003
		EQ 20MG BASE	N21299	002	Jul 03, 2003
		EQ 30MG BASE	N21299	003	Jul 03, 2003
	+	EQ 40MG BASE	N21299	004	Jul 03, 2003

PEGADEMASE BOVINE

INJECTABLE; INJECTION

ADAGEN

	+ ENZON	250 UNITS/ML	N19818	001	Mar 21, 1990
--	---------	--------------	--------	-----	--------------

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN

	+ EYETECH PHARMS	EQ 0.3MG ACID/0.09ML	N21756	001	Dec 17, 2004
--	------------------	----------------------	--------	-----	--------------

PEGVISOMANT

INJECTABLE; SUBCUTANEOUS

SOMAVERT

	+ PHARMACIA AND UPJOHN	10MG/VIAL	N21106	001	Mar 25, 2003
	+	15MG/VIAL	N21106	002	Mar 25, 2003
	+	20MG/VIAL	N21106	003	Mar 25, 2003

PEMETREXED DISODIUM

INJECTABLE; IV (INFUSION)

ALIMTA

	+ LILLY	EQ 500MG BASE/VIAL	N21462	001	Feb 04, 2004
--	---------	--------------------	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 284 (of 360)

PEMIROLAST POTASSIUM

SOLUTION/DROPS; OPHTHALMIC
ALAMAST

+ SANTEN	0.1%	N21079 001	Sep 24, 1999
----------	------	------------	--------------

PENBUTOLOL SULFATE

TABLET; ORAL
LEVATOL

+ SCHWARZ PHARMA	20MG	N18976 004	Jan 05, 1989
------------------	------	------------	--------------

PENCICLOVIR SODIUM

CREAM; TOPICAL
DENA VIR

+ NOVARTIS	1%	N20629 001	Sep 24, 1996
------------	----	------------	--------------

PENICILLAMINE

CAPSULE; ORAL
CUPRIMINE

MERCK	125MG	N19853 002	
+	250MG	N19853 001	

TABLET; ORAL
DEPEN

+ MEDPOINTE PHARM HLC	250MG	N19854 001	
-----------------------	-------	------------	--

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION
BICILLIN L-A

BC	+ KING PHARMS	600,000 UNITS/ML	N50141 001	
	+	300,000 UNITS/ML	N50141 003	
	PERMAPEN			
BC	PFIZER	600,000 UNITS/ML	N60014 001	

PENICILLIN G BENZATHINE; PENICILLIN G PROCAINE

INJECTABLE; INJECTION
BICILLIN C-R

+ KING PHARMS	150,000 UNITS/ML;150,000 UNITS/ML	N50138 002	
+	300,000 UNITS/ML;300,000 UNITS/ML	N50138 001	
	BICILLIN C-R 900/300		
+ KING PHARMS	900,000 UNITS/2ML;300,000 UNITS/2ML	N50138 003	

PENICILLIN G POTASSIUM

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM

<u>AP</u>	PFIZER	<u>20,000,000 UNITS/VIAL</u>	<u>N60074 003</u>	
<u>AP</u>	SANDOZ	<u>1,000,000 UNITS/VIAL</u>	<u>N65079 001</u>	Aug 30, 2002
<u>AP</u>		<u>5,000,000 UNITS/VIAL</u>	<u>N65079 002</u>	Aug 30, 2002
<u>AP</u>		<u>20,000,000 UNITS/VIAL</u>	<u>N65079 003</u>	Aug 30, 2002
	PENICILLIN G POTASSIUM IN PLASTIC CONTAINER			
	+ BAXTER HLTHCARE	20,000 UNITS/ML	N50638 001	Jun 25, 1990
	+	40,000 UNITS/ML	N50638 002	Jun 25, 1990
	+	60,000 UNITS/ML	N50638 003	Jun 25, 1990
	<u>PFIZERPEN</u>			
<u>AP</u>	+ PFIZER	<u>1,000,000 UNITS/VIAL</u>	<u>N60657 001</u>	
<u>AP</u>	+	<u>5,000,000 UNITS/VIAL</u>	<u>N60657 002</u>	
<u>AP</u>	+	<u>20,000,000 UNITS/VIAL</u>	<u>N60657 003</u>	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 285 (of 360)

PENICILLIN G PROCAINEINJECTABLE; INJECTION
PENICILLIN G PROCAINE

+ KING PHARMS	300,000 UNITS/ML	N60101	002	
+	600,000 UNITS/ML	N60101	001	

PENICILLIN G SODIUMINJECTABLE; IM-IV
PENICILLIN G SODIUM

+ BIOCHEMIE	5,000,000 UNITS/VIAL	N65068	001	Feb 26, 2001
-------------	----------------------	--------	-----	--------------

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN V POTASSIUM

<u>AA</u>	CLONMEL	<u>EQ 125MG BASE/5ML</u>	<u>N62981</u>	<u>001</u>	Feb 10, 1989
<u>AA</u>		<u>EQ 250MG BASE/5ML</u>	<u>N62981</u>	<u>002</u>	Feb 10, 1989
<u>AA</u>	CONSOLIDATED PHARM	<u>EQ 125MG BASE/5ML</u>	<u>N61529</u>	<u>001</u>	
<u>AA</u>		<u>EQ 250MG BASE/5ML</u>	<u>N61529</u>	<u>002</u>	

PENICILLIN-VK

<u>AA</u>	TEVA	<u>EQ 125MG BASE/5ML</u>	<u>N60456</u>	<u>001</u>	
<u>AA</u>		<u>EQ 250MG BASE/5ML</u>	<u>N60456</u>	<u>002</u>	

VEETIDS

<u>AA</u>	APOTHECON	<u>EQ 125MG BASE/5ML</u>	<u>N61410</u>	<u>001</u>	
<u>AA</u>		<u>EQ 250MG BASE/5ML</u>	<u>N61410</u>	<u>002</u>	

TABLET; ORAL

PENICILLIN V POTASSIUM

<u>AB</u>	BIOCHEMIE	<u>EQ 250MG BASE</u>	<u>N64071</u>	<u>001</u>	Nov 30, 1995
<u>AB</u>	+	<u>EQ 500MG BASE</u>	<u>N64071</u>	<u>002</u>	Nov 30, 1995
<u>AB</u>	CLONMEL	<u>EQ 250MG BASE</u>	<u>N62936</u>	<u>001</u>	Nov 25, 1988
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N62935</u>	<u>001</u>	Nov 23, 1988

PENICILLIN-VK

<u>AB</u>	TEVA	<u>EQ 250MG BASE</u>	<u>N60711</u>	<u>002</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N60711</u>	<u>003</u>	

VEETIDS

<u>AB</u>	APOTHECON	<u>EQ 250MG BASE</u>	<u>N61411</u>	<u>001</u>	
<u>AB</u>		<u>EQ 500MG BASE</u>	<u>N61411</u>	<u>002</u>	

PENTAMIDINE ISETHIONATEFOR SOLUTION; INHALATION
NEBUPENT

+ AM PHARM PARTNERS	300MG/VIAL	N19887	001	Jun 15, 1989
	600MG/VIAL	N19887	002	Mar 22, 1996

INJECTABLE; INJECTION

PENTAM

<u>AP</u>	+	AM PHARM PARTNERS	<u>300MG/VIAL</u>	<u>N19264</u>	<u>001</u>	Oct 16, 1984
-----------	---	-------------------	-------------------	---------------	------------	--------------

PENTAMIDINE ISETHIONATE

<u>AP</u>	HOSPIRA	<u>300MG/VIAL</u>	<u>N73479</u>	<u>001</u>	Jun 30, 1992
<u>AP</u>	STERIS	<u>300MG/VIAL</u>	<u>N74303</u>	<u>001</u>	Aug 17, 1995

PENTAZOCINE LACTATEINJECTABLE; INJECTION
TALWIN

+ HOSPIRA	EQ 30MG BASE/ML	N16194	001	
-----------	-----------------	--------	-----	--

PENTETATE CALCIUM TRISODIUMSOLUTION; INHALATION, INTRAVENOUS
PENTETATE CALCIUM TRISODIUM

+ HAMELN PHARMS	EQ 1GM/5ML(EQ 200MG/ML)	N21749	001	Aug 11, 2004
-----------------	-------------------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 286 (of 360)

PENTETATE ZINC TRISODIUM

SOLUTION; INHALATION, INTRAVENOUS

PENTETATE ZINC TRISODIUM

+ HAMELN PHARMS EQ 1GM/5ML(EQ 200MG/ML)

N21751 001 Aug 11, 2004

PENTOBARBITAL SODIUM

INJECTABLE; INJECTION

NEMBUTAL SODIUM

+ OVATION PHARMS 50MG/ML

N83246 001

PENTOSAN POLYSULFATE SODIUM

CAPSULE; ORAL

ELMIRON

+ ORTHO MCNEIL PHARM 100MG

N20193 001 Sep 26, 1996

PENTOSTATIN

INJECTABLE; INJECTION

NIPENT

+ SUPERGEN 10MG/VIAL

N20122 001 Oct 11, 1991

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

<u>AB</u>	BIOVAIL	<u>400MG</u>	<u>N75028</u>	<u>001</u>	Jul 20, 1998
<u>AB</u>	CLONMEL HLTHCARE	<u>400MG</u>	<u>N74877</u>	<u>001</u>	Jul 08, 1997
<u>AB</u>	IMPAX LABS	<u>400MG</u>	<u>N75093</u>	<u>001</u>	Aug 10, 1999
<u>AB</u>	MYLAN	<u>400MG</u>	<u>N74425</u>	<u>001</u>	Jul 08, 1997
<u>AB</u>	PLIVA	<u>400MG</u>	<u>N74874</u>	<u>001</u>	May 25, 1999
<u>AB</u>	PUREPAC PHARM	<u>400MG</u>	<u>N74878</u>	<u>001</u>	Jul 09, 1997
<u>AB</u>	TEVA	<u>400MG</u>	<u>N75199</u>	<u>001</u>	Sep 03, 1999
<u>AB</u>	TORPHARM	<u>400MG</u>	<u>N75191</u>	<u>001</u>	Jun 09, 1999
<u>AB</u>	WATSON LABS	<u>400MG</u>	<u>N75107</u>	<u>001</u>	Sep 04, 1998

PENTOXIL

<u>AB</u>	UPSHER SMITH	<u>400MG</u>	<u>N74962</u>	<u>001</u>	Mar 31, 1999
-----------	--------------	--------------	---------------	------------	--------------

TRENTAL

<u>AB</u>	+ AVENTIS PHARMS	<u>400MG</u>	<u>N18631</u>	<u>001</u>	Aug 30, 1984
-----------	------------------	--------------	---------------	------------	--------------

PERFLUTREN

INJECTABLE; INTRAVENOUS

DEFINITY

+ BRISTOL MYERS SQUIBB 6.52MG/ML

N21064 001 Jul 31, 2001

PERGOLIDE MESYLATE

TABLET; ORAL

PERGOLIDE MESYLATE

<u>AB</u>	IVAX PHARMS	<u>EQ 0.05MG BASE</u>	<u>N76094</u>	<u>001</u>	Sep 04, 2003
<u>AB</u>		<u>EQ 0.25MG BASE</u>	<u>N76094</u>	<u>002</u>	Sep 04, 2003
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76094</u>	<u>003</u>	Sep 04, 2003
<u>AB</u>	TEVA	<u>EQ 0.05MG BASE</u>	<u>N76061</u>	<u>001</u>	Nov 27, 2002
<u>AB</u>		<u>EQ 0.25MG BASE</u>	<u>N76061</u>	<u>002</u>	Nov 27, 2002
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N76061</u>	<u>003</u>	Nov 27, 2002
<u>PERMAX</u>					
<u>AB</u>	+ VALEANT	<u>EQ 0.05MG BASE</u>	<u>N19385</u>	<u>001</u>	Dec 30, 1988
<u>AB</u>		<u>EQ 0.25MG BASE</u>	<u>N19385</u>	<u>002</u>	Dec 30, 1988
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N19385</u>	<u>003</u>	Dec 30, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 287 (of 360)

PERINDOPRIL ERBUMINE

TABLET; ORAL

ACEON

SOLVAY PHARMS

2MG

N20184 001

Dec 30, 1993

4MG

N20184 002

Dec 30, 1993

+

8MG

N20184 003

Dec 30, 1993

PERMETHRIN

CREAM; TOPICAL

ELIMITEAB + ALLERGAN5%N19855 001

Aug 25, 1989

PERMETHRINAB ALPHARMA US PHARMS5%N74806 001

Jan 23, 1998

AB CLAY PARK5%N76369 001

Apr 21, 2003

PERPHENAZINE

CONCENTRATE; ORAL

PERPHENAZINE

+ PHARM ASSOC

16MG/5ML

N40360 001

May 25, 2001

TABLET; ORAL

PERPHENAZINEAB IVAX PHARMS2MGN89707 001

Sep 10, 1987

AB4MGN89708 001

Sep 10, 1987

AB8MGN89456 001

Sep 10, 1987

AB +16MGN89457 001

Sep 10, 1987

AB

SANDOZ

2MGN89683 001

Dec 08, 1988

AB4MGN89684 001

Dec 08, 1988

AB8MGN89685 001

Dec 08, 1988

AB16MGN89686 001

Dec 08, 1988

AB

VINTAGE PHARMS

2MGN40226 001

Dec 31, 1998

AB4MGN40226 002

Dec 31, 1998

AB8MGN40226 003

Dec 31, 1998

AB16MGN40226 004

Dec 31, 1998

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL

PHENDIMETRAZINE TARTRATEAA + SANDOZ35MGN85695 001X-TROZINEAA SHIRE RICHWOOD35MGN87394 001

Sep 22, 1982

CAPSULE, EXTENDED RELEASE; ORAL

BONTRIL

BC MALLINCKRODT

105MG

N88021 001

Sep 21, 1982

PHENDIMETRAZINE TARTRATE

BC + SANDOZ

105MG

N18074 001

X-TROZINE L.A.

BC SHIRE RICHWOOD

105MG

N87371 001

Aug 24, 1982

TABLET; ORAL

BONTRIL PDMAA + VALEANT35MGN85272 001PHENDIMETRAZINE TARTRATEAA MIKART35MGN89452 001

Oct 30, 1991

AA SANDOZ35MGN85588 001PHENELZINE SULFATE

TABLET; ORAL

NARDIL

+ PARKE DAVIS

EQ 15MG BASE

N11909 002

PRESCRIPTION DRUG PRODUCT LIST

3 - 288 (of 360)

PHENOXYBENZAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIBENZYLINE

+ WELLSPRING PHARM	10MG	N08708	001	
--------------------	------	--------	-----	--

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

ADIPEX-P

<u>AA</u> + TEVA	<u>37.5MG</u>	<u>N88023</u>	<u>001</u>	Aug 02, 1983
<u>PHENTERMINE HYDROCHLORIDE</u>				
<u>AA</u> AMBI PHARMS	<u>30MG</u>	<u>N40083</u>	<u>001</u>	Mar 07, 1997
<u>AA</u> AMIDE PHARM	<u>15MG</u>	<u>N40460</u>	<u>001</u>	Jan 14, 2003
<u>AA</u>	<u>30MG</u>	<u>N40227</u>	<u>001</u>	Jun 18, 1997
<u>AA</u>	<u>30MG</u>	<u>N40448</u>	<u>001</u>	Jan 22, 2003
<u>AA</u>	<u>37.5MG</u>	<u>N40228</u>	<u>001</u>	Jun 19, 1997
<u>AA</u> MUTUAL PHARM	<u>30MG</u>	<u>N40525</u>	<u>001</u>	Oct 23, 2003
<u>AA</u>	<u>37.5MG</u>	<u>N40527</u>	<u>001</u>	Oct 23, 2003
<u>AA</u> + SANDOZ	<u>15MG</u>	<u>N87301</u>	<u>001</u>	
<u>AA</u> +	<u>30MG</u>	<u>N86945</u>	<u>001</u>	Jul 20, 1983
<u>AA</u>	<u>30MG</u>	<u>N87190</u>	<u>001</u>	
<u>AA</u>	<u>30MG</u>	<u>N87208</u>	<u>001</u>	
<u>AA</u>	<u>30MG</u>	<u>N87223</u>	<u>001</u>	
<u>AA</u> VINTAGE PHARMS	<u>37.5MG</u>	<u>N40377</u>	<u>001</u>	Jan 04, 2002

TABLET; ORAL

ADIPEX-P

<u>AA</u> + TEVA	<u>37.5MG</u>	<u>N85128</u>	<u>001</u>	
<u>PHENTERMINE HYDROCHLORIDE</u>				
<u>AA</u> AMIDE PHARM	<u>37.5MG</u>	<u>N40190</u>	<u>001</u>	May 30, 1997
<u>AA</u> LANNETT	<u>37.5MG</u>	<u>N40555</u>	<u>001</u>	Apr 15, 2005
<u>AA</u> MUTUAL PHARM	<u>37.5MG</u>	<u>N40526</u>	<u>001</u>	Oct 23, 2003
<u>AA</u> PUREPAC PHARM	<u>37.5MG</u>	<u>N40276</u>	<u>001</u>	Nov 25, 1998
+ SANDOZ	30MG	N88605	001	Sep 28, 1987

PHENTERMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL

IONAMIN

UCB	EQ 15MG BASE	N11613	004	
+	EQ 30MG BASE	N11613	002	

PHENTOLAMINE MESYLATE

INJECTABLE; INJECTION

PHENTOLAMINE MESYLATE

<u>AP</u> BEDFORD	<u>5MG/VIAL</u>	<u>N40235</u>	<u>001</u>	Mar 11, 1998
<u>REGITINE</u>				
<u>AP</u> + NOVARTIS	<u>5MG/VIAL</u>	<u>N08278</u>	<u>003</u>	

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH VC PLAIN

<u>AA</u> + ALPHARMA US PHARMS	<u>5MG/5ML; 6.25MG/5ML</u>	<u>N88761</u>	<u>001</u>	Nov 08, 1984
<u>PROMETHAZINE VC PLAIN</u>				
<u>AA</u> MORTON GROVE	<u>5MG/5ML; 6.25MG/5ML</u>	<u>N88897</u>	<u>001</u>	Jan 04, 1985

PHENYTOIN

SUSPENSION; ORAL

DILANTIN-125

<u>AB</u> + PARKE DAVIS	<u>125MG/5ML</u>	<u>N08762</u>	<u>001</u>	
<u>PHENTYTOIN</u>				
<u>AB</u> MORTON GROVE	<u>125MG/5ML</u>	<u>N40420</u>	<u>001</u>	Apr 19, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 289 (of 360)

PHENYTOIN

SUSPENSION; ORAL

PHENYTOIN

<u>AB</u>	ALPHARMA US PHARMS	<u>125MG/5ML</u>	<u>N89892</u>	<u>001</u>	Sep 25, 1992
<u>AB</u>	TARO	<u>125MG/5ML</u>	<u>N40521</u>	<u>001</u>	Mar 08, 2004
<u>AB</u>	VISTAPHARM	<u>125MG/5ML</u>	<u>N40342</u>	<u>001</u>	Jan 31, 2001
<u>AB</u>		<u>125MG/5ML</u>	<u>N40610</u>	<u>001</u>	Aug 18, 2005

TABLET, CHEWABLE; ORAL
DILANTIN

+ PFIZER PHARMS 50MG N84427 001

PHENYTOIN SODIUM

CAPSULE; ORAL

DILANTIN

<u>AB</u>	+ PARKE DAVIS	<u>100MG EXTENDED</u>	<u>N84349</u>	<u>002</u>	
	+	30MG EXTENDED	N84349	001	

EXTENDED PHENYTOIN SODIUM

<u>AB</u>	BARR	<u>100MG EXTENDED</u>	<u>N40435</u>	<u>001</u>	Jun 20, 2003
<u>AB</u>	MYLAN	<u>100MG EXTENDED</u>	<u>N40298</u>	<u>001</u>	Dec 28, 1998
	PHENYTEK				
	+ MYLAN	200MG EXTENDED	N40298	002	Dec 06, 2001
	+	300MG EXTENDED	N40298	003	Dec 06, 2001
	PROMPT PHENYTOIN SODIUM				
BX	+ IVAX PHARMS	100MG PROMPT	N80259	001	

INJECTABLE; INJECTION

PHENYTOIN

<u>AP</u>	+ ELKINS SINN	<u>50MG/ML</u>	<u>N84307</u>	<u>001</u>	
-----------	---------------	----------------	---------------	------------	--

PHENYTOIN SODIUM

<u>AP</u>	HOSPIRA	<u>50MG/ML</u>	<u>N89521</u>	<u>001</u>	Mar 17, 1987
<u>AP</u>		<u>50MG/ML</u>	<u>N89744</u>	<u>001</u>	Dec 18, 1987

PHYTONADIONE

INJECTABLE; INJECTION

AQUAMEPHYTON

BP	+ MERCK	1MG/0.5ML	N12223	002	
BP	+	10MG/ML	N12223	001	
	PHYTONADIONE				
BP	INTL MEDICATION	1MG/0.5ML	N83722	001	
	VITAMIN K1				
BP	HOSPIRA	1MG/0.5ML	N87954	001	Jul 25, 1983
BP		10MG/ML	N87955	001	Jul 25, 1983

TABLET; ORAL

MEPHYTON

+ MERCK 5MG N10104 003

PILOCARPINE HYDROCHLORIDE

GEL; OPHTHALMIC

PILOPINE HS

+ ALCON 4% N18796 001 Oct 01, 1984

TABLET; ORAL

PILOCARPINE HYDROCHLORIDE

<u>AB</u>	COREPHARMA	<u>5MG</u>	<u>N76746</u>	<u>001</u>	Nov 16, 2004
<u>AB</u>	LANNETT	<u>5MG</u>	<u>N77220</u>	<u>001</u>	Oct 14, 2005
<u>AB</u>	ROXANE	<u>5MG</u>	<u>N76963</u>	<u>001</u>	Dec 22, 2004
	<u>SALAGEN</u>				
<u>AB</u>	MGI PHARMA INC	<u>5MG</u>	<u>N20237</u>	<u>001</u>	Mar 22, 1994
	+	7.5MG	N20237	002	Apr 18, 2003

PRESCRIPTION DRUG PRODUCT LIST

3 - 290 (of 360)

PIMECROLIMUS

CREAM; TOPICAL
ELIDEL

+ NOVARTIS	1%	N21302	001	Dec 13, 2001
------------	----	--------	-----	--------------

PIMOZIDE

TABLET; ORAL
ORAP

TEVA	1MG	N17473	003	Aug 27, 1997
+	2MG	N17473	001	Jul 31, 1984

PINDOLOL

TABLET; ORAL

PINDOLOL

<u>AB</u> GENPHARM	<u>5MG</u>	<u>N74013</u>	<u>001</u>	Sep 24, 1992
<u>AB</u>	<u>10MG</u>	<u>N74018</u>	<u>001</u>	Sep 24, 1992
<u>AB</u> IVAX PHARMS	<u>5MG</u>	<u>N73687</u>	<u>001</u>	Feb 26, 1993
<u>AB</u>	<u>10MG</u>	<u>N73687</u>	<u>002</u>	Feb 26, 1993
<u>AB</u> MARTEC	<u>5MG</u>	<u>N74474</u>	<u>001</u>	Oct 28, 1996
<u>AB</u>	<u>10MG</u>	<u>N74474</u>	<u>002</u>	Oct 28, 1996
<u>AB</u> MUTUAL PHARM	<u>5MG</u>	<u>N74063</u>	<u>001</u>	Jan 27, 1994
<u>AB</u>	<u>10MG</u>	<u>N74063</u>	<u>002</u>	Jan 27, 1994
<u>AB</u> MYLAN	<u>5MG</u>	<u>N74019</u>	<u>001</u>	Sep 03, 1992
<u>AB</u>	<u>10MG</u>	<u>N74019</u>	<u>002</u>	Sep 03, 1992
<u>AB</u> SANDOZ	<u>5MG</u>	<u>N73608</u>	<u>001</u>	Mar 29, 1993
<u>AB</u>	<u>10MG</u>	<u>N73609</u>	<u>001</u>	Mar 29, 1993
<u>AB</u> TEVA	<u>5MG</u>	<u>N73661</u>	<u>001</u>	Oct 31, 1993
<u>AB</u>	<u>5MG</u>	<u>N74123</u>	<u>001</u>	Apr 17, 1997
<u>AB</u>	<u>10MG</u>	<u>N73661</u>	<u>002</u>	Oct 31, 1993
<u>AB</u>	<u>10MG</u>	<u>N74123</u>	<u>002</u>	Apr 17, 1997
<u>AB</u> WATSON LABS	<u>5MG</u>	<u>N74437</u>	<u>001</u>	Feb 27, 1995
<u>AB</u>	<u>10MG</u>	<u>N74437</u>	<u>002</u>	Feb 27, 1995
<u>VISKEN</u>				
<u>AB</u> NOVARTIS	<u>5MG</u>	<u>N18285</u>	<u>001</u>	Sep 03, 1982
<u>AB</u> +	<u>10MG</u>	<u>N18285</u>	<u>002</u>	Sep 03, 1982

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL
ACTOS

TAKEDA PHARMS NA	EQ 15MG BASE	N21073	001	Jul 15, 1999
	EQ 30MG BASE	N21073	002	Jul 15, 1999
+	EQ 45MG BASE	N21073	003	Jul 15, 1999

PIPERACILLIN SODIUM

INJECTABLE; INJECTION
PIPERACILLIN

+ AM PHARM	EQ 2GM BASE/VIAL	N65114	001	Nov 14, 2003
+	EQ 3GM BASE/VIAL	N65114	002	Nov 14, 2003
+	EQ 4GM BASE/VIAL	N65114	003	Nov 14, 2003
+	EQ 40GM BASE/VIAL	N65157	001	Jul 12, 2004

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION
ZOSYN

+ WYETH PHARMS INC	EQ 2GM BASE/VIAL;EQ 250MG BASE/VIAL	N50684	001	Oct 22, 1993
+	EQ 3GM BASE/VIAL;EQ 375MG BASE/VIAL	N50684	002	Oct 22, 1993
+	EQ 4GM BASE/VIAL;EQ 500MG BASE/VIAL	N50684	003	Oct 22, 1993
+	EQ 36GM BASE/VIAL;EQ 4.5GM BASE/VIAL	N50684	004	Oct 22, 1993

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 291 (of 360)

PIPERACILLIN SODIUM; TAZOBACTAM SODIUM

INJECTABLE; INJECTION

ZOSYN IN PLASTIC CONTAINER

+	WYETH PHARMS INC	EQ 40MG BASE/ML;EQ 5MG BASE/ML	N50750	001	Feb 24, 1998
+		EQ 60MG BASE/ML;EQ 7.5MG BASE/ML	N50750	002	Feb 24, 1998
+		EQ 4GM BASE/100ML;EQ 500MG BASE/100ML	N50750	003	Feb 24, 1998

PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION

MAXAIR

+	3M	EQ 0.2MG BASE/INH	N20014	001	Nov 30, 1992
---	----	-------------------	--------	-----	--------------

PIROXICAM

CAPSULE; ORAL

FELDENE

<u>AB</u>	PFIZER	<u>10MG</u>	<u>N18147</u>	<u>002</u>	Apr 06, 1982
<u>AB</u>	+	<u>20MG</u>	<u>N18147</u>	<u>003</u>	Apr 06, 1982

PIROXICAM

<u>AB</u>	EGIS	<u>10MG</u>	<u>N74808</u>	<u>001</u>	Jul 08, 1997
<u>AB</u>		<u>20MG</u>	<u>N74808</u>	<u>002</u>	Jul 08, 1997
<u>AB</u>	GENPHARM	<u>10MG</u>	<u>N74043</u>	<u>001</u>	Sep 22, 1992
<u>AB</u>		<u>20MG</u>	<u>N74043</u>	<u>002</u>	Sep 22, 1992
<u>AB</u>	MEPHA	<u>10MG</u>	<u>N74116</u>	<u>001</u>	Jun 15, 1993
<u>AB</u>		<u>20MG</u>	<u>N74118</u>	<u>001</u>	Jun 15, 1993
<u>AB</u>	MUTUAL PHARM	<u>10MG</u>	<u>N73535</u>	<u>001</u>	Mar 12, 1993
<u>AB</u>		<u>20MG</u>	<u>N73536</u>	<u>001</u>	Mar 12, 1993
<u>AB</u>	MYLAN	<u>10MG</u>	<u>N74102</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>		<u>20MG</u>	<u>N74102</u>	<u>002</u>	Jul 31, 1992
<u>AB</u>	TEVA	<u>10MG</u>	<u>N73637</u>	<u>001</u>	Jan 28, 1994
<u>AB</u>		<u>10MG</u>	<u>N74131</u>	<u>001</u>	Dec 11, 1992
<u>AB</u>		<u>20MG</u>	<u>N73638</u>	<u>001</u>	Jan 28, 1994
<u>AB</u>		<u>20MG</u>	<u>N74131</u>	<u>002</u>	Dec 11, 1992
<u>AB</u>	TEVA PHARMS	<u>10MG</u>	<u>N74103</u>	<u>001</u>	Aug 28, 1992
<u>AB</u>		<u>20MG</u>	<u>N74103</u>	<u>002</u>	Aug 28, 1992
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N74287</u>	<u>001</u>	May 16, 1996
<u>AB</u>		<u>10MG</u>	<u>N74460</u>	<u>001</u>	Sep 29, 1995
<u>AB</u>		<u>20MG</u>	<u>N74287</u>	<u>002</u>	May 16, 1996
<u>AB</u>		<u>20MG</u>	<u>N74460</u>	<u>002</u>	Sep 29, 1995

PLICAMYCIN

INJECTABLE; INJECTION

MITHRACIN

+	PFIZER	2.5MG/VIAL	N50109	001	
---	--------	------------	--------	-----	--

PODOFILOX

GEL; TOPICAL

CONDYLOX

+	OCLASSEN	0.5%	N20529	001	Mar 13, 1997
---	----------	------	--------	-----	--------------

SOLUTION; TOPICAL

CONDYLOX

<u>AT</u>	+	OCLASSEN	<u>0.5%</u>	<u>N19795</u>	<u>001</u>	Dec 13, 1990
-----------	---	----------	-------------	---------------	------------	--------------

PODOFILOX

<u>AT</u>	PADDOCK	<u>0.5%</u>	<u>N75600</u>	<u>001</u>	Jan 29, 2002
-----------	---------	-------------	---------------	------------	--------------

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

GLYCOLAX

<u>AA</u>	SCHWARZ PHARMA	<u>17GM/SC00PFUL</u>	<u>N76652</u>	<u>001</u>	Jul 02, 2004
-----------	----------------	----------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 292 (of 360)

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL

MIRALAX

<u>AA</u>	+	BRAINTREE	<u>17GM/SC00PFUL</u>	<u>N20698</u>	<u>001</u>	Feb 18, 1999
-----------	---	-----------	----------------------	---------------	------------	--------------

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL

NULYTELY

<u>AA</u>	+	BRAINTREE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N19797</u>	<u>001</u>	Apr 22, 1991
-----------	---	-----------	---	---------------	------------	--------------

NULYTELY-FLAVORED

<u>AA</u>	+	BRAINTREE	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N19797</u>	<u>002</u>	Nov 18, 1994
-----------	---	-----------	---	---------------	------------	--------------

TRILYTE

<u>AA</u>		SCHWARZ PHARMA	<u>420GM/BOT;1.48GM/BOT;5.72GM/BOT;11.2GM/BOT</u>	<u>N76491</u>	<u>001</u>	Feb 05, 2004
-----------	--	----------------	---	---------------	------------	--------------

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE, ANHYDROUS

FOR SOLUTION; ORAL

COLYTE

<u>AA</u>	+	SCHWARZ PHARMA	<u>227.1GM/BOT;2.82GM/BOT;6.36GM/BOT;5.53GM/BOT;21.5GM/BOT</u>	<u>N18983</u>	<u>010</u>	Jan 31, 1989
-----------	---	----------------	--	---------------	------------	--------------

<u>AA</u>	+		<u>240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT</u>	<u>N18983</u>	<u>007</u>	Jun 12, 1987
-----------	---	--	---	---------------	------------	--------------

COLYTE WITH FLAVOR PACKS

<u>AA</u>	+	SCHWARZ PHARMA	<u>240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT</u>	<u>N18983</u>	<u>012</u>	Oct 08, 1998
-----------	---	----------------	---	---------------	------------	--------------

COLYTE-FLAVORED

<u>AA</u>	+	SCHWARZ PHARMA	<u>227.1GM/BOT;2.82GM/BOT;6.36GM/BOT;5.53GM/BOT;21.5GM/BOT</u>	<u>N18983</u>	<u>008</u>	Nov 14, 1991
-----------	---	----------------	--	---------------	------------	--------------

<u>AA</u>	+		<u>240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT</u>	<u>N18983</u>	<u>009</u>	Nov 14, 1991
-----------	---	--	---	---------------	------------	--------------

GOLYTELY

<u>AA</u>	+	BRAINTREE	<u>227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET</u>	<u>N19011</u>	<u>002</u>	Jun 02, 1992
-----------	---	-----------	---	---------------	------------	--------------

<u>AA</u>	+		<u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u>	<u>N19011</u>	<u>001</u>	Jul 13, 1984
-----------	---	--	---	---------------	------------	--------------

FOR SUSPENSION; ORAL

CO-LAV

<u>AA</u>		COPLEY PHARM	<u>240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT</u>	<u>N73428</u>	<u>001</u>	Jan 28, 1992
-----------	--	--------------	---	---------------	------------	--------------

GO-EVAC

<u>AA</u>		COPLEY PHARM	<u>236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT</u>	<u>N73433</u>	<u>001</u>	Apr 28, 1992
-----------	--	--------------	---	---------------	------------	--------------

POLYMYXIN B SULFATE

INJECTABLE; INJECTION

POLYMYXIN B SULFATE

<u>AP</u>	+	BEDFORD	<u>EQ 500,000 U BASE/VIAL</u>	<u>N60716</u>	<u>001</u>	
-----------	---	---------	-------------------------------	---------------	------------	--

<u>AP</u>		PHARMA TEK	<u>EQ 500,000 U BASE/VIAL</u>	<u>N63000</u>	<u>001</u>	Sep 30, 1994
-----------	--	------------	-------------------------------	---------------	------------	--------------

POWDER; FOR RX COMPOUNDING

POLY-RX

+	PHARMA TEK	100,000,000 UNITS/BOT	N61578	001	
---	------------	-----------------------	--------	-----	--

POLYMYXIN B SULFATE; TRIMETHOPRIM

SOLUTION/DROPS; OPHTHALMIC

TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE

<u>AT</u>		TAYLOR PHARMA	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>N65006</u>	<u>001</u>	Dec 17, 1998
-----------	--	---------------	---------------------------------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 293 (of 360)

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS; OPHTHALMIC

POLYTRIM

<u>AT</u>	+	ALLERGAN	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>N50567</u>	<u>001</u>	Oct 20, 1988
-----------	---	----------	---------------------------------------	---------------	------------	--------------

TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE

<u>AT</u>		BAUSCH AND LOMB	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>N64120</u>	<u>001</u>	Feb 14, 1997
-----------	--	-----------------	---------------------------------------	---------------	------------	--------------

<u>AT</u>		FALCON PHARMS	<u>10,000 UNITS/ML;EQ 1MG BASE/ML</u>	<u>N64211</u>	<u>001</u>	Apr 13, 1998
-----------	--	---------------	---------------------------------------	---------------	------------	--------------

POLYTHIAZIDE

TABLET; ORAL

RENESE

PFIZER

1MG

N12845 001

2MG

N12845 002

+

4MG

N12845 003

POLYTHIAZIDE; PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIZIDE

PFIZER

0.5MG;EQ 1MG BASE

N17986 001

0.5MG;EQ 2MG BASE

N17986 002

+

0.5MG;EQ 5MG BASE

N17986 003

POLYTHIAZIDE; RESERPINE

TABLET; ORAL

RENESE-R

+ PFIZER

2MG;0.25MG

N13636 001

PORACTANT ALFA

SUSPENSION; INTRATRACHEAL

CUROSURF

+ DEY

80MG/ML

N20744 001

Nov 18, 1999

PORFIMER SODIUM

INJECTABLE; INJECTION

PHOTOFRIN

+ AXCAN SCANDIPHARM

75MG/VIAL

N20451 001

Dec 27, 1995

POTASSIUM ACETATE

INJECTABLE; INJECTION

POTASSIUM ACETATE IN PLASTIC CONTAINER

+ HOSPIRA

2MEQ/ML

N18896 001

Jul 20, 1984

POTASSIUM AMINOSALICYLATE

CAPSULE; ORAL

PASKALIUM

+ GLENWOOD

500MG

N09395 004

TABLET; ORAL

PASKALIUM

+ GLENWOOD

1GM

N09395 003

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MICRO-KAB KV PHARM8MEQN18238 001MICRO-K 10AB + KV PHARM10MEQN18238 002

May 14, 1984

POTASSIUM CHLORIDEAB KV PHARM10MEQN70980 001

Feb 17, 1987

AB TEVA8MEQN73531 001

Apr 26, 1996

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 294 (of 360)

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

POTASSIUM CHLORIDE

<u>AB</u>	TEVA	<u>10MEQ</u>	<u>N73532</u>	<u>001</u>	Apr 26, 1996
-----------	------	--------------	---------------	------------	--------------

INJECTABLE; INJECTION

POTASSIUM CHLORIDE

<u>AP</u>	AM PHARM PARTNERS	<u>2MEQ/ML</u>	<u>N80225</u>	<u>001</u>	
	+	3MEQ/ML	N80225	003	

<u>AP</u>	B BRAUN	<u>2MEQ/ML</u>	<u>N85870</u>	<u>001</u>	
-----------	---------	----------------	---------------	------------	--

<u>AP</u>	BAXTER HLTHCARE	<u>2MEQ/ML</u>	<u>N85499</u>	<u>001</u>	
-----------	-----------------	----------------	---------------	------------	--

<u>AP</u>	+	HOSPIRA	<u>2MEQ/ML</u>	<u>N80205</u>	<u>001</u>
-----------	---	---------	----------------	---------------	------------

<u>AP</u>			<u>2MEQ/ML</u>	<u>N83345</u>	<u>002</u>
-----------	--	--	----------------	---------------	------------

	+		1.5MEQ/ML	N83345	001
--	---	--	-----------	--------	-----

<u>AP</u>	INTL MEDICATION	<u>2MEQ/ML</u>	<u>N83163</u>	<u>001</u>	
-----------	-----------------	----------------	---------------	------------	--

<u>AP</u>	LUITPOLD	<u>2MEQ/ML</u>	<u>N87584</u>	<u>001</u>	
-----------	----------	----------------	---------------	------------	--

<u>AP</u>		<u>2MEQ/ML</u>	<u>N87585</u>	<u>001</u>	
-----------	--	----------------	---------------	------------	--

POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	14.9MG/ML	N19904	001	Dec 26, 1989
---	-----------------	-----------	--------	-----	--------------

+		746MG/100ML	N19904	005	Dec 17, 1990
---	--	-------------	--------	-----	--------------

<u>AP</u>	HOSPIRA	<u>14.9MG/ML</u>	<u>N20161</u>	<u>005</u>	Nov 30, 1992
-----------	---------	------------------	---------------	------------	--------------

<u>AP</u>		<u>745MG/100ML</u>	<u>N20161</u>	<u>001</u>	Nov 30, 1992
-----------	--	--------------------	---------------	------------	--------------

POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>29.8MG/ML</u>	<u>N19904</u>	<u>002</u>
-----------	---	-----------------	------------------	---------------	------------

+		1.49GM/100ML	N19904	006	Dec 17, 1990
---	--	--------------	--------	-----	--------------

<u>AP</u>	+	HOSPIRA	<u>29.8MG/ML</u>	<u>N20161</u>	<u>006</u>
-----------	---	---------	------------------	---------------	------------

<u>AP</u>		<u>1.49GM/100ML</u>	<u>N20161</u>	<u>002</u>	Nov 30, 1992
-----------	--	---------------------	---------------	------------	--------------

POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>2.24GM/100ML</u>	<u>N19904</u>	<u>003</u>
-----------	---	-----------------	---------------------	---------------	------------

<u>AP</u>	+	HOSPIRA	<u>2.24GM/100ML</u>	<u>N20161</u>	<u>003</u>
-----------	---	---------	---------------------	---------------	------------

POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER

<u>AP</u>	+	BAXTER HLTHCARE	<u>2.98GM/100ML</u>	<u>N19904</u>	<u>004</u>
-----------	---	-----------------	---------------------	---------------	------------

<u>AP</u>	+	HOSPIRA	<u>2.98GM/100ML</u>	<u>N20161</u>	<u>004</u>
-----------	---	---------	---------------------	---------------	------------

POTASSIUM CHLORIDE IN PLASTIC CONTAINER

<u>AP</u>	AM PHARM PARTNERS	<u>2MEQ/ML</u>	<u>N88901</u>	<u>001</u>	Jan 25, 1985
-----------	-------------------	----------------	---------------	------------	--------------

<u>AP</u>		<u>2MEQ/ML</u>	<u>N88908</u>	<u>001</u>	Jan 25, 1985
-----------	--	----------------	---------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

K+10

BC	ALRA	10MEQ	N70999	001	Oct 22, 1987
----	------	-------	--------	-----	--------------

K+8

<u>AB</u>	ALRA	<u>8MEQ</u>	<u>N70998</u>	<u>001</u>	Jan 25, 1993
-----------	------	-------------	---------------	------------	--------------

KAON CL-10

BC	SAVAGE LABS	10MEQ	N17046	002	
----	-------------	-------	--------	-----	--

K-DUR 10

<u>AB</u>	KEY PHARMS	<u>10MEQ</u>	<u>N19439</u>	<u>002</u>	Jun 13, 1986
-----------	------------	--------------	---------------	------------	--------------

K-DUR 20

<u>AB</u>	+	KEY PHARMS	<u>20MEQ</u>	<u>N19439</u>	<u>001</u>
-----------	---	------------	--------------	---------------	------------

KLOR-CON

<u>AB</u>	+	UPSHER SMITH	<u>8MEQ</u>	<u>N19123</u>	<u>001</u>
-----------	---	--------------	-------------	---------------	------------

BC		10MEQ	N19123	002	Apr 17, 1986
----	--	-------	--------	-----	--------------

KLOR-CON M10

<u>AB</u>	UPSHER SMITH	<u>10MEQ</u>	<u>N74726</u>	<u>002</u>	Aug 09, 2000
-----------	--------------	--------------	---------------	------------	--------------

KLOR-CON M15

	UPSHER SMITH	15MEQ	N74726	003	Jun 06, 2003
--	--------------	-------	--------	-----	--------------

KLOR-CON M20

<u>AB</u>	UPSHER SMITH	<u>20MEQ</u>	<u>N74726</u>	<u>001</u>	Nov 20, 1998
-----------	--------------	--------------	---------------	------------	--------------

KLOTRIX

BC	APOTHECON	10MEQ	N17850	001	
----	-----------	-------	--------	-----	--

K-TAB

BC	ABBOTT	10MEQ	N18279	001	
----	--------	-------	--------	-----	--

POTASSIUM CHLORIDE

BC	ABBOTT	8MEQ	N18279	002	Aug 01, 1988
----	--------	------	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 295 (of 360)

POTASSIUM CHLORIDE

TABLET, EXTENDED RELEASE; ORAL
POTASSIUM CHLORIDE

AB	ANDRX PHARMS	10MEQ	N75604	001	Apr 10, 2002
AB		20MEQ	N75604	002	Apr 10, 2002
AB	COPLEY PHARM	8MEQ	N70618	001	Sep 09, 1987
AB	EURAND	20MEQ	N76368	001	Aug 18, 2004
AB	KV PHARM	20MEQ	N76044	001	Apr 05, 2002

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER					
AP	B BRAUN	150MG/100ML; 900MG/100ML	N19708	004	Sep 29, 1989
POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER					
AP	HOSPIRA	149MG/100ML; 900MG/100ML	N19686	001	Oct 17, 1988
POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER					
AP	HOSPIRA	298MG/100ML; 900MG/100ML	N19686	002	Oct 17, 1988
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER					
AP	BAXTER HLTHCARE	150MG/100ML; 900MG/100ML	N17648	001	
	SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.224%				
	BAXTER HLTHCARE	224MG/100ML; 900MG/100ML	N17648	003	
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER					
AP	BAXTER HLTHCARE	300MG/100ML; 900MG/100ML	N17648	002	

POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL
UROCIT-K

	MISSION PHARMA	5MEQ	N19071	001	Aug 30, 1985
+		10MEQ	N19071	002	Aug 31, 1992

POVIDONE-IODINE

SOLUTION/DROPS; OPHTHALMIC

	BETADINE				
+	ALCON	5%	N18634	001	Dec 17, 1986

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION

PRALIDOXIME CHLORIDE					
+	MERIDIAN MEDCL TECHN	300MG/ML	N18986	001	Apr 26, 1983
PROTOPAM CHLORIDE					
+	BAXTER HLTHCARE CORP	1GM/VIAL	N14134	001	

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL

MIRAPEX					
	BOEHRINGER INGELHEIM	0.125MG	N20667	001	Jul 01, 1997
+		0.25MG	N20667	002	Jul 01, 1997
		0.5MG	N20667	006	Feb 12, 1998
		1MG	N20667	003	Jul 01, 1997
		1.5MG	N20667	005	Jul 01, 1997

PRAMLINTIDE ACETATE

INJECTABLE; SUBCUTANEOUS

SYMLIN					
+	AMYLIN	EQ 3MG BASE/5ML (EQ 0.6MG BASE/ML)	N21332	001	Mar 16, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 296 (of 360)

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOL

BRISTOL MYERS SQUIBB

10MG

N19898 002

Oct 31, 1991

20MG

N19898 003

Oct 31, 1991

40MG

N19898 004

Mar 22, 1993

+

80MG

N19898 008

Dec 18, 2001

PRAZQUANTEL

TABLET; ORAL

BILTRICIDE

+ BAYER PHARMS

600MG

N18714 001

Dec 29, 1982

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

MINIPRESS

AB PFIZER

EQ 1MG BASE

N17442 002

AB +

EQ 2MG BASE

N17442 003

AB

EQ 5MG BASE

N17442 001

PRAZOSIN HCL

AB CLONMEL HLTHCARE

EQ 1MG BASE

N72705 001

May 16, 1989

AB

EQ 5MG BASE

N72707 001

May 16, 1989

PRAZOSIN HYDROCHLORIDE

AB IVAX PHARMS

EQ 1MG BASE

N71994 001

Sep 12, 1988

AB

EQ 2MG BASE

N71995 001

Sep 12, 1988

AB

EQ 5MG BASE

N71745 001

Sep 12, 1988

AB MYLAN

EQ 1MG BASE

N72573 001

May 16, 1989

AB

EQ 2MG BASE

N72574 001

May 16, 1989

AB

EQ 5MG BASE

N72575 001

May 16, 1989

AB SANDOZ

EQ 1MG BASE

N72576 001

May 16, 1989

AB

EQ 2MG BASE

N72577 001

May 16, 1989

AB

EQ 5MG BASE

N72578 001

May 16, 1989

AB WATSON LABS

EQ 1MG BASE

N72352 001

May 16, 1989

AB

EQ 2MG BASE

N72333 001

May 16, 1989

AB

EQ 5MG BASE

N72609 001

May 16, 1989

TABLET, EXTENDED RELEASE; ORAL

MINIPRESS XL

+ PFIZER

2.5MG

N19775 001

Jan 29, 1992

+

5MG

N19775 002

Jan 29, 1992

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLLIENT

+ DERMIK LABS

0.1%

N20279 001

Oct 29, 1993

OINTMENT; TOPICAL

DERMATOP

+ DERMIK LABS

0.1%

N19568 001

Sep 23, 1991

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

AA APOTEX

5MG/5ML

N40570 001

Aug 25, 2005

AA

15MG/5ML

N40571 001

Aug 25, 2005

AA HI TECH PHARMA

15MG/5ML

N40401 001

Feb 27, 2003

AA IVAX PHARMS

15MG/5ML

N40287 001

May 28, 1999

AA + KV PHARM

5MG/5ML

N40423 001

Oct 22, 2001

AA +

15MG/5ML

N40364 001

Apr 10, 2002

AA MORTON GROVE

15MG/5ML

N40313 001

Sep 10, 2003

AA PHARM ASSOC

15MG/5ML

N40399 001

Mar 05, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 297 (of 360)

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

<u>AA</u>	TEVA PHARMS	<u>15MG/5ML</u>	<u>N40322</u>	<u>001</u>	Jan 19, 2000
<u>AA</u>	UDL	<u>15MG/5ML</u>	<u>N40323</u>	<u>001</u>	May 13, 1999
<u>AA</u>	WE PHARMS	<u>15MG/5ML</u>	<u>N40192</u>	<u>001</u>	May 28, 1998

PRELONE

<u>AA</u>	TEVA	<u>15MG/5ML</u>	<u>N89081</u>	<u>001</u>	Feb 04, 1986
-----------	------	-----------------	---------------	------------	--------------

TABLET; ORAL

PREDNISOLONE

BX	LANNETT	5MG	N80531	002	
BX +	WATSON LABS	5MG	N80354	001	

PREDNISOLONE ACETATE

INJECTABLE; INJECTION

PREDNISOLONE ACETATE

+	STERIS	25MG/ML	N83398	001	
+		50MG/ML	N83764	001	

SUSPENSION/DROPS; OPHTHALMIC

ECONOPRED PLUS

<u>AB</u>	FALCON PHARMS	<u>1%</u>	<u>N17469</u>	<u>001</u>	
-----------	---------------	-----------	---------------	------------	--

PRED FORTE

<u>AB</u> +	ALLERGAN	<u>1%</u>	<u>N17011</u>	<u>001</u>	
-------------	----------	-----------	---------------	------------	--

PRED MILD

+	ALLERGAN	0.12%	N17100	001	
---	----------	-------	--------	-----	--

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

BLEPHAMIDE S.O.P.

+	ALLERGAN	0.2%;10%	N87748	001	Dec 03, 1986
---	----------	----------	--------	-----	--------------

SUSPENSION; OPHTHALMIC

BLEPHAMIDE

+	ALLERGAN	0.2%;10%	N12813	002	
---	----------	----------	--------	-----	--

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION; ORAL

ORAPRED

<u>AA</u> +	ASCENT PEDS	<u>EQ 15MG BASE/5ML</u>	<u>N75117</u>	<u>001</u>	Dec 14, 2000
-------------	-------------	-------------------------	---------------	------------	--------------

PEDIAPRED

<u>AA</u> +	UCB	<u>EQ 5MG BASE/5ML</u>	<u>N19157</u>	<u>001</u>	May 28, 1986
-------------	-----	------------------------	---------------	------------	--------------

PREDNISOLONE SODIUM PHOSPHATE

<u>AA</u>	HI TECH PHARMA	<u>EQ 5MG BASE/5ML</u>	<u>N75183</u>	<u>001</u>	Mar 26, 2003
<u>AA</u>	KV PHARM	<u>EQ 5MG BASE/5ML</u>	<u>N76982</u>	<u>001</u>	May 24, 2005
<u>AA</u>		<u>EQ 15MG BASE/5ML</u>	<u>N76988</u>	<u>001</u>	May 24, 2005
<u>AA</u>	MORTON GROVE	<u>EQ 5MG BASE/5ML</u>	<u>N75099</u>	<u>001</u>	Jun 28, 2002
<u>AA</u>		<u>EQ 15MG BASE/5ML</u>	<u>N76895</u>	<u>001</u>	Oct 04, 2004
<u>AA</u>	PADDOCK	<u>EQ 5MG BASE/5ML</u>	<u>N75988</u>	<u>001</u>	May 25, 2004
<u>AA</u>	PHARM ASSOC	<u>EQ 5MG BASE/5ML</u>	<u>N76123</u>	<u>001</u>	Dec 23, 2002
<u>AA</u>		<u>EQ 15MG BASE/5ML</u>	<u>N76913</u>	<u>001</u>	Apr 25, 2005
<u>AA</u>	WE PHARMS	<u>EQ 5MG BASE/5ML</u>	<u>N75181</u>	<u>001</u>	Dec 23, 2002
<u>AA</u>		<u>EQ 15MG BASE/5ML</u>	<u>N75250</u>	<u>001</u>	Jul 12, 2002

SOLUTION/DROPS; OPHTHALMIC

INFLAMASE FORTE

<u>AT</u> +	NOVARTIS	<u>EQ 0.9% PHOSPHATE</u>	<u>N80751</u>	<u>002</u>	
-------------	----------	--------------------------	---------------	------------	--

INFLAMASE MILD

<u>AT</u> +	NOVARTIS	<u>EQ 0.11% PHOSPHATE</u>	<u>N80751</u>	<u>001</u>	
-------------	----------	---------------------------	---------------	------------	--

PREDNISOLONE SODIUM PHOSPHATE

<u>AT</u>	BAUSCH AND LOMB	<u>EQ 0.11% PHOSPHATE</u>	<u>N40065</u>	<u>001</u>	Jul 29, 1994
<u>AT</u>		<u>EQ 0.9% PHOSPHATE</u>	<u>N40070</u>	<u>001</u>	Jul 29, 1994

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 298 (of 360)

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS; OPHTHALMIC

SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE

<u>AT</u>	ALCON	<u>EQ 0.23% PHOSPHATE;10%</u>	<u>N73630</u>	<u>001</u>	May 27, 1993
<u>AT</u>	BAUSCH AND LOMB	<u>EQ 0.23% PHOSPHATE;10%</u>	<u>N74449</u>	<u>001</u>	Dec 29, 1995
<u>VASOCIDIN</u>					
<u>AT</u>	+ NOVARTIS	<u>EQ 0.23% PHOSPHATE;10%</u>	<u>N18988</u>	<u>001</u>	Aug 26, 1988

PREDNISONE

SOLUTION; ORAL

PREDNISONE

+	ROXANE	5MG/5ML	N88703	001	Nov 08, 1984
PREDNISONE INTENSOL					
+	ROXANE	5MG/ML	N88810	001	Feb 20, 1985

TABLET; ORAL

PREDNISONE

<u>AB</u>	MUTUAL PHARM	<u>5MG</u>	<u>N89245</u>	<u>001</u>	Dec 04, 1985
<u>AB</u>		<u>10MG</u>	<u>N89246</u>	<u>001</u>	Dec 04, 1985
<u>AB</u>		<u>20MG</u>	<u>N89247</u>	<u>001</u>	Dec 04, 1985
<u>AB</u>	PHARM FORM	<u>5MG</u>	<u>N80209</u>	<u>001</u>	
<u>AB</u>	+ ROXANE	<u>1MG</u>	<u>N87800</u>	<u>001</u>	Apr 22, 1982
<u>AB</u>	+	<u>2.5MG</u>	<u>N87801</u>	<u>001</u>	Apr 22, 1982
<u>AB</u>	+	<u>5MG</u>	<u>N80352</u>	<u>001</u>	
<u>AB</u>	+	<u>10MG</u>	<u>N84122</u>	<u>001</u>	
<u>AB</u>	+	<u>20MG</u>	<u>N87342</u>	<u>001</u>	
<u>AB</u>	+	<u>50MG</u>	<u>N84283</u>	<u>001</u>	
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>N89983</u>	<u>001</u>	Jan 12, 1989
<u>AB</u>		<u>50MG</u>	<u>N89984</u>	<u>001</u>	Jan 12, 1989
<u>AB</u>	TRIGEN	<u>1MG</u>	<u>N40611</u>	<u>001</u>	Jun 06, 2005
<u>AB</u>		<u>5MG</u>	<u>N40362</u>	<u>002</u>	Aug 29, 2001
<u>AB</u>		<u>10MG</u>	<u>N40362</u>	<u>001</u>	Aug 29, 2001
<u>AB</u>		<u>20MG</u>	<u>N40362</u>	<u>003</u>	Jun 29, 2005
<u>AB</u>	VINTAGE PHARMS	<u>1MG</u>	<u>N40584</u>	<u>001</u>	Dec 21, 2004
<u>AB</u>		<u>2.5MG</u>	<u>N40581</u>	<u>001</u>	Dec 21, 2004
<u>AB</u>		<u>5MG</u>	<u>N40256</u>	<u>001</u>	Jul 12, 2002
<u>AB</u>		<u>10MG</u>	<u>N40256</u>	<u>002</u>	Jul 12, 2002
<u>AB</u>		<u>20MG</u>	<u>N40392</u>	<u>001</u>	Feb 12, 2003
<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>N80356</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>N85162</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>N87773</u>	<u>001</u>	Jul 13, 1982
<u>AB</u>		<u>20MG</u>	<u>N85161</u>	<u>001</u>	
<u>AB</u>		<u>20MG</u>	<u>N86813</u>	<u>001</u>	
<u>AB</u>		<u>50MG</u>	<u>N87772</u>	<u>001</u>	Jul 13, 1982
<u>AB</u>	WEST WARD	<u>2.5MG</u>	<u>N40538</u>	<u>001</u>	Jan 08, 2004
<u>AB</u>		<u>5MG</u>	<u>N80292</u>	<u>001</u>	
<u>AB</u>		<u>10MG</u>	<u>N88832</u>	<u>001</u>	Dec 04, 1985
<u>AB</u>		<u>20MG</u>	<u>N83677</u>	<u>001</u>	

PREGABALIN

CAPSULE; ORAL

LYRICA

CP PHARMS	25MG	N21446	001	Dec 30, 2004
	50MG	N21446	002	Dec 30, 2004
	75MG	N21446	003	Dec 30, 2004
	100MG	N21446	004	Dec 30, 2004
	150MG	N21446	005	Dec 30, 2004
	200MG	N21446	006	Dec 30, 2004
	225MG	N21446	007	Dec 30, 2004
+	300MG	N21446	008	Dec 30, 2004

PRESCRIPTION DRUG PRODUCT LIST

3 - 299 (of 360)

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION
CITANEST PLAIN

+ DENTSPLY PHARM	4%		N21382	001
------------------	----	--	--------	-----

PRIMAQUINE PHOSPHATE

TABLET; ORAL
PRIMAQUINE

+ SANOFI SYNTHELABO	EQ 15MG BASE		N08316	001
---------------------	--------------	--	--------	-----

PRIMIDONE

TABLET; ORAL

MYSOLINE

AB + VALEANT	50MG		N09170	003	
--------------	------	--	--------	-----	--

AB	250MG		N09170	002	
----	-------	--	--------	-----	--

PRIMIDONE

AB LANNETT	50MG		N84903	002	May 24, 2001
------------	------	--	--------	-----	--------------

AB	250MG		N84903	001	
----	-------	--	--------	-----	--

AB MUTUAL PHARM	50MG		N40626	001	Sep 29, 2005
-----------------	------	--	--------	-----	--------------

AB	250MG		N40626	002	Sep 29, 2005
----	-------	--	--------	-----	--------------

AB VINTAGE PHARMS	50MG		N40586	001	Feb 24, 2005
-------------------	------	--	--------	-----	--------------

AB	250MG		N40586	002	Feb 24, 2005
----	-------	--	--------	-----	--------------

AB WATSON LABS	250MG		N83551	001	
----------------	-------	--	--------	-----	--

PROBENECID

TABLET; ORAL

PROBALAN

AB LANNETT	500MG		N80966	001	
------------	-------	--	--------	-----	--

PROBENECID

AB IVAX PHARMS	500MG		N83740	001	May 09, 1984
----------------	-------	--	--------	-----	--------------

AB + MYLAN	500MG		N84211	002	
------------	-------	--	--------	-----	--

AB WATSON LABS	500MG		N84442	004	Mar 29, 1983
----------------	-------	--	--------	-----	--------------

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

AB IVAX PHARMS	250MG		N84604	001	
----------------	-------	--	--------	-----	--

AB	375MG		N84595	001	
----	-------	--	--------	-----	--

AB	500MG		N84606	001	
----	-------	--	--------	-----	--

AB WATSON LABS	250MG		N83287	001	
----------------	-------	--	--------	-----	--

AB	375MG		N84403	001	
----	-------	--	--------	-----	--

AB	500MG		N84280	001	
----	-------	--	--------	-----	--

PRONESTYL

AB APOTHECON	250MG		N07335	001	
--------------	-------	--	--------	-----	--

AB	375MG		N07335	004	
----	-------	--	--------	-----	--

AB +	500MG		N07335	003	
------	-------	--	--------	-----	--

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

AP HOSPIRA	100MG/ML		N89069	001	Feb 12, 1986
------------	----------	--	--------	-----	--------------

AP	500MG/ML		N89070	001	Feb 12, 1986
----	----------	--	--------	-----	--------------

AP	500MG/ML		N89537	001	Aug 25, 1987
----	----------	--	--------	-----	--------------

AP INTL MEDICATION	100MG/ML		N88636	001	Jul 31, 1984
--------------------	----------	--	--------	-----	--------------

AP	500MG/ML		N88637	001	Jul 31, 1984
----	----------	--	--------	-----	--------------

PRONESTYL

AP + APOTHECON	100MG/ML		N07335	002	
----------------	----------	--	--------	-----	--

AP +	500MG/ML		N07335	005	
------	----------	--	--------	-----	--

TABLET; ORAL

PRONESTYL

APOTHECON	250MG		N17371	001	
-----------	-------	--	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 300 (of 360)

PROCAINAMIDE HYDROCHLORIDE

TABLET; ORAL				
PRONESTYL				
	APOTHECON	375MG	N17371	002
+		500MG	N17371	003
TABLET, EXTENDED RELEASE; ORAL				
PROCANBID				
+	KING PHARMS	500MG	N20545	001 Jan 31, 1996
+		1GM	N20545	002 Jan 31, 1996
PRONESTYL-SR				
BC	APOTHECON	500MG	N87361	001

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION				
<u>NOVOCAIN</u>				
<u>AP</u>	+	HOSPIRA	<u>1%</u>	<u>N85362</u> <u>003</u>
<u>AP</u>	+		<u>2%</u>	<u>N85362</u> <u>004</u>
	+		10%	N86797 001
<u>PROCAINE HYDROCHLORIDE</u>				
<u>AP</u>		HOSPIRA	<u>1%</u>	<u>N80416</u> <u>001</u>
<u>AP</u>			<u>2%</u>	<u>N80416</u> <u>002</u>
<u>AP</u>		STERIS	<u>1%</u>	<u>N80658</u> <u>001</u>

PROCARBAZINE HYDROCHLORIDE

CAPSULE; ORAL				
MATULANE				
+	SIGMA TAU	EQ 50MG BASE	N16785	001

PROCHLORPERAZINE

SUPPOSITORY; RECTAL				
<u>COMPAZINE</u>				
<u>AB</u>	+	GLAXOSMITHKLINE	<u>25MG</u>	<u>N11127</u> <u>002</u>
			2.5MG	N11127 003
			5MG	N11127 001
<u>COMPRO</u>				
<u>AB</u>		PADDOCK	<u>25MG</u>	<u>N40246</u> <u>001</u> Jun 28, 2000
<u>PROCHLORPERAZINE</u>				
<u>AB</u>		G AND W LABS	<u>25MG</u>	<u>N40058</u> <u>001</u> Nov 24, 1993

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION				
<u>COMPAZINE</u>				
<u>AP</u>	+	GLAXOSMITHKLINE	<u>EQ 5MG BASE/ML</u>	<u>N10742</u> <u>002</u>
<u>PROCHLORPERAZINE</u>				
<u>AP</u>		BAXTER HLTHCARE	<u>EQ 5MG BASE/ML</u>	<u>N87759</u> <u>001</u> Oct 01, 1982
<u>PROCHLORPERAZINE EDISYLATE</u>				
<u>AP</u>		BAXTER HLTHCARE	<u>EQ 5MG BASE/ML</u>	<u>N89903</u> <u>001</u> Aug 29, 1989
<u>AP</u>		BEDFORD	<u>EQ 5MG BASE/ML</u>	<u>N40540</u> <u>001</u> May 28, 2004
<u>AP</u>		HOSPIRA	<u>EQ 5MG BASE/ML</u>	<u>N89703</u> <u>001</u> Apr 07, 1988
<u>AP</u>		SICOR PHARMS	<u>EQ 5MG BASE/ML</u>	<u>N40505</u> <u>001</u> May 30, 2003
<u>AP</u>		STERIS	<u>EQ 5MG BASE/ML</u>	<u>N89530</u> <u>001</u> Jul 08, 1987
SYRUP; ORAL				
COMPAZINE				
	GLAXOSMITHKLINE	EQ 5MG BASE/5ML	N11188	001

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL				
COMPAZINE				
+	GLAXOSMITHKLINE	EQ 10MG BASE	N21019	001 Oct 06, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 301 (of 360)

PROCHLORPERAZINE MALEATECAPSULE, EXTENDED RELEASE; ORAL
COMPAZINE

+ GLAXOSMITHKLINE EQ 15MG BASE N21019 002 Oct 06, 1999

TABLET; ORAL

COMPAZINEAB GLAXOSMITHKLINE EQ 5MG BASE N10571 001AB EQ 10MG BASE N10571 002AB + EQ 25MG BASE N10571 003PROCHLORPERAZINE MALEATEAB DURAMED PHARMS BARR EQ 5MG BASE N40207 001 May 01, 1997AB EQ 10MG BASE N40207 002 May 01, 1997AB IVAX PHARMS EQ 5MG BASE N40162 001 Jan 20, 1998AB EQ 10MG BASE N40162 002 Jan 20, 1998AB MYLAN EQ 5MG BASE N40185 002 Oct 28, 1996AB EQ 10MG BASE N40185 001 Oct 28, 1996AB SANDOZ EQ 5MG BASE N40101 001 Jul 19, 1996AB EQ 10MG BASE N40101 002 Jul 19, 1996AB EQ 25MG BASE N40101 003 Jul 19, 1996AB TEVA PHARMS EQ 5MG BASE N40120 001 Jul 11, 1996AB EQ 10MG BASE N40120 002 Jul 11, 1996AB TRIGEN EQ 5MG BASE N40268 001 Feb 27, 1998AB EQ 10MG BASE N40268 002 Feb 27, 1998PROCYCLIDINE HYDROCHLORIDE

TABLET; ORAL

KEMADRIN

+ MONARCH PHARMS 5MG N09818 003

PROGESTERONE

CAPSULE; ORAL

PROMETRIUM

UNIMED PHARMS 100MG N19781 001 May 14, 1998

+ 200MG N19781 002 Oct 15, 1999

GEL; VAGINAL

CRINONE

COLUMBIA RES LABS 4% N20701 001 Jul 31, 1997

+ 8% N20701 002 Jul 31, 1997

INJECTABLE; INJECTION

PROGESTERONEAO AM PHARM PARTNERS 50MG/ML N75906 001 Apr 25, 2001AO + WATSON LABS (UTAH) 50MG/ML N17362 002PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMETHAZINE HYDROCHLORIDEAP BAXTER HLTHCARE 25MG/ML N83312 001AP 50MG/ML N83312 002AP BEDFORD LABS 25MG/ML N40524 001 Mar 17, 2004AP 50MG/ML N40524 002 Mar 17, 2004AP BIONICHE PHARMA 25MG/ML N40471 001 Nov 21, 2002AP HOSPIRA 25MG/ML N40372 001 Jun 08, 2000AP 50MG/ML N40372 002 Jun 08, 2000AP 50MG/ML N83838 002AP PHARMAFORCE 25MG/ML N40515 001 Mar 19, 2003AP + SICOR PHARMS 25MG/ML N40454 001 Aug 22, 2002AP + 50MG/ML N40454 002 Aug 22, 2002AP STERIS 25MG/ML N84591 001AP 50MG/ML N80629 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 302 (of 360)

PROMETHAZINE HYDROCHLORIDE

SUPPOSITORY; RECTAL

PHENERGAN

<u>AB</u>	WYETH PHARMS INC	<u>12.5MG</u>	<u>N10926</u>	<u>002</u>	
<u>AB</u>	+	<u>25MG</u>	<u>N10926</u>	<u>001</u>	

BR	PROMETHACON	25MG	N84901	001	
	POLYMEDICA				

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	CLAY PARK	<u>12.5MG</u>	<u>N40500</u>	<u>001</u>	Jun 30, 2003
<u>AB</u>		<u>25MG</u>	<u>N40500</u>	<u>002</u>	Jun 30, 2003
<u>AB</u>	G AND W LABS	<u>12.5MG</u>	<u>N40428</u>	<u>002</u>	Mar 31, 2003
<u>AB</u>		<u>25MG</u>	<u>N40428</u>	<u>001</u>	Feb 05, 2002
<u>AB</u>	PADDOCK	<u>12.5MG</u>	<u>N40479</u>	<u>001</u>	Jun 24, 2003
<u>AB</u>		<u>25MG</u>	<u>N40479</u>	<u>002</u>	Jun 24, 2003
	PROMETHEGAN				
	G AND W LABS	50MG	N87165	001	Aug 14, 1987

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

<u>AA</u>	HI TECH PHARMA	<u>6.25MG/5ML</u>	<u>N40026</u>	<u>001</u>	Sep 25, 1998
-----------	----------------	-------------------	---------------	------------	--------------

PROMETHAZINE PLAIN

<u>AA</u>	+	<u>6.25MG/5ML</u>	<u>N87953</u>	<u>001</u>	Nov 15, 1982
	MORTON GROVE				

TABLET; ORAL

PHENERGAN

<u>AB</u>	WYETH PHARMS INC	<u>25MG</u>	<u>N07935</u>	<u>003</u>	
-----------	------------------	-------------	---------------	------------	--

PROMETHAZINE HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>25MG</u>	<u>N84234</u>	<u>001</u>	
<u>AB</u>	+	<u>50MG</u>	<u>N84176</u>	<u>001</u>	
BP	WATSON LABS	25MG	N83426	001	
BP		50MG	N83711	001	
<u>AB</u>	ZYDUS PHARMS USA	<u>25MG</u>	<u>N40596</u>	<u>002</u>	Nov 18, 2005
<u>AB</u>		<u>50MG</u>	<u>N40596</u>	<u>003</u>	Nov 18, 2005
		12.5MG	N40596	001	Nov 18, 2005

PROPAFENONE HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
RYTHMOL SR

	RELIANT PHARMS	225MG	N21416	001	Sep 04, 2003
		325MG	N21416	002	Sep 04, 2003
+		425MG	N21416	003	Sep 04, 2003

TABLET; ORAL

PROPAFENONE HYDROCHLORIDE

<u>AB</u>	KV PHARM	<u>150MG</u>	<u>N76193</u>	<u>001</u>	Feb 07, 2002
<u>AB</u>		<u>225MG</u>	<u>N76193</u>	<u>002</u>	Feb 07, 2002
<u>AB</u>		<u>300MG</u>	<u>N76193</u>	<u>003</u>	Feb 07, 2002
<u>AB</u>	MUTUAL PHARM	<u>150MG</u>	<u>N75998</u>	<u>001</u>	Nov 29, 2001
<u>AB</u>		<u>225MG</u>	<u>N75998</u>	<u>002</u>	Nov 29, 2001
<u>AB</u>		<u>300MG</u>	<u>N75998</u>	<u>003</u>	Nov 29, 2001
<u>AB</u>	PLIVA	<u>150MG</u>	<u>N76550</u>	<u>001</u>	Apr 23, 2004
<u>AB</u>		<u>225MG</u>	<u>N76550</u>	<u>002</u>	Apr 23, 2004
<u>AB</u>		<u>300MG</u>	<u>N76550</u>	<u>003</u>	Apr 23, 2004
<u>AB</u>	VINTAGE PHARMS	<u>150MG</u>	<u>N75938</u>	<u>001</u>	Oct 17, 2002
<u>AB</u>		<u>225MG</u>	<u>N75938</u>	<u>002</u>	Oct 17, 2002
<u>AB</u>		<u>300MG</u>	<u>N75938</u>	<u>003</u>	Oct 17, 2002
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>N75203</u>	<u>001</u>	Oct 24, 2000
<u>AB</u>		<u>225MG</u>	<u>N75203</u>	<u>002</u>	Oct 24, 2000
	<u>RYTHMOL</u>				
<u>AB</u>	RELIANT PHARMS	<u>150MG</u>	<u>N19151</u>	<u>001</u>	Nov 27, 1989
<u>AB</u>		<u>225MG</u>	<u>N19151</u>	<u>003</u>	Nov 20, 1992
<u>AB</u>	+	<u>300MG</u>	<u>N19151</u>	<u>002</u>	Nov 27, 1989

PRESCRIPTION DRUG PRODUCT LIST

3 - 303 (of 360)

PROPANTHELINE BROMIDE

TABLET; ORAL			
PRO-BANTHINE			
BP	+ SHIRE	7.5MG	N08732 003
BP	+	15MG	N08732 002
PROPANTHELINE BROMIDE			
BP	ROXANE	7.5MG	N80927 001
BP		15MG	N80927 002

PROPARACAINE HYDROCHLORIDE

SOLUTION; OPHTHALMIC			
<u>PROPARACAINE HYDROCHLORIDE</u>			
<u>AT</u>	TAYLOR PHARMA	<u>0.5%</u>	<u>N40277</u> <u>001</u> Mar 16, 2000
SOLUTION/DROPS; OPHTHALMIC			
<u>ALCAINE</u>			
<u>AT</u>	ALCON	<u>0.5%</u>	<u>N80027</u> <u>001</u>
<u>OPHTHAINE</u>			
<u>AT</u>	+ APOTHECON	<u>0.5%</u>	<u>N08883</u> <u>001</u>
<u>OPHTHETIC</u>			
<u>AT</u>	+ ALLERGAN	<u>0.5%</u>	<u>N12583</u> <u>001</u>
<u>PARACAINE</u>			
<u>AT</u>	OPTOPICS	<u>0.5%</u>	<u>N87681</u> <u>001</u> Aug 05, 1982
<u>PROPARACAINE HYDROCHLORIDE</u>			
<u>AT</u>	BAUSCH AND LOMB	<u>0.5%</u>	<u>N40074</u> <u>001</u> Sep 29, 1995

PROPOFOL

INJECTABLE; INJECTION			
<u>DIPRIVAN</u>			
<u>AB</u>	+ ASTRAZENECA	<u>10MG/ML</u>	<u>N19627</u> <u>002</u> Jun 11, 1996
<u>PROPOFOL</u>			
<u>AB</u>	BEDFORD	<u>10MG/ML</u>	<u>N74848</u> <u>001</u> Apr 19, 2005
<u>AB</u>	SICOR PHARMS	<u>10MG/ML</u>	<u>N75102</u> <u>001</u> Jan 04, 1999
<u>AB</u>		<u>10MG/ML</u>	<u>N75392</u> <u>001</u> Sep 19, 2000

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL			
<u>DARVON</u>			
<u>AA</u>	+ XANODYNE PHARM	<u>65MG</u>	<u>N10997</u> <u>003</u>
<u>PROPOXYPHENE HYDROCHLORIDE</u>			
<u>AA</u>	IVAX PHARMS	<u>65MG</u>	<u>N80269</u> <u>001</u>
<u>AA</u>	MYLAN	<u>65MG</u>	<u>N40569</u> <u>001</u> Dec 16, 2004
<u>AA</u>	TEVA	<u>65MG</u>	<u>N88615</u> <u>001</u> Oct 22, 1984
<u>AA</u>	WEST WARD	<u>65MG</u>	<u>N83501</u> <u>001</u>

PROPOXYPHENE NAPSYLATE

TABLET; ORAL			
DARVON-N			
	+ XANODYNE PHARM	100MG	N16862 002

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL			
INDERAL LA			
BX	WYETH PHARMS INC	80MG	N18553 002 Apr 19, 1983
BX		120MG	N18553 003 Apr 19, 1983
		60MG	N18553 004 Mar 18, 1987
	+	160MG	N18553 001 Apr 19, 1983
INNOPRAN XL			
BX	RELIANT PHARMS	80MG	N21438 001 Mar 12, 2003
BX		120MG	N21438 002 Mar 12, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 304 (of 360)

PROPRANOLOL HYDROCHLORIDE

CONCENTRATE; ORAL

PROPRANOLOL HYDROCHLORIDE INTENSOL

+	ROXANE	80MG/ML	N71388	001	May 15, 1987
---	--------	---------	--------	-----	--------------

INJECTABLE; INJECTION

PROPRANOLOL

<u>AP</u>	AM PHARM PARTNERS	<u>1MG/ML</u>	<u>N75826</u>	<u>001</u>	Aug 31, 2001
-----------	-------------------	---------------	---------------	------------	--------------

PROPRANOLOL HYDROCHLORIDE

<u>AP</u>	+	BAXTER HLTHCARE CORP	<u>1MG/ML</u>	<u>N16419</u>	<u>001</u>
-----------	---	----------------------	---------------	---------------	------------

<u>AP</u>	BEDFORD	<u>1MG/ML</u>	<u>N75792</u>	<u>001</u>	Aug 29, 2000
-----------	---------	---------------	---------------	------------	--------------

<u>AP</u>	SABEX 2002	<u>1MG/ML</u>	<u>N76400</u>	<u>001</u>	Feb 26, 2003
-----------	------------	---------------	---------------	------------	--------------

SOLUTION; ORAL

PROPRANOLOL HYDROCHLORIDE

+	ROXANE	20MG/5ML	N70979	001	May 15, 1987
---	--------	----------	--------	-----	--------------

+		40MG/5ML	N70690	001	May 15, 1987
---	--	----------	--------	-----	--------------

TABLET; ORAL

INDERAL

<u>AB</u>	WYETH PHARMS INC	<u>10MG</u>	<u>N16418</u>	<u>001</u>	
-----------	------------------	-------------	---------------	------------	--

<u>AB</u>		<u>20MG</u>	<u>N16418</u>	<u>003</u>	
-----------	--	-------------	---------------	------------	--

<u>AB</u>		<u>40MG</u>	<u>N16418</u>	<u>002</u>	
-----------	--	-------------	---------------	------------	--

<u>AB</u>		<u>60MG</u>	<u>N16418</u>	<u>009</u>	Oct 18, 1982
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	+	<u>80MG</u>	<u>N16418</u>	<u>004</u>	
-----------	---	-------------	---------------	------------	--

PROPRANOLOL HYDROCHLORIDE

<u>AB</u>	MYLAN	<u>10MG</u>	<u>N70211</u>	<u>001</u>	Nov 19, 1985
-----------	-------	-------------	---------------	------------	--------------

<u>AB</u>		<u>20MG</u>	<u>N70212</u>	<u>001</u>	Nov 19, 1985
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>40MG</u>	<u>N70213</u>	<u>001</u>	Nov 19, 1985
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>80MG</u>	<u>N70214</u>	<u>001</u>	Nov 19, 1985
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	PAR PHARM	<u>10MG</u>	<u>N70217</u>	<u>001</u>	Aug 01, 1986
-----------	-----------	-------------	---------------	------------	--------------

<u>AB</u>		<u>20MG</u>	<u>N70218</u>	<u>001</u>	Aug 01, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>40MG</u>	<u>N70219</u>	<u>001</u>	Aug 01, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>60MG</u>	<u>N70220</u>	<u>001</u>	Sep 24, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>80MG</u>	<u>N70221</u>	<u>001</u>	Apr 14, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>90MG</u>	<u>N71288</u>	<u>001</u>	Oct 22, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	PLIVA	<u>10MG</u>	<u>N71972</u>	<u>001</u>	Apr 06, 1988
-----------	-------	-------------	---------------	------------	--------------

<u>AB</u>		<u>20MG</u>	<u>N71973</u>	<u>001</u>	Apr 06, 1988
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>40MG</u>	<u>N71974</u>	<u>001</u>	Apr 06, 1988
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>60MG</u>	<u>N71975</u>	<u>001</u>	Apr 06, 1988
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>80MG</u>	<u>N71976</u>	<u>001</u>	Apr 06, 1988
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>90MG</u>	<u>N71977</u>	<u>001</u>	Apr 06, 1988
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>10MG</u>	<u>N70663</u>	<u>001</u>	Jun 13, 1986
-----------	--------	-------------	---------------	------------	--------------

<u>AB</u>		<u>20MG</u>	<u>N70664</u>	<u>001</u>	Jun 13, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>40MG</u>	<u>N70665</u>	<u>001</u>	Jun 13, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>60MG</u>	<u>N70666</u>	<u>001</u>	Oct 10, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>80MG</u>	<u>N70667</u>	<u>001</u>	Jun 13, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	TEVA	<u>40MG</u>	<u>N70234</u>	<u>001</u>	Jun 23, 1986
-----------	------	-------------	---------------	------------	--------------

<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N70175</u>	<u>001</u>	May 13, 1986
-----------	-------------	-------------	---------------	------------	--------------

<u>AB</u>		<u>10MG</u>	<u>N70548</u>	<u>001</u>	Jul 10, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>20MG</u>	<u>N70176</u>	<u>001</u>	May 13, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>40MG</u>	<u>N70177</u>	<u>001</u>	May 13, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>40MG</u>	<u>N70550</u>	<u>001</u>	Apr 11, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>60MG</u>	<u>N71791</u>	<u>001</u>	Jul 15, 1987
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>80MG</u>	<u>N70178</u>	<u>001</u>	May 13, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>80MG</u>	<u>N70551</u>	<u>001</u>	Jul 10, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>		<u>90MG</u>	<u>N71792</u>	<u>001</u>	Jul 15, 1987
-----------	--	-------------	---------------	------------	--------------

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

BD	+	CLONMEL HLTHCARE	50MG	N06188	001
----	---	------------------	------	--------	-----

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 305 (of 360)

PROPYLTHIOURACIL

TABLET; ORAL				
PROPYLTHIOURACIL				
BD	IMPAX LABS	50MG	N80159	001
BD	PUREPAC PHARM	50MG	N80172	001
BD	WEST WARD	50MG	N80154	001

PROTAMINE SULFATE

INJECTABLE; INJECTION				
PROTAMINE SULFATE				
	+ AM PHARM PARTNERS	10MG/ML	N89454	001 Apr 07, 1987

PROTRIPTYLINE HYDROCHLORIDE

TABLET; ORAL				
<u>VIVACTIL</u>				
<u>AB</u>	ODYSSEY PHARMS	<u>5MG</u>	<u>N73644</u>	<u>001</u> Aug 24, 1995
<u>AB</u>	+	<u>10MG</u>	<u>N73645</u>	<u>001</u> Aug 24, 1995

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

TABLET; ORAL				
<u>CORPHED</u>				
<u>AA</u>	SANDOZ	<u>60MG;2.5MG</u>	<u>N88602</u>	<u>001</u> Apr 11, 1985
<u>PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE</u>				
<u>AA</u>	+	<u>60MG;2.5MG</u>	<u>N88193</u>	<u>001</u> May 17, 1983

PYRAZINAMIDE

TABLET; ORAL				
<u>PYRAZINAMIDE</u>				
<u>AB</u>	+	CLONMEL HLTHCARE	<u>500MG</u>	<u>N80157</u> <u>001</u>
<u>AB</u>		MIKART	<u>500MG</u>	<u>N81319</u> <u>001</u> Jun 30, 1992

PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION				
<u>MESTINON</u>				
<u>AP</u>	+	VALEANT PHARM INTL	<u>5MG/ML</u>	<u>N09830</u> <u>001</u>
<u>REGONOL</u>				
<u>AP</u>		SABEX 2002	<u>5MG/ML</u>	<u>N17398</u> <u>001</u>
SYRUP; ORAL				
MESTINON				
	+	VALEANT PHARM INTL	60MG/5ML	N15193 001
TABLET; ORAL				
<u>MESTINON</u>				
<u>AB</u>	+	VALEANT PHARM INTL	<u>60MG</u>	<u>N09829</u> <u>002</u>
<u>PYRIDOSTIGMINE BROMIDE</u>				
<u>AB</u>		BARR	<u>60MG</u>	<u>N40512</u> <u>001</u> Oct 08, 2003
			30MG	N40512 002 Jul 20, 2005
<u>AB</u>		COREPHARMA	<u>60MG</u>	<u>N40457</u> <u>001</u> Dec 26, 2002
<u>AB</u>		IMPAX LABS	<u>60MG</u>	<u>N40502</u> <u>001</u> Apr 24, 2003
TABLET, EXTENDED RELEASE; ORAL				
MESTINON				
	+	VALEANT PHARM INTL	180MG	N11665 001

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION				
<u>PYRIDOXINE HYDROCHLORIDE</u>				
<u>AP</u>		AM PHARM PARTNERS	<u>100MG/ML</u>	<u>N80618</u> <u>001</u>
<u>AP</u>	+	STERIS	<u>100MG/ML</u>	<u>N80572</u> <u>001</u>

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 306 (of 360)

PYRIMETHAMINE

TABLET; ORAL				
DARAPRIM				
+ GLAXOSMITHKLINE	25MG	N08578	001	

PYRIMETHAMINE; SULFADOXINE

TABLET; ORAL				
FANSIDAR				
+ ROCHE	25MG;500MG	N18557	001	

QUAZEPAM

TABLET; ORAL				
DORAL				
+ MEDPOINTE PHARM HLC	15MG	N18708	001	Dec 27, 1985

QUETIAPINE FUMARATE

TABLET; ORAL				
SEROQUEL				
+ ASTRAZENECA	EQ 25MG BASE	N20639	001	Sep 26, 1997
	EQ 50MG BASE	N20639	007	Oct 04, 2005
	EQ 100MG BASE	N20639	002	Sep 26, 1997
	EQ 200MG BASE	N20639	003	Sep 26, 1997
	EQ 300MG BASE	N20639	005	Jul 26, 2000
	EQ 400MG BASE	N20639	006	Oct 04, 2005

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL				
<u>ACCUPRIL</u>				
<u>AB</u> PFIZER PHARMS	<u>EQ 5MG BASE</u>	<u>N19885</u>	<u>001</u>	Nov 19, 1991
<u>AB</u>	<u>EQ 10MG BASE</u>	<u>N19885</u>	<u>002</u>	Nov 19, 1991
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N19885</u>	<u>003</u>	Nov 19, 1991
<u>AB</u> +	<u>EQ 40MG BASE</u>	<u>N19885</u>	<u>004</u>	Nov 19, 1991
<u>QUINAPRIL</u>				
<u>AB</u> RANBAXY	<u>EQ 5MG BASE</u>	<u>N76607</u>	<u>001</u>	Dec 15, 2004
<u>AB</u>	<u>EQ 10MG BASE</u>	<u>N76607</u>	<u>002</u>	Dec 15, 2004
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76607</u>	<u>003</u>	Dec 15, 2004
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>N76607</u>	<u>004</u>	Dec 15, 2004
<u>QUINAPRIL HYDROCHLORIDE</u>				
<u>AB</u> AMIDE PHARM	<u>EQ 5MG BASE</u>	<u>N76459</u>	<u>001</u>	Dec 22, 2004
<u>AB</u>	<u>EQ 10MG BASE</u>	<u>N76459</u>	<u>002</u>	Dec 22, 2004
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76459</u>	<u>003</u>	Dec 22, 2004
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>N76459</u>	<u>004</u>	Dec 22, 2004
<u>AB</u> ANDRX PHARMS	<u>EQ 5MG BASE</u>	<u>N76049</u>	<u>001</u>	Jan 14, 2005
<u>AB</u>	<u>EQ 10MG BASE</u>	<u>N76049</u>	<u>002</u>	Jan 14, 2005
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76049</u>	<u>003</u>	Jan 14, 2005
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>N76049</u>	<u>004</u>	Jan 14, 2005
<u>AB</u> GENPHARM	<u>EQ 5MG BASE</u>	<u>N76036</u>	<u>001</u>	Jan 28, 2005
<u>AB</u>	<u>EQ 10MG BASE</u>	<u>N76036</u>	<u>002</u>	Jan 28, 2005
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76036</u>	<u>003</u>	Jan 28, 2005
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>N76036</u>	<u>004</u>	Jan 28, 2005
<u>AB</u> MYLAN	<u>EQ 5MG BASE</u>	<u>N76694</u>	<u>001</u>	Dec 23, 2004
<u>AB</u>	<u>EQ 10MG BASE</u>	<u>N76694</u>	<u>002</u>	Dec 23, 2004
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76694</u>	<u>003</u>	Dec 23, 2004
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>N76694</u>	<u>004</u>	Dec 23, 2004
<u>AB</u> SANDOZ	<u>EQ 5MG BASE</u>	<u>N76803</u>	<u>001</u>	Mar 02, 2005
<u>AB</u>	<u>EQ 10MG BASE</u>	<u>N76803</u>	<u>002</u>	Mar 02, 2005
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76803</u>	<u>003</u>	Mar 02, 2005
<u>AB</u>	<u>EQ 40MG BASE</u>	<u>N76803</u>	<u>004</u>	Mar 02, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 307 (of 360)

QUINIDINE GLUCONATEINJECTABLE; INJECTION
QUINIDINE GLUCONATE

+ LILLY 80MG/ML N07529 002 Feb 10, 1989

TABLET, EXTENDED RELEASE; ORAL
QUINIDINE GLUCONATE

+ MUTUAL PHARM 324MG N89338 001 Feb 11, 1987

QUINIDINE SULFATE

TABLET; ORAL

QUINIDINE SULFATEAB CLONMEL HLTHCARE 200MG N87011 001AB LANNETT 200MG N83743 001AB MUTUAL PHARM 100MG N81029 001 Apr 14, 1989AB 200MG N81030 001 Apr 14, 1989AB PHARM FORM 200MG N83808 001AB SANDOZ 200MG N84631 001AB 200MG N84914 001AB 300MG N88072 001 Sep 26, 1983AB 300MG N89839 001 Sep 29, 1988AB WATSON LABS 200MG N83288 001AB 200MG N85140 002AB 300MG N85583 001TABLET, EXTENDED RELEASE; ORAL
QUINIDINE SULFATE

+ TEVA PHARMS 300MG N40045 001 Jun 30, 1994

QUININE SULFATECAPSULE; ORAL
QUININE SULFATE

+ MUTUAL PHARM 324MG N21799 001 Aug 12, 2005

RABEPRAZOLE SODIUMTABLET, DELAYED RELEASE; ORAL
ACIPHEX

+ EISAI MEDCL RES 20MG N20973 002 Aug 19, 1999

RALOXIFENE HYDROCHLORIDETABLET; ORAL
EVISTA

+ LILLY 60MG N20815 001 Dec 09, 1997

RAMELTEONTABLET; ORAL
ROZEREM

+ TAKEDA GLOBAL 8MG N21782 001 Jul 22, 2005

RAMIPRIL

CAPSULE; ORAL

ALTACEAB KING PHARMS 1.25MG N19901 001 Jan 28, 1991AB 2.5MG N19901 002 Jan 28, 1991AB 5MG N19901 003 Jan 28, 1991AB + 10MG N19901 004 Jan 28, 1991RAMIPRILAB COBALT 1.25MG N76549 001 Oct 24, 2005AB 2.5MG N76549 002 Oct 24, 2005AB 5MG N76549 003 Oct 24, 2005AB 10MG N76549 004 Oct 24, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 308 (of 360)

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

<u>AB</u>	DR REDDYS LABS LTD	<u>EQ 150MG BASE</u>	<u>N75742</u>	<u>001</u>	Nov 29, 2000
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N75742</u>	<u>002</u>	Nov 29, 2000
<u>AB</u>	GENPHARM	<u>EQ 150MG BASE</u>	<u>N75564</u>	<u>001</u>	Oct 27, 2000
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N75564</u>	<u>002</u>	Oct 27, 2000
<u>AB</u>	SANDOZ	<u>EQ 150MG BASE</u>	<u>N74655</u>	<u>001</u>	Oct 22, 1997
<u>AB</u>	+	<u>EQ 300MG BASE</u>	<u>N74655</u>	<u>002</u>	Oct 22, 1997
<u>AB</u>	TEVA	<u>EQ 150MG BASE</u>	<u>N75557</u>	<u>001</u>	Oct 31, 2003
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N75557</u>	<u>002</u>	Oct 31, 2003

GRANULE, EFFERVESCENT; ORAL

ZANTAC 150

+	GLAXOSMITHKLINE	EQ 150MG BASE/PACKET	N20251	002	Mar 31, 1994
---	-----------------	----------------------	--------	-----	--------------

INJECTABLE; INJECTION

RANITIDINE

<u>AP</u>	<u>BEDFORD</u>	<u>EQ 25MG BASE/ML</u>	<u>N74764</u>	<u>001</u>	Nov 19, 2004
	<u>RANITIDINE HYDROCHLORIDE</u>				
<u>AP</u>	<u>BEN VENUE</u>	<u>EQ 25MG BASE/ML</u>	<u>N74777</u>	<u>001</u>	Mar 02, 2005
	<u>ZANTAC</u>				
<u>AP</u>	<u>+ GLAXOSMITHKLINE</u>	<u>EQ 25MG BASE/ML</u>	<u>N19090</u>	<u>001</u>	Oct 19, 1984
	ZANTAC IN PLASTIC CONTAINER				
	<u>+ GLAXOSMITHKLINE</u>	<u>EO 1MG BASE/ML</u>	<u>N19593</u>	<u>002</u>	Sep 27, 1991

SYRUP; ORAL

ZANTAC

+	GLAXOSMITHKLINE	EQ 15MG BASE/ML	N19675	001	Dec 30, 1988
---	-----------------	-----------------	--------	-----	--------------

TABLET; ORAL

RANITIDINE

<u>AB</u>	<u>RANBAXY</u>	<u>EQ 150MG BASE</u>	<u>N75439</u>	<u>001</u>	Apr 19, 2000
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N75439</u>	<u>002</u>	Apr 19, 2000
<u>RANITIDINE HYDROCHLORIDE</u>					
<u>AB</u>	<u>COBALT</u>	<u>EQ 150MG BASE</u>	<u>N77426</u>	<u>001</u>	Dec 19, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N77426</u>	<u>002</u>	Dec 19, 2005
<u>AB</u>	<u>DR REDDYS LABS INC</u>	<u>EQ 150MG BASE</u>	<u>N76705</u>	<u>001</u>	Jul 27, 2005
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N76705</u>	<u>002</u>	Jul 27, 2005
<u>AB</u>	<u>GENPHARM</u>	<u>EQ 150MG BASE</u>	<u>N74023</u>	<u>001</u>	Aug 22, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N74023</u>	<u>002</u>	Aug 22, 1997
<u>AB</u>	<u>IVAX PHARMS</u>	<u>EQ 150MG BASE</u>	<u>N75165</u>	<u>001</u>	Sep 30, 1998
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N75165</u>	<u>002</u>	Sep 30, 1998
<u>AB</u>	<u>MYLAN</u>	<u>EQ 150MG BASE</u>	<u>N74552</u>	<u>001</u>	Jul 30, 1998
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N74552</u>	<u>002</u>	Jul 30, 1998
<u>AB</u>	<u>PAR PHARM</u>	<u>EQ 150MG BASE</u>	<u>N75180</u>	<u>001</u>	Jan 28, 1999
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N75180</u>	<u>002</u>	Jan 28, 1999
<u>AB</u>	<u>SANDOZ</u>	<u>EQ 150MG BASE</u>	<u>N74467</u>	<u>001</u>	Aug 29, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N74467</u>	<u>002</u>	Aug 29, 1997
<u>AB</u>	<u>TEVA</u>	<u>EQ 150MG BASE</u>	<u>N74488</u>	<u>001</u>	Jul 31, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N74488</u>	<u>002</u>	Jul 31, 1997
<u>AB</u>	<u>TORPHARM</u>	<u>EQ 150MG BASE</u>	<u>N74680</u>	<u>001</u>	Sep 12, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N74680</u>	<u>002</u>	Sep 12, 1997
<u>AB</u>	<u>WATSON LABS</u>	<u>EQ 150MG BASE</u>	<u>N74864</u>	<u>001</u>	Oct 20, 1997
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N74864</u>	<u>002</u>	Oct 20, 1997
<u>AB</u>	<u>WOCKHARDT</u>	<u>EQ 150MG BASE</u>	<u>N75208</u>	<u>001</u>	Dec 17, 1998
<u>AB</u>		<u>EQ 300MG BASE</u>	<u>N75208</u>	<u>002</u>	Dec 17, 1998
<u>ZANTAC 150</u>					
<u>AB</u>	<u>GLAXOSMITHKLINE</u>	<u>EQ 150MG BASE</u>	<u>N18703</u>	<u>001</u>	Jun 09, 1983
<u>ZANTAC 300</u>					
<u>AB</u>	<u>+ GLAXOSMITHKLINE</u>	<u>EQ 300MG BASE</u>	<u>N18703</u>	<u>002</u>	Dec 09, 1985

PRESCRIPTION DRUG PRODUCT LIST

3 - 309 (of 360)

RANITIDINE HYDROCHLORIDE

TABLET, EFFERVESCENT; ORAL

ZANTAC 150

+ GLAXOSMITHKLINE	EQ 150MG BASE	N20251 001	Mar 31, 1994
-------------------	---------------	------------	--------------

ZANTAC 25

GLAXOSMITHKLINE	EQ 25MG BASE	N20251 003	Apr 01, 2004
-----------------	--------------	------------	--------------

REMIFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ULTIVA

ABBOTT

EQ 1MG BASE/VIAL	N20630 001	Jul 12, 1996
------------------	------------	--------------

EQ 2MG BASE/VIAL	N20630 002	Jul 12, 1996
------------------	------------	--------------

+	EQ 5MG BASE/VIAL	N20630 003	Jul 12, 1996
---	------------------	------------	--------------

REPAGLINIDE

TABLET; ORAL

PRANDIN

NOVO NORDISK INC

0.5MG	N20741 001	Dec 22, 1997
-------	------------	--------------

1MG	N20741 002	Dec 22, 1997
-----	------------	--------------

+	2MG	N20741 003	Dec 22, 1997
---	-----	------------	--------------

RESCINNAMINE

TABLET; ORAL

MODERIL

PFIZER

0.25MG	N10686 003	
--------	------------	--

+	0.5MG	N10686 006	
---	-------	------------	--

RESERPINE

TABLET; ORAL

RESERPINE

BP	SANDOZ	0.1MG	N09838 001	
----	--------	-------	------------	--

BP	+	0.25MG	N09838 002	
----	---	--------	------------	--

SERPALAN

BP	LANNETT	0.1MG	N10124 001	
----	---------	-------	------------	--

BP		0.25MG	N10124 002	
----	--	--------	------------	--

RIBAVIRIN

CAPSULE; ORAL

REBETOL

<u>AB</u>	+ SCHERING PLOUGH RES	<u>200MG</u>	<u>N20903 002</u>	Jul 25, 2001
-----------	-----------------------	--------------	-------------------	--------------

+		200MG ***Indicated for use and comarketed with interferon alfa-2b, recombinant (Intron A), as Rebetron Combination Therapy***	N20903 001	Jun 03, 1998
---	--	---	------------	--------------

RIBASPHERE

<u>AB</u>	THREE RIVERS PHARMS	<u>200MG</u>	<u>N76203 001</u>	Apr 06, 2004
-----------	---------------------	--------------	-------------------	--------------

RIBAVIRIN

<u>AB</u>	SANDOZ	<u>200MG</u>	<u>N76192 001</u>	Apr 06, 2004
-----------	--------	--------------	-------------------	--------------

<u>AB</u>	TEVA	<u>200MG</u>	<u>N76277 001</u>	Oct 04, 2004
-----------	------	--------------	-------------------	--------------

<u>AB</u>	ZYDUS PHARMS USA	<u>200MG</u>	<u>N77224 001</u>	Oct 28, 2005
-----------	------------------	--------------	-------------------	--------------

FOR SOLUTION; INHALATION

VIRAZOLE

+	VALEANT PHARM INTL	6GM/VIAL	N18859 001	Dec 31, 1985
---	--------------------	----------	------------	--------------

SOLUTION; ORAL

REBETOL

+	SCHERING	40MG/ML	N21546 001	Jul 29, 2003
---	----------	---------	------------	--------------

TABLET; ORAL

COPEGUS

<u>AB</u>	+ ROCHE	<u>200MG</u>	<u>N21511 001</u>	Dec 03, 2002
-----------	---------	--------------	-------------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 310 (of 360)

RIBAVIRIN

TABLET; ORAL

RIBAVIRIN

<u>AB</u>	TEVA	<u>200MG</u>	<u>N77053</u>	<u>001</u>	Dec 05, 2005
<u>AB</u>	THREE RIVERS PHARMS	<u>200MG</u>	<u>N77456</u>	<u>001</u>	Dec 05, 2005
		400MG	N77456	002	Dec 05, 2005
	+	600MG	N77456	003	Dec 05, 2005
<u>AB</u>	ZYDUS PHARMS USA	<u>200MG</u>	<u>N77094</u>	<u>001</u>	Dec 05, 2005

RIFABUTIN

CAPSULE; ORAL

MYCOBUTIN

	+	PHARMACIA AND UPJOHN	150MG	N50689	001	Dec 23, 1992
--	---	----------------------	-------	--------	-----	--------------

RIFAMPIN

CAPSULE; ORAL

RIFADIN

<u>AB</u>	AVENTIS PHARMS	<u>150MG</u>	<u>N62303</u>	<u>001</u>	
<u>AB</u>	+	<u>300MG</u>	<u>N50420</u>	<u>001</u>	
<u>AB</u>	<u>RIFAMPIN</u>				
<u>AB</u>	SANDOZ	<u>150MG</u>	<u>N64150</u>	<u>002</u>	Jan 02, 1998
<u>AB</u>		<u>300MG</u>	<u>N64150</u>	<u>001</u>	May 28, 1997
<u>AB</u>	VERSAPHARM	<u>150MG</u>	<u>N65028</u>	<u>001</u>	Mar 14, 2001
<u>AB</u>		<u>300MG</u>	<u>N65028</u>	<u>002</u>	Mar 14, 2001
<u>AB</u>	<u>RIMACTANE</u>				
<u>AB</u>	AMIDE PHARM	<u>300MG</u>	<u>N50429</u>	<u>001</u>	

INJECTABLE; INJECTION

RIFADIN

AP	+	AVENTIS PHARMS	600MG/VIAL	N50627	001	May 25, 1989
		RIFAMPIN				
AP		BEDFORD	600MG/VIAL	N64217	001	Oct 29, 1999

RIFAPENTINE

TABLET; ORAL

PRIFTIN

	+	AVENTIS PHARMS	150MG	N21024	001	Jun 22, 1998
--	---	----------------	-------	--------	-----	--------------

RIFAXIMIN

TABLET; ORAL

XIFAXAN

	+	SALIX PHARMS	200MG	N21361	001	May 25, 2004
--	---	--------------	-------	--------	-----	--------------

RILUZOLE

TABLET; ORAL

RILUTEK

<u>AB</u>	+	<u>AVENTIS</u>	<u>50MG</u>	<u>N20599</u>	<u>001</u>	Dec 12, 1995
		<u>RILUZOLE</u>				
<u>AB</u>		<u>IMPAX LABS</u>	<u>50MG</u>	<u>N76173</u>	<u>001</u>	Jan 29, 2003

RIMANTADINE HYDROCHLORIDE

SYRUP; ORAL

FLUMADINE

	+	FOREST LABS	50MG/5ML	N19650	001	Sep 17, 1993
--	---	-------------	----------	--------	-----	--------------

TABLET; ORAL

FLUMADINE

<u>AB</u>	+	FOREST LABS	<u>100MG</u>	<u>N19649</u>	<u>001</u>	Sep 17, 1993
		<u>RIMANTADINE HYDROCHLORIDE</u>				
<u>AB</u>		AMIDE PHARM	<u>100MG</u>	<u>N76375</u>	<u>001</u>	Jan 14, 2003
<u>AB</u>		COREPHARMA	<u>100MG</u>	<u>N75916</u>	<u>001</u>	Nov 02, 2001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 311 (of 360)

RIMANTADINE HYDROCHLORIDE

TABLET; ORAL

RIMANTADINE HYDROCHLORIDE

AB	IMPAX LABS	100MG	N76132	001	Aug 30, 2002
-----------	------------	--------------	---------------	------------	--------------

RIMEXOLONE

SUSPENSION/DROPS; OPHTHALMIC

VEXOL

+	ALCON	1%	N20474	001	Dec 30, 1994
---	-------	----	--------	-----	--------------

RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL

PROCTER AND GAMBLE

5MG

N20835 002 Apr 14, 2000

30MG

N20835 001 Mar 27, 1998

+

35MG

N20835 003 May 25, 2002

RISPERIDONE

INJECTABLE; INTRAMUSCULAR

RISPERDAL CONSTA

JANSSEN PHARMA

25MG/VIAL

N21346 001 Oct 29, 2003

37.5MG/VIAL

N21346 002 Oct 29, 2003

+

50MG/VIAL

N21346 003 Oct 29, 2003

SOLUTION; ORAL

RISPERDAL

+ JANSSEN PHARMA

1MG/ML

N20588 001 Jun 10, 1996

TABLET; ORAL

RISPERDAL

JANSSEN PHARMA

0.25MG

N20272 008 May 10, 1999

0.5MG

N20272 007 Jan 27, 1999

+

1MG

N20272 001 Dec 29, 1993

2MG

N20272 002 Dec 29, 1993

3MG

N20272 003 Dec 29, 1993

4MG

N20272 004 Dec 29, 1993

TABLET, ORALLY DISINTEGRATING; ORAL

RISPERDAL

JANSSEN PHARMA

0.5MG

N21444 001 Apr 02, 2003

+

1MG

N21444 002 Apr 02, 2003

2MG

N21444 003 Apr 02, 2003

RITODRINE HYDROCHLORIDE

INJECTABLE; INJECTION

RITODRINE HYDROCHLORIDE

+ HOSPIRA

10MG/ML

N71618 001 Feb 28, 1991

+

15MG/ML

N71619 001 Feb 28, 1991

RITODRINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER

+ HOSPIRA

30MG/100ML

N71438 001 Jan 22, 1991

RITONAVIR

CAPSULE; ORAL

NORVIR

+ ABBOTT

100MG

N20945 001 Jun 29, 1999

SOLUTION; ORAL

NORVIR

+ ABBOTT

80MG/ML

N20659 001 Mar 01, 1996

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 312 (of 360)

RIVASTIGMINE TARTRATE

CAPSULE; ORAL

EXELON

+ NOVARTIS	EQ 1.5MG BASE	N20823 003	Apr 21, 2000
	EQ 3MG BASE	N20823 004	Apr 21, 2000
	EQ 4.5MG BASE	N20823 005	Apr 21, 2000
+	EQ 6MG BASE	N20823 006	Apr 21, 2000

SOLUTION; ORAL

EXELON

+ NOVARTIS	EQ 2MG BASE/ML	N21025 001	Apr 21, 2000
------------	----------------	------------	--------------

RIZATRIPTAN BENZOATE

TABLET; ORAL

MAXALT

MERCK

	EQ 5MG BASE	N20864 001	Jun 29, 1998
+	EQ 10MG BASE	N20864 002	Jun 29, 1998

TABLET, ORALLY DISINTEGRATING; ORAL

MAXALT-MLT

MERCK

	EQ 5MG BASE	N20865 001	Jun 29, 1998
+	EQ 10MG BASE	N20865 002	Jun 29, 1998

ROCURONIUM BROMIDE

INJECTABLE; INJECTION

ZEMURON (P/F)

+ ORGANON USA INC	50MG/5ML (10MG/ML)	N20214 001	Mar 17, 1994
+	100MG/10ML (10MG/ML)	N20214 003	Mar 17, 1994

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

REQUIP

+ GLAXOSMITHKLINE	EQ 0.25MG BASE	N20658 001	Sep 19, 1997
	EQ 0.5MG BASE	N20658 002	Sep 19, 1997
	EQ 1MG BASE	N20658 003	Sep 19, 1997
	EQ 2MG BASE	N20658 004	Sep 19, 1997
	EQ 3MG BASE	N20658 006	Jan 27, 1999
	EQ 4MG BASE	N20658 007	Jan 27, 1999
	EQ 5MG BASE	N20658 005	Sep 19, 1997

ROIIVACAINE HYDROCHLORIDE MONOHYDRATE

INJECTABLE; INJECTION

NAROPIN

ASTRAZENECA

	2MG/ML	N20533 001	Sep 24, 1996
	5MG/ML	N20533 003	Sep 24, 1996
	7.5MG/ML	N20533 004	Sep 24, 1996
+	10MG/ML	N20533 005	Sep 24, 1996

ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA

SB PHARMC

	EQ 2MG BASE	N21071 002	May 25, 1999
	EQ 4MG BASE	N21071 003	May 25, 1999
+	EQ 8MG BASE	N21071 004	May 25, 1999

ROSUVASTATIN CALCIUM

TABLET; ORAL

CRESTOR

ASTRAZENECA

	5MG	N21366 002	Aug 12, 2003
	10MG	N21366 003	Aug 12, 2003
	20MG	N21366 004	Aug 12, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 313 (of 360)

ROSUVASTATIN CALCIUMTABLET; ORAL
CRESTOR

+ ASTRAZENECA 40MG N21366 005 Aug 12, 2003

RUBIDIUM CHLORIDE RB-82INJECTABLE; INJECTION
CARDIOGEN-82
BRACCO

N/A N19414 001 Dec 29, 1989

SACROSIDASESOLUTION; ORAL
SUCRAID

+ QOL MEDCL 8,500 IU/ML N20772 001 Apr 09, 1998

SAFFLOWER OIL; SOYBEAN OILINJECTABLE; INJECTION
LIPOSYN II 10%

+ HOSPIRA 5%;5% (5GM/100ML) N18997 001 Aug 27, 1984

LIPOSYN II 20%

+ HOSPIRA 10%;10% (10GM/100ML) N18991 001 Aug 27, 1984

SALMETEROL XINAFOATEAEROSOL, METERED; INHALATION
SEREVENT

+ GLAXOSMITHKLINE EQ 0.021MG BASE/INH N20236 001 Feb 04, 1994

POWDER; INHALATION
SEREVENT

+ GLAXOSMITHKLINE EQ 0.046MG BASE/INH N20692 001 Sep 19, 1997

SAMARIUM SM 153 LEXIDRONAM PENTASODIUMINJECTABLE; INJECTION
QUADRAMET

+ CYTOGEN 50mCi/ML N20570 001 Mar 28, 1997

SAQUINAVIRCAPSULE; ORAL
FORTOVASE

+ HLR 200MG N20828 001 Nov 07, 1997

SAQUINAVIR MESYLATECAPSULE; ORAL
INVIRASE

+ HLR EQ 200MG BASE N20628 001 Dec 06, 1995

TABLET; ORAL
INVIRASE

+ ROCHE EQ 500MG BASE N21785 001 Dec 17, 2004

SCOPOLAMINEFILM, EXTENDED RELEASE; TRANSDERMAL
TRANSDERM SCOP

+ NOVARTIS 1MG/72HR N17874 001

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECOBARBITAL SODIUMAA EVERYLIFE 100MGN85895 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 314 (of 360)

SECOBARBITAL SODIUM

CAPSULE; ORAL

SECONAL SODIUM

<u>AA</u>	+	RANBAXY	<u>100MG</u>	<u>N86101</u>	<u>002</u>	Oct 03, 1983
	+		<u>50MG</u>	N86101	001	Oct 03, 1983

SECRETIN SYNTHETIC HUMANFOR SOLUTION; INTRAVENOUS
CHIRHOSTIM

+	CHIRHOCLIN	16UGM/VIAL	N21256	001	Apr 09, 2004
---	------------	------------	--------	-----	--------------

SECRETIN SYNTHETIC PORCINEFOR SOLUTION; INTRAVENOUS
SECREMAX

+	CHIRHOCLIN	16UGM/VIAL	N21136	001	Apr 04, 2002
---	------------	------------	--------	-----	--------------

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

ELDEPRYL

<u>AB</u>	+	SOMERSET	<u>5MG</u>	<u>N20647</u>	<u>001</u>	May 15, 1996
-----------	---	----------	------------	---------------	------------	--------------

SELEGILINE HYDROCHLORIDE

<u>AB</u>		AAIPHARMA LLC	<u>5MG</u>	<u>N75145</u>	<u>001</u>	Sep 15, 2003
<u>AB</u>		CLONMEL HLTHCARE	<u>5MG</u>	<u>N75352</u>	<u>001</u>	Nov 30, 1998
<u>AB</u>		TORPHARM	<u>5MG</u>	<u>N75321</u>	<u>001</u>	Dec 04, 1998

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

<u>AB</u>		ALPHAPHARM	<u>5MG</u>	<u>N74866</u>	<u>001</u>	Nov 26, 1997
<u>AB</u>	+	APOTEX	<u>5MG</u>	<u>N74871</u>	<u>001</u>	Jun 06, 1997
<u>AB</u>		CLONMEL HLTHCARE	<u>5MG</u>	<u>N74641</u>	<u>001</u>	Aug 02, 1996
<u>AB</u>		ENDO PHARMS	<u>5MG</u>	<u>N74565</u>	<u>001</u>	Aug 02, 1996
<u>AB</u>		IVAX PHARMS	<u>5MG</u>	<u>N74756</u>	<u>001</u>	Nov 25, 1998
<u>AB</u>		SIEGFRIED	<u>5MG</u>	<u>N74672</u>	<u>001</u>	Apr 01, 1997
<u>AB</u>		STASON	<u>5MG</u>	<u>N74912</u>	<u>001</u>	Apr 30, 1998
<u>AB</u>		TEVA	<u>5MG</u>	<u>N74537</u>	<u>001</u>	Aug 02, 1996
<u>AB</u>			<u>5MG</u>	<u>N74744</u>	<u>001</u>	Jan 27, 1997

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

SELENIUM SULFIDE

<u>AT</u>		ALPHARMA US PHARMS	<u>2.5%</u>	<u>N84394</u>	<u>001</u>	
<u>AT</u>		CLAY PARK	<u>2.5%</u>	<u>N89996</u>	<u>001</u>	Jan 10, 1991
<u>AT</u>		MORTON GROVE	<u>2.5%</u>	<u>N88228</u>	<u>001</u>	Sep 01, 1983

SELSUN

<u>AT</u>	+	CHATTEM	<u>2.5%</u>	<u>N07936</u>	<u>001</u>	
-----------	---	---------	-------------	---------------	------------	--

SERMORELIN ACETATEINJECTABLE; INJECTION
GEREF

+	SERONO	EQ 0.05MG BASE/AMP	N19863	001	Dec 28, 1990
---	--------	--------------------	--------	-----	--------------

SERTACONAZOLE NITRATE

CREAM; TOPICAL

ERTACZO

+	JOHNSON AND JOHNSON	2%	N21385	001	Dec 10, 2003
---	---------------------	----	--------	-----	--------------

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

ZOLOFT

+	PFIZER	EQ 20MG BASE/ML	N20990	001	Dec 07, 1999
---	--------	-----------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 315 (of 360)

SERTRALINE HYDROCHLORIDE

TABLET; ORAL

ZOLOFT

PFIZER

EQ 25MG BASE

N19839 005 Mar 06, 1996

EQ 50MG BASE

N19839 001 Dec 30, 1991

+

EQ 100MG BASE

N19839 002 Dec 30, 1991

SEVELAMER HYDROCHLORIDE

TABLET; ORAL

RENAGEL

GENZYME

400MG

N21179 001 Jul 12, 2000

+

800MG

N21179 002 Jul 12, 2000

SEVOFLURANE

LIQUID; INHALATION

SEVOFLURANEAN BAXTER HLTHCARE100%N75895 001 Jul 02, 2002ULTANEAN + ABBOTT100%N20478 001 Jun 07, 1995SIBUTRAMINE HYDROCHLORIDE

CAPSULE; ORAL

MERIDIA

ABBOTT

5MG

N20632 001 Nov 22, 1997

10MG

N20632 002 Nov 22, 1997

+

15MG

N20632 003 Nov 22, 1997

SILDENAFIL CITRATE

TABLET; ORAL

REVATIO

+ PFIZER

EQ 20MG BASE

N21845 001 Jun 03, 2005

VIAGRA

PFIZER IRELAND

EQ 25MG BASE

N20895 001 Mar 27, 1998

EQ 50MG BASE

N20895 002 Mar 27, 1998

+

EQ 100MG BASE

N20895 003 Mar 27, 1998

SILVER SULFADIAZINE

CREAM; TOPICAL

SILVADENEAB + KING PHARMS1%N17381 001SSDAB BASF1%N18578 001 Feb 25, 1982

SSD AF

BX BASF

1%

N18578 003 Jul 11, 1990

THERMAZENEAB KENDALL LP1%N18810 001 Dec 23, 1985SIMVASTATIN

TABLET; ORAL

ZOCOR

MERCK

5MG

N19766 001 Dec 23, 1991

10MG

N19766 002 Dec 23, 1991

20MG

N19766 003 Dec 23, 1991

40MG

N19766 004 Dec 23, 1991

+

80MG

N19766 005 Jul 10, 1998

PRESCRIPTION DRUG PRODUCT LIST

3 - 316 (of 360)

SINCALIDE

INJECTABLE; INJECTION KINEVAC			
+ BRACCO	0.005MG/VIAL	N17697	001

SIROLIMUS

SOLUTION; ORAL RAPAMUNE			
+ WYETH PHARMS INC	1MG/ML	N21083	001 Sep 15, 1999
TABLET; ORAL RAPAMUNE			
WYETH PHARMS INC	1MG	N21110	001 Aug 25, 2000
+	2MG	N21110	002 Aug 22, 2002

SODIUM ACETATE, ANHYDROUS

INJECTABLE; INJECTION SODIUM ACETATE IN PLASTIC CONTAINER			
+ HOSPIRA	2MEQ/ML	N18893	001 May 04, 1983

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; IV (INFUSION) AMMONUL			
+ UCYCLYD	10%;10% (5GM/50ML;5GM/50ML)	N20645	001 Feb 17, 2005

SODIUM BICARBONATE

INJECTABLE; INJECTION SODIUM BICARBONATE			
HOSPIRA	0.9MEQ/ML	N77394	001 Nov 09, 2005
+	1MEQ/ML	N77394	002 Nov 09, 2005

SODIUM BICARBONATE; TARTARIC ACID

GRANULE, EFFERVESCENT; ORAL BAROS			
LAFAYETTE PHARMS	460MG/GM;420MG/GM	N18509	001 Aug 07, 1985

SODIUM CHLORIDE

INJECTABLE; INJECTION			
<u>BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP	AM PHARM PARTNERS	9MG/ML	N88911 001 Feb 07, 1985
AP	+ HOSPIRA	9MG/ML	N18800 001 Oct 29, 1982
<u>SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER</u>			
AP	B BRAUN	450MG/100ML	N19635 001 Mar 09, 1988
AP	BAXTER HLTHCARE	450MG/100ML	N18016 001
AP	HOSPIRA	450MG/100ML	N18090 001
AP		450MG/100ML	N19759 001 Jun 08, 1988
<u>SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>			
AP	AM PHARM PARTNERS	9MG/ML	N88912 001 Jan 10, 1985
AP	B BRAUN	900MG/100ML	N17464 001
AP		900MG/100ML	N19635 002 Mar 09, 1988
AP	+ BAXTER HLTHCARE	9MG/ML	N16677 004 Oct 30, 1985
AP		9MG/ML	N20178 002 Dec 07, 1992
AP		900MG/100ML	N16677 001
AP		900MG/100ML	N20178 001 Dec 07, 1992
AP	HAEMONETICS	900MG/100ML	N76316 001 Oct 27, 2004
AP	HOSPIRA	9MG/ML	N18803 001 Oct 29, 1982
AP	+	9MG/ML	N19217 001 Jul 13, 1984
AP		9MG/ML	N19465 002 Jul 15, 1985
AP		900MG/100ML	N16366 001
AP		900MG/100ML	N19465 001 Jul 15, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 317 (of 360)

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AP</u>	HOSPIRA	<u>900MG/100ML</u>	<u>N19480</u>	<u>001</u>	Sep 17, 1985
	SODIUM CHLORIDE 3% IN PLASTIC CONTAINER				
	BAXTER HLTHCARE	3GM/100ML	N19022	001	Nov 01, 1983
	SODIUM CHLORIDE 5% IN PLASTIC CONTAINER				
	BAXTER HLTHCARE	5GM/100ML	N19022	002	Nov 01, 1983
	SODIUM CHLORIDE IN PLASTIC CONTAINER				
	HOSPIRA	2.5MEQ/ML	N18897	001	Jul 20, 1984

SOLUTION FOR SLUSH; IRRIGATION

SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER

	BAXTER HLTHCARE	900MG/100ML	N19319	002	May 17, 1985
--	-----------------	-------------	--------	-----	--------------

SOLUTION; IRRIGATION

SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

<u>AT</u>	BAXTER HLTHCARE	<u>450MG/100ML</u>	<u>N17864</u>	<u>001</u>	
<u>AT</u>	HOSPIRA	<u>450MG/100ML</u>	<u>N17670</u>	<u>001</u>	
<u>AT</u>		<u>450MG/100ML</u>	<u>N18380</u>	<u>001</u>	

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

<u>AT</u>	B BRAUN	<u>900MG/100ML</u>	<u>N16733</u>	<u>001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>900MG/100ML</u>	<u>N17427</u>	<u>001</u>	
<u>AT</u>		<u>900MG/100ML</u>	<u>N17867</u>	<u>001</u>	
<u>AT</u>	HOSPIRA	<u>900MG/100ML</u>	<u>N17514</u>	<u>001</u>	
<u>AT</u>		<u>900MG/100ML</u>	<u>N18314</u>	<u>001</u>	

SODIUM CHROMATE, CR-51

INJECTABLE; INJECTION

CHROMITOPES SODIUM

	BRACCO	200uCi/ML	N13993	001	
--	--------	-----------	--------	-----	--

SODIUM CHROMATE CR 51

	MALLINCKRODT	100uCi/ML	N16708	001	
--	--------------	-----------	--------	-----	--

SODIUM FERRIC GLUCONATE COMPLEX

INJECTABLE; INJECTION

FERRLECIT

	+ WATSON PHARMS	62.5MG/5ML	N20955	001	Feb 18, 1999
--	-----------------	------------	--------	-----	--------------

SODIUM IODIDE, I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

<u>AA</u>	GE HEALTHCARE	<u>100uCi</u>	<u>N17630</u>	<u>001</u>	
<u>AA</u>	MALLINCKRODT	<u>100uCi</u>	<u>N71909</u>	<u>001</u>	Feb 28, 1989
<u>AA</u>		<u>200uCi</u>	<u>N71910</u>	<u>001</u>	Feb 28, 1989
<u>AA</u>	SYNCOR PHARMS	<u>100uCi</u>	<u>N18671</u>	<u>001</u>	May 27, 1982
<u>AA</u>		<u>200uCi</u>	<u>N18671</u>	<u>002</u>	May 27, 1982

SOLUTION; ORAL

SODIUM IODIDE I 123

	GE HEALTHCARE	2mCi/ML	N17630	002	
--	---------------	---------	--------	-----	--

SODIUM IODIDE, I-131

CAPSULE; ORAL

IODOTOPE

	+ BRACCO	1-130mCi	N10929	001	
--	----------	----------	--------	-----	--

SODIUM IODIDE I 131

	+ MALLINCKRODT	15-100uCi	N16517	002	
--	----------------	-----------	--------	-----	--

	+	0.8-100mCi	N16517	001	
--	---	------------	--------	-----	--

SOLUTION; ORAL

SODIUM IODIDE I 131

	+ MALLINCKRODT	3.5-150mCi/VIAL	N16515	001	
--	----------------	-----------------	--------	-----	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 318 (of 360)

SODIUM IODIDE, I-131

SOLUTION; ORAL

SODIUM IODIDE I 131, KIT

+	DRAXIMAGE	1-250mCi/0.25ML	N21305	002	Jan 24, 2003
+		1-500mCi/0.5ML	N21305	003	Jan 24, 2003

SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER

<u>AP</u>	BAXTER HLTHCARE	1.87GM/100ML	N16692	001	
------------------	-----------------	--------------	--------	-----	--

<u>AP</u>	HOSPIRA	1.87GM/100ML	N18249	001	
------------------	---------	--------------	--------	-----	--

SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER

<u>AP</u>	B BRAUN	1.87GM/100ML	N20004	001	Apr 21, 1992
------------------	---------	--------------	--------	-----	--------------

SODIUM LACTATE IN PLASTIC CONTAINER

HOSPIRA	5MEQ/ML	N18947	001	Sep 05, 1984
---------	---------	--------	-----	--------------

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NITROPRESS

<u>AP</u>	+ HOSPIRA	25MG/ML	N71961	001	Aug 01, 1988
------------------	-----------	---------	--------	-----	--------------

+		50MG/VIAL	N70566	001	Jun 09, 1986
---	--	-----------	--------	-----	--------------

SODIUM NITROPRUSSIDE

<u>AP</u>	SICOR PHARMS	25MG/ML	N73465	001	Mar 30, 1992
------------------	--------------	---------	--------	-----	--------------

SODIUM OXYBATE

SOLUTION; ORAL

XYREM

+	ORPHAN MEDCL	500MG/ML	N21196	001	Jul 17, 2002
---	--------------	----------	--------	-----	--------------

SODIUM PHENYLBUTYRATE

POWDER; ORAL

BUPHENYL

+	MEDICIS	3GM/TEASPOONFUL	N20573	001	Apr 30, 1996
---	---------	-----------------	--------	-----	--------------

TABLET; ORAL

BUPHENYL

+	MEDICIS	500MG	N20572	001	May 13, 1996
---	---------	-------	--------	-----	--------------

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

TABLET; ORAL

VISICOL

+	SALIX PHARMS	0.398GM;1.102GM	N21097	001	Sep 21, 2000
---	--------------	-----------------	--------	-----	--------------

SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE; SODIUM PHOSPHATE, MONOBASIC, ANHYDROUS

INJECTABLE; INJECTION

SODIUM PHOSPHATES IN PLASTIC CONTAINER

HOSPIRA	142MG/ML;276MG/ML	N18892	001	May 10, 1983
---------	-------------------	--------	-----	--------------

SODIUM PHOSPHATE, P-32

SOLUTION; INJECTION, ORAL

SODIUM PHOSPHATE P 32

MALLINCKRODT	0.67mCi/ML	N11777	001	
--------------	------------	--------	-----	--

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

KAYEXALATE

<u>AA</u>	+ SANOFI SYNTHELABO	453.6GM/BOT	N11287	001	
------------------	---------------------	-------------	--------	-----	--

KIONEX

<u>AA</u>	PADDOCK	454GM/BOT	N40029	001	Feb 06, 1998
------------------	---------	-----------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 319 (of 360)

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

<u>AA</u>	CAROLINA MEDCL	<u>454GM/BOT</u>	<u>N89910</u>	<u>001</u>	Jan 19, 1989
-----------	----------------	------------------	---------------	------------	--------------

SUSPENSION; ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

<u>AA</u>	ROXANE	<u>15GM/60ML</u>	<u>N89049</u>	<u>001</u>	Nov 17, 1986
-----------	--------	------------------	---------------	------------	--------------

SPS

<u>AA</u>	+ CAROLINA MEDCL	<u>15GM/60ML</u>	<u>N87859</u>	<u>001</u>	Dec 08, 1982
-----------	------------------	------------------	---------------	------------	--------------

SODIUM TETRADECYL SULFATE

INJECTABLE; INJECTION

SOTRADECOL

	BIONICHE PHARM USA	20MG/2ML (10MG/ML)	N40541	001	Nov 12, 2004
--	--------------------	--------------------	--------	-----	--------------

+		60MG/2ML (30MG/ML)	N40541	002	Nov 12, 2004
---	--	--------------------	--------	-----	--------------

SOLIFENACIN SUCCINATE

TABLET; ORAL

VESICARE

	ASTELLAS	5MG	N21518	001	Nov 19, 2004
--	----------	-----	--------	-----	--------------

+		10MG	N21518	002	Nov 19, 2004
---	--	------	--------	-----	--------------

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

GENOTROPIN

+	PHARMACIA AND UPJOHN	5.8MG/VIAL	N20280	006	Aug 24, 1995
---	----------------------	------------	--------	-----	--------------

+		13.8MG/VIAL	N20280	007	Oct 23, 1996
---	--	-------------	--------	-----	--------------

GENOTROPIN PRESERVATIVE FREE

	PHARMACIA AND UPJOHN	0.2MG/VIAL	N20280	001	Jan 27, 1998
--	----------------------	------------	--------	-----	--------------

		0.4MG/VIAL	N20280	002	Jan 27, 1998
--	--	------------	--------	-----	--------------

		0.6MG/VIAL	N20280	003	Jan 27, 1998
--	--	------------	--------	-----	--------------

		0.8MG/VIAL	N20280	005	Jan 27, 1998
--	--	------------	--------	-----	--------------

		1MG/VIAL	N20280	008	Jan 27, 1998
--	--	----------	--------	-----	--------------

		1.2MG/VIAL	N20280	009	Jan 27, 1998
--	--	------------	--------	-----	--------------

		1.4MG/VIAL	N20280	010	Jan 27, 1998
--	--	------------	--------	-----	--------------

		1.5MG/VIAL	N20280	004	Aug 24, 1995
--	--	------------	--------	-----	--------------

		1.6MG/VIAL	N20280	011	Jan 27, 1998
--	--	------------	--------	-----	--------------

		1.8MG/VIAL	N20280	012	Jan 27, 1998
--	--	------------	--------	-----	--------------

+		2MG/VIAL	N20280	013	Jan 27, 1998
---	--	----------	--------	-----	--------------

HUMATROPE

BX	+	LILLY	5MG/VIAL	N19640	004	Mar 08, 1987
----	---	-------	----------	--------	-----	--------------

BX			6MG/VIAL	N19640	005	Feb 04, 1999
----	--	--	----------	--------	-----	--------------

+		12MG/VIAL	N19640	006	Feb 04, 1999
---	--	-----------	--------	-----	--------------

+		24MG/VIAL	N19640	007	Feb 04, 1999
---	--	-----------	--------	-----	--------------

NORDITROPIN

BX		NOVO NORDISK INC	4MG/VIAL	N19721	001	May 08, 1995
----	--	------------------	----------	--------	-----	--------------

		5MG/1.5ML	N21148	001	Jun 20, 2000
--	--	-----------	--------	-----	--------------

		8MG/VIAL	N19721	002	May 08, 1995
--	--	----------	--------	-----	--------------

		10MG/1.5ML	N21148	002	Jun 20, 2000
--	--	------------	--------	-----	--------------

+		15MG/1.5ML	N21148	003	Jun 20, 2000
---	--	------------	--------	-----	--------------

NORDITROPIN NORDIFLEX

		NOVO NORDISK INC	5MG/1.5ML	N21148	004	Oct 01, 2004
--	--	------------------	-----------	--------	-----	--------------

		10MG/1.5ML	N21148	005	Oct 01, 2004
--	--	------------	--------	-----	--------------

		15MG/1.5ML	N21148	006	Oct 01, 2004
--	--	------------	--------	-----	--------------

NUTROPIN

BX		GENENTECH	5MG/VIAL	N20168	001	Nov 17, 1993
----	--	-----------	----------	--------	-----	--------------

+			10MG/VIAL	N20168	002	Nov 17, 1993
---	--	--	-----------	--------	-----	--------------

NUTROPIN AQ

+		GENENTECH	5MG/ML	N20522	001	Dec 29, 1995
---	--	-----------	--------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 320 (of 360)

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION					
NUTROPIN AQ PEN					
BX	+	GENENTECH	5MG/ML	N20522 002	Apr 22, 2002
NUTROPIN DEPOT					
	+	GENENTECH	13.5MG/VIAL	N21075 001	Dec 22, 1999
	+		18MG/VIAL	N21075 002	Dec 22, 1999
	+		22.5MG/VIAL	N21075 003	Dec 22, 1999
SAIZEN					
BX		SERONO	5MG/VIAL	N19764 002	Oct 08, 1996
	+		8.8MG/VIAL	N19764 003	Aug 29, 2000
SEROSTIM					
BX		SERONO	4MG/VIAL	N20604 003	Jul 25, 1997
BX			5MG/VIAL	N20604 002	Aug 23, 1996
BX			6MG/VIAL	N20604 001	Aug 23, 1996
TEV-TROPIN					
BX	+	FERRING	5MG/ML	N19774 002	Jan 04, 2002
ZORBTIVE					
	+	SERONO INC	8.8MG/VIAL	N21597 004	Dec 01, 2003

SORAFENIB TOSYLATE

TABLET; ORAL					
NEXAVAR					
	+	BAYER PHARMS	EQ 200MG BASE	N21923 001	Dec 20, 2005

SORBITOL

SOLUTION; IRRIGATION					
SORBITOL 3% IN PLASTIC CONTAINER					
		BAXTER HLTHCARE	3GM/100ML	N17863 001	
SORBITOL 3.3% IN PLASTIC CONTAINER					
		B BRAUN	3.3GM/100ML	N16741 001	

SOTALOL HYDROCHLORIDE

TABLET; ORAL					
<u>BETAPACE</u>					
<u>AB1</u>		BERLEX	<u>80MG</u>	<u>N19865 001</u>	Oct 30, 1992
<u>AB1</u>			<u>120MG</u>	<u>N19865 005</u>	Apr 20, 1994
<u>AB1</u>	+		<u>160MG</u>	<u>N19865 002</u>	Oct 30, 1992
<u>AB1</u>			<u>240MG</u>	<u>N19865 003</u>	Oct 30, 1992
<u>BETAPACE AF</u>					
<u>AB2</u>		BERLEX	<u>80MG</u>	<u>N21151 001</u>	Feb 22, 2000
<u>AB2</u>			<u>120MG</u>	<u>N21151 002</u>	Feb 22, 2000
<u>AB2</u>	+		<u>160MG</u>	<u>N21151 003</u>	Feb 22, 2000
<u>SORINE</u>					
<u>AB1</u>		UPSHER SMITH	<u>80MG</u>	<u>N75500 001</u>	Apr 27, 2001
<u>AB1</u>			<u>120MG</u>	<u>N75500 004</u>	Apr 27, 2001
<u>AB1</u>			<u>160MG</u>	<u>N75500 002</u>	Apr 27, 2001
<u>AB1</u>			<u>240MG</u>	<u>N75500 003</u>	Apr 27, 2001
<u>SOTALOL HYDROCHLORIDE</u>					
<u>AB1</u>		APOTEX	<u>80MG</u>	<u>N76140 001</u>	Sep 26, 2002
<u>AB1</u>			<u>120MG</u>	<u>N76140 002</u>	Sep 26, 2002
<u>AB1</u>			<u>160MG</u>	<u>N76140 003</u>	Sep 26, 2002
<u>AB1</u>			<u>240MG</u>	<u>N76140 004</u>	Sep 26, 2002
<u>AB1</u>		GENPHARM	<u>80MG</u>	<u>N75237 001</u>	May 01, 2000
<u>AB1</u>			<u>120MG</u>	<u>N75237 002</u>	May 01, 2000
<u>AB1</u>			<u>160MG</u>	<u>N75237 003</u>	May 01, 2000
<u>AB1</u>			<u>240MG</u>	<u>N75237 004</u>	May 01, 2000
<u>AB2</u>			<u>80MG</u>	<u>N77070 001</u>	Nov 04, 2005
<u>AB2</u>			<u>120MG</u>	<u>N77070 002</u>	Nov 04, 2005
<u>AB2</u>			<u>160MG</u>	<u>N77070 003</u>	Nov 04, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 321 (of 360)

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

<u>AB1</u>	IMPAX PHARMS	<u>80MG</u>	<u>N75663</u>	<u>001</u>	Nov 07, 2000
<u>AB1</u>		<u>120MG</u>	<u>N75663</u>	<u>002</u>	Nov 07, 2000
<u>AB1</u>		<u>160MG</u>	<u>N75663</u>	<u>003</u>	Nov 07, 2000
<u>AB1</u>		<u>240MG</u>	<u>N75663</u>	<u>004</u>	Nov 07, 2000
<u>AB1</u>	MUTUAL PHARM	<u>80MG</u>	<u>N75515</u>	<u>001</u>	Oct 15, 2001
<u>AB1</u>		<u>120MG</u>	<u>N75515</u>	<u>004</u>	Oct 15, 2001
<u>AB1</u>		<u>160MG</u>	<u>N75515</u>	<u>002</u>	Oct 15, 2001
<u>AB1</u>		<u>240MG</u>	<u>N75515</u>	<u>003</u>	Oct 15, 2001
<u>AB2</u>		<u>80MG</u>	<u>N76576</u>	<u>001</u>	Apr 08, 2004
<u>AB2</u>		<u>120MG</u>	<u>N76576</u>	<u>002</u>	Apr 08, 2004
<u>AB2</u>		<u>160MG</u>	<u>N76576</u>	<u>003</u>	Apr 08, 2004
<u>AB1</u>	MYLAN	<u>80MG</u>	<u>N75725</u>	<u>001</u>	Dec 19, 2000
<u>AB1</u>		<u>120MG</u>	<u>N75725</u>	<u>002</u>	Dec 19, 2000
<u>AB1</u>		<u>160MG</u>	<u>N75725</u>	<u>003</u>	Dec 19, 2000
<u>AB1</u>		<u>240MG</u>	<u>N75725</u>	<u>004</u>	Dec 19, 2000
<u>AB1</u>	SANDOZ	<u>80MG</u>	<u>N75366</u>	<u>001</u>	May 01, 2000
<u>AB1</u>		<u>120MG</u>	<u>N75366</u>	<u>002</u>	May 01, 2000
<u>AB1</u>		<u>160MG</u>	<u>N75366</u>	<u>003</u>	May 01, 2000
<u>AB1</u>		<u>240MG</u>	<u>N75366</u>	<u>004</u>	May 01, 2000
<u>AB1</u>	TEVA	<u>80MG</u>	<u>N75429</u>	<u>001</u>	May 01, 2000
<u>AB1</u>		<u>120MG</u>	<u>N75429</u>	<u>002</u>	May 01, 2000
<u>AB1</u>		<u>160MG</u>	<u>N75429</u>	<u>003</u>	May 01, 2000
<u>AB1</u>		<u>240MG</u>	<u>N75429</u>	<u>004</u>	May 01, 2000
<u>AB2</u>		<u>80MG</u>	<u>N76883</u>	<u>001</u>	Jul 26, 2004
<u>AB2</u>		<u>120MG</u>	<u>N76883</u>	<u>002</u>	Jul 26, 2004
<u>AB2</u>		<u>160MG</u>	<u>N76883</u>	<u>003</u>	Jul 26, 2004
<u>AB2</u>	TORPHARM	<u>80MG</u>	<u>N76214</u>	<u>001</u>	Aug 27, 2003
<u>AB2</u>		<u>120MG</u>	<u>N76214</u>	<u>002</u>	Aug 27, 2003
<u>AB2</u>		<u>160MG</u>	<u>N76214</u>	<u>003</u>	Aug 27, 2003
<u>AB1</u>	VINTAGE PHARMS	<u>80MG</u>	<u>N75563</u>	<u>001</u>	Nov 07, 2003
<u>AB1</u>		<u>120MG</u>	<u>N75563</u>	<u>002</u>	Nov 07, 2003
<u>AB1</u>		<u>160MG</u>	<u>N75563</u>	<u>003</u>	Nov 07, 2003
<u>AB1</u>		<u>240MG</u>	<u>N75563</u>	<u>004</u>	Nov 07, 2003
<u>AB1</u>	WATSON LABS	<u>80MG</u>	<u>N75238</u>	<u>001</u>	Jul 13, 2000
<u>AB1</u>		<u>120MG</u>	<u>N75238</u>	<u>002</u>	Jul 13, 2000
<u>AB1</u>		<u>160MG</u>	<u>N75238</u>	<u>003</u>	Jul 13, 2000
<u>AB1</u>		<u>240MG</u>	<u>N75238</u>	<u>004</u>	Jul 13, 2000

SOYBEAN OIL

INJECTABLE; INJECTION

	<u>INTRALIPID 10%</u>				
<u>AP</u>	+ FRESenius	<u>10%</u>	<u>N17643</u>	<u>001</u>	
	<u>INTRALIPID 20%</u>				
<u>AP</u>	+ FRESenius	<u>20%</u>	<u>N18449</u>	<u>001</u>	
<u>AP</u>	+	<u>20%</u>	<u>N20248</u>	<u>001</u>	Aug 07, 1996
	<u>INTRALIPID 30%</u>				
<u>AP</u>	+ FRESenius	<u>30%</u>	<u>N19942</u>	<u>001</u>	Dec 30, 1993
	<u>LIPOSYN III 10%</u>				
<u>AP</u>	+ HOSPIRA	<u>10%</u>	<u>N18969</u>	<u>001</u>	Sep 24, 1984
	<u>LIPOSYN III 20%</u>				
<u>AP</u>	+ HOSPIRA	<u>20%</u>	<u>N18970</u>	<u>001</u>	Sep 25, 1984
	<u>LIPOSYN III 30%</u>				
<u>AP</u>	+ HOSPIRA	<u>30%</u>	<u>N20181</u>	<u>001</u>	Jan 13, 1998
	<u>NUTRILIPID 10%</u>				
<u>AP</u>	+ B BRAUN	<u>10%</u>	<u>N19531</u>	<u>001</u>	May 28, 1993
	<u>NUTRILIPID 20%</u>				
<u>AP</u>	+ B BRAUN	<u>20%</u>	<u>N19531</u>	<u>002</u>	May 28, 1993

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 322 (of 360)

SOYBEAN OIL

INJECTABLE; INJECTION

SOYACAL 10%

<u>AP</u>	+	ALPHA THERA	<u>10%</u>	<u>N18465</u>	<u>001</u>	Jun 29, 1983
-----------	---	-------------	------------	---------------	------------	--------------

SOYACAL 20%

<u>AP</u>	+	ALPHA THERA	<u>20%</u>	<u>N18786</u>	<u>001</u>	Jun 29, 1983
-----------	---	-------------	------------	---------------	------------	--------------

SPECTINOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

TROBICIN

+	PHARMACIA AND UPJOHN	EQ 2GM BASE/VIAL	N50347	001	
---	----------------------	------------------	--------	-----	--

SPIRONOLACTONE

TABLET; ORAL

ALDACTONE

<u>AB</u>		GD SEARLE LLC	<u>25MG</u>	<u>N12151</u>	<u>009</u>	Dec 30, 1983
-----------	--	---------------	-------------	---------------	------------	--------------

<u>AB</u>			<u>50MG</u>	<u>N12151</u>	<u>008</u>	Dec 30, 1982
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>	+		<u>100MG</u>	<u>N12151</u>	<u>010</u>	Dec 30, 1983
-----------	---	--	--------------	---------------	------------	--------------

SPIRONOLACTONE

<u>AB</u>		MUTUAL PHARM	<u>25MG</u>	<u>N87265</u>	<u>001</u>	
-----------	--	--------------	-------------	---------------	------------	--

<u>AB</u>			<u>25MG</u>	<u>N89424</u>	<u>001</u>	Jul 23, 1986
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>50MG</u>	<u>N89424</u>	<u>002</u>	Aug 11, 1999
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>100MG</u>	<u>N89424</u>	<u>003</u>	Aug 11, 1999
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		MYLAN	<u>25MG</u>	<u>N40424</u>	<u>001</u>	Aug 20, 2001
-----------	--	-------	-------------	---------------	------------	--------------

<u>AB</u>			<u>25MG</u>	<u>N87086</u>	<u>001</u>	
-----------	--	--	-------------	---------------	------------	--

<u>AB</u>			<u>50MG</u>	<u>N40424</u>	<u>002</u>	Aug 20, 2001
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>100MG</u>	<u>N40424</u>	<u>003</u>	Aug 20, 2001
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		PUREPAC PHARM	<u>50MG</u>	<u>N40353</u>	<u>001</u>	Jul 29, 1999
-----------	--	---------------	-------------	---------------	------------	--------------

<u>AB</u>			<u>100MG</u>	<u>N40353</u>	<u>002</u>	Jul 29, 1999
-----------	--	--	--------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>25MG</u>	<u>N86809</u>	<u>001</u>	
-----------	--	--------	-------------	---------------	------------	--

STAVUDINE

CAPSULE; ORAL

ZERIT

	BRISTOL MYERS SQUIBB	15MG	N20412	002	Jun 24, 1994
--	----------------------	------	--------	-----	--------------

		20MG	N20412	003	Jun 24, 1994
--	--	------	--------	-----	--------------

		30MG	N20412	004	Jun 24, 1994
--	--	------	--------	-----	--------------

+		40MG	N20412	005	Jun 24, 1994
---	--	------	--------	-----	--------------

FOR SOLUTION; ORAL

ZERIT

+	BRISTOL MYERS SQUIBB	1MG/ML	N20413	001	Sep 06, 1996
---	----------------------	--------	--------	-----	--------------

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

+	PFIZER	EQ 1GM BASE/2.5ML	N60111	001	
---	--------	-------------------	--------	-----	--

+	PHARMA TEK	EQ 1GM BASE/VIAL	N64210	001	Jun 30, 1998
---	------------	------------------	--------	-----	--------------

STREPTOZOCIN

INJECTABLE; INJECTION

ZANOSAR

+	SICOR PHARMS	1GM/VIAL	N50577	001	May 07, 1982
---	--------------	----------	--------	-----	--------------

STRONTIUM CHLORIDE, SR-89

INJECTABLE; INJECTION

METASTRON

<u>AP</u>	+	GE HEALTHCARE	<u>1mCi/ML</u>	<u>N20134</u>	<u>001</u>	Jun 18, 1993
-----------	---	---------------	----------------	---------------	------------	--------------

STRONTIUM CHLORIDE SR-89

<u>AP</u>		BIO NUCLEONICS	<u>1mCi/ML</u>	<u>N75941</u>	<u>001</u>	Jan 06, 2003
-----------	--	----------------	----------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 323 (of 360)

SUCCIMER

CAPSULE; ORAL
CHEMET

+ OVATION PHARMS	100MG	N19998 002	Jan 30, 1991
------------------	-------	------------	--------------

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION

<u>ANECTINE</u>			
<u>AP</u> + SABEX 2002	<u>20MG/ML</u>	<u>N08453</u>	<u>002</u>
<u>QUELICIN</u>			
<u>AP</u> + HOSPIRA	<u>20MG/ML</u>	<u>N08845</u>	<u>006</u>
<u>QUELICIN PRESERVATIVE FREE</u>			
<u>AP</u> + HOSPIRA	<u>20MG/ML</u>	<u>N08845</u>	<u>001</u>
+	100MG/ML	N08845	004
<u>SUCCINYLCHOLINE CHLORIDE</u>			
<u>AP</u> ORGANON USA INC	<u>20MG/ML</u>	<u>N80997</u>	<u>001</u>

SUCRALFATE

SUSPENSION; ORAL
CARAFATE

+ AXCAN SCANDIPHARM	1GM/10ML	N19183 001	Dec 16, 1993
---------------------	----------	------------	--------------

TABLET; ORAL

<u>CARAFATE</u>			
<u>AB</u> + AXCAN SCANDIPHARM	<u>1GM</u>	<u>N18333</u>	<u>001</u>
<u>SUCRALFATE</u>			
<u>AB</u> RATIOPHARM	<u>1GM</u>	<u>N74415</u>	<u>001</u> Jun 08, 1998
<u>AB</u> TEVA	<u>1GM</u>	<u>N70848</u>	<u>001</u> Mar 29, 1996

SUFENTANIL CITRATE

INJECTABLE; INJECTION

<u>SUFENTA</u>			
<u>AP</u> + AKORN	<u>EQ 0.05MG BASE/ML</u>	<u>N19050</u>	<u>001</u> May 04, 1984
<u>SUFENTANIL CITRATE</u>			
<u>AP</u> BAXTER HLTHCARE	<u>EQ 0.05MG BASE/ML</u>	<u>N74413</u>	<u>001</u> Dec 15, 1995
<u>SUFENTANIL CITRATE</u>			
<u>AP</u> HOSPIRA	<u>EQ 0.05MG BASE/ML</u>	<u>N74534</u>	<u>001</u> Dec 11, 1996

SULCONAZOLE NITRATE

CREAM; TOPICAL
EXELDERM

+ WESTWOOD SQUIBB	1%	N18737 001	Feb 28, 1989
-------------------	----	------------	--------------

SULFACETAMIDE SODIUM

LOTION; TOPICAL
KLARON

+ DERMIK LABS	10%	N19931 001	Dec 23, 1996
---------------	-----	------------	--------------

OINTMENT; OPHTHALMIC

<u>CETAMIDE</u>			
<u>AT</u> + ALCON	<u>10%</u>	<u>N80021</u>	<u>001</u>
<u>SULFACETAMIDE SODIUM</u>			
<u>AT</u> ALTANA	<u>10%</u>	<u>N80029</u>	<u>001</u>

SOLUTION/DROPS; OPHTHALMIC

<u>BLEPH-10</u>			
<u>AT</u> + ALLERGAN	<u>10%</u>	<u>N80028</u>	<u>001</u>
<u>BLEPH-30</u>			
<u>AT</u> + ALLERGAN	<u>30%</u>	<u>N80028</u>	<u>002</u>
<u>ISOPTO CETAMIDE</u>			
<u>AT</u> + ALCON	<u>15%</u>	<u>N80020</u>	<u>002</u>

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 324 (of 360)

SULFACETAMIDE SODIUM

SOLUTION/DROPS; OPHTHALMIC

<u>OCUSULF-10</u>				
<u>AT</u>	MIZA PHARMS USA	<u>10%</u>	<u>N80660</u>	<u>001</u>
<u>SULF-10</u>				
<u>AT</u>	NOVARTIS	<u>10%</u>	<u>N80025</u>	<u>001</u>
<u>SULFACEL-15</u>				
<u>AT</u>	OPTOPICS	<u>15%</u>	<u>N80024</u>	<u>001</u>
<u>SULFACETAMIDE SODIUM</u>				
<u>AT</u>	AKORN	<u>10%</u>	<u>N40215</u>	<u>001</u> May 25, 1999
<u>AT</u>		<u>30%</u>	<u>N40216</u>	<u>001</u> May 25, 1999
<u>AT</u>	ALCON UNIVERSAL	<u>10%</u>	<u>N89560</u>	<u>001</u> Oct 18, 1988
<u>AT</u>	BAUSCH AND LOMB	<u>10%</u>	<u>N40066</u>	<u>001</u> Dec 28, 1994
<u>AT</u>	STERIS	<u>30%</u>	<u>N89068</u>	<u>001</u> May 05, 1987

SULFADIAZINE

TABLET; ORAL
SULFADIAZINE
+ SANDOZ

500MG N40091 001 Jul 29, 1994

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

<u>BACTRIM</u>				
<u>AP</u>	+ MUTUAL PHARM	<u>80MG/ML;16MG/ML</u>	<u>N18374</u>	<u>001</u>
<u>SULFAMETHOXAZOLE AND TRIMETHOPRIM</u>				
<u>AP</u>	BAXTER HLTHCARE	<u>80MG/ML;16MG/ML</u>	<u>N70628</u>	<u>001</u> Dec 29, 1987
<u>AP</u>	HOSPIRA	<u>80MG/ML;16MG/ML</u>	<u>N73199</u>	<u>001</u> Sep 11, 1992
<u>AP</u>	SICOR PHARMS	<u>80MG/ML;16MG/ML</u>	<u>N73303</u>	<u>001</u> Oct 31, 1991

SUSPENSION; ORAL

<u>SEPTRA</u>				
<u>AB</u>	+ MONARCH PHARMS	<u>200MG/5ML;40MG/5ML</u>	<u>N17598</u>	<u>001</u>
<u>SEPTRA GRAPE</u>				
<u>AB</u>	MONARCH PHARMS	<u>200MG/5ML;40MG/5ML</u>	<u>N17598</u>	<u>002</u> Feb 12, 1986
<u>SULFAMETHOXAZOLE AND TRIMETHOPRIM</u>				
<u>AB</u>	HI TECH PHARMA	<u>200MG/5ML;40MG/5ML</u>	<u>N74650</u>	<u>001</u> Dec 29, 1997
<u>AB</u>	TEVA	<u>200MG/5ML;40MG/5ML</u>	<u>N18812</u>	<u>002</u> Jun 10, 1983
<u>AB</u>		<u>200MG/5ML;40MG/5ML</u>	<u>N18812</u>	<u>001</u> Jan 28, 1983
<u>SULFATRIM</u>				
<u>AB</u>	ALPHARMA US PHARMS	<u>200MG/5ML;40MG/5ML</u>	<u>N18615</u>	<u>002</u> Jan 07, 1983
<u>SULFATRIM PEDIATRIC</u>				
<u>AB</u>	ALPHARMA US PHARMS	<u>200MG/5ML;40MG/5ML</u>	<u>N18615</u>	<u>001</u> Jan 07, 1983

TABLET; ORAL

<u>BACTRIM</u>				
<u>AB</u>	MUTUAL PHARM	<u>400MG;80MG</u>	<u>N17377</u>	<u>001</u>
<u>BACTRIM DS</u>				
<u>AB</u>	+ MUTUAL PHARM	<u>800MG;160MG</u>	<u>N17377</u>	<u>002</u>
<u>COTRIM</u>				
<u>AB</u>	TEVA	<u>400MG;80MG</u>	<u>N70034</u>	<u>001</u> May 16, 1985
<u>COTRIM D.S.</u>				
<u>AB</u>	TEVA	<u>800MG;160MG</u>	<u>N70048</u>	<u>001</u> Mar 18, 1985
<u>SEPTRA</u>				
<u>AB</u>	MONARCH PHARMS	<u>400MG;80MG</u>	<u>N17376</u>	<u>001</u>
<u>SEPTRA DS</u>				
<u>AB</u>	MONARCH PHARMS	<u>800MG;160MG</u>	<u>N17376</u>	<u>002</u>
<u>SULFAMETHOPRIM</u>				
<u>AB</u>	PAR PHARM	<u>400MG;80MG</u>	<u>N70022</u>	<u>001</u> Feb 15, 1985
<u>SULFAMETHOXAZOLE AND TRIMETHOPRIM</u>				
<u>AB</u>	INTERPHARM	<u>400MG;80MG</u>	<u>N76899</u>	<u>001</u> Jan 27, 2005
<u>AB</u>		<u>800MG;160MG</u>	<u>N76899</u>	<u>002</u> Jan 27, 2005
<u>AB</u>	MUTUAL PHARM	<u>400MG;80MG</u>	<u>N71016</u>	<u>001</u> Aug 25, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 325 (of 360)

SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM

<u>AB</u>	MUTUAL PHARM	<u>800MG;160MG</u>	<u>N71017</u>	<u>001</u>	Aug 25, 1986
<u>AB</u>	PLIVA	<u>400MG;80MG</u>	<u>N70215</u>	<u>001</u>	Sep 10, 1985
<u>AB</u>		<u>800MG;160MG</u>	<u>N70216</u>	<u>001</u>	Sep 10, 1985
<u>AB</u>	SANDOZ	<u>800MG;160MG</u>	<u>N70890</u>	<u>001</u>	Nov 13, 1986
<u>AB</u>	VISTA PHARMS	<u>400MG;80MG</u>	<u>N76817</u>	<u>001</u>	Oct 07, 2005
<u>AB</u>		<u>800MG;160MG</u>	<u>N76817</u>	<u>002</u>	Oct 07, 2005
<u>AB</u>	WATSON LABS	<u>400MG;80MG</u>	<u>N18852</u>	<u>001</u>	May 09, 1983

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

<u>AB</u>	SANDOZ	<u>800MG;160MG</u>	<u>N18598</u>	<u>004</u>	May 19, 1982
<u>AB</u>	TEVA	<u>800MG;160MG</u>	<u>N70037</u>	<u>001</u>	Jun 02, 1987
<u>AB</u>	WATSON LABS	<u>800MG;160MG</u>	<u>N18854</u>	<u>001</u>	May 09, 1983

SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH

<u>AB</u>	PLANTEX	<u>400MG;80MG</u>	<u>N70030</u>	<u>001</u>	Jun 02, 1987
-----------	---------	-------------------	---------------	------------	--------------

SULFASALAZINE

SUSPENSION; ORAL

AZULFIDINE

+	PHARMACIA AND UPJOHN	250MG/5ML	N86983	001	
---	----------------------	-----------	--------	-----	--

TABLET; ORAL

AZULFIDINE

<u>AB</u>	+	PHARMACIA AND UPJOHN	<u>500MG</u>	<u>N07073</u>	<u>001</u>
-----------	---	----------------------	--------------	---------------	------------

SULFASALAZINE

<u>AB</u>	MUTUAL PHARM	<u>500MG</u>	<u>N89590</u>	<u>001</u>	Oct 19, 1987
<u>AB</u>	VINTAGE PHARMS	<u>500MG</u>	<u>N40349</u>	<u>001</u>	Jan 11, 2002
<u>AB</u>	WATSON LABS	<u>500MG</u>	<u>N85828</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N87197</u>	<u>001</u>	

TABLET, DELAYED RELEASE; ORAL

AZULFIDINE EN-TABS

<u>AB</u>	+	PHARMACIA AND UPJOHN	<u>500MG</u>	<u>N07073</u>	<u>002</u>	Apr 06, 1983
-----------	---	----------------------	--------------	---------------	------------	--------------

SULFASALAZINE

<u>AB</u>	VINTAGE PHARMS	<u>500MG</u>	<u>N75339</u>	<u>001</u>	Jan 11, 2002
-----------	----------------	--------------	---------------	------------	--------------

SULFINPYRAZONE

CAPSULE; ORAL

ANTURANE

<u>AB</u>	+	NOVARTIS	<u>200MG</u>	<u>N11556</u>	<u>004</u>
-----------	---	----------	--------------	---------------	------------

SULFINPYRAZONE

<u>AB</u>	BARR	<u>200MG</u>	<u>N87666</u>	<u>001</u>	Sep 17, 1982
<u>AB</u>	IVAX PHARMS	<u>200MG</u>	<u>N87770</u>	<u>001</u>	Nov 19, 1982
<u>AB</u>	PAR PHARM	<u>200MG</u>	<u>N88934</u>	<u>001</u>	Sep 06, 1985

TABLET; ORAL

ANTURANE

<u>AB</u>	+	NOVARTIS	<u>100MG</u>	<u>N11556</u>	<u>003</u>
-----------	---	----------	--------------	---------------	------------

SULFINPYRAZONE

<u>AB</u>	BARR	<u>100MG</u>	<u>N87665</u>	<u>001</u>	Sep 17, 1982
<u>AB</u>	PAR PHARM	<u>100MG</u>	<u>N88933</u>	<u>001</u>	Sep 06, 1985

SULFISOXAZOLE

TABLET; ORAL

SULFISOXAZOLE

+	IVAX PHARMS	500MG	N80142	001	
---	-------------	-------	--------	-----	--

SULFISOXAZOLE ACETYL

SUSPENSION; ORAL

GANTRISIN PEDIATRIC

+	ROCHE	EQ 500MG BASE/5ML	N09182	004	
---	-------	-------------------	--------	-----	--

PRESCRIPTION DRUG PRODUCT LIST

3 - 326 (of 360)

SULINDAC

TABLET; ORAL				
<u>CLINORIL</u>				
<u>AB</u>	MERCK	<u>150MG</u>	<u>N17911</u>	<u>001</u>
<u>AB</u>	+	<u>200MG</u>	<u>N17911</u>	<u>002</u>
<u>SULINDAC</u>				
<u>AB</u>	MUTUAL PHARM	<u>150MG</u>	<u>N72050</u>	<u>001</u> Apr 17, 1991
<u>AB</u>		<u>200MG</u>	<u>N72051</u>	<u>001</u> Apr 17, 1991
<u>AB</u>	MYLAN	<u>150MG</u>	<u>N73038</u>	<u>001</u> Jun 22, 1993
<u>AB</u>		<u>200MG</u>	<u>N73039</u>	<u>001</u> Jun 22, 1993
<u>AB</u>	SANDOZ	<u>150MG</u>	<u>N72712</u>	<u>001</u> Aug 30, 1991
<u>AB</u>		<u>200MG</u>	<u>N72713</u>	<u>001</u> Aug 30, 1991
<u>AB</u>	TEVA	<u>150MG</u>	<u>N72972</u>	<u>001</u> Feb 28, 1992
<u>AB</u>		<u>200MG</u>	<u>N72973</u>	<u>001</u> Feb 28, 1992
<u>AB</u>	WATSON LABS	<u>150MG</u>	<u>N71891</u>	<u>001</u> Apr 03, 1990
<u>AB</u>		<u>200MG</u>	<u>N71795</u>	<u>001</u> Apr 03, 1990

SUMATRIPTAN

SPRAY; NASAL				
IMITREX				
+	GLAXOSMITHKLINE	5MG/SPRAY	N20626	001 Aug 26, 1997
+		10MG/SPRAY	N20626	002 Aug 26, 1997
+		20MG/SPRAY	N20626	003 Aug 26, 1997

SUMATRIPTAN SUCCINATE

INJECTABLE; INJECTION				
IMITREX				
+	GLAXOSMITHKLINE	EQ 6MG BASE/0.5ML	N20080	001 Dec 28, 1992
TABLET; ORAL				
IMITREX				
	GLAXOSMITHKLINE	EQ 25MG BASE	N20132	002 Jun 01, 1995
		EQ 50MG BASE	N20132	003 Jun 01, 1995
+		EQ 100MG BASE	N20132	001 Jun 01, 1995

TACRINE HYDROCHLORIDE

CAPSULE; ORAL				
COGNEX				
	FIRST HORIZON	EQ 10MG BASE	N20070	001 Sep 09, 1993
		EQ 20MG BASE	N20070	002 Sep 09, 1993
		EQ 30MG BASE	N20070	003 Sep 09, 1993
+		EQ 40MG BASE	N20070	004 Sep 09, 1993

TACROLIMUS

CAPSULE; ORAL				
PROGRAF				
	ASTELLAS	EQ 0.5MG BASE	N50708	003 Aug 24, 1998
		EQ 1MG BASE	N50708	001 Apr 08, 1994
+		EQ 5MG BASE	N50708	002 Apr 08, 1994
INJECTABLE; INJECTION				
PROGRAF				
+	ASTELLAS	EQ 5MG BASE/ML	N50709	001 Apr 08, 1994
OINTMENT; TOPICAL				
PROTOPIC				
	ASTELLAS	0.03%	N50777	001 Dec 08, 2000
+		0.1%	N50777	002 Dec 08, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 327 (of 360)

TADALAFILTABLET; ORAL
CIALIS

LILLY ICOS	5MG	N21368	001	Nov 21, 2003
	10MG	N21368	002	Nov 21, 2003
+	20MG	N21368	003	Nov 21, 2003

TALCAEROSOL, METERED; INTRAPLEURAL
SCLEROSOL

+ BRYAN	400MG/SPRAY	N20587	001	Dec 24, 1997
---------	-------------	--------	-----	--------------

POWDER; INTRAPLEURAL
TALC

+ BRYAN	5GM/BOT	N21388	001	Dec 15, 2003
---------	---------	--------	-----	--------------

TAMOXIFEN CITRATESOLUTION; ORAL
SOLTAMOX

+ SAVIENT PHARMS	EQ 10MG BASE/5ML	N21807	001	Oct 29, 2005
------------------	------------------	--------	-----	--------------

TABLET; ORAL

NOLVADEX

<u>AB</u> ASTRAZENECA	<u>EQ 10MG BASE</u>	<u>N17970</u>	<u>001</u>	
<u>AB</u> +	<u>EQ 20MG BASE</u>	<u>N17970</u>	<u>002</u>	Mar 21, 1994

TAMOXIFEN CITRATE

<u>AB</u> AEGIS PHARMS	<u>EQ 10MG BASE</u>	<u>N76398</u>	<u>001</u>	Mar 31, 2003
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76398</u>	<u>002</u>	Mar 31, 2003
<u>AB</u> ANDRX PHARMS	<u>EQ 10MG BASE</u>	<u>N76179</u>	<u>001</u>	Feb 20, 2003
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76179</u>	<u>002</u>	Feb 20, 2003
<u>AB</u> BARR	<u>EQ 10MG BASE</u>	<u>N70929</u>	<u>001</u>	Feb 20, 2003
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N70929</u>	<u>002</u>	Feb 20, 2003
<u>AB</u> IVAX PHARMS	<u>EQ 10MG BASE</u>	<u>N75740</u>	<u>001</u>	Feb 20, 2003
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N75740</u>	<u>002</u>	Feb 20, 2003
<u>AB</u> MYLAN	<u>EQ 10MG BASE</u>	<u>N74732</u>	<u>002</u>	Feb 20, 2003
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N74732</u>	<u>001</u>	Feb 20, 2003
<u>AB</u> PHARMACHEMIE	<u>EQ 20MG BASE</u>	<u>N74858</u>	<u>001</u>	Feb 20, 2003
<u>AB</u> ROXANE	<u>EQ 10MG BASE</u>	<u>N76027</u>	<u>001</u>	Feb 20, 2003
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N76027</u>	<u>002</u>	Feb 20, 2003
<u>AB</u> TEVA	<u>EQ 10MG BASE</u>	<u>N74504</u>	<u>001</u>	Apr 28, 2003
<u>AB</u>	<u>EQ 10MG BASE</u>	<u>N75797</u>	<u>001</u>	Feb 20, 2003
<u>AB</u>	<u>EQ 20MG BASE</u>	<u>N74504</u>	<u>002</u>	Apr 28, 2003

TAMSULOSIN HYDROCHLORIDECAPSULE; ORAL
FLOMAX

+ BOEHRINGER INGELHEIM	0.4MG	N20579	001	Apr 15, 1997
------------------------	-------	--------	-----	--------------

TAZAROTENECREAM; TOPICAL
AVAGE

+ ALLERGAN	0.1%	N21184	003	Sep 30, 2002
TAZORAC				
+ ALLERGAN	0.05%	N21184	001	Sep 29, 2000
+	0.1%	N21184	002	Sep 29, 2000

GEL; TOPICAL
TAZORAC

+ ALLERGAN	0.05%	N20600	001	Jun 13, 1997
+	0.1%	N20600	002	Jun 13, 1997

PRESCRIPTION DRUG PRODUCT LIST

3 - 328 (of 360)

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE; INJECTION			
MACROTEC			
BS	BRACCO	N/A	N17833 001
PULMOLITE			
BS	CIS	N/A	N17776 001
TECHNISCAN MAA			
BS	MALLINCKRODT	N/A	N17842 001
TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT			
BS	DRAXIMAGE	N/A	N17881 001 Dec 30, 1987

TECHNETIUM TC-99M APCITIDE

INJECTABLE; INJECTION			
ACUTECT			
	BERLEX LABS	N/A	N20887 001 Sep 14, 1998

TECHNETIUM TC-99M BICISATE KIT

INJECTABLE; INJECTION			
NEUROLITE			
	BRISTOL MYERS SQUIBB	N/A/VIAL	N20256 001 Nov 23, 1994

TECHNETIUM TC-99M DEPREOTIDE

INJECTABLE; INJECTION			
NEO TECT KIT			
+	BERLEX LABS	N/A	N21012 001 Aug 03, 1999

TECHNETIUM TC-99M DISOFENIN KIT

INJECTABLE; INJECTION			
HEPATOLITE			
	CIS	N/A	N18467 001 Mar 16, 1982

TECHNETIUM TC-99M EXAMETAZIME KIT

INJECTABLE; INJECTION			
CERETEC			
	GE HEALTHCARE	N/A	N19829 001 Dec 30, 1988

TECHNETIUM TC-99M GLUCEPTATE KIT

INJECTABLE; INJECTION			
TECHNISCAN GLUCEPTATE			
	DRAXIMAGE	N/A	N18272 001 Jan 27, 1982

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION			
CHOLETEC			
	BRACCO	N/A	N18963 001 Jan 21, 1987

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION			
<u>CIS-MDP</u>			
<u>AP</u>	CIS	<u>N/A</u>	<u>N18124 001</u>
<u>DRAXIMAGE MDP</u>			
<u>AP</u>	+ DRAXIMAGE	<u>N/A</u>	<u>N18035 001</u>
<u>MDP-BRACCO</u>			
<u>AP</u>	BRACCO	<u>N/A</u>	<u>N18107 001</u>
<u>TECHNETIUM TC 99M MPI MDP</u>			
<u>AP</u>	GE HEALTHCARE	<u>N/A</u>	<u>N18141 001</u>

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 329 (of 360)

TECHNETIUM TC-99M MERTIATIDE KIT

INJECTABLE; INJECTION

TECHNESCAN MAG3

MALLINCKRODT N/A

N19882 001 Jun 15, 1990

TECHNETIUM TC-99M OXIDRONATE KIT

INJECTABLE; INJECTION

TECHNESCAN

+ MALLINCKRODT N/A

N18321 001

TECHNETIUM TC-99M PENTETATE KIT

INJECTABLE; INJECTION

AN-DTPAAP CIS N/AN17714 001DTPAAP DRAXIMAGE N/AN18511 001

Dec 29, 1989

MPI DTPA KIT - CHELATEAP GE HEALTHCARE N/AN17255 001TECHNETIUM TC-99M PENTETATE KITAP GE HEALTHCARE N/AN17264 002TECHNETIUM TC-99M PYROPHOSPHATE KIT

INJECTABLE; INJECTION

CIS-PYROAP CIS N/AN19039 001

Jun 30, 1987

PHOSPHOTECAP BRACCO N/AN17680 001TECHNESCAN PYP KITAP MALLINCKRODT N/AN17538 001TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION

ULTRATAG

MALLINCKRODT N/A

N19981 001 Jun 10, 1991

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

CARDIOLITE

+ BRISTOL MYERS SQUIBB N/A

N19785 001 Dec 21, 1990

MIRALUMA

+ BRISTOL MYERS SQUIBB N/A

N19785 003 May 23, 1997

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INJECTION, ORAL

TECHNELITE

BRISTOL MYERS SQUIBB 0.0083-2.7 CI/GENERATOR

N17771 001

ULTRA-TECHNEKOW FM

MALLINCKRODT 0.25-3 CI/GENERATOR

N17243 002

TECHNETIUM TC-99M SUCCIMER KIT

INJECTABLE; INJECTION

MPI DMSA KIDNEY REAGENT

GE HEALTHCARE N/A

N17944 001 May 18, 1982

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL

AN-SULFUR COLLOIDAP CIS N/AN17858 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 330 (of 360)

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL

TECHNECOLL

<u>AP</u>	MALLINCKRODT	<u>N/A</u>	<u>N17059</u>	<u>001</u>	
-----------	--------------	------------	---------------	------------	--

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW

+	GE HEALTHCARE	N/A	N20372	001	Feb 09, 1996
---	---------------	-----	--------	-----	--------------

MYOVIEW 30ML

+	GE HEALTHCARE	N/A	N20372	002	Jul 07, 2005
---	---------------	-----	--------	-----	--------------

TEGASEROD MALEATE

TABLET; ORAL

ZELNORM

NOVARTIS

EQ 2MG BASE

N21200	001	Jul 24, 2002
--------	-----	--------------

+		EQ 6MG BASE	N21200	002	Jul 24, 2002
---	--	-------------	--------	-----	--------------

TELITHROMYCIN

TABLET; ORAL

KETEK

AVENTIS PHARMS

300MG

N21144	002	Feb 09, 2005
--------	-----	--------------

+		400MG	N21144	001	Apr 01, 2004
---	--	-------	--------	-----	--------------

TELMISARTAN

TABLET; ORAL

MICARDIS

BOEHRINGER INGELHEIM

20MG

N20850	003	Apr 04, 2000
--------	-----	--------------

40MG

N20850	001	Nov 10, 1998
--------	-----	--------------

+		80MG	N20850	002	Nov 10, 1998
---	--	------	--------	-----	--------------

TEMAZEPAM

CAPSULE; ORAL

RESTORIL

<u>AB</u>	TYCO HLTHCARE	<u>15MG</u>	<u>N18163</u>	<u>001</u>	
-----------	---------------	-------------	---------------	------------	--

<u>AB</u>	+	<u>30MG</u>	<u>N18163</u>	<u>002</u>	
-----------	---	-------------	---------------	------------	--

		7.5MG	N18163	003	Oct 25, 1991
--	--	-------	--------	-----	--------------

		22.5MG	N18163	004	Nov 02, 2004
--	--	--------	--------	-----	--------------

TEMAZEPAM

<u>AB</u>	MYLAN	<u>15MG</u>	<u>N70919</u>	<u>001</u>	Jul 07, 1986
-----------	-------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N70920</u>	<u>001</u>	Jul 10, 1986
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	PAR PHARM	<u>15MG</u>	<u>N71456</u>	<u>001</u>	Apr 21, 1987
-----------	-----------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N71457</u>	<u>001</u>	Apr 21, 1987
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	PUREPAC PHARM	<u>15MG</u>	<u>N71638</u>	<u>001</u>	Aug 07, 1987
-----------	---------------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N71620</u>	<u>001</u>	Aug 07, 1987
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>15MG</u>	<u>N71427</u>	<u>001</u>	Jan 12, 1988
-----------	--------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N71428</u>	<u>001</u>	Jan 12, 1988
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	WATSON LABS	<u>15MG</u>	<u>N71446</u>	<u>001</u>	May 21, 1993
-----------	-------------	-------------	---------------	------------	--------------

<u>AB</u>		<u>30MG</u>	<u>N71447</u>	<u>001</u>	May 21, 1993
-----------	--	-------------	---------------	------------	--------------

TEMOZOLOMIDE

CAPSULE; ORAL

TEMODAR

SCHERING

5MG

N21029	001	Aug 11, 1999
--------	-----	--------------

20MG

N21029	002	Aug 11, 1999
--------	-----	--------------

100MG

N21029	003	Aug 11, 1999
--------	-----	--------------

+		250MG	N21029	004	Aug 11, 1999
---	--	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 331 (of 360)

TENIPOSIDE

INJECTABLE; INJECTION

VUMON

+ BRISTOL MYERS SQUIBB 10MG/ML

N20119 001 Jul 14, 1992

TENOFOVIR DISOPROXIL FUMARATE

TABLET; ORAL

VIREAD

+ GILEAD 300MG

N21356 001 Oct 26, 2001

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

HYTRINAB ABBOTTEQ 1MG BASEN20347 001 Dec 14, 1994AB +EQ 2MG BASEN20347 002 Dec 14, 1994ABEQ 5MG BASEN20347 003 Dec 14, 1994ABEQ 10MG BASEN20347 004 Dec 14, 1994TERAZOSIN HYDROCHLORIDEAB MYLANEQ 1MG BASEN75140 002 Feb 11, 2000ABEQ 2MG BASEN75140 003 Feb 11, 2000ABEQ 5MG BASEN75140 001 Feb 11, 2000ABEQ 10MG BASEN75140 004 Feb 11, 2000AB MYLAN TECHNOLOGIESEQ 1MG BASEN75384 001 Dec 01, 2000ABEQ 2MG BASEN75384 002 Dec 01, 2000ABEQ 5MG BASEN75384 003 Dec 01, 2000ABEQ 10MG BASEN75384 004 Dec 01, 2000AB RANBAXYEQ 1MG BASEN76021 001 Aug 22, 2002ABEQ 2MG BASEN76021 002 Aug 22, 2002ABEQ 5MG BASEN76021 003 Aug 22, 2002ABEQ 10MG BASEN76021 004 Aug 22, 2002AB SANDOZEQ 1MG BASEN74823 001 Mar 30, 1998ABEQ 1MG BASEN75667 001 Jul 28, 2000ABEQ 2MG BASEN74823 002 Mar 30, 1998ABEQ 2MG BASEN75667 002 Jul 28, 2000ABEQ 5MG BASEN74823 003 Mar 30, 1998ABEQ 5MG BASEN75667 003 Jul 28, 2000ABEQ 10MG BASEN74823 004 Mar 30, 1998ABEQ 10MG BASEN75667 004 Jul 28, 2000AB TORPHARMEQ 1MG BASEN75498 001 Apr 12, 2001ABEQ 2MG BASEN75498 002 Apr 12, 2001ABEQ 5MG BASEN75498 003 Apr 12, 2001ABEQ 10MG BASEN75498 004 Apr 12, 2001AB TRIGENEQ 1MG BASEN75317 001 Dec 20, 2004ABEQ 2MG BASEN75317 002 Dec 20, 2004ABEQ 5MG BASEN75317 003 Dec 20, 2004ABEQ 10MG BASEN75317 004 Dec 20, 2004

TABLET; ORAL

HYTRINAB ABBOTTEQ 1MG BASEN19057 001 Aug 07, 1987AB +EQ 2MG BASEN19057 002 Aug 07, 1987ABEQ 5MG BASEN19057 003 Aug 07, 1987ABEQ 10MG BASEN19057 004 Aug 07, 1987TERAZOSIN HYDROCHLORIDEAB SANDOZEQ 1MG BASEN74315 001 Dec 31, 1998ABEQ 1MG BASEN74657 001 Apr 28, 2000ABEQ 2MG BASEN74315 002 Dec 31, 1998ABEQ 2MG BASEN74657 002 Apr 28, 2000ABEQ 5MG BASEN74315 003 Dec 31, 1998ABEQ 5MG BASEN74657 003 Apr 28, 2000ABEQ 10MG BASEN74315 004 Dec 31, 1998

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 332 (of 360)

TERAZOSIN HYDROCHLORIDE

TABLET; ORAL

TERAZOSIN HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>N74657</u>	<u>004</u>	Apr 28, 2000
-----------	--------	---------------------	---------------	------------	--------------

TERBINAFINE

GEL; TOPICAL

LAMISIL

NOVARTIS

1%

N20846 001 Apr 29, 1998

TERBINAFINE HYDROCHLORIDE

SOLUTION; TOPICAL

LAMISIL

+ NOVARTIS

1%

N20749 001 Oct 17, 1997

TABLET; ORAL

LAMISIL

+ NOVARTIS

EQ 250MG BASE

N20539 001 May 10, 1996

TERBUTALINE SULFATE

INJECTABLE; INJECTION

BRETHINE

<u>AP</u>	+ AAIPHARMA LLC	<u>1MG/ML</u>	<u>N18571</u>	<u>001</u>	
-----------	-----------------	---------------	---------------	------------	--

TERBUTALINE SULFATE

<u>AP</u>	AM PHARM	<u>1MG/ML</u>	<u>N76887</u>	<u>001</u>	May 26, 2004
-----------	----------	---------------	---------------	------------	--------------

<u>AP</u>	BEDFORD	<u>1MG/ML</u>	<u>N76770</u>	<u>001</u>	Apr 23, 2004
-----------	---------	---------------	---------------	------------	--------------

<u>AP</u>	SICOR PHARMS	<u>1MG/ML</u>	<u>N76853</u>	<u>001</u>	Jul 20, 2004
-----------	--------------	---------------	---------------	------------	--------------

TABLET; ORAL

BRETHINE

<u>AB</u>	AAIPHARMA LLC	<u>2.5MG</u>	<u>N17849</u>	<u>001</u>	
-----------	---------------	--------------	---------------	------------	--

<u>AB</u>	+	<u>5MG</u>	<u>N17849</u>	<u>002</u>	
-----------	---	------------	---------------	------------	--

TERBUTALINE SULFATE

<u>AB</u>	IMPAX LABS	<u>2.5MG</u>	<u>N75877</u>	<u>001</u>	Jun 26, 2001
-----------	------------	--------------	---------------	------------	--------------

<u>AB</u>		<u>5MG</u>	<u>N75877</u>	<u>002</u>	Jun 26, 2001
-----------	--	------------	---------------	------------	--------------

<u>AB</u>	LANNETT	<u>2.5MG</u>	<u>N77152</u>	<u>001</u>	Mar 25, 2005
-----------	---------	--------------	---------------	------------	--------------

<u>AB</u>		<u>5MG</u>	<u>N77152</u>	<u>002</u>	Mar 25, 2005
-----------	--	------------	---------------	------------	--------------

TERCONAZOLE

CREAM; VAGINAL

TERAZOL 3

<u>AB</u>	+ ORTHO MCNEIL PHARM	<u>0.8%</u>	<u>N19964</u>	<u>001</u>	Feb 21, 1991
-----------	----------------------	-------------	---------------	------------	--------------

TERAZOL 7

<u>AB</u>	+ ORTHO MCNEIL PHARM	<u>0.4%</u>	<u>N19579</u>	<u>001</u>	Dec 31, 1987
-----------	----------------------	-------------	---------------	------------	--------------

TERCONAZOLE

<u>AB</u>	ALTANA	<u>0.4%</u>	<u>N76712</u>	<u>001</u>	Feb 18, 2005
-----------	--------	-------------	---------------	------------	--------------

BX	+	0.8%	N21735	001	Oct 01, 2004
----	---	------	--------	-----	--------------

<u>AB</u>	TARO	<u>0.4%</u>	<u>N76043</u>	<u>001</u>	Jan 19, 2005
-----------	------	-------------	---------------	------------	--------------

<u>AB</u>		<u>0.8%</u>	<u>N75953</u>	<u>001</u>	Apr 06, 2004
-----------	--	-------------	---------------	------------	--------------

SUPPOSITORY; VAGINAL

TERAZOL 3

+ ORTHO MCNEIL PHARM 80MG

N19641 001 May 24, 1988

TERIPARATIDE RECOMBINANT HUMAN

INJECTABLE; SUBCUTANEOUS

FORTEO

+ LILLY EQ 0.75MG/3ML(0.25MG/ML)

N21318 001 Nov 26, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 333 (of 360)

TESTOLACTONE

TABLET; ORAL

TESLAC

+ BRISTOL MYERS SQUIBB 50MG

N16118 001

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

ANDRODERM

+ WATSON LABS 2.5MG/24HR

N20489 001

Sep 29, 1995

+ 5MG/24HR

N20489 002

May 02, 1997

GEL; TRANSDERMAL

ANDROGEL

BX + UNIMED PHARMS 1%

N21015 001

Feb 28, 2000

TESTIM

BX + AUXILIUM PHARMS 1%

N21454 001

Oct 31, 2002

INJECTABLE; INJECTION

TESTOSTERONE

+ STERIS 100MG/ML

N86417 001

Jul 07, 1983

PELLET; IMPLANTATION

TESTOSTERONE

BARTOR 75MG

N80911 001

TABLET, EXTENDED RELEASE; BUCCAL

STRIANT

+ COLUMBIA LABS 30MG

N21543 001

Jun 19, 2003

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTOSTERONEAO + PHARMACIA AND UPJOHN 100MG/MLN85635 002AO + 200MG/MLN85635 003TESTOSTERONE CYPIONATEAO PADDOCK 200MG/MLN40530 001

Jan 31, 2005

AO STERIS 100MG/MLN86029 001AO 200MG/MLN86030 001TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION

DELATESTRYLAO + SAVIENT PHARMS 200MG/MLN09165 003TESTOSTERONE ENANTHATEAO STERIS 200MG/MLN85598 001TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION

TESTOSTERONE PROPIONATE

+ STERIS 25MG/ML

N80188 001

50MG/ML

N80188 002

+ 100MG/ML

N80188 003

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

BRISTACYCLINEAB BRISTOL 250MGN61658 001AB 250MGN61888 001AB 500MGN61658 002AB 500MGN61888 002TETRACYCLINE HYDROCHLORIDEAB BARR 250MGN61837 001AB 500MGN61837 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 334 (of 360)

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

<u>AB</u>	IMPAX LABS	<u>250MG</u>	<u>N60469</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N60469</u>	<u>003</u>	
		100MG	N60469	002	
<u>AB</u>	IVAX PHARMS	<u>250MG</u>	<u>N60704</u>	<u>001</u>	
<u>AB</u>	+	<u>500MG</u>	<u>N60704</u>	<u>002</u>	
<u>AB</u>	LABS ATRAL	<u>250MG</u>	<u>N62752</u>	<u>001</u>	Aug 12, 1988
<u>AB</u>		<u>500MG</u>	<u>N62752</u>	<u>002</u>	Aug 12, 1988
<u>AB</u>	MUTUAL PHARM	<u>250MG</u>	<u>N60736</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N60736</u>	<u>002</u>	
<u>AB</u>	MYLAN	<u>250MG</u>	<u>N60783</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N60783</u>	<u>002</u>	
SUSPENSION; ORAL					
SUMYCIN					
	+	APOTHECON	125MG/5ML	N60400	001
TABLET; ORAL					
SUMYCIN					
		PAR PHARM	250MG	N61147	001
	+		500MG	N61147	004

TETRACYCLINE PHOSPHATE COMPLEX

CAPSULE; ORAL

TETREX

BRISTOL

EQ 100MG HCL

N61653 001

EQ 250MG HCL

N61653 002

EQ 250MG HCL

N61889 002

+

EQ 500MG HCL

N61653 003

EQ 500MG HCL

N61889 001

TETRAHYDROZOLINE HYDROCHLORIDE

SOLUTION; NASAL

TYZINE

+ KENWOOD LABS

0.05%

N86576 002

0.1%

N86576 001

SPRAY; NASAL

TYZINE

+ KENWOOD LABS

0.1%

N86576 003

THALIDOMIDE

CAPSULE; ORAL

THALOMID

CELGENE

50MG

N20785 001

Jul 16, 1998

100MG

N20785 002

Jan 17, 2003

+

200MG

N20785 003

Jan 17, 2003

THALLOUS CHLORIDE, TL-201

INJECTABLE; INJECTION

THALLOUS CHLORIDE TL 201

<u>AP</u>	+	BRISTOL MYERS SQUIBB	<u>1mCi/ML</u>	<u>N17806</u>	<u>001</u>	
<u>AP</u>	+	GE HEALTHCARE	<u>1mCi/ML</u>	<u>N18110</u>	<u>002</u>	Feb 27, 1996
<u>AP</u>	+	MALLINCKRODT	<u>1mCi/ML</u>	<u>N18150</u>	<u>001</u>	
<u>AP</u>		MOUNT SINAI MEDCTR	<u>1mCi/ML</u>	<u>N75569</u>	<u>001</u>	Nov 21, 2001
INJECTABLE; INTRAVENOUS						
THALLOUS CHLORIDE TL 201						
	+	BRISTOL MYERS SQUIBB	2mCi/ML	N17806	002	Oct 09, 1998

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 335 (of 360)

THEOPHYLLINECAPSULE, EXTENDED RELEASE; ORAL
THEO-24

BC	UCB	100MG	N87942	001	Aug 22, 1983
BC		200MG	N87943	001	Aug 22, 1983
BC		300MG	N87944	001	Aug 22, 1983
		400MG	N81034	001	Feb 28, 1992

THEOPHYLLINE					
BC	INWOOD LABS	100MG	N40052	001	Feb 14, 1994
BC		200MG	N40052	003	Feb 14, 1994
BC	+	300MG	N40052	004	Feb 14, 1994
		125MG	N40052	002	Feb 14, 1994

ELIXIR; ORAL

ELIXOPHYLLIN

<u>AA</u>	FOREST LABS	<u>80MG/15ML</u>	<u>N85186</u>	<u>001</u>	
<u>THEOPHYLLINE</u>					
<u>AA</u>	ALPHARMA US PHARMS	<u>80MG/15ML</u>	<u>N85863</u>	<u>001</u>	
<u>AA</u>	MORTON GROVE	<u>80MG/15ML</u>	<u>N86748</u>	<u>001</u>	
<u>AA</u>	PHARM ASSOC	<u>80MG/15ML</u>	<u>N86720</u>	<u>001</u>	
<u>AA</u>	TARO	<u>80MG/15ML</u>	<u>N89626</u>	<u>001</u>	Oct 28, 1988

INJECTABLE; INJECTION

<u>THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+	B BRAUN	<u>40MG/100ML</u>	<u>N19826</u>	<u>001</u> Aug 14, 1992
<u>THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+	B BRAUN	<u>80MG/100ML</u>	<u>N19826</u>	<u>002</u> Aug 14, 1992
<u>THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+	B BRAUN	<u>160MG/100ML</u>	<u>N19826</u>	<u>003</u> Aug 14, 1992
<u>THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+	B BRAUN	<u>320MG/100ML</u>	<u>N19826</u>	<u>006</u> Aug 14, 1992
<u>THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+	BAXTER HLTHCARE	<u>4MG/ML</u>	<u>N18649</u>	<u>007</u> Jul 26, 1982
<u>AP</u>	+		<u>40MG/100ML</u>	<u>N18649</u>	<u>001</u> Jul 26, 1982
<u>AP</u>	+		<u>80MG/100ML</u>	<u>N18649</u>	<u>002</u> Jul 26, 1982
<u>AP</u>	+		<u>160MG/100ML</u>	<u>N18649</u>	<u>003</u> Jul 26, 1982
<u>AP</u>	+		<u>200MG/100ML</u>	<u>N18649</u>	<u>004</u> Jul 26, 1982
<u>AP</u>	+		<u>320MG/100ML</u>	<u>N18649</u>	<u>006</u> Nov 13, 1985
<u>AP</u>	+		<u>400MG/100ML</u>	<u>N18649</u>	<u>005</u> Jul 26, 1982
<u>THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
<u>AP</u>	+	HOSPIRA	<u>4MG/ML</u>	<u>N19211</u>	<u>007</u> Dec 14, 1984
<u>AP</u>	+		<u>40MG/100ML</u>	<u>N19211</u>	<u>001</u> Dec 14, 1984
<u>AP</u>	+		<u>80MG/100ML</u>	<u>N19211</u>	<u>002</u> Dec 14, 1984
<u>AP</u>	+		<u>160MG/100ML</u>	<u>N19211</u>	<u>003</u> Dec 14, 1984
<u>AP</u>	+		<u>200MG/100ML</u>	<u>N19211</u>	<u>004</u> Dec 14, 1984
<u>AP</u>	+		<u>320MG/100ML</u>	<u>N19211</u>	<u>006</u> Jan 20, 1988
<u>AP</u>	+		<u>400MG/100ML</u>	<u>N19211</u>	<u>005</u> Dec 14, 1984

SOLUTION; ORAL

<u>THEOLAIR</u>					
<u>AA</u>	+	3M	<u>80MG/15ML</u>	<u>N86107</u>	<u>001</u>
<u>THEOPHYLLINE</u>					
<u>AA</u>		ROXANE	<u>80MG/15ML</u>	<u>N87449</u>	<u>001</u> Sep 15, 1983

TABLET; ORAL

QUIBRON-T					
	+	MONARCH PHARMS	300MG	N88656	001 Aug 22, 1985
THEOLAIR					
	+	3M	125MG	N86399	001
	+		250MG	N86399	002

TABLET, EXTENDED RELEASE; ORAL

QUIBRON-T/SR					
BC		MONARCH PHARMS	300MG	N87563	001 Jun 21, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 336 (of 360)

THEOPHYLLINE

TABLET, EXTENDED RELEASE; ORAL

THEOCHRON

<u>AB</u>	INWOOD LABS	<u>100MG</u>	<u>N88320</u>	<u>001</u>	Feb 21, 1985
<u>AB</u>		<u>200MG</u>	<u>N88321</u>	<u>001</u>	Feb 21, 1985
<u>AB</u>		<u>300MG</u>	<u>N87400</u>	<u>002</u>	Jan 11, 1983

THEOPHYLLINE

<u>AB</u>	INWOOD LABS	<u>450MG</u>	<u>N40034</u>	<u>001</u>	Apr 28, 1995
<u>AB</u>	+ PLIVA	<u>100MG</u>	<u>N89807</u>	<u>001</u>	Apr 30, 1990
<u>AB</u>	+	<u>200MG</u>	<u>N89808</u>	<u>001</u>	Apr 30, 1990
<u>AB</u>		<u>300MG</u>	<u>N89763</u>	<u>001</u>	Apr 30, 1990
<u>AB</u>	+	<u>450MG</u>	<u>N81236</u>	<u>001</u>	Nov 09, 1992
	T-PHYL				
BC	PURDUE FREDERICK	200MG	N88253	001	Aug 17, 1983
	UNIPHYL				
	+ PURDUE FREDERICK	400MG	N87571	001	Sep 01, 1982
	+	600MG	N40086	001	Apr 15, 1996

THIABENDAZOLE

SUSPENSION; ORAL

MINTEZOL

+	MERCK	500MG/5ML	N16097	001	
---	-------	-----------	--------	-----	--

TABLET, CHEWABLE; ORAL

MINTEZOL

+	MERCK	500MG	N16096	001	
---	-------	-------	--------	-----	--

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

THIAMINE HYDROCHLORIDE

<u>AP</u>	+	AM PHARM PARTNERS	<u>100MG/ML</u>	<u>N80556</u>	<u>001</u>	
<u>AP</u>		BAXTER HLTHCARE	<u>100MG/ML</u>	<u>N80575</u>	<u>001</u>	
<u>AP</u>		HOSPIRA	<u>100MG/ML</u>	<u>N40079</u>	<u>001</u>	May 03, 1996
<u>AP</u>		STERIS	<u>100MG/ML</u>	<u>N80571</u>	<u>001</u>	
	+		200MG/ML	N80571	002	

THIETHYLPERAZINE MALEATE

TABLET; ORAL

TORECAN

+	NOVARTIS	10MG	N12753	001	
---	----------	------	--------	-----	--

THIOGUANINE

TABLET; ORAL

THIOGUANINE

+	GLAXOSMITHKLINE	40MG	N12429	001	
---	-----------------	------	--------	-----	--

THIORIDAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

THIORIDAZINE HYDROCHLORIDE

<u>AA</u>		ALPHARMA US PHARMS	<u>100MG/ML</u>	<u>N88229</u>	<u>001</u>	Aug 23, 1983
<u>AA</u>		PHARM ASSOC	<u>30MG/ML</u>	<u>N40187</u>	<u>001</u>	Aug 28, 1997
<u>AA</u>			<u>100MG/ML</u>	<u>N40213</u>	<u>001</u>	May 29, 1998
<u>AA</u>	+	TEVA PHARMS	<u>30MG/ML</u>	<u>N89602</u>	<u>001</u>	Nov 09, 1987
<u>AA</u>	+		<u>100MG/ML</u>	<u>N89603</u>	<u>001</u>	Nov 09, 1987

THIORIDAZINE HYDROCHLORIDE INTENSOL

<u>AA</u>		ROXANE	<u>30MG/ML</u>	<u>N88941</u>	<u>001</u>	Dec 16, 1985
<u>AA</u>			<u>100MG/ML</u>	<u>N88942</u>	<u>001</u>	Dec 16, 1985

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

<u>AB</u>		IVAX PHARMS	<u>10MG</u>	<u>N88270</u>	<u>001</u>	Apr 14, 1983
-----------	--	-------------	-------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 337 (of 360)

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

<u>AB</u>	IVAX PHARMS	<u>15MG</u>	<u>N88271</u>	<u>001</u>	Apr 14, 1983
<u>AB</u>		<u>25MG</u>	<u>N88272</u>	<u>001</u>	Apr 14, 1983
<u>AB</u>	MUTUAL PHARM	<u>10MG</u>	<u>N89431</u>	<u>001</u>	Aug 01, 1986
<u>AB</u>		<u>25MG</u>	<u>N89432</u>	<u>001</u>	Aug 01, 1986
<u>AB</u>		<u>50MG</u>	<u>N88370</u>	<u>001</u>	Nov 18, 1983
<u>AB</u>		<u>50MG</u>	<u>N89433</u>	<u>001</u>	Aug 01, 1986
<u>AB</u>		<u>100MG</u>	<u>N89953</u>	<u>001</u>	Oct 07, 1988
<u>AB</u>	MYLAN	<u>10MG</u>	<u>N88001</u>	<u>001</u>	Mar 15, 1983
<u>AB</u>		<u>25MG</u>	<u>N88002</u>	<u>001</u>	Mar 15, 1983
<u>AB</u>		<u>50MG</u>	<u>N88003</u>	<u>001</u>	Mar 15, 1983
<u>AB</u>		<u>100MG</u>	<u>N88004</u>	<u>001</u>	Nov 18, 1983
<u>AB</u>	SANDOZ	<u>10MG</u>	<u>N88131</u>	<u>001</u>	Aug 30, 1983
<u>AB</u>		<u>15MG</u>	<u>N88132</u>	<u>001</u>	Aug 30, 1983
<u>AB</u>	+	<u>25MG</u>	<u>N88133</u>	<u>001</u>	Aug 30, 1983
<u>AB</u>		<u>50MG</u>	<u>N88134</u>	<u>001</u>	Aug 30, 1983
<u>AB</u>	+	<u>100MG</u>	<u>N88135</u>	<u>001</u>	Nov 20, 1984
<u>AB</u>		<u>150MG</u>	<u>N88136</u>	<u>001</u>	Sep 17, 1986
<u>AB</u>	+	<u>200MG</u>	<u>N88137</u>	<u>001</u>	Sep 17, 1986
<u>AB</u>	TEVA	<u>100MG</u>	<u>N88456</u>	<u>001</u>	May 17, 1985
<u>AB</u>	WATSON LABS	<u>10MG</u>	<u>N88476</u>	<u>001</u>	Nov 08, 1983
<u>AB</u>		<u>15MG</u>	<u>N88477</u>	<u>001</u>	Nov 08, 1983
<u>AB</u>		<u>25MG</u>	<u>N88478</u>	<u>001</u>	Nov 08, 1983
<u>AB</u>		<u>25MG</u>	<u>N88567</u>	<u>001</u>	May 11, 1984
<u>AB</u>		<u>50MG</u>	<u>N88479</u>	<u>001</u>	Nov 08, 1983
<u>AB</u>		<u>50MG</u>	<u>N88563</u>	<u>001</u>	May 11, 1984
<u>AB</u>		<u>100MG</u>	<u>N88564</u>	<u>001</u>	May 11, 1984
<u>AB</u>		<u>100MG</u>	<u>N88736</u>	<u>001</u>	Jul 24, 1984
<u>AB</u>		<u>150MG</u>	<u>N88869</u>	<u>001</u>	Jun 28, 1985
<u>AB</u>		<u>200MG</u>	<u>N88872</u>	<u>001</u>	Apr 26, 1985

THIOTEPA

INJECTABLE; INJECTION

THIOTEPA

<u>AP</u>	AM PHARM PARTNERS	<u>15MG/VIAL</u>	<u>N75698</u>	<u>001</u>	Sep 20, 2001
<u>AP</u>	BEDFORD	<u>15MG/VIAL</u>	<u>N75547</u>	<u>001</u>	Apr 02, 2001
<u>AP</u>	+	<u>15MG/VIAL</u>	<u>N75730</u>	<u>001</u>	Apr 20, 2001
	+	30MG/VIAL	N75730	002	Apr 20, 2001

THIOTHIXENE

CAPSULE; ORAL

NAVANE

<u>AB</u>	PFIZER	<u>1MG</u>	<u>N16584</u>	<u>001</u>	
<u>AB</u>		<u>2MG</u>	<u>N16584</u>	<u>002</u>	
<u>AB</u>	+	<u>5MG</u>	<u>N16584</u>	<u>003</u>	
<u>AB</u>		<u>10MG</u>	<u>N16584</u>	<u>004</u>	
		20MG	N16584	005	

THIOTHIXENE

<u>AB</u>	MYLAN	<u>1MG</u>	<u>N71090</u>	<u>001</u>	Jun 23, 1987
<u>AB</u>		<u>2MG</u>	<u>N71091</u>	<u>001</u>	Jun 23, 1987
<u>AB</u>		<u>5MG</u>	<u>N71092</u>	<u>001</u>	Jun 23, 1987
<u>AB</u>		<u>10MG</u>	<u>N71093</u>	<u>001</u>	Jun 23, 1987
<u>AB</u>	SANDOZ	<u>1MG</u>	<u>N71610</u>	<u>001</u>	Jun 24, 1987
<u>AB</u>		<u>2MG</u>	<u>N71570</u>	<u>001</u>	Jun 24, 1987
<u>AB</u>		<u>5MG</u>	<u>N71529</u>	<u>001</u>	Jun 24, 1987
<u>AB</u>		<u>10MG</u>	<u>N71530</u>	<u>001</u>	Jun 24, 1987
<u>AB</u>	WATSON LABS	<u>1MG</u>	<u>N70600</u>	<u>001</u>	Jun 05, 1987
<u>AB</u>		<u>2MG</u>	<u>N70601</u>	<u>001</u>	Jun 05, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 338 (of 360)

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

<u>AB</u>	WATSON LABS	<u>5MG</u>	<u>N70602</u>	<u>001</u>	Jun 05, 1987
<u>AB</u>		<u>10MG</u>	<u>N70603</u>	<u>001</u>	Jun 05, 1987

THIOTHIXENE HYDROCHLORIDE

CONCENTRATE; ORAL

NAVANE

<u>AA</u>	+	PFIZER	<u>EQ 5MG BASE/ML</u>	<u>N16758</u>	<u>001</u>	
<u>AA</u>			<u>THIOTHIXENE HYDROCHLORIDE</u>			
<u>AA</u>		TEVA PHARMS	<u>EQ 5MG BASE/ML</u>	<u>N71554</u>	<u>001</u>	Oct 16, 1987

INJECTABLE; INJECTION

NAVANE

+	PFIZER	EQ 10MG BASE/VIAL	N16904	002	
---	--------	-------------------	--------	-----	--

THYROTROPIN ALFA

INJECTABLE; INJECTION

THYROGEN

+	GENZYME	1.1MG/VIAL	N20898	001	Nov 30, 1998
---	---------	------------	--------	-----	--------------

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

+	CEPHALON	2MG	N20646	005	Apr 16, 1999
		4MG	N20646	001	Sep 30, 1997
		6MG	N20646	006	Nov 29, 2005
		8MG	N20646	007	Nov 29, 2005
		10MG	N20646	008	Nov 29, 2005
		12MG	N20646	002	Sep 30, 1997
+		16MG	N20646	003	Sep 30, 1997

TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TICAR

+	GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	N50497	001	
+		EQ 3GM BASE/VIAL	N50497	002	
+		EQ 20GM BASE/VIAL	N50497	004	
+		EQ 30GM BASE/VIAL	N50497	005	Apr 04, 1984

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLID

<u>AB</u>	+	ROCHE PALO	<u>250MG</u>	<u>N19979</u>	<u>002</u>	Oct 31, 1991
<u>AB</u>			<u>TICLOPIDINE HYDROCHLORIDE</u>			
<u>AB</u>		CARACO	<u>250MG</u>	<u>N75526</u>	<u>001</u>	Sep 26, 2002
<u>AB</u>		GENPHARM	<u>250MG</u>	<u>N75161</u>	<u>001</u>	Sep 13, 1999
<u>AB</u>		MYLAN	<u>250MG</u>	<u>N75316</u>	<u>001</u>	Nov 02, 1999
<u>AB</u>		PUREPAC PHARM	<u>250MG</u>	<u>N75253</u>	<u>001</u>	Aug 20, 1999
<u>AB</u>		SANDOZ	<u>250MG</u>	<u>N75318</u>	<u>001</u>	Aug 20, 1999
<u>AB</u>			<u>250MG</u>	<u>N75326</u>	<u>001</u>	Aug 20, 1999
<u>AB</u>		TEVA	<u>250MG</u>	<u>N75149</u>	<u>001</u>	Aug 20, 1999
<u>AB</u>		TORPHARM	<u>250MG</u>	<u>N75089</u>	<u>001</u>	Jul 01, 1999

TIGECYCLINE

INJECTABLE; IV (INFUSION)

TYGACIL

+	WYETH PHARMS INC	50MG/VIAL	N21821	001	Jun 15, 2005
---	------------------	-----------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 339 (of 360)

TILUDRONATE DISODIUM

TABLET; ORAL

SKELID

+	SANOFI SYNTHELABO	EQ 200MG BASE	N20707	001	Mar 07, 1997
---	-------------------	---------------	--------	-----	--------------

TIMOLOL

SOLUTION/DROPS; OPHTHALMIC

BETIMOL

+	SANTEN OY	EQ 0.25% BASE	N20439	001	Mar 31, 1995
---	-----------	---------------	--------	-----	--------------

+		EQ 0.5% BASE	N20439	002	Mar 31, 1995
---	--	--------------	--------	-----	--------------

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS; OPHTHALMIC

TIMOLOL MALEATE

<u>AB</u>	FALCON PHARMS	<u>EQ 0.25% BASE</u>	<u>N20963</u>	<u>001</u>	Oct 21, 1998
-----------	---------------	----------------------	---------------	------------	--------------

<u>AB</u>		<u>EQ 0.5% BASE</u>	<u>N20963</u>	<u>002</u>	Oct 21, 1998
-----------	--	---------------------	---------------	------------	--------------

TIMOPTIC-XE

<u>AB</u>	+	MERCK	<u>EQ 0.25% BASE</u>	<u>N20330</u>	<u>001</u>	Nov 04, 1993
-----------	---	-------	----------------------	---------------	------------	--------------

<u>AB</u>	+		<u>EQ 0.5% BASE</u>	<u>N20330</u>	<u>002</u>	Nov 04, 1993
-----------	---	--	---------------------	---------------	------------	--------------

SOLUTION/DROPS; OPHTHALMIC

ISTALOL

BT	+	ISTA PHARMS	EQ 0.5% BASE	N21516	001	Jun 04, 2004
----	---	-------------	--------------	--------	-----	--------------

TIMOLOL MALEATE

<u>AT</u>		AKORN	<u>EQ 0.25% BASE</u>	<u>N74515</u>	<u>001</u>	Mar 25, 1997
-----------	--	-------	----------------------	---------------	------------	--------------

<u>AT</u>			<u>EQ 0.5% BASE</u>	<u>N74466</u>	<u>001</u>	Mar 25, 1997
-----------	--	--	---------------------	---------------	------------	--------------

<u>AT</u>			<u>EQ 0.5% BASE</u>	<u>N74516</u>	<u>001</u>	Mar 25, 1997
-----------	--	--	---------------------	---------------	------------	--------------

<u>AT</u>		BAUSCH AND LOMB	<u>EQ 0.25% BASE</u>	<u>N74778</u>	<u>001</u>	Mar 25, 1997
-----------	--	-----------------	----------------------	---------------	------------	--------------

<u>AT</u>			<u>EQ 0.5% BASE</u>	<u>N74776</u>	<u>001</u>	Mar 25, 1997
-----------	--	--	---------------------	---------------	------------	--------------

<u>AT</u>		FALCON PHARMS	<u>EQ 0.25% BASE</u>	<u>N74261</u>	<u>001</u>	Apr 28, 1995
-----------	--	---------------	----------------------	---------------	------------	--------------

<u>AT</u>			<u>EQ 0.5% BASE</u>	<u>N74262</u>	<u>001</u>	Apr 28, 1995
-----------	--	--	---------------------	---------------	------------	--------------

<u>AT</u>		HI TECH PHARMA	<u>EQ 0.5% BASE</u>	<u>N75163</u>	<u>001</u>	Sep 10, 2002
-----------	--	----------------	---------------------	---------------	------------	--------------

<u>AT</u>		NOVEX	<u>EQ 0.25% BASE</u>	<u>N75411</u>	<u>001</u>	Sep 08, 2000
-----------	--	-------	----------------------	---------------	------------	--------------

<u>AT</u>			<u>EQ 0.5% BASE</u>	<u>N75412</u>	<u>001</u>	Sep 08, 2000
-----------	--	--	---------------------	---------------	------------	--------------

<u>AT</u>		PACIFIC PHARMA	<u>EQ 0.25% BASE</u>	<u>N74746</u>	<u>001</u>	Mar 25, 1997
-----------	--	----------------	----------------------	---------------	------------	--------------

<u>AT</u>			<u>EQ 0.5% BASE</u>	<u>N74747</u>	<u>001</u>	Mar 25, 1997
-----------	--	--	---------------------	---------------	------------	--------------

TIMOPTIC

<u>AT</u>	+	MERCK	<u>EQ 0.25% BASE</u>	<u>N18086</u>	<u>001</u>	
-----------	---	-------	----------------------	---------------	------------	--

<u>AT</u>	+		<u>EQ 0.5% BASE</u>	<u>N18086</u>	<u>002</u>	
-----------	---	--	---------------------	---------------	------------	--

TIMOPTIC IN OCUDOSE

+	MERCK	EQ 0.25% BASE	N19463	001	Nov 05, 1986
---	-------	---------------	--------	-----	--------------

+		EQ 0.5% BASE	N19463	002	Nov 05, 1986
---	--	--------------	--------	-----	--------------

TABLET; ORAL

BLOCADREN

<u>AB</u>		MERCK	<u>5MG</u>	<u>N18017</u>	<u>001</u>	
-----------	--	-------	------------	---------------	------------	--

<u>AB</u>			<u>10MG</u>	<u>N18017</u>	<u>002</u>	
-----------	--	--	-------------	---------------	------------	--

<u>AB</u>	+		<u>20MG</u>	<u>N18017</u>	<u>004</u>	
-----------	---	--	-------------	---------------	------------	--

TIMOLOL MALEATE

<u>AB</u>		MYLAN	<u>5MG</u>	<u>N72666</u>	<u>001</u>	Jun 08, 1990
-----------	--	-------	------------	---------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>N72667</u>	<u>001</u>	Jun 08, 1990
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>20MG</u>	<u>N72668</u>	<u>001</u>	Jun 08, 1990
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>		SANDOZ	<u>5MG</u>	<u>N72550</u>	<u>001</u>	Apr 13, 1989
-----------	--	--------	------------	---------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>N72551</u>	<u>001</u>	Apr 13, 1989
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>20MG</u>	<u>N72552</u>	<u>001</u>	Apr 13, 1989
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>		WATSON LABS	<u>5MG</u>	<u>N72917</u>	<u>001</u>	Jul 31, 1991
-----------	--	-------------	------------	---------------	------------	--------------

<u>AB</u>			<u>10MG</u>	<u>N72918</u>	<u>001</u>	Jul 31, 1991
-----------	--	--	-------------	---------------	------------	--------------

<u>AB</u>			<u>20MG</u>	<u>N72919</u>	<u>001</u>	Jul 31, 1991
-----------	--	--	-------------	---------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 340 (of 360)

TINIDAZOLE

TABLET; ORAL

TINDAMAX

PRESUTTI LABS

250MG

N21618 001

May 17, 2004

+

500MG

N21618 002

May 17, 2004

TINZAPARIN SODIUM

INJECTABLE; INJECTION

INNOHEP

+ PHARMION

20,000 IU/ML

N20484 001

Jul 14, 2000

TIOPRONIN

TABLET; ORAL

TIOPRONIN

+ MISSION PHARMA

100MG

N19569 001

Aug 11, 1988

TIOTROPIUM BROMIDE MONOHYDRATE

POWDER; INHALATION

SPIRIVA

+ BOEHRINGER INGELHEIM EQ 0.018 BASE/INH

N21395 001

Jan 30, 2004

TIPRANAVIR

CAPSULE; ORAL

APTIVUS

+ BOEHRINGER INGELHEIM 250MG

N21814 001

Jun 22, 2005

TIROFIBAN HYDROCHLORIDE

INJECTABLE; INJECTION

AGGRASTAT

+ MGI PHARMA INC EQ 0.05MG BASE/ML

N20913 001

May 14, 1998

+

EQ 0.25MG BASE/ML

N20912 001

May 14, 1998

TIZANIDINE HYDROCHLORIDE

CAPSULE; ORAL

ZANAFLEX

ACORDA

EQ 2MG BASE

N21447 001

Aug 29, 2002

EQ 4MG BASE

N21447 002

Aug 29, 2002

+

EQ 6MG BASE

N21447 003

Aug 29, 2002

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

<u>AB</u>	ALPHAPHARM	<u>EQ 2MG BASE</u>	<u>N76282</u>	<u>001</u>	Dec 16, 2003
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76282</u>	<u>002</u>	Dec 16, 2003
<u>AB</u>	AMIDE PHARM	<u>EQ 2MG BASE</u>	<u>N76281</u>	<u>001</u>	Oct 20, 2003
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76281</u>	<u>002</u>	Oct 20, 2003
<u>AB</u>	BARR	<u>EQ 2MG BASE</u>	<u>N76371</u>	<u>001</u>	Apr 09, 2003
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76371</u>	<u>002</u>	Apr 09, 2003
<u>AB</u>	CARACO	<u>EQ 2MG BASE</u>	<u>N76416</u>	<u>001</u>	Sep 29, 2003
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76416</u>	<u>002</u>	Sep 29, 2003
<u>AB</u>	COREPHARMA	<u>EQ 2MG BASE</u>	<u>N76347</u>	<u>001</u>	Oct 11, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76347</u>	<u>002</u>	Oct 11, 2002
<u>AB</u>	DR REDDYS LABS INC	<u>EQ 2MG BASE</u>	<u>N76286</u>	<u>001</u>	Jul 03, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76286</u>	<u>002</u>	Jul 03, 2002
<u>AB</u>	IVAX PHARMS	<u>EQ 2MG BASE</u>	<u>N76321</u>	<u>001</u>	Sep 30, 2004
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76321</u>	<u>002</u>	Sep 30, 2004
<u>AB</u>	MYLAN	<u>EQ 2MG BASE</u>	<u>N76354</u>	<u>001</u>	Mar 28, 2003
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76354</u>	<u>002</u>	Mar 28, 2003
<u>AB</u>	PUREPAC PHARM	<u>EQ 2MG BASE</u>	<u>N76283</u>	<u>001</u>	Jul 12, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76283</u>	<u>002</u>	Jul 12, 2002
<u>AB</u>	SANDOZ	<u>EQ 2MG BASE</u>	<u>N76399</u>	<u>001</u>	Nov 26, 2002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 341 (of 360)

TIZANIDINE HYDROCHLORIDE

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>EQ 4MG BASE</u>	<u>N76280</u>	<u>002</u>	Jun 27, 2002
<u>AB</u>	TEVA	<u>EQ 2MG BASE</u>	<u>N76284</u>	<u>001</u>	Jul 03, 2002
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76284</u>	<u>002</u>	Jul 03, 2002
<u>AB</u>	TORPHARM	<u>EQ 2MG BASE</u>	<u>N76533</u>	<u>001</u>	Jan 16, 2004
<u>AB</u>		<u>EQ 4MG BASE</u>	<u>N76533</u>	<u>002</u>	Jan 16, 2004
<u>ZANAFLEX</u>					
<u>AB</u>	ACORDA	<u>EQ 2MG BASE</u>	<u>N20397</u>	<u>002</u>	Feb 04, 2000
<u>AB</u>	+	<u>EQ 4MG BASE</u>	<u>N20397</u>	<u>001</u>	Nov 27, 1996

TOBRAMYCIN

OINTMENT; OPHTHALMIC

TOBREX

+	ALCON	0.3%	N50555	001	
---	-------	------	--------	-----	--

SOLUTION; INHALATION

TOBI

+	CHIRON	300MG/5ML	N50753	001	Dec 22, 1997
---	--------	-----------	--------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

AKTOB

<u>AT</u>	AKORN	<u>0.3%</u>	<u>N64096</u>	<u>001</u>	Jan 31, 1996
-----------	-------	-------------	---------------	------------	--------------

TOBRAMYCIN

<u>AT</u>	ALTANA	<u>0.3%</u>	<u>N65026</u>	<u>001</u>	Sep 11, 2001
-----------	--------	-------------	---------------	------------	--------------

<u>AT</u>	BAUSCH AND LOMB	<u>0.3%</u>	<u>N64052</u>	<u>001</u>	Nov 29, 1993
-----------	-----------------	-------------	---------------	------------	--------------

<u>AT</u>	NOVEX	<u>0.3%</u>	<u>N65087</u>	<u>001</u>	Feb 25, 2002
-----------	-------	-------------	---------------	------------	--------------

TOBREX

<u>AT</u>	ALCON	<u>0.3%</u>	<u>N62535</u>	<u>001</u>	Dec 13, 1984
-----------	-------	-------------	---------------	------------	--------------

<u>AT</u>	+	FALCON PHARMS	<u>0.3%</u>	<u>N50541</u>	<u>001</u>
-----------	---	---------------	-------------	---------------	------------

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN

	AM PHARM	EQ 1.2GM BASE/VIAL	N50789	001	Jul 13, 2004
--	----------	--------------------	--------	-----	--------------

<u>AP</u>	AM PHARM PARTNERS	<u>EQ 10MG BASE/ML</u>	<u>N65122</u>	<u>001</u>	Nov 29, 2002
-----------	-------------------	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N65122</u>	<u>002</u>	Nov 29, 2002
-----------	--	------------------------	---------------	------------	--------------

+	PHARMA TEK	EQ 1.2GM BASE/VIAL	N65013	001	Aug 17, 2001
---	------------	--------------------	--------	-----	--------------

TOBRAMYCIN (PHARMACY BULK)

<u>AP</u>	AM PHARM PARTNERS	<u>EQ 40MG BASE/ML</u>	<u>N65120</u>	<u>001</u>	Nov 29, 2002
-----------	-------------------	------------------------	---------------	------------	--------------

TOBRAMYCIN SULFATE

<u>AP</u>	APOTHECON	<u>EQ 10MG BASE/ML</u>	<u>N64021</u>	<u>001</u>	May 31, 1994
-----------	-----------	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N64021</u>	<u>002</u>	May 31, 1994
-----------	--	------------------------	---------------	------------	--------------

<u>AP</u>	BAXTER HLTHCARE	<u>EQ 40MG BASE/ML</u>	<u>N63117</u>	<u>001</u>	Apr 26, 1991
-----------	-----------------	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N63118</u>	<u>001</u>	Jul 29, 1991
-----------	--	------------------------	---------------	------------	--------------

<u>AP</u>	+	HOSPIRA	<u>EQ 10MG BASE/ML</u>	<u>N63080</u>	<u>001</u>
-----------	---	---------	------------------------	---------------	------------

<u>AP</u>		<u>EQ 10MG BASE/ML</u>	<u>N63112</u>	<u>001</u>	Apr 30, 1991
-----------	--	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N63111</u>	<u>001</u>	Apr 30, 1991
-----------	--	------------------------	---------------	------------	--------------

<u>AP</u>	+		<u>EQ 40MG BASE/ML</u>	<u>N63116</u>	<u>001</u>
-----------	---	--	------------------------	---------------	------------

<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N63161</u>	<u>001</u>	May 29, 1991
-----------	--	------------------------	---------------	------------	--------------

<u>AP</u>	MARSAM PHARMS LLC	<u>EQ 10MG BASE/ML</u>	<u>N62945</u>	<u>001</u>	Aug 09, 1989
-----------	-------------------	------------------------	---------------	------------	--------------

<u>AP</u>		<u>EQ 40MG BASE/ML</u>	<u>N62945</u>	<u>002</u>	Aug 09, 1989
-----------	--	------------------------	---------------	------------	--------------

<u>AP</u>	SICOR PHARMS	<u>EQ 40MG BASE/ML</u>	<u>N63100</u>	<u>001</u>	Jan 30, 1992
-----------	--------------	------------------------	---------------	------------	--------------

TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

+	HOSPIRA	EQ 1.2MG BASE/ML	N63081	003	Jul 31, 1990
---	---------	------------------	--------	-----	--------------

+		EQ 1.6MG BASE/ML	N63081	006	Jun 02, 1993
---	--	------------------	--------	-----	--------------

+		EQ 80MG BASE/100ML	N63081	001	Jul 31, 1990
---	--	--------------------	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 342 (of 360)

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

<u>AB</u>	IVAX PHARMS	<u>100MG</u>	<u>N18894</u>	<u>001</u>	Nov 02, 1984
<u>AB</u>		<u>250MG</u>	<u>N18894</u>	<u>002</u>	Nov 02, 1984
<u>AB</u>		<u>500MG</u>	<u>N18894</u>	<u>003</u>	Nov 02, 1984
<u>AB</u>	MUTUAL PHARM	<u>100MG</u>	<u>N71357</u>	<u>001</u>	Jul 16, 1987
<u>AB</u>		<u>250MG</u>	<u>N71358</u>	<u>001</u>	Jul 16, 1987
<u>AB</u>		<u>500MG</u>	<u>N71359</u>	<u>001</u>	Jul 16, 1987
<u>AB</u>	MYLAN	<u>250MG</u>	<u>N70259</u>	<u>001</u>	Jan 02, 1986
<u>AB</u>		<u>500MG</u>	<u>N70913</u>	<u>001</u>	Mar 17, 1986
<u>AB</u>	PAR PHARM	<u>100MG</u>	<u>N70159</u>	<u>001</u>	Jan 06, 1986
<u>AB</u>		<u>250MG</u>	<u>N70160</u>	<u>001</u>	Jan 06, 1986
<u>AB</u>		<u>500MG</u>	<u>N70161</u>	<u>001</u>	Jan 06, 1986
<u>AB</u>	SANDOZ	<u>100MG</u>	<u>N71633</u>	<u>001</u>	Dec 09, 1987
<u>AB</u>		<u>250MG</u>	<u>N70289</u>	<u>001</u>	Mar 13, 1986
<u>AB</u>		<u>500MG</u>	<u>N70290</u>	<u>001</u>	Mar 13, 1986
<u>AB</u>	WATSON LABS	<u>250MG</u>	<u>N70514</u>	<u>001</u>	Jan 09, 1986
<u>AB</u>		<u>500MG</u>	<u>N70515</u>	<u>001</u>	Jan 09, 1986

TOLINASE

<u>AB</u>	PHARMACIA AND UPJOHN	<u>100MG</u>	<u>N15500</u>	<u>002</u>	
<u>AB</u>		<u>250MG</u>	<u>N15500</u>	<u>004</u>	
<u>AB</u>	+	<u>500MG</u>	<u>N15500</u>	<u>005</u>	

TOLBUTAMIDE

TABLET; ORAL

TOLBUTAMIDE

<u>AB</u>	IVAX PHARMS	<u>500MG</u>	<u>N87093</u>	<u>001</u>	
<u>AB</u>	MYLAN	<u>500MG</u>	<u>N86445</u>	<u>001</u>	
<u>AB</u>	SANDOZ	<u>500MG</u>	<u>N86574</u>	<u>001</u>	
<u>AB</u>	+	<u>500MG</u>	<u>N86109</u>	<u>001</u>	
<u>AB</u>		<u>500MG</u>	<u>N87318</u>	<u>001</u>	

TOLCAPONE

TABLET; ORAL

TASMAR

	VALEANT	100MG	N20697	001	Jan 29, 1998
	+	200MG	N20697	002	Jan 29, 1998

TOLMETIN SODIUM

CAPSULE; ORAL

TOLECTIN DS

<u>AB</u>	+	ORTHO MCNEIL PHARM	<u>EQ 400MG BASE</u>	<u>N18084</u>	<u>001</u>
-----------	---	--------------------	----------------------	---------------	------------

TOLMETIN SODIUM

<u>AB</u>	IVAX PHARMS	<u>EQ 400MG BASE</u>	<u>N73392</u>	<u>001</u>	Jan 24, 1992
<u>AB</u>	MUTUAL PHARM	<u>EQ 400MG BASE</u>	<u>N73311</u>	<u>001</u>	Nov 27, 1991
<u>AB</u>	MYLAN	<u>EQ 400MG BASE</u>	<u>N73393</u>	<u>001</u>	May 27, 1993
<u>AB</u>	PUREPAC PHARM	<u>EQ 400MG BASE</u>	<u>N73308</u>	<u>001</u>	Jan 24, 1992
<u>AB</u>	TEVA	<u>EQ 400MG BASE</u>	<u>N73290</u>	<u>001</u>	Nov 27, 1991

TABLET; ORAL

TOLECTIN

<u>AB</u>	ORTHO MCNEIL PHARM	<u>EQ 200MG BASE</u>	<u>N17628</u>	<u>001</u>	
-----------	--------------------	----------------------	---------------	------------	--

TOLECTIN 600

<u>AB</u>	+	ORTHO MCNEIL PHARM	<u>EQ 600MG BASE</u>	<u>N17628</u>	<u>002</u> Mar 08, 1989
-----------	---	--------------------	----------------------	---------------	-------------------------

TOLMETIN SODIUM

<u>AB</u>	IVAX PHARMS	<u>EQ 600MG BASE</u>	<u>N74399</u>	<u>001</u>	Mar 28, 1996
<u>AB</u>	MUTUAL PHARM	<u>EQ 200MG BASE</u>	<u>N73310</u>	<u>001</u>	Nov 27, 1991
<u>AB</u>	MYLAN	<u>EQ 600MG BASE</u>	<u>N74473</u>	<u>001</u>	Aug 30, 1994
<u>AB</u>	PUREPAC PHARM	<u>EQ 600MG BASE</u>	<u>N73527</u>	<u>001</u>	Jun 30, 1992
<u>AB</u>	SANDOZ	<u>EQ 200MG BASE</u>	<u>N73588</u>	<u>001</u>	Jul 31, 1992

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 343 (of 360)

TOLMETIN SODIUM

TABLET; ORAL

TOLMETIN SODIUM

<u>AB</u>	SANDOZ	<u>EQ 600MG BASE</u>	<u>N74002</u>	<u>001</u>	Sep 27, 1993
-----------	--------	----------------------	---------------	------------	--------------

TOLTERODINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

DETROL LA

	PHARMACIA AND UPJOHN	2MG	N21228	001	Dec 22, 2000
+		4MG	N21228	002	Dec 22, 2000

TABLET; ORAL

DETROL

	PHARMACIA AND UPJOHN	1MG	N20771	001	Mar 25, 1998
+		2MG	N20771	002	Mar 25, 1998

TOPIRAMATE

CAPSULE; ORAL

TOPAMAX SPRINKLE

	ORTHO MCNEIL PHARM	15MG	N20844	001	Oct 26, 1998
+		25MG	N20844	002	Oct 26, 1998

TABLET; ORAL

TOPAMAX

+	ORTHO MCNEIL PHARM	25MG	N20505	004	Dec 24, 1996
		50MG	N20505	005	Dec 24, 1996
		100MG	N20505	001	Dec 24, 1996
		200MG	N20505	002	Dec 24, 1996

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

HYCAMTIN

+	GLAXOSMITHKLINE	EQ 4MG BASE/VIAL	N20671	001	May 28, 1996
---	-----------------	------------------	--------	-----	--------------

TOREMIFENE CITRATE

TABLET; ORAL

FARESTON

+	GTX INC	EQ 60MG BASE	N20497	001	May 29, 1997
---	---------	--------------	--------	-----	--------------

TORSEMIDE

INJECTABLE; INJECTION

DEMADEX

+	ROCHE	10MG/ML	N20137	002	Aug 23, 1993
---	-------	---------	--------	-----	--------------

TABLET; ORAL

DEMADEX

<u>AB</u>	ROCHE	<u>5MG</u>	<u>N20136</u>	<u>001</u>	Aug 23, 1993
<u>AB</u>		<u>10MG</u>	<u>N20136</u>	<u>002</u>	Aug 23, 1993
<u>AB</u>	+	<u>20MG</u>	<u>N20136</u>	<u>003</u>	Aug 23, 1993
<u>AB</u>		<u>100MG</u>	<u>N20136</u>	<u>004</u>	Aug 23, 1993

TORSEMIDE

<u>AB</u>	APOTEX	<u>5MG</u>	<u>N76894</u>	<u>001</u>	May 31, 2005
<u>AB</u>		<u>10MG</u>	<u>N76894</u>	<u>002</u>	May 31, 2005
<u>AB</u>		<u>20MG</u>	<u>N76894</u>	<u>003</u>	May 31, 2005
<u>AB</u>		<u>100MG</u>	<u>N76894</u>	<u>004</u>	May 31, 2005
<u>AB</u>	PAR PHARM	<u>5MG</u>	<u>N76226</u>	<u>001</u>	May 27, 2003
<u>AB</u>		<u>10MG</u>	<u>N76226</u>	<u>002</u>	May 27, 2003
<u>AB</u>		<u>20MG</u>	<u>N76226</u>	<u>003</u>	May 27, 2003
<u>AB</u>		<u>100MG</u>	<u>N76226</u>	<u>004</u>	May 27, 2003
<u>AB</u>	PLIVA PHARM IND	<u>5MG</u>	<u>N76346</u>	<u>001</u>	May 30, 2003
<u>AB</u>		<u>10MG</u>	<u>N76346</u>	<u>002</u>	May 30, 2003
<u>AB</u>		<u>20MG</u>	<u>N76346</u>	<u>003</u>	May 30, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 344 (of 360)

TORSEMIDE

TABLET; ORAL

TORSEMIDE

<u>AB</u>	PLIVA PHARM IND	<u>100MG</u>	<u>N76346</u>	<u>004</u>	Oct 19, 2004
<u>AB</u>	ROXANE	<u>5MG</u>	<u>N76943</u>	<u>001</u>	Mar 01, 2005
<u>AB</u>		<u>10MG</u>	<u>N76943</u>	<u>002</u>	Mar 01, 2005
<u>AB</u>		<u>20MG</u>	<u>N76943</u>	<u>003</u>	Mar 01, 2005
<u>AB</u>	TEVA	<u>5MG</u>	<u>N76110</u>	<u>001</u>	May 14, 2002
<u>AB</u>		<u>10MG</u>	<u>N76110</u>	<u>002</u>	May 14, 2002
<u>AB</u>		<u>20MG</u>	<u>N76110</u>	<u>003</u>	May 14, 2002
<u>AB</u>		<u>100MG</u>	<u>N76110</u>	<u>004</u>	May 14, 2002

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

<u>AB</u>	ALPHAPHARM	<u>50MG</u>	<u>N75980</u>	<u>001</u>	Nov 21, 2002
<u>AB</u>	CARACO	<u>50MG</u>	<u>N75964</u>	<u>001</u>	Jun 19, 2002
<u>AB</u>	COREPHARMA	<u>50MG</u>	<u>N76003</u>	<u>001</u>	Jun 20, 2002
<u>AB</u>	IVAX PHARMS	<u>50MG</u>	<u>N75963</u>	<u>001</u>	Jul 03, 2002
<u>AB</u>	MALLINCKRODT	<u>50MG</u>	<u>N75983</u>	<u>001</u>	Jun 25, 2002
<u>AB</u>	MUTUAL PHARM	<u>50MG</u>	<u>N76100</u>	<u>001</u>	Jun 20, 2002
<u>AB</u>	MYLAN	<u>50MG</u>	<u>N75986</u>	<u>001</u>	Jun 21, 2002
<u>AB</u>	PLIVA	<u>50MG</u>	<u>N75982</u>	<u>001</u>	Jul 01, 2002
<u>AB</u>	PUREPAC PHARM	<u>50MG</u>	<u>N75960</u>	<u>001</u>	Jun 19, 2002
<u>AB</u>	SANDOZ	<u>50MG</u>	<u>N75968</u>	<u>001</u>	Jun 25, 2002
<u>AB</u>	TEVA	<u>50MG</u>	<u>N75977</u>	<u>001</u>	Jun 19, 2002
<u>AB</u>	TORPHARM	<u>50MG</u>	<u>N75981</u>	<u>001</u>	Jul 10, 2002
<u>AB</u>	WATSON LABS	<u>50MG</u>	<u>N75962</u>	<u>001</u>	Jun 24, 2002

ULTRAM

<u>AB</u>	+ ORTHO MCNEIL PHARM	<u>50MG</u>	<u>N20281</u>	<u>002</u>	Mar 03, 1995
-----------	----------------------	-------------	---------------	------------	--------------

TABLET, EXTENDED RELEASE; ORAL

TRAMADOL HCL

	BIOVAIL LABS INTL	100MG	N21692	001	Sep 08, 2005
		200MG	N21692	002	Sep 08, 2005
+		300MG	N21692	003	Sep 08, 2005

TABLET, ORALLY DISINTEGRATING; ORAL

TRAMADOL HYDROCHLORIDE

+	BIOVAIL	50MG	N21693	001	May 05, 2005
---	---------	------	--------	-----	--------------

TRANDOLAPRIL

TABLET; ORAL

MAVIK

	ABBOTT	1MG	N20528	001	Apr 26, 1996
		2MG	N20528	002	Apr 26, 1996
+		4MG	N20528	003	Apr 26, 1996

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARKA

+	ABBOTT	1MG;240MG	N20591	003	Oct 22, 1996
+		2MG;180MG	N20591	001	Oct 22, 1996
+		2MG;240MG	N20591	004	Oct 22, 1996
+		4MG;240MG	N20591	002	Oct 22, 1996

TRANEXAMIC ACID

INJECTABLE; INJECTION

CYKLOKAPRON

+	PHARMACIA AND UPJOHN	100MG/ML	N19281	001	Dec 30, 1986
---	----------------------	----------	--------	-----	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 345 (of 360)

TRANLYCYPROMINE SULFATE

TABLET; ORAL

PARNATE

+ GLAXOSMITHKLINE	EQ 10MG BASE	N12342 003	Aug 16, 1985
-------------------	--------------	------------	--------------

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC

TRAVATAN

+ ALCON	0.004%	N21257 001	Mar 16, 2001
---------	--------	------------	--------------

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

DESYREL

<u>AB</u> APOTHECON	<u>50MG</u>	<u>N18207</u>	<u>001</u>	
<u>AB</u>	<u>100MG</u>	<u>N18207</u>	<u>002</u>	
<u>AB</u>	<u>150MG</u>	<u>N18207</u>	<u>003</u>	Mar 25, 1985
<u>AB</u> +	<u>300MG</u>	<u>N18207</u>	<u>004</u>	Nov 07, 1988

TRAZODONE HYDROCHLORIDE

<u>AB</u> BARR	<u>50MG</u>	<u>N71258</u>	<u>001</u>	Mar 25, 1987
<u>AB</u>	<u>100MG</u>	<u>N71196</u>	<u>001</u>	Mar 25, 1987
<u>AB</u>	<u>150MG</u>	<u>N71196</u>	<u>002</u>	Apr 26, 1999
<u>AB</u>	<u>300MG</u>	<u>N71196</u>	<u>003</u>	Apr 26, 1999
<u>AB</u> MUTUAL PHARM	<u>50MG</u>	<u>N73136</u>	<u>001</u>	Mar 24, 1993
<u>AB</u>	<u>100MG</u>	<u>N73137</u>	<u>001</u>	Mar 24, 1993
<u>AB</u>	<u>150MG</u>	<u>N73138</u>	<u>001</u>	Dec 22, 1995
<u>AB</u> MYLAN	<u>50MG</u>	<u>N71405</u>	<u>001</u>	Feb 27, 1991
<u>AB</u> PLIVA	<u>50MG</u>	<u>N71523</u>	<u>001</u>	Dec 11, 1987
<u>AB</u>	<u>100MG</u>	<u>N71524</u>	<u>001</u>	Dec 11, 1987
<u>AB</u>	<u>150MG</u>	<u>N71525</u>	<u>001</u>	Mar 09, 1988
<u>AB</u> PUREPAC PHARM	<u>50MG</u>	<u>N71636</u>	<u>001</u>	Apr 18, 1988
<u>AB</u>	<u>100MG</u>	<u>N71514</u>	<u>001</u>	Apr 18, 1988
<u>AB</u> SANDOZ	<u>50MG</u>	<u>N72484</u>	<u>001</u>	Apr 30, 1990
<u>AB</u>	<u>100MG</u>	<u>N72483</u>	<u>001</u>	Apr 30, 1990
<u>AB</u> TEVA	<u>50MG</u>	<u>N72192</u>	<u>001</u>	Feb 02, 1989
<u>AB</u>	<u>100MG</u>	<u>N72193</u>	<u>001</u>	Feb 02, 1989
<u>AB</u>	<u>150MG</u>	<u>N74357</u>	<u>001</u>	Apr 30, 1997
<u>AB</u> WATSON LABS	<u>50MG</u>	<u>N70857</u>	<u>001</u>	Oct 10, 1986
<u>AB</u>	<u>100MG</u>	<u>N70858</u>	<u>001</u>	Oct 10, 1986

TREPROSTINIL SODIUM

INJECTABLE; IV (INFUSION)-SC

REMODULIN

UNITED THERAP	1MG/ML	N21272 001	May 21, 2002
	2.5MG/ML	N21272 002	May 21, 2002
	5MG/ML	N21272 003	May 21, 2002
+	10MG/ML	N21272 004	May 21, 2002

TRETINOIN

CAPSULE; ORAL

VESANOID

+ ROCHE	10MG	N20438 001	Nov 22, 1995
---------	------	------------	--------------

CREAM; TOPICAL

AVITA

<u>AB</u> MYLAN BERTEK	<u>0.025%</u>	<u>N20404</u>	<u>003</u>	Jan 14, 1997
-------------------------------	----------------------	----------------------	-------------------	--------------

RENOVA

<u>AB2</u> + JOHNSON AND JOHNSON	<u>0.05%</u>	<u>N19963</u>	<u>001</u>	Dec 29, 1995
+	0.02%	N21108 001		Aug 31, 2000

RETIN-A

<u>AB</u> + JOHNSON AND JOHNSON	<u>0.025%</u>	<u>N19049</u>	<u>001</u>	Sep 16, 1988
--	----------------------	----------------------	-------------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 346 (of 360)

TRETINOIN

CREAM; TOPICAL

RETIN-A

<u>AB</u> + JOHNSON AND JOHNSON	<u>0.1%</u>	<u>N17340</u>	<u>001</u>	
<u>AB1</u> +	<u>0.05%</u>	<u>N17522</u>	<u>001</u>	

TRETINOIN

<u>AB</u> SPEAR PHARMS	<u>0.025%</u>	<u>N75264</u>	<u>001</u>	Dec 24, 1998
<u>AB</u>	<u>0.1%</u>	<u>N75213</u>	<u>001</u>	Dec 24, 1998
<u>AB1</u>	<u>0.05%</u>	<u>N75265</u>	<u>001</u>	Dec 24, 1998
<u>AB2</u>	<u>0.05%</u>	<u>N76498</u>	<u>001</u>	Sep 15, 2005

GEL; TOPICAL

AVITA

BT MYLAN BERTEK	0.025%	N20400	001	Jan 29, 1998
-----------------	--------	--------	-----	--------------

RETIN-A

<u>AB</u> + JOHNSON AND JOHNSON	<u>0.01%</u>	<u>N17955</u>	<u>001</u>	
<u>AB</u> +	<u>0.025%</u>	<u>N17579</u>	<u>002</u>	

RETIN-A MICRO

+ JOHNSON AND JOHNSON	0.04%	N20475	002	May 10, 2002
+	0.1%	N20475	001	Feb 07, 1997

TRETINOIN

<u>AB</u> SPEAR PHARMS	<u>0.01%</u>	<u>N75589</u>	<u>001</u>	Jun 11, 2002
<u>AB</u>	<u>0.025%</u>	<u>N75529</u>	<u>001</u>	Feb 22, 2000

SOLUTION; TOPICAL

RETIN-A

<u>AT</u> + JOHNSON AND JOHNSON	<u>0.05%</u>	<u>N16921</u>	<u>001</u>	
---------------------------------	--------------	---------------	------------	--

TRETINOIN

<u>AT</u> MORTON GROVE	<u>0.05%</u>	<u>N75260</u>	<u>001</u>	Jan 25, 1999
<u>AT</u> TEVA PHARMS	<u>0.05%</u>	<u>N74873</u>	<u>001</u>	Jun 19, 1998

TRIAMCINOLONE

TABLET; ORAL

ARISTOCORT

BP + ASTELLAS	4MG	N11161	007	
---------------	-----	--------	-----	--

KENACORT

BP BRISTOL MYERS SQUIBB	4MG	N11283	006	
-------------------------	-----	--------	-----	--

TRIAMCINOLONE ACETONIDE

AEROSOL, METERED; INHALATION

AZMACORT

+ KOS LIFE	0.1MG/INH	N18117	001	Apr 23, 1982
------------	-----------	--------	-----	--------------

CREAM; TOPICAL

ARISTOCORT A

<u>AT</u> ASTELLAS	<u>0.025%</u>	<u>N88818</u>	<u>001</u>	Oct 16, 1984
<u>AT</u>	<u>0.1%</u>	<u>N88819</u>	<u>001</u>	Oct 16, 1984
<u>AT</u>	<u>0.5%</u>	<u>N88820</u>	<u>001</u>	Oct 16, 1984

KENALOG

<u>AT</u> + APOTHECON	<u>0.025%</u>	<u>N11601</u>	<u>003</u>	
<u>AT</u> +	<u>0.1%</u>	<u>N11601</u>	<u>006</u>	
<u>AT</u> +	<u>0.5%</u>	<u>N83943</u>	<u>001</u>	

TRIACET

<u>AT</u> TEVA	<u>0.025%</u>	<u>N84908</u>	<u>001</u>	
<u>AT</u>	<u>0.1%</u>	<u>N84908</u>	<u>002</u>	
<u>AT</u>	<u>0.5%</u>	<u>N84908</u>	<u>003</u>	

TRIAMCINOLONE ACETONIDE

<u>AT</u> ALPHARMA US PHARMS	<u>0.1%</u>	<u>N87798</u>	<u>001</u>	Jun 04, 1982
<u>AT</u> ALTANA	<u>0.025%</u>	<u>N85692</u>	<u>001</u>	
<u>AT</u>	<u>0.1%</u>	<u>N85692</u>	<u>003</u>	
<u>AT</u>	<u>0.5%</u>	<u>N85692</u>	<u>002</u>	
<u>AT</u> CLAY PARK	<u>0.025%</u>	<u>N86415</u>	<u>001</u>	
<u>AT</u>	<u>0.1%</u>	<u>N86414</u>	<u>001</u>	

PRESCRIPTION DRUG PRODUCT LIST

3 - 347 (of 360)

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL				
<u>TRIAMCINOLONE ACETONIDE</u>				
<u>AT</u>	CLAY PARK	<u>0.5%</u>	<u>N86413</u>	<u>001</u>
<u>AT</u>	G AND W LABS	<u>0.025%</u>	<u>N89797</u>	<u>001</u> May 31, 1991
<u>AT</u>		<u>0.1%</u>	<u>N89798</u>	<u>001</u> May 31, 1991
<u>AT</u>	TARO	<u>0.025%</u>	<u>N86277</u>	<u>001</u>
<u>AT</u>		<u>0.1%</u>	<u>N40039</u>	<u>001</u> Nov 26, 1997
<u>AT</u>		<u>0.1%</u>	<u>N86276</u>	<u>001</u>
<u>AT</u>		<u>0.5%</u>	<u>N86275</u>	<u>001</u>
<u>TRIDERM</u>				
<u>AT</u>	DEL RAY LABS	<u>0.1%</u>	<u>N88042</u>	<u>001</u> Mar 19, 1984
INJECTABLE; INJECTION				
KENALOG-10				
	APOTHECON	10MG/ML	N12041	001
KENALOG-40				
	+ APOTHECON	40MG/ML	N14901	001
LOTION; TOPICAL				
<u>KENALOG</u>				
<u>AT</u>	APOTHECON	<u>0.025%</u>	<u>N84343</u>	<u>001</u>
<u>AT</u>	+	<u>0.1%</u>	<u>N84343</u>	<u>002</u>
<u>TRIAMCINOLONE ACETONIDE</u>				
<u>AT</u>	ALTANA	<u>0.025%</u>	<u>N40467</u>	<u>001</u> Apr 21, 2003
<u>AT</u>		<u>0.1%</u>	<u>N40467</u>	<u>002</u> Apr 21, 2003
<u>AT</u>	MORTON GROVE	<u>0.025%</u>	<u>N88450</u>	<u>001</u> Apr 01, 1985
<u>AT</u>		<u>0.1%</u>	<u>N88451</u>	<u>001</u> Apr 03, 1985
<u>AT</u>	TARO	<u>0.1%</u>	<u>N89129</u>	<u>001</u> Aug 14, 1986
OINTMENT; TOPICAL				
<u>KENALOG</u>				
<u>AT</u>	APOTHECON	<u>0.025%</u>	<u>N11600</u>	<u>003</u>
<u>AT</u>		<u>0.1%</u>	<u>N11600</u>	<u>001</u>
<u>TRIAMCINOLONE ACETONIDE</u>				
<u>AT</u>	ALPHARMA US PHARMS	<u>0.1%</u>	<u>N87799</u>	<u>001</u> Jun 07, 1982
<u>AT</u>	ALTANA	<u>0.025%</u>	<u>N85691</u>	<u>001</u>
<u>AT</u>		<u>0.1%</u>	<u>N85691</u>	<u>003</u>
<u>AT</u>		<u>0.5%</u>	<u>N85691</u>	<u>002</u>
<u>AT</u>	+	<u>0.025%</u>	<u>N87356</u>	<u>001</u>
<u>AT</u>	+	<u>0.1%</u>	<u>N87357</u>	<u>001</u>
<u>AT</u>	+	<u>0.5%</u>	<u>N87385</u>	<u>001</u>
<u>AT</u>	TARO	<u>0.025%</u>	<u>N40040</u>	<u>001</u> Sep 30, 1994
<u>AT</u>		<u>0.025%</u>	<u>N40374</u>	<u>001</u> Jun 05, 2001
<u>AT</u>		<u>0.1%</u>	<u>N40037</u>	<u>001</u> Sep 30, 1994
<u>AT</u>		<u>0.1%</u>	<u>N87902</u>	<u>001</u> Dec 27, 1982
<u>AT</u>		<u>0.5%</u>	<u>N40386</u>	<u>001</u> Jun 05, 2001
TRIAMCINOLONE ACETONIDE IN ABSORBASE				
	+ CAROLINA MEDCL	0.05%	N89595	001 Mar 23, 1995
PASTE; DENTAL				
<u>KENALOG IN ORABASE</u>				
<u>AT</u>	+	<u>0.1%</u>	<u>N12097</u>	<u>001</u>
<u>ORACORT</u>				
<u>AT</u>	TARO	<u>0.1%</u>	<u>N70730</u>	<u>001</u> Oct 01, 1986
<u>ORALONE</u>				
<u>AT</u>	TARO	<u>0.1%</u>	<u>N71383</u>	<u>001</u> Jul 06, 1987
SPRAY; TOPICAL				
KENALOG				
	+ APOTHECON	0.147MG/GM	N12104	001
SPRAY, METERED; NASAL				
NASACORT AQ				
	+ AVENTIS	0.055MG/SPRAY	N20468	001 May 20, 1996

PRESCRIPTION DRUG PRODUCT LIST

3 - 348 (of 360)

TRIAMCINOLONE ACETONIDE

SPRAY, METERED; NASAL NASACORT HFA				
+ AVENTIS PHARMS	0.055MG/SPRAY	N20784	001	Apr 07, 2004

TRIAMCINOLONE HEXACETONIDE

INJECTABLE; INJECTION ARISTOSPAN				
+ SABEX 2002	5MG/ML	N16466	001	
+	20MG/ML	N16466	002	

TRIAMTERENE

CAPSULE; ORAL DYRENIUM				
WELLSPRING PHARM	50MG	N13174	001	
+	100MG	N13174	002	

TRIAZOLAM

TABLET; ORAL				
<u>HALCION</u>				
<u>AB</u>	PHARMACIA AND UPJOHN	<u>0.125MG</u>	<u>N17892</u>	<u>003</u> Apr 26, 1985
<u>AB</u>	+	<u>0.25MG</u>	<u>N17892</u>	<u>001</u> Nov 15, 1982
<u>TRIAZOLAM</u>				
<u>AB</u>	ALPHAPHARM	<u>0.125MG</u>	<u>N74031</u>	<u>001</u> Mar 25, 1994
<u>AB</u>		<u>0.25MG</u>	<u>N74031</u>	<u>002</u> Mar 25, 1994
<u>AB</u>	ROXANE	<u>0.125MG</u>	<u>N74224</u>	<u>001</u> Jun 01, 1994
<u>AB</u>		<u>0.25MG</u>	<u>N74224</u>	<u>002</u> Jun 01, 1994
<u>AB</u>	WATSON LABS	<u>0.125MG</u>	<u>N74445</u>	<u>001</u> Oct 20, 1995
<u>AB</u>		<u>0.25MG</u>	<u>N74445</u>	<u>002</u> Oct 20, 1995

TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL SYPRINE				
+ MERCK	250MG	N19194	001	Nov 08, 1985

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE; ORAL				
<u>STELAZINE</u>				
<u>AA</u>	+ GLAXOSMITHKLINE	<u>EQ 10MG BASE/ML</u>	<u>N11552</u>	<u>006</u>
INJECTABLE; INJECTION				
	STELAZINE			
	+ GLAXOSMITHKLINE	EQ 2MG BASE/ML	N11552	005
TABLET; ORAL				
<u>STELAZINE</u>				
<u>AB</u>	GLAXOSMITHKLINE	<u>EQ 1MG BASE</u>	<u>N11552</u>	<u>001</u>
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N11552</u>	<u>002</u>
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>N11552</u>	<u>003</u>
<u>AB</u>	+	<u>EQ 10MG BASE</u>	<u>N11552</u>	<u>004</u>
<u>TRIFLUOPERAZINE HYDROCHLORIDE</u>				
<u>AB</u>	MYLAN	<u>EQ 1MG BASE</u>	<u>N40209</u>	<u>001</u> Jul 07, 1997
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N40209</u>	<u>002</u> Jul 07, 1997
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>N40209</u>	<u>003</u> Jul 07, 1997
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N40209</u>	<u>004</u> Jul 07, 1997
<u>AB</u>	SANDOZ	<u>EQ 1MG BASE</u>	<u>N40153</u>	<u>001</u> Oct 25, 1996
<u>AB</u>		<u>EQ 1MG BASE</u>	<u>N85785</u>	<u>001</u>
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N40153</u>	<u>002</u> Oct 25, 1996
<u>AB</u>		<u>EQ 2MG BASE</u>	<u>N85786</u>	<u>001</u>
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>N40153</u>	<u>003</u> Oct 25, 1996
<u>AB</u>		<u>EQ 5MG BASE</u>	<u>N85789</u>	<u>001</u>

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 349 (of 360)

TRIFLUOPERAZINE HYDROCHLORIDE

TABLET; ORAL

TRIFLUOPERAZINE HYDROCHLORIDE

<u>AB</u>	SANDOZ	<u>EQ 10MG BASE</u>	<u>N40153</u>	<u>004</u>	Oct 25, 1996
<u>AB</u>		<u>EQ 10MG BASE</u>	<u>N85788</u>	<u>001</u>	

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINE

<u>AT</u>	ALCON	<u>1%</u>	<u>N74311</u>	<u>001</u>	Oct 06, 1995
-----------	-------	-----------	---------------	------------	--------------

VIROPTIC

<u>AT</u>	+ MONARCH PHARMS	<u>1%</u>	<u>N18299</u>	<u>001</u>	
-----------	------------------	-----------	---------------	------------	--

TRIHEXYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

<u>AA</u>	MIKART	<u>2MG/5ML</u>	<u>N40251</u>	<u>001</u>	Sep 27, 1999
<u>AA</u>	PHARM ASSOC	<u>2MG/5ML</u>	<u>N40177</u>	<u>001</u>	Apr 17, 1997
<u>AA</u>	+ PHARM VENTURES	<u>2MG/5ML</u>	<u>N89514</u>	<u>001</u>	Apr 07, 1989

TABLET; ORAL

TRIHEXYPHENIDYL HYDROCHLORIDE

<u>AA</u>	VINTAGE PHARMS	<u>2MG</u>	<u>N40254</u>	<u>001</u>	Dec 24, 1998
<u>AA</u>		<u>5MG</u>	<u>N40254</u>	<u>002</u>	Dec 24, 1998
<u>AA</u>	WATSON LABS	<u>2MG</u>	<u>N40184</u>	<u>001</u>	Feb 06, 1998
<u>AA</u>	+	<u>2MG</u>	<u>N84363</u>	<u>001</u>	
<u>AA</u>		<u>5MG</u>	<u>N40184</u>	<u>002</u>	Feb 06, 1998
<u>AA</u>	+	<u>5MG</u>	<u>N84364</u>	<u>001</u>	
<u>AA</u>	WEST WARD	<u>2MG</u>	<u>N40337</u>	<u>002</u>	Feb 16, 2000
<u>AA</u>		<u>5MG</u>	<u>N40337</u>	<u>001</u>	Feb 16, 2000

TRIMETHADIONE

TABLET; ORAL

TRIDIONE

+	ABBOTT	150MG	N05856	009	
---	--------	-------	--------	-----	--

TRIMETHOBENZAMIDE HYDROCHLORIDE

CAPSULE; ORAL

TIGAN

<u>AB</u>	+ KING PHARMS	<u>300MG</u>	<u>N17531</u>	<u>006</u>	Dec 13, 2001
-----------	---------------	--------------	---------------	------------	--------------

TRIMETHOBENZAMIDE HYDROCHLORIDE

<u>AB</u>	AMIDE PHARM	<u>300MG</u>	<u>N76546</u>	<u>001</u>	Aug 20, 2003
<u>AB</u>	MUTUAL PHARMA	<u>300MG</u>	<u>N76570</u>	<u>001</u>	Aug 28, 2003

INJECTABLE; INJECTION

TIGAN

<u>AP</u>	+ KING PHARMS	<u>100MG/ML</u>	<u>N17530</u>	<u>001</u>	
-----------	---------------	-----------------	---------------	------------	--

TRIMETHOBENZAMIDE HYDROCHLORIDE

<u>AP</u>	HOSPIRA	<u>100MG/ML</u>	<u>N88804</u>	<u>001</u>	Apr 03, 1987
-----------	---------	-----------------	---------------	------------	--------------

TRIMETHOPRIM

TABLET; ORAL

PROLOPRIM

<u>AB</u>	MONARCH PHARMS	<u>100MG</u>	<u>N17943</u>	<u>001</u>	
-----------	----------------	--------------	---------------	------------	--

TRIMETHOPRIM

<u>AB</u>	TEVA	<u>100MG</u>	<u>N18679</u>	<u>001</u>	Jul 30, 1982
	+	<u>200MG</u>	<u>N71259</u>	<u>001</u>	Jun 18, 1987
<u>AB</u>	WATSON LABS	<u>100MG</u>	<u>N70049</u>	<u>001</u>	Jun 06, 1985

PRESCRIPTION DRUG PRODUCT LIST

3 - 350 (of 360)

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL					
PRIMSOL					
+ FSC	EQ 50MG BASE/5ML	N74973	001	Jan 24,	2000

TRIMETREXATE GLUCURONATE

INJECTABLE; INJECTION					
NEUTREXIN					
+ MEDIMMUNE ONCOLOGY	EQ 25MG BASE/VIAL	N20326	001	Dec 17,	1993
+	EQ 200MG BASE/VIAL	N20326	002	Jul 31,	1998

TRIMIPRAMINE MALEATE

CAPSULE; ORAL					
SURMONTIL					
ODYSSEY PHARMS	EQ 25MG BASE	N16792	001		
	EQ 50MG BASE	N16792	002		
+	EQ 100MG BASE	N16792	003	Sep 15,	1982

TRIPLENNAMINE HYDROCHLORIDE

TABLET; ORAL					
PBZ					
+ NOVARTIS	50MG	N05914	002		

TRIPTORELIN PAMOATE

INJECTABLE; INTRAMUSCULAR					
TRELSTAR					
+ WATSON LABS	11.25MG/VIAL	N21288	001	Jun 29,	2001
TRELSTAR DEPOT					
+ WATSON LABS	EQ 3.75MG BASE/VIAL	N20715	001	Jun 15,	2000

TROLEANDOMYCIN

CAPSULE; ORAL					
TAO					
+ PFIZER	EQ 250MG BASE	N50336	002		

TROMETHAMINE

INJECTABLE; INJECTION					
THAM					
+ HOSPIRA	3.6GM/100ML	N13025	002		

TROPICAMIDE

SOLUTION/DROPS; OPHTHALMIC					
<u>MYDRIACYL</u>					
<u>AT</u>	+	ALCON	<u>0.5%</u>	<u>N84305</u>	<u>001</u>
<u>AT</u>	+		<u>1%</u>	<u>N84306</u>	<u>001</u>
<u>TROPICACYL</u>					
<u>AT</u>		AKORN	<u>0.5%</u>	<u>N40314</u>	<u>001</u> Sep 29, 2000
<u>AT</u>			<u>1%</u>	<u>N40315</u>	<u>001</u> Sep 29, 2000
<u>TROPICAMIDE</u>					
<u>AT</u>		ALCON UNIVERSAL	<u>1%</u>	<u>N89172</u>	<u>001</u> Dec 28, 1990
<u>AT</u>		BAUSCH AND LOMB	<u>0.5%</u>	<u>N40067</u>	<u>001</u> Jul 27, 1994
<u>AT</u>			<u>1%</u>	<u>N40064</u>	<u>001</u> Jul 27, 1994
<u>AT</u>		MIZA PHARMS USA	<u>0.5%</u>	<u>N87636</u>	<u>001</u> Jul 30, 1982
<u>AT</u>			<u>1%</u>	<u>N87637</u>	<u>001</u> Aug 09, 1982

TROSPIUM CHLORIDE

TABLET; ORAL					
SANCTURA					
+ INDEVUS	20MG	N21595	001	May 28,	2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 351 (of 360)

TRYPAN BLUESOLUTION; OPHTHALMIC
VISIONBLUE

+ DORC 0.06% N21670 001 Dec 16, 2004

UNOPROSTONE ISOPROPYLSOLUTION/DROPS; OPHTHALMIC
RESCULA

+ NOVARTIS 0.15% N21214 001 Aug 03, 2000

UREAINJECTABLE; INJECTION
UREAPHIL

+ HOSPIRA 40GM/VIAL N12154 001

UREA, C-13FOR SOLUTION; ORAL
BREATHTEK UBT FOR H-PYLORI

+ MERETEK EQ 75MG /POUCHE N20586 002 May 10, 2001

UREA, C-14CAPSULE; ORAL
PYTEST

+ BALLARD MEDCL 1uCi N20617 001 May 09, 1997

PYTEST KIT

+ BALLARD MEDCL 1uCi N20617 002 May 09, 1997

UROFOLLITROPININJECTABLE; INTRAMUSCULAR, SUBCUTANEOUS
BRAVELLE

+ FERRING 75 IU/VIAL N21289 001 May 06, 2002

UROKINASEINJECTABLE; INJECTION
ABBOKINASE

+ ABBOTT 250,000 IU/VIAL N21846 001

URSODIOL

CAPSULE; ORAL

ACTIGALLAB + WATSON PHARMS 300MG N19594 002 Dec 31, 1987URSODIOLAB AMIDE PHARM 300MG N75517 001 Mar 14, 2000AB TEVA PHARMS 300MG N75592 001 May 25, 2000

TABLET; ORAL

URSO 250

AXCAN SCANDIPHARM 250MG N20675 001 Dec 10, 1997

URSO FORTE

+ AXCAN SCANDIPHARM 500MG N20675 002 Jul 21, 2004

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALTREX

GLAXOSMITHKLINE EQ 500MG BASE N20487 001 Jun 23, 1995

+ EQ 1GM BASE N20487 002 Jun 23, 1995

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 352 (of 360)

VALDECOXIB

TABLET; ORAL

BEXTRA

GD SEARLE LLC	10MG	N21341	002	Nov 16, 2001
+	20MG	N21341	003	Nov 16, 2001

VALGANCICLOVIR HYDROCHLORIDE

TABLET; ORAL

VALCYTE

+ ROCHE PALO	EQ 450MG BASE	N21304	001	Mar 29, 2001
--------------	---------------	--------	-----	--------------

VALPROATE SODIUM

INJECTABLE; INJECTION

DEPACON

<u>AP</u> + ABBOTT	<u>EQ 100MG BASE/ML</u>	<u>N20593</u>	<u>001</u>	Dec 30, 1996
--------------------	-------------------------	---------------	------------	--------------

VALPROATE SODIUM

<u>AP</u> AM PHARM	<u>EQ 100MG BASE/ML</u>	<u>N76539</u>	<u>001</u>	Jun 26, 2003
--------------------	-------------------------	---------------	------------	--------------

<u>AP</u> BEDFORD	<u>EQ 100MG BASE/ML</u>	<u>N76295</u>	<u>001</u>	Nov 14, 2002
-------------------	-------------------------	---------------	------------	--------------

VALPROIC ACID

CAPSULE; ORAL

DEPAKENE

<u>AB</u> + ABBOTT	<u>250MG</u>	<u>N18081</u>	<u>001</u>	
--------------------	--------------	---------------	------------	--

VALPROIC ACID

<u>AB</u> BANNER PHARMACAPS	<u>250MG</u>	<u>N73484</u>	<u>001</u>	Jun 29, 1993
-----------------------------	--------------	---------------	------------	--------------

<u>AB</u> SCHERER RP	<u>250MG</u>	<u>N73229</u>	<u>001</u>	Oct 29, 1991
----------------------	--------------	---------------	------------	--------------

<u>AB</u> USL PHARMA	<u>250MG</u>	<u>N70631</u>	<u>001</u>	Jun 11, 1987
----------------------	--------------	---------------	------------	--------------

SYRUP; ORAL

DEPAKENE

<u>AA</u> + ABBOTT	<u>250MG/5ML</u>	<u>N18082</u>	<u>001</u>	
--------------------	------------------	---------------	------------	--

MYPROIC ACID

<u>AA</u> MORTON GROVE	<u>250MG/5ML</u>	<u>N70868</u>	<u>001</u>	Jul 01, 1986
------------------------	------------------	---------------	------------	--------------

VALPROIC ACID

<u>AA</u> APOTEX	<u>250MG/5ML</u>	<u>N77105</u>	<u>001</u>	Jul 29, 2005
------------------	------------------	---------------	------------	--------------

<u>AA</u> HIGH TECH PHARMA	<u>250MG/5ML</u>	<u>N74060</u>	<u>001</u>	Jan 13, 1995
----------------------------	------------------	---------------	------------	--------------

<u>AA</u> PHARM ASSOC	<u>250MG/5ML</u>	<u>N75379</u>	<u>001</u>	Dec 15, 2000
-----------------------	------------------	---------------	------------	--------------

<u>AA</u> TEVA PHARMS	<u>250MG/5ML</u>	<u>N73178</u>	<u>001</u>	Aug 25, 1992
-----------------------	------------------	---------------	------------	--------------

<u>AA</u> UDL	<u>250MG/5ML</u>	<u>N75782</u>	<u>001</u>	Dec 22, 2000
---------------	------------------	---------------	------------	--------------

VALRUBICIN

SOLUTION; INTRAVESICAL

VALSTAR PRESERVATIVE FREE

+ ANTHRA	40MG/ML	N20892	001	Sep 25, 1998
----------	---------	--------	-----	--------------

VALSARTAN

TABLET; ORAL

DIOVAN

NOVARTIS	40MG	N21283	004	Aug 14, 2002
	80MG	N21283	001	Jul 18, 2001
	160MG	N21283	002	Jul 18, 2001
+	320MG	N21283	003	Jul 18, 2001

VANCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

VANCOCIN HYDROCHLORIDE

VIROPHARMA	EQ 125MG BASE	N50606	001	Apr 15, 1986
+	EQ 250MG BASE	N50606	002	Apr 15, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 353 (of 360)

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER

+	BAXTER HLTHCARE	EQ 500MG BASE/100ML	N50671	001	Apr 29, 1993
---	-----------------	---------------------	--------	-----	--------------

VANCOMYCIN HYDROCHLORIDE

<u>AP</u>	+	AM PHARM PARTNERS	<u>EQ 500MG BASE/VIAL</u>	<u>N62663</u>	<u>001</u>	Mar 17, 1987
<u>AP</u>	+		<u>EQ 1GM BASE/VIAL</u>	<u>N62663</u>	<u>002</u>	Jul 31, 1987
<u>AP</u>	+		<u>EQ 5GM BASE/VIAL</u>	<u>N62663</u>	<u>003</u>	Jun 03, 1988
	+		EQ 10GM BASE/VIAL	N62663	004	Nov 28, 1997
<u>AP</u>	+	HOSPIRA	<u>EQ 500MG BASE/VIAL</u>	<u>N62911</u>	<u>001</u>	Aug 04, 1988
<u>AP</u>	+		<u>EQ 500MG BASE/VIAL</u>	<u>N62931</u>	<u>001</u>	Oct 29, 1992
<u>AP</u>	+		<u>EQ 1GM BASE/VIAL</u>	<u>N62912</u>	<u>001</u>	Aug 04, 1988
<u>AP</u>	+		<u>EQ 1GM BASE/VIAL</u>	<u>N62933</u>	<u>001</u>	Oct 29, 1992
<u>AP</u>	+		<u>EQ 5GM BASE/VIAL</u>	<u>N63076</u>	<u>001</u>	Dec 21, 1990

VARDENAFIL HYDROCHLORIDE

TABLET; ORAL

LEVITRA

	BAYER PHARMS	2.5MG	N21400	003	Aug 19, 2003
		5MG	N21400	001	Aug 19, 2003
		10MG	N21400	002	Aug 19, 2003
+		20MG	N21400	004	Aug 19, 2003

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

<u>AP</u>		BAXTER HLTHCARE	<u>10MG/VIAL</u>	<u>N75218</u>	<u>001</u>	Aug 23, 1999
<u>AP</u>			<u>20MG/VIAL</u>	<u>N75218</u>	<u>002</u>	Aug 23, 1999
<u>AP</u>	+	BEDFORD	<u>10MG/VIAL</u>	<u>N75549</u>	<u>001</u>	Jun 13, 2000
<u>AP</u>			<u>20MG/VIAL</u>	<u>N75549</u>	<u>002</u>	Jun 13, 2000
<u>AP</u>		HOSPIRA	<u>10MG/VIAL</u>	<u>N75164</u>	<u>001</u>	Oct 21, 1999
<u>AP</u>			<u>20MG/VIAL</u>	<u>N75164</u>	<u>002</u>	Oct 21, 1999
	+		4MG/VIAL	N75558	001	Sep 11, 2001
<u>AP</u>		SICOR PHARMS	<u>10MG/VIAL</u>	<u>N74688</u>	<u>001</u>	Aug 25, 1999
<u>AP</u>			<u>20MG/VIAL</u>	<u>N74688</u>	<u>002</u>	Aug 25, 1999
<u>AP</u>		STERIS	<u>10MG/VIAL</u>	<u>N74334</u>	<u>001</u>	Aug 31, 1995
<u>AP</u>			<u>20MG/VIAL</u>	<u>N74334</u>	<u>002</u>	Aug 31, 1995

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

EFFEXOR XR

	WYETH PHARMS INC	EQ 37.5MG BASE	N20699	001	Oct 20, 1997
		EQ 75MG BASE	N20699	002	Oct 20, 1997
+		EQ 150MG BASE	N20699	004	Oct 20, 1997

TABLET; ORAL

EFFEXOR

	WYETH PHARMS INC	EQ 25MG BASE	N20151	002	Dec 28, 1993
		EQ 37.5MG BASE	N20151	006	Dec 28, 1993
+		EQ 50MG BASE	N20151	003	Dec 28, 1993
		EQ 75MG BASE	N20151	004	Dec 28, 1993
		EQ 100MG BASE	N20151	005	Dec 28, 1993

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>		MYLAN	<u>120MG</u>	<u>N75138</u>	<u>001</u>	Apr 20, 1999
<u>AB</u>			<u>180MG</u>	<u>N75138</u>	<u>002</u>	Apr 20, 1999
<u>AB</u>			<u>240MG</u>	<u>N75138</u>	<u>003</u>	Apr 20, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 354 (of 360)

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

VERELAN

<u>AB</u>	+	ELAN DRUG	<u>120MG</u>	<u>N19614</u>	<u>001</u>	May 29, 1990
<u>AB</u>	+		<u>180MG</u>	<u>N19614</u>	<u>003</u>	Jan 09, 1992
<u>AB</u>	+		<u>240MG</u>	<u>N19614</u>	<u>002</u>	May 29, 1990
	+		360MG	N19614	004	May 10, 1996
		VERELAN PM				
		ELAN DRUG	100MG	N20943	001	Nov 25, 1998
			200MG	N20943	002	Nov 25, 1998
	+		300MG	N20943	003	Nov 25, 1998

INJECTABLE; INJECTION

ISOPTIN

<u>AP</u>	+	FSC	<u>2.5MG/ML</u>	<u>N18485</u>	<u>001</u>	
		<u>VERAPAMIL HYDROCHLORIDE</u>				
<u>AP</u>		BEDFORD	<u>2.5MG/ML</u>	<u>N72888</u>	<u>001</u>	Jul 28, 1995
<u>AP</u>		HOSPIRA	<u>2.5MG/ML</u>	<u>N70577</u>	<u>001</u>	Feb 02, 1987
<u>AP</u>			<u>2.5MG/ML</u>	<u>N70737</u>	<u>001</u>	May 06, 1987
<u>AP</u>			<u>2.5MG/ML</u>	<u>N70738</u>	<u>001</u>	May 06, 1987
<u>AP</u>			<u>2.5MG/ML</u>	<u>N75136</u>	<u>001</u>	Oct 20, 1998
<u>AP</u>		INTL MEDICATION	<u>2.5MG/ML</u>	<u>N70451</u>	<u>001</u>	Dec 16, 1985
<u>AP</u>		LUITPOLD	<u>2.5MG/ML</u>	<u>N70225</u>	<u>001</u>	Nov 12, 1985
<u>AP</u>			<u>2.5MG/ML</u>	<u>N70617</u>	<u>001</u>	Nov 12, 1985

TABLET; ORAL

CALAN

<u>AB</u>		GD SEARLE LLC	<u>40MG</u>	<u>N18817</u>	<u>003</u>	Feb 23, 1988
<u>AB</u>			<u>80MG</u>	<u>N18817</u>	<u>001</u>	Sep 10, 1984
<u>AB</u>			<u>120MG</u>	<u>N18817</u>	<u>002</u>	Sep 10, 1984

ISOPTIN

<u>AB</u>		FSC	<u>40MG</u>	<u>N18593</u>	<u>003</u>	Nov 23, 1987
<u>AB</u>			<u>80MG</u>	<u>N18593</u>	<u>001</u>	Mar 08, 1982
<u>AB</u>	+		<u>120MG</u>	<u>N18593</u>	<u>002</u>	Mar 08, 1982

VERAPAMIL HYDROCHLORIDE

<u>AB</u>		MUTUAL PHARM	<u>80MG</u>	<u>N71488</u>	<u>001</u>	Jan 13, 1988
<u>AB</u>			<u>120MG</u>	<u>N71489</u>	<u>001</u>	Jan 13, 1988
<u>AB</u>		MYLAN	<u>80MG</u>	<u>N71482</u>	<u>001</u>	Feb 15, 1989
<u>AB</u>			<u>120MG</u>	<u>N71483</u>	<u>001</u>	Feb 15, 1989
<u>AB</u>		PLIVA	<u>80MG</u>	<u>N72124</u>	<u>001</u>	Jan 26, 1989
<u>AB</u>			<u>120MG</u>	<u>N72125</u>	<u>001</u>	Jan 26, 1989
<u>AB</u>		PUREPAC PHARM	<u>80MG</u>	<u>N71019</u>	<u>001</u>	Sep 24, 1986
<u>AB</u>			<u>120MG</u>	<u>N70468</u>	<u>001</u>	Sep 24, 1986
<u>AB</u>		SANDOZ	<u>40MG</u>	<u>N73168</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>			<u>80MG</u>	<u>N71423</u>	<u>001</u>	May 24, 1988
<u>AB</u>			<u>120MG</u>	<u>N71424</u>	<u>001</u>	May 25, 1988
<u>AB</u>		WATSON LABS	<u>40MG</u>	<u>N72923</u>	<u>001</u>	Jun 29, 1993
<u>AB</u>			<u>40MG</u>	<u>N72924</u>	<u>001</u>	Jun 29, 1993
<u>AB</u>			<u>80MG</u>	<u>N70855</u>	<u>001</u>	Sep 24, 1986
<u>AB</u>			<u>80MG</u>	<u>N70995</u>	<u>001</u>	Oct 01, 1986
<u>AB</u>			<u>80MG</u>	<u>N71366</u>	<u>001</u>	Oct 01, 1986
<u>AB</u>			<u>120MG</u>	<u>N70856</u>	<u>001</u>	Sep 24, 1986
<u>AB</u>			<u>120MG</u>	<u>N70994</u>	<u>001</u>	Oct 01, 1986
<u>AB</u>			<u>120MG</u>	<u>N71367</u>	<u>001</u>	Oct 01, 1986

TABLET, EXTENDED RELEASE; ORAL

COVERA-HS

BC	+	GD SEARLE LLC	180MG	N20552	001	Feb 26, 1996
BC	+		240MG	N20552	002	Feb 26, 1996
		<u>ISOPTIN SR</u>				
<u>AB</u>	+	PAR PHARM	<u>120MG</u>	<u>N19152</u>	<u>003</u>	Mar 06, 1991
<u>AB</u>	+		<u>180MG</u>	<u>N19152</u>	<u>002</u>	Dec 15, 1989
<u>AB</u>	+		<u>240MG</u>	<u>N19152</u>	<u>001</u>	Dec 16, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 355 (of 360)

VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

VERAPAMIL HYDROCHLORIDE

<u>AB</u>	IVAX PHARMS	<u>120MG</u>	<u>N73568</u>	<u>002</u>	Oct 10, 1997
<u>AB</u>		<u>180MG</u>	<u>N74330</u>	<u>001</u>	Jan 31, 1994
<u>AB</u>		<u>240MG</u>	<u>N73568</u>	<u>001</u>	Jul 31, 1992
<u>AB</u>	KALI LABS	<u>120MG</u>	<u>N75072</u>	<u>001</u>	May 25, 1999
<u>AB</u>		<u>240MG</u>	<u>N75072</u>	<u>003</u>	May 25, 1999
<u>AB</u>	MYLAN	<u>120MG</u>	<u>N74587</u>	<u>002</u>	Feb 21, 1997
<u>AB</u>		<u>180MG</u>	<u>N74587</u>	<u>003</u>	Sep 09, 1997
<u>AB</u>		<u>240MG</u>	<u>N74587</u>	<u>001</u>	Mar 23, 1996
<u>AB</u>	PLIVA	<u>240MG</u>	<u>N72922</u>	<u>001</u>	Mar 01, 1996

VERTEPORFIN

INJECTABLE; INJECTION

VISUDYNE

+	QLT	15MG/VIAL	N21119	001	Apr 12, 2000
---	-----	-----------	--------	-----	--------------

VINBLASTINE SULFATE

INJECTABLE; INJECTION

VINBLASTINE SULFATE

+	AM PHARM PARTNERS	1MG/ML	N89515	001	Apr 29, 1987
---	-------------------	--------	--------	-----	--------------

+	BEDFORD	10MG/VIAL	N89395	001	Apr 09, 1987
---	---------	-----------	--------	-----	--------------

VINCRISTINE SULFATE

INJECTABLE; INJECTION

VINCRISTINE SULFATE

<u>AP</u>	AM PHARM	<u>1MG/ML</u>	<u>N76401</u>	<u>001</u>	Oct 28, 2003
<u>AP</u>	+ AM PHARM PARTNERS	<u>1MG/ML</u>	<u>N76296</u>	<u>001</u>	Dec 20, 2002

VINCRISTINE SULFATE PFS

<u>AP</u>	MAYNE PHARMA USA	<u>1MG/ML</u>	<u>N71484</u>	<u>001</u>	Apr 19, 1988
<u>AP</u>	SICOR PHARMS	<u>1MG/ML</u>	<u>N75493</u>	<u>001</u>	Sep 01, 1999

VINORELBINE TARTRATE

INJECTABLE; INJECTION

NAVELBINE

<u>AP</u>	+ PIERRE FABRE	<u>EQ 10MG BASE/ML</u>	<u>N20388</u>	<u>001</u>	Dec 23, 1994
-----------	----------------	------------------------	---------------	------------	--------------

VINORELBINE TARTRATE

<u>AP</u>	AM PHARM	<u>EQ 10MG BASE/ML</u>	<u>N76849</u>	<u>001</u>	Apr 18, 2005
<u>AP</u>	BAXTER HLTHCARE	<u>EQ 10MG BASE/ML</u>	<u>N75992</u>	<u>001</u>	Jun 10, 2003
<u>AP</u>	BEDFORD	<u>EQ 10MG BASE/ML</u>	<u>N76461</u>	<u>001</u>	Dec 11, 2003
<u>AP</u>	MAYNE PHARMA USA	<u>EQ 10MG BASE/ML</u>	<u>N76827</u>	<u>001</u>	Jun 02, 2005
<u>AP</u>	SICOR PHARMS	<u>EQ 10MG BASE/ML</u>	<u>N76028</u>	<u>001</u>	Feb 03, 2003

VITAMIN A PALMITATE

CAPSULE; ORAL

VITAMIN A PALMITATE

<u>AA</u>	ARCUM	<u>EQ 50,000 UNITS BASE</u>	<u>N83311</u>	<u>001</u>	
<u>AA</u>		<u>EQ 50,000 UNITS BASE</u>	<u>N83321</u>	<u>001</u>	

INJECTABLE; INJECTION

AQUASOL A

+	MAYNE PHARMA USA	EQ 50,000 UNITS BASE/ML	N06823	001	
---	------------------	-------------------------	--------	-----	--

VORICONAZOLE

FOR SUSPENSION; ORAL

VFEND

+	PFIZER	200MG/5ML	N21630	001	Dec 19, 2003
---	--------	-----------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 356 (of 360)

VORICONAZOLE

INJECTABLE; IV (INFUSION)

VFEND

+	PFIZER	200MG/VIAL	N21267	001	May 24, 2002
---	--------	------------	--------	-----	--------------

TABLET; ORAL

VFEND

	PFIZER	50MG	N21266	001	May 24, 2002
--	--------	------	--------	-----	--------------

+		200MG	N21266	002	May 24, 2002
---	--	-------	--------	-----	--------------

WARFARIN SODIUM

INJECTABLE; INJECTION

COUMADIN

+	BRISTOL MYERS SQUIBB	5MG/VIAL	N09218	024	Feb 07, 1995
---	----------------------	----------	--------	-----	--------------

TABLET; ORAL

COUMADIN

<u>AB</u>	BRISTOL MYERS SQUIBB	<u>1MG</u>	<u>N09218</u>	<u>022</u>	Mar 01, 1990
-----------	----------------------	------------	---------------	------------	--------------

<u>AB</u>		<u>2MG</u>	<u>N09218</u>	<u>013</u>	
-----------	--	------------	---------------	------------	--

<u>AB</u>		<u>2.5MG</u>	<u>N09218</u>	<u>018</u>	
-----------	--	--------------	---------------	------------	--

<u>AB</u>		<u>3MG</u>	<u>N09218</u>	<u>025</u>	Nov 18, 1996
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>4MG</u>	<u>N09218</u>	<u>023</u>	Aug 24, 1993
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>5MG</u>	<u>N09218</u>	<u>007</u>	
-----------	--	------------	---------------	------------	--

<u>AB</u>		<u>6MG</u>	<u>N09218</u>	<u>026</u>	Nov 18, 1996
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>7.5MG</u>	<u>N09218</u>	<u>016</u>	
-----------	--	--------------	---------------	------------	--

<u>AB</u>	+	<u>10MG</u>	<u>N09218</u>	<u>005</u>	
-----------	---	-------------	---------------	------------	--

JANTOVEN

<u>AB</u>	USL PHARMA	<u>1MG</u>	<u>N40416</u>	<u>001</u>	Oct 02, 2003
-----------	------------	------------	---------------	------------	--------------

<u>AB</u>		<u>2MG</u>	<u>N40416</u>	<u>002</u>	Oct 02, 2003
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>2.5MG</u>	<u>N40416</u>	<u>003</u>	Oct 02, 2003
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>3MG</u>	<u>N40416</u>	<u>004</u>	Oct 02, 2003
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>4MG</u>	<u>N40416</u>	<u>005</u>	Oct 02, 2003
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>5MG</u>	<u>N40416</u>	<u>006</u>	Oct 02, 2003
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>6MG</u>	<u>N40416</u>	<u>007</u>	Oct 02, 2003
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>7.5MG</u>	<u>N40416</u>	<u>008</u>	Oct 02, 2003
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>10MG</u>	<u>N40416</u>	<u>009</u>	Oct 02, 2003
-----------	--	-------------	---------------	------------	--------------

WARFARIN SODIUM

<u>AB</u>	BARR	<u>1MG</u>	<u>N40145</u>	<u>001</u>	Mar 26, 1997
-----------	------	------------	---------------	------------	--------------

<u>AB</u>		<u>2MG</u>	<u>N40145</u>	<u>002</u>	Mar 26, 1997
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>2.5MG</u>	<u>N40145</u>	<u>003</u>	Mar 26, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>3MG</u>	<u>N40145</u>	<u>008</u>	Nov 05, 1998
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>4MG</u>	<u>N40145</u>	<u>004</u>	Mar 26, 1997
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>5MG</u>	<u>N40145</u>	<u>005</u>	Mar 26, 1997
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>6MG</u>	<u>N40145</u>	<u>009</u>	Nov 05, 1998
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>7.5MG</u>	<u>N40145</u>	<u>006</u>	Mar 26, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>10MG</u>	<u>N40145</u>	<u>007</u>	Mar 26, 1997
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	GENPHARM	<u>1MG</u>	<u>N40415</u>	<u>001</u>	Sep 27, 2004
-----------	----------	------------	---------------	------------	--------------

<u>AB</u>		<u>2MG</u>	<u>N40415</u>	<u>002</u>	Sep 27, 2004
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>2.5MG</u>	<u>N40415</u>	<u>003</u>	Sep 29, 2004
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>3MG</u>	<u>N40415</u>	<u>004</u>	Sep 27, 2004
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>4MG</u>	<u>N40415</u>	<u>005</u>	Sep 27, 2004
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>5MG</u>	<u>N40415</u>	<u>006</u>	Sep 27, 2004
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>6MG</u>	<u>N40415</u>	<u>007</u>	Sep 27, 2004
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>7.5MG</u>	<u>N40415</u>	<u>008</u>	Sep 27, 2004
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>10MG</u>	<u>N40415</u>	<u>009</u>	Sep 27, 2004
-----------	--	-------------	---------------	------------	--------------

<u>AB</u>	SANDOZ	<u>1MG</u>	<u>N40196</u>	<u>001</u>	Sep 30, 1997
-----------	--------	------------	---------------	------------	--------------

<u>AB</u>		<u>2MG</u>	<u>N40196</u>	<u>002</u>	Sep 30, 1997
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>2.5MG</u>	<u>N40196</u>	<u>003</u>	Sep 30, 1997
-----------	--	--------------	---------------	------------	--------------

<u>AB</u>		<u>3MG</u>	<u>N40196</u>	<u>008</u>	Jul 26, 2000
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>4MG</u>	<u>N40196</u>	<u>004</u>	Sep 30, 1997
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>5MG</u>	<u>N40196</u>	<u>005</u>	Sep 30, 1997
-----------	--	------------	---------------	------------	--------------

<u>AB</u>		<u>6MG</u>	<u>N40196</u>	<u>009</u>	Jul 26, 2000
-----------	--	------------	---------------	------------	--------------

PRESCRIPTION DRUG PRODUCT LIST

3 - 357 (of 360)

WARFARIN SODIUM

TABLET; ORAL

WARFARIN SODIUM

<u>AB</u>	SANDOZ	<u>7.5MG</u>	<u>N40196</u>	<u>006</u>	Sep 30, 1997
<u>AB</u>		<u>10MG</u>	<u>N40196</u>	<u>007</u>	Sep 30, 1997
<u>AB</u>	TARO	<u>1MG</u>	<u>N40301</u>	<u>002</u>	Jul 15, 1999
<u>AB</u>		<u>2MG</u>	<u>N40301</u>	<u>003</u>	Jul 15, 1999
<u>AB</u>		<u>2.5MG</u>	<u>N40301</u>	<u>004</u>	Jul 15, 1999
<u>AB</u>		<u>3MG</u>	<u>N40301</u>	<u>005</u>	Jul 15, 1999
<u>AB</u>		<u>4MG</u>	<u>N40301</u>	<u>006</u>	Jul 15, 1999
<u>AB</u>		<u>5MG</u>	<u>N40301</u>	<u>007</u>	Jul 15, 1999
<u>AB</u>		<u>6MG</u>	<u>N40301</u>	<u>008</u>	Jul 15, 1999
<u>AB</u>		<u>7.5MG</u>	<u>N40301</u>	<u>009</u>	Jul 15, 1999
<u>AB</u>		<u>10MG</u>	<u>N40301</u>	<u>001</u>	Jul 15, 1999

WATER FOR INJECTION, STERILE

LIQUID; N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

<u>AP</u>	+ HOSPIRA	<u>100%</u>	<u>N18802</u>	<u>001</u>	Oct 27, 1982
<u>STERILE WATER FOR INJECTION IN PLASTIC CONTAINER</u>					
<u>AP</u>	AM PHARM PARTNERS	<u>100%</u>	<u>N88400</u>	<u>001</u>	Jan 16, 1984
<u>AP</u>	+ B BRAUN	<u>100%</u>	<u>N19633</u>	<u>001</u>	Feb 29, 1988
<u>AP</u>	+ BAXTER HLTHCARE	<u>100%</u>	<u>N18632</u>	<u>001</u>	Jun 30, 1982
<u>AP</u>	+	<u>100%</u>	<u>N18632</u>	<u>002</u>	Apr 19, 1988
<u>AP</u>	+ HOSPIRA	<u>100%</u>	<u>N18233</u>	<u>001</u>	
<u>AP</u>	+	<u>100%</u>	<u>N18801</u>	<u>001</u>	Oct 27, 1982
<u>AP</u>	+	<u>100%</u>	<u>N19869</u>	<u>001</u>	Dec 26, 1989

WATER FOR IRRIGATION, STERILE

LIQUID; IRRIGATION

STERILE WATER

<u>AT</u>	BAXTER HLTHCARE	<u>100%</u>	<u>N17428</u>	<u>001</u>	
<u>STERILE WATER IN PLASTIC CONTAINER</u>					
<u>AT</u>	B BRAUN	<u>100%</u>	<u>N16734</u>	<u>001</u>	
<u>AT</u>	BAXTER HLTHCARE	<u>100%</u>	<u>N17866</u>	<u>001</u>	
<u>AT</u>	HOSPIRA	<u>100%</u>	<u>N17513</u>	<u>001</u>	
<u>AT</u>		<u>100%</u>	<u>N18313</u>	<u>001</u>	

XENON, XE-133

GAS; INHALATION

XENON XE 133

<u>AA</u>	BRISTOL MYERS SQUIBB	<u>10mCi/VIAL</u>	<u>N17284</u>	<u>001</u>	
<u>AA</u>		<u>20mCi/VIAL</u>	<u>N17284</u>	<u>002</u>	
<u>AA</u>	MALLINCKRODT	<u>10mCi/VIAL</u>	<u>N18327</u>	<u>001</u>	Mar 09, 1982
<u>AA</u>		<u>20mCi/VIAL</u>	<u>N18327</u>	<u>002</u>	Mar 09, 1982

ZAFIRLUKAST

TABLET; ORAL

ACCOLATE

	ASTRAZENECA	10MG	N20547	003	Sep 17, 1999
+		20MG	N20547	001	Sep 26, 1996

ZALCITABINE

TABLET; ORAL

HIVID

	ROCHE	0.375MG	N20199	001	Jun 19, 1992
+		0.75MG	N20199	002	Jun 19, 1992

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 358 (of 360)

ZALEPLONCAPSULE; ORAL
SONATA

JONES PHARMA

5MG

N20859 001

Aug 13, 1999

+

10MG

N20859 002

Aug 13, 1999

ZANAMIVIRPOWDER; INHALATION
RELENZA

+ GLAXOSMITHKLINE

5MG

N21036 001

Jul 26, 1999

ZICONOTIDEINJECTABLE; INTRATHECAL
PRIALT

+ ELAN PHARMS

100UGM/1ML (100MG/ML)

N21060 002

Dec 28, 2004

+

500UGM/20ML (25MG/ML)

N21060 001

Dec 28, 2004

+

500UGM/5ML (100MG/ML)

N21060 004

Dec 28, 2004

ZIDOVUDINECAPSULE; ORAL
RETROVIR

+ GLAXOSMITHKLINE

100MG

N19655 001

Mar 19, 1987

INJECTABLE; INJECTION
RETROVIR

+ GLAXOSMITHKLINE

10MG/ML

N19951 001

Feb 02, 1990

SYRUP; ORAL

RETROVIRAA + GLAXOSMITHKLINE50MG/5MLN19910 001

Sep 28, 1989

ZIDOVUDINEAA AUROBINDO50MG/5MLN77268 001

Sep 19, 2005

TABLET; ORAL

RETROVIRAB + GLAXOSMITHKLINE300MGN20518 002

Oct 04, 1996

ZIDOVUDINEAB AUROBINDO300MGN77267 001

Sep 19, 2005

AB RANBAXY300MGN77327 001

Sep 19, 2005

AB ROXANE300MGN76844 001

Sep 19, 2005

ZILEUTONTABLET; ORAL
ZYFLO

+ CRITICAL

600MG

N20471 003

Dec 09, 1996

ZINC ACETATECAPSULE; ORAL
GALZIN

TEVA

EQ 25MG ZINC

N20458 001

Jan 28, 1997

+

EQ 50MG ZINC

N20458 002

Jan 28, 1997

ZINC CHLORIDEINJECTABLE; INJECTION
ZINC CHLORIDE IN PLASTIC CONTAINER

+ HOSPIRA

EQ 1MG ZINC/ML

N18959 001

Jun 26, 1986

ZIPRASIDONE HYDROCHLORIDECAPSULE; ORAL
GEODON

+ PFIZER

EQ 20MG BASE

N20825 001

Feb 05, 2001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

PRESCRIPTION DRUG PRODUCT LIST

3 - 359 (of 360)

ZIPRASIDONE HYDROCHLORIDE

CAPSULE; ORAL

GEODON

PFIZER

EQ 40MG BASE

N20825 002

Feb 05, 2001

EQ 60MG BASE

N20825 003

Feb 05, 2001

EQ 80MG BASE

N20825 004

Feb 05, 2001

ZIPRASIDONE MESYLATE

INJECTABLE; INTRAMUSCULAR

GEODON

+ PFIZER

EQ 20MG BASE/ML

N20919 001

Jun 21, 2002

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)

ZOMETA

+ NOVARTIS

EQ 4MG BASE/5ML

N21223 002

Mar 07, 2003

+

EQ 4MG BASE/VIAL

N21223 001

Aug 20, 2001

ZOLMITRIPTAN

SPRAY; NASAL

ZOMIG

+ ASTRAZENECA

5MG/SPRAY

N21450 004

Sep 30, 2003

TABLET; ORAL

ZOMIG

IPR

2.5MG

N20768 001

Nov 25, 1997

+

5MG

N20768 002

Nov 25, 1997

TABLET, ORALLY DISINTEGRATING; ORAL

ZOMIG-ZMT

ASTRAZENECA

2.5MG

N21231 001

Feb 13, 2001

+

5MG

N21231 002

Sep 17, 2001

ZOLPIDEM TARTRATE

TABLET; ORAL

AMBIEN

SANOFI SYNTHELABO

5MG

N19908 001

Dec 16, 1992

+

10MG

N19908 002

Dec 16, 1992

TABLET, EXTENDED RELEASE; ORAL

AMBIEN CR

SANOFI SYNTHELABO

6.25MG

N21774 002

Sep 02, 2005

+

12.5MG

N21774 001

Sep 02, 2005

ZONISAMIDE

CAPSULE; ORAL

ZONEGRANAB DAINIPPON25MGN20789 003

Aug 22, 2003

AB50MGN20789 002

Aug 22, 2003

AB +100MGN20789 001

Mar 27, 2000

ZONISAMIDEAB ALPHAPHARM25MGN77647 001

Dec 22, 2005

AB50MGN77647 002

Dec 22, 2005

AB100MGN77647 003

Dec 22, 2005

AB APOTEX25MGN77642 001

Dec 22, 2005

AB50MGN77642 002

Dec 22, 2005

AB100MGN77642 003

Dec 22, 2005

AB BARR25MGN77639 001

Dec 22, 2005

AB50MGN77639 002

Dec 22, 2005

AB100MGN77639 003

Dec 22, 2005

AB DR REDDYS LABS LTD100MGN77645 001

Dec 22, 2005

AB MUTUAL PHARM25MGN77635 001

Dec 22, 2005

PRESCRIPTION DRUG PRODUCT LIST

3 - 360 (of 360)

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

<u>AB</u>	MUTUAL PHARM	<u>50MG</u>	<u>N77635</u>	<u>002</u>	Dec 22, 2005
<u>AB</u>		<u>100MG</u>	<u>N77635</u>	<u>003</u>	Dec 22, 2005
<u>AB</u>	MYLAN	<u>25MG</u>	<u>N77637</u>	<u>001</u>	Dec 22, 2005
<u>AB</u>		<u>50MG</u>	<u>N77637</u>	<u>002</u>	Dec 22, 2005
<u>AB</u>		<u>100MG</u>	<u>N77637</u>	<u>003</u>	Dec 22, 2005
<u>AB</u>	ROXANE	<u>25MG</u>	<u>N77648</u>	<u>001</u>	Dec 22, 2005
<u>AB</u>		<u>50MG</u>	<u>N77648</u>	<u>002</u>	Dec 22, 2005
<u>AB</u>		<u>100MG</u>	<u>N77648</u>	<u>003</u>	Dec 22, 2005
<u>AB</u>	SANDOZ	<u>25MG</u>	<u>N77644</u>	<u>001</u>	Dec 22, 2005
<u>AB</u>		<u>50MG</u>	<u>N77644</u>	<u>002</u>	Dec 22, 2005
<u>AB</u>		<u>100MG</u>	<u>N77644</u>	<u>003</u>	Dec 22, 2005
<u>AB</u>	TEVA PHARMS	<u>25MG</u>	<u>N77641</u>	<u>003</u>	Dec 22, 2005
<u>AB</u>		<u>50MG</u>	<u>N77641</u>	<u>002</u>	Dec 22, 2005
<u>AB</u>		<u>100MG</u>	<u>N77641</u>	<u>001</u>	Dec 22, 2005
<u>AB</u>	WOCKHARDT	<u>100MG</u>	<u>N77636</u>	<u>001</u>	Dec 22, 2005

OTC DRUG PRODUCT LIST

4 - 1 (of 15)

ACETAMINOPHEN

SUPPOSITORY; RECTAL

ACEPHEN

G AND W LABS	120MG	N18060	001	
	120MG	N72218	001	Mar 27, 1992
	325MG	N18060	003	Dec 18, 1986
	325MG	N72344	001	Mar 27, 1992
	650MG	N18060	002	
	650MG	N72237	001	Mar 27, 1992

ACETAMINOPHEN

ALPHARMA US PHARMS	120MG	N18337	003	Sep 12, 1983
	325MG	N18337	002	
+	650MG	N18337	001	
SUPPOSITORIA	120MG	N70607	001	Apr 06, 1987
	650MG	N70608	001	Dec 01, 1986

INFANTS' FEVERALL

ALPHARMA US PHARMS	80MG	N18337	004	Aug 26, 1992
--------------------	------	--------	-----	--------------

NEOPAP

POLYMEDICA	120MG	N16401	001	
------------	-------	--------	-----	--

TABLET, EXTENDED RELEASE; ORAL

ACETAMINOPHEN

COREPHARMA	650MG	N76200	001	Mar 19, 2002
PERRIGO	650MG	N75077	001	Feb 25, 2000

TYLENOL (CAPLET)

+	MCNEIL CONS SPECLT	650MG	N19872	001	Jun 08, 1994
---	--------------------	-------	--------	-----	--------------

TYLENOL (GELTAB)

+	MCNEIL CONS SPECLT	650MG	N19872	002	Jan 11, 2001
---	--------------------	-------	--------	-----	--------------

ACETAMINOPHEN; ASPIRIN; CAFFEINE

TABLET; ORAL

ACETAMINOPHEN, ASPIRIN AND CAFFEINE

PERRIGO	250MG;250MG;65MG	N75794	001	Nov 26, 2001
---------	------------------	--------	-----	--------------

EXCEDRIN (MIGRAINE)

+	NOVARTIS	250MG;250MG;65MG	N20802	001	Jan 14, 1998
---	----------	------------------	--------	-----	--------------

ACETAMINOPHEN; CLEMASTINE FUMARATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL

TAVIST ALLERGY/SINUS/HEADACHE

+	NOVARTIS	500MG;EQ 0.25MG BASE;30MG	N21082	001	Mar 01, 2001
---	----------	---------------------------	--------	-----	--------------

ACETAMINOPHEN; DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

DRIXORAL PLUS

+	SCHERING PLOUGH	500MG;3MG;60MG	N19453	001	May 22, 1987
---	-----------------	----------------	--------	-----	--------------

ALCOHOL; CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

AVAGARD

+	3M	61%;1%	N21074	001	Jun 07, 2001
---	----	--------	--------	-----	--------------

ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE; ORAL

FOAMCOAT

GUARDIAN DRUG	80MG;20MG	N71793	001	Sep 04, 1987
---------------	-----------	--------	-----	--------------

GAVISCON

AVENTIS PHARMS	80MG;20MG	N18685	001	Dec 09, 1983
----------------	-----------	--------	-----	--------------

+		160MG;40MG	N18685	002	Dec 09, 1983
---	--	------------	--------	-----	--------------

ANTAZOLINE PHOSPHATE; NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

VASOCON-A

+	NOVARTIS	0.5%;0.05%	N18746	002	Jul 11, 1994
---	----------	------------	--------	-----	--------------

OTC DRUG PRODUCT LIST

4 - 2 (of 15)

ASPIRIN

TABLET; ORAL		
BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN		
BAYER	500MG	N21317 001 Oct 18, 2001

AVOBENZONE; OCTINOXATE; OXYBENZONE

LOTION; TOPICAL		
SHADE UVAGUARD		
+ SCHERING PLOUGH	3%; 7.5%; 3%	N20045 001 Dec 07, 1992

BENTOQUATAM

LOTION; TOPICAL		
IVY BLOCK		
+ STAND HOMEOPATH	5%	N20532 001 Aug 26, 1996

BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL		
EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE		
+ ALZA	16MG; 240MG	N19672 001 Mar 29, 1996

BUTENAFINE HYDROCHLORIDE

CREAM; TOPICAL		
LOTRIMIN ULTRA		
+ SCHERING PLOUGH	1%	N21307 001 Dec 07, 2001

BUTOCONAZOLE NITRATE

CREAM; VAGINAL		
FEMSTAT 3		
+ BAYER	2%	N20421 001 Dec 21, 1995

CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE

TABLET, CHEWABLE; ORAL		
PEPCID COMPLETE		
+ MERCK	800MG; 10MG; 165MG	N20958 001 Oct 16, 2000

CHLORHEXIDINE GLUCONATE

AEROSOL, METERED; TOPICAL		
EXIDINE		
+ XTTRIUM	4%	N19127 001 Dec 24, 1984
CLOTH; TOPICAL		
CHLORHEXIDINE GLUCONATE		
+ SAGE PRODS	2%	N21669 001 Apr 25, 2005
SOLUTION; TOPICAL		
BRIAN CARE		
SOAPCO	4%	N71419 001 Dec 17, 1987
CHG SCRUB		
ECOLAB	4%	N19258 002 Jul 22, 1986
CIDA-STAT		
ECOLAB	2%	N19258 001 Jul 22, 1986
DYNA-HEX		
XTTRIUM	0.75%	N20111 001 Sep 11, 1997
EXIDINE		
+ XTTRIUM	2%	N19422 001 Dec 17, 1985
	4%	N19125 001 Dec 24, 1984
HIBICLENS		
+ REGENT	4%	N17768 001
HIBISTAT		
+ REGENT	0.5%	N18300 001
MICRODERM		
J AND J	4%	N72255 001 Apr 15, 1991
PREVACARE R		
J AND J	0.5%	N72292 001 Jan 28, 1992
STERI-STAT		
MATRIX MEDCL	4%	N70104 001 Jul 24, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 3 (of 15)

CHLORHEXIDINE GLUCONATE

SPONGE; TOPICAL

BIOSCRUB

GRIFFEN	4%	N19822	001	Mar 31, 1989
---------	----	--------	-----	--------------

CHLORHEXIDINE GLUCONATE

DESERET	4%	N72525	001	Oct 24, 1989
---------	----	--------	-----	--------------

KENDALL IL	4%	N19490	001	Mar 27, 1987
------------	----	--------	-----	--------------

HIBICLENS

+ REAGENT	4%	N18423	001	
-----------	----	--------	-----	--

MICRODERM

J AND J	4%	N72295	001	Feb 28, 1991
---------	----	--------	-----	--------------

PHARMASEAL SCRUB CARE

PHARMASEAL	4%	N19793	001	Dec 02, 1988
------------	----	--------	-----	--------------

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SPONGE; TOPICAL

CHLORAPREP ONE-STEP

+ MEDI FLEX INC	2%;70% (3ML)	N20832	001	Jul 14, 2000
-----------------	--------------	--------	-----	--------------

+	2%;70% (10.5ML)	N20832	004	Aug 20, 2003
---	-----------------	--------	-----	--------------

CHLORAPREP ONE-STEP FREPP

+ MEDI FLEX INC	2%;70% (1.5ML)	N20832	003	Apr 26, 2002
-----------------	----------------	--------	-----	--------------

CHLORAPREP WITH TINT

+ MEDI FLEX INC	2%;70% (26ML)	N20832	002	May 03, 2005
-----------------	---------------	--------	-----	--------------

SWAB; TOPICAL

CHLORAPREP ONE-STEP SEPP

+ MEDI FLEX INC	2%;70% (0.67ML)	N21555	001	Oct 07, 2002
-----------------	-----------------	--------	-----	--------------

CHLORAPREP SINGLE SWABSTICK

+ MEDI FLEX INC	2%;70% (1.75ML)	N21555	002	May 10, 2005
-----------------	-----------------	--------	-----	--------------

CHLORASCUB MAXI SWABSTICK

+ NICE PAK	3.15%;70% (5.1ML)	N21524	003	Jun 03, 2005
------------	-------------------	--------	-----	--------------

CHLORASCUB SWAB

+ NICE PAK	3.15%;70% (1ML)	N21524	001	Jun 03, 2005
------------	-----------------	--------	-----	--------------

CHLORASCUB SWABSTICK

+ NICE PAK	3.15%;70% (1.6ML)	N21524	002	Jun 03, 2005
------------	-------------------	--------	-----	--------------

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

+ SCHERING PLOUGH	12MG	N07638	002	
-------------------	------	--------	-----	--

CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S ADVIL ALLERGY SINUS

+ WYETH CONS	1MG/5ML;100MG/5ML;15MG/5ML	N21587	001	Feb 24, 2004
--------------	----------------------------	--------	-----	--------------

TABLET; ORAL

ADVIL ALLERGY SINUS

+ WYETH CONS	2MG;200MG;30MG	N21441	001	Dec 19, 2002
--------------	----------------	--------	-----	--------------

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

+ SCHERING PLOUGH	8MG;120MG	N18397	001	
-------------------	-----------	--------	-----	--

CIMETIDINE

TABLET; ORAL

CIMETIDINE

IVAX PHARMS	200MG	N75345	001	Jun 16, 1999
-------------	-------	--------	-----	--------------

LEINER	200MG	N74961	001	Jun 19, 1998
--------	-------	--------	-----	--------------

LEK PHARMS	200MG	N75122	002	Jun 19, 1998
------------	-------	--------	-----	--------------

PERRIGO	200MG	N75285	001	Oct 29, 1998
---------	-------	--------	-----	--------------

PHARM FORM	200MG	N74963	001	Jun 19, 1998
------------	-------	--------	-----	--------------

TORPHARM	100MG	N74948	001	Jun 19, 1998
----------	-------	--------	-----	--------------

	200MG	N74948	002	Jul 26, 2002
--	-------	--------	-----	--------------

WATSON LABS	200MG	N75425	001	Jul 29, 1999
-------------	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 4 (of 15)

CIMETIDINE

TABLET; ORAL				
TAGAMET HB				
+ GLAXOSMITHKLINE	200MG	N20238	002	Aug 21, 1996

CLEMASTINE FUMARATE

TABLET; ORAL				
CLEMASTINE FUMARATE				
PERRIGO	1.34MG	N74512	001	Nov 22, 1995
SANDOZ	1.34MG	N73458	001	Oct 31, 1993
TEVA	1.34MG	N73282	002	Dec 03, 1992
TAVIST-1				
+ NOVARTIS	1.34MG	N20925	001	Aug 21, 1992

CLOTRIMAZOLE

CREAM, TABLET; TOPICAL, VAGINAL				
GYNE-LOTTRIMIN 3 COMBINATION PACK				
+ SCHERING PLOUGH	1%, 200MG	N20526	002	Jul 29, 1996
GYNE-LOTTRIMIN COMBINATION PACK				
+ SCHERING PLOUGH	1%, 100MG	N20289	002	Apr 26, 1993
MYCELEX-7 COMBINATION PACK				
BAYER PHARMS	1%, 100MG	N20389	002	Jun 23, 1994
CREAM; VAGINAL				
CLOTRIMAZOLE				
ALPHARMA US PHARMS	1%	N74165	001	Jul 16, 1993
TARO	1%	N72641	001	Dec 04, 1995
GYNE-LOTTRIMIN				
+ SCHERING PLOUGH	1%	N18052	002	Nov 30, 1990
GYNE-LOTTRIMIN 3				
+ SCHERING PLOUGH	2%	N20574	001	Nov 24, 1998
MYCELEX-7				
BAYER PHARMS	1%	N18230	002	Dec 26, 1991
TRIVAGIZOLE 3				
TARO	2%	N21143	001	Apr 12, 2000
TABLET; VAGINAL				
GYNE-LOTTRIMIN				
+ SCHERING PLOUGH	100MG	N17717	002	Nov 30, 1990
GYNE-LOTTRIMIN 3				
+ SCHERING PLOUGH	200MG	N20525	001	Jul 29, 1996
GYNIX				
TEVA PHARMS	100MG	N73249	001	Feb 13, 1998
MYCELEX-7				
BAYER PHARMS	100MG	N18182	002	Dec 26, 1991

CROMOLYN SODIUM

SPRAY, METERED; NASAL				
CROMOLYN SODIUM				
ALPHARMA US PHARMS	5.2MG/INH	N74800	001	Jul 26, 2001
BAUSCH AND LOMB	5.2MG/SPRAY	N75702	001	Jul 03, 2001
PERRIGO	5.2MG/SPRAY	N75427	001	Dec 12, 2001
NASALCROM				
+ PHARMACIA UPJOHN	5.2MG/SPRAY	N20463	001	Jan 03, 1997

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL				
DISOPHROL				
SCHERING PLOUGH	6MG;120MG	N13483	004	Sep 13, 1982
DRIXORAL				
+ SCHERING PLOUGH	6MG;120MG	N13483	003	Sep 13, 1982

DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL				
MUCINEX DM				
ADAMS LABS INC	30MG;600MG	N21620	002	Apr 29, 2004

OTC DRUG PRODUCT LIST

4 - 5 (of 15)

DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

MUCINEX DM

+ ADAMS LABS INC 60MG;1.2GM

N21620 001 Apr 29, 2004

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

DELSYM

+ UCB EQ 30MG HBR/5ML

N18658 001 Oct 08, 1982

DIPHENHYDRAMINE CITRATE; IBUPROFEN

TABLET; ORAL

ADVIL PM

+ WYETH CONS 38MG;200MG

N21394 001 Dec 21, 2005

DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN POTASSIUM

CAPSULE; ORAL

ADVIL PM

+ WYETH CONS 25MG;200MG

N21393 001 Dec 21, 2005

DOCOSANOL

CREAM; TOPICAL

ABREVA

+ GLAXOSMITHKLINE 10%

N20941 001 Jul 25, 2000

DOXYLAMINE SUCCINATE

TABLET; ORAL

DOXYLAMINE SUCCINATE

COPLY PHARM 25MG

N88900 002 Feb 12, 1988

LNK 25MG

N40564 001 Aug 27, 2004

PERRIGO 25MG

N40167 001 Sep 18, 1996

UNISOM

+ PFIZER 25MG

N18066 001

EPINEPHRINE

AEROSOL, METERED; INHALATION

EPINEPHRINE

ARMSTRONG PHARMS 0.2MG/INH

N87907 001 May 23, 1984

PRIMATENE MIST

+ WYETH CONS 0.2MG/INH

N16126 001

EPINEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION

BRONITIN MIST

WYETH CONS 0.3MG/INH

N16126 002

FAMOTIDINE

TABLET, CHEWABLE; ORAL

FAMOTIDINE

PERRIGO 10MG

N75715 001 Aug 22, 2003

PEPCID AC

+ MERCK 10MG

N20801 001 Sep 24, 1998

TABLET; ORAL

FAMOTIDINE

DR REDDYS LABS LTD 10MG

N75758 001 Aug 17, 2001

GENPHARM 10MG

N75674 001 Dec 21, 2001

IVAX PHARMS 10MG

N75512 001 Jul 26, 2001

PERRIGO 10MG

N75400 001 Mar 18, 2005

SANDOZ 10MG

N76101 001 Oct 21, 2002

TEVA 10MG

N75312 001 May 31, 2001

TORPHARM 10MG

N75610 001 Mar 12, 2002

WATSON LABS 10MG

N75404 001 Nov 28, 2001

WOCKHARDT 10MG

N77146 001 Mar 07, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 6 (of 15)

FAMOTIDINE

TABLET; ORAL

PEPCID AC

MERCK

10MG

N20325 001

Apr 28, 1995

+

20MG

N20325 002

Sep 23, 2003

PEPCID AC (GELTAB)

MERCK

10MG

N20902 001

Aug 05, 1999

GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

MUCINEX

ADAMS LABS INC

600MG

N21282 001

Jul 12, 2002

+

1.2GM

N21282 002

Dec 18, 2002

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

MUCINEX D

ADAMS LABS INC

600MG;60MG

N21585 001

Jun 22, 2004

+

1.2GM;120MG

N21585 002

Jun 22, 2004

IBUPROFEN

CAPSULE; ORAL

IBUPROFEN

+ BANNER PHARMACAPS

200MG

N21472 001

Oct 18, 2002

+ PHARM FORM

200MG

N74782 001

Jul 06, 1998

SUSPENSION/DROPS; ORAL

CHILDREN'S MOTRIN

+ MCNEIL

40MG/ML

N20603 001

Jun 10, 1996

IBUPROFEN

PERRIGO

40MG/ML

N75217 001

Dec 16, 1998

PEDIATRIC ADVIL

+ WYETH CONS

100MG/2.5ML

N20812 001

Jan 30, 1998

SUSPENSION; ORAL

CHILDREN'S ADVIL

WYETH CONS

100MG/5ML

N20589 001

Jun 27, 1996

CHILDREN'S ADVIL-FLAVORED

WYETH CONS

100MG/5ML

N20589 002

Nov 07, 1997

CHILDREN'S ELIXSURE

TARO

100MG/5ML

N21604 001

Jan 07, 2004

CHILDREN'S IBUPROFEN

PERRIGO

100MG/5ML

N74937 001

Dec 22, 1998

CHILDREN'S MOTRIN

+ MCNEIL

100MG/5ML

N20516 001

Jun 16, 1995

IBUPROFEN

ALPHARMA US PHARMS

100MG/5ML

N74916 001

Apr 30, 1999

TABLET, CHEWABLE; ORAL

CHILDREN'S ADVIL

WYETH CONS

50MG

N20944 001

Dec 18, 1998

CHILDREN'S MOTRIN

MCNEIL

50MG

N20601 001

Nov 15, 1996

IBUPROFEN

PERRIGO

50MG

N76359 001

Jan 16, 2004

100MG

N76359 002

Jan 16, 2004

JUNIOR STRENGTH ADVIL

WYETH CONS

100MG

N20944 002

Dec 18, 1998

JUNIOR STRENGTH MOTRIN

+ MCNEIL

100MG

N20601 003

Nov 15, 1996

TABLET; ORAL

ADVIL

WYETH CONS

200MG

N18989 001

May 18, 1984

CAP-PROFEN

PERRIGO

200MG

N72097 001

Dec 08, 1987

IBUPROFEN

BASF

200MG

N75661 001

Dec 12, 2001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 7 (of 15)

IBUPROFEN

TABLET; ORAL

IBUPROFEN

DR REDDYS LABS INC	100MG	N76117	001	Nov 20, 2001
INTERPHARM	200MG	N71333	001	Feb 17, 1987
	200MG	N72199	001	May 23, 1988
IVAX PHARMS	200MG	N71144	001	Jan 20, 1987
LEINER	200MG	N74931	001	Jul 20, 1998
LNK	100MG	N76741	001	Jun 17, 2004
	200MG	N75010	001	Mar 01, 1999
	200MG	N75139	001	Mar 01, 1999
MCNEIL	200MG	N73019	001	Mar 30, 1994
MUTUAL PHARM	200MG	N71229	001	Apr 01, 1987
	200MG	N72249	001	Jan 10, 1989
MYLAN	200MG	N71870	001	May 05, 1988
NEIL	200MG	N76460	001	Nov 26, 2003
OHM	200MG	N71163	001	Jul 15, 1986
PAR PHARM	200MG	N70481	001	Sep 24, 1986
	200MG	N70985	001	Oct 02, 1987
	200MG	N71575	001	May 08, 1987
PERRIGO	200MG	N72096	001	Dec 08, 1987
	200MG	N72098	001	Dec 08, 1987
	200MG	N75995	001	Mar 14, 2002
PERRIGO R AND D	200MG	N77349	001	Jun 21, 2005
PVT FORM	200MG	N71732	001	Sep 10, 1987
	200MG	N71735	001	Sep 10, 1987
	200MG	N72299	001	Jul 01, 1988
	200MG	N73691	001	Feb 25, 1994
SANDOZ	200MG	N70733	001	Sep 19, 1986
	200MG	N71807	001	Feb 25, 1988
	200MG	N74525	001	Dec 15, 1995
	200MG	N74533	001	Dec 15, 1995
VINTAGE PHARMS	200MG	N71639	001	Feb 02, 1988
WATSON LABS	200MG	N70435	001	Mar 05, 1986
IBUPROHM				
OHM LABS	200MG	N71214	001	Dec 01, 1986
IBU-TAB 200				
ALRA	200MG	N71057	001	Aug 11, 1988
JUNIOR STRENGTH ADVIL				
WYETH CONS	100MG	N20267	002	Dec 13, 1996
JUNIOR STRENGTH IBUPROFEN				
PERRIGO	100MG	N75367	001	Apr 22, 1999
JUNIOR STRENGTH MOTRIN				
MCNEIL	100MG	N20602	001	Jun 10, 1996
MEDIPREN				
MCNEIL	200MG	N70475	001	Feb 06, 1986
	200MG	N71215	001	Jun 26, 1986
MOTRIN IB				
+ MCNEIL	200MG	N19012	003	Dec 17, 1990
MOTRIN MIGRAINE PAIN				
+ MCNEIL	200MG	N19012	004	Feb 25, 2000
PROFEN				
PVT FORM	200MG	N71265	001	Oct 15, 1986
TAB-PROFEN				
PERRIGO	200MG	N72095	001	Dec 08, 1987

IBUPROFEN POTASSIUM

CAPSULE; ORAL

ADVIL LIQUI-GELS

+ WYETH CONS	200MG	N20402	001	Apr 20, 1995
ADVIL MIGRAINE LIQUI-GELS				
+ WYETH CONS	200MG	N20402	002	Mar 16, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 8 (of 15)

IBUPROFEN POTASSIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL		
ADVIL COLD AND SINUS		
+ WYETH CONS	200MG;30MG	N21374 001 May 30, 2002

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL		
CHILDREN'S ADVIL COLD		
WYETH CONS	100MG/5ML;15MG/5ML	N21373 001 Apr 18, 2002
CHILDREN'S MOTRIN COLD		
+ MCNEIL CONS SPECLT	100MG/5ML;15MG/5ML	N21128 001 Aug 01, 2000
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE		
PERRIGO	100MG/5ML;15MG/5ML	N76478 001 Nov 05, 2003
TABLET; ORAL		
ADVIL COLD AND SINUS		
+ WYETH CONS	200MG;30MG	N19771 001 Sep 19, 1989
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE		
PHARM FORM	200MG;30MG	N75588 001 Apr 08, 2002
IBUPROHM COLD AND SINUS		
OHM LABS	200MG;30MG	N74567 001 Apr 17, 2001
SINE-AID IB		
MCNEIL CONS SPECLT	200MG;30MG	N19899 001 Dec 31, 1992

INSULIN PURIFIED PORK

INJECTABLE; INJECTION		
REGULAR ILETIN II (PORK)		
+ LILLY	100 UNITS/ML	N18344 001

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION		
HUMULIN R		
+ LILLY	100 UNITS/ML	N18780 001 Oct 28, 1982
HUMULIN R PEN		
+ LILLY	100 UNITS/ML	N18780 005 Aug 06, 1998
NOVOLIN R		
+ NOVO NORDISK INC	100 UNITS/ML	N19938 001 Jun 25, 1991
VELOSULIN BR		
+ NOVO NORDISK INC	100 UNITS/ML	N21028 001 Jul 19, 1999

INSULIN RECOMBINANT HUMAN; INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION		
HUMULIN 50/50		
+ LILLY	50 UNITS/ML;50 UNITS/ML	N20100 001 Apr 29, 1992
HUMULIN 70/30		
+ LILLY	30 UNITS/ML;70 UNITS/ML	N19717 001 Apr 25, 1989
HUMULIN 70/30 PEN		
+ LILLY	30 UNITS/ML;70 UNITS/ML	N19717 002 Aug 06, 1998
NOVOLIN 70/30		
+ NOVO NORDISK INC	30 UNITS/ML;70 UNITS/ML	N19991 001 Jun 25, 1991

INSULIN SUSP ISOPHANE BEEF/PORK

INJECTABLE; INJECTION		
NPH ILETIN I (BEEF-PORK)		
+ LILLY	100 UNITS/ML	N17936 002

INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION		
NPH ILETIN II (PORK)		
+ LILLY	100 UNITS/ML	N18345 001

INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION		
HUMULIN N		
+ LILLY	100 UNITS/ML	N18781 001 Oct 28, 1982

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 9 (of 15)

INSULIN SUSP ISOPHANE RECOMBINANT HUMAN

INJECTABLE; INJECTION				
NOVOLIN N				
NOVO NORDISK INC	100 UNITS/ML	N19959	001	Jul 01, 1991

INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN

INJECTABLE; INJECTION				
HUMULIN U				
+ LILLY	100 UNITS/ML	N19571	002	Jun 10, 1987

INSULIN ZINC SUSP PURIFIED PORK

INJECTABLE; INJECTION				
LENTE ILETIN II (PORK)				
+ LILLY	100 UNITS/ML	N18347	001	

INSULIN ZINC SUSP RECOMBINANT HUMAN

INJECTABLE; INJECTION				
HUMULIN L				
+ LILLY	100 UNITS/ML	N19377	002	Sep 30, 1985
NOVOLIN L				
NOVO NORDISK INC	100 UNITS/ML	N19965	001	Jun 25, 1991

KETOCONAZOLE

SHAMPOO; TOPICAL				
NIZORAL A-D				
+ MCNEIL CONS SPECLT	1%	N20310	001	Oct 10, 1997

KETOPROFEN

TABLET; ORAL				
ACTRON				
BAYER	12.5MG	N20499	001	Oct 06, 1995
ORUDIS KT				
+ WYETH CONS	12.5MG	N20429	001	Oct 06, 1995

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL				
LOPERAMIDE HYDROCHLORIDE				
BANNER PHARMACAPS	1MG	N21855	001	Aug 04, 2005
+	2MG	N21855	002	Aug 04, 2005
SOLUTION; ORAL				
IMODIUM A-D				
+ MCNEIL	1MG/5ML	N19487	001	Mar 01, 1988
+	1MG/7.5ML	N19487	002	Jul 08, 2004
LOPERAMIDE HYDROCHLORIDE				
DURAMED PHARMS BARR	1MG/5ML	N74991	001	Dec 29, 1997
HI TECH PHARMA	1MG/5ML	N74352	001	Nov 17, 1995
MORTON GROVE	1MG/5ML	N74730	001	Aug 28, 1997
PERRIGO	1MG/5ML	N73243	001	Jan 21, 1992
ROXANE	1MG/5ML	N73079	001	Apr 30, 1992
TEVA	1MG/5ML	N73478	001	Jun 23, 1995
TABLET, CHEWABLE; ORAL				
IMODIUM A-D				
+ MCNEIL	2MG	N20448	001	Jul 24, 1997
TABLET; ORAL				
IMODIUM A-D				
+ MCNEIL	2MG	N19860	001	Nov 22, 1989
LOPERAMIDE HYDROCHLORIDE				
LEINER	2MG	N73254	001	Jul 30, 1993
LNK	2MG	N76497	001	Jun 10, 2003
OHM LABS	2MG	N74091	001	Dec 10, 1992
PERRIGO	2MG	N75232	001	Jan 06, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 10 (of 15)

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET, CHEWABLE; ORAL				
IMODIUM ADVANCED				
+ MCNEIL	2MG;125MG	N20606	001	Jun 26, 1996
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE				
PERRIGO	2MG;125MG	N76029	001	Aug 30, 2002
TABLET; ORAL				
IMODIUM ADVANCED				
+ MCNEIL CONS SPECLT	2MG;125MG	N21140	001	Nov 30, 2000

LORATADINE

SUSPENSION; ORAL				
LORATADINE				
+ TARO	1MG/ML	N21734	001	Oct 04, 2005
SYRUP; ORAL				
CLARITIN				
+ SCHERING PLOUGH	1MG/ML	N20641	002	Nov 27, 2002
LORATADINE				
APOTEX	1MG/ML	N75565	001	Oct 05, 2004
MORTON GROVE	1MG/ML	N75815	001	Aug 20, 2004
PERRIGO	1MG/ML	N75728	001	Aug 20, 2004
RANBAXY	1MG/ML	N76529	001	Aug 20, 2004
TARO	1MG/ML	N76805	001	Aug 20, 2004
TEVA	1MG/ML	N75505	001	Nov 07, 2003
TABLET, ORALLY DISINTEGRATING; ORAL				
ALAVERT				
WYETH CONS	10MG	N21375	001	Dec 19, 2002
CLARITIN HIVES RELIEF REDITAB				
+ SCHERING PLOUGH	10MG	N20704	003	Nov 19, 2003
CLARITIN REDITABS				
+ SCHERING PLOUGH	10MG	N20704	002	Nov 27, 2002
LORATADINE				
ANDRX PHARMS	10MG	N75990	001	Nov 03, 2003
IMPAX LABS	10MG	N76011	001	Sep 29, 2003
WYETH CONS	10MG	N75822	001	Feb 10, 2003
TABLET; ORAL				
CLARITIN				
+ SCHERING PLOUGH	10MG	N19658	002	Nov 27, 2002
CLARITIN HIVES RELIEF				
+ SCHERING PLOUGH	10MG	N19658	003	Nov 19, 2003
LORATADINE				
GENPHARM	10MG	N76154	001	Aug 20, 2003
PERRIGO	10MG	N76301	001	Jun 25, 2004
RANBAXY	10MG	N76134	001	Aug 18, 2003
SANDOZ	10MG	N75209	001	Jan 21, 2003

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL				
CLARITIN-D 24 HOUR				
+ SCHERING PLOUGH	10MG;240MG	N20470	002	Nov 27, 2002
LOPATADINE AND PSEUDOEPHEDRINE SULFATE				
ANDRX PHARMS	5MG;120MG	N76208	001	Jan 28, 2004
	10MG;240MG	N75706	001	Feb 21, 2003
+ IMPAX LABS	5MG;120MG	N76050	001	Jan 30, 2003
	10MG;240MG	N75989	001	Mar 04, 2004
RANBAXY	10MG;240MG	N76557	001	Sep 22, 2004

MICONAZOLE NITRATE

CREAM, SUPPOSITORY; TOPICAL, VAGINAL				
MICONAZOLE 7 COMBINATION PACK				
G AND W LABS	2%,100MG	N76585	001	Mar 26, 2004
MICONAZOLE NITRATE COMBINATION PACK				
PERRIGO	2%,200MG	N75329	001	Apr 20, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 11 (of 15)

MICONAZOLE NITRATE

CREAM, SUPPOSITORY; TOPICAL, VAGINAL

MONISTAT 1 COMBINATION PACK

+ PERSONAL PRODS 1.2GM, 2% N21308 001 Jun 29, 2001

MONISTAT 7 COMBINATION PACK

+ ADV CARE 2%, 100MG N20288 002 Apr 26, 1993

MONISTAT-3 COMBINATION PACK

+ ADV CARE 2%, 200MG N20670 002 Apr 16, 1996

M-ZOLE 3 COMBINATION PACK

ALPHARMA US PHARMS 2%, 200MG N74926 001 Apr 16, 1999

M-ZOLE 7 DUAL PACK

ALPHARMA US PHARMS 2%, 100MG N74586 001 Jul 17, 1997

CREAM; TOPICAL, VAGINAL

MICONAZOLE 3 COMBINATION PACK

PERRIGO 2%, 4% N76357 001 Mar 30, 2004

MONISTAT 3 COMBINATION PACK

PERSONAL PRODS 2%, 4% N21261 003 Jun 17, 2003

MONISTAT 3 COMBINATION PACK (PREFILLED)

+ PERSONAL PRODS 2%, 4% N21261 001 Feb 02, 2001

CREAM; VAGINAL

MICONAZOLE 3

TARO 4% N76773 001 Mar 02, 2005

MICONAZOLE 7

ALPHARMA US PHARMS 2% N74164 001 Mar 29, 1996

MICONAZOLE NITRATE

G AND W LABS 2% N74366 001 Feb 22, 1996

PERRIGO 2% N74760 001 May 15, 1997

TARO 2% N74444 001 Jan 13, 1997

TEVA 2% N74136 001 Jan 04, 1995

TEVA PHARMS 2% N74030 001 Oct 30, 1992

MONISTAT 3

+ ADV CARE PRODS 4% N20827 001 Mar 30, 1998

MONISTAT 7

+ ADV CARE 2% N17450 002 Feb 15, 1991

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

ALPHARMA US PHARMS 100MG N73507 001 Nov 19, 1993

G AND W LABS 100MG N74414 001 Apr 30, 1997

+ PERRIGO 100MG N74395 001 Mar 20, 1997

MONISTAT 7

+ ADV CARE PRODS 100MG N18520 002 Feb 15, 1991

MINOXIDIL

SOLUTION; TOPICAL

MINOXIDIL (FOR MEN)

ALPHARMA US PHARMS 2% N74588 001 Apr 05, 1996

BAUSCH AND LOMB 2% N74643 001 Apr 09, 1996

COPLEY PHARM 2% N74500 001 May 23, 1996

HI TECH PHARMA 2% N74731 001 Dec 24, 1996

MORTON GROVE 2% N74767 001 Feb 28, 1997

NOVEX 2% N74924 001 Apr 29, 1998

PERRIGO 2% N75357 001 Jul 30, 1999

SIGHT PHARMS 2% N74743 002 Oct 18, 1996

TEVA 2% N74589 001 Apr 05, 1996

MINOXIDIL (FOR WOMEN)

NOVEX 2% N74924 002 Apr 29, 1998

PERRIGO 2% N75357 002 Jul 30, 1999

SIGHT PHARMS 2% N74743 001 Oct 18, 1996

MINOXIDIL EXTRA STRENGTH (FOR MEN)

ALPHARMA US PHARMS 5% N75518 001 Nov 17, 2000

CLAY PARK 5% N75737 001 Mar 15, 2002

MORTON GROVE 5% N75438 001 Feb 27, 2003

NOVEX 5% N75839 001 Oct 01, 2001

PERRIGO 5% N75598 001 Jun 13, 2001

TEVA 5% N75619 001 Nov 17, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 12 (of 15)

MINOXIDIL

SOLUTION; TOPICAL

ROGAINE (FOR MEN)

+ PHARMACIA AND UPJOHN 2% N19501 002 Feb 09, 1996

ROGAINE (FOR WOMEN)

+ PHARMACIA AND UPJOHN 2% N19501 003 Feb 09, 1996

ROGAINE EXTRA STRENGTH (FOR MEN)

+ PHARMACIA AND UPJOHN 5% N20834 001 Nov 14, 1997

THEROXIDIL

HARMONY LABS 5% N76239 001 Aug 24, 2004

NAPHAZOLINE HYDROCHLORIDE; PHENIRAMINE MALEATE

SOLUTION/DROPS; OPHTHALMIC

NAPHCN-A

+ ALCON 0.025%;0.3% N20226 001 Jun 08, 1994

OPCON-A

+ BAUSCH AND LOMB 0.02675%;0.315% N20065 001 Jun 08, 1994

VISINE-A

PFIZER 0.025%;0.3% N20485 001 Jan 31, 1996

NAPROXEN SODIUM

TABLET; ORAL

ALEVE

+ BAYER EQ 200MG BASE N20204 002 Jan 11, 1994

NAPROXEN SODIUM

DR REDDYS LABS INC EQ 200MG BASE N75168 001 Jul 28, 1998

LEINER EQ 200MG BASE N74635 001 Jan 13, 1997

PERRIGO EQ 200MG BASE N74661 001 Jan 13, 1997

PVT FORM EQ 200MG BASE N74789 001 Feb 27, 1997

SANDOZ EQ 200MG BASE N74646 001 Jan 13, 1997

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ALEVE COLD AND SINUS

+ BAYER EQ 200MG BASE;120MG N21076 001 Nov 29, 1999

NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

PERRIGO EQ 200MG BASE;120MG N76518 001 Mar 17, 2004

NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

HABITROL

+ NOVARTIS 7MG/24HR N20076 004 Nov 12, 1999

+ 14MG/24HR N20076 005 Nov 12, 1999

+ 21MG/24HR N20076 006 Nov 12, 1999

NICODERM CQ

+ AVENTIS PHARMS 7MG/24HR N20165 006 Aug 02, 1996

+ 14MG/24HR N20165 005 Aug 02, 1996

+ 21MG/24HR N20165 004 Aug 02, 1996

PROSTEP

+ AVEVA 11MG/24HR N19983 003 Dec 23, 1998

+ 22MG/24HR N19983 004 Dec 23, 1998

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICORETTE

+ GLAXOSMITHKLINE EQ 2MG BASE N18612 002 Feb 09, 1996

+ EQ 4MG BASE N20066 002 Feb 09, 1996

NICORETTE (MINT)

+ GLAXOSMITHKLINE EQ 2MG BASE N18612 003 Dec 23, 1998

+ EQ 4MG BASE N20066 003 Dec 23, 1998

NICORETTE (ORANGE)

+ GLAXOSMITHKLINE EQ 2MG BASE N18612 004 Sep 25, 2000

+ EQ 4MG BASE N20066 004 Sep 25, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 13 (of 15)

NICOTINE POLACRILEXGUM, CHEWING; BUCCAL
NICOTINE POLACRILEX

PERRIGO	EQ 2MG BASE	N76775	001	Sep 16, 2004
	EQ 4MG BASE	N76789	001	Sep 16, 2004
WATSON LABS	EQ 2MG BASE	N74507	001	Mar 15, 1999
	EQ 4MG BASE	N74707	001	Mar 19, 1999

NICOTINE POLACRILEX (MINT)

PERRIGO	EQ 2MG BASE	N76777	001	Sep 16, 2004
	EQ 4MG BASE	N76779	001	Sep 16, 2004
WATSON LABS	EQ 2MG BASE	N76569	001	Jul 29, 2004
	EQ 4MG BASE	N76568	002	Jul 29, 2004

NICOTINE POLACRILEX (ORANGE)

PERRIGO	EQ 2MG BASE	N76776	001	Sep 16, 2004
	EQ 4MG BASE	N76778	001	Sep 16, 2004

TROCHE/LOZENGE; ORAL

COMMIT

GLAXOSMITHKLINE CONS	EQ 2MG BASE	N21330	001	Oct 31, 2002
+	EQ 4MG BASE	N21330	002	Oct 31, 2002

NIZATIDINE

TABLET; ORAL

AXID AR

+	WYETH CONS	75MG	N20555	001	May 09, 1996
---	------------	------	--------	-----	--------------

OMEPRAZOLE MAGNESIUM

TABLET, DELAYED RELEASE; ORAL

PRILOSEC OTC

+	ASTRAZENECA	EQ 20MG BASE	N21229	001	Jun 20, 2003
---	-------------	--------------	--------	-----	--------------

OXYMETAZOLINE HYDROCHLORIDESOLUTION/DROPS; OPHTHALMIC
OCUCLEAR

SCHERING PLOUGH	0.025%	N18471	001	May 30, 1986
-----------------	--------	--------	-----	--------------

VISINE L.R.

+	PFIZER	0.025%	N19407	001	Mar 31, 1989
---	--------	--------	--------	-----	--------------

PERMETHRIN

LOTION; TOPICAL

NIX

+	INSIGHT PHARMS	1%	N19918	001	May 02, 1990
---	----------------	----	--------	-----	--------------

PERMETHRIN

ALPHARMA US PHARMS	1%	N75014	001	Mar 28, 2000
--------------------	----	--------	-----	--------------

CLAY PARK	1%	N76090	001	Dec 20, 2001
-----------	----	--------	-----	--------------

PIPERONYL BUTOXIDE; PYRETHRINS

AEROSOL; TOPICAL

RID MOUSSE

+	PFIZER	4%;EQ 0.33% BASE	N21043	001	Mar 07, 2000
---	--------	------------------	--------	-----	--------------

POTASSIUM IODIDE

SOLUTION; ORAL

THYROSHIELD

FLEMING	65MG/ML	N77218	001	Jan 12, 2005
---------	---------	--------	-----	--------------

TABLET; ORAL

IOSAT

+	ANBEX	130MG	N18664	001	Oct 14, 1982
---	-------	-------	--------	-----	--------------

THYROSAFE

+	R R REGISTRATIONS	65MG	N76350	001	Sep 10, 2002
---	-------------------	------	--------	-----	--------------

POVIDONE- IODINE

SOLUTION; TOPICAL

E-Z PREP

+	BECTON DICKINSON	10%	N19382	001	Jul 25, 1989
---	------------------	-----	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

OTC DRUG PRODUCT LIST

4 - 14 (of 15)

POVIDONE- IODINE

SOLUTION; TOPICAL

POVIDONE IODINE

+ ALLEGIANCE HLTHCARE	1%	N19522	001	Mar 31, 1989
-----------------------	----	--------	-----	--------------

SPONGE; TOPICAL

E-Z PREP

+ BECTON DICKINSON	5%	N19382	002	Jul 25, 1989
--------------------	----	--------	-----	--------------

E-Z PREP 220

+ BECTON DICKINSON	5%	N19382	003	Jul 25, 1989
--------------------	----	--------	-----	--------------

E-Z SCRUB 201

+ BECTON DICKINSON	20%	N19240	001	Nov 29, 1985
--------------------	-----	--------	-----	--------------

E-Z SCRUB 241

+ BECTON DICKINSON	10%	N19476	001	Jan 07, 1987
--------------------	-----	--------	-----	--------------

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE

+ ALZA	240MG	N20021	002	Dec 15, 1992
--------	-------	--------	-----	--------------

PSEUDOEPHEDRINE HYDROCHLORIDE

PERRIGO 120MG

N75153	001	Feb 26, 1999
--------	-----	--------------

RANBAXY 120MG

N77442	001	Sep 28, 2005
--------	-----	--------------

SUDAFED 12 HOUR

+ WARNER LAMBERT	120MG	N73585	001	Oct 31, 1991
------------------	-------	--------	-----	--------------

PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

AFRINOL

+ SCHERING PLOUGH	120MG	N18191	001	
-------------------	-------	--------	-----	--

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE

DR REDDYS LABS LTD EQ 75MG BASE

N75294	001	Mar 28, 2000
--------	-----	--------------

GENPHARM EQ 75MG BASE

N75497	001	Jan 14, 2000
--------	-----	--------------

IVAX PHARMS EQ 75MG BASE

N75296	001	Jan 14, 2000
--------	-----	--------------

LEINER EQ 75MG BASE

N75094	001	Jun 21, 1999
--------	-----	--------------

PERRIGO EQ 75MG BASE

N76195	001	Aug 30, 2002
--------	-----	--------------

RANBAXY EQ 75MG BASE

N75132	001	Jan 14, 2000
--------	-----	--------------

SANDOZ EQ 75MG BASE

N75519	001	Sep 26, 2002
--------	-----	--------------

TORPHARM EQ 75MG BASE

N75167	001	May 04, 2000
--------	-----	--------------

WATSON LABS EQ 75MG BASE

N75212	001	Jan 14, 2000
--------	-----	--------------

ZANTAC 150

+ PFIZER CONS HLTHCARE	EQ 150MG BASE	N21698	001	Aug 31, 2004
------------------------	---------------	--------	-----	--------------

ZANTAC 75

WARNER LAMBERT	EQ 75MG BASE	N20520	001	Dec 19, 1995
----------------	--------------	--------	-----	--------------

SODIUM CHLORIDE

AEROSOL, METERED; INHALATION

BRONCHO SALINE

+ BLAIREX	0.9%	N19912	001	Sep 03, 1992
-----------	------	--------	-----	--------------

SODIUM FLUORIDE; TRICLOSAN

PASTE; DENTAL

COLGATE TOTAL

+ COLGATE PALMOLIVE	0.24%;0.3%	N20231	001	Jul 11, 1997
---------------------	------------	--------	-----	--------------

SODIUM MONOFLUOROPHOSPHATE

GEL; DENTAL

EXTRA-STRENGTH AIM

+ CHESEBROUGH PONDS	1.2%	N19518	002	Aug 06, 1986
---------------------	------	--------	-----	--------------

PASTE; DENTAL

EXTRA-STRENGTH AIM

+ CHESEBROUGH PONDS	1.2%	N19518	001	Jun 03, 1987
---------------------	------	--------	-----	--------------

OTC DRUG PRODUCT LIST

4 - 15 (of 15)

TERBINAFINE HYDROCHLORIDE

CREAM; TOPICAL				
LAMISIL				
+ NOVARTIS	1%	N20980	001	Mar 09, 1999
SOLUTION; TOPICAL				
LAMISIL AT				
+ NOVARTIS	1%	N21124	001	Mar 17, 2000
SPRAY; TOPICAL				
LAMISIL AT				
+ NOVARTIS	1%	N21124	002	Mar 17, 2000

TIOCONAZOLE

OINTMENT; VAGINAL				
TIOCONAZOLE				
PERRIGO	6.5%	N75915	001	Nov 21, 2001
VAGISTAT-1				
+ NOVARTIS	6.5%	N20676	001	Feb 11, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCT LIST

5 - 1 of 5

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT 4% SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

MEDSHEP CORPORATION

N760305

Jun 30, 1978

ANTICOAGULANT CITRATE DEXTROSE SOLUTION (ACD)

INJECTABLE; INJECTION

CYTOSOL

LABORATORIES INC

N020037

Aug 26, 2003

ACD-A SOLUTION

GAMBRO BCT INC

N010228

ANDA

Feb 25, 2002

ADSOL WITH ACD-A

BAXTER HEALTHCARE

CORP FENWAL DIVISION

N000922

Aug 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION FORMULA A

HAEMONETICS CORP

N980728

ANDA

Feb 06, 2002

AS3 SOLUTION/ACD-A

GAMBRO BCT INC

N001214

May 29, 2002

ANTICOAGULANT CITRATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

BAXTER HEALTHCARE

CORP - FENWAL

DIVISION

N100855

Jun 03, 1959

FRESENIUS USA INC

N160918

Mar 17, 1978

MEDSEP CORPORATION

N110912

Sep 02, 1959

MILES INC

N710497

ANDA

N100102

Dec 14, 1961

ANTICOAGULANT CITRATE PHOSPHATE 2X DEXTROSE SOLUTION (CP2D)

INJECTABLE; INJECTION

CITRATE PHOSPHATE DOUBLE DEXTROSE/ADDITIVE SOLUTION 3

HAEMONETICS CORP

N000127

Jan 18, 2002

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION

INJECTABLE; INJECTION

NONE

FRESENIUS USA INC

N780519

Apr 23, 1980

TERUMO MEDICAL CORP

N820528

Nov 03, 1982

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION (CPDA)

INJECTABLE; INJECTION

CPDA-1 BLOOD-PACK UNIT (PL 146 PLASTIC) 250, 450, 500 ML BLOOD PACK UNITS

BAXTER HEALTHCARE

N770420

May 12, 1978

CORP

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE SOLUTION USP

INJECTABLE; INJECTION

BLOOD PACK UNIT CPDA-1 IN PLASTIC CONTAINER

BAXTER HEALTHCARE

N940404

Jul 28, 1994

CORP - FENWAL

DIVISION

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE ADENINE-1 SOLUTION

INJECTABLE; INJECTION

NONE

MEDSEP CORPORATION

N800077

Nov 06, 1980

26TH EDITION - 2006 - APPROVED DRUG PRODUCT LIST

5 - 2 of 5

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION

INJECTABLE; INJECTION

ADSOL IN PLASTIC CONTAINER

BAXTER HEALTHCARE
CORP - FENWAL
DIVISION

N900223

Dec 27, 1991

CDP BLOOD BAG UNIT IN PLASTIC CONTAINER

BAXTER HLTHCARE -
FENWAL DIVISION

N900224

Dec 27, 1991

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION (CPD)

INJECTABLE; INJECTION

MACOPRODUCTIONS SAS CDP/AS-1: MACOPHARMA LEUCOFLEX MTL1 LEUKOREDUCTION SYSTEM FOR BLOOD
COMPONENTS KNOWN AS MTL1-WB

MACOPRODUCTIONS SAS

N040083

Nov 21, 2005

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP

INJECTABLE; INJECTION

NONE

BAXTER HEALTHCARE
CORP - FENWAL
DIVISION

N170401

Dec 06, 1977

FRESENIUS USA INC

N811012

Jun 28, 1983

MEDSEP CORPORATION

N160907

May 16, 1973

MILES INC

N800222

Aug 23, 1982

TERUMO MEDICAL CORP

N160527

Jun 22, 1970

N781211

Jun 10, 1981

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-1:DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

INJECTABLE; INJECTION

ADSOL RED BLOOD CELL PRESERVATIVE SOLUTION

BAXTER HEALTHCARE
CORP - FENWAL
DIVISION

N811104

May 16, 1983

ANTICOAGULANT CITRATE PHOSPHATE DEXTROSE SOLUTION USP WITH: AS-5:DEXTROSE USP; SODIUM CHLORIDE USP; MANNITOL USP; ADENINE

INJECTABLE; INJECTION

OPTISOL RED BLOOD CELL PRESERVATIVE SOLUTION

TERUMO MEDICAL CORP

N880217

Oct 07, 1988

ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:AS-3: CITRIC ACID USP; MONOBASIC SODIUM PHOSPHATE USP; SODIUM CHLORIDE USP; ADENINE;
DEXTROSE USP; SODIUM CITRATE USP

INJECTABLE; INJECTION

AS-3 NUTRICEL ADDITIVE SYSTEM

MEDSEP CORP

0.042GM/100ML;0.276GM/100ML;
0.410GM/100ML;0.30GM/100ML;
1.10GM/100ML;0.588GM/100ML

N820915

Oct 19, 1984

ANTICOAGULANT CITRATE PHOSPHATE DOUBLE DEXTROSE SOLUTION WITH:AS-2: CITRIC ACID USP; DIBASIC SODIUM PHOSPHATE USP; SODIUM CHLORIDE USP; ADENINE;
DEXTROSE USP; SODIUM CITRATE USP

INJECTABLE; INJECTION

AS-2 NUTRICEL ADDITIVE SYSTEM

MEDSEP CORP

0.042GM/100ML;0.285GM/100ML;
0.718GM/100ML;0.017GM/100ML;
0.396GM/100ML;0.588GM/100ML

N820915

Sep 22, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCT LIST

5 - 3 of 5

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

ANTICOAGULANT ETHYLENEDIAMINE TETRAACETIC ACID (EDTA)

INJECTABLE; INJECTION

NONE

BAXTER HEATHCARE
CORP

N090480

Mar 07, 1955

ANTICOAGULANT HEPARIN SOLUTION USP

INJECTABLE; INJECTION

NONE

FRESENIUS USA INC

N770822

May 17, 1978

ANTICOAGULANT SODIUM CITRATE 4% SOLUTION

INJECTABLE; INJECTION

NONE

HAEMONETICS
CORPORATION

N980123

Mar 03, 2000

ANTICOAGULANT SODIUM CITRATE SOLUTION

INJECTABLE; INJECTION

TRICITRASOL

CYTOSOL
LABORATORIES INC

N010409

Jul 10, 2003

ANTICOAGULANT SODIUM CITRATE SOLUTION USP

INJECTABLE; INJECTION

NONE

BAXTER HEALTHCARE
CORP - FENWAL
DIVISION

N770923

Jan 20, 1978

FRESENIUS USA INC

N160702

Dec 28, 1970

TERUMO MEDICAL CORP

N781214

Feb 08, 1980

DEXTRAN 1 IN SODIUM CHLORIDE 0.6%

INJECTABLE; INJECTION

PROMIT

PHARMALINK
BASLAKEMEDEL AB

N830715

Oct 30, 1984

DEXTRAN 40, 10% IN DEXTROSE 5%

INJECTABLE; INJECTION

GENTRAN 40

BAXTER HLTHCARE IV 10GM/100ML;5GM/100ML
SYSTEMS

N840619

Feb 22, 1985

LMD IN PLASTIC CONTAINER

ABBOTT LABS 10GM/100ML;5GM/100ML

N720563

ANDA

Oct 30, 1992

NONE

ABBOTT LABS 10GM/100ML;5GM/100ML

N160375

Jul 25, 1967

B BRAUN MEDICAL INC 10GM/100ML;5GM/100ML

N160767

Apr 06, 1970

MILES INC 10GM/100ML;5GM/100ML

N160653

Sep 23, 1969

PHARMACHEM 10GM/100ML;5GM/100ML

N160836

Nov 14, 1972

RHEOMACRODEX

MEDA AB 10GM/100ML;5GM/100ML

N140716

Jan 18, 1967

26TH EDITION - 2006 - APPROVED DRUG PRODUCT LIST

5 - 4 of 5

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

DEXTRAN 40, 10% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

GENTRAN 40

BAXTER HLTHCARE IV SYSTEMS	10GM/100ML;0.9GM/100ML	N840620	Feb 22, 1985
-------------------------------	------------------------	---------	--------------

LMD IN PLASTIC CONTAINER

ABBOTT LABS	10GM/100ML;0.9GM/100ML	N720562	ANDA Oct 30, 1992
-------------	------------------------	---------	-------------------

NONE

ABBOTT LABS	10GM/100ML;0.9GM/100ML	N160375	Jul 25, 1967
-------------	------------------------	---------	--------------

MCGAW INC	10GM/100ML;0.9GM/100ML	N160767	Apr 06, 1970
-----------	------------------------	---------	--------------

MILES INC	10GM/100ML;0.9GM/100ML	N160653	Sep 23, 1969
-----------	------------------------	---------	--------------

PHARMACHEM	10GM/100ML;0.9GM/100ML	N160836	Nov 14, 1972
------------	------------------------	---------	--------------

RHEOMACRODEX

PHARMALINK	10GM/100ML;0.9GM/100ML	N140716	Jan 18, 1967
------------	------------------------	---------	--------------

BASLAKEMEDEL AB			
-----------------	--	--	--

DEXTRAN 40, 10% W/V DEXTRAN 40 IN 0.8% NACL 500 ML PVC BAGS

INJECTABLE; INJECTION

RHEOMACRODEX

PHARMALINK		N830527	Mar 27, 1985
------------	--	---------	--------------

BASLAKEMEDEL AB			
-----------------	--	--	--

DEXTRAN 40, 10% W/V DEXTRAN 40 IN 5% DEXTROSE 500 ML PVC BAGS

INJECTABLE; INJECTION

RHEOMACRODEX

PHARMALINK		N830627	Mar 27, 1985
------------	--	---------	--------------

BASLAKEMEDEL AB			
-----------------	--	--	--

DEXTRAN 70, 6% IN DEXTROSE 5%

INJECTABLE; INJECTION

MACRODEX

MEDA AB	6GM/100ML;5GM/100ML	N060826	Jun 08, 1954
---------	---------------------	---------	--------------

NONE

ABBOTT LABS	6GM/100ML;5GM/100ML	N080819	Mar 31, 1953
-------------	---------------------	---------	--------------

DEXTRAN 70, 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

GENTRAIN 70

BAXTER HLTHCARE IV SYSTEMS		N160607	Jan 26, 1970
-------------------------------	--	---------	--------------

MACRODEX

PHARMALINK	6GM/100ML;5GM/100ML	N060826	Jun 08, 1954
------------	---------------------	---------	--------------

BASLAKEMEDEL AB			
-----------------	--	--	--

NONE

MCGAW INC	6GM/100ML;0.9 GM/100ML	N090024	Aug 18, 1969
-----------	------------------------	---------	--------------

MILES INC	6GM/100ML;0.9 GM/100ML	N080716	Mar 13, 1953
-----------	------------------------	---------	--------------

DEXTRAN 70, 6% W/V DEXTRAN 70 IN 0.9% NACL IN 500 ML PVC BAGS

INJECTABLE; INJECTION

MACRODEX

PHARMALINK		N830613	Mar 27, 1985
------------	--	---------	--------------

BASLAKEMEDEL AB			
-----------------	--	--	--

DEXTRAN 70, 6% W/V DEXTRAN 70 IN 5% DEXTROSE

INJECTABLE; INJECTION

MACRODEX

PHARMALINK		N830629	Mar 27, 1985
------------	--	---------	--------------

BASLAKEMEDEL AB			
-----------------	--	--	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCT LIST

5 - 5 of 5

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

DEXTRAN 75, 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

NONE

PHARMACHEM

6GM/100ML;0.9GM/100ML

N080564

Sep 19, 1952

6GM/100ML;0.9GM/100ML

N160759

Aug 19, 1970

HETASTARCH 6% IN 0.9% NACL

INJECTABLE; INJECTION

NONE

GENSIA LABS INC

6GM/100ML;0.9GM/100ML

N740592

ANDA

Nov 12, 1998

MCGAW INC

6GM/100ML;0.9GM/100ML

N740283

ANDA

Oct 21, 1998

HETASTARCH 6% IN LACTATED ELECTROLYTE INJECTION

INJECTABLE; INJECTION

HEXTEND

BIOTIME INC

6GM/100ML

N200952

Mar 31, 1999

HETASTARCH 6% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

6% HETASTARCH IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

ABBOTT

6GM/100ML;0.9GM/100ML

N740193

ANDA

Jan 30, 1995

HESPAN

MCGAW INC

6GM/100ML;0.9GM/100ML

N160889

Jul 17, 1972

HESPAN IN PLASTIC CONTAINER

MCGAW INC

6GM/100ML;0.9GM/100ML

N890105

Apr 04, 1991

HETASTARCH 6% IN SODIUM CHLORIDE 0.9% NACL 500 ML GLASS BOTTLES

INJECTABLE; INJECTION

NONE

ABBOTT

6GM/100ML;0.9 GM/100ML

N720746

ANDA

Feb 07, 1996

LABORATORIES -

HOSPITAL PRODUCTS

DIV

PENTASTARCH 10% IN SODIUM CHLORIDE 0.9%

INJECTABLE; INJECTION

PENTASPAN

MCGAW INC

10GM/100ML;0.9GM/100ML

N841207

May 19, 1987

PENTASPAN IN PLASTIC CONTAINER

MCGAW INC

10GM/100ML;0.9GM/100ML

N890104

Apr 04, 1991

PERFLUORODECALIN; PERFLUOROTRI-N-PROPYLAMINE

INJECTABLE; INJECTION

FLUOSOL

ALPHA THERPTC

17.5GM/100ML;7.5GM/100ML

N860909

Dec 26, 1989

RED BLOOD CELL PROCESSING SOLUTION

INJECTABLE; INJECTION

REJUVESOL

CYTOSOL LABS INC

N950522

Feb 26, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 1 (of 263)

ACETAMINOPHEN

INJECTABLE; INJECTION

INJECTAPAP

ORTHO MCNEIL PHARM

100MG/ML

N17785 001

Mar 07, 1986

SUPPOSITORY; RECTAL

ACETAMINOPHEN

ABLE

120MG

N73106 001

Feb 27, 1995

325MG

N73107 001

Feb 27, 1995

650MG

N73108 001

Feb 27, 1995

ROXANE

120MG

N71010 001

May 12, 1987

650MG

N71011 001

May 12, 1987

TYLENOL

MCNEIL CONS SPECLT

120MG

N17756 002

650MG

N17756 001

ACETAMINOPHEN; ASPIRIN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE

MIKART

150MG;180MG;15MG

N81095 001

Oct 26, 1990

150MG;180MG;60MG

N81097 001

Oct 26, 1990

CODEINE, ASPIRIN, APAP FORMULA NO. 2

SCHERER LABS

150MG;180MG;15MG

N85640 001

CODEINE, ASPIRIN, APAP FORMULA NO. 3

SCHERER LABS

150MG;180MG;30MG

N85639 001

CODEINE, ASPIRIN, APAP FORMULA NO. 4

SCHERER LABS

150MG;180MG;60MG

N85638 001

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

BANCAP

FOREST PHARMS

325MG;50MG

N88889 001

Jan 16, 1986

TRIAPRIN

DUNHALL

325MG;50MG

N89268 001

Jul 02, 1987

TABLET; ORAL

BUTALBITAL AND ACETAMINOPHEN

HALSEY

325MG;50MG

N89568 001

Oct 05, 1988

WATSON LABS

325MG;50MG

N87550 001

Oct 19, 1984

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

ACETAMINOPHEN, BUTALBITAL AND CAFFEINE

GILBERT LABS

325MG;50MG;40MG

N88825 001

Dec 05, 1984

ANOQUAN

SHIRE

325MG;50MG;40MG

N87628 001

Oct 01, 1986

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

GRAHAM DM

325MG;50MG;40MG

N88743 001

Jul 18, 1985

325MG;50MG;40MG

N88765 001

Mar 27, 1985

325MG;50MG;40MG

N89067 001

Apr 19, 1985

MALLINCKRODT

325MG;50MG;40MG

N88758 001

Mar 27, 1985

FEMCET

MALLINCKRODT

325MG;50MG;40MG

N89102 001

Jun 19, 1985

MEDIGESIC PLUS

US CHEM

325MG;50MG;40MG

N89115 001

Jan 14, 1986

TRIAD

MALLINCKRODT

325MG;50MG;40MG

N89023 001

Jun 19, 1985

TABLET; ORAL

ACETAMINOPHEN, BUTALBITAL AND CAFFEINE

GILBERT LABS

325MG;50MG;40MG

N87629 001

Nov 13, 1984

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

ABLE

325MG;50MG;40MG

N40390 001

Jul 23, 2001

500MG;50MG;40MG

N40394 001

Jul 23, 2001

ESGIC

FOREST PHARMS

325MG;50MG;40MG

N89660 001

Dec 23, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 2 (of 263)

ACETAMINOPHEN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE
ABLE 325MG;50MG;40MG;30MG

N76528 001 Aug 21, 2003

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

DHC PLUS

PURDUE FREDERICK 356.4MG;30MG;16MG

N88584 001 Mar 04, 1986

SYNALGOS-DC-A

LEITNER PHARMS 356.4MG;30MG;16MG

N89166 001 May 14, 1986

ACETAMINOPHEN; CODEINE PHOSPHATE

CAPSULE; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

TEVA 300MG;30MG

N88324 001 Dec 29, 1983

ACETAMINOPHEN W/ CODEINE #2

TEVA 300MG;15MG

N88537 001 Jun 04, 1984

ACETAMINOPHEN W/ CODEINE #4

TEVA 300MG;60MG

N88599 001 Jun 01, 1984

PHENAPHEN W/ CODEINE NO. 2

ROBINS AH 325MG;15MG

N84444 001

PHENAPHEN W/ CODEINE NO. 3

ROBINS AH 325MG;30MG

N84445 001

PHENAPHEN W/ CODEINE NO. 4

ROBINS AH 325MG;60MG

N84446 001

PROVAL #3

SOLVAY 325MG;30MG

N85685 001

TYLENOL W/ CODEINE NO. 3

ORTHO MCNEIL PHARM 300MG;30MG

N87422 001

TYLENOL W/ CODEINE NO. 4

ORTHO MCNEIL PHARM 300MG;60MG

N87421 001

SOLUTION; ORAL

TYLENOL W/ CODEINE

ORTHO MCNEIL PHARM 120MG/5ML;12MG/5ML

N85057 001

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

ABLE 300MG;30MG

N40452 001

300MG;60MG

N40459 001

DURAMED PHARMS BARR 300MG;15MG

N88353 001

300MG;30MG

N88354 001

300MG;60MG

N88355 001

HALSEY 300MG;60MG

N86549 001

KV PHARM 300MG;30MG

N85288 001

300MG;60MG

N85365 001

325MG;15MG

N85364 001

325MG;45MG **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N85363 001

MIKART 300MG;60MG

N89244 001

MUTUAL PHARM 300MG;15MG

N85795 001

300MG;30MG

N85794 001

300MG;60MG

N89673 001

PUREPAC PHARM 300MG;30MG

N86681 001

300MG;60MG

N86683 001

ROXANE 500MG;15MG

N89511 001

500MG;30MG

N89512 001

500MG;60MG

N89513 001

SANDOZ 300MG;15MG

N87433 001

300MG;30MG

N85291 002

300MG;30MG

N85917 001

300MG;60MG

N85964 001

300MG;60MG

N87423 001

WARNER CHILCOTT 300MG;15MG

N85992 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 3 (of 263)

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE			
WARNER CHILCOTT	300MG;30MG	N85218	002
ACETAMINOPHEN AND CODEINE PHOSPHATE #2			
SUPERPHARM	300MG;15MG	N89183	001 Oct 18, 1985
ACETAMINOPHEN AND CODEINE PHOSPHATE #3			
PUREPAC PHARM	300MG;30MG	N89080	001 Jul 17, 1986
SUPERPHARM	300MG;30MG	N89184	001 Oct 18, 1985
	300MG;30MG	N89253	001 May 19, 1986
ACETAMINOPHEN AND CODEINE PHOSPHATE #4			
SUPERPHARM	300MG;60MG	N89185	001 Oct 18, 1985
	300MG;60MG	N89254	001 May 19, 1986
ACETAMINOPHEN AND CODEINE PHOSPHATE NO. 2			
AM THERAP	300MG;15MG	N89478	001 Mar 03, 1987
	300MG;15MG	N89481	001 Mar 03, 1987
ACETAMINOPHEN AND CODEINE PHOSPHATE NO. 3			
AM THERAP	300MG;30MG	N89479	001 Mar 03, 1987
	300MG;30MG	N89482	001 Mar 03, 1987
ACETAMINOPHEN AND CODEINE PHOSPHATE NO. 4			
AM THERAP	300MG;60MG	N89480	001 Mar 03, 1987
	300MG;60MG	N89483	001 Mar 03, 1987
ROXANE	300MG;60MG	N84667	001
ACETAMINOPHEN W/ CODEINE			
LEDERLE	300MG;30MG	N87141	001
MUTUAL PHARM	300MG;60MG	N87653	001 Apr 13, 1982
ACETAMINOPHEN W/ CODEINE NO. 2			
ROXANE	300MG;15MG	N84659	001
ACETAMINOPHEN W/ CODEINE PHOSPHATE			
HALSEY	300MG;15MG	N83871	001
	300MG;30MG	N83872	001
USL PHARMA	300MG;30MG	N87919	001 Jun 22, 1982
	300MG;60MG	N87920	001 Jun 22, 1982
VITARINE	300MG;30MG	N85676	001
WARNER CHILCOTT	300MG;60MG	N87306	001
WATSON LABS	300MG;15MG	N87277	001 May 26, 1982
	300MG;30MG	N87276	001 May 26, 1982
	300MG;60MG	N87275	001 May 26, 1982
WHITEWORTH TOWN PLSN	300MG;30MG	N84360	001
	300MG;60MG	N85607	001
APAP W/ CODEINE PHOSPHATE			
EVERYLIFE	325MG;30MG	N85217	001
CAPITAL WITH CODEINE			
CARNRICK	325MG;30MG	N83643	001
CODEINE PHOSPHATE AND ACETAMINOPHEN			
VALEANT PHARM INTL	300MG;30MG	N85896	001
EMPRACET W/ CODEINE PHOSPHATE #3			
GLAXOSMITHKLINE	300MG;30MG	N83951	001
EMPRACET W/ CODEINE PHOSPHATE #4			
GLAXOSMITHKLINE	300MG;60MG	N83951	002
PAPA-DEINE #3			
VANGARD	300MG;30MG	N88037	001 Mar 20, 1984
PAPA-DEINE #4			
VANGARD	300MG;60MG	N88715	001 Mar 20, 1984
PHENAPHEN-650 W/ CODEINE			
ROBINS AH	650MG;30MG	N85856	001
TYLENOL W/ CODEINE			
ORTHO MCNEIL PHARM	325MG;7.5MG	N85056	001
	325MG;15MG	N85056	002
	325MG;30MG	N85056	003
	325MG;60MG	N85056	004
TYLENOL W/ CODEINE NO. 1			
ORTHO MCNEIL PHARM	300MG;7.5MG	N85055	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 4 (of 263)

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

TYLENOL W/ CODEINE NO. 2

ORTHO MCNEIL PHARM 300MG;15MG

N85055 002

TYLENOL W/ CODEINE NO. 4

ORTHO MCNEIL PHARM 300MG;60MG

N85055 004

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN AND HYDROCODONE BITARTRATE

CENT PHARMS 500MG;5MG

N88898 001 Mar 27, 1985

BANCAP HC

FOREST PHARMS 500MG;5MG

N87961 001 Mar 17, 1983

CO-GESIC

CENT PHARMS 500MG;5MG

N89360 001 Mar 02, 1988

TABLET; ORAL

DURADYNE DHC

FOREST PHARMS 500MG;5MG

N87809 001 Mar 17, 1983

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

ABLE 325MG;5MG

N40478 001 Nov 08, 2002

325MG;7.5MG

N40464 001 Oct 23, 2002

325MG;10MG

N40464 002 Oct 23, 2002

500MG;5MG

N40477 001 Nov 06, 2002

500MG;7.5MG

N40490 001 May 21, 2003

500MG;10MG

N40473 001 Nov 06, 2002

650MG;7.5MG

N40474 001 Jan 02, 2003

650MG;10MG

N40476 001 Oct 23, 2002

750MG;7.5MG

N40469 001 Oct 25, 2002

BARR 500MG;5MG

N88577 001 Dec 21, 1984

HALSEY 500MG;5MG

N89554 001 Jun 12, 1987

USL PHARMA 500MG;5MG

N89290 001 May 29, 1987

500MG;5MG

N89291 001 May 29, 1987

WATSON LABS 325MG;7.5MG

N40248 001 Apr 28, 2000

NORCET

ABANA 500MG;5MG

N88871 001 May 15, 1986

TYCOLET

ORTHO MCNEIL PHARM 500MG;5MG

N89385 001 Aug 27, 1986

VICODIN

ABBOTT 500MG;5MG

N85667 001

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

OXYCODONE AND ACETAMINOPHEN

HALSEY 500MG;5MG

N89994 001 May 04, 1989

TYLOX-325

ORTHO MCNEIL PHARM 325MG;5MG

N88246 001 Nov 08, 1984

TABLET; ORAL

OXYCODONE 2.5/APAP 500

BRISTOL MYERS SQUIBB 500MG;2.5MG

N85910 001

OXYCODONE 5/APAP 500

BRISTOL MYERS SQUIBB 500MG;5MG

N85911 001

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

CAPSULE; ORAL

TYLOX

ORTHO MCNEIL PHARM 500MG;4.5MG;0.38MG

N85375 001

ACETAMINOPHEN; PROPOXYPHENE HYDROCHLORIDE

TABLET; ORAL

DARVOCET

AAIPHARMA LLC 325MG;32.5MG

N16844 001

DOLENE AP-65

LEDERLE 650MG;65MG

N85100 001

DISCONTINUED DRUG PRODUCT LIST

6 - 5 (of 263)

ACETAMINOPHEN; PROPOXYPHENE HYDROCHLORIDE

TABLET; ORAL			
PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN			
MYLAN	325MG;32MG	N83689	001

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL			
PROPACET 100			
TEVA	650MG;100MG	N70107	001 Jun 12, 1985
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN			
ABLE	650MG;100MG	N75838	001 Jul 11, 2001
HALSEY	325MG;50MG	N72105	001 May 13, 1988
	650MG;100MG	N72106	001 May 13, 1988
MUTUAL PHARM	325MG;50MG	N70115	001 Jun 12, 1985
	650MG;100MG	N70116	001 Jun 12, 1985
SUPERPHARM	650MG;100MG	N71319	001 Jan 06, 1987
TEVA	650MG;100MG	N70732	001 Jan 03, 1986
WATSON LABS	325MG;50MG	N70398	001 Dec 18, 1986
	650MG;100MG	N70399	001 Dec 18, 1986

ACETAZOLAMIDE

CAPSULE, EXTENDED RELEASE; ORAL			
DIAMOX			
DURAMED PHARMS BARR	500MG	N12945	001
TABLET; ORAL			
ACETAZOLAMIDE			
ALRA	250MG	N83320	001
ASCOT	250MG	N87686	001 Oct 20, 1982
VANGARD	250MG	N87654	001 Feb 05, 1982
WATSON LABS	250MG	N84498	002

ACETIC ACID, GLACIAL

SOLUTION/DROPS; OTIC			
ACETIC ACID			
KV PHARM	2%	N85493	001
ORLEX			
PROCTER AND GAMBLE	2%	N86845	001
VOSOL			
MEDPOINTE PHARM HLC	2%	N12179	001

ACETIC ACID, GLACIAL; ALUMINUM ACETATE

SOLUTION/DROPS; OTIC			
BOROFAIR			
PHARMAFAIR	2%;0.79%	N88606	001 Aug 21, 1985
DOMEBORO			
BAYER PHARMS	2%;0.79%	N84476	001

ACETIC ACID, GLACIAL; DESONIDE

SOLUTION/DROPS; OTIC			
TRIDESILON			
BAYER PHARMS	2%;0.05%	N17914	001

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC			
ACETIC ACID W/ HYDROCORTISONE			
KV PHARM	2%;1%	N85492	001
HYDROCORTISONE AND ACETIC ACID			
BAUSCH AND LOMB	2%;1%	N40097	001 Oct 31, 1994
ORLEX HC			
PROCTER AND GAMBLE	2%;1%	N86844	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 6 (of 263)

ACETIC ACID, GLACIAL; HYDROCORTISONE; NEOMYCIN SULFATE

SUSPENSION/DROPS; OTIC

NEO-CORT-DOME

BAYER PHARMS

2%;1%;EQ 0.35% BASE

N50238 001

ACETOHEXAMIDE

TABLET; ORAL

ACETOHEXAMIDE

USL PHARMA

250MG

N70753 001

Nov 03, 1986

500MG

N70754 001

Nov 03, 1986

DYMELOR

LILLY

250MG

N13378 002

500MG

N13378 001

ACETOPHENAZINE MALEATE

TABLET; ORAL

TINDAL

SCHERING

20MG

N12254 002

ACETRIZOATE SODIUM

SOLUTION; INTRAUTERINE

SALPIX

ORTHO MCNEIL PHARM

53%

N09008 001

ACETYLCYSTEINE

SOLUTION; INHALATION, ORAL

ACETYLCYSTEINE

ROXANE

10%

N72621 001

Sep 30, 1992

20%

N72622 001

Sep 30, 1992

MUCOSIL-10

DEY

10%

N70575 001

Oct 14, 1986

MUCOSIL-20

DEY

20%

N70576 001

Oct 14, 1986

ACETYLCYSTEINE; ISOPROTERENOL HYDROCHLORIDE

SOLUTION; INHALATION

MUCOMYST W/ ISOPROTERENOL

MEAD JOHNSON

10%;0.05%

N17366 001

ACETYLDIGITOXIN

TABLET; ORAL

ACYLANID

NOVARTIS

0.1MG

N09436 001

ACRISORCIN

CREAM; TOPICAL

AKRINOL

SCHERING

2MG/GM

N12470 001

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

LEK PHARM

200MG

N74750 001

Apr 22, 1997

ROXANE

200MG

N74570 002

Apr 22, 1997

TABLET; ORAL

ACYCLOVIR

CLONMEL HLTHCARE

400MG

N74834 001

Apr 24, 1997

800MG

N74834 002

Apr 24, 1997

LEK PHARM

400MG

N74658 001

Apr 22, 1997

800MG

N74658 002

Apr 22, 1997

TEVA

200MG **Federal Register determination
that product was not discontinued or
withdrawn for safety or efficacy
reasons**

N74556 001

Apr 22, 1997

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 7 (of 263)

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR

ABBOTT

EQ 50MG BASE/ML

N75114 001

Jul 26, 1999

ACYCLOVIR SODIUM

APOTHECON

EQ 500MG BASE/VIAL

N74897 001

Feb 27, 1998

EQ 1GM BASE/VIAL

N74897 002

Feb 27, 1998

ZOVIRAX

GLAXOSMITHKLINE

EQ 250MG BASE/VIAL

N18603 003

Aug 30, 1983

ADAPALENE

SOLUTION; TOPICAL

DIFFERIN

GALDERMA LABS LP

0.1%

N20338 001

May 31, 1996

ALATROFLOXACIN MESYLATE

INJECTABLE; INJECTION

TROVAN PRESERVATIVE FREE

PFIZER

EQ 200MG BASE/VIAL

N20760 001

Dec 18, 1997

EQ 300MG BASE/VIAL

N20760 002

Dec 18, 1997

ALBUMIN CHROMATED CR-51 SERUM

INJECTABLE; INJECTION

CHROMALBIN

ISO TEX

100uCi/VIAL

N17835 001

250uCi/VIAL

N17835 002

500uCi/VIAL

N17835 003

ALBUMIN IODINATED I-125 SERUM

INJECTABLE; INJECTION

RADIO-IODINATED (I 125) SERUM ALBUMIN (HUMAN)

BAYER PHARMS

2.5uCi/AMP

N17846 001

RADIOIODINATED SERUM ALBUMIN (HUMAN) IHSA I 125

MALLINCKRODT

6.67uCi/ML

N17844 003

10uCi/ML

N17844 001

100uCi/ML

N17844 002

ALBUMIN IODINATED I-131 SERUM

INJECTABLE; INJECTION

MEGATOPE

ISO TEX

2mCi/VIAL

N17837 003

5uCi/AMP

N17837 004

20uCi/AMP

N17837 005

ALBUTEROL SULFATE

CAPSULE; INHALATION

VENTOLIN ROTACAPS

GLAXOSMITHKLINE

EQ 0.2MG BASE

N19489 001

May 04, 1988

SOLUTION; INHALATION

ALBUTEROL SULFATE

COPLY PHARM

EQ 0.083% BASE

N73495 001

May 28, 1993

EQ 0.5% BASE

N73307 001

Nov 27, 1991

PROVENTIL

SCHERING

EQ 0.083% BASE

N19243 002

Jan 14, 1987

EQ 0.5% BASE

N19243 001

Jan 14, 1987

VENTOLIN

GLAXOSMITHKLINE

EQ 0.083% BASE

N19773 001

Apr 23, 1992

EQ 0.5% BASE

N19269 002

Jan 16, 1987

SYRUP; ORAL

ALBUTEROL SULFATE

MOVA

EQ 2MG BASE/5ML

N74302 001

Sep 30, 1994

WATSON LABS

EQ 2MG BASE/5ML

N73165 001

Apr 29, 1993

PROVENTIL

SCHERING

EQ 2MG BASE/5ML

N18062 001

Jan 19, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 8 (of 263)

ALBUTEROL SULFATESYRUP; ORAL
VENTOLIN

GLAXOSMITHKLINE EQ 2MG BASE/5ML

N19621 001 Jun 10, 1987

TABLET; ORAL
ALBUTEROL SULFATE

AM THERAP EQ 2MG BASE

N72449 001 Dec 05, 1989

EQ 4MG BASE

N72450 001 Dec 05, 1989

CLONMEL HLTHCARE EQ 2MG BASE

N72859 001 Dec 20, 1989

EQ 4MG BASE

N72860 001 Dec 20, 1989

COPLEY PHARM EQ 2MG BASE

N72966 001 Nov 22, 1991

EQ 4MG BASE

N72967 001 Nov 22, 1991

UCB EQ 2MG BASE

N73120 001 Sep 29, 1992

EQ 4MG BASE

N73121 001 Sep 29, 1992

WARNER CHILCOTT EQ 2MG BASE

N72817 001 Jan 09, 1990

EQ 4MG BASE

N72818 001 Jan 09, 1990

PROVENTIL

SCHERING EQ 2MG BASE

N17853 001 May 07, 1982

EQ 4MG BASE

N17853 002 May 07, 1982

VENTOLIN

GLAXOSMITHKLINE EQ 2MG BASE

N19112 001 Jul 10, 1986

EQ 4MG BASE

N19112 002 Jul 10, 1986

TABLET, EXTENDED RELEASE; ORAL

PROVENTIL

SCHERING EQ 4MG BASE

N19383 001 Jul 13, 1987

VOLMAX

MURO EQ 4MG BASE

N19604 002 Dec 23, 1992

EQ 8MG BASE

N19604 001 Dec 23, 1992

ALCOHOL

INJECTABLE; INJECTION

ALCOHOL 5% IN DEXTROSE 5%
MILES 5ML/100ML

N83483 001

ALCOHOL; DEXTROSE

INJECTABLE; INJECTION

ALCOHOL 5% IN DEXTROSE 5% IN WATER
BAXTER HLTHCARE 5ML/100ML; 5GM/100ML

N83256 001

ALGLUCERASE

INJECTABLE; INJECTION

CEREDASE

GENZYME 10 UNITS/ML

N20057 004 May 08, 1992

ALKAVERVIR

TABLET; ORAL

VERILOID

3M 2MG

N07336 002

3MG

N07336 003

ALLOPURINOL

TABLET; ORAL

ALLOPURINOL

MUTUAL PHARM 100MG

N70466 001 Dec 24, 1985

300MG

N70467 001 Dec 24, 1985

PUREPAC PHARM 100MG

N70579 001 Apr 14, 1986

300MG

N70580 001 Apr 14, 1986

SUPERPHARM 100MG

N70950 001 Nov 30, 1988

300MG

N70951 001 Nov 30, 1988

WATSON LABS 100MG

N18241 001 Nov 16, 1984

100MG

N18785 001 Sep 28, 1984

300MG

N18241 002 Nov 16, 1984

300MG

N18785 002 Sep 28, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 9 (of 263)

ALLOPURINOL

TABLET; ORAL
LOPURIN
ABBOTT

100MG	N18297	001	
300MG	N18297	002	

ALPHA-TOCOPHEROL; ASCORBIC ACID; BIOTIN; CHOLECALCIFEROL; CYANOCOBALAMIN; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A

INJECTABLE; INJECTION
CERNEVIT-12

BAXTER HLTHCARE	11.2 IU/VIAL;125MG/VIAL;60UGM/VIAL;200 IU/VIAL;5.5MG/VIAL;414UGM/VIAL;46MG/VIAL;17.25MG/VIAL;4.53MG/VIAL;4.14MG/VIAL;3.51MG/VIAL;3,500 IU/VIAL	N20924	001	Apr 06, 1999
-----------------	--	--------	-----	--------------

ALPRAZOLAM

SOLUTION; ORAL
ALPRAZOLAM
ROXANE

0.5MG/5ML	N74314	001	Oct 31, 1993
-----------	--------	-----	--------------

TABLET; ORAL
ALPRAZOLAM

CLONMEL HLTHCARE	0.25MG	N74174	001	Oct 19, 1993
	0.5MG	N74174	002	Oct 19, 1993
	1MG	N74174	003	Oct 19, 1993
	2MG	N74174	004	Oct 19, 1993
ROXANE	0.25MG	N74199	001	Oct 19, 1993
	0.5MG	N74199	002	Oct 19, 1993
	1MG	N74199	003	Oct 19, 1993

ALPROSTADIL

INJECTABLE; INJECTION
CAVERJECT
PFIZER

0.005MG/ML	N20755	001	Oct 31, 1997
0.01MG/ML	N20755	002	Oct 01, 1997
0.02MG/ML	N20755	003	Oct 01, 1997

EDEX

SCHWARZ PHARMA	0.005MG/VIAL	N20649	001	Jun 12, 1997
----------------	--------------	--------	-----	--------------

ALSEROXYLON

TABLET; ORAL
RAUTENSIN
NOVARTIS

2MG	N09215	001	
-----	--------	-----	--

RAUWILOID
3M

2MG	N08867	001	
-----	--------	-----	--

ALUMINUM HYDROXIDE; MAGNESIUM TRISILICATE

TABLET, CHEWABLE; ORAL
ALUMINUM HYDROXIDE AND MAGNESIUM TRISILICATE

PENNEX	80MG;20MG	N89449	001	Nov 27, 1987
--------	-----------	--------	-----	--------------

FOAMICON

NOVARTIS	80MG;20MG	N72687	001	Jun 28, 1989
----------	-----------	--------	-----	--------------

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL
AMANTADINE HYDROCHLORIDE

WATSON LABS	100MG	N71382	001	Jan 21, 1987
-------------	-------	--------	-----	--------------

SYMADINE

SOLVAY	100MG	N71000	001	Sep 04, 1986
--------	-------	--------	-----	--------------

SYMMETREL

ENDO PHARMS	100MG	N16020	001	
-------------	-------	--------	-----	--

SYRUP; ORAL

SYMMETREL	50MG/5ML	N16023	002	
-----------	----------	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 10 (of 263)

AMCINONIDE

CREAM; TOPICAL
CYCLOCORT
ASTELLAS

0.025%

N18116 001

AMDINOCILLIN

INJECTABLE; INJECTION
COACTIN
ROCHE

250MG/VIAL
500MG/VIAL
1GM/VIAL

N50565 001 Dec 21, 1984
N50565 002 Dec 21, 1984
N50565 003 Dec 21, 1984

AMIFOSTINE

INJECTABLE; INJECTION
ETHYOL

MEDIMMUNE ONCOLOGY 375MG/VIAL

N20221 002 Sep 10, 1999

AMIKACIN SULFATE

INJECTABLE; INJECTION
AMIKACIN SULFATE

ABBOTT EQ 250MG BASE/ML
EQ 250MG BASE/ML
ASTRAZENECA EQ 50MG BASE/ML
EQ 250MG BASE/ML
BAXTER HLTHCARE EQ 50MG BASE/ML
EQ 250MG BASE/ML
HOSPIRA EQ 250MG BASE/ML
MAYNE PHARMA USA EQ 50MG BASE/ML
EQ 250MG BASE/ML

N63265 001 Nov 30, 1994
N63266 001 Oct 31, 1994
N63167 001 Dec 14, 1995
N63169 001 Dec 14, 1995
N63274 001 May 18, 1992
N63275 001 May 18, 1992
N64099 001 Jun 20, 1995
N63350 001 Jul 30, 1993
N63350 002 Jul 30, 1993

AMIKIN

APOTHECON EQ 50MG BASE/ML
EQ 50MG BASE/ML
EQ 250MG BASE/ML
EQ 250MG BASE/ML

N50495 001
N62562 001 Sep 20, 1984
N50495 002
N62562 002 Sep 20, 1984

AMIKIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

APOTHECON EQ 5MG BASE/ML
EQ 10MG BASE/ML

N50618 002 Nov 30, 1987
N50618 001 Nov 30, 1987

AMINO ACIDS

INJECTABLE; INJECTION

AMINESS 5.2% ESSENTIAL AMINO ACIDS W/ HISTADINE

HOSPIRA 5.2% (5.2GM/100ML)

N18901 001 Apr 06, 1984

AMINOSYN 3.5% IN PLASTIC CONTAINER

ABBOTT 3.5% (3.5GM/100ML)
3.5% (3.5GM/100ML)

N18804 001 May 15, 1984
N18875 001 Aug 08, 1984

AMINOSYN II 3.5%

HOSPIRA 3.5% (3.5GM/100ML)

N19438 001 Apr 03, 1986

AMINOSYN II 3.5% IN PLASTIC CONTAINER

ABBOTT 3.5% (3.5GM/100ML)

N19491 001 Oct 10, 1986

AMINOSYN II 5%

HOSPIRA 5% (5GM/100ML)

N19438 002 Apr 03, 1986

AMINOSYN-HBC 7% IN PLASTIC CONTAINER

ABBOTT 7% (7GM/100ML)

N19400 001 Jul 23, 1986

BRANCHAMIN 4%

BAXTER HLTHCARE 4% (4GM/100ML)

N18678 001 Sep 28, 1984

FREAMINE 8.5%

B BRAUN 8.5% (8.5GM/100ML)

N16822 001

FREAMINE II 8.5%

B BRAUN 8.5% (8.5GM/100ML)

N16822 002

NEOPHAM 6.4%

HOSPIRA 6.4% (6.4GM/100ML)

N18792 001 Jan 17, 1984

NOVAMINE 15% SULFITE FREE IN PLASTIC CONTAINER

BAXTER HLTHCARE 15% (15GM/100ML)

N20107 001 Feb 05, 1993

DISCONTINUED DRUG PRODUCT LIST

6 - 11 (of 263)

AMINO ACIDS

INJECTABLE; INJECTION

NOVAMINE 8.5%

HOSPIRA

8.5% (8.5GM/100ML)

N17957 002

Aug 09, 1982

AMINO ACIDS; CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

ABBOTT

3.5%;36.8MG/100ML;25GM/100ML;51MG/100ML
;22.4MG/100ML;261MG/100ML;205MG/100ML

N19714 001

Sep 12, 1988

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER

ABBOTT

4.25%;36.8MG/100ML;20GM/100ML;51MG/100M
L;22.4MG/100ML;261MG/100ML;205MG/100ML

N19714 002

Sep 12, 1988

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

ABBOTT

4.25%;36.8MG/100ML;25GM/100ML;51MG/100M
L;22.4MG/100ML;261MG/100ML;205MG/100ML

N19714 004

Sep 12, 1988

AMINOSYN II 5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER

ABBOTT

5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2
2.4MG/100ML;261MG/100ML;205MG/100ML

N19714 003

Sep 12, 1988

HOSPIRA

5%;36.8MG/100ML;25GM/100ML;51MG/100ML;2
2.4MG/100ML;261MG/100ML;205MG/100ML

N19683 004

Nov 07, 1988

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

AMINOSYN 3.5% W/ DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT

3.5%;25GM/100ML

N19118 001

Oct 11, 1984

AMINOSYN 3.5% W/ DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT

3.5%;5GM/100ML

N19120 001

Oct 11, 1984

AMINOSYN 4.25% W/ DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT

4.25%;25GM/100ML

N19119 001

Oct 11, 1984

AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT

3.5%;25GM/100ML

N19505 002

Nov 07, 1986

3.5%;25GM/100ML

N19713 006

Sep 09, 1988

AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT

3.5%;5GM/100ML

N19506 001

Nov 07, 1986

3.5%;5GM/100ML

N19713 002

Sep 09, 1988

AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER

ABBOTT

4.25%;10GM/100ML

N19713 001

Sep 09, 1988

AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER

ABBOTT

4.25%;20GM/100ML

N19713 004

Sep 09, 1988

AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT

4.25%;25GM/100ML

N19504 002

Nov 07, 1986

4.25%;25GM/100ML

N19713 005

Sep 09, 1988

AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT

5%;25GM/100ML

N19565 001

Dec 17, 1986

5%;25GM/100ML

N19713 003

Sep 09, 1988

TRAVASOL 2.75% IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE

2.75%;10GM/100ML

N19520 002

Sep 23, 1988

TRAVASOL 2.75% IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE

2.75%;15GM/100ML

N19520 003

Sep 23, 1988

TRAVASOL 2.75% IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE

2.75%;20GM/100ML

N19520 004

Sep 23, 1988

TRAVASOL 2.75% IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE

2.75%;25GM/100ML

N19520 005

Sep 23, 1988

TRAVASOL 2.75% IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE

2.75%;5GM/100ML

N19520 001

Sep 23, 1988

TRAVASOL 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE

4.25%;10GM/100ML

N19520 007

Sep 23, 1988

TRAVASOL 4.25% IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE

4.25%;15GM/100ML

N19520 008

Sep 23, 1988

TRAVASOL 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE

4.25%;20GM/100ML

N19520 009

Sep 23, 1988

TRAVASOL 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE

4.25%;25GM/100ML

N19520 010

Sep 23, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 12 (of 263)

AMINO ACIDS; DEXTROSE

INJECTABLE; INJECTION

TRAVASOL 4.25% IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%;5GM/100ML

N19520 006 Sep 23, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 4.25% W/ ELECT AND ADJUSTED PHOSPHATE IN DEXTROSE 10% IN PLASTIC CONTAINER

ABBOTT 4.25%;10GM/100ML;51MG/100ML;176.5MG/100ML;22.4MG/100ML;104.5MG/100ML;205MG/100ML

N19712 002 Sep 08, 1988

HOSPIRA

4.25%;10GM/100ML;51MG/100ML;176.5MG/100ML;22.4MG/100ML;104.5MG/100ML;205MG/100ML

N19682 003 Nov 01, 1988

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT 3.5%;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML

N19564 002 Dec 16, 1986

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

ABBOTT 4.25%;25GM/100ML;51MG/100ML;22.4MG/100ML;261MG/100ML;205MG/100ML

N19564 004 Dec 16, 1986

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT 3.5%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML
3.5%;5GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML

N19564 001 Dec 16, 1986

N19712 001 Sep 08, 1988

AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER

ABBOTT 4.25%;10GM/100ML;30MG/100ML;97MG/100ML;120MG/100ML;49.3MG/100ML

N19564 003 Dec 16, 1986

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%;10GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML

N20147 002 Oct 23, 1995

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%;15GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML

N20147 003 Oct 23, 1995

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%;20GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML

N20147 004 Oct 23, 1995

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%;25GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML

N20147 005 Oct 23, 1995

TRAVASOL 2.75% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE 2.75%;5GM/100ML;51MG/100ML;261MG/100ML;216MG/100ML;112MG/100ML

N20147 001 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 10% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%;10GM/100ML;51MG/100ML;261MG/100ML;297MG/100ML;77MG/100ML

N20147 007 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 15% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%;15GM/100ML;51MG/100ML;261MG/100ML;297MG/100ML;77MG/100ML

N20147 008 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 20% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%;20GM/100ML;51MG/100ML;261MG/100ML;297MG/100ML;77MG/100ML

N20147 009 Oct 23, 1995

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 25% IN PLASTIC CONTAINER

BAXTER HLTHCARE 4.25%;25GM/100ML;51MG/100ML;261MG/100ML;297MG/100ML;77MG/100ML

N20147 010 Oct 23, 1995

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 13 (of 263)

AMINO ACIDS; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TRAVASOL 4.25% SULFITE FREE W/ ELECTROLYTES IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	4.25%;56MG/100ML;51MG/100ML;261MG/100ML;	N20147 006	Oct 23, 1995
	297MG/100ML;77MG/100ML		

AMINO ACIDS; MAGNESIUM ACETATE; PHOSPHORIC ACID; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M IN PLASTIC CONTAINER

ABBOTT	3.5%;21MG/100ML;40MG/100ML;128MG/100ML;	N18804 002	May 15, 1984
	234MG/100ML		
	3.5%;21MG/100ML;40MG/100ML;128MG/100ML;	N18875 002	Aug 08, 1984
	234MG/100ML		

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN 3.5% M

HOSPIRA	3.5%;21MG/100ML;128MG/100ML;234MG/100ML	N17789 005	
---------	---	------------	--

AMINO ACIDS; MAGNESIUM ACETATE; POTASSIUM ACETATE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M IN PLASTIC CONTAINER

ABBOTT	3.5%;32MG/100ML;128MG/100ML;222MG/100ML	N19493 001	Oct 16, 1986
	;49MG/100ML		

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATE

INJECTABLE; INJECTION

VEINAMINE 8%

HOSPIRA	8%;61MG/100ML;211MG/100ML;56MG/100ML;38	N17957 001	
	8MG/100ML		

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM CHLORIDE

INJECTABLE; INJECTION

AMINOSYN II 7% W/ ELECTROLYTES

HOSPIRA	7%;102MG/100ML;45MG/100ML;522MG/100ML;4	N19437 006	Apr 03, 1986
	10MG/100ML		

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

AMINOSYN II 3.5% M

HOSPIRA	3.5%;30MG/100ML;97MG/100ML;120MG/100ML;	N19437 007	Apr 03, 1986
	49MG/100ML		

AMINOCAPROIC ACID

INJECTABLE; INJECTION

AMINOCAPROIC ACID

AM PHARM PARTNERS	250MG/ML	N70522 001	Jun 17, 1986
BAXTER HLTHCARE	250MG/ML	N18590 001	Oct 29, 1982

AMINOPHYLLINE

ENEMA; RECTAL

SOMOPHYLLIN

FISONS	300MG/5ML	N18232 001	Apr 02, 1982
--------	-----------	------------	--------------

INJECTABLE; INJECTION

AMINOPHYLLIN

GD SEARLE LLC	25MG/ML	N87243 001	May 24, 1982
	25MG/ML	N87621 001	May 24, 1982

AMINOPHYLLINE

AM PHARM PARTNERS	25MG/ML	N84568 001	
	25MG/ML	N87200 001	
	25MG/ML	N87250 001	Jan 06, 1982

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 14 (of 263)

AMINOPHYLLINE

INJECTABLE; INJECTION

AMINOPHYLLINE

AM PHARM PARTNERS

25MG/ML

N87431 001

25MG/ML

N87886 001

Aug 30, 1983

25MG/ML

N88407 001

Jan 25, 1984

ELKINS SINN

25MG/ML

N87239 001

INTL MEDICATION

25MG/ML

N87867 001

Nov 10, 1983

25MG/ML

N87868 001

Nov 10, 1983

KING PHARMS

25MG/ML

N86606 001

PHARMA SERVE NY

25MG/ML

N87387 001

Jun 03, 1983

SMITH AND NEPHEW

25MG/ML

N88429 001

May 30, 1985

25MG/ML

N88749 001

May 30, 1985

AMINOPHYLLINE IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

HOSPIRA

100MG/100ML

N18924 001

Dec 12, 1984

200MG/100ML

N18924 002

Dec 12, 1984

400MG/100ML

N18924 003

Dec 12, 1984

500MG/100ML

N18924 004

Dec 12, 1984

SOLUTION; ORAL

AMINOPHYLLINE

MORTON GROVE

105MG/5ML

N88156 001

Dec 05, 1983

SOMOPHYLLIN

FISONS

105MG/5ML

N86466 001

SOMOPHYLLIN-DF

FISONS

105MG/5ML

N87045 001

TABLET; ORAL

AMINOPHYLLIN

GD SEARLE LLC

100MG

N02386 002

200MG

N02386 003

AMINOPHYLLINE

ASCOT

100MG

N87522 001

Feb 12, 1982

200MG

N87523 001

Feb 12, 1982

BARR

100MG

N88297 001

Aug 19, 1983

200MG

N88298 001

Aug 19, 1983

DURAMED PHARMS BARR

100MG

N88182 001

Mar 31, 1983

200MG

N88183 001

Mar 31, 1983

HALSEY

100MG

N84674 001

IMPAX LABS

100MG

N84574 001

200MG

N84576 001

KV PHARM

100MG

N85284 001

200MG

N85289 001

LANNETT

100MG

N84588 001

200MG

N84588 002

PAL PAK

100MG

N84533 001

PANRAY

100MG

N84552 001

200MG

N84552 002

PUREPAC PHARM

100MG

N84699 001

200MG

N85333 001

ROXANE

100MG

N87500 001

Feb 09, 1982

200MG

N87501 001

Feb 09, 1982

SANDOZ

100MG

N85261 003

100MG

N85262 002

200MG

N85261 002

VALEANT PHARM INTL

200MG

N84563 001

VANGARD

100MG

N88314 001

Oct 03, 1983

200MG

N88319 001

Oct 03, 1983

VINTAGE PHARMS

100MG

N85409 001

200MG

N85410 001

WATSON LABS

100MG

N85567 001

200MG

N85564 001

TABLET, DELAYED RELEASE; ORAL

AMINOPHYLLINE

IMPAX LABS

100MG

N84577 001

200MG

N84575 001

TABLICAPS

100MG

N84632 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 15 (of 263)

AMINOPHYLLINE

TABLET, DELAYED RELEASE; ORAL			
AMINOPHYLLINE			
VALE	100MG	N84531	001
	200MG	N84530	001
TABLET, EXTENDED RELEASE; ORAL			
PHYLLOCONTIN			
PURDUE FREDERICK	225MG	N86760	001

AMINOSALICYLATE SODIUM

POWDER; ORAL			
P.A.S. SODIUM			
CENTURY PHARMS	4GM/PACKET	N80947	001
SODIUM AMINOSALICYLATE			
HEXCEL	100%	N80097	001
TABLET; ORAL			
PARASAL SODIUM			
PANRAY	500MG	N06811	006
	1GM	N06811	011
TEEBACIN			
CONSOLIDATED MIDLAND	500MG	N07320	002

AMINOSALICYLATE SODIUM; AMINOSALICYLIC ACID

TABLET; ORAL			
NEOPASALATE			
MEDPOINTE PHARM HLC	846MG;112MG	N80059	002

AMINOSALICYLIC ACID

TABLET; ORAL			
PARASAL			
PANRAY	500MG	N06811	001
	1GM	N06811	002

AMINOSALICYLIC ACID RESIN COMPLEX

POWDER; ORAL			
REZIPAS			
BRISTOL MYERS SQUIBB	EQ 500MG BASE/GM	N09052	001

AMITRIPTYLINE HYDROCHLORIDE

CONCENTRATE; ORAL			
ENDEP			
ROCHE	40MG/ML	N85749	001
INJECTABLE; INJECTION			
AMITRIPTYLINE HYDROCHLORIDE			
STERIS	10MG/ML	N85594	001
ELAVIL			
ASTRAZENECA	10MG/ML	N12704	001
TABLET; ORAL			
AMITID			
BRISTOL MYERS SQUIBB	10MG	N86454	001
	25MG	N86454	002
	50MG	N86454	003
	75MG	N86454	004
	100MG	N86454	005
AMITRIL			
WARNER CHILCOTT	10MG	N83939	001
	25MG	N83937	001
	50MG	N83938	002
	75MG	N84957	001
	100MG	N85093	001
	150MG	N86295	001
AMITRIPTYLINE HCL			
UCB			
	10MG	N85864	001
	25MG	N85935	001

DISCONTINUED DRUG PRODUCT LIST

6 - 16 (of 263)

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HCL

UCB

50MG	N85936	001
75MG	N86337	001
100MG	N86336	001
150MG	N86335	001

AMITRIPTYLINE HYDROCHLORIDE

AM THERAP

25MG	N88672	001	Nov 20, 1984
50MG	N88673	001	Nov 20, 1984
75MG	N88674	001	Nov 20, 1984
100MG	N88675	001	Nov 20, 1984

COPLEY PHARM

10MG	N88421	001	Apr 30, 1984
25MG	N88422	001	Apr 30, 1984
50MG	N88423	001	Apr 30, 1984
75MG	N88424	001	Apr 30, 1984
100MG	N88425	001	Apr 30, 1984
150MG	N88426	001	Apr 30, 1984

HALSEY

10MG	N85923	001	
25MG	N85922	001	
50MG	N85925	001	
50MG	N87557	001	Mar 05, 1982
75MG	N85926	001	May 20, 1983
100MG	N85927	001	May 20, 1983

LEDERLE

10MG	N86744	001	
10MG	N87366	001	Jan 04, 1982
25MG	N86746	001	
25MG	N87367	001	May 03, 1982
50MG	N86743	001	
50MG	N87181	001	Jan 04, 1982
75MG	N86745	001	
75MG	N87369	001	Jan 04, 1982
100MG	N86747	001	
100MG	N87368	001	May 03, 1982
150MG	N87370	001	Jan 04, 1982

MUTUAL PHARM

10MG	N85744	001	
25MG	N85627	001	
50MG	N85745	001	
75MG	N85743	001	
100MG	N85742	002	May 11, 1982
150MG	N89423	001	Feb 17, 1987

PAR PHARM

10MG	N88697	001	Sep 25, 1984
25MG	N88698	001	Sep 25, 1984
50MG	N88699	001	Sep 25, 1984
75MG	N88700	001	Sep 25, 1984
100MG	N88701	001	Sep 25, 1984
150MG	N88702	001	Sep 25, 1984

PUREPAC PHARM

10MG	N88075	001	Sep 16, 1983
10MG	N88084	001	Jul 18, 1983
25MG	N88076	001	May 20, 1983
25MG	N88085	001	Jul 18, 1983
50MG	N88077	001	Sep 16, 1983
50MG	N88105	001	Jul 18, 1983
75MG	N88078	001	Sep 16, 1983
75MG	N88106	001	Jul 18, 1983
100MG	N88079	001	Sep 16, 1983
100MG	N88107	001	Jul 18, 1983

ROXANE

10MG	N86002	001
10MG	N86144	001
25MG	N85944	001
25MG	N86145	001
50MG	N85945	001
50MG	N86143	001
75MG	N86004	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 17 (of 263)

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

ROXANE	75MG	N86147	001	
	100MG	N86003	001	
	100MG	N86146	001	
	150MG	N86090	001	
	150MG	N86148	001	
SUPERPHARM	10MG	N88853	001	Nov 13, 1984
	25MG	N88854	001	Nov 13, 1984
	50MG	N88855	001	Nov 13, 1984
	75MG	N88856	001	Nov 13, 1984
	100MG	N88857	001	Nov 13, 1984
TEVA	10MG	N86610	001	
	25MG	N86859	001	
	50MG	N85032	001	
	50MG	N86857	001	
	75MG	N85030	001	
	75MG	N86860	001	
	100MG	N86854	001	
	150MG	N86853	001	
USL PHARMA	25MG	N87775	001	Feb 10, 1982
VANGARD	10MG	N87632	001	Feb 01, 1982
	50MG	N87616	001	Feb 08, 1982
	75MG	N87617	001	Feb 05, 1982
	100MG	N87639	001	Feb 08, 1982
WATSON LABS	10MG	N85816	001	
	10MG	N88620	001	Mar 02, 1984
	25MG	N85817	001	
	25MG	N88621	001	Mar 02, 1984
	50MG	N85815	001	
	50MG	N88622	001	Mar 02, 1984
	75MG	N85819	001	
	75MG	N88633	001	Mar 02, 1984
	100MG	N85820	001	
	100MG	N88634	001	Mar 02, 1984
	150MG	N85821	001	
	150MG	N88635	001	Mar 02, 1984
	10MG	N87647	001	Mar 05, 1982
	25MG	N87278	001	
ELAVIL				
ASTRAZENECA	10MG	N12703	001	
	25MG	N12703	003	
	50MG	N12703	004	
	75MG	N12703	005	
	100MG	N12703	006	
	150MG	N12703	007	
ENDEP				
ROCHE	10MG	N83639	001	
	25MG	N83639	002	
	50MG	N83639	003	
	75MG	N83639	004	
	100MG	N83639	005	
	150MG	N85303	001	

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET; ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

MUTUAL PHARM	EQ 12.5MG BASE;5MG	N70765	001	Dec 10, 1986
	EQ 25MG BASE;10MG	N70766	001	Dec 10, 1986
USL PHARMA	EQ 12.5MG BASE;5MG	N70477	001	Jan 12, 1988
	EQ 25MG BASE;10MG	N70478	001	Jan 12, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 18 (of 263)

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL

ETRAFON 2-10				
SCHERING	10MG;2MG	N14713	007	
ETRAFON 2-25				
SCHERING	25MG;2MG	N14713	004	
ETRAFON-A				
SCHERING	10MG;4MG	N14713	002	
ETRAFON-FORTE				
SCHERING	25MG;4MG	N14713	006	
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE				
IVAX PHARMS	10MG;2MG	N70935	001	Sep 11, 1986
	10MG;4MG	N70937	001	Sep 11, 1986
	25MG;2MG	N70936	001	Sep 11, 1986
	25MG;4MG	N70938	001	Sep 11, 1986
	50MG;4MG	N70939	001	Sep 12, 1986
MUTUAL PHARM	10MG;2MG	N71077	001	Nov 12, 1986
	10MG;4MG	N71078	001	Nov 12, 1986
	25MG;2MG	N70297	001	Nov 12, 1986
	25MG;4MG	N71079	001	Nov 12, 1986
	10MG;2MG	N70565	001	Sep 11, 1986
PAR PHARM	10MG;4MG	N70620	001	Sep 11, 1986
	25MG;2MG	N70621	001	Sep 11, 1986
	25MG;4MG	N70595	001	Sep 11, 1986
	50MG;4MG	N70574	001	Sep 11, 1986
	10MG;2MG	N70373	001	Aug 25, 1986
WATSON LABS	10MG;2MG	N72539	001	Feb 15, 1989
	10MG;4MG	N70375	001	Aug 25, 1986
	10MG;4MG	N72540	001	Feb 15, 1989
	25MG;2MG	N70374	001	Aug 25, 1986
	25MG;2MG	N72541	001	Feb 15, 1989
	25MG;4MG	N70376	001	Aug 25, 1986
	25MG;4MG	N72134	001	Feb 15, 1989
	50MG;4MG	N70377	001	Nov 04, 1986
	50MG;4MG	N71558	001	Mar 02, 1987
	50MG;4MG	N72135	001	Feb 15, 1989
TRIAVIL 2-10				
NEW RIVER	10MG;2MG	N14715	004	
TRIAVIL 2-25				
NEW RIVER	25MG;2MG	N14715	002	
TRIAVIL 4-10				
NEW RIVER	10MG;4MG	N14715	003	
TRIAVIL 4-25				
NEW RIVER	25MG;4MG	N14715	005	
TRIAVIL 4-50				
NEW RIVER	50MG;4MG	N14715	006	

AMLODIPINE MALEATE

TABLET; ORAL

AMVAZ

DR REDDYS LABS INC	2.5MG	N21435	001	Oct 31, 2003
	5MG	N21435	002	Oct 31, 2003
	10MG	N21435	003	Oct 31, 2003

AMMONIUM CHLORIDE

INJECTABLE; INJECTION

AMMONIUM CHLORIDE

ABBOTT	5MEQ/ML	N83130	001	
GD SEARLE LLC	3MEQ/ML	N86205	001	
AMMONIUM CHLORIDE 0.9% IN NORMAL SALINE				
MCGAW	900MG/100ML	N06580	001	
AMMONIUM CHLORIDE 2.14%				
B BRAUN	40MEQ/100ML	N85734	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 19 (of 263)

AMODIAQUINE HYDROCHLORIDE

TABLET; ORAL

CAMOQUIN HYDROCHLORIDE

PARKE DAVIS

EQ 200MG BASE

N06441 001

AMOXAPINE

TABLET; ORAL

ASENDIN

LEDERLE

25MG

N18021 001

50MG

N18021 002

100MG

N18021 003

150MG

N18021 004

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

LABS ATRAL

250MG

N62528 001

Aug 07, 1985

500MG

N62528 002

Aug 07, 1985

MYLAN

250MG

N62067 001

500MG

N62067 002

TEVA

250MG

N63030 001

Feb 28, 1989

500MG

N63031 001

Feb 28, 1989

AMOXIL

GLAXOSMITHKLINE

250MG

N50459 001

250MG

N62216 003

500MG

N50459 002

TRIMOX

APOTHECON

250MG

N62098 001

250MG

N62152 001

250MG

N63099 001

Mar 20, 1992

500MG

N62098 002

500MG

N62152 002

500MG

N63099 002

Mar 20, 1992

UTIMOX

PARKE DAVIS

250MG

N62107 001

500MG

N62107 002

WYMOX

WYETH AYERST

250MG

N62120 001

500MG

N62120 002

FOR SUSPENSION; ORAL

AMOXICILLIN

MYLAN

125MG/5ML

N62090 001

250MG/5ML

N62090 002

AMOXIL

GLAXOSMITHKLINE

50MG/ML

N50460 005

125MG/5ML

N50460 001

250MG/5ML

N50460 002

LAROTID

GLAXOSMITHKLINE

50MG/ML

N50460 006

POLYMOX

APOTHECON

125MG/5ML

N61851 001

125MG/5ML

N62323 001

250MG/5ML

N61851 002

250MG/5ML

N62323 002

TRIMOX

APOTHECON

125MG/5ML

N62099 001

125MG/5ML

N62154 001

250MG/5ML

N62099 002

250MG/5ML

N62154 002

UTIMOX

PARKE DAVIS

125MG/5ML

N62127 001

250MG/5ML

N62127 002

WYMOX

WYETH AYERST

125MG/5ML

N62131 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 20 (of 263)

AMOXICILLIN

FOR SUSPENSION; ORAL

WYMOX

WYETH AYERST

250MG/5ML

N62131 002

TABLET, CHEWABLE; ORAL

AMOXICILLIN

APOTHECON

125MG

N64131 001

May 06, 1996

250MG

N64131 002

May 06, 1996

AMOXIL

GLAXOSMITHKLINE

125MG

N50542 002

250MG

N50542 001

AMPHETAMINE ADIPATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE ADIPATE;
DEXTROAMPHETAMINE SULFATE

CAPSULE; ORAL

DELCOBESE

TEVA

1.25MG;1.25MG;1.25MG;1.25MG **Federal
Register determination that product
was not discontinued or withdrawn for
safety or efficacy reasons**

N83564 001

2.5MG;2.5MG;2.5MG;2.5MG **Federal
Register determination that product
was not discontinued or withdrawn for
safety or efficacy reasons**

N83564 002

3.75MG;3.75MG;3.75MG;3.75MG **Federal
Register determination that product
was not discontinued or withdrawn for
safety or efficacy reasons**

N83564 003

5MG;5MG;5MG;5MG **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N83564 004

TABLET; ORAL

DELCOBESE

TEVA

1.25MG;1.25MG;1.25MG;1.25MG **Federal
Register determination that product
was not discontinued or withdrawn for
safety or efficacy reasons**

N83563 004

2.5MG;2.5MG;2.5MG;2.5MG **Federal
Register determination that product
was not discontinued or withdrawn for
safety or efficacy reasons**

N83563 003

3.75MG;3.75MG;3.75MG;3.75MG **Federal
Register determination that product
was not discontinued or withdrawn for
safety or efficacy reasons**

N83563 002

5MG;5MG;5MG;5MG **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N83563 001

AMPHETAMINE RESIN COMPLEX; DEXTROAMPHETAMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL

BIPHETAMINE 12.5

UCB

EQ 6.25MG BASE;EQ 6.25MG BASE

N10093 007

BIPHETAMINE 20

UCB

EQ 10MG BASE;EQ 10MG BASE

N10093 003

BIPHETAMINE 7.5

UCB

EQ 3.75MG BASE;EQ 3.75MG BASE

N10093 009

AMPHETAMINE SULFATE

TABLET; ORAL

AMPHETAMINE SULFATE

LANNETT

5MG

N83901 001

Aug 31, 1984

10MG

N83901 002

Aug 31, 1984

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 21 (of 263)

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

ABBOTT

50MG/VIAL

N64141 001

Dec 23, 1996

AM PHARM PARTNERS

50MG/VIAL

N62728 001

Apr 13, 1987

OINTMENT; TOPICAL

FUNGIZONE

APOTHECON

3%

N50313 001

SUSPENSION; ORAL

FUNGIZONE

BRISTOL MYERS SQUIBB

100MG/ML

N50341 003

AMPICILLIN SODIUM

INJECTABLE; INJECTION

AMPICILLIN SODIUM

APOTHECON

EQ 125MG BASE/VIAL

N62860 001

Feb 05, 1988

EQ 250MG BASE/VIAL

N62860 002

Feb 05, 1988

EQ 500MG BASE/VIAL

N62860 003

Feb 05, 1988

EQ 1GM BASE/VIAL

N62860 004

Feb 05, 1988

EQ 2GM BASE/VIAL

N62860 005

Feb 05, 1988

BAXTER HLTHCARE

EQ 125MG BASE/VIAL

N62692 001

Jun 24, 1986

EQ 250MG BASE/VIAL

N62692 002

Jun 24, 1986

EQ 500MG BASE/VIAL

N62692 003

Jun 24, 1986

EQ 1GM BASE/VIAL

N62692 004

Jun 24, 1986

EQ 2GM BASE/VIAL

N62692 005

Jun 24, 1986

EQ 10GM BASE/VIAL

N62692 006

Jun 24, 1986

CONSOLIDATED PHARM

EQ 125MG BASE/VIAL

N61936 005

EQ 250MG BASE/VIAL

N61936 001

EQ 500MG BASE/VIAL

N61936 002

EQ 1GM BASE/VIAL

N61936 003

EQ 2GM BASE/VIAL

N61936 004

HANFORD GC

EQ 125MG BASE/VIAL

N63143 001

Apr 15, 1993

EQ 250MG BASE/VIAL

N63145 001

Apr 15, 1993

EQ 500MG BASE/VIAL

N63146 001

Apr 15, 1993

EQ 500MG BASE/VIAL

N63147 001

Apr 15, 1993

EQ 1GM BASE/VIAL

N62772 001

Apr 15, 1993

EQ 1GM BASE/VIAL

N63139 001

Apr 15, 1993

EQ 2GM BASE/VIAL

N63140 001

Apr 15, 1993

EQ 2GM BASE/VIAL

N63141 001

Apr 15, 1993

EQ 10GM BASE/VIAL

N63142 001

Apr 15, 1993

INTL MEDICATION

EQ 1GM BASE/VIAL

N62634 002

Jan 09, 1987

EQ 2GM BASE/VIAL

N62634 003

Jan 09, 1987

LILLY

EQ 500MG BASE/VIAL

N62565 001

Apr 04, 1985

EQ 1GM BASE/VIAL

N62565 002

Apr 04, 1985

EQ 2GM BASE/VIAL

N62565 003

Jun 24, 1986

MARSAM PHARMS LLC

EQ 125MG BASE/VIAL

N62816 001

Oct 24, 1988

EQ 250MG BASE/VIAL

N62816 002

Oct 24, 1988

EQ 500MG BASE/VIAL

N62816 003

Oct 24, 1988

EQ 1GM BASE/VIAL

N62816 004

Oct 24, 1988

EQ 2GM BASE/VIAL

N62816 005

Oct 24, 1988

EQ 10GM BASE/VIAL

N62994 001

Sep 15, 1988

OMNIPEN-N

WYETH AYERST

EQ 125MG BASE/VIAL

N60626 001

EQ 125MG BASE/VIAL

N62718 001

Dec 16, 1986

EQ 250MG BASE/VIAL

N60626 002

EQ 250MG BASE/VIAL

N62718 002

Dec 16, 1986

EQ 500MG BASE/VIAL

N60626 003

EQ 500MG BASE/VIAL

N62718 003

Dec 16, 1986

EQ 1GM BASE/VIAL

N60626 004

EQ 1GM BASE/VIAL

N62718 004

Dec 16, 1986

EQ 2GM BASE/VIAL

N60626 005

EQ 2GM BASE/VIAL

N62718 005

Dec 16, 1986

PENBRITIN-S

WYETH AYERST

EQ 125MG BASE/VIAL

N50072 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 22 (of 263)

AMPICILLIN SODIUMINJECTABLE; INJECTION
PENBRITIN-S

WYETH AYERST

EQ 250MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 4GM BASE/VIALN50072 002
N50072 003
N50072 004
N50072 005
N50072 006POLYCILLIN-N
BRISTOLEQ 125MG BASE/VIAL
EQ 250MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
EQ 2GM BASE/VIALN50309 001
N50309 002
N50309 003
N50309 004
N50309 005

TOTACILLIN-N

GLAXOSMITHKLINE

EQ 125MG BASE/VIAL
EQ 250MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 10GM BASE/VIALN60677 001
N60677 002
N60677 003
N60677 004
N62727 001
N60677 005
N62727 002
N60677 006

Dec 19, 1986

Dec 19, 1986

AMPICILLIN SODIUM; SULBACTAM SODIUMINJECTABLE; INJECTION
UNASYN

PFIZER

EQ 500MG BASE/VIAL;EQ 250MG BASE/VIAL

N50608 003

Dec 31, 1986

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMCILL

PARKE DAVIS

EQ 250MG BASE
EQ 500MG BASEN62041 001
N62041 002

AMPICILLIN

LEDERLE

EQ 250MG BASE
EQ 500MG BASEN62208 001
N62208 002

VITARINE

EQ 250MG BASE
EQ 500MG BASEN61387 001
N61387 003

AMPICILLIN TRIHYDRATE

BIOCHEMIE

EQ 250MG BASE
EQ 500MG BASEN64082 001
N64082 002

Aug 29, 1995

Aug 29, 1995

CONSOLIDATED PHARM

EQ 250MG BASE
EQ 500MG BASEN61602 001
N61602 002

IVAX PHARMS

EQ 250MG BASE
EQ 500MG BASEN60765 001
N60765 002

MYLAN

EQ 250MG BASE
EQ 500MG BASEN61755 001
N61755 002

PUREPAC PHARM

EQ 250MG BASE
EQ 500MG BASEN61853 001
N61853 002

TEVA

EQ 250MG BASE
EQ 500MG BASEN61502 001
N61502 002

OMNIPEN (AMPICILLIN)

WYETH AYERST

250MG
500MGN60624 001
N60624 002

PENBRITIN

WYETH AYERST

EQ 250MG BASE
EQ 500MG BASEN60908 001
N60908 002

PFIZERPEN-A

PFIZER

EQ 250MG BASE
EQ 500MG BASEN62050 001
N62050 002

POLYCILLIN

BRISTOL

EQ 250MG BASE
EQ 500MG BASEN50310 001
N50310 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 23 (of 263)

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

PRINCIPEN '250'

APOTHECON

EQ 250MG BASE

N50056 001

EQ 250MG BASE

N62157 002

PRINCIPEN '500'

APOTHECON

EQ 500MG BASE

N50056 002

EQ 500MG BASE

N62157 001

TOTACILLIN

GLAXOSMITHKLINE

EQ 250MG BASE

N60060 001

EQ 250MG BASE

N62212 001

EQ 500MG BASE

N60060 002

EQ 500MG BASE

N62212 002

FOR SUSPENSION; ORAL

AMCILL

PARKE DAVIS

EQ 125MG BASE/5ML

N62030 001

EQ 250MG BASE/5ML

N62030 002

AMPICILLIN TRIHYDRATE

CONSOLIDATED PHARM

EQ 125MG BASE/5ML

N61601 001

EQ 250MG BASE/5ML

N61601 002

MYLAN

EQ 125MG BASE/5ML

N61829 002

EQ 250MG BASE/5ML

N61829 001

PUREPAC PHARM

EQ 125MG BASE/5ML

N61980 001

TEVA

EQ 125MG BASE/5ML

N61370 001

EQ 250MG BASE/5ML

N61370 002

OMNIPEN (AMPICILLIN)

WYETH AYERST

100MG/ML

N60625 001

125MG/5ML

N60625 002

250MG/5ML

N60625 003

500MG/5ML

N60625 004

PENBRITIN

WYETH AYERST

EQ 100MG BASE/ML

N50019 001

EQ 125MG BASE/5ML

N50019 002

EQ 250MG BASE/5ML

N50019 003

PFIZERPEN-A

PFIZER

EQ 125MG BASE/5ML

N62049 001

EQ 250MG BASE/5ML

N62049 002

POLYCILLIN

APOTHECON

EQ 125MG BASE/5ML

N62297 001

EQ 250MG BASE/5ML

N62297 002

BRISTOL

EQ 100MG BASE/ML

N50308 004

EQ 125MG BASE/5ML

N50308 001

EQ 250MG BASE/5ML

N50308 002

EQ 500MG BASE/5ML

N50308 003

PRINCIPEN '125'

APOTHECON

EQ 125MG BASE/5ML

N60127 002

EQ 125MG BASE/5ML

N62151 001

PRINCIPEN '250'

APOTHECON

EQ 250MG BASE/5ML

N60127 001

EQ 250MG BASE/5ML

N62151 002

TOTACILLIN

GLAXOSMITHKLINE

EQ 125MG BASE/5ML

N60666 001

EQ 125MG BASE/5ML

N62223 001

EQ 250MG BASE/5ML

N60666 002

EQ 250MG BASE/5ML

N62223 002

TABLET, CHEWABLE; ORAL

POLYCILLIN

BRISTOL

EQ 125MG BASE

N50093 001

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

CAPSULE; ORAL

PRINCIPEN W/ PROBENECID

APOTHECON

EQ 389MG BASE;111MG

N50488 001

EQ 389MG BASE;111MG

N62150 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 24 (of 263)

AMPICILLIN/AMPICILLIN TRIHYDRATE; PROBENECID

FOR SUSPENSION; ORAL

POLYCILLIN-PRB

APOTHECON

EQ 3.5GM BASE/BOT;1GM/BOT

N61898 001

BRISTOL

EQ 3.5GM BASE/BOT;1GM/BOT

N50457 001

PROBAMPACIN

TEVA

EQ 3.5GM BASE/BOT;1GM/BOT

N61741 001

AMPRENAVIR

CAPSULE; ORAL

AGENERASE

GLAXOSMITHKLINE

150MG

N21007 002

Apr 15, 1999

ANILERIDINE HYDROCHLORIDE

TABLET; ORAL

LERITINE

MERCK

EQ 25MG BASE

N10585 002

ANILERIDINE PHOSPHATE

INJECTABLE; INJECTION

LERITINE

MERCK

25MG/ML

N10520 003

ANISOTROPINE METHYLBROMIDE

TABLET; ORAL

ANISOTROPINE METHYLBROMIDE

WATSON LABS

50MG

N86046 001

VALPIN 50

ENDO PHARMS

50MG

N13428 001

ARBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

GENESA

GENSIA AUTOMEDICS

0.05MG/ML

N20420 001

Sep 12, 1997

ARDEPARIN SODIUM

INJECTABLE; INJECTION

NORMIFLO

PHARMACIA AND UPJOHN

5,000 UNITS/0.5ML **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N20227 002

May 23, 1997

10,000 UNITS/0.5ML **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N20227 001

May 23, 1997

ARIPIRAZOLE

TABLET; ORAL

ABILIFY

OTSUKA

2MG

N21436 006

Nov 15, 2002

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID;
NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE
HYDROCHLORIDE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION

BEROCCA PN

ROCHE

50MG/ML;0.03MG/ML;0.0025MG/ML;7.5MG/ML;
100
IU/ML;0.2MG/ML;20MG/ML;2MG/ML;1.8MG/ML;
1.5MG/ML;1,650 IU/ML;5 IU/ML

N06071 003

Oct 10, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 25 (of 263)

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.C. 9+3

AM PHARM PARTNERS	10MG/ML;0.006MG/ML;0.5UGM/ML;1.5MG/ML;20 IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/ML;0.3MG/ML;330 UNITS/ML;1 IU/ML	N18440 002	Aug 08, 1985
-------------------	--	------------	--------------

M.V.I. -12

MAYNE PHARMA USA	10MG/ML;0.006MG/ML;0.5UGM/ML;1.5MG/ML;20 IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/ML;0.3MG/ML;330 UNITS/ML;1 IU/ML	N08809 004	Aug 08, 1985
------------------	--	------------	--------------

MVC PLUS

STERIS	10MG/ML;0.006MG/ML;0.5UGM/ML;1.5MG/ML;20 IU/ML;0.04MG/ML;4MG/ML;0.4MG/ML;0.36MG/ML;0.3MG/ML;330 UNITS/ML;1 IU/ML	N18439 002	Aug 08, 1985
--------	--	------------	--------------

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I. -12

MAYNE PHARMA USA	20MG/ML;0.006MG/ML;0.5UGM/ML;1.5MG/ML;20 IU/ML;0.6MG/ML;4MG/ML;0.4MG/ML;0.36MG/ML;0.6MG/ML;330 UNITS/ML;1 IU/ML	N08809 005	Apr 22, 2004
------------------	---	------------	--------------

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE; VITAMIN A; VITAMIN E

INJECTABLE; INJECTION

M.V.I. -12 LYOPHILIZED

ASTRAZENECA	100MG/VIAL;0.06MG/VIAL;0.005MG/VIAL;15MG/VIAL;5UGM/VIAL;0.4MG/VIAL;40MG/VIAL;4MG/VIAL;3.6MG/VIAL;3MG/VIAL;1MG/VIAL;10MG/VIAL	N18933 002	Aug 08, 1985
-------------	--	------------	--------------

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION

VITAPED

HOSPIRA	N/A, 80MG/VIAL; N/A, 0.02MG/VIAL; N/A, 0.001 MG/VIAL; 400 IU/10ML, N/A; N/A, 0.14MG/VIAL; N/A, 17MG/VIAL; N/A, 5MG/VIAL; 0.2MG/10ML, N/A; N/A, 1MG/VIAL; N/A, 1.4MG/VIAL; N/A, 1.2MG/VIAL; EQ 2, 300 UNITS BASE/10ML, N/A; 7 IU/10ML, N/A	N20176 001	Dec 29, 1993
---------	---	------------	--------------

ASPIRIN

TABLET, EXTENDED RELEASE; ORAL

8-HOUR BAYER

BAYER	650MG	N16030 001
-------	-------	------------

MEASURIN

BAYER	650MG	N16030 002
-------	-------	------------

ASPIRIN; BUTALBITAL

TABLET; ORAL

AXOTAL

SAVAGE LABS	650MG; 50MG	N88305 001	Oct 13, 1983
-------------	-------------	------------	--------------

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

WATSON LABS	325MG; 50MG; 40MG	N86231 002	Feb 12, 1985
-------------	-------------------	------------	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 26 (of 263)

ASPIRIN; BUTALBITAL; CAFFEINE

TABLET; ORAL

BUTALBITAL ASPIRIN AND CAFFEINE

QUANTUM PHARMICS	325MG;50MG;40MG	N88972	001	Jun 18, 1985
BUTALBITAL COMPOUND				
IVAX PHARMS	325MG;50MG;40MG	N85441	002	Oct 31, 1984
BUTALBITAL W/ ASPIRIN & CAFFEINE				
PHARMERAL	325MG;50MG;40MG	N87048	002	Dec 09, 1983
BUTALBITAL, ASPIRIN AND CAFFEINE				
HALSEY	325MG;50MG;40MG	N89448	001	Dec 01, 1986
WATSON LABS	325MG;50MG;40MG	N86237	002	Mar 23, 1984
FIORINAL				
WATSON PHARMS	325MG;50MG;40MG	N17534	003	Apr 16, 1986
LANORINAL				
LANNETT	325MG;50MG;40MG	N86986	002	Oct 18, 1985

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

COMPOUND 65

ALRA	389MG;32.4MG;65MG	N84553	002	Aug 17, 1983
DARVON COMPOUND				
XANODYNE PHARM	389MG;32.4MG;32MG	N10996	006	Mar 08, 1983
PROPOXYPHENE COMPOUND 65				
IVAX PHARMS	389MG;32.4MG;65MG	N83077	002	Dec 07, 1984
SANDOZ	389MG;32.4MG;65MG	N80044	002	Sep 16, 1983
PROPOXYPHENE COMPOUND-65				
SANDOZ	389MG;32.4MG;65MG	N83101	002	Jun 24, 1985
PROPOXYPHENE HYDROCHLORIDE W/ ASPIRIN AND CAFFEINE				
WATSON LABS	389MG;32.4MG;65MG	N85732	002	Sep 03, 1984

ASPIRIN; CARISOPRODOL

TABLET; ORAL

CARISOPRODOL COMPOUND

WATSON LABS	325MG;200MG	N88809	001	Oct 03, 1985
-------------	-------------	--------	-----	--------------

ASPIRIN; HYDROCODONE BITARTRATE

TABLET; ORAL

VICOPRIN

ABBOTT	500MG;5MG	N86333	001	Sep 14, 1983
--------	-----------	--------	-----	--------------

ASPIRIN; MEPROBAMATE

TABLET; ORAL

MEPRO-ASPIRIN

SANDOZ	325MG;200MG	N89127	001	Mar 02, 1987
MEPROBAMATE AND ASPIRIN				
PAR PHARM	325MG;200MG	N89126	001	Aug 19, 1986
MICRAININ				
MEDPOINTE PHARM HLC	325MG;200MG	N84978	001	
Q-GESIC				
QUANTUM PHARMICS	325MG;200MG	N88740	001	Jun 01, 1984

ASPIRIN; METHOCARBAMOL

TABLET; ORAL

METHOCARBAMOL AND ASPIRIN

MCNEIL	325MG;400MG	N89193	001	Feb 12, 1986
ROBAXISAL				
ROBINS AH	325MG;400MG	N12281	001	

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

CODOXY

HALSEY	325MG;4.5MG;0.38MG	N87464	001	Jul 01, 1982
--------	--------------------	--------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 27 (of 263)

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL			
OXYCODONE AND ASPIRIN (HALF-STRENGTH)			
ROXANE	325MG;2.25MG;0.19MG	N87742 001	Jun 04, 1982
ROXIPRIN			
ROXANE	325MG;4.5MG;0.38MG	N87743 001	Jun 04, 1982

ASPIRIN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL			
TALWIN COMPOUND			
SANOFI SYNTHELABO	325MG;EQ 12.5MG BASE	N16891 001	

ASPIRIN; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL			
DARVON W/ ASA			
XANODYNE PHARM	325MG;65MG	N10996 005	

ASPIRIN; PROPOXYPHENE NAPSYLATE

CAPSULE; ORAL			
DARVON-N W/ ASA			
AAIPHARMA LLC	325MG;100MG	N16829 001	
TABLET; ORAL			
DARVON-N W/ ASA			
AAIPHARMA LLC	325MG;100MG	N16863 001	

ATENOLOL

TABLET; ORAL			
ATENOLOL			
ABLE	25MG	N76907 001	Jul 30, 2004
	50MG	N76907 002	Jul 30, 2004
	100MG	N76907 003	Jul 30, 2004
APOTHECON			
	50MG	N73317 001	Mar 20, 1992
	100MG	N73318 001	Mar 20, 1992

ATOMOXETINE HYDROCHLORIDE

CAPSULE; ORAL			
STRATTERA			
LILLY	5MG	N21411 001	Nov 26, 2002

ATRACURIUM BESYLATE

INJECTABLE; INJECTION			
ATRACURIUM BESYLATE PRESERVATIVE FREE			
BAXTER HLTHCARE	10MG/ML	N74825 001	Sep 30, 1997

ATROPINE

INJECTABLE; INJECTION			
ATROPINE			
SOLVAY	EQ 2MG SULFATE/0.7ML	N71295 001	Jan 30, 1987

ATROPINE SULFATE

AEROSOL, METERED; INHALATION			
ATROPINE SULFATE			
US ARMY	EQ 0.36MG BASE/INH	N20056 001	Sep 19, 1990

ATROPINE SULFATE; DIFENOXIN HYDROCHLORIDE

TABLET; ORAL			
MOTOFEN HALF-STRENGTH			
VALEANT	0.025MG;0.5MG	N17744 001	

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL			
COLONALID			
MEDPOINTE PHARM HLC	0.025MG/5ML;2.5MG/5ML	N85735 001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 28 (of 263)

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

SOLUTION; ORAL			
LOMANATE			
ALPHARMA US PHARMS	0.025MG/5ML;2.5MG/5ML	N85746	001
TABLET; ORAL			
COLONAIID			
MEDPOINTE PHARM HLC	0.025MG;2.5MG	N85737	001
DI-ATRO			
MD PHARM	0.025MG;2.5MG	N85266	001
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE			
ABLE	0.025MG;2.5MG	N40395	001 Nov 27, 2000
ASCOT	0.025MG;2.5MG	N87934	001 Jul 19, 1983
HEATHER	0.025MG;2.5MG	N86798	001
INWOOD LABS	0.025MG;2.5MG	N85509	001
LEDERLE	0.025MG;2.5MG	N86950	001
MUTUAL PHARM	0.025MG;2.5MG	N85506	001
PARKE DAVIS	0.025MG;2.5MG	N87131	001
R AND S PHARMA	0.025MG;2.5MG	N85035	001
ROXANE	0.025MG;2.5MG	N86057	001
WATSON LABS	0.025MG;2.5MG	N85876	001
WEST WARD	0.025MG;2.5MG	N87765	001 Mar 15, 1982
DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE			
KV PHARM	0.025MG;2.5MG	N85659	001
PVT FORM	0.025MG;2.5MG	N85766	001
USL PHARMA	0.025MG;2.5MG	N87842	001 Mar 29, 1982
VALEANT PHARM INTL	0.025MG;2.5MG	N87195	001 Feb 16, 1982
LOGEN			
SUPERPHARM	0.025MG;2.5MG	N88962	001 May 10, 1985
LO-TROL			
VANGARD	0.025MG;2.5MG	N88009	001 Mar 25, 1983
LOW-QUEL			
HALSEY	0.025MG;2.5MG	N85211	001

ATROPINE SULFATE; MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION			
ATROPINE AND DEMEROL			
ABBOTT	0.4MG/ML;50MG/ML	N87853	001 Nov 26, 1982
	0.4MG/ML;75MG/ML	N87847	001 Nov 26, 1982
	0.4MG/ML;100MG/ML	N87848	001 Nov 26, 1982
MEPERIDINE AND ATROPINE SULFATE			
WYETH AYERST	0.4MG/ML;50MG/ML	N85121	001
	0.4MG/ML;75MG/ML	N85121	002
	0.4MG/ML;100MG/ML	N85121	003

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR			
ATNAA			
US ARMY	2.1MG/0.7ML,N/A;N/A,600MG/2ML	N21175	001 Jan 17, 2002

AZATADINE MALEATE

TABLET; ORAL			
OPTIMINE			
SCHERING	1MG	N17601	001

AZATADINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL			
TRINALIN			
SCHERING	1MG;120MG	N18506	001 Mar 23, 1982

AZATHIOPRINE

TABLET; ORAL			
IMURAN			
PROMETHEUS LABS	25MG	N16324	002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 29 (of 263)

AZATHIOPRINE SODIUMINJECTABLE; INJECTION
IMURAN

PROMETHEUS LABS EQ 100MG BASE/VIAL N17391 001

AZITHROMYCINCAPSULE; ORAL
ZITHROMAXPFIZER EQ 250MG BASE **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons** N50670 001 Nov 01, 1991AZITHROMYCIN DIHYDRATE; TROVAFLOXACIN MESYLATEFOR SUSPENSION, TABLET; ORAL
TROVAN/ZITHROMAX COMPLIANCE PAK

PFIZER EQ 1GM BASE, N/A; N/A, EQ 100MG BASE N50762 001 Dec 18, 1998

AZLOCILLIN SODIUMINJECTABLE; INJECTION
AZLIN

BAYER PHARMS	EQ 2GM BASE/VIAL	N50562	001	Sep 03, 1982
	EQ 2GM BASE/VIAL	N62388	001	Sep 08, 1982
	EQ 2GM BASE/VIAL	N62417	001	Oct 12, 1982
	EQ 3GM BASE/VIAL	N50562	002	Sep 03, 1982
	EQ 3GM BASE/VIAL	N62388	002	Sep 08, 1982
	EQ 3GM BASE/VIAL	N62417	002	Oct 12, 1982
	EQ 4GM BASE/VIAL	N50562	003	Sep 03, 1982
	EQ 4GM BASE/VIAL	N62388	003	Sep 08, 1982
	EQ 4GM BASE/VIAL	N62417	003	Oct 12, 1982

AZTREONAMINJECTABLE; INJECTION
AZACTAM IN PLASTIC CONTAINER

BRISTOL MYERS SQUIBB 10MG/ML N50632 003 May 24, 1989

BACAMPICILLIN HYDROCHLORIDETABLET; ORAL
SPECTROBID

PFIZER 800MG N50520 002 Sep 12, 1983

BACITRACININJECTABLE; INJECTION
BACITRACIN

PFIZER 50,000 UNITS/VIAL N60282 001

OINTMENT; OPHTHALMIC
BACIGUENT

PHARMACIA AND UPJOHN 500 UNITS/GM N60734 001

BACITRACIN

LILLY 500 UNITS/GM N60687 001

PHARMADERM 500 UNITS/GM N62158 001

PHARMAFAIR 500 UNITS/GM N62453 001 Mar 28, 1984

OINTMENT; TOPICAL
BACITRACIN

COMBE 500 UNITS/GM N62799 001 May 14, 1987

NASKA 500 UNITS/GM N62857 001 Nov 13, 1987

POWDER; FOR RX COMPOUNDING

BACITRACIN

PADDOCK 5,000,000 UNITS/BOT N62456 001 Jul 27, 1983

BACITRACIN ZINCPOWDER; FOR RX COMPOUNDING
ZIBA-RX

PHARMA TEK 500,000 UNITS/BOT N61737 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 30 (of 263)

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC				
ZINC BACITRACIN, NEOMYCIN SULFATE, POLYMYXIN B SULFATE & HYDROCORTISONE				
PHARMAFAIR	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	N62389	001	Jul 02, 1982
OINTMENT; TOPICAL				
NEOMYCIN & POLYMYXIN B SULFATES & BACITRACIN ZINC & HYDROCORTISONE				
PHARMAFAIR	400 UNITS/GM;1%;EQ 3.5MG UNITS/GM	N62381	001	Sep 06, 1985

BACITRACIN ZINC; LIDOCAINE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; TOPICAL				
LANABIOTIC				
COMBE	400 UNITS/GM;40MG/GM;EQ 5MG BASE/GM;5,000 UNITS/GM	N62499	001	Jun 03, 1985

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC				
BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE				
PHARMAFAIR	400 UNITS/GM;EQ 3.5MG UNITS/GM	N62386	001	Sep 09, 1982
BACITRACIN-NEOMYCIN-POLYMYXIN				
PHARMADERM	400 UNITS/GM;EQ 3.5MG UNITS/GM	N62167	001	
NEO-POLYCIN				
DOW PHARM	500 UNITS/GM;EQ 3.5MG UNITS/GM	N60647	001	
OINTMENT; TOPICAL				
BACITRACIN ZINC-NEOMYCIN SULFATE-POLYMYXIN B SULFATE				
NASKA	400 UNITS/GM;EQ 3.5MG UNITS/GM	N62833	001	Nov 09, 1987

BACITRACIN ZINC; POLYMYXIN B SULFATE

AEROSOL; TOPICAL				
POLYSPORIN				
GLAXOSMITHKLINE	10,000 UNITS/GM;2,000,000 UNITS/GM	N50167	002	Mar 01, 1985
OINTMENT; OPHTHALMIC				
OCUMYCIN				
PHARMAFAIR	500 UNITS/GM;10,000 UNITS/GM	N62430	001	Apr 08, 1983
OINTMENT; TOPICAL				
BACITRACIN ZINC-POLYMYXIN B SULFATE				
NASKA	500 UNITS/GM;10,000 UNITS/GM	N62849	001	Nov 13, 1987

BACITRACIN; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC				
BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE				
ALTANA	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	N60731	002	

BACITRACIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC				
MYCITRACIN				
PHARMACIA AND UPJOHN	500 UNITS/GM;EQ 3.5MG UNITS/GM	N61048	001	

BACITRACIN; POLYMYXIN B SULFATE

DISC; TOPICAL				
LANABIOTIC				
COMBE	500 UNITS/GM;5,000 UNITS/GM	N50598	001	Sep 22, 1986

BACLOFEN

TABLET; ORAL				
BACLOFEN				
TEVA	10MG	N73043	001	Feb 27, 1992
	20MG	N73044	001	Feb 27, 1992
USL PHARMA	10MG	N71260	001	May 06, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 31 (of 263)

BACLOFENTABLET; ORAL
BACLOFEN

USL PHARMA	20MG	N71261	001	May 06, 1988
WATSON LABS	10MG	N74698	001	Aug 20, 1996
	20MG	N73093	001	Jan 28, 1994
	20MG	N74698	002	Aug 20, 1996
LIORESAL				
NOVARTIS	10MG	N17851	001	
	20MG	N17851	003	Jan 20, 1982

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

BECLOVENT

GLAXOSMITHKLINE 0.042MG/INH N18153 001

VANCERIL

SCHERING 0.042MG/INH N17573 001

VANCERIL DOUBLE STRENGTH

SCHERING 0.084MG/INH N20486 001 Dec 24, 1996

AEROSOL, METERED; NASAL

BECONASE

GLAXOSMITHKLINE 0.042MG/INH N18584 001

VANCENASE

SCHERING 0.042MG/INH N18521 001

BECLOMETHASONE DIPROPIONATE MONOHYDRATE

SPRAY, METERED; NASAL

VANCENASE AQ

SCHERING EQ 0.042MG DIPROP/SPRAY N19589 001 Dec 23, 1987

EQ 0.084MG DIPROP/SPRAY N20469 001 Jun 26, 1996

BENDROFLUMETHIAZIDE

TABLET; ORAL

NATURETIN-10

APOTHECON 10MG N12164 003

NATURETIN-2.5

APOTHECON 2.5MG N12164 001

BENOXINATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BENOXINATE HYDROCHLORIDE

SOLA BARNES HIND 0.4% N84149 001

BENTIROMIDE

SOLUTION; ORAL

CHYMEX

SAVAGE LABS 500MG/7.5ML N18366 001 Dec 29, 1983

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

DIDREX

PHARMACIA AND UPJOHN 25MG N12427 003

BENZQUINAMIDE HYDROCHLORIDE

SUPPOSITORY; RECTAL

EMETE-CON

ROERIG EQ 100MG BASE N16818 006

BENZTHIAZIDE

TABLET; ORAL

AQUATAG

SOLVAY 25MG N16001 001

50MG N16001 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 32 (of 263)

BENZTHIAZIDE

TABLET; ORAL		
BENZTHIAZIDE		
PVT FORM	50MG	N83206 001
EXNA		
AH ROBINS INC	50MG	N12489 001
FOVANE		
PFIZER	50MG	N12128 002
URESE		
PFIZER	25MG	N12128 003

BENZTROPINE MESYLATE

TABLET; ORAL			
BENZTROPINE MESYLATE			
QUANTUM PHARMICS	0.5MG	N88514 001	Jan 31, 1984
	1MG	N88510 001	Jan 31, 1984
	2MG	N88511 001	Jan 31, 1984
USL PHARMA	0.5MG	N89211 001	Jun 14, 1988
	1MG	N89212 001	Jun 14, 1988
	2MG	N89213 001	Jun 14, 1988
COGENTIN			
MERCK	0.5MG	N09193 004	
	1MG	N09193 003	
	2MG	N09193 002	

BENZYL BENZOATE

EMULSION; TOPICAL		
BENZYL BENZOATE		
LANNETT	50%	N84535 001

BENZYL PENICILLOYL-POLYLYSINE

INJECTABLE; INJECTION		
PRE-PEN		
ALLERQUEST	60UMOLAR	N50114 001

BEPRIDIL HYDROCHLORIDE

TABLET; ORAL			
BEPADIN			
MEDPOINTE PHARM HLC	200MG	N19001 001	Dec 28, 1990
	300MG	N19001 002	Dec 28, 1990
	400MG	N19001 003	Dec 28, 1990
VASCOR			
JOHNSON AND JOHNSON	200MG	N19002 001	Dec 28, 1990
	300MG	N19002 002	Dec 28, 1990
	400MG	N19002 003	Dec 28, 1990

BETA-CAROTENE

CAPSULE; ORAL		
SOLATENE		
ROCHE	30MG	N17589 001

BETAMETHASONE

CREAM; TOPICAL		
CELESTONE		
SCHERING	0.2%	N14762 001
TABLET; ORAL		
CELESTONE		
SCHERING	0.6MG	N12657 003

BETAMETHASONE BENZOATE

CREAM; TOPICAL		
UTICORT		
PARKE DAVIS	0.025%	N16998 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 33 (of 263)

BETAMETHASONE BENZOATE

GEL; TOPICAL				
UTICORT				
PARKE DAVIS	0.025%	N17244	001	
LOTION; TOPICAL				
UTICORT				
PARKE DAVIS	0.025%	N17528	001	
OINTMENT; TOPICAL				
UTICORT				
PARKE DAVIS	0.025%	N18089	001	

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL				
BETAMETHASONE DIPROPIONATE				
CLAY PARK	EQ 0.05% BASE	N72536	001	Jan 31, 1990
	EQ 0.05% BASE	N74579	001	Nov 26, 1997
PHARMADERM	EQ 0.05% BASE	N19136	001	Jun 26, 1984
DIPROSONE				
SCHERING	EQ 0.05% BASE	N17536	001	
CREAM, AUGMENTED; TOPICAL				
DIPROLENE				
SCHERING	EQ 0.05% BASE	N19408	001	Jan 31, 1986
DISC; TOPICAL				
DIPROSONE				
SCHERING	EQ 0.1% BASE	N17829	001	
LOTION; TOPICAL				
BETAMETHASONE DIPROPIONATE				
ALPHARMA US PHARMS	EQ 0.05% BASE	N71085	001	Feb 03, 1987
PHARMADERM	EQ 0.05% BASE	N70274	001	Aug 12, 1985
DIPROSONE				
SCHERING	EQ 0.05% BASE	N17781	001	
OINTMENT; TOPICAL				
ALPHATREX				
SAVAGE LABS	EQ 0.05% BASE	N19143	001	Sep 04, 1984
BETAMETHASONE DIPROPIONATE				
CLAY PARK	EQ 0.05% BASE	N72526	001	Jan 31, 1990
PHARMADERM	EQ 0.05% BASE	N19140	001	Sep 04, 1984
DIPROSONE				
SCHERING	EQ 0.05% BASE	N17691	001	

BETAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION				
BETAMETHASONE SODIUM PHOSPHATE				
STERIS	EQ 3MG BASE/ML	N85738	001	
CELESTONE				
SCHERING	EQ 3MG BASE/ML **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N17561	001	

BETAMETHASONE VALERATE

CREAM; TOPICAL				
BETADERM				
ROACO	EQ 0.1% BASE	N18839	001	Jun 30, 1983
BETAMETHASONE VALERATE				
CLAY PARK	EQ 0.1% BASE	N70053	001	Jun 10, 1986
PHARMADERM	EQ 0.1% BASE	N18860	002	Aug 31, 1983
PHARMAFAIR	EQ 0.1% BASE	N70485	001	May 29, 1987
BETATREX				
SAVAGE LABS	EQ 0.1% BASE	N18862	001	Aug 31, 1983
VALISONE				
SCHERING	EQ 0.01% BASE	N16322	002	
	EQ 0.1% BASE	N16322	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 34 (of 263)

BETAMETHASONE VALERATE

LOTION; TOPICAL

BETAMETHASONE VALERATE

PHARMADERM EQ 0.1% BASE

N18870 001 Aug 31, 1983

PHARMAFAIR EQ 0.1% BASE

N70484 001 May 29, 1987

BETATREX

SAVAGE LABS EQ 0.1% BASE

N18867 001 Aug 31, 1983

VALISONE

SCHERING EQ 0.1% BASE

N16932 001

OINTMENT; TOPICAL

BETAMETHASONE VALERATE

CLAY PARK EQ 0.1% BASE

N71478 001 Dec 23, 1987

PHARMADERM EQ 0.1% BASE

N18864 001 Aug 31, 1983

PHARMAFAIR EQ 0.1% BASE

N70486 001 May 29, 1987

BETATREX

SAVAGE LABS EQ 0.1% BASE

N18863 001 Aug 31, 1983

VALISONE

SCHERING EQ 0.1% BASE

N16740 001

BETAXOLOL HYDROCHLORIDE; CHLORTHALIDONE

TABLET; ORAL

KERLEDEX

LOREX 5MG;12.5MG

N19807 001 Oct 30, 1992

10MG;12.5MG

N19807 002 Oct 30, 1992

BETAXOLOL HYDROCHLORIDE; PILOCARPINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETOPTIC PILO

ALCON EQ 0.25% BASE;1.75%

N20619 001 Apr 17, 1997

BETAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

HISTALOG

LILLY 50MG/ML

N09344 001

BETHANECHOL CHLORIDE

INJECTABLE; INJECTION

URECHOLINE

ODYSSEY PHARMS 5MG/ML **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N06536 001

TABLET; ORAL

BETHANECHOL CHLORIDE

ABLE

5MG

N40492 001 Jul 27, 2004

10MG

N40483 001 Jul 27, 2004

25MG

N40485 001 Jul 27, 2004

50MG

N40509 001 Jul 27, 2004

ASCOT

10MG

N88288 001 Jun 08, 1983

25MG

N88289 001 Jun 08, 1983

IVAX PHARMS

25MG

N84689 001

LANNETT

5MG

N84702 001

10MG

N84712 001

25MG

N84074 001

SANDOZ

5MG

N84353 001

10MG

N84378 001

10MG

N84379 001

25MG

N84383 001

25MG

N84384 001

WATSON LABS

5MG

N84402 001

5MG

N85230 002

5MG

N85841 001

10MG

N84408 001

10MG

N85228 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 35 (of 263)

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

WATSON LABS

10MG

N85842 001

25MG

N84441 001

25MG

N85229 001

25MG

N85839 001

50MG

N87397 001

50MG

N87444 001

MYOTONACHOL

GLENWOOD

5MG

N84188 001

10MG

N84188 003

25MG

N84188 004

URECHOLINE

ODYSSEY PHARMS

5MG

N06536 003

10MG

N06536 002

25MG

N06536 004

50MG

N06536 005

BETHANIDINE SULFATE

TABLET; ORAL

TENATHAN

ROBINS AH

10MG

N17675 001

25MG

N17675 002

BIPERIDEN LACTATE

INJECTABLE; INJECTION

AKINETON

ABBOTT

5MG/ML

N12418 002

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

APOTHECON

2.5MG;6.25MG

N75642 002

Dec 27, 2000

5MG;6.25MG

N75642 001

Dec 27, 2000

10MG;6.25MG

N75642 003

Dec 27, 2000

IVAX PHARMS

2.5MG;6.25MG

N75632 001

Sep 27, 2000

5MG;6.25MG

N75632 002

Sep 27, 2000

10MG;6.25MG

N75632 003

Sep 27, 2000

BITOLTEROL MESYLATE

AEROSOL, METERED; INHALATION

TORNALATE

SANOFI SYNTHELABO

0.37MG/INH

N18770 001

Dec 28, 1984

SOLUTION; INHALATION

TORNALATE

SANOFI SYNTHELABO

0.2%

N19548 001

Feb 19, 1992

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE

AM PHARM PARTNERS

50MG/ML

N70134 001

Apr 29, 1986

100MG/ML

N71298 001

Feb 13, 1987

ASTRAZENECA

50MG/ML

N71151 001

Aug 10, 1987

50MG/ML

N71152 001

Aug 10, 1987

50MG/ML

N71153 001

Aug 10, 1987

BAXTER HLTHCARE

50MG/ML

N70545 001

May 14, 1986

50MG/ML

N70546 001

May 14, 1986

HOSPIRA

50MG/ML

N19033 001

Apr 29, 1986

BRETYLIUM TOSYLATE IN DEXTROSE 5%

ABBOTT

200MG/100ML

N19005 002

Apr 29, 1986

400MG/100ML

N19005 003

Apr 29, 1986

800MG/100ML

N19005 001

Apr 29, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 36 (of 263)

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION

BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE	200MG/100ML	N19837	002	Apr 12, 1989
	400MG/100ML	N19837	001	Apr 12, 1989
HOSPIRA	800MG/100ML	N19008	001	Apr 29, 1986
BRETYL0L				
MAYNE PHARMA USA	50MG/ML	N17954	001	

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN

ALLERGAN	0.2%	N20613	001	Sep 06, 1996
	0.5%	N20490	001	Mar 13, 1997

BROMOCRIPTINE MESYLATE

CAPSULE; ORAL

BROMOCRIPTINE MESYLATE

LEK PHARM	EQ 5MG BASE	N75100	001	Dec 10, 1998
-----------	-------------	--------	-----	--------------

BROMODIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

AMBODRYL

PARKE DAVIS	25MG	N07984	001	
-------------	------	--------	-----	--

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP; ORAL

BROMANYL

ALPHARMA US PHARMS	12.5MG/5ML;10MG/5ML	N88343	001	Aug 15, 1984
--------------------	---------------------	--------	-----	--------------

BROMPHENIRAMINE MALEATE

ELIXIR; ORAL

BROMPHENIRAMINE MALEATE

ALPHARMA US PHARMS	2MG/5ML	N86936	001	
PHARM ASSOC	2MG/5ML	N87517	001	
USL PHARMA	2MG/5ML	N87964	001	Jan 25, 1983

INJECTABLE; INJECTION

BROMPHENIRAMINE MALEATE

STERIS	100MG/ML	N83820	001	
DIMETANE-TEN				
WYETH AYERST	10MG/ML	N11418	002	

TABLET; ORAL

BROMPHENIRAMINE MALEATE

ANABOLIC	4MG	N86187	001	
BARR	4MG	N84468	001	
IVAX PHARMS	4MG	N84351	001	
NEWTRON PHARMS	4MG	N86987	001	
PAR PHARM	4MG	N87009	001	
PIIONEER PHARMS	4MG	N88604	001	Jul 13, 1984
SANDOZ	4MG	N83215	001	
VITARINE	4MG	N85850	001	
WATSON LABS	4MG	N83123	001	
	4MG	N85769	001	

DIMETANE

WYETH CONS	4MG	N10799	003	
------------	-----	--------	-----	--

TABLET, EXTENDED RELEASE; ORAL

DIMETANE

WYETH CONS	8MG	N10799	010	Jun 10, 1983
	12MG	N10799	011	Jun 10, 1983

BROMPHENIRAMINE MALEATE; CODEINE PHOSPHATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

SYRUP; ORAL

BROMANATE DC

ALPHARMA US PHARMS	2MG/5ML;10MG/5ML;12.5MG/5ML	N88723	001	Feb 25, 1985
--------------------	-----------------------------	--------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 37 (of 263)

BROMPHENIRAMINE MALEATE; CODEINE PHOSPHATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

SYRUP; ORAL		
DIMETANE-DC		
ROBINS AH	2MG/5ML;10MG/5ML;12.5MG/5ML	N11694 006 Mar 29, 1984
MYPHETANE DC		
MORTON GROVE	2MG/5ML;10MG/5ML;12.5MG/5ML	N88904 001 Feb 21, 1985

BROMPHENIRAMINE MALEATE; DEXTROMETHORPHAN HYDROBROMIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SYRUP; ORAL		
BROMANATE DM		
ALPHARMA US PHARMS	2MG/5ML;10MG/5ML;30MG/5ML	N88722 001 Mar 07, 1985
DIMETANE-DX		
ROBINS AH	2MG/5ML;10MG/5ML;30MG/5ML	N19279 001 Aug 24, 1984

BROMPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

ELIXIR; ORAL		
BIPHETAP		
MORTON GROVE	4MG/5ML;25MG/5ML	N88687 001 Sep 26, 1984
BROMANATE		
ALPHARMA US PHARMS	4MG/5ML;25MG/5ML	N88688 001 Feb 06, 1985
DIMETAPP		
WYETH CONS	2MG/5ML;12.5MG/5ML	N13087 003 Mar 29, 1984
TABLET, EXTENDED RELEASE; ORAL		
BROMATAPP		
COPLEY PHARM	12MG;75MG	N71099 001 Jul 02, 1987
DIMETAPP		
WYETH CONS	12MG;75MG	N12436 003 May 14, 1985

BUCLIZINE HYDROCHLORIDE

TABLET; ORAL		
BUCLADIN-S		
STUART PHARMS	50MG	N10911 006

BUDESONIDE

AEROSOL, METERED; NASAL		
RHINOCORT		
ASTRAZENECA	0.032MG/INH	N20233 001 Feb 14, 1994
POWDER, METERED; INHALATION		
PULMICORT		
ASTRAZENECA	0.32MG/INH	N20441 003 Jun 24, 1997
SUSPENSION; INHALATION		
PULMICORT RESPULES		
ASTRAZENECA	1MG/2ML	N20929 003 Aug 08, 2000

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION		
BUPIVACAINE HYDROCHLORIDE KIT		
HOSPIRA	0.075%	N19978 001 Sep 03, 1992
	0.114%	N19978 002 Sep 03, 1992
	0.23%	N19978 003 Sep 03, 1992

BUPROPION HYDROCHLORIDE

TABLET; ORAL		
WELLBUTRIN		
GLAXOSMITHKLINE	50MG	N18644 001 Dec 30, 1985
TABLET, EXTENDED RELEASE; ORAL		
ZYBAN		
GLAXOSMITHKLINE	100MG	N20711 002 May 14, 1997

BUSPIRONE HYDROCHLORIDE

CAPSULE; ORAL		
BUSPAR		
BRISTOL MYERS SQUIBB	5MG	N21190 001 Dec 20, 2000
	7.5MG	N21190 002 Dec 20, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 38 (of 263)

BUSPIRONE HYDROCHLORIDECAPSULE; ORAL
BUSPAR

BRISTOL MYERS SQUIBB	10MG	N21190	003	Dec 20, 2000
	15MG	N21190	004	Dec 20, 2000

BUTABARBITAL SODIUMCAPSULE; ORAL
BUTICAPS

MEDPOINTE PHARM HLC	15MG	N85381	001
	30MG	N85381	002
	50MG	N85381	003
	100MG	N85381	004

ELIXIR; ORAL

BUTABARB

ALPHARMA US PHARMS	30MG/5ML	N85873	001
--------------------	----------	--------	-----

BUTABARBITAL SODIUM

MORTON GROVE	30MG/5ML	N85383	001
--------------	----------	--------	-----

BUTALAN

LANNETT	33.3MG/5ML	N85880	001
---------	------------	--------	-----

SARISOL

HALSEY	30MG/5ML	N84723	001
--------	----------	--------	-----

TABLET; ORAL

BUTABARBITAL SODIUM

SANDOZ	15MG	N84292	003	Feb 09, 1982
--------	------	--------	-----	--------------

	15MG	N85938	001
--	------	--------	-----

	30MG	N84272	002
--	------	--------	-----

	30MG	N85934	001
--	------	--------	-----

SOLVAY	16.2MG	N83606	001
--------	--------	--------	-----

	32.4MG	N83898	001
--	--------	--------	-----

	48.6MG	N83897	001
--	--------	--------	-----

	97.2MG	N83896	001
--	--------	--------	-----

TEVA	15MG	N88632	001	May 18, 1985
------	------	--------	-----	--------------

	30MG	N88631	001	May 01, 1985
--	------	--------	-----	--------------

WATSON LABS	15MG	N85764	001
-------------	------	--------	-----

	30MG	N85772	001
--	------	--------	-----

WHITEWORTH TOWN PLSN	15MG	N83325	002
----------------------	------	--------	-----

	30MG	N83337	001
--	------	--------	-----

BUTISOL SODIUM

MEDPOINTE PHARM HLC	15MG	N00793	002
---------------------	------	--------	-----

	100MG	N00793	005
--	-------	--------	-----

SARISOL NO. 1

HALSEY	15MG	N84719	001
--------	------	--------	-----

SARISOL NO. 2

HALSEY	30MG	N84719	002
--------	------	--------	-----

SODIUM BUTABARBITAL

IVAX PHARMS	15MG	N83484	001
-------------	------	--------	-----

	30MG	N84040	001
--	------	--------	-----

LANNETT	15MG	N85849	001
---------	------	--------	-----

	30MG	N85866	001
--	------	--------	-----

	100MG	N85881	001
--	-------	--------	-----

WEST WARD	15MG	N85418	001
-----------	------	--------	-----

	30MG	N85432	001
--	------	--------	-----

BUTOCONAZOLE NITRATECREAM; VAGINAL
FEMSTAT

ROCHE PALO	2%	N19215	001	Nov 25, 1985
------------	----	--------	-----	--------------

SUPPOSITORY; VAGINAL
FEMSTAT

ROCHE PALO	100MG	N19359	001	Nov 25, 1985
------------	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 39 (of 263)

BUTORPHANOL TARTRATESPRAY, METERED; NASAL
STADOL

BRISTOL MYERS SQUIBB 1MG/SPRAY

N19890 001 Dec 12, 1991

CAFFEINE; ERGOTAMINE TARTRATETABLET; ORAL
CAFERGOT

NOVARTIS 100MG;1MG

N06620 001

WIGRAINE

ORGANON USA INC 100MG;1MG

N86562 001

CALCIFEDIOLCAPSULE; ORAL
CALDEROL

ORGANON USA INC 0.02MG

N18312 001

0.05MG

N18312 002

CALCITONIN HUMANINJECTABLE; INJECTION
CIBACALCIN

NOVARTIS 0.5MG/VIAL

N18470 001 Oct 31, 1986

CALCITONIN, SALMONINJECTABLE; INJECTION
CALCIMARAVENTIS 200 IU/ML
400 IU/VIAL

N17769 001

N17497 001

CALCITONIN-SALMON

ASTRAZENECA 200 IU/ML

N73690 001 Apr 14, 1995

MIACALCIN

NOVARTIS 100 IU/ML

N17808 001 Jul 03, 1986

CALCITRIOLINJECTABLE; INJECTION
CALCITRIOLMAYNE PHARMA USA 0.001MG/ML
0.002MG/ML

N75816 001 Jan 16, 2004

N75816 002 Jan 16, 2004

CALCIUM ACETATECAPSULE; ORAL
PHOSLONABI EQ 84.5MG CALCIUM
EQ 169MG CALCIUM

N21160 001 Apr 02, 2001

N21160 002 Apr 02, 2001

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 37MG/100ML;5GM/100ML;31MG/100ML;120MG/100ML;330MG/100ML;88MG/100ML N18271 001

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN 35MG/100ML;5GM/100ML;30MG/100ML;74MG/100ML;640MG/100ML;500MG/100ML;74MG/100ML N18269 002 Jan 17, 1983

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

SOLUTION; INTRAPERITONEAL

DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN 29MG/100ML;2.5GM/100ML;15MG/100ML;610MG/100ML;560MG/100ML N18460 006 Jan 29, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 40 (of 263)

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDESOLUTION; INTRAPERITONEALDIALYTE W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

B BRAUN	29MG/100ML;1.5GM/100ML;15MG/100ML;610MG /100ML;560MG/100ML	N18460	001	
---------	---	--------	-----	--

DIALYTE W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

B BRAUN	29MG/100ML;4.25GM/100ML;15MG/100ML;610M G/100ML;560MG/100ML	N18460	003	
---------	--	--------	-----	--

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATESOLUTION; INTRAPERITONEALDIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER

B BRAUN	26MG/100ML;1.5GM/100ML;15MG/100ML;560MG /100ML;390MG/100ML	N18460	002	
---------	---	--------	-----	--

DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER

B BRAUN	26MG/100ML;5GM/100ML;5MG/100ML;530MG/10 0ML;450MG/100ML	N18460	008	Jan 29, 1986
---------	--	--------	-----	--------------

DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER

B BRAUN	26MG/100ML;4.25GM/100ML;15MG/100ML;560M G/100ML;390MG/100ML	N18460	004	
---------	--	--------	-----	--

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDEINJECTABLE; INJECTIONDEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER

B BRAUN	33MG/100ML;5GM/100ML;30MG/100ML;860MG/1 00ML	N18256	001	
---------	---	--------	-----	--

CALCIUM CHLORIDE; DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATEINJECTABLE; INJECTIONDEXTROSE 4% IN MODIFIED LACTATED RINGER'S IN PLASTIC CONTAINER

B BRAUN	4MG/100ML;4GM/100ML;6MG/100ML;120MG/100 ML;62MG/100ML	N19634	002	Feb 24, 1988
---------	--	--------	-----	--------------

DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER

B BRAUN	20MG/100ML;5GM/100ML;30MG/100ML;600MG/1 00ML;310MG/100ML	N17510	001	
---------	---	--------	-----	--

MILES	20MG/100ML;5GM/100ML;30MG/100ML;600MG/1 00ML;310MG/100ML	N18499	001	
-------	---	--------	-----	--

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

HOSPIRA	20MG/100ML;5GM/100ML;104MG/100ML;600MG/ 100ML;310MG/100ML	N19685	005	Oct 17, 1988
---------	--	--------	-----	--------------

	20MG/100ML;5GM/100ML;179MG/100ML;600MG/ 100ML;310MG/100ML	N19685	006	Oct 17, 1988
--	--	--------	-----	--------------

POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

HOSPIRA	20MG/100ML;5GM/100ML;254MG/100ML;600MG/ 100ML;310MG/100ML	N19685	007	Oct 17, 1988
---------	--	--------	-----	--------------

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

HOSPIRA	20MG/100ML;5GM/100ML;254MG/100ML;600MG/ 100ML;310MG/100ML	N19685	003	Oct 17, 1988
---------	--	--------	-----	--------------

POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER

HOSPIRA	20MG/100ML;5GM/100ML;104MG/100ML;600MG/ 100ML;310MG/100ML	N19685	001	Oct 17, 1988
---------	--	--------	-----	--------------

CALCIUM CHLORIDE; DEXTROSE; SODIUM CHLORIDE; SODIUM LACTATESOLUTION; INTRAPERITONEALINPERSOL-ZM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML;1.5GM/100ML;538MG/100ML;44 8MG/100ML	N19395	001	Mar 26, 1986
-----------------	--	--------	-----	--------------

INPERSOL-ZM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML;2.5GM/100ML;538MG/100ML;44 8MG/100ML	N19395	002	Mar 26, 1986
-----------------	--	--------	-----	--------------

INPERSOL-ZM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

FRESENIUS MEDCL	25.7MG/100ML;4.25GM/100ML;538MG/100ML;4 48MG/100ML	N19395	003	Mar 26, 1986
-----------------	---	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 41 (of 263)

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

TPN ELECTROLYTES IN PLASTIC CONTAINER

ABBOTT	16.5MG/ML;25.4MG/ML;74.6MG/ML;121MG/ML; 16.1MG/ML	N19399	001	Jun 16, 1986
--------	--	--------	-----	--------------

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

INJECTABLE; INJECTION

ISOLYTE E IN PLASTIC CONTAINER

B BRAUN	35MG/100ML;30MG/100ML;74MG/100ML;640MG/ 100ML;500MG/100ML;74MG/100ML	N18899	001	Oct 31, 1983
---------	---	--------	-----	--------------

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ACETATED RINGER'S IN PLASTIC CONTAINER

B BRAUN	20MG/100ML;30MG/100ML;380MG/100ML;600MG/ /100ML	N18725	001	Nov 29, 1982
---------	--	--------	-----	--------------

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

RINGER'S IN PLASTIC CONTAINER

B BRAUN	33MG/100ML;30MG/100ML;860MG/100ML	N18721	001	Nov 09, 1982
---------	-----------------------------------	--------	-----	--------------

SOLUTION; IRRIGATION

RINGER'S IN PLASTIC CONTAINER

ABBOTT	33MG/100ML;30MG/100ML;860MG/100ML	N18462	001	
--------	-----------------------------------	--------	-----	--

CALCIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

INJECTABLE; INJECTION

LACTATED RINGER'S IN PLASTIC CONTAINER

ABBOTT	20MG/100ML;30MG/100ML;600MG/100ML;310MG/ /100ML	N19485	001	Oct 24, 1985
B BRAUN	20MG/100ML;30MG/100ML;600MG/100ML;310MG/ /100ML	N18023	001	
MILES	20MG/100ML;30MG/100ML;600MG/100ML;310MG/ /100ML	N18417	001	

CALCIUM GLUCEPTATE

INJECTABLE; INJECTION

CALCIUM GLUCEPTATE

ABBOTT	EQ 90MG CALCIUM/5ML	N80001	001	
	EQ 90MG CALCIUM/5ML	N83159	001	
AM PHARM PARTNERS	EQ 90MG CALCIUM/5ML	N89373	001	Apr 30, 1987
LILLY	EQ 90MG CALCIUM/5ML	N06470	001	

CALCIUM METRIZOATE; MEGLUMINE METRIZOATE; METRIZOATE MAGNESIUM; METRIZOATE SODIUM

INJECTABLE; INJECTION

ISOPAQUE 440

GE HEALTHCARE	0.78MG/ML;75.9MG/ML;0.15MG/ML;16.6MG/ML	N16847	001	
---------------	---	--------	-----	--

CALCIUM; MEGLUMINE; METRIZOIC ACID

INJECTABLE; INJECTION

ISOPAQUE 280

GE HEALTHCARE	0.35MG/ML;140.1MG/ML;461.8MG/ML	N17506	001	
---------------	---------------------------------	--------	-----	--

CANDICIDIN

OINTMENT; VAGINAL

VANOBIID

AVENTIS PHARMS	0.6MG/GM	N61596	001	
----------------	----------	--------	-----	--

TABLET; VAGINAL

VANOBIID

AVENTIS PHARMS	3MG	N61613	001	
----------------	-----	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 42 (of 263)

CAPTOPRIL

TABLET; ORAL
CAPOTEN

PAR PHARM	37.5MG	N18343	006	Sep 17, 1986
	75MG	N18343	007	Jun 13, 1995
	150MG	N18343	004	Jun 13, 1995

CAPTOPRIL

PAR PHARM	12.5MG	N74493	001	Feb 13, 1996
	25MG	N74493	002	Feb 13, 1996
	50MG	N74493	003	Feb 13, 1996
	100MG	N74493	004	Feb 13, 1996
PUREPAC PHARM	12.5MG	N74640	001	Mar 31, 1997
	25MG	N74640	002	Mar 31, 1997
	50MG	N74640	003	Mar 31, 1997
	100MG	N74640	004	Mar 31, 1997

CAPTOPRIL; HYDROCHLOROTHIAZIDE

TABLET; ORAL
CAPTOPRIL AND HYDROCHLOROTHIAZIDE
WATSON LABS

50MG;25MG	N74832	001	Dec 29, 1997
-----------	--------	-----	--------------

CARBACHOL

SOLUTION; INTRAOCULAR
CARBACHOL
PHARMAFAIR

0.01%	N70292	001	May 21, 1986
-------	--------	-----	--------------

CARBAMAZEPINE

SUSPENSION; ORAL
CARBAMAZEPINE

TARO	100MG/5ML	N75875	001	Dec 21, 2000
------	-----------	--------	-----	--------------

TABLET; ORAL
CARBAMAZEPINE

USL PHARMA	200MG	N70300	001	May 15, 1986
WARNER CHILCOTT	200MG	N70429	001	Jan 02, 1987

CARBENICILLIN DISODIUM

INJECTABLE; INJECTION
GEOPEN

ROERIG	EQ 1GM BASE/VIAL	N50306	001	
	EQ 2GM BASE/VIAL	N50306	004	
	EQ 5GM BASE/VIAL	N50306	002	
	EQ 10GM BASE/VIAL	N50306	006	
	EQ 30GM BASE/VIAL	N50306	007	

PYOPEN

GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	N50298	001	
	EQ 2GM BASE/VIAL	N50298	002	
	EQ 5GM BASE/VIAL	N50298	003	
	EQ 10GM BASE/VIAL	N50298	006	
	EQ 20GM BASE/VIAL	N50298	007	

CARBIDOPA; LEVODOPA

TABLET; ORAL
CARBIDOPA AND LEVODOPA
SCS

10MG;100MG	N74080	001	Mar 25, 1994	
25MG;100MG	N74080	002	Mar 25, 1994	
25MG;250MG	N74080	003	Mar 25, 1994	
WATSON LABS	10MG;100MG	N73381	001	Sep 28, 1993
	25MG;100MG	N73382	001	Sep 28, 1993
	25MG;250MG	N73383	001	Sep 28, 1993

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 43 (of 263)

CARBINOXAMINE MALEATE

ELIXIR; ORAL CLISTIN MCNEIL	4MG/5ML **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N08955	001	
TABLET; ORAL CLISTIN ORTHO MCNEIL PHARM	4MG	N08915	001	

CARBOPLATIN

INJECTABLE; INJECTION CARBOPLATIN MAYNE PHARMA USA	50MG/VIAL 150MG/VIAL 450MG/VIAL	N76473 N76473 N76473	001 002 003	Oct 27, 2004 Oct 27, 2004 Oct 27, 2004
--	---------------------------------------	----------------------------	-------------------	--

CARISOPRODOL

CAPSULE; ORAL SOMA MEDPOINTE PHARM HLC	250MG	N11792	003	
TABLET; ORAL CARISOPRODOL ABLE	350MG	N40421	001	Jun 21, 2001
PIONEER PHARMS	350MG	N89390	001	Oct 13, 1988
SANDOZ	350MG	N89566	001	Aug 30, 1988
WATSON LABS	350MG	N85433	001	
RELA SCHERING	350MG	N12155	001	

CARPHENAZINE MALEATE

CONCENTRATE; ORAL PROKETAZINE WYETH AYERST	50MG/ML	N14173	001	
TABLET; ORAL PROKETAZINE WYETH AYERST	12.5MG 25MG 50MG	N12768 N12768 N12768	001 002 004	

CARPROFEN

TABLET; ORAL RIMADYL ROCHE	100MG 150MG	N18550 N18550	002 003	Dec 31, 1987 Dec 31, 1987
----------------------------------	----------------	------------------	------------	------------------------------

CARTEOLOL HYDROCHLORIDE

TABLET; ORAL CARTROL ABBOTT	2.5MG 5MG 10MG	N19204 N19204 N19204	001 002 003	Dec 28, 1988 Dec 28, 1988 Dec 28, 1988
-----------------------------------	----------------------	----------------------------	-------------------	--

CEFACLOR

CAPSULE; ORAL CECLOR LILLY	EQ 250MG BASE EQ 500MG BASE	N50521 N50521	001 002	
CEFACLOR CLONMEL HLTHCARE	EQ 250MG BASE EQ 500MG BASE	N64107 N64107	001 002	Apr 27, 1995 Apr 27, 1995
TEVA	EQ 250MG BASE EQ 500MG BASE	N64081 N64081	001 002	Sep 16, 1996 Sep 16, 1996

DISCONTINUED DRUG PRODUCT LIST

6 - 44 (of 263)

CEFACLOR

FOR SUSPENSION; ORAL

CECLOR

LILLY

EQ 125MG BASE/5ML

N50522 001

EQ 250MG BASE/5ML

N50522 002

CEFACLOR

CLONMEL HLTHCARE

EQ 125MG BASE/5ML

N64114 001

Apr 28, 1995

EQ 187MG BASE/5ML

N64115 001

Apr 28, 1995

EQ 250MG BASE/5ML

N64116 001

Apr 28, 1995

EQ 375MG BASE/5ML

N64110 001

Apr 28, 1995

IVAX PHARMS

EQ 125MG BASE/5ML

N64087 001

Apr 28, 1995

EQ 187MG BASE/5ML

N64086 001

Apr 28, 1995

EQ 250MG BASE/5ML

N64085 001

Apr 28, 1995

TABLET, EXTENDED RELEASE; ORAL

CECLOR CD

LILLY

EQ 375MG BASE

N50673 001

Jun 28, 1996

EQ 500MG BASE

N50673 002

Jun 28, 1996

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

PUREPAC PHARM

EQ 500MG BASE

N63017 001

Jan 05, 1989

TEVA

EQ 500MG BASE

N62695 001

Feb 10, 1989

DURICEF

WARNER CHILCOTT

EQ 250MG BASE

N50512 002

ULTRACEF

BRISTOL

EQ 500MG BASE

N62378 001

Mar 16, 1982

FOR SUSPENSION; ORAL

CEFADROXIL

APOTHECON

EQ 125MG BASE/5ML

N62334 001

EQ 250MG BASE/5ML

N62334 002

EQ 500MG BASE/5ML

N62334 003

TEVA

EQ 125MG BASE/5ML

N62698 001

Mar 01, 1989

EQ 250MG BASE/5ML

N62698 002

Mar 01, 1989

EQ 500MG BASE/5ML

N62698 003

Mar 01, 1989

ULTRACEF

BRISTOL

EQ 125MG BASE/5ML

N62376 001

Mar 16, 1982

EQ 250MG BASE/5ML

N62376 002

Mar 16, 1982

EQ 500MG BASE/5ML

N62376 003

Mar 16, 1982

TABLET; ORAL

ULTRACEF

APOTHECON

EQ 1GM BASE

N62390 001

Jun 10, 1982

BRISTOL

EQ 1GM BASE

N62408 001

Aug 31, 1982

CEFAMANDOLE NAFATE

INJECTABLE; INJECTION

MANDOL

LILLY

EQ 500MG BASE/VIAL

N50504 001

EQ 1GM BASE/VIAL

N50504 002

EQ 1GM BASE/VIAL

N62560 001

Sep 10, 1985

EQ 2GM BASE/VIAL

N50504 003

EQ 2GM BASE/VIAL

N62560 002

Sep 10, 1985

EQ 10GM BASE/VIAL

N50504 004

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

ANCEF

GLAXOSMITHKLINE

EQ 250MG BASE/VIAL

N50461 001

EQ 500MG BASE/VIAL

N50461 002

EQ 5GM BASE/VIAL

N50461 004

ANCEF IN DEXTROSE 5% IN

BAXTER HLTHCARE

EQ 10MG BASE/ML

N50566 003

Jun 08, 1983

EQ 20MG BASE/ML

N50566 004

Jun 08, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 45 (of 263)

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

ANCEF IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

BAXTER HLTHCARE	EQ 10MG BASE/ML	N50566	001	Jun 08, 1983
	EQ 20MG BASE/ML	N50566	002	Jun 08, 1983

CEFAZOLIN SODIUM

AM PHARM PARTNERS	EQ 500MG BASE/VIAL	N62688	002	Nov 17, 1986
	EQ 1GM BASE/VIAL	N62688	003	Nov 17, 1986
	EQ 10GM BASE/VIAL	N62688	004	Nov 17, 1986
	EQ 20GM BASE/VIAL	N62688	005	Aug 03, 1987
BEDFORD	EQ 250MG BASE/VIAL	N62894	001	Jul 21, 1988
	EQ 500MG BASE/VIAL	N62894	002	Jul 21, 1988
	EQ 1GM BASE/VIAL	N62894	003	Jul 21, 1988
	EQ 5GM BASE/VIAL	N62894	004	Jul 21, 1988
	EQ 10GM BASE/VIAL	N62894	005	Jul 21, 1988
ELKINS SINN	EQ 250MG BASE/VIAL	N62807	001	Jan 12, 1988
	EQ 500MG BASE/VIAL	N62807	002	Jan 12, 1988
	EQ 1GM BASE/VIAL	N62807	003	Jan 12, 1988
	EQ 5GM BASE/VIAL	N62807	004	Jan 12, 1988
	EQ 10GM BASE/VIAL	N62807	005	Jan 12, 1988
	EQ 20GM BASE/VIAL	N62807	006	Jan 12, 1988
MARSAM PHARMS LLC	EQ 250MG BASE/VIAL	N62988	001	Dec 29, 1989
	EQ 500MG BASE/VIAL	N62988	002	Dec 29, 1989
	EQ 1GM BASE/VIAL	N62988	003	Dec 29, 1989
	EQ 5GM BASE/VIAL	N62989	001	Dec 29, 1989
	EQ 10GM BASE/VIAL	N62989	002	Dec 29, 1989
	EQ 20GM BASE/VIAL	N62989	003	Dec 29, 1989
TEVA	EQ 250MG BASE/VIAL	N63016	001	Mar 14, 1989
	EQ 500MG BASE/VIAL	N63016	002	Mar 14, 1989
	EQ 1GM BASE/VIAL	N63016	003	Mar 14, 1989
	EQ 5GM BASE/VIAL	N63018	001	Mar 05, 1990
	EQ 10GM BASE/VIAL	N63018	002	Mar 05, 1990
KEFZOL				
LILLY	EQ 250MG BASE/VIAL	N61773	001	
	EQ 500MG BASE/VIAL	N61773	002	
	EQ 500MG BASE/VIAL	N62557	001	Sep 10, 1985
	EQ 1GM BASE/VIAL	N61773	003	
	EQ 1GM BASE/VIAL	N62557	002	Sep 10, 1985
	EQ 10GM BASE/VIAL	N61773	004	
	EQ 20GM BASE/VIAL	N61773	005	Sep 08, 1987

CEFIXIME

FOR SUSPENSION; ORAL

SUPRAX

LEDERLE	100MG/5ML	N50622	001	Apr 28, 1989
---------	-----------	--------	-----	--------------

TABLET; ORAL

SUPRAX

LEDERLE	200MG **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N50621	001	Apr 28, 1989
---------	--	--------	-----	--------------

	400MG	N50621	002	Apr 28, 1989
--	-------	--------	-----	--------------

LUPIN	400MG	N65130	001	Feb 12, 2004
-------	-------	--------	-----	--------------

CEFMENOXIME HYDROCHLORIDE

INJECTABLE; INJECTION

CEFMAX

TAP PHARM	EQ 500MG BASE/VIAL	N50571	001	Dec 30, 1987
	EQ 1GM BASE/VIAL	N50571	002	Dec 30, 1987
	EQ 2GM BASE/VIAL	N50571	003	Dec 30, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 46 (of 263)

CEFMETAZOLE SODIUMINJECTABLE; INJECTION
ZEFAZONEPHARMACIA AND UPJOHN EQ 1GM BASE/VIAL
EQ 2GM BASE/VIALN50637 001 Dec 11, 1989
N50637 002 Dec 11, 1989

ZEFAZONE IN PLASTIC CONTAINER

PHARMACIA AND UPJOHN EQ 20MG BASE/ML
EQ 40MG BASE/MLN50683 001 Dec 29, 1992
N50683 002 Dec 29, 1992CEFONICID SODIUMINJECTABLE; INJECTION
MONOCIDGLAXOSMITHKLINE EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 10GM BASE/VIALN50579 001 May 23, 1984
N50579 002 May 23, 1984
N63295 001 Jul 26, 1993
N50579 003 May 23, 1984
N50579 004 May 23, 1984CEFOPERAZONE SODIUMINJECTABLE; INJECTION
CEFOBIDPFIZER EQ 1GM BASE/VIAL
EQ 2GM BASE/VIALN63333 001 Mar 31, 1995
N63333 002 Mar 31, 1995CEFORANIDEINJECTABLE; INJECTION
PRECEFAPOTHECON 500MG/VIAL
1GM/VIAL
2GM/VIAL
10GM/VIAL
20GM/VIALN62579 001 Nov 26, 1984
N62579 002 Nov 26, 1984
N62579 003 Nov 26, 1984
N62579 004 Nov 26, 1984
N62579 005 Nov 26, 1984BRISTOL 500MG/VIAL
1GM/VIAL
2GM/VIAL
10GM/VIAL
20GM/VIALN50554 001 May 24, 1984
N50554 002 May 24, 1984
N50554 003 May 24, 1984
N50554 004 May 24, 1984
N50554 005 May 24, 1984CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CLAFORAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AVENTIS PHARMS EQ 20MG BASE/ML
EQ 40MG BASE/MLN50596 001 May 20, 1985
N50596 003 May 20, 1985CEFOTIAM HYDROCHLORIDEINJECTABLE; INJECTION
CERADON

TAKEDA EQ 1GM BASE/VIAL

N50601 001 Dec 30, 1988

CEFOXITIN SODIUMINJECTABLE; INJECTION
MEFOXINMERCK EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 10GM BASE/VIALN50517 001
N50517 002
N50517 003

MEFOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

MERCK EQ 20MG BASE/ML
EQ 40MG BASE/MLN50581 003 Sep 20, 1984
N50581 004 Sep 20, 1984

MEFOXIN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

MERCK EQ 20MG BASE/ML
EQ 40MG BASE/MLN50581 002 Sep 20, 1984
N50581 001 Sep 20, 1984

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 47 (of 263)

CEFTIRAMIDE SODIUM

INJECTABLE; INJECTION

CEFTIRAMIDE SODIUM

WYETH AYERST

EQ 1GM BASE/VIAL

N50633 002 Jan 31, 1989

EQ 2GM BASE/VIAL

N50633 003 Jan 31, 1989

EQ 10GM BASE/VIAL

N50633 005 Jan 31, 1989

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

BANAN

SANKYO

EQ 50MG BASE/5ML

N50688 002 Aug 07, 1992

EQ 100MG BASE/5ML

N50688 001 Aug 07, 1992

TABLET; ORAL

BANAN

SANKYO

EQ 100MG BASE

N50687 001 Aug 07, 1992

EQ 200MG BASE

N50687 002 Aug 07, 1992

CEFTAZIDIME

INJECTABLE; INJECTION

TAZIDIME

LILLY

1GM/VIAL

N62655 001 Nov 20, 1985

2GM/VIAL

N62655 002 Nov 20, 1985

TAZIDIME IN PLASTIC CONTAINER

LILLY

1GM/VIAL

N62739 001 Jul 10, 1986

2GM/VIAL

N62739 002 Jul 10, 1986

CEFTAZIDIME (ARGININE FORMULATION)

INJECTABLE; INJECTION

CEPTAZ

GLAXOSMITHKLINE

500MG/VIAL

N50646 001 Sep 27, 1990

PENTACEF

GLAXOSMITHKLINE

1GM/VIAL

N63322 001 Nov 07, 1995

1GM/VIAL

N64006 001 Mar 31, 1992

2GM/VIAL

N63322 002 Nov 07, 1995

2GM/VIAL

N64006 002 Mar 31, 1992

6GM/VIAL

N64008 001 Mar 31, 1992

10GM/VIAL

N64008 002 Mar 31, 1992

CEFTAZIDIME SODIUM

INJECTABLE; INJECTION

FORTAZ IN PLASTIC CONTAINER

GLAXOSMITHKLINE

EQ 10MG BASE/ML

N50634 001 Apr 28, 1989

CEFTIBUTEN DIHYDRATE

FOR SUSPENSION; ORAL

CEDAX

SHIONOGI

EQ 180MG BASE/5ML

N50686 002 Dec 20, 1995

CEFTIZOXIME SODIUM

INJECTABLE; INJECTION

CEFIZOX

ASTELLAS

EQ 500MG BASE/VIAL

N50560 001 Sep 15, 1983

CEFIZOX IN DEXTROSE 5% IN PLASTIC CONTAINER

ASTELLAS

EQ 20MG BASE/ML

N50589 001 Oct 03, 1984

EQ 40MG BASE/ML

N50589 002 Oct 03, 1984

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION

ROCEPHIN

HLR

EQ 250MG BASE/VIAL

N63239 001 Aug 13, 1993

EQ 500MG BASE/VIAL

N62654 001 Apr 30, 1987

EQ 500MG BASE/VIAL

N63239 002 Aug 13, 1993

EQ 1GM BASE/VIAL

N63239 003 Aug 13, 1993

DISCONTINUED DRUG PRODUCT LIST

6 - 48 (of 263)

CEFTRIAXONE SODIUM

INJECTABLE; INJECTION
ROCEPHIN

ROCHE	EQ 250MG BASE/VIAL	N62510	001	Mar 12, 1985
	EQ 500MG BASE/VIAL	N62510	002	Mar 12, 1985
	EQ 1GM BASE/VIAL	N62510	003	Mar 12, 1985
ROCEPHIN W/ DEXTROSE IN	PLASTIC CONTAINER			
HLR	EQ 10MG BASE/ML	N50624	001	Feb 11, 1987

CEFTRIAXONE SODIUM; LIDOCAINE

INJECTABLE; INJECTION
ROCEPHIN KIT

HLR	EQ 500MG BASE/VIAL, N/A; N/A, 1%	N50585	007	May 08, 1996
	EQ 1GM BASE/VIAL, N/A; N/A, 1%	N50585	006	May 08, 1996

CEFUROXIME SODIUM

INJECTABLE; IM-IV
KEFUROX

LILLY	EQ 750MG BASE/VIAL	N62591	001	Jan 10, 1986
-------	--------------------	--------	-----	--------------

INJECTABLE; INJECTION
KEFUROX

LILLY	EQ 1.5GM BASE/VIAL	N62591	002	Jan 10, 1986
	EQ 1.5GM BASE/VIAL	N62592	002	Jan 10, 1986
	EQ 7.5GM BASE/VIAL	N62591	003	Dec 17, 1987

KEFUROX IN PLASTIC CONTAINER

LILLY	EQ 1.5GM BASE/VIAL	N62590	002	Jan 10, 1986
-------	--------------------	--------	-----	--------------

INJECTABLE; INTRAVENOUS

KEFUROX

LILLY	EQ 750MG BASE/VIAL	N62592	001	Jan 10, 1986
-------	--------------------	--------	-----	--------------

KEFUROX IN PLASTIC CONTAINER

LILLY	EQ 750MG BASE/VIAL	N62590	001	Jan 10, 1986
-------	--------------------	--------	-----	--------------

CELLULOSE SODIUM PHOSPHATE

POWDER; ORAL
CALCIBIND

MISSION PHARMA	2.5GM/PACKET	N18757	002	Dec 28, 1982
	300GM/BOT	N18757	003	Oct 16, 1984

CEPHALEXIN

CAPSULE; ORAL
CEPHALEXIN

APOTHECON	EQ 250MG BASE	N62973	001	Nov 08, 1988
	EQ 250MG BASE	N63186	001	Dec 30, 1994
	EQ 500MG BASE	N62974	001	Nov 23, 1988
	EQ 500MG BASE	N63186	002	Dec 30, 1994
PUREPAC PHARM	EQ 250MG BASE	N62809	001	Apr 22, 1987
	EQ 500MG BASE	N62809	002	Apr 22, 1987
STEVENS J	EQ 500MG BASE	N62869	001	Mar 17, 1988
TEVA	EQ 250MG BASE	N62821	001	Feb 05, 1988
	EQ 500MG BASE	N62823	001	Feb 05, 1988
YOSHITOMI	EQ 250MG BASE	N62872	001	Jun 20, 1988
	EQ 500MG BASE	N62871	001	Jul 05, 1988

FOR SUSPENSION; ORAL

CEPHALEXIN

APOTHECON	EQ 125MG BASE/5ML	N62986	001	Apr 18, 1991
	EQ 250MG BASE/5ML	N62987	001	Jul 25, 1989
BARR	EQ 125MG BASE/5ML	N62778	001	Aug 06, 1987
TEVA	EQ 125MG BASE/5ML	N62873	001	May 23, 1988
	EQ 250MG BASE/5ML	N62867	001	Apr 15, 1988
VITARINE	EQ 125MG BASE/5ML	N62779	001	Dec 22, 1987
	EQ 250MG BASE/5ML	N62781	001	Dec 22, 1987

KEFLEX

ADVANCIS PHARM	EQ 100MG BASE/ML	N50406	003	
	EQ 125MG BASE/5ML	N50406	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 49 (of 263)

CEPHALEXIN

FOR SUSPENSION; ORAL

KEFLEX

ADVANCIS PHARM

EQ 250MG BASE/5ML

N50406 002

TABLET; ORAL

CEPHALEXIN

BARR

EQ 250MG BASE

N62826 001

Aug 17, 1987

EQ 500MG BASE

N62827 001

Aug 17, 1987

VITARINE

EQ 250MG BASE

N62863 001

Aug 11, 1988

EQ 500MG BASE

N62863 002

Aug 11, 1988

EQ 1GM BASE

N62863 003

Aug 11, 1988

KEFLET

LILLY

EQ 250MG BASE

N50440 003

Feb 26, 1987

EQ 250MG BASE

N62745 001

Dec 01, 1986

EQ 500MG BASE

N50440 001

EQ 500MG BASE

N62745 002

Dec 01, 1986

EQ 1GM BASE

N50440 002

CEPHALEXIN HYDROCHLORIDE

TABLET; ORAL

KEFTAB

LILLY

EQ 250MG BASE

N50614 001

Oct 29, 1987

EQ 333MG BASE

N50614 003

May 16, 1988

EQ 500MG BASE

N50614 002

Oct 29, 1987

CEPHALOGLYCIN

CAPSULE; ORAL

KAFOCIN

LILLY

250MG

N50219 001

CEPHALOTHIN SODIUM

INJECTABLE; INJECTION

CEPHALOTHIN

INTL MEDICATION

EQ 500MG BASE/VIAL

N62426 001

May 03, 1985

EQ 1GM BASE/VIAL

N62426 002

May 03, 1985

EQ 2GM BASE/VIAL

N62426 003

May 03, 1985

EQ 4GM BASE/VIAL

N62426 004

May 03, 1985

CEPHALOTHIN SODIUM

ABBOTT

EQ 1GM BASE/VIAL

N62547 001

Sep 11, 1985

EQ 1GM BASE/VIAL

N62548 001

Sep 11, 1985

EQ 2GM BASE/VIAL

N62547 002

Sep 11, 1985

EQ 2GM BASE/VIAL

N62548 002

Sep 11, 1985

AM PHARM PARTNERS

EQ 1GM BASE/VIAL

N62666 002

Jun 10, 1987

EQ 2GM BASE/VIAL

N62666 001

Jun 10, 1987

BRISTOL

EQ 1GM BASE/VIAL

N62464 001

May 07, 1984

EQ 2GM BASE/VIAL

N62464 002

May 07, 1984

EQ 4GM BASE/VIAL

N62464 003

May 07, 1984

CEPHALOTHIN SODIUM W/ DEXTROSE IN PLASTIC CONTAINER

BAXTER HLTHCARE

EQ 20MG BASE/ML

N62422 003

Jan 31, 1984

EQ 20MG BASE/ML

N62422 005

Jul 16, 1991

EQ 20MG BASE/ML

N62730 001

Mar 05, 1987

EQ 40MG BASE/ML

N62422 004

Jan 31, 1984

EQ 40MG BASE/ML

N62422 006

Jul 16, 1991

EQ 40MG BASE/ML

N62730 002

Mar 05, 1987

CEPHALOTHIN SODIUM W/ SODIUM CHLORIDE IN PLASTIC CONTAINER

BAXTER HLTHCARE

EQ 20MG BASE/ML

N62422 001

Jan 31, 1984

EQ 40MG BASE/ML

N62422 002

Jan 31, 1984

KEFLIN

LILLY

EQ 1GM BASE/VIAL

N50482 001

EQ 2GM BASE/VIAL

N50482 002

EQ 4GM BASE/VIAL

N50482 003

EQ 20GM BASE/VIAL

N50482 007

KEFLIN IN PLASTIC CONTAINER

LILLY

EQ 1GM BASE/VIAL

N62549 001

Sep 10, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 50 (of 263)

CEPHALOTHIN SODIUM

INJECTABLE; INJECTION

KEFLIN IN PLASTIC CONTAINER

LILLY	EQ 2GM BASE/VIAL	N62549	002	Sep 10, 1985
SEFFIN				
GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	N62435	001	Nov 15, 1983
	EQ 2GM BASE/VIAL	N62435	002	Nov 15, 1983
	EQ 10GM BASE/VIAL	N62435	003	Nov 15, 1983

CEPHAPIRIN SODIUM

INJECTABLE; INJECTION

CEFADYL

APOTHECON	EQ 500MG BASE/VIAL	N50446	005	
	EQ 500MG BASE/VIAL	N62961	001	Sep 20, 1988
	EQ 1GM BASE/VIAL	N50446	001	
	EQ 1GM BASE/VIAL	N61769	001	
	EQ 1GM BASE/VIAL	N62724	001	Dec 23, 1986
	EQ 1GM BASE/VIAL	N62961	002	Sep 20, 1988
	EQ 2GM BASE/VIAL	N50446	002	
	EQ 2GM BASE/VIAL	N61769	002	
	EQ 2GM BASE/VIAL	N62724	002	Dec 23, 1986
	EQ 2GM BASE/VIAL	N62961	003	Sep 20, 1988
	EQ 4GM BASE/VIAL	N50446	003	
	EQ 4GM BASE/VIAL	N61769	003	
	EQ 4GM BASE/VIAL	N62961	004	Sep 20, 1988
	EQ 20GM BASE/VIAL	N50446	004	

CEPHAPIRIN SODIUM

AM PHARM PARTNERS

	EQ 500MG BASE/VIAL	N62723	001	Nov 17, 1986
	EQ 1GM BASE/VIAL	N62723	002	Nov 17, 1986
	EQ 2GM BASE/VIAL	N62723	003	Nov 17, 1986
	EQ 4GM BASE/VIAL	N62723	004	Nov 17, 1986
	EQ 20GM BASE/VIAL	N62723	005	Nov 17, 1986
BAXTER HLTHCARE	EQ 500MG BASE/VIAL	N62720	001	Jul 02, 1987
	EQ 1GM BASE/VIAL	N62720	002	Jul 02, 1987
	EQ 2GM BASE/VIAL	N62720	003	Jul 02, 1987
	EQ 20GM BASE/VIAL	N62720	004	Jul 02, 1987

CEPHRADINE

CAPSULE; ORAL

CEPHRADINE

BARR	250MG	N62850	001	Apr 22, 1988
	500MG	N62851	001	Apr 22, 1988
IVAX PHARMS	250MG	N62762	001	Mar 06, 1987
	500MG	N62762	002	Mar 06, 1987
VITARINE	250MG	N62813	001	Feb 25, 1988
	500MG	N62813	002	Feb 25, 1988

VELOSEF '250'

ERSANA

VELOSEF '500'

ERSANA

FOR SUSPENSION; ORAL

CEPHRADINE

BARR	125MG/5ML	N62858	001	May 19, 1988
	250MG/5ML	N62859	001	May 19, 1988

INJECTABLE; INJECTION

VELOSEF

APOTHECON

	250MG/VIAL	N61976	001	
	500MG/VIAL	N61976	002	
	1GM/VIAL	N61976	004	
	2GM/VIAL	N61976	003	
	4GM/VIAL	N61976	005	

TABLET; ORAL

VELOSEF

BRISTOL MYERS SQUIBB 1GM

N50530 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 51 (of 263)

CERIVASTATIN SODIUM

TABLET; ORAL

BAYCOL

BAYER PHARMS

0.05MG

N20740 001

Jun 26, 1997

0.1MG

N20740 002

Jun 26, 1997

0.2MG

N20740 003

Jun 26, 1997

0.3MG

N20740 004

Jun 26, 1997

0.4MG

N20740 005

May 24, 1999

0.8MG

N20740 006

Jul 24, 2000

CERULETIDE DIETHYLAMINE

INJECTABLE; INJECTION

TYMTRAN

PHARMACIA AND UPJOHN

0.02MG/ML

N18296 001

CHENODIOL

TABLET; ORAL

CHENIX

AXCAN SCANDIPHARM

250MG

N18513 002

Jul 28, 1983

CHLOPHEDIANOL HYDROCHLORIDE

SYRUP; ORAL

ULO

3M

25MG/5ML

N12126 001

CHLORAMPHENICOL

CAPSULE; ORAL

AMPHICOL

FERRANTE

100MG

N60058 001

250MG

N60058 002

CHLORAMPHENICOL

IVAX PHARMS

250MG

N62247 001

CHLOROMYCETIN

PARKEDALE

50MG

N60591 001

100MG

N60591 003

250MG

N60591 002

MYCHEL

ARMENPHARM

250MG

N60851 001

CREAM; TOPICAL

CHLOROMYCETIN

PARKE DAVIS

1%

N50183 001

FOR SOLUTION; OPHTHALMIC

CHLOROMYCETIN

PARKEDALE

25MG/VIAL

N50143 001

INJECTABLE; INJECTION

CHLOROMYCETIN

PARKE DAVIS

250MG/ML

N50153 001

OINTMENT; OPHTHALMIC

CHLOROFAIR

PHARMAFAIR

1%

N62439 001

Apr 21, 1983

CHLOROMYCETIN

PARKEDALE

1%

N50156 001

CHLOROPTIC S.O.P.

ALLERGAN

1%

N61187 001

ECONOCHLOR

ALCON

1%

N61648 001

SOLUTION/DROPS; OPHTHALMIC

CHLORAMPHENICOL

AKORN

0.5%

N62042 001

ALCON

0.5%

N62628 001

Sep 25, 1985

CHLOROFAIR

PHARMAFAIR

0.5%

N62437 001

Apr 14, 1983

CHLOROPTIC

ALLERGAN

0.5%

N50091 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 52 (of 263)

CHLORAMPHENICOL

SOLUTION/DROPS; OPHTHALMIC

ECONOCHLOR

ALCON

0.5%

N61645 001

OPHTHOCHLOR

PARKEDALE

0.5%

N61220 001

OPTOMYCIN

OPTOPICS

0.5%

N62171 001

Mar 31, 1982

SOLUTION/DROPS; OTIC

CHLOROMYCETIN

PARKEDALE

0.5%

N50205 001

CHLORAMPHENICOL PALMITATE

SUSPENSION; ORAL

CHLOROMYCETIN PALMITATE

PARKE DAVIS

EQ 150MG BASE/5ML

N50152 001

EQ 150MG BASE/5ML

N62301 001

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL

ELKINS SINN

EQ 1GM BASE/VIAL

N62406 001

Nov 09, 1982

CHLORAMPHENICOL SODIUM SUCCINATE

GRUPPO LEPETIT

EQ 1GM BASE/VIAL

N62278 001

MYCHEL-S

ANGUS

EQ 1GM BASE/VIAL

N60132 001

CHLORAMPHENICOL; DESOXYRIBONUCLEASE; FIBRINOLYSIN

OINTMENT; TOPICAL

ELASE-CHLOROMYCETIN

PARKE DAVIS

10MG/GM;666 UNITS/GM;1 UNITS/GM

N50294 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE

FOR SUSPENSION; OPHTHALMIC

CHLOROMYCETIN HYDROCORTISONE

PARKEDALE

12.5MG/VIAL;25MG/VIAL

N50202 001

CHLORAMPHENICOL; HYDROCORTISONE ACETATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

OPHTHOCORT

PARKEDALE

10MG/GM;5MG/GM;10,000 UNITS/GM

N50201 002

CHLORAMPHENICOL; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

CHLOROMYXIN

PARKE DAVIS

1%;10,000 UNITS/GM

N50203 002

CHLORAMPHENICOL; PREDNISOLONE

OINTMENT; OPHTHALMIC

CHLOROPTIC-P S.O.P.

ALLERGAN

1%;0.5%

N61188 001

CHLORDIAZEPOXIDE

CAPSULE, EXTENDED RELEASE; ORAL

LIBRELEASE

VALEANT PHARM INTL

30MG

N17813 001

Sep 12, 1983

TABLET; ORAL

LIBRITABS

VALEANT PHARM INTL

5MG

N85482 001

10MG

N85481 001

25MG

N85488 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 53 (of 263)

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

A-POXIDE

ABBOTT

5MG

N85447 001

5MG

N85517 001

10MG

N85447 002

10MG

N85518 001

25MG

N85447 003

25MG

N85513 001

CHLORDIAZACHEL

RACHELLE

5MG

N85086 001

10MG

N84639 001

25MG

N85087 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

ASCOT

5MG

N87525 001

Jan 07, 1982

10MG

N87524 001

Jan 07, 1982

25MG

N87512 001

Jan 07, 1982

FERRANTE

5MG

N85118 001

10MG

N85119 001

25MG

N85120 001

HALSEY

5MG

N85340 001

10MG

N85339 001

25MG

N84685 001

IMPAX LABS

5MG

N86213 001

10MG

N85113 001

25MG

N86212 001

IVAX PHARMS

5MG

N83741 001

10MG

N83742 001

25MG

N83570 001

LEDERLE

5MG

N86892 001

5MG

N87234 001

10MG

N86876 001

10MG

N87037 001

25MG

N86893 001

25MG

N87231 001

MAST MM

10MG

N86217 001

MYLAN

5MG

N84886 001

10MG

N84601 001

25MG

N84887 001

PARKE DAVIS

5MG

N85163 001

10MG

N84598 001

25MG

N85164 001

PIONEER PHARMS

10MG

N89533 001

Jul 15, 1988

25MG

N89558 001

Jul 15, 1988

PUREPAC PHARM

5MG

N85155 001

10MG

N84939 002

25MG

N85144 001

ROXANE

5MG

N84706 001

10MG

N84700 001

25MG

N84705 001

SANDOZ

5MG

N84678 001

5MG

N84919 001

10MG

N84041 001

10MG

N84920 001

25MG

N84679 002

25MG

N84823 001

SUPERPHARM

5MG

N88987 001

Apr 25, 1985

10MG

N88986 001

Apr 25, 1985

25MG

N88988 001

Apr 25, 1985

TEVA

5MG

N88705 001

Jan 18, 1985

10MG

N88706 001

Jan 18, 1985

25MG

N86494 001

25MG

N88707 001

Jan 18, 1985

USL PHARMA

5MG

N84644 001

DISCONTINUED DRUG PRODUCT LIST

6 - 54 (of 263)

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL

CHLORDIAZEPOXIDE HYDROCHLORIDE

USL PHARMA	25MG	N84645	001	
VANGARD	5MG	N88129	001	Mar 28, 1983
	10MG	N88010	001	Mar 28, 1983
	25MG	N88130	001	Mar 28, 1983
WEST WARD	5MG	N85014	001	
	10MG	N85000	001	
	25MG	N85294	001	
LIBRIUM				
VALEANT PHARM INTL	5MG	N12249	002	
	10MG	N12249	001	
	25MG	N12249	003	
LYGEN				
ALRA	5MG	N85107	001	
	10MG	N85009	001	
	25MG	N85108	001	

INJECTABLE; INJECTION

LIBRIUM

VALEANT PHARM INTL	100MG/AMP	N12301	001	
--------------------	-----------	--------	-----	--

CHLORDIAZEPOXIDE; ESTROGENS, ESTERIFIED

TABLET; ORAL

MENRIUM 10-4

ROCHE	10MG;0.4MG	N14740	006	
-------	------------	--------	-----	--

MENRIUM 5-2

ROCHE	5MG;0.2MG	N14740	002	
-------	-----------	--------	-----	--

MENRIUM 5-4

ROCHE	5MG;0.4MG	N14740	004	
-------	-----------	--------	-----	--

CHLORHEXIDINE GLUCONATE

SOLUTION; TOPICAL

EXIDINE

XTTRIUM	2.5%	N19421	001	Dec 17, 1985
---------	------	--------	-----	--------------

SPONGE; TOPICAL

E-Z SCRUB

BECTON DICKINSON	4%	N73416	001	Mar 14, 2000
------------------	----	--------	-----	--------------

CHLORMERODRIN, HG-197

INJECTABLE; INJECTION

CHLORMERODRIN HG 197

BRACCO	0.6-1.4mCi/ML	N17269	001	
--------	---------------	--------	-----	--

CHLORMEZANONE

TABLET; ORAL

TRANCOPAL

SANOFI SYNTHELABO	100MG	N11467	003	
	200MG	N11467	005	

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

NESACAINE-MPF

ASTRAZENECA	2%	N09435	003	
	3%	N09435	004	

CHLOROQUINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARALEN HYDROCHLORIDE

SANOFI SYNTHELABO	EQ 40MG BASE/ML	N06002	002	
-------------------	-----------------	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 55 (of 263)

CHLOROQUINE PHOSPHATE

TABLET; ORAL

CHLOROQUINE PHOSPHATE

MD PHARM	EQ 150MG BASE	N87228	001	
PUREPAC PHARM	EQ 150MG BASE	N80886	001	
TEVA	EQ 150MG BASE	N87504	001	Jan 13, 1982
WATSON LABS	EQ 150MG BASE	N87979	001	Dec 21, 1982
	EQ 300MG BASE	N88030	001	Dec 21, 1982

CHLOROQUINE PHOSPHATE; PRIMAQUINE PHOSPHATE

TABLET; ORAL

ARALEN PHOSPHATE W/ PRIMAQUINE PHOSPHATE

SANOFI SYNTHELABO	EQ 300MG BASE;EQ 45MG BASE	N14860	002	
-------------------	----------------------------	--------	-----	--

CHLOROTHIAZIDE

TABLET; ORAL

CHLOROTHIAZIDE

ABC HOLDING	250MG	N85569	001	
LEDERLE	250MG	N86940	001	
	500MG	N86938	001	
SANDOZ	250MG	N85485	001	
WATSON LABS	250MG	N85165	001	
	250MG	N85173	001	
	250MG	N86795	001	Aug 15, 1983
	500MG	N84026	001	Sep 01, 1982
	500MG	N86796	001	Aug 15, 1983
DIURIL				
OVATION PHARMS	250MG	N11145	004	
	500MG	N11145	002	

CHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDOCLOR-150

MERCK	150MG;250MG	N16016	001	
-------	-------------	--------	-----	--

ALDOCLOR-250

MERCK	250MG;250MG	N16016	002	
-------	-------------	--------	-----	--

METHYLDOPA AND CHLOROTHIAZIDE

PAR PHARM	150MG;250MG	N70783	001	Nov 06, 1987
	250MG;250MG	N70654	001	Nov 06, 1987

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CHLOROTHIAZIDE AND RESERPINE

WEST WARD	250MG;0.125MG	N88557	001	Dec 22, 1983
	500MG;0.125MG	N88365	001	Dec 22, 1983

CHLOROTHIAZIDE W/ RESERPINE

WATSON LABS	250MG;0.125MG	N84853	001	
	500MG;0.125MG	N88151	001	Jun 09, 1983

CHLOROTHIAZIDE-RESERPINE

MYLAN	250MG;0.125MG	N87744	001	May 06, 1982
	500MG;0.125MG	N87745	001	May 06, 1982

CHLOROTRIANISENE

CAPSULE; ORAL

CHLOROTRIANISENE

BANNER PHARMACAPS	12MG	N84652	001	
-------------------	------	--------	-----	--

TACE

AVENTIS PHARMS	12MG	N08102	004	
	25MG	N11444	001	
	72MG	N16235	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 56 (of 263)

CHLORPHENESIN CARBAMATETABLET; ORAL
MAOLATE

PHARMACIA AND UPJOHN 400MG

N14217 002

CHLORPHENIRAMINE MALEATECAPSULE, EXTENDED RELEASE; ORAL
CHLORPHENIRAMINE MALEATE

SANDOZ 12MG

N70797 001

Aug 12, 1988

TELDRIN

GLAXOSMITHKLINE 8MG

N17369 001

12MG

N17369 002

INJECTABLE; INJECTION

CHLORPHENIRAMINE MALEATE

BEL MAR 10MG/ML

N80821 001

ELKINS SINN 10MG/ML

N80797 001

STERIS 10MG/ML

N83593 001

10MG/ML

N86096 001

100MG/ML

N86095 001

CHLOR-TRIMETON

SCHERING PLOUGH 10MG/ML

N08826 001

100MG/ML

N08794 001

PYRIDAMAL 100

BEL MAR 100MG/ML

N83733 001

SYRUP; ORAL

CHLORPHENIRAMINE MALEATE

PHARM ASSOC 2MG/5ML

N87520 001

Feb 10, 1982

CHLOR-TRIMETON

SCHERING 2MG/5ML

N06921 006

TABLET; ORAL

ANTAGONATE

BAYER PHARMS 4MG

N83381 001

CHLORPHENIRAMINE MALEATE

ANABOLIC 4MG

N83078 001

BELL PHARMA 4MG

N83062 001

ELKINS SINN 4MG

N80938 001

IMPAX LABS 4MG

N80809 001

IVAX PHARMS 4MG

N80779 001

KV PHARM 4MG

N87164 001

LEDERLE 4MG

N86941 001

MUTUAL PHARM 4MG

N80700 001

NEWTRON PHARMS 4MG

N86519 001

PANRAY 4MG

N83243 001

PHARMAVITE 4MG

N85104 001

PHARMERAL 4MG

N83753 001

PIONEER PHARMS 4MG

N88556 001

Jul 13, 1984

PUREPAC PHARM 4MG

N86306 001

PVT FORM 4MG

N80786 001

ROXANE 4MG

N80626 001

SANDOZ 4MG

N80961 001

VITARINE 4MG

N85837 001

WATSON LABS 4MG

N80696 001

4MG

N80791 001

4MG

N85139 001

WEST WARD 4MG

N83787 001

CHLOR-TRIMETON

SCHERING 4MG

N06921 002

KLOROMIN

HALSEY 4MG

N83629 001

PHENETRON

LANNETT 4MG

N80846 001

TABLET, EXTENDED RELEASE; ORAL

CHLOR-TRIMETON

SCHERING PLOUGH 8MG

N07638 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 57 (of 263)

CHLORPHENIRAMINE MALEATE

TABLET, EXTENDED RELEASE; ORAL				
EFIDAC 24 CHLORPHENIRAMINE MALEATE				
ALZA	16MG	N19746	002	Nov 18, 1994

CHLORPHENIRAMINE MALEATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL				
CHLOROHENIRAMINE MALEATE AND PHENYLPROPANOLAMINE HYDROCHLORIDE				
WATSON LABS	12MG;75MG	N88681	001	Sep 29, 1987
CHLORPHENIRAMINE MALEATE AND PHENYLPROPANOLAMINE HYDROCHLORIDE				
SANDOZ	12MG;75MG	N88940	001	Jan 26, 1989
COLD CAPSULE IV				
GRAHAM DM	12MG;75MG	N18793	001	Apr 25, 1985
COLD CAPSULE V				
GRAHAM DM	8MG;75MG	N18794	001	Apr 23, 1985
CONTAC 12 HOUR				
GLAXOSMITHKLINE	8MG;75MG	N18099	001	
DRIZE				
ASCHER	12MG;75MG	N88359	001	Feb 13, 1986
ORNADE				
GLAXOSMITHKLINE	12MG;75MG	N12152	004	
PHENYLPROPANOLAMINE HYDROCHLORIDE W/ CHLORPHENIRAMINE MALEATE				
CENT PHARMS	8MG;75MG	N18809	001	May 07, 1984
TABLET, EXTENDED RELEASE; ORAL				
CONTAC				
NOVARTIS	12MG;75MG	N19613	001	Jun 13, 1986
DEMAZIN				
SCHERING PLOUGH	4MG;25MG	N18556	001	May 14, 1984
TRIAMINIC-12				
NOVARTIS	12MG;75MG	N18115	001	

CHLORPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL				
CODIMAL-L.A. 12				
SCHWARZ PHARMA	12MG;120MG	N18935	001	Apr 15, 1985
ISOCLOR				
FISONS	8MG;120MG	N18747	001	Mar 06, 1986
PSEUDOEPHEDRINE HYDROCHLORIDE AND CHLORPHENIRAMINE MALEATE				
CENT PHARMS	8MG;120MG	N19428	001	Aug 02, 1988
GRAHAM DM	8MG;120MG	N18844	001	Mar 20, 1985
	12MG;120MG	N18843	001	Mar 18, 1985
KV PHARM	12MG;120MG	N71455	001	Mar 01, 1989

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL				
PENNTUSS				
FISONS	EQ 4MG MALEATE/5ML;EQ 10MG BASE/5ML	N18928	001	Aug 14, 1985

CHLORPHENIRAMINE POLISTIREX; PHENYLPROPANOLAMINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL				
CORSYM				
UCB	EQ 4MG MALEATE/5ML;EQ 37.5MG HCL/5ML	N18050	001	Jan 04, 1984

CHLORPHENTERMINE HYDROCHLORIDE

TABLET; ORAL				
PRE-SATE				
PARKE DAVIS	EQ 65MG BASE	N14696	001	

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL				
THORAZINE				
GLAXOSMITHKLINE	30MG	N11120	016	
	75MG	N11120	017	
	150MG	N11120	018	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 58 (of 263)

CHLORPROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

MORTON GROVE	30MG/ML	N87032	001	Jul 08, 1982
	100MG/ML	N87053	001	

INJECTABLE; INJECTION

CHLORPROMAZINE HYDROCHLORIDE

AM PHARM PARTNERS	25MG/ML	N84911	001	
MARSAM PHARMS LLC	25MG/ML	N89563	001	Apr 15, 1988
STERIS	25MG/ML	N80365	001	
	25MG/ML	N85591	001	
WYETH AYERST	25MG/ML	N80370	001	

SYRUP; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ALPHARMA US PHARMS	10MG/5ML	N86712	001	
--------------------	----------	--------	-----	--

TABLET; ORAL

CHLORPROMAZINE HYDROCHLORIDE

ABBOTT	10MG	N84414	001	
	25MG	N84415	001	
	50MG	N84411	001	
	100MG	N84412	001	
	200MG	N84413	001	
IVAX PHARMS	10MG	N83549	001	
	25MG	N83549	002	
	50MG	N83549	003	
	100MG	N83574	001	
	200MG	N83575	001	
KV PHARM	10MG	N85750	002	Jan 04, 1982
	25MG	N85751	001	
	50MG	N85484	001	
	100MG	N85752	001	
	200MG	N85748	002	Jan 04, 1982
LEDERLE	10MG	N84803	001	
	25MG	N84801	001	
	50MG	N84800	001	
	100MG	N84789	001	
	200MG	N84802	001	
PUREPAC PHARM	10MG	N80403	004	
	25MG	N80403	001	
	50MG	N80403	002	
	100MG	N80403	003	
	200MG	N80403	005	
PVT FORM	25MG	N80340	001	
	50MG	N80340	002	
	200MG	N80340	003	
ROXANE	10MG	N85331	001	
	25MG	N85331	002	
	50MG	N85331	003	
	100MG	N85331	004	
	200MG	N85331	005	
VANGARD	10MG	N88038	001	Aug 16, 1982
	25MG	N87645	001	
	50MG	N87646	001	
WATSON LABS	10MG	N85959	001	
	25MG	N85956	001	
	50MG	N85960	001	
	100MG	N85957	001	
	200MG	N85958	001	
WEST WARD	10MG	N87783	001	Sep 16, 1982
	25MG	N87865	001	Sep 16, 1982
	50MG	N87878	001	Sep 15, 1982
	100MG	N87884	001	Sep 15, 1982
	200MG	N87880	001	Sep 16, 1982
PROMAPAR				
PARKE DAVIS	10MG	N86886	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 59 (of 263)

CHLORPROMAZINE HYDROCHLORIDE

TABLET; ORAL

PROMAPAR

PARKE DAVIS

25MG

N84423 001

50MG

N86887 001

100MG

N86888 001

200MG

N86885 001

THORAZINE

GLAXOSMITHKLINE

10MG

N09149 002

25MG

N09149 007

50MG

N09149 013

100MG

N09149 018

200MG

N09149 020

CHLORPROPAMIDE

TABLET; ORAL

CHLORPROPAMIDE

BARR

100MG

N88812 001 Oct 19, 1984

100MG

N89446 001 Nov 17, 1986

250MG

N88813 001 Oct 19, 1984

250MG

N89447 001 Nov 17, 1986

CLONMEL HLTHCARE

100MG

N89561 001 Sep 04, 1987

250MG

N89562 001 Sep 04, 1987

DURAMED PHARMS BARR

100MG

N88918 001 Oct 16, 1984

250MG

N88919 001 Oct 16, 1984

HALSEY

100MG

N89321 001 Jan 16, 1986

250MG

N88662 001 Jan 09, 1986

IVAX PHARMS

100MG

N88840 001 Oct 25, 1984

250MG

N87353 001

SANDOZ

250MG

N84669 001

SUPERPHARM

100MG

N88694 001 Sep 17, 1984

250MG

N88695 001 Sep 17, 1984

TEVA

100MG

N88768 001 Oct 11, 1984

USL PHARMA

100MG

N88708 001 Aug 30, 1984

250MG

N88709 001 Aug 30, 1984

WATSON LABS

100MG

N86865 001 Sep 24, 1984

100MG

N88608 001 Apr 12, 1984

250MG

N86866 001

250MG

N88568 001 Apr 12, 1984

GLUCAMIDE

TEVA

250MG

N88641 001 Oct 11, 1984

CHLORPROTHIXENE

CONCENTRATE; ORAL

TARACTAN

ROCHE

100MG/5ML

N16149 002

INJECTABLE; INJECTION

TARACTAN

ROCHE

12.5MG/ML

N12487 001

TABLET; ORAL

TARACTAN

ROCHE

10MG

N12486 005

25MG

N12486 004

50MG

N12486 003

100MG

N12486 001

CHLORTETRACYCLINE HYDROCHLORIDE

OINTMENT; OPHTHALMIC

AUREOMYCIN

LEDERLE

1%

N50404 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 60 (of 263)

CHLORTHALIDONETABLET; ORAL
CHLORTHALIDONE

ABBOTT	25MG	N87364	001	
	50MG	N87384	001	
ASCOT	25MG	N87698	001	Oct 20, 1982
	50MG	N87699	001	Oct 20, 1982
CLONMEL HLTHCARE	25MG	N87451	001	
	50MG	N87450	001	
IVAX PHARMS	25MG	N87555	001	
	25MG	N88164	001	Jan 09, 1984
	50MG	N87176	001	
	50MG	N87947	001	Feb 27, 1984
KV PHARM	25MG	N87311	001	
	50MG	N87312	001	
MUTUAL PHARM	25MG	N87292	001	
	25MG	N89285	001	Jul 21, 1986
	25MG	N89738	001	Sep 19, 1988
	50MG	N87293	001	
	50MG	N89286	001	Jul 21, 1986
	50MG	N89739	001	Sep 19, 1988
PIONEER PHARMS	50MG	N89591	001	Jul 21, 1988
PUREPAC PHARM	25MG	N88139	001	Jul 16, 1986
	50MG	N88140	001	Aug 11, 1983
SANDOZ	25MG	N87380	001	
	50MG	N87118	001	
	50MG	N87381	001	
SUPERPHARM	25MG	N87473	001	Feb 09, 1983
	50MG	N87247	001	Feb 09, 1983
TEVA	50MG	N88651	001	May 30, 1985
USL PHARMA	25MG	N89051	001	Jun 01, 1987
	50MG	N89052	001	Jun 01, 1987
VANGARD	25MG	N88012	001	Jul 14, 1982
	50MG	N88073	001	Mar 25, 1983
WARNER CHILCOTT	25MG	N87515	001	Jan 24, 1983
	50MG	N87516	001	Feb 09, 1983
WATSON LABS	25MG	N87050	001	
	25MG	N87100	001	
	25MG	N87296	001	
	25MG	N87706	001	
	50MG	N87029	001	
	50MG	N87082	001	
	50MG	N87521	001	
	50MG	N87689	001	
HYGROTON				
AVENTIS PHARMS	25MG	N12283	004	
	50MG	N12283	003	
THALITONE				
MONARCH PHARMS	25MG	N19574	002	Feb 12, 1992
	25MG	N88051	001	Nov 12, 1982

CHLORTHALIDONE; CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HYDROCHLORIDE AND CHLORTHALIDONE

PAR PHARM	15MG;0.1MG	N71179	001	Dec 16, 1987
	15MG;0.2MG	N71178	001	Dec 16, 1987
	15MG;0.3MG	N71142	001	Dec 16, 1987
COMBIPRES				
BOEHRINGER INGELHEIM	15MG;0.1MG	N17503	001	
	15MG;0.2MG	N17503	002	
	15MG;0.3MG	N17503	003	Apr 10, 1984

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 61 (of 263)

CHLORTHALIDONE; METOPROLOL TARTRATECAPSULE; ORAL
LOPRESSIDONE

NOVARTIS

25MG;100MG

N19451 001

Dec 31, 1987

25MG;200MG

N19451 002

Dec 31, 1987

CHLORTHALIDONE; RESERPINE

TABLET; ORAL

DEMI-REGROTON

AVENTIS PHARMS

25MG;0.125MG

N15103 002

REGROTON

AVENTIS PHARMS

50MG;0.25MG

N15103 001

CHLORZOXAZONE

TABLET; ORAL

CHLORZOXAZONE

PIONEER PHARMS

250MG

N89592 001

Jan 06, 1989

500MG

N89948 001

Jan 06, 1989

WATSON LABS

250MG

N86901 001

250MG

N86948 001

Aug 09, 1982

500MG

N81019 001

Jul 29, 1991

PARAFLEX

ORTHO MCNEIL PHARM

250MG

N11300 003

CHOLESTYRAMINE

BAR, CHEWABLE; ORAL

CHOLYBAR

PARKE DAVIS

EQ 4GM RESIN/BAR

N71621 001

May 26, 1988

EQ 4GM RESIN/BAR

N71739 001

May 26, 1988

POWDER; ORAL

CHOLESTYRAMINE

IVAX PHARMS

EQ 4GM RESIN/PACKET

N74771 001

Jul 09, 1997

EQ 4GM RESIN/SCOOPFUL

N74771 002

Jul 09, 1997

TABLET; ORAL

QUESTRAN

APOTHECON

EQ 800MG RESIN

N73403 002

Dec 27, 1999

EQ 1GM RESIN

N73403 001

Apr 28, 1994

CHORIOGONADOTROPIN ALFA

INJECTABLE; INJECTION

OVIDREL

SERONO INC

0.25MG/VIAL

N21149 001

Sep 20, 2000

CHROMIC CHLORIDE

INJECTABLE; INJECTION

CHROMIC CHLORIDE

AM PHARM PARTNERS

EQ 0.004MG CHROMIUM/ML

N19271 001

May 05, 1987

CHYMOPAPAIN

INJECTABLE; INJECTION

CHYMODIACTIN

ABBOTT

4,000 UNITS/VIAL

N18663 002

Aug 21, 1984

10,000 UNITS/VIAL **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N18663 001

Nov 10, 1982

DISCASE

ABBOTT

12,500 UNITS/VIAL

N18625 001

Jan 18, 1984

CHYMOTRYPSIN

FOR SOLUTION; OPHTHALMIC

ALPHA CHYMAR

SOLA BARNES HIND

750 UNITS/VIAL

N11837 001

DISCONTINUED DRUG PRODUCT LIST

6 - 62 (of 263)

CHYMOTRYPSIN

FOR SOLUTION; OPHTHALMIC

CATARASE

CIBA	300 UNITS/VIAL	N16938	001
------	----------------	--------	-----

NOVARTIS	150 UNITS/VIAL	N18121	001
----------	----------------	--------	-----

ZOLYSE

ALCON	750 UNITS/VIAL	N11903	001
-------	----------------	--------	-----

CIMETIDINE

SUSPENSION; ORAL

TAGAMET HB 200

GLAXOSMITHKLINE	200MG/20ML	N20951	001	Jul 09, 1999
-----------------	------------	--------	-----	--------------

TABLET; ORAL

CIMETIDINE

IVAX PHARMS	200MG	N74401	001	May 30, 1995
-------------	-------	--------	-----	--------------

	300MG	N74401	002	May 30, 1995
--	-------	--------	-----	--------------

	400MG	N74401	003	May 30, 1995
--	-------	--------	-----	--------------

LEK PHARMS	100MG	N75122	001	Jun 19, 1998
------------	-------	--------	-----	--------------

	200MG	N74250	001	Jun 29, 1995
--	-------	--------	-----	--------------

PERRIGO	100MG	N74972	001	Jun 19, 1998
---------	-------	--------	-----	--------------

ROXANE	300MG	N74361	001	Dec 23, 1994
--------	-------	--------	-----	--------------

	400MG	N74361	002	Dec 23, 1994
--	-------	--------	-----	--------------

	800MG	N74371	001	Dec 23, 1994
--	-------	--------	-----	--------------

TAGAMET HB

GLAXOSMITHKLINE	100MG	N20238	001	Jun 19, 1995
-----------------	-------	--------	-----	--------------

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

HOSPIRA	EQ 300MG BASE/2ML	N74296	001	Mar 28, 1997
---------	-------------------	--------	-----	--------------

	EQ 300MG BASE/2ML	N74412	001	Mar 28, 1997
--	-------------------	--------	-----	--------------

	EQ 300MG BASE/2ML	N74422	001	Jan 31, 1995
--	-------------------	--------	-----	--------------

LUITPOLD	EQ 300MG BASE/2ML	N74353	001	Dec 20, 1994
----------	-------------------	--------	-----	--------------

CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

HOSPIRA	EQ 90MG BASE/100ML	N74468	005	Dec 29, 1994
---------	--------------------	--------	-----	--------------

	EQ 120MG BASE/100ML	N74468	006	Dec 29, 1994
--	---------------------	--------	-----	--------------

	EQ 180MG BASE/100ML	N74468	003	Dec 29, 1994
--	---------------------	--------	-----	--------------

	EQ 240MG BASE/100ML	N74468	004	Dec 29, 1994
--	---------------------	--------	-----	--------------

	EQ 360MG BASE/100ML	N74468	001	Dec 29, 1994
--	---------------------	--------	-----	--------------

	EQ 480MG BASE/100ML	N74468	002	Dec 29, 1994
--	---------------------	--------	-----	--------------

TAGAMET

GLAXOSMITHKLINE	EQ 300MG BASE/2ML	N17939	002
-----------------	-------------------	--------	-----

TAGAMET HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

GLAXOSMITHKLINE	EQ 6MG BASE/ML	N19434	001	Oct 31, 1985
-----------------	----------------	--------	-----	--------------

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

ROXANE	EQ 300MG BASE/5ML	N74541	001	Aug 05, 1997
--------	-------------------	--------	-----	--------------

TAGAMET

GLAXOSMITHKLINE	EQ 300MG BASE/5ML	N17924	001
-----------------	-------------------	--------	-----

CINOXACIN

CAPSULE; ORAL

CINOBAC

LILLY	250MG	N18067	001
-------	-------	--------	-----

	500MG	N18067	002
--	-------	--------	-----

CINOXACIN

TEVA	250MG	N73005	001	Feb 28, 1992
------	-------	--------	-----	--------------

	500MG	N73006	001	Feb 28, 1992
--	-------	--------	-----	--------------

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPRO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

BAYER PHARMS	200MG/100ML	N19858	001	Dec 26, 1990
--------------	-------------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 63 (of 263)

CISAPRIDE MONOHYDRATE

SUSPENSION; ORAL PROPULSID				
JANSSEN PHARMA	EQ 1MG BASE/ML	N20398	001	Sep 15, 1995
TABLET; ORAL PROPULSID				
JANSSEN PHARMA	EQ 10MG BASE	N20210	001	Jul 29, 1993
	EQ 20MG BASE	N20210	002	Dec 23, 1993
TABLET, ORALLY DISINTEGRATING; ORAL PROPULSID QUICKSOLV				
JANSSEN PHARMA	EQ 20MG BASE	N20767	001	Nov 07, 1997

CISPLATIN

INJECTABLE; INJECTION PLATINOL-AQ				
BRISTOL MYERS	0.5MG/ML	N18057	003	Jul 18, 1984

CITALOPRAM HYDROBROMIDE

TABLET; ORAL CELEXA				
FOREST LABS	EQ 60MG BASE	N20822	004	Jul 17, 1998

CITRIC ACID; UREA, C-13

TABLET, FOR SOLUTION, FOR SOLUTION; ORAL IDKIT:HP				
ORIDION	N/A, 4GM; 75MG, N/A	N21314	001	Dec 17, 2002

CLARITHROMYCIN

FOR SUSPENSION; ORAL BIAXIN				
ABBOTT	187MG/5ML	N50698	003	Sep 30, 1998

CLEMASTINE FUMARATE

SYRUP; ORAL TAVIST				
NOVARTIS	EQ 0.5MG BASE/5ML	N18675	001	Jun 28, 1985
TABLET; ORAL CLEMASTINE FUMARATE				
TEVA	1.34MG	N73282	001	Jan 31, 1992
TAVIST				
NOVARTIS	2.68MG	N17661	001	
TAVIST-1				
NOVARTIS	1.34MG	N17661	002	
	1.34MG	N17661	003	Aug 21, 1992

CLEMASTINE FUMARATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL TAVIST-D				
NOVARTIS	EQ 1MG BASE; 75MG	N18298	001	Dec 15, 1982
	1.34MG; 75MG	N18298	002	Aug 21, 1992
	1.34MG; 75MG	N20640	001	Aug 09, 1996

CLIDINIUM BROMIDE

CAPSULE; ORAL QUARZAN				
ROCHE	2.5MG	N10355	001	
	5MG	N10355	002	

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL CLEOCIN				
PHARMACIA AND UPJOHN	EQ 75MG BASE	N61809	001	
	EQ 150MG BASE	N61809	002	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 64 (of 263)

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLINDAMYCIN HYDROCHLORIDE

WATSON LABS

EQ 75MG BASE

N63082 001 Jul 31, 1991

CLINDAMYCIN PALMITATE HYDROCHLORIDE

FOR SOLUTION; ORAL

CLEOCIN

PHARMACIA AND UPJOHN EQ 75MG BASE/5ML

N61827 001

CLINDAMYCIN PHOSPHATE

CREAM; VAGINAL

CLEOCIN

PHARMACIA AND UPJOHN EQ 2% BASE

N50680 001 Aug 11, 1992

INJECTABLE; INJECTION

CLEOCIN PHOSPHATE

PHARMACIA AND UPJOHN EQ 150MG BASE/ML

N61839 001

CLINDAMYCIN PHOSPHATE

AM PHARM PARTNERS EQ 150MG BASE/ML

N62747 001 Jun 03, 1988

ASTRAZENECA EQ 150MG BASE/ML

N62928 001 Feb 13, 1989

BAXTER HLTHCARE EQ 150MG BASE/ML

N62806 001 Oct 15, 1987

EQ 150MG BASE/ML

N62953 001 Apr 21, 1988

EQ 150MG BASE/ML

N63068 001 Aug 28, 1989

BEDFORD EQ 150MG BASE/ML

N63163 001 Jun 30, 1994

BRISTOL MYERS SQUIBB EQ 150MG BASE/ML

N62908 001 Feb 01, 1989

LOCH EQ 150MG BASE/ML

N62905 001 May 09, 1988

MARSAM PHARMS LLC EQ 150MG BASE/ML

N62913 001 Oct 20, 1988

SICOR PHARMS EQ 150MG BASE/ML

N63041 001 Dec 29, 1989

EQ 150MG BASE/ML

N63282 001 May 29, 1992

SOLOPAK EQ 150MG BASE/ML

N62819 001 Mar 15, 1988

EQ 150MG BASE/ML

N62852 001 Mar 17, 1988

STERIS EQ 150MG BASE/ML

N62900 001 Jun 08, 1988

EQ 150MG BASE/ML

N63079 001 Mar 05, 1990

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%

AM PHARM PARTNERS EQ 12MG BASE/ML

N50636 001 Dec 22, 1989

CLINDAMYCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT EQ 6MG BASE/ML

N65027 001 Jun 29, 2001

EQ 12MG BASE/ML

N65027 002 Jun 29, 2001

EQ 18MG BASE/ML

N65027 003 Jun 29, 2001

BAXTER HLTHCARE EQ 6MG BASE/ML

N50648 001 Dec 29, 1989

EQ 12MG BASE/ML

N50648 002 Dec 29, 1989

EQ 900MG BASE/100ML

N50648 003 Dec 29, 1989

SOLUTION; TOPICAL

CLEOCIN T

PHARMACIA AND UPJOHN EQ 1% BASE

N62363 001 Feb 08, 1982

CLINDAMYCIN PHOSPHATE

COPLY PHARM EQ 1% BASE

N62944 001 Jan 11, 1989

TEVA EQ 1% BASE

N62930 001 Jun 28, 1989

CLIOQUINOL; NYSTATIN

OINTMENT; TOPICAL

NYSTAFORM

BAYER PHARMS 10MG/GM;100,000 UNITS/GM

N50235 001

CLOFAZIMINE

CAPSULE; ORAL

LAMPRENE

NOVARTIS

50MG

N19500 002 Dec 15, 1986

100MG

N19500 001 Dec 15, 1986

CLOFIBRATE

CAPSULE; ORAL

ATROMID-S

WYETH AYERST

500MG

N16099 002

DISCONTINUED DRUG PRODUCT LIST

6 - 65 (of 263)

CLOFIBRATE

CAPSULE; ORAL
CLOFIBRATE

SANDOZ	500MG	N72191	001	May 02, 1988
WATSON LABS	500MG	N71603	001	Sep 18, 1987

CLONAZEPAM

TABLET; ORAL
KLONOPIN

ROCHE	0.125MG	N17533	005	Apr 09, 1997
	0.25MG	N17533	006	Apr 09, 1997

CLONIDINE HYDROCHLORIDE

TABLET; ORAL
CLONIDINE HYDROCHLORIDE

AM THERAP	0.1MG	N70881	001	Jul 08, 1986
	0.2MG	N70882	001	Jul 08, 1986
	0.3MG	N70883	001	Jul 08, 1986
DURAMED PHARMS BARR	0.1MG	N71103	001	Aug 14, 1986
	0.2MG	N71102	001	Aug 14, 1986
	0.3MG	N71101	001	Aug 14, 1986
INTERPHARM	0.1MG	N71252	001	Oct 01, 1986
	0.2MG	N71253	001	Oct 01, 1986
	0.3MG	N71254	001	Oct 01, 1986
PAR PHARM	0.1MG	N70461	001	Jul 08, 1986
	0.2MG	N70460	001	Jul 08, 1986
	0.3MG	N70459	001	Jul 08, 1986
TEVA	0.1MG	N70747	001	Jul 08, 1986
	0.2MG	N70702	001	Jul 08, 1986
	0.3MG	N70659	001	Jul 08, 1986
WARNER CHILCOTT	0.1MG	N72138	001	Jun 13, 1988
	0.2MG	N72139	001	Jun 13, 1988
	0.3MG	N72140	001	Jun 13, 1988
WATSON LABS	0.1MG	N70395	001	Mar 23, 1987
	0.2MG	N70396	001	Mar 23, 1987
	0.3MG	N70397	001	Mar 23, 1987

CLORAZEPATE DIPOTASSIUM

CAPSULE; ORAL
CLORAZEPATE DIPOTASSIUM

ABLE	3.75MG	N71777	001	Jul 14, 1987
	7.5MG	N71778	001	Jul 14, 1987
	15MG	N71779	001	Jul 14, 1987
AM THERAP	3.75MG	N71429	001	Jun 23, 1987
	7.5MG	N71430	001	Jun 23, 1987
	15MG	N71431	001	Jun 23, 1987
CLONMEL HLTHCARE	3.75MG	N71742	001	Dec 14, 1987
	7.5MG	N71743	001	Dec 14, 1987
	15MG	N71744	001	Dec 14, 1987
GD SEARLE LLC	3.75MG	N71727	001	Dec 18, 1987
	7.5MG	N71728	001	Dec 18, 1987
	15MG	N71729	001	Dec 18, 1987
PUREPAC PHARM	3.75MG	N71924	001	Apr 25, 1988
	7.5MG	N71925	001	Apr 25, 1988
	15MG	N71926	001	Apr 25, 1988
QUANTUM PHARMICS	3.75MG	N71549	001	Sep 12, 1988
	7.5MG	N71550	001	Sep 12, 1988
	15MG	N71522	001	Sep 12, 1988
SANDOZ	7.5MG	N72220	001	Aug 26, 1988
USL PHARMA	3.75MG	N71242	001	Jun 23, 1987
	7.5MG	N71243	001	Jun 23, 1987
	15MG	N71244	001	Jun 23, 1987
WARNER CHILCOTT	3.75MG	N71774	001	Mar 01, 1988
	7.5MG	N71775	001	Mar 01, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 66 (of 263)

CLORAZEPATE DIPOTASSIUM

CAPSULE; ORAL

CLORAZEPATE DIPOTASSIUM

WARNER CHILCOTT	15MG	N71776	001	Mar 01, 1988
WATSON LABS	3.75MG	N71878	001	Mar 15, 1988
	7.5MG	N71879	001	Mar 15, 1988
	15MG	N71860	001	Mar 15, 1988

TRANXENE

OVATION PHARMS	3.75MG	N17105	001	
	7.5MG	N17105	002	
	15MG	N17105	003	

TABLET; ORAL

CLORAZEPATE DIPOTASSIUM

ABLE	3.75MG	N71780	001	Jun 26, 1987
	7.5MG	N71781	001	Jun 26, 1987
	15MG	N71782	001	Jun 26, 1987
AM THERAP	3.75MG	N71747	001	Jun 23, 1987
	7.5MG	N71748	001	Jun 23, 1987
	15MG	N71749	001	Jun 23, 1987
LEDERLE	3.75MG	N72013	001	Dec 15, 1987
	7.5MG	N72014	001	Dec 15, 1987
	15MG	N72015	001	Dec 15, 1987
PUREPAC PHARM	3.75MG	N72330	001	Aug 08, 1988
	7.5MG	N72331	001	Aug 08, 1988
	15MG	N72332	001	Aug 08, 1988
QUANTUM PHARMICS	3.75MG	N71730	001	Oct 26, 1987
	7.5MG	N71731	001	Oct 26, 1987
	15MG	N71702	001	Oct 26, 1987
SANDOZ	3.75MG	N72512	001	May 11, 1990
WARNER CHILCOTT	3.75MG	N71828	001	Mar 03, 1988
	7.5MG	N71829	001	Mar 03, 1988
	15MG	N71830	001	Mar 03, 1988

CLOTRIMAZOLE

CREAM; TOPICAL

LOTRIMIN

SCHERING PLOUGH	1%	N17619	001	
MYCELEX				
BAYER PHARMS	1%	N18183	001	

LOTION; TOPICAL

LOTRIMIN

SCHERING	1%	N18813	001	Feb 17, 1984
----------	----	--------	-----	--------------

SOLUTION; TOPICAL

LOTRIMIN

SCHERING PLOUGH	1%	N17613	001	
-----------------	----	--------	-----	--

TABLET; VAGINAL

MYCELEX-G

BAYER PHARMS	500MG	N19069	001	Apr 19, 1985
--------------	-------	--------	-----	--------------

CLOXACILLIN SODIUM

CAPSULE; ORAL

CLOXACILLIN SODIUM

APOTHECON	EQ 250MG BASE	N61452	001	
	EQ 500MG BASE	N61452	002	

CLOXAPEN

GLAXOSMITHKLINE	EQ 250MG BASE	N62233	001	
	EQ 500MG BASE	N62233	002	

FOR SOLUTION; ORAL

CLOXACILLIN SODIUM

TEVA	EQ 125MG BASE/5ML	N62978	001	Apr 06, 1989
TEGOPEN				
APOTHECON	EQ 125MG BASE/5ML	N50192	001	
	EQ 125MG BASE/5ML	N61453	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 67 (of 263)

CLOZAPINE

TABLET; ORAL
CLOZAPINE
SANDOZ

25MG
100MG

N74546 001 Aug 30, 1996
N74546 002 Aug 30, 1996

COBALT CHLORIDE, CO-60; CYANOCOBALAMIN; CYANOCOBALAMIN, CO-60; INTRINSIC FACTOR

N/A; N/A
RUBRATOPE-60 KIT
BRACCO

N/A;N/A;N/A;N/A

N16090 001

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENERGAN VC W/ CODEINE

WYETH AYERST 10MG/5ML;5MG/5ML;6.25MG/5ML

N08306 005 Apr 02, 1984

PHERAZINE VC W/ CODEINE

HALSEY 10MG/5ML;5MG/5ML;6.25MG/5ML

N88870 001 Mar 02, 1987

PROMETHAZINE VC W/ CODEINE

CENCI 10MG/5ML;5MG/5ML;6.25MG/5ML

N88816 001 Nov 22, 1985

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENERGAN W/ CODEINE

WYETH AYERST 10MG/5ML;6.25MG/5ML

N08306 004 Apr 02, 1984

PHERAZINE W/ CODEINE

HALSEY 10MG/5ML;6.25MG/5ML

N88739 001 Dec 23, 1988

PROMETHAZINE W/ CODEINE

CENCI 10MG/5ML;6.25MG/5ML

N88814 001 Nov 22, 1985

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL

ACTIFED W/ CODEINE

GLAXOSMITHKLINE 10MG/5ML;30MG/5ML;1.25MG/5ML

N12575 003 Apr 04, 1984

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES W/ CODEINE

CENCI 10MG/5ML;30MG/5ML;1.25MG/5ML

N89018 001 Jul 23, 1986

COLCHICINE; PROBENECID

TABLET; ORAL

COLBENEMID

MERCK 0.5MG;500MG

N12383 001

PROBEN-C

WATSON LABS 0.5MG;500MG

N85552 001

PROBENECID AND COLCHICINE

BEECHAM 0.5MG;500MG

N84321 001

IMPAX LABS 0.5MG;500MG

N83720 002

SANDOZ 0.5MG;500MG

N86130 001

PROBENECID W/ COLCHICINE

LEDERLE 0.5MG;500MG

N86954 001

WATSON LABS 0.5MG;500MG

N83221 001

COLESEVELAM HYDROCHLORIDE

CAPSULE; ORAL

WELCHOL

SANKYO 375MG

N21141 001 May 26, 2000

COLISTIN SULFATE

SUSPENSION; ORAL

COLY-MYCIN S

PARKE DAVIS EQ 25MG BASE/5ML

N50355 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 68 (of 263)

COPPERINTRAUTERINE DEVICE; INTRAUTERINE
CU-7

GD SEARLE LLC 89MG N17408 001

TATUM-T

GD SEARLE LLC 120MG N18205 001

CORTICOTROPIN

INJECTABLE; INJECTION

ACTH

PARKEDALE 25 UNITS/VIAL N08317 002

40 UNITS/VIAL N08317 004

ACTHAR

AVENTIS 25 UNITS/VIAL N07504 002

40 UNITS/VIAL N07504 003

CORTICOTROPIN

ORGANICS LAGRANGE 40 UNITS/ML N10831 001

80 UNITS/ML N10831 002

STERIS 40 UNITS/VIAL N88772 001

Nov 21, 1984

H.P. ACTHAR GEL

QUESTCOR PHARMS 40 UNITS/ML N08372 006

PURIFIED CORTROPHIN GEL

ORGANON USA INC 40 UNITS/ML N08975 001

80 UNITS/ML N08975 002

CORTICOTROPIN-ZINC HYDROXIDE

INJECTABLE; INJECTION

CORTROPHIN-ZINC

ORGANON USA INC 40 UNITS/ML N09854 001

CORTISONE ACETATE

INJECTABLE; INJECTION

CORTISONE ACETATE

PHARMACIA AND UPJOHN 25MG/ML N08126 002

STERIS 25MG/ML N83147 003

25MG/ML N85677 001

50MG/ML N83147 004

50MG/ML N85677 002

CORTONE

MERCK 25MG/ML N07110 002

50MG/ML N07110 003

TABLET; ORAL

CORTISONE ACETATE

BARR 25MG N83471 001

ELKINS SINN 25MG N80836 001

EVERYLIFE 25MG N84246 001

HEATHER 25MG N85736 001

IMPAX LABS 25MG N09458 001

INWOOD LABS 25MG N80731 001

IVAX PHARMS 25MG N80630 001

25MG N83536 001

LANNETT 25MG N80694 001

PANRAY 5MG N08284 002

25MG N08284 001

PUREPAC PHARM 25MG N80493 001

VITARINE 25MG N80333 001

WATSON LABS 25MG N85884 001

WHITEWORTH TOWN PLSN 25MG N80341 001

CORTONE

MERCK 25MG N07750 003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 69 (of 263)

CROMOLYN SODIUMCAPSULE; INHALATION
INTAL

AVENTIS 20MG N16990 001

CAPSULE; ORAL
GASTROCRON

UCB 100MG N19188 001 Dec 22, 1989

SOLUTION/DROPS; OPHTHALMIC
CROMOPTIC

KING PHARMS 4% N75088 001 Apr 27, 1999

CRYPTENAMINE ACETATESINJECTABLE; INJECTION
UNITENSEN

MEDPOINTE PHARM HLC 260CSR UNIT/ML N08814 001

CRYPTENAMINE TANNATESTABLET; ORAL
UNITENSEN

MEDPOINTE PHARM HLC 260CSR UNIT N09217 001

CUPRIC SULFATEINJECTABLE; INJECTION
CUPRIC SULFATE

AM PHARM PARTNERS EQ 0.4MG COPPER/ML N19350 001 May 05, 1987

CYANOCOBALAMININJECTABLE; INJECTION
BERUBIGEN

PHARMACIA AND UPJOHN 1MG/ML N06798 001

BETALIN 12

LILLY 0.1MG/ML N80855 001

1MG/ML N80855 002

COBAVITE

STERIS 0.1MG/ML N83013 001

1MG/ML N83064 001

CYANOCOBALAMIN

AKORN 1MG/ML N87969 001 Nov 10, 1983

AM PHARM PARTNERS 0.03MG/ML N80510 003

0.1MG/ML N80510 001

1MG/ML N80510 002

1MG/ML N83075 001

AVENTIS PHARMS 1MG/ML N80564 001

BAXTER HLTHCARE 1MG/ML N80515 002

DELL LABS 0.03MG/ML N80689 001

0.1MG/ML N80689 002

1MG/ML N80689 003

LUITPOLD 0.03MG/ML N80668 001

SOLOPAK 1MG/ML N87551 001 Feb 29, 1984

STERIS 0.1MG/ML N83120 001

1MG/ML N83120 002

WARNER CHILCOTT 1MG/ML N07085 002

WYETH AYERST 0.1MG/ML N80554 001

1MG/ML N80554 002

REDISOL

MERCK 1MG/ML N06668 010

RUBIVITE

BEL MAR 0.03MG/ML N10791 004

0.05MG/ML N10791 001

0.1MG/ML N10791 002

0.12MG/ML N10791 005

1MG/ML N10791 003

RUBRAMIN PC

BRISTOL MYERS SQUIBB 0.1MG/ML N06799 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 70 (of 263)

CYANOCOBALAMIN

INJECTABLE; INJECTION

RUBRAMIN PC

BRISTOL MYERS SQUIBB 1MG/ML

N06799 004

RUVITE

SAVAGE LABS 1MG/ML

N80570 002

VI-TWEL

BERLEX 1MG/ML

N07012 002

TABLET; ORAL

CYANOCOBALAMIN

WEST WARD 1MG

N84264 001

CYANOCOBALAMIN, CO-60

CAPSULE; ORAL

RUBRATOPE-60

BRACCO 0.5-1uCi

N16090 002

CYANOCOBALAMIN; CYANOCOBALAMIN, CO-57; CYANOCOBALAMIN, CO-58

N/A; N/A

DICOPAC KIT

GE HEALTHCARE N/A;N/A;N/A

N17406 001

CYANOCOBALAMIN; CYANOCOBALAMIN, CO-57; INTRINSIC FACTOR

N/A; N/A

CYANOCOBALAMIN CO 57 SCHILLING TEST KIT

MALLINCKRODT 0.1MG;0.5uCi;60MG

N16635 001

CYANOCOBALAMIN; TANNIC ACID; ZINC ACETATE

INJECTABLE; INJECTION

DEPINAR

ARMOUR PHARM 0.5MG/ML;2.3MG/ML;1MG/ML

N11208 001

CYCLACILLIN

FOR SUSPENSION; ORAL

CYCLAPEN-W

WYETH AYERST 125MG/5ML

N50508 001

250MG/5ML

N50508 002

500MG/5ML

N50508 003

TABLET; ORAL

CYCLACILLIN

TEVA 250MG

N62895 001

Aug 04, 1988

500MG

N62895 002

Aug 04, 1988

CYCLAPEN-W

WYETH AYERST 250MG

N50509 001

500MG

N50509 002

CYCLIZINE LACTATE

INJECTABLE; INJECTION

MAREZINE

GLAXOSMITHKLINE 50MG/ML

N09495 001

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

AK-PENTOLATE

AKORN 1%

N85555 001

CYCLOPENTOLATE HYDROCHLORIDE

SOLA BARNES HIND 1%

N84150 001

1%

N84863 001

PENTOLAIR

PHARMAFAIR 0.5%

N88643 001

Feb 09, 1987

1%

N88150 001

Feb 25, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 71 (of 263)

CYCLOPHOSPHAMIDEINJECTABLE; INJECTION
CYTOXAN

BRISTOL MYERS SQUIBB	100MG/VIAL	N12142	001	
	200MG/VIAL	N12142	002	
	500MG/VIAL	N12142	003	
	1GM/VIAL	N12142	004	Aug 30, 1982
	2GM/VIAL	N12142	005	Aug 30, 1982

NEOSAR

SICOR PHARMS	100MG/VIAL	N40015	001	Apr 29, 1993
	200MG/VIAL	N40015	002	Apr 29, 1993
	500MG/VIAL	N40015	003	Apr 29, 1993
	1GM/VIAL	N40015	004	Apr 29, 1993
	2GM/VIAL	N40015	005	Apr 29, 1993

CYCLOSPORINECAPSULE; ORAL
NEORAL

NOVARTIS	50MG	N50715	003	Jul 14, 1995
----------	------	--------	-----	--------------

CYCLOTHIAZIDETABLET; ORAL
ANHYDRON

LILLY	2MG	N13157	002	
FLUIDIL				
PHARMACIA AND UPJOHN	2MG	N18173	001	

CYCRIMINE HYDROCHLORIDETABLET; ORAL
PAGITANE

LILLY	1.25MG	N08951	001	
	2.5MG	N08951	002	

CYPROHEPTADINE HYDROCHLORIDE

SYRUP; ORAL

CYPROHEPTADINE HYDROCHLORIDE

HALSEY	2MG/5ML	N89199	001	Jul 03, 1986
MORTON GROVE	2MG/5ML	N87001	001	Nov 04, 1982
NASKA	2MG/5ML	N89021	001	Dec 21, 1987
PERIACTIN				
MERCK	2MG/5ML	N13220	002	

TABLET; ORAL

CYPROHEPTADINE HYDROCHLORIDE

ABC HOLDING	4MG	N88212	001	May 26, 1983
AM THERAP	4MG	N88798	001	Feb 15, 1985
ASCOT	4MG	N87685	001	Oct 25, 1982
DURAMED PHARMS BARR	4MG	N88232	001	Oct 25, 1983
HALSEY	4MG	N89057	001	Jul 03, 1986
KV PHARM	4MG	N86737	001	
MD PHARM	4MG	N87566	001	Nov 10, 1982
MYLAN	4MG	N86678	001	
PIONEER PHARMS	4MG	N87839	001	Feb 08, 1984
SANDOZ	4MG	N86808	001	
SUPERPHARM	4MG	N87405	001	
VITARINE	4MG	N87284	001	
WATSON LABS	4MG	N85245	001	
	4MG	N86165	001	
	4MG	N86580	001	
PERIACTIN				
MERCK	4MG	N12649	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 72 (of 263)

CYSTEINE HYDROCHLORIDE

INJECTABLE; INJECTION				
CYSTEINE HYDROCHLORIDE				
HOSPIRA	7.25%	N19523	001	Oct 22, 1986

CYTARABINE

INJECTABLE; INJECTION				
CYTARABINE				
SICOR PHARMS	100MG/VIAL	N16793	001	
	500MG/VIAL	N16793	002	
	1GM/VIAL	N16793	003	Dec 21, 1987
	2GM/VIAL	N16793	004	Dec 21, 1987

DACARBAZINE

INJECTABLE; INJECTION				
DACARBAZINE				
AM PHARM PARTNERS	100MG/VIAL	N70962	001	Aug 28, 1986
	200MG/VIAL	N70990	001	Aug 28, 1986

DALFOPRISTIN; QUINUPRISTIN

INJECTABLE; IV (INFUSION)				
SYNERCID				
KING PHARMS	420MG/VIAL;180MG/VIAL	N50748	002	Aug 24, 2000

DALTEPARIN SODIUM

INJECTABLE; INJECTION				
FRAGMIN				
PHARMACIA AND UPJOHN	7,500 IU/0.75ML	N20287	008	Apr 04, 2002

DANAPAROID SODIUM

INJECTABLE; INJECTION				
ORGARAN				
ORGANON USA INC	750 UNITS/0.6ML	N20430	001	Dec 24, 1996

DANAZOL

CAPSULE; ORAL				
DANAZOL				
AM THERAP	200MG	N71569	001	Dec 30, 1987
DANOCRINE				
SANOFI SYNTHELABO	50MG	N17557	003	
	100MG	N17557	004	
	200MG	N17557	002	

DAPSONE

GEL; TOPICAL				
ACZONE				
QLT USA	5%	N21794	001	Jul 07, 2005

DAUNORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION				
CERUBIDINE				
AVENTIS	EQ 20MG BASE/VIAL	N61876	001	
WYETH AYERST	EQ 20MG BASE/VIAL	N50484	001	

DECAMETHONIUM BROMIDE

INJECTABLE; INJECTION				
SYNCURINE				
GLAXOSMITHKLINE	1MG/ML	N06931	002	

DEMECARIUM BROMIDE

SOLUTION/DROPS; OPHTHALMIC				
HUMORSOL				
MERCK	0.125%	N11860	002	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 73 (of 263)

DEMECARIUM BROMIDESOLUTION/DROPS; OPHTHALMIC
HUMORSOL

MERCK 0.25% N11860 001

DEMECLOCYCLINE HYDROCHLORIDECAPSULE; ORAL
DECLOMYCIN

LEDERLE 150MG N50262 001

SYRUP; ORAL
DECLOMYCIN

LEDERLE 75MG/5ML N50257 001

TABLET; ORAL
DECLOMYCIN

ESP PHARMA 75MG N50261 001

DESERPIDINETABLET; ORAL
HARMONYL

ABBOTT 0.1MG N10796 001

0.25MG N10796 002

DESERPIDINE; HYDROCHLOROTHIAZIDETABLET; ORAL
ORETICYL 25

ABBOTT 0.125MG;25MG N12148 001

ORETICYL 50

ABBOTT 0.125MG;50MG N12148 003

ORETICYL FORTE

ABBOTT 0.25MG;25MG N12148 002

DESERPIDINE; METHYCLOTHIAZIDETABLET; ORAL
ENDURONYL

ABBOTT 0.25MG;5MG N12775 001

ENDURONYL FORTE

ABBOTT 0.5MG;5MG N12775 002

METHYCLOTHIAZIDE AND DESERPIDINE

WATSON LABS 0.25MG;5MG N88486 001 Aug 10, 1984

0.5MG;5MG N88452 001 Aug 10, 1984

DESIPRAMINE HYDROCHLORIDECAPSULE; ORAL
PERTOFRANE

AVENTIS 25MG N13621 001

50MG N13621 002

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

USL PHARMA 25MG N71864 001 Sep 09, 1987

50MG N71865 001 Sep 09, 1987

75MG N71866 001 Sep 09, 1987

100MG N71867 001 Sep 09, 1987

DESLANOSIDEINJECTABLE; INJECTION
CEDILANID-D

NOVARTIS 0.2MG/ML N09282 002

DESMOPRESSIN ACETATESOLUTION; NASAL
CONCENTRAID

FERRING 0.01% N19776 001 Dec 26, 1990

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 74 (of 263)

DESMOPRESSIN ACETATESPRAY, METERED; NASAL
DDAVP

AVENTIS 0.01MG/SPRAY N17922 002 Feb 06, 1989

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL - 21

DESOGEN

ORGANON USA INC 0.15MG;0.03MG N20071 001 Dec 10, 1992

ORTHO-CEPT

ORTHO MCNEIL PHARM 0.15MG;0.03MG N20301 001 Dec 14, 1992

DESOXIMETASONE

OINTMENT; TOPICAL

DESOXIMETASONE

ALTANA 0.25% N73440 001 Apr 01, 1998

TOPICORT

TARO PHARMS NORTH 0.05% N18594 001 Jan 17, 1985

DESOXYCORTICOSTERONE ACETATE

INJECTABLE; INJECTION

DOCA

ORGANON USA INC 5MG/ML N01104 001

PELLET; IMPLANTATION

PERCORTEN

NOVARTIS 125MG N05151 001

DESOXYCORTICOSTERONE PIVALATE

INJECTABLE; INJECTION

PERCORTEN

NOVARTIS 25MG/ML N08822 001

DEXAMETHASONE

AEROSOL; TOPICAL

AEROSEB-DEX

ALLERGAN HERBERT 0.01% N83296 002

DECASPRAY

MERCK 0.04% N12731 002

ELIXIR; ORAL

DECADRON

MERCK 0.5MG/5ML N12376 002

DEXAMETHASONE

ALPHARMA US PHARMS 0.5MG/5ML N88997 001 Oct 10, 1986

GEL; TOPICAL

DECADERM

MERCK 0.1% N13538 001

SUSPENSION/DROPS; OPHTHALMIC

DEXAMETHASONE

STERIS 0.1% N89170 001 May 09, 1989

TABLET; ORAL

DECADRON

MERCK 0.25MG N11664 004

1.5MG **Federal Register determination
that product was not discontinued or
withdrawn for safety or efficacy
reasons** N11664 003

4MG N11664 005

6MG N11664 006 Jul 30, 1982

DEXAMETHASONE

IMPAX LABS 0.75MG N85376 001

MUTUAL PHARM 0.25MG N84013 001

0.25MG N84764 001

0.5MG N84084 001

0.5MG N84766 001

0.75MG N84081 001

DISCONTINUED DRUG PRODUCT LIST

6 - 75 (of 263)

DEXAMETHASONE

TABLET; ORAL

DEXAMETHASONE

MUTUAL PHARM

0.75MG

N84765 001

1.5MG

N84086 001

1.5MG

N84763 001

PHOENIX LABS NY

0.75MG

N83806 001

PVT FORM

0.75MG

N83420 001

ROXANE

0.25MG

N84614 001

SANDOZ

0.75MG

N80399 001

UPSHER SMITH

0.75MG

N87534 001

1.5MG

N87533 001

WATSON LABS

0.25MG

N85455 001

0.5MG

N85458 001

0.75MG

N80968 001

0.75MG

N84457 001

0.75MG

N85818 001

1.5MG

N85456 001

1.5MG

N85840 001

WHITEWORTH TOWN PLSN

0.75MG

N84327 001

DEXONE 0.5

SOLVAY

0.5MG

N84991 001

DEXONE 0.75

SOLVAY

0.75MG

N84993 001

DEXONE 1.5

SOLVAY

1.5MG

N84990 001

DEXONE 4

SOLVAY

4MG

N84992 001

HEXADROL

ORGANON USA INC

0.5MG

N12675 004

0.75MG

N12675 007

1.5MG

N12675 009

DEXAMETHASONE ACETATE

INJECTABLE; INJECTION

DECADRON-LA

MERCK

EQ 8MG BASE/ML **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N16675 001

DEXAMETHASONE ACETATE

STERIS

EQ 8MG BASE/ML

N84315 001

EQ 16MG BASE/ML

N87711 001

May 24, 1982

DEXAMETHASONE SODIUM PHOSPHATE

AEROSOL; NASAL

DEXACORT

UCB

EQ 0.1MG PHOSPHATE/INH

N14242 001

AEROSOL, METERED; INHALATION

DEXACORT

UCB

EQ 0.1MG PHOSPHATE/INH

N13413 001

CREAM; TOPICAL

DECADRON

MERCK

EQ 0.1% PHOSPHATE

N11983 002

INJECTABLE; INJECTION

DECADRON

MERCK

EQ 4MG PHOSPHATE/ML

N12071 002

EQ 24MG PHOSPHATE/ML **Federal
Register determination that product
was not discontinued or withdrawn for
safety or efficacy reasons**

N12071 004

DEXACEN-4

CENT PHARMS

EQ 4MG PHOSPHATE/ML

N84342 001

DEXAMETHASONE

AM PHARM PARTNERS

EQ 4MG PHOSPHATE/ML

N88448 001

Jan 25, 1984

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 76 (of 263)

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE

AM PHARM PARTNERS EQ 10MG PHOSPHATE/ML N88469 001 Jan 25, 1984

DEXAMETHASONE SODIUM PHOSPHATE

AKORN EQ 4MG PHOSPHATE/ML N84493 001

AM PHARM PARTNERS EQ 4MG PHOSPHATE/ML N87065 001

BEL MAR EQ 4MG PHOSPHATE/ML N84752 001

DELL LABS EQ 4MG PHOSPHATE/ML N83161 001

INTL MEDICATION EQ 20MG PHOSPHATE/ML N88522 001 Feb 17, 1984

SICOR PHARMS EQ 4MG PHOSPHATE/ML N81125 001 Aug 31, 1990

STERIS EQ 4MG PHOSPHATE/ML N83702 001

EQ 4MG PHOSPHATE/ML N84355 001

EQ 4MG PHOSPHATE/ML N89169 001 Apr 09, 1986

EQ 10MG PHOSPHATE/ML N87668 001 Jul 01, 1982

EQ 24MG PHOSPHATE/ML N85606 001

WYETH AYERST EQ 4MG PHOSPHATE/ML N85641 001

HEXADROL

ORGANON USA INC EQ 4MG PHOSPHATE/ML N14694 002

EQ 10MG PHOSPHATE/ML N14694 003

EQ 20MG PHOSPHATE/ML N14694 004

OINTMENT; OPHTHALMIC

DECADRON

MERCK EQ 0.05% PHOSPHATE N11977 001

DEXAIR

PHARMAFAIR EQ 0.05% PHOSPHATE N88071 001 Dec 28, 1982

MAXIDEX

ALCON EQ 0.05% PHOSPHATE N83342 001

SOLUTION/DROPS; OPHTHALMIC

DEXAIR

PHARMAFAIR EQ 0.1% PHOSPHATE N88433 001 Dec 15, 1983

DEXAMETHASONE SODIUM PHOSPHATE

SOLA BARNES HIND EQ 0.1% PHOSPHATE N84170 001

EQ 0.1% PHOSPHATE N84173 001

SOLUTION/DROPS; OPHTHALMIC, OTIC

DECADRON

MERCK EQ 0.1% PHOSPHATE N11984 001

SOLUTION/DROPS; OTIC

DEXAMETHASONE SODIUM PHOSPHATE

AKORN EQ 0.1% PHOSPHATE N84855 001

DEXAMETHASONE SODIUM PHOSPHATE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

DECADRON W/ XYLOCAINE

MERCK EQ 4MG PHOSPHATE/ML;10MG/ML N13334 002

DEXAMETHASONE SODIUM PHOSPHATE; NEOMYCIN SULFATE

OINTMENT; OPHTHALMIC

NEODECADRON

MERCK EQ 0.05% PHOSPHATE;EQ 3.5MG BASE/GM N50324 001

SOLUTION/DROPS; OPHTHALMIC

NEODECADRON

MERCK EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML N50322 001

NEOMYCIN SULFATE AND DEXAMETHASONE SODIUM PHOSPHATE

BAUSCH AND LOMB EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML N64055 001 Oct 30, 1995

NEOMYCIN SULFATE-DEXAMETHASONE SODIUM PHOSPHATE

ALCON UNIVERSAL EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML N62714 001 Jul 21, 1986

PHARMAFAIR EQ 0.1% PHOSPHATE;EQ 3.5MG BASE/ML N62539 001 Jan 10, 1985

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

DEXACIDIN

NOVARTIS 0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM N62566 001 Feb 22, 1985

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 77 (of 263)

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC			
DEXASPORIN			
PHARMAFAIR	0.1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	N62411 001	May 16, 1983
SUSPENSION/DROPS; OPHTHALMIC			
DEXACIDIN			
NOVARTIS	0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N62544 001	Oct 29, 1984
DEXASPORIN			
PHARMAFAIR	0.1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N62428 001	May 18, 1983

DEXBROMPHENIRAMINE MALEATE

SYRUP; ORAL			
DISOMER			
SCHERING	2MG/5ML	N11814 002	
TABLET; ORAL			
DISOMER			
SCHERING	2MG	N11814 001	

DEXBROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE SULFATE

TABLET; ORAL			
DISOPHROL			
SCHERING	2MG;60MG	N12394 002	
TABLET, EXTENDED RELEASE; ORAL			
BROMPHERIL			
COPLEY PHARM	6MG;120MG	N89116 001	Jan 22, 1987
DISOBROM			
SANDOZ	6MG;120MG	N70770 001	Sep 30, 1991
RESPORAL			
PIONEER PHARMS	6MG;120MG	N89139 001	Jun 16, 1988

DEXCHLORPHENIRAMINE MALEATE

SYRUP; ORAL			
POLARAMINE			
SCHERING	2MG/5ML	N86837 001	Jul 19, 1982
TABLET; ORAL			
POLARAMINE			
SCHERING	2MG	N86835 001	

DEXTROAMPHETAMINE SULFATE

CAPSULE; ORAL			
DEXAMPEX			
TEVA	15MG	N85355 001	
CAPSULE, EXTENDED RELEASE; ORAL			
DEXTROAMPHETAMINE SULFATE			
ABLE	5MG	N76814 001	Aug 25, 2004
	10MG	N76814 002	Aug 25, 2004
	15MG	N76814 003	Aug 25, 2004
ELIXIR; ORAL			
DEXEDRINE			
GLAXOSMITHKLINE	5MG/5ML	N83902 001	
TABLET; ORAL			
DEXAMPEX			
TEVA	5MG	N83735 001	
	10MG	N83735 002	
DEXTROAMPHETAMINE SULFATE			
HALSEY	10MG	N83930 001	
LANNETT	5MG	N83903 001	
	10MG	N83903 003	
	15MG **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N85652 001	
MAST MM	5MG	N86521 001	
PUREPAC PHARM	5MG	N84125 001	
SANDOZ	5MG	N85370 001	

DISCONTINUED DRUG PRODUCT LIST

6 - 78 (of 263)

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL			
DEXTROAMPHETAMINE SULFATE			
SANDOZ	10MG	N85371	001
VITARINE	5MG	N84986	001
	10MG	N85892	001
FERNDEX			
FERNDAL LABS	5MG	N84001	001

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL			
PHENERGAN W/ DEXTROMETHORPHAN			
WYETH AYERST	15MG/5ML;6.25MG/5ML	N11265	002 Apr 02, 1984
PHERAZINE DM			
HALSEY	15MG/5ML;6.25MG/5ML	N88913	001 Mar 02, 1987

DEXTROSE

INJECTABLE; INJECTION			
DEXTROSE 10% IN PLASTIC CONTAINER			
B BRAUN	10GM/100ML	N18046	001
MILES	10GM/100ML	N18504	001
DEXTROSE 2.5% IN PLASTIC CONTAINER			
B BRAUN	2.5GM/100ML	N18358	001
	2.5GM/100ML	N19626	001 Feb 02, 1988
DEXTROSE 38.5% IN PLASTIC CONTAINER			
ABBOTT	38.5GM/100ML	N18923	001 Sep 19, 1984
DEXTROSE 60%			
B BRAUN	60GM/100ML	N17995	002 Sep 22, 1982
DEXTROSE 60% IN PLASTIC CONTAINER			
B BRAUN	60GM/100ML	N17995	001
BAXTER HLTHCARE	60GM/100ML	N20047	002 Jul 02, 1991
DEXTROSE 7.7% IN PLASTIC CONTAINER			
B BRAUN	7.7GM/100ML	N19626	003 Feb 02, 1988

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE

INJECTABLE; INJECTION			
ISOLYTE P W/ DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;31MG/100ML;130MG/100ML;26MG/100ML;320MG/100ML	N19025	001 Dec 27, 1984

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION			
ISOLYTE H W/ DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;30MG/100ML;97MG/100ML;220MG/100ML;140MG/100ML	N18273	001

DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION			
ISOLYTE S W/ DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;30MG/100ML;37MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML	N18274	001

DEXTROSE; POTASSIUM CHLORIDE

INJECTABLE; INJECTION			
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;37MG/100ML	N19699	001 Sep 29, 1989
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;75MG/100ML	N19699	002 Sep 29, 1989
POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;110MG/100ML	N19699	003 Sep 29, 1989
POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;220MG/100ML	N19699	005 Sep 29, 1989

DISCONTINUED DRUG PRODUCT LIST

6 - 79 (of 263)

DEXTROSE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, DIBASIC; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE M W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;150MG/100ML;130MG/100ML;280MG /100ML;91MG/100ML	N18270 001
---------	--	------------

DEXTROSE; POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.075%

B BRAUN	5GM/100ML;75MG/100ML;200MG/100ML	N18268 009
---------	----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;150MG/100ML;200MG/100ML	N18268 004
---------	-----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;220MG/100ML;200MG/100ML	N18268 005
---------	-----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;300MG/100ML;200MG/100ML	N18268 006
---------	-----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;75MG/100ML;330MG/100ML	N18268 011 Jan 18, 1986
---------	----------------------------------	-------------------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;150MG/100ML;330MG/100ML	N18268 012 Jan 18, 1986
---------	-----------------------------------	-------------------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;220MG/100ML;330MG/100ML	N18268 013 Jan 18, 1986
---------	-----------------------------------	-------------------------

DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 0.30% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;300MG/100ML;330MG/100ML	N18268 014 Jan 18, 1986
---------	-----------------------------------	-------------------------

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.075%

B BRAUN	5GM/100ML;75MG/100ML;450MG/100ML	N18268 010
---------	----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;150MG/100ML;450MG/100ML	N18268 001
---------	-----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;220MG/100ML;450MG/100ML	N18268 002
---------	-----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;300MG/100ML;450MG/100ML	N18268 003
---------	-----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER

BAXTER HLTHCARE	5GM/100ML;224MG/100ML;450MG/100ML	N18008 003
-----------------	-----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER

BAXTER HLTHCARE	5GM/100ML;300MG/100ML;450MG/100ML	N18008 001
-----------------	-----------------------------------	------------

DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER

BAXTER HLTHCARE	5GM/100ML;75MG/100ML;450MG/100ML	N18008 002
-----------------	----------------------------------	------------

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;200MG/100ML	N18386 001
---------	------------------------	------------

DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;450MG/100ML	N18229 001
---------	------------------------	------------

DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	10GM/100ML;900MG/100ML	N18047 001
---------	------------------------	------------

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER

B BRAUN	2.5GM/100ML;450MG/100ML	N18030 001
---------	-------------------------	------------

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

B BRAUN	2.5GM/100ML;900MG/100ML	N18376 001
---------	-------------------------	------------

DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

ABBOTT	3.3GM/100ML;300MG/100ML	N18055 001
--------	-------------------------	------------

DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;110MG/100ML	N18030 005
---------	-----------------------	------------

DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER

B BRAUN	5GM/100ML;200MG/100ML	N18030 004
---------	-----------------------	------------

MILES	5GM/100ML;200MG/100ML	N18399 001
-------	-----------------------	------------

DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER

ABBOTT	5GM/100ML;225MG/100ML	N19482 001 Oct 04, 1985
--------	-----------------------	-------------------------

DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER

ABBOTT	5GM/100ML;300MG/100ML	N19486 001 Oct 04, 1985
--------	-----------------------	-------------------------

MILES	5GM/100ML;300MG/100ML	N18501 001
-------	-----------------------	------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 80 (of 263)

DEXTROSE; SODIUM CHLORIDE

INJECTABLE; INJECTION

DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER			
B BRAUN	5GM/100ML;330MG/100ML	N18030 003	
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;450MG/100ML	N19484 001	Oct 04, 1985
B BRAUN	5GM/100ML;450MG/100ML	N18030 002	
MILES	5GM/100ML;450MG/100ML	N18400 001	
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER			
ABBOTT	5GM/100ML;900MG/100ML	N19483 001	Oct 04, 1985
B BRAUN	5GM/100ML;900MG/100ML	N18026 001	
MILES	5GM/100ML;900MG/100ML	N18500 001	

DEXTROTHYROXINE SODIUM

TABLET; ORAL

CHOLOXIN

ABBOTT	1MG	N12302 005	
	2MG	N12302 002	
	4MG	N12302 004	
	6MG	N12302 006	

DEZOCINE

INJECTABLE; INJECTION

DALGAN

ASTRAZENECA	5MG/ML	N19082 001	Dec 29, 1989
	10MG/ML	N19082 002	Dec 29, 1989
	15MG/ML	N19082 003	Dec 29, 1989

DIATRIZOATE MEGLUMINE

INJECTABLE; INJECTION

ANGIOVIST 282

BERLEX	60%	N87726 001	Sep 23, 1982
--------	-----	------------	--------------

CARDIOGRAFIN

BRACCO	85%	N11620 002	
--------	-----	------------	--

DIATRIZOATE MEGLUMINE

BRACCO	76%	N10040 017	
--------	-----	------------	--

HYPAQUE

GE HEALTHCARE	30%	N16403 002	
---------------	-----	------------	--

UROVIST MEGLUMINE DIU/CT

BERLEX	30%	N87739 001	Sep 23, 1982
--------	-----	------------	--------------

SOLUTION; URETERAL

UROVIST CYSTO

BERLEX	30%	N87729 001	Sep 23, 1982
--------	-----	------------	--------------

UROVIST CYSTO PEDIATRIC

BERLEX	30%	N87731 001	Sep 23, 1982
--------	-----	------------	--------------

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION

ANGIOVIST 292

BERLEX	52%;8%	N87724 001	Sep 23, 1982
--------	--------	------------	--------------

ANGIOVIST 370

BERLEX	66%;10%	N87723 001	Sep 23, 1982
--------	---------	------------	--------------

DIATRIZOATE-60

INTL MEDICATION	52%;8%	N88166 001	Jun 17, 1983
-----------------	--------	------------	--------------

HYPAQUE-M, 75%

GE HEALTHCARE	50%;25%	N10220 003	
---------------	---------	------------	--

HYPAQUE-M, 90%

GE HEALTHCARE	60%;30%	N10220 002	
---------------	---------	------------	--

MD-60

MALLINCKRODT	52%;8%	N87074 001	
--------------	--------	------------	--

MD-76

MALLINCKRODT	66%;10%	N87073 001	
--------------	---------	------------	--

RENOVIST

BRACCO	34.3%;35%	N10040 020	
--------	-----------	------------	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 81 (of 263)

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUM

INJECTABLE; INJECTION				
RENOVIST II				
BRACCO	28.5%;29.1%	N10040	019	
SOLUTION; ORAL, RECTAL				
GASTROVIST				
BERLEX	66%;10%	N87728	001	Sep 23, 1982

DIATRIZOATE SODIUM

INJECTABLE; INJECTION				
HYPAQUE				
GE HEALTHCARE	25%	N09561	003	
MD-50				
MALLINCKRODT	50%	N87075	001	
UROVIST SODIUM 300				
BERLEX	50%	N87725	001	Sep 23, 1982
SOLUTION; ORAL, RECTAL				
HYPAQUE				
GE HEALTHCARE	40%	N11386	003	
SOLUTION; URETERAL				
HYPAQUE SODIUM 20%				
GE HEALTHCARE	20%	N09561	002	

DIAZEPAM

CAPSULE, EXTENDED RELEASE; ORAL				
VALRELEASE				
ROCHE	15MG	N18179	001	
GEL; RECTAL				
DIASTAT				
VALEANT	5MG/ML (5MG/ML)	N20648	002	Jul 29, 1997
	10MG/2ML (5MG/ML)	N20648	003	Jul 29, 1997
	15MG/3ML (5MG/ML)	N20648	004	Jul 29, 1997
	20MG/4ML (5MG/ML)	N20648	005	Jul 29, 1997
INJECTABLE; INJECTION				
DIAZEPAM				
AM PHARM PARTNERS	5MG/ML	N70662	001	Jun 25, 1986
BAXTER HLTHCARE	5MG/ML	N70311	001	Dec 16, 1985
	5MG/ML	N70312	001	Dec 16, 1985
	5MG/ML	N70313	001	Dec 16, 1985
	5MG/ML	N71308	001	Jul 17, 1987
MARSAM PHARMS LLC	5MG/ML	N72371	001	Jan 29, 1993
STERIS	5MG/ML	N70911	001	Aug 28, 1986
	5MG/ML	N70912	001	Aug 28, 1986
	5MG/ML	N70930	001	Dec 01, 1986
US ARMY	5MG/ML	N20124	001	Dec 05, 1990
WARNER CHILCOTT	5MG/ML	N71613	001	Oct 22, 1987
	5MG/ML	N71614	001	Oct 22, 1987
DIZAC				
PHARMACIA AND UPJOHN	5MG/ML	N19287	001	Jun 18, 1993
VALIUM				
ROCHE	5MG/ML	N16087	001	
TABLET; ORAL				
DIAZEPAM				
CLONMEL HLTHCARE	5MG	N70227	001	Sep 26, 1985
	10MG	N70228	001	Sep 26, 1985
DURAMED PHARMS BARR	2MG	N70894	001	Aug 27, 1986
	5MG	N70895	001	Aug 27, 1986
	10MG	N70896	001	Aug 27, 1986
FERNDAL LABS	2MG	N70903	001	Apr 01, 1987
	5MG	N70904	001	Apr 01, 1987
	10MG	N70905	001	Apr 01, 1987
HALSEY	2MG	N70987	001	Aug 15, 1986
	5MG	N70996	001	Aug 15, 1986
	10MG	N70956	001	Aug 15, 1986
IVAX PHARMS	2MG	N70360	001	Sep 04, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 82 (of 263)

DIAZEPAM

TABLET; ORAL
DIAZEPAM

IVAX PHARMS	5MG	N70361	001	Sep 04, 1985
	10MG	N70362	001	Sep 04, 1985
MARTEC	10MG	N72402	001	Apr 25, 1989
PAR PHARM	10MG	N70464	001	Feb 25, 1986
PIONEER PHARMS	2MG	N70787	001	Aug 02, 1988
	5MG	N70788	001	Aug 02, 1988
	10MG	N70776	001	Aug 02, 1988
ROXANE	2MG	N70356	001	Jun 17, 1986
	5MG	N70357	001	Jun 17, 1986
	10MG	N70358	001	Jun 17, 1986
WARNER CHILCOTT	2MG	N70209	001	Sep 04, 1985
	5MG	N70210	001	Sep 04, 1985
	10MG	N70222	001	Sep 04, 1985
WATSON LABS	2MG	N70456	001	Nov 01, 1985
	5MG	N70457	001	Nov 01, 1985
	10MG	N70458	001	Nov 01, 1985
Q-PAM				
QUANTUM PHARMICS	2MG	N70423	001	Dec 12, 1985
	2MG	N72431	001	Apr 29, 1988
	5MG	N70424	001	Dec 12, 1985
	5MG	N72432	001	Apr 29, 1988
	10MG	N70425	001	Dec 12, 1985
	10MG	N72433	001	Apr 29, 1988

DIAZOXIDE

CAPSULE; ORAL
PROGLYCEM

BAKER NORTON	50MG	N17425	001	
	100MG	N17425	002	

INJECTABLE; INJECTION
DIAZOXIDE

AM PHARM PARTNERS	15MG/ML	N71519	001	Aug 26, 1987
-------------------	---------	--------	-----	--------------

DIBUCAINE HYDROCHLORIDE

INJECTABLE; INJECTION
HEAVY SOLUTION NUPERCAINE

NOVARTIS	2.5MG/ML	N06203	001	
----------	----------	--------	-----	--

DICHLORPHENAMIDE

TABLET; ORAL
DARANIDE

MERCK	50MG	N11366	001	
-------	------	--------	-----	--

DICLOFENAC POTASSIUM

TABLET; ORAL
CATAFLAM

NOVARTIS	25MG **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N20142	001	Nov 24, 1993
----------	---	--------	-----	--------------

DICLOFENAC SODIUM

SOLUTION/DROPS; OPHTHALMIC
DICLOFENAC SODIUM

FALCON PHARMS	0.1%	N20809	001	May 04, 1998
---------------	------	--------	-----	--------------

TABLET, DELAYED RELEASE; ORAL
VOLTAREN

NOVARTIS	25MG	N19201	001	Jul 28, 1988
	50MG	N19201	002	Jul 28, 1988
	75MG	N19201	003	Jul 28, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 83 (of 263)

DICLOXACILLIN SODIUMCAPSULE; ORAL
DYCILL

GLAXOSMITHKLINE	EQ 250MG BASE	N60254	002
	EQ 250MG BASE	N62238	001
	EQ 500MG BASE	N60254	003
	EQ 500MG BASE	N62238	002

PATHOCIL

WYETH AYERST	EQ 250MG BASE	N50011	002
	EQ 500MG BASE	N50011	003

Mar 28, 1983

FOR SUSPENSION; ORAL
DYNAPEN

APOTHECON	EQ 62.5MG BASE/5ML	N50337	002
-----------	--------------------	--------	-----

PATHOCIL

WYETH AYERST	EQ 62.5MG BASE/5ML	N50092	001
--------------	--------------------	--------	-----

DICUMAROLCAPSULE; ORAL
DICUMAROL

LILLY	25MG	N05509	003
	50MG	N05509	001

TABLET; ORAL

DICUMAROL

ABBOTT	25MG	N05545	003
	50MG	N05545	004
	100MG	N05545	005

DICYCLOMINE HYDROCHLORIDE

CAPSULE; ORAL

DICYCLOMINE HYDROCHLORIDE

MUTUAL PHARM	10MG	N84505	001	Oct 21, 1986
PIONEER PHARMS	10MG	N89361	001	Jan 10, 1989
WATSON LABS	10MG	N83179	001	Feb 12, 1986

INJECTABLE; INJECTION

DICYCLOMINE HYDROCHLORIDE

STERIS	10MG/ML	N80614	001	Feb 11, 1986
--------	---------	--------	-----	--------------

SYRUP; ORAL

DICYCLOMINE HYDROCHLORIDE

ALPHARMA US PHARMS	10MG/5ML	N84479	001
--------------------	----------	--------	-----

TABLET; ORAL

DICYCLOMINE HYDROCHLORIDE

MUTUAL PHARM	20MG	N84600	001	Jul 29, 1985
PIONEER PHARMS	20MG	N88585	001	Aug 20, 1986
WATSON LABS	20MG	N84361	001	Feb 06, 1986

DIDANOSINE

FOR SOLUTION; ORAL

VIDEX

BRISTOL MYERS SQUIBB	100MG/PACKET	N20155	003	Oct 09, 1991
	167MG/PACKET	N20155	004	Oct 09, 1991
	250MG/PACKET	N20155	005	Oct 09, 1991
	375MG/PACKET	N20155	006	Oct 09, 1991

DIENESTROL

CREAM; VAGINAL

DIENESTROL

ORTHO MCNEIL PHARM	0.01%	N06110	005
--------------------	-------	--------	-----

DV

AVENTIS PHARMS	0.01%	N83518	001
----------------	-------	--------	-----

ESTRAGUARD

SOLVAY	0.01%	N84436	001
--------	-------	--------	-----

SUPPOSITORY; VAGINAL

DV

AVENTIS PHARMS	0.7MG	N83517	001
----------------	-------	--------	-----

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 84 (of 263)

DIETHYLCARBAMAZINE CITRATE

TABLET; ORAL			
HETRAZAN			
LEDERLE	50MG	N06459	001

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL			
DIETHYLPROPION HCL			
UCB	25MG	N85544	001
DIETHYLPROPION HYDROCHLORIDE			
ABC HOLDING	25MG	N88267	001 Aug 25, 1983
	25MG	N88268	001 Aug 25, 1983
SANDOZ	25MG	N85916	001
TEVA	25MG	N88642	001 Sep 20, 1984
WATSON LABS	25MG	N85741	001
TENUATE			
AVENTIS PHARMS	25MG	N17668	001
TEPANIL			
3M	25MG	N11673	001
TABLET, EXTENDED RELEASE; ORAL			
TENUATE			
AVENTIS PHARMS	75MG	N17669	001
TEPANIL TEN-TAB			
3M	75MG	N17956	001

DIETHYLSTILBESTROL

INJECTABLE; INJECTION			
STILBESTROL			
BRISTOL MYERS SQUIBB	0.2MG/ML	N04056	003
	0.5MG/ML	N04056	004
	1MG/ML	N04056	005
	5MG/ML	N04056	006
SUPPOSITORY; VAGINAL			
DIETHYLSTILBESTROL			
LILLY	0.1MG	N04040	001
	0.5MG	N04040	002
STILBESTROL			
BRISTOL MYERS SQUIBB	0.1MG	N04056	001
	0.5MG	N04056	002
TABLET; ORAL			
DIETHYLSTILBESTROL			
LILLY	0.1MG	N04041	002
	0.5MG	N04041	003
	1MG	N04041	004
	5MG	N04041	005
STILBESTROL			
TABLICAPS	0.5MG	N83004	001
	1MG	N83002	001
	5MG	N83006	001
STILBETIN			
BRISTOL MYERS SQUIBB	0.1MG	N04056	007
	0.25MG	N04056	017
	0.5MG	N04056	008
	1MG	N04056	009
	5MG	N04056	010
TABLET, DELAYED RELEASE; ORAL			
DIETHYLSTILBESTROL			
LILLY	0.1MG	N04039	002
	0.25MG	N04039	005
	0.5MG	N04039	003
	1MG	N04039	004
	5MG	N04039	006
STILBESTROL			
TABLICAPS	0.5MG	N83003	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 85 (of 263)

DIETHYLSTILBESTROL

TABLET, DELAYED RELEASE; ORAL

STILBESTROL

TABLICAPS

1MG

N83005 001

5MG

N83007 001

STILBETIN

BRISTOL MYERS SQUIBB

0.1MG

N04056 011

0.5MG

N04056 012

1MG

N04056 013

5MG

N04056 014

DIETHYLSTILBESTROL DIPHOSPHATE

INJECTABLE; INJECTION

STILPHOSTROL

BAYER PHARMS

250MG/5ML

N10010 001

TABLET; ORAL

STILPHOSTROL

BAYER PHARMS

50MG

N10010 002

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL

PUREPAC PHARM

250MG

N74285 001

May 07, 1996

500MG

N74285 002

May 07, 1996

ROXANE

250MG

N73562 001

Nov 27, 1992

500MG

N73563 001

Nov 27, 1992

DIGITOXIN

INJECTABLE; INJECTION

CRYSTODIGIN

LILLY

0.2MG/ML

N84100 005

DIGOXIN

CAPSULE; ORAL

LANOXICAPS

SMITHKLINE BEECHAM

0.15MG

N18118 004

Sep 24, 1984

INJECTABLE; INJECTION

DIGOXIN

AM PHARM PARTNERS

0.25MG/ML

N83217 001

WYETH AYERST

0.25MG/ML

N84386 001

TABLET; ORAL

LANOXIN

SMITHKLINE BEECHAM

0.0625MG

N20405 001

Sep 30, 1997

0.1875MG

N20405 003

Sep 30, 1997

0.375MG

N20405 005

Sep 30, 1997

0.5MG

N20405 006

Sep 30, 1997

DIHYDROERGOTAMINE MESYLATE; HEPARIN SODIUM; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

EMBOLEX

NOVARTIS

0.5MG/0.5ML; 2,500
UNITS/0.5ML; 5.33MG/0.5ML

N18885 001

Nov 30, 1984

0.5MG/0.7ML; 5,000
UNITS/0.7ML; 7.46MG/0.7ML

N18885 002

Nov 30, 1984

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CARDIZEM SR

BIOVAIL

60MG

N19471 001

Jan 23, 1989

90MG

N19471 002

Jan 23, 1989

120MG

N19471 003

Jan 23, 1989

180MG

N19471 004

Jan 23, 1989

DILTIAZEM HYDROCHLORIDE

BIOVAIL

60MG

N74845 001

Sep 15, 1999

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 86 (of 263)

DILTIAZEM HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
DILTIAZEM HYDROCHLORIDE

BIOVAIL	90MG	N74845	002	Sep 15, 1999
	120MG	N74845	003	Sep 15, 1999
TEVA	60MG	N74079	001	Nov 30, 1993
	90MG	N74079	002	Nov 30, 1993
	120MG	N74079	003	Nov 30, 1993

TABLET; ORAL

DILTIAZEM HYDROCHLORIDE

APOTHECON	30MG	N74051	001	Mar 31, 1993
	60MG	N74051	002	Mar 31, 1993
	90MG	N74051	003	Mar 31, 1993
	120MG	N74051	004	Mar 31, 1993
TEVA	30MG	N74084	001	Feb 25, 1994
	60MG	N74084	002	Feb 25, 1994
TEVA PHARMS	30MG	N74067	001	Nov 05, 1992
	60MG	N74067	002	Nov 05, 1992
	90MG	N74067	003	Nov 05, 1992
	120MG	N74067	004	Nov 05, 1992

DILTIAZEM MALATE

TABLET, EXTENDED RELEASE; ORAL

TIAMATE

MERCK	EQ 120MG HCL	N20506	001	Oct 04, 1996
	EQ 180MG HCL	N20506	002	Oct 04, 1996
	EQ 240MG HCL	N20506	003	Oct 04, 1996

DIMENHYDRINATE

INJECTABLE; INJECTION

DIMENHYDRINATE

BAXTER HLTHCARE	50MG/ML	N84767	001
STERIS	50MG/ML	N83531	001
WYETH AYERST	50MG/ML	N84316	001

LIQUID; ORAL

DIMENHYDRINATE

ALRA	12.5MG/4ML	N80715	001
------	------------	--------	-----

TABLET; ORAL

DIMENHYDRINATE

ANABOLIC	50MG	N85985	001
HEATHER	50MG	N80841	001
WATSON LABS	50MG	N85166	001

DIMYRISTOYL LECITHIN; PERFLEXANE

INJECTABLE; INTRAVENOUS

IMAGENT

IMCOR PH	0.92MG/VIAL;0.092MG/VIAL	N21191	001	May 31, 2002
----------	--------------------------	--------	-----	--------------

DINOPROST TROMETHAMINE

INJECTABLE; INJECTION

PROSTIN F2 ALPHA

PHARMACIA AND UPJOHN	EQ 5MG BASE/ML	N17434	001
----------------------	----------------	--------	-----

DIPHEMANIL METHYLSULFATE

TABLET; ORAL

PRANTAL

SCHERING	100MG	N08114	004
----------	-------	--------	-----

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

BENADRYL

PARKE DAVIS	25MG	N05845	007
	50MG	N05845	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 87 (of 263)

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

ALRA	25MG	N80519	004	
	50MG	N80519	003	
ANABOLIC	25MG	N83634	001	
	50MG	N83275	001	
ELKINS SINN	25MG	N85701	001	
	50MG	N85701	002	
HALSEY	50MG	N87914	001	Jun 04, 1984
HEATHER	25MG	N84524	001	
	50MG	N83953	001	
IMPAX LABS	25MG	N80807	001	
	50MG	N80807	002	
IVAX PHARMS	25MG	N80762	001	
	50MG	N80762	002	
LANNETT	25MG	N80868	001	
	50MG	N80868	002	
LEDERLE	25MG	N86874	001	
	50MG	N86875	001	
MK LABS	25MG	N83087	001	
	50MG	N83087	002	
NEWTRON PHARMS	25MG	N86543	001	
	50MG	N86544	001	
PERRIGO	25MG	N83061	001	
	50MG	N83061	002	
PIONEER PHARMS	25MG	N89101	001	Dec 20, 1985
	50MG	N88880	001	Dec 20, 1985
PUREPAC PHARM	25MG	N85156	001	
	50MG	N85150	001	
PVT FORM	25MG	N83027	001	
	50MG	N83027	002	
ROXANE	50MG	N80635	001	
SANDOZ	25MG	N80845	002	
	50MG	N80845	001	
SUPERPHARM	25MG	N89040	001	May 15, 1985
	50MG	N89041	001	May 15, 1985
TEVA	25MG	N85874	001	
	50MG	N85874	002	
VALEANT PHARM INTL	25MG	N80596	001	
VANGARD	25MG	N88034	001	Oct 27, 1982
	50MG	N87630	001	
WATSON LABS	25MG	N83797	001	
	25MG	N85138	001	
	50MG	N83797	002	
	50MG	N85083	001	
WEST WARD	50MG	N83567	001	
WHITEWORTH TOWN PLSN	25MG	N83441	001	
	50MG	N80800	001	

ELIXIR; ORAL

BELIX

HALSEY	12.5MG/5ML	N86586	001	Oct 03, 1983
BENADRYL				
PARKE DAVIS	12.5MG/5ML	N05845	004	
DIBENIL				
CENCI	12.5MG/5ML	N88304	001	Dec 16, 1983
DIPHEN				
USL PHARMA	12.5MG/5ML	N84640	001	
DIPHENHYDRAMINE HYDROCHLORIDE				
BUNDY	12.5MG/5ML	N83674	001	
CENCI	12.5MG/5ML	N87941	001	Dec 17, 1982
KV PHARM	12.5MG/5ML	N85621	001	
LANNETT	12.5MG/5ML	N80939	002	
LEDERLE	12.5MG/5ML	N86937	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 88 (of 263)

DIPHENHYDRAMINE HYDROCHLORIDE

ELIXIR; ORAL

DIPHENHYDRAMINE HYDROCHLORIDE

MK LABS	12.5MG/5ML	N83088	002	
NASKA	12.5MG/5ML	N88680	001	May 31, 1985
PERRIGO	12.5MG/5ML	N83063	001	
PUREPAC PHARM	12.5MG/5ML	N83237	001	Jan 25, 1982
PVT FORM	12.5MG/5ML	N85287	001	
ROXANE	12.5MG/5ML	N80643	001	

HYDRAMINE

ALPHARMA US PHARMS	12.5MG/5ML	N80763	002	
--------------------	------------	--------	-----	--

INJECTABLE; INJECTION

BENADRYL

PARKE DAVIS	10MG/ML	N06146	001	
-------------	---------	--------	-----	--

DIPHENHYDRAMINE HYDROCHLORIDE

AM PHARM PARTNERS	10MG/ML	N87066	001	
BAXTER HLTHCARE	50MG/ML	N83183	001	
BEL MAR	10MG/ML	N80822	001	
STERIS	10MG/ML	N83533	001	
WYETH AYERST	50MG/ML	N80577	001	

DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE

AM PHARM PARTNERS	50MG/ML	N80586	002	
-------------------	---------	--------	-----	--

SYRUP; ORAL

ANTITUSSIVE

PERRIGO	12.5MG/5ML	N71292	001	Apr 10, 1987
---------	------------	--------	-----	--------------

BELDIN

HALSEY	12.5MG/5ML	N89179	001	Jun 05, 1986
--------	------------	--------	-----	--------------

BENYLIN

PARKE DAVIS	12.5MG/5ML	N06514	004	
-------------	------------	--------	-----	--

DIPHEN

MORTON GROVE	12.5MG/5ML	N70118	001	Oct 01, 1985
--------------	------------	--------	-----	--------------

DIPHENHYDRAMINE HYDROCHLORIDE

ALPHARMA US PHARMS	12.5MG/5ML	N70497	001	Apr 25, 1989
CUMBERLAND SWAN	12.5MG/5ML	N73611	001	Aug 20, 1992
HI TECH PHARMA	12.5MG/5ML	N72416	001	Sep 28, 1990

HYDRAMINE

ALPHARMA US PHARMS	12.5MG/5ML	N70205	001	Jan 28, 1986
--------------------	------------	--------	-----	--------------

SILPHEN

SILARX	12.5MG/5ML	N72646	001	Feb 27, 1992
--------	------------	--------	-----	--------------

VICKS FORMULA 44

PROCTER AND GAMBLE	12.5MG/5ML	N70524	001	Jan 14, 1987
--------------------	------------	--------	-----	--------------

DIPHENHYDRAMINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE

SOLUTION; ORAL

BENYLIN

PARKE DAVIS	12.5MG/5ML;30MG/5ML	N19014	001	Jun 11, 1985
-------------	---------------------	--------	-----	--------------

DIPHENIDOL HYDROCHLORIDE

TABLET; ORAL

VONTROL

GLAXOSMITHKLINE	EQ 25MG BASE	N16033	001	
-----------------	--------------	--------	-----	--

DIPHENYLPYRALINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

HISPRIL

GLAXOSMITHKLINE	5MG	N11945	001	
-----------------	-----	--------	-----	--

DIPYRIDAMOLE

INJECTABLE; INJECTION

IV PERSANTINE

BOEHRINGER INGELHEIM	5MG/ML	N19817	001	Dec 13, 1990
----------------------	--------	--------	-----	--------------

TABLET; ORAL

DIPYRIDAMOLE

CLONMEL HLTHCARE	25MG	N88999	001	Feb 05, 1991
------------------	------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 89 (of 263)

DIPYRIDAMOLE

TABLET; ORAL

DIPYRIDAMOLE

CLONMEL HLTHCARE

75MG

N89001 001

Feb 05, 1991

PUREPAC PHARM

50MG

N89426 001

Jul 12, 1990

75MG

N89427 001

Jul 12, 1990

DIRITHROMYCIN

TABLET, DELAYED RELEASE; ORAL

DYNABAC

LILLY RES LABS

250MG

N50678 001

Jun 19, 1995

DISOPYRAMIDE PHOSPHATE

CAPSULE; ORAL

DISOPYRAMIDE PHOSPHATE

INTERPHARM

EQ 100MG BASE

N71190 001

Jan 15, 1987

EQ 150MG BASE

N71191 001

Jan 15, 1987

MUTUAL PHARM

EQ 100MG BASE

N70351 001

Dec 17, 1985

EQ 150MG BASE

N70352 001

Dec 17, 1985

MYLAN

EQ 100MG BASE

N70138 001

Jun 14, 1985

EQ 150MG BASE

N70139 001

Jun 14, 1985

SUPERPHARM

EQ 100MG BASE

N70940 001

Feb 09, 1987

EQ 150MG BASE

N70941 001

Feb 09, 1987

WATSON LABS

EQ 100MG BASE

N70240 001

Feb 02, 1986

EQ 150MG BASE

N70241 001

Feb 02, 1986

DISULFIRAM

TABLET; ORAL

ANTABUSE

ODYSSEY PHARMS

250MG

N07883 003

500MG **Federal Register determination
that product was not discontinued or
withdrawn for safety or efficacy
reasons**

N07883 002

500MG

N88483 001

Dec 08, 1983

DISULFIRAM

PAR PHARM

250MG

N88792 001

Aug 14, 1984

500MG

N88793 001

Aug 14, 1984

WATSON LABS

250MG

N86889 001

250MG

N87973 001

Aug 05, 1983

500MG

N86890 001

500MG

N87974 001

Aug 05, 1983

DIVALPROEX SODIUM

TABLET, DELAYED RELEASE; ORAL

DEPAKOTE CP

ABBOTT

EQ 250MG BASE

N19794 001

Jul 11, 1990

EQ 500MG BASE

N19794 002

Jul 11, 1990

DOBUTAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

DOBUTAMINE HYDROCHLORIDE

ASTRAZENECA

EQ 12.5MG BASE/ML

N74098 001

Feb 21, 1995

BAXTER HLTHCARE

EQ 12.5MG BASE/ML

N74381 001

Sep 26, 1996

DOBUTREX

LILLY

EQ 12.5MG BASE/ML

N17820 002

DONEPEZIL HYDROCHLORIDE

SOLUTION; ORAL

ARICEPT

EISAI MEDCL RES

5MG/5ML

N21719 001

Oct 18, 2004

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 90 (of 263)

DOPAMINE HYDROCHLORIDEINJECTABLE; INJECTION
DOPAMINE HYDROCHLORIDE

ABBOTT	40MG/ML	N70656	001	Jan 24, 1989
	80MG/ML	N70657	001	Jan 24, 1989
AM PHARM PARTNERS	40MG/ML	N18549	001	Mar 11, 1983
	40MG/ML	N70012	001	Jun 12, 1985
	40MG/ML	N70058	001	Mar 20, 1985
	80MG/ML	N70013	001	Jun 12, 1985
	80MG/ML	N70059	001	Mar 20, 1985
	160MG/ML	N70364	001	Dec 04, 1985
ASTRAZENECA	40MG/ML	N18656	001	Jun 28, 1983
	40MG/ML	N70087	001	Oct 23, 1985
	80MG/ML	N70089	001	Oct 23, 1985
	80MG/ML	N70090	001	Oct 23, 1985
	80MG/ML	N70091	001	Oct 23, 1985
	160MG/ML	N70092	001	Oct 23, 1985
	160MG/ML	N70093	001	Oct 23, 1985
	160MG/ML	N70094	001	Oct 23, 1985
BAXTER HLTHCARE	40MG/ML	N18398	001	
	80MG/ML	N18398	002	Mar 22, 1982
SMITH AND NEPHEW	40MG/ML	N70011	001	Aug 29, 1985
	40MG/ML	N70046	001	Aug 29, 1985
	80MG/ML	N70047	001	Aug 29, 1985
WARNER CHILCOTT	40MG/ML	N18138	001	
	40MG/ML	N70558	001	Sep 20, 1985
	80MG/ML	N70559	001	Sep 20, 1985
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5%				
HOSPIRA	1.6MG/ML	N20542	001	Aug 30, 1995
INTROPIN				
MAYNE PHARMA USA	40MG/ML	N17395	001	
	80MG/ML	N17395	002	
	160MG/ML	N17395	003	

DOXACURIUM CHLORIDEINJECTABLE; INJECTION
NUROMAX

ABBOTT	EQ 1MG BASE/ML	N19946	001	Mar 07, 1991
--------	----------------	--------	-----	--------------

DOXEPIN HYDROCHLORIDECAPSULE; ORAL
DOXEPIN HYDROCHLORIDE

CLONMEL HLTHCARE	EQ 10MG BASE	N71685	001	Jan 05, 1988
	EQ 25MG BASE	N71686	001	Jan 05, 1988
	EQ 50MG BASE	N71673	001	Jan 05, 1988
	EQ 75MG BASE	N71674	001	Jan 05, 1988
	EQ 100MG BASE	N71675	001	Jan 05, 1988
	EQ 150MG BASE	N71676	001	Jan 05, 1988
MUTUAL PHARM	EQ 25MG BASE	N71502	001	Feb 18, 1988
	EQ 50MG BASE	N71653	001	Feb 18, 1988
	EQ 75MG BASE	N71654	001	Feb 18, 1988
	EQ 100MG BASE	N71521	001	Feb 18, 1988
NEW RIVER	EQ 10MG BASE	N16987	001	
	EQ 25MG BASE	N16987	002	
	EQ 50MG BASE	N16987	003	
	EQ 75MG BASE	N16987	006	
	EQ 100MG BASE	N16987	004	
PUREPAC PHARM	EQ 150MG BASE	N16987	007	Apr 13, 1987
	EQ 10MG BASE	N73054	001	Dec 28, 1990
	EQ 25MG BASE	N72109	001	Dec 28, 1990
	EQ 50MG BASE	N73055	001	Dec 28, 1990
	EQ 75MG BASE	N72386	001	Sep 08, 1988
	EQ 100MG BASE	N72110	001	Sep 08, 1988
	EQ 150MG BASE	N72387	001	Sep 08, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 91 (of 263)

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

QUANTUM PHARMICS

EQ 10MG BASE

N70972 001 Sep 29, 1987

EQ 25MG BASE

N70973 001 Sep 29, 1987

EQ 50MG BASE

N70931 001 Sep 29, 1987

EQ 75MG BASE

N70932 001 Sep 29, 1987

EQ 100MG BASE

N72375 001 Mar 15, 1989

EQ 150MG BASE

N72376 001 Mar 15, 1989

SANDOZ

EQ 25MG BASE

N70827 001 May 15, 1986

EQ 50MG BASE

N70828 001 May 15, 1986

EQ 75MG BASE

N70825 001 May 15, 1986

EQ 100MG BASE

N71562 001 Mar 02, 1987

WATSON LABS

EQ 10MG BASE

N70952 001 Mar 04, 1987

EQ 25MG BASE

N70953 001 May 15, 1986

EQ 50MG BASE

N70954 001 May 15, 1986

EQ 75MG BASE

N71763 001 Feb 09, 1988

EQ 100MG BASE

N70955 001 May 15, 1986

EQ 150MG BASE

N71764 001 Feb 09, 1988

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ADRIAMYCIN RDF

PHARMACIA AND UPJOHN

10MG/VIAL

N50467 001

20MG/VIAL

N50467 003 May 20, 1985

50MG/VIAL

N50467 002

150MG/VIAL

N50467 004 Jul 22, 1987

DOXYCYCLINE

FOR SUSPENSION; ORAL

DOXYCHEL

RACHELLE

EQ 25MG BASE/5ML

N61720 001

TABLET; ORAL

DOXYCYCLINE

PAR PHARM

EQ 75MG BASE

N65070 003 Dec 30, 2002

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE

HALSEY

EQ 50MG BASE

N62119 002 May 24, 1985

EQ 100MG BASE

N62119 001 May 24, 1985

HEATHER

EQ 50MG BASE

N62463 001 Dec 07, 1983

EQ 100MG BASE

N62463 002 Dec 07, 1983

INTERPHARM

EQ 50MG BASE

N62763 001 Sep 02, 1988

EQ 100MG BASE

N62763 002 Sep 02, 1988

MUTUAL PHARM

EQ 50MG BASE

N62418 001 Jan 28, 1983

EQ 100MG BASE

N62418 002 Jan 28, 1983

PAR PHARM

EQ 50MG BASE

N62434 001 Oct 19, 1984

EQ 100MG BASE

N62442 001 Dec 22, 1983

PVT FORM

EQ 50MG BASE

N62631 001 Jul 24, 1986

EQ 100MG BASE

N62631 002 Jul 24, 1986

RANBAXY

EQ 50MG BASE

N62479 001 Dec 23, 1983

EQ 100MG BASE

N62479 002 Dec 23, 1983

SUPERPHARM

EQ 50MG BASE

N62469 001 Oct 31, 1984

EQ 100MG BASE

N62469 002 Oct 31, 1984

WARNER CHILCOTT

EQ 50MG BASE

N62594 001 Dec 05, 1985

EQ 100MG BASE

N62594 002 Dec 05, 1985

WATSON LABS

EQ 50MG BASE

N62142 001

EQ 100MG BASE

N62142 002

DOXY-LEMMON

TEVA

EQ 50MG BASE

N62497 001 Aug 23, 1984

EQ 100MG BASE

N62497 002 Jun 15, 1984

PERIOSTAT

COLLAGENEX

EQ 20MG BASE

N50744 001 Sep 30, 1998

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 92 (of 263)

DOXYCYCLINE HYCLATE

CAPSULE, COATED PELLETS; ORAL

DOXYCYCLINE HYCLATE

PLIVA

EQ 100MG BASE

N63187 001

Jun 30, 1992

INJECTABLE; INJECTION

DOXYCHEL HYCLATE

RACHELLE

EQ 100MG BASE/VIAL

N61953 001

DOXYCYCLINE

BAXTER HLTHCARE

EQ 100MG BASE/VIAL

N62450 001

Oct 27, 1983

EQ 200MG BASE/VIAL

N62450 002

Oct 27, 1983

BEDFORD

EQ 100MG BASE/VIAL

N62569 001

Mar 09, 1988

EQ 200MG BASE/VIAL

N62569 002

Mar 09, 1988

DOXYCYCLINE HYCLATE

BAXTER HLTHCARE

EQ 100MG BASE/VIAL

N62992 001

Feb 16, 1989

EQ 200MG BASE/VIAL

N62992 002

Feb 16, 1989

VIBRAMYCIN

PFIZER

EQ 100MG BASE/VIAL

N50442 002

EQ 200MG BASE/VIAL

N50442 001

TABLET; ORAL

DOXYCYCLINE HYCLATE

HEATHER

EQ 100MG BASE

N62462 001

May 11, 1983

INTERPHARM

EQ 100MG BASE

N62764 001

Sep 02, 1988

MUTUAL PHARM

EQ 100MG BASE

N62391 001

Sep 30, 1982

SUPERPHARM

EQ 100MG BASE

N62494 001

Feb 20, 1985

TRUXTON INC

EQ 50MG BASE

N62269 003

WARNER CHILCOTT

EQ 100MG BASE

N62593 001

Aug 28, 1985

WATSON LABS

EQ 50MG BASE

N62392 001

Mar 31, 1983

EQ 100MG BASE

N62392 002

Mar 31, 1983

DOXY-LEMMON

TEVA

EQ 100MG BASE

N62581 001

Mar 15, 1985

DOXYLAMINE SUCCINATE

CAPSULE; ORAL

UNISOM

PFIZER

25MG

N19440 001

Feb 05, 1986

TABLET; ORAL

DECAPRYN

AVENTIS PHARMS

12.5MG

N06412 015

25MG

N06412 014

DOXYLAMINE SUCCINATE

QUANTUM PHARMICS

25MG

N88603 001

Aug 07, 1984

DOXY-SLEEP-AID

PAR PHARM

25MG

N70156 001

Jul 02, 1987

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

BENDECTIN

AVENTIS PHARMS

10MG;10MG **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N10598 002

DROMOSTANOLONE PROPIONATE

INJECTABLE; INJECTION

DROLBAN

LILLY

50MG/ML

N12936 001

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL

AM PHARM PARTNERS

2.5MG/ML

N70992 001

Nov 17, 1986

2.5MG/ML

N70993 001

Nov 17, 1986

ASTRAZENECA

2.5MG/ML

N72018 001

Oct 20, 1988

2.5MG/ML

N72019 001

Oct 19, 1988

2.5MG/ML

N72020 001

Oct 19, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 93 (of 263)

DROPERIDOLINJECTABLE; INJECTION
DROPERIDOL

ASTRAZENECA	2.5MG/ML	N72021	001	Oct 19, 1988
MAYNE PHARMA USA	2.5MG/ML	N71645	001	Apr 07, 1988
SMITH AND NEPHEW	2.5MG/ML	N71750	001	Sep 06, 1988
SOLOPAK	2.5MG/ML	N71754	001	Sep 06, 1988
	2.5MG/ML	N71755	001	Sep 06, 1988
STERIS	2.5MG/ML	N73520	001	Nov 27, 1991
	2.5MG/ML	N73521	001	Nov 27, 1991
	2.5MG/ML	N73523	001	Nov 27, 1991

DROPERIDOL; FENTANYL CITRATEINJECTABLE; INJECTION
FENTANYL CITRATE AND DROPERIDOL

ASTRAZENECA	2.5MG/ML;EQ 0.05MG BASE/ML	N72026	001	Apr 13, 1989
	2.5MG/ML;EQ 0.05MG BASE/ML	N72027	001	Apr 13, 1989
	2.5MG/ML;EQ 0.05MG BASE/ML	N72028	001	Apr 13, 1989
HOSPIRA	2.5MG/ML;EQ 0.05MG BASE/ML	N71982	001	May 04, 1988
INNOVAR				
AKORN MFG	2.5MG/ML;EQ 0.05MG BASE/ML	N16049	001	

DYCLONINE HYDROCHLORIDESOLUTION; TOPICAL
DYCLONE

ASTRAZENECA	0.5% **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N09925	002	
	1% **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N09925	001	

DYDROGESTERONETABLET; ORAL
GYNOREST

SOLVAY	5MG	N17388	001	
	10MG	N17388	002	

DYPHYLLINEELIXIR; ORAL
NEOTHYLLINE

TEVA	160MG/15ML	N07794	003	
------	------------	--------	-----	--

INJECTABLE; INJECTION
NEOTHYLLINE

TEVA	250MG/ML	N09088	001	
------	----------	--------	-----	--

TABLET; ORAL
NEOTHYLLINE

TEVA	200MG	N07794	001	
	400MG	N07794	002	

ECHOTHIOPHATE IODIDEFOR SOLUTION; OPHTHALMIC
PHOSPHOLINE IODIDE

AYERST	0.03%	N11963	002	
	0.06%	N11963	004	
	0.25%	N11963	003	

EDETATE CALCIUM DISODIUM

TABLET; ORAL

CALCIUM DISODIUM VERSENATE

3M	500MG	N08922	002	
----	-------	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 94 (of 263)

EDETATE DISODIUM

INJECTABLE; INJECTION				
DISODIUM EDETATE				
STERIS	150MG/ML	N84356	001	
EDETATE DISODIUM				
STERIS	150MG/ML	N80391	001	
SODIUM VERSENATE				
3M	200MG/ML	N10573	001	

EDROPHONIUM CHLORIDE

INJECTABLE; INJECTION				
EDROPHONIUM CHLORIDE				
STERIS	10MG/ML	N40044	001	Mar 20, 1996
EDROPHONIUM CHLORIDE PRESERVATIVE FREE				
STERIS	10MG/ML	N40043	001	Mar 20, 1996
REVERSOL				
ORGANON USA INC	10MG/ML	N89624	001	May 13, 1988

EFAVIRENZ

TABLET; ORAL				
SUSTIVA				
BRISTOL MYERS SQUIBB	300MG	N21360	001	Feb 01, 2002

EFLORNITHINE HYDROCHLORIDE

INJECTABLE; INJECTION				
ORNIDYL				
AVENTIS PHARMS	200MG/ML	N19879	002	Nov 28, 1990

ENALAPRIL MALEATE

TABLET; ORAL				
ENALAPRIL MALEATE				
APOTHECON	2.5MG	N75583	001	Aug 22, 2000
	5MG	N75583	002	Aug 22, 2000
	10MG	N75583	003	Aug 22, 2000
	20MG	N75583	004	Aug 22, 2000
IVAX PHARMS	2.5MG	N75482	001	Aug 22, 2000
	5MG	N75482	002	Aug 22, 2000
	10MG	N75482	003	Aug 22, 2000
	20MG	N75482	004	Aug 22, 2000

ENALAPRIL MALEATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL				
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE				
IVAX PHARMS	5MG;12.5MG	N75736	001	Mar 25, 2003
	10MG;25MG	N75736	002	Mar 25, 2003

ENFLURANE

LIQUID; INHALATION				
ENFLURANE				
ABBOTT	99.9%	N70803	001	Sep 08, 1987

ENOXACIN

TABLET; ORAL				
PENETREX				
AVENTIS	200MG	N19616	004	Dec 31, 1991
	400MG	N19616	005	Dec 31, 1991

ENOXAPARIN SODIUM

INJECTABLE; SUBCUTANEOUS				
LOVENOX (PRESERVATIVE FREE)				
AVENTIS	90MG/0.6ML (150MG/ML) **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N20164	006	Jun 02, 2000

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 95 (of 263)

EPINEPHRINEAEROSOL, METERED; INHALATION
BRONKAID MIST

STERLING 0.25MG/INH

N16803 001

INJECTABLE; INJECTION
SUS-PHRINE SULFITE-FREEFOREST LABS 1.5MG/AMP
5MG/MLN07942 003
N07942 001

Feb 05, 1999

INJECTABLE; INTRAMUSCULAR

EPI E Z PEN JR

MERIDIAN MEDCL TECHN 0.15MG/DELIVERY

N19430 004

Aug 03, 1995

EPIPEN E Z PEN

MERIDIAN MEDCL TECHN 0.3MG/DELIVERY

N19430 003

Aug 03, 1995

EPINEPHRINE BITARTRATEAEROSOL, METERED; INHALATION
MEDIHALER-EPI

3M 0.3MG/INH

N10374 003

EPINEPHRINE BITARTRATE; ETIDOCAINE HYDROCHLORIDEINJECTABLE; INJECTION
DURANEST

ASTRAZENECA 0.005MG/ML;1%

N17751 006

0.005MG/ML;1.5%

N17751 007

DENTSPLY PHARM 0.005MG/ML;1.5%

N21384 001

EPINEPHRINE BITARTRATE; PRILOCAINE HYDROCHLORIDEINJECTABLE; INJECTION
CITANEST FORTE

ASTRAZENECA 0.005MG/ML;4%

N14763 008

EPINEPHRINE; ETIDOCAINE HYDROCHLORIDEINJECTABLE; INJECTION
DURANEST

ASTRAZENECA 0.005MG/ML;0.5%

N17751 004

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ALPHACAINE HYDROCHLORIDE W/ EPINEPHRINE

CARLISLE 0.01MG/ML;2%

N84720 001

0.02MG/ML;2%

N84732 001

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

ELKINS SINN 0.01MG/ML;1%

N80406 001

0.01MG/ML;2%

N80406 002

LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE

ABBOTT 0.01MG/ML;1%

N83154 001

BEL MAR 0.01MG/ML;1%

N80820 001

0.01MG/ML;2%

N80757 001

INTL MEDICATION 0.01MG/ML;1%

N86402 001

STERIS 0.01MG/ML;1%

N80377 003

0.01MG/ML;1%

N85463 001

0.01MG/ML;2%

N80377 004

LIDOCATON

PHARMATON 0.02MG/ML;2%

N84728 001

Aug 17, 1983

XYLOCAINE W/ EPINEPHRINE

ASTRAZENECA 0.005MG/ML;1%

N10418 006

0.005MG/ML;1.5%

N10418 010

0.005MG/ML;2%

N10418 008

0.01MG/ML;2%

N06488 003

0.02MG/ML;2%

N06488 005

SOLUTION; IONTOPHORESIS

IONTOCAINE

IOMED 0.01MG/ML;2%

N20530 001

Dec 21, 1995

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 96 (of 263)

EPINEPHRINE; PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE W/ EPINEPHRINE

BEL MAR

0.02MG/ML;1%

N80758 001

0.02MG/ML;2%

N80759 001

EPLERENONE

TABLET; ORAL

INSPRA

GD SEARLE LLC

100MG

N21437 003

Sep 27, 2002

EPROSARTAN MESYLATE

TABLET; ORAL

TEVETEN

KOS LIFE

EQ 300MG BASE

N20738 004

Dec 22, 1997

ERGOCALCIFEROL

CAPSULE; ORAL

DELTALIN

LILLY

50,000 IU

N80884 001

VITAMIN D

CHASE CHEM

50,000 IU

N80747 001

EVERYLIFE

50,000 IU

N80956 001

IMPAX LABS

50,000 IU

N80951 001

LANNETT

50,000 IU

N80825 001

VITARINE

50,000 IU

N84053 001

WEST WARD

50,000 IU

N83102 001

ERGOLOID MESYLATES

CAPSULE; ORAL

HYDERGINE LC

NOVARTIS

1MG

N18706 001

Jan 18, 1983

SOLUTION; ORAL

HYDERGINE

NOVARTIS

1MG/ML

N18418 001

TABLET; ORAL

ERGOLOID MESYLATES

MUTUAL PHARM

1MG

N88891 001

Nov 01, 1985

WATSON LABS

1MG

N86433 001

May 27, 1982

1MG

N87244 001

Aug 16, 1982

GERIMAL

WATSON LABS

1MG

N88207 001

Mar 22, 1984

HYDERGINE

NOVARTIS

0.5MG

N17993 003

TABLET; SUBLINGUAL

ALKERGOT

SANDOZ

0.5MG

N85153 001

1MG

N87417 001

CIRCANOL

3M

0.5MG

N84868 001

1MG

N85809 001

DEAPRIL-ST

BRISTOL MYERS SQUIBB

1MG

N85020 002

ERGOLOID MESYLATES

KV PHARM

0.5MG

N85899 001

0.5MG

N86265 001

1MG

N85900 001

1MG

N86264 001

LEDERLE

0.5MG

N86984 001

1MG

N86985 001

MUTUAL PHARM

0.5MG

N87407 001

1MG

N87552 001

SUPERPHARM

0.5MG

N89233 001

Sep 23, 1986

1MG

N89234 001

Sep 23, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6- 97 (of 263)

ERGOLOID MESYLATES

TABLET; SUBLINGUAL				
ERGOLOID MESYLATES				
VANGARD	0.5MG	N88013	001	Sep 20, 1982
	1MG	N88014	001	Sep 20, 1982
WATSON LABS	0.5MG	N84930	001	
	1MG	N85177	001	
	1MG	N87183	001	
HYDROGENATED ERGOT ALKALOIDS				
IVAX PHARMS	0.5MG	N87186	001	

ERGOTAMINE TARTRATE

AEROSOL, METERED; INHALATION				
MEDIHALER ERGOTAMINE				
3M	0.36MG/INH	N12102	001	
TABLET; SUBLINGUAL				
ERGOSTAT				
PARKE DAVIS	2MG	N88337	001	Jun 08, 1984
WIGRETTES				
ORGANON USA INC	2MG	N86750	001	Jul 29, 1982

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL				
ERYC				
PARKE DAVIS	250MG	N62546	001	Jul 25, 1985
	250MG	N62618	001	Sep 25, 1985
ERYC 125				
PARKE DAVIS	125MG	N62648	001	Oct 24, 1985
ERYC SPRINKLES				
MAYNE PHARMA USA	125MG	N50593	001	Jul 22, 1985
GEL; TOPICAL				
EMGEL				
ALTANA	2%	N63107	001	Aug 23, 1991
OINTMENT; OPHTHALMIC				
ERYTHROMYCIN				
PHARMADERM	5MG/GM	N62446	001	Sep 26, 1983
PHARMAFAIR	5MG/GM	N62481	001	Apr 05, 1984
ILOTYCIN				
DISTA	0.5%	N50368	001	
POWDER; FOR RX COMPOUNDING				
ERYTHROMYCIN				
PADDOCK	100%	N50610	001	Nov 07, 1986
SOLUTION; TOPICAL				
C-SOLVE-2				
BIOGLAN PHARMA	2%	N62468	001	Jul 03, 1985
ERYMAX				
MERZ PHARMS	2%	N62508	002	Jul 11, 1985
ERYTHROMYCIN				
ALPHARMA US PHARMS	1.5%	N62328	001	Apr 19, 1982
	2%	N62326	001	Apr 19, 1982
	2%	N62327	001	Apr 19, 1982
	2%	N62342	001	Feb 25, 1982
	2%	N62957	001	Jul 21, 1988
BAUSCH AND LOMB	2%	N64039	001	Jan 27, 1994
LILLY	2%	N50532	001	
PHARMAFAIR	1.5%	N62485	001	Jul 11, 1984
	2%	N62616	001	Jul 25, 1985
STIEFEL	2%	N64127	001	Feb 14, 1997
ERYTHRO-STATIN				
HI TECH PHARMA	2%	N64101	001	Oct 22, 1996
SANSAC				
HEALTHPOINT	2%	N62522	001	Jan 24, 1985
STATICIN				
WESTWOOD SQUIBB	1.5%	N50526	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 98 (of 263)

ERYTHROMYCIN

SOLUTION; TOPICAL

T-STAT

WESTWOOD SQUIBB 2%

N62436 001 Mar 09, 1983

SWAB; TOPICAL

T-STAT

WESTWOOD SQUIBB 2%

N62748 001 Jul 23, 1987

TABLET, DELAYED RELEASE; ORAL

E-BASE

BARR 333MG

N63028 001 May 15, 1990

ILOTYCIN

DISTA 250MG

N61910 001

ROBIMYCIN

ROBINS AH 250MG

N61633 001

R-P MYCIN

SOLVAY 250MG

N61659 001

ERYTHROMYCIN ESTOLATE

CAPSULE; ORAL

ERYTHROMYCIN ESTOLATE

BARR EQ 125MG BASE

N62162 001

EQ 250MG BASE

N62162 002

IVAX PHARMS

EQ 250MG BASE

N62237 001

WATSON LABS

EQ 250MG BASE

N62087 001

ILOSONE

LILLY EQ 125MG BASE

N61897 001

EQ 250MG BASE

N61897 002

FOR SUSPENSION; ORAL

ILOSONE

DISTA EQ 125MG BASE/5ML

N61893 001

SUSPENSION; ORAL

ERYTHROMYCIN ESTOLATE

BARR EQ 125MG BASE/5ML

N62169 001 Oct 17, 1990

EQ 250MG BASE/5ML

N62169 002 Oct 17, 1990

LIFE LABS

EQ 250MG BASE/5ML

N62362 001 Dec 17, 1982

ILOSONE

LILLY EQ 125MG BASE/5ML

N50010 001

EQ 125MG BASE/5ML

N61894 001

EQ 250MG BASE/5ML

N50010 002

EQ 250MG BASE/5ML

N61894 002

SUSPENSION/DROPS; ORAL

ILOSONE

LILLY EQ 100MG BASE/ML

N61894 003

TABLET; ORAL

ILOSONE

LILLY EQ 500MG BASE

N61896 001

TABLET, CHEWABLE; ORAL

ILOSONE

DISTA EQ 125MG BASE

N61895 001

EQ 250MG BASE

N61895 002

ERYTHROMYCIN ESTOLATE; SULFISOXAZOLE ACETYL

SUSPENSION; ORAL

ILOSONE SULFA

LILLY EQ 125MG BASE/5ML;EQ 600MG BASE/5ML

N50599 001 Sep 29, 1989

ERYTHROMYCIN ETHYLSUCCINATE

GRANULE; ORAL

PEDIAMYCIN

ROSS LABS EQ 200MG BASE/5ML

N62305 001

SUSPENSION; ORAL

E-MYCIN E

PHARMACIA AND UPJOHN EQ 200MG BASE/5ML

N62198 001

EQ 400MG BASE/5ML

N62198 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 99 (of 263)

ERYTHROMYCIN ETHYLSUCCINATE

SUSPENSION; ORAL

ERYTHROMYCIN ETHYLSUCCINATE

DISTA	EQ 200MG BASE/5ML	N62177	001	
	EQ 400MG BASE/5ML	N62177	002	
NASKA	EQ 400MG BASE/5ML	N62674	001	Mar 10, 1987
PARKE DAVIS	EQ 200MG BASE/5ML	N62231	001	
	EQ 400MG BASE/5ML	N62231	002	
PHARMAFAIR	EQ 200MG BASE/5ML	N62559	001	Mar 15, 1985
	EQ 400MG BASE/5ML	N62558	001	Mar 15, 1985
WYAMYCIN E				
WYETH AYERST	EQ 200MG BASE/5ML	N62123	002	
	EQ 400MG BASE/5ML	N62123	001	

SUSPENSION/DROPS; ORAL

PEDIAMYCIN

ROSS LABS	EQ 100MG BASE/2.5ML	N62305	002	
-----------	---------------------	--------	-----	--

TABLET; ORAL

E.E.S. 400

ABBOTT	EQ 400MG BASE	N61905	001	
--------	---------------	--------	-----	--

ERYTHROMYCIN ETHYLSUCCINATE

BARR	EQ 400MG BASE	N62256	001	
------	---------------	--------	-----	--

TABLET, CHEWABLE; ORAL

PEDIAMYCIN

ROSS LABS	EQ 200MG BASE	N62306	001	
-----------	---------------	--------	-----	--

ERYTHROMYCIN GLUCEPTATE

INJECTABLE; INJECTION

ILOTYCIN GLUCEPTATE

DISTA	EQ 250MG BASE/VIAL	N50370	001	
	EQ 500MG BASE/VIAL	N50370	002	
	EQ 1GM BASE/VIAL	N50370	003	

ERYTHROMYCIN LACTOBIONATE

INJECTABLE; INJECTION

ERYTHROCIN

ABBOTT	EQ 500MG BASE/VIAL	N62586	001	Jan 04, 1988
	EQ 1GM BASE/VIAL	N62586	002	Jan 04, 1988

ERYTHROMYCIN

ELKINS SINN	EQ 500MG BASE/VIAL	N62563	001	Mar 28, 1985
	EQ 1GM BASE/VIAL	N62563	002	Mar 28, 1985

ERYTHROMYCIN LACTOBIONATE

AM PHARM PARTNERS	EQ 500MG BASE/VIAL	N62604	001	Nov 24, 1986
	EQ 1GM BASE/VIAL	N62604	002	Nov 24, 1986

ERYTHROMYCIN STEARATE

TABLET; ORAL

BRISTAMYCIN

BRISTOL	EQ 250MG BASE	N61304	001	
	EQ 250MG BASE	N61887	001	

ERYPAR

PARKE DAVIS	EQ 250MG BASE	N62032	001	
	EQ 500MG BASE	N62032	002	
WARNER CHILCOTT	EQ 250MG BASE	N62322	001	

ERYTHROCIN STEARATE

ABBOTT	EQ 125MG BASE	N60359	002	
--------	---------------	--------	-----	--

ERYTHROMYCIN STEARATE

BARR	EQ 500MG BASE	N63179	001	May 15, 1990
IVAX PHARMS	EQ 250MG BASE	N61461	001	
	EQ 500MG BASE	N61461	002	
LEDERLE	EQ 250MG BASE	N62089	001	
	EQ 500MG BASE	N62089	002	
PUREPAC PHARM	EQ 250MG BASE	N61743	001	
WATSON LABS	EQ 250MG BASE	N62121	002	
	EQ 500MG BASE	N62121	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 100 (of 263)

ERYTHROMYCIN STEARATE

TABLET; ORAL

ETHRIL 250

BRISTOL MYERS SQUIBB EQ 250MG BASE

N61605 001

ETHRIL 500

BRISTOL MYERS SQUIBB EQ 500MG BASE

N61605 002

PFIZER-E

PFIZER

EQ 250MG BASE

N61791 001

EQ 500MG BASE

N61791 002

WYAMYCIN S

WYETH AYERST

EQ 250MG BASE

N61675 001

EQ 500MG BASE

N61675 002

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

BREVIBLOC

BAXTER HLTHCARE CORP 100MG/ML

N19386 003 Dec 31, 1986

ESTRADIOL

FILM, EXTENDED RELEASE; TRANSDERMAL

ESCLIM

WOMEN FIRST HLTHCARE 0.025MG/24HR

N20847 001 Aug 04, 1998

0.0375MG/24HR

N20847 002 Aug 04, 1998

0.05MG/24HR

N20847 003 Aug 04, 1998

0.075MG/24HR

N20847 004 Aug 04, 1998

0.1MG/24HR

N20847 005 Aug 04, 1998

ESTRADIOL

ORTHO MCNEIL PHARM

0.05MG/24HR

N21048 001 Sep 20, 1999

0.075MG/24HR

N21048 002 Sep 20, 1999

0.1MG/24HR

N21048 003 Sep 20, 1999

FEMPATCH

PARKE DAVIS

0.025MG/24HR

N20417 001 Dec 03, 1996

VIVELLE

NOVARTIS

0.025MG/24HR

N20323 005 Aug 16, 2000

0.0375MG/24HR

N20323 001 Oct 28, 1994

0.075MG/24HR

N20323 003 Oct 28, 1994

GEL; TOPICAL

ESTROGEL

SOLVAY

0.06%

N21166 001 Feb 09, 2004

TABLET; ORAL

ESTRADIOL

CLONMEL HLTHCARE

0.5MG

N40275 001 Dec 29, 1998

1MG

N40275 002 Dec 29, 1998

2MG

N40275 003 Dec 29, 1998

ESTRADIOL CYPIONATE

INJECTABLE; INJECTION

DEPO-ESTRADIOL

PHARMACIA AND UPJOHN 1MG/ML

N85470 001

3MG/ML

N85470 002

ESTRADIOL CYPIONATE; MEDROXYPROGESTERONE ACETATE

INJECTABLE; INTRAMUSCULAR

LUNELLE

PHARMACIA AND UPJOHN 5MG/0.5ML; 25MG/0.5ML

N20874 001 Oct 05, 2000

ESTRADIOL CYPIONATE; TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

DEPO-TESTADIOL

PHARMACIA AND UPJOHN 2MG/ML; 50MG/ML

N17968 001

TESTOSTERONE CYPIONATE-ESTRADIOL CYPIONATE

STERIS

2MG/ML; 50MG/ML

N85603 001 Mar 13, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 101 (of 263)

ESTRADIOL VALERATE

INJECTABLE; INJECTION
ESTRADIOL VALERATE
STERIS

10MG/ML

N83546 001

ESTRADIOL VALERATE; TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION
DITATE-DS

SAVAGE LABS

8MG/ML;180MG/ML

N86423 001

TESTOSTERONE ENANTHATE AND ESTRADIOL VALERATE

STERIS

4MG/ML;90MG/ML

N85865 001

8MG/ML;180MG/ML

N85860 001

ESTROGENS, CONJUGATED

TABLET; ORAL
PREMARIN

WYETH PHARMS INC

2.5MG

N04782 002

ESTROGENS, CONJUGATED SYNTHETIC B

TABLET; ORAL
ENJUVIA

DURAMED

0.3MG

N21443 001

Dec 20, 2004

0.45MG

N21443 002

Dec 20, 2004

0.625MG

N21443 003

May 10, 2004

1.25MG

N21443 004

May 10, 2004

ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE

TABLET; ORAL-28

PREMPHASE (PREMARIN;CYCRIN 14/14)

WYETH PHARMS INC

0.625MG,0.625MG;N/A,5MG

N20303 002

Dec 30, 1994

PREMPRO (PREMARIN;CYCRIN)

WYETH PHARMS INC

0.625MG,0.625MG;2.5MG,2.5MG

N20303 001

Dec 30, 1994

ESTROGENS, CONJUGATED; MEPROBAMATE

TABLET; ORAL

MILPREM-200

MEDPOINTE PHARM HLC

0.45MG;200MG

N11045 002

MILPREM-400

MEDPOINTE PHARM HLC

0.45MG;400MG

N11045 001

PMB 200

WYETH AYERST

0.45MG;200MG

N10971 005

PMB 400

WYETH AYERST

0.45MG;400MG

N10971 003

ESTROGENS, ESTERIFIED

TABLET; ORAL

AMNESTROGEN

BRISTOL MYERS SQUIBB

0.3MG

N83266 001

0.625MG

N83266 002

1.25MG

N83266 003

2.5MG

N83266 004

ESTERIFIED ESTROGENS

PVT FORM

0.625MG

N83414 001

1.25MG

N83765 001

2.5MG

N85907 001

SANDOZ

1.25MG

N85302 001

ESTRATAB

SOLVAY

0.3MG

N86715 001

0.625MG

N83209 001

1.25MG

N83856 001

2.5MG

N83857 001

EVEX

ROCHE PALO

0.625MG

N84215 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 102 (of 263)

ESTROGENS, ESTERIFIED

TABLET; ORAL

EVEX

ROCHE PALO

1.25MG

N83376 002

FEMOGEN

PVT FORM

0.625MG

N85076 001

1.25MG

N85008 001

2.5MG

N85007 001

ESTRONE

INJECTABLE; INJECTION

ESTROGENIC SUBSTANCE

WYETH AYERST

2MG/ML

N83488 001

ESTRONE

STERIS

2MG/ML

N83397 001

NATURAL ESTROGENIC SUBSTANCE-ESTRONE

STERIS

2MG/ML

N85237 001

Nov 23, 1982

THEELIN

PARKEDALE

1MG/ML

N03977 001

2MG/ML

N03977 002

5MG/ML

N03977 003

ETHACRYNIC ACID

TABLET; ORAL

EDECRIN

MERCK

50MG

N16092 002

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL

MYAMBUTOL

STAT TRADE

200MG

N16320 002

500MG

N16320 004

ETHCHLORVYNOL

CAPSULE; ORAL

ETHCHLORVYNOL

BANNER PHARMACAPS

100MG

N84463 001

200MG

N84463 002

500MG

N84463 003

750MG

N84463 004

PLACIDYL

ABBOTT

100MG

N10021 004

200MG

N10021 007

500MG

N10021 002

750MG

N10021 010

ETHINAMATE

CAPSULE; ORAL

VALMID

DISTA

500MG

N09750 001

ETHINYL ESTRADIOL

TABLET; ORAL

ESTINYL

SCHERING

0.02MG

N05292 001

0.05MG

N05292 002

0.5MG

N05292 003

FEMINONE

PHARMACIA AND UPJOHN

0.05MG

N16649 001

LYNORAL

ORGANON USA INC

0.01MG

N05490 003

0.05MG

N05490 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 103 (of 263)

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE

TABLET; ORAL-28		
NORQUEST FE		
GD SEARLE LLC	0.035MG;75MG;1MG	N18926 001 Jul 18, 1986

ETHINYL ESTRADIOL; FERROUS FUMARATE; NORETHINDRONE ACETATE

TABLET; ORAL-28		
NORLESTRIN FE 1/50		
PARKE DAVIS	0.05MG;75MG;1MG	N16766 001
NORLESTRIN FE 2.5/50		
PARKE DAVIS	0.05MG;75MG;2.5MG	N16854 001

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL		
PREVEN EMERGENCY CONTRACEPTIVE KIT		
DURAMED	0.05MG;0.25MG	N20946 001 Sep 01, 1998
TABLET; ORAL-21		
ALESSE		
WYETH PHARMS INC	0.02MG;0.1MG	N20683 001 Mar 27, 1997
AVIANE-21		
DURAMED PHARMS BARR	0.02MG;0.1MG	N75796 002 Apr 30, 2001
ENPRESSE-21		
DURAMED PHARMS BARR	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	N75809 001 Jul 16, 2001
LESSINA-21		
BARR	0.02MG;0.1MG	N75803 001 Mar 20, 2002
LEVLITE		
BERLEX	0.02MG;0.1MG	N20860 001 Jul 13, 1998
LEVONORGESTREL AND ETHINYL ESTRADIOL		
BARR	0.02MG;0.1MG	N75862 001 Apr 29, 2003
LEVORA 0.15/30-21		
WATSON LABS	0.03MG;0.15MG	N73592 001 Dec 13, 1993
NORDETTE-21		
DURAMED	0.03MG;0.15MG	N18668 001 May 10, 1982
PORTIA-21		
BARR	0.03MG;0.15MG	N75866 001 May 23, 2002
TRIPHASIL-21		
WYETH PHARMS INC	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	N19192 001 Nov 01, 1984
TRIVORA-21		
WATSON LABS	0.03MG,0.04MG,0.03MG;0.05MG,0.075MG,0.125MG	N74538 001 Dec 18, 1997

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21		
MODICON 21		
ORTHO MCNEIL PHARM	0.035MG;0.5MG	N17488 001
N.E.E. 1/35 21		
LPI	0.035MG;1MG	N71541 001 Dec 14, 1987
NORCEPT-E 1/35 21		
ORTHO MCNEIL PHARM	0.035MG;1MG	N71545 001 Feb 09, 1989
ORTHO-NOVUM 1/35-21		
ORTHO MCNEIL PHARM	0.035MG;1MG	N17489 002
ORTHO-NOVUM 10/11-21		
ORTHO MCNEIL PHARM	0.035MG,0.035MG;0.5MG,1MG	N18354 001 Jan 11, 1982
ORTHO-NOVUM 7/14-21		
ORTHO MCNEIL PHARM	0.035MG,0.035MG;0.5MG,1MG	N19004 001 Apr 04, 1984
ORTHO-NOVUM 7/7/7-21		
ORTHO MCNEIL PHARM	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	N18985 001 Apr 04, 1984
OVCON-35		
WARNER CHILCOTT	0.035MG;0.4MG	N18127 001
OVCON-50		
WARNER CHILCOTT	0.05MG;1MG	N18128 001

DISCONTINUED DRUG PRODUCT LIST

6 - 104 (of 263)

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21					
TRI-NORINYL 21-DAY					
WATSON LABS	0.035MG, 0.035MG, 0.035MG; 0.5MG, 1MG, 0.5MG	N18977 001	Apr 13, 1984		
TABLET; ORAL-28					
N.E.E. 1/35 28					
LPI	0.035MG; 1MG	N71542 001	Dec 14, 1987		
NORCEPT-E 1/35 28					
ORTHO MCNEIL PHARM	0.035MG; 1MG	N71546 001	Feb 09, 1989		
ORTHO-NOVUM 7/14-28					
ORTHO MCNEIL PHARM	0.035MG, 0.035MG; 0.5MG, 1MG	N19004 002	Apr 04, 1984		

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-21					
ESTROSTEP 21					
WARNER CHILCOTT	0.02MG, 0.03MG, 0.035MG; 1MG, 1MG, 1MG	N20130 001	Oct 09, 1996		
NORLESTRIN 21 1/50					
PARKE DAVIS	0.05MG; 1MG	N16749 001			
NORLESTRIN 21 2.5/50					
PARKE DAVIS	0.05MG; 2.5MG	N16852 001			
TABLET; ORAL-28					
NORLESTRIN 28 1/50					
PARKE DAVIS	0.05MG; 1MG	N16723 001			

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-21					
ORTHO CYCLEN-21					
JOHNSON RW	0.035MG; 0.25MG	N19653 001	Dec 29, 1989		
ORTHO TRI-CYCLEN					
JOHNSON RW	0.035MG, 0.035MG, 0.035MG; 0.18MG, 0.215MG, 0.25MG	N19697 002	Jul 03, 1992		

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21					
LO/OVRAL					
WYETH PHARMS INC	0.03MG; 0.3MG	N17612 001			
OVRAL					
WYETH PHARMS INC	0.05MG; 0.5MG	N16672 001			
TABLET; ORAL-28					
OVRAL-28					
WYETH PHARMS INC	0.05MG; 0.5MG	N16806 001			

ETHOPROPAZINE HYDROCHLORIDE

TABLET; ORAL					
PARSIDOL					
PARKE DAVIS	10MG	N09078 003			
	50MG	N09078 006			
	100MG	N09078 008			

ETHOTOIN

TABLET; ORAL					
PEGANONE					
OVATION PHARMS	500MG	N10841 003			

ETHOXZOLAMIDE

TABLET; ORAL					
CARDRASE					
PHARMACIA AND UPJOHN	62.5MG	N11047 002			
	125MG	N11047 001			
ETHAMIDE					
ALLERGAN	125MG	N16144 001			

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 105 (of 263)

ETHYLESTRENOL

ELIXIR; ORAL MAXIBOLIN ORGANON USA INC	2MG/5ML	N14006	002	
TABLET; ORAL MAXIBOLIN ORGANON USA INC	2MG	N14005	002	

ETHYNODIOL DIACETATE; MESTRANOL

TABLET; ORAL-20 OVULEN GD SEARLE LLC	1MG;0.1MG	N16029	002	
TABLET; ORAL-21 OVULEN-21 GD SEARLE LLC	1MG;0.1MG	N16029	003	
TABLET; ORAL-28 OVULEN-28 GD SEARLE LLC	1MG;0.1MG	N16705	001	

ETIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION DURANEST ASTRAZENECA	0.5%	N17751	003	
	1%	N17751	005	

ETIDRONATE DISODIUM

INJECTABLE; INJECTION DIDRONEL MGI PHARMA INC	50MG/ML	N19545	001	Apr 20, 1987
---	---------	--------	-----	--------------

ETODOLAC

CAPSULE; ORAL ETODOLAC IVAX PHARMS	200MG 300MG	N74899	001	Jul 08, 1997
		N74899	002	Jul 08, 1997
LODINE WYETH PHARMS INC	200MG	N18922	002	Jan 31, 1991
TABLET; ORAL LODINE WYETH PHARMS INC	400MG 500MG	N18922	004	Jul 29, 1993
		N18922	005	Jun 28, 1996
TABLET, EXTENDED RELEASE; ORAL ETODOLAC POINT HOLDINGS	400MG	N75696	001	Jul 31, 2000
LODINE XL WYETH PHARMS INC	400MG 500MG 600MG	N20584	001	Oct 25, 1996
		N20584	003	Jan 20, 1998
		N20584	002	Oct 25, 1996

ETOPOSIDE

CAPSULE; ORAL VEPESID BRISTOL MYERS SQUIBB	100MG	N19557	002	Dec 30, 1986
INJECTABLE; INJECTION ETOPOSIDE PIERRE FABRE	20MG/ML	N74813	001	Jul 09, 1997
SICOR PHARMS	20MG/ML	N74510	001	Jun 29, 1995
STERIS	20MG/ML	N74228	001	Oct 15, 1996

ETOPOSIDE PHOSPHATE

INJECTABLE; INJECTION ETOPOPHOS PRESERVATIVE FREE BRISTOL MYERS SQUIBB	EQ 500MG BASE/VIAL EQ 1GM BASE/VIAL	N20906	001	Feb 27, 1998
		N20906	002	Feb 27, 1998

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 106 (of 263)

ETRETINATECAPSULE; ORAL
TEGISON
ROCHE10MG
25MGN19369 001 Sep 30, 1986
N19369 002 Sep 30, 1986EVANS BLUEINJECTABLE; INJECTION
EVANS BLUE
PARKE DAVIS0.5% **Federal Register determination
that product was not discontinued or
withdrawn for safety or efficacy
reasons**

N08041 001

FAMOTIDINEINJECTABLE; INJECTION
FAMOTIDINEAPOTHECON 10MG/ML
MAYNE PHARMA USA 10MG/MLN75707 001 Apr 16, 2001
N75705 001 Apr 16, 2001

FAMOTIDINE PRESERVATIVE FREE

APOTHECON 10MG/ML
MAYNE PHARMA USA 10MG/MLN75708 001 Apr 16, 2001
N75669 001 Apr 16, 2001

FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER

ABBOTT 0.4MG/ML

N75729 001 Dec 17, 2001

TABLET, ORALLY DISINTEGRATING; ORAL
PEPCID RPDMERCK 20MG
40MGN20752 001 May 28, 1998
N20752 002 May 28, 1998FENOFIBRATECAPSULE; ORAL
ANTARA (MICRONIZED)

RELIANT PHARMS INC 87MG

N21695 002 Nov 30, 2004

LIPIDIL

ABBOTT 100MG

N19304 001 Dec 31, 1993

TRICOR (MICRONIZED)

ABBOTT 67MG
134MG
200MGN19304 002 Feb 09, 1998
N19304 003 Jun 30, 1999
N19304 004 Jun 30, 1999FENOPROFEN CALCIUMCAPSULE; ORAL
FENOPROFEN CALCIUMAM THERAP EQ 200MG BASE
EQ 300MG BASEN72307 001 Aug 22, 1988
N72308 001 Aug 22, 1988HALSEY EQ 200MG BASE
EQ 300MG BASEN72355 001 Aug 17, 1988
N72356 001 Aug 17, 1988QUANTUM PHARMICS EQ 200MG BASE
EQ 300MG BASEN72214 001 Aug 17, 1988
N71738 001 Aug 17, 1988WARNER CHILCOTT EQ 200MG BASE
EQ 300MG BASEN72946 001 Apr 30, 1991
N72472 001 Apr 30, 1991WATSON LABS EQ 200MG BASE
EQ 200MG BASEN72294 001 Aug 17, 1988
N72981 001 Aug 19, 1991

EQ 300MG BASE

N72293 001 Aug 17, 1988

EQ 300MG BASE

N72982 001 Aug 19, 1991

TABLET; ORAL
FENOPROFEN CALCIUMAM THERAP EQ 600MG BASE
EQ 600MG BASEN72309 001 Aug 17, 1988
N72357 001 Aug 17, 1988

HALSEY EQ 600MG BASE

N72194 001 Aug 17, 1988

QUANTUM PHARMICS EQ 600MG BASE

N72362 001 Aug 17, 1988

USL PHARMA EQ 600MG BASE

N72165 001 Aug 17, 1988

WATSON LABS EQ 600MG BASE

NALFON

DISTA EQ 600MG BASE

N17710 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 107 (of 263)

FENTANYL CITRATE

INJECTABLE; INJECTION

FENTANYL CITRATE

ABBOTT

EQ 0.05MG BASE/ML

N70636 001 Apr 30, 1990

EQ 0.05MG BASE/ML

N70637 001 Apr 30, 1990

STERIS

EQ 0.05MG BASE/ML

N73488 001 Jun 30, 1992

FENTANYL CITRATE PRESERVATIVE FREE

MARSAM PHARMS LLC

EQ 0.05MG BASE/ML

N74917 001 Feb 03, 1998

TROCHE/LOZENGE; ORAL

FENTANYL

ANESTA

EQ 0.1MG BASE

N20195 007 Oct 30, 1995

EQ 0.2MG BASE

N20195 001 Oct 04, 1993

EQ 0.3MG BASE

N20195 002 Oct 04, 1993

EQ 0.4MG BASE

N20195 003 Oct 04, 1993

FERRIC AMMONIUM CITRATE

FOR SOLUTION; ORAL

FERRISELTZ

OTSUKA

600MG/PACKET

N20292 001 Oct 14, 1997

FERROUS CITRATE, FE-59

INJECTABLE; INJECTION

FERROUS CITRATE FE 59

MALLINCKRODT

25uCi/ML

N16729 001

FERROUS SULFATE; FOLIC ACID

CAPSULE; ORAL

FOLVRON

LEDERLE

182MG;0.33MG

N06012 003

FIBRINOGEN, I-125

INJECTABLE; INJECTION

IBRIN

GE HEALTHCARE

154uCi/VIAL

N17879 001

RADIONUCLIDE-LABELED (125 I) FIBRINOGEN (HUMAN) SENSOR

ABBOTT

140uCi/ML

N17787 001

FLECAINIDE ACETATE

TABLET; ORAL

TAMBOCOR

3M

200MG

N18830 002 Oct 31, 1985

FLUCONAZOLE

INJECTABLE; INJECTION

DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER

PFIZER

2MG/ML

N19950 005 Jul 08, 1994

DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

PFIZER

2MG/ML

N19950 004 Jul 08, 1994

FLUDEOXYGLUCOSE F-18

INJECTABLE; INJECTION

FLUDEOXYGLUCOSE F 18

DOWNSTATE CLINCL

4-40mCi/ML

N20306 001 Aug 19, 1994

4-90mCi/ML

N20306 002 Sep 25, 2001

FLUMETHASONE PIVALATE

CREAM; TOPICAL

LOCORTEN

NOVARTIS

0.03%

N16379 001

DISCONTINUED DRUG PRODUCT LIST

6 - 108 (of 263)

FLUNISOLIDE

SPRAY, METERED; NASAL
NASALIDE

IVAX RES	0.025MG/SPRAY	N18148	001	
----------	---------------	--------	-----	--

FLUOCINOLONE ACETONIDE

CREAM; TOPICAL

FLUOCET

ALPHARMA US PHARMS	0.025%	N88360	001	Jan 16, 1984
--------------------	--------	--------	-----	--------------

FLUOCINOLONE ACETONIDE

ALPHARMA US PHARMS	0.01%	N88361	001	Jan 16, 1984
--------------------	-------	--------	-----	--------------

CLAY PARK	0.01%	N86810	001	Mar 04, 1982
-----------	-------	--------	-----	--------------

	0.025%	N86811	001	Mar 04, 1982
--	--------	--------	-----	--------------

PHARMADERM	0.01%	N88047	001	Dec 16, 1982
------------	-------	--------	-----	--------------

	0.025%	N88045	001	Dec 16, 1982
--	--------	--------	-----	--------------

PHARMAFAIR	0.01%	N88499	001	Aug 02, 1984
------------	-------	--------	-----	--------------

	0.025%	N88506	001	Aug 02, 1984
--	--------	--------	-----	--------------

TARO	0.01%	N40035	001	Oct 31, 1994
------	-------	--------	-----	--------------

	0.025%	N40042	001	Oct 31, 1994
--	--------	--------	-----	--------------

USL PHARMA	0.01%	N88757	001	Feb 11, 1985
------------	-------	--------	-----	--------------

	0.025%	N88756	001	Mar 28, 1985
--	--------	--------	-----	--------------

FLUONID

ALLERGAN HERBERT	0.025%	N87156	002	Sep 06, 1984
------------------	--------	--------	-----	--------------

FLUOTREX

SAVAGE LABS	0.01%	N88174	001	May 06, 1983
-------------	-------	--------	-----	--------------

	0.025%	N88173	001	Mar 09, 1983
--	--------	--------	-----	--------------

SYNALAR-HP

MEDICIS	0.2%	N16161	002	
---------	------	--------	-----	--

GEL; TOPICAL

FLUONID

ALLERGAN HERBERT	0.025%	N87300	001	May 27, 1982
------------------	--------	--------	-----	--------------

OINTMENT; TOPICAL

FLUOCINOLONE ACETONIDE

PHARMADERM	0.025%	N88046	001	Dec 16, 1982
------------	--------	--------	-----	--------------

PHARMAFAIR	0.025%	N88507	001	Feb 27, 1984
------------	--------	--------	-----	--------------

USL PHARMA	0.025%	N88742	001	Feb 08, 1985
------------	--------	--------	-----	--------------

FLUONID

ALLERGAN HERBERT	0.025%	N87157	001	Sep 06, 1984
------------------	--------	--------	-----	--------------

FLUOTREX

SAVAGE LABS	0.025%	N88172	001	Mar 09, 1983
-------------	--------	--------	-----	--------------

SOLUTION; TOPICAL

FLUOCINOLONE ACETONIDE

ALPHARMA US PHARMS	0.01%	N87159	001	Jun 16, 1982
--------------------	-------	--------	-----	--------------

BAUSCH AND LOMB	0.01%	N40059	001	Dec 20, 1993
-----------------	-------	--------	-----	--------------

MORTON GROVE	0.01%	N88312	001	Jan 27, 1984
--------------	-------	--------	-----	--------------

PHARMADERM	0.01%	N88048	001	Dec 16, 1982
------------	-------	--------	-----	--------------

PHARMAFAIR	0.01%	N88449	001	Feb 08, 1984
------------	-------	--------	-----	--------------

FLUONID

ALLERGAN HERBERT	0.01%	N87158	001	Mar 17, 1983
------------------	-------	--------	-----	--------------

FLUOTREX

SAVAGE LABS	0.01%	N88171	001	Mar 09, 1983
-------------	-------	--------	-----	--------------

FLUOCINOLONE ACETONIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-SYNALAR

MEDICIS	0.025%;EQ 3.5MG BASE/GM	N60700	001	
---------	-------------------------	--------	-----	--

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

CLAY PARK	0.05%	N71790	001	Jul 13, 1988
-----------	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 109 (of 263)

FLUORESCEIN SODIUM

INJECTABLE; INJECTION
FUNDUSCEIN-25
NOVARTIS

25%

N17869 001

FLUOROMETHOLONE

CREAM; TOPICAL
OXYLONE

PHARMACIA AND UPJOHN 0.025%

N11748 001

FLUOROURACIL

INJECTABLE; INJECTION
ADRUCIL

PHARMACIA AND UPJOHN 50MG/ML
50MG/ML

N17959 001

N81222 001 Jun 28, 1991

FLUOROURACIL

ABIC 50MG/ML
AM PHARM PARTNERS 50MG/ML

N88929 001

Mar 04, 1986

N89152 001

Mar 21, 1986

N89428 001

Jan 12, 1987

N89519 001

Mar 12, 1987

MARCHAR 50MG/ML
SMITH AND NEPHEW 50MG/ML

N87791 001

Jan 18, 1983

N88766 001

Dec 28, 1984

N88767 001

Dec 28, 1984

N89434 001

Mar 26, 1987

SOLUTION; TOPICAL
FLUOROPLEX

ALLERGAN HERBERT 1%

N16765 001

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL
PROZAC

LILLY EQ 60MG BASE

N18936 004

Jun 15, 1999

TABLET; ORAL
PROZAC

LILLY EQ 20MG BASE

N20974 002

Mar 09, 1999

FLUOXYMESTERONE

TABLET; ORAL
ANDROID-F

VALEANT PHARM INTL 10MG

N87196 001

FLUOXYMESTERONE

VALEANT PHARM INTL 10MG

N88221 001

May 05, 1983

WATSON LABS 2MG

N88260 001

Dec 06, 1983

5MG

N88265 001

Dec 06, 1983

10MG

N88309 001

Dec 06, 1983

ORA-TESTRYL

BRISTOL MYERS SQUIBB 2MG

N11359 001

5MG

N11359 002

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION
FLUPHENAZINE DECANOATE

MAYNE PHARMA USA 25MG/ML

N74966 001

Apr 16, 1998

FLUPHENAZINE ENANTHATE

INJECTABLE; INJECTION
PROLIXIN ENANTHATE

APOTHECON 25MG/ML

N16110 001

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL
PERMITIL

SCHERING 5MG/ML

N16008 001

DISCONTINUED DRUG PRODUCT LIST

6 - 110 (of 263)

FLUPHENAZINE HYDROCHLORIDE

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

WATSON LABS	1MG	N88555	001	Dec 18, 1987
	2.5MG	N88544	001	Dec 18, 1987
	5MG	N88527	001	Dec 18, 1987
	10MG	N88550	001	Dec 18, 1987

PERMITIL

SCHERING

0.25MG	N12034	001	
2.5MG	N12034	004	
5MG	N12034	005	
10MG	N12034	006	

TABLET, EXTENDED RELEASE; ORAL

PERMITIL

SCHERING

1MG	N12419	004	
-----	--------	-----	--

FLUPREDNISOLONE

TABLET; ORAL

ALPHADROL

PHARMACIA AND UPJOHN	1.5MG	N12259	002	
----------------------	-------	--------	-----	--

FLURANDRENOLIDE

LOTION; TOPICAL

FLURANDRENOLIDE

ALPHARMA US PHARMS	0.05%	N87203	001	Apr 29, 1982
--------------------	-------	--------	-----	--------------

FLURANDRENOLIDE; NEOMYCIN SULFATE

CREAM; TOPICAL

CORDRAN-N

LILLY	0.05%;EQ 3.5MG BASE/GM	N50346	001	
-------	------------------------	--------	-----	--

ointment; TOPICAL

CORDRAN-N

LILLY	0.05%;EQ 3.5MG BASE/GM	N50345	001	
-------	------------------------	--------	-----	--

FLURAZEPAM HYDROCHLORIDE

CAPSULE; ORAL

FLURAZEPAM HYDROCHLORIDE

HALSEY	15MG	N71808	001	Jan 07, 1988
	30MG	N71809	001	Jan 07, 1988
MUTUAL PHARM	15MG	N70454	001	Aug 04, 1986
	30MG	N70455	001	Aug 04, 1986
PUREPAC PHARM	15MG	N71927	001	Sep 09, 1987
	30MG	N71551	001	Sep 09, 1987
SUPERPHARM	15MG	N71659	001	Aug 04, 1988
	30MG	N71660	001	Aug 04, 1988
USL PHARMA	15MG	N70562	001	Jul 09, 1987
	30MG	N70563	001	Jul 09, 1987
WARNER CHILCOTT	15MG	N71767	001	Dec 04, 1987
	30MG	N71768	001	Dec 04, 1987
WATSON LABS	15MG	N72368	001	Mar 30, 1989

FLURBIPROFEN

TABLET; ORAL

FLURBIPROFEN

IVAX PHARMS	50MG	N74411	001	May 31, 1995
	100MG	N74411	002	May 31, 1995

FLUVOXAMINE MALEATE

TABLET; ORAL

LUVOX

SOLVAY	25MG	N20243	001	Dec 05, 1994
	50MG	N20243	002	Dec 05, 1994
	100MG	N20243	003	Dec 05, 1994
	150MG	N20243	004	Dec 05, 1994

DISCONTINUED DRUG PRODUCT LIST

6 - 111 (of 263)

FOLIC ACID

INJECTABLE; INJECTION
FOLVITE

WYETH PHARMS INC	5MG/ML	N05897	008
------------------	--------	--------	-----

TABLET; ORAL
FOLIC ACID

ANABOLIC	1MG	N84915	001	
BARR	1MG	N89177	001	Jan 08, 1986
EVERYLIFE	1MG	N80755	001	
HALSEY	1MG	N83598	001	
IMPAX LABS	1MG	N80686	001	
IVAX PHARMS	1MG	N83000	001	
LANNETT	1MG	N80816	001	
LILLY	1MG	N06135	003	
MK LABS	1MG	N83526	001	
PHARMERAL	1MG	N84158	001	
PIONEER PHARMS	1MG	N88949	001	Sep 13, 1985
PUREPAC PHARM	1MG	N80784	001	
SANDOZ	1MG	N84472	001	
UDL	1MG	N88199	001	Mar 29, 1983
USL PHARMA	1MG	N87828	001	May 13, 1982
VALEANT PHARM INTL	1MG	N80903	001	
VANGARD	1MG	N88730	001	Mar 23, 1984
VINTAGE PHARMS	1MG	N86296	001	
WATSON LABS	1MG	N83141	001	
	1MG	N85141	002	
WHITEWORTH TOWN PLSN	1MG	N80691	002	

FOLVITE

WYETH PHARMS INC	1MG	N05897	004
------------------	-----	--------	-----

FOLLITROPIN ALFA/BETA

INJECTABLE; IM-SC
FOLLISTIM

ORGANON USA INC	75 IU/VIAL	N20582	001	Sep 29, 1997
	150 IU/VIAL	N20582	002	Sep 29, 1997

INJECTABLE; SUBCUTANEOUS
GONAL-F

SERONO INC	37.5 IU/VIAL	N20378	003	May 25, 2000
	37.5 IU/VIAL	N21765	001	Mar 25, 2004
	75 IU/VIAL	N20378	001	Sep 29, 1997
	150 IU/VIAL	N20378	002	Sep 29, 1997
	150 IU/VIAL	N21765	003	Mar 25, 2004
	1,050 IU/VIAL	N20378	004	Feb 28, 2001

FOMIVIRSEN SODIUM

INJECTABLE; INJECTION
VITRAVENE PRESERVATIVE FREE

NOVARTIS	6.6MG/ML	N20961	001	Aug 26, 1998
----------	----------	--------	-----	--------------

FURAZOLIDONE

SUSPENSION; ORAL
FUROXONE

SHIRE	50MG/15ML	N11323	002
-------	-----------	--------	-----

TABLET; ORAL
FUROXONE

SHIRE	100MG	N11270	002
-------	-------	--------	-----

FUROSEMIDE

INJECTABLE; INJECTION
FUROSEMIDE

AM PHARM PARTNERS	10MG/ML	N18507	001	Jul 30, 1982
	10MG/ML	N19036	001	Aug 13, 1984
ASTRAZENECA	10MG/ML	N70014	001	Sep 09, 1985
	10MG/ML	N70095	001	Sep 09, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 112 (of 263)

FUROSEMIDE

INJECTABLE; INJECTION FUROSEMIDE

ASTRAZENECA	10MG/ML	N70096	001	Sep 09, 1985
BAXTER HLTHCARE	10MG/ML	N18267	001	
MARSAM PHARMS LLC	10MG/ML	N74017	001	Jun 30, 1994
ORGANON USA INC	10MG/ML	N70017	001	Dec 15, 1986
SMITH AND NEPHEW	10MG/ML	N70023	001	Feb 05, 1986
	10MG/ML	N70078	001	Feb 05, 1986
STERIS	10MG/ML	N70019	001	Sep 22, 1986
	10MG/ML	N70604	001	Jan 02, 1987
WARNER CHILCOTT	10MG/ML	N18420	001	Feb 26, 1982
WYETH AYERST	10MG/ML	N18670	001	Jul 20, 1982

LASIX

AVENTIS PHARMS	10MG/ML	N16363	001	
----------------	---------	--------	-----	--

SOLUTION; ORAL

LASIX

AVENTIS PHARMS	10MG/ML	N17688	001	
----------------	---------	--------	-----	--

TABLET; ORAL

FUROSEMIDE

CLOMEL HLTHCARE	20MG	N18415	001	Jul 27, 1982
	40MG	N18415	002	Jul 27, 1982
	80MG	N18415	003	Nov 26, 1984
KALAPHARM	20MG	N18868	001	Jun 28, 1983
	40MG	N18868	002	Jun 28, 1983
MUTUAL PHARM	20MG	N70043	001	Sep 26, 1985
	40MG	N18790	001	Nov 29, 1983
	80MG	N70100	001	Jan 26, 1988
SANDOZ	40MG	N18750	002	Jul 30, 1984
SUPERPHARM	20MG	N18370	002	Jun 26, 1984
	40MG	N18370	001	Feb 10, 1983
WARNER CHILCOTT	20MG	N18419	001	Jan 31, 1983
	40MG	N18419	002	Jan 31, 1983
	80MG	N18419	003	Nov 13, 1984
WATSON LABS	20MG	N18369	001	May 14, 1982
	40MG	N18369	002	May 14, 1982
	40MG	N70413	001	Feb 26, 1986

GALLAMINE TRIETHIODIDE

INJECTABLE; INJECTION FLAXEDIL

DAVIS AND GECK	20MG/ML	N07842	001	
	100MG/ML	N07842	002	

GALLIUM CITRATE, GA-67

INJECTABLE; INJECTION GALLIUM CITRATE GA 67

GE HEALTHCARE	1mCi/ML	N17700	001	
NEOSCAN				
GE HEALTHCARE	2mCi/ML	N17655	001	

GATIFLOXACIN

INJECTABLE; INJECTION TEQUIN

BRISTOL MYERS SQUIBB	EQ 10MG /ML(200MG)	N21062	003	Dec 17, 1999
----------------------	--------------------	--------	-----	--------------

GEMFIBROZIL

CAPSULE; ORAL GEMFIBROZIL

MYLAN	300MG	N73466	001	Jan 25, 1993
PUREPAC PHARM	300MG	N72929	001	Jan 29, 1993
LOPID				
PFIZER PHARMS	200MG	N18422	001	
	300MG	N18422	002	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 113 (of 263)

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

CLONMEL HLTHCARE

600MG

N74270 001

Sep 27, 1993

PUREPAC PHARM

600MG

N74360 001

Aug 31, 1994

WATSON LABS

600MG

N74156 001

Oct 24, 1994

GENTAMICIN SULFATE

CREAM; TOPICAL

GARAMYCIN

SCHERING

EQ 0.1% BASE

N60462 001

GENTAFAIR

PHARMAFAIR

EQ 0.1% BASE

N62458 001

Sep 01, 1983

GENTAMICIN SULFATE

ALPHARMA US PHARMS

EQ 0.1% BASE

N62471 001

Sep 27, 1983

BAUSCH AND LOMB

EQ 0.1% BASE

N64056 001

Apr 29, 1994

PHARMADERM

EQ 1MG BASE/GM

N62530 001

Jul 05, 1984

INJECTABLE; INJECTION

APOGEN

KING PHARMS

EQ 10MG BASE/ML

N62289 001

EQ 40MG BASE/ML

N62289 002

BRISTAGEN

BRISTOL

EQ 40MG BASE/ML

N62288 001

GARAMYCIN

SCHERING

EQ 1MG BASE/ML

N61716 002

EQ 10MG BASE/ML

N61739 001

EQ 40MG BASE/ML

N61716 001

GENTAFAIR

PHARMAFAIR

EQ 40MG BASE/ML

N62493 001

Aug 28, 1985

GENTAMICIN

INTL MEDICATION

EQ 1MG BASE/ML

N62325 003

Jun 23, 1982

EQ 40MG BASE/ML

N62325 001

EQ 100MG BASE/100ML

N62325 004

Jun 23, 1982

GENTAMICIN SULFATE

ABBOTT

EQ 1.2MG BASE/ML

N62413 001

Aug 11, 1983

EQ 1.4MG BASE/ML

N62413 002

Aug 11, 1983

EQ 1.6MG BASE/ML

N62413 003

Aug 11, 1983

EQ 1.8MG BASE/ML

N62413 004

Aug 11, 1983

EQ 2MG BASE/ML

N62413 005

Aug 11, 1983

EQ 60MG BASE/100ML

N62413 006

Aug 11, 1983

EQ 70MG BASE/100ML

N62413 007

Aug 11, 1983

EQ 80MG BASE/100ML

N62413 008

Aug 11, 1983

EQ 90MG BASE/100ML

N62413 009

Aug 11, 1983

EQ 100MG BASE/100ML

N62413 010

Aug 11, 1983

BAXTER HLTHCARE

EQ 10MG BASE/ML

N62251 002

EQ 40MG BASE/ML

N62251 001

KALAPHARM

EQ 40MG BASE/ML

N62354 001

Apr 05, 1982

SICOR PHARMS

EQ 10MG BASE/ML

N63149 001

Nov 21, 1991

EQ 40MG BASE/ML

N63106 002

Nov 21, 1991

SOLOPAK

EQ 10MG BASE/ML

N62507 001

Jun 06, 1985

EQ 40MG BASE/ML

N62507 002

Jun 06, 1985

STERIS

EQ 10MG BASE/ML

N62318 002

EQ 40MG BASE/ML

N62318 001

WYETH AYERST

EQ 10MG BASE/ML

N62264 001

EQ 40MG BASE/ML

N62264 002

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

HOSPIRA

EQ 1.2MG BASE/ML

N62588 001

Jan 06, 1986

EQ 1.4MG BASE/ML

N62588 002

Jan 06, 1986

EQ 1.6MG BASE/ML

N62588 003

Jan 06, 1986

EQ 1.8MG BASE/ML

N62588 004

Jan 06, 1986

EQ 2MG BASE/ML

N62588 005

Jan 06, 1986

EQ 60MG BASE/100ML

N62588 006

Jan 06, 1986

EQ 70MG BASE/100ML

N62588 007

Jan 06, 1986

EQ 80MG BASE/100ML

N62588 008

Jan 06, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 114 (of 263)

GENTAMICIN SULFATE

INJECTABLE; INJECTION

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

HOSPIRA	EQ 90MG BASE/100ML	N62588	009	Jan 06, 1986
	EQ 100MG BASE/100ML	N62588	010	Jan 06, 1986

U-GENCIN

PHARMACIA AND UPJOHN	EQ 10MG BASE/ML	N62248	001	
	EQ 40MG BASE/ML	N62248	002	

INJECTABLE; INTRATHECAL

GARAMYCIN

SCHERING	EQ 2MG BASE/ML	N50505	001	
----------	----------------	--------	-----	--

OINTMENT; OPHTHALMIC

GARAMYCIN

SCHERING	EQ 0.3% BASE	N50425	001	
----------	--------------	--------	-----	--

GENTACIDIN

NOVARTIS	EQ 0.3% BASE	N62501	001	Jul 26, 1984
----------	--------------	--------	-----	--------------

GENTAFAIR

PHARMAFAIR	EQ 3MG BASE/GM	N62443	001	May 26, 1983
------------	----------------	--------	-----	--------------

OINTMENT; TOPICAL

GARAMYCIN

SCHERING	EQ 0.1% BASE	N60463	001	
----------	--------------	--------	-----	--

GENTAFAIR

PHARMAFAIR	EQ 0.1% BASE	N62444	001	May 26, 1983
------------	--------------	--------	-----	--------------

GENTAMICIN SULFATE

ALPHARMA US PHARMS	EQ 0.1% BASE	N62496	001	Mar 14, 1984
--------------------	--------------	--------	-----	--------------

BAUSCH AND LOMB	EQ 0.1% BASE	N64054	001	Apr 29, 1994
-----------------	--------------	--------	-----	--------------

PHARMADERM	EQ 0.1% BASE	N62534	001	Oct 10, 1984
------------	--------------	--------	-----	--------------

SOLUTION/DROPS; OPHTHALMIC

GARAMYCIN

SCHERING	EQ 0.3% BASE	N50039	002	
----------	--------------	--------	-----	--

GENTAFAIR

PHARMAFAIR	EQ 0.3% BASE	N62440	001	May 03, 1983
------------	--------------	--------	-----	--------------

GENTAMICIN SULFATE

ALCON UNIVERSAL	EQ 0.3% BASE	N62523	001	Nov 25, 1985
-----------------	--------------	--------	-----	--------------

PACO	EQ 3MG BASE/ML	N62932	001	Nov 07, 1988
------	----------------	--------	-----	--------------

GENTIAN VIOLET

SUPPOSITORY; VAGINAL

GVS

SAVAGE LABS	0.4%	N83513	001	
-------------	------	--------	-----	--

TAMPON; VAGINAL

GENAPAX

KEY PHARMS	5MG	N85017	001	
------------	-----	--------	-----	--

GLIPIZIDE

TABLET; ORAL

GLUCOTROL

PFIZER	2.5MG	N17783	003	May 11, 1993
--------	-------	--------	-----	--------------

GLUCAGON HYDROCHLORIDE

INJECTABLE; INJECTION

GLUCAGON

LILLY	EQ 1MG BASE/VIAL	N12122	001	
-------	------------------	--------	-----	--

	EQ 10MG BASE/VIAL	N12122	002	
--	-------------------	--------	-----	--

GLUTAMINE

FOR SOLUTION; ORAL

NUTRESTORE

NUTRITIONAL RESTART	5GM/PACKET	N21667	001	Jun 10, 2004
---------------------	------------	--------	-----	--------------

GLUTETHIMIDE

CAPSULE; ORAL

DORIDEN

AVENTIS	500MG	N09519	008	
---------	-------	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 115 (of 263)

GLUTETHIMIDETABLET; ORAL
DORIDEN

AVENTIS	250MG	N09519	002	
	500MG	N09519	005	

GLUTETHIMIDE

HALSEY	250MG	N89458	001	Oct 10, 1986
	500MG	N89459	001	Oct 10, 1986
LANNETT	250MG	N83475	001	
	500MG	N85571	001	
SANDOZ	500MG	N83234	002	
UCB	500MG	N85171	001	
VITARINE	500MG	N87297	001	
WATSON LABS	500MG	N84362	001	
	500MG	N85763	001	

GLYBURIDETABLET; ORAL
GLYBURIDE (MICRONIZED)

AVENTIS PHARMS	1.5MG	N20055	001	Apr 17, 1992
	3MG	N20055	002	Apr 17, 1992
	6MG	N20055	003	Mar 08, 2000

GLYNASE

PHARMACIA AND UPJOHN	4.5MG	N20051	003	Sep 24, 1993
----------------------	-------	--------	-----	--------------

GLYCINE

SOLUTION; IRRIGATION

GLYCINE 1.5% IN PLASTIC CONTAINER
BAXTER HLTHCARE

1.5GM/100ML	N18522	001	Feb 19, 1982
-------------	--------	-----	--------------

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

AM PHARM PARTNERS	0.2MG/ML	N88475	001	Jun 12, 1984
HOSPIRA	0.2MG/ML	N89393	001	Jun 15, 1988
SICOR PHARMS	0.2MG/ML	N81169	001	Sep 10, 1991
STERIS	0.2MG/ML	N86947	001	Jun 24, 1983

ROBINUL

ROBINS AH	0.2MG/ML	N14764	001	
-----------	----------	--------	-----	--

TABLET; ORAL

GLYCOPYRROLATE

WATSON LABS	1MG	N85562	001	
	1MG	N86902	001	
	2MG	N85563	001	
	2MG	N86178	001	
	2MG	N86900	001	

GONADORELIN ACETATE

INJECTABLE; INJECTION

LUTREPULSE KIT

FERRING	0.8MG/VIAL	N19687	001	Oct 10, 1989
	3.2MG/VIAL	N19687	002	Oct 10, 1989

GONADORELIN HYDROCHLORIDE

INJECTABLE; INJECTION

FACTREL

BAXTER HLTHCARE CORP	EQ 0.1MG BASE/VIAL	N18123	001	Sep 30, 1982
	EQ 0.2MG BASE/VIAL	N18123	002	Sep 30, 1982
	EQ 0.5MG BASE/VIAL	N18123	003	Sep 30, 1982

GONADOTROPIN, CHORIONIC

INJECTABLE; INJECTION

A.P.L.

FERRING	5,000 UNITS/VIAL	N17055	001	
---------	------------------	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 116 (of 263)

GONADOTROPIN, CHORIONIC

INJECTABLE; INJECTION

A.P.L.

FERRING	10,000 UNITS/VIAL	N17055	002	
	20,000 UNITS/VIAL	N17055	003	

CHORIONIC GONADOTROPIN

AM PHARM PARTNERS	5,000 UNITS/VIAL	N17067	001	
	15,000 UNITS/VIAL	N17067	004	
	20,000 UNITS/VIAL	N17067	003	

BEL MAR	5,000 UNITS/VIAL	N17054	001	
	10,000 UNITS/VIAL	N17054	002	

WATSON LABS (UTAH)	2,000 UNITS/VIAL	N17016	009	Dec 27, 1984
	2,000 UNITS/VIAL	N17016	011	Feb 16, 1990
	5,000 UNITS/VIAL	N17016	006	
	15,000 UNITS/VIAL	N17016	010	Feb 15, 1985
	20,000 UNITS/VIAL	N17016	004	

FOLLUTEIN

BRISTOL MYERS SQUIBB	10,000 UNITS/VIAL	N17056	001	
----------------------	-------------------	--------	-----	--

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

IPHARM	0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML	N62818	001	Oct 11, 1988
--------	--	--------	-----	--------------

STERIS	0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML	N62788	001	Jun 11, 1987
--------	--	--------	-----	--------------

NEOMYCIN SULFATE AND POLYMYXIN B SULFATE GRAMICIDIN

PHARMAFAIR	0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML	N62383	001	Aug 31, 1982
------------	--	--------	-----	--------------

NEO-POLYCIN

DOW PHARM	0.025MG/ML;EQ 1.75MG BASE/ML;10,000 UNITS/ML	N60427	001	
-----------	--	--------	-----	--

GRANISETRON HYDROCHLORIDE

INJECTABLE; INJECTION

KYTRIL

ROCHE	EQ 3MG BASE/ML	N20239	001	Dec 29, 1993
-------	----------------	--------	-----	--------------

TABLET; ORAL

KYTRIL

ROCHE	EQ 2MG BASE	N20305	002	Jun 15, 1998
-------	-------------	--------	-----	--------------

GREPAFLOXACIN HYDROCHLORIDE

TABLET; ORAL

RAXAR

OTSUKA	EQ 200MG BASE	N20695	001	Nov 06, 1997
	EQ 400MG BASE	N20695	002	May 14, 1998
	EQ 600MG BASE	N20695	003	May 14, 1998

GRISEOFULVIN, MICROCRYSTALLINE

CAPSULE; ORAL

GRISACTIN

WYETH AYERST	125MG	N50051	002	
	250MG	N50051	001	

SUSPENSION; ORAL

GRIFULVIN V

JOHNSON AND JOHNSON	125MG/5ML	N50448	001	
---------------------	-----------	--------	-----	--

TABLET; ORAL

GRIFULVIN V

J AND J	125MG	N60618	001	
	250MG	N60618	002	
	500MG	N60618	003	

GRISACTIN

WYETH AYERST	500MG	N60212	001	
--------------	-------	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 117 (of 263)

GRISEOFULVIN, ULTRAMICROCRYSTALLINE

TABLET; ORAL				
FULVICIN P/G				
SCHERING	125MG	N61996	001	
	250MG	N61996	002	
FULVICIN P/G 165				
SCHERING	165MG	N61996	003	Apr 06, 1982
FULVICIN P/G 330				
SCHERING	330MG	N61996	004	Apr 06, 1982
GRISACTIN ULTRA				
WYETH AYERST	125MG	N62178	001	
	165MG	N62438	001	Nov 17, 1983
	250MG	N62178	002	
	330MG	N62438	002	Nov 17, 1983
ULTRAGRIS-165				
PLIVA	165MG	N62645	001	Jun 30, 1992
ULTRAGRIS-330				
PLIVA	330MG	N62646	001	Jun 30, 1992

GUANABENZ ACETATE

TABLET; ORAL				
WYTENSIN				
WYETH AYERST	EQ 4MG BASE	N18587	001	Sep 07, 1982
	EQ 8MG BASE	N18587	002	Sep 07, 1982
	EQ 16MG BASE	N18587	003	Sep 07, 1982

GUANADREL SULFATE

TABLET; ORAL				
HYLOREL				
PHARMACIA AND UPJOHN	10MG	N18104	001	Dec 29, 1982
	25MG	N18104	002	Dec 29, 1982

GUANETHIDINE MONOSULFATE

TABLET; ORAL				
GUANETHIDINE MONOSULFATE				
WATSON LABS	EQ 10MG SULFATE	N86113	001	Mar 26, 1985
	EQ 25MG SULFATE	N86114	001	Mar 26, 1985
ISMELIN				
NOVARTIS	EQ 10MG SULFATE	N12329	001	
	EQ 25MG SULFATE	N12329	002	

GUANETHIDINE MONOSULFATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL				
ESIMIL				
NOVARTIS	10MG;25MG	N13553	001	

GUANFACINE HYDROCHLORIDE

TABLET; ORAL				
TENEX				
ESP PHARMA	EQ 3MG BASE	N19032	003	Nov 07, 1988

HALAZEPAM

TABLET; ORAL				
PAXIPAM				
SCHERING	20MG	N17736	003	
	40MG	N17736	004	

HALCINONIDE

CREAM; TOPICAL				
HALOG				
WESTWOOD SQUIBB	0.025%	N17818	001	
HALOG-E				
WESTWOOD SQUIBB	0.1%	N18234	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 119 (of 263)

HALOPERIDOL

TABLET; ORAL
HALOPERIDOL

SCS	1MG	N70721	001	Jun 10, 1986
	2MG	N70722	001	Jun 10, 1986
	5MG	N70723	001	Jun 10, 1986
	10MG	N70724	001	Jun 10, 1986
	20MG	N70725	001	Sep 24, 1986
WATSON LABS	0.5MG	N71571	001	Jun 03, 1988
	1MG	N70982	001	Mar 06, 1987
	1MG	N71572	001	Jun 03, 1988
	2MG	N71573	001	Jun 03, 1988
	5MG	N71374	001	Jun 03, 1988
	10MG	N71375	001	Jun 03, 1988
	10MG	N72113	001	Aug 27, 1991
	20MG	N71376	001	Jun 03, 1988
	20MG	N72353	001	Aug 27, 1991

HALOPERIDOL DECANOATE

INJECTABLE; INJECTION
HALOPERIDOL DECANOATE
MAYNE PHARMA USA

EQ 50MG BASE/ML	N75176	001	Feb 09, 2000
EQ 100MG BASE/ML	N75176	002	Feb 09, 2000

HALOPERIDOL LACTATE

CONCENTRATE; ORAL
HALDOL

ORTHO MCNEIL	EQ 2MG BASE/ML	N15922	001	
HALOPERIDOL				
ALPHARMA	EQ 2MG BASE/ML	N70318	001	Apr 11, 1986
MORTON GROVE	EQ 2MG BASE/ML	N70710	001	Mar 07, 1986
SCS	EQ 2MG BASE/ML	N70726	001	Jun 10, 1986
TEVA	EQ 2MG BASE/ML	N71015	001	Aug 25, 1987

HALOPERIDOL INTENSOL
ROXANE

EQ 2MG BASE/ML	N72045	001	Apr 12, 1988
----------------	--------	-----	--------------

INJECTABLE; INJECTION
HALOPERIDOL

AM PHARM PARTNERS	EQ 5MG BASE/ML	N71187	001	Jan 20, 1987
MARSAM PHARMS LLC	EQ 5MG BASE/ML	N72516	001	Feb 25, 1993
	EQ 5MG BASE/ML	N72517	001	Feb 25, 1993
SMITH AND NEPHEW	EQ 5MG BASE/ML	N70802	001	Dec 14, 1987
SOLOPAK	EQ 5MG BASE/ML	N70800	001	Dec 14, 1987
	EQ 5MG BASE/ML	N70801	001	Dec 14, 1987
	EQ 5MG BASE/ML	N70864	001	Dec 14, 1987
STERIS	EQ 5MG BASE/ML	N70713	001	May 17, 1988
	EQ 5MG BASE/ML	N70744	001	May 17, 1988

HALOPROGIN

CREAM; TOPICAL
HALOTEX

WESTWOOD SQUIBB	1%	N16942	001
-----------------	----	--------	-----

SOLUTION; TOPICAL
HALOTEX

WESTWOOD SQUIBB	1%	N16943	001
-----------------	----	--------	-----

HALOTHANE

LIQUID; INHALATION
FLUOTHANE

WYETH AYERST	99.99%	N11338	001
--------------	--------	--------	-----

HALOTHANE

BH	99.99%	N84977	001
----	--------	--------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 120 (of 263)

HEPARIN CALCIUM

INJECTABLE; INJECTION
CALCIPARINE

SANOFI SYNTHELABO	25,000 UNITS/ML	N18237	001
-------------------	-----------------	--------	-----

HEPARIN SODIUM

INJECTABLE; INJECTION

HEP FLUSH KIT IN PLASTIC CONTAINER

AM PHARM	10 UNITS/ML	N17029	017	Dec 05, 1985
	100 UNITS/ML	N17029	018	Dec 05, 1985

HEPARIN LOCK FLUSH

AM PHARM	100 UNITS/ML	N17651	010	
BAXTER HLTHCARE CORP	10 UNITS/ML	N17007	008	
	100 UNITS/ML	N17007	009	
HOSPIRA	100 UNITS/ML	N05264	010	
INTL MEDICATION	10 UNITS/ML	N86357	001	
	500 UNITS/ML	N86357	002	
LUITPOLD	10 UNITS/ML	N89063	001	Oct 09, 1985
	100 UNITS/ML	N89064	001	Oct 09, 1985
PARKE DAVIS	10 UNITS/ML	N17346	006	
SMITH AND NEPHEW	10 UNITS/ML	N87904	001	Apr 20, 1983
	10 UNITS/ML	N87958	001	Apr 20, 1983
	10 UNITS/ML	N88458	001	Jul 26, 1984
	10 UNITS/ML	N88580	001	Oct 25, 1984
	100 UNITS/ML	N87906	001	Apr 20, 1983
	100 UNITS/ML	N87959	001	Apr 20, 1983
	100 UNITS/ML	N88460	001	Jul 26, 1984
	100 UNITS/ML	N88581	001	Oct 25, 1984
SOLOPAK	10 UNITS/ML	N87903	001	Apr 20, 1983
	10 UNITS/ML	N88457	001	Oct 25, 1984
	100 UNITS/ML	N87905	001	Apr 20, 1983
	100 UNITS/ML	N88459	001	Jul 26, 1984
WATSON PHARMS	100 UNITS/ML	N17064	001	

HEPARIN LOCK FLUSH PRESERVATIVE FREE

AM PHARM	10 UNITS/ML	N17029	011	Sep 22, 1987
	100 UNITS/ML	N17029	012	Sep 22, 1987

HEPARIN LOCK FLUSH PRESERVATIVE FREE IN PLASTIC CONTAINER

AM PHARM	10 UNITS/ML	N17029	008	Sep 22, 1987
	100 UNITS/ML	N17029	009	Sep 22, 1987

HEPARIN SODIUM

AKORN	1,000 UNITS/ML	N17486	001	
	5,000 UNITS/ML	N17486	002	
	10,000 UNITS/ML	N17486	003	
	20,000 UNITS/ML	N17486	004	
	40,000 UNITS/ML	N17486	005	
AM PHARM	1,000 UNITS/ML	N17651	005	
	5,000 UNITS/ML	N17029	002	
	5,000 UNITS/ML	N17979	003	
	10,000 UNITS/ML	N17651	003	
	20,000 UNITS/ML	N17651	008	
AM PHARM PARTNERS	1,000 UNITS/ML	N17033	001	
BAXTER HLTHCARE CORP	1,000 UNITS/ML	N17007	001	
	2,500 UNITS/ML	N17007	007	
	5,000 UNITS/ML	N17007	002	
	5,000 UNITS/0.5ML	N17007	010	
	7,500 UNITS/ML	N17007	003	
	10,000 UNITS/ML	N17007	004	
	15,000 UNITS/ML	N17007	005	
	20,000 UNITS/ML	N17007	006	
CHAMBERLIN PARENTERL	1,000 UNITS/ML	N17130	001	
	5,000 UNITS/ML	N17130	002	
	10,000 UNITS/ML	N17130	003	
	20,000 UNITS/ML	N17130	004	
DELL LABS	1,000 UNITS/ML	N17540	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 121 (of 263)

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM

DELL LABS	5,000 UNITS/ML	N17540	002	
	10,000 UNITS/ML	N17540	003	
	20,000 UNITS/ML	N17540	004	
	40,000 UNITS/ML	N17540	005	
HOSPIRA	2,500 UNITS/ML	N88099	001	Apr 28, 1983
	10,000 UNITS/ML	N40095	001	Jul 26, 1996
LILLY	1,000 UNITS/ML	N05521	001	
	10,000 UNITS/ML	N05521	002	
	20,000 UNITS/ML	N05521	004	
LUITPOLD	1,000 UNITS/ML	N87452	001	Oct 31, 1983
MARSAM PHARMS LLC	1,000 UNITS/ML	N40007	001	Jun 07, 1996
ORGANON USA INC	1,000 UNITS/ML	N00552	008	
	5,000 UNITS/ML	N00552	009	
	10,000 UNITS/ML	N00552	010	
PARKE DAVIS	1,000 UNITS/ML	N17346	001	
	5,000 UNITS/ML	N17346	002	
	7,500 UNITS/ML	N17346	003	
	10,000 UNITS/ML	N17346	004	
	20,000 UNITS/ML	N17346	005	
PHARM SPEC	1,000 UNITS/ML	N17780	001	
	5,000 UNITS/ML	N17780	002	
	10,000 UNITS/ML	N17780	003	
	20,000 UNITS/ML	N17780	004	
	40,000 UNITS/ML	N17780	005	
SMITH AND NEPHEW	1,000 UNITS/ML	N88239	001	Jul 26, 1984
SOLOPAK	1,000 UNITS/ML	N87043	001	
	5,000 UNITS/ML	N87077	001	
	5,000 UNITS/0.5ML	N87395	001	
	10,000 UNITS/ML	N87107	001	
	10,000 UNITS/0.5ML	N87363	001	
WATSON PHARMS	1,000 UNITS/ML	N17064	002	
	2,500 UNITS/ML	N17064	015	
	3,000 UNITS/ML	N17064	016	
	4,000 UNITS/ML	N17064	017	
	5,000 UNITS/ML	N17064	003	
	6,000 UNITS/ML	N17064	018	
	7,500 UNITS/ML	N17064	019	
	10,000 UNITS/ML	N17064	004	
	20,000 UNITS/ML	N17064	005	
	40,000 UNITS/ML	N17064	006	
HEPARIN SODIUM 1,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
MCGAW	200 UNITS/100ML	N19130	001	Dec 31, 1984
HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	200 UNITS/100ML	N19042	001	Mar 29, 1985
HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5%				
HOSPIRA	10,000 UNITS/100ML	N18911	006	Jan 30, 1985
HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	2,000 UNITS/100ML	N18814	002	Jul 09, 1985
HEPARIN SODIUM 10,000 UNITS IN SODIUM CHLORIDE 0.45%				
HOSPIRA	10,000 UNITS/100ML	N18911	001	Jan 30, 1985
	10,000 UNITS/100ML	N18916	005	Jan 31, 1984
HEPARIN SODIUM 10,000 UNITS IN SODIUM CHLORIDE 0.9%				
HOSPIRA	10,000 UNITS/100ML	N18911	003	Jan 30, 1985
	10,000 UNITS/100ML	N18916	002	Jan 31, 1984
HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5%				
HOSPIRA	5,000 UNITS/100ML	N18911	007	Jan 30, 1985
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N19802	001	Jul 20, 1992
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.9%				
HOSPIRA	5,000 UNITS/100ML	N18911	005	Jan 30, 1985
	5,000 UNITS/100ML	N18916	003	Jan 31, 1984

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 122 (of 263)

HEPARIN SODIUM

INJECTABLE; INJECTION

HEPARIN SODIUM 2,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
MCGAW	200 UNITS/100ML	N19130	003	Dec 31, 1984
HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	200 UNITS/100ML	N19042	002	Mar 29, 1985
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5%				
HOSPIRA	5,000 UNITS/100ML	N18911	009	Jan 30, 1985
	10,000 UNITS/100ML	N18911	008	Jan 30, 1985
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N19134	001	Mar 29, 1985
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N19802	005	Jul 20, 1992
	10,000 UNITS/100ML	N19802	002	Jul 20, 1992
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.9%				
HOSPIRA	5,000 UNITS/100ML	N18911	004	Jan 30, 1985
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	5,000 UNITS/100ML	N19135	001	Mar 29, 1985
	5,000 UNITS/100ML	N19802	003	Jul 20, 1992
HOSPIRA	5,000 UNITS/100ML	N18916	009	Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	500 UNITS/100ML	N18609	003	Apr 28, 1982
HEPARIN SODIUM 5,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER				
MCGAW	1,000 UNITS/100ML	N19130	002	Dec 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.45%				
HOSPIRA	100 UNITS/ML	N18911	002	Jan 30, 1985
	100 UNITS/ML	N18916	004	Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.9%				
HOSPIRA	1,000 UNITS/100ML	N18916	001	Jan 31, 1984
HEPARIN SODIUM 5,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	1,000 UNITS/100ML	N19042	004	Mar 29, 1985
HEPARIN SODIUM PRESERVATIVE FREE				
MARSAM PHARMS LLC	1,000 UNITS/ML	N89464	001	Jun 03, 1986
PHARMA SERVE NY	1,000 UNITS/ML	N86129	001	
LIPO-HEPIN				
3M	1,000 UNITS/0.5ML	N17027	001	
	1,000 UNITS/ML	N17027	006	
	5,000 UNITS/0.5ML	N17027	002	
	5,000 UNITS/ML	N17027	008	
	7,500 UNITS/0.5ML	N17027	010	
	10,000 UNITS/0.5ML	N17027	003	
	10,000 UNITS/ML	N17027	009	
	15,000 UNITS/0.5ML	N17027	011	
	20,000 UNITS/0.5ML	N17027	004	
	20,000 UNITS/ML	N17027	007	
	40,000 UNITS/ML	N17027	005	
LIQUAEMIN LOCK FLUSH				
ORGANON USA INC	100 UNITS/ML	N00552	007	
LIQUAEMIN SODIUM				
ORGANON USA INC	1,000 UNITS/ML	N00552	004	
	5,000 UNITS/ML	N00552	003	
	10,000 UNITS/ML	N00552	005	
	20,000 UNITS/ML	N00552	001	
	40,000 UNITS/ML	N00552	002	
LIQUAEMIN SODIUM PRESERVATIVE FREE				
ORGANON USA INC	1,000 UNITS/ML	N00552	011	Apr 11, 1986
	5,000 UNITS/ML	N00552	012	Apr 11, 1986
	10,000 UNITS/ML	N00552	013	Apr 11, 1986
PANHEPRIN				
HOSPIRA	1,000 UNITS/ML	N05264	004	
	5,000 UNITS/ML	N05264	006	
	10,000 UNITS/ML	N05264	007	
	20,000 UNITS/ML	N05264	008	
	40,000 UNITS/ML	N05264	009	

DISCONTINUED DRUG PRODUCT LIST

6 - 123 (of 263)

HEPARIN SODIUM

INJECTABLE; INJECTION
SODIUM HEPARIN

AM PHARM PARTNERS	5,000 UNITS/ML	N17033	002	
	10,000 UNITS/ML	N17033	003	
	20,000 UNITS/ML	N17033	004	
BAXTER HLTHCARE	1,000 UNITS/ML	N17036	001	Mar 04, 1988

HETACILLIN

FOR SUSPENSION; ORAL
VERSAPEN

BRISTOL	EQ 112.5MG AMPICIL/5ML	N50060	001	
	EQ 112.5MG AMPICIL/ML	N50060	003	
	EQ 112.5MG AMPICIL/ML	N61398	001	
	EQ 225MG AMPICIL/5ML	N61398	002	

HETACILLIN POTASSIUM

CAPSULE; ORAL
VERSAPEN-K

BRISTOL	EQ 225MG AMPICIL	N61396	001	
	EQ 450MG AMPICIL	N61396	002	

HEXACHLOROPHENE

AEROSOL; TOPICAL
SEPTISOL

VESTAL LABS	0.23%	N17424	001	
TURGEX				
XTTRIUM	3%	N18375	001	

EMULSION; TOPICAL
HEXA-GERM

HUNTINGTON LABS	3%	N17411	001	
PHISOHEX				
SANOFI SYNTHELABO	3%	N08402	001	
SOY-DOME				
BAYER PHARMS	3%	N17405	001	

TURGEX
XTTRIUM

	3%	N19055	001	Nov 30, 1984
--	----	--------	-----	--------------

SOAP; TOPICAL
GAMOPHEN

ARBROOK	2%	N06270	003	
---------	----	--------	-----	--

SOLUTION; TOPICAL
DIAL

DIAL	0.25%	N17421	002	
GERMA-MEDICA				
HUNTINGTON LABS	1%	N17412	001	
GERMA-MEDICA "MG"				
HUNTINGTON LABS	0.25%	N17412	002	
SEPTI-SOFT				
CALGON	0.25%	N17460	001	
SEPTISOL				
VESTAL LABS	0.25%	N17423	001	

SPONGE; TOPICAL
E-Z SCRUB

BECTON DICKINSON	450MG	N17452	001	
HEXASCRUB				
PROF DSPLS	3%	N18363	001	
PHISO-SCRUB				
SANOFI SYNTHELABO	3%	N17446	001	
SCRUBTEAM SURGICAL SPONGEBRUSH				
3M	330MG	N17413	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 124 (of 263)

HEXAFLUORENIUM BROMIDEINJECTABLE; INJECTION
MYLAXEN

MEDPOINTE PHARM HLC 20MG/ML

N09789 003

HEXOCYCLIUM METHYLSULFATETABLET; ORAL
TRAL

ABBOTT 25MG

N10599 001

HEXYLCAINE HYDROCHLORIDESOLUTION; TOPICAL
CYCLAIN

MERCK 5%

N08472 001

HISTAMINE PHOSPHATEINJECTABLE; INJECTION
HISTAMINE PHOSPHATE
LILLY

EQ 0.1MG BASE/ML

N00734 003

EQ 0.2MG BASE/ML

N00734 002

EQ 1MG BASE/ML

N00734 001

HISTRELIN ACETATEINJECTABLE; INJECTION
SUPPRELIN

SHIRE EQ 0.2MG BASE/ML

N19836 001 Dec 24, 1991

EQ 0.5MG BASE/ML

N19836 002 Dec 24, 1991

EQ 1MG BASE/ML

N19836 003 Dec 24, 1991

HOMATROPINE METHYLBROMIDETABLET; ORAL
HOMAPIN-10

MISSION PHARMA 10MG

N86308 001

HOMAPIN-5

MISSION PHARMA 5MG

N86309 001

TABLET, CHEWABLE; ORAL
EQUIPIN

MISSION PHARMA 3MG

N86310 001

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATESYRUP; ORAL
HYDRODANE

HALSEY 1.5MG/5ML; 5MG/5ML

N88066 001 Jun 28, 1985

HYALURONIDASEINJECTABLE; INJECTION
WYDASE

BAXTER HLTHCARE 150 UNITS/ML

N06343 002

150 UNITS/VIAL **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N06343 006

1,500 UNITS/VIAL **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N06343 005

HYDRALAZINE HYDROCHLORIDEINJECTABLE; INJECTION
APRESOLINE

NOVARTIS 20MG/ML

N08303 003

HYDRALAZINE HYDROCHLORIDE

AM PHARM PARTNERS 20MG/ML

N89532 001 Aug 11, 1987

SMITH AND NEPHEW 20MG/ML

N88518 001 Apr 20, 1984

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 125 (of 263)

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE

SOLOPAK 20MG/ML

N88517 001 Aug 22, 1985

TABLET; ORAL

DRALZINE

TEVA 25MG

N84301 001

HYDRALAZINE HYDROCHLORIDE

ABC HOLDING 10MG

N88846 001 Feb 26, 1985

25MG

N88847 001 Feb 26, 1985

50MG

N88848 001 Feb 26, 1985

100MG

N88849 001 Feb 26, 1985

AMIDE PHARM 25MG

N88560 001 Oct 04, 1984

50MG

N88649 001 Oct 18, 1984

ASCOT 25MG

N88310 001 Dec 19, 1984

50MG

N88311 001 Dec 19, 1984

CLONMEL HLTHCARE 25MG

N86243 001

50MG

N86242 002

HALSEY 10MG

N89218 001 Jan 22, 1986

25MG

N89130 001 Jan 15, 1986

50MG

N89222 001 Jan 22, 1986

100MG

N89178 001 Jan 15, 1986

IMPAX LABS 25MG

N84922 001

50MG

N84923 001

IVAX PHARMS 10MG

N84443 001

25MG

N84437 001

50MG

N84469 002

100MG

N84581 001

MUTUAL PHARM 10MG

N88728 001 Apr 11, 1985

25MG

N84106 002

50MG

N84107 002

100MG

N88729 001 Apr 11, 1985

PUREPAC PHARM 25MG

N88177 001 Jul 29, 1983

50MG

N88178 001 Aug 15, 1983

QUANTUM PHARMICS 10MG

N88671 001 May 01, 1984

25MG

N88657 001 Jun 15, 1984

50MG

N88652 001 May 08, 1984

100MG

N88686 001 May 01, 1984

SANDOZ 50MG

N85088 001

SUPERPHARM 10MG

N88787 001 Aug 28, 1984

25MG

N88788 001 Aug 28, 1984

50MG

N88789 001 Aug 28, 1984

USL PHARMA 25MG

N87780 001 Mar 29, 1982

50MG

N87751 001 Mar 29, 1982

VANGARD 25MG

N87712 001

50MG

N87908 001 May 07, 1982

VITARINE 25MG

N86088 001

WATSON LABS 25MG

N85532 002 May 24, 1982

50MG

N85533 002 May 25, 1982

WEST WARD 25MG

N88240 001 May 27, 1983

50MG

N88241 001 May 27, 1983

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

APRESAZIDE

NOVARTIS 100MG;50MG

N84811 001

HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

SOLVAY 25MG;25MG

N87608 001 Feb 08, 1982

50MG;50MG

N87213 001 Feb 08, 1982

100MG;50MG

N87609 001 Feb 08, 1982

SUPERPHARM 25MG;25MG

N89200 001 Feb 09, 1987

50MG;50MG

N89201 001 Feb 09, 1987

WATSON LABS 25MG;25MG

N85457 001 Mar 04, 1982

50MG;50MG

N85446 001 Mar 04, 1982

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 126 (of 263)

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

WATSON LABS 100MG;50MG

N85440 001 Mar 04, 1982

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 100/50

IVAX PHARMS 100MG;50MG

N88358 001 Apr 10, 1984

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 25/25

IVAX PHARMS 25MG;25MG

N88356 001 Apr 10, 1984

HYDRALAZINE HYDROCHLORIDE W/ HYDROCHLOROTHIAZIDE 50/50

IVAX PHARMS 50MG;50MG

N88357 001 Apr 10, 1984

TABLET; ORAL

APRESOLINE-ESIDRIX

NOVARTIS 25MG;15MG

N12026 002

HYDRALAZINE AND HYDROCHLOROTHIAZIDE

WATSON LABS 25MG;15MG

N85827 001

HYDROCHLOROTHIAZIDE W/ HYDRALAZINE

WATSON LABS 25MG;15MG

N85373 001

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

ABC HOLDING 25MG;15MG;0.1MG

N84897 001

HYDRALAZINE HYDROCHLORIDE, HYDROCHLOROTHIAZIDE AND RESERPINE

IVAX PHARMS 25MG;15MG;0.1MG

N84291 001

HYDRALAZINE HYDROCHLORIDE-HYDROCHLOROTHIAZIDE-RESERPINE

MYLAN 25MG;15MG;0.1MG

N87085 001

HYDRALAZINE, HYDROCHLOROTHIAZIDE W/ RESERPINE

WATSON LABS 25MG;15MG;0.1MG

N85771 001

HYDRAP-ES

SANDOZ 25MG;15MG;0.1MG

N84876 001

HYDROCHLOROTHIAZIDE W/ RESERPINE AND HYDRALAZINE

WATSON LABS 25MG;15MG;0.1MG

N83770 001

HYDROSERPINE PLUS (R-H-H)

IVAX PHARMS 25MG;15MG;0.1MG

N83877 001

RESERPINE, HYDRALAZINE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

MUTUAL PHARM 25MG;15MG;0.1MG

N88570 001 Apr 10, 1984

SOLVAY 25MG;15MG;0.1MG

N88376 001 Oct 28, 1983

WATSON LABS 25MG;15MG;0.1MG

N85549 001

25MG;15MG;0.1MG

N87556 001

RESERPINE, HYDROCHLOROTHIAZIDE, AND HYDRALAZINE HYDROCHLORIDE

LEDERLE 25MG;15MG;0.1MG

N87709 001 May 13, 1982

SER-A-GEN

SOLVAY 25MG;15MG;0.1MG

N87210 001

SER-AP-ES

NOVARTIS 25MG;15MG;0.1MG

N12193 005

UNIPRES

SOLVAY 25MG;15MG;0.1MG

N85893 001

25MG;15MG;0.1MG

N86298 001

HYDRALAZINE HYDROCHLORIDE; RESERPINE

TABLET; ORAL

DRALSERP

SANDOZ 25MG;0.1MG

N84617 001

SERPASIL-APRESOLINE

NOVARTIS 25MG;0.1MG

N09296 004

50MG;0.2MG

N09296 002

HYDROCHLOROTHIAZIDE

SOLUTION; ORAL

HYDROCHLOROTHIAZIDE

MORTON GROVE 50MG/5ML

N89661 001 Jun 20, 1988

HYDROCHLOROTHIAZIDE INTENSOL

ROXANE 100MG/ML

N88588 001 Jul 02, 1984

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 127 (of 263)

HYDROCHLOROTHIAZIDE

TABLET; ORAL

ESIDRIX

NOVARTIS

100MG

N11793 009

HYDROCHLOROTHIAZIDE

ABC HOLDING

25MG

N85683 001

50MG

N83965 001

50MG

N85672 001

ALRA

25MG

N86369 001

50MG

N83554 001

ASCOT

25MG

N87539 001

Feb 03, 1982

50MG

N87540 001

Feb 03, 1982

BARR

50MG

N84771 001

CLONMEL HLTHCARE

100MG

N87060 001

ELKINS SINN

50MG

N85152 002

HEATHER

50MG

N84135 001

IMPAX LABS

25MG

N84029 001

50MG

N83607 002

100MG

N85098 001

INWOOD LABS

25MG

N84776 001

25MG

N85067 001

50MG

N84776 002

IVAX PHARMS

50MG

N84658 001

100MG

N85022 001

MAST MM

25MG

N86192 001

50MG

N86192 002

MUTUAL PHARM

25MG

N83972 001

50MG

N83972 002

100MG

N83972 003

MYLAN

25MG

N84880 001

50MG

N85112 001

PHARMERAL

25MG

N84325 001

50MG

N84324 001

PVT FORM

50MG

N86597 001

ROXANE

25MG

N85004 001

50MG

N84536 002

50MG

N85005 001

SANDOZ

25MG

N83899 001

SOLVAY

25MG

N85323 001

SUPERPHARM

25MG

N88827 001

Dec 28, 1984

50MG

N88828 001

Dec 28, 1984

100MG

N88829 001

Dec 28, 1984

TEVA

25MG

N88924 001

Feb 07, 1985

50MG

N88923 001

Feb 07, 1985

USL PHARMA

25MG

N87827 001

Apr 19, 1982

50MG

N87752 001

Apr 19, 1982

VANGARD

25MG

N87638 001

50MG

N87610 001

WARNER CHILCOTT

25MG

N87586 001

May 03, 1982

50MG

N87587 001

May 03, 1982

WATSON LABS

25MG

N83458 001

25MG

N85232 002

50MG

N83232 001

50MG

N83456 001

50MG

N85233 001

50MG

N86087 001

50MG

N86594 001

100MG

N81190 001

Jan 24, 1992

100MG

N85099 001

100MG

N87002 001

WEST WARD

25MG

N84899 001

50MG

N84878 001

WHITEWORTH TOWN PLSN

25MG

N83809 002

50MG

N83809 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 128 (of 263)

HYDROCHLOROTHIAZIDE

TABLET; ORAL			
HYDROCHLOROTHIAZIDE			
WHITEWORTH TOWN PLSN	100MG	N85347	001
HYDRO-D			
HALSEY	25MG	N86504	001
	50MG	N83891	002
HYDRODIURIL			
MERCK	25MG	N11835	003
	50MG	N11835	006
	100MG	N11835	007
ORETIC			
ABBOTT	25MG	N11971	001
ZIDE			
SOLVAY	50MG	N83925	001

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL			
AVALIDE			
SANOFI SYNTHELABO	12.5MG;75MG	N20758	001 Sep 30, 1997

HYDROCHLOROTHIAZIDE; LABETALOL HYDROCHLORIDE

TABLET; ORAL			
NORMOZIDE			
SCHERING	25MG;100MG	N19046	001 Apr 06, 1987
	25MG;200MG	N19046	002 Apr 06, 1987
	25MG;300MG	N19046	003 Apr 06, 1987
	25MG;400MG	N19046	004 Apr 06, 1987
TRANDATE HCT			
GLAXOSMITHKLINE	25MG;100MG	N19174	001 Apr 10, 1987
	25MG;200MG	N19174	002 Apr 10, 1987
	25MG;300MG	N19174	003 Apr 10, 1987
	25MG;400MG	N19174	004 Apr 10, 1987

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL			
METHYLDOPA AND HYDROCHLOROTHIAZIDE			
CLONMEL HLTHCARE	15MG;250MG	N72507	001 Jun 02, 1989
	25MG;250MG	N72508	001 Jun 02, 1989
	30MG;500MG	N72509	001 Jun 02, 1989
	50MG;500MG	N72510	001 Jun 02, 1989
IVAX PHARMS	15MG;250MG	N71458	001 Mar 08, 1988
	25MG;250MG	N71459	001 Mar 08, 1988
	30MG;500MG	N71460	001 Mar 08, 1988
	50MG;500MG	N71461	001 Mar 08, 1988
PARKE DAVIS	15MG;250MG	N71897	001 Nov 23, 1987
	25MG;250MG	N71898	001 Nov 23, 1987
	30MG;500MG	N71899	001 Nov 23, 1987
	50MG;500MG	N71900	001 Nov 23, 1987
PUREPAC PHARM	15MG;250MG	N70853	001 Oct 08, 1986
	25MG;250MG	N70688	001 Apr 24, 1986
	30MG;500MG	N70854	001 Oct 08, 1986
	50MG;500MG	N70689	001 Apr 24, 1986
SANDOZ	15MG;250MG	N70829	001 Mar 09, 1987
	25MG;250MG	N70830	001 Mar 09, 1987
TEVA	15MG;250MG	N71819	001 Apr 08, 1988
	25MG;250MG	N71820	001 Apr 08, 1988
	30MG;500MG	N71821	001 Apr 08, 1988
	50MG;500MG	N71822	001 Apr 08, 1988
WATSON LABS	15MG;250MG	N70365	001 Mar 19, 1986
	15MG;250MG	N71920	001 Aug 29, 1988
	25MG;250MG	N70366	001 Apr 16, 1986
	25MG;250MG	N71921	001 Aug 29, 1988
	30MG;500MG	N70367	001 Mar 19, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 129 (of 263)

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

METHYLDOPA AND HYDROCHLOROTHIAZIDE

WATSON LABS

30MG;500MG

N71922 001

Aug 29, 1988

50MG;500MG

N70368 001

Apr 16, 1986

50MG;500MG

N71923 001

Aug 29, 1988

HYDROCHLOROTHIAZIDE; PINDOLOL

TABLET; ORAL

VISKAZIDE

NOVARTIS

25MG;5MG

N18872 001

Jul 22, 1987

25MG;10MG

N18872 002

Jul 22, 1987

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

INDERIDE LA 120/50

WYETH AYERST

50MG;120MG

N19059 002

Jul 03, 1985

INDERIDE LA 160/50

WYETH AYERST

50MG;160MG

N19059 003

Jul 03, 1985

INDERIDE LA 80/50

WYETH AYERST

50MG;80MG

N19059 001

Jul 03, 1985

TABLET; ORAL

INDERIDE-80/25

WYETH PHARMS INC

25MG;80MG

N18031 002

PROPRANOLOL HYDROCHLORIDE & HYDROCHLOROTHIAZIDE

DURAMED PHARMS BARR

25MG;40MG

N71126 001

Mar 02, 1987

25MG;80MG

N71127 001

Mar 02, 1987

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

IVAX PHARMS

25MG;40MG

N71552 001

Dec 01, 1988

25MG;80MG

N71553 001

Dec 01, 1988

WARNER CHILCOTT

25MG;40MG

N71771 001

Jan 26, 1988

25MG;80MG

N71772 001

Jan 26, 1988

WATSON LABS

25MG;40MG

N71498 001

Dec 18, 1991

25MG;80MG

N71501 001

Dec 18, 1991

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

H.R.-50

WHITEWORTH TOWN PLSN

50MG;0.125MG

N85338 001

HYDROCHLOROTHIAZIDE W/ RESERPINE

IVAX PHARMS

25MG;0.1MG

N83572 001

25MG;0.125MG

N83571 001

50MG;0.1MG

N83568 001

50MG;0.125MG

N83573 001

PHARMERAL

25MG;0.125MG

N85421 001

50MG;0.125MG

N85420 001

ROXANE

50MG;0.125MG

N84603 001

WATSON LABS

25MG;0.125MG

N84466 001

25MG;0.125MG

N85317 001

25MG;0.125MG

N86330 002

50MG;0.125MG

N83666 001

50MG;0.125MG

N84467 001

50MG;0.125MG

N86331 001

HYDROPRES 25

MERCK

25MG;0.125MG

N11958 002

HYDROPRES 50

MERCK

50MG;0.125MG

N11958 003

HYDRO-RESERP

ABC HOLDING

50MG;0.125MG

N84714 002

Jun 29, 1982

HYDRO-SERP "25"

SANDOZ

25MG;0.125MG

N84827 001

HYDRO-SERP "50"

SANDOZ

50MG;0.125MG

N85213 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 130 (of 263)

HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

RESERPINE AND HYDROCHLOROTHIAZIDE

BARR	25MG;0.125MG	N84580	001	
	50MG;0.125MG	N84579	001	
SANDOZ	50MG;0.125MG	N88200	001	Jan 31, 1984
RESERPINE AND HYDROCHLOROTHIAZIDE-50				
WEST WARD	50MG;0.125MG	N88189	001	May 10, 1984
SERPASIL-ESIDRIX #1				
NOVARTIS	25MG;0.1MG	N11878	003	
SERPASIL-ESIDRIX #2				
NOVARTIS	50MG;0.1MG	N11878	005	

HYDROCHLOROTHIAZIDE; SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE

ASCOT	25MG;25MG	N88025	001	Nov 23, 1984
PUREPAC PHARM	25MG;25MG	N87999	001	Nov 06, 1985
SUPERPHARM	25MG;25MG	N89137	001	Aug 26, 1985

SPIRONOLACTONE W/ HYDROCHLOROTHIAZIDE

IVAX PHARMS	25MG;25MG	N87004	002	May 24, 1982
LEDERLE	25MG;25MG	N87511	001	
PARKE DAVIS	25MG;25MG	N87948	001	Feb 22, 1983
PUREPAC PHARM	25MG;25MG	N88054	001	Aug 18, 1983
UPSHER SMITH	25MG;25MG	N87553	001	
USL PHARMA	25MG;25MG	N87651	001	
VANGARD	25MG;25MG	N87655	001	
WATSON LABS	25MG;25MG	N85974	001	
	25MG;25MG	N86026	001	

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

DYAZIDE

GLAXOSMITHKLINE	25MG;50MG	N16042	002	
-----------------	-----------	--------	-----	--

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

NOVARTIS	25MG;37.5MG	N74857	001	Sep 09, 1997
VITARINE	25MG;50MG	N71737	001	Feb 12, 1988

TABLET; ORAL

TRIAMTERENE AND HYDROCHLOROTHIAZIDE

AM THERAP	50MG;75MG	N72022	001	Apr 17, 1988
QUANTUM PHARMICS	50MG;75MG	N71980	001	Apr 17, 1988

HYDROCODONE BITARTRATE; PHENYLPROPANOLAMINE HYDROCHLORIDE

SYRUP; ORAL

CODAMINE

ALPHARMA US PHARMS	5MG/5ML;25MG/5ML	N75103	001	Sep 29, 2000
--------------------	------------------	--------	-----	--------------

HYCOMINE

ENDO PHARMS	5MG/5ML;25MG/5ML	N19410	001	Aug 17, 1990
-------------	------------------	--------	-----	--------------

HYCOMINE PEDIATRIC

ENDO PHARMS	2.5MG/5ML;12.5MG/5ML	N19411	001	Aug 17, 1990
-------------	----------------------	--------	-----	--------------

HYDROCORTAMATE HYDROCHLORIDE

OINTMENT; TOPICAL

MAGNACORT

PFIZER	0.5%	N10554	001	
--------	------	--------	-----	--

HYDROCORTISONE

AEROSOL; TOPICAL

AEROSEB-HC

ALLERGAN HERBERT	0.5%	N85805	001	
------------------	------	--------	-----	--

CREAM; TOPICAL

CORT-DOME

BAYER PHARMS	0.5%	N09585	003	
	1%	N09585	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 131 (of 263)

HYDROCORTISONE

CREAM; TOPICAL				
DERMACORT				
MONARCH PHARMS	1%	N83011	002	
ELDECORT				
VALEANT PHARM INTL	1%	N80459	001	
	2.5%	N84055	001	
FLEXICORT				
WESTWOOD SQUIBB	0.5%	N87136	003	Apr 08, 1982
	1%	N87136	002	Apr 08, 1982
	2.5%	N87136	001	Apr 08, 1982
HC #1				
BAYER PHARMS	0.5%	N80438	001	
HC #4				
BAYER PHARMS	1%	N80438	002	
HC (HYDROCORTISONE)				
C AND M PHARMA	0.5%	N80482	003	
	1%	N80482	004	
H-CORT				
PHARM ASSOC	0.5%	N86823	001	
HYDROCORTISONE				
ALPHARMA US PHARMS	2.5%	N89754	001	Feb 01, 1989
ALTANA	0.5%	N80848	002	
	1%	N80848	003	
AMBIX	1%	N86080	001	
	2.5%	N86271	001	
CLAY PARK	0.5%	N84970	002	
	1%	N85026	001	
EVERYLIFE	0.5%	N80452	001	
G AND W LABS	1%	N84059	001	
IVAX PHARMS	1%	N85733	001	
NASKA	1%	N89706	001	Mar 10, 1988
PHARMADERM	1%	N88845	001	Feb 27, 1986
	2.5%	N89413	001	Dec 16, 1986
PHARMAFAIR	1%	N87838	001	Jul 28, 1982
STIEFEL	1%	N86170	001	
SYOSSET	0.5%	N85527	001	
TARO	0.5%	N86154	001	
TEVA	0.5%	N80400	002	
	1%	N80400	003	
	1%	N85191	001	
	2.5%	N80400	004	
TOPIDERM	1%	N89273	001	Feb 17, 1989
USL PHARMA	1%	N88027	001	Sep 27, 1983
	2.5%	N88029	001	Sep 27, 1983
WHITEWORTH TOWN PLSN	1%	N80496	002	
NOGENIC HC				
IVAX PHARMS	1%	N87427	001	Apr 04, 1988
NUTRACORT				
HEALTHPOINT	0.5%	N80442	002	
	1%	N80442	003	
PROCTOCORT				
MONARCH PHARMS	1%	N83011	001	
SYNACORT				
MEDICIS	0.5%	N87459	001	
GEL; TOPICAL				
NUTRACORT				
HEALTHPOINT	1%	N84698	001	
PENECORT				
ALLERGAN HERBERT	1%	N88215	001	Jun 06, 1984
INJECTABLE; INJECTION				
CORTEF				
PHARMACIA AND UPJOHN	50MG/ML	N09864	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 132 (of 263)

HYDROCORTISONE

LOTION; TOPICAL

ACTICORT

BAKER NORTON

1%

N86535 001

BALNEOL-HC

SOLVAY

1%

N88041 001 Dec 03, 1982

BETA-HC

BETA DERMAC

1%

N89495 001 Jan 25, 1988

CORT-DOME

BAYER PHARMS

0.5%

N09895 003

1%

N09895 001

DERMACORT

SOLVAY

0.5%

N84573 002

1%

N86462 001

EPICORT

BLULINE

0.5%

N83219 002

GLYCORT

HERAN

1%

N87489 001 Oct 03, 1983

H-CORT

PHARM ASSOC

0.5%

N86824 001

HYDROCORTISONE

ALPHARMA US PHARMS

0.5%

N87317 001 Jun 07, 1982

1%

N87315 001 Jun 07, 1982

CLAY PARK

0.5%

N85662 001

1%

N85663 001

MERICON

0.5%

N85282 001

1%

N85282 002 Feb 26, 1987

NASKA

1%

N89705 001 Apr 25, 1988

TARO

1%

N89024 001 Feb 12, 1986

NUTRACORT

HEALTHPOINT

0.5%

N80443 002

OINTMENT; TOPICAL

CORTRIL

PFIZER GLOBAL

1%

N09176 001

2.5%

N09176 002

HC (HYDROCORTISONE)

C AND M PHARMA

0.5%

N80481 001

1%

N80481 002

HYDROCORTISONE

ALTANA

0.5%

N80489 002

1%

N80489 003

AMBIX

1%

N86079 001

2.5%

N86272 001

CLAY PARK

0.5%

N84969 003

1%

N85028 001

NASKA

1%

N89704 001 Mar 10, 1988

PHARMADERM

1%

N88842 001 Feb 09, 1987

TARO

0.5%

N86256 001

USL PHARMA

1%

N88061 001 Sep 27, 1983

2.5%

N88039 001 Sep 27, 1983

HYTONE

DERMIK LABS

1%

N80474 003

2.5%

N80474 004

PENECORT

ALLERGAN HERBERT

2.5%

N88217 001 Jun 06, 1984

POWDER; FOR RX COMPOUNDING

H-CORT

TORCH

100%

N87834 001 Mar 29, 1982

HYDROCORTISONE

PADDOCK

100%

N88082 001 Apr 08, 1983

TABLET; ORAL

CORTRIL

PFIZER

10MG

N09127 005

20MG

N09127 003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 133 (of 263)

HYDROCORTISONE

TABLET; ORAL

HYDROCORTISONE

ANABOLIC	20MG	N83140	001	
BARR	20MG	N83999	001	
ELKINS SINN	20MG	N80624	001	
FERRANTE	10MG	N80568	001	
	20MG	N80568	002	
IMPAX LABS	20MG	N80781	001	
INWOOD LABS	20MG	N80732	001	
LANNETT	20MG	N85070	001	
PANRAY	10MG	N09659	001	
	20MG	N09659	002	
PARKE DAVIS	20MG	N84243	001	
PUREPAC PHARM	10MG	N84247	003	Aug 31, 1982
	20MG	N80395	001	
	20MG	N84247	002	
ROXANE	10MG	N88539	001	Mar 21, 1984
SANDOZ	20MG	N80642	002	
WATSON LABS	20MG	N80355	001	
WHITEWORTH TOWN PLSN	10MG	N80344	001	
	20MG	N80344	002	
TABLET; VAGINAL				
CORTRIL				
PFIPHARMECS	10MG	N09796	001	

HYDROCORTISONE ACETATE

CREAM; TOPICAL

HEMSOL-HC

ABLE	1%	N81274	001	Jun 19, 1992
HYDROCORTISONE ACETATE				
CENCI	1%	N80419	001	Jan 25, 1982
PARKE DAVIS	1%	N89914	001	Jan 03, 1989
PUREPAC PHARM	0.5%	N86050	001	
	1%	N86052	001	

INJECTABLE; INJECTION

CORTEF ACETATE

PHARMACIA AND UPJOHN	50MG/ML	N09378	002	
----------------------	---------	--------	-----	--

CORTRIL

PFIZER	25MG/ML	N09164	001	
--------	---------	--------	-----	--

HYDROCORTISONE ACETATE

AKORN	25MG/ML	N09637	001	
	50MG/ML	N09637	002	
BEL MAR	25MG/ML	N83739	001	
	50MG/ML	N83739	002	
STERIS	25MG/ML	N83128	001	
	25MG/ML	N83759	001	
	50MG/ML	N83759	002	
	50MG/ML	N85214	001	

HYDROCORTONE

MERCK	25MG/ML	N08228	001	
	50MG/ML	N08228	004	

OINTMENT; OPHTHALMIC

HYDROCORTISONE ACETATE

ALTANA	0.5%	N80828	001	
--------	------	--------	-----	--

OINTMENT; OPHTHALMIC, OTIC

HYDROCORTONE

MERCK	1.5%	N09018	003	
-------	------	--------	-----	--

OINTMENT; TOPICAL

CORTEF ACETATE

PHARMACIA AND UPJOHN	1%	N08917	002	
----------------------	----	--------	-----	--

2.5% **Federal Register determination
that product was not discontinued or
withdrawn for safety or efficacy
reasons**

N08917 001

DISCONTINUED DRUG PRODUCT LIST

6 - 134 (of 263)

HYDROCORTISONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL			
NEO-CORTEF			
PHARMACIA AND UPJOHN	1%;EQ 3.5MG BASE/GM	N61049	001
	2.5%;EQ 3.5MG BASE/GM	N61049	002
OINTMENT; OPHTHALMIC			
NEO-CORTEF			
PHARMACIA AND UPJOHN	0.5%;EQ 3.5MG BASE/GM	N60610	001
	1.5%;EQ 3.5MG BASE/GM	N60610	002
OINTMENT; TOPICAL			
NEO-CORTEF			
PHARMACIA AND UPJOHN	0.5%;EQ 3.5MG BASE/GM	N60751	001
	1%;EQ 3.5MG BASE/GM	N60751	002
	2.5%;EQ 3.5MG BASE/GM	N60751	003
SUSPENSION/DROPS; OPHTHALMIC			
COR-OTICIN			
AKORN			
	1.5%;EQ 3.5MG BASE/ML	N60188	001
NEO-CORTEF			
PHARMACIA AND UPJOHN	0.5%;EQ 3.5MG BASE/ML	N60612	002
	1.5%;EQ 3.5MG BASE/ML	N60612	001

HYDROCORTISONE ACETATE; PRAMOXINE HYDROCHLORIDE

LOTION; TOPICAL			
PRAMOSONE			
FERNDAL LABS	0.5%;1%	N83213	002

HYDROCORTISONE BUTYRATE

CREAM; TOPICAL			
LOCOID			
YAMANOUCHI	0.1%	N18795	001 Jan 07, 1983
OINTMENT; TOPICAL			
LOCOID			
YAMANOUCHI	0.1%	N19106	001 Jul 03, 1984
SOLUTION; TOPICAL			
LOCOID			
YAMANOUCHI	0.1%	N19819	001 Sep 15, 1988

HYDROCORTISONE CYPIONATE

SUSPENSION; ORAL			
CORTEF			
PHARMACIA AND UPJOHN	EQ 10MG BASE/5ML	N09900	001

HYDROCORTISONE SODIUM PHOSPHATE

INJECTABLE; INJECTION			
HYDROCORTONE			
MERCK	EQ 50MG BASE/ML	N12052	001

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION			
A-HYDROCORT			
ABBOTT			
	EQ 100MG BASE/VIAL	N85928	001
	EQ 100MG BASE/VIAL	N89577	001 Apr 11, 1989
	EQ 250MG BASE/VIAL	N89578	001 Apr 11, 1989
	EQ 500MG BASE/VIAL	N89579	001 Apr 11, 1989
	EQ 1GM BASE/VIAL	N89580	001 Apr 11, 1989
HYDROCORTISONE SODIUM SUCCINATE			
AM PHARM PARTNERS			
	EQ 100MG BASE/VIAL	N88667	001 Jun 08, 1984
	EQ 100MG BASE/VIAL	N88712	001 Jun 08, 1984
	EQ 250MG BASE/VIAL	N88668	001 Jun 08, 1984
	EQ 500MG BASE/VIAL	N88669	001 Jun 08, 1984
	EQ 1GM BASE/VIAL	N88670	001 Jun 08, 1984
BAXTER HLTHCARE			
	EQ 100MG BASE/VIAL	N86619	001
	EQ 250MG BASE/VIAL	N87567	001
	EQ 500MG BASE/VIAL	N87568	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 135 (of 263)

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

HYDROCORTISONE SODIUM SUCCINATE

BAXTER HLTHCARE	EQ 1GM BASE/VIAL	N87569	001	
INTL MEDICATION	EQ 100MG BASE/VIAL	N87532	001	Mar 19, 1982
STERIS	EQ 100MG BASE/VIAL	N84737	002	
	EQ 100MG BASE/VIAL	N84738	001	
	EQ 250MG BASE/VIAL	N84737	001	
	EQ 500MG BASE/VIAL	N84747	001	
	EQ 1GM BASE/VIAL	N84748	001	

HYDROCORTISONE; NEOMYCIN SULFATE

CREAM; TOPICAL

NEO-CORT-DOME

BAYER PHARMS	0.5%;EQ 3.5MG BASE/GM	N50237	006	Jun 05, 1984
	1%;EQ 3.5MG BASE/GM	N50237	005	Jun 05, 1984

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE

PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N62394	001	Sep 29, 1982
------------	-------------------------------------	--------	-----	--------------

OTOCORT

STERIS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N60730	002	
--------	-------------------------------------	--------	-----	--

SUSPENSION/DROPS; OPHTHALMIC

NEOMYCIN SULFATE-POLYMYXIN B SULFATE-HYDROCORTISONE

PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N62623	001	Sep 24, 1985
------------	-------------------------------------	--------	-----	--------------

SUSPENSION/DROPS; OTIC

NEOMYCIN SULFATE, POLYMYXIN B SULFATE & HYDROCORTISONE

PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N62617	001	Sep 18, 1985
------------	-------------------------------------	--------	-----	--------------

OTICAIR

PHARMAFAIR	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N62399	001	Nov 18, 1982
------------	-------------------------------------	--------	-----	--------------

OTOBIONE

SCHERING	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N61816	001	
----------	-------------------------------------	--------	-----	--

OTOCORT

STERIS	1%;EQ 3.5MG BASE/ML;10,000 UNITS/ML	N62521	001	Jul 11, 1985
--------	-------------------------------------	--------	-----	--------------

HYDROCORTISONE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

OTOBIOITIC

SCHERING	5MG/ML;EQ 10,000 UNITS BASE/ML	N62302	001	
----------	--------------------------------	--------	-----	--

PYOCIDIN

FOREST LABS	5MG/ML;EQ 10,000 UNITS BASE/ML	N61606	001	
-------------	--------------------------------	--------	-----	--

HYDROCORTISONE; TETRACYCLINE HYDROCHLORIDE

OINTMENT; OPHTHALMIC

ACHROMYCIN

LEDERLE	1.5%;1%	N50272	001	
---------	---------	--------	-----	--

HYDROCORTISONE; UREA

CREAM; TOPICAL

ALPHADERM

BIOGLAN	1%;10%	N86008	001	
---------	--------	--------	-----	--

CALMURID HC

PHARMACIA AND UPJOHN	1%;10%	N83947	001	
----------------------	--------	--------	-----	--

HYDROFLUMETHIAZIDE

TABLET; ORAL

DIUCARDIN

WYETH AYERST	50MG	N83383	001	
--------------	------	--------	-----	--

HYDROFLUMETHIAZIDE

WATSON LABS	50MG	N88031	001	Apr 06, 1983
	50MG	N88528	001	Aug 15, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 136 (of 263)

HYDROFLUMETHIAZIDE; RESERPINE

TABLET; ORAL

HYDROFLUMETHIAZIDE AND RESERPINE

USL PHARMA	50MG;0.125MG	N88195	001	Oct 26, 1983
WATSON LABS	25MG;0.125MG	N88127	001	Mar 22, 1983
	50MG;0.125MG	N88110	001	Mar 22, 1983

RESERPINE AND HYDROFLUMETHIAZIDE

IVAX PHARMS	50MG;0.125MG	N88932	001	Jan 11, 1985
PAR PHARM	50MG;0.125MG	N88907	001	Sep 20, 1985
SALUTENSIN				
SHIRE	50MG;0.125MG	N12359	003	
SALUTENSIN-DEMI				
SHIRE	25MG;0.125MG	N12359	004	

HYDROMORPHONE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

PALLADONE

PURDUE PHARMA LP	12MG	N21044	001	Sep 24, 2004
	16MG	N21044	002	Sep 24, 2004
	24MG	N21044	003	Sep 24, 2004
	32MG	N21044	004	Sep 24, 2004

HYDROXOCOBALAMIN

INJECTABLE; INJECTION

ALPHAREDISOL

MERCK	1MG/ML	N80778	001	
-------	--------	--------	-----	--

HYDROXOCOBALAMIN

AM PHARM PARTNERS	1MG/ML	N84921	001	
STERIS	1MG/ML	N85528	001	

HYDROXOMIN

BEL MAR	1MG/ML	N84629	001	
---------	--------	--------	-----	--

HYDROXYAMPHETAMINE HYDROBROMIDE

SOLUTION/DROPS; OPHTHALMIC

PAREDINE

AKORN	1%	N00004	004	
-------	----	--------	-----	--

HYDROXYPROGESTERONE CAPROATE

INJECTABLE; INJECTION

DELALUTIN

BRISTOL MYERS SQUIBB	125MG/ML	N10347	004	
	125MG/ML	N16911	001	
	250MG/ML	N10347	002	
	250MG/ML	N16911	002	

HYDROXYPROGESTERONE CAPROATE

AKORN	125MG/ML	N18004	001	
WATSON LABS	125MG/ML	N17439	001	
	250MG/ML	N17439	002	

HYDROXYSTILBAMIDINE ISETHIONATE

INJECTABLE; INJECTION

HYDROXYSTILBAMIDINE ISETHIONATE

AVENTIS PHARMS	225MG/AMP	N09166	001	
----------------	-----------	--------	-----	--

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE

BAXTER HLTHCARE	50MG/ML	N85551	002	
-----------------	---------	--------	-----	--

HYDROXYZINE HYDROCHLORIDE

ALTANA	25MG/ML	N87273	001	Apr 20, 1982
	50MG/ML	N87273	002	Apr 20, 1982
AM PHARM PARTNERS	25MG/ML	N88184	001	Mar 31, 1983
	50MG/ML	N88185	001	Mar 31, 1983

DISCONTINUED DRUG PRODUCT LIST

6 - 137 (of 263)

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

BAXTER HLTHCARE	25MG/ML	N85551	001	
HOSPIRA	50MG/ML	N86821	001	
PHARMAFAIR	25MG/ML	N88862	001	Feb 14, 1986
	25MG/ML	N89106	001	Feb 14, 1986
	50MG/ML	N88881	001	Feb 14, 1986
	50MG/ML	N89107	001	Feb 14, 1986
SMITH AND NEPHEW	25MG/ML	N87592	001	
SOLOPAK	25MG/ML	N86822	001	
	25MG/ML	N87591	001	
	50MG/ML	N87310	001	
	50MG/ML	N87593	001	
	50MG/ML	N87595	001	
	50MG/ML	N87596	001	
STERIS	25MG/ML	N85778	001	
	25MG/ML	N87274	001	
	50MG/ML	N87274	002	
WYETH AYERST	25MG/ML	N86258	001	
	50MG/ML	N86258	002	
ORGATRX				
ORGANON USA INC	25MG/ML	N87014	001	
	50MG/ML	N87014	002	

SYRUP; ORAL

HYDROXYZINE HYDROCHLORIDE

ALPHARMA US PHARMS	10MG/5ML	N88785	001	Feb 03, 1988
KV PHARM	10MG/5ML	N87730	001	Jul 01, 1982

TABLET; ORAL

ATARAX

PFIZER	10MG	N10392	001	
	25MG	N10392	004	
	50MG	N10392	006	
	100MG	N10392	005	

HYDROXYZINE HYDROCHLORIDE

ABLE	10MG	N40559	001	Jul 22, 2004
	25MG	N40562	001	Jul 22, 2004
	50MG	N40563	001	Jul 22, 2004
AMIDE PHARM	10MG	N89071	001	Jul 22, 1986
	25MG	N89072	001	Jul 22, 1986
	50MG	N89073	001	Jul 22, 1986
HALSEY	10MG	N89366	001	May 02, 1988
	25MG	N89117	001	May 02, 1988
	50MG	N89396	001	May 02, 1988
IVAX PHARMS	10MG	N87216	001	
	25MG	N87410	001	
	50MG	N87411	001	
KV PHARM	10MG	N87819	001	Jun 23, 1982
	25MG	N87820	001	Jun 23, 1982
	50MG	N87821	001	Jun 23, 1982
	100MG	N87822	001	Jun 23, 1982
MUTUAL PHARM	10MG	N88409	001	Nov 15, 1983
	25MG	N87857	001	Apr 18, 1983
	50MG	N87860	001	Apr 18, 1983
	100MG	N87862	001	Apr 18, 1983
PAR PHARM	10MG	N87602	001	Jan 22, 1982
	25MG	N87603	001	Jan 22, 1982
	50MG	N87604	001	Jan 22, 1982
PLIVA	100MG	N81054	001	Sep 25, 1995
PUREPAC PHARM	10MG	N88120	001	Sep 25, 1984
	25MG	N88121	001	Sep 25, 1984
	50MG	N88122	001	Sep 25, 1984
QUANTUM PHARMICS	10MG	N88540	001	Oct 22, 1985
	25MG	N88551	001	Oct 22, 1985

DISCONTINUED DRUG PRODUCT LIST

6 - 138 (of 263)

HYDROXYZINE HYDROCHLORIDE

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

QUANTUM PHARMICS	50MG	N88529	001	Oct 22, 1985
SANDOZ	10MG	N87246	002	
	25MG	N85247	001	
	50MG	N87245	001	
SUPERPHARM	10MG	N88794	001	Dec 05, 1984
	25MG	N88795	001	Dec 05, 1984
	50MG	N88796	001	Dec 05, 1984
USL PHARMA	10MG	N89121	001	Mar 20, 1986
	25MG	N89122	001	Mar 20, 1986
	50MG	N89123	001	Mar 20, 1986
WATSON LABS	10MG	N86827	001	
	25MG	N86829	001	
	50MG	N86836	001	

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

DURAMED PHARMS BARR	EQ 25MG HCL	N88593	001	Feb 29, 1984
	EQ 50MG HCL	N88594	001	Feb 29, 1984
	EQ 100MG HCL	N88595	001	Feb 29, 1984
PAR PHARM	EQ 25MG HCL	N87656	001	Jun 11, 1982
	EQ 25MG HCL	N89145	001	Mar 17, 1986
	EQ 50MG HCL	N87657	001	Jun 11, 1982
	EQ 50MG HCL	N89146	001	Mar 17, 1986
	EQ 100MG HCL	N87658	001	Jun 11, 1982
SANDOZ	EQ 50MG HCL	N81128	001	Jun 28, 1991
	EQ 100MG HCL	N81129	001	Jun 28, 1991
SUPERPHARM	EQ 25MG HCL	N89031	001	Jan 02, 1987
	EQ 50MG HCL	N89032	001	Jan 02, 1987
	EQ 100MG HCL	N89033	001	Jan 02, 1987
VANGARD	EQ 25MG HCL	N88392	001	Sep 19, 1983
	EQ 50MG HCL	N88393	001	Sep 19, 1983
WATSON LABS	EQ 25MG HCL	N86698	001	
	EQ 25MG HCL	N86840	001	Jul 01, 1982
	EQ 50MG HCL	N86695	001	
	EQ 50MG HCL	N86705	001	Jul 01, 1982
	EQ 50MG HCL	N87767	001	Aug 16, 1982
	EQ 100MG HCL	N86697	001	
	EQ 100MG HCL	N86728	001	Oct 05, 1982
	EQ 100MG HCL	N87790	001	Aug 16, 1982
HY-PAM "25"				
TEVA	EQ 25MG HCL	N88713	001	Mar 04, 1985

IBUPROFEN

CAPSULE; ORAL

MIDOL

BAYER	200MG **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N70626	001	Sep 02, 1987
	200MG **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N71002	001	Sep 02, 1987

SUSPENSION; ORAL

IBU

ABBOTT	100MG/5ML	N19784	001	Dec 18, 1989
SUSPENSION/DROPS; ORAL				
MOTRIN				
MCNEIL	40MG/ML	N20476	001	May 25, 1995

DISCONTINUED DRUG PRODUCT LIST

6 - 139 (of 263)

IBUPROFEN

TABLET; ORAL
ACHES-N-PAIN

LEDERLE	200MG	N71065	001	May 28, 1987
IBU				
BASF	600MG	N70088	001	Feb 08, 1985
IBUPRIN				
PLIVA	200MG	N71773	001	Jul 16, 1987
IBUPROFEN				
ABBOTT	600MG	N70556	001	Jun 14, 1985
	800MG	N71264	001	Jul 25, 1986
HALSEY	200MG	N71027	001	Sep 29, 1987
	300MG	N71028	001	Mar 23, 1987
	400MG	N71029	001	Mar 23, 1987
	600MG	N71030	001	Mar 23, 1987
	800MG	N72137	001	Feb 05, 1988
IVAX PHARMS	200MG	N71154	001	Oct 27, 1987
	200MG	N72040	001	Apr 29, 1988
	200MG	N72901	001	Dec 19, 1991
	200MG	N72903	001	Dec 19, 1991
	400MG	N71145	001	Sep 23, 1986
	600MG	N71146	001	Sep 23, 1986
	800MG	N71769	001	May 08, 1987
LEDERLE	400MG	N70629	001	Sep 19, 1986
	600MG	N70630	001	Sep 19, 1986
MCNEIL	400MG	N70081	001	Jun 16, 1986
	600MG	N70476	001	Jun 16, 1986
MUTUAL PHARM	200MG	N70493	001	Dec 24, 1985
	200MG	N70908	001	Sep 26, 1986
	200MG	N71462	001	Oct 02, 1986
	400MG	N70079	001	Jul 24, 1985
	600MG	N70080	001	Jul 24, 1985
	800MG	N71448	001	Feb 18, 1987
PAR PHARM	300MG	N70328	001	Aug 06, 1985
PUREPAC PHARM	200MG	N71122	001	Oct 03, 1986
	200MG	N71664	001	Feb 03, 1987
	300MG	N71123	001	Sep 19, 1986
	400MG	N71124	001	Sep 19, 1986
	600MG	N71125	001	Sep 19, 1986
	800MG	N71964	001	Feb 01, 1988
SANDOZ	400MG	N72064	001	Jan 14, 1988
	600MG	N72065	001	Jan 14, 1988
	800MG	N71938	001	Jan 14, 1988
SUPERPHARM	600MG	N70709	001	Apr 25, 1986
TEVA	200MG	N73141	001	May 29, 1992
	400MG	N73343	001	Jun 30, 1992
	600MG	N73344	001	Jun 30, 1992
	800MG	N73345	001	Jun 30, 1992
WATSON LABS	200MG	N71765	001	Sep 04, 1987
	200MG	N71905	001	Mar 08, 1988
	300MG	N71338	001	Dec 01, 1986
	400MG	N70038	001	Sep 06, 1985
	600MG	N70041	001	Sep 06, 1985
	800MG	N71911	001	Oct 13, 1987
IBU-TAB				
ALRA	800MG	N71965	001	Aug 11, 1988
MIDOL				
BAYER	200MG	N70591	001	Sep 02, 1987
	200MG	N71001	001	Sep 02, 1987
MOTRIN				
MCNEIL CONS SPECLT	100MG	N20418	001	Nov 16, 1994
NUPRIN				
BRISTOL MYERS	200MG	N72035	001	Feb 16, 1988
	200MG	N72036	001	Feb 16, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 140 (of 263)

IBUPROFEN

TABLET; ORAL

NUPRIN

MCNEIL

200MG

N19012 001

May 18, 1984

200MG

N19012 002

Jul 29, 1987

RUFEN

BASF

600MG

N18197 002

Mar 05, 1984

TABLET, CHEWABLE; ORAL

MOTRIN

MCNEIL CONS SPECLT

50MG

N20135 001

Nov 16, 1994

100MG

N20135 002

Nov 16, 1994

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDAMYCIN

PHARMACIA AND UPJOHN

5MG/VIAL

N50661 002

Sep 27, 1990

10MG/VIAL

N50661 001

Sep 27, 1990

20MG/VIAL

N50661 003

Apr 25, 1995

IDOXURIDINE

OINTMENT; OPHTHALMIC

STOXIL

GLAXOSMITHKLINE

0.5%

N15868 001

SOLUTION/DROPS; OPHTHALMIC

STOXIL

GLAXOSMITHKLINE

0.1%

N13934 001

IFOSFAMIDE

INJECTABLE; INJECTION

IFEX

BRISTOL MYERS SQUIBB

1GM/VIAL

N19763 001

Dec 30, 1988

3GM/VIAL

N19763 002

Dec 30, 1988

IMATINIB MESYLATE

CAPSULE; ORAL

GLEEVEC

NOVARTIS

50MG

N21335 001

May 10, 2001

100MG

N21335 002

May 10, 2001

IMIPRAMINE HYDROCHLORIDE

CONCENTRATE; ORAL

IMIPRAMINE HYDROCHLORIDE

NOVARTIS

25MG/ML

N86765 001

INJECTABLE; INJECTION

TOFRANIL

NOVARTIS

12.5MG/ML

N11838 002

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

LEDERLE

10MG

N86269 001

25MG

N86267 001

50MG

N86268 001

ROXANE

10MG

N83799 001

25MG

N83799 002

50MG

N83799 003

SANDOZ

10MG

N85200 001

25MG

N84869 002

50MG

N85133 001

TEVA

10MG

N83729 001

25MG

N83729 004

50MG

N83729 003

USL PHARMA

25MG

N87776 001

Feb 10, 1982

VANGARD

10MG

N88036 001

Nov 03, 1982

25MG

N87619 001

Feb 09, 1982

50MG

N87631 001

Jan 04, 1982

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 141 (of 263)

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

WATSON LABS

10MG

N85220 001

10MG

N85875 001

25MG

N84252 002

25MG

N85878 001

50MG

N85221 001

50MG

N85877 001

WEST WARD

25MG

N88222 001

May 26, 1983

50MG

N88223 001

May 26, 1983

JANIMINE

ABBOTT

10MG

N17895 001

25MG

N17895 002

50MG

N17895 003

PRAMINE

ALRA

10MG

N83827 001

25MG

N83827 002

50MG

N83827 003

PRESAMINE

AVENTIS

10MG

N11836 006

25MG

N11836 003

50MG

N11836 007

INAMRINONE LACTATE

INJECTABLE; INJECTION

INOCOR

SANOFI SYNTHELABO

EQ 5MG BASE/ML

N18700 001

Jul 31, 1984

INDAPAMIDE

TABLET; ORAL

INDAPAMIDE

TEVA

1.25MG

N74665 001

Apr 04, 1997

2.5MG

N74665 002

Apr 04, 1997

INDECAINIDE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

DECABID

LILLY

EQ 50MG BASE

N19693 001

Dec 29, 1989

EQ 75MG BASE

N19693 002

Dec 29, 1989

EQ 100MG BASE

N19693 003

Dec 29, 1989

INDOCYANINE GREEN

INJECTABLE; INJECTION

IC-GREEN

AKORN

10MG/VIAL

N11525 003

40MG/VIAL

N11525 004

50MG/VIAL

N11525 002

INDOMETHACIN

CAPSULE; ORAL

INDOMETHACIN

ABLE

25MG

N76666 001

Dec 17, 2003

50MG

N76666 002

Dec 17, 2003

DURAMED PHARMS BARR

25MG

N70326 001

Oct 18, 1985

50MG

N70327 001

Oct 18, 1985

HALSEY

25MG

N70782 001

Jun 03, 1987

50MG

N70635 001

Jun 03, 1987

IVAX PHARMS

25MG

N18730 001

May 04, 1984

50MG

N18730 002

May 04, 1984

MUTUAL PHARM

25MG

N70067 001

Oct 03, 1986

50MG

N70068 001

Oct 03, 1986

PARKE DAVIS

25MG

N18806 001

Nov 23, 1984

50MG

N18806 002

Nov 23, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 142 (of 263)

INDOMETHACIN

CAPSULE; ORAL
INDOMETHACIN

PIONEER PHARMS	25MG	N70813	001	Aug 11, 1986
	50MG	N70592	001	Aug 11, 1986
ROXANE	25MG	N70353	001	Jun 18, 1985
	50MG	N70354	001	Jun 18, 1985
SUPERPHARM	25MG	N70487	001	Oct 10, 1986
	50MG	N70488	001	Oct 10, 1986
WATSON LABS	25MG	N18690	001	Jul 31, 1984
	25MG	N70529	001	Oct 18, 1985
	25MG	N70784	001	Aug 20, 1986
	25MG	N72996	001	Jul 31, 1991
	50MG	N18690	002	Jul 31, 1984
	50MG	N70530	001	Oct 18, 1985
	50MG	N70785	001	Aug 20, 1986
	50MG	N71635	001	May 18, 1987
	50MG	N72997	001	Jul 31, 1991

CAPSULE, EXTENDED RELEASE; ORAL

INDOCIN SR

MERCK	75MG	N18185	001	Feb 23, 1982
-------	------	--------	-----	--------------

INDOMETHACIN

ABLE	75MG	N76114	001	Feb 06, 2002
------	------	--------	-----	--------------

SUPPOSITORY; RECTAL

INDOCIN

MERCK	50MG	N17814	001	Aug 13, 1984
-------	------	--------	-----	--------------

SUSPENSION; ORAL

INDOMETHACIN

ROXANE	25MG/5ML	N71412	001	Mar 18, 1987
--------	----------	--------	-----	--------------

INSULIN GLULISINE RECOMBINANT

INJECTABLE; SUBCUTANEOUS

APIDRA

AVENTIS PHARMS	1000 UNITS/10ML (100 UNITS/ML)	N21629	001	Apr 16, 2004
	300 UNITS/3ML (100 UNITS/ML)	N21629	002	Dec 20, 2005

INSULIN PORK

INJECTABLE; INJECTION

Iletin I

LILLY	500 UNITS/ML	N17931	001	
-------	--------------	--------	-----	--

INSULIN

NOVO NORDISK INC	40 UNITS/ML	N17926	001	
------------------	-------------	--------	-----	--

REGULAR INSULIN

NOVO NORDISK INC	100 UNITS/ML	N17926	003	
------------------	--------------	--------	-----	--

INSULIN PURIFIED BEEF

INJECTABLE; INJECTION

REGULAR Iletin II

LILLY	100 UNITS/ML	N18478	001	
-------	--------------	--------	-----	--

INSULIN PURIFIED PORK

INJECTABLE; INJECTION

REGULAR PURIFIED PORK INSULIN

NOVO NORDISK INC	100 UNITS/ML	N18381	001	
------------------	--------------	--------	-----	--

VELOSULIN

NOVO NORDISK INC	100 UNITS/ML	N18193	001	
------------------	--------------	--------	-----	--

INSULIN PURIFIED PORK; INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

INSULIN NORDISK MIXTARD (PORK)

NOVO NORDISK INC	30 UNITS/ML; 70 UNITS/ML	N18195	001	
------------------	--------------------------	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 143 (of 263)

INSULIN RECOMBINANT HUMAN

INJECTABLE; INJECTION
HUMULIN BR

LILLY	100 UNITS/ML	N19529 001	Apr 28, 1986
-------	--------------	------------	--------------

INSULIN RECOMBINANT PURIFIED HUMAN

INJECTABLE; INJECTION
NOVOLIN R

NOVO NORDISK INC	100 UNITS/ML	N18778 001	Aug 30, 1983
------------------	--------------	------------	--------------

VELOSULIN BR HUMAN			
NOVO NORDISK INC	100 UNITS/ML	N19450 001	May 30, 1986

INSULIN RECOMBINANT PURIFIED HUMAN; INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION
MIXTARD HUMAN 70/30

BAYER PHARMS	30 UNITS/ML;70 UNITS/ML	N19585 001	Mar 11, 1988
--------------	-------------------------	------------	--------------

NOVOLIN 70/30			
NOVO NORDISK INC	30 UNITS/ML;70 UNITS/ML	N19441 001	Jul 11, 1986

INSULIN SUSP ISOPHANE BEEF

INJECTABLE; INJECTION
NPH INSULIN

NOVO NORDISK INC	40 UNITS/ML	N17929 001	
	100 UNITS/ML	N17929 003	

INSULIN SUSP ISOPHANE BEEF/PORK

INJECTABLE; INJECTION
NPH ILETIN I (BEEF-PORK)

LILLY	40 UNITS/ML	N17936 001	
-------	-------------	------------	--

INSULIN SUSP ISOPHANE PURIFIED BEEF

INJECTABLE; INJECTION
NPH ILETIN II

LILLY	100 UNITS/ML	N18479 001	
-------	--------------	------------	--

INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

INSULIN INSULATARD NPH NORDISK			
NOVO NORDISK INC	100 UNITS/ML	N18194 001	

NPH PURIFIED PORK ISOPHANE INSULIN			
NOVO NORDISK INC	100 UNITS/ML	N18623 001	

INSULIN SUSP ISOPHANE SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION
INSULATARD NPH HUMAN

NOVO NORDISK INC	100 UNITS/ML	N19449 001	May 30, 1986
------------------	--------------	------------	--------------

NOVOLIN N			
NOVO NORDISK INC	100 UNITS/ML	N19065 001	Jan 23, 1985

INSULIN SUSP PROTAMINE ZINC BEEF/PORK

INJECTABLE; INJECTION

PROTAMINE ZINC & ILETIN I (BEEF-PORK)			
LILLY	40 UNITS/ML	N17932 001	
	100 UNITS/ML	N17932 002	

INSULIN SUSP PROTAMINE ZINC PURIFIED BEEF

INJECTABLE; INJECTION

PROTAMINE ZINC AND ILETIN II			
LILLY	100 UNITS/ML	N18476 001	

PROTAMINE ZINC INSULIN			
BRISTOL MYERS SQUIBB	40 UNITS/ML	N17928 001	
	100 UNITS/ML	N17928 003	

DISCONTINUED DRUG PRODUCT LIST

6 - 144 (of 263)

INSULIN SUSP PROTAMINE ZINC PURIFIED PORK

INJECTABLE; INJECTION			
PROTAMINE ZINC AND ILETIN II (PORK)			
LILLY	100 UNITS/ML	N18346	001

INSULIN ZINC SUSP BEEF

INJECTABLE; INJECTION			
LENTE INSULIN			
NOVO NORDISK INC	40 UNITS/ML	N17998	001
	100 UNITS/ML	N17998	003

INSULIN ZINC SUSP EXTENDED BEEF

INJECTABLE; INJECTION			
ULTRALENTE INSULIN			
NOVO NORDISK INC	100 UNITS/ML	N17997	003

INSULIN ZINC SUSP EXTENDED PURIFIED BEEF

INJECTABLE; INJECTION			
ULTRALENTE			
NOVO NORDISK INC	100 UNITS/ML	N18385	001

INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN

INJECTABLE; INJECTION			
HUMULIN U			
LILLY	40 UNITS/ML	N19571	001 Jun 10, 1987

INSULIN ZINC SUSP PROMPT BEEF

INJECTABLE; INJECTION			
SEMILENTE INSULIN			
NOVO NORDISK INC	100 UNITS/ML	N17996	003

INSULIN ZINC SUSP PROMPT PURIFIED PORK

INJECTABLE; INJECTION			
SEMILENTE			
NOVO NORDISK INC	100 UNITS/ML	N18382	001

INSULIN ZINC SUSP PURIFIED BEEF

INJECTABLE; INJECTION			
LENTE ILETIN II			
LILLY	100 UNITS/ML	N18477	001

INSULIN ZINC SUSP PURIFIED BEEF/PORK

INJECTABLE; INJECTION			
LENTARD			
NOVO NORDISK INC	100 UNITS/ML	N18384	001

INSULIN ZINC SUSP PURIFIED PORK

INJECTABLE; INJECTION			
LENTE			
NOVO NORDISK INC	100 UNITS/ML	N18383	001

INSULIN ZINC SUSP SEMISYNTHETIC PURIFIED HUMAN

INJECTABLE; INJECTION			
NOVOLIN L			
NOVO NORDISK INC	100 UNITS/ML	N18777	001 Aug 30, 1983

INULIN

INJECTABLE; INJECTION			
INULIN AND SODIUM CHLORIDE			
ISO TEX	100MG/ML	N02282	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 145 (of 263)

INVERT SUGAR

INJECTABLE; INJECTION			
TRAVERS 10% IN PLASTIC CONTAINER			
BAXTER HLTHCARE	10GM/100ML	N16717	001

IO CETAMIC ACID

TABLET; ORAL			
CHOLEBRINE			
MALLINCKRODT	750MG	N17129	001

IODAMIDE MEGLUMINE

INJECTABLE; INJECTION			
RENOVUE-65			
BRACCO	65%	N17902	001
RENOVUE-DIP			
BRACCO	24%	N17903	001

IODIPAMIDE MEGLUMINE

INJECTABLE; INJECTION			
CHOLOGRAFIN MEGLUMINE			
BRACCO	10.3%	N09321	007

IODIPAMIDE SODIUM

INJECTABLE; INJECTION			
CHOLOGRAFIN SODIUM			
BRACCO	20%	N09321	001

IODOHIPPURATE SODIUM, I-123

INJECTABLE; INJECTION				
NEPHROFLOW				
GE HEALTHCARE	1mCi/ML	N18289	001	Dec 28, 1984

IODOHIPPURATE SODIUM, I-131

INJECTABLE; INJECTION			
HIPPURAN I 131			
MALLINCKRODT	0.25mCi/ML	N16666	001
HIPPUTOPE			
BRACCO	1-2mCi/VIAL	N15419	002
IODOHIPPURATE SODIUM I 131			
CIS	0.2mCi/ML	N17313	001

IO DOXAMATE MEGLUMINE

INJECTABLE; INJECTION			
CHOLOVUE			
BRACCO	9.9%	N18077	001
	40.3%	N18076	001

IOFETAMINE HYDROCHLORIDE I-123

INJECTABLE; INJECTION				
SPECTAMINE				
IMP	1mCi/ML	N19432	001	Dec 24, 1987

IO HEXOL

INJECTABLE; INJECTION				
OMNIPAQUE 210				
GE HEALTHCARE	45.3%	N18956	006	Jun 30, 1989
SOLUTION; URETHRAL				
OMNIPAQUE 70				
GE HEALTHCARE	15.1%	N18956	007	Jun 01, 1994

DISCONTINUED DRUG PRODUCT LIST

6 - 146 (of 263)

IOPAMIDOL

INJECTABLE; INJECTION
IOPAMIDOL

BAXTER HLTHCARE	41%	N74629 001	Nov 06, 1996
	51%	N74629 004	Mar 31, 1998
	61%	N74629 002	Nov 06, 1996
	76%	N74629 003	Nov 06, 1996
MAYNE PHARMA USA	61%	N74734 001	Dec 10, 1996
	76%	N74734 002	Dec 10, 1996
IOPAMIDOL-300			
ABBOTT	61%	N74638 001	Apr 30, 1997
ISOVUE-128			
BRACCO	26%	N18735 005	Oct 21, 1986
ISOVUE-200			
BRACCO	41%	N20327 001	Oct 12, 1994

IOPANOIC ACID

TABLET; ORAL
TELEPAQUE

GE HEALTHCARE	500MG	N08032 001	
---------------	-------	------------	--

IOPHENDYLATE

INJECTABLE; INJECTION
PANTOPAQUE

ALCON	100%	N05319 001	
-------	------	------------	--

IOTHALAMATE MEGLUMINE; IOTHALAMATE SODIUM

INJECTABLE; INJECTION
VASCORAY

MALLINCKRODT	52%;26%	N16783 001	
--------------	---------	------------	--

IOTHALAMATE SODIUM

INJECTABLE; INJECTION
ANGIO-CONRAY

MALLINCKRODT	80%	N13319 001	
--------------	-----	------------	--

CONRAY 325			
MALLINCKRODT	54.3%	N17685 001	

IOTROLAN

INJECTABLE; INTRATHECAL
OSMOVIST 190

BERLEX	40.6%	N19580 001	Dec 07, 1989
--------	-------	------------	--------------

OSMOVIST 240			
BERLEX	51.3%	N19580 002	Dec 07, 1989

IOVERSOL

INJECTABLE; INJECTION
OPTIRAY 240

MALLINCKRODT	51%	N20923 001	May 28, 1998
--------------	-----	------------	--------------

OPTIRAY 320			
MALLINCKRODT	68%	N20923 002	May 29, 1998

IPODATE CALCIUM

GRANULE; ORAL
ORAGRAFIN CALCIUM

BRACCO	3GM/PACKET	N12968 001	
--------	------------	------------	--

IPODATE SODIUM

CAPSULE; ORAL
BILIVIST

BERLEX	500MG	N87768 001	Aug 11, 1982
--------	-------	------------	--------------

ORAGRAFIN SODIUM			
BRACCO	500MG	N12967 001	

DISCONTINUED DRUG PRODUCT LIST

6 - 147 (of 263)

IRON DEXTRAN

INJECTABLE; INJECTION
IRON DEXTRAN

AVENTIS PHARMS	EQ 50MG IRON/ML	N10787	002
----------------	-----------------	--------	-----

ISOETHARINE HYDROCHLORIDE

SOLUTION; INHALATION
BRONKOSOL

SANOFI SYNTHELABO	0.25%	N12339	009
	1%	N12339	008

ISOETHARINE HYDROCHLORIDE

ALPHARMA US PHARMS	1%	N87101	001	
ASTRAZENECA	0.062%	N87937	001	Nov 15, 1982
	0.062%	N89614	001	Jun 13, 1991
	0.125%	N87938	001	Nov 15, 1982
	0.125%	N89615	001	Jun 13, 1991
	0.167%	N88470	001	Mar 14, 1984
	0.167%	N89616	001	Jun 13, 1991
	0.2%	N88471	001	Mar 14, 1984
	0.2%	N89617	001	Jun 13, 1991
	0.25%	N88472	001	Mar 14, 1984
	0.25%	N89618	001	Jun 13, 1991
BAXTER HLTHCARE	0.08%	N88144	001	Jul 29, 1983
	0.14%	N88145	001	Mar 26, 1984
	0.25%	N88146	001	Aug 01, 1983
DEY	0.08%	N88187	001	Dec 03, 1982
	0.1%	N87389	001	
	0.17%	N87390	001	
	0.25%	N88188	001	Dec 03, 1982
	1%	N86763	001	
INTL MEDICATION	0.077%	N86651	001	
	0.08%	N86651	002	
	0.1%	N86651	003	
	0.143%	N86651	004	
	0.167%	N86651	005	
	0.2%	N86651	006	
	0.25%	N86651	007	
	1%	N86651	008	
PARKE DAVIS	0.5%	N85997	001	
	1%	N85889	001	
ROXANE	0.1%	N87396	001	
	0.125%	N87025	001	
	0.167%	N88226	001	Sep 16, 1983
	0.2%	N87324	001	
	0.25%	N88275	001	Jun 03, 1983
	1%	N86899	001	
ISOETHARINE HYDROCHLORIDE S/F				
DEY	0.08%	N89817	001	Nov 22, 1988
	0.1%	N89818	001	Nov 22, 1988
	0.17%	N89819	001	Nov 22, 1988
	0.25%	N89820	001	Nov 22, 1988
	1%	N89252	001	Sep 15, 1986

ISOETHARINE MESYLATE

AEROSOL, METERED; INHALATION
BRONKOMETER

SANOFI SYNTHELABO	0.34MG/INH	N12339	007	
ISOETHARINE MESYLATE				
ALPHARMA US PHARMS	0.34MG/INH	N87858	001	Aug 21, 1984

ISOFLUROPHATE

OINTMENT; OPHTHALMIC
FLOROPRYL

MERCK	0.025%	N10656	001
-------	--------	--------	-----

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 148 (of 263)

ISONIAZID

INJECTABLE; INJECTION

RIMIFON

ROCHE

25MG/ML

N08420 002

100MG/ML

N08420 003

SYRUP; ORAL

ISONIAZID

MIKART

50MG/5ML

N81118 001

Jul 21, 1997

LANIAZID

LANNETT

50MG/5ML

N89243 001

Feb 03, 1986

RIMIFON

ROCHE

50MG/5ML

N08420 001

TABLET; ORAL

DOW-ISONIAZID

DOW PHARM

300MG

N80330 002

HYZYD

MEDPOINTE PHARM HLC

100MG

N80134 003

300MG

N80134 004

INH

NOVARTIS

300MG

N80935 001

ISONIAZID

ANABOLIC

100MG

N84050 001

DURAMED PHARMS BARR

100MG

N88231 001

Mar 17, 1983

HALSEY

50MG

N83632 001

IMPAX LABS

100MG

N80153 001

IVAX PHARMS

100MG

N80270 001

300MG

N83610 001

LILLY

100MG

N08499 002

300MG

N08499 003

MK LABS

100MG

N80941 001

MUTUAL PHARM

100MG

N80136 001

PANRAY

50MG

N08428 001

100MG

N08428 002

300MG

N08428 003

PERRIGO

100MG

N83060 001

PHARMAVITE

100MG

N85091 001

PHOENIX LABS NY

50MG

N80368 001

100MG

N80368 002

PUREPAC PHARM

50MG

N80132 003

Jul 14, 1982

100MG

N80132 004

Jul 14, 1982

WATSON LABS

50MG

N80522 001

100MG

N80401 001

100MG

N80523 001

100MG

N85790 001

300MG

N83178 001

300MG

N85784 001

WHITEWORTH TOWN PLSN

100MG

N80120 002

NYDRAZID

BRISTOL MYERS SQUIBB

100MG

N08392 003

STANOZIDE

EVERYLIFE

100MG

N80126 001

300MG

N80126 002

ISOPROPAMIDE IODIDE

TABLET; ORAL

DARBID

GLAXOSMITHKLINE

EQ 5MG BASE

N10744 001

ISOPROTERENOL HYDROCHLORIDE

AEROSOL, METERED; INHALATION

ISOPROTERENOL HYDROCHLORIDE

3M

0.12MG/INH

N10375 004

ALPHARMA US PHARMS

0.12MG/INH

N85904 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 149 (of 263)

ISOPROTERENOL HYDROCHLORIDE

AEROSOL, METERED; INHALATION			
ISUPREL			
SANOFI SYNTHELABO	0.103MG/INH	N11178	001
DISC; INHALATION			
NORISODRINE AEROTROL			
ABBOTT	0.25%	N16814	001
INJECTABLE; INJECTION			
ISOPROTERENOL HYDROCHLORIDE			
AM PHARM PARTNERS	0.2MG/ML	N83431	001
BAXTER HLTHCARE	0.2MG/ML	N83486	001
SOLUTION; INHALATION			
AEROLONE			
LILLY	0.25%	N07245	001
ISOPROTERENOL HYDROCHLORIDE			
ARMOUR PHARM	0.031%	N87935	001
	0.062%	N87936	001
DEY	0.5%	N86764	001
PARKE DAVIS	0.25%	N85994	001
	0.5%	N85540	001
ISUPREL			
SANOFI SYNTHELABO	0.5%	N06327	002
	1%	N06327	003
VAPO-ISO			
FISONS	0.5%	N16813	001
TABLET; RECTAL, SUBLINGUAL			
ISUPREL			
SANOFI SYNTHELABO	10MG	N06328	001
	15MG	N06328	002

Nov 18, 1982

Nov 18, 1982

Jan 04, 1982

ISOPROTERENOL HYDROCHLORIDE; PHENYLEPHRINE BITARTRATE

AEROSOL, METERED; INHALATION			
DUO-MEDIHALER			
3M	0.16MG/INH;0.24MG/INH	N13296	001

ISOPROTERENOL SULFATE

AEROSOL, METERED; INHALATION			
MEDIHALER-ISO			
3M	0.08MG/INH	N10375	003
POWDER; INHALATION			
NORISODRINE			
ABBOTT	10%	N06905	003
	25%	N06905	002

ISOSORBIDE

SOLUTION; ORAL			
ISMOTIC			
ALCON	100GM/220ML	N17063	001

ISOSORBIDE DINITRATE

CAPSULE, EXTENDED RELEASE; ORAL			
ISORDIL			
WYETH AYERST	40MG	N12882	002
TABLET; ORAL			
ISOSORBIDE DINITRATE			
MUTUAL PHARM	5MG	N86166	002
	10MG	N86169	001
	20MG	N86167	001
	30MG	N87564	001
SUPERPHARM	5MG	N89190	001
	10MG	N89191	001
	20MG	N89192	001
SORBITRATE			
ASTRAZENECA	5MG	N16192	001

Jul 29, 1988

Sep 19, 1986

Sep 19, 1986

Sep 19, 1986

Sep 18, 1986

Feb 17, 1987

Feb 17, 1987

Feb 17, 1987

Apr 01, 1996

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 150 (of 263)

ISOSORBIDE DINITRATETABLET; ORAL
SORBITRATE

ASTRAZENECA

10MG

N16192 002 Apr 01, 1996

20MG

N86405 002 Aug 21, 1990

30MG

N88124 001 Aug 21, 1990

40MG

N88125 001 Aug 21, 1990

TABLET; SUBLINGUAL
ISOSORBIDE DINITRATE

MUTUAL PHARM

2.5MG

N84204 001 Sep 18, 1986

5MG

N86168 001 Sep 18, 1986

10MG

N87545 001 Sep 18, 1986

SORBITRATE

ASTRAZENECA

2.5MG

N16191 002 Apr 01, 1996

5MG

N16191 001 Apr 01, 1996

TABLET, CHEWABLE; ORAL
SORBITRATE

ASTRAZENECA

5MG

N16776 002 Apr 01, 1996

10MG

N16776 003 Apr 01, 1996

TABLET, EXTENDED RELEASE; ORAL
ISORDIL

WYETH AYERST

40MG

N12882 001 Jul 29, 1988

ISOSORBIDE MONONITRATETABLET, EXTENDED RELEASE; ORAL
ISOSORBIDE MONONITRATE

IVAX PHARMS

30MG

N75448 002 Aug 07, 2001

60MG

N75448 001 Jun 19, 2000

120MG

N75448 003 Aug 07, 2001

ISRADIPINECAPSULE; ORAL
DYNACIRC

RELIANT PHARMS

5MG

N19546 002 Dec 20, 1990

KANAMYCIN SULFATECAPSULE; ORAL
KANTREX

APOTHECON

EQ 500MG BASE

N60516 001

EQ 500MG BASE

N61911 001

EQ 500MG BASE

N62726 001 Mar 06, 1987

INJECTABLE; INJECTION
KANAMYCIN

BAXTER HLTHCARE

EQ 75MG BASE/2ML

N62324 001

EQ 500MG BASE/2ML

N62324 002

EQ 1GM BASE/3ML

N62324 003

KANAMYCIN SULFATE

AM PHARM PARTNERS

EQ 75MG BASE/2ML

N62504 001 Apr 05, 1984

EQ 500MG BASE/2ML

N62504 002 Apr 05, 1984

EQ 1GM BASE/3ML

N62504 003 Apr 05, 1984

INTL MEDICATION

EQ 500MG BASE/2ML

N62466 001 Sep 30, 1983

EQ 1GM BASE/3ML

N62466 002 Sep 30, 1983

LOCH

EQ 75MG BASE/2ML

N63021 001 Jul 31, 1992

EQ 500MG BASE/2ML

N63022 001 Jul 31, 1992

EQ 1GM BASE/3ML

N63025 001 Jul 31, 1992

PHARMAFAIR

EQ 75MG BASE/2ML

N62668 001 May 07, 1987

EQ 500MG BASE/2ML

N62672 001 May 07, 1987

EQ 1GM BASE/3ML

N62669 001 May 07, 1987

SOLOPAK

EQ 75MG BASE/2ML

N62605 003 Feb 26, 1986

EQ 500MG BASE/2ML

N62605 001 Feb 26, 1986

EQ 1GM BASE/3ML

N62605 002 Feb 26, 1986

STERIS

EQ 1GM BASE/3ML

N62520 003 May 09, 1985

WARNER CHILCOTT

EQ 1GM BASE/3ML

N63092 001 Oct 11, 1989

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 151 (of 263)

KANAMYCIN SULFATE

INJECTABLE; INJECTION

KANTREX

APOTHECON

EQ 75MG BASE/2ML

N61655 003

EQ 75MG BASE/2ML

N62564 001

Sep 21, 1984

EQ 500MG BASE/2ML

N61655 001

EQ 500MG BASE/2ML

N62564 002

Sep 21, 1984

EQ 1GM BASE/3ML

N61655 002

EQ 1GM BASE/3ML

N62564 003

Sep 21, 1984

KLEBCIL

KING PHARMS

EQ 75MG BASE/2ML

N62170 001

EQ 500MG BASE/2ML

N62170 002

EQ 1GM BASE/3ML

N62170 003

KETOCONAZOLE

CREAM; TOPICAL

NIZORAL

JANSSEN PHARMA

2%

N19084 001

Dec 31, 1985

SUSPENSION; ORAL

NIZORAL

JANSSEN PHARMA

100MG/5ML

N70767 001

Nov 07, 1986

KETOPROFEN

CAPSULE; ORAL

ORUDIS

WYETH AYERST

25MG

N18754 001

Jul 31, 1987

50MG

N18754 002

Jan 09, 1986

75MG

N18754 003

Jan 09, 1986

CAPSULE, EXTENDED RELEASE; ORAL

ORUVAIL

WYETH PHARMS INC

100MG

N19816 003

Feb 08, 1995

150MG

N19816 002

Feb 08, 1995

TABLET; ORAL

KETOPROFEN

PERRIGO

12.5MG

N75364 001

Feb 07, 2002

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

APOTHECON

15MG/ML

N75348 001

Nov 28, 2000

30MG/ML

N75348 002

Nov 28, 2000

BEDFORD

15MG/ML

N75230 002

Oct 25, 1999

30MG/ML

N75230 001

Oct 25, 1999

HOSPIRA

15MG/ML

N74801 001

Jun 05, 1997

30MG/ML

N74801 002

Jun 05, 1997

TORADOL

ROCHE PALO

15MG/ML

N19698 001

Nov 30, 1989

30MG/ML

N19698 002

Nov 30, 1989

TABLET; ORAL

KETOROLAC TROMETHAMINE

ROXANE

10MG

N74790 001

Jun 26, 1997

KRYPTON, KR-81M

GAS; INHALATION

MPI KRYPTON 81M GAS GENERATOR

GE HEALTHCARE

N/A

N18088 001

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HYDROCHLORIDE

APOTHECON

5MG/ML

N75355 001

Nov 29, 1999

NORMODYNE

SCHERING

5MG/ML

N18686 001

Aug 01, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 152 (of 263)

LABETALOL HYDROCHLORIDE

TABLET; ORAL

LABETALOL HYDROCHLORIDE

APOTHECON

100MG

N75223 001

Nov 20, 1998

200MG

N75223 002

Nov 20, 1998

300MG

N75223 003

Nov 20, 1998

NORMODYNE

SCHERING

100MG

N18687 001

Aug 31, 1987

200MG

N18687 002

Aug 01, 1984

300MG

N18687 003

Aug 01, 1984

400MG

N18687 004

Aug 01, 1984

TRANDATE

PROMETHEUS LABS

400MG

N18716 004

Aug 01, 1984

LACTULOSE

SOLUTION; ORAL

CHRONULAC

AVENTIS PHARMS

10GM/15ML

N17884 001

DUPHALAC

SOLVAY

10GM/15ML

N72372 001

Mar 22, 1989

LACTULOSE

MORTON GROVE

10GM/15ML

N71841 001

Sep 22, 1988

PACO

10GM/15ML

N73160 001

Aug 25, 1992

SOLUTION; ORAL, RECTAL

CEPHULAC

AVENTIS PHARMS

10GM/15ML

N17657 001

GENERLAC

MORTON GROVE

10GM/15ML

N71842 001

Sep 27, 1988

LACTULOSE

PACO

10GM/15ML

N72029 001

Aug 25, 1992

ROXANE

10GM/15ML

N73590 001

May 29, 1992

SOLVAY

10GM/15ML

N17906 001

PORTALAC

SOLVAY

10GM/15ML

N72374 001

Mar 22, 1989

LAMOTRIGINE

TABLET; ORAL

LAMICTAL

GLAXOSMITHKLINE

50MG

N20241 006

Dec 27, 1994

250MG

N20241 004

Dec 27, 1994

TABLET, CHEWABLE; ORAL

LAMICTAL CD

GLAXOSMITHKLINE

100MG

N20764 003

Aug 24, 1998

LEFLUNOMIDE

TABLET; ORAL

LEFLUNOMIDE

SANDOZ

10MG

N77085 001

Sep 13, 2005

20MG

N77085 002

Sep 13, 2005

LEUCOVORIN CALCIUM

FOR SOLUTION; ORAL

LEUCOVORIN CALCIUM

MAYNE PHARMA USA

EQ 60MG BASE/VIAL

N08107 003

Jan 30, 1987

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

ABIC

EQ 3MG BASE/ML

N89352 001

Jun 01, 1988

EQ 50MG BASE/VIAL

N89353 001

Jun 01, 1988

AM PHARM PARTNERS

EQ 50MG BASE/VIAL

N88939 001

Dec 01, 1986

ELKINS SINN

EQ 50MG BASE/VIAL

N70480 001

Jan 02, 1987

EQ 100MG BASE/VIAL

N81224 001

Jun 03, 1994

MAYNE PHARMA USA

EQ 3MG BASE/ML

N08107 001

EQ 50MG BASE/VIAL

N08107 002

EQ 100MG BASE/VIAL

N08107 004

May 23, 1988

DISCONTINUED DRUG PRODUCT LIST

6 - 153 (of 263)

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

PHARMACHEMIE USA EQ 100MG BASE/VIAL

N89915 001 Apr 17, 1997

LEUCOVORIN CALCIUM PRESERVATIVE FREE

HOSPIRA EQ 10MG BASE/ML

N40147 001 Jun 25, 1997

WELLCOVORIN

GLAXOSMITHKLINE EQ 5MG BASE/ML

N87439 001 Oct 19, 1982

EQ 25MG BASE/VIAL

N89833 001 Jan 23, 1989

EQ 50MG BASE/VIAL

N89465 001 Jan 23, 1989

EQ 100MG BASE/VIAL

N89834 001 Jan 23, 1989

TABLET; ORAL

LEUCOVORIN CALCIUM

PAR PHARM EQ 5MG BASE

N71600 001 Oct 14, 1987

EQ 25MG BASE

N71598 001 Oct 14, 1987

WELLCOVORIN

GLAXOSMITHKLINE EQ 5MG BASE

N18342 001 Jul 08, 1983

EQ 25MG BASE

N18342 002 Jul 08, 1983

LEUPROLIDE ACETATE

INJECTABLE; INJECTION

LUPRON DEPOT-PED

TAP PHARM 3.75MG/VIAL, 7.5MG/VIAL

N20263 003 Apr 16, 1993

7.5MG/VIAL, 7.5MG/VIAL

N20263 004 Apr 16, 1993

LEVALLORPHAN TARTRATE

INJECTABLE; INJECTION

LORFAN

ROCHE 1MG/ML

N10423 001

LEVAMISOLE HYDROCHLORIDE

TABLET; ORAL

ERGAMISOL

JANSSEN PHARMA EQ 50MG BASE

N20035 001 Jun 18, 1990

LEVOBETAXOLOL HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETAXON

ALCON EQ 0.5% BASE

N21114 001 Feb 23, 2000

LEVOPUIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

CHIROCAINE

PURDUE PHARMA LP EQ 2.5MG BASE/ML

N20997 001 Aug 05, 1999

EQ 5MG BASE/ML

N20997 002 Aug 05, 1999

EQ 7.5MG BASE/ML

N20997 003 Aug 05, 1999

LEVOCARNITINE

SOLUTION; ORAL

CARNITOR

SIGMA TAU 1GM/10ML

N18948 002 Apr 27, 1988

LEVODOPA

CAPSULE; ORAL

BENDOPA

VALEANT PHARM INTL 100MG

N16948 003

250MG

N16948 001

500MG

N16948 002

DOPAR

SHIRE 100MG

N16913 003

250MG

N16913 001

500MG

N16913 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 154 (of 263)

LEVODOPA

CAPSULE; ORAL

LARODOPA

ROCHE

100MG

N16912 002

250MG

N16912 001

500MG

N16912 006

TABLET; ORAL

DOPAR

SHIRE

250MG

N16913 004

500MG

N16913 005

LARODOPA

ROCHE

100MG

N16912 005

250MG

N16912 003

500MG

N16912 004

LEVOMEPRMAZINE

INJECTABLE; INJECTION

LEVOPROME

IMMUNEX

20MG/ML

N15865 001

LEVOMETHADYL ACETATE HYDROCHLORIDE

CONCENTRATE; ORAL

ORLAAM

ROXANE

10MG/ML

N20315 001

Jul 09, 1993

LEVONORDEFIN; MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ARESTOCAINE HYDROCHLORIDE W/ LEVONORDEFIN

SOLVAY

0.05MG/ML; 2%

N85010 001

LEVONORDEFIN; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

RAVOCAINE AND NOVOCAIN W/ NEO-COBEFRIN

EASTMAN KODAK

0.05MG/ML; 2%; 0.4%

N08592 007

LEVONORGESTREL

IMPLANT; IMPLANTATION

LEVONORGESTREL

WYETH PHARMS INC

75MG/IMPLANT

N20627 001

Aug 15, 1996

NORPLANT

POPULATION COUNCIL

36MG/IMPLANT

N19897 001

Dec 10, 1990

NORPLANT SYSTEM IN PLASTIC CONTAINER

WYETH PHARMS INC

36MG/IMPLANT

N20088 001

Dec 10, 1990

LEVOPROPOXYPHENE NAPSYLATE, ANHYDROUS

CAPSULE; ORAL

NOVRAD

LILLY

EQ 50MG BASE

N12928 006

EQ 100MG BASE

N12928 004

SUSPENSION; ORAL

NOVRAD

LILLY

EQ 50MG BASE/5ML

N12928 002

LEVORPHANOL TARTRATE

INJECTABLE; INJECTION

LEVO-DROMORAN

VALEANT PHARM INTL

2MG/ML

N08719 001

Dec 19, 1991

LIDOCAINE

AEROSOL; ORAL

XYLOCAINE

ASTRAZENECA

10%

N14394 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 155 (of 263)

LIDOCAINE

FILM, EXTENDED RELEASE; BUCCAL

LIDOCAINE

NOVEN

23MG/PATCH

N20575 001

May 21, 1996

OINTMENT; TOPICAL

ALPHACAINE

CARLISLE

5%

N84944 001

5%

N84946 001

5%

N84947 001

XYLOCAINE

ASTRAZENECA

5%

N08048 001

SOLUTION; TOPICAL

XYLOCAINE

ASTRAZENECA

5%

N14127 001

SUPPOSITORY; RECTAL

XYLOCAINE

ASTRAZENECA

100MG

N13077 001

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

ALPHACAINE HYDROCHLORIDE

CARLISLE

2%

N84721 001

LIDOCAINE HYDROCHLORIDE

ABBOTT

10%

N87980 001

Feb 02, 1983

20%

N89362 001

May 25, 1988

AKORN

1%

N85037 001

2%

N85037 002

AM PHARM PARTNERS

1%

N80390 001

1%

N80420 001

1%

N86761 001

1.5%

N80420 005

2%

N17508 001

2%

N80390 002

2%

N80420 002

2%

N80420 004

2%

N86761 002

4%

N17508 002

20%

N17508 004

BAXTER HLTHCARE

1%

N80407 001

2%

N80407 002

BEL MAR

1%

N80710 001

2%

N80760 001

DELL LABS

1%

N83387 001

2%

N83388 001

ELKINS SINN

0.5%

N85131 001

4%

N84626 001

GD SEARLE LLC

1%

N83135 001

2%

N83135 002

HOSPIRA

1.5%

N88330 001

May 17, 1984

INTL MEDICATION

1%

N17701 002

2%

N17701 001

1GM/VIAL

N18543 001

2GM/VIAL

N18543 002

MILES

1%

N80414 001

2%

N80414 002

STERIS

1%

N80377 001

1%

N83627 001

2%

N80377 002

2%

N83627 002

WYETH AYERST

1%

N83083 001

2%

N83083 002

LIDOCAINE HYDROCHLORIDE 0.1% AND DEXTROSE 5% IN PLASTIC CONTAINER

BAXTER HLTHCARE

100MG/100ML

N18461 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 156 (of 263)

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

LIDOCAINE HYDROCHLORIDE	0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	200MG/100ML	N18967	001	Mar 30, 1984
LIDOCAINE HYDROCHLORIDE	0.2% IN DEXTROSE 5% IN PLASTIC CONTAINER			
ABBOTT	200MG/100ML	N18954	001	Jul 09, 1985
LIDOCAINE HYDROCHLORIDE	0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	400MG/100ML	N18967	002	Mar 30, 1984
LIDOCAINE HYDROCHLORIDE	0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER			
B BRAUN	800MG/100ML	N18967	003	Mar 30, 1984
LIDOCAINE HYDROCHLORIDE	PRESERVATIVE FREE			
BAXTER HLTHCARE	1%	N84625	001	
	2%	N84625	002	
INTL MEDICATION	4%	N17702	002	
LIDOCATON				
PHARMATON	2%	N84727	001	Aug 17, 1983
XYLOCAINE				
ASTRAZENECA	1%	N10418	005	
	1.5%	N10418	009	
	2%	N06488	002	
	2%	N10418	007	

INJECTABLE; SPINAL

XYLOCAINE 1.5% W/ DEXTROSE 7.5%				
ASTRAZENECA	1.5%	N16297	001	
XYLOCAINE 5% W/ GLUCOSE 7.5%				
ASTRAZENECA	5%	N10496	002	Jul 07, 1982

SOLUTION; ORAL

LIDOCAINE HYDROCHLORIDE	VISCOUS			
INTL MEDICATION	2%	N86389	001	Feb 02, 1982

SOLUTION; TOPICAL

LARYNGOTRACHEAL ANESTHESIA KIT				
KENDALL IL	4%	N87931	001	Jun 10, 1983
LIDOCAINE HYDROCHLORIDE				
PACO	4%	N89688	001	Jun 30, 1989
PEDIATRIC LTA KIT				
ABBOTT	2%	N88572	001	Jul 31, 1984

LIDOCAINE; PRILOCAINE

DISC; TOPICAL

EMLA				
ASTRAZENECA	2.5%;2.5%	N20962	001	Feb 04, 1998

LINCOMYCIN HYDROCHLORIDE

CAPSULE; ORAL

LINCOCIN				
PHARMACIA AND UPJOHN	EQ 250MG BASE	N50316	001	
	EQ 500MG BASE	N50316	002	

INJECTABLE; INJECTION

LINCOMYCIN HYDROCHLORIDE				
STERIS	EQ 300MG BASE/ML	N63180	001	Apr 16, 1991

LINDANE

CREAM; TOPICAL

KWELL				
REED AND CARNRICK	1%	N06309	001	
	1%	N84218	001	

LOTION; TOPICAL

GAMENE				
SOLA BARNES HIND	1%	N84989	001	
KWELL				
REED AND CARNRICK	1%	N06309	003	
	1%	N84218	002	
SCABENE				
STIEFEL	1%	N86769	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 157 (of 263)

LINDANE

SHAMPOO; TOPICAL				
GAMENE				
SOLA BARNES HIND	1%	N84988	001	
KWELL				
REED AND CARNRICK	1%	N10718	001	
	1%	N84219	001	
SCABENE				
STIEFEL	1%	N87940	001	Apr 08, 1983

LINEZOLID

TABLET; ORAL				
ZYVOX				
PHARMACIA AND UPJOHN	400MG **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N21130	001	Apr 18, 2000

LIOTHYRONINE SODIUM

TABLET; ORAL				
LIOTHYRONINE SODIUM				
WATSON LABS	EQ 0.025MG BASE	N85755	001	Jan 25, 1982
	EQ 0.05MG BASE	N85753	001	Feb 03, 1982

LIOTRIX (T4;T3)

TABLET; ORAL				
EUTHROID-0.5				
PARKE DAVIS	0.03MG;0.0075MG	N16680	001	
EUTHROID-1				
PARKE DAVIS	0.06MG;0.015MG	N16680	002	
EUTHROID-2				
PARKE DAVIS	0.12MG;0.03MG	N16680	003	
EUTHROID-3				
PARKE DAVIS	0.18MG;0.045MG	N16680	004	
THYROLAR-5				
FOREST LABS	0.25MG;0.0625MG	N16807	006	

LITHIUM CARBONATE

CAPSULE; ORAL				
LITHIUM CARBONATE				
ABLE	150MG	N76823	001	Jun 29, 2004
	300MG	N76121	001	Sep 27, 2001
	300MG	N76823	002	Jun 29, 2004
	600MG	N76823	003	Jun 29, 2004
USL PHARMA	300MG	N72542	001	Feb 01, 1989
WATSON LABS	300MG	N70407	001	Mar 19, 1987
LITHONATE				
SOLVAY	300MG	N16782	001	
TABLET; ORAL				
ESKALITH				
GLAXOSMITHKLINE	300MG	N17971	001	
LITHANE				
BAYER PHARMS	300MG	N18833	001	Jul 18, 1985
LITHOTABS				
SOLVAY	300MG	N16980	001	
TABLET, EXTENDED RELEASE; ORAL				
LITHIUM CARBONATE				
ABLE	300MG	N76382	001	Apr 21, 2003

LITHIUM CITRATE

SYRUP; ORAL				
LITHONATE				
SOLVAY	EQ 300MG CARBONATE/5ML	N17672	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 158 (of 263)

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL			
IMODIUM			
MCNEIL CONS SPECLT	2MG	N17690 001	
LOPERAMIDE HYDROCHLORIDE			
ROXANE	2MG	N73080 001	Nov 27, 1991
SOLUTION; ORAL			
IMODIUM			
JANSSEN PHARMA	1MG/5ML	N19037 001	Jul 31, 1984
LOPERAMIDE HYDROCHLORIDE			
ALPHARMA US PHARMS	1MG/5ML	N73187 001	Sep 15, 1992
WATSON LABS	1MG/5ML	N73062 001	May 28, 1993
TABLET; ORAL			
LOPERAMIDE HYDROCHLORIDE			
ABLE	2MG	N73528 001	Nov 30, 1993
PERRIGO	2MG	N74194 001	Oct 30, 1992

LORATADINE

SYRUP; ORAL			
CLARITIN HIVES RELIEF			
SCHERING PLOUGH	1MG/ML	N20641 003	Nov 19, 2003
TABLET; ORAL			
LORATADINE			
PERRIGO	10MG	N21512 001	Jun 24, 2004

LORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL			
CLARITIN-D			
SCHERING PLOUGH	5MG;120MG	N19670 002	Nov 27, 2002

LORAZEPAM

INJECTABLE; INJECTION			
LORAZEPAM			
AKORN	2MG/ML	N74974 001	Jul 23, 1998
BAXTER HLTHCARE	2MG/ML	N74496 001	Sep 28, 1998
	4MG/ML	N74496 002	Sep 28, 1998
CLONMEL HLTHCARE	2MG/ML	N74793 001	Mar 16, 2000
	4MG/ML	N74793 002	Mar 16, 2000
MARSAM PHARMS LLC	1MG/0.5ML	N74551 003	Sep 12, 1996
	2MG/ML	N74535 001	Sep 12, 1996
	2MG/ML	N74551 001	Sep 12, 1996
	4MG/ML	N74535 002	Sep 12, 1996
	4MG/ML	N74551 002	Sep 12, 1996
TAYLOR	2MG/ML	N75025 001	Jul 23, 1998
TABLET; ORAL			
LORAZ			
QUANTUM PHARMICS	0.5MG	N70200 001	Aug 09, 1985
	1MG	N70201 001	Aug 09, 1985
	2MG	N70202 001	Aug 09, 1985
LORAZEPAM			
AM THERAP	0.5MG	N70727 001	Mar 07, 1986
	1MG	N70728 001	Mar 07, 1986
	2MG	N70729 001	Mar 07, 1986
HALSEY	0.5MG	N71434 001	Sep 01, 1987
	1MG	N71435 001	Sep 01, 1987
	2MG	N71436 001	Sep 01, 1987
MUTUAL PHARM	0.5MG	N70472 001	Dec 10, 1985
	1MG	N70473 001	Dec 10, 1985
	2MG	N70474 001	Dec 10, 1985
PAR PHARM	0.5MG	N70675 001	Dec 01, 1986
	1MG	N70676 001	Dec 01, 1986
	2MG	N70677 001	Dec 01, 1986
SUPERPHARM	0.5MG	N71245 001	Feb 09, 1987
	1MG	N71246 001	Feb 09, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 159 (of 263)

LORAZEPAM

TABLET; ORAL

LORAZEPAM

SUPERPHARM	2MG	N71247	001	Feb 09, 1987
USL PHARMA	1MG	N70539	001	Dec 22, 1986
	2MG	N70540	001	Dec 22, 1986
WARNER CHILCOTT	1MG	N71038	001	Jan 12, 1988
	2MG	N71039	001	Jan 12, 1988
WATSON LABS	0.5MG	N71086	001	Mar 23, 1987
	1MG	N71087	001	Mar 23, 1987
	2MG	N71088	001	Mar 23, 1987

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS; OPHTHALMIC

LOTEMAX

PHARMOS	0.5%	N20841	001	Mar 09, 1998
---------	------	--------	-----	--------------

LOVASTATIN; NIACIN

TABLET, EXTENDED RELEASE; ORAL

ADVICOR

KOS LIFE	20MG;750MG	N21249	002	Dec 17, 2001
----------	------------	--------	-----	--------------

LOXAPINE HYDROCHLORIDE

CONCENTRATE; ORAL

LOXITANE C

WATSON LABS	EQ 25MG BASE/ML	N17658	001	
-------------	-----------------	--------	-----	--

INJECTABLE; INJECTION

LOXITANE IM

WATSON LABS	EQ 50MG BASE/ML	N18039	001	
-------------	-----------------	--------	-----	--

LOXAPINE SUCCINATE

TABLET; ORAL

LOXITANE

WATSON LABS	EQ 10MG BASE	N17525	006	
	EQ 25MG BASE	N17525	007	
	EQ 50MG BASE	N17525	008	

LYPRESSIN

SOLUTION; NASAL

DIAPID

NOVARTIS	0.185MG/ML	N16755	001	
----------	------------	--------	-----	--

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE, MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE; SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE

INJECTABLE; INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER

B BRAUN	30MG/100ML;37MG/100ML;0.82MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML;12MG/100ML	N19006	001	Apr 04, 1984
---------	---	--------	-----	--------------

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE

INJECTABLE; INJECTION

ISOLYTE S IN PLASTIC CONTAINER

B BRAUN	30MG/100ML;37MG/100ML;370MG/100ML;530MG/100ML;500MG/100ML	N18252	001	
---------	---	--------	-----	--

SOLUTION; IRRIGATION

PHYSIOSOL IN PLASTIC CONTAINER

HOSPIRA	14MG/100ML;37MG/100ML;222MG/100ML;526MG/100ML;502MG/100ML	N18406	001	
---------	---	--------	-----	--

SYNOVALYTE IN PLASTIC CONTAINER

BAXTER HLTHCARE	30MG/100ML;37MG/100ML;368MG/100ML;526MG/100ML;502MG/100ML	N19326	001	Jan 25, 1985
-----------------	---	--------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 160 (of 263)

MANGAFODIPIR TRISODIUM

INJECTABLE; INJECTION
TESLASCAN

GE HEALTHCARE	37.9MG/ML	N20652 001	Nov 26, 1997
---------------	-----------	------------	--------------

MANGANESE CHLORIDE TETRAHYDRATE

FOR SOLUTION; ORAL
LUMENHANCE

BRACCO	3.49MG/GM	N20686 001	Dec 19, 1997
--------	-----------	------------	--------------

MANGANESE SULFATE

INJECTABLE; INJECTION
MANGANESE SULFATE

AM PHARM PARTNERS	EQ 0.1MG MANGANESE/ML	N19228 001	May 05, 1987
-------------------	-----------------------	------------	--------------

MANNITOL

INJECTABLE; INJECTION
MANNITOL 10%

HOSPIRA	10GM/100ML	N16269 002	
---------	------------	------------	--

MILES	10GM/100ML	N16472 002	
-------	------------	------------	--

MANNITOL 15%

HOSPIRA	15GM/100ML	N16269 003	
---------	------------	------------	--

MILES	15GM/100ML	N16472 005	
-------	------------	------------	--

MANNITOL 20%

HOSPIRA	20GM/100ML	N16269 004	
---------	------------	------------	--

MILES	20GM/100ML	N16472 004	
-------	------------	------------	--

MANNITOL 25%

AM PHARM PARTNERS	12.5GM/50ML	N86754 001	
-------------------	-------------	------------	--

ASTRAZENECA	12.5GM/50ML	N89239 001	May 06, 1987
-------------	-------------	------------	--------------

	12.5GM/50ML	N89240 001	May 06, 1987
--	-------------	------------	--------------

HOSPIRA	12.5GM/50ML	N16269 005	
---------	-------------	------------	--

MERCK	12.5GM/50ML	N05620 001	
-------	-------------	------------	--

STERIS	12.5GM/50ML	N87460 001	Jun 27, 1983
--------	-------------	------------	--------------

MANNITOL 5%

HOSPIRA	5GM/100ML	N16269 001	
---------	-----------	------------	--

SOLUTION; IRRIGATION

RESECTISOL

B BRAUN	5GM/100ML	N16704 002	
---------	-----------	------------	--

MANNITOL; SORBITOL

SOLUTION; IRRIGATION

SORBITOL-MANNITOL

HOSPIRA	540MG/100ML; 2.7GM/100ML	N80224 001	
---------	--------------------------	------------	--

MAPROTILINE HYDROCHLORIDE

TABLET; ORAL

LUDIOMIL

NOVARTIS	25MG	N17543 001	
----------	------	------------	--

	50MG	N17543 002	
--	------	------------	--

	75MG	N17543 003	Sep 30, 1982
--	------	------------	--------------

MAPROTILINE HYDROCHLORIDE

AM THERAP	25MG	N72129 001	Jan 14, 1988
-----------	------	------------	--------------

	50MG	N72130 001	Jan 14, 1988
--	------	------------	--------------

	75MG	N72131 001	Jan 14, 1988
--	------	------------	--------------

WATSON LABS	25MG	N71943 001	Dec 30, 1987
-------------	------	------------	--------------

	50MG	N71944 001	Dec 30, 1987
--	------	------------	--------------

	75MG	N71945 001	Dec 30, 1987
--	------	------------	--------------

MASOPROCOL

CREAM; TOPICAL

ACTINEX

UNIV AZ CANCER CTR	10%	N19940 001	Sep 04, 1992
--------------------	-----	------------	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 161 (of 263)

MAZINDOL

TABLET; ORAL
MAZANOR

WYETH AYERST	1MG	N17980	002	
	2MG	N17980	001	
SANOREX				
NOVARTIS	1MG	N17247	001	
	2MG	N17247	002	

MEBENDAZOLE

TABLET, CHEWABLE; ORAL
VERMOX

MCNEIL CONS SPECLT	100MG	N17481	001	
--------------------	-------	--------	-----	--

MEBUTAMATE

TABLET; ORAL
DORMATE

MEDPOINTE PHARM HLC	600MG	N17374	001	
---------------------	-------	--------	-----	--

MECLIZINE HYDROCHLORIDE

TABLET; ORAL
MECLIZINE HYDROCHLORIDE

ABC HOLDING	12.5MG	N85253	001	
	25MG	N85252	001	
ANABOLIC	25MG	N85891	001	
IVAX PHARMS	12.5MG	N83784	001	
KV PHARM	12.5MG	N85524	001	
	25MG	N85523	001	
SUPERPHARM	12.5MG	N89113	001	Aug 20, 1985
	25MG	N89114	001	Aug 20, 1985
UDL	12.5MG	N88256	001	Jun 13, 1983
	25MG	N88257	001	Jun 13, 1983
VANGARD	12.5MG	N87877	001	Apr 20, 1982
	25MG	N87620	001	Jan 04, 1982
WATSON LABS	12.5MG	N85195	001	
	12.5MG	N85269	001	
TABLET, CHEWABLE; ORAL				
MECLIZINE HYDROCHLORIDE				
ANABOLIC	25MG	N86392	001	
IVAX PHARMS	25MG	N84976	001	

MECLOCYCLINE SULFOSALICYLATE

CREAM; TOPICAL
MECLAN

JOHNSON AND JOHNSON	1%	N50518	001	
---------------------	----	--------	-----	--

MECLOFENAMATE SODIUM

CAPSULE; ORAL
MECLODIUM

QUANTUM PHARMICS	EQ 50MG BASE	N71380	001	Jul 14, 1987
	EQ 100MG BASE	N71381	001	Jul 14, 1987
MECLOFENAMATE SODIUM				
AM THERAP	EQ 50MG BASE	N71362	001	Feb 10, 1987
	EQ 100MG BASE	N71363	001	Feb 10, 1987
BARR	EQ 50MG BASE	N72848	001	Mar 20, 1989
	EQ 100MG BASE	N72809	001	Mar 20, 1989
PAR PHARM	EQ 50MG BASE	N72077	001	Mar 10, 1988
	EQ 100MG BASE	N72078	001	Mar 10, 1988
USL PHARMA	EQ 50MG BASE	N71007	001	Mar 25, 1988
	EQ 100MG BASE	N71008	001	Mar 25, 1988
VITARINE	EQ 50MG BASE	N71710	001	Jun 15, 1988
	EQ 100MG BASE	N71684	001	Jun 15, 1988
WATSON LABS	EQ 50MG BASE	N70400	001	Nov 25, 1986
	EQ 50MG BASE	N71640	001	Aug 11, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 162 (of 263)

MECLOFENAMATE SODIUM

CAPSULE; ORAL

MECLOFENAMATE SODIUM

WATSON LABS

EQ 100MG BASE

N70401 001

Nov 25, 1986

EQ 100MG BASE

N71641 001

Aug 11, 1987

MECLOMEN

PARKE DAVIS

EQ 50MG BASE

N18006 001

EQ 100MG BASE

N18006 002

MEDROXYPROGESTERONE ACETATE

INJECTABLE; INJECTION

DEPO-PROVERA

PHARMACIA AND UPJOHN

100MG/ML **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N12541 002

TABLET; ORAL

AMEN

AMARIN PHARMS

10MG

N83242 001

CURRETAB

SOLVAY

10MG

N85686 001

CYCRIN

ESI

2.5MG

N81239 001

Oct 30, 1992

5MG

N81240 001

Oct 30, 1992

10MG

N89386 001

Sep 09, 1987

MEFLOQUINE HYDROCHLORIDE

TABLET; ORAL

MEFLOQUINE HYDROCHLORIDE

US ARMY WALTER REED

250MG

N19578 001

May 02, 1989

MEGESTROL ACETATE

TABLET; ORAL

MEGESTROL ACETATE

USL PHARMA

20MG

N70646 001

Oct 02, 1987

40MG

N70647 001

Oct 02, 1987

MEMANTINE HYDROCHLORIDE

TABLET; ORAL

NAMENDA

FOREST LABS

15MG

N21487 003

Oct 16, 2003

20MG

N21487 004

Oct 16, 2003

MENADIOL SODIUM DIPHOSPHATE

INJECTABLE; INJECTION

KAPPADIONE

LILLY

10MG/ML

N05725 001

SYNKAYVITE

ROCHE

5MG/ML

N03718 004

10MG/ML

N03718 006

37.5MG/ML

N03718 008

TABLET; ORAL

SYNKAYVITE

ROCHE

5MG

N03718 010

MENADIONE

TABLET; ORAL

MENADIONE

LILLY

5MG

N02139 003

MENOTROPINS (FSH;LH)

INJECTABLE; INJECTION

HUMEGON

ORGANON USA INC

75 IU/VIAL;75 IU/VIAL

N20328 001

Sep 01, 1994

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 163 (of 263)

MENOTROPINS (FSH;LH)

INJECTABLE; INJECTION

HUMEGON

ORGANON USA INC	150 IU/VIAL;150 IU/VIAL	N20328	002	Sep 01, 1994
-----------------	-------------------------	--------	-----	--------------

MENOTROPINS

FERRING	75 IU/VIAL;75 IU/VIAL	N73598	001	Jan 30, 1997
	150 IU/VIAL;150 IU/VIAL	N73599	001	Jan 30, 1997

PERGONAL

SERONO	75 IU/AMP;75 IU/AMP	N17646	001	
	150 IU/AMP;150 IU/AMP	N17646	002	May 20, 1985

REPRONEX

FERRING	150 IU/VIAL;150 IU/VIAL	N21047	002	Aug 27, 1999
---------	-------------------------	--------	-----	--------------

MEPENZOLATE BROMIDE

SOLUTION; ORAL

CANTIL

AVENTIS PHARMS	25MG/5ML	N10679	004	
----------------	----------	--------	-----	--

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

DEMEROL

SANOFI SYNTHELABO	25MG/ML	N05010	007	
	50MG/ML	N05010	002	
	75MG/ML	N05010	009	
	100MG/ML	N05010	003	

MEPERIDINE HYDROCHLORIDE

ABBOTT

25MG/ML	N80388	001	
50MG/ML	N80385	001	
50MG/ML	N80387	001	
75MG/ML	N80389	001	
100MG/ML	N80386	001	

ASTRAZENECA

25MG/ML	N89781	001	Mar 31, 1989
50MG/ML	N89782	001	Mar 31, 1989
50MG/ML	N89783	001	Mar 31, 1989
50MG/ML	N89784	001	Mar 31, 1989
75MG/ML	N89785	001	Mar 31, 1989
100MG/ML	N89786	001	Mar 31, 1989
100MG/ML	N89787	001	Mar 31, 1989
100MG/ML	N89788	001	Mar 31, 1989

BAXTER HLTHCARE

25MG/ML	N88279	001	Jun 15, 1984
50MG/ML	N88280	001	Jun 15, 1984
75MG/ML	N88281	001	Jun 15, 1984
100MG/ML	N88282	001	Jun 15, 1984

INTL MEDICATION

10MG/ML	N86332	001	
---------	--------	-----	--

PARKE DAVIS

50MG/ML	N80364	002	
75MG/ML	N80364	003	
100MG/ML	N80364	001	

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

MAYNE PHARMA USA	10MG/ML	N40305	001	Mar 10, 1999
------------------	---------	--------	-----	--------------

TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

WYETH AYERST	50MG	N80454	001	
--------------	------	--------	-----	--

MEPERIDINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERGAN

BAXTER HLTHCARE CORP	25MG/ML;25MG/ML	N11730	001	
----------------------	-----------------	--------	-----	--

MEPHENTERMINE SULFATE

INJECTABLE; INJECTION

WYAMINE SULFATE

BAXTER HLTHCARE CORP	EQ 15MG BASE/ML	N08248	002	
	EQ 30MG BASE/ML	N08248	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 164 (of 263)

MEPHENYTOIN

TABLET; ORAL			
MESANTOIN			
NOVARTIS	100MG	N06008	001

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION			
ARESTOCAINE HYDROCHLORIDE			
SOLVAY	3%	N84777	002 Apr 18, 1982
MEPIVACAINE HYDROCHLORIDE			
INTL MEDICATION	1%	N87509	001 Oct 05, 1982

MEPREDNISONE

TABLET; ORAL			
BETAPAR			
SCHERING	4MG	N16053	002

MEPROBAMATE

CAPSULE; ORAL			
EQUANIL			
WYETH AYERST	400MG	N12455	002
CAPSULE, EXTENDED RELEASE; ORAL			
MEPROSPAN			
MEDPOINTE PHARM HLC	200MG	N11284	001
	400MG	N11284	002
TABLET; ORAL			
AMOSENE			
FERNDAL LABS	400MG	N84030	001
BAMATE			
ALRA	200MG	N80380	001
	400MG	N80380	002
EQUANIL			
WYETH AYERST	200MG	N10028	005
	400MG	N10028	004
MEPRIAM			
TEVA	400MG	N16069	001
MEPROBAMATE			
ANABOLIC	200MG	N84220	001
	400MG	N84589	001
BARR	600MG	N84230	001
ELKINS SINN	200MG	N15426	002
	400MG	N15426	001
HEATHER	400MG	N16928	003
	600MG	N84329	001
IMPAX LABS	200MG	N14322	002
	400MG	N14322	001
IVAX PHARMS	200MG	N15438	001
	400MG	N15438	002
	600MG	N84181	001
LANNETT	200MG	N14882	002
	400MG	N14882	001
LEDERLE	400MG	N86299	001
LEE KM	400MG	N89538	001 Nov 25, 1987
MALLARD	400MG	N15072	002
MK LABS	200MG	N14368	004
	400MG	N14368	002
MUTUAL PHARM	200MG	N80699	001
	400MG	N80699	002
MYLAN	400MG	N83618	001
PARKE DAVIS	200MG	N84744	001
	400MG	N84744	002
PERRIGO	200MG	N84546	001
	400MG	N84547	001
PHARMAVITE	400MG	N84438	001

DISCONTINUED DRUG PRODUCT LIST

6 - 165 (of 263)

MEPROBAMATE

TABLET; ORAL			
MEPROBAMATE			
PHARMERAL	400MG	N84153	001
PUREPAC PHARM	200MG	N84804	001
	400MG	N84804	002
PVT FORM	400MG	N14601	001
SOLVAY	200MG	N84435	001
STANLABS PHARM	200MG	N14474	002
	400MG	N14474	004
USL PHARMA	200MG	N87825	001
	400MG	N87826	001
VALEANT PHARM INTL	200MG	N15139	006
	400MG	N15139	005
VANGARD	400MG	N88011	001
WATSON LABS	600MG	N84274	001
	600MG	N85719	001
WEST WARD	200MG	N15417	003
	400MG	N15417	002
WHITEWORTH TOWN PLSN	200MG	N83830	001
	400MG	N83442	001
MILTOWN			
MEDPOINTE PHARM HLC	600MG	N83919	001
NEURAMATE			
HALSEY	200MG	N14359	002
	400MG	N14359	001
TRANMEP			
SOLVAY	400MG	N84369	001

Mar 18, 1982

Mar 18, 1982

Jul 14, 1982

MERSALYL SODIUM; THEOPHYLLINE

INJECTABLE; INJECTION			
MERSALYL-THEOPHYLLINE			
STERIS	100MG/ML;50MG/ML	N84875	001

MESALAMINE

SUPPOSITORY; RECTAL			
ROWASA			
SOLVAY	500MG	N19919	001

Dec 18, 1990

MESORIDAZINE BESYLATE

CONCENTRATE; ORAL			
SERENTIL			
NOVARTIS	EQ 25MG BASE/ML	N16997	001
INJECTABLE; INJECTION			
SERENTIL			
NOVARTIS	EQ 25MG BASE/ML	N16775	001
TABLET; ORAL			
SERENTIL			
NOVARTIS	EQ 10MG BASE	N16774	001
	EQ 25MG BASE	N16774	002
	EQ 50MG BASE	N16774	003
	EQ 100MG BASE	N16774	004

MESTRANOL; NORETHINDRONE

TABLET; ORAL-20			
NORINYL			
WATSON LABS	0.1MG;2MG	N13625	004
TABLET; ORAL-21			
NORINYL 1+80 21-DAY			
GD SEARLE LLC	0.08MG;1MG	N16724	001
ORTHO-NOVUM 1/50 21			
ORTHO MCNEIL PHARM	0.05MG;1MG	N12728	004
ORTHO-NOVUM 1/80 21			
ORTHO MCNEIL PHARM	0.08MG;1MG	N16715	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 166 (of 263)

MESTRANOL; NORETHINDRONE

TABLET; ORAL-21			
ORTHO-NOVUM 10-21			
ORTHO MCNEIL PHARM	0.06MG;10MG	N12728	001
ORTHO-NOVUM 2-21			
ORTHO MCNEIL PHARM	0.1MG;2MG	N12728	005
TABLET; ORAL-28			
NORINYL 1+80 28-DAY			
GD SEARLE LLC	0.08MG;1MG	N16725	001
ORTHO-NOVUM 1/80 28			
ORTHO MCNEIL PHARM	0.08MG;1MG	N16715	002

MESTRANOL; NORETHYNODREL

TABLET; ORAL			
ENOVID			
GD SEARLE LLC	0.075MG;5MG	N10976	008
	0.15MG;9.85MG	N10976	005
TABLET; ORAL-20			
ENOVID			
GD SEARLE LLC	0.075MG;5MG	N10976	004
ENOVID-E			
GD SEARLE LLC	0.1MG;2.5MG	N10976	006
TABLET; ORAL-21			
ENOVID-E 21			
GD SEARLE LLC	0.1MG;2.5MG	N10976	007

METAPROTERENOL SULFATE

SOLUTION; INHALATION				
ALUPENT				
BOEHRINGER INGELHEIM	5%	N17659	001	
METAPROTERENOL SULFATE				
ASTRAZENECA	0.4%	N71275	001	Jul 27, 1988
	0.6%	N71018	001	Jul 27, 1988
DEY	0.33%	N71806	001	Aug 05, 1988
	0.5%	N71805	001	Aug 05, 1988
	5%	N70805	001	Aug 17, 1987
MORTON GROVE	5%	N72190	001	Jun 07, 1988
PROMETA				
MURO	5%	N73340	001	Mar 30, 1992
SYRUP; ORAL				
ALUPENT				
BOEHRINGER INGELHEIM	10MG/5ML	N17571	001	
METAPROTERENOL SULFATE				
MORTON GROVE	10MG/5ML	N71656	001	Oct 13, 1987
TEVA	10MG/5ML	N72761	001	Feb 27, 1992
TEVA PHARMS	10MG/5ML	N73034	001	Aug 30, 1991
PROMETA				
MURO	10MG/5ML	N72023	001	Sep 15, 1988
TABLET; ORAL				
ALUPENT				
BOEHRINGER INGELHEIM	10MG	N15874	002	
	20MG	N15874	001	
METAPROTERENOL SULFATE				
AM THERAP	10MG	N72054	001	Jun 23, 1988
	20MG	N72055	001	Jun 23, 1988
USL PHARMA	10MG	N71013	001	Jan 25, 1988
	20MG	N71014	001	Jan 25, 1988

METARAMINOL BITARTRATE

INJECTABLE; INJECTION			
METARAMINOL BITARTRATE			
AM PHARM PARTNERS	EQ 10MG BASE/ML	N80431	001
ELKINS SINN	EQ 10MG BASE/ML	N83363	001
GD SEARLE LLC	EQ 10MG BASE/ML	N86418	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 167 (of 263)

METARAMINOL BITARTRATE

INJECTABLE; INJECTION			
METARAMINOL BITARTRATE			
GD SEARLE LLC	EQ 20MG BASE/ML	N86418	002

METAXALONE

TABLET; ORAL			
SKELAXIN			
JONES PHARMA INC	400MG	N13217	001

METFORMIN HYDROCHLORIDE

TABLET; ORAL			
GLUCOPHAGE			
BRISTOL MYERS SQUIBB	625MG	N20357	003 Nov 05, 1998
	750MG	N20357	004 Nov 05, 1998

METHACYCLINE HYDROCHLORIDE

CAPSULE; ORAL			
RONDONMYCIN			
MEDPOINTE PHARM HLC	EQ 140MG BASE	N60641	001
	EQ 280MG BASE	N60641	002
SYRUP; ORAL			
RONDONMYCIN			
MEDPOINTE PHARM HLC	EQ 70MG BASE/5ML	N60641	003

METHADONE HYDROCHLORIDE

SYRUP; ORAL			
DOLOPHINE HYDROCHLORIDE			
ROXANE	10MG/30ML	N06134	004
TABLET; ORAL			
METHADONE HYDROCHLORIDE			
SANDOZ	5MG	N40241	001 May 29, 1998
TABLET, DISPERSIBLE; ORAL			
WESTADONE			
SANDOZ	2.5MG	N17108	001
TABLET, EFFERVESCENT; ORAL			
WESTADONE			
SANDOZ	5MG	N17108	002
	10MG	N17108	003
	40MG	N17108	004

METHAMPHETAMINE HYDROCHLORIDE

TABLET; ORAL			
METHAMPEX			
TEVA	10MG	N83889	001
METHAMPHETAMINE HYDROCHLORIDE			
ABLE	5MG	N40529	001 Feb 25, 2004
REXAR	5MG	N84931	001
	10MG	N84931	002
TEVA	5MG	N86359	001
TABLET, EXTENDED RELEASE; ORAL			
DESOXYN			
OVATION PHARMS	5MG	N05378	004
	10MG	N05378	003
	15MG	N05378	005

METHANTHELINE BROMIDE

TABLET; ORAL			
BANTHINE			
SHIRE	50MG	N07390	001

DISCONTINUED DRUG PRODUCT LIST

6 - 168 (of 263)

METHARBITAL

TABLET; ORAL GEMONIL ABBOTT	100MG	N08322	001
-----------------------------------	-------	--------	-----

METHAZOLAMIDE

TABLET; ORAL METHAZOLAMIDE APPLIED ANAL	25MG 50MG	N40011 N40011	001 002	Jul 17, 1997 Jul 17, 1997
NEPTAZANE LEDERLE	25MG 50MG	N11721 N11721	002 001	Nov 25, 1991

METHDILAZINE

TABLET, CHEWABLE; ORAL TACARYL WESTWOOD SQUIBB	3.6MG	N11950	009
--	-------	--------	-----

METHDILAZINE HYDROCHLORIDE

SYRUP; ORAL METHDILAZINE HYDROCHLORIDE ALPHARMA US PHARMS	4MG/5ML	N87122	001
TACARYL WESTWOOD SQUIBB	4MG/5ML	N11950	007
TABLET; ORAL TACARYL WESTWOOD SQUIBB	8MG	N11950	006

METHICILLIN SODIUM

INJECTABLE; INJECTION STAPHICILLIN APOTHECON	EQ 900MG BASE/VIAL EQ 900MG BASE/VIAL EQ 3.6GM BASE/VIAL EQ 3.6GM BASE/VIAL EQ 5.4GM BASE/VIAL EQ 5.4GM BASE/VIAL	N50117 N61449 N50117 N61449 N50117 N61449	001 001 002 002 003 003
--	--	--	--

METHIMAZOLE

TABLET; ORAL METHIMAZOLE GENPHARM	20MG	N40350	003	Jun 07, 2001
TAPAZOLE KING PHARMS	5MG 10MG	N07517 N07517	002 004	

METHIXENE HYDROCHLORIDE

TABLET; ORAL TREST NOVARTIS	1MG	N13420	001
-----------------------------------	-----	--------	-----

METHOCARBAMOL

INJECTABLE; INJECTION METHOCARBAMOL MARSAM PHARMS LLC	100MG/ML	N89849	001	Dec 27, 1991
TABLET; ORAL DELAXIN FERNDAL LABS	500MG	N85454	001	
FORBAXIN FOREST LABS	750MG	N85136	001	
METHOCARBAMOL ABLE	500MG 750MG	N40413 N40413	001 002	Mar 17, 2003 Mar 17, 2003

DISCONTINUED DRUG PRODUCT LIST

6 - 169 (of 263)

METHOCARBAMOL

TABLET; ORAL
METHOCARBAMOL

AM THERAP	500MG	N89417	001	Feb 11, 1987
	750MG	N89418	001	Feb 11, 1987
ASCOT	500MG	N87660	001	Oct 27, 1982
	750MG	N87661	001	Oct 27, 1982
CLONMEL HLTHCARE	500MG	N85961	001	
	750MG	N85963	001	
HEATHER	500MG	N84675	001	
	750MG	N84924	001	
INWOOD LABS	500MG	N85137	001	
IVAX PHARMS	500MG	N84648	001	
	750MG	N84649	001	
KV PHARM	500MG	N85660	001	
	750MG	N85658	001	
MUTUAL PHARM	500MG	N84488	001	
	750MG	N84486	001	
MYLAN	500MG	N84259	001	
	750MG	N84323	001	
NYLOS	750MG	N85033	001	
PHARMERAL	500MG	N84231	002	
	750MG	N84471	001	
PIONEER PHARMS	500MG	N88731	001	Dec 13, 1985
	750MG	N89082	001	Dec 13, 1985
PUREPAC PHARM	500MG	N85718	001	
	750MG	N85718	002	
ROXANE	500MG	N88646	001	Feb 29, 1984
	750MG	N88647	001	Feb 29, 1984
SOLVAY	500MG	N84448	001	
	750MG	N84449	001	
SUPERPHARM	500MG	N87589	001	Jan 22, 1982
	750MG	N87590	001	Jan 22, 1982
UPSHER SMITH	500MG	N87453	001	
	750MG	N87454	001	
WATSON LABS	500MG	N83605	001	
	750MG	N83605	002	

METHOTREXATE SODIUM

INJECTABLE; INJECTION
ABITREXATE

ABIC	EQ 25MG BASE/ML	N89161	001	Mar 10, 1987
	EQ 50MG BASE/VIAL	N89354	001	Jul 17, 1987
	EQ 100MG BASE/VIAL	N89355	001	Jul 17, 1987
	EQ 250MG BASE/VIAL	N89356	001	Jul 17, 1987
FOLEX				
PHARMACIA AND UPJOHN	EQ 25MG BASE/VIAL	N87695	001	Apr 08, 1983
	EQ 50MG BASE/VIAL	N87695	002	Apr 08, 1983
	EQ 100MG BASE/VIAL	N87695	003	Apr 08, 1983
	EQ 250MG BASE/VIAL	N88954	001	Oct 24, 1985
FOLEX PFS				
PHARMACIA AND UPJOHN	EQ 25MG BASE/ML	N81242	001	Aug 23, 1991
	EQ 25MG BASE/ML	N89180	001	Jan 03, 1986
METHOTREXATE				
AM PHARM	EQ 25MG BASE/ML	N40263	001	Feb 26, 1999
METHOTREXATE LPF				
MAYNE PHARMA USA	EQ 25MG BASE/ML	N11719	007	Mar 31, 1982
METHOTREXATE PRESERVATIVE FREE				
AM PHARM PARTNERS	EQ 25MG BASE/ML	N40265	001	Feb 26, 1999
	EQ 1GM BASE/VIAL	N40266	001	Feb 26, 1999
MAYNE PHARMA USA	EQ 20MG BASE/2ML (10 MG/ML)	N11719	014	Apr 13, 2005
	EQ 500MG BASE/20ML (25 MG/ML)	N11719	013	Apr 13, 2005
	EQ 2.5GM BASE/100ML (25 MG/ML)	N11719	011	Apr 13, 2005

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 170 (of 263)

METHOTREXATE SODIUMINJECTABLE; INJECTION
METHOTREXATE SODIUM

AM PHARM PARTNERS	EQ 2.5MG BASE/ML	N89323	001	Jun 13, 1986
	EQ 20MG BASE/VIAL	N88935	001	Oct 11, 1985
	EQ 25MG BASE/ML	N89263	001	Jun 13, 1986
	EQ 25MG BASE/ML	N89322	001	Jun 13, 1986
	EQ 50MG BASE/VIAL	N88936	001	Oct 11, 1985
	EQ 100MG BASE/VIAL	N88937	001	Oct 11, 1985
MAYNE PHARMA USA	EQ 2.5MG BASE/ML	N11719	004	
	EQ 20MG BASE/VIAL	N11719	001	
	EQ 25MG BASE/ML	N11719	005	
	EQ 50MG BASE/VIAL	N11719	003	
	EQ 100MG BASE/VIAL	N11719	006	
NORBROOK	EQ 25MG BASE/ML	N88648	001	May 09, 1986
PHARMACHEMIE USA	EQ 25MG BASE/ML	N89158	001	Jul 08, 1988
METHOTREXATE SODIUM PRESERVATIVE FREE				
MAYNE PHARMA USA	EQ 1GM BASE/VIAL	N11719	009	Apr 07, 1988
MEXATE				
BRISTOL	EQ 20MG BASE/VIAL	N86358	001	
	EQ 50MG BASE/VIAL	N86358	002	
	EQ 100MG BASE/VIAL	N86358	003	
	EQ 250MG BASE/VIAL	N86358	004	
MEXATE-AQ				
BRISTOL MYERS	EQ 25MG BASE/ML	N88760	001	Feb 14, 1985
MEXATE-AQ PRESERVED				
BRISTOL MYERS SQUIBB	EQ 25MG BASE/ML	N89887	001	Apr 14, 1989

METHOXAMINE HYDROCHLORIDEINJECTABLE; INJECTION
VASOXYL

GLAXOSMITHKLINE	10MG/ML	N06772	002	
	20MG/ML	N06772	001	

METHOXSALENCAPSULE; ORAL
METHOXSALEN
SANDOZ

10MG	N87781	001	Jun 08, 1982
------	--------	-----	--------------

METHSCOPOLAMINE BROMIDETABLET; ORAL
METHSCOPOLAMINE BROMIDE
PVT FORM

2.5MG	N80970	001	
-------	--------	-----	--

METHYCLOTHIAZIDETABLET; ORAL
AQUATENSEN

MEDPOINTE PHARM HLC	5MG	N17364	001	
METHYCLOTHIAZIDE				
IVAX PHARMS	5MG	N87786	001	May 18, 1982
MYLAN	2.5MG	N87671	001	Aug 17, 1982
PAR PHARM	2.5MG	N89135	001	Feb 12, 1986
	5MG	N89136	001	Feb 12, 1986
USL PHARMA	5MG	N88745	001	Mar 21, 1985
WATSON LABS	2.5MG	N85487	001	Mar 11, 1982
	2.5MG	N88750	001	Sep 06, 1984
	5MG	N85476	001	Mar 11, 1982

METHYCLOTHIAZIDE; PARGYLINE HYDROCHLORIDETABLET; ORAL
EUTRON

ABBOTT	5MG; 25MG	N16047	001	
--------	-----------	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 171 (of 263)

METHYLCLOTHIAZIDE; RESERPINE

TABLET; ORAL				
DIUTENSEN-R				
MEDPOINTE PHARM HLC	2.5MG;0.1MG	N12708	005	

METHYLDOPA

SUSPENSION; ORAL				
ALDOMET				
MERCK	250MG/5ML	N18389	001	
TABLET; ORAL				
ALDOMET				
MERCK	125MG	N13400	003	
	250MG	N13400	001	
	500MG	N13400	002	

METHYLDOPA

ACCORD HLTH	125MG	N70070	003	Oct 15, 1985
DURAMED PHARMS BARR	250MG	N71006	001	Dec 16, 1986
	500MG	N71009	001	Dec 16, 1986
HALSEY	125MG	N71751	001	Mar 28, 1988
	250MG	N71752	001	Mar 28, 1988
	500MG	N71753	001	Mar 28, 1988
MUTUAL PHARM	125MG	N70073	001	Oct 09, 1986
	250MG	N70060	001	Oct 09, 1986
	500MG	N70074	001	Oct 09, 1986
PARKE DAVIS	125MG	N70331	001	Apr 15, 1986
	250MG	N70332	001	Apr 15, 1986
	500MG	N70333	001	Apr 15, 1986
PUREPAC PHARM	125MG	N70749	001	Feb 07, 1986
	250MG	N70750	001	Feb 07, 1986
	500MG	N70452	001	Feb 07, 1986
ROXANE	125MG	N70192	001	Apr 25, 1986
	250MG	N70193	001	Apr 25, 1986
	500MG	N70194	001	Apr 25, 1986
SUPERPHARM	250MG	N70669	001	Jun 23, 1989
	500MG	N70670	001	Jun 23, 1989
TEVA	125MG	N71105	001	Dec 05, 1986
	250MG	N71106	001	Dec 05, 1986
	500MG	N71067	001	Dec 05, 1986
WATSON LABS	125MG	N70245	001	Feb 25, 1986
	125MG	N70260	001	Jun 24, 1985
	250MG	N70246	001	Feb 25, 1986
	250MG	N70261	001	Jun 24, 1985
	500MG	N70247	001	Feb 25, 1986
	500MG	N70262	001	Jun 24, 1985

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION				
ALDOMET				
MERCK	50MG/ML	N13401	001	
METHYLDOPATE HYDROCHLORIDE				
AM PHARM PARTNERS	50MG/ML	N70652	001	Jun 03, 1986
BAXTER HLTHCARE	50MG/ML	N70291	001	Jul 01, 1986
MARSAM PHARMS LLC	50MG/ML	N71812	001	Dec 22, 1987
MAYNE PHARMA USA	50MG/ML	N70691	001	Jun 19, 1987
	50MG/ML	N70849	001	Jun 19, 1987
SMITH AND NEPHEW	50MG/ML	N70841	001	Jan 02, 1987

METHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL				
METHYLPHENIDATE HYDROCHLORIDE				
ABLE	5MG	N40404	001	Mar 29, 2001
	10MG	N40404	002	Mar 29, 2001
	20MG	N40404	003	Mar 29, 2001

DISCONTINUED DRUG PRODUCT LIST

6 - 172 (of 263)

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
METHYLPHENIDATE HYDROCHLORIDE
ABLE 20MG

N76032 001 May 09, 2001

METHYLPREDNISOLONE

TABLET; ORAL
MEDROL
PHARMACIA AND UPJOHN 2MG
METHYLPREDNISOLONE
HEATHER 4MG
SANDOZ 4MG
WATSON LABS 4MG
16MG

N11153 002
N85650 001
N87341 001
N86161 001 Feb 09, 1982
N86159 001 Feb 09, 1982

METHYLPREDNISOLONE ACETATE

ENEMA; RECTAL
MEDROL
PHARMACIA AND UPJOHN 40MG/BOT
INJECTABLE; INJECTION
METHYLPREDNISOLONE ACETATE
AKORN 40MG/ML
80MG/ML
STERIS 20MG/ML
20MG/ML
40MG/ML
40MG/ML
80MG/ML
80MG/ML
M-PREDROL
BEL MAR 40MG/ML
80MG/ML
OINTMENT; TOPICAL
MEDROL ACETATE
PHARMACIA AND UPJOHN 0.25%
1%

N18102 001
N86903 001 Oct 20, 1982
N86903 002 Oct 20, 1982
N85597 001
N87248 001
N85374 001
N85600 001
N85595 001
N86507 001
N86666 001
N87135 001
N12421 001
N12421 002

METHYLPREDNISOLONE ACETATE; NEOMYCIN SULFATE

CREAM; TOPICAL
NEO-MEDROL ACETATE
PHARMACIA AND UPJOHN 0.25%;EQ 3.5MG BASE/GM
1%;EQ 3.5MG BASE/GM

N60611 002
N60611 001

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION
A-METHAPRED
ABBOTT EQ 40MG BASE/VIAL
EQ 125MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
METHYLPREDNISOLONE
ELKINS SINN EQ 125MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
ORGANON USA INC EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
METHYLPREDNISOLONE SODIUM SUCCINATE
AM PHARM PARTNERS EQ 40MG BASE/VIAL
EQ 40MG BASE/VIAL
EQ 125MG BASE/VIAL
EQ 125MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 500MG BASE/VIAL

N89573 001 Feb 22, 1991
N89574 001 Feb 22, 1991
N89575 001 Feb 22, 1991
N89576 001 Feb 22, 1991
N86906 002
N86906 003
N86906 004
N87535 001 Jun 25, 1982
N87535 002 Jun 25, 1982
N88676 001 Jun 08, 1984
N89143 001 Mar 28, 1986
N88677 001 Jun 08, 1984
N89144 001 Mar 28, 1986
N88678 001 Jun 08, 1984
N89186 001 Mar 28, 1986
N89187 001 Mar 28, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 173 (of 263)

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

AM PHARM PARTNERS	EQ 1GM BASE/VIAL	N88679	001	Jun 08, 1984
	EQ 1GM BASE/VIAL	N89188	001	Mar 28, 1986
	EQ 1GM BASE/VIAL	N89189	001	Mar 28, 1986
ELKINS SINN	EQ 40MG BASE/VIAL	N86906	001	
	EQ 40MG BASE/VIAL	N87812	001	Feb 09, 1983
	EQ 125MG BASE/VIAL	N87813	001	Feb 09, 1983
INTL MEDICATION	EQ 500MG BASE/VIAL	N87851	001	Feb 09, 1983
	EQ 1GM BASE/VIAL	N87852	001	Feb 09, 1983
	EQ 500MG BASE/VIAL	N81267	001	Nov 30, 1992
SICOR PHARMS	EQ 1GM BASE/VIAL	N81268	001	Nov 30, 1992
	EQ 40MG BASE/VIAL	N86953	001	Jul 22, 1982
	EQ 125MG BASE/VIAL	N87030	001	Jul 22, 1982
STERIS	EQ 500MG BASE/VIAL	N88523	001	Jul 24, 1984
	EQ 1GM BASE/VIAL	N88524	001	Jul 24, 1984

METHYLPREDNISOLONE; NEOMYCIN SULFATE

OINTMENT; OPHTHALMIC

NEO-MEDROL

PHARMACIA AND UPJOHN	0.1%;EQ 3.5MG BASE/GM	N60645	001	
----------------------	-----------------------	--------	-----	--

METHYLTESTOSTERONE

CAPSULE; ORAL

METHYLTESTOSTERONE

HEATHER	10MG	N84967	001	
---------	------	--------	-----	--

TABLET; BUCCAL

ANDROID 5

VALEANT PHARM INTL	5MG	N87222	001	
--------------------	-----	--------	-----	--

ORETON

SCHERING	10MG	N80281	001	
----------	------	--------	-----	--

TABLET; BUCCAL/SUBLINGUAL

METANDREN

NOVARTIS	5MG	N03240	004	
----------	-----	--------	-----	--

	10MG	N03240	005	
--	------	--------	-----	--

METHYLTESTOSTERONE

IMPAX LABS	10MG	N84287	001	
------------	------	--------	-----	--

LILLY	10MG	N80256	001	
-------	------	--------	-----	--

PUREPAC PHARM	10MG	N80308	001	
---------------	------	--------	-----	--

	10MG	N80475	001	
--	------	--------	-----	--

PVT FORM	5MG	N83836	001	
----------	-----	--------	-----	--

TABLICAPS	10MG	N85125	001	
-----------	------	--------	-----	--

USL PHARMA	10MG	N80271	001	
------------	------	--------	-----	--

TABLET; ORAL

METANDREN

NOVARTIS	10MG	N03240	001	
----------	------	--------	-----	--

	25MG	N03240	003	
--	------	--------	-----	--

METHYLTESTOSTERONE

INWOOD LABS	10MG	N80839	001	
-------------	------	--------	-----	--

	25MG	N80973	001	
--	------	--------	-----	--

KV PHARM	10MG	N84312	001	
----------	------	--------	-----	--

LANNETT	10MG	N87092	001	Nov 05, 1982
---------	------	--------	-----	--------------

	25MG	N87111	001	Jan 27, 1983
--	------	--------	-----	--------------

LILLY	25MG	N80256	002	
-------	------	--------	-----	--

PARKE DAVIS	10MG	N84244	001	
-------------	------	--------	-----	--

	25MG	N84241	001	
--	------	--------	-----	--

PUREPAC PHARM	10MG	N80309	001	
---------------	------	--------	-----	--

	10MG	N80475	002	
--	------	--------	-----	--

	25MG	N80310	001	
--	------	--------	-----	--

	25MG	N80475	003	
--	------	--------	-----	--

PVT FORM	5MG	N80214	001	
----------	-----	--------	-----	--

	10MG	N80214	002	
--	------	--------	-----	--

	25MG	N80214	003	
--	------	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 174 (of 263)

METHYLTESTOSTERONE

TABLET; ORAL			
METHYLTESTOSTERONE			
TABLICAPS	10MG	N80313	001
	25MG	N85270	001
WATSON LABS	10MG	N80933	001
	25MG	N80931	001
WEST WARD	10MG	N84331	001
	25MG	N84331	002
	25MG	N84642	001
ORETON METHYL			
SCHERING	10MG	N03158	001
	25MG	N03158	002

METHYPRYLON

CAPSULE; ORAL			
NOLUDAR			
ROCHE	300MG	N09660	008
ELIXIR; ORAL			
NOLUDAR			
ROCHE	50MG/5ML	N09660	007
TABLET; ORAL			
NOLUDAR			
ROCHE	50MG	N09660	002
	200MG	N09660	004

METHYSERGIDE MALEATE

TABLET; ORAL			
SANSERT			
NOVARTIS	2MG	N12516	001

METOCLOPRAMIDE HYDROCHLORIDE

CONCENTRATE; ORAL			
METOCLOPRAMIDE INTENSOL			
ROXANE	EQ 10MG BASE/ML	N72995	001 Jan 30, 1992
INJECTABLE; INJECTION			
METOCLOPRAMIDE HYDROCHLORIDE			
AM PHARM PARTNERS	EQ 10MG BASE/2ML	N70293	001 Jan 24, 1986
BEDFORD	EQ 5MG BASE/ML	N72155	001 Mar 30, 1992
	EQ 5MG BASE/ML	N72244	001 Mar 30, 1992
	EQ 5MG BASE/ML	N72247	001 May 18, 1992
HOSPIRA	EQ 5MG BASE/ML	N70506	001 Jun 22, 1989
MAYNE PHARMA USA	EQ 5MG BASE/ML	N71990	001 Jan 18, 1989
NORBROOK	EQ 10MG BASE/2ML	N70892	001 Aug 26, 1988
SMITH AND NEPHEW	EQ 5MG BASE/ML	N70623	001 Mar 02, 1987
	EQ 10MG BASE/2ML	N70622	001 Mar 02, 1987
REGLAN			
BAXTER HLTHCARE CORP	EQ 10MG BASE/ML	N17862	004 May 28, 1987
SOLUTION; ORAL			
METOCLOPRAMIDE HYDROCHLORIDE			
MORTON GROVE	EQ 5MG BASE/5ML	N70949	001 Mar 06, 1987
PACO	EQ 5MG BASE/5ML	N71665	001 Dec 05, 1988
REGLAN			
ROBINS AH	EQ 5MG BASE/5ML	N18821	001 Mar 25, 1983
TABLET; ORAL			
CLOPRA			
QUANTUM PHARMICS	EQ 5MG BASE	N72384	001 Jun 02, 1988
	EQ 10MG BASE	N70294	001 Jul 29, 1985
CLOPRA-"YELLOW"			
QUANTUM PHARMICS	EQ 10MG BASE	N70632	001 Oct 28, 1985
MAXOLON			
KING PHARMS	EQ 10MG BASE	N70106	001 Mar 04, 1986
METOCLOPRAMIDE HYDROCHLORIDE			
CLONMEL HLTHCARE	EQ 10MG BASE	N72639	001 May 09, 1991

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 175 (of 263)

METOCLOPRAMIDE HYDROCHLORIDE

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

HALSEY	EQ 10MG BASE	N70906	001	Oct 28, 1986
INTERPHARM	EQ 10MG BASE	N71213	001	Sep 24, 1986
MUTUAL PHARM	EQ 5MG BASE	N71536	002	Jan 16, 1997
	EQ 10MG BASE	N70660	001	Feb 10, 1987
	EQ 10MG BASE	N71536	001	Apr 28, 1993
PAR PHARM	EQ 10MG BASE	N70342	001	Mar 25, 1986
SANDOZ	EQ 5MG BASE	N72436	001	Jun 22, 1989
	EQ 10MG BASE	N70850	001	Feb 03, 1987
SCHERING	EQ 10MG BASE	N70598	001	Feb 02, 1987
SUPERPHARM	EQ 10MG BASE	N70926	001	Jun 26, 1987
USL PHARMA	EQ 10MG BASE	N70339	001	Jul 29, 1985
WATSON LABS	EQ 10MG BASE	N70363	001	Mar 02, 1987
	EQ 10MG BASE	N70453	001	Jun 06, 1986
	EQ 10MG BASE	N70645	001	May 11, 1987

METOCURINE IODIDE

INJECTABLE; INJECTION

METUBINE IODIDE

LILLY	2MG/ML	N06632	003	
-------	--------	--------	-----	--

METOLAZONE

TABLET; ORAL

DIULO

GD SEARLE LLC	2.5MG	N18535	001	
	5MG	N18535	002	
	10MG	N18535	003	

MYKROX

UCB	0.5MG	N19532	001	Oct 30, 1987
-----	-------	--------	-----	--------------

METOPROLOL FUMARATE

TABLET, EXTENDED RELEASE; ORAL

LOPRESSOR

NOVARTIS	EQ 100MG TARTRATE	N19786	001	Dec 27, 1989
	EQ 200MG TARTRATE	N19786	002	Dec 27, 1989
	EQ 300MG TARTRATE	N19786	003	Dec 27, 1989
	EQ 400MG TARTRATE	N19786	004	Dec 27, 1989

METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE

PUREPAC PHARM	50MG	N74380	001	Jul 29, 1994
	100MG	N74380	002	Jul 29, 1994

METRIZAMIDE

INJECTABLE; INJECTION

AMIPAQUE

GE HEALTHCARE	2.5GM/VIAL	N17982	003	Sep 12, 1983
	3.75GM/VIAL	N17982	001	
	6.75GM/VIAL	N17982	002	
	13.5GM/VIAL	N17982	004	Sep 12, 1983

METRONIDAZOLE

CAPSULE; ORAL

METRONIDAZOLE

ABLE	375MG	N76505	001	Nov 13, 2003
------	-------	--------	-----	--------------

INJECTABLE; INJECTION

METRO I.V.

B BRAUN	500MG/100ML	N18674	001	Aug 31, 1982
---------	-------------	--------	-----	--------------

METRONIDAZOLE

ABBOTT	500MG/100ML	N18889	001	Nov 18, 1983
AM PHARM PARTNERS	500MG/100ML	N70071	001	Dec 03, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 176 (of 263)

METRONIDAZOLE

INJECTABLE; INJECTION

METRONIDAZOLE

ELKINS SINN	500MG/100ML	N18907	001	Mar 30, 1984
INTL MEDICATION	500MG/100ML	N70004	001	May 08, 1985
STERIS	500MG/100ML	N70042	001	Dec 20, 1984
	500MG/100ML	N70170	001	Apr 01, 1986

TABLET; ORAL

METROMIDOL

LABS AF	250MG	N74523	001	Oct 24, 1996
	500MG	N74523	002	Oct 24, 1996

METRONIDAZOLE

ABLE	250MG	N76519	001	Jun 27, 2003
	500MG	N76519	002	Jun 27, 2003
HALSEY	250MG	N70021	001	Apr 02, 1985
	500MG	N70593	001	Feb 27, 1986
MUTUAL PHARM	250MG	N18818	001	Feb 16, 1983
	500MG	N18818	002	Feb 16, 1983
SUPERPHARM	250MG	N70008	001	Dec 11, 1984
	500MG	N70009	001	Dec 11, 1984
WATSON LABS	250MG	N18599	001	Sep 17, 1982
	500MG	N18599	002	Feb 13, 1984

PROTOSTAT

ORTHO MCNEIL PHARM	250MG	N18871	001	Mar 02, 1983
	500MG	N18871	002	Mar 02, 1983

SATRIC

SAVAGE LABS	250MG	N70029	001	Mar 19, 1985
	500MG	N70731	001	Jun 08, 1987

TABLET, EXTENDED RELEASE; ORAL

METRONIDAZOLE

ABLE	750MG	N76462	001	Jun 25, 2003
------	-------	--------	-----	--------------

METRONIDAZOLE HYDROCHLORIDE

INJECTABLE; INJECTION

METRONIDAZOLE HYDROCHLORIDE

AM PHARM PARTNERS	EQ 500MG BASE/VIAL	N70295	001	Oct 15, 1985
-------------------	--------------------	--------	-----	--------------

METYRAPONE

TABLET; ORAL

METOPIRONE

NOVARTIS	250MG	N12911	001	
----------	-------	--------	-----	--

MEZLOCILLIN SODIUM MONOHYDRATE

INJECTABLE; INJECTION

MEZLIN

BAYER PHARMS	EQ 1GM BASE/VIAL	N50549	001	
	EQ 1GM BASE/VIAL	N62333	001	
	EQ 1GM BASE/VIAL	N62372	005	Jan 13, 1983
	EQ 2GM BASE/VIAL	N50549	002	
	EQ 2GM BASE/VIAL	N62333	002	
	EQ 2GM BASE/VIAL	N62372	001	May 13, 1982
	EQ 3GM BASE/VIAL	N50549	003	
	EQ 3GM BASE/VIAL	N62333	003	
	EQ 3GM BASE/VIAL	N62372	002	May 13, 1982
	EQ 3GM BASE/VIAL	N62697	001	Jan 22, 1987
	EQ 4GM BASE/VIAL	N50549	004	
	EQ 4GM BASE/VIAL	N62333	004	
	EQ 4GM BASE/VIAL	N62372	003	May 13, 1982
	EQ 4GM BASE/VIAL	N62697	002	Jan 22, 1987
	EQ 20GM BASE/VIAL	N50549	005	Mar 02, 1988
	EQ 20GM BASE/VIAL	N62372	004	Mar 02, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 177 (of 263)

MICONAZOLEINJECTABLE; INJECTION
MONISTAT

JANSSEN PHARMA 10MG/ML N18040 001

MICONAZOLE NITRATELOTION; TOPICAL
MONISTAT-DERM

JOHNSON AND JOHNSON 2% N17739 001

TAMPON; VAGINAL
MONISTAT 5

PERSONAL PRODS 100MG N18592 001 Oct 27, 1989

MIDAZOLAM HYDROCHLORIDEINJECTABLE; INJECTION
MIDAZOLAM HYDROCHLORIDE

APOTHECON EQ 1MG BASE/ML N75620 001 Nov 01, 2000

EQ 5MG BASE/ML N75620 002 Nov 01, 2000

EQ 5MG BASE/ML N75641 001 Oct 19, 2000

ASTRAZENECA EQ 5MG BASE/ML N75263 001 Jun 26, 2000

BEDFORD EQ 5MG BASE/ML N75249 001 Jun 23, 2000

BEN VENUE EQ 5MG BASE/ML N75455 001 Jun 20, 2000

VERSED

HLR EQ 1MG BASE/ML N18654 002 May 26, 1987

EQ 5MG BASE/ML N18654 001 Dec 20, 1985

SYRUP; ORAL

VERSED

ROCHE EQ 2MG BASE/ML N20942 001 Oct 15, 1998

MILRINONE LACTATE

INJECTABLE; INJECTION

PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER

SANOFI SYNTHELABO EQ 10MG BASE/100ML N20343 001 Aug 09, 1994

EQ 15MG BASE/100ML N20343 002 Aug 09, 1994

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

DYNACIN

MEDICIS EQ 50MG BASE N63066 001 Aug 14, 1990

MINOCIN

LEDERLE EQ 50MG BASE N50315 002

EQ 100MG BASE N50315 001

WYETH PHARMS INC EQ 75MG BASE N50649 003 Feb 12, 2001

INJECTABLE; INJECTION

MINOCIN

LEDERLE EQ 100MG BASE/VIAL N62139 001

WYETH PHARMS INC EQ 100MG BASE/VIAL N50444 001

SUSPENSION; ORAL

MINOCIN

LEDERLE EQ 50MG BASE/5ML N50445 001

TABLET; ORAL

MINOCYCLINE HYDROCHLORIDE

LEDERLE EQ 50MG BASE N50451 003 Aug 10, 1982

EQ 100MG BASE N50451 002 Aug 10, 1982

MINOXIDIL

TABLET; ORAL

MINODYL

QUANTUM PHARMICS 2.5MG N72153 001 Jul 13, 1988

10MG N71534 001 Mar 19, 1987

MINOXIDIL

ROYCE LABS 2.5MG N71799 001 Nov 10, 1987

10MG N71796 001 Nov 10, 1987

USL PHARMA 2.5MG N71537 001 Dec 16, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 178 (of 263)

MITOMYCININJECTABLE; INJECTION
MITOMYCIN

MAYNE PHARMA USA 20MG/VIAL

N64106 001 Nov 29, 1995

MUTAMYCIN

BRISTOL 5MG/VIAL
20MG/VIAL

N50450 001

N50450 002

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER

ABBOTT EQ 0.5MG BASE/ML

N20098 002 Jan 22, 1992

MOLINDONE HYDROCHLORIDE

CAPSULE; ORAL

MOBAN

ENDO PHARMS 5MG
10MG
25MG

N17111 001

N17111 002

N17111 003

CONCENTRATE; ORAL

MOBAN

ENDO PHARMS 20MG/ML

N17938 001

TABLET; ORAL

MOBAN

ENDO PHARMS 100MG

N17111 008

MONOCTANOIN

LIQUID; PERFUSION, BILIARY

MOCTANIN

ETHITEK 100%

N19368 001 Oct 29, 1985

MORPHINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

CLONMEL HLTHCARE 15MG

N75407 001 Jan 28, 2000

MOXALACTAM DISODIUM

INJECTABLE; INJECTION

MOXAM

LILLY EQ 250MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 10GM BASE/VIAL

N50550 001

N50550 002

N50550 003

N50550 004

N50550 008

NABILONE

CAPSULE; ORAL

CESAMET

VALEANT 1MG

N18677 001 Dec 26, 1985

NAFCILLIN SODIUM

CAPSULE; ORAL

UNIPEN

WYETH AYERST EQ 250MG BASE

N50111 001

FOR SOLUTION; ORAL

UNIPEN

WYETH AYERST EQ 250MG BASE/5ML

N50199 001

INJECTABLE; INJECTION

NAFCILLIN SODIUM

APOTHECON EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 4GM BASE/VIAL
MARSAM PHARMS LLC EQ 500MG BASE/VIAL

N61984 001

N61984 002

N61984 003

N61984 005

N62844 001 Oct 26, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 179 (of 263)

NAFCILLIN SODIUM

INJECTABLE; INJECTION

NAFCILLIN SODIUM

MARSAM PHARMS LLC

EQ 1GM BASE/VIAL

N62844 002 Oct 26, 1988

EQ 1.5GM BASE/VIAL

N62844 003 Oct 26, 1988

EQ 2GM BASE/VIAL

N62844 004 Oct 26, 1988

EQ 4GM BASE/VIAL

N62844 005 Oct 26, 1988

EQ 10GM BASE/VIAL

N63008 001 Sep 29, 1988

NALLPEN

GLAXOSMITHKLINE

EQ 500MG BASE/VIAL

N61999 001

EQ 1GM BASE/VIAL

N61999 002

EQ 1GM BASE/VIAL

N62755 001 Dec 19, 1986

EQ 2GM BASE/VIAL

N61999 003

EQ 2GM BASE/VIAL

N62755 002 Dec 19, 1986

EQ 10GM BASE/VIAL

N61999 004

UNIPEN

WYETH AYERST

EQ 500MG BASE/VIAL

N50320 001

EQ 500MG BASE/VIAL

N62717 001 Dec 16, 1986

EQ 1GM BASE/VIAL

N62717 002 Dec 16, 1986

EQ 2GM BASE/VIAL

N50320 003

EQ 2GM BASE/VIAL

N62717 004 Dec 16, 1986

EQ 4GM BASE/VIAL

N50320 004

EQ 10GM BASE/VIAL

N50320 005

EQ 20GM BASE/VIAL

N50320 006

UNIPEN IN PLASTIC CONTAINER

WYETH AYERST

EQ 1GM BASE/VIAL

N50320 002

TABLET; ORAL

UNIPEN

WYETH AYERST

EQ 500MG BASE

N50462 001

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE

AM PHARM PARTNERS

10MG/ML

N70751 001 Jul 02, 1986

20MG/ML

N70752 001 Sep 24, 1986

NALBUPHINE HYDROCHLORIDE

ABBOTT

1.5MG/ML

N20200 001 Mar 12, 1993

20MG/ML

N70917 001 Feb 03, 1989

ASTRAZENECA

10MG/ML

N72070 001 Apr 10, 1989

10MG/ML

N72071 001 Apr 10, 1989

10MG/ML

N72072 001 Apr 10, 1989

20MG/ML

N72073 001 Apr 10, 1989

20MG/ML

N72074 001 Apr 10, 1989

20MG/ML

N72075 001 Apr 10, 1989

NALIDIXIC ACID

SUSPENSION; ORAL

NEGGRAM

SANOFI SYNTHELABO

250MG/5ML

N17430 001

TABLET; ORAL

NALIDIXIC ACID

MUTUAL PHARM

250MG

N70270 001 Jun 29, 1988

500MG

N70271 001 Jun 29, 1988

1GM

N70272 001 Jun 29, 1988

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE

BAXTER HLTHCARE

0.4MG/ML

N70298 001 Sep 24, 1986

0.4MG/ML

N70299 001 Sep 24, 1986

0.4MG/ML

N70496 001 Sep 24, 1986

WYETH AYERST

0.02MG/ML

N70188 001 Sep 24, 1986

0.02MG/ML

N70189 001 Sep 24, 1986

0.4MG/ML

N70190 001 Sep 24, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 180 (of 263)

NALOXONE HYDROCHLORIDEINJECTABLE; INJECTION
NALOXONE

WYETH AYERST	0.4MG/ML	N70191	001	Sep 24, 1986
NALOXONE HYDROCHLORIDE				
AM PHARM PARTNERS	0.02MG/ML	N70648	001	Nov 17, 1986
	0.02MG/ML	N70661	001	Nov 17, 1986
	0.4MG/ML	N70649	001	Nov 17, 1986
	1MG/ML	N71604	001	Dec 16, 1988
ASTRAZENECA	0.02MG/ML	N72081	001	Apr 11, 1989
	0.02MG/ML	N72082	001	Apr 11, 1989
	0.02MG/ML	N72083	001	Apr 11, 1989
	0.02MG/ML	N72084	001	Apr 11, 1989
	0.02MG/ML	N72085	001	Apr 11, 1989
	0.4MG/ML	N72086	001	Apr 11, 1989
	0.4MG/ML	N72087	001	Apr 11, 1989
	0.4MG/ML	N72088	001	Apr 11, 1989
	0.4MG/ML	N72089	001	Apr 11, 1989
	0.4MG/ML	N72090	001	Apr 11, 1989
	1MG/ML	N72091	001	Apr 11, 1989
	1MG/ML	N72092	001	Apr 11, 1989
	1MG/ML	N72093	001	Apr 11, 1989
BAXTER HLTHCARE	0.02MG/ML	N71272	001	May 24, 1988
	1MG/ML	N71273	001	May 24, 1988
	1MG/ML	N71274	001	May 24, 1988
	1MG/ML	N71287	001	May 24, 1988
INTL MEDICATION	0.4MG/ML	N70417	001	Sep 24, 1986
	1MG/ML	N72115	001	Apr 27, 1988
MARSAM PHARMS LLC	0.4MG/ML	N71811	001	Jul 19, 1988
SMITH AND NEPHEW	0.02MG/ML	N71671	001	Nov 17, 1987
	0.4MG/ML	N71681	001	Nov 17, 1987
	0.4MG/ML	N71682	001	Nov 17, 1987
SOLOPAK	0.02MG/ML	N71672	001	Nov 17, 1987
	0.4MG/ML	N71683	001	Nov 17, 1987
STERIS	0.4MG/ML	N71339	001	Nov 18, 1987
NARCAN				
BRISTOL MYERS SQUIBB	0.4MG/ML	N71083	001	Jul 28, 1988
	1MG/ML	N71084	001	Jul 28, 1988
	1MG/ML	N71311	001	Jul 28, 1988

NANDROLONE DECANOATEINJECTABLE; INJECTION
DECA-DURABOLIN

ORGANON USA INC	50MG/ML	N13132	001	Jun 12, 1986
	100MG/ML	N13132	002	Jun 12, 1986
	200MG/ML	N13132	003	Jun 12, 1986
NANDROLONE DECANOATE				
AKORN	100MG/ML	N87519	001	Sep 28, 1983
AM PHARM PARTNERS	100MG/ML	N88290	001	Oct 03, 1983
	200MG/ML	N88317	001	Oct 14, 1983
STERIS	50MG/ML	N87598	001	Oct 06, 1983
	50MG/ML	N88554	001	Feb 10, 1986
	100MG/ML	N87599	001	Oct 06, 1983

NANDROLONE PHENPROPIONATEINJECTABLE; INJECTION
DURABOLIN

ORGANON USA INC	25MG/ML	N11891	001	
	50MG/ML	N11891	002	
NANDROLONE PHENPROPIONATE				
STERIS	25MG/ML	N86386	001	Jun 17, 1983
	50MG/ML	N87488	001	Jun 17, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 181 (of 263)

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

NAFAZAIR

PHARMAFAIR	0.1%	N88101	001	Apr 15, 1983
------------	------	--------	-----	--------------

NAPHCON FORTE

ALCON	0.1%	N80229	001	
-------	------	--------	-----	--

OPCON

BAUSCH AND LOMB	0.1%	N87506	001	
-----------------	------	--------	-----	--

NAPROXEN

TABLET; ORAL

NAPROXEN

BAXTER HLTHCARE	250MG	N74105	001	Dec 21, 1993
-----------------	-------	--------	-----	--------------

	375MG	N74105	002	Dec 21, 1993
--	-------	--------	-----	--------------

	500MG	N74105	003	Dec 21, 1993
--	-------	--------	-----	--------------

HAMILTON PHARMS	250MG	N74110	001	Oct 30, 1992
-----------------	-------	--------	-----	--------------

	375MG	N74110	002	Oct 30, 1992
--	-------	--------	-----	--------------

	500MG	N74110	003	Oct 30, 1992
--	-------	--------	-----	--------------

IVAX PHARMS	250MG	N74111	001	Feb 28, 1995
-------------	-------	--------	-----	--------------

	375MG	N74111	002	Feb 28, 1995
--	-------	--------	-----	--------------

	500MG	N74111	003	Feb 28, 1995
--	-------	--------	-----	--------------

PUREPAC PHARM	250MG	N74263	001	Dec 21, 1993
---------------	-------	--------	-----	--------------

	375MG	N74263	002	Dec 21, 1993
--	-------	--------	-----	--------------

	500MG	N74263	003	Dec 21, 1993
--	-------	--------	-----	--------------

ROXANE	250MG	N74211	001	Feb 28, 1994
--------	-------	--------	-----	--------------

	375MG	N74211	002	Feb 28, 1994
--	-------	--------	-----	--------------

	500MG	N74211	003	Feb 28, 1994
--	-------	--------	-----	--------------

NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

ABLE	EQ 250MG BASE	N76544	001	Aug 22, 2003
------	---------------	--------	-----	--------------

	EQ 500MG BASE	N76544	002	Aug 22, 2003
--	---------------	--------	-----	--------------

HAMILTON PHARMS	EQ 250MG BASE	N74106	001	Aug 31, 1993
-----------------	---------------	--------	-----	--------------

	EQ 500MG BASE	N74106	002	Aug 31, 1993
--	---------------	--------	-----	--------------

IVAX PHARMS	EQ 250MG BASE	N74230	001	Mar 14, 1995
-------------	---------------	--------	-----	--------------

	EQ 500MG BASE	N74230	002	Mar 14, 1995
--	---------------	--------	-----	--------------

PUREPAC PHARM	EQ 250MG BASE	N74319	001	Mar 20, 1995
---------------	---------------	--------	-----	--------------

	EQ 500MG BASE	N74319	002	Mar 20, 1995
--	---------------	--------	-----	--------------

ROXANE	EQ 250MG BASE	N74257	001	Dec 21, 1993
--------	---------------	--------	-----	--------------

	EQ 500MG BASE	N74257	002	Dec 21, 1993
--	---------------	--------	-----	--------------

TABLET, EXTENDED RELEASE; ORAL

NAPRELAN

STAT TRADE	EQ 750MG BASE	N20353	003	Jan 05, 1996
------------	---------------	--------	-----	--------------

NEDOCROMIL SODIUM

SOLUTION; INHALATION

TILADE

AVENTIS	0.5%	N20750	001	Oct 01, 1997
---------	------	--------	-----	--------------

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

ROXANE	50MG	N76196	001	Sep 16, 2003
--------	------	--------	-----	--------------

	100MG	N76196	002	Sep 16, 2003
--	-------	--------	-----	--------------

	150MG	N76196	003	Sep 16, 2003
--	-------	--------	-----	--------------

	200MG	N76196	004	Sep 16, 2003
--	-------	--------	-----	--------------

	250MG	N76196	005	Sep 16, 2003
--	-------	--------	-----	--------------

SERZONE

BRISTOL MYERS SQUIBB	50MG	N20152	001	Dec 22, 1994
----------------------	------	--------	-----	--------------

	100MG	N20152	002	Dec 22, 1994
--	-------	--------	-----	--------------

	150MG	N20152	003	Dec 22, 1994
--	-------	--------	-----	--------------

	200MG	N20152	004	Dec 22, 1994
--	-------	--------	-----	--------------

	250MG	N20152	005	Dec 22, 1994
--	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 182 (of 263)

NEFAZODONE HYDROCHLORIDETABLET; ORAL
SERZONE

BRISTOL MYERS SQUIBB 300MG

N20152 006 Dec 22, 1994

NEOMYCIN SULFATESOLUTION; ORAL
MYCIFRADIN

PHARMACIA AND UPJOHN EQ 87.5MG BASE/5ML

N50285 001

TABLET; ORAL
MYCIFRADIN

PHARMACIA AND UPJOHN EQ 350MG BASE

N60520 001

NEOBIOITIC

PFIZER EQ 350MG BASE

N60475 001

NEOMYCIN SULFATE

BRISTOL MYERS SQUIBB EQ 350MG BASE

N60365 001

LANNETT EQ 350MG BASE

N60607 001

LILLY EQ 350MG BASE

N60385 001

ROXANE EQ 350MG BASE

N62173 001

SANDOZ EQ 350MG BASE

N61586 001

NEOMYCIN SULFATE; POLYMYXIN B SULFATE

CREAM; TOPICAL

NEOSPORIN

GLAXOSMITHKLINE EQ 3.5MG BASE/GM;10,000 UNITS/GM

N50176 002 Jan 14, 1985

OINTMENT; OPHTHALMIC

STATROL

ALCON EQ 3.5MG BASE/GM;10,000 UNITS/GM

N50344 002

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATES

STERIS EQ 40MG BASE/ML;200,000 UNITS/ML

N62664 001 Apr 08, 1986

SOLUTION/DROPS; OPHTHALMIC

STATROL

ALCON EQ 3.5MG BASE/ML;16,250 UNITS/ML

N50456 001

EQ 3.5MG BASE/ML;16,250 UNITS/ML

N62339 001 Nov 30, 1984

NEOMYCIN SULFATE; PREDNISOLONE ACETATE

OINTMENT; OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN EQ 3.5MG BASE/GM;0.25%

N61039 002

EQ 3.5MG BASE/GM;0.5%

N61039 001

SUSPENSION/DROPS; OPHTHALMIC

NEO-DELTA-CORTEF

PHARMACIA AND UPJOHN EQ 3.5MG BASE/ML;0.25%

N61037 001

NEOMYCIN SULFATE; PREDNISOLONE SODIUM PHOSPHATE

OINTMENT; OPHTHALMIC

NEO-HYDELTRASOL

MERCK EQ 3.5MG BASE/GM;EQ 0.25% PHOSPHATE

N50378 001

NEOMYCIN SULFATE; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYTRES A

SAVAGE LABS EQ 3.5MG BASE/GM;0.1%

N62598 001 Jul 21, 1986

NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA EQ 3.5MG BASE/GM;0.1%

N62600 001 Jul 21, 1986

PHARMADERM EQ 3.5MG BASE/GM;0.1%

N62595 001 Jul 21, 1986

OINTMENT; TOPICAL

MYTRES A

SAVAGE LABS EQ 3.5MG BASE/GM;0.1%

N62609 001 May 23, 1986

NEOMYCIN SULFATE-TRIAMCINOLONE ACETONIDE

FOUGERA EQ 3.5MG BASE/GM;0.1%

N62608 001 May 23, 1986

PHARMADERM EQ 3.5MG BASE/GM;0.1%

N62607 001 May 23, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 183 (of 263)

NETILMICIN SULFATEINJECTABLE; INJECTION
NETROMYCIN

SCHERING

EQ 10MG BASE/ML

N50544 001

Feb 28, 1983

EQ 25MG BASE/ML

N50544 002

Feb 28, 1983

EQ 100MG BASE/ML

N50544 003

Feb 28, 1983

NIACIN

CAPSULE; ORAL

WAMPOCAP

MEDPOINTE PHARM HLC 500MG

N11073 003

TABLET; ORAL

NIACIN

EVERYLIFE 500MG

N83203 001

HALSEY 500MG

N83453 001

IMPAX LABS 500MG

N83115 001

IVAX PHARMS 500MG

N83180 001

MK LABS 500MG

N83525 001

PUREPAC PHARM 500MG

N83271 001

SANDOZ 500MG

N83306 001

TABLICAPS 500MG

N84237 001

WATSON LABS 500MG

N83136 001

500MG

N83305 001

500MG

N85172 001

WEST WARD 500MG

N83718 001

NICOLAR

AVENTIS 500MG

N83823 001

TABLET, EXTENDED RELEASE; ORAL

NIASPAN

KOS LIFE 375MG

N20381 001

Jul 28, 1997

NIASPAN TITRATION STARTER PACK

KOS LIFE 375MG;500MG;750MG

N20381 005

Jul 28, 1997

NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; TYROSINE

SUSPENSION; ORAL

TPN SUSPENSION

INTL MINERALS 15MG/5ML;3.75MG/5ML;600MG/5ML

N08378 003

NICLOSAMIDE

TABLET, CHEWABLE; ORAL

NICLOCIDE

BAYER PHARMS 500MG

N18669 001

May 14, 1982

NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

NICOTROL

PHARMACIA AND UPJOHN 15MG/16HR

N20536 001

Jul 03, 1996

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICOTINE POLACRILEX

WATSON LABS EQ 2MG BASE

N76568 001

Jul 29, 2004

EQ 4MG BASE

N76569 002

Jul 29, 2004

NIFEDIPINE

CAPSULE; ORAL

ADALAT

BAYER PHARMS 10MG

N19478 001

Nov 27, 1985

20MG

N19478 002

Sep 17, 1986

NIFEDIPINE

CHASE LABS NJ 10MG

N72409 001

Jul 04, 1990

20MG

N73421 001

Jun 19, 1991

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 184 (of 263)

NIFEDIPINE

TABLET, EXTENDED RELEASE; ORAL

NIFEDIPINE

MYLAN

30MG

N75108 001

Dec 17, 1999

NILUTAMIDE

TABLET; ORAL

NILANDRON

AVENTIS PHARMS

50MG

N20169 001

Sep 19, 1996

NITROFURANTOIN

CAPSULE; ORAL

NITROFURANTOIN

WATSON LABS

50MG

N84326 001

100MG

N84326 002

TABLET; ORAL

FURADANTIN

PROCTER AND GAMBLE

50MG

N08693 001

100MG

N08693 002

FURALAN

LANNETT

50MG

N80017 001

100MG

N80017 002

NITROFURANTOIN

ELKINS SINN

50MG

N80003 001

100MG

N80003 002

IVAX PHARMS

50MG

N80078 002

100MG

N80078 001

SANDOZ

50MG

N80043 001

100MG

N80043 002

WATSON LABS

50MG

N80447 001

50MG

N85797 001

100MG

N80447 002

100MG

N85796 001

WHITEWORTH TOWN PLSN

100MG

N84085 002

NITROFURANTOIN SODIUM

INJECTABLE; INJECTION

IVADANTIN

PROCTER AND GAMBLE

EQ 180MG BASE/VIAL

N12402 001

NITROFURANTOIN, MACROCRYSTALLINE

CAPSULE; ORAL

NITROFURANTOIN

MYLAN

100MG

N74967 002

Jul 09, 1997

NITROFURANTOIN MACROCRYSTALLINE

WATSON LABS

50MG

N70248 001

Jun 24, 1988

100MG

N70249 001

Jun 24, 1988

NITROFURAZONE

CREAM; TOPICAL

FURACIN

SHIRE

0.2%

N83789 001

DRESSING; TOPICAL

ACTIN-N

SHERWOOD MEDCL

0.2%

N17343 001

OINTMENT; TOPICAL

FURACIN

SHIRE

0.2%

N05795 001

NITROFURAZONE

AMBIX

0.2%

N86077 001

CLAY PARK

0.2%

N84968 001

LANNETT

0.2%

N84393 001

WENDT

0.2%

N86766 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 185 (of 263)

NITROFURAZONE

POWDER; TOPICAL			
FURACIN			
SHIRE	0.2%	N83791	001
SOLUTION; TOPICAL			
NITROFURAZONE			
CLAY PARK	0.2%	N85130	001
WENDT	0.2%	N87081	001

NITROGLYCERIN

AEROSOL; SUBLINGUAL				
NITROLINGUAL				
POHL BOSKAMP	0.4MG/SPRAY	N18705	001	Oct 31, 1985
FILM, EXTENDED RELEASE; TRANSDERMAL				
TRANSDERM-NITRO				
NOVARTIS	0.1MG/HR	N20144	001	Feb 27, 1996
	0.2MG/HR	N20144	002	Feb 27, 1996
	0.4MG/HR	N20144	003	Feb 27, 1996
	0.6MG/HR	N20144	004	Feb 27, 1996
	0.8MG/HR	N20144	005	Feb 27, 1996
INJECTABLE; INJECTION				
NITRO IV				
POHL BOSKAMP	5MG/ML	N18672	002	Aug 30, 1983
NITRO-BID				
AVENTIS PHARMS	5MG/ML	N18621	001	Jan 05, 1982
	10MG/ML	N71159	001	Feb 28, 1990
NITROGLYCERIN				
AM PHARM PARTNERS	5MG/ML	N70077	001	Dec 13, 1985
	5MG/ML	N71203	001	May 08, 1987
INTL MEDICATION	5MG/ML	N70026	001	Sep 10, 1985
LUITPOLD	5MG/ML	N71492	001	May 24, 1988
SMITH AND NEPHEW	5MG/ML	N70633	001	Jun 19, 1986
	5MG/ML	N70634	001	Jun 19, 1986
NITROGLYCERIN IN DEXTROSE 5%				
HOSPIRA	0.1MG/ML	N74083	001	Oct 26, 1994
NITROL				
RORER	0.8MG/ML	N18774	001	Jan 19, 1983
NITRONAL				
POHL BOSKAMP	1MG/ML	N18672	001	Aug 30, 1983
NITROSTAT				
PARKE DAVIS	0.8MG/ML	N18588	001	
	5MG/ML	N18588	002	Dec 23, 1983
	5MG/ML	N70863	001	Jan 08, 1987
	10MG/ML	N70871	001	Jan 08, 1987
	10MG/ML	N70872	001	Jan 08, 1987
TRIDIL				
MAYNE PHARMA USA	0.5MG/ML	N18537	002	Jun 16, 1983
	5MG/ML	N18537	001	

NONOXYNOL-9

AEROSOL; VAGINAL				
DELFIN				
PERSONAL PRODS	12.5%	N14349	002	
SPONGE; VAGINAL				
TODAY				
ALLENDAL PHARMS	1GM	N18683	001	Apr 01, 1983

NOREPINEPHRINE BITARTRATE; PROCAINE HYDROCHLORIDE; PROPOXYCAINE HYDROCHLORIDE

INJECTABLE; INJECTION				
RAVOCAINE AND NOVOCAIN W/ LEVOPHED				
EASTMAN KODAK	EQ 0.033MG BASE/ML; 2%; 0.4%	N08592	003	

DISCONTINUED DRUG PRODUCT LIST

6 - 186 (of 263)

NORETHINDRONE

TABLET; ORAL NORLUTIN			
PARKE DAVIS	5MG		N10895 002

NORETHINDRONE ACETATE

TABLET; ORAL NORLUTATE			
PARKE DAVIS	5MG		N12184 002

NORFLOXACIN

SOLUTION/DROPS; OPHTHALMIC CHIBROXIN			
MERCK	0.3%		N19757 001 Jun 17, 1991

NORGESTREL

TABLET; ORAL OVRETTE			
WYETH PHARMS INC	0.075MG		N17031 001

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL AVENTYL HYDROCHLORIDE			
LILLY	EQ 10MG BASE		N14684 001
	EQ 25MG BASE		N14684 002

NOVOBIOCIN SODIUM

CAPSULE; ORAL ALBAMYCIN			
PHARMACIA AND UPJOHN	EQ 250MG BASE		N50339 001

NYSTATIN

CREAM; TOPICAL CANDEX			
BAYER PHARMS	100,000 UNITS/GM		N61810 001
MYKINAC			
ALPHARMA US PHARMS	100,000 UNITS/GM		N62387 001 Jul 29, 1982
NILSTAT			
LEDERLE	100,000 UNITS/GM		N61445 001
NYSTATIN			
TEVA	100,000 UNITS/GM		N61966 001
LOTION; TOPICAL CANDEX			
BAYER PHARMS	100,000 UNITS/ML		N50233 001
OINTMENT; TOPICAL MYCOSTATIN			
WESTWOOD SQUIBB	100,000 UNITS/GM		N60571 001
MYKINAC			
ALPHARMA US PHARMS	100,000 UNITS/GM		N62731 001 Sep 22, 1986
NILSTAT			
LEDERLE	100,000 UNITS/GM		N61444 001
PASTILLE; ORAL MYCOSTATIN			
BRISTOL MYERS SQUIBB	200,000 UNITS		N50619 001 Apr 09, 1987
POWDER; ORAL BARSTATIN 100			
BARLAN	100%		N62489 001 Apr 27, 1988
NILSTAT			
CLONMEL HLTHCARE	100%		N50576 001 Dec 22, 1983
SUPPOSITORY; VAGINAL NYSERT			
PROCTER AND GAMBLE	100,000 UNITS		N50478 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 187 (of 263)

NYSTATIN

SUSPENSION; ORAL

MYCOSTATIN

APOTHECON

100,000 UNITS/ML

N61533 001

NYSTATIN

ALPHARMA US PHARMS

100,000 UNITS/ML

N62571 001

Oct 29, 1985

MORTON GROVE

100,000 UNITS/ML

N62835 001

Nov 19, 1987

PHARMADERM

100,000 UNITS/ML

N62518 001

Jul 06, 1984

PHARMAFAIR

100,000 UNITS/ML

N62541 001

Jan 16, 1985

ROXANE

100,000 UNITS/ML

N62832 001

Dec 27, 1991

TARO

100,000 UNITS/ML

N62876 001

Feb 29, 1988

TEVA

100,000 UNITS/ML

N62670 001

Jun 18, 1987

100,000 UNITS/ML

N62776 001

Dec 17, 1987

NYSTEX

SAVAGE LABS

100,000 UNITS/ML

N62519 001

Jul 06, 1984

TABLET; ORAL

NILSTAT

LEDERLE

500,000 UNITS

N61151 001

NYSTATIN

QUANTUM PHARMICS

500,000 UNITS

N62525 001

Oct 29, 1984

SANDOZ

500,000 UNITS

N62065 001

USL PHARMA

500,000 UNITS

N62524 001

Nov 26, 1985

WATSON LABS

500,000 UNITS

N62402 001

Dec 16, 1982

TABLET; VAGINAL

KOROSTATIN

HOLLAND RANTOS

100,000 UNITS

N61718 001

MYCOSTATIN

BRISTOL MYERS SQUIBB

100,000 UNITS

N60577 001

NILSTAT

LEDERLE

100,000 UNITS

N61325 001

NYSTATIN

FOUGERA

100,000 UNITS

N62459 001

Nov 09, 1983

PHARMADERM

100,000 UNITS

N62460 001

Nov 09, 1983

QUANTUM PHARMICS

100,000 UNITS

N62509 001

Apr 03, 1984

SANDOZ

100,000 UNITS

N61965 001

TEVA

100,000 UNITS

N62502 001

Dec 23, 1983

WATSON LABS

100,000 UNITS

N62176 001

NYSTATIN; TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

MYCOLOG-II

APOTHECON

100,000 UNITS/GM;0.1%

N60576 002

May 01, 1985

100,000 UNITS/GM;0.1%

N62606 001

May 15, 1985

MYCO-TRIACET II

TEVA

100,000 UNITS/GM;0.1%

N61954 002

Sep 20, 1985

MYTREX F

SAVAGE LABS

100,000 UNITS/GM;0.1%

N62597 001

Oct 08, 1985

NYSTATIN AND TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS

100,000 UNITS/GM;0.1%

N63010 001

Dec 20, 1988

CLAY PARK

100,000 UNITS/GM;0.1%

N62186 002

Jun 06, 1985

PHARMAFAIR

100,000 UNITS/GM;0.1%

N62657 001

Jul 30, 1986

TARO

100,000 UNITS/GM;0.1%

N62347 001

Mar 30, 1987

NYSTATIN-TRIAMCINOLONE ACETONIDE

PHARMADERM

100,000 UNITS/GM;0.1%

N62596 001

Oct 08, 1985

OINTMENT; TOPICAL

MYCOLOG-II

APOTHECON

100,000 UNITS/GM;0.1%

N60572 001

Jun 28, 1985

MYCO-TRIACET II

TEVA

100,000 UNITS/GM;0.1%

N62045 002

Nov 26, 1985

MYTREX F

SAVAGE LABS

100,000 UNITS/GM;0.1%

N62601 001

Oct 09, 1985

NYSTATIN AND TRIAMCINOLONE ACETONIDE

CLAY PARK

100,000 UNITS/GM;0.1%

N62280 002

Oct 10, 1985

PHARMAFAIR

100,000 UNITS/GM;0.1%

N62656 001

Jul 30, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 188 (of 263)

NYSTATIN; TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

NYSTATIN-TRIAMCINOLONE ACETONIDE

PHARMADERM

100,000 UNITS/GM;0.1%

N62603 001

Oct 09, 1985

OFLOXACIN

INJECTABLE; INJECTION

FLOXIN

ORTHO MCNEIL PHARM

20MG/ML

N20087 002

Mar 31, 1992

40MG/ML

N20087 003

Mar 31, 1992

FLOXIN IN DEXTROSE 5%

ORTHO MCNEIL PHARM

400MG/100ML

N20087 001

Mar 31, 1992

FLOXIN IN DEXTROSE 5% IN PLASTIC CONTAINER

ORTHO MCNEIL PHARM

4MG/ML

N20087 004

Mar 31, 1992

400MG/100ML

N20087 005

Mar 31, 1992

ORPHENADRINE CITRATE

TABLET, EXTENDED RELEASE; ORAL

NORFLEX

3M

100MG

N12157 001

ORPHENADRINE CITRATE

ASCOT

100MG

N88067 001

Apr 06, 1983

SANDOZ

100MG

N85046 001

WATSON LABS

100MG

N84303 001

ORPHENADRINE HYDROCHLORIDE

TABLET; ORAL

DISIPAL

3M

50MG

N10653 001

OXACILLIN SODIUM

CAPSULE; ORAL

BACTOCILL

GLAXOSMITHKLINE

EQ 250MG BASE

N62241 001

EQ 500MG BASE

N62241 002

OXACILLIN SODIUM

APOTHECON

EQ 250MG BASE

N61450 002

EQ 500MG BASE

N61450 001

PROSTAPHLIN

APOTHECON

EQ 500MG BASE

N50118 002

FOR SOLUTION; ORAL

OXACILLIN SODIUM

APOTHECON

EQ 250MG BASE/5ML

N61457 001

PROSTAPHLIN

APOTHECON

EQ 250MG BASE/5ML

N50194 001

INJECTABLE; INJECTION

BACTOCILL

GLAXOSMITHKLINE

EQ 500MG BASE/VIAL

N61334 009

Mar 26, 1982

EQ 1GM BASE/VIAL

N61334 006

Mar 26, 1982

EQ 1GM BASE/VIAL

N62736 001

Dec 19, 1986

EQ 2GM BASE/VIAL

N61334 007

Mar 26, 1982

EQ 2GM BASE/VIAL

N62736 002

Dec 19, 1986

EQ 4GM BASE/VIAL

N61334 008

Mar 26, 1982

EQ 10GM BASE/VIAL

N61334 010

OXACILLIN SODIUM

APOTHECON

EQ 250MG BASE/VIAL

N50195 001

EQ 500MG BASE/VIAL

N50195 002

EQ 1GM BASE/VIAL

N50195 003

EQ 2GM BASE/VIAL

N50195 004

EQ 4GM BASE/VIAL

N50195 005

ELKINS SINN

EQ 250MG BASE/VIAL

N62711 001

Feb 03, 1989

EQ 500MG BASE/VIAL

N62711 002

Feb 03, 1989

EQ 1GM BASE/VIAL

N62711 003

Feb 03, 1989

EQ 2GM BASE/VIAL

N62711 004

Feb 03, 1989

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 189 (of 263)

OXACILLIN SODIUMINJECTABLE; INJECTION
OXACILLIN SODIUM

ELKINS SINN

EQ 4GM BASE/VIAL

N62711 005

Feb 03, 1989

EQ 10GM BASE/VIAL

N62711 006

Feb 03, 1989

IBI

EQ 125MG BASE/VIAL

N62798 003

Dec 11, 1995

EQ 250MG BASE/VIAL

N62798 004

Dec 11, 1995

EQ 500MG BASE/VIAL

N62798 005

Dec 11, 1995

EQ 1GM BASE/VIAL

N62798 001

Dec 11, 1995

EQ 2GM BASE/VIAL

N62798 002

Dec 11, 1995

OXAMNIQUINECAPSULE; ORAL
VANSIL

PFIZER

250MG

N18069 001

OXAPROZIN POTASSIUMTABLET; ORAL
DAYPRO ALTA

GD SEARLE

600MG

N20776 001

Oct 17, 2002

OXAZEPAMCAPSULE; ORAL
OXAZEPAM

AM THERAP

10MG

N71955 001

Mar 03, 1988

15MG

N71956 001

Mar 03, 1988

30MG

N71957 001

Mar 03, 1988

MUTUAL PHARM

10MG

N70957 001

Aug 10, 1987

15MG

N71025 001

Aug 10, 1987

30MG

N71026 001

Aug 10, 1987

MYLAN

10MG

N71713 001

Oct 20, 1987

15MG

N71714 001

Oct 20, 1987

30MG

N71715 001

Oct 20, 1987

SERAX

ALPHARMA US PHARMS

10MG

N15539 002

15MG

N15539 004

30MG

N15539 006

ZAXOPAM

QUANTUM PHARMICS

10MG

N70650 001

Mar 01, 1988

15MG

N70640 001

Mar 01, 1988

30MG

N70641 001

Mar 01, 1988

TABLET; ORAL
OXAZEPAM

MUTUAL PHARM

15MG

N70683 001

Jan 16, 1987

PARKE DAVIS

15MG

N71508 001

Feb 02, 1987

WATSON LABS

15MG

N71494 001

Apr 21, 1987

SERAX

ALPHARMA US PHARMS

15MG

N15539 008

OXPRENOLOL HYDROCHLORIDECAPSULE; ORAL
TRASICOR

NOVARTIS

20MG

N18166 001

Dec 28, 1983

40MG

N18166 002

Dec 28, 1983

80MG

N18166 003

Dec 28, 1983

160MG

N18166 004

Dec 28, 1983

OXTRIPHYLLINESOLUTION; ORAL
CHOLEDYL

PARKE DAVIS

100MG/5ML

N09268 012

Nov 27, 1984

OXTRIPHYLLINE

MORTON GROVE

100MG/5ML

N88243 001

Dec 05, 1983

DISCONTINUED DRUG PRODUCT LIST

6 - 190 (of 263)

OXTRIPHYLLINE

SYRUP; ORAL			
CHOLEDYL			
PARKE DAVIS	50MG/5ML	N09268	011
OXTRIPHYLLINE PEDIATRIC			
MORTON GROVE	50MG/5ML	N88242	001 Dec 05, 1983
TABLET, DELAYED RELEASE; ORAL			
CHOLEDYL			
PARKE DAVIS	100MG	N09268	003
	200MG	N09268	007
OXTRIPHYLLINE			
WATSON LABS	100MG	N87866	001 Aug 25, 1983
	200MG	N87835	001 Aug 25, 1983

OXYBUTYNIN CHLORIDE

TABLET; ORAL			
OXYBUTYNIN CHLORIDE			
QUANTUM PHARMICS	5MG	N72296	001 Dec 08, 1988
USL PHARMA	5MG	N70746	001 Mar 10, 1988
WATSON LABS	5MG	N72485	001 Apr 19, 1989

OXYCODONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL			
OXYCONTIN			
PURDUE PHARMA LP	160MG	N20553	005 Mar 15, 2000
ROXICODONE			
ROXANE	10MG	N20932	001 Oct 26, 1998
	30MG	N20932	002 Oct 26, 1998

OXYMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION			
NUMORPHAN			
ENDO PHARMS	1.5MG/ML	N11707	001
SUPPOSITORY; RECTAL			
NUMORPHAN			
ENDO PHARMS	5MG	N11738	004

OXYPHENBUTAZONE

TABLET; ORAL			
OXYPHENBUTAZONE			
WATSON LABS	100MG	N88399	001 Sep 17, 1984
TANDEARIL			
NOVARTIS	100MG	N12542	004 Sep 03, 1982

OXYPHENCYCLIMINE HYDROCHLORIDE

TABLET; ORAL			
DARICON			
PFIZER	10MG	N11612	001

OXYPHENONIUM BROMIDE

TABLET; ORAL			
ANTRENYL			
NOVARTIS	5MG	N08492	002

OXYTETRACYCLINE

TABLET; ORAL			
TERRAMYCIN			
PFIZER	250MG	N50287	001

OXYTETRACYCLINE CALCIUM

SYRUP; ORAL			
TERRAMYCIN			
PFIZER	EQ 125MG BASE/5ML	N60595	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 191 (of 263)

OXYTETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL		
OXY-KESSO-TETRA		
FERRANTE	EQ 250MG BASE	N60179 001
OXYTETRACYCLINE HYDROCHLORIDE		
IMPAX LABS	EQ 250MG BASE	N60760 001
PROTER	EQ 250MG BASE	N60869 001
PUREPAC PHARM	EQ 250MG BASE	N60634 001
WEST WARD	EQ 250MG BASE	N60770 001
TERRAMYCIN		
PFIZER	EQ 125MG BASE	N50286 001
INJECTABLE; INJECTION		
TERRAMYCIN		
PFIZER	EQ 250MG BASE/VIAL	N60586 001
	EQ 500MG BASE/VIAL	N60586 002

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT; OTIC		
TERRAMYCIN W/ POLYMYXIN		
PFIZER	EQ 5MG BASE/GM;10,000 UNITS/GM	N61841 001
TABLET; VAGINAL		
TERRAMYCIN-POLYMYXIN		
PFIZER	EQ 100MG BASE;100,000 UNITS	N61009 001

OXYTOCIN

INJECTABLE; INJECTION			
OXYTOCIN			
BAXTER HLTHCARE CORP	10USP UNITS/ML	N18243	001
OXYTOCIN 10 USP UNITS IN DEXTROSE 5%			
ABBOTT	1USP UNITS/100ML	N19185	004 Mar 29, 1985
	2USP UNITS/100ML	N19185	003 Mar 29, 1985
OXYTOCIN 20 USP UNITS IN DEXTROSE 5%			
ABBOTT	2USP UNITS/100ML	N19185	002 Mar 29, 1985
OXYTOCIN 5 USP UNITS IN DEXTROSE 5%			
ABBOTT	1USP UNITS/100ML	N19185	001 Mar 29, 1985
SYNTOCINON			
NOVARTIS	10USP UNITS/ML	N18245	001
SOLUTION; NASAL			
SYNTOCINON			
NOVARTIS	40USP UNITS/ML	N12285	001

PAMIDRONATE DISODIUM

INJECTABLE; INJECTION			
AREDIA			
NOVARTIS	60MG/VIAL	N20036	003 May 06, 1993

PANCRELIPASE (AMYLASE;LIPASE;PROTEASE)

CAPSULE; ORAL			
COTAZYM			
ORGANON USA INC	30,000USP UNITS;8,000USP UNITS;30,000USP UNITS	N20580	001 Dec 09, 1996

PANCURONIUM BROMIDE

INJECTABLE; INJECTION			
PANCURONIUM BROMIDE			
ASTRAZENECA	1MG/ML	N72210	001 Mar 31, 1988
	2MG/ML	N72211	001 Mar 31, 1988
	2MG/ML	N72212	001 Mar 31, 1988
	2MG/ML	N72213	001 Mar 31, 1988
PAVULON			
ORGANON USA INC	1MG/ML	N17015	002
	2MG/ML	N17015	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 192 (of 263)

PARAMETHADIONE

CAPSULE; ORAL			
PARADIONE			
ABBOTT	150MG	N06800	003
	300MG	N06800	001
SOLUTION; ORAL			
PARADIONE			
ABBOTT	300MG/ML	N06800	002

PARAMETHASONE ACETATE

TABLET; ORAL			
HALDRONE			
LILLY	1MG	N12772	005
	2MG	N12772	006

PARGYLINE HYDROCHLORIDE

TABLET; ORAL			
EUTONYL			
ABBOTT	10MG	N13448	002
	25MG	N13448	003
	50MG	N13448	004

PAROMOMYCIN SULFATE

CAPSULE; ORAL			
HUMATIN			
PARKEDALE	EQ 250MG BASE	N60521	001
SYRUP; ORAL			
HUMATIN			
PARKE DAVIS	EQ 125MG BASE/5ML	N60522	001

PAROXETINE HYDROCHLORIDE

CAPSULE; ORAL				
PAXIL				
GLAXOSMITHKLINE	EQ 10MG BASE **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N20885	001	Oct 09, 1998
	EQ 20MG BASE **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N20885	002	Oct 09, 1998
	EQ 30MG BASE **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N20885	003	Oct 09, 1998
	EQ 40MG BASE **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N20885	004	Oct 09, 1998
TABLET; ORAL				
PAXIL				
GLAXOSMITHKLINE	EQ 50MG BASE	N20031	004	Dec 29, 1992

PEMOLINE

TABLET; ORAL				
CYLERT				
ABBOTT	18.75MG	N16832	001	
	37.5MG	N16832	002	
	75MG	N16832	003	
PEMOLINE				
AMIDE PHARM	18.75MG	N75595	001	Feb 28, 2000
	37.5MG	N75595	002	Feb 28, 2000
	75MG	N75595	003	Feb 28, 2000
MALLINCKRODT	18.75MG	N75726	003	Mar 30, 2001
	37.5MG	N75726	002	Mar 30, 2001
	75MG	N75726	001	Mar 30, 2001

DISCONTINUED DRUG PRODUCT LIST

6 - 193 (of 263)

PEMOLINE

TABLET; ORAL
PEMOLINE

SANDOZ	18.75MG	N75286 001	Dec 27, 1999
	37.5MG	N75286 002	Jun 30, 1999
	75MG	N75286 003	Jun 30, 1999
TEVA PHARMS	18.75MG	N75030 003	Feb 22, 2000
	37.5MG	N75030 001	Jan 29, 1999
	75MG	N75030 002	Jan 29, 1999
VINTAGE PHARMS	18.75MG	N75328 001	Apr 19, 2000
	37.5MG	N75328 002	Apr 19, 2000
	75MG	N75328 003	Apr 19, 2000
WATSON LABS	18.75MG	N75287 001	Jun 13, 2001
	37.5MG	N75287 002	Sep 18, 2000
	75MG	N75287 003	Sep 18, 2000

TABLET, CHEWABLE; ORAL
CYLERT

ABBOTT	37.5MG	N17703 001	
PEMOLINE			
AMIDE PHARM	37.5MG	N75678 001	Jul 26, 2000
TEVA PHARMS	37.5MG	N75555 001	Feb 18, 2000

PENBUTOLOL SULFATE

TABLET; ORAL
LEVATOL

SCHWARZ PHARMA	10MG	N18976 001	Dec 30, 1987
----------------	------	------------	--------------

PENICILLIN G BENZATHINE

INJECTABLE; INJECTION
BICILLIN L-A

WYETH AYERST	300,000 UNITS/ML	N50131 001	
--------------	------------------	------------	--

SUSPENSION; ORAL
BICILLIN

WYETH AYERST	300,000 UNITS/5ML	N50126 002	
--------------	-------------------	------------	--

TABLET; ORAL
BICILLIN

WYETH AYERST	200,000 UNITS	N50128 001	
--------------	---------------	------------	--

PENICILLIN G POTASSIUM

FOR SOLUTION; ORAL
PENICILLIN

TEVA	200,000 UNITS/5ML	N60307 002	
	400,000 UNITS/5ML	N60307 004	

PENICILLIN G POTASSIUM
MYLAN

	200,000 UNITS/5ML	N60752 003	
	250,000 UNITS/5ML	N60752 002	
	400,000 UNITS/5ML	N60752 001	
PUREPAC PHARM	250,000 UNITS/5ML	N61740 001	
	400,000 UNITS/5ML	N61740 002	

PENICILLIN-2
TEVA

	250,000 UNITS/5ML	N60307 003	
--	-------------------	------------	--

PENTIDS '200'

APOTHECON	200,000 UNITS/5ML	N62149 001	
-----------	-------------------	------------	--

PENTIDS '400'

APOTHECON	400,000 UNITS/5ML	N62149 002	
-----------	-------------------	------------	--

PFIZERPEN G

PFIZER	400,000 UNITS/5ML	N60587 001	
--------	-------------------	------------	--

INJECTABLE; INJECTION

PENICILLIN G POTASSIUM
APOTHECON

	1,000,000 UNITS/VIAL	N60362 001	
	5,000,000 UNITS/VIAL	N60362 003	
	10,000,000 UNITS/VIAL	N60362 004	
	20,000,000 UNITS/VIAL	N60362 002	
CONSOLIDATED PHARM	500,000 UNITS/VIAL	N60806 001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 194 (of 263)

PENICILLIN G POTASSIUMINJECTABLE; INJECTION
PENICILLIN G POTASSIUM

CONSOLIDATED PHARM	1,000,000 UNITS/VIAL	N60806	002	
	5,000,000 UNITS/VIAL	N60806	003	
	10,000,000 UNITS/VIAL	N60806	004	
LILLY	200,000 UNITS/VIAL	N60384	004	
	500,000 UNITS/VIAL	N60384	003	
	1,000,000 UNITS/VIAL	N60384	002	
	5,000,000 UNITS/VIAL	N60384	001	
	20,000,000 UNITS/VIAL	N60384	005	
	20,000,000 UNITS/VIAL	N60601	001	
MARSAM PHARMS LLC	1,000,000 UNITS/VIAL	N62991	001	Sep 13, 1988
	5,000,000 UNITS/VIAL	N62991	002	Sep 13, 1988
	10,000,000 UNITS/VIAL	N62991	003	Sep 13, 1988
	20,000,000 UNITS/VIAL	N62991	004	Sep 13, 1988
PARKE DAVIS	1,000,000 UNITS/VIAL	N62003	001	
	5,000,000 UNITS/VIAL	N62003	002	

TABLET; ORAL

PENICILLIN G POTASSIUM

APOTHECON	250,000 UNITS	N60392	003	
IVAX PHARMS	400,000 UNITS	N60073	004	
LILLY	250,000 UNITS	N60403	001	
MYLAN	200,000 UNITS	N60781	001	
	250,000 UNITS	N60781	002	
	400,000 UNITS	N60781	003	
	500,000 UNITS	N60781	005	
	800,000 UNITS	N60781	004	
PUREPAC PHARM	200,000 UNITS	N61588	001	
	250,000 UNITS	N61588	002	
	400,000 UNITS	N61588	003	
	400,000 UNITS	N61588	004	
TEVA	200,000 UNITS	N60306	001	
	250,000 UNITS	N60306	002	
	400,000 UNITS	N60306	003	
	500,000 UNITS	N60306	004	
	800,000 UNITS	N60413	001	
WYETH AYERST	200,000 UNITS	N60413	002	
	250,000 UNITS	N60413	003	
	400,000 UNITS	N60413	003	
PENTIDS '200'				
APOTHECON	200,000 UNITS	N62155	001	
PENTIDS '250'				
APOTHECON	250,000 UNITS	N62155	002	
PENTIDS '400'				
APOTHECON	400,000 UNITS	N60392	004	
	400,000 UNITS	N62155	003	
PENTIDS '800'				
APOTHECON	800,000 UNITS	N60392	005	
	800,000 UNITS	N62155	004	
PFIZERPEN G				
PFIZER	50,000 UNITS	N60075	001	
	100,000 UNITS	N60075	002	
	200,000 UNITS	N60075	003	
	250,000 UNITS	N60075	004	
	400,000 UNITS	N60075	005	
	800,000 UNITS	N60075	006	

PENICILLIN G PROCAINEINJECTABLE; INJECTION
DURACILLIN A.S.

LILLY	300,000 UNITS/ML	N60093	001	
PENICILLIN G PROCAINE				
CONSOLIDATED PHARM	300,000 UNITS/ML	N60800	001	
	600,000 UNITS/1.2ML	N60800	002	
PARKE DAVIS	300,000 UNITS/ML	N62029	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 195 (of 263)

PENICILLIN G PROCAINE

INJECTABLE; INJECTION
PENICILLIN G PROCAINE

PFIZER	300,000 UNITS/VIAL	N60099	001
	1,500,000 UNITS/VIAL	N60099	002
PFIZERPEN-AS			
PFIZER	300,000 UNITS/ML	N60286	001
	600,000 UNITS/ML	N60286	002

PENICILLIN G SODIUM

INJECTABLE; IM-IV
PENICILLIN G SODIUM

MARSAM PHARMS LLC	5,000,000 UNITS/VIAL	N63014	001	Sep 13, 1988
INJECTABLE; INJECTION PENICILLIN G SODIUM				
BRISTOL MYERS SQUIBB	5,000,000 UNITS/VIAL	N61935	001	
COPANOS	5,000,000 UNITS/VIAL	N61051	001	
PHARMACIA AND UPJOHN	1,000,000 UNITS/VIAL	N61046	001	

PENICILLIN V

FOR SUSPENSION; ORAL
V-CILLIN

LILLY	125MG/0.6ML	N60002	001
-------	-------------	--------	-----

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL
BEEPEN-VK

GLAXOSMITHKLINE	EQ 125MG BASE/5ML	N62270	001
	EQ 250MG BASE/5ML	N62270	002
BETAPEN-VK			
APOTHECON	EQ 125MG BASE/5ML	N61149	001
	EQ 250MG BASE/5ML	N61149	002
LEDERCILLIN VK			
LEDERLE	EQ 125MG BASE/5ML	N60136	001
	EQ 250MG BASE/5ML	N60136	002
PENAPAR-VK			
PARKE DAVIS	EQ 125MG BASE/5ML	N62002	001
	EQ 250MG BASE/5ML	N62002	002
PENICILLIN V POTASSIUM			
MYLAN	EQ 125MG BASE/5ML	N61624	002
	EQ 250MG BASE/5ML	N61624	001
PUREPAC PHARM	EQ 125MG BASE/5ML	N61758	001
	EQ 250MG BASE/5ML	N61758	002
PEN-VEE K			
WYETH AYERST	EQ 125MG BASE/5ML	N60007	001
	EQ 250MG BASE/5ML	N60007	002
PFIZERPEN VK			
PFIZER	EQ 125MG BASE/5ML	N61815	001
	EQ 250MG BASE/5ML	N61815	002
V-CILLIN K			
LILLY	EQ 125MG BASE/5ML	N60004	001
	EQ 250MG BASE/5ML	N60004	002
VEETIDS '125'			
APOTHECON	EQ 125MG BASE/5ML	N61206	001
	EQ 125MG BASE/5ML	N62153	001
VEETIDS '250'			
APOTHECON	EQ 250MG BASE/5ML	N61206	002
	EQ 250MG BASE/5ML	N62153	002
TABLET; ORAL			
BEEPEN-VK			
GLAXOSMITHKLINE	EQ 250MG BASE	N62273	001
	EQ 500MG BASE	N62273	002
BETAPEN-VK			
BRISTOL	EQ 250MG BASE	N61150	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 196 (of 263)

PENICILLIN V POTASSIUM

TABLET; ORAL

BETAPEN-VK

BRISTOL

EQ 500MG BASE

N61150 002

LEDERCILLIN VK

LEDERLE

EQ 250MG BASE

N60134 001

EQ 500MG BASE

N60134 002

PENAPAR-VK

PARKE DAVIS

EQ 250MG BASE

N62001 001

EQ 500MG BASE

N62001 002

PENICILLIN V POTASSIUM

CONSOLIDATED PHARM

EQ 250MG BASE

N61528 001

EQ 500MG BASE

N61528 002

IVAX PHARMS

EQ 125MG BASE

N60518 001

EQ 250MG BASE

N60518 002

EQ 500MG BASE

N60518 003

MYLAN

EQ 250MG BASE

N61530 001

EQ 500MG BASE

N61530 002

PUREPAC PHARM

EQ 125MG BASE

N61571 001

EQ 250MG BASE

N61571 002

EQ 500MG BASE

N61571 003

PEN-VEE K

WYETH AYERST

EQ 125MG BASE

N60006 001

EQ 250MG BASE

N60006 002

EQ 500MG BASE

N60006 003

PFIZERPEN VK

PFIZER

EQ 250MG BASE

N61836 001

EQ 500MG BASE

N61836 002

UTICILLIN VK

PHARMACIA AND UPJOHN

EQ 250MG BASE

N61651 001

EQ 500MG BASE

N61651 002

V-CILLIN K

LILLY

EQ 125MG BASE

N60003 001

EQ 250MG BASE

N60003 002

EQ 500MG BASE

N60003 003

VEETIDS '250'

APOTHECON

EQ 250MG BASE

N61164 001

EQ 250MG BASE

N62156 002

VEETIDS '500'

APOTHECON

EQ 500MG BASE

N61164 002

EQ 500MG BASE

N62156 001

PENTAGASTRIN

INJECTABLE; INJECTION

PEPTAVLON

WYETH AYERST

0.25MG/ML

N17048 001

PENTAMIDINE ISETHIONATE

INJECTABLE; INJECTION

PENTACARINAT

ARMOUR PHARM

300MG/VIAL

N73447 001

Apr 28, 1994

PENTAMIDINE ISETHIONATE

BAXTER HLTHCARE

300MG/VIAL

N73617 001

Dec 18, 1995

PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN 50

SANOFI SYNTHELABO

EQ 50MG BASE

N16732 001

PENTETATE CALCIUM TRISODIUM YB-169

INJECTABLE; INJECTION

YTTERBIUM YB 169 DTPA

3M

2mCi/ML

N17518 001

DISCONTINUED DRUG PRODUCT LIST

6 - 197 (of 263)

PENTOBARBITAL

ELIXIR; ORAL
NEMBUTAL

OVATION PHARMS	18.2MG/5ML	N83244	001	
----------------	------------	--------	-----	--

PENTOBARBITAL SODIUM

CAPSULE; ORAL

NEMBUTAL SODIUM

OVATION PHARMS	30MG	N84095	001	
	50MG	N84093	001	
	100MG	N83245	001	

PENTOBARBITAL SODIUM

LANNETT	50MG	N85937	001	
	100MG	N85915	001	
VITARINE	100MG	N83284	001	
WHITEWORTH TOWN PLSN	100MG	N83338	001	

SODIUM PENTOBARBITAL

ANABOLIC	100MG	N84590	001	
ELKINS SINN	100MG	N83368	001	
EVERYLIFE	100MG	N83259	001	
HALSEY	100MG	N84677	001	
IVAX PHARMS	50MG	N83461	001	
	100MG	N83461	002	
PARKE DAVIS	100MG	N84156	001	
PERRIGO	100MG	N84560	001	
PUREPAC PHARM	100MG	N83301	001	
VALEANT PHARM INTL	100MG	N83264	001	
WATSON LABS	100MG	N85791	001	
WYETH AYERST	100MG	N83239	001	

INJECTABLE; INJECTION

PENTOBARBITAL SODIUM

ELKINS SINN	50MG/ML	N83270	001	
-------------	---------	--------	-----	--

SODIUM PENTOBARBITAL

WYETH AYERST	50MG/ML	N83261	001	
--------------	---------	--------	-----	--

SUPPOSITORY; RECTAL

NEMBUTAL

OVATION PHARMS	30MG	N83247	001	Jan 25, 1982
	60MG	N83247	002	Jan 25, 1982
	120MG	N83247	003	Jan 25, 1982
	200MG	N83247	004	Jan 25, 1982

TABLET; ORAL

PENTOBARBITAL SODIUM

VITARINE	100MG	N83285	001	
----------	-------	--------	-----	--

SODIUM PENTOBARBITAL

ANABOLIC	100MG	N84238	001	
----------	-------	--------	-----	--

PENTOLINIUM TARTRATE

INJECTABLE; INJECTION

ANSOLYSEN

WYETH AYERST	10MG/ML	N09372	001	
--------------	---------	--------	-----	--

PERFLUBRON

LIQUID; ORAL

IMAGENT

ALLIANCE PHARM	100%	N20091	001	Aug 13, 1993
----------------	------	--------	-----	--------------

PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENE

PASTE; TOPICAL

SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS

US ARMY	50%;50%	N21084	001	Feb 17, 2000
---------	---------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 198 (of 263)

PERMETHRINLOTION; TOPICAL
NIX

GLAXOSMITHKLINE 1% N19435 001 Mar 31, 1986

PERPHENAZINE

CONCENTRATE; ORAL

TRILAFON

SCHERING 16MG/5ML N11557 001

INJECTABLE; INJECTION

TRILAFON

SCHERING 5MG/ML N11213 002

SYRUP; ORAL

TRILAFON

SCHERING 2MG/5ML N11294 002

TABLET; ORAL

TRILAFON

SCHERING 2MG N10775 001

4MG N10775 002

8MG N10775 003

16MG N10775 004

TABLET, EXTENDED RELEASE; ORAL

TRILAFON

SCHERING 8MG N11361 002

PHENACEMIDE

TABLET; ORAL

PHENURONE

ABBOTT 500MG N07707 001

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE

TABLET; ORAL

AZO GANTANOL

ROCHE 100MG;500MG N13294 001 Sep 10, 1987

PHENAZOPYRIDINE HYDROCHLORIDE; SULFAMETHOXAZOLE; TRIMETHOPRIM

TABLET; ORAL

SULFAMETHOXAZOLE AND TRIMETHOPRIM AND PENAZOPYRIDINE HYDROCHLORIDE

ABLE N/A, 200MG; 800MG, N/A; 160MG, N/A N21105 001 Jun 26, 2001

PHENAZOPYRIDINE HYDROCHLORIDE; SULFISOXAZOLE

TABLET; ORAL

AZO GANTRISIN

ROCHE 50MG;500MG N19358 001 Aug 31, 1990

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL

PHENAZINE

MAST MM 35MG N86523 001

35MG N86524 001

35MG N86525 001

PHENDIMETRAZINE TARTRATE

SANDOZ 35MG N85633 001

35MG N85694 001

35MG N85702 001

VITARINE 35MG N85634 001

35MG N85645 001

35MG N85670 001

35MG N86403 001

35MG N86408 001

35MG N86410 001

35MG N87424 001

SPRX-3

SOLVAY 35MG N85897 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 199 (of 263)

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL				
STATOBEX				
TEVA	35MG	N85507	001	
CAPSULE, EXTENDED RELEASE; ORAL				
MELFIAT-105				
NUMARK	105MG	N87487	001	Oct 13, 1982
PHENDIMETRAZINE TARTRATE				
GRAHAM DM	105MG	N87214	001	May 26, 1982
	105MG	N88020	001	Aug 16, 1982
	105MG	N88028	001	Aug 16, 1982
	105MG	N88062	001	Sep 13, 1982
	105MG	N88063	001	Sep 10, 1982
	105MG	N88111	001	Oct 18, 1982
SANDOZ	105MG	N87378	001	
SPRX-105				
NUMARK	105MG	N88024	001	Dec 22, 1982
TABLET; ORAL				
ADPHEN				
FERNDAL LABS	35MG	N83655	001	
ALPHAZINE				
SANDOZ	35MG	N85034	001	
CAM-METRAZINE				
ABC HOLDING	35MG	N83922	001	
	35MG	N85318	001	
	35MG	N85320	001	
	35MG	N85321	001	
	35MG	N85511	001	
CAMALL	35MG	N85756	001	
DI-METREX				
PVT FORM	35MG	N85698	001	
MELFIAT				
NUMARK	35MG	N83790	002	
METRA				
FOREST PHARMS	35MG	N83754	001	
PHENAZINE				
MAST MM	35MG	N87305	001	
PHENAZINE-35				
ABC HOLDING	35MG	N85512	001	
PHENDIMETRAZINE TARTRATE				
ABC HOLDING	35MG	N85761	001	
	35MG	N85941	001	Jun 27, 1983
ANABOLIC	35MG	N86020	001	
BARR	35MG	N83644	001	
	35MG	N83684	001	
	35MG	N83686	001	
	35MG	N83687	001	
	35MG	N84831	001	
	35MG	N84834	001	
	35MG	N84835	001	
FERNDAL LABS	35MG	N86834	001	Sep 15, 1983
INWOOD LABS	35MG	N84740	001	
	35MG	N84741	001	
	35MG	N84742	001	
	35MG	N84743	001	
IVAX PHARMS	35MG	N83682	001	
	35MG	N85611	001	
	35MG	N85612	001	
KV PHARM	35MG	N84138	001	
	35MG	N84141	001	
	35MG	N85525	001	
MFG CHEMISTS	35MG	N85914	001	
NUMARK	35MG	N83790	001	
PVT FORM	35MG	N85199	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 200 (of 263)

PHENDIMETRAZINE TARTRATE

TABLET; ORAL

PHENDIMETRAZINE TARTRATE

PVT FORM	35MG	N85697	001
SANDOZ	35MG	N85402	001
	35MG	N85497	001
	35MG	N85830	001
	35MG	N86365	001
	35MG	N86370	001
SOLVAY	35MG	N83993	001
USL PHARMA	35MG	N83805	001
	35MG	N84398	001
	35MG	N84399	001
VITARINE	35MG	N85519	001
	35MG	N86005	001
	35MG	N86106	001
WATSON LABS	35MG	N85767	001
	35MG	N85768	001
	35MG	N85770	001
	35MG	N85773	001
PLEGINE			
WYETH AYERST	35MG	N12248	001
STATOBEX			
TEVA	35MG	N86013	001
STATOBEX-G			
TEVA	35MG	N85095	001
X-TROZINE			
SHIRE RICHWOOD	35MG	N86550	001
	35MG	N86551	001
	35MG	N86552	001
	35MG	N86553	001
	35MG	N86554	001

PHENINDIONE

TABLET; ORAL

HEDULIN

AVENTIS PHARMS	50MG	N08767	002
----------------	------	--------	-----

PHENMETRAZINE HYDROCHLORIDE

TABLET; ORAL

PRELUDIN

BOEHRINGER INGELHEIM	25MG	N10460	005
TABLET, EXTENDED RELEASE; ORAL			
PRELUDIN			
BOEHRINGER INGELHEIM	50MG	N11752	004
	75MG	N11752	003

PHENPROCOUMON

TABLET; ORAL

LIQUAMAR

ORGANON USA INC	3MG	N11228	001
-----------------	-----	--------	-----

PHENSUXIMIDE

CAPSULE; ORAL

MILONTIN

PARKE DAVIS	500MG	N08855	004
-------------	-------	--------	-----

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

FASTIN

GLAXOSMITHKLINE	30MG	N17352	001
-----------------	------	--------	-----

OBESTIN-30

FERNDAL LABS	30MG	N87144	001
--------------	------	--------	-----

DISCONTINUED DRUG PRODUCT LIST

6 - 201 (of 263)

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

OBY-TRIM

SHIRE RICHWOOD	30MG	N87764	001	Mar 18, 1982
----------------	------	--------	-----	--------------

ONA-MAST

MAST MM	30MG	N86511	001	
	30MG	N86516	001	

PHENTERMINE HYDROCHLORIDE

ABC HOLDING	18.75MG	N88576	001	May 23, 1984
	30MG	N85411	001	
	30MG	N85417	001	
	30MG	N86732	002	
	30MG	N87215	001	
	37.5MG	N87915	001	Dec 22, 1983
	37.5MG	N87918	001	Dec 22, 1983
	37.5MG	N87930	001	Oct 14, 1983
	37.5MG	N88610	001	Jun 04, 1984
	37.5MG	N88611	001	Jun 04, 1984
	37.5MG	N88625	001	Aug 23, 1984
ABLE	15MG	N40497	001	Mar 13, 2003
	30MG	N40403	001	Aug 30, 2001
	30MG	N40427	001	Aug 30, 2001
CAMALL	15MG	N86735	001	
	30MG	N87226	001	
DURAMED PHARMS BARR	30MG	N88948	001	Apr 25, 1986
IVAX PHARMS	30MG	N86329	001	
LANNETT	30MG	N87022	001	Feb 03, 1983
SANDOZ	37.5MG	N88414	001	Oct 19, 1983
TEVA	30MG	N86911	001	
	30MG	N87126	001	
	30MG	N87777	001	Nov 01, 1985
	30MG	N88612	001	Apr 04, 1984
	30MG	N88613	001	Apr 09, 1984
	30MG	N88614	001	Apr 09, 1984
USL PHARMA	30MG	N84487	001	Apr 09, 1982
	30MG	N88430	001	Mar 27, 1984
	30MG	N88797	001	Dec 10, 1984
VITARINE	30MG	N87202	001	
	30MG	N87235	001	
WATSON LABS	30MG	N86740	001	Mar 21, 1985

TABLET; ORAL

ONA-MAST

MAST MM	8MG	N86260	001	
---------	-----	--------	-----	--

PHENTERMINE HYDROCHLORIDE

ABC HOLDING	8MG	N83923	001	
	8MG	N85319	001	
	37.5MG	N87805	001	Dec 06, 1982
	37.5MG	N88596	001	Apr 04, 1984
ABLE	37.5MG	N40402	001	Aug 30, 2001
IVAX PHARMS	8MG	N85553	001	
SANDOZ	8MG	N85671	001	
	8MG	N85689	001	
USL PHARMA	8MG	N83804	001	
	37.5MG	N88910	001	Jul 17, 1985
	37.5MG	N88917	001	Jul 17, 1985
VITARINE	8MG	N86453	001	
	8MG	N86456	001	
WATSON LABS	8MG	N85739	001	
TORA				
SOLVAY	8MG	N84035	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 202 (of 263)

PHENTERMINE RESIN COMPLEX

CAPSULE, EXTENDED RELEASE; ORAL

PHENTERMINE RESIN 30

QUANTUM PHARMICS EQ 30MG BASE

N89120 001 Feb 04, 1988

PHENYL AMINOSALICYLATE

POWDER; ORAL

PHENY-PAS-TEBAMIN

PURDUE FREDERICK 50%

N11695 002

TABLET; ORAL

PHENY-PAS-TEBAMIN

PURDUE FREDERICK 500MG

N11695 003

PHENYLBUTAZONE

CAPSULE; ORAL

AZOLID

AVENTIS 100MG

N87260 001

BUTAZOLIDIN

NOVARTIS 100MG

N08319 009

PHENYLBUTAZONE

IVAX PHARMS 100MG

N88218 001 Jun 24, 1983

MUTUAL PHARM 100MG

N88994 001 Dec 04, 1985

SANDOZ 100MG

N87774 001 Jun 16, 1982

WATSON LABS 100MG

N87756 001 Dec 17, 1982

TABLET; ORAL

AZOLID

AVENTIS 100MG

N87091 001

BUTAZOLIDIN

NOVARTIS 100MG

N08319 008

PHENYLBUTAZONE

MUTUAL PHARM 100MG

N88863 001 Dec 04, 1985

SANDOZ 100MG

N84339 001

WATSON LABS 100MG

N86151 001

100MG N87674 001 Apr 21, 1982

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PHENERGAN VC

WYETH AYERST 5MG/5ML; 6.25MG/5ML

N08604 003 Apr 02, 1984

PHERAZINE VC

HALSEY 5MG/5ML; 6.25MG/5ML

N88868 001 Mar 02, 1987

PROMETHAZINE VC PLAIN

CENCI 5MG/5ML; 6.25MG/5ML

N88815 001 Nov 22, 1985

PHENYLEPHRINE HYDROCHLORIDE; PYRILAMINE MALEATE

SOLUTION/DROPS; OPHTHALMIC

PREFRIN-A

ALLERGAN 0.12%; 0.1%

N07953 001

PHENYTOIN

SUSPENSION; ORAL

DILANTIN-30

PARKE DAVIS 30MG/5ML

N08762 002

PHENYTOIN SODIUM

CAPSULE; ORAL

DIPHENYLAN SODIUM

LANNETT 30MG PROMPT

N80857 001

100MG PROMPT N80857 002

EXTENDED PHENYTOIN SODIUM

PLIVA 100MG EXTENDED

N89441 001 Dec 18, 1986

PHENYTEX

WATSON LABS 100MG EXTENDED

N88711 001 Dec 21, 1984

DISCONTINUED DRUG PRODUCT LIST

6 - 203 (of 263)

PHENYTOIN SODIUM

CAPSULE; ORAL

PHENYTOIN SODIUM

PHARMERAL 100MG PROMPT N85435 001

WATSON LABS 100MG PROMPT N85894 001

PROMPT PHENYTOIN SODIUM

WATSON LABS 100MG PROMPT N80905 001

INJECTABLE; INJECTION

DILANTIN

PARKE DAVIS 50MG/ML N10151 001

PHENYTOIN SODIUM

AM PHARM PARTNERS 50MG/ML N89003 001 May 31, 1985

MARSAM PHARMS LLC 50MG/ML N89501 001 Oct 13, 1987

50MG/ML N89779 001 Nov 27, 1992

SMITH AND NEPHEW 50MG/ML N88519 001 Dec 19, 1984

50MG/ML N88521 001 Dec 18, 1984

SOLOPAK 50MG/ML N88520 001 Dec 17, 1984

STERIS 50MG/ML N85434 001

WARNER CHILCOTT 50MG/ML N89900 001 Mar 30, 1990

PHYTONADIONE

INJECTABLE; INJECTION

KONAKION

ROCHE 1MG/0.5ML N11745 001

10MG/ML N11745 003

PHYTONADIONE

GLAXOSMITHKLINE 1MG/0.5ML N84060 001

10MG/ML N84060 002

VITAMIN K1

HOSPIRA 10MG/ML N87956 001 Jul 25, 1983

PILOCARPINE

INSERT, EXTENDED RELEASE; OPHTHALMIC

OCUSERT PILO-20

AKORN 5MG N17431 001

OCUSERT PILO-40

AKORN 11MG N17548 001

PINACIDIL

CAPSULE, EXTENDED RELEASE; ORAL

PINDAC

LEO PHARM 12.5MG N19456 001 Dec 28, 1989

25MG N19456 002 Dec 28, 1989

PINDOLOL

TABLET; ORAL

PINDOLOL

PUREPAC PHARM 5MG N74125 001 Apr 28, 1993

10MG N74125 002 Apr 28, 1993

PIPECURONIUM BROMIDE

INJECTABLE; INJECTION

ARDUAN

ORGANON USA INC 10MG/VIAL N19638 001 Jun 26, 1990

PIPERACETAZINE

TABLET; ORAL

QUIDE

DOW PHARM 10MG N13615 001

25MG N13615 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 204 (of 263)

PIPERACILLIN SODIUM

INJECTABLE; INJECTION

PIPRACIL

WYETH PHARMS INC

EQ 2GM BASE/VIAL

N50545 002

EQ 2GM BASE/VIAL

N62750 001

Oct 13, 1987

EQ 3GM BASE/VIAL

N50545 003

EQ 3GM BASE/VIAL

N62750 002

Oct 13, 1987

EQ 4GM BASE/VIAL

N50545 004

EQ 4GM BASE/VIAL

N62750 003

Oct 13, 1987

EQ 40GM BASE/VIAL

N50545 006

Sep 30, 1985

PIPERAZINE CITRATE

SYRUP; ORAL

ANTEPAR

GLAXOSMITHKLINE

EQ 500MG BASE/5ML

N09102 001

BRYREL

SANOFI SYNTHELABO

EQ 500MG BASE/5ML

N17796 001

MULTIFUGE

BLULINE

EQ 500MG BASE/5ML

N09452 001

PIPERAZINE CITRATE

ALPHARMA US PHARMS

EQ 500MG BASE/5ML

N80774 001

LANNETT

EQ 500MG BASE/5ML

N80963 001

LUITPOLD

EQ 500MG BASE/5ML

N80671 001

VERMIDOL

SOLVAY

EQ 500MG BASE/5ML

N80992 001

TABLET; ORAL

ANTEPAR

GLAXOSMITHKLINE

EQ 500MG BASE

N09102 003

PIPERAZINE CITRATE

IMPAX LABS

EQ 250MG BASE

N80874 001

PIPOBROMAN

TABLET; ORAL

VERCYTE

ABBOTT

10MG

N16245 001

25MG

N16245 002

PIRBUTEROL ACETATE

AEROSOL, METERED; INHALATION

MAXAIR

3M

EQ 0.2MG BASE/INH

N19009 001

Dec 30, 1986

PIROXICAM

CAPSULE; ORAL

PIROXICAM

IVAX PHARMS

10MG

N74148 001

Jun 03, 1996

20MG

N74148 002

Jun 03, 1996

ROXANE

10MG

N73651 001

Feb 26, 1993

20MG

N73651 002

Feb 26, 1993

SCS

10MG

N74036 001

May 29, 1992

20MG

N74036 002

May 29, 1992

POLYESTRADIOL PHOSPHATE

INJECTABLE; INJECTION

ESTRADURIN

WYETH AYERST

40MG/AMP

N10753 001

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE

SOLUTION; ORAL

OCL

HOSPIRA

6GM/100ML;75MG/100ML;168MG/100ML;146MG/100ML;1.29GM/100ML

N19284 001

Apr 30, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 205 (of 263)

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE;
SODIUM SULFATE, ANHYDROUSFOR SOLUTION; ORAL
COLYTE

SCHWARZ PHARMA	120GM/PACKET;1.49GM/PACKET;3.36GM/PACKET;2.92GM/PACKET;11.36GM/PACKET	N18983	005	Oct 26, 1984
	227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET	N18983	004	Oct 26, 1984
	360GM/PACKET;4.47GM/PACKET;10.08GM/PACKET;8.76GM/PACKET;34.08GM/PACKET	N18983	006	Oct 26, 1984

FOR SUSPENSION; ORAL
COLOVAGE

DYNAPHARM	227.1GM/PACKET;2.82GM/PACKET;6.36GM/PACKET;5.53GM/PACKET;21.5GM/PACKET	N71320	001	Apr 20, 1988
-----------	--	--------	-----	--------------

E-Z-EM PREP LYTE
E Z EM

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	N71278	001	Nov 21, 1988
--	--	--------	-----	--------------

GLYCOPREP
GOLDLINE

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	N72319	001	Dec 23, 1988
--	--	--------	-----	--------------

PEG-LYTE
SANDOZ

	236GM/BOT;2.97GM/BOT;6.74GM/BOT;5.86GM/BOT;22.74GM/BOT	N73098	001	Aug 31, 1993
--	--	--------	-----	--------------

POLYMYXIN B SULFATEINJECTABLE; INJECTION
AEROSPORIN

GLAXOSMITHKLINE	EQ 500,000 U BASE/VIAL	N62036	001	
-----------------	------------------------	--------	-----	--

POWDER; FOR RX COMPOUNDING
POLYMYXIN B SULFATE

PADDOCK	100,000,000 UNITS/BOT	N62455	001	Jul 27, 1983
---------	-----------------------	--------	-----	--------------

POTASSIUM AMINOSALICYLATE

POWDER; ORAL

POTASSIUM AMINOSALICYLATE
HEXCEL

100%	N80098	001	
------	--------	-----	--

POTASSIUM CHLORIDECAPSULE, EXTENDED RELEASE; ORAL
K-LEASE

SAVAGE LABS	8MEQ	N73398	001	Jan 28, 1992
	10MEQ	N72427	001	Mar 28, 1990

FOR SUSPENSION, EXTENDED RELEASE; ORAL
MICRO-K LS

KV PHARM	20MEQ/PACKET	N19561	003	Aug 26, 1988
----------	--------------	--------	-----	--------------

INJECTABLE; INJECTION
POTASSIUM CHLORIDE

AKORN	2MEQ/ML	N88286	001	Sep 05, 1985
-------	---------	--------	-----	--------------

AM PHARM PARTNERS	2MEQ/ML	N80204	001	
-------------------	---------	--------	-----	--

	2MEQ/ML	N84290	001	
--	---------	--------	-----	--

	2MEQ/ML	N86713	001	
--	---------	--------	-----	--

	2MEQ/ML	N86714	001	
--	---------	--------	-----	--

	2MEQ/ML	N87787	001	Apr 20, 1982
--	---------	--------	-----	--------------

	2MEQ/ML	N87817	001	Oct 20, 1982
--	---------	--------	-----	--------------

	2MEQ/ML	N87885	001	Feb 03, 1983
--	---------	--------	-----	--------------

BAXTER HLTHCARE	2MEQ/ML	N80203	001	
-----------------	---------	--------	-----	--

GD SEARLE LLC	1MEQ/ML	N86219	001	
---------------	---------	--------	-----	--

	2MEQ/ML	N86219	002	
--	---------	--------	-----	--

	2MEQ/ML	N86220	002	
--	---------	--------	-----	--

	3MEQ/ML	N86219	003	
--	---------	--------	-----	--

	3MEQ/ML	N86220	001	
--	---------	--------	-----	--

	4MEQ/ML	N86219	004	
--	---------	--------	-----	--

HOSPIRA	1MEQ/ML	N80205	003	
---------	---------	--------	-----	--

	1MEQ/ML	N83345	003	
--	---------	--------	-----	--

	2.4MEQ/ML	N80205	004	
--	-----------	--------	-----	--

	3.2MEQ/ML	N80205	005	
--	-----------	--------	-----	--

DISCONTINUED DRUG PRODUCT LIST

6 - 206 (of 263)

POTASSIUM CHLORIDE

INJECTABLE; INJECTION POTASSIUM CHLORIDE

LILLY	2MEQ/ML	N07865	002	
LUITPOLD	2MEQ/ML	N80221	001	
	2MEQ/ML	N80736	001	
MILES	1MEQ/ML	N80195	002	
	2MEQ/ML	N80195	001	
	3MEQ/ML	N80195	003	
	4MEQ/ML	N80195	004	
PHARMA SERVE NY	2MEQ/ML	N86297	001	
	2MEQ/ML	N87362	001	Mar 08, 1983
STERIS	2MEQ/ML	N86208	001	
	2MEQ/ML	N89163	001	Mar 10, 1988
	2MEQ/ML	N89421	001	Jan 02, 1987
	3MEQ/ML	N86210	001	

TABLET, EXTENDED RELEASE; ORAL

KAON CL				
SAVAGE LABS	6.7MEQ	N17046	001	
SLOW-K				
NOVARTIS	8MEQ	N17476	002	
TEN-K				
NOVARTIS	10MEQ	N19381	001	Apr 16, 1986

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

POTASSIUM CHLORIDE 0.037% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	37MG/100ML;900MG/100ML	N19708	001	Sep 29, 1989
POTASSIUM CHLORIDE 0.075% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	75MG/100ML;900MG/100ML	N19708	002	Sep 29, 1989
POTASSIUM CHLORIDE 0.11% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	110MG/100ML;900MG/100ML	N19708	003	Sep 29, 1989
POTASSIUM CHLORIDE 0.22% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	220MG/100ML;900MG/100ML	N19708	005	Sep 29, 1989
POTASSIUM CHLORIDE 0.3% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
B BRAUN	300MG/100ML;900MG/100ML	N19708	006	Sep 29, 1989
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER				
B BRAUN	75MG/100ML;900MG/100ML	N18722	001	Nov 09, 1982
BAXTER HLTHCARE	75MG/100ML;900MG/100ML	N17648	004	
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER				
B BRAUN	150MG/100ML;900MG/100ML	N18722	002	Nov 09, 1982
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.22% IN PLASTIC CONTAINER				
B BRAUN	220MG/100ML;900MG/100ML	N18722	003	Nov 09, 1982
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER				
B BRAUN	300MG/100ML;900MG/100ML	N18722	004	Nov 09, 1982

POTASSIUM CHLORIDE; SODIUM CHLORIDE; TROMETHAMINE

INJECTABLE; INJECTION

THAM-E				
HOSPIRA	370MG/VIAL;1.75GM/VIAL;36GM/VIAL	N13025	001	

POTASSIUM CITRATE

FOR SOLUTION; ORAL POTASSIUM CITRATE

MISSION PHARMA	10MEQ/PACKET	N19647	002	Oct 13, 1988
	20MEQ/PACKET	N19647	001	Oct 13, 1988

POTASSIUM IODIDE

SOLUTION; ORAL POTASSIUM IODIDE

ROXANE	1GM/ML	N18551	001	Feb 19, 1982
--------	--------	--------	-----	--------------

TABLET; ORAL THYRO-BLOCK

MEDPOINTE PHARM HLC	130MG	N18307	001	
---------------------	-------	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 207 (of 263)

POTASSIUM PERCHLORATE

CAPSULE; ORAL PERCHLORACAP MALLINCKRODT	200MG	N17551	001	
---	-------	--------	-----	--

PRALIDOXIME CHLORIDE

INJECTABLE; INJECTION PRALIDOXIME CHLORIDE BAXTER HLTHCARE CORP	300MG/ML	N18799	001	Dec 13, 1982
TABLET; ORAL PROTOPAM CHLORIDE WYETH AYERST	500MG	N14122	002	

PRAMIPEXOLE DIHYDROCHLORIDE

TABLET; ORAL MIRAPEX BOEHRINGER INGELHEIM	1.25MG	N20667	004	Jul 01, 1997
---	--------	--------	-----	--------------

PRAZEPAM

CAPSULE; ORAL CENTRAX PARKE DAVIS	5MG 10MG 20MG	N18144 N18144 N18144	001 002 003	May 10, 1982
PRAZEPAM USL PHARMA	5MG 10MG	N70427 N70428	001 001	Nov 06, 1987 Nov 06, 1987
TABLET; ORAL CENTRAX PARKE DAVIS	10MG	N17415	001	

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL PRAZOSIN HYDROCHLORIDE AM THERAP	EQ 1MG BASE EQ 2MG BASE EQ 5MG BASE	N72782 N72783 N72784	001 001 001	May 16, 1989 May 16, 1989 May 16, 1989
CLONMEL HLTHCARE PUREPAC PHARM	EQ 2MG BASE EQ 1MG BASE EQ 2MG BASE EQ 5MG BASE	N72706 N72991 N72921 N72992	001 001 001 001	May 16, 1989 May 16, 1989 May 16, 1989 May 16, 1989

PREDNISOLONE

CREAM; TOPICAL METI-DERM SCHERING	0.5%	N10209	002	
SYRUP; ORAL PRELONE MURO	5MG/5ML	N89654	001	Jan 17, 1989
TABLET; ORAL CORTALONE HALSEY	1MG 2.5MG 5MG	N80304 N80304 N80304	003 002 001	
DELTA-CORTEF PHARMACIA AND UPJOHN	5MG	N09987	004	
FERNISOLONE-P FERNDAL LABS	5MG	N83941	001	
PREDNISOLONE BARR	5MG	N84426	002	
BUNDY	5MG	N83675	001	
ELKINS SINN	5MG	N80625	001	
EVERYLIFE	1MG 2.5MG	N84439 N84439	001 002	

DISCONTINUED DRUG PRODUCT LIST

6 - 208 (of 263)

PREDNISOLONE

TABLET; ORAL
PREDNISOLONE

EVERYLIFE	5MG	N84439	003	
FERRANTE	2.5MG	N80562	001	
	5MG	N80562	002	
HEATHER	5MG	N80326	001	
IMPAX LABS	5MG	N80780	001	
INWOOD LABS	5MG	N80748	001	
IVAX PHARMS	5MG	N80378	001	
MARSHALL PHARMA	5MG	N80307	001	
PANRAY	1MG	N80351	001	
	5MG	N80351	002	
PERRIGO	5MG	N84542	001	
PHOENIX LABS NY	5MG	N80322	001	
PUREPAC PHARM	5MG	N80325	001	
PVT FORM	5MG	N80211	001	
ROXANE	5MG	N80327	002	
SANDOZ	5MG	N80339	001	
	5MG	N84773	001	
SPERTI	1MG	N80358	001	
	2.5MG	N80358	002	
	5MG	N80358	003	
SUPERPHARM	5MG	N88892	001	Feb 26, 1985
TABLICAPS	5MG	N85170	001	
TEVA	5MG	N80398	001	
UDL	5MG	N87987	001	Jan 18, 1983
VALEANT PHARM INTL	5MG	N80236	001	
VITARINE	5MG	N80534	001	
WATSON LABS	5MG	N85085	002	
	5MG	N85415	001	
	5MG	N85416	001	
WEST WARD	5MG	N80324	001	
WHITEWORTH TOWN PLSN	5MG	N80342	001	
STERANE				
PFIZER	5MG	N09996	001	

PREDNISOLONE ACETATE

INJECTABLE; INJECTION
METICORTEZONE

SCHERING	25MG/ML	N10255	002	
PREDNISOLONE ACETATE				
AKORN	25MG/ML	N83032	001	
	50MG/ML	N84492	001	
BEL MAR	25MG/ML	N83738	001	
	50MG/ML	N83738	002	
CENT PHARMS	25MG/ML	N84717	001	
	50MG/ML	N84717	002	
STERIS	25MG/ML	N83654	001	
	40MG/ML	N83767	001	
	50MG/ML	N85781	001	
STERANE				
PFIZER	25MG/ML	N11446	001	
SUSPENSION/DROPS; OPHTHALMIC				
ECONOPRED				
ALCON	0.125%	N17468	001	

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC
CETAPRED

ALCON	0.25%;10%	N87771	001	Aug 06, 1993
METIMYD				
SCHERING	0.5%;10%	N10210	002	Sep 09, 1984

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 209 (of 263)

PREDNISOLONE ACETATE; SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC

PREDSULFAIR

PHARMAFAIR

0.5%;10%

N88032 001

Apr 15, 1983

VASOCIDIN

NOVARTIS

0.5%;10%

N88791 001

Oct 05, 1984

SUSPENSION; OPHTHALMIC

ISOPTO CETAPRED

ALCON

0.25%;10%

N87547 001

SUSPENSION/DROPS; OPHTHALMIC

METIMYD

SCHERING

0.5%;10%

N10210 001

PREDAMIDE

AKORN

0.5%;10%

N88059 001

Jul 29, 1983

PREDSULFAIR

PHARMAFAIR

0.5%;10%

N88007 001

Apr 19, 1983

PREDSULFAIR II

PHARMAFAIR

0.2%;10%

N88837 001

Dec 24, 1985

SULPHRIN

BAUSCH AND LOMB

0.5%;10%

N88089 001

Dec 28, 1982

PREDNISOLONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

HYDELTRASOL

MERCK

EQ 20MG PHOSPHATE/ML

N11583 002

PREDNISOLONE SODIUM PHOSPHATE

STERIS

EQ 20MG PHOSPHATE/ML

N80517 001

OINTMENT; OPHTHALMIC, OTIC

HYDELTRASOL

MERCK

EQ 0.25% PHOSPHATE

N11028 001

SOLUTION/DROPS; OPHTHALMIC

METRETON

SCHERING

EQ 0.5% PHOSPHATE

N83834 001

PREDAIR

PHARMAFAIR

EQ 0.11% PHOSPHATE

N88415 001

Feb 29, 1984

PREDAIR FORTE

PHARMAFAIR

EQ 0.9% PHOSPHATE

N88165 001

Mar 28, 1983

PREDNISOLONE SODIUM PHOSPHATE

AKORN

EQ 0.11% PHOSPHATE

N83358 001

EQ 0.9% PHOSPHATE

N83358 002

ALCON UNIVERSAL

EQ 0.11% PHOSPHATE

N81043 001

Oct 24, 1991

EQ 0.9% PHOSPHATE

N81044 001

Oct 24, 1991

SOLA BARNES HIND

EQ 0.11% PHOSPHATE

N84171 001

EQ 0.9% PHOSPHATE

N84168 001

EQ 0.9% PHOSPHATE

N84169 001

EQ 0.9% PHOSPHATE

N84172 001

PREDNISOLONE SODIUM PHOSPHATE; SULFACETAMIDE SODIUM

SOLUTION/DROPS; OPHTHALMIC

SULSTER

AKORN

EQ 0.23% PHOSPHATE;10%

N74511 001

Jul 30, 1996

PREDNISOLONE TEBUTATE

INJECTABLE; INJECTION

HYDELTRA-TBA

MERCK

20MG/ML

N10562 001

PREDNISOLONE TEBUTATE

STERIS

20MG/ML

N83362 001

Feb 17, 1984

PREDNISONE

SOLUTION; ORAL

PREDNISONE

MORTON GROVE

5MG/5ML

N89726 001

Aug 02, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 210 (of 263)

PREDNISONE

SYRUP; ORAL				
LIQUID PRED				
MURO	5MG/5ML	N87611	002	Sep 07, 1982
TABLET; ORAL				
CORTAN				
HALSEY	20MG	N87480	001	
DELTA-DOME				
BAYER PHARMS	5MG	N80293	001	
DELTASONE				
PHARMACIA AND UPJOHN	2.5MG	N09986	005	
	5MG	N09986	002	
	10MG	N09986	006	
	20MG	N09986	007	
	50MG	N09986	008	
FERNISONE				
FERNDAL LABS	5MG	N83364	001	
METICORTEN				
SCHERING	1MG	N09766	002	
	5MG	N09766	001	
ORASONE				
SOLVAY	1MG	N83009	001	
	5MG	N83009	002	
	10MG	N83009	003	
	20MG	N83009	004	
	50MG	N85999	001	
PARACORT				
PARKE DAVIS	5MG	N10962	002	
PREDNICEN-M				
SCHWARZ PHARMA	5MG	N84655	001	
PREDNISONE				
AM THERAP	5MG	N89387	001	Nov 06, 1986
	10MG	N89388	001	Nov 06, 1986
	20MG	N89389	001	Nov 06, 1986
BUNDY	5MG	N83676	001	
DURAMED PHARMS BARR	5MG	N88394	001	Oct 04, 1983
	10MG	N88395	001	Oct 04, 1983
	20MG	N88396	001	Oct 04, 1983
ELKINS SINN	5MG	N80491	001	
	20MG	N85811	001	
EVERYLIFE	1MG	N84440	001	
	2.5MG	N84440	002	
	5MG	N84440	003	
FERRANTE	2.5MG	N80563	001	
	5MG	N80563	002	
HALSEY	5MG	N80300	001	
HEATHER	5MG	N80320	001	
	10MG	N84341	001	
	20MG	N84417	001	
	20MG	N85543	001	
	50MG	N86946	001	
IMPAX LABS	5MG	N80782	001	
INTERPHARM	5MG	N89597	001	Oct 05, 1987
	10MG	N89598	001	Oct 05, 1987
	20MG	N89599	001	Oct 05, 1987
INWOOD LABS	1MG	N80328	001	
	2.5MG	N80306	001	
	5MG	N80279	001	
IVAX PHARMS	5MG	N80283	001	
	10MG	N84133	001	
	20MG	N84134	001	
KV PHARM	5MG	N84236	001	
LANNETT	5MG	N80514	001	
	20MG	N84275	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 211 (of 263)

PREDNISONE

TABLET; ORAL
PREDNISONE

LEDERLE	5MG	N86968	001	
MARSHALL PHARMA	5MG	N80301	001	
MUTUAL PHARM	5MG	N80701	001	
	10MG	N86595	001	
	20MG	N84634	001	
	50MG	N86596	001	
NYLOS	5MG	N85115	001	
PANRAY	1MG	N80350	001	
	2.5MG	N80350	002	
	5MG	N80350	003	
PERRIGO	5MG	N83059	001	
PHARMAVITE	5MG	N84662	002	
PHOENIX LABS NY	5MG	N80321	001	
	20MG	N83807	001	
PUREPAC PHARM	5MG	N80353	001	
	10MG	N86062	001	
	20MG	N86061	001	
PVT FORM	20MG	N85151	001	
REXALL	5MG	N80232	001	
ROXANE	20MG	N17109	001	
	25MG	N87833	001	May 04, 1982
SANDOZ	5MG	N80336	002	
	5MG	N84774	001	
	20MG	N85813	001	
SCHERER LABS	5MG	N80371	001	
SPERTI	1MG	N80359	001	
	2.5MG	N80359	002	
	5MG	N80359	003	
SUPERPHARM	5MG	N88865	001	Oct 25, 1984
	10MG	N88866	001	Oct 25, 1984
	20MG	N88867	001	Oct 25, 1984
TEVA	5MG	N80397	001	
UDL	5MG	N87984	001	Jan 18, 1983
	10MG	N87985	001	Jan 18, 1983
	20MG	N87986	001	Jan 18, 1983
UPSHER SMITH	5MG	N87471	001	
	20MG	N87470	001	
VALEANT PHARM INTL	5MG	N80237	001	
VANGARD	5MG	N87682	001	Jan 15, 1982
	20MG	N87701	001	Jan 15, 1982
VITARINE	5MG	N80334	001	
	5MG	N80506	001	
WATSON LABS	5MG	N85084	002	
	50MG	N86867	001	
WEST WARD	50MG	N88465	001	Jun 01, 1984
WHITEWORTH TOWN PLSN	2.5MG	N84913	001	
	5MG	N80343	001	
	10MG	N89028	001	Jul 24, 1986
	20MG	N84913	002	
SERVISONE				
LEDERLE	5MG	N80223	001	

PRILOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION
CITANEST

ASTRAZENECA	1%	N14763	004
	2%	N14763	005
	3%	N14763	003
CITANEST PLAIN			
ASTRAZENECA	4%	N14763	007

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 212 (of 263)

PRIMIDONESUSPENSION; ORAL
MYSOLINE

XCEL PHARMS 250MG/5ML

N10401 001

TABLET; ORAL
PRIMIDONE

WATSON LABS 250MG

N85052 001

PROBENECIDTABLET; ORAL
BENEMID

MERCK 500MG

N07898 004

PROBENECID

LEDERLE 500MG

N86917 001

WATSON LABS 500MG

N86150 002

Apr 23, 1982

PROBUCOLTABLET; ORAL
LORELCO

AVENTIS PHARMS 250MG

N17535 001

500MG

N17535 002

Jul 06, 1988

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

ASCOT 250MG

N87542 001

Jan 08, 1982

375MG

N87697 001

Mar 01, 1983

500MG

N87543 001

Jan 08, 1982

LANNETT

250MG

N83693 001

500MG

N84696 001

LEDERLE

250MG

N86942 001

375MG

N86952 001

500MG

N86943 001

ROXANE

250MG

N88989 001

Apr 26, 1985

500MG

N88990 001

Apr 26, 1985

SANDOZ

250MG

N89219 001

Jul 01, 1986

375MG

N89220 001

Jul 01, 1986

500MG

N89221 001

Jul 01, 1986

VANGARD

250MG

N87643 001

Jun 01, 1982

500MG

N87875 001

Jun 01, 1982

WATSON LABS

250MG

N83795 001

250MG

N85167 001

375MG

N87020 001

500MG

N84357 001

500MG

N87021 001

PROCAN

PARKE DAVIS

250MG

N85804 001

375MG

N87502 001

500MG

N85079 001

PROCAPAN

PANRAY

250MG

N83553 002

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

AM PHARM PARTNERS 100MG/ML

N89415 001

Nov 17, 1986

500MG/ML

N89416 001

Nov 17, 1986

BAXTER HLTHCARE

100MG/ML

N89029 001

Apr 17, 1986

500MG/ML

N89030 001

Apr 17, 1986

PHARMAFAIR

100MG/ML

N88824 001

Nov 20, 1985

500MG/ML

N88830 001

Nov 20, 1985

SMITH AND NEPHEW

100MG/ML

N88530 001

Mar 04, 1985

500MG/ML

N88531 001

Mar 04, 1985

SOLOPAK

500MG/ML

N88532 001

Mar 04, 1985

STERIS

100MG/ML

N87079 001

500MG/ML

N87080 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 213 (of 263)

PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HYDROCHLORIDE

WARNER CHILCOTT	100MG/ML	N89528	001	May 03, 1988
	500MG/ML	N89529	001	May 03, 1988

TABLET, EXTENDED RELEASE; ORAL

PROCAINAMIDE HYDROCHLORIDE

COPLEY PHARM	500MG	N88974	001	Jul 22, 1985
	750MG	N89438	001	Mar 23, 1987
	1GM	N40111	001	Dec 13, 1996
INWOOD LABS	500MG	N89840	001	Mar 06, 1989
PLIVA	250MG	N88958	001	Dec 02, 1985
	500MG	N88959	001	Dec 02, 1985
SANDOZ	250MG	N89369	001	Aug 14, 1987
	500MG	N89284	001	Jun 23, 1986
	500MG	N89370	001	Jan 09, 1987
	750MG	N89371	001	Aug 14, 1987
WATSON LABS	250MG	N88533	001	Dec 03, 1984
	250MG	N89026	001	Oct 22, 1985
	500MG	N88534	001	Dec 03, 1984
	500MG	N89027	001	Oct 22, 1985
	750MG	N88535	001	Nov 03, 1984
	750MG	N89042	001	Oct 22, 1985
	1GM	N89520	001	Jan 15, 1987

PROCAN SR

PARKE DAVIS	250MG	N86468	001	
PARKEDALE	500MG	N86065	001	
	750MG	N87510	001	Apr 01, 1982
	1GM	N88489	001	Jan 16, 1985

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE

AM PHARM PARTNERS	1%	N80384	002	
	1%	N80421	001	
	2%	N80384	003	
	2%	N80421	002	
BEL MAR	1%	N80711	001	
	2%	N80756	001	
ELKINS SINN	1%	N83315	001	
	2%	N83315	002	
GD SEARLE LLC	1%	N86202	001	
	2%	N86202	002	
MILES	1%	N80415	001	
	2%	N80415	002	
STERIS	1%	N83535	001	
	2%	N80658	002	
	2%	N83535	002	

PROCAINE HYDROCHLORIDE; TETRACYCLINE HYDROCHLORIDE

INJECTABLE; INJECTION

ACHROMYCIN

LEDERLE	40MG/VIAL;100MG/VIAL	N50276	001	
	40MG/VIAL;250MG/VIAL	N50276	003	

TETRACYN

PFIZER	40MG/VIAL;100MG/VIAL	N60285	002	
	40MG/VIAL;250MG/VIAL	N60285	003	

PROCAINE MERETHOXYLLINE; THEOPHYLLINE

INJECTABLE; INJECTION

DICURIN PROCAINE

LILLY	100MG/ML;50MG/ML	N08869	001	
-------	------------------	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 214 (of 263)

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

PROCHLORPERAZINE

ABLE	2.5MG	N40407	001	Jul 11, 2001
	5MG	N40407	002	Jul 11, 2001
	25MG	N40407	003	Jul 11, 2001

PROCHLORPERAZINE EDISYLATE

CONCENTRATE; ORAL

COMPAZINE

GLAXOSMITHKLINE	EQ 10MG BASE/ML	N11276	001	
-----------------	-----------------	--------	-----	--

PROCHLORPERAZINE

ALPHARMA US PHARMS	EQ 10MG BASE/ML	N87153	001	Jun 08, 1982
--------------------	-----------------	--------	-----	--------------

PROCHLORPERAZINE EDISYLATE

MORTON GROVE	EQ 10MG BASE/ML	N88598	001	Oct 25, 1984
--------------	-----------------	--------	-----	--------------

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

BAXTER HLTHCARE	EQ 5MG BASE/ML	N89523	001	May 03, 1988
-----------------	----------------	--------	-----	--------------

MARSAM PHARMS LLC	EQ 5MG BASE/ML	N89675	001	Dec 05, 1988
-------------------	----------------	--------	-----	--------------

SMITH AND NEPHEW	EQ 5MG BASE/ML	N89251	001	Dec 04, 1986
------------------	----------------	--------	-----	--------------

STERIS	EQ 5MG BASE/ML	N89605	001	Jul 08, 1987
--------	----------------	--------	-----	--------------

	EQ 5MG BASE/ML	N89606	001	Jul 08, 1987
--	----------------	--------	-----	--------------

WYETH AYERST	EQ 5MG BASE/ML	N86348	001	
--------------	----------------	--------	-----	--

SYRUP; ORAL

PROCHLORPERAZINE EDISYLATE

ALPHARMA US PHARMS	EQ 5MG BASE/5ML	N87154	001	Sep 01, 1982
--------------------	-----------------	--------	-----	--------------

MORTON GROVE	EQ 5MG BASE/5ML	N88597	001	Oct 25, 1984
--------------	-----------------	--------	-----	--------------

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

COMPAZINE

GLAXOSMITHKLINE	EQ 10MG BASE	N11000	001	
-----------------	--------------	--------	-----	--

	EQ 15MG BASE	N11000	002	
--	--------------	--------	-----	--

	EQ 30MG BASE	N11000	003	
--	--------------	--------	-----	--

	EQ 75MG BASE	N11000	004	
--	--------------	--------	-----	--

TABLET; ORAL

PROCHLORPERAZINE

WATSON LABS	EQ 5MG BASE	N85580	001	
-------------	-------------	--------	-----	--

	EQ 10MG BASE	N85178	001	
--	--------------	--------	-----	--

	EQ 25MG BASE	N85579	001	
--	--------------	--------	-----	--

PROCHLORPERAZINE MALEATE

DURAMED PHARMS BARR	EQ 5MG BASE	N89484	001	Jan 20, 1987
---------------------	-------------	--------	-----	--------------

	EQ 10MG BASE	N89485	001	Jan 20, 1987
--	--------------	--------	-----	--------------

	EQ 25MG BASE	N89486	001	Jan 20, 1987
--	--------------	--------	-----	--------------

PROCYCLIDINE HYDROCHLORIDE

TABLET; ORAL

KEMADRIN

MONARCH PHARMS	2MG	N09818	005	
----------------	-----	--------	-----	--

PROGESTERONE

CAPSULE; ORAL

PROMETRIUM

UNIMED PHARMS	300MG	N19781	003	Oct 15, 1999
---------------	-------	--------	-----	--------------

INJECTABLE; INJECTION

PROGESTERONE

LILLY	25MG/ML	N09238	002	
-------	---------	--------	-----	--

	50MG/ML	N09238	001	
--	---------	--------	-----	--

INSERT, EXTENDED RELEASE; INTRAUTERINE

PROGESTASERT

ALZA	38MG	N17553	001	
------	------	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 215 (of 263)

PROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

SPARINE

WYETH AYERST

30MG/ML

N10942 001

100MG/ML

N10942 004

INJECTABLE; INJECTION

PROMAZINE HYDROCHLORIDE

STERIS

25MG/ML

N84510 001

50MG/ML

N84517 001

SPARINE

BAXTER HLTHCARE CORP

25MG/ML

N10349 008

50MG/ML

N10349 006

SYRUP; ORAL

SPARINE

WYETH AYERST

10MG/5ML

N10942 003

TABLET; ORAL

SPARINE

WYETH AYERST

10MG

N10348 006

25MG

N10348 001

50MG

N10348 002

100MG

N10348 003

200MG

N10348 004

PROMETHAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PHENERGAN

WYETH AYERST

25MG/ML

N08857 002

50MG/ML

N08857 003

PROMETHAZINE HYDROCHLORIDE

ABBOTT

25MG/ML

N84223 001

50MG/ML

N84222 001

AKORN

25MG/ML

N83955 002

50MG/ML

N83955 001

MARSAM PHARMS LLC

25MG/ML

N89463 001

May 02, 1988

50MG/ML

N89477 001

May 02, 1988

STERIS

25MG/ML

N83532 001

50MG/ML

N83532 002

ZIPAN-25

ALTANA

25MG/ML

N83997 001

ZIPAN-50

ALTANA

50MG/ML

N83997 002

SUPPOSITORY; RECTAL

PHENERGAN

WYETH PHARMS INC

50MG

N11689 001

PROMETHACON

POLYMEDICA

50MG

N84902 001

PROMETHAZINE HYDROCHLORIDE

ABLE

12.5MG

N40504 001

Apr 11, 2003

25MG

N40504 002

Apr 11, 2003

50MG

N40449 001

Feb 27, 2003

SYRUP; ORAL

MYMETHAZINE FORTIS

USL PHARMA

25MG/5ML

N87996 001

Jan 18, 1983

PHENERGAN FORTIS

WYETH AYERST

25MG/5ML

N08381 003

PHENERGAN PLAIN

WYETH AYERST

6.25MG/5ML

N08381 004

Apr 18, 1984

PROMETH FORTIS

ALPHARMA US PHARMS

25MG/5ML

N84772 001

PROMETH PLAIN

ALPHARMA US PHARMS

6.25MG/5ML

N85953 001

PROMETHAZINE

CENCI

6.25MG/5ML

N89013 001

Sep 20, 1985

PROMETHAZINE HYDROCHLORIDE

KV PHARM

6.25MG/5ML

N85388 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 216 (of 263)

PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETHAZINE HYDROCHLORIDE

KV PHARM 25MG/5ML N85385 001

PHARM ASSOC 6.25MG/5ML N87518 001

WHITEWORTH TOWN PLSN 6.25MG/5ML N86395 001

TABLET; ORAL

PHENERGAN

WYETH PHARMS INC 12.5MG N07935 002

50MG N07935 004

PROMETHAZINE HYDROCHLORIDE

ABBOTT 12.5MG N84160 001

25MG N84166 001

50MG N84539 001

ABLE 12.5MG N40558 001 Jul 01, 2004

25MG N40558 002 Jul 01, 2004

50MG N40558 003 Jul 01, 2004

IMPAX LABS 25MG N84214 002 Jul 07, 1982

IVAX PHARMS 12.5MG N83604 001

25MG N83603 001

50MG N83613 001

LANNETT 12.5MG N80949 001

25MG N80949 002

50MG N80949 003

MUTUAL PHARM 12.5MG N84555 001

25MG N84554 001

50MG N84557 001

PVT FORM 12.5MG N83214 001

25MG N83658 001

SANDOZ 12.5MG N84233 001

25MG N85146 001

50MG N85146 002

TABLICAPS 12.5MG N84080 001

25MG N84027 001

TEVA 25MG N89109 001 Sep 10, 1985

WATSON LABS 12.5MG N83401 001

12.5MG N83712 001

12.5MG N85986 001

25MG N83204 001

25MG N85684 001

50MG N83403 001

50MG N85664 001

REMSED

BRISTOL MYERS SQUIBB 25MG N83176 002

50MG N83176 001

PROPANTHELINE BROMIDE

INJECTABLE; INJECTION

PRO-BANTHINE

GD SEARLE LLC 30MG/VIAL N08843 001

TABLET; ORAL

PROPANTHELINE BROMIDE

ASCOT 15MG N87663 001 Oct 25, 1982

HEATHER 15MG N85780 001

IMPAX LABS 15MG N84541 002

MYLAN 15MG N83706 001

PAR PHARM 15MG N88377 001 Dec 08, 1983

PVT FORM 15MG N80977 001

SANDOZ 15MG N80928 001

TABLICAPS 15MG N84428 001

WATSON LABS 15MG N83029 002

15MG N83151 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 217 (of 263)

PROPARACAINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

KAINAIR

PHARMAFAIR 0.5%

N88087 001 Jun 07, 1983

PROPARACAINE HYDROCHLORIDE

SOLA BARNES HIND 0.5%

N84144 001

0.5%

N84151 001

PROPIOLACTONE

SOLUTION; IRRIGATION

BETAPRONE

FOREST LABS N/A

N11657 001

PROPIOMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

LARGON

BAXTER HLTHCARE CORP 20MG/ML

N12382 002

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

ASTRAZENECA 10MG/ML

N19627 001 Oct 02, 1989

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON

XANODYNE PHARM 32MG

N10997 001

DOLENE

CLONMEL HLTHCARE 65MG

N80530 001

KESSO-GESIC

MK LABS 65MG

N83544 001

PROPHENE 65

HALSEY 65MG

N83538 002

PROPOXYPHENE HYDROCHLORIDE

ALRA 65MG

N83184 001

ANABOLIC 65MG

N83185 001

IMPAX LABS 65MG

N83317 001

IVAX PHARMS 32MG

N83597 001

MUTUAL PHARM 65MG

N83186 001

MYLAN 32MG

N83528 001

65MG

N83299 001

PUREPAC PHARM 65MG

N83278 001

PVT FORM 32MG

N83464 001

65MG

N83113 001

ROXANE 32MG

N83089 001

65MG

N83089 002

SANDOZ 32MG

N84014 001

65MG

N83125 002

65MG

N83688 001

65MG

N83870 002

65MG

N86495 001

VALEANT PHARM INTL 65MG

N80783 001

WATSON LABS 65MG

N80908 002

65MG

N85190 001

WHITEWORTH TOWN PLSN 65MG

N84551 001

PROPOXYPHENE HYDROCHLORIDE 65

WARNER CHILCOTT 65MG

N83786 001

PROPOXYPHENE NAPSYLATE

SUSPENSION; ORAL

DARVON-N

AAIPHARMA LLC 50MG/5ML

N16861 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 218 (of 263)

PROPRANOLOL HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
PROPRANOLOL HYDROCHLORIDE

INWOOD LABS	60MG	N72499	001	Apr 11, 1989
	80MG	N72500	001	Apr 11, 1989
	120MG	N72501	001	Apr 11, 1989
	160MG	N72502	001	Apr 11, 1989

INJECTABLE; INJECTION
PROPRANOLOL HYDROCHLORIDE

SMITH AND NEPHEW	1MG/ML	N70135	001	Apr 15, 1986
	1MG/ML	N70137	001	Apr 15, 1986
SOLOPAK	1MG/ML	N70136	001	Apr 15, 1986

SOLUTION; ORAL
PROPRANOLOL HYDROCHLORIDE

MORTON GROVE	20MG/5ML	N71984	001	Mar 03, 1989
	40MG/5ML	N71985	001	Mar 03, 1989

SUSPENSION; ORAL

INDERAL

WYETH AYERST	10MG/ML	N19536	001	Dec 12, 1986
--------------	---------	--------	-----	--------------

TABLET; ORAL

INDERAL

WYETH PHARMS INC	90MG	N16418	010	Oct 18, 1982
------------------	------	--------	-----	--------------

PROPRANOLOL HYDROCHLORIDE

CLONMEL HLTHCARE	10MG	N70125	001	Jul 30, 1985
	20MG	N70126	001	Jul 30, 1985
	40MG	N70127	001	Jul 30, 1985
	60MG	N71495	001	Dec 31, 1987
	80MG	N70128	001	Jul 30, 1985
	90MG	N71496	001	Dec 31, 1987
DURAMED PHARMS BARR	10MG	N70306	001	Sep 09, 1985
	20MG	N70307	001	Sep 09, 1985
	40MG	N70308	001	Sep 09, 1985
	60MG	N70309	001	Oct 01, 1986
	80MG	N70310	001	Sep 09, 1985
	90MG	N71327	001	Oct 01, 1986
INTERPHARM	10MG	N71368	001	May 05, 1987
	20MG	N71369	001	May 05, 1987
	40MG	N71370	001	May 05, 1987
	80MG	N71371	001	May 05, 1987
IVAX PHARMS	10MG	N72063	001	Jul 29, 1988
	20MG	N72066	001	Jul 29, 1988
	40MG	N72067	001	Jul 29, 1988
	60MG	N72068	001	Jul 29, 1988
	80MG	N72069	001	Jul 29, 1988
LEDERLE	10MG	N72117	001	Jun 23, 1988
	20MG	N72118	001	Jun 23, 1988
	40MG	N72119	001	Jun 23, 1988
	80MG	N72120	001	Jun 23, 1988
MUTUAL PHARM	10MG	N70319	001	Oct 22, 1985
	20MG	N70320	001	Oct 22, 1985
	40MG	N70103	001	Oct 22, 1985
	60MG	N70321	001	Sep 24, 1986
	80MG	N70322	001	Aug 04, 1986
MYLAN	60MG	N72275	001	Jun 09, 1989
PUREPAC PHARM	10MG	N70814	001	Nov 03, 1986
	20MG	N70815	001	Nov 03, 1986
	40MG	N70816	001	Nov 03, 1986
	60MG	N70817	001	Nov 03, 1986
	80MG	N70757	001	Nov 03, 1986
ROXANE	10MG	N70516	001	Jul 07, 1986
	20MG	N70517	001	Jul 07, 1986
	40MG	N70518	001	Jul 07, 1986
	60MG	N70519	001	Sep 24, 1986
	80MG	N70520	001	Jul 07, 1986
	90MG	N70521	001	Sep 24, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 219 (of 263)

PROPRANOLOL HYDROCHLORIDE

TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE

SANDOZ	10MG	N71658	001	Jul 05, 1988	
	20MG	N71687	001	Jul 05, 1988	
	40MG	N71688	001	Jul 05, 1988	
	60MG	N72197	001	Jul 05, 1988	
	80MG	N71689	001	Jul 05, 1988	
	90MG	N72198	001	Jul 05, 1988	
	SCHERING	10MG	N70120	001	Aug 06, 1985
		20MG	N70121	001	Aug 06, 1985
		40MG	N70122	001	Aug 06, 1985
		60MG	N70123	001	Oct 29, 1986
80MG		N70124	001	Aug 06, 1985	
SUPERPHARM	10MG	N71515	001	Jun 08, 1988	
	20MG	N71516	001	Jun 08, 1988	
	40MG	N71517	001	Jun 08, 1988	
	80MG	N71518	001	Jun 08, 1988	
TEVA	10MG	N70232	001	Oct 07, 1987	
	20MG	N70233	001	Jun 23, 1986	
WARNER CHILCOTT	10MG	N70438	001	Sep 15, 1986	
	20MG	N70439	001	Sep 15, 1986	
	40MG	N70440	001	Sep 15, 1986	
	60MG	N70441	001	Sep 24, 1986	
WATSON LABS	80MG	N70442	001	Sep 15, 1986	
	10MG	N70140	001	Jul 30, 1985	
	10MG	N70378	001	Mar 19, 1987	
	20MG	N70141	001	Jul 30, 1985	
	20MG	N70379	001	Mar 19, 1987	
	20MG	N70549	001	Apr 11, 1986	
	40MG	N70142	001	Jul 30, 1985	
	40MG	N70380	001	Mar 19, 1987	
	60MG	N70143	001	Jan 15, 1987	
	60MG	N70381	001	Mar 19, 1987	
	60MG	N71098	001	Oct 06, 1986	
	80MG	N70144	001	Jul 30, 1985	
	80MG	N70382	001	Mar 19, 1987	
	90MG	N71183	001	Oct 06, 1986	

PROPYLIODONE

SUSPENSION; INTRATRACHEAL

DIONOSIL AQUEOUS

GLAXOSMITHKLINE	50%	N09309	001
-----------------	-----	--------	-----

DIONOSIL OILY

GLAXOSMITHKLINE	60%	N09309	002
-----------------	-----	--------	-----

PROPYLTHIOURACIL

TABLET; ORAL

PROPYLTHIOURACIL

ABBOTT	50MG	N84075	001
ANABOLIC	50MG	N80285	001
HALSEY	50MG	N80015	001
IVAX PHARMS	50MG	N80215	001
LANNETT	50MG	N80016	001
LILLY	50MG	N06213	001
MUTUAL PHARM	50MG	N83982	001
PERRIGO	50MG	N84543	001
TABLICAPS	50MG	N80840	001
WATSON LABS	50MG	N80932	001
	50MG	N85201	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 220 (of 263)

PROTAMINE SULFATEINJECTABLE; INJECTION
PROTAMINE SULFATE

BAXTER HLTHCARE	10MG/ML	N89474	001	Nov 05, 1986
	10MG/ML	N89475	001	Nov 05, 1986
LILLY	10MG/ML	N06460	002	
PHARMACIA AND UPJOHN	50MG/VIAL	N07413	001	
	250MG/VIAL	N07413	002	Aug 02, 1984

PROTEIN HYDROLYSATEINJECTABLE; INJECTION
AMINOSOL 5%

ABBOTT	5%	N05932	012	Jan 31, 1985
HYPROTIGEN 5%				
B BRAUN	5%	N06170	003	Jan 10, 1984

PROTIRELININJECTABLE; INJECTION
THYPINONE

ABBOTT	0.5MG/ML	N17638	001	
THYREL TRH				
FERRING	0.5MG/ML	N18087	001	

PROTOKYLOL HYDROCHLORIDETABLET; ORAL
VENTAIRE

AVENTIS PHARMS	2MG	N83459	001	
----------------	-----	--------	-----	--

PROTRIPTYLINE HYDROCHLORIDETABLET; ORAL
VIVACTIL

ODYSSEY PHARMS	5MG	N16012	001	
	10MG	N16012	002	

PSEUDOEPHEDRINE HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
NOVAFED

AVENTIS PHARMS	120MG	N17603	001	
----------------	-------	--------	-----	--

SUDAFED 12 HOUR

GLAXOSMITHKLINE	120MG **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N17941	002	
-----------------	--	--------	-----	--

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDECAPSULE, EXTENDED RELEASE; ORAL
ACTIFED

GLAXOSMITHKLINE	120MG;5MG	N18996	001	Jun 17, 1985
-----------------	-----------	--------	-----	--------------

TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES

KV PHARM	120MG;5MG	N71798	001	Mar 16, 1989
----------	-----------	--------	-----	--------------

SYRUP; ORAL

ACTAHIST

CENCI	30MG/5ML;1.25MG/5ML	N88344	001	Feb 09, 1984
-------	---------------------	--------	-----	--------------

HISTAFED

CENCI	30MG/5ML;1.25MG/5ML	N88283	001	Apr 20, 1984
-------	---------------------	--------	-----	--------------

MYFED

USL PHARMA	30MG/5ML;1.25MG/5ML	N88116	001	Mar 04, 1983
------------	---------------------	--------	-----	--------------

TRILITRON

NEWTRON PHARMS	30MG/5ML;1.25MG/5ML	N88474	001	Feb 12, 1985
----------------	---------------------	--------	-----	--------------

TABLET; ORAL

ALLERFED

PVT FORM	60MG;2.5MG	N88860	001	Jan 31, 1985
----------	------------	--------	-----	--------------

TRILITRON

NEWTRON PHARMS	60MG;2.5MG	N88515	001	Jan 09, 1985
----------------	------------	--------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 221 (of 263)

PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

TABLET; ORAL		
TRIPHED		
TEVA	60MG;2.5MG	N88630 001 May 17, 1984
TRIPROLIDINE AND PSEUDOEPHEDRINE		
WATSON LABS	60MG;2.5MG	N88318 002 Jan 13, 1984
WEST WARD	60MG;2.5MG	N88117 001 Apr 19, 1983
TRIPROLIDINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE		
IVAX PHARMS	60MG;2.5MG	N85273 001 Dec 12, 1984
SUPERPHARM	60MG;2.5MG	N88578 001 Feb 21, 1985
TABLET, EXTENDED RELEASE; ORAL		
TRIPROLIDINE AND PSEUDOEPHEDRINE HYDROCHLORIDES		
KV PHARM	120MG;5MG	N72758 001 Nov 25, 1991

PSEUDOEPHEDRINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL		
PSEUDO-12		
UCB	EQ 60MG HCL/5ML	N19401 001 Jun 19, 1987

PYRIDOSTIGMINE BROMIDE

TABLET; ORAL		
PYRIDOSTIGMINE BROMIDE		
SOLVAY	30MG	N89572 001 Nov 27, 1990
US ARMY	30MG	N20414 001 Feb 05, 2003

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION		
HEXA-BETALIN		
LILLY	100MG/ML	N80854 001
PYRIDOXINE HYDROCHLORIDE		
AKORN	100MG/ML	N87967 001 Oct 01, 1982
BEL MAR	100MG/ML	N80761 001
DELL LABS	50MG/ML	N83771 001
	100MG/ML	N83772 001
ELKINS SINN	100MG/ML	N80581 001
LUITPOLD	100MG/ML	N80669 001
STERIS	100MG/ML	N83760 001

PYRILAMINE MALEATE

TABLET; ORAL		
PYRILAMINE MALEATE		
IMPAX LABS	25MG	N80808 001
WATSON LABS	25MG	N85231 001

PYRITHIONE ZINC

LOTION; TOPICAL		
HEAD & SHOULDERS CONDITIONER		
PROCTER AND GAMBLE	0.3%	N19412 002 Mar 10, 1986

PYRVINIUM PAMOATE

SUSPENSION; ORAL		
POVAN		
PARKE DAVIS	EQ 50MG BASE/5ML	N11964 001
TABLET; ORAL		
POVAN		
PARKE DAVIS	EQ 50MG BASE	N12485 002

QUAZEPAM

TABLET; ORAL		
DORAL		
MEDPOINTE PHARM HLC	7.5MG	N18708 003 Feb 26, 1987

DISCONTINUED DRUG PRODUCT LIST

6 - 222 (of 263)

QUETIAPINE FUMARATE

TABLET; ORAL SEROQUEL				
ASTRAZENECA	EQ 150MG BASE	N20639	004	Dec 20, 1998

QUINESTROL

TABLET; ORAL ESTROVIS				
PARKE DAVIS	0.1MG	N16768	002	
	0.2MG	N16768	003	

QUINETHAZONE

TABLET; ORAL HYDROMOX				
LEDERLE	50MG	N13264	001	

QUINETHAZONE; RESERPINE

TABLET; ORAL HYDROMOX R				
LEDERLE	50MG;0.125MG	N13927	001	

QUINIDINE GLUCONATE

TABLET; ORAL QUINACT				
BERLEX	266MG	N85978	001	
	400MG	N86099	001	
TABLET, EXTENDED RELEASE; ORAL				
DURAQUIN				
WARNER CHILCOTT	330MG	N17917	001	
QUINAGLUTE				
BERLEX	324MG	N16647	001	
QUINALAN				
LANNETT	324MG	N88081	001	Feb 10, 1986
QUINATIME				
WATSON LABS	324MG	N87448	001	
QUINIDINE GLUCONATE				
ASCOT	324MG	N88582	001	Jun 17, 1985
HALSEY	324MG	N89476	001	Apr 10, 1987
ROXANE	324MG	N88431	001	Jan 06, 1984
SANDOZ	324MG	N89894	001	Dec 15, 1988
SUPERPHARM	324MG	N89164	001	Nov 21, 1985
WATSON LABS	324MG	N87785	001	Jan 24, 1983
	324MG	N87810	001	Sep 29, 1982

QUINIDINE POLYGALACTURONATE

TABLET; ORAL CARDIOQUIN				
PURDUE FREDERICK	275MG	N11642	002	

QUINIDINE SULFATE

CAPSULE; ORAL CIN-QUIN				
SOLVAY	200MG	N85296	001	
	300MG	N85297	001	
QUINIDINE SULFATE				
LILLY	200MG	N85103	001	
TABLET; ORAL				
CIN-QUIN				
SOLVAY	100MG	N85299	001	
	200MG	N84932	001	
	300MG	N85298	001	
QUINIDINE SULFATE				
BARR	200MG	N84177	001	
ELKINS SINN	200MG	N83622	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 223 (of 263)

QUINIDINE SULFATE

TABLET; ORAL

QUINIDINE SULFATE

EVERYLIFE	200MG	N83439	001	
HALSEY	200MG	N83583	001	
IMPAX LABS	200MG	N83347	001	
IVAX PHARMS	200MG	N84549	001	
KING PHARMS	200MG	N85175	001	
KV PHARM	200MG	N85276	001	
LEDERLE	200MG	N86176	001	
LILLY	200MG	N85038	001	
MUTUAL PHARM	300MG	N81031	001	Apr 14, 1989
PERRIGO	200MG	N85322	001	
PHARMAVITE	200MG	N84627	001	
PUREPAC PHARM	200MG	N84003	001	
ROXANE	200MG	N83640	001	
	300MG	N85632	001	
SCHERER LABS	200MG	N85068	001	
SUPERPHARM	200MG	N88973	001	Apr 10, 1985
USL PHARMA	200MG	N87837	001	Apr 14, 1982
VALEANT PHARM INTL	200MG	N83393	001	
VANGARD	200MG	N87909	001	Jul 13, 1982
VINTAGE PHARMS	200MG	N83963	001	
WARNER CHILCOTT	200MG	N83879	001	
WATSON LABS	100MG	N85584	001	
WEST WARD	200MG	N83862	001	
WHITEWORTH TOWN PLSN	200MG	N85444	001	
QUINORA				
KEY PHARMS	200MG	N83576	001	
SCHERING	300MG	N85222	001	
TABLET, EXTENDED RELEASE; ORAL				
QUINIDEX				
WYETH PHARMS INC	300MG	N12796	002	

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE; ORAL

ACIPHEX

EISAI MEDCL RES	10MG **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N20973	001	May 29, 2002
-----------------	---	--------	-----	--------------

RANITIDINE BISMUTH CITRATE

TABLET; ORAL

TRITEC

GLAXOSMITHKLINE	400MG	N20559	001	Aug 08, 1996
-----------------	-------	--------	-----	--------------

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

ZANTAC 150

GLAXOSMITHKLINE	EQ 150MG BASE	N20095	001	Mar 08, 1994
-----------------	---------------	--------	-----	--------------

ZANTAC 300

GLAXOSMITHKLINE	EQ 300MG BASE	N20095	002	Mar 08, 1994
-----------------	---------------	--------	-----	--------------

INJECTABLE; INJECTION

ZANTAC IN PLASTIC CONTAINER

GLAXOSMITHKLINE	EQ 50MG BASE/100ML	N19593	001	Dec 17, 1986
-----------------	--------------------	--------	-----	--------------

TABLET; ORAL

RANITIDINE

RANBAXY	EQ 75MG BASE	N75254	001	Jan 14, 2000
---------	--------------	--------	-----	--------------

RANITIDINE HYDROCHLORIDE

BOEHRINGER INGELHEIM	EQ 150MG BASE	N74662	001	Aug 29, 1997
----------------------	---------------	--------	-----	--------------

	EQ 300MG BASE	N74662	002	Aug 29, 1997
--	---------------	--------	-----	--------------

RANBAXY	EQ 150MG BASE	N75000	001	Jan 30, 1998
---------	---------------	--------	-----	--------------

	EQ 300MG BASE	N75000	002	Jan 30, 1998
--	---------------	--------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 224 (of 263)

RANITIDINE HYDROCHLORIDE

TABLET, EFFERVESCENT; ORAL
ZANTAC 75

WARNER LAMBERT	EQ 75MG BASE **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N20745 001	Feb 26, 1998
----------------	---	------------	--------------

RAPACURONIUM BROMIDE

INJECTABLE; INJECTION
RAPLON

ORGANON USA INC	100MG/VIAL	N20984 001	Aug 18, 1999
	200MG/VIAL	N20984 002	Aug 18, 1999

RAUWOLFIA SERPENTINA

TABLET; ORAL

HIWOLFIA

BOWMAN PHARMS	50MG	N09276 003	
	50MG	N09276 005	
	100MG	N09276 004	

HYSERPIN

PHYS PRODS VA	50MG	N10581 001	
---------------	------	------------	--

KOGLUCOID

PANRAY	50MG	N09278 001	
	100MG	N09278 002	

RAUDIXIN

APOTHECON	50MG	N08842 001	
	100MG	N08842 002	

RAUSERPIN

FERNDAL LABS	50MG	N09926 002	
	100MG	N09926 004	

RAUVAL

PAL PAK	50MG	N09108 002	
	100MG	N09108 004	

RAUWOLFIA SERPENTINA

BUNDY	50MG	N09477 001	
	100MG	N09477 002	

HALSEY	50MG	N80498 001	
--------	------	------------	--

	100MG	N80498 002	
--	-------	------------	--

IMPAX LABS	50MG	N09273 001	
------------	------	------------	--

	100MG	N09273 002	
--	-------	------------	--

IVAX PHARMS	50MG	N11521 001	
-------------	------	------------	--

	100MG	N11521 002	
--	-------	------------	--

PUREPAC PHARM	50MG	N80842 001	
---------------	------	------------	--

	100MG	N80842 002	
--	-------	------------	--

PVT FORM	50MG	N80583 001	
----------	------	------------	--

	100MG	N80583 002	
--	-------	------------	--

SOLVAY	50MG	N80500 001	
--------	------	------------	--

	100MG	N80500 002	
--	-------	------------	--

TABLICAPS	50MG	N83867 001	
-----------	------	------------	--

	100MG	N83444 001	
--	-------	------------	--

VALEANT PHARM INTL	50MG	N09668 001	
--------------------	------	------------	--

	100MG	N09668 002	
--	-------	------------	--

WATSON LABS	50MG	N80907 001	
-------------	------	------------	--

	100MG	N80914 001	
--	-------	------------	--

WOLFINA

FOREST PHARMS	50MG	N09255 008	
---------------	------	------------	--

	100MG	N09255 006	
--	-------	------------	--

RESCINNAMINE

CAPSULE; ORAL

CINNASIL

PANRAY	0.5MG	N84736 001	
--------	-------	------------	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 225 (of 263)

RESERPINE

ELIXIR; ORAL				
SERPASIL				
NOVARTIS	0.2MG/4ML	N09115	005	
INJECTABLE; INJECTION				
SANDRIL				
LILLY	2.5MG/ML	N10012	001	
SERPASIL				
NOVARTIS	2.5MG/ML	N09434	002	
TABLET; ORAL				
HISERPIA				
BOWMAN PHARMS	0.1MG	N09631	002	
	0.25MG	N09631	004	
RAU-SED				
BRISTOL MYERS SQUIBB	0.1MG	N09357	001	
	0.25MG	N09357	004	
	0.5MG	N09357	006	
	1MG	N09357	008	
RESERPINE				
BARR	0.25MG	N80721	002	
BELL PHARMA	0.1MG	N83058	001	
	0.25MG	N83058	002	
BUNDY	0.1MG	N09663	001	
	0.25MG	N09663	003	
ELKINS SINN	0.1MG	N83145	001	
	0.25MG	N83145	002	
EVERYLIFE	0.1MG	N10441	001	
	0.25MG	N10441	002	
	0.5MG	N10441	003	
	1MG	N10441	004	
HALSEY	0.1MG	N80457	002	
	0.25MG	N80457	001	
	1MG	N80457	003	
IMPAX LABS	0.1MG	N09627	001	
	0.25MG	N09627	002	
IVAX PHARMS	0.1MG	N11185	001	
	0.25MG	N11185	002	
MARSHALL PHARMA	0.1MG	N80492	001	
	0.25MG	N80492	002	
MK LABS	0.1MG	N80525	002	
	0.25MG	N80525	001	
MYLAN	1MG	N84974	001	
PHARMAVITE	0.25MG	N84663	001	
PUREPAC PHARM	0.1MG	N80753	002	
	0.25MG	N80753	001	
PVT FORM	0.1MG	N86117	001	
	0.25MG	N80582	001	
	0.25MG	N85775	001	
	1MG	N80582	002	
REXALL	0.25MG	N80637	001	
ROXANE	0.1MG	N09859	001	
	0.25MG	N09859	002	
SOLVAY	0.25MG	N80446	001	
TABLICAPS	0.25MG	N85207	001	
TEVA	0.1MG	N89020	001	Mar 07, 1985
	0.25MG	N89019	001	Mar 07, 1985
VALEANT PHARM INTL	0.1MG	N09667	001	
	0.25MG	N09667	002	
WATSON LABS	0.1MG	N80679	001	
	0.25MG	N80393	001	
	0.25MG	N85401	001	
	1MG	N80749	001	
WEST WARD	0.1MG	N80975	001	
	0.25MG	N80975	002	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 226 (of 263)

RESERPINETABLET; ORAL
RESERPINE

WEST WARD	1MG	N80975	003
WHITEWORTH TOWN PLSN	0.1MG	N80723	001
	0.25MG	N80723	002
	1MG	N80723	003
SANDRIL			
LILLY	0.1MG	N09376	004
	0.25MG	N09376	001
SERPANRAY			
PANRAY	0.1MG	N09391	001
	0.25MG	N09391	002
	1MG	N09391	004
SERPASIL			
NOVARTIS	0.1MG	N09115	001
	0.25MG	N09115	003
	1MG	N09115	004
SERPATE			
VALE	0.1MG	N09453	001
	0.25MG	N09453	002
SERPIVITE			
VITARINE	0.25MG	N09645	002

RESERPINE; TRICHLORMETHIAZIDE

TABLET; ORAL

METATENSIN #2			
AVENTIS PHARMS	0.1MG;2MG	N12972	001
METATENSIN #4			
AVENTIS PHARMS	0.1MG;4MG	N12972	002
NAQUIVAL			
SCHERING	0.1MG;4MG	N12265	003
TRICHLORMETHIAZIDE W/ RESERPINE			
WATSON LABS	0.1MG;4MG	N85248	001

RIBAVIRIN

TABLET; ORAL

COPEGUS			
ROCHE	400MG	N21511	002 Jun 21, 2005

RISPERIDONE

TABLET; ORAL

RISPERDAL

JANSSEN PHARMA	5MG	N20272	005 Dec 29, 1993
TABLET, ORALLY DISINTEGRATING; ORAL			
RISPERDAL			
JANSSEN PHARMA	3MG	N21444	004 Dec 23, 2004
	4MG	N21444	005 Dec 23, 2004

RITODRINE HYDROCHLORIDE

INJECTABLE; INJECTION

RITODRINE HYDROCHLORIDE

AM PHARM PARTNERS	10MG/ML	N71188	001 Jul 23, 1987
	15MG/ML	N71189	001 Jul 23, 1987
YUTOPAR			
ASTRAZENECA	10MG/ML	N18580	001
	15MG/ML	N18580	002
TABLET; ORAL			
YUTOPAR			
ASTRAZENECA	10MG	N18555	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 227 (of 263)

RITONAVIRCAPSULE; ORAL
NORVIR
ABBOTT

100MG

N20680 001 Mar 01, 1996

ROCURONIUM BROMIDEINJECTABLE; INJECTION
ZEMURON
ORGANON USA INC

10MG/ML

N20214 002 Mar 17, 1994

ROFECOXIBSUSPENSION; ORAL
VIOXX

MERCK

12.5MG/5ML
25MG/5MLN21052 001 May 20, 1999
N21052 002 May 20, 1999TABLET; ORAL
VIOXX

MERCK

12.5MG
25MG
50MGN21042 001 May 20, 1999
N21042 002 May 20, 1999
N21042 003 Feb 25, 2000ROSE BENGAL SODIUM, I-131INJECTABLE; INJECTION
ROBENGATOPE
BRACCO0.5mCi/VIAL
1mCi/VIAL
2mCi/VIALN16224 001
N16224 002
N16224 003SODIUM ROSE BENGAL I 131
SORIN

0.5mCi/ML

N17318 001

SAFFLOWER OILINJECTABLE; INJECTION
LIPOSYN 10%
ABBOTT
LIPOSYN 20%
ABBOTT10% (10GM/100ML)
20% (20GM/100ML)N18203 001
N18614 001SARALASIN ACETATEINJECTABLE; INJECTION
SARENIN

PROCTER AND GAMBLE

EQ 0.6MG BASE/ML

N18009 001

SECOBARBITAL SODIUMCAPSULE; ORAL
SECOBARBITAL SODIUMIVAX PHARMS
LANNETT100MG
50MG
100MGN85869 001
N85909 001
N85903 001

PARKE DAVIS

100MG

N84762 001

PUREPAC PHARM

100MG

N85867 001

VALEANT PHARM INTL

100MG

N85477 001

VITARINE

100MG

N85898 001

100MG

N86273 001

WHITEWORTH TOWN PLSN

100MG

N85798 001

WYETH AYERST

100MG

N86390 001

SODIUM SECOBARBITAL

ANABOLIC

100MG

N84422 001

BARR

100MG

N84225 001

HALSEY

100MG

N84676 001

KV PHARM

100MG

N85285 001

PERRIGO

100MG

N84561 001

WATSON LABS

100MG

N85792 001

WEST WARD

100MG

N84926 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 228 (of 263)

SECOBARBITAL SODIUM

INJECTABLE; INJECTION			
SECOBARBITAL SODIUM			
ELKINS SINN	100MG/VIAL	N83281	001
SECONAL SODIUM			
LILLY	50MG/ML	N07392	002
SODIUM SECOBARBITAL			
WYETH AYERST	50MG/ML	N83262	001
SUPPOSITORY; RECTAL			
SECONAL SODIUM			
LILLY	30MG	N86530	001
	60MG	N86530	002
	120MG	N86530	003
	200MG	N86530	004

SECRETIN

INJECTABLE; INJECTION			
SECRETIN-FERRING			
FERRING	75CU/VIAL	N18290	001

SELEGILINE HYDROCHLORIDE

TABLET; ORAL			
SELEGILINE HYDROCHLORIDE			
SOMERSET	5MG	N19334	001 Jun 05, 1989

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL			
EXSEL			
ALLERGAN HERBERT	2.5%	N83892	001
SELENIUM SULFIDE			
IVAX PHARMS	2.5%	N85777	001
TARO	2.5%	N86209	001

SELENOMETHIONINE, SE-75

INJECTABLE; INJECTION			
SELENOMETHIONINE SE 75			
CIS	500uCi/ML	N17322	001
GE HEALTHCARE	250uCi/ML	N17257	001
MALLINCKRODT	100uCi/ML	N17098	001
SETHOTOPE			
BRACCO	85-550uCi/ML	N17047	001

SERACTIDE ACETATE

INJECTABLE; INJECTION			
ACTHAR GEL-SYNTHETIC			
ARMOUR PHARM	40 UNITS/ML	N17861	001
	80 UNITS/ML	N17861	002

SERMORELIN ACETATE

INJECTABLE; INJECTION			
GEREF			
SERONO	EQ 0.5MG BASE/VIAL	N20443	001 Sep 26, 1997
	EQ 1MG BASE/VIAL	N20443	002 Sep 26, 1997

SERTRALINE HYDROCHLORIDE

TABLET; ORAL			
ZOLOFT			
PFIZER	EQ 150MG BASE **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N19839	003 Dec 30, 1991
	EQ 200MG BASE **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N19839	004 Dec 30, 1991

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 229 (of 263)

SEVELAMER HYDROCHLORIDE

CAPSULE; ORAL				
RENAGEL				
GENZYME	403MG	N20926	001	Oct 30, 1998

SILVER SULFADIAZINE

DRESSING; TOPICAL				
SILDAFLO				
FRANKLIN PHARMS	1%	N19608	001	Nov 30, 1989

SIMETHICONE-CELLULOSE

SUSPENSION; ORAL				
SONORX				
BRACCO	7.5MG/ML	N20773	001	Oct 29, 1998

SIROLIMUS

TABLET; ORAL				
RAPAMUNE				
WYETH PHARMS INC	5MG	N21110	003	Feb 23, 2004

SODIUM BENZOATE; SODIUM PHENYLACETATE

SOLUTION; ORAL				
UCEPHAN				
B BRAUN	100MG/ML;100MG/ML	N19530	001	Dec 23, 1987

SODIUM BICARBONATE

INJECTABLE; INJECTION				
SODIUM BICARBONATE IN PLASTIC CONTAINER				
ABBOTT	0.9MEQ/ML	N19443	001	Jun 03, 1986
	1MEQ/ML	N19443	002	Jun 03, 1986

SODIUM CHLORIDE

INJECTABLE; INJECTION				
BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
AM PHARM PARTNERS	9MG/ML	N88909	001	Feb 07, 1985
SODIUM CHLORIDE				
ABBOTT	20GM/100ML	N17013	001	
B BRAUN	20GM/100ML	N17038	001	
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
B BRAUN	450MG/100ML	N18184	001	
MILES	450MG/100ML	N18503	001	
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER				
ABBOTT	9MG/ML	N19218	001	Jul 13, 1984
MILES	900MG/100ML	N18502	001	
SODIUM CHLORIDE 23.4% IN PLASTIC CONTAINER				
AM PHARM PARTNERS	234MG/ML	N19329	001	Apr 22, 1987
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER				
B BRAUN	3GM/100ML	N19635	003	Mar 09, 1988
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER				
B BRAUN	5GM/100ML	N19635	004	Mar 09, 1988
SOLUTION; IRRIGATION				
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER				
BAXTER HLTHCARE	450MG/100ML	N18497	001	Feb 19, 1982
SODIUM CHLORIDE IN PLASTIC CONTAINER				
MILES	900MG/100ML	N18247	001	

SODIUM CHROMATE, CR-51

INJECTABLE; INJECTION				
CHROMITOPHE SODIUM				
BRACCO	2mCi/VIAL	N13993	002	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 230 (of 263)

SODIUM FLUORIDE, F-18INJECTABLE; INTRAVENOUS
FLUORINE F-18

GE HEALTHCARE

2mCi/ML **Federal Register
determination that product was not
discontinued or withdrawn for safety
or efficacy reasons**

N17042 001

SODIUM IODIDE, I-123

CAPSULE; ORAL

SODIUM IODIDE I 123

SYNCOR PHARMS

400uCi

N18671 003

May 27, 1982

SODIUM IODIDE, I-131

CAPSULE; ORAL

IODOTOPE

BRACCO

1-150mCi

N10929 003

SODIUM IODIDE I 131

CIS

50uCi

N17316 001

100uCi

N17316 002

MALLINCKRODT

0.8-100mCi

N16515 002

SOLUTION; ORAL

IODOTOPE

BRACCO

7-106mCi/BOT

N10929 002

SODIUM IODIDE I 131

CIS

50mCi/ML

N17315 001

SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER

B BRAUN

1.87GM/100ML

N18186 001

SODIUM NITROPRUSSIDE

INJECTABLE; INJECTION

NIPRIDE

ROCHE

50MG/VIAL

N17546 001

NITROPRESS

ABBOTT

50MG/VIAL

N18450 001

50MG/VIAL

N71555 001

Nov 16, 1987

SODIUM NITROPRUSSIDE

AM PHARM PARTNERS

50MG/VIAL

N70031 001

Jan 17, 1985

BAXTER HLTHCARE

50MG/VIAL

N18581 001

Jul 28, 1982

SODIUM PHOSPHATE, P-32

SOLUTION; INJECTION, ORAL

PHOSPHOTOPE

BRACCO

1-8mCi/VIAL

N10927 001

SODIUM PHOSPHATE P 32

MALLINCKRODT

1.5mCi/VIAL

N11777 002

SODIUM POLYSTYRENE SULFONATE

POWDER; ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

MORTON GROVE

453.6GM/BOT

N88786 001

Sep 11, 1984

SUSPENSION; ORAL, RECTAL

SODIUM POLYSTYRENE SULFONATE

MORTON GROVE

15GM/60ML

N88717 001

Sep 11, 1984

ROXANE

15GM/60ML

N88453 001

Nov 17, 1983

SODIUM SUCCINATE

INJECTABLE; INJECTION

SODIUM SUCCINATE

ELKINS SINN

30%

N80516 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 231 (of 263)

SODIUM TETRADECYL SULFATEINJECTABLE; INJECTION
SOTRADECOL

ELKINS SINN

1% **Federal Register determination
that product was not discontinued or
withdrawn for safety or efficacy
reasons**

N05970 004

3% **Federal Register determination
that product was not discontinued or
withdrawn for safety or efficacy
reasons**

N05970 005

SODIUM THIOSULFATEINJECTABLE; INJECTION
SODIUM THIOSULFATE

US ARMY

250MG/ML

N20166 001

Feb 14, 1992

SOMATREMINJECTABLE; INJECTION
PROTROPIN

GENENTECH

5MG/VIAL

N19107 001

Oct 17, 1985

10MG/VIAL

N19107 002

Oct 24, 1989

SOMATROPININJECTABLE; INJECTION
ASELLACRIN 10

SERONO

10 IU/VIAL

N17726 001

ASELLACRIN 2

SERONO

2 IU/VIAL

N17726 002

Jul 21, 1983

CRESCORMON

GENENTECH

4 IU/VIAL

N17992 001

SOMATROPIN RECOMBINANTINJECTABLE; INJECTION
BIO-TROPIN

FERRING

4.8MG/VIAL

N19774 001

May 25, 1995

HUMATROPE

LILLY

2MG/VIAL

N19640 001

Jun 23, 1987

SAIZEN

SERONO

6MG/VIAL

N19764 001

Oct 08, 1996

SEROSTIM

SERONO

8.8MG/VIAL

N20604 004

Sep 06, 2001

ZORBTIVE

SERONO INC

4MG/VIAL

N21597 001

Dec 01, 2003

5MG/VIAL

N21597 002

Dec 01, 2003

6MG/VIAL

N21597 003

Dec 01, 2003

INJECTABLE; SUBCUTANEOUS

SEROSTIM LQ

SERONO

6MG/0.5ML

N20604 005

Feb 11, 2005

SORBITOL

SOLUTION; IRRIGATION

SORBITOL 3% IN PLASTIC CONTAINER

BAXTER HLTHCARE

3GM/100ML

N18512 001

May 27, 1982

SOTALOL HYDROCHLORIDE

TABLET; ORAL

BETAPACE

BERLEX

320MG

N19865 004

Oct 30, 1992

BETAPACE AF

BERLEX

40MG

N21151 006

Apr 02, 2003

60MG

N21151 007

Apr 02, 2003

100MG

N21151 005

Mar 14, 2003

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 232 (of 263)

SOYBEAN OIL

INJECTABLE; INJECTION

TRAVAMULSION 10%

BAXTER HLTHCARE 10%

N18660 001 Feb 26, 1982

TRAVAMULSION 20%

BAXTER HLTHCARE 20%

N18758 001 Feb 15, 1983

SPARFLOXACIN

TABLET; ORAL

ZAGAM

MYLAN 200MG

N20677 001 Dec 19, 1996

SPECTINOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION

TROBICIN

PHARMACIA AND UPJOHN EQ 4GM BASE/VIAL

N50347 002

SPIRAPRIL HYDROCHLORIDE

TABLET; ORAL

RENORMAX

SCHERING 3MG

N20240 001 Dec 29, 1994

6MG

N20240 002 Dec 29, 1994

12MG

N20240 003 Dec 29, 1994

24MG

N20240 004 Dec 29, 1994

SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE

ASCOT 25MG

N87687 001 Oct 20, 1982

IVAX PHARMS 25MG

N87108 001

LEDERLE 25MG

N87634 001

PUREPAC PHARM 25MG

N87998 001 Oct 14, 1983

25MG

N88053 001 Aug 25, 1983

SUPERPHARM 25MG

N89364 001 Nov 07, 1986

UPSHER SMITH 25MG

N87554 001

VANGARD 25MG

N87648 001 Feb 01, 1982

WARNER CHILCOTT 25MG

N87952 001 Nov 18, 1982

WATSON LABS 25MG

N86898 002 Mar 02, 1982

25MG

N87078 001

STANOSZOLOL

TABLET; ORAL

WINSTROL

OVATION PHARMS 2MG

N12885 001 May 14, 1984

STAVUDINE

CAPSULE; ORAL

ZERIT

BRISTOL MYERS SQUIBB 5MG

N20412 001 Jun 24, 1994

CAPSULE, EXTENDED RELEASE; ORAL

ZERIT XR

BRISTOL MYERS SQUIBB 37.5MG

N21453 001 Dec 31, 2002

50MG

N21453 002 Dec 31, 2002

75MG

N21453 003 Dec 31, 2002

100MG

N21453 004 Dec 31, 2002

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION

STREPTOMYCIN SULFATE

COPANOS EQ 500MG BASE/ML

N60684 001

LILLY EQ 1GM BASE/VIAL

N60107 001

EQ 1GM BASE/2ML

N60404 001

EQ 5GM BASE/VIAL

N60107 002

DISCONTINUED DRUG PRODUCT LIST

6 - 233 (of 263)

STREPTOMYCIN SULFATE

INJECTABLE; INJECTION
STREPTOMYCIN SULFATE
PFIZER

EQ 1GM BASE/VIAL
EQ 5GM BASE/VIAL

N60076 001
N60076 002

SUCCINYLCHOLINE CHLORIDE

INJECTABLE; INJECTION
ANECTINE

SABEX 2002

50MG/ML
500MG/VIAL
1GM/VIAL

N08453 003
N08453 001
N08453 004

QUELICIN PRESERVATIVE FREE

HOSPIRA

50MG/ML

N08845 002

SUCCINYLCHOLINE CHLORIDE

INTL MEDICATION

100MG/VIAL

N85400 001 Feb 04, 1982

SUCOSTRIN

APOTHECON

20MG/ML
100MG/ML

N08847 001
N08847 003

SUFENTANIL CITRATE

INJECTABLE; INJECTION
SUFENTANIL CITRATE

STERIS

EQ 0.05MG BASE/ML

N74406 001 Dec 15, 1995

SULCONAZOLE NITRATE

SOLUTION; TOPICAL
EXELDERM

WESTWOOD SQUIBB

1%

N18738 001 Aug 30, 1985

SULFACETAMIDE SODIUM

OINTMENT; OPHTHALMIC
BLEPH-10

ALLERGAN

10%

N84015 001

SODIUM SULAMYD

SCHERING

10%

N05963 002

SULFAIR 10

PHARMAFAIR

10%

N88000 001 Dec 22, 1982

SOLUTION/DROPS; OPHTHALMIC
OCUSULF-30

MIZA PHARMS USA

30%

N80660 002

SODIUM SULAMYD

SCHERING

10%
30%

N05963 001
N05963 003

SODIUM SULFACETAMIDE

AKORN

10%
15%
30%
10%
10%
30%
30%

N83021 001
N83021 002
N83021 003
N84143 001
N84145 001
N84146 001
N84147 001

SULF-15

NOVARTIS

15%

N89047 001 Oct 31, 1995

SULFACETAMIDE SODIUM

PHARMAFAIR

10%

N88947 001 May 17, 1985

SULFAIR 10

PHARMAFAIR

10%

N87949 001 Dec 13, 1982

SULFAIR FORTE

PHARMAFAIR

30%

N88385 001 Oct 13, 1983

SULFAIR-15

PHARMAFAIR

15%

N88186 001 May 25, 1983

SULTEN-10

BAUSCH AND LOMB

10%

N87818 001 Feb 03, 1983

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 234 (of 263)

SULFACYTINE

TABLET; ORAL			
RENOQUID			
GLENWOOD	250MG	N17569	001

SULFADIAZINE

TABLET; ORAL			
SULFADIAZINE			
ABBOTT	300MG	N04125	005
EVERYLIFE	500MG	N80088	001
IMPAX LABS	500MG	N80081	001
LANNETT	500MG	N80084	001
LEDERLE	500MG	N04054	001
LILLY	500MG	N04122	002

SULFADIAZINE SODIUM

INJECTABLE; INJECTION			
SULFADIAZINE SODIUM			
LEDERLE	250MG/ML	N04054	002

SULFADIAZINE; SULFAMERAZINE

SUSPENSION; ORAL			
SULFONAMIDES DUPLEX			
LILLY	250MG/5ML; 250MG/5ML	N06317	007

SULFAMETER

TABLET; ORAL			
SULLA			
BERLEX	500MG	N16000	002

SULFAMETHIZOLE

TABLET; ORAL			
MICROSUL			
FOREST PHARMS	1GM	N86012	001
PROKLAR			
FOREST PHARMS	500MG	N80273	001
THIOSULFIL			
WYETH AYERST	250MG	N08565	001
	500MG	N08565	004

SULFAMETHOXAZOLE

SUSPENSION; ORAL			
GANTANOL			
ROCHE	500MG/5ML	N13664	002
TABLET; ORAL			
GANTANOL			
ROCHE	500MG	N12715	002
GANTANOL -DS			
ROCHE	1GM	N12715	003
SULFAMETHOXAZOLE			
ASCOT	500MG	N87662	001
BARR	500MG	N87189	001
HEATHER	500MG	N86163	001
SANDOZ	500MG	N85844	001
WATSON LABS	500MG	N85053	001
	1GM	N86000	001
UROBAK			
SHIONOGI	500MG	N87307	001

Oct 20, 1982
Jul 25, 1983

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION			
SEPTRA			
MONARCH PHARMS	80MG/ML; 16MG/ML	N18452	001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 235 (of 263)

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AM PHARM PARTNERS	80MG/ML;16MG/ML	N70223	001	Dec 29, 1987
BAXTER HLTHCARE	80MG/ML;16MG/ML	N70627	001	Dec 29, 1987
BEDFORD	80MG/ML;16MG/ML	N72383	001	Apr 29, 1992
STERIS	80MG/ML;16MG/ML	N71556	001	Dec 29, 1987

SUSPENSION; ORAL

BACTRIM

MUTUAL PHARM	200MG/5ML;40MG/5ML	N17560	001	
--------------	--------------------	--------	-----	--

BACTRIM PEDIATRIC

MUTUAL PHARM	200MG/5ML;40MG/5ML	N17560	002	
--------------	--------------------	--------	-----	--

SULFAMETHOXAZOLE AND TRIMETHOPRIM

TEVA	200MG/5ML;40MG/5ML	N70028	001	Jun 02, 1987
------	--------------------	--------	-----	--------------

SULMEPRIM

USL PHARMA	200MG/5ML;40MG/5ML	N70063	001	Aug 01, 1986
------------	--------------------	--------	-----	--------------

SULMEPRIM PEDIATRIC

USL PHARMA	200MG/5ML;40MG/5ML	N70064	001	Aug 01, 1986
------------	--------------------	--------	-----	--------------

TRIMETH/SULFA

ALPHARMA US PHARMS	200MG/5ML;40MG/5ML	N72289	001	May 23, 1988
--------------------	--------------------	--------	-----	--------------

	200MG/5ML;40MG/5ML	N72398	001	May 23, 1988
--	--------------------	--------	-----	--------------

NASKA	200MG/5ML;40MG/5ML	N72399	001	May 23, 1988
-------	--------------------	--------	-----	--------------

TABLET; ORAL

SULFAMETHOPRIM-DS

PAR PHARM	800MG;160MG	N70032	001	Feb 15, 1985
-----------	-------------	--------	-----	--------------

SULFAMETHOXAZOLE AND TRIMETHOPRIM

HEATHER	400MG;80MG	N18946	001	Aug 10, 1984
---------	------------	--------	-----	--------------

	800MG;160MG	N18946	002	Aug 10, 1984
--	-------------	--------	-----	--------------

INTERPHARM	400MG;80MG	N71299	001	Oct 27, 1987
------------	------------	--------	-----	--------------

	800MG;160MG	N71300	001	Oct 27, 1987
--	-------------	--------	-----	--------------

MARTEC	400MG;80MG	N72408	001	Dec 07, 1988
--------	------------	--------	-----	--------------

MUTUAL PHARM	400MG;80MG	N70006	001	Nov 14, 1984
--------------	------------	--------	-----	--------------

ROXANE	400MG;80MG	N72768	001	Aug 30, 1991
--------	------------	--------	-----	--------------

SANDOZ	400MG;80MG	N18598	003	May 19, 1982
--------	------------	--------	-----	--------------

	400MG;80MG	N70889	001	Nov 13, 1986
--	------------	--------	-----	--------------

TEVA	400MG;80MG	N18242	001	
------	------------	--------	-----	--

	800MG;160MG	N18242	002	
--	-------------	--------	-----	--

USL PHARMA	400MG;80MG	N70203	001	Nov 08, 1985
------------	------------	--------	-----	--------------

	800MG;160MG	N70204	001	Nov 08, 1985
--	-------------	--------	-----	--------------

WATSON LABS	400MG;80MG	N70002	001	Nov 07, 1984
-------------	------------	--------	-----	--------------

	800MG;160MG	N70000	001	Nov 07, 1984
--	-------------	--------	-----	--------------

SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH

MARTEC	800MG;160MG	N72417	001	Dec 07, 1988
--------	-------------	--------	-----	--------------

MUTUAL PHARM	800MG;160MG	N70007	001	Nov 14, 1984
--------------	-------------	--------	-----	--------------

ROXANE	800MG;160MG	N72769	001	Aug 30, 1991
--------	-------------	--------	-----	--------------

SULFATRIM-DS

SUPERPHARM	800MG;160MG	N70066	001	Jun 24, 1985
------------	-------------	--------	-----	--------------

SULFATRIM-SS

SUPERPHARM	400MG;80MG	N70065	002	Jun 24, 1985
------------	------------	--------	-----	--------------

UROPLUS DS

SHIONOGI	800MG;160MG	N71816	001	Sep 28, 1987
----------	-------------	--------	-----	--------------

UROPLUS SS

SHIONOGI	400MG;80MG	N71815	001	Sep 28, 1987
----------	------------	--------	-----	--------------

SULFANILAMIDE

CREAM; VAGINAL

AVC

PHARMELLE	15%	N06530	003	Jan 27, 1987
-----------	-----	--------	-----	--------------

SULFANILAMIDE

TEVA	15%	N88718	001	Sep 19, 1985
------	-----	--------	-----	--------------

SUPPOSITORY; VAGINAL

AVC

PHARMELLE	1.05GM	N06530	004	Jan 27, 1987
-----------	--------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 236 (of 263)

SULFAPHENAZOLE

SUSPENSION; ORAL

SULFABID

PURDUE FREDERICK

500MG/5ML

N13093 001

TABLET; ORAL

SULFABID

PURDUE FREDERICK

500MG

N13092 002

SULFAPYRIDINE

TABLET; ORAL

SULFAPYRIDINE

LILLY

500MG

N00159 001

SULFASALAZINE

SUSPENSION; ORAL

AZULFIDINE

PHARMACIA AND UPJOHN

250MG/5ML

N18605 001

TABLET; ORAL

S.A.S. -500

SOLVAY

500MG

N83450 001

SULFASALAZINE

CLONMEL HLTHCARE

500MG

N80197 001

SANDOZ

500MG

N86184 001

SUPERPHARM

500MG

N89339 001

Oct 26, 1987

WATSON LABS

500MG

N84964 001

TABLET, DELAYED RELEASE; ORAL

SULFASALAZINE

WATSON LABS

500MG

N88052 001

May 24, 1983

SULFINPYRAZONE

CAPSULE; ORAL

SULFINPYRAZONE

VANGARD

200MG

N88666 001

Feb 17, 1984

TABLET; ORAL

SULFINPYRAZONE

IVAX PHARMS

100MG

N87769 001

Jun 01, 1982

WATSON LABS

100MG

N87667 001

May 26, 1982

SULFISOXAZOLE

TABLET; ORAL

GANTRISIN

ROCHE

500MG

N06525 001

SOSOL

MK LABS

500MG

N80036 001

SOXAZOLE

ALRA

500MG

N80366 001

SULFALAR

PARKE DAVIS

500MG

N84955 001

SULFISOXAZOLE

BARR

500MG

N84031 001

HEATHER

500MG

N80189 001

IMPAX LABS

500MG

N80109 001

LANNETT

500MG

N80085 001

LEDERLE

500MG

N87649 001

PHARMERAL

500MG

N84385 001

PUREPAC PHARM

500MG

N80087 001

ROXANE

500MG

N80082 001

SANDOZ

500MG

N85628 001

VALEANT PHARM INTL

500MG

N80268 002

VITARINE

500MG

N87332 001

WATSON LABS

500MG

N85534 001

WEST WARD

500MG

N80379 001

SULSOXIN

SOLVAY

500MG

N80040 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 237 (of 263)

SULFISOXAZOLE ACETYL

EMULSION; ORAL LIPO GANTRISIN ROCHE	EQ 1GM BASE/5ML	N09182	009
SYRUP; ORAL GANTRISIN ROCHE	EQ 500MG BASE/5ML	N09182	002

SULFISOXAZOLE DIOLAMINE

INJECTABLE; INJECTION GANTRISIN ROCHE	EQ 400MG BASE/ML	N06917	001
OINTMENT; OPHTHALMIC GANTRISIN ROCHE	EQ 4% BASE	N08414	002
SOLUTION/DROPS; OPHTHALMIC GANTRISIN ROCHE	EQ 4% BASE	N07757	002
SULFISOXAZOLE DIOLAMINE SOLA BARNES HIND	EQ 4% BASE	N84148	001

SULFOXONE SODIUM

TABLET, DELAYED RELEASE; ORAL DIASONE SODIUM ABBOTT	165MG	N06044	003
---	-------	--------	-----

SULFUR

POWDER; TOPICAL BENSULFOID POYTHRESS	33.32%	N02918	001
--	--------	--------	-----

SULINDAC

TABLET; ORAL SULINDAC CLONMEL HLTHCARE	150MG	N73261	001	Sep 06, 1991
	200MG	N73262	001	Sep 06, 1991
WARNER CHILCOTT	150MG	N72710	001	Mar 25, 1991
	200MG	N72711	001	Mar 25, 1991

SUPROFEN

SOLUTION/DROPS; OPHTHALMIC PROFENAL ALCON	1%	N19387	001	Dec 23, 1988
---	----	--------	-----	--------------

SUTILAINS

OINTMENT; TOPICAL TRAVASE ABBOTT	82,000 UNITS/GM **Federal Register determination that product was not discontinued or withdrawn for safety or efficacy reasons**	N12828	001
--	---	--------	-----

TALBUTAL

TABLET; ORAL LOTUSATE SANOFI SYNTHELABO	120MG	N09410	005
---	-------	--------	-----

TAMOXIFEN CITRATE

TABLET; ORAL TAMOXIFEN CITRATE PHARMACHEMIE	EQ 10MG BASE	N74539	001	Mar 31, 2003
---	--------------	--------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 238 (of 263)

TECHNETIUM TC-99M ALBUMIN AGGREGATED

INJECTABLE; INJECTION		
TC 99M-LUNGAGGREGATE		
GE HEALTHCARE	5mCi/ML	N17848 001

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE; INJECTION		
A-N STANNOUS AGGREGATED ALBUMIN		
SYNCOR PHARMS	N/A	N17916 001
AN-MAA		
CIS	N/A	N17792 001
LUNGAGGREGATE REAGENT		
GE HEALTHCARE	N/A	N17838 001
TECHNETIUM TC 99M MAA		
GE HEALTHCARE	N/A	N17773 001

TECHNETIUM TC-99M ALBUMIN COLLOID KIT

INJECTABLE; INJECTION			
MICROLITE			
CIS	N/A	N18263 001	Mar 25, 1983

TECHNETIUM TC-99M ALBUMIN KIT

INJECTABLE; INJECTION		
TECHNETIUM TC 99M HSA		
GE HEALTHCARE	N/A	N17775 001

TECHNETIUM TC-99M ALBUMIN MICROSPHERES KIT

INJECTABLE; INJECTION		
INSTANT MICROSPHERES		
3M	N/A	N17832 001

TECHNETIUM TC-99M ETIDRONATE KIT

INJECTABLE; INJECTION		
CINTICHEM TECHNETIUM 99M HEDSPA		
GE HEALTHCARE	N/A	N17653 001
MPI STANNOUS DIPHOSPHONATE		
GE HEALTHCARE	N/A	N17667 001
OSTEOSCAN		
MALLINCKRODT	N/A	N17454 001
TECHNETIUM TC 99M DIPHOSPHONATE-TIN KIT		
GE HEALTHCARE	N/A	N17562 001

TECHNETIUM TC-99M FERMENTETATE KIT

INJECTABLE; INJECTION		
RENOTEC		
BRACCO	N/A	N17045 001

TECHNETIUM TC-99M GLUCEPTATE KIT

INJECTABLE; INJECTION		
GLUCOSCAN		
BRISTOL MYERS SQUIBB	N/A	N17907 001

TECHNETIUM TC-99M LIDOFENIN KIT

INJECTABLE; INJECTION			
TECHNISCAN HIDA			
DRAXIMAGE	N/A	N18489 001	Oct 31, 1986

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION			
AMERSCAN MDP KIT			
GE HEALTHCARE	N/A	N18335 001	Aug 05, 1982
OSTEOLITE			
CIS	N/A	N17972 001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 239 (of 263)

TECHNETIUM TC-99M POLYPHOSPHATE KIT

INJECTABLE; INJECTION			
SODIUM POLYPHOSPHATE-TIN KIT			
GE HEALTHCARE	N/A	N17664	001

TECHNETIUM TC-99M PYRO/TRIMETA PHOSPHATES KIT

INJECTABLE; INJECTION			
PYROLITE			
CIS	N/A	N17684	001

TECHNETIUM TC-99M RED BLOOD CELL KIT

INJECTABLE; INJECTION			
RBC-SCAN			
CADEMA	N/A	N20063	001 Jun 11, 1992

TECHNETIUM TC-99M SODIUM PERTECHNETATE

SOLUTION; INJECTION, ORAL			
SODIUM PERTECHNETATE TC 99M			
CIS	12mCi/ML	N17321	001
	24mCi/ML	N17321	002
	48mCi/ML	N17321	003
GE HEALTHCARE	2-100mCi/ML	N17471	001
MALLINCKRODT	10-60mCi/ML	N17725	001

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR

SOLUTION; INJECTION, ORAL			
MINITEC			
BRACCO	0.22-2.22 CI/GENERATOR	N17339	001
TECHNETIUM TC 99M GENERATOR			
GE HEALTHCARE	830-16600mCi/GENERATOR	N17693	001

TECHNETIUM TC-99M SULFUR COLLOID

SOLUTION; INJECTION, ORAL			
TECHNETIUM TC 99M SULFUR COLLOID			
GE HEALTHCARE	4mCi/ML	N17456	001
SOLUTION; ORAL			
TECHNETIUM TC 99M SULFUR COLLOID			
MALLINCKRODT	3mCi/ML	N17724	001

TECHNETIUM TC-99M SULFUR COLLOID KIT

SOLUTION; INJECTION, ORAL			
TECHNETIUM TC 99M TSC			
GE HEALTHCARE	N/A	N17784	001
TESULOID			
BRACCO	N/A	N16923	001

TECHNETIUM TC-99M TEBOROXIME KIT

INJECTABLE; INJECTION			
CARDIOTEC			
BRACCO	N/A	N19928	001 Dec 19, 1990

TEMAZEPAM

CAPSULE; ORAL			
TEMAZ			
QUANTUM PHARMICS	15MG	N70564	001 Oct 15, 1985
	30MG	N70547	001 Oct 15, 1985
TEMAZEPAM			
DURAMED PHARMS BARR	15MG	N71708	001 Sep 29, 1988
	30MG	N71709	001 Sep 29, 1988
MUTUAL PHARM	15MG	N71174	001 Jul 10, 1986
	30MG	N71175	001 Jul 10, 1986
USL PHARMA	15MG	N70489	001 Jul 07, 1986
	30MG	N70490	001 Jul 07, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 240 (of 263)

TEMAZEPAM

CAPSULE; ORAL
TEMAZEPAM

WATSON LABS	15MG	N70383	001	Mar 23, 1987
	30MG	N70384	001	Mar 23, 1987

TERAZOSIN HYDROCHLORIDE

CAPSULE; ORAL
TERAZOSIN HYDROCHLORIDE

IVAX PHARMS	EQ 1MG BASE	N75614	002	Jan 30, 2001
	EQ 2MG BASE	N75614	001	Jan 30, 2001
	EQ 5MG BASE	N75614	003	Jan 30, 2001
	EQ 10MG BASE	N75614	004	Jan 30, 2001

TABLET; ORAL
TERAZOSIN HYDROCHLORIDE

IVAX PHARMS	EQ 1MG BASE	N74530	001	Apr 21, 2000
	EQ 2MG BASE	N74530	002	Apr 21, 2000
	EQ 5MG BASE	N74530	003	Apr 21, 2000
	EQ 10MG BASE	N74530	004	Apr 21, 2000
TEVA	EQ 1MG BASE	N74446	001	May 18, 2000
	EQ 2MG BASE	N74446	002	May 18, 2000
	EQ 5MG BASE	N74446	003	May 18, 2000
	EQ 10MG BASE	N74446	004	May 18, 2000

TERBINAFINE HYDROCHLORIDE

CREAM; TOPICAL
LAMISIL

NOVARTIS	1%	N20192	001	Dec 30, 1992
----------	----	--------	-----	--------------

TERBUTALINE SULFATE

AEROSOL, METERED; INHALATION

BRETHAIRE

NOVARTIS	0.2MG/INH	N18762	001	Aug 17, 1984
----------	-----------	--------	-----	--------------

BRICANYL

AVENTIS PHARMS	0.2MG/INH	N18000	001	Mar 19, 1985
----------------	-----------	--------	-----	--------------

INJECTABLE; INJECTION

BRICANYL

AVENTIS PHARMS	1MG/ML	N17466	001	
----------------	--------	--------	-----	--

TABLET; ORAL

BRICANYL

AVENTIS PHARMS	2.5MG	N17618	001	
	5MG	N17618	002	

TERIPARATIDE ACETATE

INJECTABLE; INJECTION

PARATHAR

AVENTIS	200 UNITS/VIAL	N19498	001	Dec 23, 1987
---------	----------------	--------	-----	--------------

TESTOLACTONE

INJECTABLE; INJECTION

TESLAC

BRISTOL MYERS SQUIBB	100MG/ML	N16119	001	
----------------------	----------	--------	-----	--

TABLET; ORAL

TESLAC

BRISTOL MYERS SQUIBB	250MG	N16118	002	
----------------------	-------	--------	-----	--

TESTOSTERONE

FILM, EXTENDED RELEASE; TRANSDERMAL

TESTODERM

ALZA	4MG/24HR	N19762	001	Oct 12, 1993
	6MG/24HR	N19762	002	Oct 12, 1993

TESTODERM TTS

ALZA	5MG/24HR	N20791	001	Dec 18, 1997
------	----------	--------	-----	--------------

DISCONTINUED DRUG PRODUCT LIST

6 - 241 (of 263)

TESTOSTERONE

INJECTABLE; INJECTION
TESTOSTERONE
STERIS

25MG/ML
50MG/ML

N86420 001 May 10, 1983
N86419 001 Aug 23, 1983

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION
DEPO-TESTOSTERONE
PHARMACIA AND UPJOHN
TESTOSTERONE CYPIONATE
STERIS

50MG/ML
100MG/ML
200MG/ML

N85635 001
N84401 001
N84401 002

TESTOSTERONE ENANTHATE

INJECTABLE; INJECTION
DELATESTRYL
SAVIENT PHARMS
TESTOSTERONE ENANTHATE
STERIS

200MG/ML
100MG/ML
100MG/ML
200MG/ML

N09165 001
N83667 001
N85599 001
N83667 002

TESTOSTERONE PROPIONATE

INJECTABLE; INJECTION
TESTOSTERONE PROPIONATE
BEL MAR

25MG/ML
50MG/ML
100MG/ML
25MG/ML
50MG/ML
25MG/ML
50MG/ML
100MG/ML

N80741 001
N80742 001
N80743 001
N80276 001
N80254 002
N85490 001
N85490 002
N83595 003

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL
ACHROMYCIN V
CLONMEL HLTHCARE

250MG
500MG

N50278 003
N50278 001

CYCLOPAR
WARNER CHILCOTT

250MG
250MG
250MG
500MG
500MG

N61725 001
N62175 001
N62332 001
N61725 002
N62332 002

PANMYCIN
PHARMACIA AND UPJOHN

250MG

N60347 001

RETET
SOLVAY

250MG
500MG

N61443 001
N61443 002

ROBITET
WYETH AYERST

250MG
500MG

N61734 001
N61734 002

SUMYCIN
APOTHECON

100MG
125MG
250MG
500MG

N60429 002
N60429 004
N60429 001
N60429 003

TETRACHEL
ANGUS

250MG
500MG

N60343 001
N60343 003

TETRACYCLINE HYDROCHLORIDE
ABBOTT

250MG

N61802 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 242 (of 263)

TETRACYCLINE HYDROCHLORIDE

CAPSULE; ORAL

TETRACYCLINE HYDROCHLORIDE

ABBOTT	500MG	N61802	002	
ELKINS SINN	250MG	N60059	001	
FERRANTE	125MG	N60173	001	
	250MG	N60173	002	
HEATHER	250MG	N61148	001	
	500MG	N61148	002	
MAST MM	250MG	N62085	001	
PUREPAC PHARM	250MG	N60290	001	
	500MG	N60290	002	
PVT FORM	250MG	N62686	001	Jul 24, 1986
	500MG	N62686	002	Jul 24, 1986
ROXANE	500MG	N61214	002	
SANDOZ	250MG	N61471	001	
SUPERPHARM	250MG	N62540	001	Mar 21, 1985
	500MG	N62540	002	Mar 21, 1985
VALEANT PHARM INTL	250MG	N60471	001	
	500MG	N60471	002	
WARNER CHILCOTT	250MG	N62300	001	
	500MG	N62300	002	
WATSON LABS	250MG	N62103	001	
	250MG	N62343	001	
	500MG	N62103	002	
	500MG	N62343	002	
WEST WARD	250MG	N60768	001	
	500MG	N60768	002	
WYETH AYERST	250MG	N61685	001	
	500MG	N61685	002	

TETRACYN

PFIPHARMECS	250MG	N60082	003
	500MG	N60082	004

FIBER, EXTENDED RELEASE; PERIODONTAL

ACTISITE

ALZA	12.7MG/FIBER	N50653	001	Mar 25, 1994
------	--------------	--------	-----	--------------

FOR SOLUTION; TOPICAL

TOPICYCLINE

SHIRE	2.2MG/ML	N50493	001
-------	----------	--------	-----

INJECTABLE; INJECTION

ACHROMYCIN

LEDERLE	250MG/VIAL	N50273	002
	500MG/VIAL	N50273	003

TETRACYN

PFIZER	250MG/VIAL	N60096	001
	500MG/VIAL	N60096	002

OINTMENT; OPHTHALMIC

ACHROMYCIN

STORZ	10MG/GM	N50266	001
-------	---------	--------	-----

SUSPENSION; ORAL

ACHROMYCIN V

LEDERLE	125MG/5ML	N50263	002
---------	-----------	--------	-----

TETRACYCLINE HYDROCHLORIDE

ALPHARMA US PHARMS	125MG/5ML	N60633	001
FERRANTE	125MG/5ML	N60174	001
PROTER	125MG/5ML	N60446	001
PUREPAC PHARM	125MG/5ML	N60291	001

TETRACYN

PFIPHARMECS	125MG/5ML	N60095	001
-------------	-----------	--------	-----

TETRAMED

IVAX PHARMS	125MG/5ML	N61468	001
-------------	-----------	--------	-----

SUSPENSION/DROPS; OPHTHALMIC

ACHROMYCIN

STORZ	1%	N50268	001
-------	----	--------	-----

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 243 (of 263)

TETRACYCLINE HYDROCHLORIDE

TABLET; ORAL

PANMYCIN

PHARMACIA AND UPJOHN	250MG	N61705	001
	500MG	N61705	002

SUMYCIN

PAR PHARM	50MG	N61147	003
	100MG	N61147	002

TETRACYCLINE PHOSPHATE COMPLEX

CAPSULE; ORAL

TETREX

BRISTOL	EQ 250MG HCL	N50212	002
	EQ 500MG HCL	N50212	003

THALLOUS CHLORIDE, TL-201

INJECTABLE; INJECTION

THALLOUS CHLORIDE TL 201

BRACCO	1mCi/ML	N18548	001	Dec 30, 1982
--------	---------	--------	-----	--------------

THEOPHYLLINE

CAPSULE; ORAL

BRONKODYL

SANOFI SYNTHELABO	100MG	N85264	001
	200MG	N85264	002

ELIXOPHYLLIN

FOREST LABS	100MG	N85545	001	Jul 31, 1984
	200MG	N83921	001	Jul 31, 1984

SOMOPHYLLIN-T

FISONS	100MG	N87155	001	Feb 25, 1985
	200MG	N87155	002	Feb 25, 1985
	250MG	N87155	003	Feb 25, 1985

THEOPHYLLINE

KV PHARM	100MG	N85263	001
	200MG	N85263	002

SCHERER RP	100MG	N84731	002	Nov 07, 1986
	200MG	N84731	001	Nov 07, 1986
	250MG	N84731	003	Nov 07, 1986

CAPSULE, EXTENDED RELEASE; ORAL

AEROLATE III

FLEMING PHARMS	65MG	N85075	003	Nov 24, 1986
----------------	------	--------	-----	--------------

AEROLATE JR

FLEMING PHARMS	130MG	N85075	002	Nov 24, 1986
----------------	-------	--------	-----	--------------

AEROLATE SR

FLEMING PHARMS	260MG	N85075	001	Nov 24, 1986
----------------	-------	--------	-----	--------------

ELIXOPHYLLIN SR

FOREST LABS	125MG	N86826	001	Jan 29, 1985
	250MG	N86826	002	Jan 29, 1985

SLO-BID

AVENTIS	50MG	N88269	001	Jan 31, 1985
	75MG	N89539	001	May 10, 1989
	100MG	N87892	001	Jan 31, 1985
	125MG	N89540	001	May 10, 1989
	200MG	N87893	001	Jan 31, 1985
	300MG	N87894	001	Jan 31, 1985

SLO-PHYLLIN

AVENTIS	60MG	N85206	001	May 24, 1982
	125MG	N85203	001	May 24, 1982
	250MG	N85205	001	May 24, 1982

SOMOPHYLLIN-CRT

GRAHAM DM	50MG	N87763	001	Feb 27, 1985
	100MG	N87194	001	
	200MG	N88382	001	Feb 27, 1985
	250MG	N87193	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 244 (of 263)

THEOPHYLLINECAPSULE, EXTENDED RELEASE; ORAL
SOMOPHYLLIN-CRT

GRAHAM DM	300MG	N88383	001	Feb 27, 1985
THEOBID				
WHITBY	260MG	N85983	001	Mar 20, 1985
THEOBID JR.				
WHITBY	130MG	N87854	001	Mar 20, 1985
THEOCLEAR L.A.-130				
SCHWARZ PHARMA	130MG	N86569	001	May 27, 1982
THEOCLEAR L.A.-260				
SCHWARZ PHARMA	260MG	N86569	002	May 27, 1982
THEO-DUR				
SCHERING	50MG	N88022	001	Sep 10, 1985
	75MG	N88015	001	Sep 10, 1985
	125MG	N88016	001	Sep 10, 1985
	200MG	N87995	001	Sep 10, 1985

THEOPHYLLINE

CENT PHARMS

	125MG	N88654	001	Feb 12, 1985
	250MG	N88689	001	Feb 12, 1985
MAYNE PHARMA USA	100MG	N89976	001	Jan 04, 1995
	200MG	N89977	001	Jan 04, 1995
	300MG	N89932	001	Jan 04, 1995
SANDOZ	260MG	N87462	001	May 11, 1982

THEOPHYLLINE-SR

SCHERER RP

THEOPHYL-SR

ORTHO MCNEIL PHARM

	125MG	N86480	001	Feb 08, 1985
	250MG	N86471	001	Feb 08, 1985
THEOVENT				
SCHERING	125MG	N87010	001	Jan 31, 1985
	250MG	N87910	001	Jan 31, 1985

ELIXIR; ORAL

ELIXOMIN

CENCI

LANOPHYLLIN

LANNETT

THEOLIXIR

PANRAY

THEOPHYL-225

ORTHO MCNEIL PHARM

THEOPHYLLINE

ALPHARMA US PHARMS

CENCI

HALSEY

PERRIGO

ROXANE

INJECTABLE; INJECTION

THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN

THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN

THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN

THEOPHYLLINE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN

THEOPHYLLINE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN

SOLUTION; ORAL

AEROLATE

FLEMING PHARMS

	40MG/100ML	N19083	001	Nov 07, 1984
	80MG/100ML	N19083	002	Nov 07, 1984
	160MG/100ML	N19083	003	Nov 07, 1984
	200MG/100ML	N19212	001	Nov 07, 1984
	200MG/100ML	N19826	004	Aug 14, 1992
	4MG/ML	N19212	003	Nov 07, 1984
	400MG/100ML	N19212	002	Nov 07, 1984
	400MG/100ML	N19826	005	Aug 14, 1992
	150MG/15ML	N89141	001	Dec 03, 1986

DISCONTINUED DRUG PRODUCT LIST

6 - 245 (of 263)

THEOPHYLLINE

SUSPENSION; ORAL				
ELIXICON				
FOREST LABS	100MG/5ML	N85502	001	
SYRUP; ORAL				
ACCURBRON				
AVENTIS PHARMS	150MG/15ML	N88746	001	Nov 22, 1985
AQUAPHYLLIN				
FERNDAL LABS	80MG/15ML	N87917	001	Jan 18, 1983
SLO-PHYLLIN				
AVENTIS	80MG/15ML	N85187	001	
THEOCLEAR-80				
CENT PHARMS	80MG/15ML	N87095	001	Mar 01, 1982
THEOPHYLLINE				
ALPHARMA US PHARMS	80MG/15ML	N86001	001	
	150MG/15ML	N86545	001	
TABLET; ORAL				
SLO-PHYLLIN				
AVENTIS	100MG	N85202	001	
	200MG	N85204	001	
THEOCLEAR-100				
CENT PHARMS	100MG	N85353	002	
THEOCLEAR-200				
CENT PHARMS	200MG	N85353	001	
THEOPHYL-225				
ORTHO MCNEIL PHARM	225MG	N84726	001	
TABLET, CHEWABLE; ORAL				
THEOPHYL				
ORTHO MCNEIL PHARM	100MG	N86506	001	Sep 12, 1985
TABLET, EXTENDED RELEASE; ORAL				
DURAPHYL				
FOREST LABS	100MG	N88503	001	Apr 03, 1985
	200MG	N88504	001	Apr 03, 1985
	300MG	N88505	001	Apr 03, 1985
LABID				
PROCTER AND GAMBLE	250MG	N87225	001	
SUSTAIRE				
ROERIG	100MG	N85665	001	
	300MG	N85665	002	
THEO-DUR				
SCHERING	100MG	N85328	001	
	200MG	N86998	001	
	300MG	N85328	002	
	450MG	N89131	001	Jun 25, 1986
THEOLAIR-SR				
3M	200MG	N88369	001	Jul 16, 1987
	250MG	N86363	002	Jul 16, 1987
	300MG	N88364	001	Jul 16, 1987
	500MG	N89132	001	Jul 16, 1987
THEOPHYLLINE				
ABLE	300MG	N40548	001	Apr 30, 2004
	400MG	N40543	001	Apr 27, 2004
	450MG	N40546	001	Apr 30, 2004
	600MG	N40539	001	Apr 27, 2004
UNI-DUR				
SCHERING	400MG	N89822	001	Jan 04, 1995
	600MG	N89823	001	Jan 04, 1995

THEOPHYLLINE SODIUM GLYCINATE

ELIXIR; ORAL				
SYNOHYLLATE				
CENT PHARMS	EQ 165MG BASE/15ML	N06333	008	
TABLET; ORAL				
ASBRON				
NOVARTIS	EQ 150MG BASE	N85148	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 246 (of 263)

THIAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

BETALIN S

LILLY

100MG/ML

N80853 001

THIAMINE HYDROCHLORIDE

AKORN

100MG/ML

N87968 001

Oct 01, 1982

AM PHARM PARTNERS

100MG/ML

N80509 001

BEL MAR

100MG/ML

N80718 001

200MG/ML

N80712 001

DELL LABS

100MG/ML

N83775 001

LUITPOLD

100MG/ML

N80667 001

PARKE DAVIS

100MG/ML

N80770 001

STERIS

100MG/ML

N83534 001

200MG/ML

N83534 002

WYETH AYERST

100MG/ML

N80553 001

THIAMYLAL SODIUM

INJECTABLE; INJECTION

SURITAL

PARKEDALE

1GM/VIAL

N07600 003

5GM/VIAL

N07600 005

10GM/VIAL

N07600 009

THIETHYLPERAZINE MALATE

INJECTABLE; INJECTION

TORECAN

NOVARTIS

5MG/ML

N12754 002

THIETHYLPERAZINE MALEATE

SUPPOSITORY; RECTAL

TORECAN

NOVARTIS

10MG

N13247 001

THIOPENTAL SODIUM

SUSPENSION; RECTAL

PENTOTHAL

ABBOTT

400MG/GM

N11679 001

THIORIDAZINE

SUSPENSION; ORAL

MELLARIL-S

NOVARTIS

EQ 25MG HCL/5ML

N17923 001

EQ 100MG HCL/5ML

N17923 002

THIORIDAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

MELLARIL

NOVARTIS

30MG/ML

N11808 012

100MG/ML

N11808 018

THIORIDAZINE HYDROCHLORIDE

ALPHARMA US PHARMS

30MG/ML

N87766 001

Apr 26, 1983

HI TECH PHARMA

30MG/ML

N40125 001

Aug 16, 1996

100MG/ML

N40126 001

Aug 16, 1996

MORTON GROVE

30MG/ML

N88258 001

Jul 25, 1983

100MG/ML

N88227 001

Jul 05, 1983

SANDOZ

30MG/ML

N88307 001

Nov 23, 1983

100MG/ML

N88308 001

Nov 23, 1983

TABLET; ORAL

MELLARIL

NOVARTIS

10MG

N11808 003

15MG

N11808 016

25MG

N11808 006

50MG

N11808 011

100MG

N11808 009

DISCONTINUED DRUG PRODUCT LIST

6 - 247 (of 263)

THIORIDAZINE HYDROCHLORIDE

TABLET; ORAL

MELLARIL

NOVARTIS

150MG

N11808 017

200MG

N11808 015

THIORIDAZINE HYDROCHLORIDE

IVAX PHARMS

50MG

N88194 001 Apr 14, 1983

100MG

N88273 001 Oct 03, 1983

MUTUAL PHARM

10MG

N88375 001 Nov 18, 1983

15MG

N88461 001 Nov 18, 1983

25MG

N87264 001 Nov 18, 1983

100MG

N88379 001 Nov 16, 1983

150MG

N88737 001 Sep 26, 1984

200MG

N88738 001 Oct 16, 1984

MYLAN

10MG

N88332 001 Jun 27, 1983

25MG

N88333 001 Jun 27, 1983

50MG

N88334 001 Jun 27, 1983

100MG

N88335 001 Nov 18, 1983

PAR PHARM

10MG

N88351 001 Dec 05, 1983

15MG

N88352 001 Dec 05, 1983

25MG

N88336 001 Dec 05, 1983

50MG

N88322 001 Dec 05, 1983

100MG

N88480 001 Dec 29, 1983

150MG

N89764 001 Feb 09, 1988

200MG

N89765 001 Feb 09, 1988

ROXANE

10MG

N88663 001 Mar 15, 1984

25MG

N88664 001 Mar 15, 1984

50MG

N88665 001 Mar 15, 1984

100MG

N89048 001 Feb 26, 1985

SUPERPHARM

10MG

N89103 001 Jul 02, 1985

25MG

N89104 001 Jul 02, 1985

50MG

N89105 001 Jul 02, 1985

TEVA

10MG

N88493 001 May 17, 1985

WATSON LABS

10MG

N88412 001 Sep 12, 1983

10MG

N88561 001 May 11, 1984

15MG

N88345 001 Jul 28, 1983

15MG

N88562 001 May 11, 1984

25MG

N88296 001 Jul 28, 1983

25MG

N88755 001 Jul 24, 1984

50MG

N88323 001 Jul 28, 1983

100MG

N88284 001 Aug 25, 1983

150MG

N88410 001 Mar 05, 1984

200MG

N88381 001 Mar 14, 1984

WEST WARD

10MG

N88658 001 Mar 26, 1984

15MG

N88659 001 Mar 26, 1984

25MG

N88660 001 Mar 26, 1984

50MG

N88661 001 Mar 26, 1984

THIOTEPA

INJECTABLE; INJECTION

THIOPLEX

IMMUNEX

15MG/VIAL

N20058 001 Dec 22, 1994

THIOTEPA

IMMUNEX

15MG/VIAL

N11683 001

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

AM THERAP

1MG

N71884 001 Aug 12, 1987

2MG

N71885 001 Aug 12, 1987

5MG

N71886 001 Aug 12, 1987

10MG

N71887 001 Aug 12, 1987

20MG

N72200 001 Dec 17, 1987

WATSON LABS

2MG

N71626 001 Jun 25, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 248 (of 263)

THIOTHIXENE

CAPSULE; ORAL

THIOTHIXENE

WATSON LABS

5MG

N71627 001

Jun 25, 1987

10MG

N71628 001

Jun 25, 1987

THIOTHIXENE HYDROCHLORIDE

CONCENTRATE; ORAL

THIOTHIXENE HYDROCHLORIDE

ALPHARMA US PHARMS

EQ 5MG BASE/ML

N70969 001

Oct 16, 1987

PACO

EQ 1MG BASE/ML

N71917 001

Sep 20, 1989

EQ 5MG BASE/ML

N71939 001

Dec 16, 1988

TEVA

EQ 5MG BASE/ML

N71184 001

Jun 22, 1987

THIOTHIXENE HYDROCHLORIDE INTENSOL

ROXANE

EQ 5MG BASE/ML

N73494 001

Jun 30, 1992

INJECTABLE; INJECTION

NAVANE

PFIZER

EQ 2MG BASE/ML

N16904 001

THYROGLOBULIN

TABLET; ORAL

PROLOID

PARKE DAVIS

16MG

N02245 009

32MG

N02245 005

65MG

N02245 002

100MG

N02245 008

130MG

N02245 010

200MG

N02245 007

325MG

N02245 004

THYROGLOBULIN

IMPAX LABS

64.8MG

N80151 001

THYROTROPIN

INJECTABLE; INJECTION

THYTROPAR

AVENTIS

10 IU/VIAL

N08682 001

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

CEPHALON

20MG

N20646 004

Sep 30, 1997

TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TICAR

GLAXOSMITHKLINE

EQ 3GM BASE/VIAL

N62690 001

Dec 19, 1986

EQ 6GM BASE/VIAL

N50497 003

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLID

ROCHE PALO

125MG

N19979 001

Mar 24, 1993

TICLOPIDINE HYDROCHLORIDE

WATSON LABS

250MG

N75309 001

Apr 26, 2000

TIMOLOL MALEATE

SOLUTION/DROPS; OPHTHALMIC

TIMOLOL MALEATE

AKORN

EQ 0.25% BASE

N74465 001

Mar 25, 1997

FOUGERA

EQ 0.25% BASE

N74667 001

Mar 25, 1997

EQ 0.5% BASE

N74668 001

Mar 25, 1997

TABLET; ORAL

TIMOLOL MALEATE

QUANTUM PHARMICS

5MG

N72466 001

May 19, 1989

DISCONTINUED DRUG PRODUCT LIST

6 - 249 (of 263)

TIMOLOL MALEATE

TABLET; ORAL

TIMOLOL MALEATE

QUANTUM PHARMICS	10MG	N72467	001	May 19, 1989
	20MG	N72468	001	May 19, 1989
TEVA	5MG	N72648	001	Jun 16, 1993
	10MG	N72649	001	Jun 16, 1993
	20MG	N72650	001	Jun 16, 1993
USL PHARMA	5MG	N72001	001	Apr 11, 1989
	10MG	N72002	001	Apr 11, 1989
	20MG	N72003	001	Apr 11, 1989
WATSON LABS	5MG	N72269	001	Apr 11, 1989
	10MG	N72270	001	Apr 11, 1989
	20MG	N72271	001	Apr 11, 1989

TIOCONAZOLE

CREAM; TOPICAL

TZ-3

PFIZER	1%	N18682	001	Feb 18, 1983
--------	----	--------	-----	--------------

TOBRAMYCIN

SOLUTION/DROPS; OPHTHALMIC

TOBRAMYCIN

ALCON UNIVERSAL	0.3%	N63176	001	May 25, 1994
-----------------	------	--------	-----	--------------

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

NEBCIN

LILLY	EQ 10MG BASE/ML	N50477	005	
	EQ 10MG BASE/ML	N62008	004	
	EQ 10MG BASE/ML	N62707	001	Apr 29, 1987
	EQ 40MG BASE/ML	N62008	001	
	EQ 1.2GM BASE/VIAL	N50519	001	

TOBRAMYCIN SULFATE

APOTHECON	EQ 40MG BASE/ML	N64026	001	May 31, 1994
ASTRAZENECA	EQ 10MG BASE/ML	N63119	001	Oct 31, 1994
	EQ 40MG BASE/ML	N63120	001	Oct 31, 1994
	EQ 40MG BASE/ML	N63121	001	Oct 31, 1994
	EQ 40MG BASE/ML	N63122	001	Oct 31, 1994
BAXTER HLTHCARE	EQ 10MG BASE/ML	N63113	001	Apr 26, 1991
	EQ 10MG BASE/ML	N63128	001	Nov 27, 1991
	EQ 40MG BASE/ML	N63127	001	Nov 27, 1991

TOCAINIDE HYDROCHLORIDE

TABLET; ORAL

TONOCARD

ASTRAZENECA	400MG	N18257	001	Nov 09, 1984
	600MG	N18257	002	Nov 09, 1984

TOLAZAMIDE

TABLET; ORAL

TOLAZAMIDE

BARR	100MG	N70162	001	Jan 14, 1986
	250MG	N70163	001	Jan 14, 1986
	500MG	N70164	001	Jan 14, 1986
DURAMED PHARMS BARR	100MG	N70165	001	Jan 10, 1986
	250MG	N70166	001	Jan 10, 1986
	500MG	N70167	001	Jan 10, 1986
INTERPHARM	250MG	N71270	001	Sep 23, 1986
	500MG	N71271	001	Sep 23, 1986
SUPERPHARM	250MG	N70763	001	Jun 16, 1986
	500MG	N70764	001	Jun 16, 1986
USL PHARMA	100MG	N71355	001	Jan 11, 1988
	250MG	N70168	001	Apr 02, 1986

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 250 (of 263)

TOLAZAMIDETABLET; ORAL
TOLAZAMIDE

USL PHARMA	500MG	N70169	001	Apr 02, 1986
WATSON LABS	100MG	N70242	001	Aug 01, 1986
	100MG	N70513	001	Jan 09, 1986
	250MG	N70243	001	Aug 01, 1986
	500MG	N70244	001	Aug 01, 1986

TOLAZOLINE HYDROCHLORIDEINJECTABLE; INJECTION
PRISCOLINE
NOVARTIS

25MG/ML	N06403	005	Feb 22, 1985
---------	--------	-----	--------------

TOLBUTAMIDETABLET; ORAL
ORINASE

PHARMACIA AND UPJOHN	250MG	N10670	002	
	500MG	N10670	001	

TOLBUTAMIDE

ALRA	500MG	N86141	001	
ASCOT	500MG	N87541	001	Mar 01, 1983
BARR	500MG	N87121	001	
CLONMEL HLTHCARE	500MG	N86926	001	
PARKE DAVIS	500MG	N86047	001	
PUREPAC PHARM	500MG	N88950	001	Jun 17, 1985
SANDOZ	500MG	N12678	001	
SUPERPHARM	500MG	N88893	001	Nov 19, 1984
VANGARD	500MG	N87876	001	Apr 20, 1982
WATSON LABS	250MG	N89110	001	May 29, 1987
	500MG	N89111	001	May 29, 1987

TOLBUTAMIDE SODIUMINJECTABLE; INJECTION
ORINASE DIAGNOSTIC

PHARMACIA AND UPJOHN	EQ 1GM BASE/VIAL	N12095	001	
----------------------	------------------	--------	-----	--

TOLMETIN SODIUMCAPSULE; ORAL
TOLMETIN SODIUM

SANDOZ	EQ 400MG BASE	N73462	001	Apr 30, 1992
TEVA	EQ 400MG BASE	N73519	001	May 29, 1992

TABLET; ORAL
TOLMETIN SODIUM

TEVA	EQ 600MG BASE	N74729	001	Feb 27, 1997
------	---------------	--------	-----	--------------

TOPIRAMATE

CAPSULE; ORAL

TOPAMAX SPRINKLE

ORTHO MCNEIL PHARM	50MG	N20844	003	Oct 26, 1998
--------------------	------	--------	-----	--------------

TABLET; ORAL

TOPAMAX

ORTHO MCNEIL PHARM	300MG	N20505	003	Dec 24, 1996
	400MG	N20505	006	Dec 24, 1996

TRAMADOL HYDROCHLORIDE

TABLET; ORAL

TRAMADOL HYDROCHLORIDE

ASTA	50MG	N75974	001	Jul 12, 2002
------	------	--------	-----	--------------

ULTRAM

ORTHO MCNEIL PHARM	100MG	N20281	001	Mar 03, 1995
--------------------	-------	--------	-----	--------------

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 251 (of 263)

TRANEXAMIC ACIDTABLET; ORAL
CYKLOKAPRON

PHARMACIA AND UPJOHN 500MG

N19280 001 Dec 30, 1986

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

TRAZODONE HYDROCHLORIDE

AM THERAP

50MG

N71139 001 Oct 29, 1986

100MG

N71140 001 Oct 29, 1986

MYLAN

100MG

N71406 001 Feb 27, 1991

QUANTUM PHARMICS

100MG

N70921 001 Dec 01, 1986

USL PHARMA

50MG

N70491 001 Apr 29, 1987

100MG

N70492 001 Apr 29, 1987

WATSON LABS

50MG

N71112 001 Nov 17, 1986

100MG

N71113 001 Nov 17, 1986

TRIALODINE

QUANTUM PHARMICS

50MG

N70942 001 Dec 01, 1986

TRETINOIN

SWAB; TOPICAL

RETIN-A

JOHNSON AND JOHNSON 0.05%

N16921 002

TRIAMCINOLONE

TABLET; ORAL

ARISTOCORT

ASTELLAS

1MG

N11161 009

2MG

N11161 004

8MG

N11161 011

16MG

N11161 010

KENACORT

BRISTOL MYERS SQUIBB

1MG

N11283 003

2MG

N11283 008

8MG

N11283 010

TRIAMCINOLONE

BARR

2MG

N84286 001

2MG

N84318 001

4MG

N84267 001

4MG

N84319 001

8MG

N84268 001

8MG

N84320 001

IMPAX LABS

4MG

N84340 001

IVAX PHARMS

4MG

N83750 001

MYLAN

2MG

N84406 001

PUREPAC PHARM

2MG

N84020 002

4MG

N84020 003

ROXANE

2MG

N84708 001

4MG

N84709 001

8MG

N84707 001

SANDOZ

4MG

N85601 001

TEVA

4MG

N84775 001

WATSON LABS

4MG

N84270 001

4MG

N85834 001

TRIAMCINOLONE ACETONIDE

AEROSOL, METERED; NASAL

NASACORT

AVENTIS

0.055MG/INH

N19798 001 Jul 11, 1991

CREAM; TOPICAL

ARISTOCORT

ASTELLAS

0.025%

N83017 003

0.1%

N83016 004

0.5%

N83015 002

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 252 (of 263)

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

ARISTOCORT A

ASTELLAS

0.025%

N83017 004

0.1%

N83016 005

0.5%

N83015 003

FLUTEX

IVAX PHARMS

0.025%

N85539 001

0.1%

N85539 002

0.5%

N85539 003

KENALOG-H

APOTHECON

0.1%

N86240 001

TRIAECORT

SOLVAY

0.1%

N87113 001

TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS

0.025%

N87797 001

Jun 07, 1982

AMBIX

0.025%

N87932 001

May 09, 1983

MORTON GROVE

0.025%

N88094 001

Sep 01, 1983

0.1%

N88095 001

Sep 01, 1983

0.5%

N88096 001

Sep 01, 1983

PHARMADERM

0.025%

N87990 001

Jul 07, 1983

0.1%

N87991 001

Jul 07, 1983

0.5%

N87992 001

Jul 07, 1983

PHARMAFAIR

0.025%

N87921 001

Aug 10, 1982

0.1%

N87912 001

Aug 10, 1982

0.5%

N87922 001

Aug 10, 1982

TARO

0.025%

N40038 001

Oct 26, 1994

TOPIDERM

0.025%

N89274 001

Feb 21, 1989

0.1%

N89275 001

Feb 21, 1989

0.5%

N89276 001

Feb 21, 1989

TRIAECX

IVAX PHARMS

0.025%

N87430 001

Nov 01, 1988

0.1%

N87429 001

Nov 01, 1988

0.5%

N87428 001

Nov 01, 1988

TRYMEX

SAVAGE LABS

0.025%

N88196 001

Mar 25, 1983

0.1%

N88197 001

Mar 25, 1983

0.5%

N88198 001

Mar 25, 1983

GEL; TOPICAL

ARISTOGEL

ASTELLAS

0.1%

N83380 001

INJECTABLE; INJECTION

TRIAMCINOLONE ACETONIDE

PARNELL

3MG/ML

N19503 001

Oct 16, 1987

STERIS

40MG/ML

N85825 001

LOTION; TOPICAL

KENALOG

BRISTOL MYERS SQUIBB

0.025%

N11602 003

0.1%

N11602 001

TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS

0.025%

N87191 001

Sep 08, 1982

0.1%

N87192 001

Sep 08, 1982

OINTMENT; TOPICAL

ARISTOCORT

ASTELLAS

0.1%

N80750 004

0.5%

N80745 002

ARISTOCORT A

ASTELLAS

0.1%

N80750 003

0.1%

N88780 001

Oct 01, 1984

0.5%

N80745 003

0.5%

N88781 001

Oct 05, 1984

FLUTEX

IVAX PHARMS

0.025%

N87375 001

Nov 01, 1988

0.1%

N87377 001

Nov 01, 1988

0.5%

N87376 001

Nov 01, 1988

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 253 (of 263)

TRIAMCINOLONE ACETONIDE

OINTMENT; TOPICAL

KENALOG

APOTHECON

0.5%

N83944 001

TRIAMCINOLONE ACETONIDE

ALPHARMA US PHARMS

0.5%

N89913 001 Dec 23, 1988

G AND W LABS

0.025%

N89795 001 Dec 23, 1988

0.1%

N89796 001 Dec 23, 1988

MORTON GROVE

0.025%

N88090 001 Sep 01, 1983

0.1%

N88091 001 Sep 01, 1983

0.5%

N88092 001 Sep 01, 1983

PHARMADERM

0.025%

N88692 001 Aug 02, 1984

0.1%

N88690 001 Aug 02, 1984

TRYMEX

SAVAGE LABS

0.025%

N88693 001 Aug 02, 1984

0.1%

N88691 001 Aug 02, 1984

SPRAY, METERED; NASAL

TRI-NASAL

COLLEGIUM

0.05MG/SPRAY

N20120 001 Feb 04, 2000

TRIAMCINOLONE DIACETATE

INJECTABLE; INJECTION

ARISTOCORT

SABEX 2002

25MG/ML

N11685 003

40MG/ML

N12802 001

TRIAMCINOLONE DIACETATE

AKORN

25MG/ML

N85122 001

40MG/ML

N86394 001

STERIS

40MG/ML

N84072 001

40MG/ML

N85529 001

SYRUP; ORAL

ARISTOCORT

ASTELLAS

2MG/5ML

N11960 004

KENACORT

BRISTOL MYERS SQUIBB

EQ 4MG BASE/5ML

N12515 001

TRIAZOLAM

TABLET; ORAL

HALCION

PHARMACIA AND UPJOHN

0.5MG

N17892 002 Nov 15, 1982

TRICHLORMETHIAZIDE

TABLET; ORAL

METAHYDRIN

AVENTIS PHARMS

2MG

N12594 001 Jun 16, 1988

4MG

N12594 002 Jun 16, 1988

NAQUA

SCHERING

2MG

N12265 001

4MG

N12265 002

TRICHLOREX

LANNETT

4MG

N83436 001

4MG

N85630 001

TRICHLORMAS

MAST MM

4MG

N86259 001

TRICHLORMETHIAZIDE

ABC HOLDING

4MG

N85568 001

IMPAX LABS

4MG

N83967 001

PAR PHARM

2MG

N87007 001

4MG

N87005 001

SANDOZ

4MG

N86171 001

WATSON LABS

2MG

N83847 001

2MG

N86458 001

4MG

N83462 001

4MG

N83855 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 254 (of 263)

TRICHLORMETHIAZIDE

TABLET; ORAL				
TRICHLORMETHIAZIDE				
WATSON LABS	4MG	N85962	001	

TRICLOFOS SODIUM

SOLUTION; ORAL				
TRICLOS				
AVENTIS PHARMS	1.5GM/15ML	N16830	001	
TABLET; ORAL				
TRICLOS				
AVENTIS PHARMS	750MG	N16809	002	

TRIDIHEXETHYL CHLORIDE

INJECTABLE; INJECTION				
PATHILON				
LEDERLE	10MG/ML	N09729	001	
TABLET; ORAL				
PATHILON				
LEDERLE	25MG	N09489	005	

TRIFLUOPERAZINE HYDROCHLORIDE

CONCENTRATE; ORAL				
TRIFLUOPERAZINE HYDROCHLORIDE				
MORTON GROVE	EQ 10MG BASE/ML	N88143	001	Jul 26, 1983
SANDOZ	EQ 10MG BASE/ML	N85787	001	Apr 15, 1982
TABLET; ORAL				
TRIFLUOPERAZINE HYDROCHLORIDE				
DURAMED PHARMS BARR	EQ 1MG BASE	N88967	001	Apr 23, 1985
	EQ 2MG BASE	N88968	001	Apr 23, 1985
	EQ 5MG BASE	N88969	001	Apr 23, 1985
	EQ 10MG BASE	N88970	001	Apr 23, 1985
IVAX PHARMS	EQ 1MG BASE	N87612	001	Nov 19, 1982
	EQ 2MG BASE	N87613	001	Nov 19, 1982
	EQ 5MG BASE	N87328	001	Nov 19, 1982
	EQ 10MG BASE	N87614	001	Nov 19, 1982
WATSON LABS	EQ 1MG BASE	N85975	001	Jun 23, 1988
	EQ 2MG BASE	N85976	001	Jun 23, 1988
	EQ 5MG BASE	N85973	001	Jun 23, 1988
	EQ 10MG BASE	N88710	001	Jun 23, 1988

TRIFLUPROMAZINE

SUSPENSION; ORAL				
VESPRIN				
APOTHECON	EQ 50MG HCL/5ML	N11491	004	

TRIFLUPROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION				
VESPRIN				
APOTHECON	3MG/ML	N11325	005	
	10MG/ML	N11325	004	
	20MG/ML	N11325	001	
TABLET; ORAL				
VESPRIN				
BRISTOL MYERS SQUIBB	10MG	N11123	001	
	25MG	N11123	002	
	50MG	N11123	003	

TRIHXYPHENIDYL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL				
ARTANE				
LEDERLE	5MG	N06773	010	
	5MG	N12947	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 255 (of 263)

TRIHEXYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL				
ARTANE				
LEDERLE	2MG/5ML	N06773	009	
TABLET; ORAL				
ARTANE				
LEDERLE	2MG	N06773	005	
	5MG	N06773	003	
TREMIN				
SCHERING	2MG	N80381	001	
	5MG	N80381	003	
TRIHEXYPHENIDYL HYDROCHLORIDE				
NYLOS	5MG	N85622	001	
VANGARD	2MG	N88035	001	Jul 30, 1982
WATSON LABS	2MG	N85117	001	
	5MG	N85105	001	

TRILOSTANE

CAPSULE; ORAL				
MODRASTANE				
BIOENVISION	30MG	N18719	002	Dec 31, 1984
	60MG	N18719	001	Dec 31, 1984

TRIMEPAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL				
TEMARIL				
ALLERGAN HERBERT	EQ 5MG BASE	N11316	004	
SYRUP; ORAL				
TEMARIL				
ALLERGAN HERBERT	EQ 2.5MG BASE/5ML	N11316	003	
TRIMEPAZINE TARTRATE				
ALPHARMA US PHARMS	EQ 2.5MG BASE/5ML	N85015	001	Feb 18, 1982
MORTON GROVE	EQ 2.5MG BASE/5ML	N88285	001	Apr 11, 1985
TABLET; ORAL				
TEMARIL				
ALLERGAN HERBERT	EQ 2.5MG BASE	N11316	001	

TRIMETHADIONE

CAPSULE; ORAL				
TRIDIONE				
ABBOTT	300MG	N05856	005	
SOLUTION; ORAL				
TRIDIONE				
ABBOTT	200MG/5ML	N05856	002	

TRIMETHAPHAN CAMSYLATE

INJECTABLE; INJECTION				
ARFONAD				
ROCHE	50MG/ML	N08983	001	

TRIMETHOBENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION				
TRIMETHOBENZAMIDE HYDROCHLORIDE				
SMITH AND NEPHEW	100MG/ML	N88960	001	Apr 04, 1986
	100MG/ML	N89043	001	Apr 04, 1986
SOLOPAK	100MG/ML	N89094	001	Apr 04, 1986
STERIS	100MG/ML	N86577	001	Oct 19, 1982
	100MG/ML	N87939	001	Dec 28, 1982

TRIMETHOPRIM

TABLET; ORAL				
PROLOPRIM				
MONARCH PHARMS	200MG	N17943	003	Jul 14, 1982

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 256 (of 263)

TRIMETHOPRIM

TABLET; ORAL				
TRIMETHOPRIM				
MUTUAL PHARM	100MG	N70494	001	Jan 22, 1986
	200MG	N70495	001	Sep 24, 1986
TRIMPEX				
ROCHE	100MG	N17952	001	
TRIMPEX 200				
ROCHE	200MG	N17952	002	Nov 09, 1982

TRIMETHOPRIM HYDROCHLORIDE

SOLUTION; ORAL				
PRIMSOL				
FSC	EQ 25MG BASE/5ML	N74374	001	Jun 23, 1995

TRIMIPRAMINE MALEATE

CAPSULE; ORAL				
TRIMIPRAMINE MALEATE				
USL PHARMA	EQ 25MG BASE	N71283	001	Dec 08, 1987
	EQ 50MG BASE	N71284	001	Dec 08, 1987
	EQ 100MG BASE	N71285	001	Dec 08, 1987

TRIOXSALEN

TABLET; ORAL				
TRISORALEN				
VALEANT PHARM INTL	5MG	N12697	001	

TRIPLENNAMINE CITRATE

ELIXIR; ORAL				
PBZ				
NOVARTIS	EQ 25MG HCL/5ML	N05914	004	

TRIPLENNAMINE HYDROCHLORIDE

TABLET; ORAL				
PBZ				
NOVARTIS	25MG	N83149	001	
TRIPLENNAMINE HYDROCHLORIDE				
ANABOLIC	50MG	N83037	001	
BARR	50MG	N80744	001	
HEATHER	50MG	N83989	001	
IMPAX LABS	50MG	N80785	001	
LANNETT	50MG	N83557	001	
NYLOS	50MG	N85412	001	
PARKE DAVIS	25MG	N83625	001	
	50MG	N83626	001	
WATSON LABS	50MG	N80713	001	
	50MG	N80790	001	
	50MG	N85188	001	
TABLET, EXTENDED RELEASE; ORAL				
PBZ-SR				
NOVARTIS	50MG	N10533	002	
	100MG	N10533	001	

TRIPLE SULFA (SULFABENZAMIDE;SULFACETAMIDE;SULFATHIAZOLE)

CREAM; VAGINAL				
GYNE-SULF				
G AND W LABS	3.7%;2.86%;3.42%	N88607	001	Jun 09, 1986
SULTRIN				
ORTHO MCNEIL PHARM	3.7%;2.86%;3.42%	N05794	001	
TRIPLE SULFA				
ALPHARMA US PHARMS	3.7%;2.86%;3.42%	N87864	001	Sep 01, 1982
CLAY PARK	3.7%;2.86%;3.42%	N87285	001	Nov 15, 1982
FOUGERA	3.7%;2.86%;3.42%	N86424	001	

DISCONTINUED DRUG PRODUCT LIST

6 - 257 (of 263)

TRIPLE SULFA (SULFABENZAMIDE;SULFACETAMIDE;SULFATHIAZOLE)

CREAM; VAGINAL				
TRYUL				
SAVAGE LABS	3.7%;2.86%;3.42%	N87887	001	Jul 23, 1982
VAGILIA				
TEVA	3.7%;2.86%;3.42%	N88821	001	Nov 09, 1987
TABLET; VAGINAL				
SULTRIN				
ORTHO MCNEIL PHARM	184MG;143.75MG;172.5MG	N05794	002	
TRIPLE SULFA				
FOUGERA	184MG;143.75MG;172.5MG	N88463	001	Jan 03, 1985
PHARMADERM	184MG;143.75MG;172.5MG	N88462	001	Jan 03, 1985

TRIPROLIDINE HYDROCHLORIDE

SYRUP; ORAL				
ACTIDIL				
GLAXOSMITHKLINE	1.25MG/5ML	N11496	002	Jul 01, 1983
MYIDYL				
USL PHARMA	1.25MG/5ML	N87963	001	Jan 18, 1983
TRIPROLIDINE HYDROCHLORIDE				
ALPHARMA US PHARMS	1.25MG/5ML	N85940	001	
HALSEY	1.25MG/5ML	N88735	001	Jan 17, 1985
PHARM ASSOC	1.25MG/5ML	N87514	001	Feb 10, 1982
TABLET; ORAL				
ACTIDIL				
GLAXOSMITHKLINE	2.5MG	N11110	002	Jul 01, 1983
TRIPROLIDINE HYDROCHLORIDE				
VITARINE	2.5MG	N85610	001	
WATSON LABS	2.5MG	N85094	001	

TRISULFAPYRIMIDINES (SULFADIAZINE;SULFAMERAZINE;SULFAMETHAZINE)

SUSPENSION; ORAL				
LANTRISUL				
LANNETT	167MG/5ML;167MG/5ML;167MG/5ML	N80123	002	
NEOTRIZINE				
LILLY	167MG/5ML;167MG/5ML;167MG/5ML	N06317	012	
SULFALOID				
FOREST PHARMS	167MG/5ML;167MG/5ML;167MG/5ML	N80100	001	
SULFOSE				
WYETH AYERST	167MG/5ML;167MG/5ML;167MG/5ML	N80013	002	
TERFONYL				
BRISTOL MYERS SQUIBB	167MG/5ML;167MG/5ML;167MG/5ML	N06904	002	
TRIPLE SULFA				
ALPHARMA US PHARMS	167MG/5ML;167MG/5ML;167MG/5ML	N80280	001	
TRIPLE SULFAS				
LEDERLE	167MG/5ML;167MG/5ML;167MG/5ML	N06920	003	
TABLET; ORAL				
NEOTRIZINE				
LILLY	167MG;167MG;167MG	N06317	011	
SULFALOID				
FOREST PHARMS	167MG;167MG;167MG	N80099	001	
SULFA-TRIPLE #2				
IMPAX LABS	167MG;167MG;167MG	N80079	001	
SULFOSE				
WYETH AYERST	167MG;167MG;167MG	N80013	001	
TERFONYL				
BRISTOL MYERS SQUIBB	167MG;167MG;167MG	N06904	001	
TRIPLE SULFA				
PUREPAC PHARM	167MG;167MG;167MG	N80086	001	
TRIPLE SULFAS				
LEDERLE	167MG;167MG;167MG	N06920	002	
TRIPLE SULFOID				
PAL PAK	167MG;167MG;167MG	N80094	001	

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 258 (of 263)

TROGLITAZONETABLET; ORAL
PRELAY

SANKYO	200MG	N20719	001	Jan 29, 1997
	300MG	N20719	003	Aug 04, 1997
	400MG	N20719	002	Jan 29, 1997

REZULIN

PFIZER PHARMS	200MG	N20720	001	Jan 29, 1997
	300MG	N20720	003	Aug 04, 1997
	400MG	N20720	002	Jan 29, 1997

TROLAMINE POLYPEPTIDE OLEATE CONDENSATESOLUTION/DROPS; OTIC
CERUMENEX

PURDUE FREDERICK	10%	N11340	002	
------------------	-----	--------	-----	--

TROLEANDOMYCINSUSPENSION; ORAL
TAO

PFIZER	EQ 125MG BASE/5ML	N50332	001	
--------	-------------------	--------	-----	--

TROPICAMIDESOLUTION/DROPS; OPHTHALMIC
MYDRIACYL

ALCON	0.5%	N12111	002	
	1%	N12111	004	

MYDRIAFAIR

PHARMAFAIR	0.5%	N88274	001	Sep 16, 1983
	1%	N88230	001	Sep 16, 1983

TROPICAMIDE

AKORN	1%	N88447	001	Aug 28, 1985
STERIS	0.5%	N89171	001	Dec 28, 1990

TROVAFLOXACIN MESYLATETABLET; ORAL
TROVAN

PFIZER	EQ 100MG BASE	N20759	001	Dec 18, 1997
	EQ 200MG BASE	N20759	002	Dec 18, 1997

TUBOCURARINE CHLORIDEINJECTABLE; INJECTION
TUBOCURARINE CHLORIDE

BRISTOL MYERS SQUIBB	3MG/ML	N05657	001	
HOSPIRA	3MG/ML	N06095	001	
LILLY	3MG/ML	N06325	001	

TYROPANOATE SODIUMCAPSULE; ORAL
BILOPAQUE

GE HEALTHCARE	750MG	N13731	001	
---------------	-------	--------	-----	--

UNDECOYLUM CHLORIDE; UNDECOYLUM CHLORIDE IODINE COMPLEXSOLUTION; TOPICAL
VIRAC REX

CHESEBROUGH PONDS	0.5%;1.8%	N11914	001	
-------------------	-----------	--------	-----	--

URACIL MUSTARDCAPSULE; ORAL
URACIL MUSTARD

SHIRE	1MG	N12892	001	
-------	-----	--------	-----	--

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 259 (of 263)

UREA

INJECTABLE; INJECTION

STERILE UREA

HOSPIRA

40GM/VIAL

N17698 001

UREA, C-13

FOR SOLUTION; ORAL

HELICOSOL

METABOLIC SOLUTIONS 125MG/VIAL

N21092 001 Dec 17, 1999

MERETEK UBT KIT (W/ PRANACTIN)

MERETEK 125MG/VIAL

N20586 001 Sep 17, 1996

PYLORI-CHEK BREATH TEST

DXS DEVICES 100MG/VIAL

N20900 001 Feb 04, 1999

UROFOLLITROPIN

INJECTABLE; INTRAMUSCULAR

METRODIN

SERONO

75 IU/AMP

N19415 002 Sep 18, 1986

150 IU/AMP

N19415 003 Sep 18, 1986

INJECTABLE; SUBCUTANEOUS

FERTINEX

SERONO

75 IU/AMP

N19415 005 Aug 23, 1996

150 IU/AMP

N19415 004 Aug 23, 1996

UROKINASE

INJECTABLE; INJECTION

ABBOKINASE

ABBOTT

5,000 IU/VIAL

N21846 003

9,000 IU/VIAL

N21846 002

URSODIOL

CAPSULE; ORAL

ACTIGALL

WATSON PHARMS

150MG

N19594 001 Dec 31, 1987

VALPROIC ACID

CAPSULE; ORAL

VALPROIC ACID

PAR PHARM

250MG

N70431 001 Feb 28, 1986

SCHERER RP

250MG

N70195 001 Jul 02, 1987

VALSARTAN

CAPSULE; ORAL

DIOVAN

NOVARTIS

80MG

N20665 001 Dec 23, 1996

160MG

N20665 002 Dec 23, 1996

VANCOMYCIN HYDROCHLORIDE

FOR SOLUTION; ORAL

VANCOCIN HYDROCHLORIDE

VIROPHARMA

EQ 250MG BASE/5ML

N61667 002 Jul 13, 1983

EQ 500MG BASE/6ML

N61667 001

VANCOLED

LEDERLE

EQ 250MG BASE/5ML

N63321 002 Oct 15, 1993

EQ 500MG BASE/6ML

N63321 003 Oct 15, 1993

INJECTABLE; INJECTION

VANCOCIN HYDROCHLORIDE

VIROPHARMA

EQ 500MG BASE/VIAL

N60180 001

EQ 500MG BASE/VIAL

N62476 001 Mar 15, 1984

EQ 500MG BASE/VIAL

N62716 001 Mar 13, 1987

EQ 500MG BASE/VIAL

N62812 001 Nov 17, 1987

EQ 1GM BASE/VIAL

N60180 002 Mar 21, 1986

EQ 1GM BASE/VIAL

N62476 002 Mar 21, 1986

EQ 1GM BASE/VIAL

N62716 002 Mar 13, 1987

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 260 (of 263)

VANCOMYCIN HYDROCHLORIDE

INJECTABLE; INJECTION
 VANCOCIN HYDROCHLORIDE
 VIROPHARMA

EQ 1GM BASE/VIAL
 EQ 10GM BASE/VIAL

N62812 002 Nov 17, 1987
 N62812 003 Nov 17, 1987

VANCOLED

BAXTER HLTHCARE

EQ 500MG BASE/VIAL
 EQ 1GM BASE/VIAL
 EQ 2GM BASE/VIAL
 EQ 5GM BASE/VIAL
 EQ 10GM BASE/VIAL

N62682 001 Jul 22, 1986
 N62682 002 Mar 30, 1988
 N62682 003 May 11, 1988
 N62682 004 May 11, 1988
 N62682 005 May 11, 1988

VANCOMYCIN HYDROCHLORIDE

BAXTER HLTHCARE

EQ 500MG BASE/VIAL
 EQ 1GM BASE/VIAL

N62879 001 Aug 02, 1988
 N62879 002 Aug 02, 1988

VANCOR

PHARMACIA AND UPJOHN

EQ 500MG BASE/VIAL
 EQ 1GM BASE/VIAL

N62956 001 Aug 01, 1988
 N62956 002 Aug 01, 1988

VASOPRESSIN TANNATE

INJECTABLE; INJECTION
 PITRESSIN TANNATE
 PARKE DAVIS

5PRESSOR UNITS/ML

N03402 001

VECURONIUM BROMIDE

INJECTABLE; INJECTION
 NORCURON

ORGANON USA INC

10MG/VIAL
 20MG/VIAL

N18776 002 Apr 30, 1984
 N18776 003 Jan 03, 1992

VENLAFAXINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
 EFFEXOR XR

WYETH PHARMS INC

EQ 100MG BASE

N20699 003 Oct 20, 1997

TABLET; ORAL
 EFFEXOR

WYETH PHARMS INC

EQ 12.5MG BASE

N20151 001 Dec 28, 1993

VERAPAMIL HYDROCHLORIDE

INJECTABLE; INJECTION
 CALAN

GD SEARLE LLC

2.5MG/ML
 2.5MG/ML

N18925 001 Mar 30, 1984
 N19038 001 Mar 30, 1984

VERAPAMIL HYDROCHLORIDE

AM PHARM PARTNERS

2.5MG/ML

N70348 001 May 01, 1986

HOSPIRA

2.5MG/ML

N70739 001 May 06, 1987

2.5MG/ML

N70740 001 May 06, 1987

MARSAM PHARMS LLC

2.5MG/ML

N72233 001 Feb 26, 1993

2.5MG/ML

N73485 001 Sep 27, 1993

SMITH AND NEPHEW

2.5MG/ML

N70696 001 Jul 31, 1987

2.5MG/ML

N70697 001 Jul 31, 1987

SOLOPAK

2.5MG/ML

N70695 001 Jul 31, 1987

TABLET; ORAL
 CALAN

GD SEARLE LLC

160MG

N18817 004 Feb 23, 1988

VERAPAMIL HYDROCHLORIDE

CLONMEL HLTHCARE

80MG
 120MG

N71880 001 Apr 05, 1988
 N71881 001 Apr 05, 1988

MUTUAL PHARM

80MG

N70482 001 Sep 24, 1986

120MG

N70483 001 Sep 24, 1986

PLIVA

40MG

N72751 001 Feb 23, 1996

WARNER CHILCOTT

80MG

N70340 001 Sep 24, 1986

120MG

N70341 001 Sep 24, 1986

WATSON LABS

40MG

N72799 001 Apr 28, 1989

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 261 (of 263)

VERATRUM VIRIDETABLET; ORAL
VERTAVIS

MEDPOINTE PHARM HLC 130CSR UNIT N05691 002

VIDARABINEINJECTABLE; INJECTION
VIRA-A

PARKEDALE EQ 187.4MG BASE/ML N50523 001

OINTMENT; OPHTHALMIC
VIRA-A

PARKEDALE 3% N50486 001

VINBLASTINE SULFATEINJECTABLE; INJECTION
VELBAN

LILLY 10MG/VIAL N12665 001

VINBLASTINE SULFATE

AM PHARM PARTNERS 10MG/VIAL N89011 001 Nov 18, 1985

MAYNE PHARMA USA 10MG/VIAL N89565 001 Aug 18, 1987

VINCRISTINE SULFATEINJECTABLE; INJECTION
ONCOVIN

LILLY 1MG/VIAL N14103 001

1MG/ML N14103 003 Mar 07, 1984

5MG/VIAL N14103 002

VINCASAR PFS

SICOR PHARMS 1MG/ML N71426 001 Jul 17, 1987

VINCREX

BRISTOL MYERS SQUIBB 5MG/VIAL N70867 001 Jul 12, 1988

VINCRISTINE SULFATE

ABIC 1MG/ML N70873 001 Feb 19, 1987

AM PHARM PARTNERS 1MG/ML N70411 001 Sep 10, 1986

MAYNE PHARMA USA 1MG/VIAL N71559 001 Apr 11, 1988

2MG/VIAL N71560 001 Apr 11, 1988

5MG/VIAL N71561 001 Apr 11, 1988

VIOMYCIN SULFATEINJECTABLE; INJECTION
VIOCIN SULFATE

PFIZER EQ 1GM BASE/VIAL N61086 001

EQ 5GM BASE/VIAL N61086 002

VITAMIN ACAPSULE; ORAL
AQUASOL A

ASTRAZENECA 25,000USP UNITS N83080 002

50,000USP UNITS N83080 001

VITAMIN A

BANNER PHARMACAPS 50,000USP UNITS N83973 001

CHASE CHEM 50,000 IU N83351 001

EVERYLIFE 50,000 IU N83134 001

IMPAX LABS 50,000USP UNITS N80952 001

WEST WARD 50,000USP UNITS N80985 001

VITAMIN A PALMITATECAPSULE; ORAL
AFAXIN

STERLING WINTHROP EQ 50,000 UNITS BASE N83187 001

ALPHALIN

LILLY EQ 50,000 UNITS BASE N80883 001

DEL-VI-A

DEL RAY LABS EQ 50,000 UNITS BASE N80830 001

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 262 (of 263)

VITAMIN A PALMITATE

CAPSULE; ORAL

VI-DOM-A

BAYER PHARMS

EQ 50,000 UNITS BASE

N80972 001

VITAMIN A

BANNER PHARMACAPS

EQ 50,000 UNITS BASE

N80702 001

BRISTOL MYERS SQUIBB

EQ 50,000 UNITS BASE

N80860 001

CHASE CHEM

EQ 50,000 UNITS BASE

N80746 001

ELKINS SINN

EQ 50,000 UNITS BASE

N83207 001

EVERYLIFE

EQ 50,000 UNITS BASE

N85479 001

IMPAX LABS

EQ 50,000 UNITS BASE

N80943 001

IVAX PHARMS

EQ 50,000 UNITS BASE

N83114 001

MK LABS

EQ 50,000 UNITS BASE

N80953 001

WEST WARD

EQ 50,000 UNITS BASE

N80955 001

WHARTON LABS

EQ 50,000 UNITS BASE

N83035 001

VITAMIN A PALMITATE

BANNER PHARMACAPS

EQ 50,000 UNITS BASE

N83190 001

TEVA

EQ 25,000 UNITS BASE

N83457 002

INJECTABLE; INJECTION

VITAMIN A PALMITATE

BANNER PHARMACAPS

EQ 50,000 UNITS BASE

N83457 001

WHARTON LABS

EQ 50,000 UNITS BASE

N80967 001

VITAMIN A SOLUBILIZED

BANNER PHARMACAPS

EQ 50,000 UNITS BASE

N83665 001

TEVA

EQ 50,000 UNITS BASE

N83948 001

INJECTABLE; INJECTION

VITAMIN A PALMITATE

BEL MAR

EQ 50,000 UNITS BASE/ML

N83981 001

WARFARIN POTASSIUM

TABLET; ORAL

ATHROMBIN-K

PURDUE FREDERICK

2MG

N11771 007

5MG

N11771 004

10MG

N11771 005

25MG

N11771 006

WARFARIN SODIUM

INJECTABLE; INJECTION

COUMADIN

BRISTOL MYERS SQUIBB

50MG/VIAL

N09218 020

75MG/VIAL

N09218 012

TABLET; ORAL

ATHROMBIN

PURDUE FREDERICK

5MG

N11771 003

10MG

N11771 002

25MG

N11771 001

PANWARFIN

ABBOTT

2MG

N17020 001

2.5MG

N17020 002

5MG

N17020 003

7.5MG

N17020 004

10MG

N17020 005

WARFARIN SODIUM

USL PHARMA

2MG

N88719 001

2.5MG

N88720 001

5MG

N88721 001

WATSON LABS

2MG

N86123 001

2.5MG

N86120 001

5MG

N86119 001

7.5MG

N86118 001

10MG

N86122 001

Jun 27, 1985

Aug 06, 1985

Jul 02, 1985

Aug 17, 1982

Aug 17, 1982

Aug 17, 1982

Aug 17, 1982

Aug 17, 1982

26TH EDITION - 2006 - APPROVED DRUG PRODUCTS LIST

DISCONTINUED DRUG PRODUCT LIST

6 - 263 (of 263)

WATER FOR INJECTION, STERILE

LIQUID; N/A

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER

AM PHARM PARTNERS 100%

N89099 001 Dec 29, 1987

100%

N89100 001 Dec 29, 1987

STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

B BRAUN 100%

N19077 001 Mar 02, 1984

WATER FOR IRRIGATION, STERILE

LIQUID; IRRIGATION

STERILE WATER IN PLASTIC CONTAINER

MILES 100%

N18246 001

XENON, XE-127

GAS; INHALATION

XENON XE 127

MALLINCKRODT 5mCi/VIAL

N18536 001 Oct 01, 1982

10mCi/VIAL

N18536 002 Oct 01, 1982

XENON, XE-133

GAS; INHALATION

XENON XE 133

GE HEALTHCARE 1 CI/AMP

N17256 002

10mCi/VIAL

N17687 002

20mCi/VIAL

N17687 003

GEN ELECTRIC 5-100 CI/CYLINDER

N17550 001

0.25-5 CI/AMP

N17550 003

XENON XE 133-V.S.S.

GE HEALTHCARE 10mCi/VIAL

N17687 001

INJECTABLE; INJECTION

XENON XE 133

BRISTOL MYERS SQUIBB 6.3mCi/ML

N17283 001

GE HEALTHCARE 1.3-1.7 CI/AMP

N17256 001

SOLUTION; INHALATION, INJECTION

XENEISOL

MALLINCKRODT 18-25mCi/AMP

N17262 002

XYLOSE

POWDER; ORAL

XYLO-PFAN

SAVAGE LABS 25GM/BOT

N17605 001

XYLOSE

LYNE 25GM/BOT

N18856 001 Mar 26, 1987

ZICONOTIDE

INJECTABLE; INTRATHECAL

PRIALT

ELAN PHARMS 200UGM/2ML (100MG/ML)

N21060 003 Dec 28, 2004

ZIDOVUDINE

TABLET; ORAL

RETROVIR

GLAXOSMITHKLINE 200MG

N20518 001 Dec 19, 1995

ZILEUTON

TABLET; ORAL

ZYFLO

CRITICAL 300MG

N20471 001 Dec 09, 1996

ZINC SULFATE

INJECTABLE; INJECTION

ZINC SULFATE

AM PHARM PARTNERS EQ 1MG ZINC/ML

N19229 002 May 05, 1987

ORPHAN PRODUCT DESIGNATIONS AND APPROVALS LIST

The list of List of Orphan Designations and Approvals is available at:
<http://www.fda.gov/orphan/designat/list.htm>

**DRUG PRODUCTS WHICH MUST DEMONSTRATE *IN VIVO* BIOAVAILABILITY
ONLY IF PRODUCT FAILS TO ACHIEVE ADEQUATE DISSOLUTION**

ACETAMINOPHEN; ASPIRIN;
BUTALBITAL
CAPSULE OR TABLET; ORAL
160-165MG; 160-165MG; 50MG

ASPIRIN; CAFFEINE;
CARISOPRODOL
TABLET; ORAL
160MG; 32MG; 200MG

ACETAMINOPHEN; ASPIRIN;
BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG; 325MG; 50MG

ASPIRIN; CAFFEINE;
CARISOPRODOL;
CODEINE PHOSPHATE
TABLET; ORAL
160MG; 32MG;
200MG; 16MG

ACETAMINOPHEN; ASPIRIN;
BUTALBITAL; CAFFEINE
CAPSULE OR TABLET; ORAL
160-165MG; 160-165MG; 50MG; 40MG

ASPIRIN;
CARISOPRODOL
TABLET; ORAL
325MG; 200MG

ACETAMINOPHEN; ASPIRIN;
BUTALBITAL; CAFFEINE
CAPSULE OR TABLET; ORAL
325MG; 325MG; 50MG; 40MG

ASPIRIN;
CARISOPRODOL;
CODEINE PHOSPHATE
TABLET; ORAL
325MG; 200MG; 16MG

ACETAMINOPHEN; BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG; 50MG
650MG; 50MG

ASPIRIN; MEPROBAMATE
TABLET; ORAL
325MG; 200MG

ACETAMINOPHEN; BUTALBITAL;
CAFFEINE
CAPSULE OR TABLET; ORAL
325MG; 50MG; 40MG
650MG; 50MG; 40MG
500MG; 50MG; 40MG

ASPIRIN;
METHOCARBAMOL
TABLET; ORAL
325MG; 400MG

AMINOPHYLLINE
TABLET; ORAL
100MG; 200MG

CHLOROTHIAZIDE
TABLET; ORAL
250MG

ASPIRIN; BUTALBITAL
CAPSULE OR TABLET; ORAL
325MG; 50MG
650MG; 50MG

HYDROXYZINE
HYDROCHLORIDE
TABLET; ORAL
10MG; 25MG;
50MG; 100MG

ASPIRIN; BUTALBITAL; CAFFEINE
CAPSULE OR TABLET; ORAL
325MG; 50MG; 40MG
650MG; 50MG; 40MG

PREDNISONE
TABLET; ORAL
1MG; 2.5MG;
5MG; 10MG;
20MG; 25MG; 50MG

APPENDIX A - PRODUCT NAME INDEX

A - 1

**** 8 ****

8-MOP, METHOXSALEN

**** A ****

A/T/S, ERYTHROMYCIN
 ABBOKINASE, UROKINASE
 ABELCET, AMPHOTERICIN B
 ABILIFY, ARIPIRAZOLE
 ABRAXANE, PACLITAXEL
 ABREVA, DOCOSANOL (OTC)
 ACCOLATE, ZAFIRLUKAST
 ACCUNEB, ALBUTEROL SULFATE
 ACCUPRIL, QUINAPRIL HYDROCHLORIDE
 ACCURETIC, HYDROCHLOROTHIAZIDE
 ACCUTANE, ISOTRETINOIN
 ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 ACEON, PERINDOPRIL ERBUMINE
 ACEPHEN, ACETAMINOPHEN (OTC)
 ACETADOTE, ACETYLCYSTEINE
 ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE, ACETAMINOPHEN
 ACETAMINOPHEN W/ CODEINE NO. 3, ACETAMINOPHEN
 ACETAMINOPHEN W/ CODEINE PHOSPHATE #3, ACETAMINOPHEN
 ACETAMINOPHEN W/ CODEINE, ACETAMINOPHEN
 ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
 ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAMINOPHEN, BUTALBITAL, AND CAFFEINE, ACETAMINOPHEN
 ACETAMINOPHEN, BUTALBITAL, CAFFEINE, AND CODEINE PHOSPHATE, ACETAMINOPHEN
 ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
 ACETASOL HC, ACETIC ACID, GLACIAL
 ACETASOL, ACETIC ACID, GLACIAL
 ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
 ACETAZOLAMIDE, ACETAZOLAMIDE
 ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
 ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE, ACETIC ACID, GLACIAL
 ACETIC ACID, ACETIC ACID, GLACIAL
 ACETOHEXAMIDE, ACETOHEXAMIDE
 ACETYLCYSTEINE, ACETYLCYSTEINE
 ACILAC, LACTULOSE
 ACIPHEX, RABEPRAZOLE SODIUM
 ACLOVATE, ALCLOMETASONE DIPROPIONATE
 ACTHREL, CORTICORELIN OVINE TRIFLUTATE
 ACTIGALL, URSODIOL
 ACTIQ, FENTANYL CITRATE
 ACTIVELLA, ESTRADIOL
 ACTONEL WITH CALCIUM (COPACKAGED), CALCIUM CARBONATE
 ACTONEL, RISEDRONATE SODIUM
 ACTOPLUS MET, METFORMIN
 ACTOS, PIOGLITAZONE HYDROCHLORIDE
 ACTRON, KETOPROFEN (OTC)
 ACULAR LS, KETOROLAC TROMETHAMINE
 ACULAR PRESERVATIVE FREE, KETOROLAC TROMETHAMINE
 ACULAR, KETOROLAC TROMETHAMINE
 ACUTECT, TECHNETIUM TC-99M APCITIDE
 ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE, ACYCLOVIR SODIUM
 ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 ACYCLOVIR, ACYCLOVIR
 ACYCLOVIR, ACYCLOVIR SODIUM
 ADAGEN, PEGADEMASE BOVINE
 ADALAT CC, NIFEDIPINE
 ADDERALL 10, AMPHETAMINE ASPARTATE
 ADDERALL 12.5, AMPHETAMINE ASPARTATE
 ADDERALL 15, AMPHETAMINE ASPARTATE

APPENDIX A - PRODUCT NAME INDEX

A - 2

**** A ****

ADDERALL 20, AMPHETAMINE ASPARTATE
ADDERALL 30, AMPHETAMINE ASPARTATE
ADDERALL 5, AMPHETAMINE ASPARTATE
ADDERALL 7.5, AMPHETAMINE ASPARTATE
ADDERALL XR 10, AMPHETAMINE ASPARTATE
ADDERALL XR 15, AMPHETAMINE ASPARTATE
ADDERALL XR 20, AMPHETAMINE ASPARTATE
ADDERALL XR 25, AMPHETAMINE ASPARTATE
ADDERALL XR 30, AMPHETAMINE ASPARTATE
ADDERALL XR 5, AMPHETAMINE ASPARTATE
ADENOCARD, ADENOSINE
ADENOSCAN, ADENOSINE
ADENOSINE, ADENOSINE
ADIPEX-P, PHENTERMINE HYDROCHLORIDE
ADRIAMYCIN PFS, DOXORUBICIN HYDROCHLORIDE
ADRUCIL, FLUOROURACIL
ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
ADVICOR, LOVASTATIN
ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL COLD AND SINUS, IBUPROFEN (OTC)
ADVIL COLD AND SINUS, IBUPROFEN POTASSIUM (OTC)
ADVIL LIQUI-GELS, IBUPROFEN POTASSIUM (OTC)
ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN POTASSIUM (OTC)
ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ADVIL, IBUPROFEN (OTC)
AEROBID, FLUNISOLIDE
AFRINOL, PSEUDOEPHEDRINE SULFATE (OTC)
AGENERASE, AMPRENAVIR
AGGRASTAT, TIROFIBAN HYDROCHLORIDE
AGGRENOX, ASPIRIN
AGRYLIN, ANAGRELIDE HYDROCHLORIDE
A-HYDROCORT, HYDROCORTISONE SODIUM SUCCINATE
AKBETA, LEVOBUNOLOL HYDROCHLORIDE
AKINETON, BIPERIDEN HYDROCHLORIDE
AKNE-MYCIN, ERYTHROMYCIN
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
AKPRO, DIPIVEFRIN HYDROCHLORIDE
AKTOB, TOBRAMYCIN
ALA-CORT, HYDROCORTISONE
ALAMAST, PEMIROLAST POTASSIUM
ALA-SCALP, HYDROCORTISONE
ALAVERT, LORATADINE (OTC)
ALBALON, NAPHAZOLINE HYDROCHLORIDE
ALBENZA, ALBENDAZOLE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALBUTEROL, ALBUTEROL
ALCAINE, PROPARACAIN HYDROCHLORIDE
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
ALCOHOL 10% AND DEXTROSE 5%, ALCOHOL
ALCOHOL 5% AND DEXTROSE 5%, ALCOHOL
ALCOHOL 5% IN D5-W, ALCOHOL
ALDACTAZIDE, HYDROCHLOROTHIAZIDE
ALDACTONE, SPIRONOLACTONE
ALDARA, IMIQUIMOD
ALDORIL 15, HYDROCHLOROTHIAZIDE
ALDORIL 25, HYDROCHLOROTHIAZIDE
ALDORIL D30, HYDROCHLOROTHIAZIDE
ALDORIL D50, HYDROCHLOROTHIAZIDE
ALESSE, ETHINYL ESTRADIOL
ALEVE COLD AND SINUS, NAPROXEN SODIUM (OTC)
ALEVE, NAPROXEN SODIUM (OTC)

APPENDIX A - PRODUCT NAME INDEX

A - 3

**** A ****

ALFENTA, ALFENTANIL HYDROCHLORIDE
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
ALIMTA, PEMETREXED DISODIUM
ALINIA, NITAZOXANIDE
ALKERAN, MELPHALAN
ALKERAN, MELPHALAN HYDROCHLORIDE
ALLAY, ACETAMINOPHEN
ALLEGRA D 24 HOUR, FEXOFENADINE HYDROCHLORIDE
ALLEGRA, FEXOFENADINE HYDROCHLORIDE
ALLEGRA-D 12 HOUR, FEXOFENADINE HYDROCHLORIDE
ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
ALLOPURINOL, ALLOPURINOL
ALOCRIL, NEDOCROMIL SODIUM
ALOMIDE, LODOXAMIDE TROMETHAMINE
ALOPRIM, ALLOPURINOL SODIUM
ALORA, ESTRADIOL
ALOXI, PALONOSETRON HYDROCHLORIDE
ALPHAGAN P, BRIMONIDINE TARTRATE
ALPHATREX, BETAMETHASONE DIPROPIONATE
ALPRAZOLAM, ALPRAZOLAM
ALPROSTADIL, ALPROSTADIL
ALREX, LOTEPREDNOL ETABONATE
ALTACE, RAMIPRIL
ALTOPREV, LOVASTATIN
ALUPENT, METAPROTERENOL SULFATE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMARYL, GLIMEPIRIDE
AMBENYL, BROMODIPHENHYDRAMINE HYDROCHLORIDE
AMBIEN CR, ZOLPIDEM TARTRATE
AMBIEN, ZOLPIDEM TARTRATE
AMBISOME, AMPHOTERICIN B
AMCINONIDE, AMCINONIDE
AMERGE, NARATRIPTAN HYDROCHLORIDE
A-METHAPRED, METHYLPREDNISOLONE SODIUM SUCCINATE
AMICAR, AMINOCAPROIC ACID
AMIDATE, ETOMIDATE
AMIKACIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, AMIKACIN SULFATE
AMIKACIN SULFATE, AMIKACIN SULFATE
AMIKIN, AMIKACIN SULFATE
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOCAPROIC ACID, AMINOCAPROIC ACID
AMINOCAPROIC, AMINOCAPROIC ACID
AMINOHIPPURATE SODIUM, AMINOHIPPURATE SODIUM
AMINOPHYLLINE DYE FREE, AMINOPHYLLINE
AMINOPHYLLINE IN SODIUM CHLORIDE 0.45%, AMINOPHYLLINE
AMINOPHYLLINE, AMINOPHYLLINE
AMINOSYN 10% (PH6), AMINO ACIDS
AMINOSYN 10%, AMINO ACIDS
AMINOSYN 3.5% M, AMINO ACIDS
AMINOSYN 3.5%, AMINO ACIDS
AMINOSYN 5%, AMINO ACIDS
AMINOSYN 7% (PH6), AMINO ACIDS
AMINOSYN 7% W/ ELECTROLYTES, AMINO ACIDS
AMINOSYN 7%, AMINO ACIDS
AMINOSYN 8.5% (PH6), AMINO ACIDS
AMINOSYN 8.5% W/ ELECTROLYTES, AMINO ACIDS
AMINOSYN 8.5%, AMINO ACIDS
AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 10% W/ ELECTROLYTES, AMINO ACIDS
AMINOSYN II 10%, AMINO ACIDS
AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS

APPENDIX A - PRODUCT NAME INDEX

A - 4

**** A ****

AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 3.5% M IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 7%, AMINO ACIDS
AMINOSYN II 8.5% W/ ELECTROLYTES, AMINO ACIDS
AMINOSYN II 8.5%, AMINO ACIDS
AMINOSYN-HBC 7%, AMINO ACIDS
AMINOSYN-HF 8%, AMINO ACIDS
AMINOSYN-PF 10%, AMINO ACIDS
AMINOSYN-PF 7%, AMINO ACIDS
AMINOSYN-RF 5.2%, AMINO ACIDS
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMLEXANOX, AMLEXANOX
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE
AMMONIUM LACTATE, AMMONIUM LACTATE
AMMONUL, SODIUM BENZOATE
AMNESTEEM, ISOTRETINOIN
AMOXAPINE, AMOXAPINE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN PEDIATRIC, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
AMOXIL, AMOXICILLIN
AMPHADASE, HYALURONIDASE
AMPHOTEC, AMPHOTERICIN B
AMPHOTERICIN B, AMPHOTERICIN B
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
AMPICILLIN SODIUM, AMPICILLIN SODIUM
AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
AMRINONE LACTATE, INAMRINONE LACTATE
AMRINONE, INAMRINONE LACTATE
ANADROL-50, OXYMETHOLONE
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ANAPROX DS, NAPROXEN SODIUM
ANAPROX, NAPROXEN SODIUM
ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
ANCEF, CEFAZOLIN SODIUM
ANCOBON, FLUCYTOSINE
ANDRODERM, TESTOSTERONE
ANDROGEL, TESTOSTERONE
ANDROID 10, METHYLTESTOSTERONE
ANDROID 25, METHYLTESTOSTERONE
AN-DTPA, TECHNETIUM TC-99M PENTETATE KIT
ANECTINE, SUCCINYLCHOLINE CHLORIDE
ANESTACON, LIDOCAINE HYDROCHLORIDE
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
ANEXSIA 7.5/650, ACETAMINOPHEN
ANEXSIA, ACETAMINOPHEN
ANGELIQ, DROSPIRENONE
ANGIOMAX, BIVALIRUDIN
ANSAID, FLURBIPROFEN
ANSPOR, CEPHRADINE
AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT

APPENDIX A - PRODUCT NAME INDEX

A - 5

**** A ****

ANTABUSE, DISULFIRAM
ANTARA (MICRONIZED), FENOFIBRATE
ANTIVERT, MECLIZINE HYDROCHLORIDE
ANTIZOL, FOMEPIZOLE
ANTURANE, SULFINPYRAZONE
ANUSOL HC, HYDROCORTISONE
ANZEMET, DOLASETRON MESYLATE MONOHYDRATE
APHTHASOL, AMLEXANOX
APOKYN, APOMORPHINE HYDROCHLORIDE
APRESAZIDE, HYDRALAZINE HYDROCHLORIDE
APRESOLINE, HYDRALAZINE HYDROCHLORIDE
APTIVUS, TIPRANAVIR
AQUAMEPHYTON, PHYTONADIONE
AQUASOL A, VITAMIN A PALMITATE
ARALEN, CHLOROQUINE PHOSPHATE
ARAMINE, METARAMINOL BITARTRATE
ARANELLE, ETHINYL ESTRADIOL
ARAVA, LEFLUNOMIDE
AREDIA, PAMIDRONATE DISODIUM
ARESTIN, MINOCYCLINE HYDROCHLORIDE
ARGATROBAN, ARGATROBAN
ARICEPT ODT, DONEPEZIL HYDROCHLORIDE
ARICEPT, DONEPEZIL HYDROCHLORIDE
ARIMIDEX, ANASTROZOLE
ARISTOCORT A, TRIAMCINOLONE ACETONIDE
ARISTOCORT, TRIAMCINOLONE
ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
ARIXTRA, FONDAPARINUX SODIUM
AROMASIN, EXEMESTANE
ARRANON, NELARABINE
ARTHROTEC, DICLOFENAC SODIUM
ASACOL, MESALAMINE
ASMANEX TWISTHALER, MOMETASONE FUROATE
ASPIRIN AND CAFFEINE W/ BUTALBITAL, ASPIRIN
ASTELIN, AZELASTINE HYDROCHLORIDE
ASTRAMORPH PF, MORPHINE SULFATE
ATACAND HCT, CANDESARTAN CILEXETIL
ATACAND, CANDESARTAN CILEXETIL
ATARAX, HYDROXYZINE HYDROCHLORIDE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
ATIVAN, LORAZEPAM
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
ATRIDOX, DOXYCYCLINE HYCLATE
ATROPEN, ATROPINE
ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE
ATROVENT HFA, IPRATROPIUM BROMIDE
ATROVENT, IPRATROPIUM BROMIDE
AUGMENTIN '125', AMOXICILLIN
AUGMENTIN '200', AMOXICILLIN
AUGMENTIN '250', AMOXICILLIN
AUGMENTIN '400', AMOXICILLIN
AUGMENTIN '500', AMOXICILLIN
AUGMENTIN '875', AMOXICILLIN
AUGMENTIN ES-600, AMOXICILLIN
AUGMENTIN XR, AMOXICILLIN
AVAGARD, ALCOHOL (OTC)
AVAGE, TAZAROTENE
AVALIDE, HYDROCHLOROTHIAZIDE
AVANDAMET, METFORMIN HYDROCHLORIDE
AVANDARYL, GLIMEPIRIDE
AVANDIA, ROSIGLITAZONE MALEATE
AVAPRO, IRBESARTAN

APPENDIX A - PRODUCT NAME INDEX

A - 6

**** A ****

AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN HYDROCHLORIDE
AVELOX, MOXIFLOXACIN HYDROCHLORIDE
AVENTYL HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
AVIANE-28, ETHINYL ESTRADIOL
AVINZA, MORPHINE SULFATE
AVITA, TRETINOIN
AVODART, DUTASTERIDE
AXERT, ALMOTRIPTAN MALATE
AXID AR, NIZATIDINE (OTC)
AXID, NIZATIDINE
AYGESTIN, NORETHINDRONE ACETATE
AZACTAM IN PLASTIC CONTAINER, AZTREONAM
AZACTAM, AZTREONAM
AZASAN, AZATHIOPRINE
AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
AZATHIOPRINE, AZATHIOPRINE
AZDONE, ASPIRIN
AZELEX, AZELAIC ACID
AZITHROMYCIN, AZITHROMYCIN
AZMACORT, TRIAMCINOLONE ACETONIDE
AZOPT, BRINZOLAMIDE
AZULFIDINE EN-TABS, SULFASALAZINE
AZULFIDINE, SULFASALAZINE

**** B ****

BACIIM, BACITRACIN
BACI-RX, BACITRACIN
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
BACITRACIN, BACITRACIN
BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN
BACITRACIN-NEOMYCIN-POLYMYXIN, BACITRACIN ZINC
BACLOFEN, BACLOFEN
BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, WATER FOR INJECTION, STERILE
BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM
BACTOCILL, OXACILLIN SODIUM
BACTRIM DS, SULFAMETHOXAZOLE
BACTRIM, SULFAMETHOXAZOLE
BACTROBAN, MUPIROCIN
BACTROBAN, MUPIROCIN CALCIUM
BAL, DIMERCAPROL
BALZIVA-21, ETHINYL ESTRADIOL
BALZIVA-28, ETHINYL ESTRADIOL
BARACLUDE, ENTECAVIR
BAROS, SODIUM BICARBONATE
BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN, ASPIRIN (OTC)
BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
BENADRYL PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
BENADRYL, DIPHENHYDRAMINE HYDROCHLORIDE
BENZAEPRIIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENZAEPRIIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BENICAR HCT, HYDROCHLOROTHIAZIDE
BENICAR, OLMESARTAN MEDOXOMIL
BENOQUIN, MONOBENZONE
BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
BENTYL, DICYCLOMINE HYDROCHLORIDE
BENZACLIN, BENZOYL PEROXIDE
BENZAMYCIN PAK, BENZOYL PEROXIDE
BENZAMYCIN, BENZOYL PEROXIDE
BENZONATATE, BENZONATATE
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BETA-2, ISOETHARINE HYDROCHLORIDE
BETADINE, POVIDONE-IODINE

APPENDIX A - PRODUCT NAME INDEX

A - 7

**** B ****

BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
BETAPACE AF, SOTALOL HYDROCHLORIDE
BETAPACE, SOTALOL HYDROCHLORIDE
BETA-VAL, BETAMETHASONE VALERATE
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
BETAXOLOL, BETAXOLOL HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
BETIMOL, TIMOLOL
BETOPTIC S, BETAXOLOL HYDROCHLORIDE
BETOPTIC, BETAXOLOL HYDROCHLORIDE
BEXTRA, VALDECOXIB
BIAXIN XL, CLARITHROMYCIN
BIAXIN, CLARITHROMYCIN
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
BICILLIN C-R, PENICILLIN G BENZATHINE
BICILLIN L-A, PENICILLIN G BENZATHINE
BICNU, CARMUSTINE
BIDIL, HYDRALAZINE HYDROCHLORIDE
BILTRICIDE, PRAZIQUANTEL
BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BLENOXANE, BLEOMYCIN SULFATE
BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
BLEOMYCIN, BLEOMYCIN SULFATE
BLEPH-10, SULFACETAMIDE SODIUM
BLEPH-30, SULFACETAMIDE SODIUM
BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
BLEPHAMIDE, PREDNISOLONE ACETATE
BLOCADREN, TIMOLOL MALEATE
BONIVA, IBANDRONATE SODIUM
BONTRIL PDM, PHENDIMETRAZINE TARTRATE
BONTRIL, PHENDIMETRAZINE TARTRATE
BRANCHAMIN 4% IN PLASTIC CONTAINER, AMINO ACIDS
BRAVELLE, UROFOLLITROPIN
BREATHTEK UBT FOR H-PYLORI, UREA, C-13
BRETHINE, TERBUTALINE SULFATE
BRETILIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER, BRETILIUM TOSYLATE
BRETILIUM TOSYLATE IN PLASTIC CONTAINER, BRETILIUM TOSYLATE
BRETILIUM TOSYLATE, BRETILIUM TOSYLATE
BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVIBLOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVIBLOC, ESMOLOL HYDROCHLORIDE
BREVICON 21-DAY, ETHINYL ESTRADIOL
BREVICON 28-DAY, ETHINYL ESTRADIOL
BREVITAL SODIUM, METHOHEXITAL SODIUM
BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
BRISTACYCLINE, TETRACYCLINE HYDROCHLORIDE
BROMFED-DM, BROMPHENIRAMINE MALEATE
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
BROMPHENIRAMINE MALEATE, BROMPHENIRAMINE MALEATE
BRONCHO SALINE, SODIUM CHLORIDE (OTC)
BRONITIN MIST, EPINEPHRINE BITARTRATE (OTC)
BSS PLUS, CALCIUM CHLORIDE
BSS, CALCIUM CHLORIDE
BUCET, ACETAMINOPHEN
BUMETANIDE, BUMETANIDE
BUMEX, BUMETANIDE
BUPHENYL, SODIUM PHENYLBUTYRATE
BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 8

**** B ****

BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
BUPIVACAINE, BUPIVACAINE HYDROCHLORIDE
BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPAR, BUSPIRONE HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
BUSULFEX, BUSULFAN
BUTABARBITAL, BUTABARBITAL SODIUM
BUTAL COMPOUND, ASPIRIN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN, AND CAFFEINE WITH CODEINE PHOSPHATE, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN
BUTALBITAL, APAP, AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
BUTAPAP, ACETAMINOPHEN
BUTISOL SODIUM, BUTABARBITAL SODIUM
BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
BYETTA, EXENATIDE SYNTHETIC

**** C ****

CABERGOLINE, CABERGOLINE
CADUET, AMLODIPINE BESYLATE
CAFCIT, CAFFEINE CITRATE
CAFERGOT, CAFFEINE
CALAN, VERAPAMIL HYDROCHLORIDE
CALCIJEX, CALCITRIOL
CALCITRIOL, CALCITRIOL
CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM
CAMILA, NORETHINDRONE
CAMPRAL, ACAMPROSATE CALCIUM
CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
CANASA, MESALAMINE
CANCIDAS, CASPOFUNGIN ACETATE
CANTIL, MEPENZOLATE BROMIDE
CAPASTAT SULFATE, CAPREOMYCIN SULFATE
CAPITAL AND CODEINE, ACETAMINOPHEN
CAPITROL, CHLOROXINE
CAPOTEN, CAPTOPRIL
CAPOZIDE 25/15, CAPTOPRIL
CAPOZIDE 25/25, CAPTOPRIL
CAPOZIDE 50/15, CAPTOPRIL
CAPOZIDE 50/25, CAPTOPRIL
CAP-PROFEN, IBUPROFEN (OTC)
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
CAPTOPRIL, CAPTOPRIL
CARAC, FLUOROURACIL
CARAFATE, SUCRALFATE
CARBAMAZEPINE, CARBAMAZEPINE
CARBASTAT, CARBACHOL
CARBATROL, CARBAMAZEPINE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARBILEV, CARBIDOPA
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
CARBOCAINE W/ NEO-COBEFRIN, LEVONORDEFIN
CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
CARBOPLATIN, CARBOPLATIN
CARDENE SR, NICARDIPINE HYDROCHLORIDE
CARDENE, NICARDIPINE HYDROCHLORIDE
CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82

APPENDIX A - PRODUCT NAME INDEX

A - 9

**** C ****

CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
CARDIZEM, DILTIAZEM HYDROCHLORIDE
CARDURA XL, DOXAZOSIN MESYLATE
CARDURA, DOXAZOSIN MESYLATE
CARISOPRODOL AND ASPIRIN, ASPIRIN
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
CARISOPRODOL, CARISOPRODOL
CARMOL HC, HYDROCORTISONE ACETATE
CARNITOR, LEVOCARNITINE
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
CARTIA XT, DILTIAZEM HYDROCHLORIDE
CASODEX, BICALUTAMIDE
CATAFLAM, DICLOFENAC POTASSIUM
CATAPRES, CLONIDINE HYDROCHLORIDE
CATAPRES-TTS-1, CLONIDINE
CATAPRES-TTS-2, CLONIDINE
CATAPRES-TTS-3, CLONIDINE
CAVERJECT, ALPROSTADIL
CEDAX, CEFTIBUTEN DIHYDRATE
CEENU, LOMUSTINE
CEFACLOL, CEFACLOL
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFAZOLIN AND DEXTROSE, CEFAZOLIN SODIUM
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFAZOLIN, CEFAZOLIN SODIUM
CEFIZOX IN PLASTIC CONTAINER, CEFTIZOXIME SODIUM
CEFIZOX, CEFTIZOXIME SODIUM
CEFOBID IN PLASTIC CONTAINER, CEFOPERAZONE SODIUM
CEFOBID, CEFOPERAZONE SODIUM
CEFOTAN IN PLASTIC CONTAINER, CEFOTETAN DISODIUM
CEFOTAN, CEFOTETAN DISODIUM
CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER, CEFOTAXIME SODIUM
CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER, CEFOTAXIME SODIUM
CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
CEFOTAXIME, CEFOTAXIME SODIUM
CEFOXITIN, CEFOXITIN SODIUM
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFPROZIL, CEFPROZIL
CEFTAZIDIME SODIUM IN PLASTIC CONTAINER, CEFTAZIDIME SODIUM
CEFTAZIDIME, CEFTAZIDIME
CEFTIN, CEFUROXIME AXETIL
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
CEFTRIAXONE, CEFTRIAXONE SODIUM
CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEFUROXIME SODIUM, CEFUROXIME SODIUM
CEFUROXIME, CEFUROXIME SODIUM
CEFZIL, CEFPROZIL
CELEBREX, CELECOXIB
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CELESTONE, BETAMETHASONE
CELEXA, CITALOPRAM HYDROBROMIDE
CELLCEPT, MYCOPHENOLATE MOFETIL
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
CELONTIN, METHSUXIMIDE
CENESTIN, ESTROGENS, CONJUGATED SYNTHETIC A
CENTANY, MUPIROCIN
CEPHALEXIN, CEPHALEXIN
CEPHRADINE, CEPHRADINE
CEPTAZ, CEFTAZIDIME (ARGININE FORMULATION)

APPENDIX A - PRODUCT NAME INDEX

A - 10

**** C ****

CEREBYX, FOSPHENYTOIN SODIUM
CEREDASE, ALGLUCERASE
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
CEREZYME, IMIGLUCERASE
CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
CERVIDIL, DINOPROSTONE
CETACORT, HYDROCORTISONE
CETAMIDE, SULFACETAMIDE SODIUM
CETROTIDE, CETRORELIX
CHEMET, SUCCIMER
CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)
CHILDREN'S ADVIL, IBUPROFEN
CHILDREN'S ADVIL, IBUPROFEN (OTC)
CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)
CHILDREN'S ELIXSURE, IBUPROFEN (OTC)
CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
CHILDREN'S MOTRIN, IBUPROFEN (OTC)
CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN
CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE
CHLORAMPHENICOL, CHLORAMPHENICOL
CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB SWAB, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)
CHLOROMYCETIN, CHLORAMPHENICOL SODIUM SUCCINATE
CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
CHLOROTHIAZIDE, CHLOROTHIAZIDE
CHLORPROMAZINE HYDROCHLORIDE INTENSOL, CHLORPROMAZINE HYDROCHLORIDE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CHLORPROPAMIDE, CHLORPROPAMIDE
CHLORTHALIDONE, CHLORTHALIDONE
CHLOR-TRIMETON, CHLORPHENIRAMINE MALEATE (OTC)
CHLORZOXAZONE, CHLORZOXAZONE
CHOLAC, LACTULOSE
CHOLEDYL SA, OXTRIPHYLLINE
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
CHOLESTYRAMINE, CHOLESTYRAMINE
CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
CHOLOGRAFIN MEGLUMINE, IODIPAMIDE MEGLUMINE
CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE
CHROMITOPES SODIUM, SODIUM CHROMATE, CR-51
CIALIS, TADALAFIL
CICLOPIROX, CICLOPIROX
CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)
CILOSTAZOL, CILOSTAZOL
CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, CIMETIDINE HYDROCHLORIDE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CIMETIDINE, CIMETIDINE
CIMETIDINE, CIMETIDINE (OTC)
CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 11

**** C ****

CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
CIPRO XR, CIPROFLOXACIN
CIPRO, CIPROFLOXACIN
CIPRO, CIPROFLOXACIN HYDROCHLORIDE
CIPRODEX, CIPROFLOXACIN
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
CISPLATIN, CISPLATIN
CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CITANEST FORTE, EPINEPHRINE BITARTRATE
CITANEST PLAIN, PRILOCAINE HYDROCHLORIDE
CLADRIBINE, CLADRIBINE
CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER, CEFOTAXIME SODIUM
CLAFORAN, CEFOTAXIME SODIUM
CLARAVIS, ISOTRETINOIN
CLARINEX D 24 HOUR, DESLORATADINE
CLARINEX, DESLORATADINE
CLARITHROMYCIN EXTENDED RELEASE, CLARITHROMYCIN
CLARITHROMYCIN, CLARITHROMYCIN
CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)
CLARITIN HIVES RELIEF, LORATADINE (OTC)
CLARITIN REDITABS, LORATADINE (OTC)
CLARITIN, LORATADINE (OTC)
CLARITIN-D 24 HOUR, LORATADINE (OTC)
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLEOCIN T, CLINDAMYCIN PHOSPHATE
CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
CLEOCIN, CLINDAMYCIN PHOSPHATE
CLIMARA PRO, ESTRADIOL
CLIMARA, ESTRADIOL
CLINDA-DERM, CLINDAMYCIN PHOSPHATE
CLINDAGEL, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLINDESSE, CLINDAMYCIN PHOSPHATE
CLINDETS, CLINDAMYCIN PHOSPHATE
CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 2.75/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 2.75/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 2.75/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 4.25/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 4.25/20 SULFITE-FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS

APPENDIX A - PRODUCT NAME INDEX

A - 12

**** C ****

CLINIMIX E 4.25/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 4.25/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/15 SULFITE-FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/20 SULFITE-FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINIMIX E 5/35 SULFITE-FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
CLINORIL, SULINDAC
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOBEX, CLOBETASOL PROPIONATE
CLODERM, CLOCORTOLONE PIVALATE
CLOFIBRATE, CLOFIBRATE
CLOLAR, CLOFARABINE
CLOMID, CLOMIPHENE CITRATE
CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
CLORPRES, CHLORTHALIDONE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOTRIMAZOLE, CLOTRIMAZOLE
CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
CLOXACILLIN SODIUM, CLOXACILLIN SODIUM
CLOXAPEN, CLOXACILLIN SODIUM
CLOZAPINE, CLOZAPINE
CLOZARIL, CLOZAPINE
CODEPREX, CHLORPHENIRAMINE POLISTIREX
CODRIX, ACETAMINOPHEN
COGENTIN, BENZTROPINE MESYLATE
CO-GESIC, ACETAMINOPHEN
COGNEX, TACRINE HYDROCHLORIDE
CO-LAV, POLYETHYLENE GLYCOL 3350
COLAZAL, BALSALAZIDE DISODIUM
COLESTID, COLESTIPOL HYDROCHLORIDE
COLGATE TOTAL, SODIUM FLUORIDE (OTC)
COLISTIMETHATE, COLISTIMETHATE SODIUM
COLOCORT, HYDROCORTISONE
COL-PROBENECID, COLCHICINE
COLY-MYCIN M, COLISTIMETHATE SODIUM
COLY-MYCIN S, COLISTIN SULFATE
COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
COLYTE, POLYETHYLENE GLYCOL 3350
COLYTE-FLAVORED, POLYETHYLENE GLYCOL 3350
COMBIPATCH, ESTRADIOL
COMBIVENT, ALBUTEROL SULFATE
COMBIVIR, LAMIVUDINE
COMBUNOX, IBUPROFEN
COMMIT, NICOTINE POLACRILEX (OTC)
COMPAZINE, PROCHLORPERAZINE
COMPAZINE, PROCHLORPERAZINE EDISYLATE
COMPAZINE, PROCHLORPERAZINE MALEATE
COMPRO, PROCHLORPERAZINE
COMTAN, ENTACAPONE
CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
CONDYLOX, PODOFILOX

APPENDIX A - PRODUCT NAME INDEX

A - 13

**** C ****

CONRAY 30, IOTHALAMATE MEGLUMINE
CONRAY 400, IOTHALAMATE SODIUM
CONRAY 43, IOTHALAMATE MEGLUMINE
CONRAY, IOTHALAMATE MEGLUMINE
CONSTILAC, LACTULOSE
CONSTULOSE, LACTULOSE
COPAXONE, GLATIRAMER ACETATE
COPEGUS, RIBAVIRIN
COPPER T MODEL TCU 380A, COPPER
CORDARONE, AMIODARONE HYDROCHLORIDE
CORDRAN SP, FLURANDRENOLIDE
CORDRAN, FLURANDRENOLIDE
COREG, CARVEDILOL
CORGARD, NADOLOL
CORLOPAM, FENOLDOPAM MESYLATE
CORMAX, CLOBETASOL PROPIONATE
CORPHED, PSEUDOEPHEDRINE HYDROCHLORIDE
CORTEF, HYDROCORTISONE
CORTENEMA, HYDROCORTISONE
CORTIFOAM, HYDROCORTISONE ACETATE
CORTISONE ACETATE, CORTISONE ACETATE
CORTISPORIN, BACITRACIN ZINC
CORTISPORIN, HYDROCORTISONE
CORTISPORIN, HYDROCORTISONE ACETATE
CORTROSYN, COSYNTROPIN
CORVERT, IBUTILIDE FUMARATE
CORZIDE, BENDROFLUMETHIAZIDE
COSMEGEN, DACTINOMYCIN
COSOPT, DORZOLAMIDE HYDROCHLORIDE
COTRIM D.S., SULFAMETHOXAZOLE
COTRIM, SULFAMETHOXAZOLE
COUMADIN, WARFARIN SODIUM
COVERA-HS, VERAPAMIL HYDROCHLORIDE
COZAAR, LOSARTAN POTASSIUM
CRESTOR, ROSUVASTATIN CALCIUM
CRINONE, PROGESTERONE
CRIXIVAN, INDINAVIR SULFATE
CROLOM, CROMOLYN SODIUM
CROMOLYN SODIUM, CROMOLYN SODIUM
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
CROTAN, CROTAMITON
CRYSELLE, ETHINYL ESTRADIOL
C-SOLVE-2, ERYTHROMYCIN
CUBICIN, DAPTOMYCIN
CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE
CUPRIMINE, PENICILLAMINE
CUROSURF, PORACTANT ALFA
CUTIVATE, FLUTICASONE PROPIONATE
CYANOCOBALAMIN, CYANOCOBALAMIN
CYCLESSA, DESOGESTREL
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
CYCLOCORT, AMCINONIDE
CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
CYCLOSPORINE, CYCLOSPORINE
CYKLOKAPRON, TRANEXAMIC ACID
CYMBALTA, DULOXETINE HYDROCHLORIDE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
CYSTADANE, BETAINE, ANHYDROUS
CYSTAGON, CYSTEAMINE BITARTRATE
CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE

APPENDIX A - PRODUCT NAME INDEX

A - 14

**** C ****

CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
CYTADREN, AMINOGLUTETHIMIDE
CYTARABINE, CYTARABINE
CYTOMEL, LIOTHYRONINE SODIUM
CYTOSAR-U, CYTARABINE
CYTOTEC, MISOPROSTOL
CYTOVENE IV, GANCICLOVIR SODIUM
CYTOVENE, GANCICLOVIR
CYTOXAN, CYCLOPHOSPHAMIDE

**** D ****

D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
DACARBAZINE, DACARBAZINE
DALMANE, FLURAZEPAM HYDROCHLORIDE
DANAZOL, DANAZOL
DANTRIUM, DANTROLENE SODIUM
DANTROLENE SODIUM, DANTROLENE SODIUM
DAPSONE, DAPSONE
DARAPRIM, PYRIMETHAMINE
DARVOCET A500, ACETAMINOPHEN
DARVOCET-N 100, ACETAMINOPHEN
DARVOCET-N 50, ACETAMINOPHEN
DARVON COMPOUND-65, ASPIRIN
DARVON, PROPOXYPHENE HYDROCHLORIDE
DARVON-N, PROPOXYPHENE NAPSYLATE
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DAUNOXOME, DAUNORUBICIN CITRATE
DAYPRO, OXAPROZIN
DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
DDAVP, DESMOPRESSIN ACETATE
DECADRON, DEXAMETHASONE
DECLOMYCIN, DEMECLOCYCLINE HYDROCHLORIDE
DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE
DEFINITY, PERFLUTREN
DELATESTRYL, TESTOSTERONE ENANTHATE
DELESTROGEN, ESTRADIOL VALERATE
Delflex W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
DEMADEX, TORSEMIDE
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
DEMEROL, MEPERIDINE HYDROCHLORIDE
DEMSE, METYROSINE
DEMULEN 1/35-21, ETHINYL ESTRADIOL
DEMULEN 1/35-28, ETHINYL ESTRADIOL
DEMULEN 1/50-21, ETHINYL ESTRADIOL
DEMULEN 1/50-28, ETHINYL ESTRADIOL
DENAVID, PENCICLOVIR SODIUM
DENDRID, IDOXURIDINE
DEPAON, VALPROATE SODIUM
DEPAKENE, VALPROIC ACID

APPENDIX A - PRODUCT NAME INDEX

A - 15

**** D ****

DEPAKOTE ER, DIVALPROEX SODIUM
DEPAKOTE, DIVALPROEX SODIUM
DEPEN, PENICILLAMINE
DEPOCYT, CYTARABINE
DEPODUR, MORPHINE SULFATE
DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
DERMABET, BETAMETHASONE VALERATE
DERMA-SMOOTHIE/FS, FLUOCINOLONE ACETONIDE
DERMATOP E EMOLLIENT, PREDNICARBATE
DERMATOP, PREDNICARBATE
DESFERAL, DEFEROXAMINE MESYLATE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
DESMOPRESSIN ACETATE PRESERVATIVE FREE, DESMOPRESSIN ACETATE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DESOGEN, DESOGESTREL
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DESONIDE, DESONIDE
DESOWEN, DESONIDE
DESOXIMETASONE, DESOXIMETASONE
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
DESYREL, TRAZODONE HYDROCHLORIDE
DETROL LA, TOLTERODINE TARTRATE
DETROL, TOLTERODINE TARTRATE
DEXAMETHASONE INTENSOL, DEXAMETHASONE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DEXAMETHASONE, DEXAMETHASONE
DEXAMETHASONE, DEXAMETHASONE SODIUM PHOSPHATE
DEXASPORIN, DEXAMETHASONE
DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
DEXEDRINE, DEXTROAMPHETAMINE SULFATE
DEXFERRUM, IRON DEXTRAN
DEXRAZOXANE, DEXRAZOXANE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE ASPARTATE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 25%, DEXTROSE
DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND ELECTROLYTE NO 75 IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND ELECTROLYTE NO.48 IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 16

**** D ****

DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN ACETATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 60% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSTAT, DEXTROAMPHETAMINE SULFATE
DIABETA, GLYBURIDE
DIABINESE, CHLORPROPAMIDE
DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIAMOX, ACETAZOLAMIDE
DIAMOX, ACETAZOLAMIDE SODIUM
DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
DIASTAT ACUDIAL, DIAZEPAM
DIASTAT, DIAZEPAM
DIAZEPAM INTENSOL, DIAZEPAM
DIAZEPAM, DIAZEPAM
DIBENZYLINE, PHENOXYBENZAMINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 17

**** D ****

DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE-FREE), DICYCLOMINE HYDROCHLORIDE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DIDANOSINE, DIDANOSINE
DIDREX, BENZPHETAMINE HYDROCHLORIDE
DIDRONEL, ETIDRONATE DISODIUM
DIFFERIN, ADAPALENE
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
DIFLUCAN IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
DIFLUCAN, FLUCONAZOLE
DIFLUNISAL, DIFLUNISAL
DIGOXIN PEDIATRIC, DIGOXIN
DIGOXIN, DIGOXIN
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
DILACOR XR, DILTIAZEM HYDROCHLORIDE
DILANTIN, PHENYTOIN
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
DILATRATE-SR, ISOSORBIDE DINITRATE
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
DILAUDID-HP, HYDROMORPHONE HYDROCHLORIDE
DILOR, DYPHYLLINE
DILOR-400, DYPHYLLINE
DILT-CD, DILTIAZEM HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIMENHYDRINATE, DIMENHYDRINATE
DIMETHYL SULFOXIDE, DIMETHYL SULFOXIDE
DIOVAN HCT, HYDROCHLOROTHIAZIDE
DIOVAN, VALSARTAN
DIPENTUM, OLSALAZINE SODIUM
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE, ATROPINE SULFATE
DIPIVEFRIN HYDROCHLORIDE, DIPIVEFRIN HYDROCHLORIDE
DIPRIVAN, PROPOFOL
DIPROLENE AF, BETAMETHASONE DIPROPIONATE
DIPROLENE, BETAMETHASONE DIPROPIONATE
DIPYRIDAMOLE, DIPYRIDAMOLE
DISOPHROL, DEXBROMPHENIRAMINE MALEATE (OTC)
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
DISPERMOX, AMOXICILLIN
DITROPAN XL, OXYBUTYNIN CHLORIDE
DITROPAN, OXYBUTYNIN CHLORIDE
DIUPRES-250, CHLOROTHIAZIDE
DIUPRES-500, CHLOROTHIAZIDE
DIURIL, CHLOROTHIAZIDE
DIURIL, CHLOROTHIAZIDE SODIUM
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOLOBID, DIFLUNISAL
DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
DOPRAM, DOXAPRAM HYDROCHLORIDE
DORAL, QUAZEPAM
DORYX, DOXYCYCLINE HYCLATE

APPENDIX A - PRODUCT NAME INDEX

A - 18

**** D ****

DOSTINEX, CABERGOLINE
DOVONEX, CALCIPOTRIENE
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXIL, DOXORUBICIN HYDROCHLORIDE
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
DOXY 100, DOXYCYCLINE HYCLATE
DOXY 200, DOXYCYCLINE HYCLATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
DRAXIMAGE MDP, TECHNETIUM TC-99M MEDRONATE KIT
DRICORT, HYDROCORTISONE ACETATE
DRISDOL, ERGOCALCIFEROL
DRIXORAL PLUS, ACETAMINOPHEN (OTC)
DRIXORAL, DEXBROMPHENIRAMINE MALEATE (OTC)
DROPERIDOL, DROPERIDOL
DROXIA, HYDROXYUREA
DTIC-DOME, DACARBAZINE
DTPA, TECHNETIUM TC-99M PENTETATE KIT
DUAC, BENZOYL PEROXIDE
DUOCAINE, BUPIVACAINE HYDROCHLORIDE
DUONEB, ALBUTEROL SULFATE
DURACLON, CLONIDINE HYDROCHLORIDE
DURAGESIC-100, FENTANYL
DURAGESIC-12, FENTANYL
DURAGESIC-25, FENTANYL
DURAGESIC-50, FENTANYL
DURAGESIC-75, FENTANYL
DURAMORPH PF, MORPHINE SULFATE
DURICEF, CEFADROXIL/CEFADROXIL HEMIHYDRATE
DUVOID, BETHANECHOL CHLORIDE
DYAZIDE, HYDROCHLOROTHIAZIDE
DYNACIN, MINOCYCLINE HYDROCHLORIDE
DYNACIRC CR, ISRADIPINE
DYNACIRC, ISRADIPINE
DYNA-HEX, CHLORHEXIDINE GLUCONATE (OTC)
DYRENIUM, TRIAMTERENE

**** E ****

E.E.S. 200, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
E-BASE, ERYTHROMYCIN
EC-NAPROSYN, NAPROXEN
ECONAZOLE NITRATE, ECONAZOLE NITRATE
ECONOPRED PLUS, PREDNISOLONE ACETATE
EDECRIN, ETHACRYNATE SODIUM
EDECRIN, ETHACRYNIC ACID
EDETATE DISODIUM, EDETATE DISODIUM
EDEX, ALPROSTADIL
EDROPHONIUM CHLORIDE, EDROPHONIUM CHLORIDE
EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
EFFEXOR, VENLAFAXINE HYDROCHLORIDE
EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE, BROMPHENIRAMINE MALEATE (OTC)
EFUDEX, FLUOROURACIL
E-GLADES, ERYTHROMYCIN
ELDEPRYL, SELEGILINE HYDROCHLORIDE
ELESTAT, EPINASTINE HYDROCHLORIDE
ELIDEL, PIMECROLIMUS
ELIGARD, LEUPROLIDE ACETATE

APPENDIX A - PRODUCT NAME INDEX

A - 19

**** E ****

ELIMITE, PERMETHRIN
ELIXOPHYLLIN, THEOPHYLLINE
ELLEENCE, EPIRUBICIN HYDROCHLORIDE
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE
ELMIRON, PENTOSAN POLYSULFATE SODIUM
ELOCON, MOMETASONE FUROATE
ELOXATIN, OXALIPLATIN
EMADINE, EMEDASTINE DIFUMARATE
EMBELINE E, CLOBETASOL PROPIONATE
EMBELINE, CLOBETASOL PROPIONATE
EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
EMEND, APREPITANT
EMETE-CON, BENZQUINAMIDE HYDROCHLORIDE
EMLA, LIDOCAINE
EMTRIVA, EMTRICITABINE
E-MYCIN, ERYTHROMYCIN
ENABLEX, DARIFENACIN HYDROBROMIDE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ENALAPRILAT, ENALAPRILAT
ENDOSOL EXTRA, CALCIUM CHLORIDE
ENDRATE, EDETATE DISODIUM
ENDURON, METHYLCLOTHIAZIDE
ENFLURANE, ENFLURANE
ENLON, EDROPHONIUM CHLORIDE
ENLON-PLUS, ATROPINE SULFATE
ENPRESSE-28, ETHINYL ESTRADIOL
ENTOCORT EC, BUDESONIDE
ENULOSE, LACTULOSE
EPIFOAM, HYDROCORTISONE ACETATE
EPINEPHRINE, EPINEPHRINE (OTC)
EPIPEN JR., EPINEPHRINE
EPIPEN, EPINEPHRINE
EPITOL, CARBAMAZEPINE
EPIVIR, LAMIVUDINE
EPIVIR-HBV, LAMIVUDINE
EPZICOM, ABACAVIR SULFATE
EQUAGESIC, ASPIRIN
EQUETRO, CARBAMAZEPINE
ERGOLOID MESYLATES, ERGOLOID MESYLATES
ERGOMAR, ERGOTAMINE TARTRATE
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ERRIN, NORETHINDRONE
ERTACZO, SERTACONAZOLE NITRATE
ERYC, ERYTHROMYCIN
ERYCETTE, ERYTHROMYCIN
ERYDERM, ERYTHROMYCIN
ERYGEL, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERY-TAB, ERYTHROMYCIN
ERYTHRA-DERM, ERYTHROMYCIN
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
ERYTHROMYCIN ESTOLATE, ERYTHROMYCIN ESTOLATE
ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
ERYTHROMYCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN, ERYTHROMYCIN
ERYZOLE, ERYTHROMYCIN ETHYLSUCCINATE
ESGIC-PLUS, ACETAMINOPHEN
ESIDRIX, HYDROCHLOROTHIAZIDE
ESKALITH CR, LITHIUM CARBONATE

APPENDIX A - PRODUCT NAME INDEX

A - 20

**** E ****

ESKALITH, LITHIUM CARBONATE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
E-SOLVE 2, ERYTHROMYCIN
ESTAZOLAM, ESTAZOLAM
ESTRACE, ESTRADIOL
ESTRADERM, ESTRADIOL
ESTRADIOL AND NORGESTIMATE, ESTRADIOL
ESTRADIOL CYPIONATE, ESTRADIOL CYPIONATE
ESTRADIOL VALERATE, ESTRADIOL VALERATE
ESTRADIOL, ESTRADIOL
ESTRASORB, ESTRADIOL HEMIHYDRATE
ESTRING, ESTRADIOL
ESTROGEL, ESTRADIOL
ESTRONE, ESTRONE
ESTROPIPATE, ESTROPIPATE
ESTROSTEP FE, ETHINYL ESTRADIOL
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
ETHAMOLIN, ETHANOLAMINE OLEATE
ETHIODOL, ETHIODIZED OIL
ETHMOZINE, MORICIZINE HYDROCHLORIDE
ETHOSUXIMIDE, ETHOSUXIMIDE
ETHRANE, ENFLURANE
ETHYOL, AMIFOSTINE
ETIDRONATE DISODIUM, ETIDRONATE DISODIUM
ETODOLAC, ETODOLAC
ETOMIDATE, ETOMIDATE
ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
ETOPOSIDE, ETOPOSIDE
EULEXIN, FLUTAMIDE
EURAX, CROTAMITON
EVALOSE, LACTULOSE
EVISTA, RALOXIFENE HYDROCHLORIDE
EVOCLIN, CLINDAMYCIN PHOSPHATE
EVOXAC, CEVIMELINE HYDROCHLORIDE
EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
EXELDERM, SULCONAZOLE NITRATE
EXELON, RIVASTIGMINE TARTRATE
EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)
EXJADE, DEFERASIROX
EXOSURF NEONATAL, CETYL ALCOHOL
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
EXTRANEAL, ICODEXTRIN
EXTRA-STRENGTH AIM, SODIUM MONOFLUOROPHOSPHATE (OTC)
E-Z PREP 220, POVIDONE-IODINE (OTC)
E-Z PREP, POVIDONE-IODINE (OTC)
E-Z SCRUB 201, POVIDONE-IODINE (OTC)
E-Z SCRUB 241, POVIDONE-IODINE (OTC)

**** F ****

FACTIVE, GEMIFLOXACIN MESYLATE
FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK), FAMOTIDINE
FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE (OTC)
FAMVIR, FAMCICLOVIR
FANSIDAR, PYRIMETHAMINE
FARESTON, TOREMIFENE CITRATE
FASLODEX, FULVESTRANT
FAZACLO ODT, CLOZAPINE
FELBATOL, FELBAMATE
FELDENE, PIROXICAM
FELODIPINE, FELODIPINE

APPENDIX A - PRODUCT NAME INDEX

A - 21

**** F ****

FEMARA, LETROZOLE
FEMHRT, ETHINYL ESTRADIOL
FEMRING, ESTRADIOL ACETATE
FEMSTAT 3, BUTOCONAZOLE NITRATE (OTC)
FEMTRACE, ESTRADIOL ACETATE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FENOFIBRATE, FENOFIBRATE
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
FENTANYL CITRATE, FENTANYL CITRATE
FENTANYL, FENTANYL
FERIDEX I.V., FERUMOXIDES
FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FINACEA, AZELAIC ACID
FIORICET W/ CODEINE, ACETAMINOPHEN
FIORICET, ACETAMINOPHEN
FIORINAL W/CODEINE NO 3, ASPIRIN
FIORINAL, ASPIRIN
FLAGYL ER, METRONIDAZOLE
FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
FLAGYL I.V., METRONIDAZOLE HYDROCHLORIDE
FLAGYL, METRONIDAZOLE
FLAREX, FLUOROMETHOLONE ACETATE
FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLEXERIL, CYCLOBENZAPRINE HYDROCHLORIDE
FLOLAN, EPOPROSTENOL SODIUM
FLOMAX, TAMSULOSIN HYDROCHLORIDE
FLONASE, FLUTICASONE PROPIONATE
FLORINEF, FLUDROCORTISONE ACETATE
FLORONE E, DIFLORASONE DIACETATE
FLORONE, DIFLORASONE DIACETATE
FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
FLOVENT DISKUS 50, FLUTICASONE PROPIONATE
FLOVENT HFA, FLUTICASONE PROPIONATE
FLOVENT, FLUTICASONE PROPIONATE
FLOXIN OTIC, OFLOXACIN
FLOXIN, OFLOXACIN
FLOXURIDINE, FLOXURIDINE
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
FLUCONAZOLE, FLUCONAZOLE
FLUDARA, FLUDARABINE PHOSPHATE
FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
FLUDEOXYGLUCOSE F 18, FLUDEOXYGLUCOSE F-18
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
FLUMADINE, RIMANTADINE HYDROCHLORIDE
FLUMAZENIL, FLUMAZENIL
FLUNISOLIDE, FLUNISOLIDE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUOR-OP, FLUOROMETHOLONE
FLUOROPLEX, FLUOROURACIL
FLUOROURACIL, FLUOROURACIL
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 22

**** F ****

FLUOXYMESTERONE, FLUOXYMESTERONE
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM
FLURBIPROFEN, FLURBIPROFEN
FLUTAMIDE, FLUTAMIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
FLUXID, FAMOTIDINE
FML FORTE, FLUOROMETHOLONE
FML, FLUOROMETHOLONE
FML-S, FLUOROMETHOLONE
FOAMCOAT, ALUMINUM HYDROXIDE (OTC)
FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
FOLICET, FOLIC ACID
FOLLISTIM AQ, FOLLITROPIN ALFA/BETA
FORADIL, FORMOTEROL FUMARATE
FORANE, ISOFLURANE
FORTAMET, METFORMIN HYDROCHLORIDE
FORTAZ IN PLASTIC CONTAINER, CEFTAZIDIME SODIUM
FORTAZ, CEFTAZIDIME
FORTEO, TERIPARATIDE RECOMBINANT HUMAN
FORTICAL, CALCITONIN SALMON RECOMBINANT
FORTOVASE, SAQUINAVIR
FOSAMAX PLUS D, ALENDRONATE SODIUM
FOSAMAX, ALENDRONATE SODIUM
FOSCARNET SODIUM, FOSCARNET SODIUM
FOSCAVIR, FOSCARNET SODIUM
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
FOSRENOL, LANTHANUM CARBONATE
FRAGMIN, DALTEPARIN SODIUM
FREAMINE HBC 6.9%, AMINO ACIDS
FREAMINE III 10%, AMINO ACIDS
FREAMINE III 3% W/ ELECTROLYTES, AMINO ACIDS
FREAMINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS
FREAMINE III 8.5%, AMINO ACIDS
FROVA, FROVATRIPTAN SUCCINATE
FS SHAMPOO, FLUOCINOLONE ACETONIDE
FUDR, FLOXURIDINE
FULVICIN-U/F, GRISEOFULVIN, MICROCRYSTALLINE
FUNGIZONE, AMPHOTERICIN B
FURADANTIN, NITROFURANTOIN
FUROSEMIDE, FUROSEMIDE
FUZEON, ENFUVIRTIDE

**** G ****

GABAPENTIN, GABAPENTIN
GABITRIL, TIAGABINE HYDROCHLORIDE
GALLIUM CITRATE GA 67, GALLIUM CITRATE, GA-67
GALZIN, ZINC ACETATE
GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
GANCICLOVIR, GANCICLOVIR
GANIRELIX ACETATE INJECTION, GANIRELIX ACETATE
GANITE, GALLIUM NITRATE
GANTRISIN PEDIATRIC, SULFISOXAZOLE ACETYL
GASTROCROM, CROMOLYN SODIUM
GASTROGRAFIN, DIATRIZOATE MEGLUMINE
GASTROMARK, FERUMOXASIL
GAVISCON, ALUMINUM HYDROXIDE (OTC)

APPENDIX A - PRODUCT NAME INDEX

A - 23

**** G ****

GEMFIBROZIL, GEMFIBROZIL
 GEMZAR, GEMCITABINE HYDROCHLORIDE
 GENCEPT 10/11-21, ETHINYL ESTRADIOL
 GENCEPT 10/11-28, ETHINYL ESTRADIOL
 GENERLAC, LACTULOSE
 GENGRAF, CYCLOSPORINE
 GENOPTIC, GENTAMICIN SULFATE
 GENOTROPIN PRESERVATIVE FREE, SOMATROPIN RECOMBINANT
 GENOTROPIN, SOMATROPIN RECOMBINANT
 GENTACIDIN, GENTAMICIN SULFATE
 GENTAK, GENTAMICIN SULFATE
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 GENTAMICIN, GENTAMICIN SULFATE
 GEN-XENE, CLORAZEPATE DIPOTASSIUM
 GEOCILLIN, CARBENICILLIN INDANYL SODIUM
 GEODON, ZIPRASIDONE HYDROCHLORIDE
 GEODON, ZIPRASIDONE MESYLATE
 GEREFF, SERMORELIN ACETATE
 GERIMAL, ERGOLOID MESYLATES
 GLEEVEC, IMATINIB MESYLATE
 GLIADEL, CARMUSTINE
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
 GLIPIZIDE, GLIPIZIDE
 GLOFIL-125, IOTHALAMATE SODIUM, I-125
 GLUCAGEN, GLUCAGON HYDROCHLORIDE RECOMBINANT
 GLUCAGON, GLUCAGON RECOMBINANT
 GLUCOPHAGE XR, METFORMIN HYDROCHLORIDE
 GLUCOPHAGE, METFORMIN HYDROCHLORIDE
 GLUCOTROL XL, GLIPIZIDE
 GLUCOTROL, GLIPIZIDE
 GLUCOVANCE, GLYBURIDE
 GLUMETZA, METFORMIN HYDROCHLORIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
 GLYCOLAX, POLYETHYLENE GLYCOL 3350
 GLYCOPYRROLATE, GLYCOPYRROLATE
 GLYNASE, GLYBURIDE
 GLYSET, MIGLITOL
 GO-EVAC, POLYETHYLENE GLYCOL 3350
 GOLYTELY, POLYETHYLENE GLYCOL 3350
 GONAL-F RFF PEN, FOLLITROPIN ALFA/BETA
 GONAL-F RFF, FOLLITROPIN ALFA/BETA
 GONAL-F, FOLLITROPIN ALFA/BETA
 GRIFULVIN V, GRISEOFULVIN, MICROCRYSTALLINE
 GRISEOFULVIN, GRISEOFULVIN, MICROCRYSTALLINE
 GRIS-PEG, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
 GUANABENZ ACETATE, GUANABENZ ACETATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE
 GYNAZOLE-1, BUTOCONAZOLE NITRATE
 GYNE-LOTRIMIN 3 COMBINATION PACK, CLOTRIMAZOLE (OTC)
 GYNE-LOTRIMIN 3, CLOTRIMAZOLE (OTC)
 GYNE-LOTRIMIN COMBINATION PACK, CLOTRIMAZOLE (OTC)
 GYNE-LOTRIMIN, CLOTRIMAZOLE (OTC)
 GYNIX, CLOTRIMAZOLE (OTC)
 GYNODIOL, ESTRADIOL

**** H ****

H.P. ACTHAR GEL, CORTICOTROPIN

APPENDIX A - PRODUCT NAME INDEX

A - 24

**** H ****

HABITROL, NICOTINE (OTC)
HALCION, TRIAZOLAM
HALDOL, HALOPERIDOL DECANOATE
HALDOL, HALOPERIDOL LACTATE
HALFLYTELY, BISACODYL
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HALOG, HALCINONIDE
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL LACTATE, HALOPERIDOL LACTATE
HALOPERIDOL, HALOPERIDOL
HALOPERIDOL, HALOPERIDOL LACTATE
HALOTESTIN, FLUOXYMESTERONE
HALOTHANE, HALOTHANE
HECTOROL, DOXERCALCIFEROL
HELIDAC, BISMUTH SUBSALICYLATE
HEMABATE, CARBOPROST TROMETHAMINE
HEPARIN LOCK FLUSH IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN LOCK FLUSH, HEPARIN SODIUM
HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM
HEPATAMINE 8%, AMINO ACIDS
HEPATASOL 8%, AMINO ACIDS
HEPATOLITE, TECHNETIUM TC-99M DISOFENIN KIT
HEPFLUSH-10, HEPARIN SODIUM
HEP-LOCK U/P, HEPARIN SODIUM
HEP-LOCK, HEPARIN SODIUM
HEPSERA, ADEFOVIR DIPIVOXIL
HEPTALAC, LACTULOSE
HERPLEX, IDOXURIDINE
HEXABRIX, IOXAGLATE MEGLUMINE
HEXADROL, DEXAMETHASONE
HEXALEN, ALTRETAMINE
HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)
HI-COR, HYDROCORTISONE
HIPREX, METHENAMINE HIPPURATE
HIVID, ZALCITABINE
HMS, MEDRYSONE
HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG PEN, INSULIN LISPRO RECOMBINANT
HUMALOG, INSULIN LISPRO RECOMBINANT
HUMATIN, PAROMOMYCIN SULFATE
HUMATROPE, SOMATROPIN RECOMBINANT
HUMULIN 50/50, INSULIN RECOMBINANT HUMAN (OTC)
HUMULIN 70/30 PEN, INSULIN RECOMBINANT HUMAN (OTC)
HUMULIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
HUMULIN L, INSULIN ZINC SUSP RECOMBINANT HUMAN (OTC)
HUMULIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
HUMULIN R PEN, INSULIN RECOMBINANT HUMAN (OTC)

APPENDIX A - PRODUCT NAME INDEX

A - 25

**** H ****

HUMULIN R, INSULIN RECOMBINANT HUMAN
HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
HUMULIN U, INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN (OTC)
HYCAMTIN, TOPOTECAN HYDROCHLORIDE
HYCODAN, HOMATROPINE METHYLBROMIDE
HYDASE, HYALURONIDASE
HYDERGINE, ERGOLOID MESYLATES
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
HYDREA, HYDROXYUREA
HYDROCET, ACETAMINOPHEN
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
HYDROCODONE COMPOUND, HOMATROPINE METHYLBROMIDE
HYDROCORTISONE ACETATE 1% AND PRAMOXINE HYDROCHLORIDE 1%, HYDROCORTISONE ACETATE
HYDROCORTISONE ACETATE, HYDROCORTISONE ACETATE
HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
HYDROCORTISONE, HYDROCORTISONE
HYDROCORTONE, HYDROCORTISONE
HYDROFLUMETHIAZIDE, HYDROFLUMETHIAZIDE
HYDROGENATED ERGOT ALKALOIDS, ERGOLOID MESYLATES
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
HYDRO-RIDE, AMILORIDE HYDROCHLORIDE
HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
HYDROXYUREA, HYDROXYUREA
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN
HYPAQUE, DIATRIZOATE MEGLUMINE
HYPAQUE, DIATRIZOATE SODIUM
HYPAQUE-76, DIATRIZOATE MEGLUMINE
HYPAQUE-CYSTO, DIATRIZOATE MEGLUMINE
HYPERSTAT, DIAZOXIDE
HY-PHEN, ACETAMINOPHEN
HYTONE, HYDROCORTISONE
HYTRIN, TERAZOSIN HYDROCHLORIDE
HYZAAR, HYDROCHLOROTHIAZIDE

**** I ****

IBU, IBUPROFEN
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
IBUPROHM, IBUPROFEN
IBUPROHM, IBUPROFEN (OTC)
IBU-TAB 200, IBUPROFEN (OTC)
IBU-TAB, IBUPROFEN
IC-GREEN, INDOCYANINE GREEN
IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
IFEX/MESNEX KIT, IFOSFAMIDE
IFOSFAMIDE, IFOSFAMIDE
IFOSFAMIDE/MESNA KIT, IFOSFAMIDE
ILETIN II, INSULIN PURIFIED PORK
IMDUR, ISOSORBIDE MONONITRATE

APPENDIX A - PRODUCT NAME INDEX

A - 26

**** I ****

IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
IMITREX, SUMATRIPTAN
IMITREX, SUMATRIPTAN SUCCINATE
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM ADVANCED, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM, LOPERAMIDE HYDROCHLORIDE
IMURAN, AZATHIOPRINE
INAPSINE, DROPERIDOL
INCRELEX, MECASERMIN RECOMBINANT
INDAPAMIDE, INDAPAMIDE
INDERAL LA, PROPRANOLOL HYDROCHLORIDE
INDERAL, PROPRANOLOL HYDROCHLORIDE
INDERIDE-40/25, HYDROCHLOROTHIAZIDE
INDICLOR, INDIUM IN 111 CHLORIDE
INDIUM IN 111 CHLORIDE, INDIUM IN 111 CHLORIDE
INDIUM IN-111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
INDOCIN I.V., INDOMETHACIN SODIUM
INDOCIN SR, INDOMETHACIN
INDOCIN, INDOMETHACIN
INDO-LEMMON, INDOMETHACIN
INDOMETHACIN, INDOMETHACIN
INDOMETHEGAN, INDOMETHACIN
INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
INFASURF PRESERVATIVE FREE, CALFACTANT
INFED, IRON DEXTRAN
INFLAMASE FORTE, PREDNISOLONE SODIUM PHOSPHATE
INFLAMASE MILD, PREDNISOLONE SODIUM PHOSPHATE
INFUMORPH, MORPHINE SULFATE
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
INFUVITE PEDIATRIC, ASCORBIC ACID
INNOFEM, ESTRADIOL
INNOHEP, TINZAPARIN SODIUM
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
INOMAX, NITRIC OXIDE
INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INSPIRA, EPLERENONE
INTAL, CROMOLYN SODIUM
INTEGRILIN, EPTIFIBATIDE
INTRALIPID 10%, SOYBEAN OIL
INTRALIPID 20%, SOYBEAN OIL
INTRALIPID 30%, SOYBEAN OIL
INVAGESIC FORTE, ASPIRIN
INVAGESIC, ASPIRIN
INVANZ, ERTAPENEM SODIUM
INVERSINE, MECAMYLAMINE HYDROCHLORIDE
INVIRASE, SAQUINAVIR MESYLATE
IOBENGUANE SULFATE I 131, IOBENGUANE SULFATE I 131
IODOTOPE, SODIUM IODIDE, I-131
IONAMIN, PHENTERMINE RESIN COMPLEX
IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IONOSOL T AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IOPAMIDOL-200 IN PLASTIC CONTAINER, IOPAMIDOL
IOPAMIDOL-200, IOPAMIDOL
IOPAMIDOL-250 IN PLASTIC CONTAINER, IOPAMIDOL
IOPAMIDOL-250, IOPAMIDOL
IOPAMIDOL-300 IN PLASTIC CONTAINER, IOPAMIDOL
IOPAMIDOL-300, IOPAMIDOL
IOPAMIDOL-370 IN PLASTIC CONTAINER, IOPAMIDOL
IOPAMIDOL-370, IOPAMIDOL

APPENDIX A - PRODUCT NAME INDEX

A - 27

**** I ****

IOPIDINE, APRACLONIDINE HYDROCHLORIDE
 IOSAT, POTASSIUM IODIDE (OTC)
 IPLEX, MECASERMIN RINFABATE RECOMBINANT
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 IPRIVASK, DESIRUDIN RECOMBINANT
 IQUIX, LEVOFLOXACIN
 IRESSA, GEFITINIB
 IRRIGATING SOLUTION G IN PLASTIC CONTAINER, CITRIC ACID
 ISMO, ISOSORBIDE MONONITRATE
 ISOCAINE HYDROCHLORIDE W/ LEVONORDEFIN, LEVONORDEFIN
 ISOCAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE
 ISOFLURANE, ISOFLURANE
 ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE E IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE S IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 ISONIAZID, ISONIAZID
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 ISOPTIN SR, VERAPAMIL HYDROCHLORIDE
 ISOPTIN, VERAPAMIL HYDROCHLORIDE
 ISOPTO CETAMIDE, SULFACETAMIDE SODIUM
 ISORDIL, ISOSORBIDE DINITRATE
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 ISOTONIC GENTAMICIN SULFATE IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 ISOVUE-200, IOPAMIDOL
 ISOVUE-250, IOPAMIDOL
 ISOVUE-300, IOPAMIDOL
 ISOVUE-370, IOPAMIDOL
 ISOVUE-M 200, IOPAMIDOL
 ISOVUE-M 300, IOPAMIDOL
 ISRADIPINE, ISRADIPINE
 ISTALOL, TIMOLOL MALEATE
 ISUPREL, ISOPROTERENOL HYDROCHLORIDE
 ITRACONAZOLE, ITRACONAZOLE
 IVY BLOCK, BENTOQUATAM (OTC)

**** J ****

JANTOVEN, WARFARIN SODIUM
 JEANATOPE, ALBUMIN IODINATED I-125 SERUM
 JUNEL 1.5/30, ETHINYL ESTRADIOL
 JUNEL 1/20, ETHINYL ESTRADIOL
 JUNEL FE 1.5/30, ETHINYL ESTRADIOL
 JUNEL FE 1/20, ETHINYL ESTRADIOL
 JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
 JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
 JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)

**** K ****

K+10, POTASSIUM CHLORIDE
 K+8, POTASSIUM CHLORIDE
 KADIAN, MORPHINE SULFATE
 KALETRA, LOPINAVIR
 KANAMYCIN, KANAMYCIN SULFATE
 KANTREX, KANAMYCIN SULFATE
 KAON CL-10, POTASSIUM CHLORIDE
 KARIVA, DESOGESTREL
 KAYEXALATE, SODIUM POLYSTYRENE SULFONATE

APPENDIX A - PRODUCT NAME INDEX

A - 28

**** K ****

K-DUR 10, POTASSIUM CHLORIDE
K-DUR 20, POTASSIUM CHLORIDE
KEFLEX, CEPHALEXIN
KELNOR, ETHINYL ESTRADIOL
KEMADRIN, PROCYCLIDINE HYDROCHLORIDE
KEMSTRO, BACLOFEN
KENACORT, TRIAMCINOLONE
KENALOG IN ORABASE, TRIAMCINOLONE ACETONIDE
KENALOG, TRIAMCINOLONE ACETONIDE
KENALOG-10, TRIAMCINOLONE ACETONIDE
KENALOG-40, TRIAMCINOLONE ACETONIDE
KEPPRA, LEVETIRACETAM
KERLONE, BETAXOLOL HYDROCHLORIDE
KETALAR, KETAMINE HYDROCHLORIDE
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
KETEK, TELITHROMYCIN
KETOCONAZOLE, KETOCONAZOLE
KETOPROFEN, KETOPROFEN
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
KETOZOLE, KETOCONAZOLE
KINEVAC, SINCALIDE
KIONEX, SODIUM POLYSTYRENE SULFONATE
KLARON, SULFACETAMIDE SODIUM
KLONOPIN RAPIDLY DISINTEGRATING, CLONAZEPAM
KLONOPIN, CLONAZEPAM
KLOR-CON M10, POTASSIUM CHLORIDE
KLOR-CON M15, POTASSIUM CHLORIDE
KLOR-CON M20, POTASSIUM CHLORIDE
KLOR-CON, POTASSIUM CHLORIDE
KLOTRIX, POTASSIUM CHLORIDE
K-TAB, POTASSIUM CHLORIDE
KYTRIL, GRANISETRON HYDROCHLORIDE

**** L ****

LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LAC-HYDRIN, AMMONIUM LACTATE
LACRISERT, HYDROXYPROPYL CELLULOSE
LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
LACTULOSE, LACTULOSE
LAMICTAL CD, LAMOTRIGINE
LAMICTAL, LAMOTRIGINE
LAMISIL AT, TERBINAFINE HYDROCHLORIDE (OTC)
LAMISIL, TERBINAFINE
LAMISIL, TERBINAFINE HYDROCHLORIDE
LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
LANIAZID, ISONIAZID
LANORINAL, ASPIRIN
LANOXICAPS, DIGOXIN
LANOXIN PEDIATRIC, DIGOXIN
LANOXIN, DIGOXIN
LANTUS, INSULIN GLARGINE RECOMBINANT
LARIAM, MEFLUQUINE HYDROCHLORIDE
LAROTID, AMOXICILLIN
LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
LASIX, FUROSEMIDE
LAXILOSE, LACTULOSE
LEFLUNOMIDE, LEFLUNOMIDE
LENTE ILETIN II (PORK), INSULIN ZINC SUSP PURIFIED PORK (OTC)
LESCOL XL, FLUVASTATIN SODIUM
LESCOL, FLUVASTATIN SODIUM
LESSINA-28, ETHINYL ESTRADIOL
LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM

APPENDIX A - PRODUCT NAME INDEX

A - 29

**** L ****

LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LEUKERAN, CHLORAMBUCIL
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
LEUSTATIN, CLADRIBINE
LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
LEVAQUIN, LEVOFLOXACIN
LEVATOL, PENBUTOLOL SULFATE
LEVEMIR, INSULIN DETEMIR RECOMBINANT
LEVITRA, VARDENAFIL HYDROCHLORIDE
LEVLITE, ETHINYL ESTRADIOL
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
LEVOCARNITINE, LEVOCARNITINE
LEVO-DROMORAN, LEVORPHANOL TARTRATE
LEVOFLOXACIN, LEVOFLOXACIN
LEVOLET, LEVOTHYROXINE SODIUM
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LEVOPHED, NOREPINEPHRINE BITARTRATE
LEVORA 0.15/30-28, ETHINYL ESTRADIOL
LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
LEVO-T, LEVOTHYROXINE SODIUM
LEVOTHROID, LEVOTHYROXINE SODIUM
LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
LEVOXYL, LEVOTHYROXINE SODIUM
LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE
LEXAPRO, ESCITALOPRAM OXALATE
LEXIVA, FOSAMPRENAVIR CALCIUM
LEXXEL, ENALAPRIL MALEATE
LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
LIDEX, FLUOCINONIDE
LIDEX-E, FLUOCINONIDE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5%, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5%, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.8% IN DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE, EPINEPHRINE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
LIDOCAINE, LIDOCAINE
LIDOCATON, EPINEPHRINE
LIDODERM, LIDOCAINE
LIDOPEN, LIDOCAINE HYDROCHLORIDE
LIDOSITE TOPICAL SYSTEM KIT, EPINEPHRINE
LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
LIMBITROL DS, AMITRIPTYLINE HYDROCHLORIDE
LIMBITROL, AMITRIPTYLINE HYDROCHLORIDE
LINCOCIN, LINCOMYCIN HYDROCHLORIDE
LINDANE, LINDANE
LIORESAL, BACLOFEN
LIOTHYRONINE SODIUM, LIOTHYRONINE SODIUM
LIPITOR, ATORVASTATIN CALCIUM
LIPOSYN II 10%, SAFFLOWER OIL
LIPOSYN II 20%, SAFFLOWER OIL

APPENDIX A - PRODUCT NAME INDEX

A - 30

**** L ****

LIPOSYN III 10%, SOYBEAN OIL
LIPOSYN III 20%, SOYBEAN OIL
LIPOSYN III 30%, SOYBEAN OIL
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LITHIUM CARBONATE, LITHIUM CARBONATE
LITHIUM CITRATE, LITHIUM CITRATE
LITHOBID, LITHIUM CARBONATE
LITHOSTAT, ACETOHYDROXAMIC ACID
LIVOSTIN, LEVOCABASTINE HYDROCHLORIDE
LO/OVRAL-28, ETHINYL ESTRADIOL
LOCHOLEST LIGHT, CHOLESTYRAMINE
LOCHOLEST, CHOLESTYRAMINE
LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
LOCOID, HYDROCORTISONE BUTYRATE
LODINE, ETODOLAC
LODOSYN, CARBIDOPA
LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL
LOESTRIN 21 1/20, ETHINYL ESTRADIOL
LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL
LOESTRIN FE 1/20, ETHINYL ESTRADIOL
LOMOTIL, ATROPINE SULFATE
LONITEN, MINOXIDIL
LONOX, ATROPINE SULFATE
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOPID, GEMFIBROZIL
LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
LOPRESSOR, METOPROLOL TARTRATE
LOPROX, CICLOPIROX
LOPURIN, ALLOPURINOL
LORABID, LORACARBEF
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE, LORATADINE (OTC)
LORAZEPAM INTENSOL, LORAZEPAM
LORAZEPAM PRESERVATIVE FREE, LORAZEPAM
LORAZEPAM, LORAZEPAM
LORCET-HD, ACETAMINOPHEN
LORTAB, ACETAMINOPHEN
LOTEMAX, LOTEPREDNOL ETABONATE
LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
LOTENSIN, BENAZEPRIL HYDROCHLORIDE
LOTREL, AMLODIPINE BESYLATE
LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
LOTRISONE, BETAMETHASONE DIPROPIONATE
LOTRONEX, ALOSETRON HYDROCHLORIDE
LOVASTATIN, LOVASTATIN
LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
LOVENOX, ENOXAPARIN SODIUM
LOW-OGESTREL-21, ETHINYL ESTRADIOL
LOW-OGESTREL-28, ETHINYL ESTRADIOL
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
LOXAPINE, LOXAPINE SUCCINATE
LOXITANE, LOXAPINE SUCCINATE
LOZOL, INDAPAMIDE
LTA II KIT, LIDOCAINE HYDROCHLORIDE
LUFYLLIN, DYPHYLLINE
LUMIGAN, BIMATOPROST
LUNESTA, ESZOPICLONE
LUPRON DEPOT, LEUPROLIDE ACETATE
LUPRON DEPOT-3, LEUPROLIDE ACETATE
LUPRON DEPOT-4, LEUPROLIDE ACETATE
LUPRON DEPOT-PED, LEUPROLIDE ACETATE

APPENDIX A - PRODUCT NAME INDEX

A - 31

**** L ****

LUPRON, LEUPROLIDE ACETATE
LUVERIS, LUTROPIN ALFA
LUXIQ, BETAMETHASONE VALERATE
LYMPHAZURIN, ISOSULFAN BLUE
LYOPHILIZED CYTOXAN, CYCLOPHOSPHAMIDE
LYRICA, PREGABALIN
LYSODREN, MITOTANE

**** M ****

M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID
M.V.I. ADULT, ASCORBIC ACID
M.V.I. PEDIATRIC, ASCORBIC ACID
M.V.I.-12 (WITHOUT VITAMIN K), ASCORBIC ACID
MACROBID, NITROFURANTOIN
MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
MACROTEC, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
MACUGEN, PEGAPTANIB SODIUM
MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE
MAGNESIUM SULFATE, MAGNESIUM SULFATE
MAGNEVIST, GADOPENTETATE DIMEGLUMINE
MALARONE PEDIATRIC, ATOVAQUONE
MALARONE, ATOVAQUONE
MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE
MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER, MANNITOL
MANNITOL 10%, MANNITOL
MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%, MANNITOL
MANNITOL 15%, MANNITOL
MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 20%, MANNITOL
MANNITOL 25%, MANNITOL
MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%, MANNITOL
MANNITOL 5%, MANNITOL
MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE
MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE
MARCAINE, BUPIVACAINE HYDROCHLORIDE
MARINOL, DRONABINOL
MARPLAN, ISOCARBOXAZID
MATULANE, PROCARBAZINE HYDROCHLORIDE
MAVIK, TRANDOLAPRIL
MAXAIR, PIRBUTEROL ACETATE
MAXALT, RIZATRIPTAN BENZOATE
MAXALT-MLT, RIZATRIPTAN BENZOATE
MAXAQUIN, LOMEFLOXACIN HYDROCHLORIDE
MAXIDEX, DEXAMETHASONE
MAXIPIME, CEFEPIME HYDROCHLORIDE (ARGININE FORMULATION)
MAXITROL, DEXAMETHASONE
MAXZIDE, HYDROCHLOROTHIAZIDE
MAXZIDE-25, HYDROCHLOROTHIAZIDE
MD-76R, DIATRIZOATE MEGLUMINE
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
MDP-BRACCO, TECHNETIUM TC-99M MEDRONATE KIT
MEBENDAZOLE, MEBENDAZOLE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
MEDIPREN, IBUPROFEN (OTC)
MEDROL, METHYLPREDNISOLONE

APPENDIX A - PRODUCT NAME INDEX

A - 32

**** M ****

MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MEFLOQUINE, MEFLOQUINE HYDROCHLORIDE
MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
MEFOXIN, CEFOXITIN SODIUM
MEGACE ES, MEGESTROL ACETATE
MEGACE, MEGESTROL ACETATE
MEGATOPE, ALBUMIN IODINATED I-131 SERUM
MEGESTROL ACETATE, MEGESTROL ACETATE
MENEST, ESTROGENS, ESTERIFIED
MENOPUR, LUTEINIZING HORMONE
MENOSTAR, ESTRADIOL
MENTAX, BUTENAFINE HYDROCHLORIDE
MENTAX-TC, BUTENAFINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MEPHYTON, PHYTONADIONE
MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFIN, LEVONORDEFIN
MEPIVACAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE
MEPROBAMATE, MEPROBAMATE
MEPRON, ATOVAQUONE
MERCAPTOPYRINE, MERCAPTOPYRINE
MERCAPTOPYRINE, MERCAPTOPYRINE ANHYDROUS
MERIDIA, SIBUTRAMINE HYDROCHLORIDE
MERREM I.V., MEROPENEM
MESALAMINE, MESALAMINE
MESNA, MESNA
MESNEX, MESNA
MESTINON, PYRIDOSTIGMINE BROMIDE
METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
METADATE ER, METHYLPHENIDATE HYDROCHLORIDE
METAGLIP, GLIPIZIDE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
METARAMINOL BITARTRATE, METARAMINOL BITARTRATE
METASTRON, STRONTIUM CHLORIDE, SR-89
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHADOSE, METHADONE HYDROCHLORIDE
METHAZOLAMIDE, METHAZOLAMIDE
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
METHERGINE, METHYLERGONOVINE MALEATE
METHIMAZOLE, METHIMAZOLE
METHOCARBAMOL AND ASPIRIN, ASPIRIN
METHOCARBAMOL, METHOCARBAMOL
METHOTREXATE PRESERVATIVE FREE, METHOTREXATE
METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
METHOTREXATE SODIUM, METHOTREXATE SODIUM
METHOTREXATE, METHOTREXATE SODIUM
METHYLCLOTHIAZIDE, METHYLCLOTHIAZIDE
METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METHYLDOPA, METHYLDOPA
METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE
METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HCL, METHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METHYLTESTOSTERONE, METHYLTESTOSTERONE
METIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 33

**** M ****

METOLAZONE, METOLAZONE
METOPIRONE, METYRAPONE
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
METROCREAM, METRONIDAZOLE
METROGEL, METRONIDAZOLE
METROGEL-VAGINAL, METRONIDAZOLE
METROLOTION, METRONIDAZOLE
METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE
METRONIDAZOLE, METRONIDAZOLE
METVIXIA, METHYL AMINOLEVULINATE HYDROCHLORIDE
MEVACOR, LOVASTATIN
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
MEXITIL, MEXILETINE HYDROCHLORIDE
MIACALCIN, CALCITONIN, SALMON
MICARDIS HCT, HYDROCHLOROTHIAZIDE
MICARDIS, TELMISARTAN
MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
MICONAZOLE 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE, MICONAZOLE NITRATE
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
MICORT-HC, HYDROCORTISONE ACETATE
MICRODERM, CHLORHEXIDINE GLUCONATE (OTC)
MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL
MICROGESTIN FE 1/20, ETHINYL ESTRADIOL
MICRO-K 10, POTASSIUM CHLORIDE
MICRO-K, POTASSIUM CHLORIDE
MICRONASE, GLYBURIDE
MICRONOR, NORETHINDRONE
MICROZIDE, HYDROCHLOROTHIAZIDE
MIDAMOR, AMILORIDE HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MIFEPREX, MIFEPRISTONE
MIGERGOT, CAFFEINE
MIGRANAL, DIHYDROERGOTAMINE MESYLATE
MILOPHENE, CLOMIPHENE CITRATE
MILRINONE LACTATE IN 5% DEXTROSE IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE, MILRINONE LACTATE
MILRINONE LACTOSE IN 5% DEXTROSE, MILRINONE LACTATE
MILTOWN, MEPROBAMATE
MINIPRESS XL, PRAZOSIN HYDROCHLORIDE
MINIPRESS, PRAZOSIN HYDROCHLORIDE
MINIRIN, DESMOPRESSIN ACETATE
MINITRAN, NITROGLYCERIN
MINIZIDE, POLYTHIAZIDE
MINOCIN, MINOCYCLINE HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL, MINOXIDIL
MINTEZOL, THIABENDAZOLE
MIOCHOL, ACETYLCHOLINE CHLORIDE
MIOCHOL-E, ACETYLCHOLINE CHLORIDE
MIOSTAT, CARBACHOL
MIRADON, ANISINDIONE
MIRALAX, POLYETHYLENE GLYCOL 3350
MIRALUMA, TECHNETIUM TC-99M SESTAMIBI KIT

APPENDIX A - PRODUCT NAME INDEX

A - 34

**** M ****

MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE
MIRCETTE, DESOGESTREL
MIRENA, LEVONORGESTREL
MIRTAZAPINE, MIRTAZAPINE
MISOPROSTOL, MISOPROSTOL
MITHRACIN, PLICAMYCIN
MITOMYCIN, MITOMYCIN
MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER, MIVACURIUM CHLORIDE
MIVACRON, MIVACURIUM CHLORIDE
MOBAN, MOLINDONE HYDROCHLORIDE
MOBIC, MELOXICAM
MODERIL, RESCINNAMINE
MODICON 28, ETHINYL ESTRADIOL
MODURETIC 5-50, AMILORIDE HYDROCHLORIDE
MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
MOMETASONE FUROATE, MOMETASONE FUROATE
MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3, MICONAZOLE NITRATE
MONISTAT 3, MICONAZOLE NITRATE (OTC)
MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 7, MICONAZOLE NITRATE (OTC)
MONISTAT-3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT-DERM, MICONAZOLE NITRATE
MONODOX, DOXYCYCLINE
MONOKET, ISOSORBIDE MONONITRATE
MONOPRIL, FOSINOPRIL SODIUM
MONOPRIL-HCT, FOSINOPRIL SODIUM
MONUROL, FOSFOMYCIN TROMETHAMINE
MORPHINE SULFATE, MORPHINE SULFATE
MOTOFEN, ATROPINE SULFATE
MOTRIN IB, IBUPROFEN (OTC)
MOTRIN MIGRAINE PAIN, IBUPROFEN (OTC)
MOTRIN, IBUPROFEN
MPI DMSA KIDNEY REAGENT, TECHNETIUM TC-99M SUCCIMER KIT
MPI DTPA KIT - CHELATE, TECHNETIUM TC-99M PENTETATE KIT
MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
MS CONTIN, MORPHINE SULFATE
MUCINEX D, GUAIFENESIN (OTC)
MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
MUCINEX, GUAIFENESIN (OTC)
MUCOMYST, ACETYLCYSTEINE
MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
MULTIHANCE, GADOBENATE DIMEGLUMINE
MUPIROCIN, MUPIROCIN
MUSE, ALPROSTADIL
MUSTARGEN, MECHLORETHAMINE HYDROCHLORIDE
MUTAMYCIN, MITOMYCIN
MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
MYBANIL, BROMODIPHENHYDRAMINE HYDROCHLORIDE
MYCAMINE, MICAFUNGIN SODIUM
MYCELEX, CLOTRIMAZOLE
MYCELEX-7 COMBINATION PACK, CLOTRIMAZOLE (OTC)
MYCELEX-7, CLOTRIMAZOLE (OTC)
MYCOBUTIN, RIFABUTIN
MYCODONE, HOMATROPINE METHYLBROMIDE
MYCOSTATIN, NYSTATIN
MYDRIACYL, TROPICAMIDE
MYFORTIC, MYCOPHENOLIC ACID
MYKACET, NYSTATIN
MYLARAMINE, DEXCHLORPHENIRAMINE MALEATE
MYLERAN, BUSULFAN
MYLOTARG, GEMTUZUMAB OZOGAMICIN

APPENDIX A - PRODUCT NAME INDEX

A - 35

**** M ****

MYMETHASONE, DEXAMETHASONE
MYOVIEW 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
MYOVIEW, TECHNETIUM TC-99M TETROFOSMIN KIT
MYPHETANE DX, BROMPHENIRAMINE MALEATE
MYPROIC ACID, VALPROIC ACID
MYSOLINE, PRIMIDONE
MYTELASE, AMBENONIUM CHLORIDE
MYTOZYTREX, MITOMYCIN
M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
M-ZOLE 7 DUAL PACK, MICONAZOLE NITRATE (OTC)

**** N ****

NABUMETONE, NABUMETONE
NADOLOL, NADOLOL
NAFAZAIR, NAPHAZOLINE HYDROCHLORIDE
NAFCILLIN SODIUM, NAFCILLIN SODIUM
NAFTIN, NAFTIFINE HYDROCHLORIDE
NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
NALFON 200, FENOPROFEN CALCIUM
NALFON, FENOPROFEN CALCIUM
NALIDIXIC ACID, NALIDIXIC ACID
NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM
NALOXONE HYDROCHLORIDE AND PENTAZOCAINE, NALOXONE HYDROCHLORIDE
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
NAMENDA, MEMANTINE HYDROCHLORIDE
NANDROLONE DECANOATE, NANDROLONE DECANOATE
NAPHAZOLINE HYDROCHLORIDE, NAPHAZOLINE HYDROCHLORIDE
NAPHCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
NAPRAPAC 250 (COPACKAGED), LANSOPRAZOLE
NAPRAPAC 375 (COPACKAGED), LANSOPRAZOLE
NAPRAPAC 500 (COPACKAGED), LANSOPRAZOLE
NAPRELAN, NAPROXEN SODIUM
NAPROSYN, NAPROXEN
NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
NAPROXEN, NAPROXEN
NARCAN, NALOXONE HYDROCHLORIDE
NARDIL, PHENELZINE SULFATE
NAROPIN, ROPIVACAINE HYDROCHLORIDE MONOHYDRATE
NASACORT AQ, TRIAMCINOLONE ACETONIDE
NASACORT HFA, TRIAMCINOLONE ACETONIDE
NASALCROM, CROMOLYN SODIUM (OTC)
NASAREL, FLUNISOLIDE
NASCOBAL, CYANOCOBALAMIN
NASONEX, MOMETASONE FUROATE MONOHYDRATE
NATACYN, NATAMYCIN
NATRECOR, NESIRITIDE RECOMBINANT
NATURETIN-5, BENDROFLUMETHIAZIDE
NAVANE, THIOTHIXENE
NAVANE, THIOTHIXENE HYDROCHLORIDE
NAVELBINE, VINORELBINE TARTRATE
NEBUPENT, PENTAMIDINE ISETHIONATE
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
NEGGRAM, NALIDIXIC ACID
NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
NEO TECT KIT, TECHNETIUM TC-99M DEPREOTIDE
NEO-FRADIN, NEOMYCIN SULFATE
NEOMYCIN AND POLYMYXIN B SULFATE AND BACITRACIN ZINC, BACITRACIN ZINC
NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN

APPENDIX A - PRODUCT NAME INDEX

A - 36

**** N ****

NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
NEOMYCIN SULFATE, NEOMYCIN SULFATE
NEOPAP, ACETAMINOPHEN (OTC)
NEORAL, CYCLOSPORINE
NEO-RX, NEOMYCIN SULFATE
NEOSAR, CYCLOPHOSPHAMIDE
NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
NEOSPORIN, BACITRACIN ZINC
NEOSPORIN, GRAMICIDIN
NEPHRAMINE 5.4%, AMINO ACIDS
NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
NEURONTIN, GABAPENTIN
NEUTREXIN, TRIMETREXATE GLUCURONATE
NEVANAC, NEPAFENAC
NEXAVAR, SORAFENIB TOSYLATE
NEXIUM IV, ESOMEPRAZOLE SODIUM
NEXIUM, ESOMEPRAZOLE MAGNESIUM
NIACIN, NIACIN
NIACOR, NIACIN
NIASPAN, NIACIN
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NICODERM CQ, NICOTINE (OTC)
NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
NICORETTE (ORANGE), NICOTINE POLACRILEX (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
NICOTINE POLACRILEX (MINT), NICOTINE POLACRILEX (OTC)
NICOTINE POLACRILEX (ORANGE), NICOTINE POLACRILEX (OTC)
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
NICOTINE, NICOTINE
NICOTROL, NICOTINE
NIFEDIPINE, NIFEDIPINE
NILANDRON, NILUTAMIDE
NILSTAT, NYSTATIN
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
NIMBEX, CISATRACURIUM BESYLATE
NIMOTOP, NIMODIPINE
NIPENT, PENTOSTATIN
NIRAVAM, ALPRAZOLAM
NITRO-DUR, NITROGLYCERIN
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NITROFURAZONE, NITROFURAZONE
NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
NITROGLYCERIN, NITROGLYCERIN
NITROLINGUAL PUMPSPRAY, NITROGLYCERIN
NITROPRESS, SODIUM NITROPRUSSIDE
NITROSTAT, NITROGLYCERIN
NIX, PERMETHRIN (OTC)
NIZATIDINE, NIZATIDINE
NIZORAL A-D, KETOCONAZOLE (OTC)
NIZORAL, KETOCONAZOLE
NOLVADEX, TAMOXIFEN CITRATE
NORCO, ACETAMINOPHEN
NORDETTE-28, ETHINYL ESTRADIOL
NORDITROPIN NORDIFLEX, SOMATROPIN RECOMBINANT
NORDITROPIN, SOMATROPIN RECOMBINANT
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
NORETHIN 1/35E-21, ETHINYL ESTRADIOL
NORETHIN 1/35E-28, ETHINYL ESTRADIOL
NORETHIN 1/50M-21, MESTRANOL
NORETHIN 1/50M-28, MESTRANOL

APPENDIX A - PRODUCT NAME INDEX

A - 37

**** N ****

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL (7/14), ETHINYL ESTRADIOL
NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
NORETHINDRONE AND MESTRANOL, MESTRANOL
NORFLEX, ORPHENADRINE CITRATE
NORGESIC FORTE, ASPIRIN
NORGESIC, ASPIRIN
NORINYL 1+35 21-DAY, ETHINYL ESTRADIOL
NORINYL 1+35 28-DAY, ETHINYL ESTRADIOL
NORINYL 1+50 21-DAY, MESTRANOL
NORINYL 1+50 28-DAY, MESTRANOL
NORITATE, METRONIDAZOLE
NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
NOROXIN, NORFLOXACIN
NORPACE CR, DISOPYRAMIDE PHOSPHATE
NORPACE, DISOPYRAMIDE PHOSPHATE
NORPLANT II, LEVONORGESTREL
NORPRAMIN, DESIPRAMINE HYDROCHLORIDE
NOR-QD, NORETHINDRONE
NORTREL 0.5/35-21, ETHINYL ESTRADIOL
NORTREL 0.5/35-28, ETHINYL ESTRADIOL
NORTREL 1/35-21, ETHINYL ESTRADIOL
NORTREL 1/35-28, ETHINYL ESTRADIOL
NORTREL 7/7/7, ETHINYL ESTRADIOL
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
NORVASC, AMLODIPINE BESYLATE
NORVIR, RITONAVIR
NOVAMINE 11.4%, AMINO ACIDS
NOVAMINE 15%, AMINO ACIDS
NOVANTRONE, MITOXANTRONE HYDROCHLORIDE
NOVOCAIN, PROCAINE HYDROCHLORIDE
NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
NOVOLIN L, INSULIN ZINC SUSP RECOMBINANT HUMAN (OTC)
NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)
NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT
NOVOLOG, INSULIN ASPART RECOMBINANT
NOVOTHYROX, LEVOTHYROXINE SODIUM
NPH ILETIN I (BEEF-PORK), INSULIN SUSP ISOPHANE BEEF/PORK (OTC)
NPH ILETIN II (PORK), INSULIN SUSP ISOPHANE PURIFIED PORK (OTC)
NUBAIN, NALBUPHINE HYDROCHLORIDE
NULYTELY, POLYETHYLENE GLYCOL 3350
NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350
NUMORPHAN, OXYMORPHONE HYDROCHLORIDE
NUTRACORT, HYDROCORTISONE
NUTRILIPID 10%, SOYBEAN OIL
NUTRILIPID 20%, SOYBEAN OIL
NUTROPIN AQ PEN, SOMATROPIN RECOMBINANT
NUTROPIN AQ, SOMATROPIN RECOMBINANT
NUTROPIN DEPOT, SOMATROPIN RECOMBINANT
NUTROPIN, SOMATROPIN RECOMBINANT
NUVARING, ETHINYL ESTRADIOL
NYDRAZID, ISONIAZID
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
NYSTATIN, NYSTATIN
NYSTATIN-TRIAMCINOLONE ACETONIDE, NYSTATIN
NYSTOP, NYSTATIN

**** 0 ****

APPENDIX A - PRODUCT NAME INDEX

A - 38

**** 0 ****

OCTOCAINE, EPINEPHRINE
OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
OCUCLEAR, OXYMETAZOLINE HYDROCHLORIDE (OTC)
OCUFEN, FLURBIPROFEN SODIUM
OCUFLOX, OFLOXACIN
OCUPRESS, CARTEOLOL HYDROCHLORIDE
OCUSULF-10, SULFACETAMIDE SODIUM
OFLOXACIN, OFLOXACIN
OGEN .625, ESTROPIPATE
OGEN 1.25, ESTROPIPATE
OGEN 2.5, ESTROPIPATE
OGEN 5, ESTROPIPATE
OGEN, ESTROPIPATE
OGESTREL 0.5/50-21, ETHINYL ESTRADIOL
OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
OLUX, CLOBETASOL PROPIONATE
OMACOR, OMEGA-3-ACID ETHYL ESTERS
OMEPRAZOLE, OMEPRAZOLE
OMNICEF, CEFDINIR
OMNIPAQUE 140, IOHEXOL
OMNIPAQUE 180, IOHEXOL
OMNIPAQUE 240, IOHEXOL
OMNIPAQUE 300, IOHEXOL
OMNIPAQUE 350, IOHEXOL
OMNISCAN, GADODIAMIDE
OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
OPHTHAINE, PROPARACAINE HYDROCHLORIDE
OPHTHETIC, PROPARACAINE HYDROCHLORIDE
OPTICROM, CROMOLYN SODIUM
OPTIMARK IN PLASTIC CONTAINER, GADOVERSETAMIDE
OPTIMARK, GADOVERSETAMIDE
OPTIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
OPTIRAY 160, IOVERSOL
OPTIRAY 240, IOVERSOL
OPTIRAY 300, IOVERSOL
OPTIRAY 320, IOVERSOL
OPTIRAY 350, IOVERSOL
OPTISON, ALBUMIN HUMAN
OPTIVAR, AZELASTINE HYDROCHLORIDE
ORABASE HCA, HYDROCORTISONE ACETATE
ORACORT, TRIAMCINOLONE ACETONIDE
ORALONE, TRIAMCINOLONE ACETONIDE
ORAMORPH SR, MORPHINE SULFATE
ORAP, PIMOZIDE
ORAPRED, PREDNISOLONE SODIUM PHOSPHATE
ORAQIX, LIDOCAINE
ORETIC, HYDROCHLOROTHIAZIDE
ORFADIN, NITISINONE
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
ORPHENGESIC FORTE, ASPIRIN
ORPHENGESIC, ASPIRIN
ORTHO CYCLEN-28, ETHINYL ESTRADIOL
ORTHO EVRA, ETHINYL ESTRADIOL
ORTHO TRI-CYCLEN LO, ETHINYL ESTRADIOL
ORTHO TRI-CYCLEN, ETHINYL ESTRADIOL
ORTHO-CEPT, DESOGESTREL
ORTHO-EST, ESTROPIPATE
ORTHO-NOVUM 1/35-28, ETHINYL ESTRADIOL
ORTHO-NOVUM 1/50 28, MESTRANOL
ORTHO-NOVUM 10/11-28, ETHINYL ESTRADIOL

APPENDIX A - PRODUCT NAME INDEX

A - 39

**** O ****

ORTHO-NOVUM 7/7/7-28, ETHINYL ESTRADIOL
ORUDIS KT, KETOPROFEN (OTC)
ORUVAIL, KETOPROFEN
ORVATEN, MIDODRINE HYDROCHLORIDE
OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL
OSMITROL 10% IN WATER, MANNITOL
OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL
OSMITROL 15% IN WATER, MANNITOL
OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL
OSMITROL 20% IN WATER, MANNITOL
OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL
OSMITROL 5% IN WATER, MANNITOL
OTICAIR, HYDROCORTISONE
OVCON-35, ETHINYL ESTRADIOL
OVCON-50, ETHINYL ESTRADIOL
OVIDE, MALATHION
OVIDREL, CHORIOGONADOTROPIN ALFA
OXACILLIN SODIUM, OXACILLIN SODIUM
OXANDRIN, OXANDROLONE
OXAPROZIN, OXAPROZIN
OXAZEPAM, OXAZEPAM
OXILAN-300, IOXILAN
OXILAN-350, IOXILAN
OXISTAT, OXICONAZOLE NITRATE
OXSORALEN, METHOXSALEN
OXSORALEN-ULTRA, METHOXSALEN
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
OXYCET, ACETAMINOPHEN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE AND ASPIRIN, ASPIRIN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
OXYCONTIN, OXYCODONE HYDROCHLORIDE
OXYTOCIN, OXYTOCIN
OXYTROL, OXYBUTYNIN

**** P ****

PACERONE, AMIODARONE HYDROCHLORIDE
PACLITAXEL, PACLITAXEL
PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
PAMINE FORTE, METHSCOPOLAMINE BROMIDE
PAMINE, METHSCOPOLAMINE BROMIDE
PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
PANCURONIUM, PANCURONIUM BROMIDE
PANDEL, HYDROCORTISONE PROBUTATE
PANIXINE DISPERDOSE, CEPHALEXIN
PANRETIN, ALITRETINOIN
PARACAINE, PROPARACAINE HYDROCHLORIDE
PARAFON FORTE DSC, CHLORZOXAZONE
PARAPLATIN, CARBOPLATIN
PARCOPA, CARBIDOPA
PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
PARLODEL, BROMOCRIPTINE MESYLATE
PARNATE, TRANYLCPROMINE SULFATE
PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PASER, AMINOSALICYLIC ACID
PASKALIUM, POTASSIUM AMINOSALICYLATE
PATANOL, OLOPATADINE HYDROCHLORIDE
PAXIL CR, PAROXETINE HYDROCHLORIDE
PAXIL, PAROXETINE HYDROCHLORIDE
PBZ, TRIPELENNAMINE HYDROCHLORIDE
PCE, ERYTHROMYCIN

APPENDIX A - PRODUCT NAME INDEX

A - 40

**** P ****

PEDIAMYCIN 400, ERYTHROMYCIN ETHYLSUCCINATE
PEDIAMYCIN, ERYTHROMYCIN ETHYLSUCCINATE
PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
PEDIATRIC ADVIL, IBUPROFEN (OTC)
PEDIATRIC LTA KIT, LIDOCAINE HYDROCHLORIDE
PEDIAZOLE, ERYTHROMYCIN ETHYLSUCCINATE
PEDIOTIC, HYDROCORTISONE
PEGANONE, ETHOTOIN
PENECORT, HYDROCORTISONE
PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
PENICILLIN G SODIUM, PENICILLIN G SODIUM
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM
PENICILLIN-VK, PENICILLIN V POTASSIUM
PENLAC, CICLOPIROX
PENTAM, PENTAMIDINE ISETHIONATE
PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
PENTASA, MESALAMINE
PENTAZOCINE AND NALOXONE HYDROCHLORIDES, NALOXONE HYDROCHLORIDE
PENTAZOCINE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
PENTETATE CALCIUM TRISODIUM, PENTETATE CALCIUM TRISODIUM
PENTETATE ZINC TRISODIUM, PENTETATE ZINC TRISODIUM
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
PENTOXIFYLLINE, PENTOXIFYLLINE
PENTOXIL, PENTOXIFYLLINE
PEPCID AC (GELTAB), FAMOTIDINE (OTC)
PEPCID AC, FAMOTIDINE (OTC)
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
PEPCID PRESERVATIVE FREE, FAMOTIDINE
PEPCID, FAMOTIDINE
PERCOCET, ACETAMINOPHEN
PERCODAN, ASPIRIN
PERCODAN-DEMI, ASPIRIN
PERGOLIDE MESYLATE, PERGOLIDE MESYLATE
PERIDEX, CHLORHEXIDINE GLUCONATE
PERIOCHIP, CHLORHEXIDINE GLUCONATE
PERIOGARD, CHLORHEXIDINE GLUCONATE
PERIOSTAT, DOXYCYCLINE HYCLATE
PERMAPEN, PENICILLIN G BENZATHINE
PERMAX, PERGOLIDE MESYLATE
PERMETHRIN, PERMETHRIN
PERMETHRIN, PERMETHRIN (OTC)
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
PERPHENAZINE, PERPHENAZINE
PERSANTINE, DIPYRIDAMOLE
PEXEVA, PAROXETINE MESYLATE
PFIZERPEN, PENICILLIN G POTASSIUM
PHARMASEAL SCRUB CARE, CHLORHEXIDINE GLUCONATE (OTC)
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENERGAN, PROMETHAZINE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
PHENYTOIN, PHENYTOIN
PHENYTEK, PHENYTOIN SODIUM
PHENYTOIN SODIUM, PHENYTOIN SODIUM
PHENYTOIN, PHENYTOIN
PHENYTOIN, PHENYTOIN SODIUM
PHISOHEX, HEXACHLOROPHENE
PHOSLO GELCAPS, CALCIUM ACETATE
PHOSLO, CALCIUM ACETATE
PHOSPHOCOL P32, CHROMIC PHOSPHATE, P-32
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE

APPENDIX A - PRODUCT NAME INDEX

A - 41

**** P ****

PHOSPHOTEC, TECHNETIUM TC-99M PYROPHOSPHATE KIT
PHOTOFRIN, PORFIMER SODIUM
PHRENILIN FORTE, ACETAMINOPHEN
PHRENILIN WITH CAFFEINE AND CODEINE, ACETAMINOPHEN
PHRENILIN, ACETAMINOPHEN
PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PHYTONADIONE, PHYTONADIONE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
PILOPINE HS, PILOCARPINE HYDROCHLORIDE
PINDOLOL, PINDOLOL
PIPERACILLIN, PIPERACILLIN SODIUM
PIROXICAM, PIROXICAM
PITOCIN, OXYTOCIN
PLAN B, LEVONORGESTREL
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
PLASMA-LYTE 56 IN PLASTIC CONTAINER, MAGNESIUM ACETATE TETRAHYDRATE
PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PLASMA-LYTE R IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PLATINOL, CISPLATIN
PLATINOL-AQ, CISPLATIN
PLAVIX, CLOPIDOGREL BISULFATE
PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE
PLENAXIS, ABARELIX
PLENDIL, FELODIPINE
PLETAL, CILOSTAZOL
PODOFILOX, PODOFILOX
POLOCAINE W/ LEVONORDEFIN, LEVONORDEFIN
POLOCAINE, MEPIVACAINE HYDROCHLORIDE
POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
POLY-PRED, NEOMYCIN SULFATE
POLY-RX, POLYMYXIN B SULFATE
POLYSPORIN, BACITRACIN ZINC
POLYTRIM, POLYMYXIN B SULFATE
PONSTEL, MEFENAMIC ACID
PORTIA-28, ETHINYL ESTRADIOL
POTASSIUM ACETATE IN PLASTIC CONTAINER, POTASSIUM ACETATE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE

A - 42

**** P ****

[illegible]

APPENDIX A - PRODUCT NAME INDEX

A - 43

**** P ****

POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM
CHLORIDE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM
CHLORIDE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
POVIDONE IODINE, POVIDONE-IODINE (OTC)
PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE
PRAMOSONE, HYDROCORTISONE ACETATE
PRANDIN, REPAGLINIDE
PRAVACHOL, PRAVASTATIN SODIUM
PRAVIGARD PAC (COPACKAGED), ASPIRIN
PRAZOSIN HCL, PRAZOSIN HYDROCHLORIDE
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
PRECEDEX, DEXMEDETOMIDINE
PRECOSE, ACARBOSE
PRED FORTE, PREDNISOLONE ACETATE
PRED MILD, PREDNISOLONE ACETATE
PRED-G, GENTAMICIN SULFATE
PREDNISOLONE ACETATE, PREDNISOLONE ACETATE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISOLONE, PREDNISOLONE
PREDNISONE INTENSOL, PREDNISONE
PREDNISONE, PREDNISONE
PREFEST, ESTRADIOL
PREGNYL, GONADOTROPIN, CHORIONIC
PRELONE, PREDNISOLONE
PREMARIN, ESTROGENS, CONJUGATED
PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
PREMPHASE 14/14, ESTROGENS, CONJUGATED
PREMPRO, ESTROGENS, CONJUGATED
PRE-OP II, HEXACHLOROPHENE
PRE-OP, HEXACHLOROPHENE
PREPIDIL, DINOPROSTONE
PREVACARE R, CHLORHEXIDINE GLUCONATE (OTC)
PREVACID IV, LANSOPRAZOLE
PREVACID, LANSOPRAZOLE
PREVALITE, CHOLESTYRAMINE
PREVIFEM, ETHINYL ESTRADIOL
PREVPAC, AMOXICILLIN
PRIALT, ZICONOTIDE
PRIFTIN, RIFAPENTINE
PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
PRILOSEC, OMEPRAZOLE
PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
PRIMACOR, MILRINONE LACTATE

APPENDIX A - PRODUCT NAME INDEX

A - 44

**** P ****

PRIMAQUINE, PRIMAQUINE PHOSPHATE
PRIMATENE MIST, EPINEPHRINE (OTC)
PRIMAXIN, CILASTATIN SODIUM
PRIMIDONE, PRIMIDONE
PRIMSOL, TRIMETHOPRIM HYDROCHLORIDE
PRINCIPEN, AMPICILLIN/AMPICILLIN TRIHYDRATE
PRINIVIL, LISINAPRIL
PRINZIDE, HYDROCHLOROTHIAZIDE
PROAIR HFA, ALBUTEROL SULFATE
PROAMATINE, MIDODRINE HYDROCHLORIDE
PROBALAN, PROBENECID
PRO-BANTHINE, PROPANTHELINE BROMIDE
PROBENECID AND COLCHICINE, COLCHICINE
PROBENECID, PROBENECID
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROCAINE HYDROCHLORIDE, PROCAINE HYDROCHLORIDE
PROCALAMINE, AMINO ACIDS
PROCANBID, PROCAINAMIDE HYDROCHLORIDE
PROCARDIA XL, NIFEDIPINE
PROCARDIA, NIFEDIPINE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROCHLORPERAZINE, PROCHLORPERAZINE
PROCHLORPERAZINE, PROCHLORPERAZINE EDISYLATE
PROCTOFOAM HC, HYDROCORTISONE ACETATE
PROFEN, IBUPROFEN (OTC)
PROFERDEX, IRON DEXTRAN
PROGESTERONE, PROGESTERONE
PROGLYCEM, DIAZOXIDE
PROGRAF, TACROLIMUS
PROHANCE MULTIPACK, GADOTERIDOL
PROHANCE, GADOTERIDOL
PROLIXIN DECANOATE, FLUPHENAZINE DECANOATE
PROLIXIN, FLUPHENAZINE HYDROCHLORIDE
PROLOPRIM, TRIMETHOPRIM
PROMETH VC PLAIN, PHENYLEPHRINE HYDROCHLORIDE
PROMETH VC W/ CODEINE, CODEINE PHOSPHATE
PROMETH W/ CODEINE, CODEINE PHOSPHATE
PROMETH W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
PROMETHACON, PROMETHAZINE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN HYDROBROMIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
PROMETHAZINE VC PLAIN, PHENYLEPHRINE HYDROCHLORIDE
PROMETHAZINE VC W/ CODEINE, CODEINE PHOSPHATE
PROMETHAZINE W/ CODEINE, CODEINE PHOSPHATE
PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
PROMETRIUM, PROGESTERONE
PROMPT PHENYTOIN SODIUM, PHENYTOIN SODIUM
PRONESTYL, PROCAINAMIDE HYDROCHLORIDE
PRONESTYL-SR, PROCAINAMIDE HYDROCHLORIDE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE
PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
PROPECIA, FINASTERIDE
PROPINE, DIPIVEFRIN HYDROCHLORIDE
PROPOFOL, PROPOFOL
PROPOXYPHENE COMPOUND 65, ASPIRIN
PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPOXYPHENE HYDROCHLORIDE, PROPOXYPHENE HYDROCHLORIDE
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

APPENDIX A - PRODUCT NAME INDEX

A - 45

**** P ****

PROPRANOLOL HYDROCHLORIDE INTENSOL, PROPRANOLOL HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
PROPRANOLOL, PROPRANOLOL HYDROCHLORIDE
PROPYLTHIOURACIL, PROPYLTHIOURACIL
PROQUIN XR, CIPROFLOXACIN HYDROCHLORIDE
PROSCAR, FINASTERIDE
PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS
PROSOM, ESTAZOLAM
PROSTEP, NICOTINE (OTC)
PROSTIN E2, DINOPROSTONE
PROSTIN VR PEDIATRIC, ALPROSTADIL
PROTAMINE SULFATE, PROTAMINE SULFATE
PROTONIX IV, PANTOPRAZOLE SODIUM
PROTONIX, PANTOPRAZOLE SODIUM
PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE
PROTOPIC, TACROLIMUS
PROVENTIL, ALBUTEROL
PROVENTIL-HFA, ALBUTEROL SULFATE
PROVERA, MEDROXYPROGESTERONE ACETATE
PROVIGIL, MODAFINIL
PROVOCHOLINE, METHACHOLINE CHLORIDE
PROZAC WEEKLY, FLUOXETINE HYDROCHLORIDE
PROZAC, FLUOXETINE HYDROCHLORIDE
PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
PSORCON, DIFLORASONE DIACETATE
PULMICORT RESPULES, BUDESONIDE
PULMICORT, BUDESONIDE
PULMOLITE, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
PURINETHOL, MERCAPTOPURINE
PYRAZINAMIDE, PYRAZINAMIDE
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
PYTEST KIT, UREA, C-14
PYTEST, UREA, C-14

**** Q ****

QUADRAMET, SAMARIUM SM 153 LEXIDRONAM PENTASODIUM
QUELICIN PRESERVATIVE FREE, SUCCINYLCHOLINE CHLORIDE
QUELICIN, SUCCINYLCHOLINE CHLORIDE
QUESTRAN LIGHT, CHOLESTYRAMINE
QUESTRAN, CHOLESTYRAMINE
QUIBRON-T, THEOPHYLLINE
QUIBRON-T/SR, THEOPHYLLINE
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
QUINAPRIL, QUINAPRIL HYDROCHLORIDE
QUINARETIC, HYDROCHLOROTHIAZIDE
QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
QUINIDINE SULFATE, QUINIDINE SULFATE
QUININE SULFATE, QUININE SULFATE
QUIXIN, LEVOFLOXACIN
QVAR 40, BECLOMETHASONE DIPROPIONATE
QVAR 80, BECLOMETHASONE DIPROPIONATE

**** R ****

RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)
RAMIPRIL, RAMIPRIL
RANICLOR, CEFACLOX
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE, RANITIDINE HYDROCHLORIDE
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
RAPAMUNE, SIROLIMUS

APPENDIX A - PRODUCT NAME INDEX

A - 46

**** R ****

RAZADYNE ER, GALANTAMINE HYDROBROMIDE
RAZADYNE, GALANTAMINE HYDROBROMIDE
REBETOL, RIBAVIRIN
REFLUDAN, LEPIRUDIN RECOMBINANT
REGITINE, PHENTOLAMINE MESYLATE
REGLAN ODT, METOCLOPRAMIDE HYDROCHLORIDE
REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
REGONOL, PYRIDOSTIGMINE BROMIDE
REGULAR ILETIN II (PORK), INSULIN PURIFIED PORK (OTC)
RELAFEN, NABUMETONE
RELENZA, ZANAMIVIR
RELPAK, ELETRIPTAN HYDROBROMIDE
REMERON SOLTAB, MIRTAZAPINE
REMERON, MIRTAZAPINE
REMODULIN, TREPROSTINIL SODIUM
RENACIDIN, CITRIC ACID
RENAGEL, SEVELAMER HYDROCHLORIDE
RENAMIN W/O ELECTROLYTES, AMINO ACIDS
RENESE, POLYTHIAZIDE
RENESE-R, POLYTHIAZIDE
RENO-30, DIATRIZOATE MEGLUMINE
RENO-60, DIATRIZOATE MEGLUMINE
RENOCAL-76, DIATRIZOATE MEGLUMINE
RENO-DIP, DIATRIZOATE MEGLUMINE
RENOGRAFIN-60, DIATRIZOATE MEGLUMINE
RENOGRAFIN-76, DIATRIZOATE MEGLUMINE
RENOVA, TRETINOIN
REPRONEX, LUTEINIZING HORMONE
REQUIP, ROPINIROLE HYDROCHLORIDE
RESCRIPTOR, DELAVIRDINE MESYLATE
RESCULA, UNOPROSTONE ISOPROPYL
RESECTISOL IN PLASTIC CONTAINER, MANNITOL
RESERPINE, RESERPINE
RESTASIS, CYCLOSPORINE
RESTORIL, TEMAZEPAM
RETIN-A MICRO, TRETINOIN
RETIN-A, TRETINOIN
RETISERT, FLUOCINOLONE ACETONIDE
RETROVIR, ZIDOVUDINE
REVATIO, SILDENAFIL CITRATE
REVEX, NALMEFENE HYDROCHLORIDE
REV-EYES, DAPIPRAZOLE HYDROCHLORIDE
REVIA, NALTREXONE HYDROCHLORIDE
REVLIMID, LENALIDOMIDE
REYATAZ, ATAZANAVIR SULFATE
R-GENE 10, ARGININE HYDROCHLORIDE
RHINOCORT, BUDESONIDE
RIBASPHERE, RIBAVIRIN
RIBAVIRIN, RIBAVIRIN
RID MOUSSE, PIPERONYL BUTOXIDE (OTC)
RIDAURA, AURANOFIN
RIFADIN, RIFAMPIN
RIFAMATE, ISONIAZID
RIFAMPIN AND ISONIAZID, ISONIAZID
RIFAMPIN, RIFAMPIN
RIFATER, ISONIAZID
RILUTEK, RILUZOLE
RILUZOLE, RILUZOLE
RIMACTANE, RIFAMPIN
RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
RIMSO-50, DIMETHYL SULFOXIDE
RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
RIOMET, METFORMIN HYDROCHLORIDE
RISPERDAL CONSTA, RISPERIDONE

APPENDIX A - PRODUCT NAME INDEX

A - 47

**** R ****

RISPERDAL, RISPERIDONE
RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
RITALIN, METHYLPHENIDATE HYDROCHLORIDE
RITALIN-SR, METHYLPHENIDATE HYDROCHLORIDE
RITODRINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, RITODRINE HYDROCHLORIDE
RITODRINE HYDROCHLORIDE, RITODRINE HYDROCHLORIDE
ROBAXIN, METHOCARBAMOL
ROBAXIN-750, METHOCARBAMOL
ROBINUL FORTE, GLYCOPYRROLATE
ROBINUL, GLYCOPYRROLATE
ROCALTROL, CALCITRIOL
ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
ROCEPHIN, CEFTRIAXONE SODIUM
ROGAINE (FOR MEN), MINOXIDIL (OTC)
ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
ROMAZICON, FLUMAZENIL
ROWASA, MESALAMINE
ROXICET 5/500, ACETAMINOPHEN
ROXICET, ACETAMINOPHEN
ROXICODONE, OXYCODONE HYDROCHLORIDE
ROXILOX, ACETAMINOPHEN
ROZEREM, RAMELTEON
RUBEX, DOXORUBICIN HYDROCHLORIDE
RUBRAMIN PC, CYANOCOBALAMIN
RUBRATOPE-57 KIT, COBALT CHLORIDE, CO-57
RUBRATOPE-57, CYANOCOBALAMIN, CO-57
RYTHMOL SR, PROPAFENONE HYDROCHLORIDE
RYTHMOL, PROPAFENONE HYDROCHLORIDE

**** S ****

SAIZEN, SOMATROPIN RECOMBINANT
SALAGEN, PILOCARPINE HYDROCHLORIDE
SALURON, HYDROFLUMETHIAZIDE
SANCTURA, TROSPIMUM CHLORIDE
SANDIMMUNE, CYCLOSPORINE
SANDOSTATIN LAR, OCTREOTIDE ACETATE
SANDOSTATIN, OCTREOTIDE ACETATE
SARAFEM, FLUOXETINE HYDROCHLORIDE
SCANDONEST L, LEVONORDEFIN
SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
SCLEROSOL, TALC
SEASONALE, ETHINYL ESTRADIOL
SECOBARBITAL SODIUM, SECOBARBITAL SODIUM
SECONAL SODIUM, SECOBARBITAL SODIUM
SECREMAX, SECRETIN SYNTHETIC PORCINE
SECTRAL, ACEBUTOLOL HYDROCHLORIDE
SEDAPAP, ACETAMINOPHEN
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SELENIUM SULFIDE, SELENIUM SULFIDE
SELSUN, SELENIUM SULFIDE
SEMPREX-D, ACRIVASTINE
SENSIPAR, CINACALCET HYDROCHLORIDE
SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
SEPTOCAINE, ARTICAIN HYDROCHLORIDE
SEPTRA DS, SULFAMETHOXAZOLE
SEPTRA GRAPE, SULFAMETHOXAZOLE
SEPTRA, SULFAMETHOXAZOLE
SEREVENT, SALMETEROL XINAFOATE
SEROMYCIN, CYCLOSERINE
SEROPHENE, CLOMIPHENE CITRATE
SEROQUEL, QUETIAPINE FUMARATE
SEROSTIM, SOMATROPIN RECOMBINANT

APPENDIX A - PRODUCT NAME INDEX

A - 48

**** S ****

SERPALAN, RESERPINE
SEVOFLURANE, SEVOFLURANE
SHADE UVAGUARD, AVOBENZONE (OTC)
SILVADENE, SILVER SULFADIAZINE
SINE-AID IB, IBUPROFEN (OTC)
SINEMET CR, CARBIDOPA
SINEMET, CARBIDOPA
SINEQUAN, DOXEPIN HYDROCHLORIDE
SINGULAIR, MONTELUKAST SODIUM
SINOGRAPHIN, DIATRIZOATE MEGLUMINE
SKELAXIN, METAXALONE
SKELID, TILUDRONATE DISODIUM
SODIUM ACETATE IN PLASTIC CONTAINER, SODIUM ACETATE, ANHYDROUS
SODIUM BICARBONATE, SODIUM BICARBONATE
SODIUM BUTABARBITAL, BUTABARBITAL SODIUM
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.224%, POTASSIUM CHLORIDE
SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHROMATE CR 51, SODIUM CHROMATE, CR-51
SODIUM IODIDE I 123, SODIUM IODIDE, I-123
SODIUM IODIDE I 131, KIT, SODIUM IODIDE, I-131
SODIUM IODIDE I 131, SODIUM IODIDE, I-131
SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
SODIUM P.A.S., AMINOSALICYLATE SODIUM
SODIUM PHOSPHATE P 32, SODIUM PHOSPHATE, P-32
SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
SOLAGE, MEQUINOL
SOLARAZE, DICLOFENAC SODIUM
SOLTAMOX, TAMOXIFEN CITRATE
SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
SOMA COMPOUND W/ CODEINE, ASPIRIN
SOMA COMPOUND, ASPIRIN
SOMA, CARISOPRODOL
SOMAVERT, PEGVISOMANT
SONATA, ZALEPLON
SONAZINE, CHLORPROMAZINE HYDROCHLORIDE
SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
SORIATANE, ACITRETIN
SORINE, SOTALOL HYDROCHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SOTRADECOL, SODIUM TETRADECYL SULFATE
SOTRET, ISOTRETINOIN
SOYACAL 10%, SOYBEAN OIL
SOYACAL 20%, SOYBEAN OIL
SPECTAZOLE, ECONAZOLE NITRATE
SPECTRACEF, CEFDITOREN PIVOXIL
SPECTROBID, BACAMPICILLIN HYDROCHLORIDE
SPIRIVA, TIOTROPIUM BROMIDE MONOHYDRATE
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
SPIRONOLACTONE, SPIRONOLACTONE
SPORANOX, ITRACONAZOLE

APPENDIX A - PRODUCT NAME INDEX

A - 49

**** S ****

SPRINTEC, ETHINYL ESTRADIOL
SPS, SODIUM POLYSTYRENE SULFONATE
SSD AF, SILVER SULFADIAZINE
SSD, SILVER SULFADIAZINE
STADOL PRESERVATIVE FREE, BUTORPHANOL TARTRATE
STADOL, BUTORPHANOL TARTRATE
STALEVO 100, CARBIDOPA
STALEVO 150, CARBIDOPA
STALEVO 50, CARBIDOPA
STARLIX, NATEGLINIDE
STELAZINE, TRIFLUOPERAZINE HYDROCHLORIDE
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, WATER FOR INJECTION, STERILE
STERILE WATER IN PLASTIC CONTAINER, WATER FOR IRRIGATION, STERILE
STERILE WATER, WATER FOR IRRIGATION, STERILE
STERI-STAT, CHLORHEXIDINE GLUCONATE (OTC)
STIE-CORT, HYDROCORTISONE
STIMATE, DESMOPRESSIN ACETATE
STRATTERA, ATOMOXETINE HYDROCHLORIDE
STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
STRIANT, TESTOSTERONE
STRIFON FORTE DSC, CHLORZOXAZONE
STROMECTOL, IVERMECTIN
STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE, SR-89
SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE
SUBOXONE, BUPRENORPHINE HYDROCHLORIDE
SUBUTEX, BUPRENORPHINE HYDROCHLORIDE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
SUCRAID, SACROSIDASE
SUCRALFATE, SUCRALFATE
SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
SUFENTA, SUFENTANIL CITRATE
SUFENTANIL CITRATE, SUFENTANIL CITRATE
SULAR, NISOLDIPINE
SULF-10, SULFACETAMIDE SODIUM
SULFACEL-15, SULFACETAMIDE SODIUM
SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
SULFADIAZINE, SULFADIAZINE
SULFAMETHOPRIM, SULFAMETHOXAZOLE
SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH, SULFAMETHOXAZOLE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SULFAMYLON, MAFENIDE ACETATE
SULFASALAZINE, SULFASALAZINE
SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
SULFATRIM, SULFAMETHOXAZOLE
SULFENTANIL CITRATE, SUFENTANIL CITRATE
SULFINPYRAZONE, SULFINPYRAZONE
SULFISOXAZOLE, SULFISOXAZOLE
SULINDAC, SULINDAC
SUMYCIN, TETRACYCLINE HYDROCHLORIDE
SUPRANE, DESFLURANE
SUPRAX, CEFIXIME
SURMONTIL, TRIMIPRAMINE MALEATE
SURVANTA, BERACTANT
SUSTIVA, EFAVIRENZ
SYMBYAX, FLUOXETINE HYDROCHLORIDE
SYMLIN, PRAMLINTIDE ACETATE
SYMMETREL, AMANTADINE HYDROCHLORIDE
SYNACORT, HYDROCORTISONE
SYNALAR, FLUOCINOLONE ACETONIDE
SYNALGOS-DC, ASPIRIN
SYNAREL, NAFARELIN ACETATE
SYNERA, LIDOCAINE

APPENDIX A - PRODUCT NAME INDEX

A - 50

**** S ****

SYNERCID, DALFOPRISTIN
 SYNTHROID, LEVOTHYROXINE SODIUM
 SYPRINE, TRIENTINE HYDROCHLORIDE

**** T ****

TAB-PROFEN, IBUPROFEN (OTC)
 TAGAMET HB, CIMETIDINE (OTC)
 TAGAMET, CIMETIDINE
 TALACEN, ACETAMINOPHEN
 TALC, TALC
 TALWIN NX, NALOXONE HYDROCHLORIDE
 TALWIN, PENTAZOCINE LACTATE
 TAMBOCOR, FLECAINIDE ACETATE
 TAMIFLU, OSELTAMIVIR PHOSPHATE
 TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 TAO, TROLEANDOMYCIN
 TARCEVA, ERLOTINIB HYDROCHLORIDE
 TARGRETIN, BEXAROTENE
 TARKA, TRANDOLAPRIL
 TASMAR, TOLCAPONE
 TAVIST ALLERGY/SINUS/HEADACHE, ACETAMINOPHEN (OTC)
 TAVIST-1, CLEMASTINE FUMARATE (OTC)
 TAXOL, PACLITAXEL
 TAXOTERE, DOCETAXEL
 TAZICEF, CEFTAZIDIME
 TAZORAC, TAZAROTENE
 TAZTIA XT, DILTIAZEM HYDROCHLORIDE
 TECHNECOLL, TECHNETIUM TC-99M SULFUR COLLOID KIT
 TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
 TECHNISCAN GLUCEPTATE, TECHNETIUM TC-99M GLUCEPTATE KIT
 TECHNISCAN MAA, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
 TECHNISCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
 TECHNISCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
 TECHNISCAN, TECHNETIUM TC-99M OXIDRONATE KIT
 TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
 TECHNETIUM TC 99M MPI MDP, TECHNETIUM TC-99M MEDRONATE KIT
 TECHNETIUM TC-99M PENTETATE KIT, TECHNETIUM TC-99M PENTETATE KIT
 TECZEM, DILTIAZEM MALATE
 TEGRETOL, CARBAMAZEPINE
 TEGRETOL-XR, CARBAMAZEPINE
 TEMAZEPAM, TEMAZEPAM
 TEMODAR, TEMOZOLOMIDE
 TEMOVATE E, CLOBETASOL PROPIONATE
 TEMOVATE, CLOBETASOL PROPIONATE
 TENCON, ACETAMINOPHEN
 TENEX, GUANFACINE HYDROCHLORIDE
 TENORETIC 100, ATENOLOL
 TENORETIC 50, ATENOLOL
 TENORMIN, ATENOLOL
 TENSILON PRESERVATIVE FREE, EDROPHONIUM CHLORIDE
 TENSILON, EDROPHONIUM CHLORIDE
 TENUATE DOSPAN, DIETHYLPROPION HYDROCHLORIDE
 TENUATE, DIETHYLPROPION HYDROCHLORIDE
 TEQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER, GATIFLOXACIN
 TEQUIN, GATIFLOXACIN
 TERAZOL 3, TERCONAZOLE
 TERAZOL 7, TERCONAZOLE
 TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 TERCONAZOLE, TERCONAZOLE
 TERIL, CARBAMAZEPINE
 TERRA-CORTRIL, HYDROCORTISONE ACETATE
 TERRAMYCIN W/ POLYMYXIN B SULFATE, OXYTETRACYCLINE HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 51

**** T ****

TERRAMYCIN, LIDOCAINE HYDROCHLORIDE
TERRAMYCIN, OXYTETRACYCLINE HYDROCHLORIDE
TESLAC, TESTOLACTONE
TESSALON, BENZONATATE
TESTIM, TESTOSTERONE
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
TESTOSTERONE PROPIONATE, TESTOSTERONE PROPIONATE
TESTOSTERONE, TESTOSTERONE
TESTRED, METHYLTESTOSTERONE
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
TETREX, TETRACYCLINE PHOSPHATE COMPLEX
TEVETEN HCT, EPROSARTAN MESYLATE
TEVETEN, EPROSARTAN MESYLATE
TEV-TROPIN, SOMATROPIN RECOMBINANT
TEXACORT, HYDROCORTISONE
THALITONE, CHLORTHALIDONE
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE, TL-201
THALOMID, THALIDOMIDE
THAM, TROMETHAMINE
THEO-24, THEOPHYLLINE
THEOCHRON, THEOPHYLLINE
THEOLAIR, THEOPHYLLINE
THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THEOPHYLLINE, THEOPHYLLINE
THERMAZENE, SILVER SULFADIAZINE
THEROXIDIL, MINOXIDIL (OTC)
THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
THIOGUANINE, THIOGUANINE
THIORIDAZINE HYDROCHLORIDE INTENSOL, THIORIDAZINE HYDROCHLORIDE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
THIOTEPA, THIOTEPA
THIOTHIXENE HYDROCHLORIDE, THIOTHIXENE HYDROCHLORIDE
THIOTHIXENE, THIOTHIXENE
THORAZINE, CHLORPROMAZINE
THORAZINE, CHLORPROMAZINE HYDROCHLORIDE
THYROGEN, THYROTROPIN ALFA
THYROLAR-0.25, LIOETHYRONINE SODIUM
THYROLAR-0.5, LIOETHYRONINE SODIUM
THYROLAR-1, LIOETHYRONINE SODIUM
THYROLAR-2, LIOETHYRONINE SODIUM
THYROLAR-3, LIOETHYRONINE SODIUM
THYROSAFE, POTASSIUM IODIDE (OTC)
THYROSHIELD, POTASSIUM IODIDE (OTC)
TIAZAC, DILTIAZEM HYDROCHLORIDE
TICAR, TICARCILLIN DISODIUM
TICLID, TICLOPIDINE HYDROCHLORIDE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
TIKOSYN, DOFETILIDE
TILADE, NEDOCROMIL SODIUM
TIMENTIN IN PLASTIC CONTAINER, CLAVULANATE POTASSIUM
TIMENTIN, CLAVULANATE POTASSIUM
TIMOLIDE 10-25, HYDROCHLOROTHIAZIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
TIMOPTIC, TIMOLOL MALEATE
TIMOPTIC-XE, TIMOLOL MALEATE
TINDAMAX, TINIDAZOLE

APPENDIX A - PRODUCT NAME INDEX

A - 52

**** T ****

TIOCONAZOLE, TIOCONAZOLE (OTC)
TIOPRONIN, TIOPRONIN
TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE
TIS-U-SOL, MAGNESIUM SULFATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOBI, TOBRAMYCIN
TOBRADEX, DEXAMETHASONE
TOBRAMYCIN (PHARMACY BULK), TOBRAMYCIN SULFATE
TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TOBRAMYCIN, TOBRAMYCIN
TOBRAMYCIN, TOBRAMYCIN SULFATE
TOBRASONE, FLUOROMETHOLONE ACETATE
TOBREX, TOBRAMYCIN
TOFRANIL, IMIPRAMINE HYDROCHLORIDE
TOFRANIL-PM, IMIPRAMINE PAMOATE
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TOLECTIN 600, TOLMETIN SODIUM
TOLECTIN DS, TOLMETIN SODIUM
TOLECTIN, TOLMETIN SODIUM
TOLINASE, TOLAZAMIDE
TOLMETIN SODIUM, TOLMETIN SODIUM
TOPAMAX SPRINKLE, TOPIRAMATE
TOPAMAX, TOPIRAMATE
TOPICORT LP, DESOXIMETASONE
TOPICORT, DESOXIMETASONE
TOPOSAR, ETOPOSIDE
TOPROL-XL, METOPROLOL SUCCINATE
TORADOL, KETOROLAC TROMETHAMINE
TORECAN, THIETHYLPERAZINE MALEATE
TORSEMIDE, TORSEMIDE
T-PHYL, THEOPHYLLINE
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
TRACLEER, BOSENTAN
TRACRIUM PRESERVATIVE FREE, ATRACURIUM BESYLATE
TRACRIUM, ATRACURIUM BESYLATE
TRAMADOL HCL, TRAMADOL HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRANDATE, LABETALOL HYDROCHLORIDE
TRANMEP, MEPROBAMATE
TRANSDERM SCOP, SCOPOLAMINE
TRANXENE SD, CLORAZEPATE DIPOTASSIUM
TRANXENE, CLORAZEPATE DIPOTASSIUM
TRASYLOL, APROTININ BOVINE
TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 10% W/O ELECTROLYTES, AMINO ACIDS
TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 3.5% W/ ELECTROLYTES, AMINO ACIDS
TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 5.5% W/ ELECTROLYTES, AMINO ACIDS
TRAVASOL 5.5% W/O ELECTROLYTES, AMINO ACIDS
TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
TRAVASOL 8.5% W/ ELECTROLYTES, AMINO ACIDS
TRAVASOL 8.5% W/O ELECTROLYTES, AMINO ACIDS
TRAVATAN, TRAVOPROST
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRECATOR-SC, ETHIONAMIDE
TRELSTAR DEPOT, TRIPTORELIN PAMOATE
TRELSTAR, TRIPTORELIN PAMOATE

APPENDIX A - PRODUCT NAME INDEX

A - 53

**** T ****

TRENTAL, PENTOXIFYLLINE
TRETINOIN, TRETINOIN
Trexall, METHOTREXATE SODIUM
TRIACET, TRIAMCINOLONE ACETONIDE
TRIACIN-C, CODEINE PHOSPHATE
TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRIAZOLAM, TRIAZOLAM
TRICOR, FENOFIBRATE
TRIDERM, TRIAMCINOLONE ACETONIDE
TRIDESILON, DESONIDE
TRIDIONE, TRIMETHADIONE
TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
TRIFLURIDINE, TRIFLURIDINE
TRIGLIDE, FENOFIBRATE
TRIHXYPHENIDYL HYDROCHLORIDE, TRIHXYPHENIDYL HYDROCHLORIDE
TRILEPTAL, OXCARBAZEPINE
TRI-LUMA, FLUOCINOLONE ACETONIDE
TRILYTE, POLYETHYLENE GLYCOL 3350
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
TRIMETHOPRIM, TRIMETHOPRIM
TRIMOX, AMOXICILLIN
TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
TRIOSTAT, LIOTHYRONINE SODIUM
TRIPHASIL-28, ETHINYL ESTRADIOL
TRI-PREVIFEM, ETHINYL ESTRADIOL
TRIPROlidine HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
TRISENOX, ARSENIC TRIOXIDE
TRI-SPRINTEC, ETHINYL ESTRADIOL
TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
TRIVORA-28, ETHINYL ESTRADIOL
TRIZIVIR, ABACAVIR SULFATE
TROBICIN, SPECTINOMYCIN HYDROCHLORIDE
TROPAMINE 10%, AMINO ACIDS
TROPAMINE, AMINO ACIDS
TROPICACYL, TROPICAMIDE
TROPICAMIDE, TROPICAMIDE
TRUPHYLLINE, AMINOPHYLLINE
TRUSOPT, DORZOLAMIDE HYDROCHLORIDE
TRUVADA, EMTRICITABINE
TUSSIGON, HOMATROPINE METHYLBROMIDE
TUSSIONEX, CHLORPHENIRAMINE POLISTIREX
TWINJECT 0.3, EPINEPHRINE
TWINJECT, EPINEPHRINE
TYGACIL, TIGECYCLINE
TYLENOL (CAPLET), ACETAMINOPHEN (OTC)
TYLENOL (GELTAB), ACETAMINOPHEN (OTC)
TYLENOL W/ CODEINE NO. 3, ACETAMINOPHEN
TYLOX, ACETAMINOPHEN
TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

**** U ****

U-CORT, HYDROCORTISONE ACETATE
ULTANE, SEVOFLURANE
ULTIVA, REMIFENTANIL HYDROCHLORIDE
ULTRACET, ACETAMINOPHEN
ULTRAM, TRAMADOL HYDROCHLORIDE
ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
ULTRAVATE, HALOBETASOL PROPIONATE
ULTRAVIST (PHARMACY BULK), IOPROMIDE

APPENDIX A - PRODUCT NAME INDEX

A - 54

**** U ****

ULTRAVIST 150, IOPROMIDE
 ULTRAVIST 240, IOPROMIDE
 ULTRAVIST 300, IOPROMIDE
 ULTRAVIST 370, IOPROMIDE
 UNASYN, AMPICILLIN SODIUM
 UNIPHYL, THEOPHYLLINE
 UNIRETIC, HYDROCHLOROTHIAZIDE
 UNISOM, DOXYLAMINE SUCCINATE (OTC)
 UNITHROID, LEVOTHYROXINE SODIUM
 UNIVASC, MOEXIPRIL HYDROCHLORIDE
 UREAPHIL, UREA
 URECHOLINE, BETHANECHOL CHLORIDE
 UREX, METHENAMINE HIPPURATE
 URISPAS, FLAVOXATE HYDROCHLORIDE
 UROCIT-K, POTASSIUM CITRATE
 UROLOGIC G IN PLASTIC CONTAINER, CITRIC ACID
 UROXATRAL, ALFUZOSIN HYDROCHLORIDE
 URSO 250, URSODIOL
 URSO FORTE, URSODIOL
 URSODIOL, URSODIOL
 UVADEX, METHOXSALEN

**** V ****

VAGIFEM, ESTRADIOL
 VAGISTAT-1, TIOCONAZOLE (OTC)
 VALCYTE, VALGANCICLOVIR HYDROCHLORIDE
 VALIUM, DIAZEPAM
 VALNAC, BETAMETHASONE VALERATE
 VALPROATE SODIUM, VALPROATE SODIUM
 VALPROIC ACID, VALPROIC ACID
 VALSTAR PRESERVATIVE FREE, VALRUBICIN
 VALTREX, VALACYCLOVIR HYDROCHLORIDE
 VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
 VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VANDAZOLE, METRONIDAZOLE
 VANIQA, EFLORNITHINE HYDROCHLORIDE
 VANOS, FLUOCINONIDE
 VANTAS, HISTRELIN ACETATE
 VANTIN, CEFPODOXIME PROXETIL
 VAPRISOL, CONIVAPTAN HYDROCHLORIDE
 VASERETIC, ENALAPRIL MALEATE
 VASOCIDIN, PREDNISOLONE SODIUM PHOSPHATE
 VASOCON, NAPHAZOLINE HYDROCHLORIDE
 VASOCON-A, ANTAZOLINE PHOSPHATE (OTC)
 VASOTEC, ENALAPRIL MALEATE
 VASOTEC, ENALAPRILAT
 VECURONIUM BROMIDE, VECURONIUM BROMIDE
 VEETIDS, PENICILLIN V POTASSIUM
 VELCADE, BORTEZOMIB
 VELIVET, DESOGESTREL
 VELOSEF '125', CEPHRADINE
 VELOSEF '250', CEPHRADINE
 VELOSEF, CEPHRADINE
 VELOSULIN BR, INSULIN RECOMBINANT HUMAN (OTC)
 VENOFER, IRON SUCROSE
 VENTAVIS, ILOPROST
 VENTOLIN HFA, ALBUTEROL SULFATE
 VENTOLIN, ALBUTEROL
 VEPESID, ETOPOSIDE
 VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
 VERELAN PM, VERAPAMIL HYDROCHLORIDE
 VERELAN, VERAPAMIL HYDROCHLORIDE

APPENDIX A - PRODUCT NAME INDEX

A - 55

**** V ****

VESANOID, TRETINOIN
VESICARE, SOLIFENACIN SUCCINATE
VEXOL, RIMEXOLONE
VFEND, VORICONAZOLE
VIADUR, LEUPROLIDE ACETATE
VIAGRA, SILDENAFIL CITRATE
VIBISONE, CYANOCOBALAMIN
VIBRAMYCIN, DOXYCYCLINE
VIBRAMYCIN, DOXYCYCLINE CALCIUM
VIBRAMYCIN, DOXYCYCLINE HYCLATE
VIBRA-TABS, DOXYCYCLINE HYCLATE
VICODIN ES, ACETAMINOPHEN
VICODIN HP, ACETAMINOPHEN
VICODIN, ACETAMINOPHEN
VICOPROFEN, HYDROCODONE BITARTRATE
VIDAZA, AZACITIDINE
VIDEX EC, DIDANOSINE
VIDEX, DIDANOSINE
VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
VINBLASTINE SULFATE, VINBLASTINE SULFATE
VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
VINCRISTINE SULFATE, VINCRISTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
VIRACEPT, NELFINAVIR MESYLATE
VIRAMUNE, NEVIRAPINE
VIRAZOLE, RIBAVIRIN
VIREAD, TENOFOVIR DISOPROXIL FUMARATE
VIRILON, METHYLTESTOSTERONE
VIROPTIC, TRIFLURIDINE
VISICOL, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
VISINE-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
VISIONBLUE, TRYPAN BLUE
VISIPAQUE 270, IODIXANOL
VISIPAQUE 320, IODIXANOL
VISKEN, PINDOLOL
VISTARIL, HYDROXYZINE HYDROCHLORIDE
VISTARIL, HYDROXYZINE PAMOATE
VISTIDE, CIDOFOVIR
VISUDYNE, VERTEPORFIN
VITAMIN A PALMITATE, VITAMIN A PALMITATE
VITAMIN D, ERGOCALCIFEROL
VITAMIN K1, PHYTONADIONE
VITRASE, HYALURONIDASE
VITRASERT, GANCICLOVIR
VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE
VIVELLE, ESTRADIOL
VIVELLE-DOT, ESTRADIOL
VOLTAREN, DICLOFENAC SODIUM
VOLTAREN-XR, DICLOFENAC SODIUM
VOSOL HC, ACETIC ACID, GLACIAL
VOSPIRE ER, ALBUTEROL SULFATE
VUMON, TENIPOSIDE
VYTORIN, EZETIMIBE

**** W ****

WARFARIN SODIUM, WARFARIN SODIUM
WELCHOL, COLESEVELAM HYDROCHLORIDE
WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
WELLBUTRIN XL, BUPROPION HYDROCHLORIDE
WELLBUTRIN, BUPROPION HYDROCHLORIDE
WESTCORT, HYDROCORTISONE VALERATE
WYGESIC, ACETAMINOPHEN

APPENDIX A - PRODUCT NAME INDEX

A - 56

**** X ****

XALATAN, LATANOPROST
 XANAX XR, ALPRAZOLAM
 XANAX, ALPRAZOLAM
 XELODA, CAPECITABINE
 XENICAL, ORLISTAT
 XENON XE 133, XENON, XE-133
 XIBROM, BROMFENAC SODIUM
 XIFAXAN, RIFAXIMIN
 XOPENEX HFA, LEVALBUTEROL TARTRATE
 XOPENEX, LEVALBUTEROL HYDROCHLORIDE
 X-TROZINE L.A., PHENDIMETRAZINE TARTRATE
 X-TROZINE, PHENDIMETRAZINE TARTRATE
 XYLOCAINE 4% PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 XYLOCAINE, LIDOCAINE HYDROCHLORIDE
 XYREM, SODIUM OXYBATE

**** Y ****

YASMIN, DROSPIRENONE

**** Z ****

ZADITOR, KETOTIFEN FUMARATE
 ZANAFLEX, TIZANIDINE HYDROCHLORIDE
 ZANOSAR, STREPTOZOCIN
 ZANTAC 150, RANITIDINE HYDROCHLORIDE
 ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)
 ZANTAC 25, RANITIDINE HYDROCHLORIDE
 ZANTAC 300, RANITIDINE HYDROCHLORIDE
 ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)
 ZANTAC IN PLASTIC CONTAINER, RANITIDINE HYDROCHLORIDE
 ZANTAC, RANITIDINE HYDROCHLORIDE
 ZARONTIN, ETHOSUXIMIDE
 ZAROXOLYN, METOLAZONE
 ZAVESCA, MIGLUSTAT
 ZEBETA, BISOPROLOL FUMARATE
 ZEGERID, OMEPRAZOLE
 ZELNORM, TEGASEROD MALEATE
 ZEMPLAR, PARICALCITOL
 ZEMURON (P/F), ROCURONIUM BROMIDE
 ZERIT, STAVUDINE
 ZESTORETIC, HYDROCHLOROTHIAZIDE
 ZESTRIL, LISINAPRIL
 ZETIA, EZETIMIBE
 ZIAC, BISOPROLOL FUMARATE
 ZIAGEN, ABACAVIR SULFATE
 ZIDOVUDINE, ZIDOVUDINE
 ZINACEF IN PLASTIC CONTAINER, CEFUROXIME SODIUM
 ZINACEF, CEFUROXIME SODIUM
 ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE
 ZINECARD, DEXRAZOXANE HYDROCHLORIDE
 ZITHROMAX, AZITHROMYCIN
 ZMAX, AZITHROMYCIN
 ZOCOR, SIMVASTATIN
 ZOFRAN IN PLASTIC CONTAINER, ONDANSETRON HYDROCHLORIDE
 ZOFRAN ODT, ONDANSETRON
 ZOFRAN PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
 ZOFRAN, ONDANSETRON HYDROCHLORIDE
 ZOLADEX, GOSERELIN ACETATE
 ZOLOFT, SERTRALINE HYDROCHLORIDE
 ZOMETA, ZOLEDRONIC ACID
 ZOMIG, ZOLMITRIPTAN

APPENDIX A - PRODUCT NAME INDEX

A - 57

**** Z ****

ZOMIG-ZMT, ZOLMITRIPTAN
ZONALON, DOXEPIN HYDROCHLORIDE
ZONEGRAN, ZONISAMIDE
ZONISAMIDE, ZONISAMIDE
ZORBTIVE, SOMATROPIN RECOMBINANT
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
ZOSYN, PIPERACILLIN SODIUM
ZOVIA 1/35E-21, ETHINYL ESTRADIOL
ZOVIA 1/35E-28, ETHINYL ESTRADIOL
ZOVIA 1/50E-21, ETHINYL ESTRADIOL
ZOVIA 1/50E-28, ETHINYL ESTRADIOL
ZOVIRAX, ACYCLOVIR
ZOVIRAX, ACYCLOVIR SODIUM
ZYBAN, BUPROPION HYDROCHLORIDE
ZYFLO, ZILEUTON
ZYLET, LOTEPREDNOL ETABONATE
ZYLOPRIM, ALLOPURINOL
ZYMAR, GATIFLOXACIN
ZYPREXA ZYDIS, OLANZAPINE
ZYPREXA, OLANZAPINE
ZYTEC, CETIRIZINE HYDROCHLORIDE
ZYTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE
ZYVOX, LINEZOLID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** 3 ****

3M

- * 3M HEALTH CARE INC
AVAGARD, ALCOHOL (OTC)
- * 3M PHARMACEUTICALS INC
ALDARA, IMIQUIMOD
CALCIUM DISODIUM VERSENATE, EDETATE CALCIUM DISODIUM
MAXAIR, PIRBUTEROL ACETATE
METROGEL-VAGINAL, METRONIDAZOLE
MINITRAN, NITROGLYCERIN
NORFLEX, ORPHENADRINE CITRATE
NORGESIC FORTE, ASPIRIN
NORGESIC, ASPIRIN
PROVENTIL-HFA, ALBUTEROL SULFATE
QVAR 40, BECLOMETHASONE DIPROPIONATE
QVAR 80, BECLOMETHASONE DIPROPIONATE
TAMBOCOR, FLECAINIDE ACETATE
THEOLAIR, THEOPHYLLINE

AAIPHARMA LLC

- * AAIPHARMA LLC
ACYCLOVIR, ACYCLOVIR
AZASAN, AZATHIOPRINE
BRETHINE, TERBUTALINE SULFATE
ETODOLAC, ETODOLAC
KETOCONAZOLE, KETOCONAZOLE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

AB GENERICS

- * AB GENERICS LP
MORPHINE SULFATE, MORPHINE SULFATE

ABBOTT

- * ABBOTT LABORATORIES
AKINETON, BIPERIDEN HYDROCHLORIDE
BIAXIN XL, CLARITHROMYCIN
BIAXIN, CLARITHROMYCIN
CYCLOSPORINE, CYCLOSPORINE
DEPAKOTE ER, DIVALPROEX SODIUM
DEPAKOTE, DIVALPROEX SODIUM
GENGRAF, CYCLOSPORINE
KALETRA, LOPINAVIR
MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER, MIVACURIUM CHLORIDE
MIVACRON, MIVACURIUM CHLORIDE
NIMBEX PRESERVATIVE FREE, CISATRACURIUM BESYLATE
NIMBEX, CISATRACURIUM BESYLATE
NORVIR, RITONAVIR
OMNICEF, CEFDINIR
SYNTHROID, LEVOTHYROXINE SODIUM
TRICOR, FENOFIBRATE
ULTANE, SEVOFLURANE
ULTIVA, REMIFENTANIL HYDROCHLORIDE
ZEMPLAR, PARICALCITOL
- * ABBOTT LABORATORIES HOSP PRODUCTS DIV
CALCIJEX, CALCITRIOL
- * ABBOTT LABORATORIES PHARMACEUTICAL PRODUCTS DIV
ABBOKINASE, UROKINASE
BIAXIN, CLARITHROMYCIN
DEPACON, VALPROATE SODIUM
DEPAKENE, VALPROIC ACID
DEPAKOTE, DIVALPROEX SODIUM
DILAUDID, HYDROMORPHONE HYDROCHLORIDE
DILAUDID-HP, HYDROMORPHONE HYDROCHLORIDE
E.E.S. 200, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S. 400, ERYTHROMYCIN ETHYLSUCCINATE
E.E.S., ERYTHROMYCIN ETHYLSUCCINATE
E-MYCIN, ERYTHROMYCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** A **

* ABBOTT LABORATORIES PHARMACEUTICAL PRODUCTS DIV

ENDURON, METHYCLOTHIAZIDE
ERYDERM, ERYTHROMYCIN
ERYPED, ERYTHROMYCIN ETHYLSUCCINATE
ERY-TAB, ERYTHROMYCIN
ERYTHROCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN, ERYTHROMYCIN
HYTRIN, TERAZOSIN HYDROCHLORIDE
K-TAB, POTASSIUM CHLORIDE
MAVIK, TRANDOLAPRIL
MERIDIA, SIBUTRAMINE HYDROCHLORIDE
NORVIR, RITONAVIR
ORETIC, HYDROCHLOROTHIAZIDE
PCE, ERYTHROMYCIN
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PROSOM, ESTAZOLAM
TARKA, TRANDOLAPRIL
TRICOR, FENOFIBRATE
TRIDIONE, TRIMETHADIONE
VICODIN ES, ACETAMINOPHEN
VICODIN HP, ACETAMINOPHEN
VICODIN, ACETAMINOPHEN
VICOPROFEN, HYDROCODONE BITARTRATE
ZEMPLAR, PARICALCITOL

ABRIKA PHARMS

* ABRIKA PHARMACEUTICALS LLLP
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
ASPARTATE
ISRADIPINE, ISRADIPINE

ACCORD HLTH

* ACCORD HEALTH CARE INC
METHYLDOPA, METHYLDOPA

ACORDA

* ACORDA THERAPEUTICS INC
ZANAFLEX, TIZANIDINE HYDROCHLORIDE

ACS DOBFAR

* ACS DOBFAR SPA
CEFTAZIDIME, CEFTAZIDIME

ACTELION

* ACTELION LTD
TRACLEER, BOSENTAN

ACTELION PHARMS

* ACTELION PHARMACEUTICALS US INC
ZAVESCA, MIGLUSTAT

ADAMS LABS INC

* ADAMS LABORATORIES INC DBA ADAMS RESPIRATORY THERAPEUTICS
MUCINEX D, GUAIFENESIN (OTC)
MUCINEX DM, DEXTROMETHORPHAN HYDROBROMIDE (OTC)
MUCINEX, GUAIFENESIN (OTC)

ADV CARE

* ADVANCED CARE PRODUCTS DIV ORTHO PHARMACEUTICAL CORP
MONISTAT 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 7, MICONAZOLE NITRATE (OTC)
MONISTAT-3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)

ADV CARE PRODS

* ADVANCED CARE PRODUCTS
MONISTAT 3, MICONAZOLE NITRATE (OTC)
MONISTAT 7, MICONAZOLE NITRATE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

ADV MAGNETICS

* ADVANCED MAGNETICS INC
FERIDEX I.V., FERUMOXIDES
GASTROMARK, FERUMOXASIL

ADVANCIS PHARM

* ADVANCIS PHARMACEUTICAL CORP
KEFLEX, CEPHALEXIN

AEGIS PHARMS

* AEGIS PHARMACEUTICALS INC
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE

AESGEN

* AESGEN INC
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM

AGIS INDS

* AGIS INDUSTRIES 1983 LTD
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
MOMETASONE FUROATE, MOMETASONE FUROATE

AGOURON

* AGOURON PHARMACEUTICALS INC
RESCRIPTOR, DELAVIRDINE MESYLATE
VIRACEPT, NELFINAVIR MESYLATE

AKORN

* AKORN INC
AKBETA, LEVOBUNOLOL HYDROCHLORIDE
AKPENTOLATE, CYCLOPENTOLATE HYDROCHLORIDE
AKPRO, DIPIVEFRIN HYDROCHLORIDE
AKTOB, TOBRAMYCIN
ALFENTA, ALFENTANIL HYDROCHLORIDE
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
BAL, DIMERCAPROL
BETAXOLOL, BETAXOLOL HYDROCHLORIDE
CROMOLYN SODIUM, CROMOLYN SODIUM
ENDOSOL EXTRA, CALCIUM CHLORIDE
ERYTHROMYCIN, ERYTHROMYCIN
GENTAK, GENTAMICIN SULFATE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
IC-GREEN, INDOCYANINE GREEN
INAPSINE, DROPERIDOL
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
NEOMYCIN AND POLYMYXIN B SULFATE AND BACITRACIN ZINC, BACITRACIN ZINC
PAREMYD, HYDROXYAMPHETAMINE HYDROBROMIDE
SUFENTA, SUFENTANIL CITRATE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
TIMOLOL MALEATE, TIMOLOL MALEATE
TROPICACYL, TROPICAMIDE

AKORN MFG

* AKORN MANUFACTURING INC
SUBLIMAZE PRESERVATIVE FREE, FENTANYL CITRATE

AKYMA PHARMS

* AKYMA PHARMACEUTICALS LLC
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE

ALAMO PHARMS

* ALAMO PHARMACEUTICALS LLC
FAZACLO ODT, CLOZAPINE

ALARA PHARM

* ALARA PHARMACEUTICAL CORPORATION
LEVO-T, LEVOTHYROXINE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** A **

ALCON

- * ALCON INC
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
CILOXAN, CIPROFLOXACIN HYDROCHLORIDE
CIPRO HC, CIPROFLOXACIN HYDROCHLORIDE
CIPRODEX, CIPROFLOXACIN
NEVANAC, NEPAFENAC
OFLOXACIN, OFLOXACIN
OLOPATADINE HYDROCHLORIDE, OLOPATADINE HYDROCHLORIDE
SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
TRAVATAN, TRAVOPROST
VIGAMOX, MOXIFLOXACIN HYDROCHLORIDE
- * ALCON LABORATORIES INC
ALCAINE, PROPARACAINE HYDROCHLORIDE
ALOMIDE, LODOXAMIDE TROMETHAMINE
AZOPT, BRINZOLAMIDE
BETADINE, POVIDONE-IODINE
BETOPTIC S, BETAXOLOL HYDROCHLORIDE
BETOPTIC, BETAXOLOL HYDROCHLORIDE
BSS PLUS, CALCIUM CHLORIDE
BSS, CALCIUM CHLORIDE
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
CETAMIDE, SULFACETAMIDE SODIUM
CROMOLYN SODIUM, CROMOLYN SODIUM
CYCLOGYL, CYCLOPENTOLATE HYDROCHLORIDE
CYCLOMYDRIL, CYCLOPENTOLATE HYDROCHLORIDE
DENDRID, IDOXURIDINE
EMADINE, EMEDASTINE DIFUMARATE
FLAREX, FLUOROMETHOLONE ACETATE
IOPIDINE, APRACLOLONIDINE HYDROCHLORIDE
ISOPTO CETAMIDE, SULFACETAMIDE SODIUM
MAXIDEX, DEXAMETHASONE
MAXITROL, DEXAMETHASONE
MIOSTAT, CARBACHOL
MYDRIACYL, TROPICAMIDE
NAPHCN-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
NATACYN, NATAMYCIN
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
PATANOL, OLOPATADINE HYDROCHLORIDE
PILOPINE HS, PILOCARPINE HYDROCHLORIDE
TOBRADEX, DEXAMETHASONE
TOBRASONE, FLUOROMETHOLONE ACETATE
TOBREX, TOBRAMYCIN
TRIFLURIDINE, TRIFLURIDINE
VEXOL, RIMEXOLONE

ALCON UNIVERSAL

- * ALCON UNIVERSAL LTD
CYCLOPENTOLATE HYDROCHLORIDE, CYCLOPENTOLATE HYDROCHLORIDE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
TROPICAMIDE, TROPICAMIDE

ALLEGIANCE HLTHCARE

- * ALLEGIANCE HEALTHCARE CORP
POVIDONE IODINE, POVIDONE-IODINE (OTC)

ALLERGAN

- * ALLERGAN
ACULAR LS, KETOROLAC TROMETHAMINE
ALPHAGAN P, BRIMONIDINE TARTRATE
BLEPH-10, SULFACETAMIDE SODIUM
BLEPH-30, SULFACETAMIDE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

- * ALLERGAN
GENOPTIC, GENTAMICIN SULFATE
- * ALLERGAN INC
ACULAR PRESERVATIVE FREE, KETOROLAC TROMETHAMINE
ACULAR, KETOROLAC TROMETHAMINE
ALBALON, NAPHAZOLINE HYDROCHLORIDE
ALOCRIL, NEDOCROMIL SODIUM
ALPHAGAN P, BRIMONIDINE TARTRATE
AVAGE, TAZAROTENE
AZELEX, AZELAIC ACID
ELESTAT, EPINASTINE HYDROCHLORIDE
ELIMITE, PERMETHRIN
LUMIGAN, BIMATOPROST
OCUFLOX, OFLOXACIN
OPTICROM, CROMOLYN SODIUM
POLYTRIM, POLYMYXIN B SULFATE
RESTASIS, CYCLOSPORINE
TAZORAC, TAZAROTENE
ZYMAR, GATIFLOXACIN
- * ALLERGAN PHARMACEUTICAL
BETAGAN, LEVOBUNOLOL HYDROCHLORIDE
BLEPHAMIDE S.O.P., PREDNISOLONE ACETATE
BLEPHAMIDE, PREDNISOLONE ACETATE
FML FORTE, FLUOROMETHOLONE
FML, FLUOROMETHOLONE
FML-S, FLUOROMETHOLONE
HERPLEX, IDOXURIDINE
HMS, MEDRYSONE
OCUFEN, FLURBIPROFEN SODIUM
OPHTHETIC, PROPARACAINE HYDROCHLORIDE
POLY-PRED, NEOMYCIN SULFATE
PRED FORTE, PREDNISOLONE ACETATE
PRED MILD, PREDNISOLONE ACETATE
PRED-G, GENTAMICIN SULFATE
PROPINE, DIPIVEFRIN HYDROCHLORIDE

ALLERGAN HERBERT

- * ALLERGAN HERBERT DIV ALLERGAN INC
PENECORT, HYDROCORTISONE
- * ALLERGAN HERBERT SKIN CARE DIV ALLERGAN INC
FLUOROPLEX, FLUOROURACIL
PENECORT, HYDROCORTISONE

ALPHA THERA

- * ALPHA THERAPEUTIC CORP
SOYACAL 10%, SOYBEAN OIL
SOYACAL 20%, SOYBEAN OIL

ALPHAPHARM

- * ALPHAPHARM PARTY LTD
ALPRAZOLAM, ALPRAZOLAM
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
BACLOFEN, BACLOFEN
CILOSTAZOL, CILOSTAZOL
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLONAZEPAM, CLONAZEPAM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
GLIPIZIDE, GLIPIZIDE
INDAPAMIDE, INDAPAMIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NAPROXEN, NAPROXEN
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** A **

- * ALPHAPHARM PARTY LTD
 - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - TRIAZOLAM, TRIAZOLAM
 - ZONISAMIDE, ZONISAMIDE
- * ALPHAPHARM PTY LTD
 - ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
 - PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE

ALPHARMA US PHARMS

- * ALPHARMA US PHARMACEUTICALS DIVISION
 - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 - ACETAMINOPHEN, ACETAMINOPHEN (OTC)
 - ACETASOL HC, ACETIC ACID, GLACIAL
 - ACETASOL, ACETIC ACID, GLACIAL
 - ACYCLOVIR, ACYCLOVIR
 - ALBUTEROL SULFATE, ALBUTEROL SULFATE
 - AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
 - AMINOPHYLLINE DYE FREE, AMINOPHYLLINE
 - BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
 - CAPITAL AND CODEINE, ACETAMINOPHEN
 - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
 - CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 - CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 - CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
 - CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
 - CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
 - CONSTULOSE, LACTULOSE
 - CROMOLYN SODIUM, CROMOLYN SODIUM
 - CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
 - CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 - DEXAMETHASONE, DEXAMETHASONE
 - ENULOSE, LACTULOSE
 - ERYTHROMYCIN ESTOLATE, ERYTHROMYCIN ESTOLATE
 - ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
 - FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 - FLUOCINONIDE, FLUOCINONIDE
 - FLUOXETINE, FLUOXETINE HYDROCHLORIDE
 - HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
 - HYDROCODONE COMPOUND, HOMATROPINE METHYLBROMIDE
 - HYDROCORTISONE, HYDROCORTISONE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - IBUPROFEN, IBUPROFEN
 - IBUPROFEN, IBUPROFEN (OTC)
 - INFANTS' FEVERALL, ACETAMINOPHEN (OTC)
 - IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 - KADIAN, MORPHINE SULFATE
 - LIDOCAINE HYDROCHLORIDE VISCOUS, LIDOCAINE HYDROCHLORIDE
 - LINDANE, LINDANE
 - METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 - MICONAZOLE 7, MICONAZOLE NITRATE (OTC)
 - MICONAZOLE NITRATE, MICONAZOLE NITRATE
 - MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 - MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 - MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 - MYKACET, NYSTATIN
 - M-ZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
 - M-ZOLE 7 DUAL PACK, MICONAZOLE NITRATE (OTC)
 - NYSTATIN, NYSTATIN
 - PERMETHRIN, PERMETHRIN
 - PERMETHRIN, PERMETHRIN (OTC)
 - PHENYTOIN, PHENYTOIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* ALPHARMA US PHARMACEUTICALS DIVISION
PROMETH VC PLAIN, PHENYLEPHRINE HYDROCHLORIDE
PROMETH VC W/ CODEINE, CODEINE PHOSPHATE
PROMETH W/ CODEINE, CODEINE PHOSPHATE
PROMETH W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
SELENIUM SULFIDE, SELENIUM SULFIDE
SULFATRIM PEDIATRIC, SULFAMETHOXAZOLE
SULFATRIM, SULFAMETHOXAZOLE
THEOPHYLLINE, THEOPHYLLINE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
TRIACIN-C, CODEINE PHOSPHATE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
VALNAC, BETAMETHASONE VALERATE

ALRA

* ALRA LABORATORIES INC
CHOLAC, LACTULOSE
CONSTILAC, LACTULOSE
ERYZOLE, ERYTHROMYCIN ETHYLSUCCINATE
GEN-XENE, CLORAZEPATE DIPOTASSIUM
IBU-TAB 200, IBUPROFEN (OTC)
IBU-TAB, IBUPROFEN
K+10, POTASSIUM CHLORIDE
K+8, POTASSIUM CHLORIDE

ALTANA

* ALTANA INC
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
AMCINONIDE, AMCINONIDE
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
BACITRACIN, BACITRACIN
BACITRACIN-NEOMYCIN-POLYMYXIN, BACITRACIN ZINC
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CHLORAMPHENICOL, CHLORAMPHENICOL
CICLOPIROX, CICLOPIROX
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CUTIVATE, FLUTICASONE PROPIONATE
DESONIDE, DESONIDE
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
ECONAZOLE NITRATE, ECONAZOLE NITRATE
ERYTHROMYCIN, ERYTHROMYCIN
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
HYDROCORTISONE, HYDROCORTISONE
KETOCONAZOLE, KETOCONAZOLE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
METRONIDAZOLE, METRONIDAZOLE
MOMETASONE FUROATE, MOMETASONE FUROATE
MUPIROCIN, MUPIROCIN
NYSTATIN, NYSTATIN
OXISTAT, OXICONAZOLE NITRATE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
TEMOVATE E, CLOBETASOL PROPIONATE
TEMOVATE, CLOBETASOL PROPIONATE
TERCONAZOLE, TERCONAZOLE
TOBRAMYCIN, TOBRAMYCIN
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

ALTANA PHARMA

* ALTANA PHARMA AG
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE

ALZA

* ALZA CORP
CONCERTA, METHYLPHENIDATE HYDROCHLORIDE
DITROPAN XL, OXYBUTYNIN CHLORIDE
DITROPAN, OXYBUTYNIN CHLORIDE
DOXIL, DOXORUBICIN HYDROCHLORIDE
DURAGESIC-100, FENTANYL
DURAGESIC-12, FENTANYL
DURAGESIC-25, FENTANYL
DURAGESIC-50, FENTANYL
DURAGESIC-75, FENTANYL
EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE, BROMPHENIRAMINE
MALEATE (OTC)
VIADUR, LEUPROLIDE ACETATE

AM BIOSCIENCE

* AMERICAN BIOSCIENCE INC
ABRAXANE, PACLITAXEL

AM PHARM

* AMERICAN PHARMACEUTICAL PARTNERS INC
ADENOSINE, ADENOSINE
AZITHROMYCIN, AZITHROMYCIN
CARBOPLATIN, CARBOPLATIN
CLADRIBINE, CLADRIBINE
CYTARABINE, CYTARABINE
DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DIMENHYDRINATE, DIMENHYDRINATE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
FLUMAZENIL, FLUMAZENIL
FLUOROURACIL, FLUOROURACIL
HEPARIN LOCK FLUSH, HEPARIN SODIUM
HEPARIN SODIUM IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM
HEPFLUSH-10, HEPARIN SODIUM
LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
PIPERACILLIN, PIPERACILLIN SODIUM
TERBUTALINE SULFATE, TERBUTALINE SULFATE
TOBRAMYCIN, TOBRAMYCIN SULFATE
VALPROATE SODIUM, VALPROATE SODIUM
VINCRISTINE SULFATE, VINCRISTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE

AM PHARM PARTNERS

* AMERICAN PHARMACEUTICAL PARTNERS INC
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
BACITRACIN, BACITRACIN
BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
CALCITRIOL, CALCITRIOL
CARBOPLATIN, CARBOPLATIN
CEFAZOLIN SODIUM, CEFZOLIN SODIUM
CEFOTAXIME, CEFOTAXIME SODIUM
CEFOXITIN, CEFOTAXIN SODIUM
CEFUROXIME SODIUM, CEFUROXIME SODIUM
CEFUROXIME, CEFUROXIME SODIUM
CHLORAMPHENICOL SODIUM SUCCINATE, CHLORAMPHENICOL SODIUM SUCCINATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** A **

* AMERICAN PHARMACEUTICAL PARTNERS INC
 CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
 CISPLATIN, CISPLATIN
 CLINDAMYCIN PHOSPHATE IN DEXTROSE 5%, CLINDAMYCIN PHOSPHATE
 CYANOCOBALAMIN, CYANOCOBALAMIN
 DACARBAZINE, DACARBAZINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 DOXY 100, DOXYCYCLINE HYCLATE
 DOXY 200, DOXYCYCLINE HYCLATE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FOLIC ACID, FOLIC ACID
 FUROSEMIDE, FUROSEMIDE
 GENTAMICIN SULFATE, GENTAMICIN SULFATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 IFOSFAMIDE, IFOSFAMIDE
 IOPAMIDOL-250, IOPAMIDOL
 IOPAMIDOL-300, IOPAMIDOL
 IOPAMIDOL-370, IOPAMIDOL
 KANAMYCIN, KANAMYCIN SULFATE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 MAGNESIUM SULFATE, MAGNESIUM SULFATE
 MANNITOL 25%, MANNITOL
 MESNA, MESNA
 METARAMINOL BITARTRATE, METARAMINOL BITARTRATE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 NEBUPENT, PENTAMIDINE ISETHIONATE
 OXYTOCIN, OXYTOCIN
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PENTAM, PENTAMIDINE ISETHIONATE
 POTASSIUM CHLORIDE IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROGESTERONE, PROGESTERONE
 PROPRANOLOL, PROPRANOLOL HYDROCHLORIDE
 PROTAMINE SULFATE, PROTAMINE SULFATE
 PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, WATER FOR INJECTION, STERILE
 THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 THIOTEPA, THIOTEPA
 TOBRAMYCIN (PHARMACY BULK), TOBRAMYCIN SULFATE
 TOBRAMYCIN, TOBRAMYCIN SULFATE
 VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
 VIBISONE, CYANOCOBALAMIN
 VINBLASTINE SULFATE, VINBLASTINE SULFATE
 VINCRISTINE SULFATE, VINCRISTINE SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** A **

AMBI PHARMS

- * AMBI PHARMACEUTICALS INC
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE

AMERSHAM

- * AMERSHAM CORP SUB RADIOCHEMICAL CENTER
INDICLOR, INDIUM IN 111 CHLORIDE

AMGEN

- * AMGEN INC
SENSIPAR, CINACALCET HYDROCHLORIDE

AMIDE PHARM

- * AMIDE PHARMACEUTICAL INC
ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE, ACETAMINOPHEN
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
CARISOPRODOL AND ASPIRIN, ASPIRIN
CARISOPRODOL, CARISOPRODOL
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
CHLORZOXAZONE, CHLORZOXAZONE
CILOSTAZOL, CILOSTAZOL
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DANTROLENE SODIUM, DANTROLENE SODIUM
DIGOXIN, DIGOXIN
GLYBURIDE, GLYBURIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
HOMATROPINE METHYLBROMIDE AND HYDROCODONE BITARTRATE, HOMATROPINE METHYLBROMIDE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NALOXONE HYDROCHLORIDE AND PENTAZOCAINE, NALOXONE HYDROCHLORIDE
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
QUINARETIC, HYDROCHLOROTHIAZIDE
RIMACTANE, RIFAMPIN
RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
URSODIOL, URSODIOL

AMPHASTAR

- * AMPHASTAR PHARMACEUTICALS INC
CORTROSYN, COSYNTROPIN
DUOCAINE, BUPIVACAINE HYDROCHLORIDE

AMPHASTAR PHARM

- * AMPHASTAR PHARMACEUTICAL INC
AMPHADASE, HYALURONIDASE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

AMYLIN

- * AMYLIN PHARMACEUTICALS INC
BYETTA, EXENATIDE SYNTHETIC
SYMLIN, PRAMLINTIDE ACETATE

ANABOLIC

- * ANABOLIC INC
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN

ANABOLIC LABS

- * ANABOLIC LABORATORIES INC
BUTALBITAL, ACETAMINOPHEN, CAFFEINE AND CODEINE PHOSPHATE, ACETAMINOPHEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** A **

ANBEX

- * ANBEX INC
IOSAT, POTASSIUM IODIDE (OTC)

ANDRX

- * ANDRX CORP
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE

ANDRX LABS LLC

- * ANDRX LABS LLC
ALTOPREV, LOVASTATIN
FORTAMET, METFORMIN HYDROCHLORIDE

ANDRX PHARMS

- * ANDRX PHARMACEUTICALS INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
CARTIA XT, DILTIAZEM HYDROCHLORIDE
CLARITHROMYCIN, CLARITHROMYCIN
CODRIX, ACETAMINOPHEN
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
KETOPROFEN, KETOPROFEN
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE, LORATADINE (OTC)
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NAPROXEN SODIUM, NAPROXEN SODIUM
NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE, ETHINYL ESTRADIOL
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PREVIFEM, ETHINYL ESTRADIOL
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
TAZTIA XT, DILTIAZEM HYDROCHLORIDE
TRI-PREVIFEM, ETHINYL ESTRADIOL
- * ANDRX PHARMACEUTICALS LLC
CODRIX, ACETAMINOPHEN
ETODOLAC, ETODOLAC
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN

ANGELINI

- * ANGELINI PHARMACEUTICALS INC
REV-EYES, DAPIPRAZOLE HYDROCHLORIDE

ANTHRA

- * ANTHRA PHARMACEUTICALS INC
VALSTAR PRESERVATIVE FREE, VALRUBICIN

APOTEX

- * APOTEX CORP
ACYCLOVIR, ACYCLOVIR
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CARBAMAZEPINE, CARBAMAZEPINE
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE
EDETATE DISODIUM, EDETATE DISODIUM
ETODOLAC, ETODOLAC
FAMOTIDINE PRESERVATIVE FREE (PHARMACY BULK), FAMOTIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

- * APOTEX CORP
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUMAZENIL, FLUMAZENIL
FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
GABAPENTIN, GABAPENTIN
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL LACTATE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LEFLUNOMIDE, LEFLUNOMIDE
LISINOPRIL, LISINOPRIL
LITHIUM CARBONATE, LITHIUM CARBONATE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MILRINONE LACTATE IN 5% DEXTROSE IN PLASTIC CONTAINER, MILRINONE LACTATE
MILRINONE LACTATE, MILRINONE LACTATE
PREDNISOLONE, PREDNISOLONE
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
TORSEMIDE, TORSEMIDE
VALPROIC ACID, VALPROIC ACID
ZONISAMIDE, ZONISAMIDE
- * APOTEX INC
ALLOPURINOL, ALLOPURINOL
CILOSTAZOL, CILOSTAZOL
DESMOPRESSIN ACETATE (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LORATADINE, LORATADINE (OTC)
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
OXAPROZIN, OXAPROZIN

APOTHECON

- * APOTHECON INC DIV BRISTOL MYERS SQUIBB
ACYCLOVIR, ACYCLOVIR
AMIKIN, AMIKACIN SULFATE
CAPOZIDE 25/15, CAPTOPRIL
CAPOZIDE 25/25, CAPTOPRIL
CAPOZIDE 50/15, CAPTOPRIL
CAPOZIDE 50/25, CAPTOPRIL
CAPTOPRIL, CAPTOPRIL
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFAZOLIN SODIUM, CEFZOLIN SODIUM
CEPHALEXIN, CEPHALEXIN
CORGARD, NADOLOL
DESYREL, TRAZODONE HYDROCHLORIDE
DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
FUNGIZONE, AMPHOTERICIN B
KANTREX, KANAMYCIN SULFATE
KENALOG IN ORABASE, TRIAMCINOLONE ACETONIDE
KENALOG, TRIAMCINOLONE ACETONIDE
KENALOG-10, TRIAMCINOLONE ACETONIDE
KENALOG-40, TRIAMCINOLONE ACETONIDE
KLOTRIX, POTASSIUM CHLORIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MUCOMYST, ACETYLCYSTEINE
MYCOSTATIN, NYSTATIN
NATURETIN-5, BENDROFLUMETHIAZIDE
OPHTHAINE, PROPARACAINE HYDROCHLORIDE
PRINCIPEN, AMPICILLIN/AMPICILLIN TRIHYDRATE
PROLIXIN, FLUPHENAZINE HYDROCHLORIDE
PRONESTYL, PROCAINAMIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** A **

* APOTHECON INC DIV BRISTOL MYERS SQUIBB
PRONESTYL-SR, PROCAINAMIDE HYDROCHLORIDE
STADOL PRESERVATIVE FREE, BUTORPHANOL TARTRATE
STADOL, BUTORPHANOL TARTRATE
SUMYCIN, TETRACYCLINE HYDROCHLORIDE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TRIMOX, AMOXICILLIN
VEETIDS, PENICILLIN V POTASSIUM
VELOSEF '125', CEPHRADINE
VELOSEF '250', CEPHRADINE
VELOSEF, CEPHRADINE

APOTHEKERNES

* APOTHEKERNES LABORATORIUM A/S
BACITRACIN, BACITRACIN

APPLIED ANAL

* APPLIED ANALYTICAL INDUSTRIES INC
ESTRADIOL, ESTRADIOL

ARCUM

* ARCUM PHARMACEUTICAL CORP
VITAMIN A PALMITATE, VITAMIN A PALMITATE

ARMSTRONG PHARMS

* ARMSTRONG PHARMACEUTICALS INC
ALBUTEROL, ALBUTEROL
EPINEPHRINE, EPINEPHRINE (OTC)

ASCENT PEDS

* ASCENT PEDIATRICS INC
ORAPRED, PREDNISOLONE SODIUM PHOSPHATE

ASCHER

* BF ASCHER AND CO INC
HY-PHEN, ACETAMINOPHEN

ASTELLAS

* ASTELLAS PHARMA US INC
ADENOCARD, ADENOSINE
ADENOSCAN, ADENOSINE
AMBISOME, AMPHOTERICIN B
ARISTOCORT A, TRIAMCINOLONE ACETONIDE
ARISTOCORT, TRIAMCINOLONE
CEFIZOX IN PLASTIC CONTAINER, CEFTIZOXIME SODIUM
CEFIZOX, CEFTIZOXIME SODIUM
CYCLOCORT, AMCINONIDE
MYCAMINE, MICA FUNGIN SODIUM
PROGRAF, TACROLIMUS
PROTOPIC, TACROLIMUS
VAPRISOL, CONIVAPTAN HYDROCHLORIDE
VESICARE, SOLIFENACIN SUCCINATE

ASTRAZENECA

* ASTRAZENECA LP
ASTRAMORPH PF, MORPHINE SULFATE
EMLA, LIDOCAINE
ENTOCORT EC, BUDESONIDE
FOSCAVIR, FOSCARNET SODIUM
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
NAROPIN, ROPIVACAINE HYDROCHLORIDE MONOHYDRATE
NESACAINE, CHLOROPROCAINE HYDROCHLORIDE
NESACAINE-MPF, CHLOROPROCAINE HYDROCHLORIDE
NEXIUM IV, ESOMEPRAZOLE SODIUM
NEXIUM, ESOMEPRAZOLE MAGNESIUM
POLOCAINE, MEPIVACAINE HYDROCHLORIDE
POLOCAINE-MPF, MEPIVACAINE HYDROCHLORIDE
PRILOSEC OTC, OMEPRAZOLE MAGNESIUM (OTC)
PRILOSEC, OMEPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

- * ASTRAZENECA LP
 - PULMICORT RESPULES, BUDESONIDE
 - RHINOCORT, BUDESONIDE
 - SENSORCAINE, BUPIVACAINE HYDROCHLORIDE
 - SEROQUEL, QUETIAPINE FUMARATE
 - TENORMIN, ATENOLOL
 - TOPROL-XL, METOPROLOL SUCCINATE
 - XYLOCAINE 4% PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 - XYLOCAINE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 - XYLOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
 - XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
 - XYLOCAINE, LIDOCAINE HYDROCHLORIDE
- * ASTRAZENECA PHARMACEUTICALS LP
 - ATACAND HCT, CANDESARTAN CILEXETIL
 - ATACAND, CANDESARTAN CILEXETIL
 - CEFOTAN IN PLASTIC CONTAINER, CEFOTETAN DISODIUM
 - CEFOTAN, CEFOTETAN DISODIUM
 - CRESTOR, ROSUVASTATIN CALCIUM
 - FASLODEX, FULVESTRANT
 - LEXXEL, ENALAPRIL MALEATE
 - NOLVADEX, TAMOXIFEN CITRATE
 - PLENDIL, FELODIPINE
 - PULMICORT, BUDESONIDE
 - TENORETIC 100, ATENOLOL
 - TENORETIC 50, ATENOLOL
 - TENORMIN, ATENOLOL
 - ZOMIG, ZOLMITRIPTAN
 - ZOMIG-ZMT, ZOLMITRIPTAN
- * ASTRAZENECA UK LTD
 - ACCOLATE, ZAFIRLUKAST
 - ARIMIDEX, ANASTROZOLE
 - CASODEX, BICALUTAMIDE
 - DIPRIVAN, PROPOFOL
 - IRESSA, GEFITINIB
 - MERREM I.V., MEROPENEM
 - ZESTORETIC, HYDROCHLOROTHIAZIDE
 - ZESTRIL, LISINAPRIL
 - ZOLADEX, GOSERELIN ACETATE

AUROBINDO

- * AUROBINDO PHARMA LTD
 - AMOXICILLIN, AMOXICILLIN
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - MIRTAZAPINE, MIRTAZAPINE
 - ZIDOVUDINE, ZIDOVUDINE

AUROBINDO PHARMA LTD

- * AUROBINDO PHARMA LTD INC
 - CEPHALEXIN, CEPHALEXIN
 - MIRTAZAPINE, MIRTAZAPINE

AUROSAL PHARMS

- * AUROSAL PHARMACEUTICALS LLC
 - AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE

AUXILIUM PHARMS

- * AUXILIUM PHARMACEUTICALS
 - TESTIM, TESTOSTERONE

AVENTIS

- * AVENTIS PHARMACEUTICAL PRODUCTS INC
 - DDAVP (NEEDS NO REFRIGERATION), DESMOPRESSIN ACETATE
 - DDAVP, DESMOPRESSIN ACETATE
 - LOVENOX (PRESERVATIVE FREE), ENOXAPARIN SODIUM
 - LOVENOX, ENOXAPARIN SODIUM
 - LOZOL, INDAPAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** A ****

* AVENTIS PHARMACEUTICAL PRODUCTS INC
NASACORT AQ, TRIAMCINOLONE ACETONIDE
RILUTEK, RILUZOLE

AVENTIS BEHRING

* AVENTIS BEHRING LLC
FLURBIPROFEN SODIUM, FLURBIPROFEN SODIUM

AVENTIS PHARMS

* AVENTIS PHARMACEUTICALS INC
A/T/S, ERYTHROMYCIN
ALLEGRA D 24 HOUR, FEXOFENADINE HYDROCHLORIDE
ALLEGRA, FEXOFENADINE HYDROCHLORIDE
ALLEGRA-D 12 HOUR, FEXOFENADINE HYDROCHLORIDE
AMARYL, GLIMEPIRIDE
ANZEMET, DOLASETRON MESYLATE MONOHYDRATE
ARAVA, LEFLUNOMIDE
CANTIL, MEPENZOLATE BROMIDE
CLAFORAN IN DEXTROSE 5% IN PLASTIC CONTAINER, CEFOTAXIME SODIUM
CLAFORAN, CEFOTAXIME SODIUM
CLOMID, CLOMIPHENE CITRATE
DIABETA, GLYBURIDE
GAVISCON, ALUMINUM HYDROXIDE (OTC)
HIPREX, METHENAMINE HIPPURATE
KETEK, TELITHROMYCIN
LANTUS, INSULIN GLARGINE RECOMBINANT
LASIX, FUROSEMIDE
NASACORT HFA, TRIAMCINOLONE ACETONIDE
NICODERM CQ, NICOTINE (OTC)
NILANDRON, NILUTAMIDE
NORPRAMIN, DESIPRAMINE HYDROCHLORIDE
PRIFTIN, RIFAPENTINE
RIFADIN, RIFAMPIN
RIFAMATE, ISONIAZID
RIFATER, ISONIAZID
TAXOTERE, DOCETAXEL
TENUATE DOSPAN, DIETHYLPROPION HYDROCHLORIDE
TENUATE, DIETHYLPROPION HYDROCHLORIDE
TRENTAL, PENTOXIFYLLINE

AVEVA

* AVEVA DRUG DELIVERY SYSTEMS INC
PROSTEP, NICOTINE (OTC)

AXCAN SCANDIPHARM

* AXCAN SCANDIPHARM INC
BENTYL PRESERVATIVE FREE, DICYCLOMINE HYDROCHLORIDE
BENTYL, DICYCLOMINE HYDROCHLORIDE
CANASA, MESALAMINE
CARAFATE, SUCRALFATE
PHOTOFRIN, PORFIMER SODIUM
URSO 250, URSODIOL
URSO FORTE, URSODIOL

AYERST

* AYERST LABORATORIES INC
PHOSPHOLINE IODIDE, ECHOTHIOPHATE IODIDE

B BRAUN

* B BRAUN MEDICAL INC
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ALCOHOL 10% AND DEXTROSE 5%, ALCOHOL
ALCOHOL 5% AND DEXTROSE 5%, ALCOHOL
BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER, BRETYLIUM TOSYLATE
CEFAZOLIN AND DEXTROSE, CEFZOLIN SODIUM
CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER, CEFOTAXIME SODIUM
CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER, CEFOTAXIME SODIUM
CEFTRIAXONE AND DEXTROSE IN DUPLEX CONTAINER, CEFTRIAXONE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* B BRAUN MEDICAL INC
 CEFUROXIME AND DEXTROSE IN DUPLEX CONTAINER, CEFUROXIME SODIUM
 DEXTROSE 10% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% IN HALF-STRENGTH LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN ACETATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% IN LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIALYTE LM/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIALYTE LM/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIALYTE LM/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DOPAMINE HYDROCHLORIDE AND DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE AND DEXTROSE 5%, DOPAMINE HYDROCHLORIDE
 FREAMINE HBC 6.9%, AMINO ACIDS
 FREAMINE III 10%, AMINO ACIDS
 FREAMINE III 3% W/ ELECTROLYTES, AMINO ACIDS
 FREAMINE III 8.5% W/ ELECTROLYTES, AMINO ACIDS
 FREAMINE III 8.5%, AMINO ACIDS
 GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
 HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPATAMINE 8%, AMINO ACIDS
 ISOLYTE E IN DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE E IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE H IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE M IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE P IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE R IN DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 ISOLYTE S IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 ISOLYTE S IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 ISOLYTE S PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE
 HYDROCHLORIDE
 MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 10% W/ DEXTROSE 5% IN DISTILLED WATER, MANNITOL
 MANNITOL 10%, MANNITOL
 MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 15% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.45%, MANNITOL
 MANNITOL 15%, MANNITOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* B BRAUN MEDICAL INC
 MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 20%, MANNITOL
 MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL
 MANNITOL 5% W/ DEXTROSE 5% IN SODIUM CHLORIDE 0.12%, MANNITOL
 MANNITOL 5%, MANNITOL
 METRO I.V. IN PLASTIC CONTAINER, METRONIDAZOLE
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 NEPHRAMINE 5.4%, AMINO ACIDS
 NUTRILIPID 10%, SOYBEAN OIL
 NUTRILIPID 20%, SOYBEAN OIL
 PHYSIOLYTE IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.037% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.075% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.11% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** B ****

* B BRAUN MEDICAL INC
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.15% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.22% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 3.3% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 0.3% IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PROCALAMINE, AMINO ACIDS
 RESECTISOL IN PLASTIC CONTAINER, MANNITOL
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM LACTATE 1/6 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
 SORBITOL 3.3% IN PLASTIC CONTAINER, SORBITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, WATER FOR INJECTION, STERILE
 STERILE WATER IN PLASTIC CONTAINER, WATER FOR IRRIGATION, STERILE
 THEOPHYLLINE 0.04% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.08% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 THEOPHYLLINE 0.16% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* B BRAUN MEDICAL INC
THEOPHYLLINE 0.32% AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
TROPAMINE 10%, AMINO ACIDS
TROPAMINE, AMINO ACIDS

BAKER NORTON

* BAKER NORTON PHARMACEUTICALS INC
PACLITAXEL, PACLITAXEL
PROGLYCEM, DIAZOXIDE

BALLARD MEDCL

* BALLARD MEDICAL PRODUCTS INC
PYTEST KIT, UREA, C-14
PYTEST, UREA, C-14

BANNER PHARMACAPS

* BANNER PHARMACAPS INC
BENZONATATE, BENZONATATE
CLOFIBRATE, CLOFIBRATE
ETHOSUXIMIDE, ETHOSUXIMIDE
IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
VALPROIC ACID, VALPROIC ACID
VITAMIN D, ERGOCALCIFEROL

BARR

* BARR LABORATORIES INC
ACETOHEXAMIDE, ACETOHEXAMIDE
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ARANELLE, ETHINYL ESTRADIOL
BALZIVA-21, ETHINYL ESTRADIOL
BALZIVA-28, ETHINYL ESTRADIOL
CAMILA, NORETHINDRONE
CEPHALEXIN, CEPHALEXIN
CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CHLORZOXAZONE, CHLORZOXAZONE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CLARAVIS, ISOTRETINOIN
CLONAZEPAM, CLONAZEPAM
DANAZOL, DANAZOL
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
ASPARTATE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DIAZEPAM, DIAZEPAM
DIDANOSINE, DIDANOSINE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE
E-BASE, ERYTHROMYCIN
ERRIN, NORETHINDRONE
ERYTHROMYCIN ETHYLSUCCINATE AND SULFISOXAZOLE ACETYL, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN STEARATE, ERYTHROMYCIN STEARATE
ERYTHROMYCIN, ERYTHROMYCIN
ESTRADIOL AND NORGESTIMATE, ESTRADIOL
ESTRADIOL, ESTRADIOL
ESTROPIPATE, ESTROPIPATE
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
FEXOFENADINE HYDROCHLORIDE AND PSEUDOEPHEDRINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

* BARR LABORATORIES INC
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
FLUTAMIDE, FLUTAMIDE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
GENCEPT 10/11-21, ETHINYL ESTRADIOL
GENCEPT 10/11-28, ETHINYL ESTRADIOL
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROXYUREA, HYDROXYUREA
HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
ISONIAZID, ISONIAZID
JUNEL 1.5/30, ETHINYL ESTRADIOL
JUNEL 1/20, ETHINYL ESTRADIOL
JUNEL FE 1.5/30, ETHINYL ESTRADIOL
JUNEL FE 1/20, ETHINYL ESTRADIOL
KARIVA, DESOGESTREL
KELNOR, ETHINYL ESTRADIOL
LEFLUNOMIDE, LEFLUNOMIDE
LESSINA-28, ETHINYL ESTRADIOL
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
LITHIUM CARBONATE, LITHIUM CARBONATE
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
MEFLOQUINE, MEFLOQUINE HYDROCHLORIDE
MEGESTROL ACETATE, MEGESTROL ACETATE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
MIRTAZAPINE, MIRTAZAPINE
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
NIACIN, NIACIN
NORETHINDRONE ACETATE, NORETHINDRONE ACETATE
NORTREL 0.5/35-21, ETHINYL ESTRADIOL
NORTREL 0.5/35-28, ETHINYL ESTRADIOL
NORTREL 1/35-21, ETHINYL ESTRADIOL
NORTREL 1/35-28, ETHINYL ESTRADIOL
NORTREL 7/7/7, ETHINYL ESTRADIOL
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PORTIA-28, ETHINYL ESTRADIOL
PROPRANLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
SPRINTEC, ETHINYL ESTRADIOL
SULFINPYRAZONE, SULFINPYRAZONE
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
Trexall, METHOTREXATE SODIUM
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRI-SPRINTEC, ETHINYL ESTRADIOL
WARFARIN SODIUM, WARFARIN SODIUM
ZONISAMIDE, ZONISAMIDE
* BARR PHARMACEUTICALS
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM

BARRIER

* BARRIER THERAPEUTICS
SOLAGE, MEQUINOL

BARTOR

* BARTOR PHARMACAL CO
TESTOSTERONE, TESTOSTERONE

BASF

* BASF CORP
IBU, IBUPROFEN
IBUPROFEN, IBUPROFEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

* BASF CORP
IBUPROFEN, IBUPROFEN (OTC)
LOPURIN, ALLOPURINOL
SSD AF, SILVER SULFADIAZINE
SSD, SILVER SULFADIAZINE

BAUSCH AND LOMB

* BAUSCH AND LOMB INC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALREX, LOTEPIEDNOL ETABONATE
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LOTEMAX, LOTEPIEDNOL ETABONATE
OFLOXACIN, OFLOXACIN
OPCON-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
RETISERT, FLUOCINOLONE ACETONIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
VITRASERT, GANCICLOVIR
ZYLET, LOTEPIEDNOL ETABONATE

* BAUSCH AND LOMB PHARMACEUTICALS INC
ACETIC ACID 2% IN AQUEOUS ALUMINUM ACETATE, ACETIC ACID, GLACIAL
BACITRACIN ZINC AND POLYMYXIN B SULFATE, BACITRACIN ZINC
BETAXOLOL HYDROCHLORIDE, BETAXOLOL HYDROCHLORIDE
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CROLOM, CROMOLYN SODIUM
CROMOLYN SODIUM, CROMOLYN SODIUM
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
DEXASPORIN, DEXAMETHASONE
DIPIVEFRIN HYDROCHLORIDE, DIPIVEFRIN HYDROCHLORIDE
ERYTHROMYCIN, ERYTHROMYCIN
FLUNISOLIDE, FLUNISOLIDE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
NAFAZAIR, NAPHAZOLINE HYDROCHLORIDE
NEOMYCIN AND POLYMYXIN B SULFATES AND BACITRACIN ZINC, BACITRACIN ZINC
NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE, BACITRACIN ZINC
NYSTATIN, NYSTATIN
OPTIPRANOLOL, METIPRANOLOL HYDROCHLORIDE
OTICAIR, HYDROCORTISONE
PENTOLAIR, CYCLOPENTOLATE HYDROCHLORIDE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
SULFACETAMIDE SODIUM AND PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
TIMOLOL MALEATE, TIMOLOL MALEATE
TOBRAMYCIN AND DEXAMETHASONE, DEXAMETHASONE
TOBRAMYCIN, TOBRAMYCIN
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
TROPICAMIDE, TROPICAMIDE

BAXTER HLTHCARE

* BAXTER HEALTHCARE CORP
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ADENOSINE, ADENOSINE
AMINOACETIC ACID 1.5% IN PLASTIC CONTAINER, GLYCINE
ANCEF IN PLASTIC CONTAINER, CEFAZOLIN SODIUM
BACTOCILL IN PLASTIC CONTAINER, OXACILLIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP
 BRANCHAMIN 4% IN PLASTIC CONTAINER, AMINO ACIDS
 CARDIOPLEGIC IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 CEFTAZIDIME SODIUM IN PLASTIC CONTAINER, CEFTAZIDIME SODIUM
 CEFTRIAXONE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
 CLINIMIX 2.75/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 2.75/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 4.25/5 SULFITE FREE IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/10 SULFITE FREE IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/15 SULFITE FREE IN DEXTROSE 15% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/20 SULFITE FREE IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/25 SULFITE FREE IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX 5/35 SULFITE FREE IN DEXTROSE 35% IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 2.75/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 2.75/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 2.75/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 4.25/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 4.25/20 SULFITE-FREE W/ ELECT IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 4.25/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 4.25/5 SULFITE-FREE W/ ELECT IN DEXTROSE 5% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/10 SULFITE-FREE W/ ELECT IN DEXTROSE 10% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/15 SULFITE-FREE W/ ELECT IN DEXTROSE 15% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/20 SULFITE-FREE W/ ELECT IN 20% DEXTROSE W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/25 SULFITE-FREE W/ ELECT IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CLINIMIX E 5/35 SULFITE-FREE W/ ELECT IN DEXTROSE 35% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS
 CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
 DEXTROSE 10% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND ELECTROLYTE NO 75 IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% AND ELECTROLYTE NO.48 IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5% IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 10MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 15MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 20MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 30MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 40MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ (K), DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.2% AND POTASSIUM CHLORIDE 5MEQ, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 15MEQ IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.33% AND POTASSIUM CHLORIDE 5MEQ IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 5%, SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 20MEQ (K) IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 60% IN PLASTIC CONTAINER, DEXTROSE
 DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE
 DIANEAL 137 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL 137 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL 137 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-1 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-1 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-1 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-1 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DIANEAL PD-2 W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
 EXTRANEAL, ICODextrin
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 HEPARIN SODIUM 1,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 2,000 UNITS AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPARIN SODIUM 25,000 UNITS AND DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
 HEPATASOL 8%, AMINO ACIDS
 IRRIGATING SOLUTION G IN PLASTIC CONTAINER, CITRIC ACID
 ISOTONIC GENTAMICIN SULFATE IN PLASTIC CONTAINER, GENTAMICIN SULFATE
 LACTATED RINGER'S AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.2% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.4% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE 0.8% AND DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
 MESNEX, MESNA
 MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 NALLPEN IN PLASTIC CONTAINER, NAFCILLIN SODIUM
 NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN
 OSMITROL 10% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 10% IN WATER, MANNITOL
 OSMITROL 15% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 15% IN WATER, MANNITOL
 OSMITROL 20% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 20% IN WATER, MANNITOL
 OSMITROL 5% IN WATER IN PLASTIC CONTAINER, MANNITOL
 OSMITROL 5% IN WATER, MANNITOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP
 PENICILLIN G POTASSIUM IN PLASTIC CONTAINER, PENICILLIN G POTASSIUM
 PLASMA-LYTE 148 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 PLASMA-LYTE 148 IN WATER IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PLASMA-LYTE 56 AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 PLASMA-LYTE 56 IN PLASTIC CONTAINER, MAGNESIUM ACETATE TETRAHYDRATE
 PLASMA-LYTE A IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE
 PLASMA-LYTE M AND DEXTROSE 5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 PLASMA-LYTE R IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER,
 CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER,
 DEXTROSE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PREMASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 PREMASOL 6% IN PLASTIC CONTAINER, AMINO ACIDS
 RENAMIN W/O ELECTROLYTES, AMINO ACIDS
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 SEVOFLURANE, SEVOFLURANE
 SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, POTASSIUM
 CHLORIDE
 SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.224%, POTASSIUM CHLORIDE
 SODIUM CHLORIDE 0.9% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 0.9% IN STERILE PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 3% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM CHLORIDE 5% IN PLASTIC CONTAINER, SODIUM CHLORIDE
 SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
 SORBITOL 3% IN PLASTIC CONTAINER, SORBITOL
 STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, WATER FOR INJECTION, STERILE
 STERILE WATER IN PLASTIC CONTAINER, WATER FOR IRRIGATION, STERILE
 STERILE WATER, WATER FOR IRRIGATION, STERILE
 THEOPHYLLINE AND DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
 TIS-U-SOL IN PLASTIC CONTAINER, MAGNESIUM SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

- * BAXTER HEALTHCARE CORP
 - TIS-U-SOL, MAGNESIUM SULFATE
 - TRAVASOL 10% IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 10% W/O ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 3.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 3.5% W/ ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 5.5% IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 5.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 5.5% W/ ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 5.5% W/O ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 8.5% IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 8.5% SULFITE FREE W/ ELECTROLYTES IN PLASTIC CONTAINER, AMINO ACIDS
 - TRAVASOL 8.5% W/ ELECTROLYTES, AMINO ACIDS
 - TRAVASOL 8.5% W/O ELECTROLYTES, AMINO ACIDS
 - VANCOCIN HYDROCHLORIDE IN PLASTIC CONTAINER, VANCOMYCIN HYDROCHLORIDE
- * BAXTER HEALTHCARE CORP ANESTHESIA AND CRITICAL CARE
 - ACYCLOVIR IN SODIUM CHLORIDE 0.9% PRESERVATIVE FREE, ACYCLOVIR SODIUM
 - ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
 - AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
 - ATRAURIUM BESYLATE, ATRACURIUM BESYLATE
 - CEFOXITIN, CEFOTAXIME SODIUM
 - CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
 - CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 - DEXAMETHASONE, DEXAMETHASONE SODIUM PHOSPHATE
 - DIAZEPAM, DIAZEPAM
 - DIGOXIN, DIGOXIN
 - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 - DIPYRIDAMOLE, DIPYRIDAMOLE
 - DURAMORPH PF, MORPHINE SULFATE
 - ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
 - FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 - FAMOTIDINE, FAMOTIDINE
 - FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
 - FLUMAZENIL, FLUMAZENIL
 - FUROSEMIDE, FUROSEMIDE
 - HEPARIN SODIUM, HEPARIN SODIUM
 - HEP-LOCK U/P, HEPARIN SODIUM
 - HEP-LOCK, HEPARIN SODIUM
 - INFUMORPH, MORPHINE SULFATE
 - KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 - MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 - MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
 - MILRINONE LACTATE, MILRINONE LACTATE
 - MILRINONE LACTOSE IN 5% DEXTROSE, MILRINONE LACTATE
 - MITOMYCIN, MITOMYCIN
 - PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 - PROCHLORPERAZINE, PROCHLORPERAZINE EDISYLATE
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - SUFENTANIL CITRATE, SUFENTANIL CITRATE
 - SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 - THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
 - TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
 - VECURIUM BROMIDE, VECURIUM BROMIDE
 - VINORELBINE TARTRATE, VINORELBINE TARTRATE
- * BAXTER HEALTHCARE INTERNATIONAL SPECIALTY THERAPIES DIV
 - FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
 - PROSOL 20% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

BAXTER HLTHCARE CORP

- * BAXTER HEALTHCARE CORP ANESTHESIA CRITICAL CARE
 - AMRINONE LACTATE, INAMRINONE LACTATE
 - ATIVAN, LORAZEPAM
 - ATRAURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
 - ATRAURIUM BESYLATE, ATRACURIUM BESYLATE
 - BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BAXTER HEALTHCARE CORP ANESTHESIA CRITICAL CARE
BREVILOC IN PLASTIC CONTAINER, ESMOLOL HYDROCHLORIDE
BREVILOC, ESMOLOL HYDROCHLORIDE
DOPRAM, DOXAPRAM HYDROCHLORIDE
ENLON, EDROPHONIUM CHLORIDE
ENLON-PLUS, ATROPINE SULFATE
ETHRANE, ENFLURANE
FORANE, ISOFLURANE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MILRINONE LACTATE, MILRINONE LACTATE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
PROTOPAM CHLORIDE, PRALIDOXIME CHLORIDE
REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
REVEX, NALMEFENE HYDROCHLORIDE
ROBAXIN, METHOCARBAMOL
ROBINUL, GLYCOPYRROLATE
SUPRANE, DESFLURANE

BAYER

* BAYER HEALTHCARE LLC
ACTRON, KETOPROFEN (OTC)
ALEVE COLD AND SINUS, NAPROXEN SODIUM (OTC)
ALEVE, NAPROXEN SODIUM (OTC)
BAYER EXTRA STRENGTH ASPIRIN FOR MIGRAINE PAIN, ASPIRIN (OTC)
FEMSTAT 3, BUTOCONAZOLE NITRATE (OTC)

BAYER PHARMS

* BAYER PHARMACEUTICALS CORP
ADALAT CC, NIFEDIPINE
AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER, MOXIFLOXACIN HYDROCHLORIDE
AVELOX, MOXIFLOXACIN HYDROCHLORIDE
BILTRICIDE, PRAZIQUANTEL
CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER, CIPROFLOXACIN
CIPRO XR, CIPROFLOXACIN
CIPRO, CIPROFLOXACIN
CIPRO, CIPROFLOXACIN HYDROCHLORIDE
DTIC-DOME, DACARBAZINE
LEVITRA, VARDENAFIL HYDROCHLORIDE
MYCELEX, CLOTRIMAZOLE
MYCELEX-7 COMBINATION PACK, CLOTRIMAZOLE (OTC)
MYCELEX-7, CLOTRIMAZOLE (OTC)
NEXAVAR, SORAFENIB TOSYLATE
NIMOTOP, NIMODIPINE
PRECOSE, ACARBOSE
TRASYLOL, APROTININ BOVINE

BECTON DICKINSON

* BECTON DICKINSON AND CO
E-Z PREP 220, POVIDONE-IODINE (OTC)
E-Z PREP, POVIDONE-IODINE (OTC)
E-Z SCRUB 201, POVIDONE-IODINE (OTC)
E-Z SCRUB 241, POVIDONE-IODINE (OTC)

BEDFORD

* BEDFORD LABORATORIES DIV BEN VENUE LABORATORIES INC
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
ACETYLCYSTEINE, ACETYLCYSTEINE
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ADENOSINE, ADENOSINE
ALPROSTADIL, ALPROSTADIL
AMIKACIN SULFATE, AMIKACIN SULFATE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMRINONE, INAMRINONE LACTATE
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BEDFORD LABORATORIES DIV BEN VENUE LABORATORIES INC
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 AZATHIOPRINE SODIUM, AZATHIOPRINE SODIUM
 BLEOMYCIN, BLEOMYCIN SULFATE
 BUMETANIDE, BUMETANIDE
 BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE
 BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
 BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
 CARBOPLATIN, CARBOPLATIN
 CERUBIDINE, DAUNORUBICIN HYDROCHLORIDE
 CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE
 CISPLATIN, CISPLATIN
 CLADRIBINE, CLADRIBINE
 CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CYCLOSPORINE, CYCLOSPORINE
 CYTARABINE, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DESMOPRESSIN ACETATE PRESERVATIVE FREE, DESMOPRESSIN ACETATE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXRAZOXANE, DEXRAZOXANE HYDROCHLORIDE
 DICYCLOMINE HYDROCHLORIDE (PRESERVATIVE-FREE), DICYCLOMINE HYDROCHLORIDE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 ETOMIDATE, ETOMIDATE
 ETOPOSIDE, ETOPOSIDE
 FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE
 FLOXURIDINE, FLOXURIDINE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 GANCICLOVIR SODIUM, GANCICLOVIR SODIUM
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEVOCARNITINE, LEVOCARNITINE
 LORAZEPAM, LORAZEPAM
 MESNA, MESNA
 METHOTREXATE PRESERVATIVE FREE, METHOTREXATE
 METHOTREXATE SODIUM, METHOTREXATE SODIUM
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE
 MITOMYCIN, MITOMYCIN
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE (PRESERVATIVE FREE), OCTREOTIDE ACETATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 OFLOXACIN, OFLOXACIN
 PACLITAXEL, PACLITAXEL
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PHENTOLAMINE MESYLATE, PHENTOLAMINE MESYLATE
 POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROPOFOL, PROPOFOL
 PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
 RANITIDINE, RANITIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

* BEDFORD LABORATORIES DIV BEN VENUE LABORATORIES INC
RIFAMPIN, RIFAMPIN
TERBUTALINE SULFATE, TERBUTALINE SULFATE
THIOTEPA, THIOTEPA
VALPROATE SODIUM, VALPROATE SODIUM
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VINBLASTINE SULFATE, VINBLASTINE SULFATE
VINOELBINE TARTRATE, VINOELBINE TARTRATE

BEDFORD LABS

* BEDFORD LABORATORIES
ALLOPURINOL SODIUM, ALLOPURINOL SODIUM
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
CARBOPLATIN, CARBOPLATIN
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
FLUMAZENIL, FLUMAZENIL
LORAZEPAM PRESERVATIVE FREE, LORAZEPAM
METOPROLOL TARTRATE, METOPROLOL TARTRATE
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
* BEDFORD LABORATORIES INC
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE

BELCHER

* BELCHER PHARMACEUTICALS INC
CEPHALEXIN, CEPHALEXIN

BEN VENUE

* BEN VENUE LABORATORIES INC
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
FAMOTIDINE PRESERVATIVE FREE, FAMOTIDINE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

BERLEX

* BERLEX INC
ANGELIQ, DROSPIRENONE
BETAPACE AF, SOTALOL HYDROCHLORIDE
BETAPACE, SOTALOL HYDROCHLORIDE
CLIMARA PRO, ESTRADIOL
CLIMARA, ESTRADIOL
FLUDARA, FLUDARABINE PHOSPHATE
LEVITE, ETHINYL ESTRADIOL
MAGNEVIST, GADOPENTETATE DIMEGLUMINE
MIRENA, LEVONORGESTREL
REFLUDAN, LEPIDURIN RECOMBINANT
ULTRAVIST (PHARMACY BULK), IOPROMIDE
ULTRAVIST 150, IOPROMIDE
ULTRAVIST 240, IOPROMIDE
ULTRAVIST 300, IOPROMIDE
ULTRAVIST 370, IOPROMIDE
YASMIN, DROSPIRENONE

BERLEX LABS

* BERLEX LABORATORIES INC
ACUTECT, TECHNETIUM TC-99M APCITIDE
MENOSTAR, ESTRADIOL
NEO TECT KIT, TECHNETIUM TC-99M DEPREOTIDE

BIO NUCLEONICS

* BIO NUCLEONICS INC
STRONTIUM CHLORIDE SR-89, STRONTIUM CHLORIDE, SR-89

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

BIOCHEMIE

- * BIOCHEMIE GMBH
AMOXICILLIN, AMOXICILLIN
PENICILLIN G SODIUM, PENICILLIN G SODIUM
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM

BIOGLAN PHARMS CORP

- * BIOGLAN PHARMS CORP DBA DOAK DERMATOLOGICS
SOLARAZE, DICLOFENAC SODIUM

BIONICHE (CANADA)

- * BIONICHE PHARMA (CANADA) LTD
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
DIMETHYL SULFOXIDE, DIMETHYL SULFOXIDE
EDETATE DISODIUM, EDETATE DISODIUM
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
MILRINONE LACTATE, MILRINONE LACTATE

BIONICHE PHARM USA

- * BIONICHE PHARMA USA INC
SOTRADECOL, SODIUM TETRADECYL SULFATE

BIONICHE PHARMA

- * BIONICHE PHARMA
CYANOCOBALAMIN, CYANOCOBALAMIN
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

BIOVAIL

- * BIOVAIL CORP INTERNATIONAL
TIAZAC, DILTIAZEM HYDROCHLORIDE
- * BIOVAIL LABORATORIES INC
ATIVAN, LORAZEPAM
CARDIZEM CD, DILTIAZEM HYDROCHLORIDE
CARDIZEM, DILTIAZEM HYDROCHLORIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
GLUMETZA, METFORMIN HYDROCHLORIDE
ISORDIL, ISOSORBIDE DINITRATE
NIFEDIPINE, NIFEDIPINE
PENTOXIFYLLINE, PENTOXIFYLLINE
TECZEM, DILTIAZEM MALATE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

BIOVAIL LABS INTL

- * BIOVAIL LABORATORIES INTERNATIONAL SRL
CARDIZEM LA, DILTIAZEM HYDROCHLORIDE
CARDIZEM, DILTIAZEM HYDROCHLORIDE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
TRAMADOL HCL, TRAMADOL HYDROCHLORIDE
VASERETIC, ENALAPRIL MALEATE
VASOTEC, ENALAPRIL MALEATE
VASOTEC, ENALAPRILAT

BLAIREX

- * BLAIREX LABORATORIES INC
BRONCHO SALINE, SODIUM CHLORIDE (OTC)

BOCA PHARMA

- * BOCA PHARMACAL INC
HYDROCORTISONE ACETATE 1% AND PRAMOXINE HYDROCHLORIDE 1%, HYDROCORTISONE ACETATE

BOEHRINGER INGELHEIM

- * BOEHRINGER INGELHEIM
CATAPRES, CLONIDINE HYDROCHLORIDE
CATAPRES-TTS-1, CLONIDINE
CATAPRES-TTS-2, CLONIDINE
CATAPRES-TTS-3, CLONIDINE
MEXITIL, MEXILETINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

- * BOEHRINGER INGELHEIM
MICARDIS HCT, HYDROCHLOROTHIAZIDE
MICARDIS, TELMISARTAN
MIRAPEX, PRAMIPEXOLE DIHYDROCHLORIDE
- * BOEHRINGER INGELHEIM PHARMACEUTICALS INC
AGGRENEX, ASPIRIN
ALUPENT, METAPROTERENOL SULFATE
APTIVUS, TIPRANAVIR
ATROVENT HFA, IPRATROPIUM BROMIDE
ATROVENT, IPRATROPIUM BROMIDE
COMBIVENT, ALBUTEROL SULFATE
FLOMAX, TAMSULOSIN HYDROCHLORIDE
MOBIC, MELOXICAM
PERSANTINE, DIPYRIDAMOLE
SPIRIVA, TIOTROPIUM BROMIDE MONOHYDRATE
VIRAMUNE, NEVIRAPINE

BRACCO

- * BRACCO DIAGNOSTICS INC
CARDIOGEN-82, RUBIDIUM CHLORIDE RB-82
CHOLETEC, TECHNETIUM TC-99M MEBROFENIN KIT
CHOLOGRAFIN MEGLUMINE, IODIPAMIDE MEGLUMINE
CHROMITOPES SODIUM, SODIUM CHROMATE, CR-51
CYSTOGRAFIN DILUTE, DIATRIZOATE MEGLUMINE
CYSTOGRAFIN, DIATRIZOATE MEGLUMINE
GASTROGRAFIN, DIATRIZOATE MEGLUMINE
IODOTOPE, SODIUM IODIDE, I-131
ISOVUE-200, IOPAMIDOL
ISOVUE-250, IOPAMIDOL
ISOVUE-300, IOPAMIDOL
ISOVUE-370, IOPAMIDOL
ISOVUE-M 200, IOPAMIDOL
ISOVUE-M 300, IOPAMIDOL
KINEVAC, SINCALIDE
MACROTEC, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
MDP-BRACCO, TECHNETIUM TC-99M MEDRONATE KIT
MULTIHANCE MULTIPACK, GADOBENATE DIMEGLUMINE
MULTIHANCE, GADOBENATE DIMEGLUMINE
PHOSPHOTEC, TECHNETIUM TC-99M PYROPHOSPHATE KIT
PROHANCE MULTIPACK, GADOTERIDOL
PROHANCE, GADOTERIDOL
RENO-30, DIATRIZOATE MEGLUMINE
RENO-60, DIATRIZOATE MEGLUMINE
RENOCAL-76, DIATRIZOATE MEGLUMINE
RENO-DIP, DIATRIZOATE MEGLUMINE
RENOGRAFIN-60, DIATRIZOATE MEGLUMINE
RENOGRAFIN-76, DIATRIZOATE MEGLUMINE
RUBRATOPE-57 KIT, COBALT CHLORIDE, CO-57
RUBRATOPE-57, CYANOCOBALAMIN, CO-57
SINOGRAPHIN, DIATRIZOATE MEGLUMINE

BRADLEY PHARMS

- * BRADLEY PHARMACEUTICALS INC
PAMINE FORTE, METHSCOPOLAMINE BROMIDE
PAMINE, METHSCOPOLAMINE BROMIDE
ZONALON, DOXEPIN HYDROCHLORIDE

BRAINTREE

- * BRAINTREE LABORATORIES INC
AXID, NIZATIDINE
GOLYTELY, POLYETHYLENE GLYCOL 3350
HALFLYTELY, BISACODYL
MIRALAX, POLYETHYLENE GLYCOL 3350
NULYTELY, POLYETHYLENE GLYCOL 3350
NULYTELY-FLAVORED, POLYETHYLENE GLYCOL 3350

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** B ****

BRIGHTON PHARMS INC

* BRIGHTON PHARMACEUTICALS INC
BROMFED-DM, BROMPHENIRAMINE MALEATE

BRIGHTSTONE

* BRIGHTSTONE PHARMA INC
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

BRISTOL

* BRISTOL LABORATORIES INC DIV BRISTOL MYERS CO
BICNU, CARMUSTINE
BRISTACYCLINE, TETRACYCLINE HYDROCHLORIDE
LYSODREN, MITOTANE
TETREX, TETRACYCLINE PHOSPHATE COMPLEX

BRISTOL MYERS

* BRISTOL MYERS CO
MUTAMYCIN, MITOMYCIN
PLATINOL, CISPLATIN
PLATINOL-AQ, CISPLATIN
QUESTRAN LIGHT, CHOLESTYRAMINE
QUESTRAN, CHOLESTYRAMINE

BRISTOL MYERS SQUIBB

* BRISTOL MYERS SQUIBB
AZACTAM, AZTREONAM
BARACLUDE, ENTECAVIR
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.075% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.224% IN PLASTIC CONTAINER, DEXTROSE
DEXTROSE 5% AND POTASSIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
GLUCOVANCE, GLYBURIDE
MEGACE, MEGESTROL ACETATE
MONOPRIL-HCT, FOSINOPRIL SODIUM
PRAVACHOL, PRAVASTATIN SODIUM
TEQUIN, GATIFLOXACIN

* BRISTOL MYERS SQUIBB CO
AZACTAM IN PLASTIC CONTAINER, AZTREONAM
CEENU, LOMUSTINE
DROXIA, HYDROXYUREA
ESTRACE, ESTRADIOL
GLUCOPHAGE XR, METFORMIN HYDROCHLORIDE
HYDREA, HYDROXYUREA
KENACORT, TRIAMCINOLONE
METAGLIP, GLIPIZIDE
PARAPLATIN, CARBOPLATIN
PRAVIGARD PAC (COPACKAGED), ASPIRIN
PROLIXIN DECANOATE, FLUPHENAZINE DECANOATE
REYATAZ, ATAZANAVIR SULFATE
RUBEX, DOXORUBICIN HYDROCHLORIDE
RUBRAMIN PC, CYANOCOBALAMIN
SUSTIVA, EFAVIRENZ
TESLAC, TESTOLACTONE
VEPESID, ETOPOSIDE
VIDEX EC, DIDANOSINE

* BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
BLENOXANE, BLEOMYCIN SULFATE
BUSPAR, BUSPIRONE HYDROCHLORIDE
CEFZIL, CEFPROZIL
CYTOXAN, CYCLOPHOSPHAMIDE
DOVONEX, CALCIPOTRIENE
ESTRACE, ESTRADIOL
ETOPOPHOS PRESERVATIVE FREE, ETOPOSIDE PHOSPHATE
GLUCOPHAGE, METFORMIN HYDROCHLORIDE
IFEX/MESNEX KIT, IFOSFAMIDE
LYOPHILIZED CYTOXAN, CYCLOPHOSPHAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** B **

- * BRISTOL MYERS SQUIBB CO PHARMACEUTICAL RESEARCH INSTITUTE
MAXIPIME, CEFEPIME HYDROCHLORIDE (ARGININE FORMULATION)
MONOPRIL, FOSINOPRIL SODIUM
PARAPLATIN, CARBOPLATIN
TAXOL, PACLITAXEL
TEQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER, GATIFLOXACIN
TEQUIN, GATIFLOXACIN
VIDEX, DIDANOSINE
VUMON, TENIPOSIDE
ZERIT, STAVUDINE
- * BRISTOL MYERS SQUIBB MEDICAL IMAGING
DEFINITY, PERFLUTREN
GALLIUM CITRATE GA 67, GALLIUM CITRATE, GA-67
NEUROLITE, TECHNETIUM TC-99M BICISATE KIT
TECHNELITE, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE, TL-201
XENON XE 133, XENON, XE-133
- * BRISTOL MYERS SQUIBB PHARMA CO
CARDIOLITE, TECHNETIUM TC-99M SESTAMIBI KIT
COUMADIN, WARFARIN SODIUM
LODOSYN, CARBIDOPA
MIRALUMA, TECHNETIUM TC-99M SESTAMIBI KIT
SINEMET CR, CARBIDOPA
SINEMET, CARBIDOPA
SUSTIVA, EFAVIRENZ

BRYAN

- * BRYAN CORP
SCLEROSOL, TALC
TALC, TALC

BUNDY

- * CM BUNDY CO
BUTABARBITAL, BUTABARBITAL SODIUM
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE

C AND M PHARMA

- * C AND M PHARMACAL INC
HI-COR, HYDROCORTISONE

CANYON

- * CANYON PHARMACEUTICALS INC
IPRIVASK, DESIRUDIN RECOMBINANT

CARACO

- * CARACO PHARMACEUTICAL LABORATORIES LTD
CARBAMAZEPINE, CARBAMAZEPINE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLONAZEPAM, CLONAZEPAM
CLOZAPINE, CLOZAPINE
DIGOXIN, DIGOXIN
FLURBIPROFEN, FLURBIPROFEN
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MIRTAZAPINE, MIRTAZAPINE
OXAPROZIN, OXAPROZIN
PAROMOMYCIN SULFATE, PAROMOMYCIN SULFATE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

CARLSBAD

- * CARLSBAD TECHNOLOGY INC
ACYCLOVIR, ACYCLOVIR
CEFACTOR, CEFACLOX
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** C ****

* CARLSBAD TECHNOLOGY INC
DICLOFENAC SODIUM, DICLOFENAC SODIUM
FAMOTIDINE, FAMOTIDINE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
LOVASTATIN, LOVASTATIN

CAROLINA MEDCL

* CAROLINA MEDICAL PRODUCTS CO
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
HYDROCORTISONE IN ABSORBASE, HYDROCORTISONE
ISONIAZID, ISONIAZID
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
SPS, SODIUM POLYSTYRENE SULFONATE
TRIAMCINOLONE ACETONIDE IN ABSORBASE, TRIAMCINOLONE ACETONIDE

CEDAR PHARMS

* CEDAR PHARMACEUTICALS LLC
METHIMAZOLE, METHIMAZOLE

CELGENE

* CELGENE CORP
REVLIMID, LENALIDOMIDE
THALOMID, THALIDOMIDE

CEPH INTL

* CEPH INTERNATIONAL CORP
CEFACLOX, CEFACLOX
CEPHALEXIN, CEPHALEXIN

CEPHALON

* CEPHALON INC
ACTIQ, FENTANYL CITRATE
GABITRIL, TIAGABINE HYDROCHLORIDE
PROVIGIL, MODAFINIL
TRISENOX, ARSENIC TRIOXIDE

CHATTEM

* CHATTEM INC
SELSUN, SELENIUM SULFIDE

CHESEBROUGH PONDS

* CHESEBROUGH PONDS INC
EXTRA-STRENGTH AIM, SODIUM MONOFLUOROPHOSPHATE (OTC)

CHIRHOCLIN

* CHIRHOCLIN INC
CHIRHOSTIM, SECRETIN SYNTHETIC HUMAN
SECREMAX, SECRETIN SYNTHETIC PORCINE

CHIRON

* CHIRON CORP
TOBI, TOBRAMYCIN

CIS

* CIS US INC
AN-DTPA, TECHNETIUM TC-99M PENTETATE KIT
AN-SULFUR COLLOID, TECHNETIUM TC-99M SULFUR COLLOID KIT
CIS-MDP, TECHNETIUM TC-99M MEDRONATE KIT
CIS-PYRO, TECHNETIUM TC-99M PYROPHOSPHATE KIT
HEPATOLITE, TECHNETIUM TC-99M DISOFENIN KIT
IOBENGUANE SULFATE I 131, IOBENGUANE SULFATE I 131
PULMOLITE, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

CLAY PARK

* CLAY PARK LABORATORIES INC
AMMONIUM LACTATE, AMMONIUM LACTATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
DESOXIMETASONE, DESOXIMETASONE
ECONAZOLE NITRATE, ECONAZOLE NITRATE
ERYTHROMYCIN, ERYTHROMYCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** C ****

- * CLAY PARK LABORATORIES INC
 - FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
 - GENTAMICIN, GENTAMICIN SULFATE
 - HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
 - HYDROCORTISONE, HYDROCORTISONE
 - KETOCONAZOLE, KETOCONAZOLE
 - MESALAMINE, MESALAMINE
 - MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 - MOMETASONE FUROATE, MOMETASONE FUROATE
 - MUPIROCI, MUPIROCI
 - NYSTATIN, NYSTATIN
 - PERMETHRIN, PERMETHRIN
 - PERMETHRIN, PERMETHRIN (OTC)
 - PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 - SELENIUM SULFIDE, SELENIUM SULFIDE
 - TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
 - TRIDESILON, DESONIDE
- * CLAY PARK LABS INC
 - DESOXIMETASONE, DESOXIMETASONE

CLINTEC NUTR

- * CLINTEC NUTRITION CO SUB CLINIQUE
 - CLINISOL 15% SULFITE FREE IN PLASTIC CONTAINER, AMINO ACIDS

CLONMEL

- * CLONMEL HEALTHCARE
 - ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
 - AMOXICILLIN, AMOXICILLIN
 - AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
 - PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM

CLONMEL HLTHCARE

- * CLONMEL HEALTHCARE LTD
 - ACYCLOVIR, ACYCLOVIR
 - ATENOLOL, ATENOLOL
 - CAPTOPRIL, CAPTOPRIL
 - CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 - CIMETIDINE, CIMETIDINE
 - CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 - DIAZEPAM, DIAZEPAM
 - DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 - DIPYRIDAMOLE, DIPYRIDAMOLE
 - DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 - FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 - GLYBURIDE (MICRONIZED), GLYBURIDE
 - HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - INDOMETHACIN, INDOMETHACIN
 - KETOPROFEN, KETOPROFEN
 - METHOTREXATE SODIUM, METHOTREXATE SODIUM
 - NAPROXEN, NAPROXEN
 - PENTOXIFYLLINE, PENTOXIFYLLINE
 - PRazosin HCL, PRAZOSIN HYDROCHLORIDE
 - PROPYLTHIOURACIL, PROPYLTHIOURACIL
 - PYRAZINAMIDE, PYRAZINAMIDE
 - QUINIDINE SULFATE, QUINIDINE SULFATE
 - SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

COBALT

- * COBALT PHARMACEUTICALS INC
 - CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
 - CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 - METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 - RAMIPRIL, RAMIPRIL
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** C **

COLGATE

- * COLGATE ORAL PHARMACEUTICALS INC
ORABASE HCA, HYDROCORTISONE ACETATE
PERIOGARD, CHLORHEXIDINE GLUCONATE

COLGATE PALMOLIVE

- * COLGATE PALMOLIVE
COLGATE TOTAL, SODIUM FLUORIDE (OTC)

COLLAGENEX PHARMS

- * COLLAGENEX PHARMACEUTICALS INC
PERIOSTAT, DOXYCYCLINE HYCLATE

COLUMBIA LABS

- * COLUMBIA LABORATORIES INC
STRIANT, TESTOSTERONE

COLUMBIA RES LABS

- * COLUMBIA RESEARCH LABORATORIES INC
CRINONE, PROGESTERONE

CONNETICS

- * CONNETICS CORP
EVOCLIN, CLINDAMYCIN PHOSPHATE
LUXIQ, BETAMETHASONE VALERATE
OLUX, CLOBETASOL PROPIONATE
SORIATANE, ACITRETIN

CONSOLIDATED PHARM

- * CONSOLIDATED PHARMACEUTICAL GROUP INC
AMOXICILLIN, AMOXICILLIN
PENICILLIN V POTASSIUM, PENICILLIN V POTASSIUM

CONTROLLED THERAP

- * CONTROLLED THERAPEUTICS (SCOTLAND) LTD
CERVIDIL, DINOPROSTONE

COOK IMAGING

- * COOK IMAGING CORP
IOPAMIDOL-200, IOPAMIDOL
IOPAMIDOL-250, IOPAMIDOL
IOPAMIDOL-300, IOPAMIDOL
IOPAMIDOL-370, IOPAMIDOL

COPLEY PHARM

- * COPLEY PHARMACEUTICAL INC
CO-LAV, POLYETHYLENE GLYCOL 3350
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
GO-EVAC, POLYETHYLENE GLYCOL 3350
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
NABUMETONE, NABUMETONE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

COREPHARMA

- * COREPHARMA LLC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CARISOPRODOL, CARISOPRODOL
CILOSTAZOL, CILOSTAZOL
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
ASPARTATE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
GLIMEPIRIDE, GLIMEPIRIDE
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
GLYBURIDE, GLYBURIDE
GLYCOPYRROLATE, GLYCOPYRROLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** C ****

* COREPHARMA LLC
LEVOCARNITINE, LEVOCARNITINE
METHENAMINE HIPPURATE, METHENAMINE HIPPURATE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

COTHERIX

* COTHERIX INC
VENTAVIS, ILOPROST

CP PHARMS

* CP PHARMACEUTICALS CV
LYRICA, PREGABALIN

CRITICAL

* CRITICAL THERAPEUTICS INC
ZYFLO, ZILEUTON

CUBIST

* CUBIST PHARMACEUTICALS INC
CUBICIN, DAPTOMYCIN

CUMBERLAND PHARMS

* CUMBERLAND PHARMACEUTICALS INC
ACETADOTE, ACETYLCYSTEINE

CYTOGEN

* CYTOGEN CORP
QUADRAMET, SAMARIUM SM 153 LEXIDRONAM PENTASODIUM

DAIICHI

* DAIICHI MEDICAL RESEARCH INC
EVOXAC, CEVIMELINE HYDROCHLORIDE
* DAIICHI PHARMACEUTICAL CORP
FLOXIN OTIC, OFLOXACIN

DAINIPPON

* DAINIPPON PHARMACEUTICAL USA CORP
ZONEGRAN, ZONISAMIDE

DANCO LABS LLC

* DANCO LABORATORIES LLC
MIFEPREX, MIFEPRISTONE

DAVIS AND GECK

* DAVIS AND GECK DIV AMERICAN CYANAMID CO
PRE-OP II, HEXACHLOROPHENE
PRE-OP, HEXACHLOROPHENE

DEL RAY LABS

* DEL RAY LABORATORIES INC
ALA-CORT, HYDROCORTISONE
ALA-SCALP, HYDROCORTISONE
TRIDERM, TRIAMCINOLONE ACETONIDE

DELL LABS

* DELL LABORATORIES INC
LIDOCAINE HYDROCHLORIDE W/ EPINEPHRINE, EPINEPHRINE

DENTSPLY PHARM

* DENTSPLY PHARMACEUTICAL
CITANEST FORTE, EPINEPHRINE BITARTRATE
CITANEST PLAIN, PRILOCAINE HYDROCHLORIDE
ORAQIX, LIDOCAINE
POLOCAINE W/ LEVONORDEFIN, LEVONORDEFIN
POLOCAINE, MEPIVACAINE HYDROCHLORIDE
XYLOCAINE W/ EPINEPHRINE, EPINEPHRINE
XYLOCAINE, LIDOCAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** D ****

DEPROCO

- * DEPROCO INC
LIGNOSPAN FORTE, EPINEPHRINE BITARTRATE
LIGNOSPAN STANDARD, EPINEPHRINE BITARTRATE
SCANDONEST L, LEVONORDEFIN
SCANDONEST PLAIN, MEPIVACAINE HYDROCHLORIDE
SEPTOCAINE, ARTICAINE HYDROCHLORIDE

DERMIK LABS

- * DERMILABORATORIES DIV AVENTIS PHARMACEUTICALS INC
BENZAOLIN, BENZOYL PEROXIDE
BENZAMYCIN PAK, BENZOYL PEROXIDE
BENZAMYCIN, BENZOYL PEROXIDE
CARAC, FLUOROURACIL
DERMATOP, PREDNICARBATE
HYTONE, HYDROCORTISONE
KLARON, SULFACETAMIDE SODIUM
NORITATE, METRONIDAZOLE
PSORCON, DIFLORASONE DIACETATE
- * DERMILABORATORIES INC
DERMATOP E EMOLLIENT, PREDNICARBATE
PENLAC, CICLOPIROX
- * DERMILABORATORIES INC SUB RORER
HYTONE, HYDROCORTISONE

DESERET

- * DESERET MEDICAL INC DIV BECTON DICKINSON AND CO
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

DEXCEL LTD

- * DEXCEL LTD
DICLOFENAC SODIUM, DICLOFENAC SODIUM
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE

DEXCEL PHARMA

- * DEXCEL PHARMA TECHNOLOGIES LTD
PERIOCHIP, CHLORHEXIDINE GLUCONATE

DEY

- * DEY LP
ACCUNEB, ALBUTEROL SULFATE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
CROMOLYN SODIUM, CROMOLYN SODIUM
CUROSURF, PORACTANT ALFA
DUONEB, ALBUTEROL SULFATE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE

DHL

- * DHL LABORATORIES INC
DEXTRORSE 5% IN PLASTIC CONTAINER, DEXTRORSE

DORC

- * DORC INTERNATIONAL BV
VISIONBLUE, TRYPAN BLUE

DOW PHARM SCI

- * DOW PHARMACEUTICAL SCIENCES
CLOBEX, CLOBETASOL PROPIONATE

DPT

- * DPT LABORATORIES INC
EMBELINE, CLOBETASOL PROPIONATE

DR REDDYS LABS INC

- * DR REDDYS LABORATORIES INC
FLUCONAZOLE, FLUCONAZOLE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** D **

* DR REDDYS LABORATORIES INC
IBUPROFEN, IBUPROFEN (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE

DR REDDYS LABS LTD

* DR REDDYS LABORATORIES LTD
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE (OTC)
GLIMEPIRIDE, GLIMEPIRIDE
NIZATIDINE, NIZATIDINE
OXAPROZIN, OXAPROZIN
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
ZONISAMIDE, ZONISAMIDE

DRAXIMAGE

* DRAXIMAGE INC
DRAXIMAGE MDP, TECHNETIUM TC-99M MEDRONATE KIT
DTPA, TECHNETIUM TC-99M PENTETATE KIT
SODIUM IODIDE I 131, KIT, SODIUM IODIDE, I-131
TECHNISCAN GLUCEPTATE, TECHNETIUM TC-99M GLUCEPTATE KIT
TECHNETIUM TC 99M ALBUMIN AGGREGATED KIT, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

DURAMED

* DURAMED PHARMACEUTICALS INC
CENESTIN, ESTROGENS, CONJUGATED SYNTHETIC A
NORDETTE-28, ETHINYL ESTRADIOL
PLAN B, LEVONORGESTREL
PREFEST, ESTRADIOL
REVIA, NALTREXONE HYDROCHLORIDE
SEASONALE, ETHINYL ESTRADIOL

DURAMED PHARMS BARR

* DURAMED PHARMACEUTICALS INC SUB BARR LABORATORIES INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
AVIANE-28, ETHINYL ESTRADIOL
AYGESTIN, NORETHINDRONE ACETATE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
COPPER T MODEL TCU 380A, COPPER
CRYSELLE, ETHINYL ESTRADIOL
DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
DIAMOX, ACETAZOLAMIDE
DIAMOX, ACETAZOLAMIDE SODIUM
ENPRESSE-28, ETHINYL ESTRADIOL
ESTROPIPATE, ESTROPIPATE
GYNODIOL, ESTRADIOL
HYDROXYUREA, HYDROXYUREA
ISONIAZID, ISONIAZID
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
METHYLPREDNISOLONE, METHYLPREDNISOLONE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VELIVET, DESOGESTREL
ZEBETA, BISOPROLOL FUMARATE
ZIAC, BISOPROLOL FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** D **

DUSA

- * DUSA PHARMACEUTICALS INC
LEVULAN, AMINOLEVULINIC ACID HYDROCHLORIDE

EASTMAN KODAK

- * EASTMAN KODAK CO
CARBOCAINE W/ NEO-COBEFRIN, LEVONORDEFIN
CARBOCAINE, MEPIVACAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE

ECOLAB

- * ECOLAB INC
CHG SCRUB, CHLORHEXIDINE GLUCONATE (OTC)
CIDA-STAT, CHLORHEXIDINE GLUCONATE (OTC)

EDWARDS LIFESCIENCES

- * EDWARDS LIFESCIENCES RESEARCH MEDICAL INC
RIMS0-50, DIMETHYL SULFOXIDE

EGIS

- * EGIS PHARMACEUTICALS
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
PIROXICAM, PIROXICAM

EGIS PHARMS

- * EGIS PHARMACEUTICALS LTD
CAPTOPRIL, CAPTOPRIL

EISAI MEDCL RES

- * EISAI MEDICAL RESEARCH INC
ACIPHEX, RABEPRAZOLE SODIUM
ARICEPT ODT, DONEPEZIL HYDROCHLORIDE
ARICEPT, DONEPEZIL HYDROCHLORIDE

ELAN DRUG

- * ELAN DRUG DELIVERY INC
VERELAN PM, VERAPAMIL HYDROCHLORIDE
VERELAN, VERAPAMIL HYDROCHLORIDE

ELAN PHARM

- * ELAN PHARMACEUTICAL RESEARCH CORP
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
KETOPROFEN, KETOPROFEN
NIFEDIPINE, NIFEDIPINE

ELAN PHARMS

- * ELAN PHARMACEUTICALS INC
PRIALT, ZICONOTIDE

ELKINS SINN

- * ELKINS SINN DIV AH ROBINS CO INC
DEXAMETHASONE, DEXAMETHASONE SODIUM PHOSPHATE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
PANCURONIUM, PANCURONIUM BROMIDE
PHENYTOIN, PHENYTOIN SODIUM

EMPI

- * EMPI INC
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE

ENCYSIVE

- * ENCYSIVE PHARMACEUTICALS INC
ARGATROBAN, ARGATROBAN

ENDO LABS

- * ENDO LABORATORIES INC DIV DUPONT MERCK PHARMACEUTICAL CO
CAPTOPRIL, CAPTOPRIL
- * ENDO LABORATORIES LLC
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** E **

ENDO PHARMS

* ENDO PHARMACEUTICALS INC
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CIMETIDINE, CIMETIDINE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
ETODOLAC, ETODOLAC
FROVA, FROVATRIPTAN SUCCINATE
GLIPIZIDE, GLIPIZIDE
HYCODAN, HOMATROPINE METHYLBROMIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
MOBAN, MOLINDONE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
NARCAN, NALOXONE HYDROCHLORIDE
NUBAIN, NALBUPHINE HYDROCHLORIDE
NUMORPHAN, OXYMORPHONE HYDROCHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PERCOCET, ACETAMINOPHEN
PERCODAN, ASPIRIN
PERCODAN-DEMI, ASPIRIN
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SYMMETREL, AMANTADINE HYDROCHLORIDE

ENZON

* ENZON INC
ABELCET, AMPHOTERICIN B
ADAGEN, PEGADEMASE BOVINE

ESP PHARMA

* ESP PHARMA INC
BUSULFEX, BUSULFAN
CARDENE, NICARDIPINE HYDROCHLORIDE
DECILOMYCIN, DEMECLOCYCLINE HYDROCHLORIDE
ISMO, ISOSORBIDE MONONITRATE
SECTRAL, ACEBUTOLOL HYDROCHLORIDE
TENEX, GUANFACINE HYDROCHLORIDE

ESPRIT PHARMA

* ESPRIT PHARMA INC
PROQUIN XR, CIPROFLOXACIN HYDROCHLORIDE

EURAND

* EURAND AMERICA INC
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE

EVERYLIFE

* EVERYLIFE
HYDROCORTISONE, HYDROCORTISONE
SECOBARBITAL SODIUM, SECOBARBITAL SODIUM

EXCELLIUM

* EXCELLIUM PHARMACEUTICAL INC
FUROSEMIDE, FUROSEMIDE

EYETECH PHARMS

* EYETECH PHARMACEUTICALS INC
MACUGEN, PEGAPTANIB SODIUM

FALCON PHARMS

* FALCON PHARMACEUTICALS INC
TIMOLOL MALEATE, TIMOLOL MALEATE
* FALCON PHARMACEUTICALS LTD
DIPIVEFRIN HYDROCHLORIDE, DIPIVEFRIN HYDROCHLORIDE
ECONOPRED PLUS, PREDNISOLONE ACETATE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
MAXITROL, DEXAMETHASONE
METIPRANOLOL, METIPRANOLOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** F ****

- * FALCON PHARMACEUTICALS LTD
TIMOLOL MALEATE, TIMOLOL MALEATE
TOBREX, TOBRAMYCIN
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

FAULDING

- * FAULDING PHARMACEUTICALS SUB FH FAULDING AND CO LTD
DORYX, DOXYCYCLINE HYCLATE

FERNDALE LABS

- * FERNDALE LABORATORIES INC
HYDROCORTISONE ACETATE, HYDROCORTISONE ACETATE
LOCOID LIPOCREAM, HYDROCORTISONE BUTYRATE
LOCOID, HYDROCORTISONE BUTYRATE
MICORT-HC, HYDROCORTISONE ACETATE
PRAMOSONE, HYDROCORTISONE ACETATE
STRIFON FORTE DSC, CHLORZOXAZONE

FERRING

- * FERRING PHARMACEUTICALS INC
ACTHREL, CORTICORELIN OVINE TRIFLUTATE
BRAVELLE, UROFOLLITROPIN
MENOPUR, LUTEINIZING HORMONE
MINIRIN, DESMOPRESSIN ACETATE
REPRONEX, LUTEINIZING HORMONE
TEV-TROPIN, SOMATROPIN RECOMBINANT

FH FAULDING CO LTD

- * FH FAULDING CO LTD TRADING MAYNE PHARMA INTERNATIONAL
DORYX, DOXYCYCLINE HYCLATE

FIRST HORIZON

- * FIRST HORIZON PHARMACEUTICAL CORP
COGNEX, TACRINE HYDROCHLORIDE
FURADANTIN, NITROFURANTOIN
PONSTEL, MEFENAMIC ACID
ROBINUL FORTE, GLYCOPYRROLATE
ROBINUL, GLYCOPYRROLATE
SULAR, NISOLDIPINE

FLEMING

- * FLEMING AND CO PHARMACEUTICALS
THYROSHIELD, POTASSIUM IODIDE (OTC)

FLEMINGTON PHARM

- * FLEMINGTON PHARMACEUTICAL CORP
NIFEDIPINE, NIFEDIPINE

FOREST LABS

- * FOREST LABORATORIES INC
AMBENYL, BROMODIPHENHYDRAMINE HYDROCHLORIDE
CAMPRAL, ACAMPROSATE CALCIUM
CELEXA, CITALOPRAM HYDROBROMIDE
COMBUNOX, IBUPROFEN
ELIXOPHYLLIN, THEOPHYLLINE
FLUMADINE, RIMANTADINE HYDROCHLORIDE
LEXAPRO, ESCITALOPRAM OXALATE
NAMENDA, MEMANTINE HYDROCHLORIDE
TESSALON, BENZONATATE
THYROLAR-0.25, LIOETHYRONINE SODIUM
THYROLAR-0.5, LIOETHYRONINE SODIUM
THYROLAR-1, LIOETHYRONINE SODIUM
THYROLAR-2, LIOETHYRONINE SODIUM
THYROLAR-3, LIOETHYRONINE SODIUM

FOUGERA

- * E FOUGERA DIV ALTANA INC
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** F **

- * E FOUGERA DIV ALTANA INC
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
ERYTHROMYCIN, ERYTHROMYCIN
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
HYDROCORTISONE, HYDROCORTISONE
LIDOCAINE, LIDOCAINE
NEOMYCIN AND POLYMYXIN B SULFATES AND DEXAMETHASONE, DEXAMETHASONE
NITROGLYCERIN, NITROGLYCERIN
NYSTATIN, NYSTATIN
NYSTATIN-TRIAMCINOLONE ACETONIDE, NYSTATIN

FRESENIUS

- * FRESENIUS KABI DEUTSCHLAND GMBH
INTRALIPID 10%, SOYBEAN OIL
INTRALIPID 20%, SOYBEAN OIL
INTRALIPID 30%, SOYBEAN OIL
- * FRESENIUS USA INC
INPERSOL-LC/LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
INPERSOL-LC/LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

FRESENIUS MEDCL

- * FRESENIUS MEDICAL CARE NORTH AMERICA
Delflex W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 1.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 1.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 2.5% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 2.5% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 4.25% LOW MAGNESIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex W/ DEXTROSE 4.25% LOW MAGNESIUM LOW CALCIUM IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex-LM W/ DEXTROSE 1.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex-LM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex-LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER, CALCIUM CHLORIDE
Delflex-LM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

FSC

- * FSC LABORATORIES INC
ISOPTIN, VERAPAMIL HYDROCHLORIDE
PRIMSOL, TRIMETHOPRIM HYDROCHLORIDE

G AND W LABS

- * G AND W LABORATORIES INC
ACEPHEN, ACETAMINOPHEN (OTC)
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
INDOMETHEGAN, INDOMETHACIN
MICONAZOLE 7 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
MIGERGOT, CAFFEINE
PROCHLORPERAZINE, PROCHLORPERAZINE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROMETHEGAN, PROMETHAZINE HYDROCHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TRUPHYLLINE, AMINOPHYLLINE

GALDERMA LABS

- * GALDERMA LABORATORIES INC
CLOBEX, CLOBETASOL PROPIONATE

GALDERMA LABS LP

- * GALDERMA LABORATORIES L P
CLOBEX, CLOBETASOL PROPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** G ****

* GALDERMA LABORATORIES LP
CLINDAGEL, CLINDAMYCIN PHOSPHATE
DESOWEN, DESONIDE
DIFFERIN, ADAPALENE
FS SHAMPOO, FLUOCINOLONE ACETONIDE
METROCREAM, METRONIDAZOLE
METROGEL, METRONIDAZOLE
METROLOTION, METRONIDAZOLE
TRI-LUMA, FLUOCINOLONE ACETONIDE

GALLEN LTD

* GALLEN LTD
FEMRING, ESTRADIOL ACETATE

GD SEARLE LLC

* GD SEARLE LLC
ALDACTAZIDE, HYDROCHLOROTHIAZIDE
ALDACTONE, SPIRONOLACTONE
ARTHROTEC, DICLOFENAC SODIUM
BEXTRA, VALDECOXIB
CALAN, VERAPAMIL HYDROCHLORIDE
CELEBREX, CELECOXIB
COVERA-HS, VERAPAMIL HYDROCHLORIDE
CYTOTEC, MISOPROSTOL
DAYPRO, OXAPROZIN
DEMULEN 1/35-21, ETHINYL ESTRADIOL
DEMULEN 1/35-28, ETHINYL ESTRADIOL
DEMULEN 1/50-21, ETHINYL ESTRADIOL
DEMULEN 1/50-28, ETHINYL ESTRADIOL
FLAGYL ER, METRONIDAZOLE
FLAGYL I.V. RTU IN PLASTIC CONTAINER, METRONIDAZOLE
FLAGYL I.V., METRONIDAZOLE HYDROCHLORIDE
FLAGYL, METRONIDAZOLE
INSpra, EPLERENONE
LOMOTIL, ATROPINE SULFATE
NORPACE CR, DISOPYRAMIDE PHOSPHATE
NORPACE, DISOPYRAMIDE PHOSPHATE
SYNAREL, NAFARELIN ACETATE

GE HEALTHCARE

* GE HEALTHCARE
CERETEC, TECHNETIUM TC-99M EXAMETAZIME KIT
HYPAQUE, DIATRIZOATE MEGLUMINE
HYPAQUE, DIATRIZOATE SODIUM
HYPAQUE-76, DIATRIZOATE MEGLUMINE
HYPAQUE-CYSTO, DIATRIZOATE MEGLUMINE
INDIUM IN-111 OXYQUINOLINE, INDIUM IN-111 OXYQUINOLINE
METASTRON, STRONTIUM CHLORIDE, SR-89
MPI DMSA KIDNEY REAGENT, TECHNETIUM TC-99M SUCCIMER KIT
MPI DTPA KIT - CHELATE, TECHNETIUM TC-99M PENTETATE KIT
MPI INDIUM DTPA IN 111, INDIUM IN-111 PENTETATE DISODIUM
MYOVUE 30ML, TECHNETIUM TC-99M TETROFOSMIN KIT
MYOVUE, TECHNETIUM TC-99M TETROFOSMIN KIT
OMNIPAQUE 140, IOHEXOL
OMNIPAQUE 180, IOHEXOL
OMNIPAQUE 240, IOHEXOL
OMNIPAQUE 300, IOHEXOL
OMNIPAQUE 350, IOHEXOL
OMNISCAN, GADODIAMIDE
OPTISON, ALBUMIN HUMAN
SODIUM IODIDE I 123, SODIUM IODIDE, I-123
TECHNETIUM TC 99M MPI MDP, TECHNETIUM TC-99M MEDRONATE KIT
TECHNETIUM TC-99M PENTETATE KIT, TECHNETIUM TC-99M PENTETATE KIT
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE, TL-201
VISIPAQUE 270, IODIXANOL
VISIPAQUE 320, IODIXANOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** G **

GEDEON RICHTER USA

* GEDEON RICHTER USA INC
FLUCONAZOLE, FLUCONAZOLE

GENENTECH

* GENENTECH INC
NUTROPIN AQ PEN, SOMATROPIN RECOMBINANT
NUTROPIN AQ, SOMATROPIN RECOMBINANT
NUTROPIN DEPOT, SOMATROPIN RECOMBINANT
NUTROPIN, SOMATROPIN RECOMBINANT

GENPHARM

* GENPHARM INC
ACYCLOVIR, ACYCLOVIR
ALBUTEROL, ALBUTEROL
AMNESTEEM, ISOTRETINOIN
AZATHIOPRINE, AZATHIOPRINE
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CLARITHROMYCIN, CLARITHROMYCIN
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ETIDRONATE DISODIUM, ETIDRONATE DISODIUM
ETODOLAC, ETODOLAC
ETOPOSIDE, ETOPOSIDE
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE (OTC)
FLUCONAZOLE, FLUCONAZOLE
FLUTAMIDE, FLUTAMIDE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
LORATADINE, LORATADINE (OTC)
LOVASTATIN, LOVASTATIN
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHIMAZOLE, METHIMAZOLE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NIZATIDINE, NIZATIDINE
NOVOTHYROX, LEVOTHYROXINE SODIUM
OXAPROZIN, OXAPROZIN
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
WARFARIN SODIUM, WARFARIN SODIUM
* GENPHARM PHARMACEUTICALS INC
ATENOLOL, ATENOLOL
LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
PINDOLOL, PINDOLOL
PIROXICAM, PIROXICAM

GENTA

* GENTA INC
GANITE, GALLIUM NITRATE

GENZYME

* GENZYME CORP
CEREDASE, ALGLUCERASE
CEREZYME, IMIGLUCERASE
CLOLAR, CLOFARABINE
HECTOROL, DOXERCALCIFEROL
LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
RENAGEL, SEVELAMER HYDROCHLORIDE
THYROGEN, THYROTROPIN ALFA

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT**** G ******GILEAD**

* GILEAD SCIENCES INC
 DAUNOXOME, DAUNORUBICIN CITRATE
 EMTRIVA, EMTRICITABINE
 HEPSERA, ADEFOVIR DIPIVOXIL
 TRUVADA, EMTRICITABINE
 VIREAD, TENOFOVIR DISOPROXIL FUMARATE
 VISTIDE, CIDOFOVIR

GLADES PHARMS

* GLADES PHARMACEUTICALS INC
 E-GLADES, ERYTHROMYCIN

GLAXOSMITHKLINE

* GLAXOSMITHKLINE
 ABREVA, DOCOSANOL (OTC)
 ACLOVATE, ALCLOMETASONE DIPROPIONATE
 ADVAIR DISKUS 100/50, FLUTICASONE PROPIONATE
 ADVAIR DISKUS 250/50, FLUTICASONE PROPIONATE
 ADVAIR DISKUS 500/50, FLUTICASONE PROPIONATE
 AGENERASE, AMPRENAVIR
 ALBENZA, ALBENDAZOLE
 ALKERAN, MELPHALAN
 ALKERAN, MELPHALAN HYDROCHLORIDE
 AMERGE, NARATRIPTAN HYDROCHLORIDE
 AMOXIL, AMOXICILLIN
 ANCEF, CEFAZOLIN SODIUM
 ANSPOR, CEPHRADINE
 ARIXTRA, FONDAPARINUX SODIUM
 ARRANON, NELARABINE
 AUGMENTIN '125', AMOXICILLIN
 AUGMENTIN '200', AMOXICILLIN
 AUGMENTIN '250', AMOXICILLIN
 AUGMENTIN '400', AMOXICILLIN
 AUGMENTIN '500', AMOXICILLIN
 AUGMENTIN '875', AMOXICILLIN
 AUGMENTIN ES-600, AMOXICILLIN
 AUGMENTIN XR, AMOXICILLIN
 AVODART, DUTASTERIDE
 BACTOCILL, OXACILLIN SODIUM
 BACTROBAN, MUPIROCIN
 BACTROBAN, MUPIROCIN CALCIUM
 BECONASE AQ, BECLOMETHASONE DIPROPIONATE MONOHYDRATE
 CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
 CEFTIN, CEFUROXIME AXETIL
 CEPTAZ, CEFTAZIDIME (ARGININE FORMULATION)
 CLOXAPEN, CLOXACILLIN SODIUM
 COMBIVIR, LAMIVUDINE
 COMPAZINE, PROCHLORPERAZINE
 COMPAZINE, PROCHLORPERAZINE EDISYLATE
 COMPAZINE, PROCHLORPERAZINE MALEATE
 COREG, CARVEDILOL
 DARAPRIM, PYRIMETHAMINE
 DEXEDRINE, DEXTROAMPHETAMINE SULFATE
 DYAZIDE, HYDROCHLOROTHIAZIDE
 EPIVIR, LAMIVUDINE
 EPIVIR-HBV, LAMIVUDINE
 ESKALITH CR, LITHIUM CARBONATE
 ESKALITH, LITHIUM CARBONATE
 EXOSURF NEONATAL, CETYL ALCOHOL
 FLOLAN, EPOPROSTENOL SODIUM
 FLONASE, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 100, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 250, FLUTICASONE PROPIONATE
 FLOVENT DISKUS 50, FLUTICASONE PROPIONATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** G ****

* GLAXOSMITHKLINE
FLOVENT HFA, FLUTICASONE PROPIONATE
FLOVENT, FLUTICASONE PROPIONATE
FORTAZ IN PLASTIC CONTAINER, CEFTAZIDIME SODIUM
FORTAZ, CEFTAZIDIME
HYCANTIN, TOPOTECAN HYDROCHLORIDE
IMITREX, SUMATRIPTAN
IMITREX, SUMATRIPTAN SUCCINATE
LAMICTAL CD, LAMOTRIGINE
LAMICTAL, LAMOTRIGINE
LANOXIN PEDIATRIC, DIGOXIN
LANOXIN, DIGOXIN
LAROTID, AMOXICILLIN
LEXIVA, FOSAMPRENAVIR CALCIUM
LOTRONEX, ALOSETRON HYDROCHLORIDE
MALARONE PEDIATRIC, ATOVAQUONE
MALARONE, ATOVAQUONE
MEPRON, ATOVAQUONE
MYLERAN, BUSULFAN
NICORETTE (MINT), NICOTINE POLACRILEX (OTC)
NICORETTE (ORANGE), NICOTINE POLACRILEX (OTC)
NICORETTE, NICOTINE POLACRILEX (OTC)
PARNATE, TRANLYCYPROMINE SULFATE
PAXIL CR, PAROXETINE HYDROCHLORIDE
PAXIL, PAROXETINE HYDROCHLORIDE
RELAFEN, NABUMETONE
RELENZA, ZANAMIVIR
REQUIP, ROPINIROLE HYDROCHLORIDE
RETROVIR, ZIDOVUDINE
SEREVENT, SALMETEROL XINAFOATE
STELAZINE, TRIFLUOPERAZINE HYDROCHLORIDE
TAGAMET HB, CIMETIDINE (OTC)
TAGAMET, CIMETIDINE
THIOGUANINE, THIOGUANINE
THORAZINE, CHLORPROMAZINE
THORAZINE, CHLORPROMAZINE HYDROCHLORIDE
TICAR, TICARCILLIN DISODIUM
TIMENTIN IN PLASTIC CONTAINER, CLAVULANATE POTASSIUM
TIMENTIN, CLAVULANATE POTASSIUM
TRIZIVIR, ABACAVIR SULFATE
VALTREX, VALACYCLOVIR HYDROCHLORIDE
VENTOLIN HFA, ALBUTEROL SULFATE
VENTOLIN, ALBUTEROL
WELLBUTRIN SR, BUPROPION HYDROCHLORIDE
WELLBUTRIN, BUPROPION HYDROCHLORIDE
ZANTAC 150, RANITIDINE HYDROCHLORIDE
ZANTAC 25, RANITIDINE HYDROCHLORIDE
ZANTAC 300, RANITIDINE HYDROCHLORIDE
ZANTAC IN PLASTIC CONTAINER, RANITIDINE HYDROCHLORIDE
ZANTAC, RANITIDINE HYDROCHLORIDE
ZIAGEN, ABACAVIR SULFATE
ZINACEF IN PLASTIC CONTAINER, CEFUROXIME SODIUM
ZINACEF, CEFUROXIME SODIUM
ZOFRAN IN PLASTIC CONTAINER, ONDANSETRON HYDROCHLORIDE
ZOFRAN ODT, ONDANSETRON
ZOFRAN PRESERVATIVE FREE, ONDANSETRON HYDROCHLORIDE
ZOFRAN, ONDANSETRON HYDROCHLORIDE
ZOVIRAX, ACYCLOVIR
ZOVIRAX, ACYCLOVIR SODIUM
ZYBAN, BUPROPION HYDROCHLORIDE

GLAXOSMITHKLINE CONS

* GLAXOSMITHKLINE CONSUMER HEALTHCARE
COMMIT, NICOTINE POLACRILEX (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** G ****

GLENMARK PHARMA

* GLENMARK PHARMACEUTICALS INC USA
NILSTAT, NYSTATIN

GLENWOOD

* GLENWOOD INC
PASKALIUM, POTASSIUM AMINOSALICYLATE

GOLDLINE

* GOLDLINE LABORATORIES INC
CLOZAPINE, CLOZAPINE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE

GRAHAM CHEM

* GRAHAM CHEMICAL CO
LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE, LIDOCAINE
MEPIVACAINE HYDROCHLORIDE W/ LEVONORDEFIN, LEVONORDEFIN
MEPIVACAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE

GRIFFEN

* KW GRIFFEN CO
BIOSCRUB, CHLORHEXIDINE GLUCONATE (OTC)

GTX INC

* GTX INC
FARESTON, TOREMIFENE CITRATE

GUARDIAN DRUG

* GUARDIAN DRUG CO INC
FOAMCOAT, ALUMINUM HYDROXIDE (OTC)

GUERBET

* GUERBET LLC
OXILAN-300, IOXILAN
OXILAN-350, IOXILAN

HAEMONETICS

* HAEMONETICS CORP
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

HALOCARBON

* HALOCARBON LABORATORIES DIV HALOCARBON PRODUCTS CORP
HALOTHANE, HALOTHANE

HALOCARBON PRODS

* HALOCARBON PRODUCTS CORP
ISOFLURANE, ISOFLURANE

HALOZYME THERAP

* HALOZYME THERAPEUTICS INC
HYLENEX RECOMBINANT, HYALURONIDASE RECOMBINANT HUMAN

HAMELN PHARMS

* HAMELN PHARMACEUTICALS GMBH
PENTETATE CALCIUM TRISODIUM, PENTETATE CALCIUM TRISODIUM
PENTETATE ZINC TRISODIUM, PENTETATE ZINC TRISODIUM

HAMPTON LAINE LLC

* HAMPTON LAINE LLC
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE

HANFORD GC

* GC HANFORD MANUFACTURING CO
AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM
CEFAZOLIN SODIUM, CEFZOLIN SODIUM
CEFUROXIME SODIUM, CEFUROXIME SODIUM

HARMONY LABS

* HARMONY LABORATORIES
THEROXIDIL, MINOXIDIL (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** H **

HARVEST PHARMS

- * HARVEST PHARMACEUTICALS INC
ERGOMAR, ERGOTAMINE TARTRATE

HEALTHPOINT

- * HEALTHPOINT LTD
AKNE-MYCIN, ERYTHROMYCIN
CETACORT, HYDROCORTISONE
CLODERM, CLOCORTOLONE PIVALATE
CORMAX, CLOBETASOL PROPIONATE
ECONAZOLE NITRATE, ECONAZOLE NITRATE
EMBELINE E, CLOBETASOL PROPIONATE
EMBELINE, CLOBETASOL PROPIONATE
NUTRACORT, HYDROCORTISONE

HELSINN HLTHCARE

- * HELSINN HEALTHCARE SA
ALOXI, PALONOSETRON HYDROCHLORIDE

HERCON LABS

- * HERCON LABORATORIES CORP
NITROGLYCERIN, NITROGLYCERIN

HEYL CHEMISCH

- * HEYL CHEMISCH PHARMAZEUTISCHE FABRIK
RADIOGARDASE (PRUSSIAN BLUE), FERRIC HEXACYANOFERRATE(II)

HI TECH PHARMA

- * HI TECH PHARMACAL CO INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACYCLOVIR, ACYCLOVIR
ALBUTEROL SULFATE, ALBUTEROL SULFATE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
LACTULOSE, LACTULOSE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
OFLOXACIN, OFLOXACIN
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISOLONE, PREDNISOLONE
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
PROMETHAZINE HYDROCHLORIDE AND DEXTROMETHORPHAN HYDROBROMIDE, DEXTROMETHORPHAN
HYDROBROMIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
TIMOLOL MALEATE, TIMOLOL MALEATE

HIGH TECH PHARMA

- * HIGH TECHNOLOGY PHARMACAL CO INC
VALPROIC ACID, VALPROIC ACID

HIKMA

- * HIKMA FARMACEUTICA LDA
CEFOTAXIME, CEFOTAXIME SODIUM
- * HIKMA PHARMACEUTICALS
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
GLYBURIDE (MICRONIZED), GLYBURIDE
NAPROXEN SODIUM, NAPROXEN SODIUM

HIKMA FARMACEUTICA

- * HIKMA FARMACEUTICA PORTUGAL LDA
CEFAZOLIN SODIUM, CEFAZOLIN SODIUM
CEFUROXIME, CEFUROXIME SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** H ****

* HIKMA FARMACEUTICA PORTUGAL LDA
FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE

HIKMA PHARMS

* HIKMA PHARMACEUTICALS
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN

HILL DERMAC

* HILL DERMACEUTICALS INC
DERMA-SMOOTH/FS, FLUOCINOLONE ACETONIDE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE

HITECH PHARMA

* HITECH PHARMACAL CORP
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE

HLR

* HLR TECHNOLOGY
ACCUTANE, ISOTRETINOIN
FORTOVASE, SAQUINAVIR
INVIRASE, SAQUINAVIR MESYLATE
ROCEPHIN W/ DEXTROSE IN PLASTIC CONTAINER, CEFTRIAXONE SODIUM
ROCEPHIN, CEFTRIAXONE SODIUM
ROMAZICON, FLUMAZENIL
XELODA, CAPECITABINE
XENICAL, ORLISTAT

HOLOPACK INTL

* HOLOPACK INTERNATIONAL CORP
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

HOSPIRA

* HOSPIRA INC
ACETAZOLAMIDE SODIUM, ACETAZOLAMIDE SODIUM
ACETIC ACID 0.25% IN PLASTIC CONTAINER, ACETIC ACID, GLACIAL
ACETYLCYSTEINE, ACETYLCYSTEINE
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
A-HYDROCORT, HYDROCORTISONE SODIUM SUCCINATE
ALCOHOL 5% IN D5-W, ALCOHOL
ALFENTANIL, ALFENTANIL HYDROCHLORIDE
A-METHAPRED, METHYLPREDNISOLONE SODIUM SUCCINATE
AMIDATE, ETOMIDATE
AMIKACIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, AMIKACIN SULFATE
AMIKACIN SULFATE, AMIKACIN SULFATE
AMINOCAPROIC ACID IN PLASTIC CONTAINER, AMINOCAPROIC ACID
AMINOCAPROIC ACID, AMINOCAPROIC ACID
AMINOPHYLLINE IN SODIUM CHLORIDE 0.45%, AMINOPHYLLINE
AMINOPHYLLINE, AMINOPHYLLINE
AMINOSYN 10% (PH6), AMINO ACIDS
AMINOSYN 10%, AMINO ACIDS
AMINOSYN 3.5% M, AMINO ACIDS
AMINOSYN 3.5%, AMINO ACIDS
AMINOSYN 5%, AMINO ACIDS
AMINOSYN 7% (PH6), AMINO ACIDS
AMINOSYN 7% W/ ELECTROLYTES, AMINO ACIDS
AMINOSYN 7%, AMINO ACIDS
AMINOSYN 8.5% (PH6), AMINO ACIDS
AMINOSYN 8.5% W/ ELECTROLYTES, AMINO ACIDS
AMINOSYN 8.5%, AMINO ACIDS
AMINOSYN II 10% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 10% W/ ELECTROLYTES, AMINO ACIDS
AMINOSYN II 10%, AMINO ACIDS
AMINOSYN II 15% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 3.5% IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 3.5% IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 3.5% M IN DEXTROSE 5% IN PLASTIC CONTAINER, AMINO ACIDS
AMINOSYN II 3.5% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** H **

* HOSPIRA INC

AMINOSYN II 4.25% IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS

AMINOSYN II 4.25% IN DEXTROSE 20% IN PLASTIC CONTAINER, AMINO ACIDS

AMINOSYN II 4.25% IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS

AMINOSYN II 4.25% M IN DEXTROSE 10% IN PLASTIC CONTAINER, AMINO ACIDS

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 20% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS

AMINOSYN II 4.25% W/ ELECTROLYTES IN DEXTROSE 25% W/ CALCIUM IN PLASTIC CONTAINER, AMINO ACIDS

AMINOSYN II 5% IN DEXTROSE 25% IN PLASTIC CONTAINER, AMINO ACIDS

AMINOSYN II 7%, AMINO ACIDS

AMINOSYN II 8.5% W/ ELECTROLYTES, AMINO ACIDS

AMINOSYN II 8.5%, AMINO ACIDS

AMINOSYN-HBC 7%, AMINO ACIDS

AMINOSYN-HF 8%, AMINO ACIDS

AMINOSYN-PF 10%, AMINO ACIDS

AMINOSYN-PF 7%, AMINO ACIDS

AMINOSYN-RF 5.2%, AMINO ACIDS

AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE

AMMONIUM CHLORIDE IN PLASTIC CONTAINER, AMMONIUM CHLORIDE

AMRINONE, INAMRINONE LACTATE

ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE

ATRACURIUM BESYLATE, ATRACURIUM BESYLATE

ATROPINE SULFATE ANSYR PLASTIC SYRINGE, ATROPINE SULFATE

BACTERIOSTATIC SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE

BACTERIOSTATIC WATER FOR INJECTION IN PLASTIC CONTAINER, WATER FOR INJECTION, STERILE

BRETYLIUM TOSYLATE IN DEXTROSE 5% IN PLASTIC CONTAINER, BRETYLIUM TOSYLATE

BRETYLIUM TOSYLATE IN PLASTIC CONTAINER, BRETYLIUM TOSYLATE

BUMETANIDE, BUMETANIDE

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE

BUPIVACAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE

BUPIVACAINE, BUPIVACAINE HYDROCHLORIDE

BUPRENORPHINE HYDROCHLORIDE, BUPRENORPHINE HYDROCHLORIDE

BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE

BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE

CALCIUM CHLORIDE 10% IN PLASTIC CONTAINER, CALCIUM CHLORIDE

CARBOCAINE, MEPIVACAINE HYDROCHLORIDE

CHLOROPROCAINE HYDROCHLORIDE, CHLOROPROCAINE HYDROCHLORIDE

CHROMIC CHLORIDE IN PLASTIC CONTAINER, CHROMIC CHLORIDE

CIMETIDINE HYDROCHLORIDE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, CIMETIDINE HYDROCHLORIDE

CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE

CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE

CORLOPAM, FENOLDOPAM MESYLATE

CUPRIC CHLORIDE IN PLASTIC CONTAINER, CUPRIC CHLORIDE

DEFEROXAMINE MESYLATE, DEFEROXAMINE MESYLATE

DEMEROL, MEPERIDINE HYDROCHLORIDE

DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE

DEXTROSE 10% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 20% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 25%, DEXTROSE

DEXTROSE 30% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 40% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE

DEXTROSE 5% AND RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE

DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 50% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 60% IN PLASTIC CONTAINER, DEXTROSE

DEXTROSE 70% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** H **

* HOSPIRA INC
DIAZEPAM, DIAZEPAM
DIGOXIN PEDIATRIC, DIGOXIN
DIGOXIN, DIGOXIN
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE IN DEXTROSE 5%, DOBUTAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, DOPAMINE HYDROCHLORIDE
DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
DROPERIDOL, DROPERIDOL
EDROPHONIUM CHLORIDE, EDROPHONIUM CHLORIDE
ENALAPRILAT, ENALAPRILAT
ENDRATE, EDETATE DISODIUM
ERYTHROCIN, ERYTHROMYCIN LACTOBIONATE
ETOPOSIDE, ETOPOSIDE
FAMOTIDINE, FAMOTIDINE
FENTANYL CITRATE PRESERVATIVE FREE, FENTANYL CITRATE
FENTANYL CITRATE, FENTANYL CITRATE
FLUCONAZOLE IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FUROSEMIDE, FUROSEMIDE
GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, GENTAMICIN SULFATE
GENTAMICIN SULFATE, GENTAMICIN SULFATE
GLYCINE 1.5% IN PLASTIC CONTAINER, GLYCINE
HALOTHANE, HALOTHANE
HEPARIN LOCK FLUSH IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN LOCK FLUSH, HEPARIN SODIUM
HEPARIN SODIUM 1,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 10,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 12,500 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 12,500 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 2,000 UNITS IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 20,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN DEXTROSE 5% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM 25,000 UNITS IN SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, HEPARIN SODIUM
HEPARIN SODIUM PRESERVATIVE FREE, HEPARIN SODIUM
HEPARIN SODIUM, HEPARIN SODIUM
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IONOSOL B AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IONOSOL MB AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IONOSOL T AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
IOPAMIDOL-200 IN PLASTIC CONTAINER, IOPAMIDOL
IOPAMIDOL-200, IOPAMIDOL
IOPAMIDOL-250 IN PLASTIC CONTAINER, IOPAMIDOL
IOPAMIDOL-250, IOPAMIDOL
IOPAMIDOL-300 IN PLASTIC CONTAINER, IOPAMIDOL
IOPAMIDOL-300, IOPAMIDOL
IOPAMIDOL-370 IN PLASTIC CONTAINER, IOPAMIDOL
IOPAMIDOL-370, IOPAMIDOL
ISOFLURANE, ISOFLURANE
ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
ISUPREL, ISOPROTERENOL HYDROCHLORIDE
KETAMINE HYDROCHLORIDE, KETAMINE HYDROCHLORIDE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
LEVOPHED, NOREPINEPHRINE BITARTRATE
LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.2% IN DEXTROSE 5%, LIDOCAINE HYDROCHLORIDE
LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** H **

* HOSPIRA INC

LIDOCAINE HYDROCHLORIDE 0.4% IN DEXTROSE 5%, LIDOCAINE HYDROCHLORIDE

LIDOCAINE HYDROCHLORIDE 0.8% IN DEXTROSE 5% IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE

LIDOCAINE HYDROCHLORIDE 5% AND DEXTROSE 7.5%, LIDOCAINE HYDROCHLORIDE

LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE, EPINEPHRINE

LIDOCAINE HYDROCHLORIDE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE IN PLASTIC CONTAINER, LIDOCAINE HYDROCHLORIDE

LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE

LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE

LIPOSYN II 10%, SAFFLOWER OIL

LIPOSYN II 20%, SAFFLOWER OIL

LIPOSYN III 10%, SOYBEAN OIL

LIPOSYN III 20%, SOYBEAN OIL

LIPOSYN III 30%, SOYBEAN OIL

LORAZEPAM, LORAZEPAM

LTA II KIT, LIDOCAINE HYDROCHLORIDE

MAGNESIUM SULFATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MAGNESIUM SULFATE

MAGNESIUM SULFATE IN PLASTIC CONTAINER, MAGNESIUM SULFATE

MAGNESIUM SULFATE, MAGNESIUM SULFATE

MANGANESE CHLORIDE IN PLASTIC CONTAINER, MANGANESE CHLORIDE

MANNITOL 10% IN PLASTIC CONTAINER, MANNITOL

MANNITOL 15% IN PLASTIC CONTAINER, MANNITOL

MANNITOL 20% IN PLASTIC CONTAINER, MANNITOL

MANNITOL 25%, MANNITOL

MANNITOL 5% IN PLASTIC CONTAINER, MANNITOL

MARCAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE

MARCAINE HYDROCHLORIDE W/ EPINEPHRINE, BUPIVACAINE HYDROCHLORIDE

MARCAINE HYDROCHLORIDE, BUPIVACAINE HYDROCHLORIDE

MARCAINE, BUPIVACAINE HYDROCHLORIDE

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE

METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE

METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE

METOPROLOL TARTRATE, METOPROLOL TARTRATE

METRONIDAZOLE IN PLASTIC CONTAINER, METRONIDAZOLE

MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE

MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE

MILRINONE LACTATE, MILRINONE LACTATE

MORPHINE SULFATE, MORPHINE SULFATE

NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE

NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE

NITROGLYCERIN IN DEXTROSE 5%, NITROGLYCERIN

NITROGLYCERIN, NITROGLYCERIN

NITROPRESS, SODIUM NITROPRUSSIDE

NORMOSOL-M AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE

NORMOSOL-R AND DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE

NORMOSOL-R IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE

NOVAMINE 11.4%, AMINO ACIDS

NOVAMINE 15%, AMINO ACIDS

NOVOCAIN, PROCAINE HYDROCHLORIDE

PANCURONIUM BROMIDE, PANCURONIUM BROMIDE

PEDIATRIC LTA KIT, LIDOCAINE HYDROCHLORIDE

PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE

PHENYTOIN SODIUM, PHENYTOIN SODIUM

PHYSIOSOL IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE

PHYSIOSOL PH 7.4 IN PLASTIC CONTAINER, MAGNESIUM CHLORIDE

PLEGISOL IN PLASTIC CONTAINER, CALCIUM CHLORIDE

POTASSIUM ACETATE IN PLASTIC CONTAINER, POTASSIUM ACETATE

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE

POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** H **

* HOSPIRA INC
 POTASSIUM CHLORIDE 10MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 15MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN DEXTROSE 5% IN SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 20MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 30MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND LACTATED RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN DEXTROSE 5% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 40MEQ IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 40MEQ IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, POTASSIUM CHLORIDE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.225% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.3% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE 5MEQ IN DEXTROSE 5% AND SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, DEXTROSE
 POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
 PRECEDEX, DEXMEDETOMIDINE
 PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 PROCAINE HYDROCHLORIDE, PROCAINE HYDROCHLORIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 QUELICIN PRESERVATIVE FREE, SUCCINYLCHOLINE CHLORIDE
 QUELICIN, SUCCINYLCHOLINE CHLORIDE
 RINGER'S IN PLASTIC CONTAINER, CALCIUM CHLORIDE
 RITODRINE HYDROCHLORIDE IN DEXTROSE 5% IN PLASTIC CONTAINER, RITODRINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** H ****

* HOSPIRA INC
RITODRINE HYDROCHLORIDE, RITODRINE HYDROCHLORIDE
SODIUM ACETATE IN PLASTIC CONTAINER, SODIUM ACETATE, ANHYDROUS
SODIUM BICARBONATE, SODIUM BICARBONATE
SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM CHLORIDE IN PLASTIC CONTAINER, SODIUM CHLORIDE
SODIUM LACTATE 0.167 MOLAR IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM LACTATE IN PLASTIC CONTAINER, SODIUM LACTATE
SODIUM PHOSPHATES IN PLASTIC CONTAINER, SODIUM PHOSPHATE, DIBASIC, HEPTAHYDRATE
SORBITOL-MANNITOL IN PLASTIC CONTAINER, MANNITOL
STERILE WATER FOR INJECTION IN PLASTIC CONTAINER, WATER FOR INJECTION, STERILE
STERILE WATER IN PLASTIC CONTAINER, WATER FOR IRRIGATION, STERILE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SUFENTANIL CITRATE, SUFENTANIL CITRATE
TALWIN, PENTAZOCINE LACTATE
TAZICEF, CEFTAZIDIME
THAM, TROMETHAMINE
THEOPHYLLINE IN DEXTROSE 5% IN PLASTIC CONTAINER, THEOPHYLLINE
THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
TOBRAMYCIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, TOBRAMYCIN SULFATE
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE
TPN ELECTROLYTES IN PLASTIC CONTAINER, CALCIUM CHLORIDE
TRACRIUM PRESERVATIVE FREE, ATRACURIUM BESYLATE
TRACRIUM, ATRACURIUM BESYLATE
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE
UREAPHIL, UREA
UROLOGIC G IN PLASTIC CONTAINER, CITRIC ACID
VANCOMYCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
VITAMIN K1, PHYTONADIONE
ZINC CHLORIDE IN PLASTIC CONTAINER, ZINC CHLORIDE

IBI

* ISTITUTO BIOCHIMICO ITALIANO SPA
AMPICILLIN SODIUM, AMPICILLIN SODIUM

IMPAX LABS

* IMPAX LABORATORIES INC
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
DANTROLENE SODIUM, DANTROLENE SODIUM
DEMECLOCYCLINE HYDROCHLORIDE, DEMECLOCYCLINE HYDROCHLORIDE
FENOFIBRATE (MICRONIZED), FENOFIBRATE
FLUDROCORTISONE ACETATE, FLUDROCORTISONE ACETATE
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE, LORATADINE (OTC)
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
METHYLTESTOSTERONE, METHYLTESTOSTERONE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
OMEPRazole, OMEPRazole
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PENTOXIFYLLINE, PENTOXIFYLLINE
PROPYLTHIOURACIL, PROPYLTHIOURACIL
PYRIDOSTIGMINE BROMIDE, PYRIDOSTIGMINE BROMIDE
RILUZOLE, RILUZOLE
RIMANTADINE HYDROCHLORIDE, RIMANTADINE HYDROCHLORIDE
TERBUTALINE SULFATE, TERBUTALINE SULFATE
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** I **

IMPAX PHARMS

- * IMPAX PHARMACEUTICALS
 FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
 MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
 ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
 OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
 SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE

INALCO

- * INALCO SPA
 LACTULOSE, LACTULOSE

INDEVUS

- * INDEVUS PHARMACEUTICALS INC
 SANCTURA, TROSPIUM CHLORIDE

INGRAM PHARM

- * INGRAM PHARMACEUTICAL CO
 DRICORT, HYDROCORTISONE ACETATE
 HYDROCORTISONE, HYDROCORTISONE

INO

- * INO THERAPEUTICS INC
 INOMAX, NITRIC OXIDE

INSIGHT PHARMS

- * INSIGHT PHARMACEUTICALS CORP
 NIX, PERMETHRIN (OTC)

INSMED

- * INSMED INC
 IPLEX, MECASERMIN RINFABATE RECOMBINANT

INSTITUTO BIOCHEMICO

- * INSTITUTO BIOCHEMICO ITALIANO SPA
 AMPICILLIN SODIUM, AMPICILLIN SODIUM

INSTITUTO BIOCHIMICO

- * INSTITUTO BIOCHIMICO ITALIANO SPA
 AMPICILLIN AND SULBACTAM, AMPICILLIN SODIUM

INTENDIS

- * INTENDIS INC
 FINACEA, AZELAIC ACID

INTERPHARM

- * INTERPHARM INC
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 NAPROXEN, NAPROXEN
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

INTL MEDICATED

- * INTERNATIONAL MEDICATED SYSTEMS LTD
 BUPIVACAINE HYDROCHLORIDE PRESERVATIVE FREE, BUPIVACAINE HYDROCHLORIDE
 MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
 MILRINONE LACTATE, MILRINONE LACTATE

INTL MEDICATION

- * INTERNATIONAL MEDICATION SYSTEM
 AMINOPHYLLINE, AMINOPHYLLINE
 BRETILIUM TOSYLATE, BRETILIUM TOSYLATE
 DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 FUROSEMIDE, FUROSEMIDE
 ISOPROTERENOL HYDROCHLORIDE, ISOPROTERENOL HYDROCHLORIDE
 LARYNG-O-JET KIT, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE PRESERVATIVE FREE, LIDOCAINE HYDROCHLORIDE
 LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
 MANNITOL 25%, MANNITOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** I **

- * INTERNATIONAL MEDICATION SYSTEM
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
NALOXONE HYDROCHLORIDE, NALOXONE HYDROCHLORIDE
PHYTONADIONE, PHYTONADIONE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
- * INTERNATIONAL MEDICATION SYSTEMS LTD
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
FUROSEMIDE, FUROSEMIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE

INTL MEDICATION SYS

- * INTERNATIONAL MEDICATION SYSTEMS LTD
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
LORAZEPAM, LORAZEPAM

INVAGEN PHARMS

- * INVAGEN PHARMACEUTICALS INC
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
GLIMEPIRIDE, GLIMEPIRIDE

INWOOD LABS

- * INWOOD LABORATORIES INC SUB FOREST LABORATORIES INC
CARBAMAZEPINE, CARBAMAZEPINE
INDOMETHACIN, INDOMETHACIN
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
THEOCHRON, THEOPHYLLINE
THEOPHYLLINE, THEOPHYLLINE

IPR

- * IPR PHARMACEUTICALS INC
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
ZOMIG, ZOLMITRIPTAN

ISO TEX

- * ISO TEX DIAGNOSTICS INC
JEANATOPE, ALBUMIN IODINATED I-125 SERUM
MEGATOPE, ALBUMIN IODINATED I-131 SERUM

ISTA PHARMS

- * ISTA PHARMACEUTICALS
ISTALOL, TIMOLOL MALEATE
VITRASE, HYALURONIDASE
XIBROM, BROMFENAC SODIUM

IVAX PHARMS

- *
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
- * IVAX PHARMACEUTICALS INC
ACYCLOVIR, ACYCLOVIR
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALBUTEROL, ALBUTEROL
ALLAY, ACETAMINOPHEN
ALPRAZOLAM, ALPRAZOLAM
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
BACLOFEN, BACLOFEN
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BRIMONIDINE TARTRATE, BRIMONIDINE TARTRATE
BUMETANIDE, BUMETANIDE
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
CAPTOPRIL, CAPTOPRIL
CEFACLOX, CEFACLOX
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEPHALEXIN, CEPHALEXIN
CILOSTAZOL, CILOSTAZOL
CIMETIDINE, CIMETIDINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** I **

* IVAX PHARMACEUTICALS INC
 CIMETIDINE, CIMETIDINE (OTC)
 CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLARITHROMYCIN, CLARITHROMYCIN
 CLOZAPINE, CLOZAPINE
 CROMOLYN SODIUM, CROMOLYN SODIUM
 C-SOLVE-2, ERYTHROMYCIN
 CYCLOSPORINE, CYCLOSPORINE
 CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
 DIAZEPAM, DIAZEPAM
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE, ATROPINE SULFATE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 ESTAZOLAM, ESTAZOLAM
 ETODOLAC, ETODOLAC
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUOXETINE, FLUOXETINE HYDROCHLORIDE
 FLUTAMIDE, FLUTAMIDE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FUROSEMIDE, FUROSEMIDE
 GABAPENTIN, GABAPENTIN
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GUANABENZ ACETATE, GUANABENZ ACETATE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE, HOMATROPINE METHYLBROMIDE
 HYDROGENATED ERGOT ALKALOIDS, ERGOLOID MESYLATES
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 IBUPROFEN, IBUPROFEN (OTC)
 INDAPAMIDE, INDAPAMIDE
 INDOMETHACIN, INDOMETHACIN
 IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METHOCARBAMOL AND ASPIRIN, ASPIRIN
 METHYLCLOTHIAZIDE, METHYLCLOTHIAZIDE
 METHYLDOPA, METHYLDOPA
 METRONIDAZOLE, METRONIDAZOLE
 MIRTAZAPINE, MIRTAZAPINE
 MISOPROSTOL, MISOPROSTOL
 NABUMETONE, NABUMETONE
 NADOLOL, NADOLOL
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 OXAPROZIN, OXAPROZIN
 OXAZEPAM, OXAZEPAM
 PACLITAXEL, PACLITAXEL
 PERGOLIDE MESYLATE, PERGOLIDE MESYLATE
 PERPHENAZINE, PERPHENAZINE
 PINDOLOL, PINDOLOL
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** I **

- * IVAX PHARMACEUTICALS INC
 - PROBENECID AND COLCHICINE, COLCHICINE
 - PROBENECID, PROBENECID
 - PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
 - PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
 - PROMPT PHENYTOIN SODIUM, PHENYTOIN SODIUM
 - PROPOXYPHENE HYDROCHLORIDE, PROPOXYPHENE HYDROCHLORIDE
 - PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
 - RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
 - RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
 - SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
 - SULFINPYRAZONE, SULFINPYRAZONE
 - SULFISOXAZOLE, SULFISOXAZOLE
 - TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
 - TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
 - THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
 - TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
 - TOLAZAMIDE, TOLAZAMIDE
 - TOLBUTAMIDE, TOLBUTAMIDE
 - TOLMETIN SODIUM, TOLMETIN SODIUM
 - TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
 - TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 - VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

IVAX RES

- * IVAX RESEARCH INC
 - NASAREL, FLUNISOLIDE
 - PROAIR HFA, ALBUTEROL SULFATE

J AND J

- * JOHNSON AND JOHNSON CONSUMER PRODUCTS INC
 - ERYCETTE, ERYTHROMYCIN
 - GRIFULVIN V, GRISEOFULVIN, MICROCRYSTALLINE
- * JOHNSON AND JOHNSON MEDICAL INC
 - MICRODERM, CHLORHEXIDINE GLUCONATE (OTC)
 - PREVACARE R, CHLORHEXIDINE GLUCONATE (OTC)

JACOBUS

- * JACOBUS PHARMACEUTICAL CO
 - DAPSONE, DAPSONE
 - PASER, AMINOSALICYLIC ACID

JANSSEN

- * JANSSEN PHARMACEUTICA INC
 - RAZADYNE ER, GALANTAMINE HYDROBROMIDE

JANSSEN PHARMA

- * JANSSEN PHARMACEUTICA PRODUCTS LP
 - NIZORAL, KETOCONAZOLE
 - RAZADYNE, GALANTAMINE HYDROBROMIDE
 - RISPERDAL CONSTA, RISPERIDONE
 - RISPERDAL, RISPERIDONE
 - SPORANOX, ITRACONAZOLE

JDS PHARMS

- * JDS PHARMACEUTICALS LLC
 - LITHOBID, LITHIUM CARBONATE

JOHN O BUTLER CO

- * JOHN O BUTLER CO
 - CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE

JOHNSON AND JOHNSON

- * JOHNSON AND JOHNSON CONSUMER COMPANIES INC
 - CENTANY, MUPIROCIN
 - MONISTAT-DERM, MICONAZOLE NITRATE
 - RENOVA, TRETINOIN
 - RETIN-A MICRO, TRETINOIN
 - RETIN-A, TRETINOIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** J **

- * JOHNSON AND JOHNSON CONSUMER COMPANIES INC
SPECTAZOLE, ECONAZOLE NITRATE
- * JOHNSON AND JOHNSON CONSUMER PRODUCTS WORLDWIDE
ERTACZO, SERTACONAZOLE NITRATE

JOHNSON RW

- * RW JOHNSON PHARMACEUTICAL RESEARCH INSTITUTE
ORTHO CYCLEN-28, ETHINYL ESTRADIOL
ORTHO TRI-CYCLEN, ETHINYL ESTRADIOL

JONES PHARMA

- * JONES PHARMA INC SUB KING PHARMACEUTICALS INC
LEVOXYL, LEVOTHYROXINE SODIUM
METHIMAZOLE, METHIMAZOLE
SONATA, ZALEPLON
TRIOSTAT, LIOTHYRONINE SODIUM
TUSSIGON, HOMATROPINE METHYLBROMIDE

JONES PHARMA INC

- * JONES PHARMA INC
SKELAXIN, METAXALONE

JVL

- * JVL CORP
METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE

KALI LABS

- * KALI LABORATORIES INC
CAPTOPRIL, CAPTOPRIL
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLONAZEPAM, CLONAZEPAM
GLIPIZIDE, GLIPIZIDE
LEFLUNOMIDE, LEFLUNOMIDE
METRONIDAZOLE, METRONIDAZOLE
NYSTATIN, NYSTATIN
TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

KENDALL IL

- * KENDALL CO
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

KENDALL LP

- * KENDALL CO LP
THERMAZENE, SILVER SULFADIAZINE

KENWOOD LABS

- * KENWOOD LABORATORIES DIV BRADLEY PHARMACEUTICALS
CARMOL HC, HYDROCORTISONE ACETATE
TYZINE, TETRAHYDROZOLINE HYDROCHLORIDE

KEY PHARMS

- * KEY PHARMACEUTICALS INC SUB SCHERING PLOUGH CORP
K-DUR 10, POTASSIUM CHLORIDE
K-DUR 20, POTASSIUM CHLORIDE
NITRO-DUR, NITROGLYCERIN

KIEL

- * KIEL LABORATORIES INC
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE

KING PHARMS

- * KING PHARMACEUTICALS INC
ALTACE, RAMIPRIL
BICILLIN C-R 900/300, PENICILLIN G BENZATHINE
BICILLIN C-R, PENICILLIN G BENZATHINE
BICILLIN L-A, PENICILLIN G BENZATHINE
BREVITAL SODIUM, METHOHEXITAL SODIUM
CORZIDE, BENDROFLUMETHIAZIDE
CYTOMEL, LIOTHYRONINE SODIUM
DELESTROGEN, ESTRADIOL VALERATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** K **

* KING PHARMACEUTICALS INC
FLORINEF, FLUDROCORTISONE ACETATE
HUMATIN, PAROMOMYCIN SULFATE
INTAL, CROMOLYN SODIUM
LORABID, LORACARBEF
PENICILLIN G PROCAINE, PENICILLIN G PROCAINE
PROCANBID, PROCAINAMIDE HYDROCHLORIDE
SILVADENE, SILVER SULFADIAZINE
SYNERCID, DALFOPRISTIN
TIGAN, TRIMETHOBENZAMIDE HYDROCHLORIDE
TILADE, NEDOCROMIL SODIUM

KOS LIFE

* KOS LIFE SCIENCES INC
ADVICOR, LOVASTATIN
AZMACORT, TRIAMCINOLONE ACETONIDE
NIASPAN, NIACIN
TEVETEN HCT, EPROSARTAN MESYLATE
TEVETEN, EPROSARTAN MESYLATE

KREMERS URBAN

* KREMERS URBAN CO
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
NITROGLYCERIN, NITROGLYCERIN

KREMERS URBAN DEV

* KREMERS URBAN DEVELOPMENT CO
OMEPRazole, OMEPRazole

KRKA DD NOVO MESTO

* KRKA DD NOVO MESTO
ENALAPRIL MALEATE, ENALAPRIL MALEATE

KV PHARM

* KV PHARMACEUTICAL CO
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BROMPHENIRAMINE MALEATE, BROMPHENIRAMINE MALEATE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CLINDESSE, CLINDAMYCIN PHOSPHATE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
GYNAZOLE-1, BUTOCONAZOLE NITRATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
MICRO-K 10, POTASSIUM CHLORIDE
MICRO-K, POTASSIUM CHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISOLONE, PREDNISOLONE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE

LABS ATRAL

* LABORATORIOS ATRAL SARL
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE

LAFAYETTE PHARMS

* LAFAYETTE PHARMACEUTICALS INC
BAROS, SODIUM BICARBONATE

LANNETT

* LANNETT CO INC
ACETAZOLAMIDE, ACETAZOLAMIDE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** L ****

- * LANNETT CO INC
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
LANIAZID, ISONIAZID
LANORINAL, ASPIRIN
METHOCARBAMOL, METHOCARBAMOL
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PREDNISOLONE, PREDNISOLONE
PRIMIDONE, PRIMIDONE
PROBALAN, PROBENECID
QUINIDINE SULFATE, QUINIDINE SULFATE
SERPALAN, RESERPINE
SODIUM P.A.S., AMINOSALICYLATE SODIUM
- * LANNETT HOLDINGS INC
BACLOFEN, BACLOFEN
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
DANAZOL, DANAZOL
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
DOXYCYCLINE, DOXYCYCLINE
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
TERBUTALINE SULFATE, TERBUTALINE SULFATE

LEINER

- * LEINER HEALTH PRODUCTS INC
CIMETIDINE, CIMETIDINE (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)

LEITNER PHARMS

- * LEITNER PHARMACEUTICALS LLC
EQUAGESIC, ASPIRIN
SYNALGOS-DC, ASPIRIN
WYGESIC, ACETAMINOPHEN

LEK PHARMS

- * LEK PHARMACEUTICALS D D
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
CIMETIDINE, CIMETIDINE
CIMETIDINE, CIMETIDINE (OTC)
ENALAPRIL MALEATE, ENALAPRIL MALEATE
LISINOPRIL, LISINOPRIL
OMEPRazole, OMEPRazole

LIGAND

- * LIGAND PHARMACEUTICALS INC
AVINZA, MORPHINE SULFATE
PANRETIN, ALITRETINOIN
TARGRETIN, BEXAROTENE

LILLY

- * ELI LILLY AND CO
ALIMTA, PEMETREXED DISODIUM
CAPASTAT SULFATE, CAPREOMYCIN SULFATE
CYMBALTA, DULOXETINE HYDROCHLORIDE
EVISTA, RALOXIFENE HYDROCHLORIDE
FORTEO, TERIPARATIDE RECOMBINANT HUMAN
GEMZAR, GEMCITABINE HYDROCHLORIDE
GLUCAGON, GLUCAGON RECOMBINANT
HUMALOG MIX 50/50, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG MIX 75/25, INSULIN LISPRO PROTAMINE RECOMBINANT
HUMALOG PEN, INSULIN LISPRO RECOMBINANT
HUMALOG, INSULIN LISPRO RECOMBINANT
HUMATROPE, SOMATROPIN RECOMBINANT
HUMULIN 50/50, INSULIN RECOMBINANT HUMAN (OTC)
HUMULIN 70/30 PEN, INSULIN RECOMBINANT HUMAN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** L **

- * ELI LILLY AND CO
 - HUMULIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
 - HUMULIN L, INSULIN ZINC SUSP RECOMBINANT HUMAN (OTC)
 - HUMULIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
 - HUMULIN R PEN, INSULIN RECOMBINANT HUMAN (OTC)
 - HUMULIN R, INSULIN RECOMBINANT HUMAN
 - HUMULIN R, INSULIN RECOMBINANT HUMAN (OTC)
 - HUMULIN U, INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN (OTC)
 - Iletin II, INSULIN PURIFIED PORK
 - LENTE Iletin II (PORK), INSULIN ZINC SUSP PURIFIED PORK (OTC)
 - NPH Iletin I (BEEF-PORK), INSULIN SUSP ISOPHANE BEEF/PORK (OTC)
 - NPH Iletin II (PORK), INSULIN SUSP ISOPHANE PURIFIED PORK (OTC)
 - PROZAC WEEKLY, FLUOXETINE HYDROCHLORIDE
 - PROZAC, FLUOXETINE HYDROCHLORIDE
 - QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
 - REGULAR Iletin II (PORK), INSULIN PURIFIED PORK (OTC)
 - SEROMYCIN, CYCLOSERINE
 - STRATTERA, ATOMOXETINE HYDROCHLORIDE
 - SYMBYAX, FLUOXETINE HYDROCHLORIDE
 - ZYPREXA ZYDIS, OLANZAPINE
 - ZYPREXA, OLANZAPINE
- * LILLY RESEARCH LABORATORIES DIV ELI LILLY AND CO
 - PROZAC, FLUOXETINE HYDROCHLORIDE
 - SARAFEM, FLUOXETINE HYDROCHLORIDE

LILLY ICOS

- * LILLY ICOS LLC
 - CIALIS, TADALAFIL

LLOYD

- * LLOYD INC
 - LEVOTHROID, LEVOTHYROXINE SODIUM

LNK

- * LNK INTERNATIONAL INC
 - DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 - DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
 - IBUPROFEN, IBUPROFEN (OTC)
 - LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 - METRONIDAZOLE, METRONIDAZOLE

LOCH

- * LOCH PHARMACEUTICALS INC
 - FOLIC ACID, FOLIC ACID

LOREX

- * LOREX PHARMACEUTICALS
 - KERLONE, BETAXOLOL HYDROCHLORIDE

LUITPOLD

- * LUITPOLD PHARMACEUTICALS INC
 - ACETYLCYSTEINE, ACETYLCYSTEINE
 - AMINOCAPROIC ACID, AMINOCAPROIC ACID
 - AMINOPHYLLINE, AMINOPHYLLINE
 - BRETYLIUM TOSYLATE, BRETYLIUM TOSYLATE
 - CALCITRIOL, CALCITRIOL
 - CYANOCOBALAMIN, CYANOCOBALAMIN
 - DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 - DEXFERRUM, IRON DEXTRAN
 - DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 - DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 - DROPERIDOL, DROPERIDOL
 - FUROSEMIDE, FUROSEMIDE
 - GLYCOPYRROLATE, GLYCOPYRROLATE
 - HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 - HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 - LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 - LEVOCARNITINE, LEVOCARNITINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** L ****

* LUITPOLD PHARMACEUTICALS INC
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
MANNITOL 25%, MANNITOL
METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE
NITROGLYCERIN, NITROGLYCERIN
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
VENOFER, IRON SUCROSE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

LUPIN

* LUPIN LTD
CEFOTAXIME SODIUM, CEFOTAXIME SODIUM
CEFPROZIL, CEFPROZIL
CEFTRIAXONE, CEFTRIAXONE SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEPHALEXIN, CEPHALEXIN
LISINOPRIL, LISINOPRIL
SUPRAX, CEFIXIME

LYNE

* LYNE LABORATORIES INC
CALCITRIOL, CALCITRIOL
LEVOCARNITINE, LEVOCARNITINE

MALLINCKRODT

* MALLINCKRODT CHEMICAL INC
ANEXSIA 7.5/650, ACETAMINOPHEN
ANEXSIA, ACETAMINOPHEN
BONTRIL, PHENDIMETRAZINE TARTRATE
BUCET, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
HYDROCET, ACETAMINOPHEN
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
LORCET-HD, ACETAMINOPHEN
LORTAB, ACETAMINOPHEN
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHADOSE, METHADONE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
TENCON, ACETAMINOPHEN

* MALLINCKRODT INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ANEXSIA 5/325, ACETAMINOPHEN
ANEXSIA 7.5/325, ACETAMINOPHEN
ANEXSIA, ACETAMINOPHEN
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE ASPARTATE
DEXTROAMPHETAMINE SULFATE, DEXTROAMPHETAMINE SULFATE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHYLIN ER, METHYLPHENIDATE HYDROCHLORIDE
METHYLIN, METHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
OPTIMARK IN PLASTIC CONTAINER, GADOVERSETAMIDE
OPTIMARK, GADOVERSETAMIDE
OXYCET, ACETAMINOPHEN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** M **

* MALLINCKRODT MEDICAL INC
CONRAY 30, IOTHALAMATE MEGLUMINE
CONRAY 400, IOTHALAMATE SODIUM
CONRAY 43, IOTHALAMATE MEGLUMINE
CONRAY, IOTHALAMATE MEGLUMINE
CYSTO-CONRAY II, IOTHALAMATE MEGLUMINE
GALLIUM CITRATE GA 67, GALLIUM CITRATE, GA-67
HEXABRIX, IOXAGLATE MEGLUMINE
INDIUM IN 111 CHLORIDE, INDIUM IN 111 CHLORIDE
MD-76R, DIATRIZOATE MEGLUMINE
MD-GASTROVIEW, DIATRIZOATE MEGLUMINE
OCTREOSCAN, INDIUM IN-111 PENTETREOTIDE KIT
OPTIRAY 160, IOVERSOL
OPTIRAY 240, IOVERSOL
OPTIRAY 300, IOVERSOL
OPTIRAY 320, IOVERSOL
OPTIRAY 350, IOVERSOL
PHOSPHOCOL P32, CHROMIC PHOSPHATE, P-32
SODIUM CHROMATE CR 51, SODIUM CHROMATE, CR-51
SODIUM IODIDE I 123, SODIUM IODIDE, I-123
SODIUM IODIDE I 131, SODIUM IODIDE, I-131
SODIUM PHOSPHATE P 32, SODIUM PHOSPHATE, P-32
TECHNECOLL, TECHNETIUM TC-99M SULFUR COLLOID KIT
TECHNESCAN MAA, TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT
TECHNESCAN MAG3, TECHNETIUM TC-99M MERTIATIDE KIT
TECHNESCAN PYP KIT, TECHNETIUM TC-99M PYROPHOSPHATE KIT
TECHNESCAN, TECHNETIUM TC-99M OXIDRONATE KIT
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE, TL-201
ULTRATAG, TECHNETIUM TC-99M RED BLOOD CELL KIT
ULTRA-TECHNEKOW FM, TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATOR
XENON XE 133, XENON, XE-133

MALLINCKRODT BAKER

* MALLINCKRODT BAKER INC
METHYLIN, METHYLPHENIDATE HYDROCHLORIDE

MARSAM PHARMS LLC

* MARSAM PHARMACEUTICALS LLC
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
CEFACLOL, CEFACLOL
CEFUROXIME SODIUM, CEFUROXIME SODIUM
DIAZEPAM, DIAZEPAM
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
ETOPOSIDE, ETOPOSIDE
HEPARIN SODIUM, HEPARIN SODIUM
ISOFLURANE, ISOFLURANE
OXACILLIN SODIUM, OXACILLIN SODIUM
TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE

MARSHALL PHARMA

* MARSHALL PHARMACAL CORP
SODIUM BUTABARBITAL, BUTABARBITAL SODIUM

MARTEC

* MARTEC PHARMACEUTICALS INC
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
PINDOLOL, PINDOLOL
* MARTEC SCIENTIFIC INC
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
NIFEDIPINE, NIFEDIPINE

MATRIX MEDCL

* MATRIX MEDICAL CORP
STERI-STAT, CHLORHEXIDINE GLUCONATE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** M ****

MAYNE PHARMA USA

* MAYNE PHARMA USA INC
ACETYLCYSTEINE, ACETYLCYSTEINE
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AQUASOL A, VITAMIN A PALMITATE
ATRACURIUM BESYLATE PRESERVATIVE FREE, ATRACURIUM BESYLATE
ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
BLEOMYCIN, BLEOMYCIN SULFATE
BUTORPHANOL TARTRATE PRESERVATIVE FREE, BUTORPHANOL TARTRATE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CARBOPLATIN, CARBOPLATIN
CYTARABINE, CYTARABINE
DACARBAZINE, DACARBAZINE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
ENALAPRILAT, ENALAPRILAT
ERYC, ERYTHROMYCIN
FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
FUODR, FLOXURIDINE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
M.V.I. ADULT (PHARMACY BULK PACKAGE), ASCORBIC ACID
M.V.I. ADULT, ASCORBIC ACID
M.V.I. PEDIATRIC, ASCORBIC ACID
M.V.I.-12 (WITHOUT VITAMIN K), ASCORBIC ACID
METHOTREXATE PRESERVATIVE FREE, METHOTREXATE SODIUM
METHOTREXATE, METHOTREXATE SODIUM
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MILRINONE LACTATE, MILRINONE LACTATE
NALBUPHINE HYDROCHLORIDE, NALBUPHINE HYDROCHLORIDE
PACLITAXEL, PACLITAXEL
PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
VINCRISTINE SULFATE PFS, VINCRISTINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE

MAYRAND

* MAYRAND INC
SEDAPAP, ACETAMINOPHEN

MCNEIL

* MCNEIL CONSUMER PRODUCTS CO DIV MCNEILAB INC
CHILDREN'S MOTRIN, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN (OTC)
IMODIUM A-D, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM ADVANCED, LOPERAMIDE HYDROCHLORIDE (OTC)
JUNIOR STRENGTH MOTRIN, IBUPROFEN (OTC)
MEDIPREN, IBUPROFEN (OTC)
MOTRIN IB, IBUPROFEN (OTC)
MOTRIN MIGRAINE PAIN, IBUPROFEN (OTC)

MCNEIL CONS SPECLT

* MCNEIL CONSUMER AND SPECIALTY PHARMACEUTICALS DIV MCNEIL PCC INC
CHILDREN'S MOTRIN COLD, IBUPROFEN (OTC)
FLEXERIL, CYCLOBENZAPRINE HYDROCHLORIDE
IMODIUM ADVANCED, LOPERAMIDE HYDROCHLORIDE (OTC)
IMODIUM, LOPERAMIDE HYDROCHLORIDE
MOTRIN, IBUPROFEN
NIZORAL A-D, KETOCONAZOLE (OTC)
NIZORAL, KETOCONAZOLE
SINE-AID IB, IBUPROFEN (OTC)
TYLENOL (CAPLET), ACETAMINOPHEN (OTC)
TYLENOL (GELTAB), ACETAMINOPHEN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** M **

MEAD JOHNSON

- * MEAD JOHNSON AND CO
CAFCIT, CAFFEINE CITRATE

MEDI FLEX INC

- * MEDI FLEX HOSP PRODUCTS INC
CHLORAPREP ONE-STEP FREPP, CHLORHEXIDINE GLUCONATE (OTC)
CHLORAPREP ONE-STEP SEPP, CHLORHEXIDINE GLUCONATE (OTC)
CHLORAPREP ONE-STEP, CHLORHEXIDINE GLUCONATE (OTC)
CHLORAPREP SINGLE SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
CHLORAPREP WITH TINT, CHLORHEXIDINE GLUCONATE (OTC)

MEDICINES CO

- * THE MEDICINES CO
ANGIOMAX, BIVALIRUDIN

MEDICIS

- * MEDICIS PHARMACEUTICAL CORP
BUPHENYL, SODIUM PHENYLBUTYRATE
DYNACIN, MINOCYCLINE HYDROCHLORIDE
LIDEX, FLUOCINONIDE
LIDEX-E, FLUOCINONIDE
LOPROX, CICLOPIROX
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
SYNACORT, HYDROCORTISONE
SYNALAR, FLUOCINOLONE ACETONIDE
VANOS, FLUOCINONIDE

MEDIMMUNE ONCOLOGY

- * MEDIMMUNE ONCOLOGY INC
ETHYOL, AMIFOSTINE
NEUTREXIN, TRIMETREXATE GLUCURONATE

MEDPOINTE

- * MEDPOINTE HEALTHCARE INC
FELBATOL, FELBAMATE

MEDPOINTE PHARM HLC

- * MEDPOINTE PHARMACEUTICALS MEDPOINTE HEALTHCARE INC
ASTELIN, AZELASTINE HYDROCHLORIDE
BUTISOL SODIUM, BUTABARBITAL SODIUM
DEPEN, PENICILLAMINE
DORAL, QUAZEPAM
LUFYLLIN, DYPHYLLINE
MILTOWN, MEPROBAMATE
OPTIVAR, AZELASTINE HYDROCHLORIDE
SOMA COMPOUND W/ CODEINE, ASPIRIN
SOMA COMPOUND, ASPIRIN
SOMA, CARISOPRODOL
VOSOL HC, ACETIC ACID, GLACIAL

MEDTRONIC

- * MEDTRONIC INC
LIORESAL, BACLOFEN

MEPHA

- * MEPHA AG
PIROXICAM, PIROXICAM

MERCK

- * MERCK AND CO INC
ALDORIL 15, HYDROCHLOROTHIAZIDE
ALDORIL 25, HYDROCHLOROTHIAZIDE
ALDORIL D30, HYDROCHLOROTHIAZIDE
ALDORIL D50, HYDROCHLOROTHIAZIDE
AMINOHIPPURATE SODIUM, AMINOHIPPURATE SODIUM
AQUAMEPHYTON, PHYTONADIONE
ARAMINE, METARAMINOL BITARTRATE
CUPRIMINE, PENICILLAMINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** M ****

* MERCK AND CO INC
DECADRON, DEXAMETHASONE
DEMSER, METYROSINE
DIUPRES-250, CHLOROTHIAZIDE
DIUPRES-500, CHLOROTHIAZIDE
DIURIL, CHLOROTHIAZIDE
DOLOBID, DIFLUNISAL
EDECRIN, ETHACRYNATE SODIUM
EDECRIN, ETHACRYNIC ACID
EMEND, APREPITANT
FOSAMAX PLUS D, ALENDRONATE SODIUM
FOSAMAX, ALENDRONATE SODIUM
HYDROCORTONE, HYDROCORTISONE
HYZAAR, HYDROCHLOROTHIAZIDE
INVANZ, ERTAPENEM SODIUM
MAXALT, RIZATRIPTAN BENZOATE
MAXALT-MLT, RIZATRIPTAN BENZOATE
MEFOXIN, CEFOXITIN SODIUM
MEPHYTON, PHYTONADIONE
MIDAMOR, AMILORIDE HYDROCHLORIDE
MINTEZOL, THIABENDAZOLE
PRIMAXIN, CILASTATIN SODIUM
PROSCAR, FINASTERIDE
SINGULAIR, MONTELUKAST SODIUM
SYPRINE, TRIENTINE HYDROCHLORIDE
TIMOLIDE 10-25, HYDROCHLOROTHIAZIDE
* MERCK RESEARCH LABORATORIES DIV MERCK CO INC
BLOCADREN, TIMOLOL MALEATE
CLINORIL, SULINDAC
COSOPT, DORZOLAMIDE HYDROCHLORIDE
COZAAR, LOSARTAN POTASSIUM
CRIXIVAN, INDINAVIR SULFATE
INDOCIN, INDOMETHACIN
LACRISERT, HYDROXYPROPYL CELLULOSE
MEFOXIN IN PLASTIC CONTAINER, CEFOXITIN SODIUM
MEVACOR, LOVASTATIN
MODURETIC 5-50, AMILORIDE HYDROCHLORIDE
NOROXIN, NORFLOXACIN
PEPCID AC (GELTAB), FAMOTIDINE (OTC)
PEPCID AC, FAMOTIDINE (OTC)
PEPCID COMPLETE, CALCIUM CARBONATE (OTC)
PEPCID PRESERVATIVE FREE IN PLASTIC CONTAINER, FAMOTIDINE
PEPCID PRESERVATIVE FREE, FAMOTIDINE
PEPCID, FAMOTIDINE
PRINIVIL, LISINOPRIL
PRINZIDE, HYDROCHLOROTHIAZIDE
PROPECIA, FINASTERIDE
SINGULAIR, MONTELUKAST SODIUM
STROMECTOL, IVERMECTIN
TIMOPTIC IN OCUDOSE, TIMOLOL MALEATE
TIMOPTIC, TIMOLOL MALEATE
TIMOPTIC-XE, TIMOLOL MALEATE
TRUSOPT, DORZOLAMIDE HYDROCHLORIDE
ZOCOR, SIMVASTATIN

MERCK RES

* MERCK RESEARCH LABORATORIES
CANCIDAS, CASPOFUNGIN ACETATE

MERETEK

* MERETEK DIAGNOSTICS INC
BREATHTEK UBT FOR H-PYLORI, UREA, C-13

MERIDIAN MEDCL TECHN

* MERIDIAN MEDICAL TECHNOLOGIES INC
ATROPEN, ATROPINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** M **

* MERIDIAN MEDICAL TECHNOLOGIES INC
EPIPEN JR., EPINEPHRINE
EPIPEN, EPINEPHRINE
LIDOPEN, LIDOCAINE HYDROCHLORIDE
MORPHINE SULFATE, MORPHINE SULFATE
PRALIDOXIME CHLORIDE, PRALIDOXIME CHLORIDE

MERZ PHARMS

* MERZ PHARMACEUTICALS LLC
ERYGEL, ERYTHROMYCIN
NAFTIN, NAFTIFINE HYDROCHLORIDE

METHAPHARM

* METHAPHARM INC
PROVOCHOLINE, METHACHOLINE CHLORIDE

MGI PHARMA INC

* MGI PHARMA INC
AGGRASTAT, TIROFIBAN HYDROCHLORIDE
GLIADEL, CARMUSTINE
HEXALEN, ALTRETAMINE
SALAGEN, PILOCARPINE HYDROCHLORIDE

MIKART

* MIKART INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAMINOPHEN, ASPIRIN, AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAMINOPHEN, BUTALBITAL, AND CAFFEINE, ACETAMINOPHEN
ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE, ACETAMINOPHEN
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMINOCAPROIC ACID, AMINOCAPROIC ACID
AMINOCAPROIC, AMINOCAPROIC ACID
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTAPAP, ACETAMINOPHEN
CARBINOXAMINE MALEATE, CARBINOXAMINE MALEATE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
ESGIC-PLUS, ACETAMINOPHEN
ETHOSUXIMIDE, ETHOSUXIMIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
ISONIAZID, ISONIAZID
METHAZOLAMIDE, METHAZOLAMIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PYRAZINAMIDE, PYRAZINAMIDE
TRIHENYPHENIDYL HYDROCHLORIDE, TRIHENYPHENIDYL HYDROCHLORIDE

MILEX

* MILEX PRODUCTS INC
MILOPHENE, CLOMIPHENE CITRATE

MILLENNIUM PHARMS

* MILLENNIUM PHARMACEUTICALS INC
VELCADE, BORTEZOMIB

MINRAD

* MINRAD INC
ENFLURANE, ENFLURANE
ISOFLURANE, ISOFLURANE

MISSION PHARMA

* MISSION PHARMACAL CO
FOLICET, FOLIC ACID
LITHOSTAT, ACETOHYDROXAMIC ACID
TIOPRONIN, TIOPRONIN
UROCIT-K, POTASSIUM CITRATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** M **

MIZA PHARMS USA

- * MIZA PHARMACEUTICALS USA INC
OCUSULF-10, SULFACETAMIDE SODIUM
TROPICAMIDE, TROPICAMIDE

MONARCH PHARMS

- * MONARCH PHARMACEUTICALS INC
CORTISPORIN, BACITRACIN ZINC
CORTISPORIN, HYDROCORTISONE
CORTISPORIN, HYDROCORTISONE ACETATE
KEMADRIN, PROCYCLIDINE HYDROCHLORIDE
MENEST, ESTROGENS, ESTERIFIED
NEOSPORIN G.U. IRRIGANT, NEOMYCIN SULFATE
NEOSPORIN, BACITRACIN ZINC
NEOSPORIN, GRAMICIDIN
PEDIOTIC, HYDROCORTISONE
POLYSPORIN, BACITRACIN ZINC
PROLOPRIM, TRIMETHOPRIM
QUIBRON-T, THEOPHYLLINE
QUIBRON-T/SR, THEOPHYLLINE
SEPTRA DS, SULFAMETHOXAZOLE
SEPTRA GRAPE, SULFAMETHOXAZOLE
SEPTRA, SULFAMETHOXAZOLE
THALITONE, CHLORTHALIDONE
VIROPTIC, TRIFLURIDINE

MORTON GROVE

- * MORTON GROVE ACQUISITION CORP
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
- * MORTON GROVE PHARMACEUTICALS INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETIC ACID, ACETIC ACID, GLACIAL
ALBUTEROL SULFATE, ALBUTEROL SULFATE
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
CARBAMAZEPINE, CARBAMAZEPINE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CROMOLYN SODIUM, CROMOLYN SODIUM
CYCLOSPORINE, CYCLOSPORINE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
ERYTHROMYCIN, ERYTHROMYCIN
FUROSEMIDE, FUROSEMIDE
GENERLAC, LACTULOSE
HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
LACTULOSE, LACTULOSE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LINDANE, LINDANE
LITHIUM CITRATE, LITHIUM CITRATE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LORATADINE, LORATADINE (OTC)
MEGESTROL ACETATE, MEGESTROL ACETATE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MYBANIL, BROMODIPHENHYDRAMINE HYDROCHLORIDE
MYCODONE, HOMATROPINE METHYLBROMIDE
MYLAMINE, DEXCHLORPHENIRAMINE MALEATE
MYMETHASONE, DEXAMETHASONE
MYPHETANE DX, BROMPHENIRAMINE MALEATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** M ****

* MORTON GROVE PHARMACEUTICALS INC
MYPROIC ACID, VALPROIC ACID
NYSTATIN, NYSTATIN
PHENTYTOIN, PHENYTOIN
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISOLONE, PREDNISOLONE
PROMETHAZINE PLAIN, PROMETHAZINE HYDROCHLORIDE
PROMETHAZINE VC PLAIN, PHENYLEPHRINE HYDROCHLORIDE
PROMETHAZINE VC W/ CODEINE, CODEINE PHOSPHATE
PROMETHAZINE W/ CODEINE, CODEINE PHOSPHATE
PROMETHAZINE W/ DEXTROMETHORPHAN, DEXTROMETHORPHAN HYDROBROMIDE
SELENIUM SULFIDE, SELENIUM SULFIDE
THEOPHYLLINE, THEOPHYLLINE
TRETINOIN, TRETINOIN
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TRIPROLIDINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE

MOUNT SINAI MEDCTR

* MOUNT SINAI MEDCTR
THALLOUS CHLORIDE TL 201, THALLOUS CHLORIDE, TL-201

MSP SINGAPORE

* MSP SINGAPORE CO LLC
VYTORIN, EZETIMIBE
ZETIA, EZETIMIBE

MUTUAL PHARM

* MUTUAL PHARMACEUTICAL CO INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAZOLAMIDE, ACETAZOLAMIDE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALLOPURINOL, ALLOPURINOL
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
BACTRIM DS, SULFAMETHOXAZOLE
BACTRIM, SULFAMETHOXAZOLE
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CARISOPRODOL, CARISOPRODOL
CHLORZOXAZONE, CHLORZOXAZONE
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE ASPARTATE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
ERGOLOID MESYLATES, ERGOLOID MESYLATES
FAMOTIDINE, FAMOTIDINE
FELODIPINE, FELODIPINE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
FOLIC ACID, FOLIC ACID
GABAPENTIN, GABAPENTIN
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
INDOMETHACIN, INDOMETHACIN
ISONIAZID, ISONIAZID
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** M **

* MUTUAL PHARMACEUTICAL CO INC
LORAZEPAM, LORAZEPAM
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRONIDAZOLE, METRONIDAZOLE
MINOXIDIL, MINOXIDIL
NYSTATIN, NYSTATIN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
OXYCODONE AND ASPIRIN, ASPIRIN
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PINDOLOL, PINDOLOL
PIROXICAM, PIROXICAM
PREDNISONE, PREDNISONE
PRIMIDONE, PRIMIDONE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
QUINIDINE GLUCONATE, QUINIDINE GLUCONATE
QUINIDINE SULFATE, QUINIDINE SULFATE
QUININE SULFATE, QUININE SULFATE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
SPIRONOLACTONE, SPIRONOLACTONE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SULFASALAZINE, SULFASALAZINE
SULINDAC, SULINDAC
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
TOLAZAMIDE, TOLAZAMIDE
TOLMETIN SODIUM, TOLMETIN SODIUM
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
ZONISAMIDE, ZONISAMIDE

MUTUAL PHARMA

* MUTUAL PHARMACAL CO
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
KETOCONAZOLE, KETOCONAZOLE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
TRIMETHOBENZAMIDE HYDROCHLORIDE, TRIMETHOBENZAMIDE HYDROCHLORIDE

MYLAN

* MYLAN LABORATORIES INC
ACYCLOVIR, ACYCLOVIR
AMLODIPINE BESYLATE, AMLODIPINE BESYLATE
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
CAPTOPRIL, CAPTOPRIL
ETODOLAC, ETODOLAC
IBUPROFEN, IBUPROFEN
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
* MYLAN PHARMACEUTICALS INC
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
ACYCLOVIR, ACYCLOVIR
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALLOPURINOL, ALLOPURINOL
ALPRAZOLAM, ALPRAZOLAM
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** M ****

* MYLAN PHARMACEUTICALS INC
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BROMOCRIPTINE MESYLATE, BROMOCRIPTINE MESYLATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
CARBIDOPA AND LEVODOPA, CARBIDOPA
CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
CHLOROTHIAZIDE, CHLOROTHIAZIDE
CHLORPROPAMIDE, CHLORPROPAMIDE
CHLORTHALIDONE, CHLORTHALIDONE
CIMETIDINE, CIMETIDINE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
CLORPRES, CHLORTHALIDONE
CLOZAPINE, CLOZAPINE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
CYSTAGON, CYSTEAMINE BITARTRATE
DIAZEPAM, DIAZEPAM
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ERYTHROMYCIN ETHYLSUCCINATE, ERYTHROMYCIN ETHYLSUCCINATE
ERYTHROMYCIN STEARATE, ERYTHROMYCIN STEARATE
ESTRADIOL, ESTRADIOL
ESTROPIPATE, ESTROPIPATE
ETODOLAC, ETODOLAC
EXTENDED PHENYTOIN SODIUM, PHENYTOIN SODIUM
FAMOTIDINE, FAMOTIDINE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
FLUCONAZOLE, FLUCONAZOLE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
FLURBIPROFEN, FLURBIPROFEN
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
FUROSEMIDE, FUROSEMIDE
GEMFIBROZIL, GEMFIBROZIL
GLIMEPIRIDE, GLIMEPIRIDE
GLIPIZIDE, GLIPIZIDE
GLYBURIDE (MICRONIZED), GLYBURIDE
GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
HALOPERIDOL, HALOPERIDOL
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
INDAPAMIDE, INDAPAMIDE
INDOMETHACIN, INDOMETHACIN
KETOCONAZOLE, KETOCONAZOLE
KETOPROFEN, KETOPROFEN
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** M ****

* MYLAN PHARMACEUTICALS INC
LEVOTHYROXINE SODIUM, LEVOTHYROXINE SODIUM
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
LORAZEPAM, LORAZEPAM
LOVASTATIN, LOVASTATIN
LOXAPINE, LOXAPINE SUCCINATE
MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE
MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
MERCAPTOPYRINE, MERCAPTOPYRINE ANHYDROUS
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
METHYLCLOTHIAZIDE, METHYLCLOTHIAZIDE
METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METHYLDOPA, METHYLDOPA
METOLAZONE, METOLAZONE
METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NADOLOL, NADOLOL
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN, NAPROXEN
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NIZATIDINE, NIZATIDINE
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
OMEPRazole, OMEPRazole
OXAPROZIN, OXAPROZIN
PACLITAXEL, PACLITAXEL
PENTOXIFYLLINE, PENTOXIFYLLINE
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
PHENYTEK, PHENYTOIN SODIUM
PINDOLOL, PINDOLOL
PIROXICAM, PIROXICAM
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
PROBENECID, PROBENECID
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPOXYPHENE HYDROCHLORIDE, PROPOXYPHENE HYDROCHLORIDE
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
SPIRONOLACTONE, SPIRONOLACTONE
SULINDAC, SULINDAC
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
TEMAZEPAM, TEMAZEPAM
TETRACYCLINE HYDROCHLORIDE, TETRACYCLINE HYDROCHLORIDE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
THIOXIXENE, THIOXIXENE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TOLMETIN SODIUM, TOLMETIN SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** M ****

* MYLAN PHARMACEUTICALS INC
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
ZONISAMIDE, ZONISAMIDE

MYLAN BERTEK

* MYLAN BERTEK PHARMACEUTICALS INC
AVITA, TRETINOIN
MAXZIDE, HYDROCHLOROTHIAZIDE
MAXZIDE-25, HYDROCHLOROTHIAZIDE
MENTAX, BUTENAFINE HYDROCHLORIDE
MENTAX-TC, BUTENAFINE HYDROCHLORIDE
SULFAMYLON, MAFENIDE ACETATE

MYLAN TECHNOLOGIES

* MYLAN TECHNOLOGIES INC
ESTRADIOL, ESTRADIOL
FENTANYL, FENTANYL
NITROGLYCERIN, NITROGLYCERIN
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

NABI

* NABI BIOPHARMACEUTICALS
ALOPRIM, ALLOPURINOL SODIUM
PHOSLO GELCAPS, CALCIUM ACETATE
PHOSLO, CALCIUM ACETATE

NEIL

* NEIL LABORATORIES INC
CARISOPRODOL, CARISOPRODOL
IBUPROFEN, IBUPROFEN (OTC)

NEPHRON

* NEPHRON CORP
ALBUTEROL SULFATE, ALBUTEROL SULFATE
BETA-2, ISOETHARINE HYDROCHLORIDE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
* NEPHRON PHARMACEUTICALS CORP
ALBUTEROL SULFATE, ALBUTEROL SULFATE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE

NEW RIVER

* NEW RIVER PHARMACEUTICALS INC
PROFERDEX, IRON DEXTRAN

NICE PAK

* NICE PAK PRODUCTS INC
CHLORASCRUB MAXI SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB SWAB, CHLORHEXIDINE GLUCONATE (OTC)
CHLORASCRUB SWABSTICK, CHLORHEXIDINE GLUCONATE (OTC)

NITROMED

* NITROMED INC
BIDIL, HYDRALAZINE HYDROCHLORIDE

NORTH SHORE LIJ

* NORTH SHORE LIJ RESEARCH INSTITUTE
FLUDEOXYGLUCOSE F18, FLUDEOXYGLUCOSE F-18

NOVARTIS

* NOVARTIS CONSUMER HEALTH INC
DENA VIR, PENCICLOVIR SODIUM
EXCEDRIN (MIGRAINE), ACETAMINOPHEN (OTC)
HABITROL, NICOTINE (OTC)
LAMISIL AT, TERBINA FINE HYDROCHLORIDE (OTC)
LAMISIL, TERBINA FINE
LAMISIL, TERBINA FINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** N ****

* NOVARTIS CONSUMER HEALTH INC
LAMISIL, TERBINAFINE HYDROCHLORIDE (OTC)
TAVIST ALLERGY/SINUS/HEADACHE, ACETAMINOPHEN (OTC)
TRANSDERM SCOP, SCOPOLAMINE
VAGISTAT-1, TIOCONAZOLE (OTC)

* NOVARTIS PHARMACEUTICALS CORP
ANTURANE, SULFINPYRAZONE
APRESAZIDE, HYDRALAZINE HYDROCHLORIDE
APRESOLINE, HYDRALAZINE HYDROCHLORIDE
AREDIA, PAMIDRONATE DISODIUM
CAFERGOT, CAFFEINE
CARBASTAT, CARBACHOL
CATAFLAM, DICLOFENAC POTASSIUM
CLOZARIL, CLOZAPINE
COMBIPATCH, ESTRADIOL
CYTADREN, AMINOGLUTETHIMIDE
DESFERAL, DEFEROXAMINE MESYLATE
DIOVAN HCT, HYDROCHLOROTHIAZIDE
DIOVAN, VALSARTAN
ELIDEL, PIMECROLIMUS
ENABLEX, DARIFENACIN HYDROBROMIDE
ESIDRIX, HYDROCHLOROTHIAZIDE
ESTRADERM, ESTRADIOL
EXELON, RIVASTIGMINE TARTRATE
EXJADE, DEFERASIROX
FAMVIR, FAMCICLOVIR
FEMARA, LETROZOLE
FLUOR-OP, FLUOROMETHOLONE
FOCALIN XR, DEXMETHYLPHENIDATE HYDROCHLORIDE
FOCALIN, DEXMETHYLPHENIDATE HYDROCHLORIDE
FORADIL, FORMOTEROL FUMARATE
GENTACIDIN, GENTAMICIN SULFATE
GLEEVEC, IMATINIB MESYLATE
HYDERGINE, ERGOLOID MESYLATES
INFLAMASE FORTE, PREDNISOLONE SODIUM PHOSPHATE
INFLAMASE MILD, PREDNISOLONE SODIUM PHOSPHATE
LAMISIL, TERBINAFINE HYDROCHLORIDE
LESCOL XL, FLUVASTATIN SODIUM
LESCOL, FLUVASTATIN SODIUM
LIVOSTIN, LEVOCABASTINE HYDROCHLORIDE
LOPRESSOR HCT, HYDROCHLOROTHIAZIDE
LOPRESSOR, METOPROLOL TARTRATE
LOTENSIN HCT, BENAZEPRIL HYDROCHLORIDE
LOTENSIN, BENAZEPRIL HYDROCHLORIDE
LOTREL, AMLODIPINE BESYLATE
METHERGINE, METHYLERGONOVINE MALEATE
METOPIRONE, METYRAPONE
MIACALCIN, CALCITONIN, SALMON
MIOCHOL, ACETYLCHOLINE CHLORIDE
MIOCHOL-E, ACETYLCHOLINE CHLORIDE
MYFORTIC, MYCOPHENOLIC ACID
NEORAL, CYCLOSPORINE
OCUPRESS, CARTEOLOL HYDROCHLORIDE
PARLODEL, BROMOCRIPTINE MESYLATE
PBZ, TRIPELENNAMINE HYDROCHLORIDE
REGITINE, PHENTOLAMINE MESYLATE
RESCULA, UNOPROSTONE ISOPROPYL
RITALIN LA, METHYLPHENIDATE HYDROCHLORIDE
RITALIN, METHYLPHENIDATE HYDROCHLORIDE
RITALIN-SR, METHYLPHENIDATE HYDROCHLORIDE
SANDIMMUNE, CYCLOSPORINE
SANDOSTATIN LAR, OCTREOTIDE ACETATE
SANDOSTATIN, OCTREOTIDE ACETATE
STARLIX, NATEGLINIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** N ****

* NOVARTIS PHARMACEUTICALS CORP
SULF-10, SULFACETAMIDE SODIUM
TAVIST-1, CLEMASTINE FUMARATE (OTC)
TEGRETOL, CARBAMAZEPINE
TEGRETOL-XR, CARBAMAZEPINE
TORECAN, THIETHYLPERAZINE MALEATE
TRILEPTAL, OXCARBAZEPINE
VASOCIDIN, PREDNISOLONE SODIUM PHOSPHATE
VASOCON, NAPHAZOLINE HYDROCHLORIDE
VASOCON-A, ANTAZOLINE PHOSPHATE (OTC)
VISKEN, PINDOLOL
VIVELLE, ESTRADIOL
VIVELLE-DOT, ESTRADIOL
VOLTAREN, DICLOFENAC SODIUM
VOLTAREN-XR, DICLOFENAC SODIUM
ZADITOR, KETOTIFEN FUMARATE
ZELNORM, TEGASEROD MALEATE
ZOMETA, ZOLEDRONIC ACID

NOVAVAX

* NOVAVAX INC
ESTRASORB, ESTRADIOL HEMIHYDRATE

NOVEN

* NOVEN PHARMACEUTICALS INC
LIDOCAINE, LIDOCAINE

NOVEX

* NOVEX PHARMA
ALBUTEROL SULFATE, ALBUTEROL SULFATE
BETAXOLOL, BETAXOLOL HYDROCHLORIDE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CARTEOLOL HYDROCHLORIDE, CARTEOLOL HYDROCHLORIDE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
CROMOLYN SODIUM, CROMOLYN SODIUM
CYCLOSPORINE, CYCLOSPORINE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LACTULOSE, LACTULOSE
LEVOBUNOLOL HYDROCHLORIDE, LEVOBUNOLOL HYDROCHLORIDE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
OFLOXACIN, OFLOXACIN
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
TOBRAMYCIN, TOBRAMYCIN

NOVO NORDISK

* NOVO NORDISK PHARMACEUTICALS INC
GLUCAGEN, GLUCAGON HYDROCHLORIDE RECOMBINANT

NOVO NORDISK INC

* NOVO NORDISK INC
ACTIVELLA, ESTRADIOL
INNOFEM, ESTRADIOL
LEVEMIR, INSULIN DETEMIR RECOMBINANT
NORDITROPIN NORDIFLEX, SOMATROPIN RECOMBINANT
NORDITROPIN, SOMATROPIN RECOMBINANT
NOVOLIN 70/30, INSULIN RECOMBINANT HUMAN (OTC)
NOVOLIN L, INSULIN ZINC SUSP RECOMBINANT HUMAN (OTC)
NOVOLIN N, INSULIN SUSP ISOPHANE RECOMBINANT HUMAN (OTC)
NOVOLIN R, INSULIN RECOMBINANT HUMAN (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** N ****

* NOVO NORDISK INC
NOVOLOG MIX 70/30, INSULIN ASPART PROTAMINE RECOMBINANT
NOVOLOG, INSULIN ASPART RECOMBINANT
PRANDIN, REPAGLINIDE
VAGIFEM, ESTRADIOL
VELOSULIN BR, INSULIN RECOMBINANT HUMAN (OTC)

NOVOCOL

* NOVOCOL PHARMACEUTICAL INC
ISOCAINE HYDROCHLORIDE W/ LEVONORDEFIN, LEVONORDEFIN
ISOCAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE

OCLASSEN

* OCLASSEN PHARMACEUTICALS INC
CONDYLOX, PODOFILOX
CORDRAN SP, FLURANDRENOLIDE
CORDRAN, FLURANDRENOLIDE
MONODOX, DOXYCYCLINE

ODYSSEY PHARMS

* ODYSSEY PHARMACEUTICALS INC
ANTABUSE, DISULFIRAM
NYSTATIN, NYSTATIN
SURMONTIL, TRIMIPRAMINE MALEATE
URECHOLINE, BETHANECHOL CHLORIDE
VIVACTIL, PROTRIPTYLINE HYDROCHLORIDE
VOSPIRE ER, ALBUTEROL SULFATE

OHM

* OHM CORP
IBUPROFEN, IBUPROFEN (OTC)

OHM LABS

* OHM LABORATORIES INC
CHLORZOXAZONE, CHLORZOXAZONE
IBUPROFEN, IBUPROFEN
IBUPROHM COLD AND SINUS, IBUPROFEN (OTC)
IBUPROHM, IBUPROFEN
IBUPROHM, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

ONY

* ONY INC
INFASURF PRESERVATIVE FREE, CALFACTANT

OPTOPICS

* OPTOPICS LABORATORIES CORP
PARACAINE, PROPARACAINE HYDROCHLORIDE
SULFACEL-15, SULFACETAMIDE SODIUM

ORAPHARMA

* ORAPHARMA INC
ARESTIN, MINOCYCLINE HYDROCHLORIDE

ORCHID HLTHCARE

* ORCHID HEALTHCARE
CEFAZOLIN SODIUM, CEFZOLIN SODIUM
CEFAZOLIN, CEFZOLIN SODIUM
CEFPROZIL, CEFPROZIL
CEFTRIAXONE, CEFTRIAXONE SODIUM
CEPHALEXIN, CEPHALEXIN

ORGANON USA INC

* ORGANON USA INC
CYCLESSA, DESOGESTREL
DESOGEN, DESOGESTREL
FOLLISTIM AQ, FOLLITROPIN ALFA/BETA
GANIRELIX ACETATE INJECTION, GANIRELIX ACETATE
HEXADROL, DEXAMETHASONE
MIRCETTE, DESOGESTREL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** 0 **

* ORGANON USA INC
NUVARING, ETHINYL ESTRADIOL
PREGNYL, GONADOTROPIN, CHORIONIC
REMERON SOLTAB, MIRTAZAPINE
REMERON, MIRTAZAPINE
SUCCINYLCHOLINE CHLORIDE, SUCCINYLCHOLINE CHLORIDE
ZEMURON (P/F), ROCURONIUM BROMIDE

ORION

* ORION CORP
COMTAN, ENTACAPONE

ORION PHARMA INC

* ORION PHARMACEUTICALS INC
STALEVO 100, CARBIDOPA
STALEVO 150, CARBIDOPA
STALEVO 50, CARBIDOPA

ORPHAN MEDCL

* ORPHAN MEDICAL INC
ANTIZOL, FOMEPIZOLE
CYSTADANE, BETAINE, ANHYDROUS
XYREM, SODIUM OXYBATE

ORTHO BIOTECH

* ORTHO BIOTECH PRODUCTS LP
LEUSTATIN, CLADRIBINE

ORTHO MCNEIL

* ORTHO MCNEIL PHARMACEUTICAL
HALDOL, HALOPERIDOL DECANOATE
HALDOL, HALOPERIDOL LACTATE

ORTHO MCNEIL PHARM

* ALZA CORP
URISPAS, FLAVOXATE HYDROCHLORIDE
* ORTHO MCNEIL PHARMACEUTICAL INC
AXERT, ALMOTRIPTAN MALATE
ELMIRON, PENTOSAN POLYSULFATE SODIUM
FLOXIN, OFLOXACIN
LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER, LEVOFLOXACIN
LEVAQUIN, LEVOFLOXACIN
MICRONOR, NORETHINDRONE
MODICON 28, ETHINYL ESTRADIOL
MONISTAT 3, MICONAZOLE NITRATE
ORTHO EVRA, ETHINYL ESTRADIOL
ORTHO TRI-CYCLEN LO, ETHINYL ESTRADIOL
ORTHO-CEPT, DESOGESTREL
ORTHO-NOVUM 1/35-28, ETHINYL ESTRADIOL
ORTHO-NOVUM 1/50 28, MESTRANOL
ORTHO-NOVUM 10/11-28, ETHINYL ESTRADIOL
ORTHO-NOVUM 7/7/7-28, ETHINYL ESTRADIOL
PARAFON FORTE DSC, CHLORZOXAZONE
TERAZOL 3, TERCONAZOLE
TERAZOL 7, TERCONAZOLE
TOLECTIN 600, TOLMETIN SODIUM
TOLECTIN DS, TOLMETIN SODIUM
TOLECTIN, TOLMETIN SODIUM
TOPAMAX SPRINKLE, TOPIRAMATE
TOPAMAX, TOPIRAMATE
TYLENOL W/ CODEINE NO. 3, ACETAMINOPHEN
TYLOX, ACETAMINOPHEN
ULTRACET, ACETAMINOPHEN
ULTRAM, TRAMADOL HYDROCHLORIDE

OSCIENT

* OSCIENT PHARMACEUTICALS
FACTIVE, GEMIFLOXACIN MESYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** 0 ****

OSI PHARMS

* OSI PHARMACEUTICALS INC
TARCEVA, ERLOTINIB HYDROCHLORIDE

OSMOTICA PHARM

* OSMOTICA PHARMACEUTICAL CORP
NIFEDIPINE, NIFEDIPINE

OTSUKA

* OTSUKA AMERICA PHARMACEUTICALS INC
ABILIFY, ARIPIPRAZOLE
* OTSUKA PHARMACEUTICAL CO LTD
ABILIFY, ARIPIPRAZOLE
PLETAL, CILOSTAZOL

OVATION PHARMS

* OVATION PHARMACEUTICALS INC
CHEMET, SUCCIMER
COGENTIN, BENZTROPINE MESYLATE
COSMEGEN, DACTINOMYCIN
DESOXYN, METHAMPHETAMINE HYDROCHLORIDE
DIURIL, CHLOROTHIAZIDE SODIUM
INDOCIN I.V., INDOMETHACIN SODIUM
MUSTARGEN, MECHLORETHAMINE HYDROCHLORIDE
NEMBUTAL SODIUM, PENTOBARBITAL SODIUM
PEGANONE, ETHOTOIN
TRANXENE SD, CLORAZEPATE DIPOTASSIUM
TRANXENE, CLORAZEPATE DIPOTASSIUM

OXFORD PHARM

* OXFORD PHARMACEUTICAL SERVICES INC
MARPLAN, ISOCARBOXAZID

PACIFIC PHARMA

* PACIFIC PHARMA
TIMOLOL MALEATE, TIMOLOL MALEATE
* PACIFIC PHARMA INC
TIMOLOL MALEATE, TIMOLOL MALEATE

PADDOCK

* PADDOCK LABORATORIES INC
AMMONIUM LACTATE, AMMONIUM LACTATE
CLINDA-DERM, CLINDAMYCIN PHOSPHATE
CLOTRIMAZOLE, CLOTRIMAZOLE
COLISTIMETHATE, COLISTIMETHATE SODIUM
COLOCORT, HYDROCORTISONE
COMPRO, PROCHLORPERAZINE
DIHYDROERGOTAMINE MESYLATE, DIHYDROERGOTAMINE MESYLATE
ERYTHRA-DERM, ERYTHROMYCIN
FLAVOXATE HYDROCHLORIDE, FLAVOXATE HYDROCHLORIDE
KIONEX, SODIUM POLYSTYRENE SULFONATE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
NYSTATIN, NYSTATIN
NYSTOP, NYSTATIN
PODOFILOX, PODOFILOX
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE

PAR PHARM

* PAR PHARMACEUTICAL INC
ALLOPURINOL, ALLOPURINOL
AMILORIDE HYDROCHLORIDE, AMILORIDE HYDROCHLORIDE
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CABERGOLINE, CABERGOLINE
CAPOTEN, CAPTOPRIL
CARISOPRODOL AND ASPIRIN, ASPIRIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** P ****

* PAR PHARMACEUTICAL INC
CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
CHLORPROPAMIDE, CHLORPROPAMIDE
CHLORZOXAZONE, CHLORZOXAZONE
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
CHOLESTYRAMINE, CHOLESTYRAMINE
CLOMIPHENE CITRATE, CLOMIPHENE CITRATE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
DEXAMETHASONE, DEXAMETHASONE
DIAZEPAM, DIAZEPAM
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
DOXYCYCLINE, DOXYCYCLINE
FENOFIBRATE, FENOFIBRATE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
GLIMEPIRIDE, GLIMEPIRIDE
HALOPERIDOL, HALOPERIDOL
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDRA-ZIDE, HYDRALAZINE HYDROCHLORIDE
HYDROFLUMETHIAZIDE, HYDROFLUMETHIAZIDE
HYDRO-RIDE, AMILORIDE HYDROCHLORIDE
HYDROXYUREA, HYDROXYUREA
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
INDOMETHACIN, INDOMETHACIN
ISOPTIN SR, VERAPAMIL HYDROCHLORIDE
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LISINOPRIL, LISINOPRIL
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
MEGACE ES, MEGESTROL ACETATE
MEGESTROL ACETATE, MEGESTROL ACETATE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
METHOCARBAMOL AND ASPIRIN, ASPIRIN
METHOCARBAMOL, METHOCARBAMOL
METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METHYLDOPA, METHYLDOPA
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRONIDAZOLE, METRONIDAZOLE
MINOXIDIL, MINOXIDIL
NYSTATIN, NYSTATIN
OFLOXACIN, OFLOXACIN
ORPHENGESIC FORTE, ASPIRIN
ORPHENGESIC, ASPIRIN
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
SULFAMETHOPRIM, SULFAMETHOXAZOLE
SULFINPYRAZONE, SULFINPYRAZONE
SUMYCIN, TETRACYCLINE HYDROCHLORIDE
TEMAZEPAM, TEMAZEPAM
TOLAZAMIDE, TOLAZAMIDE
TORSEMIDE, TORSEMIDE

PARENTA PHARMS

* PARENTA PHARMACEUTICALS INC
DIAZEPAM, DIAZEPAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** P ****

PARKE DAVIS

- * PARKE DAVIS DIV WARNER LAMBERT CO
BENADRYL PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
BENADRYL, DIPHENHYDRAMINE HYDROCHLORIDE
CELONTIN, METHSUXIMIDE
CEREBYX, FOSPHENYTOIN SODIUM
CHOLEDYL SA, OXTRIPHYLLINE
DILANTIN, PHENYTOIN SODIUM
DILANTIN-125, PHENYTOIN
NARDIL, PHENELZINE SULFATE
NEURONTIN, GABAPENTIN
ZARONTIN, ETHOSUXIMIDE
- * PARKE DAVIS PHARMACEUTICAL RESEARCH DIV WARNER LAMBERT CO
ZARONTIN, ETHOSUXIMIDE

PARKEDALE

- * PARKEDALE PHARMACEUTICALS INC
CHLOROMYCETIN, CHLORAMPHENICOL SODIUM SUCCINATE
COLY-MYCIN M, COLISTIMETHATE SODIUM
COLY-MYCIN S, COLISTIN SULFATE
KETALAR, KETAMINE HYDROCHLORIDE
PITOCIN, OXYTOCIN

PEDINOL

- * PEDINOL PHARMACAL INC
GRIS-PEG, GRISEOFULVIN, ULTRAMICROCRYSTALLINE
NALFON 200, FENOPROFEN CALCIUM
NALFON, FENOPROFEN CALCIUM

PERRIGO

- * L PERRIGO CO
ACETAMINOPHEN, ACETAMINOPHEN (OTC)
ACETAMINOPHEN, ASPIRIN AND CAFFEINE, ACETAMINOPHEN (OTC)
CAP-PROFEN, IBUPROFEN (OTC)
CHILDREN'S IBUPROFEN, IBUPROFEN (OTC)
CROMOLYN SODIUM, CROMOLYN SODIUM (OTC)
DOXYLAMINE SUCCINATE, DOXYLAMINE SUCCINATE (OTC)
FAMOTIDINE, FAMOTIDINE (OTC)
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE, LOPERAMIDE HYDROCHLORIDE (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
LORATADINE, LORATADINE (OTC)
MICONAZOLE 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE, NAPROXEN SODIUM (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
TAB-PROFEN, IBUPROFEN (OTC)
TIOCONAZOLE, TIOCONAZOLE (OTC)
- * PERRIGO CO
CIMETIDINE, CIMETIDINE (OTC)
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
FAMOTIDINE, FAMOTIDINE
IBUPROFEN, IBUPROFEN (OTC)
JUNIOR STRENGTH IBUPROFEN, IBUPROFEN (OTC)
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
MICONAZOLE NITRATE COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)
NICOTINE POLACRILEX (MINT), NICOTINE POLACRILEX (OTC)
NICOTINE POLACRILEX (ORANGE), NICOTINE POLACRILEX (OTC)
NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** P ****

PERRIGO R AND D

- * PERRIGO R AND D CO
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
NAPROXEN, NAPROXEN

PERSONAL PRODS

- * PERSONAL PRODUCTS CO
MONISTAT 1 COMBINATION PACK, MICONAZOLE NITRATE (OTC)
MONISTAT 3 COMBINATION PACK (PREFILLED), MICONAZOLE NITRATE (OTC)
MONISTAT 3 COMBINATION PACK, MICONAZOLE NITRATE (OTC)

PFIZER

- * PFIZER AGRICULTURAL DIV
ANTIVERT, MECLIZINE HYDROCHLORIDE
DEPO-SUBQ PROVERA 104, MEDROXYPROGESTERONE ACETATE
NORVASC, AMLODIPINE BESYLATE
- * PFIZER CENTRAL RESEARCH
DIFLUCAN, FLUCONAZOLE
ZITHROMAX, AZITHROMYCIN
- * PFIZER CHEMICALS DIV PFIZER INC
DIFLUCAN, FLUCONAZOLE
ZITHROMAX, AZITHROMYCIN
- * PFIZER INC
CADUET, AMLODIPINE BESYLATE
CARDURA XL, DOXAZOSIN MESYLATE
DIFLUCAN IN DEXTROSE 5% IN PLASTIC CONTAINER, FLUCONAZOLE
DIFLUCAN IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
DIFLUCAN IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
EMETE-CON, BENZQUINAMIDE HYDROCHLORIDE
GEODON, ZIPRASIDONE HYDROCHLORIDE
GEODON, ZIPRASIDONE MESYLATE
GLUCOTROL XL, GLIPIZIDE
GLUCOTROL, GLIPIZIDE
LIPITOR, ATORVASTATIN CALCIUM
LITHIUM CARBONATE, LITHIUM CARBONATE
MINIPRESS XL, PRAZOSIN HYDROCHLORIDE
MINIZIDE, POLYTHIAZIDE
NAVANE, THIOTHIXENE
NAVANE, THIOTHIXENE HYDROCHLORIDE
PROCARDIA, NIFEDIPINE
RENESE, POLYTHIAZIDE
REVATIO, SILDENAFIL CITRATE
RID MOUSSE, PIPERONYL BUTOXIDE (OTC)
UNASYN, AMPICILLIN SODIUM
VFEND, VORICONAZOLE
VISINE-A, NAPHAZOLINE HYDROCHLORIDE (OTC)
ZITHROMAX, AZITHROMYCIN
ZYTEC, CETIRIZINE HYDROCHLORIDE
ZYTEC-D 12 HOUR, CETIRIZINE HYDROCHLORIDE
- * PFIZER LABORATORIES DIV PFIZER INC
CARDURA, DOXAZOSIN MESYLATE
CEFOBID IN PLASTIC CONTAINER, CEFOPERAZONE SODIUM
CEFOBID, CEFOPERAZONE SODIUM
DIABINESE, CHLORPROPAMIDE
FELDENE, PIROXICAM
GEOCILLIN, CARBENICILLIN INDANYL SODIUM
MINIPRESS, PRAZOSIN HYDROCHLORIDE
MITHRACIN, PLICAMYCIN
MODERIL, RESCINNAMINE
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
PERMAPEN, PENICILLIN G BENZATHINE
PFIZERPEN, PENICILLIN G POTASSIUM
PROCARDIA XL, NIFEDIPINE
RENESE-R, POLYTHIAZIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** P ****

- * PFIZER LABORATORIES DIV PFIZER INC
SINEQUAN, DOXEPIN HYDROCHLORIDE
SPECTROBID, BACAMPICILLIN HYDROCHLORIDE
STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
TERRA-CORTRIL, HYDROCORTISONE ACETATE
TERRAMYCIN W/ POLYMYXIN B SULFATE, OXYTETRACYCLINE HYDROCHLORIDE
TERRAMYCIN, LIDOCAINE HYDROCHLORIDE
TERRAMYCIN, OXYTETRACYCLINE HYDROCHLORIDE
UNASYN, AMPICILLIN SODIUM
UNISOM, DOXYLAMINE SUCCINATE (OTC)
VIBRAMYCIN, DOXYCYCLINE
VIBRAMYCIN, DOXYCYCLINE CALCIUM
VIBRAMYCIN, DOXYCYCLINE HYCLATE
VIBRA-TABS, DOXYCYCLINE HYCLATE
VISINE L.R., OXYMETAZOLINE HYDROCHLORIDE (OTC)
VISTARIL, HYDROXYZINE HYDROCHLORIDE
VISTARIL, HYDROXYZINE PAMOATE
- * PFIZER PHARMACEUTICALS INC
TAO, TROLEANDOMYCIN
ZOLOFT, SERTRALINE HYDROCHLORIDE
- * PFIZER PHARMACEUTICALS PRODUCTION CORP LTD
TIKOSYN, DOFETILIDE

PFIZER CONS HLTHCARE

- * PFIZER CONSUMER HEALTHCARE DIV WARNER LAMBERT CO
ZANTAC 150, RANITIDINE HYDROCHLORIDE (OTC)

PFIZER GLOBAL

- * PFIZER GLOBAL RESEARCH DEVELOPMENT
ZMAX, AZITHROMYCIN

PFIZER INC

- * PFIZER INC
ELLENC, EPIRUBICIN HYDROCHLORIDE

PFIZER IRELAND

- * PFIZER IRELAND PHARMACEUTICALS
RELPA, ELETRIPTAN HYDROBROMIDE
VIAGRA, SILDENAFIL CITRATE

PFIZER PHARMS

- * PFIZER PHARMACEUTICALS LTD
ACCUPRIL, QUINAPRIL HYDROCHLORIDE
ACCURETIC, HYDROCHLOROTHIAZIDE
DILANTIN, PHENYTOIN
LOPID, GEMFIBROZIL
NEURONTIN, GABAPENTIN
NITROSTAT, NITROGLYCERIN

PHARM ASSOC

- * PHARMACEUTICAL ASSOC INC
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
PERPHENAZINE, PERPHENAZINE
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
- * PHARMACEUTICAL ASSOC INC DIV BEACH PRODUCTS
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
ETHOSUXIMIDE, ETHOSUXIMIDE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
HALOPERIDOL, HALOPERIDOL LACTATE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
LACTULOSE, LACTULOSE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

- * PHARMACEUTICAL ASSOC INC DIV BEACH PRODUCTS
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PREDNISOLONE, PREDNISOLONE
PROMETHAZINE HYDROCHLORIDE AND CODEINE PHOSPHATE, CODEINE PHOSPHATE
THEOPHYLLINE, THEOPHYLLINE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
VALPROIC ACID, VALPROIC ACID

PHARM FORM

- * PHARMACEUTICAL FORMULATIONS INC
CIMETIDINE, CIMETIDINE (OTC)
FOLIC ACID, FOLIC ACID
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE, IBUPROFEN (OTC)
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
PREDNISONE, PREDNISONE
QUINIDINE SULFATE, QUINIDINE SULFATE

PHARM SPEC

- * PHARMACEUTICAL SPECIALIST ASSOC
GENTAMICIN SULFATE, GENTAMICIN SULFATE

PHARM VENTURES

- * PHARMACEUTICAL VENTURES LTD
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

PHARMA SERVE NY

- * PHARMA SERVE INC SUB TORIGIAN LABORATORIES
AMINOPHYLLINE, AMINOPHYLLINE

PHARMA TEK

- * PHARMA TEK INC
AMPHOTERICIN B, AMPHOTERICIN B
BACIIM, BACITRACIN
BACI-RX, BACITRACIN
COLISTIMETHATE, COLISTIMETHATE SODIUM
HYDROCORTISONE ACETATE, HYDROCORTISONE ACETATE
HYDROCORTISONE, HYDROCORTISONE
NEO-RX, NEOMYCIN SULFATE
POLYMYXIN B SULFATE, POLYMYXIN B SULFATE
POLY-RX, POLYMYXIN B SULFATE
STREPTOMYCIN SULFATE, STREPTOMYCIN SULFATE
TOBRAMYCIN, TOBRAMYCIN SULFATE

PHARMACHEMIE

- * PHARMACHEMIE BV
CARBOPLATIN, CARBOPLATIN
CISPLATIN, CISPLATIN
DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
ETOPOSIDE, ETOPOSIDE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE

PHARMACHEMIE USA

- * PHARMACHEMIE USA INC
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM

PHARMACIA

- * PHARMACIA CORP
MAXAQUIN, LOMEFLOXACIN HYDROCHLORIDE

PHARMACIA AND UPJOHN

- * PHARMACIA AND UPJOHN
XANAX XR, ALPRAZOLAM
- * PHARMACIA AND UPJOHN CO
ADRIAMYCIN PFS, DOXORUBICIN HYDROCHLORIDE
ANSAID, FLURBIPROFEN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

* PHARMACIA AND UPJOHN CO
 AROMASIN, EXEMESTANE
 AZULFIDINE EN-TABS, SULFASALAZINE
 AZULFIDINE, SULFASALAZINE
 BACITRACIN, BACITRACIN
 CAMPTOSAR, IRINOTECAN HYDROCHLORIDE
 CAVERJECT, ALPROSTADIL
 CLEOCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLEOCIN PHOSPHATE IN DEXTROSE 5% IN PLASTIC CONTAINER, CLINDAMYCIN PHOSPHATE
 CLEOCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
 CLEOCIN T, CLINDAMYCIN PHOSPHATE
 CLEOCIN, CLINDAMYCIN PALMITATE HYDROCHLORIDE
 CLEOCIN, CLINDAMYCIN PHOSPHATE
 COLESTID, COLESTIPOL HYDROCHLORIDE
 CORTEF, HYDROCORTISONE
 CORTISONE ACETATE, CORTISONE ACETATE
 CORVERT, IBUTILIDE FUMARATE
 CYKLOKAPRON, TRANEXAMIC ACID
 DEPO-ESTRADIOL, ESTRADIOL CYPIONATE
 DEPO-MEDROL, METHYLPREDNISOLONE ACETATE
 DEPO-PROVERA, MEDROXYPROGESTERONE ACETATE
 DEPO-TESTOSTERONE, TESTOSTERONE CYPIONATE
 DETROL LA, TOLTERODINE TARTRATE
 DETROL, TOLTERODINE TARTRATE
 DIDREX, BENZPHETAMINE HYDROCHLORIDE
 DOSTINEX, CABERGOLINE
 EMCYT, ESTRAMUSTINE PHOSPHATE SODIUM
 ESTRING, ESTRADIOL
 FLAVORED COLESTID, COLESTIPOL HYDROCHLORIDE
 FLORONE E, DIFLORASONE DIACETATE
 FLORONE, DIFLORASONE DIACETATE
 FRAGMIN, DALTEPARIN SODIUM
 GENOTROPIN PRESERVATIVE FREE, SOMATROPIN RECOMBINANT
 GENOTROPIN, SOMATROPIN RECOMBINANT
 GLYNASE, GLYBURIDE
 GLYSET, MIGLITOL
 HALCION, TRIAZOLAM
 HALOTESTIN, FLUOXYMESTERONE
 HEMABATE, CARBOPROST TROMETHAMINE
 HEPARIN SODIUM, HEPARIN SODIUM
 IDAMYCIN PFS, IDARUBICIN HYDROCHLORIDE
 LINCOCIN, LINCOMYCIN HYDROCHLORIDE
 LONITEN, MINOXIDIL
 MEDROL, METHYLPREDNISOLONE
 MICRONASE, GLYBURIDE
 MYCOBUTIN, RIFABUTIN
 NICOTROL, NICOTINE
 OGEN .625, ESTROPIPATE
 OGEN 1.25, ESTROPIPATE
 OGEN 2.5, ESTROPIPATE
 OGEN 5, ESTROPIPATE
 OGEN, ESTROPIPATE
 PREPIDIL, DINOPROSTONE
 PROSTIN E2, DINOPROSTONE
 PROSTIN VR PEDIATRIC, ALPROSTADIL
 PROVERA, MEDROXYPROGESTERONE ACETATE
 PSORCON, DIFLORASONE DIACETATE
 R-GENE 10, ARGININE HYDROCHLORIDE
 ROGAINE (FOR MEN), MINOXIDIL (OTC)
 ROGAINE (FOR WOMEN), MINOXIDIL (OTC)
 ROGAINE EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 SOLU-CORTEF, HYDROCORTISONE SODIUM SUCCINATE
 SOLU-MEDROL, METHYLPREDNISOLONE SODIUM SUCCINATE
 SOMAVERT, PEGVISOMANT

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** P ****

* PHARMACIA AND UPJOHN CO
TOLINASE, TOLAZAMIDE
TROBICIN, SPECTINOMYCIN HYDROCHLORIDE
VANTIN, CEFPODOXIME PROXETIL
XALATAN, LATANOPROST
XANAX, ALPRAZOLAM
ZINECARD, DEXRAZOXANE HYDROCHLORIDE
ZYVOX, LINEZOLID

PHARMACIA UPJOHN

* PHARMACIA AND UPJOHN CONSUMER HEALTHCARE
NASALCROM, CROMOLYN SODIUM (OTC)

PHARMADERM

* PHARMADERM DIV ALTANA INC
BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE, BACITRACIN

PHARMAFORCE

* PHARMAFORCE INC
CYCLOSPORINE, CYCLOSPORINE
ESMOLOL HYDROCHLORIDE, ESMOLOL HYDROCHLORIDE
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
FOSCARNET SODIUM, FOSCARNET SODIUM
LIOETHYRONINE SODIUM, LIOETHYRONINE SODIUM
NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN, GRAMICIDIN
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE, HYDROCORTISONE
NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
OFLOXACIN, OFLOXACIN
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE

PHARMASEAL

* PHARMASEAL DIV BAXTER HEALTHCARE CORP
PHARMASEAL SCRUB CARE, CHLORHEXIDINE GLUCONATE (OTC)

PHARMATEK

* PHARMATEK INC
NEO-FRADIN, NEOMYCIN SULFATE

PHARMATON

* PHARMATON LTD
LIDOCATON, EPINEPHRINE

PHARMAX

* PHARMAX GROUP INC
FOLIC ACID, FOLIC ACID

PHARMERAL

* PHARMERAL INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN

PHARMEX PRODS

* PHARMEX PRODUCTS INC
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE

PHARMION

* PHARMION CORP
INNOHEP, TINZAPARIN SODIUM
VIDAZA, AZACITIDINE

PHOTOCURE ASA

* PHOTOCURE ASA
METVIXIA, METHYL AMINOLEVULINATE HYDROCHLORIDE

PIERRE FABRE

* PIERRE FABRE MEDICAMENT
NAVELBINE, VINORELBINE TARTRATE

PLANTEX

* PLANTEX USA INC DIV IKAPHARM INC
SULFAMETHOXAZOLE AND TRIMETHOPRIM SINGLE STRENGTH, SULFAMETHOXAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** P ****

PLIVA

* PLIVA INC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALBUTEROL, ALBUTEROL
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
AZITHROMYCIN, AZITHROMYCIN
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CARBAMAZEPINE, CARBAMAZEPINE
CARBOPLATIN, CARBOPLATIN
CHLORPROPAMIDE, CHLORPROPAMIDE
CHLORTHALIDONE, CHLORTHALIDONE
CIMETIDINE, CIMETIDINE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
CYCLOSPORINE, CYCLOSPORINE
CYPROHEPTADINE HYDROCHLORIDE, CYPROHEPTADINE HYDROCHLORIDE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DEXCHLORPHENIRAMINE MALEATE, DEXCHLORPHENIRAMINE MALEATE
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
FLUCONAZOLE, FLUCONAZOLE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
FLURBIPROFEN, FLURBIPROFEN
GLIPIZIDE, GLIPIZIDE
HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN
INDOMETHACIN, INDOMETHACIN
KETOCONAZOLE, KETOCONAZOLE
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
METHYLDOPA, METHYLDOPA
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METRONIDAZOLE, METRONIDAZOLE
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN, NAPROXEN
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
PENTOXIFYLLINE, PENTOXIFYLLINE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
THEOPHYLLINE, THEOPHYLLINE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

PLIVA PHARM IND

* PLIVA PHARMACEUTICAL INDUSTRY INC
TORSEMIDE, TORSEMIDE

POHL BOSKAMP

* G POHL BOSKAMP GMBH AND CO
NITROLINGUAL PUMPSPRAY, NITROGLYCERIN

POLYMEDICA

* POLYMEDICA INDUSTRIES INC
ANESTACON, LIDOCAINE HYDROCHLORIDE
NEOPAP, ACETAMINOPHEN (OTC)
PROMETHACON, PROMETHAZINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** P **

POPULATION COUNCIL

- * POPULATION COUNCIL CENTER FOR BIOMEDICAL RESEARCH
NORPLANT II, LEVONORGESTREL

PRAECIS

- * PRAECIS PHARMACEUTICALS INC
PLENAXIS, ABARELIX

PRESUTTI LABS

- * PRESUTTI LABORATORIES
TINDAMAX, TINIDAZOLE

PRIMAPHARM

- * PRIMAPHARM INC
HYDASE, HYALURONIDASE

PROCTER AND GAMBLE

- * PROCTER AND GAMBLE CO
DANTRUM, DANTROLENE SODIUM
MACRODANTIN, NITROFURANTOIN, MACROCRYSTALLINE
- * PROCTER AND GAMBLE PHARMACEUTICALS INC
ACTONEL WITH CALCIUM (COPACKAGED), CALCIUM CARBONATE
- * PROCTER AND GAMBLE PHARMACEUTICALS INC SUB PROCTER AND GAMBLE CO
ACTONEL, RISEDRONATE SODIUM
ASACOL, MESALAMINE
DANTRUM, DANTROLENE SODIUM
DIDRONEL, ETIDRONATE DISODIUM
MACROBID, NITROFURANTOIN

PROMETHEUS LABS

- * PROMETHEUS LABORATORIES INC
HELIDAC, BISMUTH SUBSALICYLATE
IMURAN, AZATHIOPRINE
MERCAPTOPURINE, MERCAPTOPURINE
RIDAURA, AURANOFIN
TRANDATE, LABETALOL HYDROCHLORIDE
ZYLOPRIM, ALLOPURINOL

PURDUE FREDERICK

- * PURDUE FREDERICK CO
MS CONTIN, MORPHINE SULFATE
T-PHYL, THEOPHYLLINE
UNIPHYL, THEOPHYLLINE

PURDUE PHARMA LP

- * PURDUE PHARMA LP
OXYCONTIN, OXYCODONE HYDROCHLORIDE
SPECTRACEF, CEFEDITOREN PIVOXIL

PUREPAC PHARM

- * PUREPAC PHARMACEUTICAL CO
ACYCLOVIR, ACYCLOVIR
ALPRAZOLAM, ALPRAZOLAM
AMPICILLIN TRIHYDRATE, AMPICILLIN/AMPICILLIN TRIHYDRATE
ASPIRIN AND CAFFEINE W/ BUTALBITAL, ASPIRIN
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
CARBAMAZEPINE, CARBAMAZEPINE
CARBIDOPA AND LEVODOPA, CARBIDOPA
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
DIAZEPAM, DIAZEPAM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DIPYRIDAMOLE, DIPYRIDAMOLE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
ETODOLAC, ETODOLAC
FAMOTIDINE, FAMOTIDINE
FENOPROFEN CALCIUM, FENOPROFEN CALCIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** P ****

* PUREPAC PHARMACEUTICAL CO
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
GABAPENTIN, GABAPENTIN
GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
INDAPAMIDE, INDAPAMIDE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
LORAZEPAM, LORAZEPAM
LOVASTATIN, LOVASTATIN
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NAPROXEN, NAPROXEN
NIFEDIPINE, NIFEDIPINE
OXAPROZIN, OXAPROZIN
OXAZEPAM, OXAZEPAM
PENTOXIFYLLINE, PENTOXIFYLLINE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PROPYLTHIOURACIL, PROPYLTHIOURACIL
SPIRONOLACTONE, SPIRONOLACTONE
TEMAZEPAM, TEMAZEPAM
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOLMETIN SODIUM, TOLMETIN SODIUM
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE

PVT FORM

* PRIVATE FORMULATIONS INC
IBUPROFEN, IBUPROFEN (OTC)
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
PROFEN, IBUPROFEN (OTC)

QLT

* QLT INC
VISUDYNE, VERTEPORFIN

QLT USA

* QLT USA INC
ATRIDOX, DOXYCYCLINE HYCLATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
ELIGARD, LEUPROLIDE ACETATE
ERYTHROMYCIN AND BENZOYL PEROXIDE, BENZOYL PEROXIDE
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
KETOCONAZOLE, KETOCONAZOLE
LIDOCAINE AND PRILOCAINE, LIDOCAINE
MOMETASONE FUROATE, MOMETASONE FUROATE

QOL MEDCL

* QOL MEDICAL LLC
ELLIOTTS B SOLUTION, CALCIUM CHLORIDE
ETHAMOLIN, ETHANOLAMINE OLEATE
GLOFIL-125, IOTHALAMATE SODIUM, I-125
NASCOBAL, CYANOCOBALAMIN
SUCRAID, SACROSIDASE

QUESTCOR PHARMS

* QUESTCOR PHARMACEUTICALS INC
H.P. ACTHAR GEL, CORTICOTROPIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** R ****

R R REGISTRATIONS

* R AND R REGISTRATIONS
THYROSAFE, POTASSIUM IODIDE (OTC)

RANBAXY

* RANBAXY LABORATORIES LTD
ACYCLOVIR, ACYCLOVIR
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
CARBILEV, CARBIDOPA
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CEFPODOXIME PROXETIL, CEFPODOXIME PROXETIL
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEPHALEXIN, CEPHALEXIN
CLARITHROMYCIN, CLARITHROMYCIN
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
DISPERMOX, AMOXICILLIN
DOXYCYCLINE, DOXYCYCLINE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ETODOLAC, ETODOLAC
FENOFIBRATE, FENOFIBRATE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLUCONAZOLE, FLUCONAZOLE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
GABAPENTIN, GABAPENTIN
GANCICLOVIR, GANCICLOVIR
GLIMEPIRIDE, GLIMEPIRIDE
LORATADINE AND PSEUDOEPHEDRINE SULFATE, LORATADINE (OTC)
LORATADINE, LORATADINE (OTC)
LORAZEPAM, LORAZEPAM
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
OFLOXACIN, OFLOXACIN
PANIXINE DISPERDOSE, CEPHALEXIN
PSEUDOEPHEDRINE HYDROCHLORIDE, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
QUINAPRIL, QUINAPRIL HYDROCHLORIDE
RANICLOR, CEFACLOL
RANITIDINE, RANITIDINE HYDROCHLORIDE
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
SOTRET, ISOTRETINOIN
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
ZIDOVUDINE, ZIDOVUDINE

* RANBAXY PHARMACEUTICALS INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAMINOPHEN W/ CODEINE PHOSPHATE #3, ACETAMINOPHEN
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
AVENTYL HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
CEFACLOL, CEFACLOL
CEFADROXIL, CEFADROXIL/CEFADROXIL HEMIHYDRATE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINOPRIL, LISINOPRIL
PENTAZOCINE AND NALOXONE HYDROCHLORIDES, NALOXONE HYDROCHLORIDE
RIOMET, METFORMIN HYDROCHLORIDE
SECONAL SODIUM, SECOBARBITAL SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** R **

* RANBAXY PHARMACEUTICALS INC
SOTRET, ISOTRETINOIN

RATIOPHARM

* RATIOPHARM GMBH AND CO ARZNEIMITTEL
SUCRALFATE, SUCRALFATE

RECKITT BENCKISER

* RECKITT BENCKISER PHARMACEUTICALS INC
BUPRENEX, BUPRENORPHINE HYDROCHLORIDE
SUBOXONE, BUPRENORPHINE HYDROCHLORIDE
SUBUTEX, BUPRENORPHINE HYDROCHLORIDE

REGENT

* REGENT MEDICAL AMERICAS LLC
HIBICLENS, CHLORHEXIDINE GLUCONATE (OTC)
HIBISTAT, CHLORHEXIDINE GLUCONATE (OTC)

RELIANT PHARMS

* RELIANT PHARMACEUTICALS LLC
AXID, NIZATIDINE
DYNACIRC CR, ISRADIPINE
DYNACIRC, ISRADIPINE
INNOPRAN XL, PROPRANOLOL HYDROCHLORIDE
RYTHMOL SR, PROPAFENONE HYDROCHLORIDE
RYTHMOL, PROPAFENONE HYDROCHLORIDE

RELIANT PHARMS INC

* RELIANT PHARMACEUTICALS INC
ANTARA (MICRONIZED), FENOFIBRATE
OMACOR, OMEGA-3-ACID ETHYL ESTERS

RESPIRARE

* RESPIRARE LLC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
CROMOLYN SODIUM, CROMOLYN SODIUM
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

RHODIA

* RHODIA LTD
ISOFLURANE, ISOFLURANE

ROCHE

* HOFFMANN LA ROCHE INC
BONIVA, IBANDRONATE SODIUM
BUMEX, BUMETANIDE
COPEGUS, RIBAVIRIN
DEMADEX, TORSEMIDE
FANSIDAR, PYRIMETHAMINE
FUZEON, ENFUVIRTIDE
GANTRISIN PEDIATRIC, SULFISOXAZOLE ACETYL
HIVID, ZALCITABINE
INVIRASE, SAQUINAVIR MESYLATE
KLONOPIN RAPIDLY DISINTEGRATING, CLONAZEPAM
KLONOPIN, CLONAZEPAM
KYTRIL, GRANISETRON HYDROCHLORIDE
LARIAM, MEFLOROQUINE HYDROCHLORIDE
ROCALTRON, CALCITRIOL
TAMIFLU, OSELTAMIVIR PHOSPHATE
VALIUM, DIAZEPAM
VESANOID, TRETINOIN

ROCHE PALO

* ROCHE PALO ALTO LLC
AEROBID, FLUNISOLIDE
ANAPROX DS, NAPROXEN SODIUM
ANAPROX, NAPROXEN SODIUM
CARDENE SR, NICARDIPINE HYDROCHLORIDE
CARDENE, NICARDIPINE HYDROCHLORIDE
CELLCEPT, MYCOPHENOLATE MOFETIL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** R ****

* ROCHE PALO ALTO LLC
CELLCEPT, MYCOPHENOLATE MOFETIL HYDROCHLORIDE
CYTOVENE IV, GANCICLOVIR SODIUM
CYTOVENE, GANCICLOVIR
EC-NAPROSYN, NAPROXEN
NAPROSYN, NAPROXEN
TICLID, TICLOPIDINE HYDROCHLORIDE
TORADOL, KETOROLAC TROMETHAMINE
VALCYTE, VALGANCICLOVIR HYDROCHLORIDE

ROERIG

* ROERIG DIV PFIZER INC
ATARAX, HYDROXYZINE HYDROCHLORIDE

ROMARK

* ROMARK LABORATORIES
ALINIA, NITAZOXANIDE

ROSS LABS

* ROSS LABORATORIES DIV ABBOTT LABORATORIES INC
PEDIAMYCIN 400, ERYTHROMYCIN ETHYLSUCCINATE
PEDIAMYCIN, ERYTHROMYCIN ETHYLSUCCINATE
PEDIAZOLE, ERYTHROMYCIN ETHYLSUCCINATE
SURVANTA, BERACTANT

ROXANE

* ROXANE LABORATORIES INC
ACETAMINOPHEN W/ CODEINE NO. 3, ACETAMINOPHEN
ACETAMINOPHEN W/ CODEINE, ACETAMINOPHEN
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALPRAZOLAM, ALPRAZOLAM
AMINOPHYLLINE, AMINOPHYLLINE
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
AZATHIOPRINE, AZATHIOPRINE
BUTORPHANOL TARTRATE, BUTORPHANOL TARTRATE
CALCITRIOL, CALCITRIOL
CHLORPROMAZINE HYDROCHLORIDE INTENSOL, CHLORPROMAZINE HYDROCHLORIDE
CILOSTAZOL, CILOSTAZOL
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLARITHROMYCIN, CLARITHROMYCIN
CLOTRIMAZOLE, CLOTRIMAZOLE
CROMOLYN SODIUM, CROMOLYN SODIUM
CYCLOPHOSPHAMIDE, CYCLOPHOSPHAMIDE
DEXAMETHASONE INTENSOL, DEXAMETHASONE
DEXAMETHASONE, DEXAMETHASONE
DIAZEPAM INTENSOL, DIAZEPAM
DIAZEPAM, DIAZEPAM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DIGOXIN, DIGOXIN
DIPHENOXYLATE HYDROCHLORIDE AND ATROPINE SULFATE, ATROPINE SULFATE
DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
FLECAINIDE ACETATE, FLECAINIDE ACETATE
FLUCONAZOLE, FLUCONAZOLE
FUROSEMIDE, FUROSEMIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
HYDROXYUREA, HYDROXYUREA
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE
LACTULOSE, LACTULOSE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
LEVORPHANOL TARTRATE, LEVORPHANOL TARTRATE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
LIDOCAINE VISCOUS, LIDOCAINE HYDROCHLORIDE
LITHIUM CARBONATE, LITHIUM CARBONATE
LITHIUM CITRATE, LITHIUM CITRATE
LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** R ****

* ROXANE LABORATORIES INC
LORAZEPAM INTENSOL, LORAZEPAM
LORAZEPAM, LORAZEPAM
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MEGESTROL ACETATE, MEGESTROL ACETATE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MEPROBAMATE, MEPROBAMATE
MERCAPTOPYRINE, MERCAPTOPYRINE
METHADONE HYDROCHLORIDE INTENSOL, METHADONE HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHOTREXATE SODIUM, METHOTREXATE SODIUM
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METOLAZONE, METOLAZONE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NAPROXEN, NAPROXEN
PILOCARPINE HYDROCHLORIDE, PILOCARPINE HYDROCHLORIDE
PREDNISONE INTENSOL, PREDNISONE
PREDNISONE, PREDNISONE
PROPANTHELINE BROMIDE, PROPANTHELINE BROMIDE
PROPRANOLOL HYDROCHLORIDE INTENSOL, PROPRANOLOL HYDROCHLORIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
ROXICET 5/500, ACETAMINOPHEN
ROXICET, ACETAMINOPHEN
ROXILOX, ACETAMINOPHEN
SODIUM POLYSTYRENE SULFONATE, SODIUM POLYSTYRENE SULFONATE
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
THEOPHYLLINE, THEOPHYLLINE
THIORIDAZINE HYDROCHLORIDE INTENSOL, THIORIDAZINE HYDROCHLORIDE
TORSEMIDE, TORSEMIDE
TRIAZOLAM, TRIAZOLAM
ZIDOVUDINE, ZIDOVUDINE
ZONISAMIDE, ZONISAMIDE

RXELITE

* RXELITE
FLUOXETINE, FLUOXETINE HYDROCHLORIDE
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

SABEX 2002

* SABEX 2002 INC
ANECTINE, SUCCINYLCHOLINE CHLORIDE
ARISTOSPAN, TRIAMCINOLONE HEXACETONIDE
DIGOXIN, DIGOXIN
FENOLDOPAM MESYLATE, FENOLDOPAM MESYLATE
FLUMAZENIL, FLUMAZENIL
HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
HALOPERIDOL, HALOPERIDOL LACTATE
INFUVITE ADULT, ALPHA-TOCOPHEROL ACETATE
INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE), ASCORBIC ACID
INFUVITE PEDIATRIC, ASCORBIC ACID
ISONIAZID, ISONIAZID
KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
REGONOL, PYRIDOSTIGMINE BROMIDE

SAGE PRODS

* SAGE PRODUCTS INC
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE (OTC)

SALIX PHARMS

* SALIX PHARMACEUTICALS INC
ANUSOL HC, HYDROCORTISONE
COLAZAL, BALSALAZIDE DISODIUM
VISICOL, SODIUM PHOSPHATE, DIBASIC, ANHYDROUS
XIFAXAN, RIFAXIMIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

SANDOZ

* SANDOZ INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALLOPURINOL, ALLOPURINOL
ALPRAZOLAM, ALPRAZOLAM
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMOXAPINE, AMOXAPINE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
AMPICILLIN SODIUM, AMPICILLIN SODIUM
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
ATENOLOL, ATENOLOL
AZITHROMYCIN, AZITHROMYCIN
BENAZEPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, BENAZEPRIL HYDROCHLORIDE
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
BUMETANIDE, BUMETANIDE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
BUTAL COMPOUND, ASPIRIN
CAFERGOT, CAFFEINE
CAPTOPRIL, CAPTOPRIL
CARBIDOPA AND LEVODOPA, CARBIDOPA
CARBOPLATIN, CARBOPLATIN
CARISOPRODOL AND ASPIRIN, ASPIRIN
CARISOPRODOL, CARISOPRODOL
CARISOPRODOL, ASPIRIN AND CODEINE PHOSPHATE, ASPIRIN
CEFPROZIL, CEFPROZIL
CEFTRIAZONE, CEFTRIAZONE SODIUM
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CHLORPROPAMIDE, CHLORPROPAMIDE
CHLORZOXAZONE, CHLORZOXAZONE
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
CHOLESTYRAMINE, CHOLESTYRAMINE
CILOSTAZOL, CILOSTAZOL
CIMETIDINE, CIMETIDINE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
CLARITHROMYCIN EXTENDED RELEASE, CLARITHROMYCIN
CLARITHROMYCIN, CLARITHROMYCIN
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
CORPHED, PSEUDOEPHEDRINE HYDROCHLORIDE
CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
CYCLOSPORINE, CYCLOSPORINE
DESIPRAMINE HYDROCHLORIDE, DESIPRAMINE HYDROCHLORIDE
DEXTRAMP SACCARATE, AMP ASPARTATE, DEXTRAMP SULFATE AND AMP SULFATE, AMPHETAMINE
ASPARTATE
DIAZEPAM, DIAZEPAM
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
DIFLUNISAL, DIFLUNISAL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** S **

* SANDOZ INC
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE, ATROPINE SULFATE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ETODOLAC, ETODOLAC
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FLECAINIDE ACETATE, FLECAINIDE ACETATE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUOXETINE, FLUOXETINE HYDROCHLORIDE
 FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FLUTAMIDE, FLUTAMIDE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GABAPENTIN, GABAPENTIN
 GEMFIBROZIL, GEMFIBROZIL
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GUANABENZ ACETATE, GUANABENZ ACETATE
 HALOPERIDOL, HALOPERIDOL
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 IMIPRAMINE HYDROCHLORIDE, IMIPRAMINE HYDROCHLORIDE
 INDAPAMIDE, INDAPAMIDE
 INDOCIN SR, INDOMETHACIN
 INDOMETHACIN, INDOMETHACIN
 INVAGESIC FORTE, ASPIRIN
 INVAGESIC, ASPIRIN
 ISONIAZID, ISONIAZID
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 ITRACONAZOLE, ITRACONAZOLE
 KETOPROFEN, KETOPROFEN
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEFLUNOMIDE, LEFLUNOMIDE
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINAPRIL, LISINAPRIL
 LOCHOLEST LIGHT, CHOLESTYRAMINE
 LOCHOLEST, CHOLESTYRAMINE
 LONOX, ATROPINE SULFATE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LORATADINE, LORATADINE (OTC)
 LORAZEPAM, LORAZEPAM
 LOVASTATIN, LOVASTATIN
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

* SANDOZ INC
MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
MEFLOQUINE HYDROCHLORIDE, MEFLOQUINE HYDROCHLORIDE
MEPROBAMATE, MEPROBAMATE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
METHAZOLAMIDE, METHAZOLAMIDE
METHIMAZOLE, METHIMAZOLE
METHOCARBAMOL, METHOCARBAMOL
METHYCLOTHIAZIDE, METHYCLOTHIAZIDE
METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
METHYLDOPA, METHYLDOPA
METHYLPREDNISOLONE, METHYLPREDNISOLONE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
METOLAZONE, METOLAZONE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
METRONIDAZOLE, METRONIDAZOLE
MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
MIDODRINE HYDROCHLORIDE, MIDODRINE HYDROCHLORIDE
MIRTAZAPINE, MIRTAZAPINE
NABUMETONE, NABUMETONE
NADOLOL, NADOLOL
NAFCILLIN SODIUM, NAFCILLIN SODIUM
NALTREXONE HYDROCHLORIDE, NALTREXONE HYDROCHLORIDE
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN SODIUM, NAPROXEN SODIUM (OTC)
NAPROXEN, NAPROXEN
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
NITROFURANTOIN (MONOHYDRATE/MACROCRYSTALS), NITROFURANTOIN
NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
NIZATIDINE, NIZATIDINE
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
NYDRAZID, ISONIAZID
OMEPRazole, OMEPRazole
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN
OXACILLIN SODIUM, OXACILLIN SODIUM
OXAPROZIN, OXAPROZIN
OXAZEPAM, OXAZEPAM
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PENICILLIN G POTASSIUM, PENICILLIN G POTASSIUM
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
PERPHENAZINE, PERPHENAZINE
PHENDIMETRAZINE TARTRATE, PHENDIMETRAZINE TARTRATE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PINDOLOL, PINDOLOL
PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
PREDNISONE, PREDNISONE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
PSEUDOEPHEDRINE HYDROCHLORIDE AND TRIPROLIDINE HYDROCHLORIDE, PSEUDOEPHEDRINE
HYDROCHLORIDE
QUINAPRIL HYDROCHLORIDE, QUINAPRIL HYDROCHLORIDE
QUINIDINE SULFATE, QUINIDINE SULFATE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
RESERPINE, RESERPINE
RIBAVIRIN, RIBAVIRIN
RIFAMPIN, RIFAMPIN
SONAZINE, CHLORPROMAZINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

* SANDOZ INC
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
SPIRONOLACTONE, SPIRONOLACTONE
SULFADIAZINE, SULFADIAZINE
SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SULINDAC, SULINDAC
TEMAZEPAM, TEMAZEPAM
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
THIOTHIXENE, THIOTHIXENE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIMOLOL MALEATE, TIMOLOL MALEATE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TOLMETIN SODIUM, TOLMETIN SODIUM
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRIFLUOPERAZINE HYDROCHLORIDE, TRIFLUOPERAZINE HYDROCHLORIDE
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
WARFARIN SODIUM, WARFARIN SODIUM
ZONISAMIDE, ZONISAMIDE

SANKYO

* SANKYO PHARMA INC
BENICAR HCT, HYDROCHLOROTHIAZIDE
BENICAR, OLMESARTAN MEDOXOMIL
WELCHOL, COLESEVELAM HYDROCHLORIDE

SANO

* SANO CORP
NICOTINE, NICOTINE

SANOFI SYN RES

* SANOFI SYNTHELABO RESEARCH DIV SANOFI SYNTHELABO INC
UROXATRAL, ALFUZOSIN HYDROCHLORIDE

SANOFI SYNTHELABO

* SANOFI SYNTHELABO INC
AMBIEN CR, ZOLPIDEM TARTRATE
AMBIEN, ZOLPIDEM TARTRATE
ARALEN, CHLOROQUINE PHOSPHATE
AVALIDE, HYDROCHLOROTHIAZIDE
AVAPRO, IRBESARTAN
DEMEROL, MEPERIDINE HYDROCHLORIDE
DRISDOL, ERGOCALCIFEROL
ELOXATIN, OXALIPLATIN
KAYEXALATE, SODIUM POLYSTYRENE SULFONATE
MYTELASE, AMBENONIUM CHLORIDE
NEGGRAM, NALIDIXIC ACID
PHISOHEX, HEXACHLOROPHENE
PLAQUENIL, HYDROXYCHLOROQUINE SULFATE
PLAVIX, CLOPIDOGREL BISULFATE
PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER, MILRINONE LACTATE
PRIMACOR, MILRINONE LACTATE
PRIMAQUINE, PRIMAQUINE PHOSPHATE
SKELID, TILUDRONATE DISODIUM
TALACEN, ACETAMINOPHEN
TALWIN NX, NALOXONE HYDROCHLORIDE

SANTARUS

* SANTARUS INC
ZEGERID, OMEPRAZOLE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

SANTEN

* SANTEN INC
ALAMAST, PEMIROLAST POTASSIUM
IQUIX, LEVOFLOXACIN
QUIXIN, LEVOFLOXACIN

SANTEN OY

* SANTEN OY
BETIMOL, TIMOLOL

SAVAGE LABS

* SAVAGE LABORATORIES INC DIV ALTANA INC
ALPHATREX, BETAMETHASONE DIPROPIONATE
DILOR, DYPHYLLINE
DILOR-400, DYPHYLLINE
ETHIODOL, ETHIODIZED OIL
KAON CL-10, POTASSIUM CHLORIDE
PANDEL, HYDROCORTISONE PROBUTATE

SAVIENT PHARMS

* SAVIENT PHARMACEUTICALS INC
DELATESTRYL, TESTOSTERONE ENANTHATE
OXANDRIN, OXANDROLONE
SOLTAMOX, TAMOXIFEN CITRATE

SB PHARMCO

* SB PHARMCO PUERTO RICO INC
AVANDAMET, METFORMIN HYDROCHLORIDE
AVANDARYL, GLIMEPIRIDE
AVANDIA, ROSIGLITAZONE MALEATE

SCHERER LABS

* SCHERER LABORATORIES INC
MEPROBAMATE, MEPROBAMATE

SCHERER RP

* RP SCHERER CORP
NIFEDIPINE, NIFEDIPINE
* RP SCHERER NORTH AMERICA
NIFEDIPINE, NIFEDIPINE
VALPROIC ACID, VALPROIC ACID
* RP SCHERER NORTH AMERICA DIV RP SCHERER CORP
DIPHENOXYLATE HYDROCHLORIDE W/ ATROPINE SULFATE, ATROPINE SULFATE

SCHERING

* SCHERING
ASMANEX TWISTHALER, MOMETASONE FUROATE
CLARINEX, DESLORATADINE
* SCHERING CORP
CLARINEX D 24 HOUR, DESLORATADINE
CLARINEX, DESLORATADINE
DIPROLENE AF, BETAMETHASONE DIPROPIONATE
GUANIDINE HYDROCHLORIDE, GUANIDINE HYDROCHLORIDE
INTEGRILIN, EPTIFIBATIDE
LOTRISONE, BETAMETHASONE DIPROPIONATE
REBETOL, RIBAVIRIN
TEMODAR, TEMOZOLOMIDE
* SCHERING CORP SUB SCHERING PLOUGH CORP
CELESTONE SOLUSPAN, BETAMETHASONE ACETATE
CELESTONE, BETAMETHASONE
DIPROLENE, BETAMETHASONE DIPROPIONATE
ELOCON, MOMETASONE FUROATE
EULEXIN, FLUTAMIDE
FULVICIN-U/F, GRISEOFULVIN, MICROCRYSTALLINE
HYPERSTAT, DIAZOXIDE
MIRADON, ANISINDIONE
PROVENTIL, ALBUTEROL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

SCHERING PLOUGH

- * SCHERING PLOUGH CORP
CLARINEX, DESLORATADINE
IMDUR, ISOSORBIDE MONONITRATE
- * SCHERING PLOUGH HEALTHCARE PRODUCTS INC
AFRINOL, PSEUDOEPHEDRINE SULFATE (OTC)
CHLOR-TRIMETON, CHLORPHENIRAMINE MALEATE (OTC)
CLARITIN HIVES RELIEF REDITAB, LORATADINE (OTC)
CLARITIN HIVES RELIEF, LORATADINE (OTC)
CLARITIN REDITABS, LORATADINE (OTC)
CLARITIN, LORATADINE (OTC)
CLARITIN-D 24 HOUR, LORATADINE (OTC)
DISOPHROL, DEXBROMPHENIRAMINE MALEATE (OTC)
DRIXORAL PLUS, ACETAMINOPHEN (OTC)
DRIXORAL, DEXBROMPHENIRAMINE MALEATE (OTC)
GYNE-LOTRIMIN 3 COMBINATION PACK, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN 3, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN COMBINATION PACK, CLOTRIMAZOLE (OTC)
GYNE-LOTRIMIN, CLOTRIMAZOLE (OTC)
LOTRIMIN ULTRA, BUTENAFINE HYDROCHLORIDE (OTC)
NASONEX, MOMETASONE FUROATE MONOHYDRATE
OCUCLEAR, OXYMETAZOLINE HYDROCHLORIDE (OTC)
SHADE UVAGUARD, AVOBENZONE (OTC)

SCHERING PLOUGH RES

- * SCHERING PLOUGH RESEARCH INSTITUTE
LOTRISONE, BETAMETHASONE DIPROPIONATE
REBETOL, RIBAVIRIN

SCHWARZ

- * SCHWARZ GMBH
MONOKET, ISOSORBIDE MONONITRATE

SCHWARZ PHARMA

- * SCHWARZ PHARMA AG
EDEX, ALPROSTADIL
GLYCOLAX, POLYETHYLENE GLYCOL 3350
- * SCHWARZ PHARMA INC
AZDONE, ASPIRIN
CO-GESIC, ACETAMINOPHEN
COLYTE WITH FLAVOR PACKS, POLYETHYLENE GLYCOL 3350
COLYTE, POLYETHYLENE GLYCOL 3350
COLYTE-FLAVORED, POLYETHYLENE GLYCOL 3350
CORTIFOAM, HYDROCORTISONE ACETATE
DILATRATE-SR, ISOSORBIDE DINITRATE
EPIFOAM, HYDROCORTISONE ACETATE
FLUXID, FAMOTIDINE
KEMSTRO, BACLOFEN
LEVATOL, PENBUTOLOL SULFATE
NIRAVAM, ALPRAZOLAM
PARCOPA, CARBIDOPA
PROCTOFOAM HC, HYDROCORTISONE ACETATE
REGLAN ODT, METOCLOPRAMIDE HYDROCHLORIDE
REGLAN, METOCLOPRAMIDE HYDROCHLORIDE
ROBAXIN, METHOCARBAMOL
ROBAXIN-750, METHOCARBAMOL
TRILYTE, POLYETHYLENE GLYCOL 3350
UNIRETIC, HYDROCHLOROTHIAZIDE
UNIVASC, MOEXIPRIL HYDROCHLORIDE

SCIOS

- * SCIOS INC
NATRECOR, NESIRITIDE RECOMBINANT

SCS

- * SCS PHARMACEUTICALS
ATENOLOL, ATENOLOL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

SEPRACOR

- * SEPRACOR INC
XOPENEX, LEVALBUTEROL HYDROCHLORIDE
- * SEPRACOR PHARMACEUTICALS
LUNESTA, ESZOPICLONE
XOPENEX HFA, LEVALBUTEROL TARTRATE

SEPTODONT

- * SEPTODONT INC
OCTOCAINE, EPINEPHRINE

SERONO

- * SERONO LABORATORIES INC
GEREF, SERMORELIN ACETATE
SAIZEN, SOMATROPIN RECOMBINANT
SEROPHENE, CLOMIPHENE CITRATE
SEROSTIM, SOMATROPIN RECOMBINANT

SERONO INC

- * SERONO INC
CETROTIDE, CETRORELIX
GONAL-F RFF PEN, FOLLITROPIN ALFA/BETA
GONAL-F RFF, FOLLITROPIN ALFA/BETA
GONAL-F, FOLLITROPIN ALFA/BETA
LUVERIS, LUTROPIN ALFA
NOVANTRONE, MITOXANTRONE HYDROCHLORIDE
OVIDREL, CHORIOGONADOTROPIN ALFA
ZORBTIVE, SOMATROPIN RECOMBINANT

SHIONOGI

- * SHIONOGI USA INC
CEDAX, CEFTIBUTEN DIHYDRATE

SHIRE

- * SHIRE DEVELOPMENT INC
ADDERALL 10, AMPHETAMINE ASPARTATE
ADDERALL 12.5, AMPHETAMINE ASPARTATE
ADDERALL 15, AMPHETAMINE ASPARTATE
ADDERALL 20, AMPHETAMINE ASPARTATE
ADDERALL 30, AMPHETAMINE ASPARTATE
ADDERALL 5, AMPHETAMINE ASPARTATE
ADDERALL 7.5, AMPHETAMINE ASPARTATE
ADDERALL XR 10, AMPHETAMINE ASPARTATE
ADDERALL XR 15, AMPHETAMINE ASPARTATE
ADDERALL XR 20, AMPHETAMINE ASPARTATE
ADDERALL XR 25, AMPHETAMINE ASPARTATE
ADDERALL XR 30, AMPHETAMINE ASPARTATE
ADDERALL XR 5, AMPHETAMINE ASPARTATE
AGRYLIN, ANAGRELIDE HYDROCHLORIDE
CARBATROL, CARBAMAZEPINE
EQUETRO, CARBAMAZEPINE
ETHMOZINE, MORICIZINE HYDROCHLORIDE
FOSRENOL, LANTHANUM CARBONATE
PENTASA, MESALAMINE
PROAMATINE, MIDODRINE HYDROCHLORIDE
PRO-BANTHINE, PROPANTHELINE BROMIDE
SALURON, HYDROFLUMETHIAZIDE

SHIRE RICHWOOD

- * SHIRE RICHWOOD INC
DEXTROSTAT, DEXTROAMPHETAMINE SULFATE
X-TROZINE L.A., PHENDIMETRAZINE TARTRATE
X-TROZINE, PHENDIMETRAZINE TARTRATE

SICOR PHARMS

- * SICOR PHARMACEUTICALS INC
ACYCLOVIR SODIUM, ACYCLOVIR SODIUM
ACYCLOVIR, ACYCLOVIR SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** S **

* SICOR PHARMACEUTICALS INC
 ADENOSINE, ADENOSINE
 ADRUCIL, FLUOROURACIL
 ALPROSTADIL, ALPROSTADIL
 AMIKACIN SULFATE, AMIKACIN SULFATE
 AMINOPHYLLINE, AMINOPHYLLINE
 AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
 AMPHOTERICIN B, AMPHOTERICIN B
 ATRACURIUM BESYLATE, ATRACURIUM BESYLATE
 BLEOMYCIN SULFATE, BLEOMYCIN SULFATE
 BLEOMYCIN, BLEOMYCIN SULFATE
 BUMETANIDE, BUMETANIDE
 CALCITRIOL, CALCITRIOL
 CARBOPLATIN, CARBOPLATIN
 CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
 CISPLATIN, CISPLATIN
 CYTOSAR-U, CYTARABINE
 DACARBAZINE, DACARBAZINE
 DAUNORUBICIN HYDROCHLORIDE, DAUNORUBICIN HYDROCHLORIDE
 DESMOPRESSIN ACETATE, DESMOPRESSIN ACETATE
 DEXAMETHASONE SODIUM PHOSPHATE, DEXAMETHASONE SODIUM PHOSPHATE
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
 DOPAMINE HYDROCHLORIDE, DOPAMINE HYDROCHLORIDE
 DOXORUBICIN HYDROCHLORIDE, DOXORUBICIN HYDROCHLORIDE
 ENALAPRILAT, ENALAPRILAT
 ERYTHROMYCIN LACTOBIONATE, ERYTHROMYCIN LACTOBIONATE
 ETOPOSIDE, ETOPOSIDE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER, FLUCONAZOLE
 FLUCONAZOLE IN SODIUM CHLORIDE 0.9%, FLUCONAZOLE
 FLUDARABINE PHOSPHATE, FLUDARABINE PHOSPHATE
 FLUMAZENIL, FLUMAZENIL
 FLUOROURACIL, FLUOROURACIL
 FLUPHENAZINE DECANOATE, FLUPHENAZINE DECANOATE
 HALOPERIDOL DECANOATE, HALOPERIDOL DECANOATE
 HALOPERIDOL, HALOPERIDOL LACTATE
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE PFS, IDARUBICIN HYDROCHLORIDE
 IDARUBICIN HYDROCHLORIDE, IDARUBICIN HYDROCHLORIDE
 IFOSFAMIDE/MESNA KIT, IFOSFAMIDE
 LEUCOVORIN CALCIUM PRESERVATIVE FREE, LEUCOVORIN CALCIUM
 LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
 LEUPROLIDE ACETATE, LEUPROLIDE ACETATE
 LEVOCARNITINE, LEVOCARNITINE
 MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
 MESNA, MESNA
 METHYLDOPATE HYDROCHLORIDE, METHYLDOPATE HYDROCHLORIDE
 METHYLPREDNISOLONE ACETATE, METHYLPREDNISOLONE ACETATE
 METHYLPREDNISOLONE SODIUM SUCCINATE, METHYLPREDNISOLONE SODIUM SUCCINATE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 NEOSAR, CYCLOPHOSPHAMIDE
 NOREPINEPHRINE BITARTRATE, NOREPINEPHRINE BITARTRATE
 OCTREOTIDE ACETATE, OCTREOTIDE ACETATE
 PAMIDRONATE DISODIUM, PAMIDRONATE DISODIUM
 PANCURONIUM BROMIDE, PANCURONIUM BROMIDE
 PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
 PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
 PROPOFOL, PROPOFOL
 SODIUM NITROPRUSSIDE, SODIUM NITROPRUSSIDE
 SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
 TERBUTALINE SULFATE, TERBUTALINE SULFATE
 THIOTEPA, THIOTEPA
 TOBRAMYCIN SULFATE, TOBRAMYCIN SULFATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

* SICOR PHARMACEUTICALS INC
TOPOSAR, ETOPOSIDE
VECURONIUM BROMIDE, VECURONIUM BROMIDE
VINCRIStINE SULFATE PFS, VINCRIStINE SULFATE
VINORELBINE TARTRATE, VINORELBINE TARTRATE
ZANOSAR, STREPTOZOcIN

SIEGFRIED

* SIEGFRIED LTD
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

SIGHT PHARMS

* SIGHT PHARMACEUTICALS INC
MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
MINOXIDIL (FOR WOMEN), MINOXIDIL (OTC)

SIGMA TAU

* SIGMA TAU PHARMACEUTICALS INC
CARNITOR, LEVOCARNITINE
MATULANE, PROCARBAZINE HYDROCHLORIDE

SILARX

* SILARX PHARMACEUTICALS INC
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
CLEMAStINE FUMARATE, CLEMAStINE FUMARATE
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
HALOPERIDOL, HALOPERIDOL LACTATE
METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE

SIRIUS LABS

* SIRIUS LABORATORIES INC
TEXACORT, HYDROCORTISONE

SKINMEDICA

* SKINMEDICA INC
VANIQA, EFLORNITHINE HYDROCHLORIDE

SKYEPHARMA

* SKYEPHARMA INC
DEPOCYT, CYTARABINE
DEPODUR, MORPHINE SULFATE
TRIGLIDE, FENOFIBRATE

SMITHKLINE BEECHAM

* SMITHKLINE BEECHAM CORP DBA GLAXOSMITHKLINE
EPZICOM, ABACAVIR SULFATE
LANOXICAPS, DIGOXIN
LANOXIN, DIGOXIN
LEUKERAN, CHLORAMBUCIL
WELLBUTRIN XL, BUPROPION HYDROCHLORIDE

SOAPCO

* SOAPCO INC
BRIAN CARE, CHLORHEXIDINE GLUCONATE (OTC)

SOLVAY

* SOLVAY PHARMACEUTICALS
CORTENEMA, HYDROCORTISONE
ESTROGEL, ESTRADIOL
ROWASA, MESALAMINE
TRANMEP, MEPROBAMATE

SOLVAY PHARMS

* SOLVAY PHARMACEUTICALS INC
ACEON, PERINDOPRIL ERBUMINE

SOMERSET

* SOMERSET PHARMACEUTICALS INC
ELDEPRYL, SELEGILINE HYDROCHLORIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** S **

SPEAR PHARMS

* SPEAR PHARMACEUTICALS INC
TRETINOIN, TRETINOIN

SPECTRUM PHARMS

* SPECTRUM PHARMACEUTICALS INC
CARBOPLATIN, CARBOPLATIN

STAND HOMEOPATH

* STANDARD HOMEOPATHIC CO
IVY BLOCK, BENTOQUATAM (OTC)

STAR PHARMS FL

* STAR PHARMACEUTICALS INC
VIRILON, METHYLTESTOSTERONE

STASON

* STASON INDUSTRIAL CORP
ACYCLOVIR, ACYCLOVIR
CAPTOPRIL, CAPTOPRIL
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE

STAT TRADE

* STAT TRADE INC
MYAMBUTOL, ETHAMBUTOL HYDROCHLORIDE
NAPRELAN, NAPROXEN SODIUM

STERIS

* STERIS LABORATORIES INC
BROMPHENIRAMINE MALEATE, BROMPHENIRAMINE MALEATE
CYANOCOBALAMIN, CYANOCOBALAMIN
DIAZEPAM, DIAZEPAM
DIMENHYDRINATE, DIMENHYDRINATE
DIPHENHYDRAMINE HYDROCHLORIDE PRESERVATIVE FREE, DIPHENHYDRAMINE HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
DOBUTAMINE HYDROCHLORIDE, DOBUTAMINE HYDROCHLORIDE
DOXAPRAM HYDROCHLORIDE, DOXAPRAM HYDROCHLORIDE
ESTRADIOL CYPIONATE, ESTRADIOL CYPIONATE
ESTRADIOL VALERATE, ESTRADIOL VALERATE
ESTRONE, ESTRONE
FLUOROURACIL, FLUOROURACIL
HALOPERIDOL, HALOPERIDOL LACTATE
HYDROMORPHONE HYDROCHLORIDE, HYDROMORPHONE HYDROCHLORIDE
HYDROXOCOBALAMIN, HYDROXOCOBALAMIN
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
LORAZEPAM, LORAZEPAM
MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE, MEPERIDINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
MEPIVACAINE HYDROCHLORIDE, MEPIVACAINE HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MORPHINE SULFATE, MORPHINE SULFATE
NANDROLONE DECANOATE, NANDROLONE DECANOATE
ORPHENADRINE CITRATE, ORPHENADRINE CITRATE
PENTAMIDINE ISETHIONATE, PENTAMIDINE ISETHIONATE
PREDNISOLONE ACETATE, PREDNISOLONE ACETATE
PROCAINE HYDROCHLORIDE, PROCAINE HYDROCHLORIDE
PROCHLORPERAZINE EDISYLATE, PROCHLORPERAZINE EDISYLATE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PYRIDOXINE HYDROCHLORIDE, PYRIDOXINE HYDROCHLORIDE
SULFACETAMIDE SODIUM, SULFACETAMIDE SODIUM
TESTOSTERONE CYPIONATE, TESTOSTERONE CYPIONATE
TESTOSTERONE ENANTHATE, TESTOSTERONE ENANTHATE
TESTOSTERONE PROPIONATE, TESTOSTERONE PROPIONATE
TESTOSTERONE, TESTOSTERONE
THIAMINE HYDROCHLORIDE, THIAMINE HYDROCHLORIDE
VECURONIUM BROMIDE, VECURONIUM BROMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** S ****

STEVENS J

- * JEROME STEVENS PHARMACEUTICALS INC
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
CEPHALEXIN, CEPHALEXIN
DIGOXIN, DIGOXIN
METHOCARBAMOL AND ASPIRIN, ASPIRIN
ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE, ASPIRIN
UNITHROID, LEVOTHYROXINE SODIUM

STIEFEL

- * STIEFEL LABORATORIES INC
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
CLINDETS, CLINDAMYCIN PHOSPHATE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DUAC, BENZOYL PEROXIDE
ERYTHROMYCIN, ERYTHROMYCIN
GRISEOFULVIN, GRISEOFULVIN, MICROCRYSTALLINE
STIE-CORT, HYDROCORTISONE

SUMMERS

- * SUMMERS LABORATORIES INC
CROTAN, CROTAMITON

SUN PHARM INDS (IN)

- * SUN PHARMACEUTICAL INDUSTRIES LTD
CEPHALEXIN, CEPHALEXIN
ORTHO-EST, ESTROPIPATE

SUPERGEN

- * SUPERGEN INC
ETOPOSIDE, ETOPOSIDE
MITOMYCIN, MITOMYCIN
MYTOZYTREX, MITOMYCIN
NIPENT, PENTOSTATIN
PACLITAXEL, PACLITAXEL

SUPPOSITORIA

- * SUPPOSITORIA LABORATORIES INC
ACETAMINOPHEN, ACETAMINOPHEN (OTC)

SWEDISH ORPHAN

- * SWEDISH ORPHAN AB
ORFADIN, NITISINONE

SYNCOR PHARMS

- * SYNCOR PHARMACEUTICALS INC
SODIUM IODIDE I 123, SODIUM IODIDE, I-123

SYNTHON PHARMS

- * SYNTHON PHARMACEUTICALS LTD
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
PEXEVA, PAROXETINE MESYLATE

SYOSSET

- * SYOSSET LABORATORIES INC
E-SOLVE 2, ERYTHROMYCIN

TABLICAPS

- * TABLICAPS INC
FOLIC ACID, FOLIC ACID
MEPROBAMATE, MEPROBAMATE
METHOCARBAMOL, METHOCARBAMOL

TAKEDA GLOBAL

- * TAKEDA GLOBAL RESEARCH DEVELOPMENT CENTER INC
ACTOPLUS MET, METFORMIN
ROZEREM, RAMELTEON

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** T ****

TAKEDA PHARMS NA

* TAKEDA PHARMACEUTICALS NORTH AMERICA INC
ACTOS, PIOGLITAZONE HYDROCHLORIDE

TANON

* TANON PHARMACEUTICALS LLC
NYSTATIN, NYSTATIN

TAP PHARM

* TAP PHARMACEUTICAL PRODUCTS INC
LUPRON DEPOT, LEUPROLIDE ACETATE
LUPRON DEPOT-3, LEUPROLIDE ACETATE
LUPRON DEPOT-4, LEUPROLIDE ACETATE
LUPRON DEPOT-PED, LEUPROLIDE ACETATE
LUPRON, LEUPROLIDE ACETATE
NAPRAPAC 250 (COPACKAGED), LANSOPRAZOLE
NAPRAPAC 375 (COPACKAGED), LANSOPRAZOLE
NAPRAPAC 500 (COPACKAGED), LANSOPRAZOLE
PREVACID IV, LANSOPRAZOLE
PREVACID, LANSOPRAZOLE
PREVPAC, AMOXICILLIN

TARGACEPT

* TARGACEPT INC
INVERSINE, MECAMYLAMINE HYDROCHLORIDE

TARO

* TARO PHARMACEUTICAL INDUSTRIES LTD
ACETAZOLAMIDE, ACETAZOLAMIDE
CARBAMAZEPINE, CARBAMAZEPINE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ETODOLAC, ETODOLAC
FLUCONAZOLE, FLUCONAZOLE
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE
KETOCONAZOLE, KETOCONAZOLE
LORATADINE, LORATADINE (OTC)
PHENYTOIN, PHENYTOIN
TERIL, CARBAMAZEPINE

* TARO PHARMACEUTICALS INC
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLOTRIMAZOLE, CLOTRIMAZOLE
CLOTRIMAZOLE, CLOTRIMAZOLE (OTC)
DESOXIMETASONE, DESOXIMETASONE
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE, FLUOCINONIDE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
WARFARIN SODIUM, WARFARIN SODIUM

* TARO PHARMACEUTICALS USA INC
ACETIC ACID, ACETIC ACID, GLACIAL
ALCLOMETASONE DIPROPIONATE, ALCLOMETASONE DIPROPIONATE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMMONIUM LACTATE, AMMONIUM LACTATE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
CHILDREN'S ELIXSURE, IBUPROFEN (OTC)
CICLOPIROX, CICLOPIROX
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CLOBETASOL PROPIONATE (EMOLLIENT), CLOBETASOL PROPIONATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** T **

* TARO PHARMACEUTICALS USA INC
CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
CLOTRIMAZOLE, CLOTRIMAZOLE
DERMABET, BETAMETHASONE VALERATE
DESONIDE, DESONIDE
DIFLORASONE DIACETATE, DIFLORASONE DIACETATE
ECONAZOLE NITRATE, ECONAZOLE NITRATE
FLUOCINOLONE ACETONIDE, FLUOCINOLONE ACETONIDE
FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
FLUOCINONIDE, FLUOCINONIDE
FLUOROURACIL, FLUOROURACIL
GENTAMICIN SULFATE, GENTAMICIN SULFATE
HALOBETASOL PROPIONATE, HALOBETASOL PROPIONATE
HYDROCORTISONE AND ACETIC ACID, ACETIC ACID, GLACIAL
HYDROCORTISONE, HYDROCORTISONE
KETOZOLE, KETOCONAZOLE
LIDOCAINE, LIDOCAINE
LORATADINE, LORATADINE (OTC)
MICONAZOLE 3, MICONAZOLE NITRATE (OTC)
MOMETASONE FUROATE, MOMETASONE FUROATE
MUPIROCIN, MUPIROCIN
NITROFURAZONE, NITROFURAZONE
NYSTATIN AND TRIAMCINOLONE ACETONIDE, NYSTATIN
NYSTATIN, NYSTATIN
ORACORT, TRIAMCINOLONE ACETONIDE
ORALONE, TRIAMCINOLONE ACETONIDE
TERCONAZOLE, TERCONAZOLE
TERIL, CARBAMAZEPINE
THEOPHYLLINE, THEOPHYLLINE
TRIAMCINOLONE ACETONIDE, TRIAMCINOLONE ACETONIDE
TRIVAGIZOLE 3, CLOTRIMAZOLE (OTC)
U-CORT, HYDROCORTISONE ACETATE

TARO PHARM INDS

* TARO PHARMACEUTICAL INDUSTRIES LTD
AMCINONIDE, AMCINONIDE
CARBAMAZEPINE, CARBAMAZEPINE
CLINDAMYCIN PHOSPHATE, CLINDAMYCIN PHOSPHATE
ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
ETODOLAC, ETODOLAC
FLUTICASONE PROPIONATE, FLUTICASONE PROPIONATE
HYDROCORTISONE BUTYRATE, HYDROCORTISONE BUTYRATE

TARO PHARMS NORTH

* TARO PHARMACEUTICALS NORTH AMERICA INC
OVIDE, MALATHION
TOPICORT LP, DESOXIMETASONE
TOPICORT, DESOXIMETASONE

TAYLOR

* TAYLOR PHARMACEUTICALS
LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
NAPHAZOLINE HYDROCHLORIDE, NAPHAZOLINE HYDROCHLORIDE

TAYLOR PHARMA

* TAYLOR PHARMACAL CO
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
MIDAZOLAM HYDROCHLORIDE, MIDAZOLAM HYDROCHLORIDE
PROPARACAINE HYDROCHLORIDE, PROPARACAINE HYDROCHLORIDE
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE, POLYMYXIN B SULFATE

TECHNILAB

* TECHNILAB INC
ACILAC, LACTULOSE
LAXILOSE, LACTULOSE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** T ****

TEIKOKU PHARMA USA

* TEIKOKU PHARMA USA INC
LIDODERM, LIDOCAINE

TERCICA

* TERCICA INC
INCRELEX, MECASERMIN RECOMBINANT

TEVA

* TEVA NEUROSCIENCE INC
COPAXONE, GLATIRAMER ACETATE

* TEVA PHARMACEUTICALS USA INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACYCLOVIR, ACYCLOVIR
ADIPEX-P, PHENTERMINE HYDROCHLORIDE
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALPRAZOLAM, ALPRAZOLAM
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
AMOXICILLIN AND CLAVULANATE POTASSIUM, AMOXICILLIN
AMOXICILLIN PEDIATRIC, AMOXICILLIN
AMOXICILLIN, AMOXICILLIN
ATENOLOL, ATENOLOL
AZITHROMYCIN, AZITHROMYCIN
BENAZEPRIL HYDROCHLORIDE, BENAZEPRIL HYDROCHLORIDE
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETA-VAL, BETAMETHASONE VALERATE
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BUPROPION HYDROCHLORIDE, BUPROPION HYDROCHLORIDE
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CALCITRIOL, CALCITRIOL
CAPTOPRIL AND HYDROCHLOROTHIAZIDE, CAPTOPRIL
CAPTOPRIL, CAPTOPRIL
CARBIDOPA AND LEVODOPA, CARBIDOPA
CEFACLOX, CEFACLOX
CEFPROXIL, CEFPROXIL
CEFUROXIME AXETIL, CEFUROXIME AXETIL
CEFUROXIME, CEFUROXIME SODIUM
CEPHALEXIN, CEPHALEXIN
CEPHRADINE, CEPHRADINE
CHLORHEXIDINE GLUCONATE, CHLORHEXIDINE GLUCONATE
CHLORZOXAZONE, CHLORZOXAZONE
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
CHOLESTYRAMINE, CHOLESTYRAMINE
CILOSTAZOL, CILOSTAZOL
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CIMETIDINE, CIMETIDINE
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CLARITHROMYCIN, CLARITHROMYCIN
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE (OTC)
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
CLOFIBRATE, CLOFIBRATE
CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CLOTRIMAZOLE, CLOTRIMAZOLE
CLOXACILLIN SODIUM, CLOXACILLIN SODIUM
CLOZAPINE, CLOZAPINE
COTRIM D.S., SULFAMETHOXAZOLE
COTRIM, SULFAMETHOXAZOLE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DICLOXACILLIN SODIUM, DICLOXACILLIN SODIUM
DIFLUNISAL, DIFLUNISAL

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** T **

* TEVA PHARMACEUTICALS USA INC
 DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 ENALAPRIL MALEATE AND HYDROCHLOROTHIAZIDE, ENALAPRIL MALEATE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 EPITOL, CARBAMAZEPINE
 ESTAZOLAM, ESTAZOLAM
 ETODOLAC, ETODOLAC
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FENOFIBRATE (MICRONIZED), FENOFIBRATE
 FENOFIBRATE, FENOFIBRATE
 FEXOFENADINE HYDROCHLORIDE, FEXOFENADINE HYDROCHLORIDE
 FLUCONAZOLE, FLUCONAZOLE
 FLUOCINONIDE EMULSIFIED BASE, FLUOCINONIDE
 FLUOCINONIDE, FLUOCINONIDE
 FLUOXETINE HYDROCHLORIDE, FLUOXETINE HYDROCHLORIDE
 FLUOXETINE, FLUOXETINE HYDROCHLORIDE
 FLURBIPROFEN, FLURBIPROFEN
 FLUTAMIDE, FLUTAMIDE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 GABAPENTIN, GABAPENTIN
 GALZIN, ZINC ACETATE
 GEMFIBROZIL, GEMFIBROZIL
 GLIMEPIRIDE, GLIMEPIRIDE
 GLIPIZIDE, GLIPIZIDE
 GLYBURIDE (MICRONIZED), GLYBURIDE
 GLYBURIDE AND METFORMIN HYDROCHLORIDE, GLYBURIDE
 GLYBURIDE, GLYBURIDE
 HYDROCODONE BITARTRATE AND IBUPROFEN, HYDROCODONE BITARTRATE
 INDAPAMIDE, INDAPAMIDE
 INDO-LEMMON, INDOMETHACIN
 INDOMETHACIN, INDOMETHACIN
 ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
 KETOCONAZOLE, KETOCONAZOLE
 KETOPROFEN, KETOPROFEN
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVOFLOXACIN, LEVOFLOXACIN
 LISINOPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 LISINOPRIL, LISINOPRIL
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE
 LOPERAMIDE HYDROCHLORIDE, LOPERAMIDE HYDROCHLORIDE (OTC)
 LORATADINE, LORATADINE (OTC)
 LOVASTATIN, LOVASTATIN
 MEGESTROL ACETATE, MEGESTROL ACETATE
 MESALAMINE, MESALAMINE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE, METRONIDAZOLE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL (FOR MEN), MINOXIDIL (OTC)
 MINOXIDIL EXTRA STRENGTH (FOR MEN), MINOXIDIL (OTC)
 MIRTAZAPINE, MIRTAZAPINE
 MOEXIPRIL HYDROCHLORIDE, MOEXIPRIL HYDROCHLORIDE
 MUPIROCIN, MUPIROCIN
 NABUMETONE, NABUMETONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** T ****

* TEVA PHARMACEUTICALS USA INC
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN, NAPROXEN
NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
NEOMYCIN SULFATE, NEOMYCIN SULFATE
NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
NIFEDIPINE, NIFEDIPINE
NIZATIDINE, NIZATIDINE
NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
NYSTATIN, NYSTATIN
OFLOXACIN, OFLOXACIN
ORAP, PIMOZIDE
OXACILLIN SODIUM, OXACILLIN SODIUM
OXAPROZIN, OXAPROZIN
OXYCODONE HYDROCHLORIDE, OXYCODONE HYDROCHLORIDE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PENICILLIN-VK, PENICILLIN V POTASSIUM
PENTOXIFYLLINE, PENTOXIFYLLINE
PERGOLIDE MESYLATE, PERGOLIDE MESYLATE
PINDOLOL, PINDOLOL
PIROXICAM, PIROXICAM
POTASSIUM CHLORIDE, POTASSIUM CHLORIDE
PRELONE, PREDNISOLONE
PROPOXYPHENE COMPOUND 65, ASPIRIN
PROPOXYPHENE HYDROCHLORIDE, PROPOXYPHENE HYDROCHLORIDE
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
PURINETHOL, MERCAPTOPURINE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RIBAVIRIN, RIBAVIRIN
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SUCRALFATE, SUCRALFATE
SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SULINDAC, SULINDAC
TAMOXIFEN CITRATE, TAMOXIFEN CITRATE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TOLMETIN SODIUM, TOLMETIN SODIUM
TORSEMIDE, TORSEMIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRIAKET, TRIAMCINOLONE ACETONIDE
TRIMETHOPRIM, TRIMETHOPRIM

TEVA PHARMS

* TEVA PHARMACEUTICALS USA
ACYCLOVIR, ACYCLOVIR
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
AMIODARONE HYDROCHLORIDE, AMIODARONE HYDROCHLORIDE
ATENOLOL, ATENOLOL
BETAMETHASONE DIPROPIONATE, BETAMETHASONE DIPROPIONATE
BETAMETHASONE VALERATE, BETAMETHASONE VALERATE
BISOPROLOL FUMARATE, BISOPROLOL FUMARATE
CAPTOPRIL, CAPTOPRIL
CEFPROZIL, CEFPROZIL
CHOLESTYRAMINE LIGHT, CHOLESTYRAMINE
CHOLESTYRAMINE, CHOLESTYRAMINE
CIMETIDINE HYDROCHLORIDE, CIMETIDINE HYDROCHLORIDE
CLEMASTINE FUMARATE, CLEMASTINE FUMARATE
CLOBETASOL PROPIONATE, CLOBETASOL PROPIONATE
DESONIDE, DESONIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** T ****

* TEVA PHARMACEUTICALS USA
DICLOFENAC SODIUM, DICLOFENAC SODIUM
DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
ETHOSUXIMIDE, ETHOSUXIMIDE
EVALOSE, LACTULOSE
FLUOCINONIDE, FLUOCINONIDE
FLUPHENAZINE HYDROCHLORIDE, FLUPHENAZINE HYDROCHLORIDE
GABAPENTIN, GABAPENTIN
GLIPIZIDE AND METFORMIN HYDROCHLORIDE, GLIPIZIDE
GUANABENZ ACETATE, GUANABENZ ACETATE
GYNIX, CLOTRIMAZOLE (OTC)
HALOPERIDOL, HALOPERIDOL LACTATE
HEPTALAC, LACTULOSE
HYDROCORTISONE VALERATE, HYDROCORTISONE VALERATE
HYDROCORTISONE, HYDROCORTISONE
HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
LEFLUNOMIDE, LEFLUNOMIDE
LIDOCAINE HYDROCHLORIDE, LIDOCAINE HYDROCHLORIDE
MEBENDAZOLE, MEBENDAZOLE
MEGESTROL ACETATE, MEGESTROL ACETATE
METHAZOLAMIDE, METHAZOLAMIDE
METOPROLOL TARTRATE, METOPROLOL TARTRATE
MICONAZOLE NITRATE, MICONAZOLE NITRATE (OTC)
NADOLOL, NADOLOL
NAPROXEN SODIUM, NAPROXEN SODIUM
NAPROXEN, NAPROXEN
PIROXICAM, PIROXICAM
PREDNISOLONE, PREDNISOLONE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
QUINIDINE SULFATE, QUINIDINE SULFATE
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
THIOTHIXENE HYDROCHLORIDE, THIOTHIXENE HYDROCHLORIDE
TRETINOIN, TRETINOIN
URSODIOL, URSODIOL
VALPROIC ACID, VALPROIC ACID
VANDAZOLE, METRONIDAZOLE
ZONISAMIDE, ZONISAMIDE

THERAKOS

* THERAKOS INC
UVADEX, METHOXSALEN

THREE RIVERS PHARMS

* THREE RIVERS PHARMACEUTICALS LLC
AMPHOTEC, AMPHOTERICIN B
RIBASPHERE, RIBAVIRIN
RIBAVIRIN, RIBAVIRIN

TORPHARM

* TORPHARM INC
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
CAPTOPRIL, CAPTOPRIL
CARBIDOPA AND LEVODOPA, CARBIDOPA
CIMETIDINE, CIMETIDINE
CIMETIDINE, CIMETIDINE (OTC)
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
CLONAZEPAM, CLONAZEPAM
CYCLOSPORINE, CYCLOSPORINE
DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
DILT-CD, DILTIAZEM HYDROCHLORIDE
DILTIAZEM HYDROCHLORIDE, DILTIAZEM HYDROCHLORIDE
DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
ENALAPRIL MALEATE, ENALAPRIL MALEATE
ETODOLAC, ETODOLAC
FAMOTIDINE, FAMOTIDINE
FAMOTIDINE, FAMOTIDINE (OTC)

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** T ****

* TORPHARM INC
FLUCONAZOLE, FLUCONAZOLE
FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
GEMFIBROZIL, GEMFIBROZIL
GLIPIZIDE, GLIPIZIDE
KETOCONAZOLE, KETOCONAZOLE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
NIZATIDINE, NIZATIDINE
OMEPRAZOLE, OMEPRAZOLE
PAROXETINE HYDROCHLORIDE, PAROXETINE HYDROCHLORIDE
PENTOXIFYLLINE, PENTOXIFYLLINE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
SELEGILINE HYDROCHLORIDE, SELEGILINE HYDROCHLORIDE
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE
TICLOPIDINE HYDROCHLORIDE, TICLOPIDINE HYDROCHLORIDE
TIZANIDINE HYDROCHLORIDE, TIZANIDINE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

TRIGEN

* TRIGEN LABORATORIES INC
FOLIC ACID, FOLIC ACID
INDAPAMIDE, INDAPAMIDE
METHYLPREDNISOLONE, METHYLPREDNISOLONE
PREDNISONE, PREDNISONE
PROCHLORPERAZINE MALEATE, PROCHLORPERAZINE MALEATE
TERAZOSIN HYDROCHLORIDE, TERAZOSIN HYDROCHLORIDE

TRUXTON INC

* TRUXTON INC
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE

TYCO HLTHCARE

* TYCO HEALTHCARE GROUP LP
ANAFRANIL, CLOMIPRAMINE HYDROCHLORIDE
PAMELOR, NORTRIPTYLINE HYDROCHLORIDE
RESTORIL, TEMAZEPAM
TOFRANIL, IMIPRAMINE HYDROCHLORIDE
TOFRANIL-PM, IMIPRAMINE PAMOATE

UCB

* UCB PHARMA INC
CODEPREX, CHLORPHENIRAMINE POLISTIREX
DELSYM, DEXTROMETHORPHAN POLISTIREX (OTC)
DIPENTUM, OLSALAZINE SODIUM
GASTROCROM, CROMOLYN SODIUM
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
IONAMIN, PHENTERMINE RESIN COMPLEX
KEPPRA, LEVETIRACETAM
LORTAB, ACETAMINOPHEN
METADATE CD, METHYLPHENIDATE HYDROCHLORIDE
METADATE ER, METHYLPHENIDATE HYDROCHLORIDE
METHYLPHENIDATE HCL, METHYLPHENIDATE HYDROCHLORIDE
PEDIAPRED, PREDNISOLONE SODIUM PHOSPHATE
SEMPREX-D, ACRIVASTINE
THEO-24, THEOPHYLLINE
TUSSIONEX, CHLORPHENIRAMINE POLISTIREX
ZAROXOLYN, METOLAZONE

UCYCLYD

* UCYCLYD PHARMA INC
AMMONUL, SODIUM BENZOATE

UDL

* UDL LABORATORIES INC
ALBUTEROL SULFATE, ALBUTEROL SULFATE
HALOPERIDOL LACTATE, HALOPERIDOL LACTATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** U ****

* UDL LABORATORIES INC
METHADONE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
PREDNISOLONE, PREDNISOLONE
VALPROIC ACID, VALPROIC ACID

ULURU

* ULURU INC
AMLEXANOX, AMLEXANOX
APHTHASOL, AMLEXANOX

UNIGENE

* UNIGENE LABORATORIES
FORTICAL, CALCITONIN SALMON RECOMBINANT

UNIMED

* UNIMED INC
MARINOL, DRONABINOL

UNIMED PHARMS

* UNIMED PHARMACEUTICALS INC
ANADROL-50, OXYMETHOLONE
ANDROGEL, TESTOSTERONE
PROMETRIUM, PROGESTERONE

UNIQUE PHARM LABS

* UNIQUE PHARMACEUTICAL LABORATORIES
CIPROFLOXACIN, CIPROFLOXACIN HYDROCHLORIDE
FLUCONAZOLE, FLUCONAZOLE

UNITED GUARDIAN

* UNITED GUARDIAN INC
RENACIDIN, CITRIC ACID

UNITED THERAP

* UNITED THERAPEUTICS CORP
REMODULIN, TREPROSTINIL SODIUM

UPSHER SMITH

* UPSHER SMITH LABORATORIES INC
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
KLOR-CON M10, POTASSIUM CHLORIDE
KLOR-CON M15, POTASSIUM CHLORIDE
KLOR-CON M20, POTASSIUM CHLORIDE
KLOR-CON, POTASSIUM CHLORIDE
NIACOR, NIACIN
NYSTATIN, NYSTATIN
ORVATEN, MIDODRINE HYDROCHLORIDE
PACERONE, AMIODARONE HYDROCHLORIDE
PENTOXIL, PENTOXIFYLLINE
PREVALITE, CHOLESTYRAMINE
SORINE, SOTALOL HYDROCHLORIDE

US SURGCL

* UNITED STATES SURGICAL DIV TYCO HEALTHCARE GROUP LP
LYMPHAZURIN, ISOSULFAN BLUE

USL PHARMA

* USL PHARMA INC
AMANTADINE HYDROCHLORIDE, AMANTADINE HYDROCHLORIDE
BACLOFEN, BACLOFEN
BENZTROPINE MESYLATE, BENZTROPINE MESYLATE
CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CHLORPROMAZINE HYDROCHLORIDE, CHLORPROMAZINE HYDROCHLORIDE
CLOFIBRATE, CLOFIBRATE
ESTRADIOL, ESTRADIOL
FLUOXYMESTERONE, FLUOXYMESTERONE
JANTOVEN, WARFARIN SODIUM
MEDROXYPROGESTERONE ACETATE, MEDROXYPROGESTERONE ACETATE
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
VALPROIC ACID, VALPROIC ACID

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** V ****

VALEANT

* VALEANT PHARMACEUTICALS INTERNATIONAL
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ANCOBON, FLUCYTOSINE
BONTRIL PDM, PHENDIMETRAZINE TARTRATE
D.H.E. 45, DIHYDROERGOTAMINE MESYLATE
DIASTAT ACUDIAL, DIAZEPAM
DIASTAT, DIAZEPAM
FLUOROURACIL, FLUOROURACIL
MIGRANAL, DIHYDROERGOTAMINE MESYLATE
MOTOFEN, ATROPINE SULFATE
MYSOLINE, PRIMIDONE
PERMAX, PERGOLIDE MESYLATE
PHRENILIN FORTE, ACETAMINOPHEN
PHRENILIN WITH CAFFEINE AND CODEINE, ACETAMINOPHEN
PHRENILIN, ACETAMINOPHEN
TASMAR, TOLCAPONE

VALEANT PHARM INTL

* VALEANT PHARMACEUTICALS INTERNATIONAL
8-MOP, METHOXSALEN
ANDROID 10, METHYLTESTOSTERONE
ANDROID 25, METHYLTESTOSTERONE
BENOQUIN, MONOBENZONE
DALMANE, FLURAZEPAM HYDROCHLORIDE
DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
EFUDEX, FLUOROURACIL
LEVO-DROMORAN, LEVORPHANOL TARTRATE
LIBRIUM, CHLORDIAZEPOXIDE HYDROCHLORIDE
LIMBITROL DS, AMITRIPTYLINE HYDROCHLORIDE
LIMBITROL, AMITRIPTYLINE HYDROCHLORIDE
MESTINON, PYRIDOSTIGMINE BROMIDE
OXSORALEN, METHOXSALEN
OXSORALEN-ULTRA, METHOXSALEN
TENSILON PRESERVATIVE FREE, EDROPHONIUM CHLORIDE
TENSILON, EDROPHONIUM CHLORIDE
TESTRED, METHYLTESTOSTERONE
VIRAZOLE, RIBAVIRIN

VALERA

* VALERA PHARMACEUTICALS INC
VANTAS, HISTRELIN ACETATE

VATRING PHARMS

* VATRING PHARMACEUTICALS LP
UREX, METHENAMINE HIPPURATE

VERNALIS

* VERNALIS RD LTD
APOKYN, APOMORPHINE HYDROCHLORIDE

VERSAPHARM

* VERSAPHARM INC
RIFAMPIN, RIFAMPIN

VERUS PHARMS

* VERUS PHARMACEUTICALS INC
TWINJECT 0.3, EPINEPHRINE
TWINJECT, EPINEPHRINE

VINTAGE

* VINTAGE PHARMACEUTICALS LLC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETIC ACID, ACETIC ACID, GLACIAL
LEVOLET, LEVOTHYROXINE SODIUM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** V **

VINTAGE PHARMS

* VINTAGE PHARMACEUTICALS INC
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAMINOPHEN, BUTALBITAL, CAFFEINE, AND CODEINE PHOSPHATE, ACETAMINOPHEN
ALLOPURINOL, ALLOPURINOL
AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
BACLOFEN, BACLOFEN
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
CARISOPRODOL, CARISOPRODOL
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
FUROSEMIDE, FUROSEMIDE
HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
HYDROCORTISONE, HYDROCORTISONE
HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
IBUPROFEN, IBUPROFEN
IBUPROFEN, IBUPROFEN (OTC)
LACTULOSE, LACTULOSE
MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
METHOCARBAMOL, METHOCARBAMOL
METHYLPREDNISOLONE, METHYLPREDNISOLONE
NYSTATIN, NYSTATIN
OXYBUTYNIN CHLORIDE, OXYBUTYNIN CHLORIDE
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PERPHENAZINE, PERPHENAZINE
PHENTERMINE HYDROCHLORIDE, PHENTERMINE HYDROCHLORIDE
PREDNISONE, PREDNISONE
PRIMIDONE, PRIMIDONE
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN, ACETAMINOPHEN
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SULFASALAZINE, SULFASALAZINE
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE

VIROPHARMA

* VIROPHARMA INC
VANCOCIN HYDROCHLORIDE, VANCOMYCIN HYDROCHLORIDE

VISTA PHARMS

* VISTA PHARMACEUTICALS INC
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE

VISTAPHARM

* VISTAPHARM INC
LACTULOSE, LACTULOSE
METOCLOPRAMIDE, METOCLOPRAMIDE HYDROCHLORIDE
NYSTATIN, NYSTATIN
PHENYTOIN, PHENYTOIN

VIVUS

* VIVUS INC
MUSE, ALPROSTADIL

VYTERIS

* VYTERIS INC
LIDOSITE TOPICAL SYSTEM KIT, EPINEPHRINE

WARNER CHILCOTT

* WARNER CHILCOTT BERMUDA LTD
CHOLEDYL SA, OXTRIPHYLLINE
DORYX, DOXYCYCLINE HYCLATE
* WARNER CHILCOTT INC
CARBAMAZEPINE, CARBAMAZEPINE
DURICEF, CEFADROXIL/CEFADROXIL HEMIHYDRATE
ERYC, ERYTHROMYCIN

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** W ****

* WARNER CHILCOTT INC
ESTRACE, ESTRADIOL
ESTROSTEP FE, ETHINYL ESTRADIOL
FEMHRT, ETHINYL ESTRADIOL
FEMTRACE, ESTRADIOL ACETATE
FLURBIPROFEN, FLURBIPROFEN
LOESTRIN 21 1.5/30, ETHINYL ESTRADIOL
LOESTRIN 21 1/20, ETHINYL ESTRADIOL
LOESTRIN FE 1.5/30, ETHINYL ESTRADIOL
LOESTRIN FE 1/20, ETHINYL ESTRADIOL
OVCON-35, ETHINYL ESTRADIOL
OVCON-50, ETHINYL ESTRADIOL

WARNER LAMBERT

* WARNER LAMBERT CO
SUDAFED 12 HOUR, PSEUDOEPHEDRINE HYDROCHLORIDE (OTC)
ZANTAC 75, RANITIDINE HYDROCHLORIDE (OTC)

WARRICK PHARMS

* WARRICK PHARMACEUTICALS
CROMOLYN SODIUM, CROMOLYN SODIUM
IPRATROPIUM BROMIDE, IPRATROPIUM BROMIDE

WATSON LABS

* WATSON LABORATORIES
ANAGRELIDE HYDROCHLORIDE, ANAGRELIDE HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
LEVONORGESTREL AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
MICROGESTIN FE 1.5/30, ETHINYL ESTRADIOL
MICROGESTIN FE 1/20, ETHINYL ESTRADIOL
NORCO, ACETAMINOPHEN
OXAPROZIN, OXAPROZIN
OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
PENTAZOCINE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPAFENONE HYDROCHLORIDE, PROPAFENONE HYDROCHLORIDE
TRAMADOL HYDROCHLORIDE, TRAMADOL HYDROCHLORIDE

* WATSON LABORATORIES INC
ACEBUTOLOL HYDROCHLORIDE, ACEBUTOLOL HYDROCHLORIDE
ACETAMINOPHEN AND CODEINE PHOSPHATE, ACETAMINOPHEN
ACETAZOLAMIDE, ACETAZOLAMIDE
ACETOHEXAMIDE, ACETOHEXAMIDE
ACYCLOVIR, ACYCLOVIR
ALBUTEROL SULFATE, ALBUTEROL SULFATE
ALLOPURINOL, ALLOPURINOL
ALORA, ESTRADIOL
ALPRAZOLAM, ALPRAZOLAM
AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, AMILORIDE HYDROCHLORIDE
AMOXAPINE, AMOXAPINE
ANDRODERM, TESTOSTERONE
ATENOLOL AND CHLORTHALIDONE, ATENOLOL
ATENOLOL, ATENOLOL
BACLOFEN, BACLOFEN
BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE, BISOPROLOL FUMARATE
BREVICON 21-DAY, ETHINYL ESTRADIOL
BREVICON 28-DAY, ETHINYL ESTRADIOL
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, APAP, AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE, ASPIRIN
CAPTOPRIL, CAPTOPRIL
CARISOPRODOL, CARISOPRODOL
CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
CHLORDIAZEPOXIDE HYDROCHLORIDE, CHLORDIAZEPOXIDE HYDROCHLORIDE
CHLORPROPAMIDE, CHLORPROPAMIDE
CHLORZOXAZONE, CHLORZOXAZONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** W **

* WATSON LABORATORIES INC
 CIMETIDINE, CIMETIDINE
 CIMETIDINE, CIMETIDINE (OTC)
 CITALOPRAM HYDROBROMIDE, CITALOPRAM HYDROBROMIDE
 CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
 CLOMIPRAMINE HYDROCHLORIDE, CLOMIPRAMINE HYDROCHLORIDE
 CLONAZEPAM, CLONAZEPAM
 CLONIDINE HYDROCHLORIDE, CLONIDINE HYDROCHLORIDE
 CLORAZEPATE DIPOTASSIUM, CLORAZEPATE DIPOTASSIUM
 COL-PROBENECID, COLCHICINE
 CORDRAN, FLURANDRENOLIDE
 CYCLOBENZAPRINE HYDROCHLORIDE, CYCLOBENZAPRINE HYDROCHLORIDE
 DESOGESTREL AND ETHINYL ESTRADIOL, DESOGESTREL
 DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE, AMPHETAMINE
 ASPARTATE
 DIAZEPAM, DIAZEPAM
 DICLOFENAC POTASSIUM, DICLOFENAC POTASSIUM
 DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
 DIFLUNISAL, DIFLUNISAL
 DILACOR XR, DILTIAZEM HYDROCHLORIDE
 DIPHENHYDRAMINE HYDROCHLORIDE, DIPHENHYDRAMINE HYDROCHLORIDE
 DIPYRIDAMOLE, DIPYRIDAMOLE
 DISOPYRAMIDE PHOSPHATE, DISOPYRAMIDE PHOSPHATE
 DOXAZOSIN MESYLATE, DOXAZOSIN MESYLATE
 DOXEPIN HYDROCHLORIDE, DOXEPIN HYDROCHLORIDE
 DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
 DOXYCYCLINE, DOXYCYCLINE
 ENALAPRIL MALEATE, ENALAPRIL MALEATE
 ERGOLOID MESYLATES, ERGOLOID MESYLATES
 ESTAZOLAM, ESTAZOLAM
 ESTRADIOL, ESTRADIOL
 ESTROPIRATE, ESTROPIRATE
 ETODOLAC, ETODOLAC
 FAMOTIDINE, FAMOTIDINE
 FAMOTIDINE, FAMOTIDINE (OTC)
 FENOPROFEN CALCIUM, FENOPROFEN CALCIUM
 FLUOXETINE, FLUOXETINE HYDROCHLORIDE
 FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
 FLUVOXAMINE MALEATE, FLUVOXAMINE MALEATE
 FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE, FOSINOPRIL SODIUM
 FOSINOPRIL SODIUM, FOSINOPRIL SODIUM
 FUROSEMIDE, FUROSEMIDE
 GEMFIBROZIL, GEMFIBROZIL
 GERIMAL, ERGOLOID MESYLATES
 GLIPIZIDE, GLIPIZIDE
 GUANABENZ ACETATE, GUANABENZ ACETATE
 GUANFACINE HYDROCHLORIDE, GUANFACINE HYDROCHLORIDE
 HALOPERIDOL, HALOPERIDOL
 HYDRALAZINE HYDROCHLORIDE, HYDRALAZINE HYDROCHLORIDE
 HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 HYDROCODONE BITARTRATE AND ACETAMINOPHEN, ACETAMINOPHEN
 HYDROXYCHLOROQUINE SULFATE, HYDROXYCHLOROQUINE SULFATE
 HYDROXYZINE HYDROCHLORIDE, HYDROXYZINE HYDROCHLORIDE
 HYDROXYZINE PAMOATE, HYDROXYZINE PAMOATE
 IBUPROFEN, IBUPROFEN
 IBUPROFEN, IBUPROFEN (OTC)
 INDAPAMIDE, INDAPAMIDE
 ISONIAZID, ISONIAZID
 ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
 KETOROLAC TROMETHAMINE, KETOROLAC TROMETHAMINE
 LABETALOL HYDROCHLORIDE, LABETALOL HYDROCHLORIDE
 LEVORA 0.15/30-28, ETHINYL ESTRADIOL
 LISINOPRIL, LISINOPRIL
 LORAZEPAM, LORAZEPAM

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

** W **

* WATSON LABORATORIES INC
 LOW-OGESTREL-21, ETHINYL ESTRADIOL
 LOW-OGESTREL-28, ETHINYL ESTRADIOL
 LOXAPINE SUCCINATE, LOXAPINE SUCCINATE
 LOXITANE, LOXAPINE SUCCINATE
 MAPROTILINE HYDROCHLORIDE, MAPROTILINE HYDROCHLORIDE
 MECLIZINE HYDROCHLORIDE, MECLIZINE HYDROCHLORIDE
 MECLOFENAMATE SODIUM, MECLOFENAMATE SODIUM
 MEPERIDINE HYDROCHLORIDE, MEPERIDINE HYDROCHLORIDE
 MEPROBAMATE, MEPROBAMATE
 METAPROTERENOL SULFATE, METAPROTERENOL SULFATE
 METHOCARBAMOL, METHOCARBAMOL
 METHYCLOTHIAZIDE, METHYCLOTHIAZIDE
 METHYLDOPA AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
 METHYLDOPA, METHYLDOPA
 METHYLPHENIDATE HYDROCHLORIDE, METHYLPHENIDATE HYDROCHLORIDE
 METHYLPREDNISOLONE, METHYLPREDNISOLONE
 METOCLOPRAMIDE HYDROCHLORIDE, METOCLOPRAMIDE HYDROCHLORIDE
 METOLAZONE, METOLAZONE
 METOPROLOL TARTRATE, METOPROLOL TARTRATE
 METRONIDAZOLE, METRONIDAZOLE
 MEXILETINE HYDROCHLORIDE, MEXILETINE HYDROCHLORIDE
 MICROZIDE, HYDROCHLOROTHIAZIDE
 MINOCYCLINE HYDROCHLORIDE, MINOCYCLINE HYDROCHLORIDE
 MINOXIDIL, MINOXIDIL
 MIRTAZAPINE, MIRTAZAPINE
 MORPHINE SULFATE, MORPHINE SULFATE
 NALIDIXIC ACID, NALIDIXIC ACID
 NAPROXEN SODIUM, NAPROXEN SODIUM
 NAPROXEN, NAPROXEN
 NEFAZODONE HYDROCHLORIDE, NEFAZODONE HYDROCHLORIDE
 NICARDIPINE HYDROCHLORIDE, NICARDIPINE HYDROCHLORIDE
 NICOTINE POLACRILEX (MINT), NICOTINE POLACRILEX (OTC)
 NICOTINE POLACRILEX, NICOTINE POLACRILEX (OTC)
 NITROFURANTOIN, NITROFURANTOIN, MACROCRYSTALLINE
 NIZATIDINE, NIZATIDINE
 NORCO, ACETAMINOPHEN
 NORETHIN 1/35E-21, ETHINYL ESTRADIOL
 NORETHIN 1/35E-28, ETHINYL ESTRADIOL
 NORETHIN 1/50M-21, MESTRANOL
 NORETHIN 1/50M-28, MESTRANOL
 NORETHINDRONE AND ETHINYL ESTRADIOL (10/11), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL (7/14), ETHINYL ESTRADIOL
 NORETHINDRONE AND ETHINYL ESTRADIOL, ETHINYL ESTRADIOL
 NORETHINDRONE AND MESTRANOL, MESTRANOL
 NORINYL 1+35 21-DAY, ETHINYL ESTRADIOL
 NORINYL 1+35 28-DAY, ETHINYL ESTRADIOL
 NORINYL 1+50 21-DAY, MESTRANOL
 NORINYL 1+50 28-DAY, MESTRANOL
 NORTRIPTYLINE HYDROCHLORIDE, NORTRIPTYLINE HYDROCHLORIDE
 OGESTREL 0.5/50-21, ETHINYL ESTRADIOL
 OGESTREL 0.5/50-28, ETHINYL ESTRADIOL
 OXAZEPAM, OXAZEPAM
 OXYCODONE AND ACETAMINOPHEN, ACETAMINOPHEN
 OXYCODONE AND ASPIRIN, ASPIRIN
 PENTAZOCINE AND NALOXONE HYDROCHLORIDES, NALOXONE HYDROCHLORIDE
 PENTOXIFYLLINE, PENTOXIFYLLINE
 PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE, AMITRIPTYLINE HYDROCHLORIDE
 PINDOLOL, PINDOLOL
 PIROXICAM, PIROXICAM
 PRAZOSIN HYDROCHLORIDE, PRAZOSIN HYDROCHLORIDE
 PREDNISOLONE, PREDNISOLONE
 PREDNISONE, PREDNISONE
 PRIMIDONE, PRIMIDONE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** W ****

- * WATSON LABORATORIES INC
PROBENECID, PROBENECID
PROCAINAMIDE HYDROCHLORIDE, PROCAINAMIDE HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
PROPOXYPHENE HYDROCHLORIDE AND ACETAMINOPHEN, ACETAMINOPHEN
PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
PROPRANOLOL HYDROCHLORIDE, PROPRANOLOL HYDROCHLORIDE
QUINIDINE SULFATE, QUINIDINE SULFATE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
RANITIDINE, RANITIDINE HYDROCHLORIDE (OTC)
SOTALOL HYDROCHLORIDE, SOTALOL HYDROCHLORIDE
SPIRONOLACTONE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
SULFAMETHOXAZOLE AND TRIMETHOPRIM DOUBLE STRENGTH, SULFAMETHOXAZOLE
SULFAMETHOXAZOLE AND TRIMETHOPRIM, SULFAMETHOXAZOLE
SULFASALAZINE, SULFASALAZINE
SULINDAC, SULINDAC
TEMAZEPAM, TEMAZEPAM
THIORIDAZINE HYDROCHLORIDE, THIORIDAZINE HYDROCHLORIDE
THIOTHIXENE, THIOTHIXENE
TIMOLOL MALEATE, TIMOLOL MALEATE
TOLAZAMIDE, TOLAZAMIDE
TOLBUTAMIDE, TOLBUTAMIDE
TRAZODONE HYDROCHLORIDE, TRAZODONE HYDROCHLORIDE
TRELSTAR DEPOT, TRIPTORELIN PAMOATE
TRELSTAR, TRIPTORELIN PAMOATE
TRIAMTERENE AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
TRIAZOLAM, TRIAZOLAM
TRIHEXYPHENIDYL HYDROCHLORIDE, TRIHEXYPHENIDYL HYDROCHLORIDE
TRIMETHOPRIM, TRIMETHOPRIM
TRI-NORINYL 28-DAY, ETHINYL ESTRADIOL
TRIVORA-28, ETHINYL ESTRADIOL
VERAPAMIL HYDROCHLORIDE, VERAPAMIL HYDROCHLORIDE
ZOVIA 1/35E-21, ETHINYL ESTRADIOL
ZOVIA 1/35E-28, ETHINYL ESTRADIOL
ZOVIA 1/50E-21, ETHINYL ESTRADIOL
ZOVIA 1/50E-28, ETHINYL ESTRADIOL
- * WATSON LABS INC
LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE

WATSON LABS (UTAH)

- * WATSON LABORATORIES INC
CHORIONIC GONADOTROPIN, GONADOTROPIN, CHORIONIC
INFED, IRON DEXTRAN
NOR-QD, NORETHINDRONE
OXYTROL, OXYBUTYNIN
PROGESTERONE, PROGESTERONE

WATSON PHARMS

- * WATSON PHARMACEUTICALS INC
ACTIGALL, URSODIOL
FERRLECIT, SODIUM FERRIC GLUCONATE COMPLEX
FIORICET W/ CODEINE, ACETAMINOPHEN
FIORICET, ACETAMINOPHEN
FIORINAL W/CODEINE NO 3, ASPIRIN
FIORINAL, ASPIRIN

WE PHARMS

- * WE PHARMACEUTICALS INC
PREDNISOLONE SODIUM PHOSPHATE, PREDNISOLONE SODIUM PHOSPHATE
PREDNISOLONE, PREDNISOLONE

WEILL MEDCL COLL

- * WEILL MEDICAL COLLEGE CORNELL UNIV
FLUDEOXYGLUCOSE F 18, FLUDEOXYGLUCOSE F-18

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** W ****

WELLSPRING PHARM

- * WELLSPRING PHARMACEUTICAL CORP
DIBENZYLIN, PHENOXYBENZAMINE HYDROCHLORIDE
DUVOID, BETHANECHOL CHLORIDE
DYRENIUM, TRIAMTERENE

WEST WARD

- * WEST WARD INC
ERGOTAMINE TARTRATE AND CAFFEINE, CAFFEINE
- * WEST WARD PHARMACEUTICAL CORP
AMINOPHYLLINE, AMINOPHYLLINE
BUTALBITAL, ACETAMINOPHEN AND CAFFEINE, ACETAMINOPHEN
BUTALBITAL, ACETAMINOPHEN, AND CAFFEINE WITH CODEINE PHOSPHATE, ACETAMINOPHEN
BUTALBITAL, ASPIRIN AND CAFFEINE, ASPIRIN
CAPTOPRIL, CAPTOPRIL
CARISOPRODOL, CARISOPRODOL
CHLOROQUINE PHOSPHATE, CHLOROQUINE PHOSPHATE
CHLOROTHIAZIDE, CHLOROTHIAZIDE
CORTISONE ACETATE, CORTISONE ACETATE
DICYCLOMINE HYDROCHLORIDE, DICYCLOMINE HYDROCHLORIDE
DOXYCYCLINE HYCLATE, DOXYCYCLINE HYCLATE
ETHAMBUTOL HYDROCHLORIDE, ETHAMBUTOL HYDROCHLORIDE
FLURAZEPAM HYDROCHLORIDE, FLURAZEPAM HYDROCHLORIDE
FOLIC ACID, FOLIC ACID
HYDROCORTISONE, HYDROCORTISONE
ISONIAZID, ISONIAZID
ISOSORBIDE DINITRATE, ISOSORBIDE DINITRATE
ISOSORBIDE MONONITRATE, ISOSORBIDE MONONITRATE
LISINAPRIL AND HYDROCHLOROTHIAZIDE, HYDROCHLOROTHIAZIDE
LISINAPRIL, LISINAPRIL
LITHIUM CARBONATE, LITHIUM CARBONATE
METHOCARBAMOL, METHOCARBAMOL
PREDNISONE, PREDNISONE
PROPOXYPHENE HYDROCHLORIDE, PROPOXYPHENE HYDROCHLORIDE
PROPYLTHIOURACIL, PROPYLTHIOURACIL
TRIHENYPHENIDYL HYDROCHLORIDE, TRIHENYPHENIDYL HYDROCHLORIDE

WESTWARD

- * WESTWARD PHARMACEUTICAL CORP
NAPROXEN, NAPROXEN
RIFAMPIN AND ISONIAZID, ISONIAZID

WESTWOOD SQUIBB

- * WESTWOOD SQUIBB PHARMACEUTICALS INC
CAPITROL, CHLOROXINE
EURAX, CROTAMITON
EXELDERM, SULCONAZOLE NITRATE
HALOG, HALCINONIDE
LAC-HYDRIN, AMMONIUM LACTATE
MYCOSTATIN, NYSTATIN
ULTRAVATE, HALOBETASOL PROPIONATE
WESTCORT, HYDROCORTISONE VALERATE

WOCKHARDT

- * WOCKHARDT AMERICAS INC
CAPTOPRIL, CAPTOPRIL
ENALAPRIL MALEATE, ENALAPRIL MALEATE
FAMOTIDINE, FAMOTIDINE
RANITIDINE HYDROCHLORIDE, RANITIDINE HYDROCHLORIDE
- * WOCKHARDT LTD
BETHANECHOL CHLORIDE, BETHANECHOL CHLORIDE
CEFUROXIME AXETIL, CEFUROXIME AXETIL
FAMOTIDINE, FAMOTIDINE (OTC)
NIACIN, NIACIN
ZONISAMIDE, ZONISAMIDE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** W ****

WYETH CONS

* WYETH CONSUMER HEALTHCARE
ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
ADVIL COLD AND SINUS, IBUPROFEN (OTC)
ADVIL COLD AND SINUS, IBUPROFEN POTASSIUM (OTC)
ADVIL LIQUI-GELS, IBUPROFEN POTASSIUM (OTC)
ADVIL MIGRAINE LIQUI-GELS, IBUPROFEN POTASSIUM (OTC)
ADVIL PM, DIPHENHYDRAMINE CITRATE (OTC)
ADVIL PM, DIPHENHYDRAMINE HYDROCHLORIDE (OTC)
ADVIL, IBUPROFEN (OTC)
ALAVERT, LORATADINE (OTC)
AXID AR, NIZATIDINE (OTC)
BRONITIN MIST, EPINEPHRINE BITARTRATE (OTC)
CHILDREN'S ADVIL ALLERGY SINUS, CHLORPHENIRAMINE MALEATE (OTC)
CHILDREN'S ADVIL COLD, IBUPROFEN (OTC)
CHILDREN'S ADVIL, IBUPROFEN
CHILDREN'S ADVIL, IBUPROFEN (OTC)
CHILDREN'S ADVIL-FLAVORED, IBUPROFEN (OTC)
JUNIOR STRENGTH ADVIL, IBUPROFEN (OTC)
LORATADINE, LORATADINE (OTC)
ORUDIS KT, KETOPROFEN (OTC)
PEDIATRIC ADVIL, IBUPROFEN (OTC)
PRIMATENE MIST, EPINEPHRINE (OTC)

WYETH PHARMS INC

* WYETH PHARMACEUTICALS INC
ALESSE, ETHINYL ESTRADIOL
CORDARONE, AMIODARONE HYDROCHLORIDE
EFFEXOR XR, VENLAFAXINE HYDROCHLORIDE
EFFEXOR, VENLAFAXINE HYDROCHLORIDE
INDERAL LA, PROPRANOLOL HYDROCHLORIDE
INDERAL, PROPRANOLOL HYDROCHLORIDE
INDERIDE-40/25, HYDROCHLOROTHIAZIDE
LO/OVRAL-28, ETHINYL ESTRADIOL
LODINE, ETODOLAC
MINOCIN, MINOCYCLINE HYDROCHLORIDE
MYLOTARG, GEMTUZUMAB OZOGAMICIN
ORUVAIL, KETOPROFEN
PHENERGAN, PROMETHAZINE HYDROCHLORIDE
PREMARIN, ESTROGENS, CONJUGATED
PREMPHASE 14/14, ESTROGENS, CONJUGATED
PREMPRO, ESTROGENS, CONJUGATED
PROTONIX IV, PANTOPRAZOLE SODIUM
PROTONIX, PANTOPRAZOLE SODIUM
RAPAMUNE, SIROLIMUS
TRECATOR-SC, ETHIONAMIDE
TRIPHASIL-28, ETHINYL ESTRADIOL
TYGACIL, TIGECYCLINE
ZOSYN IN PLASTIC CONTAINER, PIPERACILLIN SODIUM
ZOSYN, PIPERACILLIN SODIUM

X GEN PHARMS

* X GEN PHARMACEUTICALS INC
NYSTATIN, NYSTATIN

XANODYNE PHARM

* XANODYNE PHARMACAL INC
AMICAR, AMINOCAPROIC ACID
CALCITRIOL, CALCITRIOL
DARVOCET A500, ACETAMINOPHEN
DARVOCET-N 100, ACETAMINOPHEN
DARVOCET-N 50, ACETAMINOPHEN
DARVON COMPOUND-65, ASPIRIN
DARVON, PROPOXYPHENE HYDROCHLORIDE
DARVON-N, PROPOXYPHENE NAPSYLATE

APPENDIX B - PRODUCT NAME SORTED BY APPLICANT

**** X ****

- * XANODYNE PHARMACAL INC
DOLOPHINE HYDROCHLORIDE, METHADONE HYDROCHLORIDE
DURACLON, CLONIDINE HYDROCHLORIDE
LEUCOVORIN CALCIUM, LEUCOVORIN CALCIUM
ORAMORPH SR, MORPHINE SULFATE
ROXICODONE, OXYCODONE HYDROCHLORIDE

XTTRIUM

- * XTTRIUM LABORATORIES INC
DYNA-HEX, CHLORHEXIDINE GLUCONATE (OTC)
EXIDINE, CHLORHEXIDINE GLUCONATE (OTC)

YUNG SHIN PHARM

- * YUNG SHIN PHARMACEUTICAL INDUSTRIAL CO LTD
CEPHALEXIN, CEPHALEXIN

ZAMBON

- * ZAMBON CORP
MONUROL, FOSFOMYCIN TROMETHAMINE

ZARS

- * ZARS INC
SYNERA, LIDOCAINE

ZENITH GOLDLINE

- * ZENITH GOLDLINE LABORATORIES INC
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
- * ZENITH GOLDLINE PHARMACEUTICALS
BUSPIRONE HYDROCHLORIDE, BUSPIRONE HYDROCHLORIDE
- * ZENITH GOLDLINE PHARMACEUTICALS INC
NIZATIDINE, NIZATIDINE

ZILA

- * ZILA PHARMACEUTICALS INC
PERIDEX, CHLORHEXIDINE GLUCONATE

ZLB BEHRING

- * ZLB BEHRING LLC
STIMATE, DESMOPRESSIN ACETATE

ZYDUS PHARMS USA

- * ZYDUS PHARMACEUTICALS USA INC
ATENOLOL, ATENOLOL
CLINDAMYCIN HYDROCHLORIDE, CLINDAMYCIN HYDROCHLORIDE
METFORMIN HYDROCHLORIDE, METFORMIN HYDROCHLORIDE
PROMETHAZINE HYDROCHLORIDE, PROMETHAZINE HYDROCHLORIDE
RIBAVIRIN, RIBAVIRIN

APPENDIX C

UNIFORM TERMS

DOSAGE FORMS

AEROSOL	LOTION/SHAMPOO
AEROSOL, FOAM	N/A
AEROSOL, METERED	OIL
CAPSULE	OIL/DROPS
CAPSULE, DELAYED REL PELLETS	OINTMENT
CAPSULE, DELAYED RELEASE	OINTMENT, AUGMENTED
CAPSULE, EXTENDED RELEASE	PASTE
CLOTH	PATCH
CONCENTRATE	PELLET
CREAM	POWDER
CREAM, AUGMENTED	POWDER, EXTENDED RELEASE
ELIXIR	POWDER, METERED
EMULSION	RING
ENEMA	SHAMPOO
FILM, EXTENDED RELEASE	SOLUTION
FOR SOLUTION	SOLUTION FOR SLUSH
FOR SUSPENSION	SOLUTION/DROPS
FOR SUSPENSION, DELAYED RELEASE	SOLUTION, GEL FORMING/DROPS
FOR SUSPENSION, EXTENDED RELEASE	SPONGE
GAS	SPRAY
GEL	SPRAY, METERED
GEL, AUGMENTED	SUPPOSITORY
GEL, METERED	SUSPENSION
GRANULE	SUSPENSION/DROPS
GRANULE, DELAYED RELEASE	SUSPENSION, EXTENDED RELEASE
GRANULE, EFFERVESCENT	SWAB
GUM, CHEWING	SYRUP
IMPLANT	TABLET
INHALANT	TABLET, CHEWABLE
INJECTABLE	TABLET, COATED PARTICLES
INJECTABLE, LIPID COMPLEX	TABLET, DELAYED RELEASE
INJECTABLE, LIPOSOMAL	TABLET, DELAYED RELEASE, ORALLY DISINTEGRATING
INSERT	TABLET, EFFERVESCENT
INSERT, EXTENDED RELEASE	TABLET, EXTENDED RELEASE
INTRAUTERINE DEVICE	TABLET, FOR SOLUTION
JELLY	TABLET, ORALLY DISINTEGRATING
LIQUID	TAPE
LIQUID, EXTENDED RELEASE	TROCHE/LOZENGE
LOTION	
LOTION, AUGMENTED	

NOTE: TERMS COMPRISE CURRENTLY MARKETED PRODUCTS

APPENDIX C

UNIFORM TERMS

ROUTES OF ADMINISTRATION

BUCCAL	IONTOPHORESIS
DENTAL	IRRIGATION
ENDOCERVICAL	IV (INFUSION)
EPIDURAL	IV (INFUSION)-SC
FOR RX COMPOUNDING	N/A
IM-IV	NASAL
IM-IV-SC	OPHTHALMIC
IMPLANTATION	ORAL
IM-SC	ORAL-21
INHALATION	ORAL-28
INJECTION	OTIC
INTRACRANIAL	PERFUSION, CARDIAC
INTRALYMPHATIC	PERIODONTAL
INTRAMUSCULAR	RECTAL
INTRAOCULAR	SPINAL
INTRAPERITONEAL	SUBCUTANEOUS
INTRAPLEURAL	SUBLINGUAL
INTRATHECAL	TOPICAL
INTRATRACHEAL	TRANSDERMAL
INTRAUTERINE	TRANSMUCOSAL
INTRAVENOUS	URETERAL
INTRAVESICAL	URETHRAL
INTRAUTERINE	VAGINAL

NOTE: TERMS COMPRISE CURRENTLY MARKETED PRODUCTS

ABBREVIATIONS

AMP	AMPULE	IU	INTERNATIONAL UNITS
AMPICIL	AMPICILLIN	KIU	KALLIKREIN INHIBITOR UNITS
APPROX	APPROXIMATELY	MCI	MILLICURIE
BOT	BOTTLE	MEQ	MILLIEQUIVALENT
CI	CURIE	MG	MILLIGRAM
CSR	CAROTID SINUS REFLEX	ML	MILLILITER
CU	CLINICAL UNITS	N/A	NOT APPLICABLE
DIPROP	DIPROPIONATE	PPM	PARTS PER MILLION
EQ	EQUIVALENT TO	REL	RELEASE
ELECT	ELECTROLYTE	SQ CM	SQUARE CENTIMETER
ER	EXTENDED RELEASE	U	UNITS
GM	GRAM	UCI	MICROCURIE
HBR	HYDROBROMIDE	UGM	MICROGRAM
HCL	HYDROCHLORIDE	UMOLAR	MICROMOLAR
HR	HOUR	USP	UNITED STATES
INH	INHALATION		PHARMACOPEIA

PATENT AND EXCLUSIVITY INFORMATION ADDENDUM

This *Addendum* identifies drugs that qualify under the Drug Price Competition and Patent Term Restoration Act (1984 Amendments) for periods of exclusivity, during which abbreviated new drug applications (ANDAs) and applications described in Section 505(b)(2) of the Federal Food, Drug, and Cosmetic Act (the Act) for those drug products may, in some instances, not be submitted or made effective as described below, and provides patent information concerning the listed drug products. Those drugs that have qualified for Orphan Drug Exclusivity pursuant to Section 527 of the Act and those drugs that have qualified for Pediatric Exclusivity pursuant to Section 505A are also included in this *Addendum*. This section is arranged in alphabetical order by active ingredient name followed the trade name. Active ingredient headings for multiple ingredient (combination) drug products are arranged alphabetically. For an explanation of the codes used in the *Addendum*, see the *Patent and Exclusivity Terms* Section. Exclusivity prevents the submission or effective approval of ANDAs or applications described in Section 505(b)(2) of the Act. It does not prevent the submission or approval of a second 505(b)(1) application except in the case of Orphan Drug exclusivity. Applications qualifying for periods of exclusivity are:

- (1) A new drug application approved after September 24, 1984, for a drug product all active ingredients (including any ester or salt of the active ingredient) of which had never been approved in any other new drug application under Section 505 (b) of the Act. No subsequent ANDA or application described in Section 505(b)(2) of the Act for the same drug may be *submitted* for a period of *five years* from the date of approval of the original application, except that such an application may be *submitted* after *four years* if it contains a certification that a patent claiming the drug is invalid or will not be infringed by the product for which approval is sought.
- (2) A new drug application approved after September 24, 1984, for a drug product containing an active ingredient (including any ester or salt of that active ingredient) that has been approved in an earlier new drug application and that includes reports of new clinical investigations (other than bioavailability studies). Such investigations must have been conducted or sponsored by the applicant and must have been essential to approval of the application. If these requirements are met, the approval of a subsequent ANDA or an application described in Section 505(b)(2) of the Act may not be *made effective* for the same drug or use, if for a new indication, before the expiration of *three years* from the date of approval of the original application. If an applicant has exclusivity for a new application or 505(b)(2) application for the drug product with indications or use, this does not preclude the approval of an ANDA or 505(b)(2) application not covered by the exclusivity.
- (3) A supplement to a new drug application for a drug containing a previously approved active ingredient including (any ester or salt of the active ingredient) approved after September 24, 1984, that contains reports of new clinical investigations (other than bioavailability studies) essential to the approval of the supplement and conducted or sponsored by the applicant. The approval of a subsequent ANDA or 505(b)(2) application for a change approved in the supplement may not be

made effective for three years from the date of approval of the original supplement.

The Act requires that patent information be filed with all newly submitted Section 505(b) drug applications. No NDA may be approved after September 24, 1984, without the submission of patent information to the Agency. Effective August 18, 2003, this information must be filed using FDA Form 3524a "Patent Information Submitted with the Filing of an NDA, Amendment or Supplement".

Upon approval, patent numbers and expiration dates, in addition to certain other information on appropriate patents claiming drug products that are the subject of approved applications, will be published on a daily basis in the Electronic Orange Book, <http://www.fda.gov/cder/ob/default.htm>. The Addendum and the monthly Cumulative Supplements to the annual edition will list patent information on a monthly basis.

Effective August 18, 2003 this, information must be submitted to the agency upon approval on FDA Form 3542 "Patent Information Submitted Upon and After Approval of an NDA or Supplement". Patent information on unapproved applications or on patents beyond the scope of the Act (i.e., process or manufacturing patents) will not be published. FDA form 3542 will be the only form used for the purposes of this publication.

The patents that FDA regards as covered by the statutory provisions for submission of patent information are: patents that claim the active ingredient(s); drug product patents which include formulation/composition patents; use patents for a particular approved indication or method of using the product; and certain other patents as detailed on FDA Form 3542. This information, as provided by the sponsor on FDA form 3542, will be published as described above.

Since all parts of this publication are subject to changes, additions, or deletions, the Electronic Orange Book, updated daily, should be consulted for the most recent patent and exclusivity information. The Addendum and the monthly Cumulative Supplements to the annual edition will list patent and exclusivity information on a monthly basis.

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ABACAVIR SULFATE; ZIAGEN</u>					
020977 001	5034394	Dec 18, 2011		D-40	Aug 02, 2007
	5034394*PED	Jun 18, 2012			
	5089500	Jun 26, 2009	U-248		
	5089500*PED	Dec 26, 2009			
	6294540	May 14, 2018	U-65		
	6294540*PED	Nov 14, 2018	U-65		
<u>ABACAVIR SULFATE; ZIAGEN</u>					
020978 001	5034394	Dec 18, 2011		D-40	Aug 02, 2007
	5034394*PED	Jun 18, 2012			
	5089500	Jun 26, 2009	U-248		
	5089500*PED	Dec 26, 2009			
	6294540	May 14, 2018	U-65		
	6294540*PED	Nov 14, 2018	U-65		
<u>ABACAVIR SULFATE; LAMIVUDINE; EPZICOM</u>					
021652 001	5034394	Dec 18, 2011	DS DP		
	5047407	Nov 17, 2009	DS DP	U-257	
	5089500	Jun 26, 2009		U-257	
	5905082	May 18, 2016	DS DP		
	6294540	May 14, 2018	DS DP	U-257	
	6417191	Mar 28, 2016	DP	U-257	
<u>ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE; TRIZIVIR</u>					
021205 001	4724232	Sep 17, 2005		U-248	
	4818538	Sep 17, 2005	DP		
	4828838	Sep 17, 2005	DP		
	4833130	Sep 17, 2005		U-248	
	4837208	Sep 17, 2005		U-248	
	5034394	Jun 26, 2009	DS DP		
	5034394*PED	Dec 26, 2009			
	5047407	Nov 17, 2009	DS DP	U-248	
	5047407*PED	May 17, 2010			
	5089500	Jun 26, 2009		U-248	
	5089500*PED	Dec 26, 2009		U-248	
	5905082	May 18, 2016	DS DP	U-248	
	5905082*PED	Nov 18, 2016			
	6294540	May 14, 2018	DS DP	U-65	
	6294540*PED	Nov 14, 2018		U-65	
	6417191	Mar 28, 2016	DP	U-248	
<u>ABARELIX; PLENAXIS</u>					
021320 001	5843901	Dec 01, 2015	DS DP	NCE	Nov 25, 2008
	5968895	Dec 11, 2016	DP		
	6180608	Dec 11, 2016	DP	U-549	
	6423686	Jun 07, 2015	DS		
	6455499	Jun 07, 2015		U-549	
	6699833	Dec 11, 2016	DP		
<u>ACAMPROSATE CALCIUM; CAMPRAL</u>					
021431 001				NCE	Jul 29, 2009
<u>ACARBOSE; PRECOSE</u>					
020482 001	4904769	Feb 27, 2007			
<u>ACARBOSE; PRECOSE</u>					
020482 002	4904769	Feb 27, 2007			
<u>ACARBOSE; PRECOSE</u>					
020482 004	4904769	Feb 27, 2007			
<u>ACETAMINOPHEN; TYLENOL (CAPLET)</u>					
019872 001	4820522	Jul 27, 2007			
	4968509	Nov 06, 2007			
	5004613	Jul 27, 2007			
<u>ACETAMINOPHEN; ASPIRIN; CAFFEINE; EXCEDRIN (MIGRAINE)</u>					
020802 001	4943565	Jul 24, 2007			
	5972916	Jul 14, 2017	U-296		
<u>ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE; TRAMADOL HCL AND ACETAMINOPHEN</u>					
076475 001				PC	Oct 18, 2005
<u>ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE; ULTRACET</u>					
021123 001	5336691	Aug 09, 2011			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ACETYLCHOLINE CHLORIDE; MIOCHOL-E</u>					
020213 001	6261546	Apr 29, 2019	U-506		
<u>ACETYLCYSTEINE; ACETADOTE</u>					
021539 001				ODE	Jan 23, 2011
<u>ACRIVASTINE; PSEUDOEPHEDRINE HYDROCHLORIDE; SEMPRES-D</u>					
019806 001	4650807	Mar 26, 2008	U-93		
<u>ACYCLOVIR; ZOVIRAX</u>					
021478 001	4963555	Oct 16, 2007		NDF	Dec 30, 2005
<u>ADAPALENE; DIFFERIN</u>					
020338 001	4717720	May 31, 2010			
	RE34440	May 31, 2010	U-275		
<u>ADAPALENE; DIFFERIN</u>					
020380 001	4717720	May 31, 2010			
	RE34440	May 31, 2010	U-275		
<u>ADAPALENE; DIFFERIN</u>					
020748 001	4717720	May 31, 2010			
	RE34440	May 31, 2010	U-275		
<u>ADEFOVIR DIPIVOXIL; HEPSERA</u>					
021449 001	4724233	Apr 21, 2006	U-470	NCE	Sep 20, 2007
	4808716	Apr 25, 2006			
	5663159	Sep 02, 2014	DS DP U-470		
	6451340	Jul 23, 2018	DS DP U-470		
	6635278	Dec 15, 2018	DS DP		
<u>ADENOSINE; ADENOSCAN</u>					
020059 001	5070877	May 18, 2009	U-116		
	5731296	Mar 24, 2015	U-221		
<u>ALATROFLOXACIN MESYLATE; TROVAN PRESERVATIVE FREE</u>					
020760 001	5164402	Nov 17, 2009	U-282		
	5763454	Jun 15, 2015	U-282		
	6080756	Jul 05, 2016			
	6194429	Jul 23, 2018			
<u>ALATROFLOXACIN MESYLATE; TROVAN PRESERVATIVE FREE</u>					
020760 002	5164402	Nov 17, 2009	U-282		
	5763454	Jun 15, 2015	U-282		
	6080756	Jul 05, 2016			
	6194429	Jul 23, 2018			
<u>ALBUMIN HUMAN; OPTISON</u>					
020899 001	5529766	Jun 25, 2013	U-505		
	5558094	Feb 28, 2012	U-505		
	5573751	Apr 25, 2012			
	6723303	Apr 20, 2021	DP		
<u>ALBUTEROL SULFATE; ACCUNEB</u>					
020949 001	6702997	Dec 28, 2021	U-558		
<u>ALBUTEROL SULFATE; ACCUNEB</u>					
020949 002	6702997	Dec 28, 2021	U-558		
<u>ALBUTEROL SULFATE; PROAIR HFA</u>					
021457 001	5605674	Feb 25, 2014	DP	NP	Oct 29, 2007
	5695743	Jul 06, 2010	DP		
	5766573	Nov 28, 2009	U-491		
	6352684	Nov 28, 2009	DP	U-356	
<u>ALBUTEROL SULFATE; PROVENTIL-HFA</u>					
020503 001	5225183	Jul 06, 2010			
	5439670	Jul 06, 2010			
	5605674	Feb 25, 2014			
	5695743	Jul 06, 2010	U-491		
	5766573	Jun 16, 2015			
	6352684	Nov 28, 2009			
<u>ALBUTEROL SULFATE; VENTOLIN HFA</u>					
020983 001	6131566	Apr 14, 2015	DP	U-589	
	6315173	Dec 23, 2017	DP		
	6510969	Dec 23, 2017	DP		
	6532955	Apr 14, 2015	DP	U-590	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ALBUTEROL SULFATE; VOLMAX</u>					
019604 001	4777049	Oct 11, 2005			
<u>ALBUTEROL SULFATE; VOLMAX</u>					
019604 002	4777049	Oct 11, 2005			
<u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE; COMBIVENT</u>					
020291 001	5603918	Jun 09, 2015			
<u>ALBUTEROL SULFATE; IPRATROPIUM BROMIDE; DUONEB</u>					
020950 001	6632842	Dec 28, 2021	U-532		
<u>ALCOHOL; CHLORHEXIDINE GLUCONATE; AVAGARD</u>					
021074 001	5897031	Jun 21, 2016			
<u>ALENDRONATE SODIUM; FOSAMAX</u>					
020560 001	4621077	Aug 06, 2007	U-114	M-51	Dec 21, 2008
	4621077*PED	Feb 06, 2008	U-114		
	5358941	Dec 02, 2012			
	5358941*PED	Jun 02, 2013			
	5681590	Dec 02, 2012			
	5681590*PED	Jun 02, 2013			
	5849726	Jun 06, 2015			
	5849726*PED	Dec 06, 2015			
	6008207	Jun 06, 2015	U-303		
	6008207*PED	Dec 06, 2015	U-303		
	6090410	Dec 02, 2012			
	6090410*PED	Jun 02, 2013			
	6194004	Dec 02, 2012			
	6194004*PED	Jun 02, 2013			
<u>ALENDRONATE SODIUM; FOSAMAX</u>					
020560 002	4621077	Aug 06, 2007	U-114	M-51	Dec 21, 2008
	4621077*PED	Feb 06, 2008	U-114		
	5358941	Dec 02, 2012			
	5358941*PED	Jun 02, 2013			
	5681590	Dec 02, 2012			
	5681590*PED	Jun 02, 2013			
	5849726	Jun 06, 2015			
	5849726*PED	Dec 06, 2015			
	6008207	Jun 06, 2015	U-303		
	6008207*PED	Jun 06, 2015	U-303		
	6090410	Dec 02, 2012			
	6090410*PED	Jun 02, 2013			
<u>ALENDRONATE SODIUM; FOSAMAX</u>					
020560 003	4621077	Aug 06, 2007	U-114	M-51	Dec 21, 2008
	4621077*PED	Feb 06, 2008	U-114		
	5358941	Dec 02, 2012			
	5358941*PED	Jun 02, 2013			
	5681590	Dec 02, 2012			
	5681590*PED	Jun 02, 2013			
	5849726	Jun 06, 2015			
	5849726*PED	Dec 06, 2015			
	6008207	Jun 06, 2015	U-303		
	6008207*PED	Dec 06, 2015	U-303		
	6090410	Dec 02, 2012			
	6090410*PED	Jun 02, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ALENDRONATE SODIUM; FOSAMAX</u>					
020560 004	4621077	Aug 06, 2007	U-114	M-51	Dec 21, 2008
	4621077*PED	Feb 06, 2008	U-114		
	5358941	Dec 02, 2012			
	5358941*PED	Jun 02, 2013			
	5681590	Dec 02, 2012			
	5681590*PED	Jun 02, 2013			
	5849726	Jun 06, 2015			
	5849726*PED	Dec 16, 2015			
	5994329	Jul 17, 2018			
	5994329*PED	Jan 17, 2019			
	6008207	Jun 06, 2015			
	6008207*PED	Dec 06, 2015			
	6015801	Jul 17, 2018	U-353		
	6015801*PED	Jan 17, 2019	U-353		
	6090410	Dec 02, 2012			
	6090410*PED	Jun 02, 2013			
	6225294	Jul 17, 2018			
	6225294*PED	Jan 17, 2019			
<u>ALENDRONATE SODIUM; FOSAMAX</u>					
020560 005	4621077	Aug 06, 2007	U-114	D-87	Apr 16, 2007
	4621077*PED	Feb 06, 2008	U-114	M-51	Dec 21, 2008
	5358941	Dec 02, 2012			
	5358941*PED	Jun 02, 2013			
	5681590	Dec 02, 2012			
	5681590*PED	Jun 02, 2013			
	5849726	Jun 06, 2015			
	5849726*PED	Dec 06, 2015			
	5994329	Jul 17, 2018			
	5994329*PED	Jan 17, 2019			
	6008207	Jun 06, 2015			
	6008207*PED	Dec 06, 2015			
	6015801	Jul 17, 2018	U-353		
	6015801*PED	Jan 17, 2019	U-353		
	6090410	Dec 02, 2012			
	6090410*PED	Jun 02, 2013			
	6225294	Jul 17, 2018			
	6225294*PED	Jan 17, 2019			
<u>ALENDRONATE SODIUM; FOSAMAX</u>					
021575 001	4621077	Aug 06, 2007		D-87	Apr 16, 2007
	4621077*PED	Feb 06, 2008		M-51	Dec 21, 2008
	5462932	May 17, 2014			
	5994329	Jul 17, 2018			
	5994329*PED	Jan 17, 2019			
	6015801	Jul 17, 2018			
	6015801*PED	Jan 17, 2019			
	6225294	Jul 17, 2018			
	6225294*PED	Jan 17, 2019			
<u>ALENDRONATE SODIUM; CHOLECALCIFEROL; FOSAMAX PLUS D</u>					
021762 001	4621077	Aug 06, 2007	U-648		
	4621077*PED	Feb 06, 2008			
	5358941	Dec 02, 2012	DP		
	5358941*PED	Jun 02, 2013			
	5681590	Dec 02, 2012	DP		
	5681590*PED	Jun 02, 2013			
	5994329	Jul 17, 2018	U-647		
	5994329*PED	Jan 17, 2019			
	6090410	Dec 02, 2012	DP		
	6090410*PED	Jun 02, 2013	DP		
<u>ALFUZOSIN HYDROCHLORIDE; UROXATRAL</u>					
021287 001	4661491	May 27, 2006		NCE	Jun 12, 2008
	6149940	Aug 22, 2017			
<u>ALITRETINOIN; PANRETIN</u>					
020886 001	5932622	Aug 03, 2016	U-562	ODE	Feb 02, 2006
<u>ALMOTRIPTAN MALATE; AXERT</u>					
021001 001	5565447	May 07, 2015		NCE	May 07, 2006
<u>ALMOTRIPTAN MALATE; AXERT</u>					
021001 002	5565447	May 07, 2015		NCE	May 07, 2006

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ALOSETRON HYDROCHLORIDE; LOTRONEX</u>					
021107 001	5360800	Jan 13, 2013	DS DP U-405		
<u>ALOSETRON HYDROCHLORIDE; LOTRONEX</u>					
021107 002	5360800	Jan 13, 2013	DS DP U-405		
<u>ALPRAZOLAM; NIRAVAM</u>					
021726 001	6024981	Apr 09, 2018	DP		
	6221392	Apr 09, 2018	DP		
<u>ALPRAZOLAM; NIRAVAM</u>					
021726 002	6024981	Apr 09, 2018	DP		
	6221392	Apr 09, 2018	DP		
<u>ALPRAZOLAM; NIRAVAM</u>					
021726 003	6024981	Apr 09, 2018	DP		
	6221392	Apr 09, 2018	DP		
<u>ALPRAZOLAM; NIRAVAM</u>					
021726 004	6024981	Apr 09, 2018	DP		
	6221392	Apr 09, 2018	DP		
<u>ALPRAZOLAM; XANAX</u>					
018276 004	5061494	Dec 12, 2008			
<u>ALPRAZOLAM; XANAX XR</u>					
021434 001				NDF	Jan 17, 2006
<u>ALPRAZOLAM; XANAX XR</u>					
021434 002				NDF	Jan 17, 2006
<u>ALPRAZOLAM; XANAX XR</u>					
021434 003				NDF	Jan 17, 2006
<u>ALPRAZOLAM; XANAX XR</u>					
021434 004				NDF	Jan 17, 2006
<u>ALPROSTADIL; CAVERJECT</u>					
020379 001	5741523	Apr 21, 2015			
<u>ALPROSTADIL; CAVERJECT</u>					
020379 002	5741523	Apr 21, 2015			
<u>ALPROSTADIL; CAVERJECT</u>					
020379 003	5741523	Apr 21, 2015			
<u>ALPROSTADIL; CAVERJECT</u>					
020379 004	5741523	Apr 21, 2015			
<u>ALPROSTADIL; MUSE</u>					
020700 001	4801587	Jan 31, 2007	U-155		
	5242391	Sep 07, 2010	U-155		
	5474535	Dec 12, 2012	U-155		
<u>ALPROSTADIL; MUSE</u>					
020700 002	4801587	Jan 31, 2007	U-155		
	5242391	Sep 07, 2010	U-155		
	5474535	Dec 12, 2012	U-155		
<u>ALPROSTADIL; MUSE</u>					
020700 003	4801587	Jan 31, 2007	U-155		
	5242391	Sep 07, 2010	U-155		
	5474535	Dec 12, 2012	U-155		
<u>ALPROSTADIL; MUSE</u>					
020700 004	4801587	Jan 31, 2007	U-155		
	5242391	Sep 07, 2010	U-155		
	5474535	Dec 12, 2012	U-155		
<u>AMIFOSTINE; ETHYOL</u>					
020221 001	5424471	Jul 31, 2012		ODE	Jun 24, 2006
	5591731	Jul 31, 2012			
	5994409	Dec 08, 2017	U-305		
<u>AMIFOSTINE; ETHYOL</u>					
020221 002	5424471	Jul 31, 2012		ODE	Jun 24, 2006
	5591731	Jul 31, 2012			
	5994409	Dec 08, 2017	U-305		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
AMINOLEVULINIC ACID HYDROCHLORIDE; LEVULAN						
020965 001	5079262	Jul 28, 2009	U-289			
	5211938	May 18, 2010		U-289		
	5422093	Jul 28, 2009		U-289		
	5954703	Oct 31, 2017		U-289		
	6710066	Jul 28, 2009		U-289		
AMLEXANOX; AMLEXANOX						
021727 001				NDF	Sep 29, 2007	
AMLEXANOX; APHTHASOL						
020511 001	5362737	Nov 08, 2011	U-243			
AMLODIPINE BESYLATE; NORVASC						
019787 001	4572909	Jul 31, 2006	DS DP	U-683	I-472	Sep 28, 2008
	4572909*PED	Jan 31, 2007				NPP
	4879303	Mar 25, 2007	DS DP		PED	Jul 08, 2007
	4879303*PED	Sep 25, 2007				
AMLODIPINE BESYLATE; NORVASC						
019787 002	4572909	Jul 31, 2006	DS DP	U-683	I-472	Sep 28, 2008
	4572909*PED	Jan 31, 2007				NPP
	4879303	Mar 25, 2007	DS DP		PED	Jul 08, 2007
	4879303*PED	Sep 25, 2007				
AMLODIPINE BESYLATE; NORVASC						
019787 003	4572909	Jul 31, 2006	DS DP	U-683	I-472	Sep 28, 2008
	4572909*PED	Jan 31, 2007				NPP
	4879303	Mar 25, 2007	DS DP		PED	Jul 08, 2007
	4879303*PED	Sep 25, 2007				
AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET						
021540 001	4572909	Jul 31, 2006	DS DP	U-3	NC	Jan 30, 2007
	4572909*PED	Jan 31, 2007				
	4681893	Sep 24, 2009	DS DP	U-161		
	4681893*PED	Mar 24, 2010				
	4879303	Mar 25, 2007	DS DP			
	4879303*PED	Sep 25, 2007				
	5273995	Dec 28, 2010	DS DP	U-162		
	5273995*PED	Jun 28, 2011				
	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018		U-552		
AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET						
021540 002	4572909	Jul 31, 2006	DS DP	U-3	NC	Jan 30, 2007
	4572909*PED	Jan 31, 2007				
	4681893	Sep 24, 2009	DS DP	U-161		
	4681893*PED	Mar 24, 2010				
	4879303	Mar 25, 2007	DS DP			
	4879303*PED	Sep 25, 2007				
	5273995	Dec 28, 2010	DS DP	U-162		
	5273995*PED	Jun 28, 2011				
	5686104	Nov 11, 2014		DP	U-213	
	5686104*PED	May 11, 2015				
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
	6126971*PED	Jul 19, 2013				
	6455574	Aug 11, 2018		U-552		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET</u>					
021540 003	4572909	Jul 31, 2006	DS DP U-3	NC	Jan 30, 2007
	4572909*PED	Jan 31, 2007			
	4681893	Sep 24, 2009	DS DP U-161		
	4681893*PED	Mar 24, 2010			
	4879303	Mar 25, 2007	DS DP		
	4879303*PED	Sep 25, 2007			
	5273995	Dec 28, 2010	DS DP U-162		
	5273995*PED	Jun 28, 2011			
	5686104	Nov 11, 2014	DP U-213		
	5686104*PED	May 11, 2015			
	5969156	Jul 08, 2016	DS		
	5969156*PED	Jan 08, 2017			
	6126971	Jan 19, 2013	DP		
	6126971*PED	Jul 19, 2013			
	6455574	Aug 11, 2018	U-552		
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET</u>					
021540 004	4572909	Jul 31, 2006	DS DP U-3	NC	Jan 30, 2007
	4572909*PED	Jan 31, 2007			
	4681893	Sep 24, 2009	DS DP U-161		
	4681893*PED	Mar 24, 2010			
	4879303	Mar 25, 2007	DS DP		
	4879303*PED	Sep 25, 2007			
	5273995	Dec 28, 2010	DS DP U-162		
	5273995*PED	Jun 28, 2011			
	5686104	Nov 11, 2014	DP U-213		
	5686104*PED	May 11, 2015			
	5969156	Jul 08, 2016	DS		
	5969156*PED	Jan 08, 2017			
	6126971	Jan 19, 2013	DP		
	6126971*PED	Jul 19, 2013			
	6455574	Aug 11, 2018	U-552		
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET</u>					
021540 005	4572909	Jul 31, 2006	DS DP U-3	NC	Jan 30, 2007
	4572909*PED	Jan 31, 2007			
	4681893	Sep 24, 2009	DS DP U-161		
	4681893*PED	Mar 24, 2010			
	4879303	Mar 25, 2007	DS DP		
	4879303*PED	Sep 25, 2007			
	5273995	Dec 28, 2010	DS DP U-162		
	5273995*PED	Jun 28, 2011			
	5686104	Nov 11, 2014	DP U-213		
	5686104*PED	May 11, 2015			
	5969156	Jul 08, 2016	DS		
	5969156*PED	Jan 08, 2017			
	6126971	Jan 19, 2013	DP		
	6126971*PED	Jul 19, 2013			
	6455574	Aug 11, 2018	U-552		
<u>AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET</u>					
021540 006	4572909	Jul 31, 2006	DS DP U-3	NC	Jan 30, 2007
	4572909*PED	Jan 31, 2007			
	4681893	Sep 24, 2009	DS DP U-161		
	4681893*PED	Mar 24, 2010			
	4879303	Mar 25, 2007	DS DP		
	4879303*PED	Sep 25, 2007			
	5273995	Dec 28, 2010	DS DP U-162		
	5273995*PED	Jun 28, 2011			
	5686104	Nov 11, 2014	DP U-213		
	5686104*PED	May 11, 2015			
	5969156	Jul 08, 2016	DS		
	5969156*PED	Jan 08, 2017			
	6126971	Jan 19, 2013	DP		
	6126971*PED	Jul 19, 2013			
	6455574	Aug 11, 2018	U-552		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET									
021540 007	4572909	Jul 31, 2006	DS	DP	U-3	NC	Jan 30, 2007		
	4572909*PED	Jan 31, 2007							
	4681893	Sep 24, 2009	DS	DP	U-161				
	4681893*PED	Mar 24, 2010							
	4879303	Mar 25, 2007	DS	DP					
	4879303*PED	Sep 25, 2007							
	5273995	Dec 28, 2010	DS	DP	U-162				
	5273995*PED	Jun 28, 2011							
	5686104	Nov 11, 2014		DP	U-213				
	5686104*PED	May 11, 2015							
	5969156	Jul 08, 2016	DS						
	5969156*PED	Jan 08, 2017							
	6126971	Jan 19, 2013		DP					
	6126971*PED	Jul 19, 2013							
	6455574	Aug 11, 2018			U-552				
AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET									
021540 008	4572909	Jul 31, 2006	DS	DP	U-3	NC	Jan 30, 2007		
	4572909*PED	Jan 31, 2007							
	4681893	Sep 24, 2009	DS	DP	U-161				
	4681893*PED	Mar 24, 2010							
	4879303	Mar 25, 2007	DS	DP					
	4879303*PED	Sep 25, 2007							
	5273995	Dec 28, 2010	DS	DP	U-162				
	5273995*PED	Jun 28, 2011							
	5686104	Nov 11, 2014		DP	U-213				
	5686104*PED	May 11, 2015							
	5969156	Jul 08, 2016	DS						
	5969156*PED	Jan 08, 2017							
	6126971	Jan 19, 2013		DP					
	6126971*PED	Jul 19, 2013							
	6455574	Aug 11, 2018			U-552				
AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET									
021540 009	4572909	Jul 31, 2006	DS	DP	U-3	NC	Jan 30, 2007		
	4572909*PED	Jan 31, 2007							
	4681893	Sep 24, 2009	DS	DP	U-161				
	4681893*PED	Mar 24, 2010							
	4879303	Mar 25, 2007	DS	DP					
	4879303*PED	Sep 25, 2007							
	5273995	Dec 28, 2010	DS	DP	U-162				
	5273995*PED	Jun 28, 2011							
	5686104	Nov 11, 2014		DP	U-213				
	5686104*PED	May 11, 2015							
	5969156	Jul 08, 2016	DS						
	5969156*PED	Jan 08, 2017							
	6126971	Jan 19, 2013		DP					
	6126971*PED	Jul 19, 2013							
	6455574	Aug 11, 2018			U-552				
AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET									
021540 010	4572909	Jul 31, 2006	DS	DP	U-3	NC	Jan 30, 2007		
	4572909*PED	Jan 31, 2007							
	4681893	Sep 24, 2009	DS	DP	U-161				
	4681893*PED	Mar 24, 2010							
	4879303	Mar 25, 2007	DS	DP					
	4879303*PED	Sep 25, 2007							
	5273995	Dec 28, 2010	DS	DP	U-162				
	5273995*PED	Jun 28, 2011							
	5686104	Nov 11, 2014		DP	U-213				
	5686104*PED	May 11, 2015							
	5969156	Jul 08, 2016	DS						
	5969156*PED	Jan 08, 2017							
	6126971	Jan 19, 2013		DP					
	6126971*PED	Jul 19, 2013							
	6455574	Aug 11, 2018			U-552				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
AMLODIPINE BESYLATE; ATORVASTATIN CALCIUM; CADUET					
021540 011	4572909	Jul 31, 2006	DS DP U-3	NC	Jan 30, 2007
	4572909*PED	Jan 31, 2007			
	4681893	Sep 24, 2009	DS DP U-161		
	4681893*PED	Mar 24, 2010			
	4879303	Mar 25, 2007	DS DP		
	4879303*PED	Sep 25, 2007			
	5273995	Dec 28, 2010	DS DP U-162		
	5273995*PED	Jun 28, 2011			
	5686104	Nov 11, 2014	DP U-213		
	5686104*PED	May 11, 2015			
	5969156	Jul 08, 2016	DS		
	5969156*PED	Jan 08, 2017			
	6126971	Jan 19, 2013	DP		
	6126971*PED	Jul 19, 2013			
6455574	Aug 11, 2018		U-552		
AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE; LOTREL					
020364 002	4572909	Jul 31, 2006			
	4879303	Mar 25, 2007			
	6162802	Dec 19, 2017			
				U-367	
AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE; LOTREL					
020364 003	4572909	Jul 31, 2006			
	4879303	Mar 25, 2007			
	6162802	Dec 19, 2017			
				U-367	
AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE; LOTREL					
020364 004	4572909	Jul 31, 2006			
	4879303	Mar 25, 2007			
	6162802	Dec 19, 2017			
				U-367	
AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE; LOTREL					
020364 005	4572909	Jul 31, 2006		PED	Dec 20, 2005
	4879303	Mar 25, 2007			
	6162802	Dec 19, 2017			
				U-367	
AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL 10					
011522 007	6384020	Jul 06, 2020			
	6384020*PED	Jan 06, 2021			
AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL 12.5					
011522 012	6384020	Jul 06, 2020			
	6384020*PED	Jan 06, 2021			
AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL 15					
011522 013	6384020	Jul 06, 2020			
	6384020*PED	Jan 06, 2021			
AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL 20					
011522 008	6384020	Jul 06, 2020			
	6384020*PED	Jan 06, 2021			
AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL 30					
011522 010	6384020	Jul 06, 2020			
	6384020*PED	Jan 06, 2021			
AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL 5					
011522 009	6384020	Jul 06, 2020			
	6384020*PED	Jan 06, 2021			
AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL 7.5					
011522 011	6384020	Jul 06, 2020			
	6384020*PED	Jan 06, 2021			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL XR 10</u>					
021303 001	6322819	Oct 21, 2018		NPP	Jul 21, 2008
	6322819*PED	Apr 21, 2019		NPP	Aug 11, 2007
	6605300	Oct 21, 2018		PED	Jan 21, 2009
	6605300*PED	Apr 21, 2019			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL XR 15</u>					
021303 006	6322819	Oct 21, 2018		NPP	Jul 21, 2008
	6322819*PED	Apr 21, 2019		NPP	Aug 11, 2007
	6605300	Oct 21, 2018		PED	Jan 21, 2009
	6605300*PED	Apr 21, 2019			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL XR 20</u>					
021303 002	6322819	Oct 21, 2018		NPP	Jul 21, 2008
	6322819*PED	Apr 21, 2019		NPP	Aug 11, 2007
	6605300	Oct 21, 2018		PED	Jan 21, 2009
	6605300*PED	Apr 21, 2019			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL XR 25</u>					
021303 004	6322819	Oct 21, 2018		NPP	Jul 21, 2008
	6322819*PED	Apr 21, 2019		NPP	Aug 11, 2007
	6605300	Oct 21, 2018		PED	Jan 21, 2009
	6605300*PED	Apr 21, 2019			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL XR 30</u>					
021303 003	6322819	Oct 21, 2018		NPP	Jul 21, 2008
	6322819*PED	Apr 21, 2019		NPP	Aug 11, 2007
	6605300	Oct 21, 2018		PED	Jan 21, 2009
	6605300*PED	Apr 21, 2019			
<u>AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE; ADDERALL XR 5</u>					
021303 005	6322819	Oct 21, 2018		NPP	Jul 21, 2008
	6322819*PED	Apr 21, 2019		NPP	Aug 11, 2007
	6605300	Oct 21, 2018		PED	Jan 21, 2009
	6605300*PED	Apr 21, 2019			
<u>AMPRENAVIR; AGENERASE</u>					
021007 001	5585397	Dec 17, 2013			
	5646180	Jul 08, 2014		U-257	
	5723490	Mar 03, 2015		U-257	
	6730679	Nov 11, 2017	DP		
<u>AMPRENAVIR; AGENERASE</u>					
021007 002	5585397	Dec 17, 2013			
	5646180	Jul 08, 2014		U-257	
	5723490	Mar 03, 2015		U-257	
	6730679	Nov 11, 2017	DP		
<u>AMPRENAVIR; AGENERASE</u>					
021039 001	5585397	Dec 17, 2013			
	5646180	Jul 08, 2014		U-257	
	5723490	Mar 03, 2015		U-257	
<u>ANAGRELIDE HYDROCHLORIDE; AGRYLIN</u>					
020333 001				M-39	Dec 10, 2007
				PED	Jun 10, 2008
<u>ANAGRELIDE HYDROCHLORIDE; AGRYLIN</u>					
020333 002				M-39	Dec 10, 2007
				PED	Jun 10, 2008
<u>ANASTROZOLE; ARIMIDEX</u>					
020541 001	RE36617	Dec 27, 2009		I-363	Sep 05, 2005
<u>APOMORPHINE HYDROCHLORIDE; APOKYN</u>					
021264 001				NCE	Apr 20, 2009
				ODE	Apr 20, 2011

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>APOMORPHINE HYDROCHLORIDE; APOKYN</u>					
021264	002			NCE	Apr 20, 2009
				ODE	Apr 20, 2011
<u>APRACLONIDINE HYDROCHLORIDE; IOPIDINE</u>					
019779	001	5212196	May 18, 2010	U-120	
<u>APREPITANT; EMEND</u>					
021549	001	5538982	Jul 23, 2013	I-475	Oct 28, 2008
		5719147	Jun 29, 2012	NCE	Mar 26, 2008
		6048859	Jun 29, 2012		
		6096742	Jul 01, 2018		
		6235735	Jun 29, 2012		
<u>APREPITANT; EMEND</u>					
021549	002	5538982	Jul 23, 2013	I-475	Oct 28, 2008
		5719147	Jun 29, 2012	NCE	Mar 26, 2008
		6048859	Jun 29, 2012		
		6096742	Jul 01, 2018		
		6235735	Jun 29, 2012		
<u>ARBUTAMINE HYDROCHLORIDE; GENESA</u>					
020420	001	5108363	Apr 28, 2009	U-220	
		5234404	Aug 10, 2010	U-220	
		5395970	Mar 07, 2012		
<u>ARGATROBAN; ARGATROBAN</u>					
020883	001	5214052	Jun 30, 2014		
<u>ARIPIRAZOLE; ABILIFY</u>					
021436	001	5006528	Oct 20, 2014	DS DP	I-437 Sep 29, 2007
					I-401 Aug 28, 2006
				NCE	Nov 15, 2007
<u>ARIPIRAZOLE; ABILIFY</u>					
021436	002	5006528	Oct 20, 2014	DS DP	I-437 Sep 29, 2007
					I-401 Aug 28, 2006
				NCE	Nov 15, 2007
<u>ARIPIRAZOLE; ABILIFY</u>					
021436	003	5006528	Oct 20, 2014	DS DP	I-437 Sep 29, 2007
					I-401 Aug 28, 2006
				NCE	Nov 15, 2007
<u>ARIPIRAZOLE; ABILIFY</u>					
021436	004	5006528	Oct 20, 2014	DS DP	I-437 Sep 29, 2007
					I-401 Aug 28, 2006
				NCE	Nov 15, 2007
<u>ARIPIRAZOLE; ABILIFY</u>					
021436	005	5006528	Oct 20, 2014	DS DP	I-437 Sep 29, 2007
					I-401 Aug 28, 2006
				NCE	Nov 15, 2007
<u>ARIPIRAZOLE; ABILIFY</u>					
021436	006	5006528	Oct 20, 2014	DS DP	I-437 Sep 29, 2007
					I-401 Aug 28, 2006
				NCE	Nov 15, 2007
<u>ARIPIRAZOLE; ABILIFY</u>					
021713	001	5006528	Oct 20, 2014	DS DP	I-437 Sep 29, 2007
					I-401 Aug 28, 2006
				NCE	Nov 15, 2007
<u>ARSENIC TRIOXIDE; TRISENOX</u>					
021248	001	6723351	Nov 10, 2018	U-573	NCE Sep 25, 2005
		6855339	Nov 10, 2018	U-617	ODE Sep 25, 2007
		6861076	Nov 10, 2018	U-617	
		6884439	Nov 10, 2018	U-651	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE HYDROCHLORIDE; VITAMIN A; VITAMIN E; M.V.I.-12 (WITHOUT VITAMIN K)</u>					
008809 006				ODE	Sep 09, 2011
<u>ASPIRIN; DIPYRIDAMOLE; AGGRENOLX</u>					
020884 001	6015577	Jan 18, 2017	U-302		
<u>ASPIRIN; PRAVASTATIN SODIUM; PRAVIGARD PAC (COPACKAGED)</u>					
021387 001	4346227	Oct 20, 2005			
	4346227*PED	Apr 20, 2006			
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM; PRAVIGARD PAC (COPACKAGED)</u>					
021387 002	4346227	Oct 20, 2005			
	4346227*PED	Apr 20, 2006			
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM; PRAVIGARD PAC (COPACKAGED)</u>					
021387 003	4346227	Oct 20, 2005			
	4346227*PED	Apr 20, 2006			
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM; PRAVIGARD PAC (COPACKAGED)</u>					
021387 004	4346227	Oct 20, 2005			
	4346227*PED	Apr 20, 2006			
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM; PRAVIGARD PAC (COPACKAGED)</u>					
021387 005	4346227	Oct 20, 2005			
	4346227*PED	Apr 20, 2006			
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>ASPIRIN; PRAVASTATIN SODIUM; PRAVIGARD PAC (COPACKAGED)</u>					
021387 006	4346227	Oct 20, 2005			
	4346227*PED	Apr 20, 2006			
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>ATAZANAVIR SULFATE; REYATAZ</u>					
021567 001	5849911	Apr 09, 2017	D-89		Jul 06, 2007
	6087383	Dec 21, 2018	NCE		Jun 20, 2008
<u>ATAZANAVIR SULFATE; REYATAZ</u>					
021567 002	5849911	Apr 09, 2017	D-89		Jul 06, 2007
	6087383	Dec 21, 2018	NCE		Jun 20, 2008

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ATAZANAVIR SULFATE; REYATAZ</u>					
021567 003	5849911	Apr 09, 2017		D-89	Jul 06, 2007
	6087383	Dec 21, 2018		NCE	Jun 20, 2008
<u>ATOMOXETINE HYDROCHLORIDE; STRATTERA</u>					
021411 001	5658590	Jan 11, 2015	U-494	NCE	Nov 26, 2007
	5658590*PED	Jul 11, 2015	U-494	PED	May 26, 2008
<u>ATOMOXETINE HYDROCHLORIDE; STRATTERA</u>					
021411 002	5658590	Jan 11, 2015	U-494	NCE	Nov 26, 2007
	5658590*PED	Jul 11, 2015	U-494	PED	May 26, 2008
<u>ATOMOXETINE HYDROCHLORIDE; STRATTERA</u>					
021411 003	5658590	Jan 11, 2015	U-494	NCE	Nov 26, 2007
	5658590*PED	Jul 11, 2015	U-494	PED	May 26, 2008
<u>ATOMOXETINE HYDROCHLORIDE; STRATTERA</u>					
021411 004	5658590	Jan 11, 2015	U-494	NCE	Nov 26, 2007
	5658590*PED	Jul 11, 2015	U-494	PED	May 26, 2008
<u>ATOMOXETINE HYDROCHLORIDE; STRATTERA</u>					
021411 005	5658590	Jan 11, 2015	U-494	NCE	Nov 26, 2007
	5658590*PED	Jul 11, 2015	U-494	PED	May 26, 2008
<u>ATOMOXETINE HYDROCHLORIDE; STRATTERA</u>					
021411 006	5658590	Jan 11, 2015	U-494	NCE	Nov 26, 2007
	5658590*PED	Jul 11, 2015	U-494	PED	May 26, 2008
<u>ATOMOXETINE HYDROCHLORIDE; STRATTERA</u>					
021411 007	5658590	Jan 11, 2015	U-494	NCE	Nov 26, 2007
	5658590*PED	Jul 11, 2015		PED	May 26, 2008
<u>ATOMOXETINE HYDROCHLORIDE; STRATTERA</u>					
021411 008	5658590	Jan 11, 2015	U-494	NCE	Nov 26, 2007
	5658590*PED	Jul 11, 2015		PED	May 26, 2008
<u>ATORVASTATIN CALCIUM; LIPITOR</u>					
020702 001	4681893	Sep 24, 2009	DS DP	U-161	I-471 Sep 21, 2008
	4681893*PED	Mar 24, 2010		U-161	I-434 Jul 30, 2007
	5273995	Dec 28, 2010	DS DP	U-162	I-350 Oct 18, 2005
	5273995*PED	Jun 28, 2011		U-162	M-36 Jul 30, 2007
	5686104	Nov 11, 2014		DP U-213	PED Apr 18, 2006
	5686104*PED	May 11, 2015		U-213	
	5969156	Jul 08, 2016	DS		
	5969156*PED	Jan 08, 2017			
	6126971	Jan 19, 2013		DP	
	6126971*PED	Jul 19, 2013			
<u>ATORVASTATIN CALCIUM; LIPITOR</u>					
020702 002	4681893	Sep 24, 2009	DS DP	U-161	I-471 Sep 21, 2008
	4681893*PED	Mar 24, 2010		U-161	I-434 Jul 30, 2007
	5273995	Dec 28, 2010	DS DP	U-162	I-350 Oct 18, 2005
	5273995*PED	Jun 28, 2011		U-162	M-36 Jul 30, 2007
	5686104	Nov 11, 2014		DP U-213	PED Apr 18, 2006
	5686104*PED	May 11, 2015		U-213	
	5969156	Jul 08, 2016	DS		
	5969156*PED	Jan 08, 2017			
	6126971	Jan 19, 2013		DP	
	6126971*PED	Jul 19, 2013			
<u>ATORVASTATIN CALCIUM; LIPITOR</u>					
020702 003	4681893	Sep 24, 2009	DS DP	U-161	I-471 Sep 21, 2008
	4681893*PED	Mar 24, 2010		U-161	I-434 Jul 30, 2007
	5273995	Dec 28, 2010	DS DP	U-162	I-350 Oct 18, 2005
	5273995*PED	Jun 28, 2011		U-162	M-36 Jul 30, 2007
	5686104	Nov 11, 2014		DP U-213	PED Apr 18, 2006
	5686104*PED	May 11, 2015		U-213	
	5969156	Jul 08, 2016	DS		
	5969156*PED	Jan 08, 2017			
	6126971	Jan 19, 2013		DP	
	6126971*PED	Jul 19, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
ATORVASTATIN CALCIUM; LIPITOR						
020702 004	4681893	Sep 24, 2009	DS DP	U-161	I-471	Sep 21, 2008
	4681893*PED	Mar 24, 2010		U-161	I-434	Jul 30, 2007
	5273995	Dec 28, 2010	DS DP	U-162	I-350	Oct 18, 2005
	5273995*PED	Jun 28, 2011		U-162		
	5686104	Nov 11, 2014		DP U-213	M-36	Jul 30, 2007
	5686104*PED	May 11, 2015		U-213	PED	Apr 18, 2006
	5969156	Jul 08, 2016	DS			
	5969156*PED	Jan 08, 2017				
	6126971	Jan 19, 2013		DP		
6126971*PED	Jul 19, 2013					
ATOVAQUONE; MEPRON						
020259 001	4981874	Aug 15, 2009		U-69		
	4981874*PED	Feb 15, 2010		U-69		
	5053432	Oct 01, 2008				
	5053432*PED	Apr 01, 2009				
ATOVAQUONE; MEPRON						
020500 001	4981874	Aug 15, 2009		U-69	ODE	Jan 05, 2006
	4981874*PED	Feb 15, 2010		U-69	PED	Jul 05, 2006
	5053432	Oct 01, 2008				
	5053432*PED	Apr 01, 2009				
	6649659	Jul 10, 2016	DS DP	U-69		
	6649659*PED	Jan 10, 2017				
ATOVAQUONE; PROGUANIL HYDROCHLORIDE; MALARONE						
021078 001	5053432	Oct 01, 2008				
	5053432*PED	Apr 01, 2009				
	6166046	Nov 25, 2013		U-406		
	6166046*PED	May 25, 2014				
	6291488	Nov 25, 2013		U-406		
	6291488*PED	May 25, 2014				
ATOVAQUONE; PROGUANIL HYDROCHLORIDE; MALARONE PEDIATRIC						
021078 002	5053432	Oct 01, 2008			NPP	Dec 02, 2006
	5053432*PED	Apr 01, 2009			PED	Jun 02, 2007
	6166046	Nov 25, 2013				
	6166046*PED	May 25, 2014				
	6291488	Nov 25, 2013		U-406		
	6291488*PED	May 25, 2014				
ATROPINE SULFATE; EDROPHONIUM CHLORIDE; ENLON-PLUS						
019677 001	4952586	Aug 28, 2007		U-54		
ATROPINE SULFATE; EDROPHONIUM CHLORIDE; ENLON-PLUS						
019678 001	4952586	Aug 28, 2007		U-54		
AZACITIDINE; VIDAZA						
050794 001					ODE	May 19, 2011
AZELAIC ACID; FINACEA						
021470 001	4713394	Jan 17, 2006		U-492	NDF	Dec 24, 2005
	6534070	Nov 18, 2018				
AZELASTINE HYDROCHLORIDE; ASTELIN						
020114 001	5164194	Nov 01, 2010		U-207		
	5164194*PED	May 01, 2011				
AZELASTINE HYDROCHLORIDE; OPTIVAR						
021127 001	5164194	Nov 01, 2010				
	5164194*PED	May 01, 2011				
BACLOFEN; KEMSTRO						
021589 001	6024981	Apr 09, 2018		DP		
	6221392	Apr 09, 2018		DP		
BACLOFEN; KEMSTRO						
021589 002	6024981	Apr 09, 2018		DP		
	6221392	Apr 09, 2018		DP		
BALSALAZIDE DISODIUM; COLAZAL						
020610 001	4412992	Jul 08, 2006			NCE	Jul 18, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BECLOMETHASONE DIPROPIONATE; QVAR 40</u>					
020911 002	5605674	Feb 25, 2014			
	5683677	Nov 04, 2014			
	5695743	Jul 06, 2010			
	5766573	Nov 28, 2009		U-356	
	5776432	Jul 07, 2015			
	6352684	Nov 28, 2009			
<u>BECLOMETHASONE DIPROPIONATE; QVAR 80</u>					
020911 001	5605674	Feb 25, 2014			
	5683677	Nov 04, 2014			
	5695743	Jul 06, 2010			
	5766573	Nov 28, 2009		U-356	
	5776432	Jul 07, 2015			
	6352684	Nov 28, 2009			
<u>BENZAEPRIIL HYDROCHLORIDE; LOTENSIN</u>					
019851 001				M-30 PED	Mar 02, 2007 Sep 02, 2007
<u>BENZAEPRIIL HYDROCHLORIDE; LOTENSIN</u>					
019851 002				M-30 PED	Mar 02, 2007 Sep 02, 2007
<u>BENZAEPRIIL HYDROCHLORIDE; LOTENSIN</u>					
019851 003				M-30 PED	Mar 02, 2007 Sep 02, 2007
<u>BENZAEPRIIL HYDROCHLORIDE; LOTENSIN</u>					
019851 004				M-30 PED	Mar 02, 2007 Sep 02, 2007
<u>BERACTANT; SURVANTA</u>					
020032 001	4397839	Jul 01, 2005			
<u>BETAMETHASONE DIPROPIONATE; DIPROLENE</u>					
019716 001	4775529	May 21, 2007			
<u>BETAMETHASONE VALERATE; LUXIQ</u>					
020934 001	6126920	Mar 01, 2016		U-484	
<u>BETAXOLOL HYDROCHLORIDE; BETOPTIC S</u>					
019845 001	4911920	Mar 27, 2007			
<u>BETAXOLOL HYDROCHLORIDE; PILOCARPINE HYDROCHLORIDE; BETOPTIC PILO</u>					
020619 001	5635172	Jun 03, 2014		U-191	
<u>BEXAROTENE; TARGRETIN</u>					
021055 001	5466861	Nov 14, 2012		ODE	Dec 29, 2006
	5780676	Jul 14, 2015		U-509	
	5962731	Oct 05, 2016		U-475	
	6043279	Apr 22, 2012		U-509	
	6320074	Apr 22, 2012	DS	U-509	
<u>BEXAROTENE; TARGRETIN</u>					
021056 001	5466861	Nov 14, 2012		ODE	Dec 29, 2006
	5780676	Jul 14, 2015		U-510	
	5962731	Oct 05, 2016			
	6043279	Apr 22, 2012		U-510	
	6320074	Apr 22, 2012	DS	U-510	
<u>BICALUTAMIDE; CASODEX</u>					
020498 001	4636505	Oct 01, 2008			
<u>BIMATOPROST; LUMIGAN</u>					
021275 001	5688819	Sep 21, 2012		NCE	Mar 16, 2006
	6403649	Sep 21, 2012		U-446	
<u>BISACODYL; POLYETHYLENE GLYCOL; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; HALFLYELY</u>					
021551 001				NP	May 10, 2007
<u>BIVALIRUDIN; ANGIOMAX</u>					
020873 001	5196404	Mar 23, 2010		I-458 NCE	Jun 13, 2008 Dec 15, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
BORTEZOMIB; VELCADE					
021602 001	5780454	Oct 28, 2014	DS DP	I-452	Mar 25, 2008
	6083903	Oct 28, 2014		NCE	May 13, 2008
	6297217	Oct 28, 2014		U-515	
	6617317	Oct 28, 2014		U-515	May 13, 2010
	6713446	Jan 25, 2022			
	6747150	Oct 28, 2014			
	6958319	Jan 25, 2022			
BOSENTAN; TRACLEER					
021290 001	5292740	Nov 20, 2015		NCE	Nov 20, 2006
				ODE	Nov 20, 2008
BOSENTAN; TRACLEER					
021290 002	5292740	Nov 20, 2015		NCE	Nov 20, 2006
				ODE	Nov 20, 2008
BRIMONIDINE TARTRATE; ALPHAGAN P					
021262 001	5424078	Jun 13, 2012	DP		
	5424078*PED	Dec 13, 2012			
	6562873	Jul 10, 2021			
	6562873*PED	Jan 10, 2022			
	6627210	Jul 18, 2021			
	6627210*PED	Jan 18, 2022			
	6641834	Jul 28, 2021			
	6641834*PED	Jan 28, 2022			
	6673337	Jul 26, 2021			
	6673337*PED	Jan 26, 2022			
BRIMONIDINE TARTRATE; ALPHAGAN P					
021770 001	5424078	Jun 13, 2012	DP	NP	Aug 19, 2008
	6562873	Jul 10, 2021	DP		
	6627210	Jul 18, 2021	DP		
	6641834	Jul 28, 2021	DP		
	6673337	Jul 26, 2021	DP		
BRINZOLAMIDE; AZOPT					
020816 001	5240923	Aug 31, 2010		U-224	
	5378703	Apr 01, 2012		U-224	
	5461081	Oct 24, 2012		U-225	
BROMFENAC SODIUM; XIBROM					
021664 001				NP	Mar 24, 2008
BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE; EFIDAC 24 PSEUDOEPHEDRINE HCL/BROMPHENIRAMINE MALEATE					
019672 001	4662880	Mar 14, 2006			
	4801461	Mar 14, 2006			
	4810502	Mar 14, 2006			
BUDESONIDE; ENTOCORT EC					
021324 001	5643602	Jul 01, 2014	DP	U-655	I-454
	5643602*PED	Jan 01, 2015			Apr 29, 2008
	6423340	Nov 15, 2010			
	6423340*PED	May 11, 2011			
BUDESONIDE; PULMICORT					
020441 002	4668218	Apr 11, 2006			
	4668218*PED	Oct 11, 2006			
	4907583	Mar 13, 2007			
	4907583*PED	Sep 13, 2007			
BUDESONIDE; PULMICORT					
020441 003	4668218	Apr 11, 2006			
	4668218*PED	Oct 11, 2006			
	4907583	Mar 13, 2007			
	4907583*PED	Sep 13, 2007			
BUDESONIDE; PULMICORT RESPULES					
020929 001	4787536	Feb 27, 2006			
	4787536*PED	Aug 27, 2006			
	6598603	Dec 23, 2018		U-529	
	6598603*PED	Jun 23, 2019			
	6899099	Dec 23, 2018		U-645	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
BUDESONIDE; PULMICORT RESPULES					
020929 002	4787536	Feb 27, 2006	U-529		
	4787536*PED	Aug 27, 2006			
	6598603	Dec 23, 2018			
	6598603*PED	Jun 23, 2019			
BUDESONIDE; PULMICORT RESPULES					
020929 003	4787536	Feb 27, 2006	U-529		
	4787536*PED	Aug 27, 2006			
	6598603	Dec 23, 2018			
	6598603*PED	Jun 23, 2019			
BUDESONIDE; RHINOCORT					
020746 001	6291445	Apr 29, 2017	DP	U-557	
	6291445*PED	Oct 29, 2017			
	6686346	Apr 29, 2017			
BUDESONIDE; RHINOCORT					
020746 002	6291445	Apr 29, 2017	DP	U-557	
	6291445*PED	Oct 29, 2017			
	6686346	Apr 29, 2017			
BUPRENORPHINE HYDROCHLORIDE; SUBUTEX					
020732 002				NDF	Oct 08, 2005
				ODE	Oct 08, 2009
BUPRENORPHINE HYDROCHLORIDE; SUBUTEX					
020732 003				NDF	Oct 08, 2005
				ODE	Oct 08, 2009
BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE; SUBOXONE					
020733 001				NDF	Oct 08, 2005
				ODE	Oct 08, 2009
BUPRENORPHINE HYDROCHLORIDE; NALOXONE HYDROCHLORIDE; SUBOXONE					
020733 002				NDF	Oct 08, 2005
				ODE	Oct 08, 2009
BUPROPION HYDROCHLORIDE; WELLBUTRIN SR					
020358 001	5358970	Aug 12, 2013			
	5427798	Aug 12, 2013			
	5731000	Aug 12, 2013			
	5763493	Aug 12, 2013			
BUPROPION HYDROCHLORIDE; WELLBUTRIN SR					
020358 002	5358970	Aug 12, 2013			
	5427798	Aug 12, 2013			
	5731000	Aug 12, 2013			
	5763493	Aug 12, 2013			
BUPROPION HYDROCHLORIDE; WELLBUTRIN SR					
020358 003	5358970	Aug 12, 2013			
	5427798	Aug 12, 2013			
	5731000	Aug 12, 2013			
	5763493	Aug 12, 2013			
BUPROPION HYDROCHLORIDE; WELLBUTRIN SR					
020358 004	5358970	Aug 12, 2013			
	5427798	Aug 12, 2013			
	5731000	Aug 12, 2013			
	5763493	Aug 12, 2013			
BUPROPION HYDROCHLORIDE; WELLBUTRIN XL					
021515 001	6096341	Oct 30, 2018			
	6143327	Oct 30, 2018			
BUPROPION HYDROCHLORIDE; WELLBUTRIN XL					
021515 002	6096341	Oct 30, 2018			
	6143327	Oct 30, 2018			
BUPROPION HYDROCHLORIDE; ZYBAN					
020711 002	5358970	Aug 12, 2013			
	5427798	Aug 12, 2013			
	5731000	Aug 12, 2013			
	5763493	Aug 12, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BUPROPION HYDROCHLORIDE; ZYBAN</u>					
020711 003	5358970	Aug 12, 2013			
	5427798	Aug 12, 2013			
	5731000	Aug 12, 2013			
	5763493	Aug 12, 2013			
<u>BUSPIRONE HYDROCHLORIDE; BUSPAR</u>					
021190 001	5015646	May 14, 2008			
	5015646*PED	Nov 14, 2008			
<u>BUSPIRONE HYDROCHLORIDE; BUSPAR</u>					
021190 002	5015646	May 14, 2008			
	5015646*PED	Nov 14, 2008			
<u>BUSPIRONE HYDROCHLORIDE; BUSPAR</u>					
021190 003	5015646	May 14, 2008			
	5015646*PED	Nov 14, 2008			
<u>BUSPIRONE HYDROCHLORIDE; BUSPAR</u>					
021190 004	5015646	May 14, 2008			
	5015646*PED	Nov 14, 2008			
<u>BUSULFAN; BUSULFEX</u>					
020954 001	5430057	Sep 30, 2013	U-263	ODE	Feb 04, 2006
	5430057*PED	Mar 30, 2014	U-263	PED	Aug 04, 2006
	5559148	May 24, 2015	U-264		
	5559148*PED	Nov 24, 2015	U-264		
<u>BUTENAFINE HYDROCHLORIDE; MENTAX</u>					
020524 001	5021458	Oct 18, 2010	U-177		
<u>BUTENAFINE HYDROCHLORIDE; MENTAX-TC</u>					
021408 001				NP	Oct 17, 2005
<u>BUTOCONAZOLE NITRATE; GYNAZOLE-1</u>					
019881 001	5266329	Nov 30, 2010	U-457		
<u>CABERGOLINE; DOSTINEX</u>					
020664 001	4526892	Dec 29, 2005			
<u>CAFFEINE CITRATE; CAFKIT</u>					
020793 001				ODE	Sep 21, 2006
<u>CALCIPOTRIENE; DOVONEX</u>					
020273 001	4866048	Dec 29, 2007			
<u>CALCIPOTRIENE; DOVONEX</u>					
020554 001	4866048	Dec 29, 2007			
	5763426	Jun 09, 2015	DS DP		
<u>CALCIPOTRIENE; DOVONEX</u>					
020611 001	4866048	Dec 29, 2007			
	5763426	Jun 09, 2015	DS DP		
<u>CALCITONIN SALMON RECOMBINANT; FORTICAL</u>					
021406 001	6440392	Feb 02, 2021	DP	U-227	
<u>CALCITONIN, SALMON; MIACALCIN</u>					
020313 002	5733569	Mar 31, 2015		U-227	
	5759565	Mar 31, 2015			
<u>CALCITRIOL; CALCIJEX</u>					
018874 001	6051567	Aug 02, 2019			
	6051567*PED	Feb 02, 2020			
	6265392	Aug 02, 2019			
	6265392*PED	Feb 02, 2020			
	6274169	Aug 02, 2019			
	6274169*PED	Feb 02, 2020			
<u>CALCITRIOL; CALCIJEX</u>					
018874 002	6051567	Aug 02, 2019			
	6051567*PED	Feb 02, 2020			
	6265392	Aug 02, 2019			
	6265392*PED	Feb 02, 2020			
	6274169	Aug 02, 2019			
	6274169*PED	Feb 02, 2020			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CALCIUM ACETATE; PHOSLO</u>					
019976 001	4870105	Apr 07, 2007	U-381		
<u>CALCIUM ACETATE; PHOSLO</u>					
021160 001	4870105	Apr 07, 2007	U-381		
<u>CALCIUM ACETATE; PHOSLO</u>					
021160 002	4870105	Apr 07, 2007	U-381		
	6576665	Apr 03, 2021			
<u>CALCIUM ACETATE; PHOSLO GELCAPS</u>					
021160 003	4870105	Apr 07, 2007	U-381		
	6576665	Apr 03, 2021			
<u>CALCIUM CARBONATE; FAMOTIDINE; MAGNESIUM HYDROXIDE; PEPCID COMPLETE</u>					
020958 001	5229137	May 16, 2012	U-349		
	5229137*PED	Nov 16, 2012			
	5817340	Dec 01, 2012			
	5817340*PED	Jun 01, 2013			
	5989588	Sep 30, 2015	U-349		
	5989588*PED	Mar 30, 2016	U-349		
<u>CALCIUM CARBONATE; RISEDRONATE SODIUM; ACTONEL WITH CALCIUM (COPACKAGED)</u>					
021823 001	5583122	Dec 10, 2013	DS DP	U-353	
	5994329	Jul 17, 2018		U-353	
	6015801	Jul 17, 2018		U-353	
	6096342	Nov 21, 2011	DP		
	6165513	Jun 10, 2018	DP		
	6432932	Jul 17, 2018		U-595	
	6465443	Aug 14, 2018	DP		
<u>CANDESARTAN CILEXETIL; ATACAND</u>					
020838 001	5196444	Jun 04, 2012	DS DP	U-3	I-456 May 18, 2008
	5196444	Jun 04, 2012	DS DP	U-660	I-455 May 18, 2008
	5534534	Jul 09, 2013	DP		I-448 Feb 22, 2008
	5705517	Apr 18, 2011	DS DP	U-660	M-21 Sep 13, 2005
<u>CANDESARTAN CILEXETIL; ATACAND</u>					
020838 002	5196444	Jun 04, 2012	DS DP	U-3	I-456 May 18, 2008
	5196444	Jun 04, 2012	DS DP	U-660	I-455 May 18, 2008
	5534534	Jul 09, 2013	DP		I-448 Feb 22, 2008
	5705517	Apr 18, 2011	DS DP	U-660	M-21 Sep 13, 2005
<u>CANDESARTAN CILEXETIL; ATACAND</u>					
020838 003	5196444	Jun 04, 2012	DS DP	U-660	I-456 May 18, 2008
	5196444	Jun 04, 2012	DS DP	U-3	I-455 May 18, 2008
	5534534	Jul 09, 2013	DP		I-448 Feb 22, 2008
	5705517	Apr 18, 2011	DS DP	U-660	M-21 Sep 13, 2005
<u>CANDESARTAN CILEXETIL; ATACAND</u>					
020838 004	5196444	Jun 04, 2012	DS DP	U-660	I-456 May 18, 2008
	5196444	Jun 04, 2012	DS DP	U-3	I-455 May 18, 2008
	5534534	Jul 09, 2013	DP		I-448 Feb 22, 2008
	5705517	Apr 18, 2011	DS DP	U-660	M-21 Sep 13, 2005
<u>CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE; ATACAND HCT</u>					
021093 001	5196444	Jun 04, 2012		U-3	
	5534534	Jul 09, 2013			
	5703110	Apr 18, 2011		U-3	
	5705517	Apr 18, 2011		U-3	
	5721263	Feb 24, 2015		U-3	
	5958961	Jun 06, 2014		U-3	
<u>CANDESARTAN CILEXETIL; HYDROCHLOROTHIAZIDE; ATACAND HCT</u>					
021093 002	5196444	Jun 04, 2012		U-3	
	5534534	Jul 09, 2013			
	5703110	Apr 18, 2011		U-3	
	5705517	Apr 18, 2011		U-3	
	5721263	Feb 24, 2015		U-3	
	5958961	Jun 06, 2014		U-3	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CAPECITABINE; XELODA</u>					
020896 001	4966891 5472949	Jan 13, 2011 Dec 14, 2013	U-272 U-271	I-461	Jun 15, 2008
<u>CAPECITABINE; XELODA</u>					
020896 002	4966891 5472949	Jan 13, 2011 Dec 14, 2013	U-272 U-271	I-461	Jun 15, 2008
<u>CAPTOPRIL; CAPOTEN</u>					
018343 001	5238924	Aug 24, 2010	U-92		
<u>CAPTOPRIL; CAPOTEN</u>					
018343 002	5238924	Aug 24, 2010	U-92		
<u>CAPTOPRIL; CAPOTEN</u>					
018343 003	5238924	Aug 24, 2010	U-92		
<u>CAPTOPRIL; CAPOTEN</u>					
018343 004	5238924	Aug 24, 2010	U-92		
<u>CAPTOPRIL; CAPOTEN</u>					
018343 005	5238924	Aug 24, 2010	U-92		
<u>CAPTOPRIL; CAPOTEN</u>					
018343 006	5238924	Aug 24, 2010	U-92		
<u>CAPTOPRIL; CAPOTEN</u>					
018343 007	5238924	Aug 24, 2010	U-92		
<u>CARBAMAZEPINE; CARBATROL</u>					
020712 001	5326570 5912013	Jul 05, 2011 Jun 15, 2016	U-215 U-277		
<u>CARBAMAZEPINE; CARBATROL</u>					
020712 002	5326570 5912013	Jul 05, 2011 Jun 15, 2016	U-215 U-277		
<u>CARBAMAZEPINE; CARBATROL</u>					
020712 003	5326570 5912013	Jul 05, 2011 Jun 15, 2016	U-215 U-277		
<u>CARBAMAZEPINE; EQUETRO</u>					
021710 001	5326570 5912013 6977253	Jul 23, 2011 Jun 15, 2016 May 19, 2024	DP DP U-693	NP	Dec 10, 2007
<u>CARBAMAZEPINE; EQUETRO</u>					
021710 002	5326570 5912013 6977253	Jul 23, 2011 Jun 15, 2016 May 19, 2024	DP DP U-693	NP	Dec 10, 2007
<u>CARBAMAZEPINE; EQUETRO</u>					
021710 003	5326570 5912013 6977253	Jul 23, 2011 Jun 15, 2016 May 19, 2024	DP DP U-693	NP	Dec 10, 2007
<u>CARBAMAZEPINE; TEGRETOL-XR</u>					
020234 001	5284662 RE34990	Feb 08, 2011 Jul 29, 2007			
<u>CARBAMAZEPINE; TEGRETOL-XR</u>					
020234 002	5284662 RE34990	Feb 08, 2011 Jul 29, 2007			
<u>CARBAMAZEPINE; TEGRETOL-XR</u>					
020234 003	5284662 RE34990	Feb 08, 2011 Jul 29, 2007			
<u>CARBIDOPA; ENTACAPONE; LEVODOPA; STALEVO 100</u>					
021485 002	4963590 5112861 5135950 5446194 6500867 6797732	Nov 27, 2007 May 12, 2009 Oct 31, 2010 Oct 19, 2013 Jun 29, 2020 Jun 29, 2020	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE		
<u>CARBIDOPA; ENTACAPONE; LEVODOPA; STALEVO 150</u>										
021485 003	4963590	Nov	27,	2007	DS	DP				
	5112861	May	12,	2009						
	5135950	Oct	31,	2010						
	5446194	Oct	19,	2013						
	6500867	Jun	29,	2020						
	6797732	Jun	29,	2020						
<u>CARBIDOPA; ENTACAPONE; LEVODOPA; STALEVO 50</u>										
021485 001	4963590	Nov	27,	2007	DS	DP				
	5112861	May	12,	2009						
	5135950	Oct	31,	2010						
	5446194	Oct	19,	2013						
	6500867	Jun	29,	2020						
	6797732	Jun	29,	2020						
<u>CARBIDOPA; LEVODOPA; SINEMET CR</u>										
019856 001	4832957	Jun	16,	2006						
	4900755	Jun	16,	2006						
<u>CARBIDOPA; LEVODOPA; SINEMET CR</u>										
019856 002	4832957	Jun	16,	2006						
	4900755	Jun	16,	2006						
<u>CARMUSTINE; GLIADEL</u>										
020637 001	4757128	Aug	01,	2006			I-382	Feb	25,	
	4789724	Aug	01,	2006						ODE
<u>CARVEDILOL; COREG</u>										
020297 001	4503067	Mar	05,	2007	U-3	I-388		Mar	27,	
	5760069	Jun	07,	2015						U-233
	5902821	Feb	07,	2016						U-313
<u>CARVEDILOL; COREG</u>										
020297 002	4503067	Mar	05,	2007	U-3	I-388		Mar	27,	
	5760069	Jun	07,	2015						U-233
	5902821	Feb	07,	2016						U-313
<u>CARVEDILOL; COREG</u>										
020297 003	4503067	Mar	05,	2007	U-3	I-388		Mar	27,	
	5760069	Jun	07,	2015						U-233
	5902821	Feb	07,	2016						U-313
<u>CARVEDILOL; COREG</u>										
020297 004	4503067	Mar	05,	2007	U-3	I-388		Mar	27,	
	5760069	Jun	07,	2015						U-233
	5902821	Feb	07,	2016						U-313
<u>CASPOFUNGIN ACETATE; CANCIDAS</u>										
021227 001	5378804	Mar	16,	2013	DS	DP	I-438	Sep	29,	
	5514650	Mar	16,	2013						U-607
	5792746	Mar	16,	2013						U-607
	5952300	Mar	28,	2017						NCE
	6136783	Mar	28,	2017						U-607
<u>CASPOFUNGIN ACETATE; CANCIDAS</u>										
021227 002	5378804	Mar	16,	2013	DS	DP	I-438	Sep	29,	
	5514650	Mar	16,	2013						U-607
	5792746	Mar	16,	2013						U-607
	5952300	Mar	28,	2017						NCE
	6136783	Mar	28,	2017						U-607
<u>CEFIDITOREN PIVOXIL; SPECTRACEF</u>										
021222 001	4839350	Jun	13,	2006			I-364	Aug	21,	
	4918068	Jun	13,	2006						NCE
	5958915	Oct	14,	2016						
<u>CELECOXIB; CELEBREX</u>										
020998 001	5466823	Nov	30,	2013	DS	DP	I-466	Jul	29,	
	5563165	Nov	30,	2013						
	5760068	Jun	02,	2015						U-299
	5760068	Jun	02,	2015						U-672
	5972986	Oct	14,	2017						U-299

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
CELECOXIB; CELEBREX									
020998 002	5466823	Nov 30, 2013	DS	DP	U-672 U-299 U-299	I-466		Jul 29, 2008	
	5563165	Nov 30, 2013							
	5760068	Jun 02, 2015							
	5760068	Jun 02, 2015							
	5972986	Oct 14, 2017							
CELECOXIB; CELEBREX									
020998 003	5466823	Nov 30, 2013	DS	DP	U-672 U-299 U-299	I-466		Jul 29, 2008	
	5563165	Nov 30, 2013							
	5760068	Jun 02, 2015							
	5760068	Jun 02, 2015							
	5972986	Oct 14, 2017							
CERIVASTATIN SODIUM; BAYCOL									
020740 001	5006530	Jun 26, 2011							
	5177080	Nov 26, 2011							
CERIVASTATIN SODIUM; BAYCOL									
020740 002	5006530	Jun 26, 2011							
	5177080	Nov 26, 2011							
CERIVASTATIN SODIUM; BAYCOL									
020740 003	5006530	Jun 26, 2011							
	5177080	Nov 26, 2011							
CERIVASTATIN SODIUM; BAYCOL									
020740 004	5006530	Jun 26, 2011							
	5177080	Nov 26, 2011							
CERIVASTATIN SODIUM; BAYCOL									
020740 005	5006530	Jun 26, 2011							
	5177080	Nov 26, 2011							
CERIVASTATIN SODIUM; BAYCOL									
020740 006	5006530	Jun 26, 2011							
	5177080	Jan 26, 2011							
CETIRIZINE HYDROCHLORIDE; ZYRTEC									
019835 001	4525358	Jun 25, 2007				NPP PED		Oct 21, 2005 Apr 21, 2006	
	4525358*PED	Dec 25, 2007							
CETIRIZINE HYDROCHLORIDE; ZYRTEC									
019835 002	4525358	Jun 25, 2007				NPP PED		Oct 21, 2005 Apr 21, 2006	
	4525358*PED	Dec 25, 2007							
CETIRIZINE HYDROCHLORIDE; ZYRTEC									
020346 001	4525358	Jun 25, 2007				NPP PED		Oct 21, 2005 Apr 21, 2006	
	4525358*PED	Dec 25, 2007							
CETIRIZINE HYDROCHLORIDE; ZYRTEC									
021621 001	4525358	Jun 25, 2007	DS	DP	U-565				
	4525358*PED	Dec 25, 2007							
	6455533	Jul 02, 2018							
CETIRIZINE HYDROCHLORIDE; ZYRTEC									
021621 002	4525358	Jun 25, 2007	DS	DP	U-565				
	4525358*PED	Dec 25, 2007							
	6455533	Jul 02, 2018							
CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE; ZYRTEC-D 12 HOUR									
021150 001	4525358	Jun 25, 2007			U-295 U-295 U-295				
	4525358*PED	Dec 25, 2007							
	6469009	Jul 13, 2019							
	6489329	Apr 08, 2016							
CETRORELIX; CETROTIDE									
021197 001	4800191	Jul 17, 2007			U-426 U-426	NCE		Aug 11, 2005	
	5198533	Jul 17, 2007							
	6319192	Apr 23, 2018							
	6863891	Feb 19, 2013							
CETRORELIX; CETROTIDE									
021197 002	4800191	Jul 17, 2007			U-426 U-426	NCE		Aug 11, 2005	
	5198533	Jul 17, 2007							
	6319192	Apr 23, 2018							
	6863891	Feb 19, 2013							

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CETYL ALCOHOL; COLFOSCERIL PALMITATE; TYLOXAPOL; EXOSURF NEONATAL</u>					
020044 001	4826821	May 02, 2006			
	5110806	May 02, 2006			
<u>CEVIMELINE HYDROCHLORIDE; EVOXAC</u>					
020989 002	4855290	Aug 30, 2009			
	5340821	Jul 07, 2013	U-309		
	5580880	Jun 06, 2015	U-310		
<u>CHLORHEXIDINE GLUCONATE; CHLORHEXIDINE GLUCONATE</u>					
021669 001				NDF	Apr 25, 2008
<u>CHLORHEXIDINE GLUCONATE; PERIOCHIP</u>					
020774 001	5002769	Mar 16, 2006			
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL; CHLORAPREP ONE-STEP</u>					
020832 001	5538353	Aug 25, 2015	DP		
	5690958	Sep 30, 2016	DP		
	5752363	Apr 22, 2017	DP		
	5772346	Apr 22, 2017	DP		
	6536975	Nov 10, 2020	DP		
	D386849	Nov 25, 2011	DP		
	D396911	Aug 11, 2012	DP		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL; CHLORAPREP ONE-STEP SEPP</u>					
021555 001	5690958	Sep 30, 2016	DP	NP	Oct 07, 2005
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL; CHLORAPREP WITH TINT</u>					
020832 002	5538353	Aug 25, 2015	DP	NP	May 03, 2008
	5690958	Sep 30, 2016	DP		
	6536975	Nov 10, 2020	DP		
	6729786	Mar 14, 2023	DP		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL; CHLORASCUB MAXI SWABSTICK</u>					
021524 003	D468424	Jan 07, 2017		NP	Jun 03, 2008
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL; CHLORASCUB SWAB</u>					
021524 001				NP	Jun 03, 2008
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL; CHLORASCUB SWABSTICK</u>					
021524 002				NP	Jun 03, 2008
<u>CHLORPHENIRAMINE MALEATE; EFIDAC 24 CHLORPHENIRAMINE MALEATE</u>					
019746 002	4857330	Aug 15, 2006			
<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE; ADVIL ALLERGY SINUS</u>					
021441 001				NP	Dec 19, 2005
<u>CHLORPHENIRAMINE MALEATE; IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE; CHILDREN'S ADVIL ALLERGY SINUS</u>					
021587 001				NP	Feb 24, 2007
<u>CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX; TUSSIONEX</u>					
019111 001	4762709	Aug 09, 2005			
<u>CHLORTHALIDONE; THALITONE</u>					
019574 001	4933360	Jun 12, 2007			
<u>CHORIOGONADOTROPIN ALFA; OVIDREL</u>					
021149 001	5767251	Jun 16, 2015			
<u>CHORIOGONADOTROPIN ALFA; OVIDREL</u>					
021149 002	5767251	Jun 16, 2015	DS		
	6706681	Mar 16, 2021	DP		
<u>CICLOPIROX; LOPROX</u>					
021159 001				NDF	Feb 28, 2006
<u>CICLOPIROX; PENLAC</u>					
021022 001	4957730	Sep 18, 2007		U-379	
<u>CIDOFOVIR; VISTIDE</u>					
020638 001	5142051	Jun 26, 2010			
<u>CINACALCET HYDROCHLORIDE; SENSIPAR</u>					
021688 001	6011068	Dec 14, 2016	DS DP	NCE	Mar 08, 2009
	6031003	Dec 14, 2016		U-559	
	6211244	Oct 23, 2015	DS DP	U-560	Mar 08, 2011
	6313146	Dec 14, 2016	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CINACALCET HYDROCHLORIDE; SENSIPAR</u>					
021688 002	6011068	Dec 14, 2016	DS DP	NCE	Mar 08, 2009
	6031003	Dec 14, 2016		ODE	Mar 08, 2011
	6211244	Oct 23, 2015	DS DP U-560		
	6313146	Dec 14, 2016	DS DP		
<u>CINACALCET HYDROCHLORIDE; SENSIPAR</u>					
021688 003	6011068	Dec 14, 2016	DS DP	NCE	Mar 08, 2009
	6031003	Dec 14, 2016		ODE	Mar 08, 2011
	6211244	Oct 23, 2015	DS DP U-559		
	6313146	Dec 14, 2016	DS DP U-560		
<u>CIPROFLOXACIN; CIPRO</u>					
019847 001	4808583	Feb 28, 2006		I-421	Mar 25, 2007
	4808583*PED	Aug 28, 2006		PED	Sep 25, 2007
<u>CIPROFLOXACIN; CIPRO</u>					
020780 001	5695784	Dec 09, 2014		I-421	Mar 25, 2007
	5695784*PED	Jun 09, 2015		PED	Sep 25, 2007
	6136347	Jan 06, 2013	U-362		
	6136347*PED	Jul 06, 2013			
<u>CIPROFLOXACIN; CIPRO</u>					
020780 002	5695784	Dec 09, 2014		I-421	Mar 25, 2007
	5695784*PED	Jun 09, 2015		PED	Sep 25, 2007
	6136347	Jan 06, 2013	U-362		
	6136347*PED	Jul 06, 2013			
<u>CIPROFLOXACIN; CIPRO IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
019857 001	4808583	Feb 28, 2006		I-421	Mar 25, 2007
	4808583*PED	Aug 28, 2006		PED	Sep 25, 2007
	4957922	Sep 18, 2007			
	4957922*PED	Mar 18, 2008			
<u>CIPROFLOXACIN; CIPRO IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER</u>					
019858 001	4808583	Feb 28, 2006			
	4808583*PED	Aug 28, 2006			
	4957922	Sep 18, 2007			
	4957922*PED	Mar 18, 2008			
<u>CIPROFLOXACIN HYDROCHLORIDE; CIPRO</u>					
019537 001	5286754	Feb 15, 2011		I-421	Mar 25, 2007
	5286754*PED	Aug 15, 2011		PED	Sep 25, 2007
<u>CIPROFLOXACIN HYDROCHLORIDE; CIPRO</u>					
019537 002	5286754	Feb 15, 2011		I-421	Mar 25, 2007
	5286754*PED	Aug 15, 2011		PED	Sep 25, 2007
<u>CIPROFLOXACIN HYDROCHLORIDE; CIPRO</u>					
019537 003	5286754	Feb 15, 2011		I-421	Mar 25, 2007
	5286754*PED	Aug 15, 2011		PED	Sep 25, 2007
<u>CIPROFLOXACIN HYDROCHLORIDE; CIPRO</u>					
019537 004	5286754	Feb 15, 2011		I-421	Mar 25, 2007
	5286754*PED	Aug 15, 2011		PED	Sep 25, 2007
<u>CIPROFLOXACIN HYDROCHLORIDE; PROQUIN XR</u>					
021744 001	5972389	Sep 19, 2016	DP U-663	NDF	May 19, 2008
	6340475	Sep 19, 2016	DP U-663		
	6488962	Jun 20, 2020	DP		
	6635280	Sep 19, 2016	DP U-663		
<u>CIPROFLOXACIN HYDROCHLORIDE; HYDROCORTISONE; CIPRO HC</u>					
020805 001	4844902	Feb 11, 2008	DP		
	5843930	Jul 06, 2015		U-646	
	5965549	Jun 06, 2015	DP		
<u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE; CIPRO XR</u>					
021473 001				NDF	Dec 13, 2005
				NC	Dec 13, 2005
				PED	Jun 13, 2006
				PED	Jun 13, 2006

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE; CIPRO XR</u>					
021473 002				I-416	Aug 28, 2006
				NS	Aug 28, 2006
				NDF	Dec 13, 2005
				NC	Dec 13, 2005
				PED	Feb 28, 2007
				PED	Feb 28, 2007
				PED	Jun 13, 2006
				PED	Jun 13, 2006
<u>CIPROFLOXACIN; DEXAMETHASONE; CIPRODEX</u>					
021537 001	4844902	Feb 11, 2008		NC	Jul 18, 2006
	6284804	Aug 10, 2020			
	6359016	Aug 10, 2020			
<u>CISAPRIDE MONOHYDRATE; PROPULSID</u>					
020210 001	4962115	Oct 09, 2007	U-79		
<u>CISAPRIDE MONOHYDRATE; PROPULSID</u>					
020210 002	4962115	Oct 09, 2007	U-79		
<u>CISAPRIDE MONOHYDRATE; PROPULSID</u>					
020398 001	4962115	Oct 09, 2007	U-79		
<u>CISAPRIDE MONOHYDRATE; PROPULSID QUICKSOLV</u>					
020767 001	4962115	Oct 09, 2007	U-79		
	5648093	Jul 15, 2014			
<u>CISATRACURIUM BESYLATE; NIMBEX</u>					
020551 001	5453510	Sep 26, 2012	U-127		
<u>CISATRACURIUM BESYLATE; NIMBEX PRESERVATIVE FREE</u>					
020551 002	5453510	Sep 26, 2012	U-127		
<u>CISATRACURIUM BESYLATE; NIMBEX PRESERVATIVE FREE</u>					
020551 003	5453510	Sep 26, 2012	U-127		
<u>CLOBETASOL PROPIONATE; CLOBEX</u>					
021535 001	6106848	Sep 22, 2017		NDF	Jul 24, 2006
<u>CLOBETASOL PROPIONATE; CLOBEX</u>					
021644 001				NDF	Feb 05, 2007
<u>CLOBETASOL PROPIONATE; OLUX</u>					
021142 001	6126920	Oct 03, 2017	U-484	I-374	Dec 20, 2005
<u>CLOFARABINE; CLOLAR</u>					
021673 001	5384310	May 23, 2009	DS DP	NCE	Dec 28, 2009
	5661136	Aug 26, 2014	U-626	ODE	Dec 28, 2011
				PED	Jun 28, 2012
				PED	Jun 28, 2010
<u>CLOPIDOGREL BISULFATE; PLAVIX</u>					
020839 001	4847265	Nov 17, 2011			
	5576328	Jan 31, 2014	U-432		
	6429210	Jun 10, 2019			
	6504030	Jun 10, 2019			
<u>CLOZAPINE; CLOZARIL</u>					
019758 001				I-380	Dec 18, 2005
<u>CLOZAPINE; CLOZARIL</u>					
019758 002				I-380	Dec 18, 2005
<u>COLESEVELAM HYDROCHLORIDE; WELCHOL</u>					
021141 001	5607669	Jun 10, 2014	U-323		
	5679717	Apr 29, 2014	U-323		
	5693675	Dec 02, 2014			
	5917007	Apr 29, 2014	U-323		
	5919832	Jun 10, 2014			
	6066678	Jun 10, 2014	U-323		
	6433026	Jun 10, 2014			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>COLESEVELAM HYDROCHLORIDE; WELCHOL</u>					
021176 001	5607669	Jun 10, 2014	U-323		
	5679717	Apr 29, 2014	U-323		
	5693675	Dec 02, 2014			
	5917007	Apr 29, 2014	U-323		
	5919832	Jun 10, 2014			
	6066678	Jun 10, 2014	U-323		
	6433026	Jun 10, 2014			
	6784254	Jun 10, 2014	DP		
<u>COLESTIPOL HYDROCHLORIDE; COLESTID</u>					
020222 001	5490987	Feb 13, 2013	DP		
<u>CONIVAPTAN HYDROCHLORIDE; VAPRISOL</u>					
021697 001				NCE	Dec 29, 2010
<u>CYCLOBENZAPRINE HYDROCHLORIDE; FLEXERIL</u>					
017821 001				D-78	Feb 03, 2006
<u>CYTARABINE; DEPOCYT</u>					
021041 001				ODE	Apr 01, 2006
<u>DALTEPARIN SODIUM; FRAGMIN</u>					
020287 001				I-414	Dec 10, 2006
<u>DALTEPARIN SODIUM; FRAGMIN</u>					
020287 003				I-414	Dec 10, 2006
<u>DALTEPARIN SODIUM; FRAGMIN</u>					
020287 004				I-414	Dec 10, 2006
<u>DANAPAROID SODIUM; ORGARAN</u>					
020430 001	5164377	Oct 03, 2010			
<u>DAPSONE; ACZONE</u>					
021794 001				NDF	Jul 07, 2008
<u>DAPTOMYCIN; CUBICIN</u>					
021572 001	5912226	Jun 15, 2016		NCE	Sep 12, 2008
	6468967	Sep 24, 2019			
	6852689	Sep 24, 2019	U-282		
<u>DAPTOMYCIN; CUBICIN</u>					
021572 002	5912226	Jun 15, 2016		NCE	Sep 12, 2008
	6468967	Sep 24, 2019			
	6852689	Sep 24, 2019	U-282		
<u>DARIFENACIN HYDROBROMIDE; ENABLEX</u>					
021513 001	5096890	Mar 13, 2010	DS DP	U-631	Dec 22, 2009
	6106864	Aug 21, 2016	DP	U-630	
<u>DARIFENACIN HYDROBROMIDE; ENABLEX</u>					
021513 002	5096890	Mar 13, 2010	DS DP	U-631	Dec 22, 2009
	6106864	Aug 21, 2016	DP	U-630	
<u>DEFERASIROX; EXJADE</u>					
021882 001				NCE	Nov 02, 2010
				ODE	Nov 02, 2012
<u>DELAVIDINE MESYLATE; RESCRIPTOR</u>					
020705 001	5563142	Oct 08, 2013			
<u>DELAVIDINE MESYLATE; RESCRIPTOR</u>					
020705 002	5563142	Oct 08, 2013			
	6177101	Jun 07, 2019			
<u>DESFLURANE; SUPRANE</u>					
020118 001	4762856	Feb 02, 2007		U-67	
<u>DESIRUDIN RECOMBINANT; IPRIVASK</u>					
021271 001	4801576	Jan 31, 2006		NCE	Apr 04, 2008
	5733874	Mar 31, 2015			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DES Loratadine; CLARINEX</u>					
021165 001	4659716	Apr 21, 2006	U-427	NCE	Dec 21, 2006
	4659716*PED	Oct 21, 2006	U-427	PED	Jun 21, 2007
	4863931	Sep 15, 2008			
	4863931*PED	Mar 15, 2009			
	6100274	Jul 07, 2019			
	6100274*PED	Jan 07, 2020			
<u>DES Loratadine; CLARINEX</u>					
021300 001	4659716	Apr 21, 2006	DP U-611	NCE	Dec 21, 2006
	4659716*PED	Oct 21, 2006		PED	Jun 21, 2007
	6514520	Jun 01, 2018	DP		
<u>DES Loratadine; CLARINEX</u>					
021312 001	4659716	Apr 21, 2006	DP U-427	NCE	Dec 21, 2006
	4659716*PED	Oct 21, 2006	U-427	PED	Jun 21, 2007
	4863931	Sep 15, 2008			
	4863931*PED	Mar 15, 2009			
	6100274	Jul 07, 2019	DP		
	6100274*PED	Jan 07, 2020			
<u>DES Loratadine; CLARINEX</u>					
021312 002	4659716	Apr 21, 2006	DP U-427	NCE	Dec 21, 2006
	4659716*PED	Oct 21, 2006	DP	PED	Jun 21, 2007
	6100274	Jul 07, 2019	DP		
	6100274*PED	Jan 07, 2020			
<u>DES Loratadine; Pseudoephedrine Sulfate; CLARINEX D 24 HOUR</u>					
021605 001	4659716	Apr 21, 2006	DP U-644	NCE	Dec 21, 2006
	4659716*PED	Oct 21, 2006		NC	Mar 03, 2008
	6100274	Jul 07, 2019	DP	PED	Jun 21, 2007
	6100274*PED	Jan 07, 2020			
<u>DES Mopressin Acetate; DDAVP</u>					
017922 001	5500413	Jun 29, 2013			
	5674850	Dec 23, 2013			
	5763407	Jun 29, 2013			
<u>DES Mopressin Acetate; DDAVP</u>					
017922 002	5500413	Jun 29, 2013			
	5674850	Dec 23, 2013			
	5763407	Jun 29, 2013			
<u>DES Mopressin Acetate; DDAVP</u>					
018938 001	5500413	Jun 29, 2013			
	5763407	Jun 29, 2013			
<u>DES Mopressin Acetate; DDAVP</u>					
018938 002	5500413	Jun 29, 2013			
	5763407	Jun 29, 2013			
<u>DES Mopressin Acetate; DDAVP</u>					
019955 001	5047398	Sep 10, 2008			
	5500413	Jun 29, 2013			
	5674850	Dec 23, 2013			
	5763407	Jun 29, 2013			
<u>DES Mopressin Acetate; DDAVP</u>					
019955 002	5047398	Sep 10, 2008			
	5500413	Jun 29, 2013			
	5674850	Dec 23, 2013			
	5763407	Jun 29, 2013			
<u>DES Mopressin Acetate; DDAVP (NEEDS NO REFRIGERATION)</u>					
017922 003	5482931	Jun 29, 2013			
	5500413	Jun 29, 2013			
	5674850	Dec 23, 2013			
	5763407	Jun 29, 2013			
<u>DES Mopressin Acetate; DES Mopressin Acetate</u>					
076470 001				PC	Dec 28, 2005
<u>DES Mopressin Acetate; DES Mopressin Acetate</u>					
076470 002				PC	Dec 28, 2005
<u>DES Ogestrel; Ethinyl Estradiol; Mircette</u>					
020713 001	RE35724	Oct 20, 2008			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEXMEDETOMIDINE; PRECEDEX</u>					
021038 001	4910214	Jul 15, 2008		U-421	
	6716867	Mar 31, 2019		U-572	
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; FOCALIN</u>					
021278 001	5908850	Dec 04, 2015		U-422	
	6355656	Dec 04, 2015			
	6528530	Dec 04, 2015	DS DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; FOCALIN</u>					
021278 002	5908850	Dec 04, 2015		U-422	
	6355656	Dec 04, 2015			
	6528530	Dec 04, 2015	DS DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; FOCALIN</u>					
021278 003	5908850	Dec 04, 2015		U-422	
	6355656	Dec 04, 2015			
	6528530	Dec 04, 2015	DS DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; FOCALIN XR</u>					
021802 001	5837284	Dec 04, 2015	DP	NDF	May 26, 2008
	5908850	Dec 04, 2015		U-678	
	6228398	Nov 01, 2019	DP	U-676	
	6528530	Dec 04, 2015	DP		
	6635284	Dec 04, 2015	DP	U-677	
	6730325	Nov 01, 2019	DP	U-676	
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; FOCALIN XR</u>					
021802 002	5837284	Dec 04, 2015	DP	NDF	May 26, 2008
	5908850	Dec 04, 2015		U-678	
	6228398	Nov 01, 2019	DP	U-676	
	6528530	Dec 04, 2015	DP		
	6635284	Dec 04, 2015	DP	U-677	
	6730325	Nov 01, 2019	DP	U-676	
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE; FOCALIN XR</u>					
021802 003	5837284	Dec 04, 2015	DP	NDF	May 26, 2008
	5908850	Dec 04, 2015		U-678	
	6228398	Nov 01, 2019	DP	U-676	
	6528530	Dec 04, 2015	DP		
	6635284	Dec 04, 2015	DP	U-677	
	6730325	Nov 01, 2019	DP	U-676	
<u>DEXRAZOXANE HYDROCHLORIDE; DEXRAZOXANE</u>					
076068 001				PC	Aug 27, 2005
<u>DEXRAZOXANE HYDROCHLORIDE; DEXRAZOXANE</u>					
076068 002				PC	Oct 19, 2005
<u>DEXRAZOXANE HYDROCHLORIDE; ZINECARD</u>					
020212 001	4963551	Dec 21, 2007			
	5242901	Sep 07, 2010		U-339	
<u>DEXRAZOXANE HYDROCHLORIDE; ZINECARD</u>					
020212 002	4963551	Dec 21, 2007			
	5242901	Sep 07, 2010		U-339	
<u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN; MUCINEX DM</u>					
021620 001	6372252	Apr 28, 2020	DP		
	6955821	Apr 28, 2020	DP	U-685	
<u>DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN; MUCINEX DM</u>					
021620 002	6372252	Apr 28, 2020	DP		
	6955821	Apr 28, 2020	DP	U-685	
<u>DEXTROMETHORPHAN POLISTIREX; DELSYM</u>					
018658 001	5980882	Apr 16, 2017	DP		
<u>DIAZEPAM; DIASTAT</u>					
020648 001	5462740	Sep 17, 2013			
<u>DIAZEPAM; DIASTAT</u>					
020648 002	5462740	Sep 17, 2013			
<u>DIAZEPAM; DIASTAT</u>					
020648 003	5462740	Sep 17, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DIAZEPAM; DIASTAT</u>					
020648 004	5462740	Sep 17, 2013			
<u>DIAZEPAM; DIASTAT</u>					
020648 005	5462740	Sep 17, 2013			
<u>DICLOFENAC SODIUM; DICLOFENAC SODIUM</u>					
020809 001	5603929	Nov 16, 2014	U-239		
	5653972	Nov 16, 2014	U-239		
<u>DICLOFENAC SODIUM; SOLARAZE</u>					
021005 001	5639738	Jun 17, 2014	U-402		
	5792753	Aug 11, 2015			
	5852002	Jun 17, 2014	U-402		
	5914322	Aug 11, 2015			
	5929048	Jul 27, 2016	U-402		
	5985850	Nov 16, 2016			
<u>DICLOFENAC SODIUM; VOLTAREN</u>					
020037 001	4829088	Apr 14, 2007			
	4960799	Oct 03, 2007			
<u>DICLOFENAC SODIUM; MISOPROSTOL; ARTHROTEC</u>					
020607 001	5601843	Feb 11, 2014			
	5698225	May 03, 2010	U-392		
<u>DICLOFENAC SODIUM; MISOPROSTOL; ARTHROTEC</u>					
020607 002	5601843	Feb 11, 2014			
	5698225	May 03, 2010	U-392		
<u>DIDANOSINE; VIDEX</u>					
020154 002	4861759	Aug 29, 2006	U-248		
	4861759*PED	Mar 01, 2007	U-248		
	5254539	Aug 29, 2006	U-248		
	5254539*PED	Mar 01, 2007	U-248		
	5616566	Aug 29, 2006	U-180		
	5616566*PED	Mar 01, 2007	U-180		
	5880106	Jul 22, 2011			
	5880106*PED	Jan 22, 2012			
<u>DIDANOSINE; VIDEX</u>					
020154 003	4861759	Aug 29, 2006	U-248		
	4861759*PED	Mar 01, 2007	U-248		
	5254539	Aug 29, 2006	U-248		
	5254539*PED	Mar 01, 2007	U-248		
	5616566	Aug 29, 2006	U-180		
	5616566*PED	Mar 01, 2007	U-180		
	5880106	Jul 22, 2011			
	5880106*PED	Jan 22, 2012			
<u>DIDANOSINE; VIDEX</u>					
020154 004	4861759	Aug 29, 2006	U-248		
	4861759*PED	Mar 01, 2007	U-248		
	5254539	Aug 29, 2006	U-248		
	5254539*PED	Mar 01, 2007	U-248		
	5616566	Aug 29, 2006	U-180		
	5616566*PED	Mar 01, 2007	U-180		
	5880106	Jul 22, 2011			
	5880106*PED	Jan 22, 2012			
<u>DIDANOSINE; VIDEX</u>					
020154 005	4861759	Aug 29, 2006	U-248		
	4861759*PED	Mar 01, 2007	U-248		
	5254539	Aug 29, 2006	U-248		
	5254539*PED	Mar 01, 2007	U-248		
	5616566	Aug 29, 2006	U-180		
	5616566*PED	Mar 01, 2007	U-180		
	5880106	Jul 22, 2011			
	5880106*PED	Jan 22, 2012			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DIDANOSINE; VIDEX</u>					
020154 006	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
	5616566	Aug 29, 2006		U-180	
	5616566*PED	Mar 01, 2007		U-180	
	5880106	Jul 22, 2011			
	5880106*PED	Jan 22, 2012			
<u>DIDANOSINE; VIDEX</u>					
020155 003	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
	5616566	Aug 29, 2006		U-180	
	5616566*PED	Mar 01, 2007		U-180	
<u>DIDANOSINE; VIDEX</u>					
020155 004	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
	5616566	Aug 29, 2006		U-180	
	5616566*PED	Mar 01, 2007		U-180	
<u>DIDANOSINE; VIDEX</u>					
020155 005	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
	5616566	Aug 29, 2006		U-180	
	5616566*PED	Mar 01, 2007		U-180	
<u>DIDANOSINE; VIDEX</u>					
020155 006	4861759	Aug 29, 2006		U-52	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
	5616566	Aug 29, 2006		U-180	
	5616566*PED	Mar 01, 2007		U-180	
<u>DIDANOSINE; VIDEX</u>					
020156 001	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
	5616566	Aug 29, 2006		U-180	
	5616566*PED	Mar 01, 2007		U-180	
<u>DIDANOSINE; VIDEX EC</u>					
021183 001	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
<u>DIDANOSINE; VIDEX EC</u>					
021183 002	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
<u>DIDANOSINE; VIDEX EC</u>					
021183 003	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
<u>DIDANOSINE; VIDEX EC</u>					
021183 004	4861759	Aug 29, 2006		U-248	
	4861759*PED	Mar 01, 2007		U-248	
	5254539	Aug 29, 2006		U-248	
	5254539*PED	Mar 01, 2007		U-248	
<u>DIHYDROERGOTAMINE MESYLATE; MIGRANAL</u>					
020148 001	5169849	Dec 08, 2009		U-227	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
DILTIAZEM HYDROCHLORIDE; CARDIZEM CD					
020062 001	4894240	Jan 16, 2007	U-3 U-107		
	5002776	Mar 26, 2008			
	5286497	May 20, 2011			
	5364620	Nov 14, 2011			
	5439689	Aug 08, 2012			
	5470584	May 20, 2011			
DILTIAZEM HYDROCHLORIDE; CARDIZEM CD					
020062 002	4894240	Jan 16, 2007	U-3 U-107		
	5002776	Mar 26, 2008			
	5286497	May 20, 2011			
	5364620	Nov 14, 2011			
	5439689	Aug 08, 2012			
	5470584	May 20, 2011			
DILTIAZEM HYDROCHLORIDE; CARDIZEM CD					
020062 003	4894240	Jan 16, 2007	U-3 U-107		
	5002776	Mar 26, 2008			
	5286497	May 20, 2011			
	5364620	Nov 14, 2011			
	5439689	Aug 08, 2012			
	5470584	May 20, 2011			
DILTIAZEM HYDROCHLORIDE; CARDIZEM CD					
020062 004	4894240	Jan 16, 2007	U-3 U-107		
	5002776	Mar 26, 2008			
	5286497	May 20, 2011			
	5364620	Nov 14, 2011			
	5439689	Aug 08, 2012			
	5470584	May 20, 2011			
DILTIAZEM HYDROCHLORIDE; CARDIZEM LA					
021392 001	5288505	Jun 26, 2011	DP	I-133	Apr 09, 2007
	5529791	Jun 25, 2013		NDF	Feb 06, 2006
	6923984	Feb 25, 2021			
DILTIAZEM HYDROCHLORIDE; CARDIZEM LA					
021392 002	5288505	Jun 26, 2011	DP	I-133	Apr 09, 2007
	5529791	Jun 25, 2013		NDF	Feb 06, 2006
	6923984	Feb 25, 2021			
DILTIAZEM HYDROCHLORIDE; CARDIZEM LA					
021392 003	5288505	Jun 26, 2011	DP	I-133	Apr 09, 2007
	5529791	Jun 25, 2013		NDF	Feb 06, 2006
	6923984	Feb 25, 2021			
DILTIAZEM HYDROCHLORIDE; CARDIZEM LA					
021392 004	5288505	Jun 26, 2011	DP	I-133	Apr 09, 2007
	5529791	Jun 25, 2013		NDF	Feb 06, 2006
	6923984	Feb 25, 2021			
DILTIAZEM HYDROCHLORIDE; CARDIZEM LA					
021392 005	5288505	Jun 26, 2011	DP	I-133	Apr 09, 2007
	5529791	Jun 25, 2013		NDF	Feb 06, 2006
	6923984	Feb 25, 2021			
DILTIAZEM HYDROCHLORIDE; CARDIZEM LA					
021392 006	5288505	Jun 26, 2011	DP	I-133	Apr 09, 2007
	5529791	Jun 25, 2013		NDF	Feb 06, 2006
	6923984	Feb 25, 2021			
DILTIAZEM HYDROCHLORIDE; DILACOR XR					
020092 001	4839177	Dec 09, 2006			
	5422123	Jun 06, 2012			
DILTIAZEM HYDROCHLORIDE; DILACOR XR					
020092 002	4839177	Dec 09, 2006			
	5422123	Jun 06, 2012			
DILTIAZEM HYDROCHLORIDE; DILACOR XR					
020092 003	4839177	Dec 09, 2006			
	5422123	Jun 06, 2012			
DILTIAZEM HYDROCHLORIDE; DILTIAZEM HCL					
020939 001	5288505	Jun 26, 2011			
	5529791	Jun 25, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DILTIAZEM HYDROCHLORIDE; DILTIAZEM HCL</u>					
020939 002	5288505	Jun 26, 2011			
	5529791	Jun 25, 2013			
<u>DILTIAZEM HYDROCHLORIDE; DILTIAZEM HCL</u>					
020939 003	5288505	Jun 26, 2011			
	5529791	Jun 25, 2013			
<u>DILTIAZEM HYDROCHLORIDE; DILTIAZEM HCL</u>					
020939 004	5288505	Jun 26, 2011			
	5529791	Jun 25, 2013			
<u>DILTIAZEM HYDROCHLORIDE; TIAZAC</u>					
020401 001	5529791	Jun 25, 2013			
<u>DILTIAZEM HYDROCHLORIDE; TIAZAC</u>					
020401 002	5529791	Jun 25, 2013			
<u>DILTIAZEM HYDROCHLORIDE; TIAZAC</u>					
020401 003	5529791	Jun 25, 2013			
<u>DILTIAZEM HYDROCHLORIDE; TIAZAC</u>					
020401 004	5529791	Jun 25, 2013			
<u>DILTIAZEM HYDROCHLORIDE; TIAZAC</u>					
020401 005	5529791	Jun 25, 2013			
<u>DILTIAZEM HYDROCHLORIDE; TIAZAC</u>					
020401 006	5529791	Jun 25, 2013			
<u>DILTIAZEM MALATE; TIAMATE</u>					
020506 001	4880631	Sep 24, 2007			
	4968507	Jul 25, 2006			
<u>DILTIAZEM MALATE; TIAMATE</u>					
020506 002	4880631	Sep 24, 2007			
	4968507	Jul 25, 2006			
<u>DILTIAZEM MALATE; TIAMATE</u>					
020506 003	4880631	Sep 24, 2007			
	4968507	Jul 25, 2006			
<u>DILTIAZEM MALATE; ENALAPRIL MALEATE; TECZEM</u>					
020507 001	4880631	Sep 24, 2007			
	4968507	Jul 25, 2006			
	4983598	Jan 08, 2008			
<u>DIMYRISTOYL LECITHIN; PERFLEXANE; IMAGENT</u>					
021191 001	5605673	Feb 25, 2014		NCE	May 31, 2007
	5626833	May 16, 2014	U-458		
	5639443	Jun 17, 2014			
	5695741	Dec 09, 2014	U-458		
	5720938	Feb 24, 2015			
	5798091	Aug 25, 2015	U-458		
	6280704	Jul 30, 2013			
	6280705	Jul 30, 2013			
	6287539	Jul 30, 2013			
<u>DINOPROSTONE; CERVIDIL</u>					
020411 001	4931288	Jan 16, 2007			
	5269321	Dec 14, 2010	U-110		
<u>DIPHENHYDRAMINE CITRATE; IBUPROFEN; ADVIL PM</u>					
021394 001				NC	Dec 21, 2008
<u>DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN POTASSIUM; ADVIL PM</u>					
021393 001				NC	Dec 21, 2008
<u>DIVALPROEX SODIUM; DEPAKOTE</u>					
018723 001	4988731	Jan 29, 2008			
	5212326	Jan 29, 2008			
<u>DIVALPROEX SODIUM; DEPAKOTE</u>					
018723 002	4988731	Jan 29, 2008			
	5212326	Jan 29, 2008			
<u>DIVALPROEX SODIUM; DEPAKOTE</u>					
018723 003	4988731	Jan 29, 2008			
	5212326	Jan 29, 2008			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DIVALPROEX SODIUM; DEPAKOTE</u>					
019680 001	4988731	Jan 29, 2008			
	5212326	Jan 29, 2008			
<u>DIVALPROEX SODIUM; DEPAKOTE CP</u>					
019794 001	4988731	Jan 29, 2008			
	5212326	Jan 29, 2008			
<u>DIVALPROEX SODIUM; DEPAKOTE CP</u>					
019794 002	4988731	Jan 29, 2008			
	5212326	Jan 29, 2008			
<u>DIVALPROEX SODIUM; DEPAKOTE ER</u>					
021168 001	4913906	Apr 03, 2007		I-482	Dec 06, 2008
	4988731	Jan 29, 2008			
	5212326	Jan 29, 2008	DS DP		
	6419953	Dec 18, 2018			
	6511678	Dec 18, 2018			
	6528090	Dec 18, 2018		DP	
	6528091	Dec 18, 2018		U-106	
	6713086	Dec 18, 2018		DP U-579	
	6720004	Dec 18, 2018		DP	
<u>DIVALPROEX SODIUM; DEPAKOTE ER</u>					
021168 002	4913906	Apr 03, 2007		I-482	Dec 06, 2008
	4988731	Jan 29, 2008			
	5212326	Jan 29, 2008	DS DP		
	6511678	Dec 18, 2018			
	6528090	Dec 18, 2018		DP	
	6713086	Dec 18, 2018		DP U-579	
	6720004	Dec 18, 2018		DP	
<u>DOCETAXEL; TAXOTERE</u>					
020449 001	4814470	May 14, 2010		I-436	Aug 18, 2007
	5438072	Nov 22, 2013		I-429	May 19, 2007
	5698582	Jul 03, 2012		I-379	Nov 27, 2005
	5714512	Jul 03, 2012			
<u>DOCOSANOL; ABREVA</u>					
020941 001	4874794	Apr 28, 2014		NCE	Jul 25, 2005
<u>DOFETILIDE; TIKOSYN</u>					
020931 001	4959366	Sep 25, 2012	DS DP	U-652	
	6124363	Oct 09, 2018			
<u>DOFETILIDE; TIKOSYN</u>					
020931 002	4959366	Sep 25, 2012	DS DP	U-652	
	6124363	Oct 09, 2018			
<u>DOFETILIDE; TIKOSYN</u>					
020931 003	4959366	Sep 25, 2012	DS DP	U-652	
	6124363	Oct 09, 2018			
<u>DOLASETRON MESYLATE MONOHYDRATE; ANZEMET</u>					
020623 001	4906755	Jul 02, 2011			
<u>DOLASETRON MESYLATE MONOHYDRATE; ANZEMET</u>					
020623 002	4906755	Jul 02, 2011			
<u>DOLASETRON MESYLATE MONOHYDRATE; ANZEMET</u>					
020624 001	4906755	Jul 02, 2011			
<u>DOLASETRON MESYLATE MONOHYDRATE; ANZEMET</u>					
020624 002	4906755	Jul 02, 2011			
<u>DOLASETRON MESYLATE MONOHYDRATE; ANZEMET</u>					
020624 003	4906755	Jul 02, 2011			
<u>DONEPEZIL HYDROCHLORIDE; ARICEPT</u>					
020690 001	4895841	Nov 25, 2010			
	5985864	Dec 30, 2016			
	6140321	Dec 30, 2016			
	6245911	Dec 01, 2018			
	6372760	Mar 31, 2019			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DONEPEZIL HYDROCHLORIDE; ARICEPT</u>					
020690 002	4895841	Nov 25, 2010			
	5985864	Dec 30, 2016			
	6140321	Dec 30, 2016			
	6245911	Dec 01, 2018			
	6372760	Mar 31, 2019			
<u>DORZOLAMIDE HYDROCHLORIDE; TRUSOPT</u>					
020408 001	4797413	Apr 28, 2008	DS DP U-103	M-38	Apr 15, 2007
	4797413*PED	Oct 28, 2008		PED	Oct 15, 2007
<u>DORZOLAMIDE HYDROCHLORIDE; TIMOLOL MALEATE; COSOPT</u>					
020869 001	4797413	Apr 28, 2008		U-103	
	4797413*PED	Oct 28, 2008			
	6248735	Apr 17, 2011			
	6316443	Apr 17, 2011	DP	U-561	
<u>DOXAZOSIN MESYLATE; CARDURA XL</u>					
021269 001				NDF	Feb 22, 2008
<u>DOXAZOSIN MESYLATE; CARDURA XL</u>					
021269 002	4837111	Mar 21, 2008	DP	NDF	Feb 22, 2008
<u>DOXERCALCIFEROL; HECTOROL</u>					
020862 001	5602116	Feb 11, 2014		U-278	Apr 23, 2007
	5707980	Aug 17, 2010		U-278	
	5861386	Aug 02, 2008		U-278	
	5869473	Aug 02, 2008		U-278	
	6903083	Jul 18, 2021	DS DP		
<u>DOXERCALCIFEROL; HECTOROL</u>					
020862 002	5869473	Aug 02, 2008		U-278	Apr 23, 2007
	6903083	Jul 18, 2021	DS DP		
<u>DOXERCALCIFEROL; HECTOROL</u>					
021027 001	5602116	Feb 11, 2014		U-321	
	5707980	Aug 17, 2010		U-321	
	6903083	Jul 18, 2021	DS DP		
<u>DOXORUBICIN HYDROCHLORIDE; DOXIL</u>					
050718 001				ODE	Jun 28, 2006
<u>DRONABINOL; MARINOL</u>					
018651 001	6703418	Feb 26, 2011		U-563	
<u>DRONABINOL; MARINOL</u>					
018651 002	6703418	Feb 26, 2011		U-563	
<u>DRONABINOL; MARINOL</u>					
018651 003	6703418	Feb 26, 2011		U-563	
<u>DROSPIRENONE; ESTRADIOL; ANGELIQ</u>					
021355 002	6933395	Aug 11, 2017	DS	NP	Sep 28, 2008
<u>DROSPIRENONE; ETHINYL ESTRADIOL; YASMIN</u>					
021098 001	5569652	Oct 29, 2013		U-1	
	6787531	Aug 31, 2020		DP	
	6933395	Aug 11, 2017	DS		
<u>DULOXETINE HYDROCHLORIDE; CYMBALTA</u>					
021427 001	5023269	Jun 11, 2008	DS DP	I-470	Sep 03, 2007
	5508276	Jul 18, 2014	DP	NCE	Aug 03, 2009
<u>DULOXETINE HYDROCHLORIDE; CYMBALTA</u>					
021427 002	5023269	Jun 11, 2008	DS DP	I-470	Sep 03, 2007
	5508276	Jul 18, 2014	DP	NCE	Aug 03, 2009
<u>DULOXETINE HYDROCHLORIDE; CYMBALTA</u>					
021427 004	5023269	Jun 11, 2008	DS DP	I-470	Sep 03, 2007
	5508276	Jul 18, 2014	DP	NCE	Aug 03, 2009
<u>DUTASTERIDE; AVODART</u>					
021319 001	5565467	Oct 15, 2013		NCE	Nov 20, 2006
	5846976	Sep 17, 2013		U-476	
	5998427	Sep 17, 2013	DS	U-477	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EFAVIRENZ; SUSTIVA</u>					
020972 001	5519021	May 21, 2013			
	5663169	Sep 02, 2014		U-257	
	5811423	Aug 07, 2012		U-256	
	6238695	Apr 06, 2019			
	6555133	Apr 06, 2019		U-248	
	6639071	Feb 14, 2018	DS		
	6939964	Jan 20, 2018	DS		
<u>EFAVIRENZ; SUSTIVA</u>					
020972 002	5519021	May 21, 2013			
	5663169	Sep 02, 2014		U-257	
	5811423	Aug 07, 2012		U-256	
	6238695	Apr 06, 2019			
	6555133	Apr 06, 2019		U-248	
	6639071	Feb 14, 2018	DS		
	6939964	Jan 20, 2018	DS		
<u>EFAVIRENZ; SUSTIVA</u>					
020972 003	5519021	May 21, 2013			
	5663169	Sep 02, 2014		U-257	
	5811423	Aug 07, 2012		U-256	
	6238695	Apr 06, 2019			
	6555133	Apr 06, 2019		U-248	
	6639071	Feb 14, 2018	DS		
	6939964	Jan 20, 2018	DS		
<u>EFAVIRENZ; SUSTIVA</u>					
021360 001	5519021	May 21, 2013			
	5663169	Sep 02, 2014			
	5811423	Aug 07, 2012		U-256	
	6639071	Feb 14, 2018	DS		
	6939964	Jan 20, 2018	DS		
<u>EFAVIRENZ; SUSTIVA</u>					
021360 002	5519021	May 21, 2013			
	5663169	Sep 02, 2014		U-248	
	5811423	Aug 07, 2012		U-256	
	6639071	Feb 14, 2018	DS		
	6939964	Jan 20, 2018	DS		
<u>EFLORNITHINE HYDROCHLORIDE; VANIQA</u>					
021145 001	5648394	Jul 15, 2014		U-334	
<u>ELETRIPTAN HYDROBROMIDE; RELPAX</u>					
021016 001	5545644	Aug 13, 2013		NCE	Dec 26, 2007
<u>ELETRIPTAN HYDROBROMIDE; RELPAX</u>					
021016 002	5545644	Aug 13, 2013		NCE	Dec 26, 2007
<u>EMEDASTINE DIFUMARATE; EMADINE</u>					
020706 001	4430343	Aug 14, 2005		U-403	
	5441958	Dec 08, 2013		U-404	
<u>EMTRICITABINE; EMTRIVA</u>					
021500 001	5210085	May 11, 2010		NCE	Jul 02, 2008
	5814639	Sep 29, 2015			
	5914331	Sep 29, 2015			
	6642245	Nov 04, 2020		U-541	
	6703396	Mar 09, 2021	DS DP		
<u>EMTRICITABINE; EMTRIVA</u>					
021896 001	5210085	May 11, 2010		U-257	Sep 27, 2008
	5814639	Sep 29, 2015	DS DP	NDF	
	5914331	Sep 29, 2015	DS	NCE	Jul 02, 2008
	6642245	Nov 04, 2020		U-257	
	6703396	Mar 09, 2021	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE; TRUVADA</u>								
021752 001	5210085	May 11, 2010				U-541	NCE	Jul 02, 2008
	5210085	May 11, 2010			U-248	NCE		
	5814639	Sep 29, 2015	DS	DP				
	5914331	Sep 29, 2015	DS	DP	U-248			
	5922695	Jul 25, 2017	DS		U-541			
	5922695	Jul 25, 2017	DS		U-248			
	5935946	Jul 25, 2017	DS	DP	U-541			
	5977089	Jul 25, 2017	DS	DP	U-541			
	5977089	Jul 25, 2017	DS	DP	U-248			
	6043230	Jul 25, 2017		DP	U-248			
	6043230	Jul 25, 2017		DP	U-541			
	6642245	Nov 04, 2020			U-541			
	6642245	Nov 04, 2020			U-248			
	6703396	Mar 09, 2021	DS	DP				
<u>ENALAPRIL MALEATE; FELODIPINE; LEXXEL</u>								
020668 001	4703038	Oct 07, 2005				U-3		
	4803081	Apr 03, 2007						
	4803081*PED	Oct 03, 2007						
<u>ENALAPRIL MALEATE; FELODIPINE; LEXXEL</u>								
020668 002	4703038	Oct 07, 2005				U-3		
	4803081	Apr 03, 2007						
	4803081*PED	Oct 03, 2007						
<u>ENFUVIRTIDE; FUZEON</u>								
021481 001	5464933	Jun 07, 2013					NCE	Mar 13, 2008
	6133418	Jun 07, 2013						
	6475491	Jun 07, 2015			U-248			
<u>ENOXAPARIN SODIUM; LOVENOX</u>								
020164 009	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENOXAPARIN SODIUM; LOVENOX (PRESERVATIVE FREE)</u>								
020164 001	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENOXAPARIN SODIUM; LOVENOX (PRESERVATIVE FREE)</u>								
020164 002	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENOXAPARIN SODIUM; LOVENOX (PRESERVATIVE FREE)</u>								
020164 003	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENOXAPARIN SODIUM; LOVENOX (PRESERVATIVE FREE)</u>								
020164 004	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENOXAPARIN SODIUM; LOVENOX (PRESERVATIVE FREE)</u>								
020164 005	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENOXAPARIN SODIUM; LOVENOX (PRESERVATIVE FREE)</u>								
020164 006	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENOXAPARIN SODIUM; LOVENOX (PRESERVATIVE FREE)</u>								
020164 007	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENOXAPARIN SODIUM; LOVENOX (PRESERVATIVE FREE)</u>								
020164 008	5389618	Feb 14, 2012	DS	DP	U-545			
	RE38743	Feb 14, 2012	DS	DP	U-545			
<u>ENTACAPONE; COMTAN</u>								
020796 001	4963590	Nov 27, 2007						
	5112861	May 12, 2009						
	5135950	Oct 31, 2010						
	5446194	Oct 19, 2013	DS					
	6599530	Sep 14, 2018			DP			
<u>ENTECAVIR; BARACLUDE</u>								
021797 001	5206244	Oct 18, 2010	DS			NCE	Mar 29, 2010	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ENTECAVIR; BARACLUDE</u>					
021797 002	5206244	Oct 18, 2010	DS	NCE	Mar 29, 2010
<u>ENTECAVIR; BARACLUDE</u>					
021798 001	5206244	Oct 18, 2010	DS	NCE	Mar 29, 2010
<u>EPINASTINE HYDROCHLORIDE; ELESTAT</u>					
021565 001				NCE	Oct 15, 2008
<u>EPINEPHRINE; LIDOCAINE HYDROCHLORIDE; LIDOCAINE HCL AND EPINEPHRINE</u>					
021486 001				NP	Oct 26, 2007
<u>EPINEPHRINE; LIDOCAINE HYDROCHLORIDE; LIDOSITE TOPICAL SYSTEM KIT</u>					
021504 001	5246418	Sep 30, 2013	DS DP	NP	May 04, 2007
	5873850	Sep 30, 2013	DS DP		
	6377847	Sep 30, 2013	DS DP		
	6385488	Sep 30, 2013	DS DP		
	6629968	Jun 30, 2020	DS DP		
	6635045	Jun 29, 2021	DS DP		
	6862473	Sep 30, 2013	DP		
<u>EPIRUBICIN HYDROCHLORIDE; ELLENCE</u>					
050778 001				ODE	Sep 15, 2006
<u>EPLERENONE; INSPRA</u>					
021437 001	4559332	Apr 09, 2006	DS DP	I-407	Oct 07, 2006
	6410054	Dec 08, 2019		NCE	Sep 27, 2007
	6410054	Dec 08, 2019			
	6410524	Nov 05, 2019			
	6495165	Dec 08, 2019			
	6495165	Dec 08, 2019			
	6534093	Dec 08, 2019			
	6534093	Dec 08, 2019			
	6558707	Dec 08, 2019	DP		
	6747020	Nov 05, 2019			
	6863902	Apr 10, 2020	DP		
<u>EPLERENONE; INSPRA</u>					
021437 002	4559332	Apr 09, 2006	DS DP	I-407	Oct 07, 2006
	6410054	Dec 08, 2019		NCE	Sep 27, 2007
	6410054	Dec 08, 2019			
	6410524	Nov 05, 2019			
	6495165	Dec 08, 2019			
	6495165	Dec 08, 2019			
	6534093	Dec 08, 2019			
	6534093	Dec 08, 2019			
	6558707	Dec 08, 2019	DP		
	6747020	Nov 05, 2019			
	6863902	Apr 10, 2020	DP		
<u>EPLERENONE; INSPRA</u>					
021437 003	4559332	Apr 09, 2006	DS DP	I-407	Oct 07, 2006
	6410054	Dec 08, 2019		NCE	Sep 27, 2007
	6410054	Dec 08, 2019			
	6410524	Nov 05, 2019			
	6495165	Dec 08, 2019			
	6495165	Dec 08, 2019			
	6534093	Dec 08, 2019			
	6534093	Dec 08, 2019			
	6558707	Dec 08, 2019	DP		
	6747020	Nov 05, 2019			
	6863902	Apr 10, 2020	DP		
<u>EPOPROSTENOL SODIUM; FLOLAN</u>					
020444 001	4883812	May 12, 2006		U-185	ODE
					Apr 14, 2007
<u>EPOPROSTENOL SODIUM; FLOLAN</u>					
020444 002	4883812	May 12, 2006		U-185	ODE
					Apr 14, 2007
<u>EPROSARTAN MESYLATE; TEVETEN</u>					
020738 004	5185351	Feb 09, 2010		U-3	
	5656650	Aug 12, 2014		U-3	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EPROSARTAN MESYLATE; TEVETEN</u>					
020738 005	5185351	Feb 09, 2010	U-3		
	5656650	Aug 12, 2014	U-3		
<u>EPROSARTAN MESYLATE; TEVETEN</u>					
020738 006	5185351	Feb 09, 2010	U-3		
	5656650	Aug 12, 2014	U-3		
<u>EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE; TEVETEN HCT</u>					
021268 001	5185351	Feb 09, 2010	U-3		
	5656650	Aug 12, 2014	U-3		
<u>EPROSARTAN MESYLATE; HYDROCHLOROTHIAZIDE; TEVETEN HCT</u>					
021268 002	5185351	Feb 09, 2010	U-3		
	5656650	Aug 12, 2014	U-3		
<u>EPTIFIBATIDE; INTEGRILIN</u>					
020718 001	5686570	Nov 11, 2014			
	5747447	May 05, 2015			
	5756451	Nov 11, 2014			
	5807825	Sep 15, 2015	U-244		
	5968902	Jun 02, 2015	U-453		
<u>EPTIFIBATIDE; INTEGRILIN</u>					
020718 002	5686570	Nov 11, 2014			
	5747447	May 05, 2015			
	5756451	Nov 11, 2014			
	5807825	Sep 15, 2015	U-244		
	5968902	Jun 02, 2015	U-453		
<u>ERLOTINIB HYDROCHLORIDE; TARCEVA</u>					
021743 001	5747498	Mar 30, 2015	DS DP U-659	I-473	Nov 02, 2008
	6900221	Nov 09, 2020	DS DP U-659	NCE	Nov 18, 2009
<u>ERLOTINIB HYDROCHLORIDE; TARCEVA</u>					
021743 002	5747498	Mar 30, 2015	DS DP U-659	I-473	Nov 02, 2008
	6900221	Nov 09, 2020	DS DP U-659	NCE	Nov 18, 2009
<u>ERLOTINIB HYDROCHLORIDE; TARCEVA</u>					
021743 003	5747498	Mar 30, 2015	DS DP U-659	I-473	Nov 02, 2008
	6900221	Nov 09, 2020	DS DP U-659	NCE	Nov 18, 2009
<u>ERTAPENEM SODIUM; INVANZ</u>					
021337 001	5478820	Feb 02, 2013	DS DP U-160	I-476	Oct 14, 2008
	5478820*PED	Aug 02, 2013		NPP	May 18, 2008
	5652233	Feb 02, 2013	DS DP U-160	NCE	Nov 21, 2006
	5652233*PED	Aug 02, 2013		PED	Nov 18, 2008
	5952323	May 15, 2017	DP	PED	May 21, 2007
	5952323*PED	Nov 15, 2017			
<u>ESCITALOPRAM OXALATE; LEXAPRO</u>					
021323 001	6916941	Jul 25, 2022	DS DP	I-366	Aug 29, 2005
	6916941*PED	Jan 25, 2023	DS DP	I-251	Dec 18, 2006
	RE34712	Jun 08, 2009		NP	Aug 14, 2005
	RE34712*PED	Dec 08, 2009		PED	Mar 01, 2006
				PED	Feb 14, 2006
<u>ESCITALOPRAM OXALATE; LEXAPRO</u>					
021323 002	6916941	Jul 25, 2022	DS DP	I-366	Aug 29, 2005
	6916941*PED	Jan 25, 2023	DS DP	I-251	Dec 18, 2006
	RE34712	Jun 08, 2009		NP	Aug 14, 2005
	RE34712*PED	Dec 08, 2009		PED	Mar 01, 2006
				PED	Feb 14, 2006
<u>ESCITALOPRAM OXALATE; LEXAPRO</u>					
021323 003	6916941	Jul 25, 2022	DS DP	I-366	Aug 29, 2005
	6916941*PED	Jan 25, 2023	DS DP	I-251	Dec 18, 2006
	RE34712	Jun 08, 2009		NP	Aug 14, 2005
	RE34712*PED	Dec 08, 2009		PED	Mar 01, 2006
				PED	Feb 14, 2006

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESCITALOPRAM OXALATE; LEXAPRO</u>					
021365 001	RE34712	Jun 08, 2009		I-366	Aug 29, 2005
	RE34712*PED	Dec 08, 2009		I-251	Dec 18, 2006
				NP	Aug 14, 2005
				PED	Mar 01, 2006
				PED	Feb 14, 2006
<u>ESMOLOL HYDROCHLORIDE; BREVIBLOC</u>					
019386 002	5017609	May 21, 2008			
	5017609*PED	Nov 21, 2008			
<u>ESMOLOL HYDROCHLORIDE; BREVIBLOC</u>					
019386 003	5017609	May 21, 2008			
	5017609*PED	Nov 21, 2008			
<u>ESMOLOL HYDROCHLORIDE; BREVIBLOC</u>					
019386 006	6310094	Jan 12, 2021			
	6310094*PED	Jul 12, 2021			
	6528540	Jan 12, 2021			
	6528540*PED	Jul 12, 2021			
<u>ESMOLOL HYDROCHLORIDE; BREVIBLOC</u>					
019386 007	6310094	Jan 12, 2021			
	6310094*PED	Jul 12, 2021			
	6528540	Jan 12, 2021			
	6528540*PED	Jul 12, 2021			
<u>ESMOLOL HYDROCHLORIDE; BREVIBLOC DOUBLE STRENGTH IN PLASTIC CONTAINER</u>					
019386 005	6310094	Jan 12, 2021			
	6310094*PED	Jul 12, 2021			
	6528540	Jan 12, 2021			
	6528540*PED	Jul 12, 2021			
<u>ESMOLOL HYDROCHLORIDE; BREVIBLOC IN PLASTIC CONTAINER</u>					
019386 004	6310094	Jan 12, 2021			
	6310094*PED	Jul 12, 2021			
	6528540	Jan 12, 2021			
	6528540*PED	Jul 12, 2021			
<u>ESOMEPRAZOLE MAGNESIUM; NEXIUM</u>					
021153 001	4738974	Apr 19, 2006	DS	DP	U-373
	4738974	Apr 19, 2006	DS	DP	U-635
	4738974*PED	Oct 19, 2006			U-373
	4786505	Apr 20, 2007			U-373
	4786505*PED	Oct 20, 2007			U-373
	4853230	Apr 20, 2007			U-373
	4853230*PED	Oct 20, 2007			U-373
	5690960	Nov 25, 2014			U-373
	5690960*PED	May 25, 2015			U-373
	5714504	Feb 03, 2015			U-373
	5714504*PED	Aug 03, 2015			U-373
	5877192	May 27, 2014			U-373
	5877192*PED	Nov 27, 2014			U-373
	5900424	May 04, 2016			U-373
	5900424*PED	Nov 04, 2016			U-373
	6147103	Oct 09, 2018			
	6147103*PED	Apr 09, 2019			
	6166213	Oct 09, 2018			
	6166213*PED	Apr 09, 2019			
	6191148	Oct 09, 2018			
	6191148*PED	Apr 09, 2019			
	6369085	May 25, 2018			
	6369085*PED	Nov 25, 2018			
	6428810	Nov 03, 2019			U-469
	6428810*PED	May 03, 2020			U-469
	6875872	May 27, 2014	DS		
	6875872*PED	Nov 27, 2014			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESOMEPRAZOLE MAGNESIUM; NEXIUM</u>					
021153 002	4738974	Apr 19, 2006	DS DP	U-373	
	4738974	Apr 19, 2006	DS DP	U-635	
	4738974*PED	Oct 19, 2006		U-373	
	4786505	Apr 20, 2007		U-373	
	4786505*PED	Oct 20, 2007		U-373	
	4853230	Apr 20, 2007		U-373	
	4853230*PED	Oct 20, 2007		U-373	
	5690960	Nov 25, 2014		U-373	
	5690960*PED	May 25, 2015		U-373	
	5714504	Feb 03, 2015		U-373	
	5714504*PED	Aug 03, 2015		U-373	
	5877192	May 27, 2014		U-373	
	5877192*PED	Nov 27, 2014		U-373	
	5900424	May 04, 2016		U-373	
	5900424*PED	Nov 04, 2016		U-373	
	6147103	Oct 09, 2018			
	6147103*PED	Apr 09, 2019			
	6166213	Oct 09, 2018			
	6166213*PED	Apr 09, 2019			
	6191148	Oct 09, 2018			
	6191148*PED	Apr 09, 2019			
	6369085	May 25, 2018			
	6369085*PED	Nov 25, 2018			
	6428810	Nov 03, 2019		U-469	
	6428810*PED	May 03, 2020		U-469	
	6875872	May 27, 2014	DS		
	6875872*PED	Nov 27, 2014			
<u>ESOMEPRAZOLE SODIUM; NEXIUM IV</u>					
021689 001	5877192	May 27, 2014		U-643	NE Mar 31, 2008
	5877192*PED	Nov 27, 2014			
	6143771	May 27, 2014	DP	U-643	
<u>ESOMEPRAZOLE SODIUM; NEXIUM IV</u>					
021689 002	5877192	May 27, 2014		U-643	NE Mar 31, 2008
	5877192*PED	Nov 27, 2014			
	6143771	May 27, 2014	DP	U-643	
<u>ESTRADIOL; ALORA</u>					
020655 001	5122383	May 17, 2011			
	5164190	Dec 11, 2010			
	5212199	May 17, 2011			
	5227169	May 17, 2011			
<u>ESTRADIOL; ALORA</u>					
020655 002	5122383	May 17, 2011			
	5164190	Dec 11, 2010			
	5212199	May 17, 2011			
	5227169	May 17, 2011			
<u>ESTRADIOL; ALORA</u>					
020655 003	5122383	May 17, 2011			
	5164190	Dec 11, 2010			
	5212199	May 17, 2011			
	5227169	May 17, 2011			
<u>ESTRADIOL; ALORA</u>					
020655 004	5122383	May 17, 2011			
	5164190	Dec 11, 2010			
	5212199	May 17, 2011			
	5227169	May 17, 2011			
<u>ESTRADIOL; CLIMARA</u>					
020375 001	5223261	Jun 29, 2010			
<u>ESTRADIOL; CLIMARA</u>					
020375 002	5223261	Jun 29, 2010			
<u>ESTRADIOL; CLIMARA</u>					
020375 003	5223261	Jun 29, 2010			
<u>ESTRADIOL; CLIMARA</u>					
020375 004	5223261	Jun 29, 2010			
<u>ESTRADIOL; CLIMARA</u>					
020375 005	5223261	Jun 29, 2010			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESTRADIOL; CLIMARA</u>					
020375 006	5223261	Jun 29, 2010			
<u>ESTRADIOL; ESCLIM</u>					
020847 001	4842864	Mar 25, 2008			
<u>ESTRADIOL; ESCLIM</u>					
020847 002	4842864	Mar 25, 2008			
<u>ESTRADIOL; ESCLIM</u>					
020847 003	4842864	Mar 25, 2008			
<u>ESTRADIOL; ESCLIM</u>					
020847 004	4842864	Mar 25, 2008			
<u>ESTRADIOL; ESCLIM</u>					
020847 005	4842864	Mar 25, 2008			
<u>ESTRADIOL; ESTRING</u>					
020472 001	4871543	Jun 12, 2007	U-306		
	5188835	Jun 12, 2007	U-306		
<u>ESTRADIOL; ESTROGEL</u>					
021166 001				NDF	Feb 09, 2007
<u>ESTRADIOL; FEMPATCH</u>					
020417 001	4906463	Mar 06, 2007			
	5006342	Apr 09, 2008			
<u>ESTRADIOL; MENOSTAR</u>					
021674 001	5223261	Jun 29, 2010	DP	U-594	NP Jun 08, 2007
	5891868	Nov 21, 2017	DP	U-594	
	6692763	Nov 21, 2017	DP	U-594	
<u>ESTRADIOL; VIVELLE</u>					
020323 001	4814168	Mar 04, 2008			
	4994267	Mar 04, 2008			
	4994278	Mar 04, 2008			
	5300291	Apr 05, 2011			
<u>ESTRADIOL; VIVELLE</u>					
020323 002	4814168	Mar 04, 2008			
	4994267	Mar 04, 2008			
	4994278	Mar 04, 2008			
	5300291	Apr 05, 2011			
<u>ESTRADIOL; VIVELLE</u>					
020323 003	4814168	Mar 04, 2008			
	4994267	Mar 04, 2008			
	4994278	Mar 04, 2008			
	5300291	Apr 05, 2011			
<u>ESTRADIOL; VIVELLE</u>					
020323 004	4814168	Mar 04, 2008			
	4994267	Mar 04, 2008			
	4994278	Mar 04, 2008			
	5300291	Apr 05, 2011			
<u>ESTRADIOL; VIVELLE</u>					
020323 005	4814168	Mar 04, 2008			
	4994267	Mar 04, 2008			
	4994278	Mar 04, 2008			
	5300291	Apr 05, 2011			
<u>ESTRADIOL; VIVELLE-DOT</u>					
020538 005	5474783	Dec 12, 2012			
	5656286	Aug 12, 2014			
	5958446	Dec 12, 2012			
	6024976	Jan 07, 2014			
<u>ESTRADIOL; VIVELLE-DOT</u>					
020538 006	5474783	Dec 12, 2012			
	5656286	Aug 12, 2014			
	5958446	Dec 12, 2012			
	6024976	Jan 07, 2014			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESTRADIOL; VIVELLE-DOT</u>					
020538 007	5474783	Dec 12, 2012			
	5656286	Aug 12, 2014			
	5958446	Dec 12, 2012			
	6024976	Jan 07, 2014			
<u>ESTRADIOL; VIVELLE-DOT</u>					
020538 008	5474783	Dec 12, 2012			
	5656286	Aug 12, 2014			
	5958446	Dec 12, 2012			
	6024976	Jan 07, 2014			
<u>ESTRADIOL ACETATE; FEMRING</u>					
021367 001	5855906	Dec 19, 2015	U-508	NP	Mar 20, 2006
<u>ESTRADIOL ACETATE; FEMRING</u>					
021367 002	5855906	Dec 19, 2015	U-508	NP	Mar 20, 2006
<u>ESTRADIOL ACETATE; FEMTRACE</u>					
021633 001	6962908	Dec 21, 2021	DP	NDF	Aug 20, 2007
<u>ESTRADIOL ACETATE; FEMTRACE</u>					
021633 002	6962908	Dec 21, 2021	DP	NDF	Aug 20, 2007
<u>ESTRADIOL ACETATE; FEMTRACE</u>					
021633 003	6962908	Dec 21, 2021	DP	NDF	Aug 20, 2007
<u>ESTRADIOL HEMIHYDRATE; ESTRASORB</u>					
021371 001	5629021	Jan 31, 2015	DP	NDF	Oct 09, 2006
<u>ESTRADIOL; LEVONORGESTREL; CLIMARA PRO</u>					
021258 001	5252334	Oct 12, 2010	DP	I-483	Dec 27, 2008
	5393529	Feb 28, 2012	DP	NC	Nov 21, 2006
	5676968	Oct 14, 2014	DP		
	5770219	Oct 12, 2010	DP		
<u>ESTRADIOL; NORETHINDRONE ACETATE; ACTIVELLA</u>					
020907 001	RE36247	May 02, 2006			
<u>ESTRADIOL; NORETHINDRONE ACETATE; COMBIPATCH</u>					
020870 001	5474783	Dec 12, 2012			
	5656286	Aug 12, 2014			
	5958446	Dec 12, 2012			
	6024976	Jan 07, 2014			
<u>ESTRADIOL; NORETHINDRONE ACETATE; COMBIPATCH</u>					
020870 002	5474783	Dec 12, 2012			
	5656286	Aug 12, 2014			
	5958446	Dec 12, 2012			
	6024976	Jan 07, 2014			
<u>ESTRADIOL; NORGESTIMATE; PREFEST</u>					
021040 001	5108995	Apr 28, 2009	U-311		
	5382573	Jan 17, 2012			
	6747019	Mar 20, 2020	U-311		
<u>ESTROGENS, CONJUGATED; PREMARIN</u>					
004782 001	5210081	Feb 26, 2012			
<u>ESTROGENS, CONJUGATED; PREMARIN</u>					
004782 002	5210081	Feb 26, 2012			
<u>ESTROGENS, CONJUGATED; PREMARIN</u>					
004782 003	5210081	Feb 26, 2012		D-82	Jul 16, 2006
<u>ESTROGENS, CONJUGATED; PREMARIN</u>					
004782 004	5210081	Feb 26, 2012			
<u>ESTROGENS, CONJUGATED; PREMARIN</u>					
004782 005	5210081	Feb 26, 2012			
<u>ESTROGENS, CONJUGATED; PREMARIN</u>					
004782 006				D-82	Jul 16, 2006
<u>ESTROGENS, CONJUGATED SYNTHETIC A; CENESTIN</u>					
020992 001	5908638	Jul 26, 2015			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ESTROGENS, CONJUGATED SYNTHETIC A; CENESTIN					
020992 002	5908638	Jul 26, 2015		D-85	Feb 05, 2007
ESTROGENS, CONJUGATED SYNTHETIC A; CENESTIN					
020992 003	5908638	Jul 26, 2015		D-85	Feb 05, 2007
ESTROGENS, CONJUGATED SYNTHETIC A; CENESTIN					
020992 004	5908638	Jul 26, 2015		D-85	Feb 05, 2007
ESTROGENS, CONJUGATED SYNTHETIC B; ENJUVIA					
021443 001	6660726	Mar 08, 2021	DS DP U-284	NP	Dec 20, 2007
	6660726	Mar 08, 2021	DS DP U-196		
	6855703	Feb 12, 2021	DS DP U-284		
ESTROGENS, CONJUGATED SYNTHETIC B; ENJUVIA					
021443 002	6660726	Mar 08, 2021	DS DP U-284	NP	Dec 20, 2007
	6660726	Mar 08, 2021	DS DP U-196		
	6855703	Feb 12, 2021	DS DP U-284		
ESTROGENS, CONJUGATED SYNTHETIC B; ENJUVIA					
021443 003	6660726	Mar 08, 2021	DS DP U-284	NP	May 10, 2007
	6660726	Mar 08, 2021	DS DP U-196		
	6855703	Feb 12, 2021	DS DP U-284		
ESTROGENS, CONJUGATED SYNTHETIC B; ENJUVIA					
021443 004	6660726	Mar 08, 2021	DS DP U-284	NP	May 10, 2007
	6660726	Mar 08, 2021	DS DP U-196		
	6855703	Feb 12, 2021	DS DP U-284		
ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE; PREMPHASE (PREMARIN;CYCRIN 14/14)					
020303 002	5210081	Feb 26, 2012			
ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE; PREMPHASE 14/14					
020527 002	5547948	Jan 17, 2015			
	RE36247	May 02, 2006			
ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE; PREMPRO					
020527 001	5547948	Jan 17, 2015		D-79	Mar 12, 2006
	RE36247	May 02, 2006	U-284		
ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE; PREMPRO					
020527 003	5547948	Jan 17, 2015		D-79	Mar 12, 2006
	RE36247	May 02, 2006	U-284		
ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE; PREMPRO					
020527 004	5547948	Jan 17, 2015		D-81	Jun 04, 2006
	RE36247	May 02, 2006		D-79	Mar 12, 2006
				NP	Mar 12, 2006
ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE; PREMPRO					
020527 005	5547948	Jan 17, 2015		D-81	Jun 04, 2006
	RE36247	May 02, 2006		NS	Jun 04, 2006
ESTROGENS, CONJUGATED; MEDROXYPROGESTERONE ACETATE; PREMPRO (PREMARIN;CYCRIN)					
020303 001	5210081	Feb 26, 2012			
	RE36247	May 02, 2006	U-284		
ESTROGENS, CONJUGATED; MEPROBAMATE; PMB 200					
010971 005	5210081	Feb 26, 2012			
ESTROGENS, CONJUGATED; MEPROBAMATE; PMB 400					
010971 003	5210081	Feb 26, 2012			
ESZOPICLONE; LUNESTA					
021476 001	6319926	Jan 16, 2012		NCE	Dec 15, 2009
	6444673	Jan 16, 2012	DS DP		
	6864257	Aug 30, 2012	U-629		
ESZOPICLONE; LUNESTA					
021476 002	6319926	Jan 16, 2012		NCE	Dec 15, 2009
	6444673	Jan 16, 2012	DS DP		
	6864257	Aug 30, 2012	U-629		
ESZOPICLONE; LUNESTA					
021476 003	6319926	Jan 16, 2012		NCE	Dec 15, 2009
	6444673	Jan 16, 2012	DS DP		
	6864257	Aug 30, 2012	U-629		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ETHINYL ESTRADIOL; ETONOGESTREL; NUVARING</u>					
021187 001	5989581	Apr 08, 2018			
<u>ETHINYL ESTRADIOL; LEVONORGESTREL; PREVEN EMERGENCY CONTRACEPTIVE KIT</u>					
020946 001	6156742	Dec 05, 2020	U-374		
<u>ETHINYL ESTRADIOL; LEVONORGESTREL; SEASONALE</u>					
021544 001	5898032	Jun 23, 2017	U-1	NP	Sep 05, 2006
<u>ETHINYL ESTRADIOL; NORELGESTROMIN; ORTHO EVRA</u>					
021180 001	5876746 5972377	Jun 07, 2015 Jun 07, 2015	U-514 U-514		
<u>ETHINYL ESTRADIOL; NORETHINDRONE; OVCON-35</u>					
021490 001	6667050	Jun 12, 2021	DP U-1	NDF	Nov 14, 2006
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE; ESTROSTEP 21</u>					
020130 001	4962098 5010070	Oct 09, 2007 Apr 23, 2008	U-112		
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE; ESTROSTEP FE</u>					
020130 002	4962098 5010070	Oct 09, 2007 Apr 23, 2008	U-112		
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE; FEMHRT</u>					
021065 001	5208225	May 04, 2010	U-283		
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE; FEMHRT</u>					
021065 002	5208225	May 04, 2010	U-283		
<u>ETHINYL ESTRADIOL; NORGESTIMATE; ORTHO CYCLEN-21</u>					
019653 001				M-41 PED	May 13, 2008 Nov 13, 2008
<u>ETHINYL ESTRADIOL; NORGESTIMATE; ORTHO CYCLEN-28</u>					
019653 002				M-41 PED	May 13, 2008 Nov 13, 2008
<u>ETHINYL ESTRADIOL; NORGESTIMATE; ORTHO TRI-CYCLEN</u>					
019697 001				M-41 PED	May 13, 2008 Nov 13, 2008
<u>ETHINYL ESTRADIOL; NORGESTIMATE; ORTHO TRI-CYCLEN</u>					
019697 002				M-41 PED	May 13, 2008 Nov 13, 2008
<u>ETHINYL ESTRADIOL; NORGESTIMATE; ORTHO TRI-CYCLEN LO</u>					
021241 001	6214815 6214815*PED	Jun 09, 2019 Dec 09, 2019	U-112	NP PED	Aug 22, 2005 Feb 22, 2006
<u>ETODOLAC; LODINE XL</u>					
020584 001	4966768 4966768*PED	Oct 30, 2007 Apr 30, 2008			
<u>ETODOLAC; LODINE XL</u>					
020584 002	4966768 4966768*PED	Oct 30, 2007 Apr 30, 2008			
<u>ETODOLAC; LODINE XL</u>					
020584 003	4966768 4966768*PED	Oct 30, 2007 Apr 30, 2008			
<u>ETOPOSIDE PHOSPHATE; ETOPOPHOS PRESERVATIVE FREE</u>					
020457 001	5041424 RE35524	Aug 20, 2008 May 17, 2010	U-135		
<u>ETOPOSIDE PHOSPHATE; ETOPOPHOS PRESERVATIVE FREE</u>					
020906 001	5041424 RE35524	Aug 20, 2008 May 17, 2010	U-135		
<u>ETOPOSIDE PHOSPHATE; ETOPOPHOS PRESERVATIVE FREE</u>					
020906 002	5041424 RE35524	Aug 20, 2008 May 17, 2010	U-135		
<u>EXEMESTANE; AROMASIN</u>					
020753 001	4808616 4904650	Apr 01, 2011 Jul 07, 2006	DS DP DS DP	U-658 ODE	Oct 21, 2006

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE		
EXENATIDE SYNTHETIC; BYETTA										
021773 001	5424286	May 24, 2013	DP		U-653	NCE		Apr 28, 2010		
	6858576	Jan 06, 2017							U-656	
	6872700	Jan 14, 2020							U-654	
	6902744	Jan 14, 2020								
	6956026	Jan 07, 2018							U-687	
EXENATIDE SYNTHETIC; BYETTA										
021773 002	5424286	May 24, 2013	DP		U-653	NCE		Apr 28, 2010		
	6858576	Jan 06, 2017							U-656	
	6872700	Jan 14, 2020							U-654	
	6902744	Jan 14, 2020								
	6956026	Jan 07, 2018							U-687	
EZETIMIBE; ZETIA										
021445 001	5846966	Sep 21, 2013			U-474	NCE		Oct 25, 2007		
	RE37721	Jun 16, 2015							U-473	
EZETIMIBE; SIMVASTATIN; VYTORIN										
021687 001	4444784	Dec 23, 2005	DS	DP	U-592	NCE		Oct 25, 2007		
	5846966	Sep 21, 2013							DP	U-593
	RE37721	Jun 16, 2015							DS	DP
EZETIMIBE; SIMVASTATIN; VYTORIN										
021687 002	4444784	Dec 23, 2005	DS	DP	U-592	NCE		Oct 25, 2007		
	5846966	Sep 21, 2013							DP	U-593
	RE37721	Jun 16, 2015							DS	DP
EZETIMIBE; SIMVASTATIN; VYTORIN										
021687 003	4444784	Dec 23, 2005	DS	DP	U-592	NCE		Oct 25, 2007		
	5846966	Sep 21, 2013							DP	U-593
	RE37721	Jun 16, 2015							DS	DP
EZETIMIBE; SIMVASTATIN; VYTORIN										
021687 004	4444784	Dec 23, 2005	DS	DP	U-592	NCE		Oct 25, 2007		
	5846966	Sep 21, 2013							DP	U-593
	RE37721	Jun 16, 2015							DS	DP
FAMCICLOVIR; FAMVIR										
020363 001	5246937	Sep 21, 2010			U-96					
	5840763	Sep 01, 2015							U-96	
	5866581	Oct 04, 2014							U-96	
	5916893	Sep 01, 2015							U-96	
	6124304	Oct 04, 2014							U-96	
FAMCICLOVIR; FAMVIR										
020363 002	5246937	Sep 21, 2010			U-96					
	5840763	Sep 01, 2015							U-96	
	5866581	Oct 04, 2014							U-96	
	5916893	Sep 01, 2015							U-96	
	6124304	Oct 04, 2014							U-96	
FAMCICLOVIR; FAMVIR										
020363 003	5246937	Sep 21, 2010			U-96					
	5840763	Sep 01, 2015							U-96	
	5866581	Oct 04, 2014							U-96	
	5916893	Sep 01, 2015							U-96	
	6124304	Oct 04, 2014							U-96	
FAMOTIDINE; FLUXID										
021712 001	6024981	Apr 09, 2018			DP					
	6221392	Apr 09, 2018							DP	
FAMOTIDINE; FLUXID										
021712 002	6024981	Apr 09, 2018			DP					
	6221392	Apr 09, 2018							DP	
FAMOTIDINE; PEPCID AC										
020325 001	5854267	Dec 29, 2015			U-267					
	5854267*PED	Jun 29, 2016							U-267	
FAMOTIDINE; PEPCID AC										
020325 002						NP		Sep 23, 2006		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FAMOTIDINE; PEPCID AC</u>					
020801 001	5075114	Dec 24, 2008			
	5075114*PED	Jun 24, 2009			
	5667794	May 02, 2015			
	5667794*PED	Nov 02, 2015			
	5854267	Dec 29, 2015	U-267		
	5854267*PED	Jun 29, 2016			
<u>FAMOTIDINE; PEPCID AC (GELTAB)</u>					
020902 001	4820524	Feb 20, 2007			
	4820524*PED	Aug 20, 2007			
	5854267	Dec 29, 2015	U-368		
	5854267*PED	Jun 29, 2016	U-368		
<u>FELBAMATE; FELBATOL</u>					
020189 001	4978680	Sep 26, 2009	U-83		
	5082861	Sep 26, 2009	U-83		
<u>FELBAMATE; FELBATOL</u>					
020189 002	4978680	Sep 26, 2009	U-83		
	5082861	Sep 26, 2009	U-83		
<u>FELBAMATE; FELBATOL</u>					
020189 003	4978680	Sep 26, 2009	U-83		
	5082861	Sep 26, 2009	U-83		
<u>FELODIPINE; PLENDIL</u>					
019834 001	4803081	Apr 03, 2007			
	4803081*PED	Oct 03, 2007			
<u>FELODIPINE; PLENDIL</u>					
019834 002	4803081	Apr 03, 2007			
	4803081*PED	Oct 03, 2007			
<u>FELODIPINE; PLENDIL</u>					
019834 004	4803081	Apr 03, 2007			
	4803081*PED	Oct 03, 2007			
<u>FENOFIBRATE; ANTARA (MICRONIZED)</u>					
021695 001	4800079	Aug 10, 2007	DP	M-47	Oct 19, 2008
<u>FENOFIBRATE; ANTARA (MICRONIZED)</u>					
021695 002	4800079	Aug 10, 2007	DP		
<u>FENOFIBRATE; ANTARA (MICRONIZED)</u>					
021695 003	4800079	Aug 10, 2007	DP	M-47	Oct 19, 2008
<u>FENOFIBRATE; FENOFIBRATE</u>					
076433 001				PC	May 16, 2006
<u>FENOFIBRATE; FENOFIBRATE</u>					
076433 002				PC	May 16, 2006
<u>FENOFIBRATE; FENOFIBRATE</u>					
076635 001				PC	May 16, 2006
<u>FENOFIBRATE; FENOFIBRATE</u>					
076635 003				PC	May 16, 2006
<u>FENOFIBRATE; TRICOR</u>					
021203 001	4895726	Jan 19, 2009			
	6074670	Jan 09, 2018			
	6277405	Jan 09, 2018			
	6589552	Jan 09, 2018			
	6652881	Jan 09, 2018	DP		
<u>FENOFIBRATE; TRICOR</u>					
021203 003	4895726	Jan 19, 2009			
	6074670	Jan 09, 2018			
	6277405	Jan 09, 2018			
	6589552	Jan 09, 2018			
	6652881	Jan 09, 2018	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FENOFIBRATE; TRICOR</u>					
021656 001	5145684	Jan 25, 2011	DP U-615		
	6277405	Jan 09, 2018	DS		
	6375986	Sep 21, 2020	DP U-615		
	6652881	Jan 09, 2018	DS		
<u>FENOFIBRATE; TRICOR</u>					
021656 002	5145684	Jan 25, 2011	DP U-615		
	6277405	Jan 09, 2018	DS		
	6375986	Sep 21, 2020	DP U-615		
	6652881	Jan 09, 2018	DS		
<u>FENOFIBRATE; TRICOR (MICRONIZED)</u>					
019304 002	4895726	Jan 19, 2009			
<u>FENOFIBRATE; TRICOR (MICRONIZED)</u>					
019304 003	4895726	Jan 19, 2009			
<u>FENOFIBRATE; TRICOR (MICRONIZED)</u>					
019304 004	4895726	Jan 19, 2009			
<u>FENOFIBRATE; TRIGLIDE</u>					
021350 001	6696084	Sep 11, 2021	DS DP U-680		
<u>FENOFIBRATE; TRIGLIDE</u>					
021350 002	6696084	Sep 11, 2021	DS DP U-680		
<u>FENOLDOPAM MESYLATE; CORLOPAM</u>					
019922 001				I-422	Apr 01, 2007
<u>FENTANYL; DURAGESIC-100</u>					
019813 001				NPP PED	May 20, 2006 Nov 20, 2006
<u>FENTANYL; DURAGESIC-12</u>					
019813 005				NPP PED	May 20, 2006 Nov 20, 2006
<u>FENTANYL; DURAGESIC-25</u>					
019813 004				NPP PED	May 20, 2006 Nov 20, 2006
<u>FENTANYL; DURAGESIC-50</u>					
019813 003				NPP PED	May 20, 2006 Nov 20, 2006
<u>FENTANYL; DURAGESIC-75</u>					
019813 002				NPP PED	May 20, 2006 Nov 20, 2006
<u>FENTANYL CITRATE; ACTIQ</u>					
020747 001	4863737	Sep 05, 2006			
<u>FENTANYL CITRATE; ACTIQ</u>					
020747 002	4863737	Sep 05, 2006			
<u>FENTANYL CITRATE; ACTIQ</u>					
020747 003	4863737	Sep 05, 2006			
<u>FENTANYL CITRATE; ACTIQ</u>					
020747 004	4863737	Sep 05, 2006			
<u>FENTANYL CITRATE; ACTIQ</u>					
020747 005	4863737	Sep 05, 2006			
<u>FENTANYL CITRATE; ACTIQ</u>					
020747 006	4863737	Sep 05, 2006			
<u>FERRIC HEXACYANOFERRATE(II); RADIOGARDASE (PRUSSIAN BLUE)</u>					
021626 001				NCE ODE	Oct 02, 2008 Oct 02, 2010

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FERUMOXIDES; FERIDEX I.V.</u>					
020416 001	4770183	Sep 13, 2005	U-145		
	4827945	May 09, 2006	U-144		
	4951675	Aug 28, 2007	U-143		
	5055288	Oct 08, 2008			
	5102652	Feb 06, 2009			
	5219554	Jun 15, 2010			
	5248492	Sep 28, 2010			
<u>FERUMOXSYL; GASTROMARK</u>					
020410 001	4770183	Sep 13, 2005	U-169		
	4827945	May 09, 2006	U-170		
	4951675	Sep 13, 2005	U-169		
	5055288	Oct 08, 2008			
	5069216	May 09, 2006	U-171		
	5219554	Jun 15, 2010			
<u>FEXOFENADINE HYDROCHLORIDE; ALLEGRA</u>					
020625 001	5578610	Nov 26, 2013	U-192	M-25	May 12, 2006
	5578610*PED	May 26, 2014	U-192	PED	Nov 12, 2006
	5738872	Feb 28, 2015			
	5738872*PED	Aug 28, 2015			
	5855912	Feb 28, 2015			
	5855912*PED	Aug 28, 2015			
	5932247	Feb 28, 2015			
	5932247*PED	Aug 28, 2015			
	6037353	Mar 14, 2017	U-138		
	6037353*PED	Sep 14, 2017	U-138		
	6113942	Feb 28, 2015			
	6113942*PED	Aug 28, 2015			
	6187791	May 11, 2012	U-138		
	6187791*PED	Nov 11, 2012	U-138		
	6399632	May 11, 2012	U-468		
	6399632*PED	Nov 11, 2012	U-468		
<u>FEXOFENADINE HYDROCHLORIDE; ALLEGRA</u>					
020872 001	5578610	Nov 26, 2013	DS DP	U-139	M-25
	5578610	Nov 26, 2013	DS DP	U-684	
	5578610*PED	May 26, 2014		U-139	PED
	5855912	Feb 28, 2015		DP	
	5855912*PED	Aug 28, 2015			
	5932247	Feb 28, 2015		DP	
	5932247*PED	Aug 28, 2015			
	6037353	Mar 14, 2017		U-684	
	6037353	Mar 14, 2017		U-138	
	6037353*PED	Sep 14, 2017		U-138	
	6113942	Feb 28, 2015	DP		
	6113942*PED	Aug 28, 2015			
	6187791	May 11, 2012		U-684	
	6187791	May 11, 2012		U-138	
	6187791*PED	Nov 11, 2012		U-138	
	6399632	May 11, 2012		U-468	
	6399632	May 11, 2012		U-684	
	6399632*PED	Nov 11, 2012		U-468	
<u>FEXOFENADINE HYDROCHLORIDE; ALLEGRA</u>					
020872 002	5578610	Nov 26, 2013	DS DP	U-139	M-25
	5578610	Nov 26, 2013	DS DP	U-684	
	5578610*PED	May 26, 2014		U-139	PED
	5855912	Feb 28, 2015		DP	
	5855912*PED	Aug 28, 2015			
	5932247	Feb 28, 2015		DP	
	5932247*PED	Aug 28, 2015			
	6037353	Mar 14, 2017		U-138	
	6037353	Mar 14, 2017		U-684	
	6037353*PED	Sep 14, 2017		U-138	
	6113942	Feb 28, 2015	DP		
	6113942*PED	Aug 28, 2015			
	6187791	May 11, 2012		U-684	
	6187791	May 11, 2012		U-138	
	6187791*PED	Nov 11, 2012		U-138	
	6399632	May 11, 2012		U-684	
	6399632	May 11, 2012		U-468	
	6399632*PED	Nov 11, 2012		U-468	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FEXOFENADINE HYDROCHLORIDE; ALLEGRA</u>					
020872 004	5578610	Nov 26, 2013	DS DP	U-684	D-101 Oct 13, 2008
	5578610	Nov 26, 2013	DS DP	U-139	M-25 May 12, 2006
	5578610*PED	May 26, 2014		U-139	PED Nov 12, 2006
	5855912	Feb 28, 2015	DP		
	5855912*PED	Aug 28, 2015			
	5932247	Feb 28, 2015	DP		
	5932247*PED	Aug 28, 2015			
	6037353	Mar 14, 2017		U-138	
	6037353	Mar 14, 2017		U-684	
	6037353*PED	Sep 14, 2017		U-138	
	6113942	Feb 28, 2015	DP		
	6113942*PED	Aug 28, 2015			
	6187791	May 11, 2012		U-684	
	6187791	May 11, 2012		U-138	
	6187791*PED	Nov 11, 2012		U-138	
	6399632	May 11, 2012		U-468	
	6399632	May 11, 2012		U-684	
	6399632*PED	Nov 11, 2012		U-468	
<u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE; ALLEGRA D 24 HOUR</u>					
021704 001	5578610	Nov 26, 2013	DS DP	U-612	
	5578610*PED	May 26, 2014			
	6004582	May 29, 2018	DP		
	6037353	Mar 17, 2017		U-612	
	6037353*PED	Sep 14, 2017			
	6187791	May 11, 2012		U-612	
	6187791*PED	Nov 11, 2012			
	6399632	May 11, 2012		U-612	
	6399632*PED	Nov 11, 2012			
	6613357	Dec 25, 2020	DP	U-612	
<u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE; ALLEGRA-D 12 HOUR</u>					
020786 001	5578610	Nov 26, 2013			
	5578610*PED	May 26, 2014			
	5855912	Feb 28, 2015			
	5855912*PED	Aug 28, 2015			
	6037353	Mar 14, 2017		U-138	
	6037353*PED	Sep 14, 2017		U-138	
	6039974	Jul 31, 2018			
	6113942	Feb 28, 2015			
	6113942*PED	Aug 28, 2015			
	6187791	May 11, 2012		U-138	
	6187791*PED	Nov 11, 2012		U-138	
	6399632	May 11, 2012		U-468	
	6399632*PED	Nov 11, 2012		U-468	
<u>FINASTERIDE; PROPECIA</u>					
020788 001	4760071	Jun 19, 2006			
	5547957	Oct 15, 2013		U-236	
	5571817	Nov 05, 2013		U-259	
	5886184	Nov 19, 2012			
<u>FINASTERIDE; PROSCAR</u>					
020180 001	4760071	Jun 19, 2006	DS DP	U-262	
	5886184	Nov 19, 2012	DS		
	5942519	Oct 23, 2018		U-280	
	6046183	Mar 20, 2011	DP	U-577	
<u>FLUNISOLIDE; AEROBID</u>					
018340 001	4933168	Jun 12, 2007			
<u>FLUNISOLIDE; NASALIDE</u>					
018148 001	4933168	Jun 12, 2007			
<u>FLUNISOLIDE; NASAREL</u>					
020409 001	4782047	May 22, 2006			
	4933168	Jun 12, 2007			
	4983595	May 22, 2006			
<u>FLUOCINOLONE ACETONIDE; FLUOCINOLONE ACETONIDE</u>					
021930 001				NP	Nov 09, 2008

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUOCINOLONE ACETONIDE; RETISERT</u>					
021737 001				NDF	Apr 08, 2008
				ODE	Apr 08, 2012
<u>FLUOCINONIDE; VANOS</u>					
021758 001	6765001	Dec 21, 2021	DP	NP	Feb 11, 2008
<u>FLUOROURACIL; CARAC</u>					
020985 001	4690825	Oct 04, 2005			
	6670335	Jun 02, 2021	DP U-68		
<u>FLUOXETINE HYDROCHLORIDE; PROZAC</u>					
018936 001				I-362	Jul 29, 2005
				NPP	Jan 03, 2006
				PED	Jul 03, 2006
<u>FLUOXETINE HYDROCHLORIDE; PROZAC</u>					
018936 003				I-362	Jul 29, 2005
				NPP	Jan 03, 2006
				PED	Jul 03, 2006
<u>FLUOXETINE HYDROCHLORIDE; PROZAC</u>					
018936 004				I-362	Jul 29, 2005
				NPP	Jan 03, 2006
				PED	Jul 03, 2006
<u>FLUOXETINE HYDROCHLORIDE; PROZAC</u>					
018936 006				I-362	Jul 29, 2005
				NPP	Jan 03, 2006
				PED	Jul 03, 2006
<u>FLUOXETINE HYDROCHLORIDE; PROZAC</u>					
020101 001				I-362	Jul 29, 2005
				NPP	Jan 03, 2006
				PED	Jul 03, 2006
<u>FLUOXETINE HYDROCHLORIDE; PROZAC</u>					
020974 001				I-362	Jul 29, 2005
				NPP	Jan 03, 2006
				PED	Jul 03, 2006
<u>FLUOXETINE HYDROCHLORIDE; PROZAC</u>					
020974 002				I-362	Jul 29, 2005
				NPP	Jan 03, 2006
				PED	Jul 03, 2006
<u>FLUOXETINE HYDROCHLORIDE; PROZAC WEEKLY</u>					
021235 001	5910319	May 29, 2017		U-396	
	5985322	May 29, 2017		U-397	
<u>FLUOXETINE HYDROCHLORIDE; SARAFEM</u>					
018936 007	4971998	Nov 20, 2007		U-338	
	4971998*PED	May 20, 2008		U-338	
<u>FLUOXETINE HYDROCHLORIDE; SARAFEM</u>					
018936 008	4971998	Nov 20, 2007		U-338	
	4971998*PED	May 20, 2008		U-338	
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE; SYMBYAX</u>					
021520 002	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5945416	Mar 24, 2017	DS DP		
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE; SYMBYAX</u>					
021520 003	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5945416	Mar 24, 2017	DS DP		
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE; SYMBYAX</u>					
021520 004	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5945416	Mar 24, 2017	DS DP		
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE; SYMBYAX</u>					
021520 005	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5945416	Mar 24, 2017	DS DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; CUTIVATE</u>					
021152 001				NDF	Mar 31, 2008
				PED	Sep 30, 2008
<u>FLUTICASONE PROPIONATE; FLONASE</u>					
020121 001				M-24	May 01, 2006
				PED	Nov 01, 2006
				PED	Nov 23, 2005
<u>FLUTICASONE PROPIONATE; FLOVENT DISKUS 100</u>					
020833 002	6536427	Mar 01, 2011	DP		
	6536427*PED	Sep 01, 2011	DP		
<u>FLUTICASONE PROPIONATE; FLOVENT DISKUS 250</u>					
020833 003	6536427	Mar 01, 2011	DP		
	6536427*PED	Sep 01, 2011	DP		
<u>FLUTICASONE PROPIONATE; FLOVENT DISKUS 50</u>					
020833 001	6536427	Mar 01, 2011	DP		
	6536427*PED	Sep 01, 2011	DP		
<u>FLUTICASONE PROPIONATE; FLOVENT HFA</u>					
021433 001	6170717	Dec 23, 2017	DP	NP	May 14, 2007
	6253762	Apr 14, 2015	DP	U-582	
	6315173	Dec 23, 2017	DP		
	6510969	Dec 23, 2017	DP		
	6546928	Apr 14, 2015	DP	U-583	
	6743413	Jun 01, 2021	DP	U-581	
<u>FLUTICASONE PROPIONATE; FLOVENT HFA</u>					
021433 002	6170717	Dec 23, 2017	DP	NP	May 14, 2007
	6253762	Apr 14, 2015	DP	U-582	
	6315173	Dec 23, 2017	DP		
	6510969	Dec 23, 2017	DP		
	6546928	Apr 14, 2015	DP	U-583	
	6743413	Jun 01, 2021	DP	U-581	
<u>FLUTICASONE PROPIONATE; FLOVENT HFA</u>					
021433 003	6170717	Dec 23, 2017	DP	NP	May 14, 2007
	6253762	Apr 14, 2015	DP	U-582	
	6315173	Dec 23, 2017	DP		
	6510969	Dec 23, 2017	DP		
	6546928	Apr 14, 2015	DP	U-583	
	6743413	Jun 01, 2021	DP	U-581	
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE; ADVAIR DISKUS 100/50</u>					
021077 001	4992474	Feb 12, 2008		M-33	Apr 21, 2007
	5126375	Feb 12, 2008			
	5225445	Feb 12, 2008		U-211	
	5270305	Sep 07, 2010		U-387	
	6536427	Mar 01, 2011	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE; ADVAIR DISKUS 250/50</u>					
021077 002	4992474	Feb 12, 2008		I-410	Nov 17, 2006
	5126375	Feb 12, 2008			
	5225445	Feb 12, 2008		U-211	
	5270305	Sep 07, 2010		U-387	
	6536427	Mar 01, 2011	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE; ADVAIR DISKUS 500/50</u>					
021077 003	4992474	Feb 12, 2008		U-211	
	5126375	Feb 12, 2008			
	5225445	Feb 12, 2008		U-211	
	5270305	Sep 07, 2010		U-387	
	6536427	Mar 01, 2011	DP		
<u>FLUVASTATIN SODIUM; LESCOL</u>					
020261 001	5354772	Oct 11, 2011	U-413	I-394	May 27, 2006
	5354772	Oct 11, 2011	U-109	PED	Nov 27, 2006
	5354772*PED	Apr 11, 2012			
	5356896	Dec 12, 2011			
	5356896*PED	Jun 12, 2012			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUVASTATIN SODIUM; LESCOL</u>					
020261 002	5354772	Oct 11, 2011	U-413	I-394	May 27, 2006
	5354772	Oct 11, 2011	U-109	PED	Nov 27, 2006
	5354772*PED	Apr 11, 2012			
	5356896	Dec 12, 2011			
	5356896*PED	Jun 12, 2012			
<u>FLUVASTATIN SODIUM; LESCOL XL</u>					
021192 001	5354772	Oct 11, 2011	U-413	I-394	May 27, 2006
	5354772	Oct 11, 2011	U-109	PED	Nov 27, 2006
	5354772*PED	Apr 11, 2012			
	5356896	Dec 12, 2011			
	5356896*PED	Jun 12, 2012			
	6242003	Apr 13, 2020			
	6242003*PED	Oct 13, 2020			
<u>FOLLITROPIN ALFA/BETA; FOLLISTIM</u>					
020582 001	5270057	Mar 20, 2011			
	5767251	Jun 16, 2015			
<u>FOLLITROPIN ALFA/BETA; FOLLISTIM</u>					
020582 002	5270057	Mar 20, 2011			
	5767251	Jun 16, 2015			
<u>FOLLITROPIN ALFA/BETA; FOLLISTIM AQ</u>					
021211 001	5767251	Jun 16, 2015	DS		
	5929028	Jan 14, 2018	DP U-567		
<u>FOLLITROPIN ALFA/BETA; FOLLISTIM AQ</u>					
021211 002	5767251	Jun 16, 2015	DS		
	5929028	Jan 14, 2018	DP U-567		
<u>FOLLITROPIN ALFA/BETA; GONAL-F</u>					
020378 001	5767251	Jun 16, 2015	DS	ODE	May 24, 2007
<u>FOLLITROPIN ALFA/BETA; GONAL-F</u>					
020378 002	5767251	Jun 16, 2015	DS	ODE	May 24, 2007
<u>FOLLITROPIN ALFA/BETA; GONAL-F</u>					
020378 003	5767251	Jun 16, 2015	DS	ODE	May 24, 2007
<u>FOLLITROPIN ALFA/BETA; GONAL-F</u>					
020378 004	5767067	Jun 16, 2015	DS	ODE	May 24, 2007
	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA; GONAL-F</u>					
020378 005	5767067	Jun 16, 2015	DS	ODE	May 24, 2007
	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA; GONAL-F</u>					
021765 001	5767251	Jun 16, 2015	DS	NP	Mar 25, 2007
<u>FOLLITROPIN ALFA/BETA; GONAL-F</u>					
021765 003	5767251	Jun 16, 2015	DS	NP	Mar 25, 2007
<u>FOLLITROPIN ALFA/BETA; GONAL-F RFF</u>					
021765 002	5767067	Jun 16, 2015	DS	NP	Mar 25, 2007
	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA; GONAL-F RFF PEN</u>					
021684 001	5767067	Jun 16, 2015	DS		
	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA; GONAL-F RFF PEN</u>					
021684 002	5767067	Jun 16, 2015	DS		
	5767251	Jun 16, 2015	DS		
<u>FOLLITROPIN ALFA/BETA; GONAL-F RFF PEN</u>					
021684 003	5767067	Jun 16, 2015	DS		
	5767251	Jun 16, 2015	DS		
<u>FOMEPIZOLE; ANTIZOL</u>					
020696 001				ODE	Dec 08, 2007

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FOMIVIRSEN SODIUM; VITRAVENE PRESERVATIVE FREE					
020961 001	5264423	Nov 23, 2010	U-522		
	5276019	Jan 04, 2011	U-522		
	5442049	Aug 15, 2012			
	5595978	Aug 15, 2012	U-522		
FONDAPARINUX SODIUM; ARIXTRA					
021345 001				I-457	May 26, 2008
				I-427	May 31, 2007
				I-426	May 31, 2007
				I-397	Jun 17, 2006
				NCE	Dec 07, 2006
FONDAPARINUX SODIUM; ARIXTRA					
021345 002				I-457	May 26, 2008
				I-427	May 31, 2007
				I-426	May 31, 2007
				I-397	Jun 17, 2006
				NS	May 31, 2007
				NCE	Dec 07, 2006
FONDAPARINUX SODIUM; ARIXTRA					
021345 003				I-457	May 26, 2008
				I-427	May 31, 2007
				I-426	May 31, 2007
				I-397	Jun 17, 2006
				NS	May 31, 2007
				NCE	Dec 07, 2006
FONDAPARINUX SODIUM; ARIXTRA					
021345 004				I-457	May 26, 2008
				I-427	May 31, 2007
				I-426	May 31, 2007
				I-397	Jun 17, 2006
				NS	May 31, 2007
				NCE	Dec 07, 2006
FORMOTEROL FUMARATE; FORADIL					
020831 001	6488027	Mar 08, 2019		NPP	Jun 27, 2006
				NCE	Feb 16, 2006
FOSAMPRENAVIR CALCIUM; LEXIVA					
021548 001	6436989	Dec 24, 2017	U-257	NE	Oct 20, 2006
	6514953	Jul 15, 2019	U-257		
FOSINOPRIL SODIUM; MONOPRIL					
019915 002	5006344	Jul 10, 2009		NPP	May 27, 2006
	5006344*PED	Jan 10, 2010		PED	Nov 27, 2006
FOSINOPRIL SODIUM; MONOPRIL					
019915 003	5006344	Jul 10, 2009		NPP	May 27, 2006
	5006344*PED	Jan 10, 2010		PED	Nov 27, 2006
FOSINOPRIL SODIUM; MONOPRIL					
019915 004	5006344	Jul 10, 2009		NPP	May 27, 2006
	5006344*PED	Jan 10, 2010		PED	Nov 27, 2006
FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE; MONOPRIL-HCT					
020286 001	5006344	Jul 10, 2009			
	5006344*PED	Jan 10, 2010			
FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE; MONOPRIL-HCT					
020286 002	5006344	Jul 10, 2009			
	5006344*PED	Jan 10, 2010			
FOSPHENYTOIN SODIUM; CEREBYX					
020450 001	4925860	Aug 05, 2007			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FROVATRIPTAN SUCCINATE; FROVA</u>					
021006 001	5464864	Nov 07, 2012	U-436	NCE	Nov 08, 2006
	5616603	Apr 01, 2014	U-436		
	5637611	Jun 10, 2014	U-436		
	5827871	Oct 27, 2015	U-436		
	5962501	Dec 16, 2013	U-436		
<u>FULVESTRANT; FASLODEX</u>					
021344 001	4659516	Oct 01, 2005		NCE	Apr 25, 2007
	6774122	Jan 09, 2021	U-596		
<u>GABAPENTIN; NEURONTIN</u>					
020235 001	4894476	May 02, 2008			
	4894476*PED	Nov 02, 2008			
	6054482	Apr 25, 2017			
	6054482*PED	Oct 25, 2017			
<u>GABAPENTIN; NEURONTIN</u>					
020235 002	4894476	May 02, 2008			
	4894476*PED	Nov 02, 2008			
	6054482	Apr 25, 2017			
	6054482*PED	Oct 25, 2017			
<u>GABAPENTIN; NEURONTIN</u>					
020235 003	4894476	May 02, 2008			
	4894476*PED	Nov 02, 2008			
	6054482	Apr 25, 2017			
	6054482*PED	Oct 25, 2017			
<u>GABAPENTIN; NEURONTIN</u>					
020882 001	4894476	May 02, 2008			
	4894476*PED	Nov 02, 2008			
	6054482	Apr 25, 2017			
	6054482*PED	Oct 25, 2017			
<u>GABAPENTIN; NEURONTIN</u>					
020882 002	4894476	May 02, 2008			
	4894476*PED	Nov 02, 2008			
	6054482	Apr 25, 2017			
	6054482*PED	Oct 25, 2017			
<u>GABAPENTIN; NEURONTIN</u>					
021129 001	4894476	May 02, 2008			
	4894476*PED	Nov 02, 2008			
	6054482	Apr 25, 2017			
	6054482*PED	Oct 25, 2017			
<u>GADOBENATE DIMEGLUMINE; MULTIANCE</u>					
021357 001				NCE	Nov 23, 2009
<u>GADOBENATE DIMEGLUMINE; MULTIANCE</u>					
021357 002				NCE	Nov 23, 2009
<u>GADOBENATE DIMEGLUMINE; MULTIANCE</u>					
021357 003				NCE	Nov 23, 2009
<u>GADOBENATE DIMEGLUMINE; MULTIANCE</u>					
021357 004				NCE	Nov 23, 2009
<u>GADOBENATE DIMEGLUMINE; MULTIANCE MULTIPACK</u>					
021358 001				NCE	Nov 23, 2009
<u>GADOBENATE DIMEGLUMINE; MULTIANCE MULTIPACK</u>					
021358 002				NCE	Nov 23, 2009
<u>GADODIAMIDE; OMNISCAN</u>					
020123 001	4687659	May 04, 2007		U-76	
	5362475	Nov 08, 2011	DS		
<u>GADOPENTETATE DIMEGLUMINE; MAGNEVIST</u>					
019596 001	5362475	Nov 08, 2011			
	5560903	Oct 01, 2013			
<u>GADOPENTETATE DIMEGLUMINE; MAGNEVIST</u>					
021037 001	5362475	Nov 08, 2011			
	5560903	Oct 01, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GADOTERIDOL; PROHANCE</u>					
020131 001	4885363	Dec 05, 2006			
	5474756	Dec 12, 2012		U-480	
	5846519	Dec 08, 2015			
	6143274	Dec 12, 2012		U-480	
<u>GADOTERIDOL; PROHANCE MULTIPACK</u>					
021489 001	4885363	Dec 05, 2006			
	5474756	Dec 12, 2012		U-536	
	5846519	Dec 08, 2015			
	6143274	Dec 12, 2012		U-536	
<u>GADOVERSETAMIDE; OPTIMARK</u>					
020937 001	5130120	Jul 14, 2009			
	5137711	Jul 14, 2009			
<u>GADOVERSETAMIDE; OPTIMARK</u>					
020975 001	5130120	Jul 14, 2009			
	5137711	Jul 14, 2009			
<u>GADOVERSETAMIDE; OPTIMARK IN PLASTIC CONTAINER</u>					
020976 001	5130120	Jul 14, 2009			
	5137711	Jul 14, 2009			
<u>GALANTAMINE HYDROBROMIDE; RAZADYNE</u>					
021169 001	4663318	Dec 14, 2008		U-322	NCE
	6099863	Jun 06, 2017			
	6358527	Jun 06, 2017	DP	U-322	
<u>GALANTAMINE HYDROBROMIDE; RAZADYNE</u>					
021169 002	4663318	Dec 14, 2008		U-322	NCE
	6099863	Jun 06, 2017			
	6358527	Jun 06, 2017	DP	U-322	
<u>GALANTAMINE HYDROBROMIDE; RAZADYNE</u>					
021169 003	4663318	Dec 14, 2008		U-322	NCE
	6099863	Jun 06, 2017			
	6358527	Jun 06, 2017	DP	U-322	
<u>GALANTAMINE HYDROBROMIDE; RAZADYNE</u>					
021224 001	4663318	Dec 14, 2008		U-322	NCE
<u>GALANTAMINE HYDROBROMIDE; RAZADYNE ER</u>					
021615 001	4663318	Dec 14, 2008		U-322	NDF
					NCE
<u>GALANTAMINE HYDROBROMIDE; RAZADYNE ER</u>					
021615 002	4663318	Dec 14, 2008		U-322	NDF
					NCE
<u>GALANTAMINE HYDROBROMIDE; RAZADYNE ER</u>					
021615 003	4663318	Dec 14, 2008		U-322	NDF
					NCE
<u>GANCICLOVIR; VITRASERT</u>					
020569 001	5378475	Jan 03, 2012			
<u>GANIRELIX ACETATE; GANIRELIX ACETATE INJECTION</u>					
021057 001	4801577	Feb 05, 2007			
	5767082	Jun 16, 2015			
<u>GATIFLOXACIN; TEQUIN</u>					
021061 001	4980470	Dec 15, 2009	DS	I-432	Jun 30, 2007
	5880283	Dec 05, 2015		I-329	Oct 17, 2005
<u>GATIFLOXACIN; TEQUIN</u>					
021061 002	4980470	Dec 15, 2009	DS	I-432	Jun 30, 2007
	5880283	Dec 05, 2015		I-329	Oct 17, 2005
<u>GATIFLOXACIN; TEQUIN</u>					
021062 003	4980470	Dec 15, 2009	DS	I-432	Jun 30, 2007
	5880283	Dec 05, 2015		I-329	Oct 17, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE				
GATIFLOXACIN; TEQUIN											
021062 004	4980470	Dec 15, 2009	DS	U-603	I-432	Jun 30, 2007					
	5880283	Dec 05, 2015			I-329	Oct 17, 2005					
GATIFLOXACIN; TEQUIN											
021678 001	4980470	Dec 15, 2009	DS		I-329	Oct 17, 2005					
	5880283	Dec 05, 2015	DS								
	6589955	May 09, 2022	DS								
GATIFLOXACIN; TEQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER											
021062 001	4980470	Dec 15, 2009	DS		U-603	I-432	Jun 30, 2007				
	5880283	Dec 05, 2015				I-329	Oct 17, 2005				
GATIFLOXACIN; TEQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER											
021062 002	4980470	Dec 15, 2009	DS	U-603		I-432	Jun 30, 2007				
	5880283	Dec 05, 2015				I-329	Oct 17, 2005				
GATIFLOXACIN; ZYMAR											
021493 001	4980470	Dec 15, 2009	DS			DP	NDF	Mar 28, 2006			
	5880283	Dec 05, 2015									
	6333045	Aug 20, 2019							DP		
GEFITINIB; IRESSA											
021399 001	5457105	Jan 19, 2013	U-146		NCE	May 05, 2008					
	5616582	Jan 19, 2013									
	5770599	Apr 26, 2016									
GEMCITABINE HYDROCHLORIDE; GEMZAR											
020509 001	4808614	May 15, 2010		DS			U-146	I-428	May 19, 2007		
	4808614*PED	Nov 15, 2010						M-40	Apr 26, 2008		
	5464826	Nov 07, 2012						PED	Oct 26, 2008		
	5464826*PED	May 07, 2013						PED	Nov 19, 2007		
GEMCITABINE HYDROCHLORIDE; GEMZAR											
020509 002	4808614	May 15, 2010		DS			U-146	I-428	May 19, 2007		
	4808614*PED	Nov 15, 2010	M-40		Apr 26, 2008						
	5464826	Nov 07, 2012	PED		Oct 26, 2008						
	5464826*PED	May 07, 2013	PED		Nov 19, 2007						
GEMIFLOXACIN MESYLATE; FACTIVE											
021158 001	5633262	Jun 15, 2015	DS	DP	U-608	NCE	Apr 04, 2008				
	5776944	Jun 15, 2015									
	5962468	Jun 15, 2015									
	6262071	Sep 21, 2019									
	6331550	Sep 21, 2019									
	6340689	Sep 14, 2019									
	6455540	Sep 21, 2019									
	6723734	Mar 20, 2018									
	6803376	Sep 21, 2019									
	6803376	Sep 21, 2019									
GEMTUZUMAB OZOGAMICIN; MYLOTARG											
021174 001	4970198	Nov 30, 2007	U-320	ODE	May 17, 2007						
	5079233	Jan 07, 2009									
	5585089	Dec 17, 2013									
	5606040	Feb 25, 2014									
	5693762	Dec 02, 2014									
	5739116	Apr 14, 2015									
	5767285	Jun 16, 2015									
	5773001	Jun 30, 2015									
GLATIRAMER ACETATE; COPAXONE											
020622 001	5981589	May 24, 2014				DS	U-441	U-441			
	6054430	May 24, 2014									
	6342476	May 24, 2014									
	6362161	May 24, 2014									
	6620847	May 24, 2014									
	6939539	May 24, 2014									

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GLATIRAMER ACETATE; COPAXONE</u>					
020622 002	5981589	May 24, 2014			
	6054430	May 24, 2014			
	6342476	May 24, 2014		U-441	
	6362161	May 24, 2014		U-441	
	6620847	May 24, 2014	DS		
	6939539	May 24, 2014	DS		
<u>GLIMEPIRIDE; AMARYL</u>					
020496 001	4379785*PED	Oct 06, 2005			
<u>GLIMEPIRIDE; AMARYL</u>					
020496 002	4379785*PED	Oct 06, 2005			
<u>GLIMEPIRIDE; AMARYL</u>					
020496 003	4379785*PED	Oct 06, 2005			
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE; AVANDARYL</u>					
021700 001	5002953	Sep 17, 2011	DS DP	U-690	
	5741803	Apr 21, 2015	DS DP	U-690	
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE; AVANDARYL</u>					
021700 002	5002953	Sep 17, 2011	DS DP	U-690	
	5741803	Apr 21, 2015	DS DP	U-690	
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE; AVANDARYL</u>					
021700 003	5002953	Sep 17, 2011	DS DP	U-690	
	5741803	Apr 21, 2015	DS DP	U-690	
<u>GLIPIZIDE; GLUCOTROL XL</u>					
020329 001	5024843	Sep 05, 2009			
	5091190	Sep 05, 2009		U-111	
	5545413	Jul 02, 2008		U-111	
	5591454	Jan 07, 2014		U-150	
<u>GLIPIZIDE; GLUCOTROL XL</u>					
020329 002	5024843	Sep 05, 2009			
	5091190	Sep 05, 2009		U-111	
	5545413	Jul 02, 2008		U-111	
	5591454	Jan 07, 2014		U-150	
<u>GLIPIZIDE; GLUCOTROL XL</u>					
020329 003	5024843	Sep 05, 2009			
	5091190	Sep 05, 2009		U-111	
	5545413	Jul 02, 2008		U-111	
	5591454	Jan 07, 2014		U-111	
<u>GLIPIZIDE; METFORMIN HYDROCHLORIDE; METAGLIP</u>					
021460 001				NC	Oct 21, 2005
<u>GLIPIZIDE; METFORMIN HYDROCHLORIDE; METAGLIP</u>					
021460 002				NC	Oct 21, 2005
<u>GLIPIZIDE; METFORMIN HYDROCHLORIDE; METAGLIP</u>					
021460 003				NC	Oct 21, 2005
<u>GLUTAMINE; NUTRESTORE</u>					
021667 001				NCE	Jun 10, 2009
				ODE	Jun 10, 2011
<u>GLYBURIDE; GLYNASE</u>					
020051 001	4735805	Mar 11, 2007			
	4916163	Apr 10, 2007			
<u>GLYBURIDE; GLYNASE</u>					
020051 002	4735805	Mar 11, 2007			
	4916163	Apr 10, 2007			
<u>GLYBURIDE; GLYNASE</u>					
020051 003	4735805	Mar 11, 2007			
	4916163	Apr 10, 2007			
<u>GLYBURIDE; GLYNASE</u>					
020051 004	4735805	Mar 11, 2007			
	4916163	Apr 10, 2007			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GLYBURIDE; METFORMIN HYDROCHLORIDE; GLUCOVANCE</u>					
021178 001	6303146	Jul 14, 2019	U-412	I-368	Sep 30, 2005
	6303146*PED	Jan 14, 2020	U-412	M-37	Mar 15, 2007
				PED	Sep 15, 2007
				PED	Mar 30, 2006
<u>GLYBURIDE; METFORMIN HYDROCHLORIDE; GLUCOVANCE</u>					
021178 002	6303146	Jul 14, 2019	U-412	I-368	Sep 30, 2005
	6303146*PED	Jan 14, 2020	U-412	M-37	Mar 15, 2007
				PED	Sep 15, 2007
				PED	Mar 30, 2006
<u>GLYBURIDE; METFORMIN HYDROCHLORIDE; GLUCOVANCE</u>					
021178 003	6303146	Jul 14, 2019	U-412	I-368	Sep 30, 2005
	6303146*PED	Jan 14, 2020	U-412	M-37	Mar 15, 2007
				PED	Sep 15, 2007
				PED	Mar 30, 2006
<u>GOSERELIN ACETATE; ZOLADEX</u>					
019726 001	4767628	Aug 30, 2005			
	5366734	Nov 22, 2011			
<u>GOSERELIN ACETATE; ZOLADEX</u>					
020578 001	4767628	Aug 30, 2005			
	5366734	Nov 22, 2011			
<u>GRANISETRON HYDROCHLORIDE; KYTRIL</u>					
020239 001	4886808	Dec 29, 2007	U-89	I-369	Aug 16, 2005
	5952340	Sep 14, 2016	U-519		
	6294548	May 04, 2019			
<u>GRANISETRON HYDROCHLORIDE; KYTRIL</u>					
020239 002	4886808	Dec 29, 2007	U-89	I-369	Aug 16, 2005
	5952340	Sep 14, 2016	U-519		
	6294548	May 04, 2019			
<u>GRANISETRON HYDROCHLORIDE; KYTRIL</u>					
020239 003	4886808	Dec 29, 2007	DS DP U-89	I-369	Aug 16, 2005
<u>GRANISETRON HYDROCHLORIDE; KYTRIL</u>					
020239 004	4886808	Dec 29, 2007	U-89	I-369	Aug 16, 2005
	5952340	Sep 14, 2016	U-519		
	6294548	May 04, 2019			
<u>GRANISETRON HYDROCHLORIDE; KYTRIL</u>					
020305 001	4886808	Dec 29, 2007	U-105		
<u>GRANISETRON HYDROCHLORIDE; KYTRIL</u>					
020305 002	4886808	Dec 20, 2007	U-105		
<u>GRANISETRON HYDROCHLORIDE; KYTRIL</u>					
021238 001	4886808	Dec 29, 2007	U-105		
<u>GREPAFLOXACIN HYDROCHLORIDE; RAXAR</u>					
020695 001	5563138	Oct 08, 2013			
<u>GUAIFENESIN; MUCINEX</u>					
021282 001	6372252	Apr 28, 2020	U-489		
	6955821	Apr 28, 2020	DP U-489		
<u>GUAIFENESIN; MUCINEX</u>					
021282 002	6372252	Apr 28, 2020	U-489		
	6955821	Apr 28, 2020	DP U-489		
<u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE; MUCINEX D</u>					
021585 001	6372252	Apr 28, 2020	DP		
	6955821	Apr 28, 2020	DP U-686		
<u>GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE; MUCINEX D</u>					
021585 002	6372252	Apr 28, 2020	DP		
	6955821	Apr 28, 2020	DP U-686		
<u>HISTRELIN ACETATE; VANTAS</u>					
021732 001	5266325	Nov 30, 2010	DP	NDF	Oct 12, 2007
	5292515	Mar 08, 2011	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYALURONIDASE; AMPHADASE</u>					
021665 001				NCE	Oct 26, 2009
<u>HYALURONIDASE; HYDASE</u>					
021716 001				NCE	Oct 25, 2010
<u>HYALURONIDASE; VITRASE</u>					
021640 001				NCE	May 05, 2009
<u>HYALURONIDASE; VITRASE</u>					
021640 002				NCE	May 05, 2009
<u>HYDRALAZINE HYDROCHLORIDE; ISOSORBIDE DINITRATE; BIDIL</u>					
020727 001	4868179	Apr 22, 2007	U-71	NC	Jun 23, 2008
	6465463	Sep 08, 2020	U-71		
	6784177	Sep 08, 2020	U-71		
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN; AVALIDE</u>					
020758 001	5270317	Sep 30, 2011			
	5270317*PED	Mar 30, 2012			
	5994348	Jun 07, 2015			
	5994348*PED	Dec 07, 2015			
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN; AVALIDE</u>					
020758 002	5270317	Sep 30, 2011			
	5270317*PED	Mar 30, 2012			
	5994348	Jun 07, 2015			
	5994348*PED	Dec 07, 2015			
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN; AVALIDE</u>					
020758 003	5270317	Sep 30, 2011			
	5270317*PED	Mar 30, 2012			
	5994348	Jun 07, 2015			
	5994348*PED	Dec 07, 2015			
<u>HYDROCHLOROTHIAZIDE; IRBESARTAN; AVALIDE</u>					
020758 004	5270317	Sep 30, 2011	DS DP		
	5270317*PED	Mar 30, 2012			
	5994348	Jun 07, 2015		DP	
	5994348*PED	Dec 07, 2015			
<u>HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM; HYZAAR</u>					
020387 001	5138069	Aug 11, 2009	DS	I-415	Sep 30, 2006
	5138069*PED	Feb 11, 2010			
	5153197	Oct 06, 2009	DP	U-538	
	5153197	Oct 06, 2009	DP	U-3	
	5153197*PED	Apr 06, 2010		U-538	
	5153197*PED	Apr 06, 2010		U-3	
	5608075	Mar 04, 2014			
	5608075*PED	Sep 04, 2014			
<u>HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM; HYZAAR</u>					
020387 002	5138069	Aug 11, 2009	DS	I-415	Sep 30, 2006
	5138069*PED	Feb 11, 2010			
	5153197	Oct 06, 2009	DP	U-3	
	5153197	Oct 06, 2009	DP	U-538	
	5153197*PED	Apr 06, 2010		U-538	
	5153197*PED	Apr 06, 2010		U-3	
	5608075	Mar 04, 2014			
	5608075*PED	Sep 04, 2014			
<u>HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM; HYZAAR</u>					
020387 003	5138069	Aug 11, 2009	DS	I-415	Sep 30, 2006
	5138069*PED	Feb 11, 2010			
	5153197	Oct 08, 2009	DP	U-538	
	5153197	Oct 08, 2009	DP	U-3	
	5153197*PED	Apr 06, 2010			
<u>HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE; UNIRETIC</u>					
020729 001	4743450	Feb 24, 2007			
<u>HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE; UNIRETIC</u>					
020729 002	4743450	Feb 24, 2007			
<u>HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE; UNIRETIC</u>					
020729 003	4743450	Feb 24, 2007			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL; BENICAR HCT</u>					
021532 002	5616599 6878703	Apr 25, 2016 Nov 19, 2021	DS DP	U-500 U-3	NCE NC
					Apr 25, 2007 Jun 05, 2006
<u>HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL; BENICAR HCT</u>					
021532 003	5616599 6878703	Apr 25, 2016 Nov 19, 2021	DS DP	U-500 U-3	NCE NC
					Apr 25, 2007 Jun 05, 2006
<u>HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL; BENICAR HCT</u>					
021532 005	5616599 6878703	Apr 25, 2016 Nov 19, 2021	DS DP	U-500 U-3	NCE NC
					Apr 25, 2007 Jun 05, 2006
<u>HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE; ACCURETIC</u>					
020125 001	4743450 4743450*PED	Feb 24, 2007 Aug 24, 2007			
<u>HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE; ACCURETIC</u>					
020125 002	4743450 4743450*PED	Feb 24, 2007 Aug 24, 2007			
<u>HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE; ACCURETIC</u>					
020125 003	4743450 4743450*PED	Feb 24, 2007 Aug 24, 2007			
<u>HYDROCHLOROTHIAZIDE; TELMISARTAN; MICARDIS HCT</u>					
021162 001	5591762 6358986	Jan 07, 2014 Jan 10, 2020		U-3	
<u>HYDROCHLOROTHIAZIDE; TELMISARTAN; MICARDIS HCT</u>					
021162 002	5591762 6358986	Jan 07, 2014 Jan 10, 2020		U-3	
<u>HYDROCHLOROTHIAZIDE; VALSARTAN; DIOVAN HCT</u>					
020818 001	5399578 6294197	Mar 21, 2012 Jun 18, 2017		U-3 U-3	
<u>HYDROCHLOROTHIAZIDE; VALSARTAN; DIOVAN HCT</u>					
020818 002	5399578 6294197	Mar 21, 2012 Jun 18, 2017		U-3 U-3	
<u>HYDROCHLOROTHIAZIDE; VALSARTAN; DIOVAN HCT</u>					
020818 003	5399578 6294197	Mar 21, 2012 Jun 18, 2017		U-3 U-3	
<u>HYDROCODONE BITARTRATE; IBUPROFEN; VICOPROFEN</u>					
020716 001	6348216 6599531	Jun 10, 2017 Jun 10, 2017			
<u>HYDROCORTISONE BUTYRATE; LOCOID LIPOCREAM</u>					
020769 001	5635497	Jun 03, 2014			
<u>HYDROCORTISONE PROBUTATE; PANDEL</u>					
020453 001	4794106	Dec 27, 2005			
<u>HYDROMORPHONE HYDROCHLORIDE; PALLADONE</u>					
021044 001	5958452 5965161 5968551 6294195 6335033 6706281 6743442	Nov 04, 2014 Nov 04, 2014 Dec 24, 2011 Dec 24, 2011 Nov 04, 2014 Nov 04, 2014 Nov 04, 2014	DP DP DP DP DP DP DP	U-616 U-616 U-616	NDF
					Sep 24, 2007
<u>HYDROMORPHONE HYDROCHLORIDE; PALLADONE</u>					
021044 002	5958452 5965161 5968551 6294195 6335033 6706281 6743442	Nov 04, 2014 Nov 04, 2014 Dec 24, 2011 Dec 24, 2011 Nov 04, 2014 Nov 04, 2014 Nov 04, 2014	DP DP DP DP DP DP DP	U-616 U-616 U-616	NDF
					Sep 24, 2007

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROMORPHONE HYDROCHLORIDE; PALLADONE</u>					
021044 003	5958452	Nov 04, 2014	DP	NDF	Sep 24, 2007
	5965161	Nov 04, 2014	DP U-616		
	5968551	Dec 24, 2011	DP		
	6294195	Dec 24, 2011	DP	U-616	
	6335033	Nov 04, 2014	DP U-616		
	6706281	Nov 04, 2014	DP U-616		
	6743442	Nov 04, 2014	DP		
<u>HYDROMORPHONE HYDROCHLORIDE; PALLADONE</u>					
021044 004	5958452	Nov 04, 2014	DP	NDF	Sep 24, 2007
	5965161	Nov 04, 2014	DP U-616		
	5968551	Dec 24, 2011	DP		
	6294195	Dec 24, 2011	DP	U-616	
	6335033	Nov 04, 2014	DP U-616		
	6706281	Nov 04, 2014	DP U-616		
	6743442	Nov 04, 2014	DP		
<u>IBANDRONATE SODIUM; BONIVA</u>					
021455 001	4927814	Jul 09, 2007		NCE	May 16, 2008
	6143326	Apr 21, 2017			
	6294196	Oct 07, 2019	DP		
<u>IBANDRONATE SODIUM; BONIVA</u>					
021455 002	4927814	Jul 09, 2007	DS DP	D-96 NS NCE	Mar 24, 2008
	6294196	Oct 07, 2019	DP		Mar 24, 2008
<u>IBUPROFEN; CHILDREN'S ADVIL</u>					
019833 002	4788220	Nov 29, 2005			
	4788220*PED	May 29, 2006			
<u>IBUPROFEN; CHILDREN'S ADVIL</u>					
020589 001	4788220	Jul 08, 2007			
	4788220*PED	Jan 08, 2008			
<u>IBUPROFEN; CHILDREN'S ELIXSURE</u>					
021604 001				NP	Jan 07, 2007
<u>IBUPROFEN; CHILDREN'S MOTRIN</u>					
020516 001	5374659	Dec 20, 2011			
	5374659*PED	Jun 20, 2012			
<u>IBUPROFEN; CHILDREN'S MOTRIN</u>					
020601 001	5215755	Jun 01, 2010			
	5215755*PED	Dec 01, 2010			
<u>IBUPROFEN; CHILDREN'S MOTRIN</u>					
020603 001	5374659	Dec 20, 2011			
	5374659*PED	Jun 20, 2012			
<u>IBUPROFEN; IBUPROFEN</u>					
021472 001	6251426	Jun 25, 2018			
<u>IBUPROFEN; JUNIOR STRENGTH MOTRIN</u>					
020601 003	5215755	Jun 01, 2010			
	5215755*PED	Dec 01, 2010			
<u>IBUPROFEN; MOTRIN</u>					
019842 001	5374659	Dec 20, 2011			
	5374659*PED	Jun 20, 2012			
<u>IBUPROFEN; MOTRIN</u>					
020135 001	5215755	Jun 01, 2010			
	5215755*PED	Dec 01, 2010			
	5320855	Jun 14, 2011			
	5320855*PED	Dec 14, 2011			
<u>IBUPROFEN; MOTRIN</u>					
020135 002	5215755	Jun 01, 2010			
	5215755*PED	Dec 01, 2010			
	5320855	Jun 14, 2011			
	5320855*PED	Dec 14, 2011			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IBUPROFEN POTASSIUM; PSEUDOEPHEDRINE HYDROCHLORIDE; ADVIL COLD AND SINUS</u>					
021374 001	5071643	Dec 10, 2008			
	5071643*PED	Jun 10, 2009			
	5360615	Dec 10, 2008			
	5360615*PED	Jun 10, 2009			
<u>IBUPROFEN; OXYCODONE HYDROCHLORIDE; COMBUNOX</u>					
021378 001				NC	Nov 26, 2007
<u>IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE; CHILDREN'S MOTRIN COLD</u>					
021128 001	6211246	Jun 10, 2019			
<u>IBUTILIDE FUMARATE; CORVERT</u>					
020491 001	5155268	Dec 28, 2009			
<u>ICODEXTRIN; EXTRANEAL</u>					
021321 001	4761237	Aug 02, 2005	U-495	I-445	Dec 17, 2007
	4886789	Dec 12, 2006	U-495	NCE	Dec 20, 2007
	6077836	Jun 20, 2017	U-495	ODE	Dec 20, 2009
	6248726	Jun 19, 2018	U-495		
<u>IFOSFAMIDE; IFEX</u>					
019763 001	4882452	Mar 03, 2008			
<u>IFOSFAMIDE; IFEX</u>					
019763 002	4882452	Mar 03, 2008			
<u>IFOSFAMIDE; MESNA; IFEX/MESNEX KIT</u>					
019763 003	4882452	Mar 03, 2008			
<u>IFOSFAMIDE; MESNA; IFEX/MESNEX KIT</u>					
019763 004	4882452	Mar 03, 2008			
<u>ILOPROST; VENTAVIS</u>					
021779 001				M-43	Aug 24, 2008
				NCE	Dec 29, 2009
				ODE	Dec 29, 2011
<u>IMATINIB MESYLATE; GLEEVEC</u>					
021335 001	5521184	Jan 04, 2015		I-392	May 20, 2006
	6894051	May 23, 2019	DS DP U-649	I-376	Dec 20, 2005
				NCE	May 10, 2006
				ODE	Feb 01, 2009
				ODE	May 10, 2008
<u>IMATINIB MESYLATE; GLEEVEC</u>					
021335 002	5521184	Jan 04, 2015		I-392	May 20, 2006
	6894051	May 23, 2019	DS DP U-649	I-376	Dec 20, 2005
				NCE	May 10, 2006
				ODE	Feb 01, 2009
				ODE	May 10, 2008
<u>IMATINIB MESYLATE; GLEEVEC</u>					
021588 001	5521184	Jan 04, 2015		I-392	May 20, 2006
	6894051	May 23, 2019	DS DP U-649	I-376	Dec 20, 2005
				NCE	May 10, 2006
				ODE	Feb 01, 2009
				ODE	May 10, 2008
<u>IMATINIB MESYLATE; GLEEVEC</u>					
021588 002	5521184	Jan 04, 2015		I-392	May 20, 2006
	6894051	May 23, 2019	DS DP U-649	I-376	Dec 20, 2005
				NCE	May 10, 2006
				ODE	Feb 01, 2009
				ODE	May 10, 2008
<u>IMIGLUCERASE; CEREZYME</u>					
020367 001	5549892	Aug 27, 2013		U-252	
<u>IMIGLUCERASE; CEREZYME</u>					
020367 002	5549892	Aug 27, 2013		U-252	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IMIQUIMOD; ALDARA</u>					
020723 001	4689338	Aug 25, 2009	U-172	I-433	Jul 14, 2007
	5238944	Aug 24, 2010		I-420	Mar 02, 2007
<u>INDINAVIR SULFATE; CRIXIVAN</u>					
020685 001	5413999	May 09, 2012	U-132		
	6645961	Mar 04, 2018	DP		
	6689761	Feb 10, 2021	U-554		
<u>INDINAVIR SULFATE; CRIXIVAN</u>					
020685 003	5413999	May 09, 2012	U-132		
	6645961	Mar 04, 2018	DP		
	6689761	Feb 10, 2021	U-554		
<u>INDINAVIR SULFATE; CRIXIVAN</u>					
020685 005	5413999	May 09, 2012	U-132		
	6645961	Mar 04, 2018	DP		
	6689761	Feb 10, 2021	U-554		
<u>INDINAVIR SULFATE; CRIXIVAN</u>					
020685 006	5413999	May 09, 2012	U-132		
	6645961	Mar 04, 2018	DP		
	6689761	Feb 10, 2021	U-554		
<u>INSULIN ASPART PROTAMINE RECOMBINANT; INSULIN ASPART RECOMBINANT; NOVOLOG MIX 70/30</u>					
021172 001	5547930	Sep 28, 2013		M-48	Nov 21, 2008
	5618913	Apr 08, 2014		PED	Dec 07, 2005
	5618913*PED	Oct 08, 2014			
	5834422	Sep 28, 2013	U-471		
	5840680	Sep 28, 2013	U-471		
	5866538	Jun 19, 2017	DP		
	5948751	Jun 19, 2017			
<u>INSULIN ASPART RECOMBINANT; NOVOLOG</u>					
020986 001	5618913	Apr 08, 2014		M-44	Sep 13, 2008
	5618913*PED	Oct 08, 2014		PED	Mar 13, 2009
	5866538	Jun 20, 2017		PED	Dec 07, 2005
	5866538*PED	Dec 20, 2017			
<u>INSULIN DETEMIR RECOMBINANT; LEVEMIR</u>					
021536 001	5750497	May 12, 2015	DS DP U-668	NCE	Jun 16, 2010
	5866538	Jun 20, 2017	DP		
	6011007	Feb 02, 2014	DS DP U-668		
	6869930	Feb 02, 2014	DS DP U-668		
<u>INSULIN GLARGINE RECOMBINANT; LANTUS</u>					
021081 001	5656722	Sep 12, 2014		D-80	May 01, 2006
	5656722*PED	Mar 12, 2015		PED	Oct 20, 2005
<u>INSULIN GLULISINE RECOMBINANT; APIDRA</u>					
021629 001	6221633	Jun 18, 2018	DS DP U-471	NCE	Apr 16, 2009
<u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT; HUMALOG MIX 50/50</u>					
021018 001	5461031	Jun 26, 2014			
	5474978	Jun 16, 2014			
	5514646	May 07, 2013			
	5747642	Jun 16, 2014			
<u>INSULIN LISPRO PROTAMINE RECOMBINANT; INSULIN LISPRO RECOMBINANT; HUMALOG MIX 75/25</u>					
021017 001	5461031	Jun 16, 2014			
	5474978	Jun 16, 2014			
	5514646	May 07, 2013			
	5747642	Jun 16, 2014			
<u>INSULIN LISPRO RECOMBINANT; HUMALOG</u>					
020563 001	5474978	Jun 16, 2014	U-534	D-86	Jun 02, 2007
	5514646	May 07, 2013	U-534		
<u>INSULIN LISPRO RECOMBINANT; HUMALOG PEN</u>					
020563 002	5474978	Jun 16, 2014	U-534	D-86	Jun 02, 2007
	5514646	May 07, 2013	U-534		
<u>IODIXANOL; VISIPAQUE 270</u>					
020351 001	5349085	Sep 20, 2011			
<u>IODIXANOL; VISIPAQUE 270</u>					
020808 001	5349085	Sep 20, 2011			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IODIXANOL; VISIPAQUE 320</u>					
020351 002	5349085	Sep 20, 2011			
<u>IODIXANOL; VISIPAQUE 320</u>					
020808 002	5349085	Sep 20, 2011			
<u>IOXILAN; OXILAN-300</u>					
020316 001	4954348	Dec 21, 2009			
<u>IOXILAN; OXILAN-350</u>					
020316 002	4954348	Dec 21, 2009			
<u>IPRATROPIUM BROMIDE; ATROVENT HFA</u>					
021527 001	5676930	Oct 14, 2014	DP	NP	Nov 17, 2007
	5683677	Nov 04, 2014	DP		
	5695743	Jul 06, 2010	DP		
	5766573	Nov 28, 2009			
	6739333	May 26, 2020	DP		
<u>IRBESARTAN; AVAPRO</u>					
020757 001	5270317	Sep 30, 2011		I-373 PED	Sep 17, 2005
	5270317*PED	Mar 30, 2012			Mar 17, 2006
	6342247	Jun 07, 2015			
	6342247*PED	Dec 07, 2015			
<u>IRBESARTAN; AVAPRO</u>					
020757 002	5270317	Sep 30, 2011		I-373 PED	Sep 17, 2005
	5270317*PED	Mar 30, 2012			Mar 17, 2006
	6342247	Jun 07, 2015			
	6342247*PED	Dec 07, 2015			
<u>IRBESARTAN; AVAPRO</u>					
020757 003	5270317	Sep 30, 2011		I-373 PED	Sep 17, 2005
	5270317*PED	Mar 30, 2012			Mar 17, 2006
	6342247	Jun 07, 2015			
	6342247*PED	Dec 07, 2015			
<u>IRINOTECAN HYDROCHLORIDE; CAMPTOSAR</u>					
020571 001	4604463	Aug 20, 2007		M-34 PED	Jun 24, 2007
	4604463*PED	Feb 20, 2008			Dec 24, 2007
	6403569	Apr 28, 2020	U-449		
	6403569*PED	Oct 28, 2020			
	6794370	May 01, 2020	U-606		
	6794370*PED	Nov 01, 2020			
<u>IRON DEXTRAN; DEXFERRUM</u>					
040024 001	5624668	Sep 29, 2015			
<u>IRON SUCROSE; VENOFER</u>					
021135 001				I-474 I-459	Oct 17, 2008
					Jun 17, 2008
<u>IRON SUCROSE; VENOFER</u>					
021135 002				I-474 I-459	Oct 17, 2008
					Jun 17, 2008
<u>IRON SUCROSE; VENOFER</u>					
021135 003				I-474 I-459	Oct 17, 2008
					Jun 17, 2008
<u>ISOTRETINOIN; ACCUTANE</u>					
018662 002				PED	Nov 02, 2005
<u>ISOTRETINOIN; ACCUTANE</u>					
018662 003				PED	Nov 02, 2005
<u>ISOTRETINOIN; ACCUTANE</u>					
018662 004				PED	Nov 02, 2005
<u>ISRADIPINE; DYNACIRC CR</u>					
020336 001	4816263	Oct 02, 2007	U-3		
	4946687	Aug 07, 2007			
	4950486	Aug 21, 2007			
	5030456	Jul 09, 2008			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>ISRADIPINE; DYNACIRC CR</u>						
020336 002	4816263	Oct 02, 2007	U-3			
	4946687	Aug 07, 2007				
	4950486	Aug 21, 2007				
	5030456	Jul 09, 2008				
<u>ITRACONAZOLE; ITRACONAZOLE</u>						
076104 001				PC	Aug 08, 2005	
<u>ITRACONAZOLE; SPORANOX</u>						
020083 001	5633015	May 27, 2014				
<u>ITRACONAZOLE; SPORANOX</u>						
020657 001	5707975	Jan 13, 2015				
	6407079	Jun 18, 2019				
<u>ITRACONAZOLE; SPORANOX</u>						
020966 001	6407079	Jun 18, 2019				
<u>KETOCONAZOLE; NIZORAL A-D</u>						
020310 001	5456851	Apr 07, 2014				
<u>KETOROLAC TROMETHAMINE; ACULAR</u>						
019700 001	5110493	May 05, 2009	U-75 U-75			
	5110493*PED	Nov 05, 2009				
<u>KETOROLAC TROMETHAMINE; ACULAR LS</u>						
021528 001	5110493	May 05, 2009		NP	May 30, 2006	
	5110493*PED	Nov 05, 2009				
<u>KETOTIFEN FUMARATE; ZADITOR</u>						
021066 001	6774137	Jan 13, 2019	DP DP	U-599		
	6776982	Jan 13, 2019				
	6777429	Jan 13, 2019				
<u>LAMIVUDINE; EPIVIR</u>						
020564 001	5047407	Nov 17, 2009				
	5047407*PED	May 17, 2010				
	5905082	May 18, 2016				
	5905082*PED	Nov 18, 2016				
<u>LAMIVUDINE; EPIVIR</u>						
020564 003	5047407	Nov 17, 2009				
	5047407*PED	May 17, 2010				
	5905082	May 18, 2016				
	5905082*PED	Nov 18, 2016				
	6180639	Jan 30, 2018		U-248		
	6180639*PED	Jul 30, 2018		U-248		
<u>LAMIVUDINE; EPIVIR</u>						
020596 001	5047407	Nov 17, 2009				
	5047407*PED	May 17, 2010				
	6004968	Mar 20, 2018				
	6004968*PED	Sep 20, 2018				
<u>LAMIVUDINE; EPIVIR-HBV</u>						
021003 001	5047407	Nov 17, 2009				
	5047407*PED	May 17, 2010				
	5532246	Jul 02, 2013		U-250		
	5532246*PED	Jan 02, 2014		U-250		
	5905082	May 18, 2016				
	5905082*PED	Nov 18, 2016				
<u>LAMIVUDINE; EPIVIR-HBV</u>						
021004 001	5047407	Nov 17, 2009				
	5047407*PED	May 17, 2010				
	5532246	Jul 02, 2013		U-250		
	5532246*PED	Jan 02, 2014		U-250		
	6004968	Mar 20, 2018				
	6004968*PED	Sep 20, 2018				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
LAMIVUDINE; ZIDOVUDINE; COMBIVIR					
020857 001	4724232	Sep 17, 2005			
	4818538	Sep 17, 2005			
	4828838	Sep 17, 2005			
	4833130	Sep 17, 2005			
	4837208	Sep 17, 2005			
	5047407	Nov 17, 2009			
	5047407*PED	May 17, 2010			
	5859021	May 15, 2012	U-248		
	5905082	May 18, 2016	U-248		
	5905082*PED	Nov 18, 2016	U-248		
6113920	Oct 23, 2017	U-257			
LAMOTRIGINE; LAMICTAL					
020241 001	4602017	Jul 22, 2008	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL					
020241 002	4602017	Jul 22, 2008	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL					
020241 003	4602017	Jul 22, 2008	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL					
020241 004	4602017	Jul 22, 2008	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL					
020241 005	4602017	Jul 22, 2008	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL					
020241 006	4602017	Jul 22, 2008	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL CD					
020764 001	4602017 5698226	Jul 22, 2008 Jan 29, 2012	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL CD					
020764 002	4602017 5698226	Jul 22, 2008 Jan 29, 2012	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL CD					
020764 003	4602017 5698226	Jul 22, 2008 Jan 29, 2012	U-106	I-404 I-387 ODE	Jun 20, 2006 Jan 17, 2006 Aug 24, 2005
LAMOTRIGINE; LAMICTAL CD					
020764 004	4602017 5698226	Jul 22, 2008 Jan 29, 2012	U-106	I-404 I-387 ODE	Jun 10, 2006 Jan 17, 2006 Aug 24, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LANSOPRAZOLE; PREVACID</u>					
020406 001	4628098	May 10, 2009		M-12	Jul 31, 2005
	4689333	Jul 29, 2005	U-126	NPP	Jun 17, 2007
	5013743	Feb 12, 2010	U-452		
	5026560	Jun 25, 2008			
	5045321	Sep 03, 2008			
	5093132	Sep 03, 2008			
	5433959	Sep 03, 2008			
<u>LANSOPRAZOLE; PREVACID</u>					
020406 002	4628098	May 10, 2009		M-12	Jul 31, 2005
	4689333	Jul 29, 2005	U-126	NPP	Jun 17, 2007
	5013743	Feb 12, 2010	U-452		
	5026560	Jun 25, 2008			
	5045321	Sep 03, 2008			
	5093132	Sep 03, 2008			
	5433959	Sep 03, 2008			
<u>LANSOPRAZOLE; PREVACID</u>					
021281 001	4628098	May 10, 2009		M-12	Aug 30, 2005
	4689333	Jul 29, 2005	U-126	NPP	Jun 17, 2007
	5013743	Feb 12, 2010	U-452		
	5026560	Jun 25, 2008			
	5045321	Sep 03, 2008			
	5093132	Sep 03, 2008			
	5433959	Sep 03, 2008			
<u>LANSOPRAZOLE; PREVACID</u>					
021281 002	4628098	May 10, 2009		M-12	Aug 30, 2005
	4689333	Jul 29, 2005	U-126	NPP	Jun 17, 2007
	5013743	Feb 12, 2010	U-452		
	5026560	Jun 25, 2008			
	5045321	Sep 03, 2008			
	5093132	Sep 03, 2008			
	5433959	Sep 03, 2008			
<u>LANSOPRAZOLE; PREVACID</u>					
021428 001	4628098	May 10, 2009		M-12	Aug 30, 2005
	4689333	Jul 29, 2005	U-126	NPP	Jun 17, 2007
	5013743	Feb 12, 2010	U-452		
	5026560	Jun 25, 2008			
	5045321	Sep 03, 2008			
	5093132	Sep 03, 2008			
	5433959	Sep 03, 2008			
	5464632	Nov 07, 2012			
	6123962	Feb 13, 2007			
	6328994	May 17, 2019			
<u>LANSOPRAZOLE; PREVACID</u>					
021428 002	4628098	May 10, 2009		M-12	Aug 30, 2005
	4689333	Jul 29, 2005	U-126	NPP	Jun 17, 2007
	5013743	Feb 12, 2010	U-452		
	5026560	Jun 25, 2008			
	5045321	Sep 03, 2008			
	5093132	Sep 03, 2008			
	5433959	Sep 03, 2008			
	5464632	Nov 07, 2012			
	6123962	Feb 13, 2007			
	6328994	May 17, 2019			
<u>LANSOPRAZOLE; PREVACID IV</u>					
021566 001				NDF	May 27, 2007
<u>LANTHANUM CARBONATE; FOSRENOL</u>					
021468 001	5968976	Mar 19, 2016	DP U-613	NCE	Oct 26, 2009
<u>LANTHANUM CARBONATE; FOSRENOL</u>					
021468 002	5968976	Mar 19, 2016	DP U-613	NCE	Oct 26, 2009
<u>LANTHANUM CARBONATE; FOSRENOL</u>					
021468 003				NCE	Oct 26, 2009
<u>LANTHANUM CARBONATE; FOSRENOL</u>					
021468 004				NCE	Oct 26, 2009

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE				
<u>LATANOPROST; XALATAN</u>											
020597 001	4599353	Jul	28, 2006		U-260	I-375	Dec	20, 2005			
	5296504	Mar	22, 2011		U-260						
	5422368	Mar	22, 2011		U-260						
	6429226	Sep	06, 2009		U-260						
<u>LEFLUNOMIDE; ARAVA</u>											
020905 001						I-395	Jun	13, 2006			
						M-32	Mar	05, 2007			
						PED	Sep	05, 2007			
						PED	Dec	13, 2006			
<u>LEFLUNOMIDE; ARAVA</u>											
020905 002						I-395	Jun	13, 2006			
						M-32	Mar	05, 2007			
						PED	Sep	05, 2007			
						PED	Dec	13, 2006			
<u>LEFLUNOMIDE; ARAVA</u>											
020905 003						I-395	Jun	13, 2006			
						M-32	Mar	05, 2007			
						PED	Sep	05, 2007			
						PED	Dec	13, 2006			
<u>LENALIDOMIDE; REVLIMID</u>											
021880 001						NCE	Dec	27, 2010			
<u>LENALIDOMIDE; REVLIMID</u>											
021880 002						NCE	Dec	27, 2010			
<u>LEPIRUDIN RECOMBINANT; REFLUDAN</u>											
020807 001	5180668	Jan	19, 2010								
<u>LETROZOLE; FEMARA</u>											
020726 001	4978672	Jun	03, 2011		U-203	I-481	Dec	28, 2008			
						I-446	Oct	29, 2007			
<u>LEUPROLIDE ACETATE; ELIGARD</u>											
021343 001	4938763	Oct	03, 2008								
	5278201	Jan	11, 2011								
	5324519	Oct	20, 2011								
	5599552	Feb	04, 2014								
	5733950	Oct	03, 2008								
	5739176	Oct	03, 2008								
<u>LEUPROLIDE ACETATE; ELIGARD</u>											
021379 001	4938763	Oct	03, 2008			NP	Jul	24, 2005			
	5278201	Jan	11, 2011								
	5324519	Oct	20, 2011								
	5599552	Feb	04, 2014								
	5733950	Oct	03, 2008								
	5739176	Oct	03, 2008								
<u>LEUPROLIDE ACETATE; ELIGARD</u>											
021488 001	4938763	Oct	03, 2008			NP	Feb	13, 2006			
	5278201	Jan	11, 2011								
	5324519	Oct	20, 2011								
	5599552	Feb	04, 2014								
	5733950	Oct	03, 2008								
	5739176	Oct	03, 2008								
<u>LEUPROLIDE ACETATE; ELIGARD</u>											
021731 001	4938763	Oct	03, 2008	DP	U-621	NP	Dec	14, 2007			
	5278201	Jan	11, 2011	DP							
	5324519	Jun	28, 2011	DP							
	5599552	Feb	04, 2014	DP	U-621						
	5739176	Oct	03, 2008	DP	U-621						
	6395293	Sep	28, 2013	DP							
	6565874	Oct	28, 2018	DP	U-621						
	6626870	Mar	27, 2020	DP							
	6773714	Oct	28, 2018		U-621						
	RE37950	Oct	03, 2008	DP	U-621						

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEUPROLIDE ACETATE; LUPRON DEPOT</u>					
019732 001	4677191	Jul 03, 2005			
	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5575987	Sep 02, 2013			
	5631020	May 20, 2014			
	5716640	Sep 02, 2013			
	6036976	Dec 13, 2016			
<u>LEUPROLIDE ACETATE; LUPRON DEPOT</u>					
020011 001	4677191	Jul 03, 2005			
	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5575987	Sep 02, 2013			
	5631021	May 20, 2014			
	5716640	Sep 02, 2013			
<u>LEUPROLIDE ACETATE; LUPRON DEPOT</u>					
020517 001	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5480656	Jan 02, 2013			
	5575987	Nov 19, 2013			
	5631020	May 20, 2014			
	5643607	Jan 02, 2013			
	5716640	Sep 02, 2013			
	5814342	Feb 01, 2011			
	6036976	Dec 13, 2016			
<u>LEUPROLIDE ACETATE; LUPRON DEPOT-3</u>					
020708 001	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5480656	Jan 02, 2013			
	5575987	Nov 19, 2013			
	5631020	May 20, 2014			
	5643607	Jan 02, 2013			
	5716640	Sep 02, 2013			
	5814342	Feb 01, 2011			
	6036976	Dec 13, 2016			
<u>LEUPROLIDE ACETATE; LUPRON DEPOT-4</u>					
020517 002	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5480656	Jan 02, 2013			
	5575987	Nov 19, 2013			
	5631020	May 20, 2014			
	5643607	Jan 02, 2013			
	5716640	Sep 02, 2013			
	5814342	Feb 01, 2011			
	6036976	Dec 13, 2016			
<u>LEUPROLIDE ACETATE; LUPRON DEPOT-PED</u>					
020263 002	4677191	Jul 03, 2005			
	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5575987	Sep 02, 2013			
	5631020	May 20, 2014			
	5716640	Sep 02, 2013			
	6036976	Dec 13, 2016			
<u>LEUPROLIDE ACETATE; LUPRON DEPOT-PED</u>					
020263 003	4677191	Jul 03, 2005			
	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5575987	Sep 02, 2013			
	5631020	May 20, 2014			
	5716640	Sep 02, 2013			
020263 003	6036976	Dec 13, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEUPROLIDE ACETATE; LUPRON DEPOT-PED</u>					
020263 004	4677191	Jul 03, 2005			
	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5575987	Sep 02, 2013			
	5631020	May 20, 2014			
	5716640	Sep 02, 2013			
	6036976	Dec 13, 2013			
<u>LEUPROLIDE ACETATE; LUPRON DEPOT-PED</u>					
020263 005	4677191	Jul 03, 2005			
	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5575987	Sep 02, 2013			
	5631020	May 20, 2014			
	5716640	Sep 02, 2013			
	6036976	Dec 13, 2016			
<u>LEUPROLIDE ACETATE; LUPRON DEPOT-PED</u>					
020263 006	4677191	Jul 03, 2005			
	4728721	May 01, 2006			
	4849228	Jul 18, 2006			
	5575987	Sep 02, 2013			
	5631020	May 20, 2014			
	5716640	Sep 02, 2013			
	6036976	Dec 13, 2016			
<u>LEUPROLIDE ACETATE; VIADUR</u>					
021088 001	5728396	Jan 30, 2017		U-316	
	5932547	Jun 13, 2017			
	5985305	Jan 30, 2017			
	6113938	Jul 24, 2018			
	6124261	Jun 13, 2017			
	6132420	Jan 30, 2017			
	6156331	Jan 30, 2017			
	6235712	Jun 13, 2017			
	6375978	Dec 17, 2018			
	6395292	Jan 30, 2017			
<u>LEVALBUTEROL HYDROCHLORIDE; XOPENEX</u>					
020837 001	5362755	Nov 08, 2011		U-332	
	5547994	Aug 20, 2013		U-332	
	5760090	Jan 05, 2010		U-332	
	5844002	Jan 05, 2010		U-332	
	6083993	Jan 05, 2010		U-332	
	6451289	Mar 21, 2021			
<u>LEVALBUTEROL HYDROCHLORIDE; XOPENEX</u>					
020837 002	5362755	Nov 08, 2011		U-332	
	5547994	Aug 20, 2013		U-332	
	5760090	Jan 05, 2010		U-332	
	5844002	Jan 05, 2010		U-332	
	6083993	Jan 05, 2010		U-332	
	6451289	Mar 21, 2021			
<u>LEVALBUTEROL HYDROCHLORIDE; XOPENEX</u>					
020837 003	5362755	Nov 08, 2011		U-332	
	5547994	Aug 20, 2013		U-332	
	5760090	Jan 05, 2010		U-332	
	5844002	Jan 05, 2010		U-332	
	6083993	Jan 05, 2010		U-332	
	6451289	Mar 21, 2021			
<u>LEVALBUTEROL HYDROCHLORIDE; XOPENEX</u>					
020837 004	5362755	Nov 08, 2011		U-332	
	5547994	Aug 20, 2013		U-332	
	5760090	Jan 05, 2010		U-332	
	5844002	Jan 05, 2010		U-5	
	6083993	Jan 05, 2010		U-332	
	6451289	Mar 21, 2021	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVALBUTEROL TARTRATE; XOPENEX HFA</u>					
021730 001	5225183	Jul 06, 2010	DP	NP	Mar 11, 2008
	5362755	Nov 08, 2011	U-636		
	5439670	Jul 06, 2010	DP		
	5547994	Aug 20, 2013	U-636		
	5605674	Feb 25, 2014	DP		
	5695743	Jul 06, 2010	DP	U-636	
	5760090	Jan 05, 2010	U-636		
	5836299	Nov 17, 2017	DP		
	5844002	Jan 05, 2010	U-636		
	6083993	Jan 05, 2010	U-636		
	6352684	Nov 28, 2009	DP		
<u>LEVETIRACETAM; KEPPRA</u>					
021035 001	4943639	Jul 14, 2008	DS	NPP	Jun 21, 2008
<u>LEVETIRACETAM; KEPPRA</u>					
021035 002	4943639	Jul 14, 2008	DS	NPP	Jun 21, 2008
<u>LEVETIRACETAM; KEPPRA</u>					
021035 003	4943639	Jul 14, 2008	DS	NPP	Jun 21, 2008
<u>LEVETIRACETAM; KEPPRA</u>					
021505 001	4943639	Jul 14, 2008	DS	NPP	Jun 21, 2008
<u>LEVOBETAXOLOL HYDROCHLORIDE; BETAXON</u>					
021114 001	4911920	Mar 27, 2007	U-369		
	5540918	Jul 30, 2013			
<u>LEVOPUIVACAINE HYDROCHLORIDE; CHIROCAINE</u>					
020997 001	5708011	Oct 13, 2014	U-276		
<u>LEVOPUIVACAINE HYDROCHLORIDE; CHIROCAINE</u>					
020997 002	5708011	Oct 13, 2014	U-276		
<u>LEVOPUIVACAINE HYDROCHLORIDE; CHIROCAINE</u>					
020997 003	5708011	Oct 13, 2014	U-276		
<u>LEVOCARNITINE; CARNITOR</u>					
020182 001	6335369	Jan 18, 2021	U-433	ODE	Dec 15, 2006
	6429230	Jan 18, 2021	U-433		
	6696493	Jan 18, 2021	U-433		
<u>LEVOFLOXACIN; IQUIX</u>					
021571 001	5053407	Dec 20, 2010	DS DP U-600	NP	Mar 01, 2007
<u>LEVOFLOXACIN; LEVAQUIN</u>					
020634 001	5053407	Dec 20, 2010		D-83 D-100 I-393 I-372	Oct 23, 2006 Aug 04, 2008 May 23, 2006 Oct 30, 2005
<u>LEVOFLOXACIN; LEVAQUIN</u>					
020634 002	5053407	Dec 20, 2010		D-83 D-100 I-393 I-372	Oct 23, 2006 Aug 04, 2008 May 23, 2006 Oct 30, 2005
<u>LEVOFLOXACIN; LEVAQUIN</u>					
020634 003	5053407	Dec 20, 2010		D-83 D-100 I-393 I-372	Oct 23, 2006 Aug 04, 2008 May 23, 2006 Oct 30, 2005
<u>LEVOFLOXACIN; LEVAQUIN</u>					
020635 001	5053407	Dec 20, 2010		D-83 D-100 I-393 I-372	Oct 23, 2006 Aug 04, 2008 May 23, 2006 Oct 30, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOFLOXACIN; LEVAQUIN</u>					
020635 004	5053407	Dec 20, 2010		D-83 D-100 I-393 I-372	Oct 23, 2006 Aug 04, 2008 May 23, 2006 Oct 30, 2005
<u>LEVOFLOXACIN; LEVAQUIN</u>					
021721 001				D-100	Aug 04, 2008
<u>LEVOFLOXACIN; LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
020635 002	5053407	Dec 20, 2010		D-83 D-100 I-393 I-372	Oct 23, 2006 Aug 04, 2008 May 23, 2006 Oct 30, 2005
<u>LEVOFLOXACIN; LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
020635 003	5053407	Dec 20, 2010		D-83 D-100 I-393 I-372	Oct 23, 2006 Aug 04, 2008 May 23, 2006 Oct 30, 2005
<u>LEVOFLOXACIN; LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
020635 005	5053407	Dec 20, 2010		D-83 D-100 I-393 I-372	Oct 23, 2006 Aug 04, 2008 May 23, 2006 Oct 30, 2005
<u>LEVOFLOXACIN; QUIXIN</u>					
021199 001	5053407	Dec 20, 2010			
<u>LEVONORGESTREL; MIRENA</u>					
021225 001	5785053	Dec 05, 2015	DP		
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 001	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 002	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 003	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 004	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 005	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 006	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 007	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 008	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 009	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 010	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVO-T</u>					
021342 011	6399101	Mar 30, 2020			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 001	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 002	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 003	6555581	Feb 15, 2022			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 004	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 005	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 006	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 007	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 008	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 009	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 010	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 011	6555581	Feb 15, 2022			
<u>LEVOTHYROXINE SODIUM; LEVOXYL</u>					
021301 012	6555581	Feb 15, 2022			
<u>LIDOCAINE; LIDOCAINE</u>					
020575 001	5234957	Feb 27, 2011			
	5332576	Jul 26, 2011			
	5446070	Feb 27, 2011			
<u>LIDOCAINE; LIDOCAINE</u>					
020575 002	5234957	Feb 27, 2011			
	5332576	Jul 26, 2011			
	5446070	Feb 27, 2011			
<u>LIDOCAINE; LIDODERM</u>					
020612 001	5411738	May 02, 2012		ODE	Mar 19, 2006
	5589180	Mar 17, 2009	U-485		
	5601838	May 02, 2012	U-488		
	5709869	Mar 17, 2009	U-485		
	5827529	Oct 27, 2015	U-486		
<u>LIDOCAINE; PRILOCAINE; ORAQIX</u>					
021451 001	6031007	Apr 01, 2017	DP U-553	NDF	Dec 19, 2006
<u>LIDOCAINE; TETRACAINE; SYNERA</u>					
021623 001	5658583	Jul 28, 2015	DP		
	5919479	Jul 28, 2015	DP		
	6306431	Jul 28, 2015	DP		
	6465006	Jul 28, 2015	DP		
	6546281	Jul 28, 2015	DP		
	6780426	Jul 28, 2015	DP		
<u>LINEZOLID; ZYVOX</u>					
021130 001	5688792	Nov 18, 2014	DS U-319	I-431	Jun 23, 2007
	5688792*PED	May 18, 2015		I-402	Jul 22, 2006
	6514529	Mar 15, 2021	DP	NPP	Dec 19, 2005
	6514529*PED	Sep 15, 2021		PED	Dec 23, 2007
	6559305	Jan 29, 2021	DS	PED	Jan 22, 2007
	6559305*PED	Jul 29, 2021		PED	Jun 19, 2006
				PED	Oct 18, 2005
<u>LINEZOLID; ZYVOX</u>					
021130 002	5688792	Nov 18, 2014	DS U-319	I-431	Jun 23, 2007
	5688792*PED	May 18, 2015		I-402	Jul 22, 2006
	6514529	Mar 15, 2021	DP	NPP	Dec 19, 2005
	6514529*PED	Sep 15, 2021		PED	Dec 23, 2007
	6559305	Jan 29, 2021	DS	PED	Jan 22, 2007
	6559305*PED	Jul 29, 2021		PED	Jun 19, 2006
				PED	Oct 18, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LINEZOLID; ZYVOX</u>					
021131 001	5688792	Nov 18, 2014	DS	I-431	Jun 23, 2007
	5688792*PED	May 18, 2015		I-402	Jul 22, 2006
	6559305	Jan 29, 2021		NPP	Dec 19, 2005
	6559305*PED	Jul 29, 2021		PED	Dec 23, 2007
				PED	Jan 22, 2007
				PED	Jun 19, 2006
				PED	Oct 18, 2005
<u>LINEZOLID; ZYVOX</u>					
021132 001	5688792	Nov 18, 2014	DS	I-431	Jun 23, 2007
	5688792*PED	May 18, 2015		I-402	Jul 22, 2006
	6559305	Jan 29, 2021	DS	NPP	Dec 19, 2005
	6559305*PED	Jul 29, 2021		PED	Dec 23, 2007
				PED	Jan 22, 2007
				PED	Jun 19, 2006
				PED	Oct 18, 2005
<u>LISINAPRIL; PRINIVIL</u>					
019558 001				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; PRINIVIL</u>					
019558 002				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; PRINIVIL</u>					
019558 003				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; PRINIVIL</u>					
019558 004				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; PRINIVIL</u>					
019558 006				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; ZESTRIL</u>					
019777 001				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; ZESTRIL</u>					
019777 002				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; ZESTRIL</u>					
019777 003				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; ZESTRIL</u>					
019777 004				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; ZESTRIL</u>					
019777 005				NPP	May 29, 2006
				PED	Nov 29, 2006
<u>LISINAPRIL; ZESTRIL</u>					
019777 006				NPP	May 29, 2006
<u>LODOXAMIDE TROMETHAMINE; ALOMIDE</u>					
020191 001	5457126	Oct 10, 2012		U-117	
<u>LOMEFLOXACIN HYDROCHLORIDE; MAXAQUIN</u>					
020013 001	4528287	Feb 21, 2006		U-36	
<u>LOPERAMIDE HYDROCHLORIDE; IMODIUM A-D</u>					
020448 001	5489436	Feb 06, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LOPERAMIDE HYDROCHLORIDE; SIMETHICONE; IMODIUM ADVANCED</u>					
020606 001	5248505	Jul 28, 2010			
	5489436	Feb 06, 2013			
	5612054	Sep 28, 2010			
	5679376	Oct 21, 2014			
	5716641	May 21, 2012	U-226		
<u>LOPERAMIDE HYDROCHLORIDE; SIMETHICONE; IMODIUM ADVANCED</u>					
021140 001	5248505	Sep 28, 2010	U-434		
	5612054	Sep 28, 2010			
	6103260	Jul 17, 2017			
<u>LOPINAVIR; RITONAVIR; KALETRA</u>					
021226 001	5541206	Jul 30, 2013	U-348	D-99	Apr 29, 2008
	5541206*PED	Jan 30, 2014		PED	Oct 29, 2008
	5635523	Jun 30, 2014	U-352		
	5635523*PED	Dec 30, 2014			
	5648497	Jul 15, 2014			
	5648497*PED	Jan 15, 2015			
	5674882	Oct 07, 2014	U-344		
	5674882*PED	Apr 07, 2015			
	5846987	Dec 29, 2012	U-350		
	5846987*PED	Jun 29, 2013			
	5886036	Dec 29, 2012	U-345		
	5886036*PED	Jun 29, 2013			
	5914332	Dec 13, 2015	U-351		
	5914332*PED	Jun 13, 2016			
	6037157	Jun 26, 2016	U-346		
	6037157*PED	Dec 26, 2016			
	6232333	Nov 07, 2017			
	6232333*PED	May 07, 2018			
	6284767	Feb 14, 2016	U-401		
	6458818	Nov 07, 2017			
	6458818*PED	May 07, 2018			
	6521651	Nov 01, 2017			
	6521651*PED	May 01, 2018			
	6703403	Jun 26, 2016	U-257		
	6703403*PED	Dec 26, 2016			
<u>LOPINAVIR; RITONAVIR; KALETRA</u>					
021251 001	5541206	Jul 30, 2013	U-348	D-99	Apr 29, 2008
	5541206*PED	Jan 30, 2014		PED	Oct 29, 2008
	5635523	Jun 03, 2014	U-352		
	5635523*PED	Dec 03, 2014			
	5648497	Jul 15, 2014			
	5648497*PED	Jan 15, 2015			
	5674882	Oct 07, 2014	U-344		
	5674882*PED	Apr 07, 2015			
	5846987	Dec 29, 2012	U-350		
	5846987*PED	Jun 29, 2013			
	5886036	Dec 29, 2012	U-345		
	5886036*PED	Jun 29, 2013			
	5914332	Dec 13, 2005	U-351		
	5914332*PED	Jun 13, 2006			
	6037157	Jun 26, 2016	U-346		
	6037157*PED	Dec 26, 2016			
	6284767	Feb 14, 2016	U-401		
	6703403	Jun 26, 2016	U-257		
	6703403*PED	Dec 26, 2016			
	6911214	Nov 28, 2021	DP		
	6911214*PED	May 28, 2022			
<u>LOPINAVIR; RITONAVIR; KALETRA</u>					
021906 001	5541206	Jul 30, 2013	DS	DP	U-688
	5635523	Jun 03, 2014			U-688
	5648497	Jul 15, 2014	DS	DP	
	5674882	Oct 07, 2014			U-688
	5846987	Dec 29, 2012			U-688
	5886036	Dec 29, 2012		DP	
	5914332	Dec 13, 2015	DS	DP	U-688
	6037157	Jun 26, 2016			U-688
	6284767	Feb 14, 2016		DP	U-688
	6703403	Jun 26, 2016			U-688

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LORATADINE; CLARITIN</u>					
019658 002	4863931	Sep 15, 2008			
	4863931*PED	Mar 15, 2009			
<u>LORATADINE; CLARITIN</u>					
020641 002	4863931	Sep 15, 2008			
	4863931*PED	Mar 15, 2009			
	6132758	Jun 01, 2018			
<u>LORATADINE; CLARITIN REDITABS</u>					
020704 002	4863931	Sep 15, 2008			
	4863931*PED	Mar 15, 2009			
<u>LORATADINE; PSEUDOEPHEDRINE SULFATE; CLARITIN-D</u>					
019670 002	4863931	Sep 15, 2008			
	4863931*PED	Mar 15, 2009			
<u>LORATADINE; PSEUDOEPHEDRINE SULFATE; CLARITIN-D 24 HOUR</u>					
020470 002	4863931	Sep 15, 2008			
	4863931*PED	Mar 15, 2009			
	5314697	Oct 23, 2012			
<u>LOSARTAN POTASSIUM; COZAAR</u>					
020386 001	5138069	Aug 11, 2009		I-383	Sep 17, 2005
	5138069*PED	Feb 11, 2010		NPP	Mar 11, 2007
	5153197	Oct 06, 2009	U-3	PED	Sep 11, 2007
	5153197*PED	Apr 06, 2010	U-3		
	5210079	May 11, 2010	U-496		
	5210079*PED	Nov 11, 2010	U-496		
	5608075	Mar 04, 2014			
	5608075*PED	Sep 04, 2014			
<u>LOSARTAN POTASSIUM; COZAAR</u>					
020386 002	5138069	Aug 11, 2009		I-383	Sep 17, 2005
	5138069*PED	Feb 11, 2010		NPP	Mar 11, 2007
	5153197	Oct 06, 2009	U-3	PED	Sep 11, 2007
	5153197*PED	Apr 06, 2010	U-3		
	5210079	May 11, 2010	U-496		
	5210079*PED	Nov 11, 2010	U-496		
	5608075	Mar 04, 2014			
	5608075*PED	Sep 04, 2014			
<u>LOSARTAN POTASSIUM; COZAAR</u>					
020386 003	5138069	Aug 11, 2009		I-383	Sep 17, 2005
	5138069*PED	Feb 11, 2010		NPP	Mar 11, 2007
	5153197	Oct 06, 2009	U-3	PED	Sep 11, 2007
	5153197*PED	Apr 06, 2010	U-3		
	5210079	May 11, 2010	U-496		
	5210079*PED	Nov 11, 2010	U-496		
	5608075	Mar 04, 2014			
	5608075*PED	Sep 04, 2014			
<u>LOTEPREDNOL ETABONATE; ALREX</u>					
020803 001	4996335	Mar 09, 2012			
	5540930	Oct 25, 2013			
	5747061	Oct 25, 2013	DP	U-576	
<u>LOTEPREDNOL ETABONATE; LOTEMAX</u>					
020583 001	4996335	Mar 09, 2012			
	5540930	Oct 25, 2013			
	5747061	Oct 25, 2013	DP	U-575	
<u>LOTEPREDNOL ETABONATE; LOTEMAX</u>					
020841 001	4996335	Feb 26, 2008			
	5540930	Oct 25, 2013			
<u>LOVASTATIN; ALTOPREV</u>					
021316 001	5916595	Dec 12, 2017			
	6080778	Mar 23, 2018		U-456	
	6485748	Dec 12, 2017	DP		
<u>LOVASTATIN; ALTOPREV</u>					
021316 002	5916595	Dec 12, 2017			
	6080778	Mar 23, 2018		U-456	
	6485748	Dec 12, 2017	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LOVASTATIN; ALTOPREV</u>					
021316 003	5916595	Dec 12, 2017			
	6080778	Mar 23, 2018		U-456	
	6485748	Dec 12, 2017	DP		
<u>LOVASTATIN; ALTOPREV</u>					
021316 004	5916595	Dec 12, 2017			
	6080778	Mar 23, 2018		U-456	
	6485748	Dec 12, 2017	DP		
<u>LOVASTATIN; MEVACOR</u>					
019643 002				PED	Aug 14, 2005
<u>LOVASTATIN; MEVACOR</u>					
019643 003				PED	Aug 14, 2005
<u>LOVASTATIN; MEVACOR</u>					
019643 004				PED	Aug 14, 2005
<u>LOVASTATIN; NIACIN; ADVICOR</u>					
021249 001	6080428	May 27, 2017		U-447	
	6129930	Sep 20, 2013		U-448	
	6406715	Oct 31, 2017		U-450	
	6676967	Sep 20, 2013		U-548	
	6746691	Sep 20, 2013		U-586	
	6818229	Feb 15, 2014	DP		
<u>LOVASTATIN; NIACIN; ADVICOR</u>					
021249 002	6080428	May 27, 2017		U-447	
	6129930	Sep 20, 2013		U-448	
	6406715	Oct 31, 2017		U-450	
	6676967	Sep 20, 2013		U-548	
	6746691	Sep 20, 2013		U-586	
	6818229	Feb 15, 2014	DP		
<u>LOVASTATIN; NIACIN; ADVICOR</u>					
021249 003	6080428	May 27, 2017		U-447	
	6129930	Sep 20, 2013		U-448	
	6406715	Oct 31, 2017		U-450	
	6676967	Sep 20, 2013		U-548	
	6746691	Sep 20, 2013		U-586	
<u>LUTROPIN ALFA; LUVERIS</u>					
021322 001	5650390	Jul 22, 2014	DP	ODE	Oct 08, 2011
	5767251	Jun 16, 2015	DS		
<u>MANGAFODIPIR TRISODIUM; TESLASCAN</u>					
020652 001	4933456	Nov 27, 2011			
	4992554	Feb 12, 2008			
	5091169	Feb 25, 2009			
	5223243	Jun 29, 2010		U-237	
<u>MASOPROCOL; ACTINEX</u>					
019940 001	4695590	Apr 17, 2008			
	5008294	Apr 15, 2008		U-68	
<u>MECASERMIN RECOMBINANT; INCRELEX</u>					
021839 001	5681814	Oct 28, 2014	DP	NCE	Aug 30, 2010
	5824642	Oct 20, 2015		ODE	Aug 30, 2012
	6207640	Apr 07, 2014		U-681	
<u>MECASERMIN RINFABATE RECOMBINANT; IPLEX</u>					
021884 001				NP	Dec 12, 2008
				ODE	Dec 12, 2012
<u>MEDROXYPROGESTERONE ACETATE; DEPO-SUBQ PROVERA 104</u>					
021583 001	6495534	May 15, 2020	DP	I-451	Mar 25, 2008
				NP	Dec 17, 2007
<u>MEGESTROL ACETATE; MEGACE</u>					
020264 001	5338732	Aug 16, 2011			
<u>MEGESTROL ACETATE; MEGACE ES</u>					
021778 001	5145684	Jan 25, 2011	DP	U-673	
	6592903	Sep 21, 2020	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MELOXICAM; MOBIC</u>					
020938 001				I-469	Aug 11, 2008
				I-430	Jul 16, 2007
				PED	Feb 11, 2009
				PED	Jan 16, 2008
				PED	Oct 13, 2005
<u>MELOXICAM; MOBIC</u>					
020938 002				I-469	Aug 11, 2008
				I-430	Jul 16, 2007
				PED	Feb 11, 2009
				PED	Jan 16, 2008
				PED	Oct 13, 2005
<u>MELOXICAM; MOBIC</u>					
021530 001	6184220	Mar 25, 2019	DP	I-469	Jun 01, 2007
	6184220*PED	Sep 25, 2019		I-430	Jul 16, 2007
				PED	Jan 16, 2008
				PED	Dec 01, 2007
				PED	Oct 13, 2005
<u>MELPHALAN HYDROCHLORIDE; ALKERAN</u>					
020207 001	4997651	Nov 18, 2008			
<u>MEMANTINE HYDROCHLORIDE; NAMENDA</u>					
021487 001	5061703	Apr 11, 2010	U-539	NCE	Oct 16, 2008
	5614560	Mar 25, 2014	U-539		
<u>MEMANTINE HYDROCHLORIDE; NAMENDA</u>					
021487 002	5061703	Apr 11, 2010	U-539	NCE	Oct 16, 2008
	5614560	Mar 25, 2014	U-539		
<u>MEMANTINE HYDROCHLORIDE; NAMENDA</u>					
021487 003	5061703	Apr 11, 2010	U-539	NCE	Oct 16, 2008
	5614560	Mar 25, 2014	U-539		
<u>MEMANTINE HYDROCHLORIDE; NAMENDA</u>					
021487 004	5061703	Apr 11, 2010	U-539	NCE	Oct 16, 2008
	5614560	Mar 25, 2014	U-539		
<u>MEMANTINE HYDROCHLORIDE; NAMENDA</u>					
021627 001				NCE	Oct 16, 2008
<u>MENOTROPINS (FSH; LH); MENOPUR</u>					
021663 001				NP	Oct 29, 2007
<u>MEQUINOL; TRETINOIN; SOLAGE</u>					
020922 001	5194247	Mar 16, 2010	U-294		
	5470567	Mar 19, 2010	U-294		
	6353029	Aug 24, 2020			
<u>MESALAMINE; ASACOL</u>					
019651 001	5541170	Jul 30, 2013	U-141		
	5541171	Jul 30, 2013	U-141		
<u>MESALAMINE; CANASA</u>					
021252 002				D-92	Nov 05, 2007
				NS	Nov 05, 2007
<u>MESNA; MESNEX</u>					
019884 001	5696172	Oct 06, 2013			
<u>MESNA; MESNEX</u>					
020855 001	5252341	Jul 16, 2011			
	5262169	Jul 16, 2011			
<u>METAXALONE; SKELAXIN</u>					
013217 001	6407128	Dec 03, 2021	U-189		
	6683102	Dec 03, 2021	U-189		
<u>METAXALONE; SKELAXIN</u>					
013217 003	6407128	Dec 03, 2021	U-189		
	6683102	Dec 03, 2021	U-189		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
METFORMIN HYDROCHLORIDE; FORTAMET									
021574 001	6099859	Mar	20, 2018	DP			NP	Apr	27, 2007
	6495162	Mar	20, 2018	DP					
	6790459	Mar	17, 2021		U-604				
	6866866	Mar	17, 2021	DP					
METFORMIN HYDROCHLORIDE; FORTAMET									
021574 002	6099859	Mar	20, 2018	DP			NP	Apr	27, 2007
	6495162	Mar	20, 2018	DP					
	6790459	Mar	17, 2021		U-604				
	6866866	Mar	17, 2021	DP					
METFORMIN HYDROCHLORIDE; GLUCOPHAGE XR									
021202 001	6475521	Mar	19, 2018				U-542		
	6660300	Mar	19, 2018						
METFORMIN HYDROCHLORIDE; GLUCOPHAGE XR									
021202 004	6475521	Mar	19, 2018				U-542		
	6660300	Mar	19, 2018						
METFORMIN HYDROCHLORIDE; GLUMETZA									
021748 001	6340475	Sep	19, 2016	DS	DP	U-669	NP	Jun	03, 2008
	6488962	Jun	20, 2020	DS	DP				
	6723340	Oct	25, 2021	DS	DP				
METFORMIN HYDROCHLORIDE; GLUMETZA									
021748 002	6340475	Sep	19, 2016	DS	DP	U-669	NP	Jun	03, 2008
	6488962	Jun	20, 2020	DS	DP				
	6723340	Oct	25, 2021	DS	DP				
METFORMIN HYDROCHLORIDE; RIOMET									
021591 001	6890957	Sep	14, 2023		DP				
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE; AVANDAMET									
021410 001	5002953	Sep	17, 2011	DS	DP	U-691			
	5002953	Sep	17, 2011	DS	DP	U-690			
	5002953	Sep	17, 2011	DS	DP	U-493			
	5002953*PED	Mar	17, 2012						
	5741803	Apr	21, 2015			U-493			
	5741803*PED	Oct	21, 2015						
	5965584	Jun	19, 2016			U-493			
	6166042	Jun	19, 2016			U-493			
	6288095	Feb	11, 2017			U-493			
	6288095*PED	Aug	11, 2017						
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE; AVANDAMET									
021410 002	5002953	Sep	17, 2011	DS	DP	U-691			
	5002953	Sep	17, 2011	DS	DP	U-690			
	5002953	Sep	17, 2011	DS	DP	U-493			
	5002953*PED	Mar	17, 2012						
	5741803	Apr	21, 2015			U-493			
	5741803*PED	Oct	21, 2015						
	5965584	Jun	19, 2016			U-493			
	6166042	Jun	19, 2016			U-493			
	6288095	Feb	11, 2017			U-493			
	6288095*PED	Aug	11, 2017						
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE; AVANDAMET									
021410 003	5002953	Sep	17, 2011	DS	DP	U-493			
	5002953	Sep	17, 2011	DS	DP	U-690			
	5002953	Sep	17, 2011	DS	DP	U-691			
	5002953*PED	Mar	17, 2012						
	5741803	Apr	21, 2015			U-493			
	5741803*PED	Oct	21, 2015						
	5965584	Jun	19, 2016			U-493			
	6166042	Jun	19, 2016			U-493			
	6288095	Feb	11, 2017			U-493			
	6288095*PED	Aug	11, 2017						
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE; AVANDAMET									
021410 004	5002953	Sep	17, 2011	DS	DP	U-690			
	5002953	Sep	17, 2011	DS	DP	U-691			
	5002953	Sep	17, 2011	DS	DP	U-493			
	5002953*PED	Mar	17, 2012						
	5741803	Apr	21, 2015	DS	DP	U-493			
	5741803*PED	Oct	21, 2015						

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE; AVANDAMET</u>					
021410 005	5002953	Sep 17, 2011	DS DP	U-690	
	5002953	Sep 17, 2011	DS DP	U-493	
	5002953	Sep 17, 2011	DS DP	U-691	
	5002953*PED	Mar 17, 2012			
	5741803	Apr 21, 2015	DS DP	U-493	
	5741803*PED	Apr 21, 2015			
<u>METFORMIN; PIOGLITAZONE HYDROCHLORIDE; ACTOPLUS MET</u>					
021842 001	4687777	Jan 17, 2011	DS		
	5965584	Jun 19, 2016		DP	U-679
	6166042	Jun 19, 2016			U-679
	6166043	Jun 19, 2016			U-679
	6172090	Jun 19, 2016			U-679
<u>METFORMIN; PIOGLITAZONE HYDROCHLORIDE; ACTOPLUS MET</u>					
021842 002	4687777	Jan 17, 2011	DS		
	5965584	Jun 19, 2016		DP	U-679
	6166042	Jun 19, 2016			U-679
	6166043	Jun 19, 2016			U-679
	6172090	Jun 19, 2016			U-679
<u>METHOXSALEN; UVADEX</u>					
020969 001	4845075	Apr 11, 2009			
	4999375	Mar 12, 2008			
	5036102	Jul 30, 2008			U-273
<u>METHYL AMINOLEVULINATE HYDROCHLORIDE; METVIXIA</u>					
021415 001				NE	Jul 27, 2007
<u>METHYLPHENIDATE HYDROCHLORIDE; CONCERTA</u>					
021121 001	6919373	Jul 31, 2017		U-666	D-94
	6919373*PED	Jan 31, 2018			NPP
	6930129	Jul 31, 2017		U-666	PED
	6930129*PED	Jan 31, 2018			
<u>METHYLPHENIDATE HYDROCHLORIDE; CONCERTA</u>					
021121 002	6919373	Jul 31, 2017		U-666	D-94
	6919373*PED	Jan 31, 2018			NPP
	6930129	Jul 31, 2017		U-666	PED
	6930129*PED	Jan 31, 2018			
<u>METHYLPHENIDATE HYDROCHLORIDE; CONCERTA</u>					
021121 003	6919373	Jul 31, 2017		U-666	D-94
	6919373*PED	Jan 31, 2018			NPP
	6930129	Jul 31, 2017		U-666	PED
	6930129*PED	Jan 31, 2018			
<u>METHYLPHENIDATE HYDROCHLORIDE; CONCERTA</u>					
021121 004	6919373	Jul 31, 2017		U-666	D-94
	6919373*PED	Jan 31, 2018			NPP
	6930129	Jul 31, 2017		U-666	PED
	6930129*PED	Jan 31, 2018			
<u>METHYLPHENIDATE HYDROCHLORIDE; METADATE CD</u>					
021259 001	6344215	Oct 27, 2020			
<u>METHYLPHENIDATE HYDROCHLORIDE; METADATE CD</u>					
021259 002	6344215	Oct 27, 2020			
<u>METHYLPHENIDATE HYDROCHLORIDE; METADATE CD</u>					
021259 003	6344215	Oct 27, 2020			
<u>METHYLPHENIDATE HYDROCHLORIDE; RITALIN LA</u>					
021284 001	5837284	Dec 04, 2015		DP	
	6228398	Nov 01, 2019		DP	U-472
	6635284	Dec 04, 2015		DP	U-591
<u>METHYLPHENIDATE HYDROCHLORIDE; RITALIN LA</u>					
021284 002	5837284	Dec 04, 2015		DP	
	6228398	Nov 01, 2019		DP	U-472
	6635284	Dec 04, 2015		DP	U-591
<u>METHYLPHENIDATE HYDROCHLORIDE; RITALIN LA</u>					
021284 003	5837284	Dec 04, 2015		DP	
	6228398	Nov 01, 2019		DP	U-472
	6635284	Dec 04, 2015		DP	U-591

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>METHYLPHENIDATE HYDROCHLORIDE; RITALIN LA</u>					
021284 004	5837284	Dec 04, 2015	DP		
	6228398	Nov 01, 2019	DP		
	6635284	Dec 04, 2015	DP	U-591	
<u>METOCLOPRAMIDE HYDROCHLORIDE; REGLAN ODT</u>					
021793 001	6024981	Apr 09, 2018	DP		
	6221392	Apr 09, 2018	DP		
<u>METOCLOPRAMIDE HYDROCHLORIDE; REGLAN ODT</u>					
021793 002	6024981	Apr 09, 2018	DP		
	6221392	Apr 09, 2018	DP		
<u>METOPROLOL FUMARATE; LOPRESSOR</u>					
019786 003	4892739	Jan 09, 2007			
<u>METOPROLOL FUMARATE; LOPRESSOR</u>					
019786 004	4892739	Jan 09, 2007			
<u>METOPROLOL SUCCINATE; TOPROL-XL</u>					
019962 001	4927640	May 22, 2007	DP		
	4957745	Sep 18, 2007	DP	U-107	
	5001161	Sep 18, 2007	DP		
	5081154	Sep 18, 2007	DS		
<u>METOPROLOL SUCCINATE; TOPROL-XL</u>					
019962 002	4927640	May 22, 2007	DP		
	4957745	Sep 18, 2007	DP	U-107	
	5001161	Sep 18, 2007	DP		
	5081154	Sep 18, 2007	DS		
<u>METOPROLOL SUCCINATE; TOPROL-XL</u>					
019962 003	4927640	May 22, 2007	DP		
	4957745	Sep 18, 2007	DP	U-107	
	5001161	Sep 18, 2007	DP		
	5081154	Sep 18, 2007	DS		
<u>METOPROLOL SUCCINATE; TOPROL-XL</u>					
019962 004	4927640	May 22, 2007	DP		
	4957745	Sep 18, 2007	DP	U-107	
	5001161	Sep 18, 2007	DP		
	5081154	Sep 18, 2007	DS		
<u>METOPROLOL SUCCINATE; TOPROL-XL</u>					
019962 004	4927640	May 22, 2007	DP		
	4957745	Sep 18, 2007	DP	U-107	
	5001161	Sep 18, 2007	DP	U-107	
	5081154	Sep 18, 2007	DS	U-107	
<u>METRONIDAZOLE; METROGEL</u>					
019737 001	4837378	Jun 06, 2006			
<u>METRONIDAZOLE; METROGEL</u>					
021789 001				NP	Jun 30, 2008
<u>METRONIDAZOLE; METROGEL-VAGINAL</u>					
020208 001	4837378	Jun 06, 2006			
	5536743	Jul 16, 2013		U-137	
	5840744	Jan 15, 2008		U-370	
<u>MICAFUNGIN SODIUM; MYCAMINE</u>					
021506 002	5376634	Dec 27, 2011	DS DP	NCE	Mar 16, 2010
	6107458	Sep 29, 2015	DS DP	U-650	
	6265536	Sep 29, 2015	DS DP	U-650	
	6774104	Jan 08, 2021	DP	U-650	
<u>MICONAZOLE NITRATE; MONISTAT 1 COMBINATION PACK</u>					
021308 001	5514698	Mar 21, 2014		D-90	Oct 01, 2007
	6153635	Nov 28, 2020			
<u>MIFEPRISTONE; MIFEPREX</u>					
020687 001				NCE	Sep 28, 2005
<u>MIGLITOL; GLYSET</u>					
020682 001	4639436	Jan 27, 2009		U-111	
<u>MIGLITOL; GLYSET</u>					
020682 002	4639436	Jan 27, 2009		U-111	
<u>MIGLITOL; GLYSET</u>					
020682 003	4639436	Jan 27, 2009		U-111	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MIGLUSTAT; ZAVESCA</u>					
021348 001	5472969	May 13, 2013		NCE	Jul 31, 2008
	5525616	Jun 11, 2013		ODE	Jul 31, 2010
<u>MIRTAZAPINE; MIRTAZAPINE</u>					
076689 003				PC	Feb 28, 2006
<u>MIRTAZAPINE; MIRTAZAPINE</u>					
076901 003				PC	Feb 28, 2006
<u>MIRTAZAPINE; REMERON SOLTAB</u>					
021208 001	5178878	Jan 12, 2010			
<u>MIRTAZAPINE; REMERON SOLTAB</u>					
021208 002	5178878	Jan 12, 2010			
<u>MIRTAZAPINE; REMERON SOLTAB</u>					
021208 003	5178878	Jan 12, 2010			
<u>MITOXANTRONE HYDROCHLORIDE; NOVANTRONE</u>					
019297 001	4820738	Apr 11, 2006	U-98	ODE	Oct 13, 2007
<u>MITOXANTRONE HYDROCHLORIDE; NOVANTRONE</u>					
019297 002	4820738	Apr 11, 2006		ODE	Oct 13, 2007
<u>MITOXANTRONE HYDROCHLORIDE; NOVANTRONE</u>					
019297 003	4820738	Apr 11, 2006		ODE	Oct 13, 2007
<u>MIVACURIUM CHLORIDE; MIVACRON</u>					
020098 001	4761418	Jan 22, 2006			
<u>MIVACURIUM CHLORIDE; MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
020098 002	4761418	Jan 22, 2006			
<u>MODAFINIL; PROVIGIL</u>					
020717 001	4927855	May 22, 2007	U-255	I-449	Jan 23, 2007
	RE37516	Oct 06, 2014	U-255	ODE	Dec 24, 2005
<u>MODAFINIL; PROVIGIL</u>					
020717 002	4927855	May 22, 2007	U-255	I-449	Jan 23, 2007
	RE37516	Oct 06, 2014	U-255	ODE	Dec 24, 2005
<u>MOEXIPRIL HYDROCHLORIDE; UNIVASC</u>					
020312 001	4743450	Feb 24, 2007			
<u>MOEXIPRIL HYDROCHLORIDE; UNIVASC</u>					
020312 002	4743450	Feb 24, 2007			
<u>MOMETASONE FUROATE; ASMANEX TWISTHALER</u>					
021067 001	5394868	Jun 25, 2012	DP	NP	Mar 30, 2008
	5687710	Nov 18, 2014	DP		
	5829434	Nov 03, 2015	DP		
	5889015	Jan 27, 2014		U-645	
	6057307	Jan 27, 2014	DP	U-645	
	6240918	Feb 20, 2017	DP		
	6365581	Jan 27, 2014		U-645	
	6503537	Mar 17, 2018	DP		
	6677322	Jan 27, 2014		U-645	
	6949532	Jan 27, 2014		U-645	
<u>MOMETASONE FUROATE; ELOCON</u>					
019625 001	4808610	Oct 02, 2006			
	4808610*PED	Apr 02, 2007			
<u>MOMETASONE FUROATE; ELOCON</u>					
019796 001	4775529	May 21, 2007			
	4775529*PED	Nov 21, 2007			
<u>MOMETASONE FUROATE; MOMETASONE FUROATE</u>					
077180 001				PC	Nov 19, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MOMETASONE FUROATE MONOHYDRATE; NASONEX</u>					
020762 001	5837699	Jan 27, 2014	DP	U-625	I-443
	5837699*PED	Jul 27, 2014			I-360
	6127353	Oct 03, 2017	DS DP		PED
	6127353*PED	Apr 03, 2018			
	6723713	Jan 27, 2014		U-625	
	6723713*PED	Jul 27, 2014			
<u>MONTELUKAST SODIUM; SINGULAIR</u>					
020829 002	5565473	Feb 03, 2012	DS DP	U-675	I-465
	5565473	Feb 03, 2012	DS DP	U-228	I-378
	5565473*PED	Aug 03, 2012		U-228	M-49
					Nov 30, 2008
<u>MONTELUKAST SODIUM; SINGULAIR</u>					
020830 001	5565473	Feb 03, 2012	DS DP	U-675	I-465
	5565473	Feb 03, 2012	DS DP	U-228	I-378
	5565473*PED	Aug 03, 2012		U-228	M-49
					Nov 30, 2008
<u>MONTELUKAST SODIUM; SINGULAIR</u>					
020830 002	5565473	Feb 03, 2012	DS DP	U-675	I-465
	5565473	Feb 03, 2012	DS DP	U-228	I-378
	5565473*PED	Aug 03, 2012		U-228	M-49
					Nov 30, 2008
<u>MONTELUKAST SODIUM; SINGULAIR</u>					
021409 001	5565473	Feb 03, 2012	DS DP	U-675	I-465
	5565473*PED	Aug 03, 2012			I-378
					M-49
					NDF
					PED
					Aug 27, 2008
					Dec 31, 2005
					Nov 30, 2008
					Jul 26, 2005
					Jan 26, 2006
<u>MORPHINE SULFATE; AVINZA</u>					
021260 001	6066339	Nov 25, 2017			
<u>MORPHINE SULFATE; AVINZA</u>					
021260 002	6066339	Nov 25, 2017			
<u>MORPHINE SULFATE; AVINZA</u>					
021260 003	6066339	Nov 25, 2017			
<u>MORPHINE SULFATE; AVINZA</u>					
021260 004	6066339	Nov 25, 2017			
<u>MORPHINE SULFATE; DEPODUR</u>					
021671 001	5723147	Mar 03, 2015	DP	U-584	
	5807572	Sep 15, 2015	DP		
	5891467	Jan 31, 2017	DP		
	5931089	Jul 14, 2015		U-584	
	5962016	Jan 31, 2017	DP	U-584	
	5997899	Sep 01, 2016	DP		
	6071534	Feb 18, 2008			
	6171613	Oct 01, 2016	DP		
	6193998	Sep 01, 2016	DP		
	6241999	Sep 01, 2016	DP		
<u>MORPHINE SULFATE; DEPODUR</u>					
021671 002	5723147	Mar 03, 2015	DP	U-584	
	5807572	Sep 15, 2015	DP		
	5891467	Jan 31, 2017	DP		
	5931089	Jul 14, 2015		U-584	
	5962016	Jan 31, 2017	DP	U-584	
	5997899	Sep 01, 2016	DP		
	6071534	Feb 18, 2008			
	6171613	Oct 01, 2016	DP		
	6193998	Sep 01, 2016	DP		
	6241999	Sep 01, 2016	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MORPHINE SULFATE; DEPODUR</u>					
021671 003	5723147	Mar 03, 2015	DP	U-584	
	5807572	Sep 15, 2015	DP		
	5891467	Jan 31, 2017	DP		
	5931089	Jul 14, 2015		U-584	
	5962016	Jan 31, 2017	DP	U-584	
	5997899	Sep 01, 2016	DP		
	6071534	Feb 18, 2008			
	6171613	Oct 01, 2016	DP		
	6193998	Sep 01, 2016	DP		
	6241999	Sep 01, 2016	DP		
<u>MORPHINE SULFATE; KADIAN</u>					
020616 001	5202128	Apr 13, 2010			
	5378474	Mar 23, 2010			
<u>MORPHINE SULFATE; KADIAN</u>					
020616 002	5202128	Apr 13, 2010			
	5378474	Mar 23, 2010			
<u>MORPHINE SULFATE; KADIAN</u>					
020616 003	5202128	Apr 13, 2010			
	5378474	Mar 23, 2010			
<u>MOXIFLOXACIN HYDROCHLORIDE; AVELOX</u>					
021085 001	4990517	Jun 30, 2009		U-298	I-479
	5607942	Mar 04, 2014		U-298	I-477
	5849752	Dec 05, 2016		U-298	I-385
<u>MOXIFLOXACIN HYDROCHLORIDE; AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER</u>					
021277 001	4990517	Jun 30, 2009		U-298	I-479
	5607942	Mar 04, 2014		U-298	I-477
	5849752	Dec 05, 2016		U-298	I-385
<u>MOXIFLOXACIN HYDROCHLORIDE; VIGAMOX</u>					
021598 001	4990517	Jun 30, 2009		NDF	Apr 15, 2006
	4990517*PED	Dec 30, 2009		PED	Oct 15, 2006
	5607942	Mar 04, 2014			
	5607942*PED	Sep 04, 2014			
	6716830	Sep 20, 2019	DP		
	6716830*PED	Mar 29, 2020			
<u>NAPROXEN SODIUM; NAPRELAN</u>					
020353 001	5637320	Jun 10, 2014			
<u>NAPROXEN SODIUM; NAPRELAN</u>					
020353 002	5637320	Jun 10, 2014			
<u>NAPROXEN SODIUM; NAPRELAN</u>					
020353 003	5637320	Jun 10, 2014			
<u>NARATRIPTAN HYDROCHLORIDE; AMERGE</u>					
020763 001	4997841	Jul 07, 2010		U-232	
<u>NARATRIPTAN HYDROCHLORIDE; AMERGE</u>					
020763 002	4997841	Jul 07, 2010		U-232	
<u>NATEGLINIDE; STARLIX</u>					
021204 001	5463116	Oct 21, 2012		I-411	Oct 20, 2006
	5488150	Jan 30, 2013		NCE	Dec 22, 2005
	6559188	Sep 17, 2019			
	6641841	Nov 14, 2017	DP	U-214	
	6844008	Nov 14, 2017	DP	U-214	
	6878749	Sep 15, 2020	DP		
	RE34878	Sep 08, 2009			
<u>NATEGLINIDE; STARLIX</u>					
021204 002	5463116	Oct 21, 2012		I-411	Oct 20, 2006
	5488150	Jan 30, 2013		NCE	Dec 22, 2005
	6559188	Sep 17, 2019			
	6641841	Nov 14, 2017	DP	U-214	
	6844008	Nov 14, 2017	DP	U-214	
	6878749	Sep 15, 2020	DP		
	RE34878	Sep 08, 2009			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NEDOCROMIL SODIUM; ALOCRI</u>					
021009 001	4474787	Oct 02, 2006	U-304		
	RE38628	Aug 22, 2012	U-304		
<u>NEDOCROMIL SODIUM; TILADE</u>					
019660 001	4474787	Oct 02, 2006			
	4760072	Jul 26, 2005			
<u>NELARABINE; ARRANON</u>					
021877 001	5424295	Jun 13, 2012	DS DP	NCE	Oct 28, 2010
	5492897	Feb 20, 2013		ODE	Oct 28, 2012
	5747472	Feb 20, 2013			
	5821236	Feb 20, 2013			
<u>NELFINAVIR MESYLATE; VIRACEPT</u>					
020778 001	5484926	Oct 07, 2013			
	5484926*PED	Apr 07, 2014			
	5952343	Oct 07, 2013	U-257		
	5952343*PED	Apr 07, 2014	U-257		
	6162812	Oct 07, 2013	U-248		
	6162812*PED	Apr 07, 2014	U-248		
<u>NELFINAVIR MESYLATE; VIRACEPT</u>					
020779 001	5484926	Oct 07, 2013			
	5484926*PED	Apr 07, 2014			
	5952343	Oct 07, 2013	U-257		
	5952343*PED	Apr 07, 2014	U-257		
	6162812	Oct 07, 2013	U-248		
	6162812*PED	Apr 07, 2014	U-248		
<u>NELFINAVIR MESYLATE; VIRACEPT</u>					
021503 001	5484926	Oct 07, 2013			
	5484926*PED	Apr 07, 2014			
	5952343	Oct 07, 2013	U-257		
	5952343*PED	Apr 07, 2014	U-257		
	6162812	Oct 07, 2013	U-248		
	6162812*PED	Apr 07, 2014	U-248		
<u>NEPAFENAC; NEVANAC</u>					
021862 001	5475034	Jun 06, 2014	U-100	NCE	Aug 19, 2010
<u>NESIRITIDE RECOMBINANT; NATRECOR</u>					
020920 001	5114923	May 19, 2009		NCE	Aug 10, 2006
<u>NEVIRAPINE; VIRAMUNE</u>					
020636 001	5366972	Nov 22, 2011	U-167		
	5366972*PED	May 22, 2012			
<u>NEVIRAPINE; VIRAMUNE</u>					
020933 001	5366972	Nov 22, 2011			
	5366972*PED	May 22, 2012			
<u>NIACIN; NIASPAN</u>					
020381 001	6080428	May 27, 2017	U-331		
	6129930	Sep 20, 2013	U-354		
	6406715	Oct 31, 2017	U-450		
	6746691	Sep 20, 2013	U-586		
	6818229	Feb 15, 2014	DP		
<u>NIACIN; NIASPAN</u>					
020381 002	6080428	May 27, 2017	U-331		
	6129930	Sep 20, 2013	U-354		
	6406715	Oct 31, 2017	U-450		
	6676967	Sep 20, 2013	U-548		
	6746691	Sep 20, 2013	U-586		
	6818229	Feb 15, 2014	DP		
<u>NIACIN; NIASPAN</u>					
020381 003	6080428	May 27, 2017	U-331		
	6129930	Sep 20, 2013	U-354		
	6406715	Oct 31, 2017	U-450		
	6676967	Sep 20, 2013	U-548		
	6746691	Sep 20, 2013	U-586		
	6818229	Feb 15, 2014	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
NIACIN; NIASPAN					
020381 004	6080428	May 27, 2017	DP	U-331	
	6129930	Sep 20, 2013		U-354	
	6406715	Oct 31, 2017		U-450	
	6676967	Sep 20, 2013		U-548	
	6746691	Sep 20, 2013		U-586	
	6818229	Feb 15, 2014			
NIACIN; NIASPAN TITRATION STARTER PACK					
020381 005	6080428	May 27, 2017		U-331	
	6129930	Sep 20, 2013		U-354	
	6406715	Oct 31, 2017		U-450	
	6746691	Sep 20, 2013		U-586	
NICARDIPINE HYDROCHLORIDE; CARDENE					
019734 001	4880823	Nov 14, 2006			
	5164405	Nov 17, 2009			
NICARDIPINE HYDROCHLORIDE; CARDENE SR					
020005 001	5198226	Mar 30, 2010			
NICARDIPINE HYDROCHLORIDE; CARDENE SR					
020005 002	5198226	Mar 30, 2010			
NICARDIPINE HYDROCHLORIDE; CARDENE SR					
020005 003	5198226	Mar 30, 2010			
NICOTINE; HABITROL					
020076 004	5016652	May 21, 2008			
	5834011	May 01, 2007		U-355	
NICOTINE; HABITROL					
020076 005	5016652	May 21, 2008			
	5834011	May 01, 2007		U-355	
NICOTINE; HABITROL					
020076 006	5016652	May 21, 2008			
	5834011	May 01, 2007		U-355	
NICOTINE; NICODERM CQ					
020165 004	5004610	Jun 14, 2008			
	5342623	Jun 14, 2008			
	5344656	Jun 14, 2008			
	5364630	Jun 14, 2008			
	5462745	Jun 14, 2008			
	5508038	Apr 16, 2013			
	5633008	Jun 14, 2008		U-389	
	6165497	Jun 14, 2008		U-388	
NICOTINE; NICODERM CQ					
020165 005	5004610	Jun 14, 2008			
	5342623	Jun 14, 2008			
	5344656	Jun 14, 2008			
	5364630	Jun 14, 2008			
	5462745	Jun 14, 2008			
	5508038	Apr 16, 2013			
	5633008	Jun 14, 2008		U-389	
	6165497	Jun 14, 2008		U-388	
NICOTINE; NICODERM CQ					
020165 006	5004610	Jun 14, 2008			
	5342623	Jun 14, 2008			
	5344656	Jun 14, 2008			
	5364630	Jun 14, 2008			
	5462745	Jun 14, 2008			
	5508038	Apr 16, 2013			
	5633008	Jun 14, 2008		U-389	
	6165497	Jun 14, 2008		U-388	
NICOTINE; NICOTROL					
020385 001	5656255	Aug 12, 2014			
NICOTINE; NICOTROL					
020536 001	5501236	Jun 08, 2010			
	6098632	Jun 08, 2010			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NICOTINE; NICOTROL</u>					
020714 001	4793366	Dec 27, 2005			
	4800903	Jan 31, 2006			
	5167242	Jun 08, 2010			
	5400808	Jun 08, 2010			
	5501236	Jun 08, 2010			
	6098632	Jun 08, 2010			
<u>NICOTINE; PROSTEP</u>					
019983 003	4946853	Aug 07, 2007		U-56	
<u>NICOTINE; PROSTEP</u>					
019983 004	4946853	Aug 07, 2007		U-56	
<u>NICOTINE POLACRILEX; COMMIT</u>					
021330 001	5110605	Aug 21, 2010		NDF	Oct 31, 2005
<u>NICOTINE POLACRILEX; COMMIT</u>					
021330 002	5110605	Aug 21, 2010		NDF	Oct 31, 2005
<u>NIFEDIPINE; ADALAT CC</u>					
020198 001	4892741	Jun 08, 2008			
	5264446	Nov 23, 2010			
<u>NIFEDIPINE; ADALAT CC</u>					
020198 002	4892741	Jun 08, 2008			
	5264446	Nov 23, 2010			
<u>NIFEDIPINE; ADALAT CC</u>					
020198 003	4892741	Jun 08, 2008			
	5264446	Nov 23, 2010			
<u>NIFEDIPINE; PROCARDIA XL</u>					
019684 001	5264446	Nov 23, 2010			
<u>NIFEDIPINE; PROCARDIA XL</u>					
019684 002	5264446	Nov 23, 2010			
<u>NIFEDIPINE; PROCARDIA XL</u>					
019684 003	5264446	Nov 23, 2010			
<u>NISOLDIPINE; SULAR</u>					
020356 001	4703038	Oct 07, 2005		U-3	
	4892741	Jun 08, 2008			
<u>NISOLDIPINE; SULAR</u>					
020356 002	4703038	Oct 07, 2005		U-3	
	4892741	Jun 08, 2008			
<u>NISOLDIPINE; SULAR</u>					
020356 003	4703038	Oct 07, 2005		U-3	
	4892741	Jun 08, 2008			
<u>NISOLDIPINE; SULAR</u>					
020356 004	4703038	Oct 07, 2005		U-3	
	4892741	Jun 08, 2008			
<u>NITAZOXANIDE; ALINIA</u>					
021497 001	5387598	Feb 07, 2012	DP	U-524	I-460 Jun 16, 2008
	5578621	Nov 26, 2013	DP	U-525	NDF Jul 21, 2007
	5968961	May 07, 2017	DP		NCE Nov 22, 2007
	6020353	Sep 18, 2014	DS DP		ODE Nov 22, 2009
<u>NITAZOXANIDE; ALINIA</u>					
021498 001	5387598	Feb 07, 2012		U-524	I-460 Jun 16, 2008
	5578621	Sep 08, 2014		U-525	NPP Jul 21, 2007
	5965590	Jul 03, 2017		U-523	NCE Nov 22, 2007
	5968961	May 07, 2017			ODE Nov 22, 2009
	6020353	Sep 08, 2014			ODE Nov 22, 2009
	6117894	May 07, 2017			ODE Nov 22, 2009
<u>NITISINONE; ORFADIN</u>					
021232 001	5006158	Apr 09, 2008		NCE	Jan 18, 2007
	5550165	Aug 27, 2013		ODE	Jan 18, 2009

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>NITISINONE; ORFADIN</u>					
021232 002	5006158	Apr 09, 2008		NCE	Jan 18, 2007
	5550165	Aug 27, 2013		ODE	Jan 18, 2009
<u>NITISINONE; ORFADIN</u>					
021232 003	5006158	Apr 09, 2008		NCE	Jan 18, 2007
	5550165	Aug 27, 2013		ODE	Jan 18, 2009
<u>NITRIC OXIDE; INOMAX</u>					
020845 002	5485827	Jan 23, 2013	U-297	ODE	Dec 23, 2006
	5873359	Jan 23, 2013	U-297		
<u>NITRIC OXIDE; INOMAX</u>					
020845 003	5485827	Jan 23, 2013	U-297	ODE	Dec 23, 2006
	5873359	Jan 23, 2013	U-297		
<u>NITROFURANTOIN; NITROFURANTOIN, MACROCRYSTALLINE; MACROBID</u>					
020064 001	4772473	Jun 16, 2006			
	4798725	Jun 16, 2006			
<u>NITROGLYCERIN; NITRO-DUR</u>					
020145 001	5186938	Feb 16, 2010			
<u>NITROGLYCERIN; NITRO-DUR</u>					
020145 002	5186938	Feb 16, 2010			
<u>NITROGLYCERIN; NITRO-DUR</u>					
020145 003	5186938	Feb 16, 2010			
<u>NITROGLYCERIN; NITRO-DUR</u>					
020145 004	5186938	Feb 16, 2010			
<u>NITROGLYCERIN; NITRO-DUR</u>					
020145 005	5186938	Feb 16, 2010			
<u>NITROGLYCERIN; NITRO-DUR</u>					
020145 006	5186938	Feb 16, 2010			
<u>NITROGLYCERIN; NITROLINGUAL PUMPSPRAY</u>					
018705 002	5186925	Feb 16, 2010			
<u>NITROGLYCERIN; NITROSTAT</u>					
021134 001	6500456	Sep 16, 2018			
<u>NITROGLYCERIN; NITROSTAT</u>					
021134 002	6500456	Sep 16, 2018			
<u>NITROGLYCERIN; NITROSTAT</u>					
021134 003	6500456	Sep 16, 2018			
<u>NIZATIDINE; AXID</u>					
021494 001	6930119	Jul 17, 2022	DP	NDF	May 25, 2007
<u>OCTREOTIDE ACETATE; OCTREOTIDE ACETATE</u>					
076330 001				PC	Nov 20, 2005
<u>OCTREOTIDE ACETATE; OCTREOTIDE ACETATE</u>					
076330 002				PC	Nov 20, 2005
<u>OCTREOTIDE ACETATE; OCTREOTIDE ACETATE (PRESERVATIVE FREE)</u>					
076313 001				PC	Oct 02, 2005
<u>OCTREOTIDE ACETATE; OCTREOTIDE ACETATE (PRESERVATIVE FREE)</u>					
076313 002				PC	Oct 02, 2005
<u>OCTREOTIDE ACETATE; OCTREOTIDE ACETATE (PRESERVATIVE FREE)</u>					
076313 003				PC	Oct 02, 2005
<u>OCTREOTIDE ACETATE; SANDOSTATIN</u>					
019667 001	5753618	May 19, 2015			
<u>OCTREOTIDE ACETATE; SANDOSTATIN</u>					
019667 002	5753618	May 19, 2015			
<u>OCTREOTIDE ACETATE; SANDOSTATIN</u>					
019667 003	5753618	May 19, 2015			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OCTREOTIDE ACETATE; SANDOSTATIN</u>					
019667 004	5753618	May 19, 2015			
<u>OCTREOTIDE ACETATE; SANDOSTATIN</u>					
019667 005	5753618	May 19, 2015			
<u>OCTREOTIDE ACETATE; SANDOSTATIN LAR</u>					
021008 001	5538739	Jul 23, 2013		ODE	Nov 25, 2005
	5639480	Jun 17, 2014			
	5688530	Nov 18, 2014	U-268		
	5922338	Jul 13, 2016			
	5922682	Jul 13, 2016			
<u>OCTREOTIDE ACETATE; SANDOSTATIN LAR</u>					
021008 002	5538739	Jul 23, 2013		ODE	Nov 25, 2005
	5639480	Jun 17, 2014			
	5688530	Nov 18, 2014	U-268		
	5922338	Jul 13, 2016			
	5922682	Jul 13, 2016			
<u>OCTREOTIDE ACETATE; SANDOSTATIN LAR</u>					
021008 003	5538739	Jul 23, 2013		ODE	Nov 25, 2005
	5639480	Jun 17, 2014			
	5688530	Nov 18, 2014	U-268		
	5922338	Jul 13, 2016			
	5922682	Jul 13, 2016			
<u>OFLOXACIN; FLOXIN OTIC</u>					
020799 001	5401741	Mar 27, 2012	U-407	D-84	Sep 30, 2006
<u>OLANZAPINE; ZYPREXA</u>					
020592 001	5229382	Apr 23, 2011	DS DP U-547	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS DP U-149	I-400	Jul 10, 2006
<u>OLANZAPINE; ZYPREXA</u>					
020592 002	5229382	Apr 23, 2011	DS DP U-149	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS DP U-547	I-400	Jul 10, 2006
<u>OLANZAPINE; ZYPREXA</u>					
020592 003	5229382	Apr 23, 2011	DS DP U-149	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS DP U-547	I-400	Jul 10, 2006
<u>OLANZAPINE; ZYPREXA</u>					
020592 004	5229382	Apr 23, 2011	DS DP U-149	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS DP U-547	I-400	Jul 10, 2006
<u>OLANZAPINE; ZYPREXA</u>					
020592 005	5229382	Apr 23, 2011	DS DP U-547	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS DP U-149	I-400	Jul 10, 2006
<u>OLANZAPINE; ZYPREXA</u>					
020592 006	5229382	Apr 23, 2011	DS DP U-149	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS DP U-547	I-400	Jul 10, 2006
<u>OLANZAPINE; ZYPREXA</u>					
021253 001	5229382	Apr 23, 2011	DS DP U-571	NP NDF	Mar 29, 2007 Mar 29, 2007
<u>OLANZAPINE; ZYPREXA ZYDIS</u>					
021086 001	5229382	Apr 23, 2011	U-324	I-417 I-400	Jan 14, 2007 Jul 10, 2006
<u>OLANZAPINE; ZYPREXA ZYDIS</u>					
021086 002	5229382	Apr 23, 2011	U-324	I-417 I-400	Jan 14, 2007 Jul 10, 2006
<u>OLANZAPINE; ZYPREXA ZYDIS</u>					
021086 003	5229382	Apr 23, 2011	U-324	I-417 I-400	Jan 14, 2007 Jul 10, 2006
<u>OLANZAPINE; ZYPREXA ZYDIS</u>					
021086 004	5229382	Apr 23, 2011	U-324	I-417 I-400	Jan 14, 2007 Jul 10, 2006

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLMESARTAN MEDOXOMIL; BENICAR</u>					
021286 001	5616599 6878703	Apr 25, 2016 Nov 19, 2021	DS DP U-500 U-3	NCE	Apr 25, 2007
<u>OLMESARTAN MEDOXOMIL; BENICAR</u>					
021286 003	5616599 6878703	Apr 25, 2016 Nov 19, 2021	DS DP U-500 U-3	NCE	Apr 25, 2007
<u>OLMESARTAN MEDOXOMIL; BENICAR</u>					
021286 004	5616599 6878703	Apr 25, 2016 Nov 19, 2021	DS DP U-500 U-3	NCE	Apr 25, 2007
<u>OLOPATADINE HYDROCHLORIDE; OLOPATADINE HYDROCHLORIDE</u>					
021545 001				NP	Dec 22, 2007
<u>OLOPATADINE HYDROCHLORIDE; PATANOL</u>					
020688 001	4871865 4923892 5116863 5641805	Oct 03, 2006 May 08, 2007 Dec 18, 2010 Jun 06, 2015		U-174 U-184	
<u>OMEGA-3-ACID ETHYL ESTERS; OMACOR</u>					
021654 001	5502077 5656667 5698594	Mar 26, 2013 Aug 12, 2014 Dec 16, 2009	DS DS DS	NCE	Nov 10, 2009
<u>OMEPRAZOLE; PRILOSEC</u>					
019810 001	4786505 4786505*PED 4853230 4853230*PED 6147103 6147103*PED 6150380 6150380*PED 6166213 6166213*PED 6191148 6191148*PED	Apr 20, 2007 Oct 20, 2007 Apr 20, 2007 Oct 20, 2007 Oct 09, 2018 Apr 09, 2019 Nov 10, 2018 May 10, 2019 Oct 09, 2018 Apr 09, 2019 Oct 09, 2018 Apr 09, 2019	U-108 U-108 U-108 U-108	M-19 PED	Jul 12, 2005 Jan 12, 2006
<u>OMEPRAZOLE; PRILOSEC</u>					
019810 002	4786505 4786505*PED 4853230 4853230*PED 6147103 6147103*PED 6150380 6150380*PED 6166213 6166213*PED 6191148 6191148*PED	Apr 20, 2007 Oct 20, 2007 Apr 20, 2007 Oct 20, 2007 Oct 09, 2018 Apr 09, 2019 Nov 10, 2018 May 10, 2019 Oct 09, 2018 Apr 09, 2019 Oct 09, 2018 Apr 09, 2019	U-108 U-108 U-108 U-108	M-19 PED	Jul 12, 2005 Jan 12, 2006
<u>OMEPRAZOLE; PRILOSEC</u>					
019810 003	4786505 4786505*PED 4853230 4853230*PED 6147103 6147103*PED 6150380 6150380*PED 6166213 6166213*PED 6191148 6191148*PED	Apr 20, 2007 Oct 20, 2007 Apr 20, 2007 Oct 20, 2007 Oct 09, 2018 Apr 09, 2019 Nov 10, 2018 May 10, 2019 Oct 09, 2018 Apr 09, 2019 Oct 09, 2018 Apr 09, 2019	U-108 U-108 U-108 U-108	M-19 PED	Jul 12, 2005 Jan 12, 2006
<u>OMEPRAZOLE; ZEGERID</u>					
021636 001	5840737 6489346 6645988 6699885 6780882	Jul 16, 2016 Jul 16, 2016 Jul 16, 2016 Jul 16, 2016 Jul 16, 2016		U-588 U-588 U-588	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
OMEPRAZOLE; ZEGERID					
021706 001	5840737	Jul 16, 2016	DS	U-624	NP Dec 21, 2007
	5840737	Jul 16, 2016	DS	U-623	
	6489346	Jul 16, 2016	DS	U-623	
	6489346	Jul 16, 2016	DS DP	U-624	
	6645988	Jul 16, 2016	DS DP		
	6699885	Jul 16, 2016		U-624	
	6699885	Jul 16, 2016		U-623	
	6780882	Jul 16, 2016	DS DP		
OMEPRAZOLE MAGNESIUM; PRILOSEC OTC					
021229 001	4786505	Apr 20, 2007		RT0	Jun 20, 2006
	4786505*PED	Oct 20, 2007			
	4853230	Apr 20, 2007			
	4853230*PED	Oct 20, 2007			
	5690960	Nov 25, 2014			
	5753265	Jun 07, 2015			
	5817338	Oct 06, 2015			
	5900424	May 04, 2016			
	6403616	Nov 15, 2019			
	6428810	Nov 03, 2019			
ONDANSETRON; ZOFTRAN ODT					
020781 001	4695578*PED	Jul 25, 2005		U-330	
	4753789	Jun 24, 2006			
	4753789*PED	Dec 24, 2006			
	5578628*PED	Aug 16, 2005			
	5955488	Nov 14, 2015			
	5955488*PED	May 14, 2016			
	6063802	Nov 14, 2015			
	6063802*PED	May 14, 2016			
ONDANSETRON; ZOFTRAN ODT					
020781 002	4695578*PED	Jul 25, 2005		U-330	
	4753789	Jun 24, 2006			
	4753789*PED	Dec 24, 2006			
	5578628*PED	Aug 16, 2005			
	5955488	Nov 14, 2015			
	5955488*PED	May 14, 2016			
	6063802	Nov 14, 2015			
	6063802*PED	May 14, 2016			
ONDANSETRON HYDROCHLORIDE; ZOFTRAN					
020007 001	4695578*PED	Jul 25, 2005		U-44	D-98 Mar 25, 2008
	4753789	Jun 24, 2006			D-97 Mar 25, 2008
	4753789*PED	Dec 24, 2006			PED Sep 25, 2008
	5578628*PED	Aug 16, 2005			PED Sep 25, 2008
ONDANSETRON HYDROCHLORIDE; ZOFTRAN					
020103 001	4695578*PED	Jul 25, 2005		U-44	
	4753789	Jun 24, 2006			
	4753789*PED	Dec 24, 2006			
	5344658	Sep 06, 2011			
	5344658*PED	Mar 06, 2012			
	5578628*PED	Aug 16, 2005			
ONDANSETRON HYDROCHLORIDE; ZOFTRAN					
020103 002	4695578*PED	Jul 25, 2005		U-44	
	4753789	Jun 24, 2006			
	4753789*PED	Dec 24, 2006			
	5344658	Sep 06, 2011			
	5344658*PED	Mar 06, 2012			
	5578628*PED	Aug 16, 2005			
ONDANSETRON HYDROCHLORIDE; ZOFTRAN					
020103 003	4695578*PED	Jul 25, 2005		U-44	
	4753789	Jun 24, 2006			
	4753789*PED	Dec 24, 2006			
	5344658	Sep 06, 2011			
	5344658*PED	Mar 06, 2012			
	5578628*PED	Aug 16, 2005			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE		
ONDANSETRON HYDROCHLORIDE; ZOFRAN							
020605 001	4695578*PED	Jul 25, 2005	U-44	DP	U-44		
	4753789	Jun 24, 2006					
	4753789*PED	Dec 24, 2006					
	5578628*PED	Aug 16, 2005					
	5854270	Nov 20, 2015					
	5854270*PED	May 20, 2016					
ONDANSETRON HYDROCHLORIDE; ZOFRAN IN PLASTIC CONTAINER							
020403 001	4695578*PED	Jul 25, 2005	U-44				
	4753789	Jun 24, 2006					
	4753789*PED	Dec 24, 2006					
	5578628*PED	Aug 16, 2005					
ONDANSETRON HYDROCHLORIDE; ZOFRAN PRESERVATIVE FREE							
020007 003	4695578*PED	Jul 25, 2005	U-44	D-98	Mar 25, 2008		
	4753789	Jun 24, 2006		D-97	Mar 25, 2008		
	4753789*PED	Dec 24, 2006		PED	Sep 25, 2008		
	5578628*PED	Aug 16, 2005		PED	Sep 25, 2008		
ORLISTAT; XENICAL							
020766 001	4598089	Jun 18, 2009		M-29	Dec 12, 2006		
	4598089*PED	Dec 18, 2009		PED	Jun 12, 2007		
	6004996	Jan 06, 2018					
	6004996*PED	Jul 06, 2018					
OSELTAMIVIR PHOSPHATE; TAMIFLU							
021087 001	5763483	Dec 27, 2016		I-480	Dec 21, 2008		
	5763483*PED	Jun 27, 2017		PED	Jun 21, 2009		
	5866601	Feb 02, 2016					
	5866601*PED	Aug 02, 2016					
	5952375	Feb 02, 2016					
	5952375*PED	Aug 02, 2016					
OSELTAMIVIR PHOSPHATE; TAMIFLU							
021246 001	5763483	Dec 27, 2016	U-376	I-480	Dec 21, 2008		
	5763483*PED	Jun 27, 2017		PED	Jun 21, 2009		
	5866601	Feb 02, 2016					
	5866601*PED	Aug 02, 2016					
	5952375	Feb 02, 2016					
	5952375*PED	Aug 02, 2016					
OXALIPLATIN; ELOXATIN							
021492 001	5290961	Jan 12, 2013		I-441	Nov 04, 2007		
	5338874	Apr 07, 2013		I-425	Jan 09, 2007		
	5420319	Apr 07, 2013		NCE	Aug 09, 2007		
OXALIPLATIN; ELOXATIN							
021492 002	5290961	Jan 12, 2013		I-441	Nov 04, 2007		
	5338874	Apr 07, 2013		I-425	Jan 09, 2007		
	5420319	Apr 07, 2013		NCE	Aug 09, 2007		
OXALIPLATIN; ELOXATIN							
021759 001	5338874	Apr 07, 2013	DS	I-441	Nov 04, 2007		
	5420319	Apr 07, 2013	DS			NCE	Aug 09, 2007
	5716988	Aug 07, 2015	DP				
OXALIPLATIN; ELOXATIN							
021759 002	5338874	Apr 07, 2013	DS	I-441	Nov 04, 2007		
	5420319	Apr 07, 2013	DS			NCE	Aug 09, 2007
	5716988	Aug 07, 2015	DP				
OXANDROLONE; OXANDRIN							
013718 001	5872147	Dec 05, 2017	U-585	M-42	Jun 20, 2008		
	6090799	Jul 18, 2017				U-585	
	6576659	Dec 05, 2017				U-585	
	6670351	Oct 20, 2012				U-585	
	6828313	Dec 05, 2017				U-585	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OXANDROLONE; OXANDRIN</u>					
013718 002	5872147	Dec 05, 2017	U-585	M-42	Jun 20, 2008
	6090799	Jul 18, 2017	U-585		
	6576659	Dec 05, 2017	U-585		
	6670351	Oct 20, 2012	U-585		
	6828313	Dec 05, 2017	U-585		
<u>OXAPROZIN POTASSIUM; DAYPRO ALTA</u>					
020776 001	6030643	May 16, 2017	U-497	NE	Oct 17, 2005
<u>OXCARBAZEPINE; TRILEPTAL</u>					
021014 001				I-478	Oct 28, 2008
				PED	Apr 28, 2009
				PED	Jul 14, 2005
<u>OXCARBAZEPINE; TRILEPTAL</u>					
021014 002				I-478	Oct 28, 2008
				PED	Apr 28, 2009
				PED	Jul 14, 2005
<u>OXCARBAZEPINE; TRILEPTAL</u>					
021014 003				I-478	Oct 28, 2008
				PED	Apr 28, 2009
				PED	Jul 14, 2005
<u>OXCARBAZEPINE; TRILEPTAL</u>					
021285 001				I-478	Oct 28, 2008
				PED	Apr 28, 2009
				PED	Jul 14, 2005
<u>OXYBUTYNIN; OXYTROL</u>					
021351 002	5164190	Dec 11, 2010		NDF	Feb 26, 2006
	5601839	Apr 26, 2015			
	5834010	Apr 26, 2015			
	6743441	Apr 26, 2020	DP U-318		
<u>OXYBUTYNIN CHLORIDE; DITROPAN</u>					
017577 001				M-28	Apr 15, 2006
				PED	Oct 15, 2006
<u>OXYBUTYNIN CHLORIDE; DITROPAN</u>					
018211 001				M-28	Apr 15, 2006
				PED	Oct 15, 2006
<u>OXYBUTYNIN CHLORIDE; DITROPAN XL</u>					
020897 001	5674895	May 22, 2015		NPP	Apr 15, 2006
	5674895*PED	Nov 22, 2015		PED	Oct 15, 2006
	5840754	May 22, 2015			
	5840754*PED	Nov 22, 2015			
	5912268	May 22, 2015			
	5912268*PED	Nov 22, 2015			
	6124355	May 22, 2015		U-378	
	6124355*PED	Nov 22, 2015		U-378	
	6262115	May 22, 2015		U-393	
	6262115*PED	Nov 22, 2015		U-393	
	6919092	May 22, 2015	DP	U-667	
	6919092*PED	Nov 22, 2015			
<u>OXYBUTYNIN CHLORIDE; DITROPAN XL</u>					
020897 002	5674895	May 22, 2015		NPP	Apr 15, 2006
	5674895*PED	Nov 22, 2015		PED	Oct 15, 2006
	5840754	May 22, 2015			
	5840754*PED	Nov 22, 2015			
	5912268	May 22, 2015			
	5912268*PED	Nov 22, 2015			
	6124355	May 22, 2015		U-378	
	6124355*PED	Nov 22, 2015		U-378	
	6262115	May 22, 2015		U-393	
	6262115*PED	Nov 22, 2015		U-393	
	6919092	May 22, 2015	DP	U-667	
	6919092*PED	Nov 22, 2015			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OXYBUTYNIN CHLORIDE; DITROPAN XL</u>					
020897 003	5674895	May 22, 2015		NPP	Apr 15, 2006
	5674895*PED	Nov 22, 2015		PED	Oct 15, 2006
	5840754	May 22, 2015			
	5840754*PED	Nov 22, 2015			
	5912268	May 22, 2015			
	5912268*PED	Nov 22, 2015			
	6124355	May 22, 2015	U-378		
	6124355*PED	Nov 22, 2015	U-378		
	6262115	May 22, 2015	U-393		
	6262115*PED	Nov 22, 2015	U-393		
	6919092	May 22, 2015	DP U-667		
	6919092*PED	Nov 22, 2015			
<u>OXYCODONE HYDROCHLORIDE; OXYCODONE HCL</u>					
075923 001				PC	Dec 04, 2005
<u>OXYCODONE HYDROCHLORIDE; OXYCODONE HCL</u>					
075923 002				PC	Dec 04, 2005
<u>OXYCODONE HYDROCHLORIDE; OXYCODONE HCL</u>					
075923 003				PC	Dec 04, 2005
<u>OXYCODONE HYDROCHLORIDE; OXYCODONE HCL</u>					
075923 004				PC	Dec 04, 2005
<u>OXYCODONE HYDROCHLORIDE; OXYCONTIN</u>					
020553 001	4861598	Aug 29, 2006			
	4970075	Aug 29, 2006			
	5266331	Oct 26, 2007			
	5508042	Apr 16, 2013	U-443		
	5549912	Oct 26, 2007			
	5656295	Oct 26, 2007	U-443		
<u>OXYCODONE HYDROCHLORIDE; OXYCONTIN</u>					
020553 002	4861598	Aug 29, 2006			
	4970075	Aug 29, 2006			
	5266331	Oct 26, 2007			
	5508042	Apr 16, 2013	U-443		
	5549912	Oct 26, 2007			
	5656295	Oct 26, 2007	U-443		
<u>OXYCODONE HYDROCHLORIDE; OXYCONTIN</u>					
020553 003	4861598	Aug 29, 2006			
	4970075	Aug 29, 2006			
	5266331	Oct 26, 2007			
	5508042	Apr 16, 2013	U-443		
	5549912	Oct 26, 2007			
	5656295	Oct 26, 2007	U-443		
<u>OXYCODONE HYDROCHLORIDE; OXYCONTIN</u>					
020553 004	4861598	Aug 29, 2006			
	4970075	Aug 29, 2006			
	5266331	Oct 26, 2007			
	5508042	Apr 16, 2013	U-443		
	5549912	Oct 26, 2007			
	5656295	Oct 26, 2007	U-443		
<u>OXYCODONE HYDROCHLORIDE; OXYCONTIN</u>					
020553 005	4861598	Aug 29, 2006			
	4970075	Aug 29, 2006			
	5266331	Oct 26, 2007			
	5508042	Apr 16, 2013	U-443		
	5549912	Oct 26, 2007			
	5656295	Oct 26, 2007	U-443		
<u>PACLITAXEL; ABRAXANE</u>					
021660 001	5439686	Feb 22, 2013	DP	NP	Jan 07, 2008
	5498421	Mar 12, 2013	DP U-634		
	6096331	Feb 22, 2013	DP U-633		
	6506405	Feb 22, 2013	DP U-633		
	6537579	Feb 22, 2013	U-632		
	6749868	Feb 22, 2013	DP		
	6753006	Feb 22, 2013	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PALONOSETRON HYDROCHLORIDE; ALOXI</u>					
021372 001	5202333	Apr 13, 2010	U-528	NCE	Jul 25, 2008
<u>PAMIDRONATE DISODIUM; AREDIA</u>					
020036 001	4711880	Jul 29, 2005			
<u>PAMIDRONATE DISODIUM; AREDIA</u>					
020036 003	4711880	Jul 29, 2005			
<u>PAMIDRONATE DISODIUM; AREDIA</u>					
020036 004	4711880	Jul 29, 2005			
<u>PANTOPRAZOLE SODIUM; PROTONIX</u>					
020987 001	4758579 5997903	Jul 19, 2010 Dec 07, 2016			
<u>PANTOPRAZOLE SODIUM; PROTONIX</u>					
020987 002	4758579 5997903	Jul 19, 2010 Dec 07, 2016			
<u>PANTOPRAZOLE SODIUM; PROTONIX IV</u>					
020988 001	4758579 6780881	Jul 19, 2010 Nov 17, 2021	DP	I-444	Dec 06, 2007
<u>PARICALCITOL; ZEMPLAR</u>					
020819 001	5246925 5246925*PED 5587497 5587497*PED 6136799 6136799*PED 6361758 6361758*PED	Apr 17, 2012 Oct 17, 2012 Dec 24, 2013 Jun 24, 2014 Apr 08, 2018 Oct 08, 2019 Apr 08, 2018 Oct 08, 2018	U-314 DP	M-31 PED	Mar 31, 2007 Oct 01, 2007
<u>PARICALCITOL; ZEMPLAR</u>					
020819 002	5246925 5246925*PED 5587497 5587497*PED 6136799 6136799*PED 6361758 6361758*PED	Apr 17, 2012 Oct 17, 2012 Dec 24, 2013 Jun 24, 2014 Apr 08, 2018 Oct 08, 2019 Apr 08, 2018 Oct 08, 2018	U-314 DP	M-31 PED	Mar 31, 2007 Oct 01, 2007
<u>PARICALCITOL; ZEMPLAR</u>					
021606 001	5246925 5246925*PED 5587497 5587497*PED	Apr 17, 2012 Oct 17, 2012 Dec 24, 2013 Jun 24, 2014	U-671 DS	NDF	May 26, 2008
<u>PARICALCITOL; ZEMPLAR</u>					
021606 002	5246925 5246925*PED 5587497 5587497*PED	Apr 17, 2012 Oct 17, 2012 Dec 24, 2013 Jun 24, 2014	U-671 DS	NDF	May 26, 2008
<u>PARICALCITOL; ZEMPLAR</u>					
021606 003	5246925 5246925*PED 5587497 5587497*PED	Apr 17, 2012 Oct 17, 2012 Dec 24, 2013 Jun 24, 2014	U-671 DS	NDF	May 26, 2008

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020031 001	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6113944	Dec 14, 2014			
	6113944*PED	Jun 14, 2015			
	6121291	Mar 17, 2017		U-431	
	6121291	Mar 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-431	
	6133289	May 19, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020031 002	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6113944	Dec 14, 2014			
	6113944*PED	Jun 14, 2015			
	6121291	Mar 17, 2017		U-286	
	6121291	Mar 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-431	
	6133289	May 19, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020031 003	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6113944	Dec 14, 2014			
	6113944*PED	Jun 14, 2015			
	6121291	Mar 17, 2017		U-286	
	6121291	Mar 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-431	
	6133289	May 19, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020031 004	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6113944	Dec 14, 2014			
	6113944*PED	Jun 14, 2015			
	6121291	Mar 17, 2017		U-431	
	6121291	Mar 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-431	
	6133289	May 15, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020031 005	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6113944	Dec 14, 2014			
	6113944*PED	Jun 14, 2015			
	6121291	Mar 17, 2017		U-431	
	6121291	Mar 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-286	
	6133289	May 15, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020710 001	4721723	Dec 29, 2006			
	4721723*PED	Jun 29, 2007			
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5811436	Sep 22, 2015			
	5811436*PED	Mar 22, 2016			
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6121291	Mar 17, 2017		U-286	
	6121291	Mar 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-286	
	6133289	May 19, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020885 001	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6063927	Apr 23, 2019			
	6063927*PED	Oct 23, 2019			
	6080759	May 19, 2015			
	6080759*PED	Nov 19, 2015			
	6121291	Mar 17, 2017		U-286	
	6121291	Mar 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-431	
	6133289	May 19, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
	6172233	Jan 15, 2018			
	6172233*PED	Jul 15, 2018			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020885 002	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6063927	Apr 23, 2019			
	6063927*PED	Oct 23, 2019			
	6080759	May 19, 2015			
	6080759*PED	Nov 19, 2015			
	6121291	Mar 17, 2017		U-431	
	6121291	Mar 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-431	
	6133289	May 19, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
	6172233	Jan 15, 2018			
	6172233*PED	Jul 15, 2018			
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020885 003	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6063927	Apr 23, 2019			
	6063927*PED	Oct 23, 2019			
	6080759	May 19, 2015			
	6080759*PED	Nov 19, 2015			
	6121291	Mar 17, 2017		U-286	
	6121291	Mar 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-286	
	6133289	May 19, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
	6172233	Jan 15, 2018			
	6172233*PED	Jul 15, 2018			
<u>PAROXETINE HYDROCHLORIDE; PAXIL</u>					
020885 004	4721723	Dec 29, 2006		U-12	
	4721723*PED	Jun 29, 2007		U-12	
	5789449	Jan 06, 2009		U-285	
	5789449*PED	Jul 06, 2009		U-285	
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6062927	Apr 23, 2019			
	6063927*PED	Oct 23, 2019			
	6080759	May 19, 2015			
	6080759*PED	Nov 19, 2015			
	6121291	Mar 17, 2017		U-286	
	6121291	Mar 17, 2017		U-431	
	6121291*PED	Sep 17, 2017		U-286	
	6121291*PED	Sep 17, 2017		U-431	
	6133289	May 19, 2015		U-358	
	6133289*PED	Nov 19, 2015		U-358	
	6172233	Jan 15, 2018			
	6172233*PED	Jul 15, 2018			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PAROXETINE HYDROCHLORIDE; PAXIL CR</u>					
020936 001	4721723	Dec 29, 2006		D-91	Jan 27, 2007
	4721723*PED	Jun 29, 2007		I-405	Aug 28, 2006
	4839177	Jun 13, 2006		PED	Aug 12, 2005
	4839177*PED	Dec 13, 2006			
	5422123	Jun 06, 2012			
	5422123*PED	Dec 06, 2012			
	5789449	Jan 06, 2009	U-286		
	5789449*PED	Jul 06, 2009	U-286		
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6121291	Mar 17, 2017	U-286		
	6121291*PED	Sep 17, 2017	U-286		
	6133289	May 19, 2015	U-286		
	6133289*PED	Nov 19, 2015	U-286		
	6548084	Jul 19, 2016			
	6548084*PED	Jan 19, 2017			
<u>PAROXETINE HYDROCHLORIDE; PAXIL CR</u>					
020936 002	4721723	Dec 29, 2006		D-91	Jan 27, 2007
	4721723*PED	Jun 29, 2007		I-405	Aug 28, 2006
	4839177	Jun 13, 2006		PED	Aug 12, 2005
	4839177*PED	Dec 13, 2006			
	5422123	Jun 06, 2012			
	5422123*PED	Dec 06, 2012			
	5789449	Jan 06, 2009	U-286		
	5789449*PED	Jul 06, 2009	U-286		
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6121291	Mar 17, 2017	U-286		
	6121291*PED	Sep 17, 2017	U-286		
	6133289	May 19, 2015	U-286		
	6133289*PED	Nov 19, 2015	U-286		
	6548084	Jul 19, 2016			
	6548084*PED	Jan 19, 2017			
<u>PAROXETINE HYDROCHLORIDE; PAXIL CR</u>					
020936 003	4721723	Dec 29, 2006		D-91	Jan 27, 2007
	4721723*PED	Jun 29, 2007		I-405	Aug 28, 2006
	4839177	Jun 13, 2006		PED	Aug 12, 2005
	4839177*PED	Dec 13, 2006			
	5422123	Jun 06, 2012			
	5422123*PED	Dec 06, 2012			
	5789449	Jan 06, 2009	U-286		
	5789449*PED	Jul 06, 2009	U-286		
	5872132	May 19, 2015			
	5872132*PED	Nov 19, 2015			
	5900423	May 19, 2015			
	5900423*PED	Nov 19, 2015			
	6121291	Mar 17, 2017	U-286		
	6121291*PED	Sep 17, 2017	U-286		
	6133289	May 19, 2015	U-286		
	6133289*PED	Nov 19, 2015	U-286		
	6548084	Jul 19, 2016			
	6548084*PED	Jan 19, 2017			
<u>PAROXETINE MESYLATE; PEXEVA</u>					
021299 001	5874447	Jun 10, 2017	U-518		
	5874447	Jun 10, 2017	U-46		
	5874447	Jun 10, 2017	U-286		
	6703408	Oct 21, 2022	DP		
<u>PAROXETINE MESYLATE; PEXEVA</u>					
021299 002	5874447	Jun 10, 2017	U-46		
	5874447	Jun 10, 2017	U-518		
	5874447	Jun 10, 2017	U-286		
	6703408	Oct 21, 2022	DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE			
PAROXETINE MESYLATE; PEVEVA										
021299 003	5874447	Jun	10,	2017	DP	U-286				
	5874447	Jun	10,	2017		U-46				
	5874447	Jun	10,	2017		U-518				
	6703408	Oct	21,	2022						
PAROXETINE MESYLATE; PEVEVA										
021299 004	5874447	Jun	10,	2017	DP	U-286				
	5874447	Jun	10,	2017		U-46				
	5874447	Jun	10,	2017		U-518				
	6703408	Oct	21,	2022						
PEGAPTANIB SODIUM; MACUGEN										
021756 001	5919455	Oct	27,	2013	DS	NCE	Dec	17,	2009	
	5932462	Aug	03,	2016	DS					
	6011020	Jan	04,	2017	DS					
	6051698	Oct	17,	2012	DS					
	6113906	Oct	27,	2013	DS					
	6147204	Jun	11,	2010	DS					
	6426335	Jun	11,	2010	U-622					
PEGVISOMANT; SOMAVERT										
021106 001	5350836	Sep	27,	2011	U-507	NCE	Mar	25,	2008	
	5681809	Sep	27,	2011	U-507					
	5849535	Sep	21,	2015	ODE					
	5958879	Sep	27,	2011	U-507					
	6057292	Sep	21,	2015	U-507					
	6583115	Sep	27,	2011	U-507					
PEGVISOMANT; SOMAVERT										
021106 002	5350836	Sep	27,	2011	U-507	NCE	Mar	25,	2008	
	5681809	Sep	27,	2011	U-507					
	5849535	Sep	21,	2015	ODE					
	5958879	Sep	27,	2011	U-507					
	6057292	Sep	21,	2015	U-507					
	6583115	Sep	27,	2011	U-507					
PEGVISOMANT; SOMAVERT										
021106 003	5350836	Sep	27,	2011	U-507	NCE	Mar	25,	2008	
	5681809	Sep	27,	2011	U-507					
	5849535	Sep	21,	2015	ODE					
	5958879	Sep	27,	2011	U-507					
	6057292	Sep	21,	2015	U-507					
	6583115	Sep	27,	2011	U-507					
PEMETREXED DISODIUM; ALIMTA										
021462 001	5217974	Mar	29,	2011	DS DP	U-551	NCE	Feb	04,	2009
	5344932	Sep	06,	2011		ODE				
PEMIROLAST POTASSIUM; ALAMAST										
021079 001	5034230	Dec	23,	2008		U-184				
	5034230*PED	Jun	23,	2009		U-184				
PENCICLOVIR SODIUM; DENAVIR										
020629 001	5075445	Sep	24,	2010		NPP	Oct	10,	2006	
	5840763	Sep	01,	2015						U-501
	5866581	Sep	04,	2014						U-501
	5916893	Sep	01,	2015						U-501
	6124304	Sep	04,	2014						U-501
	6469015	Oct	22,	2019						U-501
	6573378	Sep	24,	2010						
	6579981	Jun	17,	2020						U-501
PENTETATE CALCIUM TRISODIUM; PENTETATE CALCIUM TRISODIUM										
021749 001						NCE	Aug	11,	2009	
						ODE				
PENTETATE ZINC TRISODIUM; PENTETATE ZINC TRISODIUM										
021751 001						NCE	Aug	11,	2009	
						ODE				
PENTOSAN POLYSULFATE SODIUM; ELMIRON										
020193 001	5180715	Jan	19,	2010		U-159				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PERFLUOROPOLYMETHYLISOPROPYL ETHER; POLYTETRAFLUOROETHYLENE; SKIN EXPOSURE REDUCTION PASTE AGAINST CHEMICAL WARFARE AGENTS</u>					
021084 001	5607979	May 30, 2015			
<u>PERFLUTREN; DEFINITY</u>					
021064 001	5547656	Apr 05, 2011		NCE	Jul 31, 2006
	5769080	Jul 20, 2010			
	6033645	Jun 19, 2016		U-665	
	6146657	Dec 22, 2009	DS		
	6528039	Apr 05, 2011	DS		
	6773696	Apr 05, 2011	DS		
<u>PERGOLIDE MESYLATE; PERMAX</u>					
019385 001	4797405	Oct 26, 2007			
	5114948	Oct 19, 2009			
<u>PERGOLIDE MESYLATE; PERMAX</u>					
019385 002	4797405	Oct 26, 2007			
	5114948	Oct 19, 2009			
<u>PERGOLIDE MESYLATE; PERMAX</u>					
019385 003	4797405	Oct 26, 2007			
	5114948	Oct 19, 2009			
<u>PERINDOPRIL ERBUMINE; ACEON</u>					
020184 001	4508729	Aug 21, 2006		I-468	Aug 23, 2008
	5162362	Nov 10, 2009	DS DP U-531		
<u>PERINDOPRIL ERBUMINE; ACEON</u>					
020184 002	4508729	Aug 21, 2006		I-468	Aug 23, 2008
	5162362	Nov 10, 2009	DS DP U-531		
<u>PERINDOPRIL ERBUMINE; ACEON</u>					
020184 003	4508729	Aug 21, 2006		I-468	Aug 23, 2008
	5162362	Nov 10, 2009	DS DP U-531		
<u>PIMECROLIMUS; ELIDEL</u>					
021302 001	5912238	Jun 15, 2016		NCE	Dec 13, 2006
	5912238*PED	Dec 15, 2016		PED	Jun 13, 2007
	6352998	Oct 26, 2015			
	6352998*PED	Apr 26, 2016			
	6423722	Jun 26, 2018			
	6423722*PED	Dec 26, 2018			
<u>PIOGLITAZONE HYDROCHLORIDE; ACTOS</u>					
021073 001	4687777	Jan 17, 2011		M-35	Nov 26, 2006
	5965584	Jun 19, 2016			
	6150383	Jun 19, 2016	U-417		
	6150384	Jun 19, 2016	U-418		
	6166042	Jun 19, 2016	U-419		
	6166043	Jun 19, 2016	U-414		
	6172090	Jun 19, 2016	U-415		
	6211205	Jun 19, 2016	U-416		
	6271243	Jun 19, 2016	U-410		
	6303640	Jun 19, 2016	U-411		
	6329404	Aug 09, 2016	U-425		
		Jun 19, 2016	U-430		
<u>PIOGLITAZONE HYDROCHLORIDE; ACTOS</u>					
021073 002	4687777	Jan 17, 2011		M-35	Nov 26, 2006
	5965584	Jun 19, 2016			
	6150383	Jun 19, 2016	U-417		
	6150384	Jun 19, 2016	U-418		
	6166042	Jun 19, 2016	U-419		
	6166043	Jun 19, 2016	U-414		
	6172090	Jun 19, 2016	U-415		
	6211205	Jun 19, 2016	U-416		
	6271243	Jun 19, 2016	U-410		
	6303640	Jun 19, 2016	U-411		
	6329404	Aug 09, 2016	U-425		
		Jun 19, 2016	U-430		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PIOGLITAZONE HYDROCHLORIDE; ACTOS</u>					
021073 003	4687777	Jan 17, 2011		M-35	Nov 26, 2006
	5965584	Jun 19, 2016	U-417		
	6150383	Jun 19, 2016	U-418		
	6150384	Jun 19, 2016	U-419		
	6166042	Jun 19, 2016	U-414		
	6166043	Jun 19, 2016	U-415		
	6172090	Jun 19, 2016	U-416		
	6211205	Jun 19, 2016	U-410		
	6271243	Jun 19, 2016	U-411		
	6303640	Aug 09, 2016	U-425		
	6329404	Jun 19, 2016	U-430		
<u>PODOFILOX; CONDYLOX</u>					
020529 001	5057616	Oct 15, 2008			
<u>POLYETHYLENE GLYCOL 3350; MIRALAX</u>					
020698 001	5710183	Jul 14, 2015	U-265		
	6048901	Apr 20, 2019	U-343		
<u>PORFIMER SODIUM; PHOTOFRIN</u>					
020451 001	4932934	Jun 12, 2007	U-128	I-391	Aug 01, 2006
	5145863	Dec 15, 2009	U-129	ODE	Aug 01, 2010
	5438071	Aug 01, 2012			
<u>POTASSIUM CHLORIDE; K-DUR 10</u>					
019439 002	4863743	Sep 05, 2006	U-99		
<u>POTASSIUM CHLORIDE; K-DUR 20</u>					
019439 001	4863743	Sep 05, 2006	U-99		
<u>PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX</u>					
020667 001	4843086	Nov 23, 2007	U-231		
	4886812	Mar 25, 2011			
<u>PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX</u>					
020667 002	4843086	Nov 23, 2007	U-231		
	4886812	Mar 25, 2011			
<u>PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX</u>					
020667 003	4843086	Nov 23, 2007	U-231		
	4886812	Mar 25, 2011			
<u>PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX</u>					
020667 004	4843086	Nov 23, 2007	U-231		
	4886812	Mar 25, 2011			
<u>PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX</u>					
020667 005	4843086	Nov 23, 2007	U-231		
	4886812	Mar 25, 2011			
<u>PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX</u>					
020667 006	4843086	Nov 23, 2007	U-231		
	4886812	Mar 25, 2011			
<u>PRAMLINTIDE ACETATE; SYMLIN</u>					
021332 001	5175145	Dec 29, 2009	U-637	NCE	Mar 16, 2010
	5686411	Nov 11, 2014	DS DP U-638		
	5814600	Sep 29, 2015	U-639		
	5998367	Mar 08, 2011	DS DP		
	6114304	Sep 05, 2017	U-640		
	6410511	Jan 09, 2018	DP		
	6608029	Sep 07, 2013	U-641		
	6610824	Mar 08, 2011	DS		
<u>PRAVASTATIN SODIUM; PRAVACHOL</u>					
019898 002	4346227	Oct 20, 2005		I-370	Oct 29, 2005
	4346227*PED	Apr 20, 2006		PED	Apr 29, 2006
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PRAVASTATIN SODIUM; PRAVACHOL</u>					
019898 003	4346227	Oct 20, 2005		I-370	Oct 29, 2005
	4346227*PED	Apr 20, 2006		PED	Apr 29, 2006
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>PRAVASTATIN SODIUM; PRAVACHOL</u>					
019898 004	4346227	Oct 20, 2005		I-370	Oct 29, 2005
	4346227*PED	Apr 20, 2006		PED	Apr 29, 2006
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>PRAVASTATIN SODIUM; PRAVACHOL</u>					
019898 008	4346227	Oct 20, 2005		I-370	Oct 29, 2005
	4346227*PED	Apr 20, 2006		PED	Apr 29, 2006
	5030447	Jul 09, 2008			
	5030447*PED	Jan 09, 2009			
	5180589	Jul 09, 2008			
	5180589*PED	Jan 09, 2009			
	5622985	Apr 22, 2014	U-335		
	5622985*PED	Oct 22, 2014	U-335		
<u>PREGABALIN; LYRICA</u>					
021446 001	5563175	Oct 08, 2013		U-661	Dec 30, 2009
	6001876	Jul 16, 2017		U-55	
	6197819	Mar 06, 2018	DS DP		
<u>PREGABALIN; LYRICA</u>					
021446 002	5563175	Oct 08, 2013		U-661	Dec 30, 2009
	6001876	Jul 16, 2017		U-55	
	6197819	Mar 06, 2018	DS DP		
<u>PREGABALIN; LYRICA</u>					
021446 003	5563175	Oct 08, 2013		U-661	Dec 30, 2009
	6001876	Jul 16, 2017		U-55	
	6197819	Mar 06, 2018	DS DP		
<u>PREGABALIN; LYRICA</u>					
021446 004	5563175	Oct 08, 2013		U-661	Dec 30, 2009
	6001876	Jul 16, 2017		U-55	
	6197819	Mar 06, 2018	DS DP		
<u>PREGABALIN; LYRICA</u>					
021446 005	5563175	Oct 08, 2013		U-661	Dec 30, 2009
	6001876	Jul 16, 2017		U-55	
	6197819	Mar 06, 2018	DS DP		
<u>PREGABALIN; LYRICA</u>					
021446 006	5563175	Oct 08, 2013		U-661	Dec 30, 2009
	6001876	Jul 16, 2017		U-55	
	6197819	Mar 06, 2018	DS DP		
<u>PREGABALIN; LYRICA</u>					
021446 007	5563175	Oct 08, 2013		U-661	Dec 30, 2009
	6001876	Jul 16, 2017		U-55	
	6197819	Mar 06, 2018	DS DP		
<u>PREGABALIN; LYRICA</u>					
021446 008	5563175	Oct 08, 2013		U-661	Dec 30, 2009
	6001876	Jul 16, 2017		U-55	
	6197819	Mar 06, 2018	DS DP		
<u>PROCAINAMIDE HYDROCHLORIDE; PROCANBID</u>					
020545 001	5656296	Aug 12, 2014			
<u>PROCAINAMIDE HYDROCHLORIDE; PROCANBID</u>					
020545 002	5656296	Aug 12, 2014			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PROGESTERONE; CRINONE</u>					
020701 001	5543150	Sep 15, 2013	U-209		
<u>PROGESTERONE; CRINONE</u>					
020701 002	5543150	Sep 15, 2013	U-209		
<u>PROPAFENONE HYDROCHLORIDE; RYTHMOL SR</u>					
021416 001	5681588	Oct 28, 2014		NDF	Sep 04, 2006
<u>PROPAFENONE HYDROCHLORIDE; RYTHMOL SR</u>					
021416 002	5681588	Oct 28, 2014		NDF	Sep 04, 2006
<u>PROPAFENONE HYDROCHLORIDE; RYTHMOL SR</u>					
021416 003	5681588	Oct 28, 2014		NDF	Sep 04, 2006
<u>PROPOFOL; DIPRIVAN</u>					
019627 002	5714520	Mar 22, 2015			
	5714520*PED	Sep 22, 2015			
	5731355	Mar 22, 2015	U-217		
	5731355*PED	Sep 22, 2015	U-217		
	5731356	Mar 22, 2015	U-218		
	5731356*PED	Sep 22, 2015	U-218		
	5908869	Mar 22, 2015	U-270		
	5908869*PED	Sep 22, 2015	U-270		
<u>PROPRANOLOL HYDROCHLORIDE; INDERAL</u>					
019536 001	4600708	Jul 19, 2005			
<u>PROPRANOLOL HYDROCHLORIDE; INNOPRAN XL</u>					
021438 001	6500454	Dec 31, 2022		NP	Mar 12, 2006
<u>PROPRANOLOL HYDROCHLORIDE; INNOPRAN XL</u>					
021438 002	6500454	Dec 31, 2022		NP	Mar 12, 2006
<u>PSEUDOEPHEDRINE HYDROCHLORIDE; EFIDAC 24 PSEUDOEPHEDRINE HCL</u>					
020021 002	4801461	Mar 14, 2006			
<u>QUETIAPINE FUMARATE; SEROQUEL</u>					
020639 001	4879288	Sep 26, 2011	DS DP U-550	I-419 I-418	Jan 12, 2007 Jan 12, 2007
<u>QUETIAPINE FUMARATE; SEROQUEL</u>					
020639 002	4879288	Sep 26, 2011	DS DP U-550	I-419 I-418	Jan 12, 2007 Jan 12, 2007
<u>QUETIAPINE FUMARATE; SEROQUEL</u>					
020639 003	4879288	Sep 26, 2011	DS DP U-550	I-419 I-418	Jan 12, 2007 Jan 12, 2007
<u>QUETIAPINE FUMARATE; SEROQUEL</u>					
020639 004	4879288	Sep 26, 2011	DS DP U-550	I-419 I-418	Jan 12, 2007 Jan 12, 2007
<u>QUETIAPINE FUMARATE; SEROQUEL</u>					
020639 005	4879288	Sep 26, 2011	DS DP U-550	I-419 I-418	Jan 12, 2007 Jan 12, 2007
<u>QUETIAPINE FUMARATE; SEROQUEL</u>					
020639 006				I-419 I-418	Jan 12, 2007 Jan 12, 2007
<u>QUETIAPINE FUMARATE; SEROQUEL</u>					
020639 007				I-419 I-418	Jan 12, 2007 Jan 12, 2007
<u>QUINAPRIL HYDROCHLORIDE; ACCUPRIL</u>					
019885 001	4743450	Feb 24, 2007			
	4743450*PED	Aug 24, 2007			
	5684016	Nov 04, 2014	U-210		
	5684016*PED	May 04, 2015	U-210		
	5747504*PED	Oct 10, 2005	U-210		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>QUINAPRIL HYDROCHLORIDE; ACCUPRIL</u>					
019885 002	4743450	Feb 24, 2007			
	4743450*PED	Aug 24, 2007			
	5684016	Nov 04, 2014		U-210	
	5684016*PED	May 04, 2015		U-210	
	5747504*PED	Oct 10, 2005		U-210	
<u>QUINAPRIL HYDROCHLORIDE; ACCUPRIL</u>					
019885 003	4743450	Feb 24, 2007			
	4743450*PED	Aug 24, 2007			
	5684016	Nov 04, 2014		U-210	
	5684016*PED	May 04, 2015		U-210	
	5747504*PED	Oct 10, 2005		U-210	
<u>QUINAPRIL HYDROCHLORIDE; ACCUPRIL</u>					
019885 004	4743450	Feb 24, 2007			
	4743450*PED	Aug 24, 2007			
	5684016	Nov 04, 2014		U-210	
	5684016*PED	May 04, 2015		U-210	
	5747504*PED	Oct 10, 2005		U-210	
<u>QUININE SULFATE; QUININE SULFATE</u>					
021799 001				ODE	Aug 12, 2012
<u>RABEPRAZOLE SODIUM; ACIPHEX</u>					
020973 001	5035899	Apr 04, 2009		U-385	
	5045552	May 08, 2013		U-385	
<u>RABEPRAZOLE SODIUM; ACIPHEX</u>					
020973 002	5035899	Apr 04, 2009		U-385	
	5045552	May 08, 2013		U-385	
<u>RALOXIFENE HYDROCHLORIDE; EVISTA</u>					
020815 001	5393763	Jul 28, 2012		U-114	
	5457117	Jul 28, 2012		U-114	
	5478847	Mar 02, 2014		U-114	
	5811120	Mar 02, 2014			
	5972383	Mar 02, 2014		U-287	
	6458811	Mar 10, 2017			
	6797719	Mar 10, 2017	DP		
	6894064	Mar 10, 2017	DP	U-657	
	6906086	Jul 28, 2012		U-662	
	6906086	Jul 28, 2012		U-657	
<u>RAMELTEON; ROZEREM</u>					
021782 001	6034239	Mar 06, 2017	DS DP	U-674	NCE Jul 22, 2010
<u>RAMIPRIL; ALTACE</u>					
019901 001	5061722	Oct 29, 2008			
	5403856	Apr 04, 2012		U-71	
<u>RAMIPRIL; ALTACE</u>					
019901 002	5061722	Oct 19, 2008			
	5403856	Apr 04, 2012		U-71	
<u>RAMIPRIL; ALTACE</u>					
019901 003	5061722	Oct 19, 2008			
	5403856	Apr 04, 2012		U-71	
<u>RAMIPRIL; ALTACE</u>					
019901 004	5061722	Oct 19, 2008			
	5403856	Apr 04, 2012		U-71	
<u>RANITIDINE BISMUTH CITRATE; TRITEC</u>					
020559 001	5008256	Jul 17, 2009			
	5256684	Oct 26, 2010		U-199	
	5403830	Apr 04, 2012		U-200	
	5407688	Apr 04, 2012		U-201	
	5456925	Oct 10, 2012			
	5601848	Feb 11, 2014		U-202	
	5629297	May 13, 2014		U-186	
<u>RANITIDINE HYDROCHLORIDE; ZANTAC</u>					
019675 001	5068249	Nov 26, 2008			
	5068249*PED	May 26, 2009			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RANITIDINE HYDROCHLORIDE; ZANTAC 150</u>					
020095 001	5028432	Feb 22, 2010			
	5028432*PED	Aug 22, 2010			
<u>RANITIDINE HYDROCHLORIDE; ZANTAC 150</u>					
020251 001	5102665	Jun 23, 2009			
	5102665*PED	Dec 23, 2009			
<u>RANITIDINE HYDROCHLORIDE; ZANTAC 150</u>					
020251 002	5102665	Jun 23, 2009			
	5102665*PED	Dec 23, 2009			
<u>RANITIDINE HYDROCHLORIDE; ZANTAC 150</u>					
021698 001				NP	Aug 31, 2007
<u>RANITIDINE HYDROCHLORIDE; ZANTAC 300</u>					
020095 002	5028432	Feb 22, 2010			
	5028432*PED	Aug 22, 2010			
<u>RAPACURONIUM BROMIDE; RAPLON</u>					
020984 001	5418226	Apr 14, 2013			
<u>RAPACURONIUM BROMIDE; RAPLON</u>					
020984 002	5418226	Apr 14, 2013			
<u>REMIFENTANIL HYDROCHLORIDE; ULTIVA</u>					
020630 001	5019583	Feb 15, 2009			
	5019583*PED	Aug 15, 2009			
	5466700	Aug 30, 2013	U-156		
	5466700*PED	Mar 01, 2014	U-156		
<u>REMIFENTANIL HYDROCHLORIDE; ULTIVA</u>					
020630 002	5019583	Feb 15, 2009			
	5019583*PED	Aug 15, 2009			
	5466700	Aug 30, 2013	U-156		
	5466700*PED	Mar 01, 2014			
<u>REMIFENTANIL HYDROCHLORIDE; ULTIVA</u>					
020630 003	5019583	Feb 15, 2009			
	5019583*PED	Aug 15, 2009			
	5466700	Aug 30, 2013	U-156		
	5466700*PED	Mar 01, 2014	U-156		
<u>REPAGLINIDE; PRANDIN</u>					
020741 001	5312924	Sep 05, 2006	U-214	I-367	Oct 21, 2005
	6143769	Sep 05, 2006			
	6677358	Jun 12, 2018	DS DP U-546		
	RE37035	Mar 14, 2009			
<u>REPAGLINIDE; PRANDIN</u>					
020741 002	5312924	Sep 05, 2006	U-214	I-367	Oct 21, 2005
	6143769	Sep 05, 2006			
	6677358	Jun 12, 2018	DS DP U-546		
	RE37035	Mar 14, 2009			
<u>REPAGLINIDE; PRANDIN</u>					
020741 003	5312924	Sep 05, 2006	U-214	I-367	Oct 21, 2005
	6143769	Sep 05, 2006			
	6677358	Jun 12, 2018	DS DP U-546		
	RE37035	Mar 14, 2009			
<u>RIBAVIRIN; COPEGUS</u>					
021511 001				I-447 NP	Feb 25, 2008 Dec 03, 2005
<u>RIBAVIRIN; COPEGUS</u>					
021511 002				I-447 NP	Feb 25, 2008 Dec 03, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RIBAVIRIN; REBETOL</u>					
020903 001	5767097	Jan 23, 2016	U-235	ODE	Jul 29, 2010
	5767097*PED	Jul 23, 2016	U-235	PED	Jan 29, 2011
	5914128	Dec 22, 2017			
	5914128*PED	Jun 22, 2018			
	6051252	Dec 22, 2017			
	6051252*PED	Jun 22, 2018			
	6063772	Jan 23, 2016	U-375		
	6063772*PED	Jul 23, 2016	U-375		
	6172046	Sep 21, 2017	U-377		
	6172046*PED	Mar 21, 2018	U-377		
	6177074	Nov 01, 2016	U-454		
	6177074*PED	May 01, 2017	U-454		
	6335032	Dec 22, 2017			
	6335032*PED	Jun 22, 2018			
	6337090	Dec 22, 2017			
	6337090*PED	Jun 22, 2018			
	6461605	Nov 01, 2016	U-478		
	6461605*PED	May 01, 2017	U-478		
	6472373	Sep 21, 2017	U-479		
	6472373*PED	Mar 21, 2018	U-479		
	6524570	Nov 01, 2016	U-499		
	6524570*PED	May 01, 2017	U-499		
<u>RIBAVIRIN; REBETOL</u>					
020903 002	5767097	Jan 23, 2016	U-235	ODE	Jul 29, 2010
	5767097*PED	Jul 23, 2016	U-235	PED	Jan 29, 2011
	5914128	Dec 22, 2017			
	5914128*PED	Jun 22, 2018			
	6051252	Dec 22, 2017			
	6051252*PED	Jun 22, 2018			
	6063772	Jan 23, 2016	U-375		
	6063772*PED	Jul 23, 2016	U-375		
	6172046	Sep 21, 2017	U-377		
	6172046*PED	Mar 21, 2018	U-377		
	6177074	Nov 01, 2016	U-454		
	6177074*PED	May 01, 2017	U-454		
	6335032	Dec 22, 2017			
	6335032*PED	Jun 22, 2018			
	6337090	Dec 22, 2017			
	6337090*PED	Jun 22, 2018			
	6461605	Nov 01, 2016	U-478		
	6461605*PED	May 01, 2017	U-478		
	6472373	Sep 21, 2017	U-479		
	6472373*PED	Mar 21, 2018	U-479		
	6524570	Nov 01, 2016	U-499		
	6524570*PED	May 01, 2017	U-499		
<u>RIBAVIRIN; REBETOL</u>					
021546 001	5767097	Jan 23, 2016	U-521	NDF	Jul 29, 2006
	5767097*PED	Jul 23, 2016	U-521	ODE	Jul 29, 2010
	6063772	Jan 23, 2016	U-521	PED	Jan 29, 2011
	6063772*PED	Jul 23, 2016	U-521	PED	Jan 29, 2007
	6172046	Sep 21, 2017	U-521		
	6172046*PED	Mar 21, 2018	U-521		
	6461605	Nov 01, 2016	U-521		
	6461605*PED	May 01, 2017	U-521		
	6472373	Sep 21, 2017	U-521		
	6472373*PED	Mar 21, 2018	U-521		
<u>RIBAVIRIN; VIRAZOLE</u>					
018859 001	6150337	Nov 21, 2017	U-400		
<u>RIFAXIMIN; XIFAXAN</u>					
021361 001				NCE	May 25, 2009
<u>RILUZOLE; RILUTEK</u>					
020599 001	5527814	Jun 18, 2013			
<u>RIMEXOLONE; VEXOL</u>					
020474 001	4686214	Jul 22, 2008	U-100		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE		
<u>RISEDRONATE SODIUM; ACTONEL</u>										
020835 001	5583122	Dec	10,	2013	U-222					
	6096342	Nov	22,	2011						
	6165513	Jun	10,	2018						
<u>RISEDRONATE SODIUM; ACTONEL</u>										
020835 002	5583122	Dec	10,	2013	U-222					
	6096342	Nov	22,	2011						
	6165513	Jun	10,	2018						
<u>RISEDRONATE SODIUM; ACTONEL</u>										
020835 003	5583122	Dec	10,	2013	U-222					
	5994329	Jul	17,	2018						U-353
	6015801	Jul	17,	2018						
	6096342	Nov	22,	2011	U-595					
	6165513	Jun	10,	2018						
	6432932	Jul	17,	2018						
	6465443	Aug	14,	2018						
					DP					
<u>RISPERIDONE; RISPERDAL</u>										
020272 001	4804663	Dec	29,	2007	U-90	I-413 I-412	Dec 04,	2006		
<u>RISPERIDONE; RISPERDAL</u>										
020272 002	4804663	Dec	29,	2007	U-90	I-413 I-412	Dec 04,	2006		
<u>RISPERIDONE; RISPERDAL</u>										
020272 003	4804663	Dec	29,	2007	U-90	I-413 I-412	Dec 04,	2006		
<u>RISPERIDONE; RISPERDAL</u>										
020272 004	4804663	Dec	29,	2007	U-90	I-413 I-412	Dec 04,	2006		
<u>RISPERIDONE; RISPERDAL</u>										
020272 005	4804663	Dec	29,	2007	U-90	I-413 I-412	Dec 04,	2006		
<u>RISPERIDONE; RISPERDAL</u>										
020272 007	4804663	Dec	29,	2007	U-90	I-413 I-412	Dec 04,	2006		
<u>RISPERIDONE; RISPERDAL</u>										
020272 008	4804663	Dec	29,	2007	U-90	I-413 I-412	Dec 04,	2006		
<u>RISPERIDONE; RISPERDAL</u>										
020588 001	4804663	Dec	29,	2007	U-90	I-413 I-412	Dec 04,	2006		
	5453425	Jul	11,	2014						
	5616587	Jul	11,	2014						
<u>RISPERIDONE; RISPERDAL</u>										
021444 001	4804663	Dec	29,	2007	DS DP DP DP	U-516	I-413 I-412	Dec 04,	2006	
	5648093	Jul	15,	2014						
	6224905	Jun	10,	2017						
<u>RISPERIDONE; RISPERDAL</u>										
021444 002	4804663	Dec	29,	2007	DS DP DP DP	U-516	I-413 I-412	Dec 04,	2006	
	5648093	Jul	15,	2014						
	6224905	Jun	10,	2017						
<u>RISPERIDONE; RISPERDAL</u>										
021444 003	4804663	Dec	29,	2007	DS DP DP DP	U-516	I-413 I-412	Dec 04,	2006	
	5648093	Jul	15,	2014						
	6224905	Jun	10,	2017						
<u>RISPERIDONE; RISPERDAL</u>										
021444 004	4804663	Dec	29,	2007	DS DP DP DP	U-543	I-413 I-412	Dec 04,	2006	
	5648093	Jul	15,	2014						
	6224905	Jun	10,	2017						

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>RISPERIDONE; RISPERDAL</u>									
021444 005	4804663	Dec	29, 2007	DS	DP	U-543	I-413	Dec	04, 2006
	5648093	Jul	15, 2014		DP		I-412	Dec	04, 2006
	6224905	Jun	10, 2017		DP				
<u>RISPERIDONE; RISPERDAL CONSTA</u>									
021346 001	4804663	Dec	29, 2007			U-543	NDF	Oct	29, 2006
	5688801	Nov	18, 2014						
	5770231	Nov	19, 2013						
	5792477	May	02, 2017						
	5916598	May	02, 2017						
	5965168	Nov	19, 2013						
	6110503	May	02, 2017						
	6110921	Nov	19, 2013						
	6194006	Dec	30, 2018						
	6264987	May	19, 2020						
	6368632	Nov	19, 2013						
	6379703	Dec	30, 2018						
	6379704	May	19, 2020						
	6403114	May	02, 2017						
	6534092	May	19, 2020						
	6544559	Nov	19, 2013						
	6596316	Dec	30, 2008						
<u>RISPERIDONE; RISPERDAL CONSTA</u>									
021346 002	4804663	Dec	29, 2007			U-543	NDF	Oct	29, 2006
	5688801	Nov	18, 2014						
	5770231	Nov	19, 2013						
	5792477	May	02, 2017						
	5916598	May	02, 2017						
	5965168	Nov	19, 2013						
	6110503	May	02, 2017						
	6110921	Nov	19, 2013						
	6194006	Dec	30, 2018						
	6264987	May	19, 2020						
	6368632	Nov	19, 2013						
	6379703	Dec	30, 2018						
	6379704	May	19, 2020						
	6403114	May	02, 2017						
	6534092	May	19, 2020						
	6544559	Nov	19, 2013						
	6596316	Dec	30, 2008						
<u>RISPERIDONE; RISPERDAL CONSTA</u>									
021346 003	4804663	Dec	29, 2007			U-543	NDF	Oct	29, 2006
	5688801	Nov	18, 2014						
	5770231	Nov	19, 2013						
	5792477	May	02, 2017						
	5916598	May	02, 2017						
	5965168	Nov	19, 2013						
	6110503	May	02, 2017						
	6110921	Nov	19, 2013						
	6194006	Dec	30, 2018						
	6264987	May	19, 2020						
	6368632	Nov	19, 2013						
	6379703	Dec	30, 2018						
	6379704	May	19, 2020						
	6403114	May	02, 2017						
	6534092	May	19, 2020						
	6544559	Nov	19, 2013						
	6596316	Dec	30, 2008						

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RITONAVIR; NORVIR</u>					
020659 001	5484801	Jan 28, 2014			
	5484801*PED	Jul 28, 2014			
	5541206	Jul 30, 2013		U-140	
	5541206*PED	Jan 30, 2014			
	5635523	Jun 03, 2014		U-190	
	5635523*PED	Dec 03, 2014			
	5648497	Jul 15, 2014			
	5648497*PED	Jan 15, 2015			
	5674882	Oct 07, 2014			
	5674882*PED	Apr 07, 2015			
	5846987	Dec 29, 2012		U-190	
	5846987*PED	Jun 29, 2013			
	5886036	Dec 29, 2012			
	5886036*PED	Jun 29, 2013			
	6037157	Jun 26, 2016			
	6037157*PED	Dec 26, 2016			
	6703403	Jun 26, 2016		U-564	
	6703403*PED	Dec 26, 2016			
<u>RITONAVIR; NORVIR</u>					
020680 001	5541206	Jul 30, 2013		U-140	
	5541206*PED	Jan 30, 2014			
	5635523	Jun 03, 2014		U-190	
	5635523*PED	Dec 03, 2014			
	5648497	Jul 15, 2014			
	5648497*PED	Jan 15, 2015			
	5846987	Dec 29, 2012		U-190	
	5846987*PED	Jun 29, 2013			
	5948436	Sep 13, 2013			
	5948436*PED	Mar 13, 2014			
<u>RITONAVIR; NORVIR</u>					
020945 001	5541206	Jul 30, 2013		U-348	
	5541206*PED	Jan 30, 2014			
	5635523	Jun 03, 2014		U-347	
	5635523*PED	Dec 03, 2014			
	5648497	Jul 15, 2014			
	5648497*PED	Jan 15, 2015			
	5846987	Dec 29, 2012		U-347	
	5846987*PED	Jun 29, 2013			
	6232333	Nov 07, 2017			
	6232333*PED	May 07, 2018			
	6703403	Jun 26, 2016		U-564	
	6703403*PED	Dec 26, 2016			
<u>RIVASTIGMINE TARTRATE; EXELON</u>					
020823 003	4948807	Aug 14, 2007		U-322	
	5602176	Feb 11, 2014		U-322	
<u>RIVASTIGMINE TARTRATE; EXELON</u>					
020823 004	4948807	Aug 14, 2007		U-322	
	5602176	Feb 11, 2014		U-322	
<u>RIVASTIGMINE TARTRATE; EXELON</u>					
020823 005	4948807	Aug 14, 2007		U-322	
	5602176	Feb 11, 2014		U-322	
<u>RIVASTIGMINE TARTRATE; EXELON</u>					
020823 006	4948807	Aug 14, 2007		U-322	
	5602176	Feb 11, 2014		U-322	
<u>RIVASTIGMINE TARTRATE; EXELON</u>					
021025 001	4948807	Aug 14, 2007		U-322	
	5602176	Feb 11, 2014		U-322	
<u>RIZATRIPTAN BENZOATE; MAXALT</u>					
020864 001	5298520	Jun 29, 2012		U-240	
	5602162	Feb 11, 2014			
<u>RIZATRIPTAN BENZOATE; MAXALT</u>					
020864 002	5298520	Jun 29, 2012		U-240	
	5602162	Feb 11, 2014			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE					
<u>RIZATRIPTAN BENZOATE; MAXALT-MLT</u>													
020865 001	5298520	Jun	29, 2012	U-240									
	5457895	Oct	01, 2013										
	5602162	Feb	11, 2014		U-240								
<u>RIZATRIPTAN BENZOATE; MAXALT-MLT</u>													
020865 002	5298520	Jun	29, 2012	U-240									
	5457895	Oct	01, 2013										
	5602162	Feb	11, 2014		U-240								
<u>ROCURONIUM BROMIDE; ZEMURON</u>													
020214 002	4894369	Apr	13, 2008										
<u>ROCURONIUM BROMIDE; ZEMURON (P/F)</u>													
020214 001	4894369	Apr	13, 2008										
<u>ROFECOXIB; VIOXX</u>													
021042 001	5474995	Jun	24, 2013	DS	DP	U-602	I-423	Mar	26, 2007				
	5474995*PED	Dec	24, 2013							M-27	Aug	06, 2006	
	5691374	May	18, 2015	PED	Feb	06, 2007							
	5691374*PED	Nov	18, 2015				PED			Oct	11, 2005		
	6063811	May	06, 2017	U-602									
	6063811*PED	Nov	06, 2017										
	6239173	Jun	24, 2013	DS	DP	U-602							
	6239173*PED	Dec	24, 2013										
<u>ROFECOXIB; VIOXX</u>													
021042 002	5474995	Jun	24, 2013	DS	DP	U-602	I-423	Mar	26, 2007				
	5474995*PED	Dec	24, 2013							M-27	Aug	06, 2006	
	5691374	May	18, 2015	PED	Feb	06, 2007							
	5691374*PED	Nov	18, 2015				PED			Oct	11, 2005		
	6063811	May	06, 2017	U-602									
	6063811*PED	Nov	06, 2017										
	6239173	Jun	24, 2013	DS	DP	U-602							
	6239173*PED	Dec	24, 2013										
<u>ROFECOXIB; VIOXX</u>													
021042 003	5474995	Jun	24, 2013	DS	DP	U-602	I-423	Mar	26, 2007				
	5474995*PED	Dec	24, 2013							M-27	Aug	06, 2006	
	5691374	May	18, 2015	PED	Feb	06, 2007							
	5691374*PED	Nov	18, 2015				PED			Oct	11, 2005		
	6063811	May	06, 2017	U-602									
	6063811*PED	Nov	06, 2017										
	6239173	Jun	24, 2013	DS	DP	U-602							
	6239173*PED	Dec	24, 2013										
<u>ROFECOXIB; VIOXX</u>													
021052 001	5474995	Jun	24, 2013	U-266	M-27	Aug	06, 2006						
	5474995*PED	Dec	24, 2013					PED	Feb	06, 2007			
	5691374	May	18, 2015								PED	Oct	11, 2005
	5691374*PED	Nov	18, 2015										
	6063811	May	06, 2017										
	6063811*PED	Nov	06, 2017										
	6239173	Jun	24, 2013										
	6239173*PED	Dec	24, 2013										
<u>ROFECOXIB; VIOXX</u>													
021052 002	5474995	Jun	24, 2013	U-266	M-27	Aug	06, 2006						
	5474995*PED	Dec	24, 2013					PED	Feb	06, 2007			
	5691374	May	18, 2015								PED	Oct	11, 2005
	5691374*PED	Nov	18, 2015										
	6063811	May	06, 2017										
	6063811*PED	Nov	06, 2017										
	6239173	Jun	24, 2013										
	6239173*PED	Dec	24, 2013										
<u>ROPINIROLE HYDROCHLORIDE; REQUIP</u>													
020658 001	4452808	Dec	07, 2007	DS	DP	I-464	May	04, 2008					
	4824860	May	19, 2008										
<u>ROPINIROLE HYDROCHLORIDE; REQUIP</u>													
020658 002	4452808	Dec	07, 2007	DS	DP	I-464	May	04, 2008					
	4824860	May	19, 2008										
<u>ROPINIROLE HYDROCHLORIDE; REQUIP</u>													
020658 003	4452808	Dec	07, 2007	DS	DP	I-464	May	04, 2008					
	4824860	May	19, 2008										

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ROPINIROLE HYDROCHLORIDE; REQUIP</u>					
020658 004	4452808 4824860	Dec 07, 2007 May 19, 2008	DS DP U-212	I-464	May 04, 2008
<u>ROPINIROLE HYDROCHLORIDE; REQUIP</u>					
020658 005	4452808 4824860	Dec 07, 2007 May 19, 2008	DS DP U-212	I-464	May 04, 2008
<u>ROPINIROLE HYDROCHLORIDE; REQUIP</u>					
020658 006	4452808 4824860	Dec 07, 2007 May 19, 2008	DS DP	I-464	May 04, 2008
<u>ROPINIROLE HYDROCHLORIDE; REQUIP</u>					
020658 007	4452808 4824860	Dec 07, 2007 May 19, 2008	DS DP	I-464	May 04, 2008
<u>ROPIVACAINE HYDROCHLORIDE MONOHYDRATE; NAROPIN</u>					
020533 001	4870086	Sep 24, 2010			
<u>ROPIVACAINE HYDROCHLORIDE MONOHYDRATE; NAROPIN</u>					
020533 003	4870086	Sep 24, 2010			
<u>ROPIVACAINE HYDROCHLORIDE MONOHYDRATE; NAROPIN</u>					
020533 004	4870086	Sep 24, 2010			
<u>ROPIVACAINE HYDROCHLORIDE MONOHYDRATE; NAROPIN</u>					
020533 005	4870086	Sep 24, 2010			
<u>ROSIGLITAZONE MALEATE; AVANDIA</u>					
021071 002	5002953	Sep 17, 2011	DS DP U-329	I-453	Feb 28, 2008
	5002953	Sep 17, 2011	DS DP U-628	I-384	Feb 27, 2006
	5002953*PED	Mar 17, 2012		PED	Aug 27, 2006
	5741803	Apr 21, 2015	DS DP U-628		
	5741803	Apr 21, 2015	DS DP U-329		
	5741803*PED	Oct 21, 2015			
	6288095	Feb 11, 2017		U-420	
	6288095*PED	Aug 11, 2017			
<u>ROSIGLITAZONE MALEATE; AVANDIA</u>					
021071 003	5002953	Sep 17, 2011	DS DP U-329	I-453	Feb 28, 2008
	5002953	Sep 17, 2011	DS DP U-628	I-384	Feb 27, 2006
	5002953*PED	Mar 17, 2012		PED	Aug 27, 2006
	5741803	Apr 21, 2015	DS DP U-628		
	5741803	Apr 21, 2015	DS DP U-329		
	5741803*PED	Oct 21, 2015			
	6288095	Feb 11, 2017		U-420	
	6288095*PED	Aug 11, 2017			
<u>ROSIGLITAZONE MALEATE; AVANDIA</u>					
021071 004	5002953	Sep 17, 2011	DS DP U-628	I-453	Feb 28, 2008
	5002953	Sep 17, 2011	DS DP U-329	I-384	Feb 27, 2006
	5002953*PED	Mar 17, 2012		PED	Aug 27, 2006
	5741803	Apr 21, 2015	DS DP U-628		
	5741803	Apr 21, 2015	DS DP U-329		
	5741803*PED	Oct 21, 2015			
	6288095	Feb 11, 2017		U-420	
	6288095*PED	Aug 11, 2017			
<u>ROSUVASTATIN CALCIUM; CRESTOR</u>					
021366 002	6316460	Aug 04, 2020		NCE	Aug 12, 2008
	6589959	Dec 23, 2019			
	6858618	Dec 17, 2021			
	RE37314	Jun 12, 2012	U-618		
<u>ROSUVASTATIN CALCIUM; CRESTOR</u>					
021366 003	6316460	Aug 04, 2020		NCE	Aug 12, 2008
	6589959	Dec 23, 2019			
	6858618	Dec 17, 2021			
	RE37314	Jun 12, 2012	U-618		
<u>ROSUVASTATIN CALCIUM; CRESTOR</u>					
021366 004	6316460	Aug 04, 2020		NCE	Aug 12, 2008
	6589959	Dec 23, 2019			
	6858618	Dec 17, 2021			
	RE37314	Jun 12, 2012	U-618		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>ROSUVASTATIN CALCIUM; CRESTOR</u>							
021366 005	6316460	Aug 04, 2020			NCE	Aug 12, 2008	
	6589959	Dec 23, 2019					
	6858618	Dec 17, 2021	U-618				
	RE37314	Jun 12, 2012					
<u>SALMETEROL XINAFOATE; SEREVENT</u>							
020236 001	4992474	Feb 12, 2008			U-182		
	5126375	Feb 12, 2008					
	5225445	Feb 12, 2008					
<u>SALMETEROL XINAFOATE; SEREVENT</u>							
020692 001	4992474	Feb 12, 2008			U-211		
	5126375	Feb 12, 2008					
	5225445	Feb 12, 2008					
<u>SAMARIUM SM 153 LEXIDRONAM PENTASODIUM; QUADRAMET</u>							
020570 001	4898724	Feb 06, 2007		U-187			
<u>SAQUINAVIR; FORTOVASE</u>							
020828 001	5196438	Nov 19, 2010					
	6008228	Jun 06, 2015					
	6352717	Nov 16, 2019					
<u>SAQUINAVIR MESYLATE; INVIRASE</u>							
020628 001	5196438	Nov 19, 2010					
<u>SECRETIN SYNTHETIC HUMAN; CHIRHOSTIM</u>							
021256 001					NCE	Apr 09, 2009	
					ODE	Apr 04, 2009	
<u>SECRETIN SYNTHETIC PORCINE; SECREMAX</u>							
021136 001					I-408	Nov 01, 2005	
					ODE	Apr 04, 2009	
<u>SERTACONAZOLE NITRATE; ERTACZO</u>							
021385 001					NCE	Dec 10, 2008	
<u>SERTRALINE HYDROCHLORIDE; ZOLOFT</u>							
019839 001	4536518	Dec 30, 2005		U-12	I-261	Feb 07, 2006	
	4536518*PED	Jun 30, 2006		U-12			
	4962128	Nov 02, 2009		U-152			
	4962128*PED	May 02, 2010		U-152			
	5248699	Aug 13, 2012					
	5248699*PED	Feb 13, 2013					
	5744501	Jan 06, 2009	U-461				
	5789449	Jan 06, 2009	U-460				
<u>SERTRALINE HYDROCHLORIDE; ZOLOFT</u>							
019839 002	4536518	Dec 30, 2005		U-12	I-261	Feb 07, 2006	
	4536518*PED	Jun 30, 2006		U-12			
	4962128	Nov 02, 2009		U-152			
	4962128*PED	May 02, 2010		U-152			
	5248699	Aug 13, 2012					
	5248699*PED	Feb 13, 2013					
	5744501	Jan 06, 2009	U-461				
	5789449	Jan 06, 2009	U-460				
<u>SERTRALINE HYDROCHLORIDE; ZOLOFT</u>							
019839 003	4536518	Dec 30, 2005		U-12	I-261	Feb 07, 2006	
	4536518*PED	Jun 30, 2006		U-12			
	4962128	Nov 02, 2009		U-152			
	4962128*PED	May 02, 2010		U-152			
	5248699	Aug 13, 2012					
	5248699*PED	Feb 13, 2013					
	5744501	Jan 06, 2009	U-461				
	5789449	Jan 06, 2009	U-460				

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SERTRALINE HYDROCHLORIDE; ZOLOFT</u>					
019839 004	4536518	Dec 30, 2005	U-12	I-261	Feb 07, 2006
	4536518*PED	Jun 30, 2006	U-12		
	4962128	Nov 02, 2009	U-152		
	4962128*PED	May 02, 2010	U-152		
	5248699	Aug 13, 2012			
	5248699*PED	Feb 13, 2013			
	5744501	Jan 06, 2009	U-461		
	5789449	Jan 06, 2009	U-460		
<u>SERTRALINE HYDROCHLORIDE; ZOLOFT</u>					
019839 005	4536518	Dec 30, 2005	U-12	I-261	Feb 07, 2006
	4536518*PED	Jun 30, 2006	U-12		
	4962128	Nov 02, 2009	U-152		
	4962128*PED	May 02, 2010	U-152		
	5248699	Aug 13, 2012			
	5248699*PED	Feb 13, 2013			
	5744501	Jan 06, 2009	U-461		
	5789449	Jan 06, 2009	U-460		
<u>SERTRALINE HYDROCHLORIDE; ZOLOFT</u>					
020990 001	4536518	Dec 30, 2005	U-286	I-261	Feb 07, 2006
	4536518*PED	Jun 30, 2006	U-286		
	5248699	Aug 13, 2012			
	5248699*PED	Feb 13, 2013			
	5744501	Jan 06, 2009	U-461		
	5789449	Jan 06, 2009	U-460		
	6727283	Oct 11, 2019	DP U-580		
<u>SEVELAMER HYDROCHLORIDE; RENAGEL</u>					
020926 001	5496545	Aug 11, 2013	U-246		
	5667775	Sep 16, 2014	U-246		
	6509013	Aug 11, 2013			
<u>SEVELAMER HYDROCHLORIDE; RENAGEL</u>					
021179 001	5496545	Aug 11, 2013	U-246		
	5667775	Sep 16, 2014	U-246		
	6509013	Aug 11, 2013			
	6733780	Oct 18, 2020	DP		
<u>SEVELAMER HYDROCHLORIDE; RENAGEL</u>					
021179 002	5496545	Aug 11, 2013	U-246		
	5667775	Sep 16, 2014	U-246		
	6509013	Aug 13, 2013			
	6733780	Oct 18, 2020	DP		
<u>SEVOFLURANE; SEVOFLURANE</u>					
075895 001				PC	Mar 22, 2006
<u>SEVOFLURANE; ULTANE</u>					
020478 001	5990176	Jan 27, 2017			
	5990176*PED	Jul 27, 2017			
	6074668	Jan 09, 2018			
	6074668*PED	Jul 09, 2018			
	6288127	Jan 27, 2017			
	6288127*PED	Jul 27, 2017			
	6444859	Jan 27, 2017			
	6444859*PED	Jul 27, 2017			
<u>SIBUTRAMINE HYDROCHLORIDE; MERIDIA</u>					
020632 001	4746680	Jun 11, 2007		U-439	
	4746680*PED	Dec 11, 2007			
	4929629	May 29, 2007			
	4929629*PED	Nov 29, 2007			
	5436272	Jul 25, 2012			
	5436272*PED	Jan 25, 2013			
<u>SIBUTRAMINE HYDROCHLORIDE; MERIDIA</u>					
020632 002	4746680	Jun 11, 2007		U-439	
	4746680*PED	Dec 11, 2007			
	4929629	May 29, 2007			
	4929629*PED	Nov 29, 2007			
	5436272	Jul 25, 2012			
	5436272*PED	Jan 25, 2013			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
SIBUTRAMINE HYDROCHLORIDE; MERIDIA								
020632 003	4746680	Jun	11, 2007	U-439				
	4746680*PED	Dec	11, 2007					
	4929629	May	29, 2007					
	4929629*PED	Nov	29, 2007					
	5436272	Jul	25, 2012					
	5436272*PED	Jan	25, 2013					
SILDENAFIL CITRATE; REVATIO								
021845 001	5250534	Mar	27, 2012	DS	DP	NP	Jun	03, 2008
SILDENAFIL CITRATE; VIAGRA								
020895 001	5250534	Mar	27, 2012	U-155				
	6469012	Oct	22, 2019					
SILDENAFIL CITRATE; VIAGRA								
020895 002	5250534	Mar	27, 2012	U-155				
	6469012	Oct	22, 2019					
SILDENAFIL CITRATE; VIAGRA								
020895 003	5250534	Mar	27, 2012	U-155				
	6469012	Oct	22, 2019					
SIMVASTATIN; ZOCOR								
019766 001	4444784	Dec	23, 2005	U-59	I-390		Apr	16, 2006
	4444784*PED	Jun	23, 2006	U-59	I-350	PED	Oct	18, 2005
							Apr	18, 2006
SIMVASTATIN; ZOCOR								
019766 002	4444784	Dec	23, 2005	U-59	I-390		Apr	16, 2006
	4444784*PED	Jun	23, 2006	U-59	I-350	PED	Oct	18, 2005
							Apr	18, 2006
SIMVASTATIN; ZOCOR								
019766 003	4444784	Dec	23, 2005	U-59	I-390		Apr	16, 2006
	4444784*PED	Jun	23, 2006	U-59	I-350	PED	Oct	18, 2005
							Apr	18, 2006
SIMVASTATIN; ZOCOR								
019766 004	4444784	Dec	23, 2005	U-59	I-390		Apr	16, 2006
	4444784*PED	Jun	23, 2006	U-59	I-350	PED	Oct	18, 2005
							Apr	18, 2006
SIMVASTATIN; ZOCOR								
019766 005	4444784	Dec	23, 2005	U-59	I-390		Apr	16, 2006
	4444784*PED	Jun	23, 2006	U-59	I-350	PED	Oct	18, 2005
							Apr	18, 2006
SINCALIDE; KINEVAC								
017697 001	6803046	Aug	16, 2022	DP				
SIROLIMUS; RAPAMUNE								
021083 001	5100899	Jun	06, 2009	U-290	I-386		Apr	11, 2006
	5100899*PED	Dec	06, 2009		NPP		Mar	11, 2008
	5212155	May	18, 2010	U-291	PED		Sep	11, 2008
	5212155*PED	Nov	18, 2010		PED		Oct	11, 2006
	5308847	May	03, 2011	U-292				
	5308847*PED	Nov	03, 2011					
	5403833	Apr	04, 2012	U-293				
	5403833*PED	Oct	04, 2012					
	5536729	Sep	30, 2013	DP				
	5536729*PED	Mar	30, 2014					

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>SIROLIMUS; RAPAMUNE</u>								
021110 001	5100899	Jun 06, 2009		U-290	I-386	Apr 11, 2006		
	5100899*PED	Dec 06, 2009			NPP	Mar 11, 2008		
	5212155	May 18, 2010		U-291	PED	Sep 11, 2008		
	5212155*PED	Nov 18, 2010			PED	Oct 11, 2006		
	5308847	May 03, 2011		U-292				
	5308847*PED	Nov 03, 2011						
	5403833	Apr 04, 2012		U-293				
	5403833*PED	Oct 04, 2012						
	5989591	Mar 11, 2018	DP					
5989591*PED	Sep 11, 2018							
<u>SIROLIMUS; RAPAMUNE</u>								
021110 002	5100899	Jun 06, 2009		U-290	I-386	Apr 11, 2006		
	5100899*PED	Dec 06, 2009			NPP	Mar 11, 2008		
	5212155	May 18, 2010		U-291	PED	Sep 11, 2008		
	5212155*PED	Nov 18, 2010			PED	Oct 11, 2006		
	5308847	May 03, 2011		U-292				
	5308847*PED	Nov 03, 2011						
	5403833	Apr 04, 2012		U-293				
	5403833*PED	Oct 04, 2012						
	5989591	Mar 11, 2018	DP					
5989591*PED	Sep 11, 2018							
<u>SIROLIMUS; RAPAMUNE</u>								
021110 003	5100899	Jun 06, 2009		U-290	I-386	Apr 11, 2006		
	5100899*PED	Dec 06, 2009			NPP	Mar 11, 2008		
	5212155	May 18, 2010		U-291	PED	Sep 11, 2008		
	5212155*PED	Nov 18, 2010			PED	Oct 11, 2006		
	5403833	Apr 04, 2012		U-293				
	5403833*PED	Oct 04, 2012						
	5989591	Mar 11, 2018	DP					
	5989591*PED	Sep 11, 2018						
<u>SODIUM BENZOATE; SODIUM PHENYLACETATE; AMMONUL</u>								
020645 001					NDF	Feb 17, 2008		
					ODE	Feb 17, 2012		
<u>SODIUM FERRIC GLUCONATE COMPLEX; FERRLECIT</u>								
020955 001					NPP	Aug 13, 2007		
					PED	Feb 13, 2008		
<u>SODIUM FLUORIDE; TRICLOSAN; COLGATE TOTAL</u>								
020231 001	4894220	Jan 30, 2007		DP				
	5037635	Jan 30, 2007		DP				
	5156835	Oct 20, 2009		DP				
	5288480	Jan 30, 2007		DP				
	5344641	Jan 30, 2007		DP				
	5538715	Jan 30, 2007		DP				
	5776435	Jan 30, 2007		DP				
<u>SODIUM OXYBATE; XYREM</u>								
021196 001	6780889	Dec 22, 2019	DP		NCE	Jul 17, 2007		
					ODE	Nov 18, 2012		
					ODE	Jul 17, 2009		
<u>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE; VISICOL</u>								
021097 001	5616346	May 18, 2013		U-359				
<u>SOLIFENACIN SUCCINATE; VESICARE</u>								
021518 001	6017927	Dec 27, 2015	DS DP		NCE	Nov 19, 2009		
<u>SOLIFENACIN SUCCINATE; VESICARE</u>								
021518 002	6017927	Dec 27, 2015	DS DP		NCE	Nov 19, 2009		
<u>SOMATROPIN RECOMBINANT; GENOTROPIN</u>								
020280 006	4968299	Jun 28, 2008			ODE	Jul 25, 2008		
					ODE	Jun 20, 2007		
<u>SOMATROPIN RECOMBINANT; GENOTROPIN</u>								
020280 007	4968299	Jun 28, 2008			ODE	Jul 25, 2008		
					ODE	Jun 20, 2007		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 001	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015	DP		
	6152897	Nov 20, 2018			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 002	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015	DP		
	6152897	Nov 20, 2018			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 003	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015	DP		
	6152897	Nov 20, 2018			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 004	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
				ODE	Jun 20, 2007
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 005	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015	DP		
	6152897	Nov 20, 2018			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 008	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015	DP		
	6152897	Nov 20, 2018			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 009	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015	DP		
	6152897	Nov 20, 2018			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 010	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 011	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 012	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015			
SOMATROPIN RECOMBINANT; GENOTROPIN PRESERVATIVE FREE					
020280 013	4968299	Jun 28, 2008	DP	ODE	Jul 25, 2008
	5435076	Apr 16, 2013		ODE	Jun 20, 2007
	5716338	Feb 10, 2015			
SOMATROPIN RECOMBINANT; HUMATROPE					
019640 001				I-398	Jul 25, 2006
				M-45	Oct 12, 2008
SOMATROPIN RECOMBINANT; HUMATROPE					
019640 004				I-398	Jul 25, 2006
				M-45	Oct 12, 2008
SOMATROPIN RECOMBINANT; HUMATROPE					
019640 005				I-398	Jul 25, 2006
				M-45	Oct 12, 2008
SOMATROPIN RECOMBINANT; HUMATROPE					
019640 006				I-398	Jul 25, 2006
				M-45	Oct 12, 2008

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOMATROPIN RECOMBINANT; HUMATROPE</u>					
019640 007				I-398 M-45	Jul 25, 2006 Oct 12, 2008
<u>SOMATROPIN RECOMBINANT; NORDITROPIN</u>					
019721 001	5633352	May 27, 2014			
<u>SOMATROPIN RECOMBINANT; NORDITROPIN</u>					
019721 002	5633352	May 27, 2014			
<u>SOMATROPIN RECOMBINANT; NORDITROPIN</u>					
021148 001	5633352	May 27, 2014	U-340	I-439	Nov 01, 2007
	5849700	Dec 15, 2015			
	5849704	Dec 15, 2015			
<u>SOMATROPIN RECOMBINANT; NORDITROPIN</u>					
021148 002	5633352	May 27, 2014	U-340	I-439	Nov 01, 2007
	5849700	Dec 15, 2015			
	5849704	Dec 15, 2015			
<u>SOMATROPIN RECOMBINANT; NORDITROPIN</u>					
021148 003	5633352	May 27, 2014	U-340	I-439	Nov 01, 2007
	5849700	Dec 15, 2015			
	5849704	Dec 15, 2015			
<u>SOMATROPIN RECOMBINANT; NORDITROPIN NORDIFLEX</u>					
021148 004				I-439	Nov 01, 2007
<u>SOMATROPIN RECOMBINANT; NORDITROPIN NORDIFLEX</u>					
021148 005				I-439	Nov 01, 2007
<u>SOMATROPIN RECOMBINANT; NORDITROPIN NORDIFLEX</u>					
021148 006				I-439	Nov 01, 2007
<u>SOMATROPIN RECOMBINANT; NUTROPIN</u>					
020168 001	5096885	Mar 17, 2009	DP	I-462	Jun 28, 2008
<u>SOMATROPIN RECOMBINANT; NUTROPIN</u>					
020168 002	5096885	Mar 17, 2009	DP	I-462	Jun 28, 2008
<u>SOMATROPIN RECOMBINANT; NUTROPIN AQ</u>					
020522 001	5763394	Jun 09, 2015	DP	I-462 M-50	Jun 28, 2008 Nov 18, 2008
<u>SOMATROPIN RECOMBINANT; NUTROPIN AQ PEN</u>					
020522 002	5763394	Jun 09, 2015	DP	I-462 M-50	Jun 28, 2008 Nov 18, 2008
<u>SOMATROPIN RECOMBINANT; NUTROPIN DEPOT</u>					
021075 001	5654010	Aug 05, 2014	U-340		
	5656297	Jul 25, 2014			
	5912015	Mar 12, 2012			
	6051259	Dec 02, 2012			
<u>SOMATROPIN RECOMBINANT; NUTROPIN DEPOT</u>					
021075 002	5654010	Aug 05, 2014	U-340		
	5656297	Jul 25, 2014			
	5912015	Mar 12, 2012			
	6051259	Dec 02, 2012			
<u>SOMATROPIN RECOMBINANT; NUTROPIN DEPOT</u>					
021075 003	5654010	Aug 05, 2014	U-340		
	5656297	Jul 25, 2014			
	5912015	Mar 12, 2012			
	6051259	Dec 02, 2012			
<u>SOMATROPIN RECOMBINANT; SAIZEN</u>					
019764 001				I-440	Aug 26, 2007
<u>SOMATROPIN RECOMBINANT; SAIZEN</u>					
019764 002				I-440	Aug 26, 2007
<u>SOMATROPIN RECOMBINANT; SAIZEN</u>					
019764 003				I-440	Aug 26, 2007

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOMATROPIN RECOMBINANT; ZORBTIVE</u>					
021597 001				NP ODE	Dec 01, 2006 Dec 01, 2010
<u>SOMATROPIN RECOMBINANT; ZORBTIVE</u>					
021597 002				NP ODE	Dec 01, 2006 Dec 01, 2010
<u>SOMATROPIN RECOMBINANT; ZORBTIVE</u>					
021597 003				NP ODE	Dec 01, 2006 Dec 01, 2010
<u>SOMATROPIN RECOMBINANT; ZORBTIVE</u>					
021597 004				NP ODE	Dec 01, 2006 Dec 01, 2010
<u>SORAFENIB TOSYLATE; NEXAVAR</u>					
021923 001				NCE	Dec 20, 2010
<u>SPARFLOXACIN; ZAGAM</u>					
020677 001	4795751	Feb 04, 2010	U-160		
<u>STAVUDINE; ZERIT</u>					
020412 001	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-94 U-94		
<u>STAVUDINE; ZERIT</u>					
020412 002	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-94 U-94		
<u>STAVUDINE; ZERIT</u>					
020412 003	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-94 U-94		
<u>STAVUDINE; ZERIT</u>					
020412 004	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-94 U-94		
<u>STAVUDINE; ZERIT</u>					
020412 005	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-94 U-94		
<u>STAVUDINE; ZERIT</u>					
020413 001	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008			
<u>STAVUDINE; ZERIT XR</u>					
021453 001	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-248 U-248	NDF PED	Dec 31, 2005 Jul 01, 2006
<u>STAVUDINE; ZERIT XR</u>					
021453 002	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-248 U-248	NDF PED	Dec 31, 2005 Jul 01, 2006
<u>STAVUDINE; ZERIT XR</u>					
021453 003	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-248	NDF PED	Dec 31, 2005 Jul 01, 2006
<u>STAVUDINE; ZERIT XR</u>					
021453 004	4978655 4978655*PED	Jun 24, 2008 Dec 24, 2008	U-248 U-248	NDF PED	Dec 31, 2005 Jul 01, 2006
<u>SUMATRIPTAN; IMITREX</u>					
020626 001	4816470 4816470*PED 5037845 5037845*PED 5307953 5307953*PED 5554639 5554639*PED 5705520 5705520*PED	Dec 28, 2006 Jun 28, 2007 Aug 06, 2008 Feb 06, 2009 Dec 02, 2012 Jun 02, 2013 Sep 10, 2013 Mar 10, 2014 Dec 10, 2011 Jun 10, 2012	U-72 U-232 U-232		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SUMATRIPTAN; IMITREX</u>					
020626 002	4816470	Dec 28, 2006		U-72	
	4816470*PED	Jun 28, 2007			
	5037845	Aug 06, 2008			
	5037845*PED	Feb 06, 2009			
	5307953	Dec 02, 2012			
	5307953*PED	Jun 02, 2013			
	5554639	Sep 10, 2013		U-232	
	5554639*PED	Mar 10, 2014			
	5705520	Dec 10, 2011		U-232	
	5705520*PED	Jun 10, 2012			
<u>SUMATRIPTAN; IMITREX</u>					
020626 003	4816470	Dec 28, 2006		U-72	
	4816470*PED	Jun 28, 2007			
	5037845	Aug 06, 2008			
	5037845*PED	Feb 06, 2009			
	5307953	Dec 02, 2012			
	5307953*PED	Jun 02, 2013			
	5554639	Sep 10, 2013		U-232	
	5554639*PED	Mar 10, 2014			
	5705520	Dec 10, 2011		U-232	
	5705520*PED	Jun 10, 2012			
<u>SUMATRIPTAN SUCCINATE; IMITREX</u>					
020080 001	4816470	Dec 28, 2006		U-72	
	4816470*PED	Jun 28, 2007			
	5037845	Aug 06, 2008		U-72	
	5037845*PED	Feb 06, 2009			
<u>SUMATRIPTAN SUCCINATE; IMITREX</u>					
020132 001	4816470	Dec 28, 2006		U-72	
	4816470*PED	Jun 28, 2007			
	5037845	Aug 06, 2008		U-72	
	5037845*PED	Feb 06, 2009			
	5863559	Jan 26, 2016		U-72	
	5863559*PED	Jul 26, 2016			
	6020001	Mar 02, 2012		U-444	
	6020001*PED	Sep 02, 2012			
	6368627	Mar 02, 2012		U-444	
	6368627*PED	Sep 02, 2012			
<u>SUMATRIPTAN SUCCINATE; IMITREX</u>					
020132 002	4816470	Dec 28, 2006		U-72	
	4816470*PED	Jun 28, 2007			
	5037845	Aug 06, 2008		U-72	
	5037845*PED	Feb 06, 2009			
	5863559	Jan 26, 2016		U-72	
	5863559*PED	Jul 26, 2016			
	6020001	Mar 02, 2012		U-444	
	6020001*PED	Sep 02, 2012			
	6368627	Mar 02, 2012		U-444	
	6368627*PED	Sep 02, 2012			
<u>SUMATRIPTAN SUCCINATE; IMITREX</u>					
020132 003	4816470	Dec 28, 2006		U-72	
	4816470*PED	Jun 28, 2007			
	5037845	Aug 06, 2008		U-72	
	5037845*PED	Feb 06, 2009			
	5863559	Jan 26, 2016		U-72	
	5863559*PED	Jul 26, 2016			
	6020001	Mar 02, 2012		U-444	
	6020001*PED	Sep 02, 2012			
	6368627	Mar 02, 2012		U-444	
	6368627*PED	Sep 02, 2012			
<u>TACRINE HYDROCHLORIDE; COGNEX</u>					
020070 001	4816456	Sep 09, 2007		U-82	
<u>TACRINE HYDROCHLORIDE; COGNEX</u>					
020070 002	4816456	Sep 09, 2007		U-82	
<u>TACRINE HYDROCHLORIDE; COGNEX</u>					
020070 003	4816456	Sep 09, 2007		U-82	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TACRINE HYDROCHLORIDE; COGNEX</u>					
020070 004	4816456	Sep 09, 2007	U-82		
<u>TADALAFIL; CIALIS</u>					
021368 001	5859006	Jan 12, 2016	DS DP	NCE	Nov 21, 2008
	6140329	Jul 11, 2016	DP U-155		
	6821975	Nov 19, 2020	DS DP U-533		
	6821975	Nov 19, 2020	DS DP U-614		
	6943166	Apr 26, 2020	U-614		
<u>TADALAFIL; CIALIS</u>					
021368 002	5859006	Jan 12, 2016	DS DP	NCE	Nov 21, 2008
	6140329	Jul 11, 2016	DP U-155		
	6821975	Nov 19, 2020	DS DP U-614		
	6821975	Nov 19, 2020	DS DP U-533		
	6943166	Apr 26, 2020	U-614		
<u>TADALAFIL; CIALIS</u>					
021368 003	5859006	Jan 12, 2016	DS DP	NCE	Nov 21, 2008
	6140329	Jul 11, 2016	DP U-155		
	6821975	Nov 19, 2020	DS DP U-533		
	6821975	Nov 19, 2020	DS DP U-614		
	6943166	Apr 26, 2020	U-614		
<u>TAMOXIFEN CITRATE; NOLVADEX</u>					
017970 001				M-20 PED	Aug 30, 2005 Mar 01, 2006
<u>TAMOXIFEN CITRATE; SOLTAMOX</u>					
021807 001	6127425	Jun 26, 2018	DP		
<u>TAMSULOSIN HYDROCHLORIDE; FLOMAX</u>					
020579 001	4703063	Oct 27, 2009			
	4772475	Feb 27, 2006			
	4868216	Sep 19, 2006	U-181		
<u>TAZAROTENE; AVAGE</u>					
021184 003	5089509	Jun 13, 2011	U-481	NP	Sep 30, 2005
<u>TAZAROTENE; TAZORAC</u>					
020600 001	5089509	Jun 13, 2011	U-481		
	5089509	Jun 13, 2011	U-193		
	5914334	Jun 07, 2014	U-517		
	6258830	Jun 07, 2014	U-517		
<u>TAZAROTENE; TAZORAC</u>					
020600 002	5089509	Jun 13, 2011	U-481		
	5089509	Jun 13, 2011	U-193		
	5914334	Jun 07, 2014	U-517		
	6258830	Jun 07, 2014	U-517		
<u>TAZAROTENE; TAZORAC</u>					
021184 001	5089509	Jun 13, 2011	U-481		
<u>TAZAROTENE; TAZORAC</u>					
021184 002	5089509	Jun 13, 2011	U-481		
<u>TECHNETIUM TC-99M APCITIDE; ACUTECT</u>					
020887 001	5443815	Aug 22, 2012			
	5508020	Apr 16, 2013			
	5645815	Jul 08, 2014			
<u>TECHNETIUM TC-99M BICISATE KIT; NEUROLITE</u>					
020256 001	5279811	Nov 23, 2008	U-101		
	5431900	Jul 11, 2012	U-336		
<u>TECHNETIUM TC-99M EXAMETAZIME KIT; CERETEC</u>					
019829 001	4789736	Dec 06, 2005			
<u>TECHNETIUM TC-99M RED BLOOD CELL KIT; ULTRATAG</u>					
019981 001	4755375	Jul 05, 2005	U-51		
<u>TECHNETIUM TC-99M SESTAMIBI KIT; CARDIOLITE</u>					
019785 001	4885100	Sep 11, 2007			
	4894445	Jan 16, 2007	U-337		
	4988827	Jan 29, 2008			
	5324824	Jan 16, 2007			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE				
TECHNETIUM TC-99M SESTAMIBI KIT; MIRALUMA									
019785 003	4885100	Sep 11, 2007	U-337						
	4894445	Jan 16, 2007							
	4988827	Jan 29, 2008							
	5324824	Jan 16, 2007							
TECHNETIUM TC-99M TEBOROXIME KIT; CARDIOTEC									
019928 001	6056941	Jul 28, 2019	DP			I-396	Feb 28, 2006		
TECHNETIUM TC-99M TETROFOSMIN KIT; MYOVUE									
020372 001	5045302	Feb 09, 2010						I-396	Feb 28, 2006
TECHNETIUM TC-99M TETROFOSMIN KIT; MYOVUE 30ML									
020372 002									
TEGASEROD MALEATE; ZELNORM									
021200 001	5510353	Apr 26, 2013	U-466	I-435 NCE	Aug 21, 2007 Jul 24, 2007				
TEGASEROD MALEATE; ZELNORM									
021200 002	5510353	Apr 26, 2013	U-466	I-435 NCE	Aug 21, 2007 Jul 24, 2007				
TELITHROMYCIN; KETEK									
021144 001	5635485 D459798	Apr 21, 2015 Sep 24, 2015	DS DP DP	U-578	NCE	Apr 01, 2009			
TELITHROMYCIN; KETEK									
021144 002	5635485 D459798	Apr 21, 2015 Sep 24, 2015	DS DP DP	U-578	NCE	Apr 01, 2009			
TELMISARTAN; MICARDIS									
020850 001	5591762 6358986	Jan 07, 2014 Jan 10, 2020		U-3					
TELMISARTAN; MICARDIS									
020850 002	5591762 6358986	Jan 07, 2014 Jan 10, 2020		U-3					
TELMISARTAN; MICARDIS									
020850 003	5591762 6358986	Jan 07, 2014 Jan 10, 2020		U-3					
TEMAZEPAM; RESTORIL									
018163 003	5030632 5211954 5326758	Jul 09, 2008 May 18, 2010 Jul 09, 2008		U-70 U-70					
TEMOZOLOMIDE; TEMODAR									
021029 001	5260291 5260291*PED	Aug 11, 2013 Feb 11, 2014	DS DP	U-619	I-450 ODE ODE PED	Mar 15, 2008 Mar 15, 2012 Aug 11, 2006 Feb 11, 2007			
TEMOZOLOMIDE; TEMODAR									
021029 002	5260291 5260291*PED	Aug 11, 2013 Feb 11, 2014	DS DP	U-619	I-450 ODE ODE PED	Mar 15, 2008 Mar 15, 2012 Aug 11, 2006 Feb 11, 2007			
TEMOZOLOMIDE; TEMODAR									
021029 003	5260291 5260291*PED	Aug 11, 2013 Feb 11, 2014	DS DP	U-619	I-450 ODE ODE PED	Mar 15, 2008 Mar 15, 2012 Aug 11, 2006 Feb 11, 2007			
TEMOZOLOMIDE; TEMODAR									
021029 004	5260291 5260291*PED	Aug 11, 2013 Feb 11, 2014	DS DP	U-619	I-450 ODE ODE PED	Mar 15, 2008 Mar 15, 2012 Aug 11, 2006 Feb 11, 2007			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TENOFOVIR DISOPROXIL FUMARATE; VIREAD</u>					
021356 001	5922695	Jul 25, 2017	DS	U-248	NPP
	5935946	Jul 25, 2017	DS DP	U-248	NCE
	5977089	Jul 25, 2017	DS DP	U-248	
	6043230	Jul 25, 2017		U-248	
<u>TERAZOSIN HYDROCHLORIDE; HYTRIN</u>					
019057 001	5212176	Jun 29, 2010			
	5294615	Apr 29, 2013		U-165	
	5294615	Apr 29, 2013		U-3	
	5412095	Apr 29, 2013			
<u>TERAZOSIN HYDROCHLORIDE; HYTRIN</u>					
019057 002	5212176	Jun 29, 2010			
	5294615	Apr 29, 2013		U-165	
	5294615	Apr 29, 2013		U-3	
	5412095	Apr 29, 2013			
<u>TERAZOSIN HYDROCHLORIDE; HYTRIN</u>					
019057 003	5212176	Jun 29, 2010			
	5294615	Apr 29, 2013		U-165	
	5294615	Apr 29, 2013		U-3	
	5412095	Apr 29, 2013			
<u>TERAZOSIN HYDROCHLORIDE; HYTRIN</u>					
019057 004	5212176	Jun 29, 2010			
	5294615	Apr 29, 2013		U-3	
	5294615	Apr 29, 2013		U-165	
	5412095	Apr 29, 2013			
<u>TERAZOSIN HYDROCHLORIDE; HYTRIN</u>					
020347 001	5212176	Jun 29, 2010			
	5294615	Apr 29, 2013		U-165	
	5294615	Apr 29, 2013		U-3	
	5412095	Apr 29, 2013			
<u>TERAZOSIN HYDROCHLORIDE; HYTRIN</u>					
020347 002	5212176	Jun 29, 2010			
	5294615	Apr 29, 2013		U-165	
	5294615	Apr 29, 2013		U-3	
	5412095	Apr 29, 2013			
<u>TERAZOSIN HYDROCHLORIDE; HYTRIN</u>					
020347 003	5212176	Jun 29, 2010			
	5294615	Apr 29, 2013		U-3	
	5294615	Apr 29, 2013		U-165	
	5412095	Apr 29, 2013			
<u>TERAZOSIN HYDROCHLORIDE; HYTRIN</u>					
020347 004	5212176	Jun 29, 2010			
	5294615	Apr 29, 2013		U-165	
	5294615	Apr 29, 2013		U-3	
	5412095	Apr 29, 2013			
<u>TERBINAFINE; LAMISIL</u>					
020846 001	4755534	Dec 30, 2006		U-445	
	5681849	Oct 28, 2014	DP		
	5856355	May 18, 2012	DP	U-504	
	5856355	May 18, 2012	DP	U-540	
	5856355	May 18, 2012	DP	U-502	
	6005001	May 18, 2012	DP	U-504	
	6005001	May 18, 2012	DP	U-540	
	6005001	May 18, 2012	DP	U-502	
	6121314	May 18, 2012	DP	U-502	
	6121314	May 18, 2012	DP	U-540	
	6121314	May 18, 2012	DP	U-504	
<u>TERBINAFINE HYDROCHLORIDE; LAMISIL</u>					
020192 001	4755534	Dec 30, 2006		U-73	
<u>TERBINAFINE HYDROCHLORIDE; LAMISIL</u>					
020539 001	4755534	Dec 30, 2006		U-73	
<u>TERBINAFINE HYDROCHLORIDE; LAMISIL</u>					
020749 001	4755534	Dec 30, 2006			
	6121314	May 18, 2012		U-502	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TERBINAFINE HYDROCHLORIDE; LAMISIL</u>					
020980 001	4755534	Dec 30, 2006	U-73		
<u>TERBINAFINE HYDROCHLORIDE; LAMISIL AT</u>					
021124 001	4755534	Dec 30, 2006	U-73		
	5681849	Oct 28, 2014			
	6121314	May 18, 2012	U-504		
<u>TERBINAFINE HYDROCHLORIDE; LAMISIL AT</u>					
021124 002	4755534	Dec 30, 2006	U-73		
	5681849	Oct 28, 2014			
	6121314	May 18, 2012	U-504		
<u>TERIPARATIDE RECOMBINANT HUMAN; FORTEO</u>					
021318 001	6770623	Dec 08, 2018	DP U-597	NP	Nov 26, 2005
<u>TESTOSTERONE; ANDRODERM</u>					
020489 001	4849224	Nov 12, 2007			
	4855294	Sep 06, 2008			
	4863970	Nov 14, 2006			
	4983395	Nov 12, 2007			
	5152997	Dec 11, 2010	U-490		
	5164190	Dec 11, 2010			
<u>TESTOSTERONE; ANDRODERM</u>					
020489 002	4849224	Nov 12, 2007			
	4855294	Sep 06, 2008			
	4863970	Nov 14, 2006			
	4983395	Nov 12, 2007			
	5152997	Dec 11, 2010	U-490		
	5164190	Dec 11, 2010			
<u>TESTOSTERONE; ANDROGEL</u>					
021015 001	6503894	Aug 30, 2020	U-490		
<u>TESTOSTERONE; STRIANT</u>					
021543 001	6248358	Aug 23, 2019	U-527	NP	Jun 19, 2006
<u>TESTOSTERONE; TESTIM</u>					
021454 001	5023252	Jun 11, 2008	U-483	NP	Oct 31, 2005
<u>TESTOSTERONE; TESTODERM</u>					
019762 001	5840327	Aug 15, 2016			
<u>TESTOSTERONE; TESTODERM</u>					
019762 002	5840327	Aug 15, 2016			
<u>TESTOSTERONE; TESTODERM TTS</u>					
020791 001	6348210	Nov 10, 2019	U-440		
<u>THALIDOMIDE; THALOMID</u>					
020785 001	6045501	Aug 28, 2018	U-371	ODE	Jul 16, 2005
	6315720	Oct 23, 2020	U-442		
	6561976	Aug 28, 2018	U-371		
	6561977	Oct 23, 2020	U-371		
	6755784	Sep 23, 2020	U-371		
	6869399	Oct 23, 2020	U-371		
	6908432	Aug 28, 2018	U-371		
<u>THALIDOMIDE; THALOMID</u>					
020785 002	6045501	Aug 28, 2018	U-371	ODE	Jul 16, 2005
	6315720	Oct 23, 2020	U-442		
	6561976	Aug 28, 2018	U-371		
	6561977	Oct 23, 2020	U-371		
	6755784	Sep 23, 2020	U-371		
	6869399	Oct 23, 2020	U-371		
	6908432	Aug 28, 2018	U-371		
<u>THALIDOMIDE; THALOMID</u>					
020785 003	6045501	Aug 28, 2018	U-371	ODE	Jul 16, 2005
	6315720	Oct 23, 2020	U-442		
	6561976	Aug 28, 2018	U-371		
	6561977	Oct 23, 2020	U-371		
	6755784	Sep 23, 2020	U-371		
	6869399	Oct 23, 2020	U-371		
	6908432	Aug 28, 2018	U-371		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>THYROTROPIN ALFA; THYROGEN</u>					
020898 001	5840566	Nov 24, 2015		ODE	Nov 30, 2005
	6365127	Nov 24, 2015	DS DP U-556		
<u>TIAGABINE HYDROCHLORIDE; GABITRIL</u>					
020646 001	5010090	Sep 30, 2011			
	5354760	Mar 24, 2012			
	5866590	Apr 29, 2016			
	5958951	Jun 10, 2017			
<u>TIAGABINE HYDROCHLORIDE; GABITRIL</u>					
020646 002	5010090	Sep 30, 2011			
	5354760	Mar 24, 2012			
	5866590	Apr 29, 2016			
	5958951	Jun 10, 2017			
<u>TIAGABINE HYDROCHLORIDE; GABITRIL</u>					
020646 003	5010090	Sep 30, 2011			
	5354760	Mar 24, 2012			
	5866590	Apr 29, 2016			
	5958951	Jun 10, 2017			
<u>TIAGABINE HYDROCHLORIDE; GABITRIL</u>					
020646 004	5010090	Sep 30, 2011			
	5354760	Mar 24, 2012			
	5866590	Apr 29, 2016			
	5958951	Jun 10, 2017			
<u>TIAGABINE HYDROCHLORIDE; GABITRIL</u>					
020646 005	5010090	Sep 30, 2011			
	5354760	Mar 24, 2012			
	5866590	Apr 29, 2016			
	5958951	Jun 10, 2017			
<u>TIGECYCLINE; TYGACIL</u>					
021821 001	5494903	Aug 13, 2012	DS DP	NCE	Jun 15, 2010
	5529990	Jun 25, 2013	U-282		
<u>TILUDRONATE DISODIUM; SKELID</u>					
020707 001	4876248	Jan 30, 2010			
	4980171	Apr 06, 2009			
<u>TIMOLOL; BETIMOL</u>					
020439 001	5231095	Jul 27, 2010			
<u>TIMOLOL; BETIMOL</u>					
020439 002	5231095	Jul 27, 2010			
<u>TIMOLOL MALEATE; TIMOLOL MALEATE</u>					
020963 001	6174524	Mar 26, 2019			
<u>TIMOLOL MALEATE; TIMOLOL MALEATE</u>					
020963 002	6174524	Mar 26, 2019			
<u>TIMOLOL MALEATE; TIMOPTIC-XE</u>					
020330 001	4861760	Sep 25, 2006			
<u>TIMOLOL MALEATE; TIMOPTIC-XE</u>					
020330 002	4861760	Sep 25, 2006			
<u>TINIDAZOLE; TINDAMAX</u>					
021618 001				NCE	May 17, 2009
				ODE	May 17, 2011
<u>TINIDAZOLE; TINDAMAX</u>					
021618 002				NCE	May 17, 2009
				ODE	May 17, 2011
<u>TINZAPARIN SODIUM; INNOHEP</u>					
020484 001				NCE	Jul 14, 2005
<u>TIOTROPIUM BROMIDE MONOHYDRATE; SPIRIVA</u>					
021395 001	5610163	Mar 11, 2014	DS DP U-566	NCE	Jan 30, 2009
	6777423	Sep 24, 2021	DS DP		
	6908928	Nov 25, 2022	DS DP U-566		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE		
TIPRANAVIR; APTIVUS							
021814 001	5852195	Dec 22, 2015	DS	NCE	Jun 22, 2010		
	6147095	Oct 29, 2019					
	6169181	May 06, 2014	DS				
	6231887	Jul 27, 2018	DP				
TIROFIBAN HYDROCHLORIDE; AGGRASTAT							
020912 001	5292756	May 14, 2012		U-230			
	5658929	Sep 27, 2010					
	5733919	Oct 23, 2016					
	5880136	Sep 27, 2010		U-254			
	5965581	Oct 23, 2016					
	5972967	Oct 23, 2016					
	5978698	Oct 08, 2017					
	6136794	Jan 29, 2019					
TIROFIBAN HYDROCHLORIDE; AGGRASTAT							
020913 001	5292756	May 14, 2012		U-230			
	5658929	Sep 27, 2010					
	5733919	Oct 23, 2016					
	5880136	Sep 27, 2010		U-254			
	5965581	Oct 23, 2016					
	5972967	Oct 23, 2016					
	5978698	Oct 08, 2017					
	6136794	Jan 29, 2019					
TIZANIDINE HYDROCHLORIDE; ZANAFLEX							
021447 001	6455557	Nov 28, 2021					
TIZANIDINE HYDROCHLORIDE; ZANAFLEX							
021447 002	6455557	Nov 28, 2021					
TIZANIDINE HYDROCHLORIDE; ZANAFLEX							
021447 003	6455557	Nov 28, 2021					
TOLCAPONE; TASMAR							
020697 001	5236952	Jan 29, 2012		U-219			
	5476875	Dec 19, 2012					
TOLCAPONE; TASMAR							
020697 002	5236952	Jan 29, 2012		U-219			
	5476875	Dec 19, 2012					
TOLTERODINE TARTRATE; DETROL							
020771 001	5382600	Mar 25, 2012					
	5382600*PED	Sep 25, 2012					
TOLTERODINE TARTRATE; DETROL							
020771 002	5382600	Mar 25, 2012					
	5382600*PED	Sep 25, 2012					
TOLTERODINE TARTRATE; DETROL LA							
021228 001	5382600	Mar 25, 2012		M-46	Oct 21, 2008		
	5382600*PED	Sep 25, 2012					
	6630162	Nov 11, 2019	DP	U-544			
	6630162*PED	May 11, 2020					
	6770295	Aug 26, 2019	DP	U-544			
	6770295*PED	Feb 26, 2020					
	6911217	Aug 26, 2019	DP	U-544			
	6911217*PED	Feb 26, 2020	DP	U-544			
TOLTERODINE TARTRATE; DETROL LA							
021228 002	5382600	Mar 25, 2012		M-46	Oct 21, 2008		
	5382600*PED	Sep 25, 2012					
	6630162	Nov 11, 2019	DP	U-544			
	6630162*PED	May 11, 2020					
	6770295	Aug 26, 2019	DP	U-544			
	6770295*PED	Feb 26, 2020					
	6911217	Aug 26, 2019	DP	U-544			
	6911217*PED	Feb 26, 2020	DP	U-544			
TOPIRAMATE; TOPAMAX							
020505 001	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008
	6503884	Oct 13, 2015			U-598	I-41	Aug 11, 2007
						ODE	Aug 28, 2008

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
TOPIRAMATE; TOPAMAX									
020505 002	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006		
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008		
	6503884	Oct 13, 2015			U-598	I-41	Aug 11, 2007		
						ODE	Aug 28, 2008		
TOPIRAMATE; TOPAMAX									
020505 003	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006		
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008		
	6503884	Oct 13, 2015				I-41	Aug 11, 2007		
						ODE	Aug 28, 2008		
TOPIRAMATE; TOPAMAX									
020505 004	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006		
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008		
	6503884	Oct 13, 2015			U-598	I-41	Aug 11, 2007		
						ODE	Aug 28, 2008		
TOPIRAMATE; TOPAMAX									
020505 005	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006		
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008		
	6503884	Oct 13, 2015			U-598	I-41	Aug 11, 2007		
						ODE	Aug 28, 2008		
TOPIRAMATE; TOPAMAX									
020505 006	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006		
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008		
	6503884	Oct 13, 2015			U-598	I-41	Aug 11, 2007		
						ODE	Aug 28, 2008		
TOPIRAMATE; TOPAMAX SPRINKLE									
020844 001	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006		
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008		
	6503884	Oct 13, 2015			U-598	I-41	Aug 11, 2007		
						ODE	Aug 28, 2008		
TOPIRAMATE; TOPAMAX SPRINKLE									
020844 002	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006		
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008		
	6503884	Oct 13, 2015			U-598	I-41	Aug 11, 2007		
						ODE	Aug 28, 2008		
TOPIRAMATE; TOPAMAX SPRINKLE									
020844 003	4513006	Sep 26, 2008	DS	DP	U-83	D-88	Dec 16, 2006		
	5998380	Oct 13, 2015			U-598	I-467	Jun 29, 2008		
	6503884	Oct 13, 2015			U-598	I-41	Aug 11, 2007		
						ODE	Aug 28, 2008		
TOPOTECAN HYDROCHLORIDE; Hycamtin									
020671 001	5004758	May 28, 2010							
	5004758*PED	Nov 28, 2010							
TOREMIFENE CITRATE; FARESTON									
020497 001	4696949	Sep 29, 2009			U-196				
TORSEMIDE; DEMADEX									
020136 001	RE34672	Aug 11, 2006							
TORSEMIDE; DEMADEX									
020136 002	RE34672	Aug 11, 2006							
TORSEMIDE; DEMADEX									
020136 003	RE34672	Aug 11, 2006							
TORSEMIDE; DEMADEX									
020136 004	RE34672	Aug 11, 2006							
TORSEMIDE; DEMADEX									
020137 002	4861786	Jul 08, 2007							
	RE34672	Aug 11, 2006							

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TRAMADOL HYDROCHLORIDE; TRAMADOL HCL</u>					
021692 001	6254887	May 10, 2014	DP	NP	Sep 08, 2008
<u>TRAMADOL HYDROCHLORIDE; TRAMADOL HCL</u>					
021692 002	6254887	May 10, 2014	DP	NP	Sep 08, 2008
<u>TRAMADOL HYDROCHLORIDE; TRAMADOL HCL</u>					
021692 003	6254887	May 10, 2014	DP	NP	Sep 08, 2008
<u>TRAMADOL HYDROCHLORIDE; TRAMADOL HCL</u>					
021693 001	5464632	Mar 22, 2013	DP		
	6106861	Dec 05, 2017	DP		
<u>TRAMADOL HYDROCHLORIDE; ULTRAM</u>					
020281 001	6339105	Oct 12, 2019		U-435	
	6339105*PED	Apr 12, 2020		U-435	
<u>TRAMADOL HYDROCHLORIDE; ULTRAM</u>					
020281 002	6339105	Oct 12, 2019		U-435	
	6339105*PED	Apr 12, 2020		U-435	
<u>TRANDOLAPRIL; MAVIK</u>					
020528 001	4933361	Jun 12, 2007			
	5744496	Apr 28, 2015		U-229	
<u>TRANDOLAPRIL; MAVIK</u>					
020528 002	4933361	Jun 12, 2007			
	5744496	Apr 28, 2015		U-229	
<u>TRANDOLAPRIL; MAVIK</u>					
020528 003	4933361	Jun 12, 2007			
	5744496	Apr 28, 2015		U-229	
<u>TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE; TARKA</u>					
020591 001	4933361	Jun 12, 2007			
	5721244	Feb 24, 2015			
<u>TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE; TARKA</u>					
020591 002	4933361	Jun 12, 2007			
	5721244	Feb 24, 2015			
<u>TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE; TARKA</u>					
020591 003	4933361	Jun 12, 2007			
	5721244	Feb 24, 2015			
<u>TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE; TARKA</u>					
020591 004	4933361	Jun 12, 2007			
	5721244	Feb 24, 2015			
<u>TRAVOPROST; TRAVATAN</u>					
021257 001	5510383	Aug 03, 2013	DP	U-383	Feb 13, 2006
	5631287	Dec 22, 2014		U-382	Mar 16, 2006
	5849792	Dec 22, 2014	DP	U-383	
	5889052	Dec 02, 2014	DP	U-383	
	6011062	Dec 22, 2014	DP		
<u>TREPROSTINIL SODIUM; REMODULIN</u>					
021272 001	5153222	Oct 06, 2009		U-455	May 21, 2007
	6765117	Oct 24, 2017	DS	ODE	May 21, 2009
<u>TREPROSTINIL SODIUM; REMODULIN</u>					
021272 002	5153222	Oct 06, 2009		U-455	May 21, 2007
	6765117	Oct 24, 2017	DS	ODE	May 21, 2009
<u>TREPROSTINIL SODIUM; REMODULIN</u>					
021272 003	5153222	Oct 06, 2009		U-455	May 21, 2007
	6765117	Oct 24, 2017	DS	ODE	May 21, 2009
<u>TREPROSTINIL SODIUM; REMODULIN</u>					
021272 004	5153222	Oct 06, 2009		U-455	May 21, 2007
	6765117	Oct 24, 2017	DS	ODE	May 21, 2009
<u>TRETINOIN; AVITA</u>					
020404 003	4971800	Nov 20, 2007		U-178	Jun 04, 2008
	5045317	Sep 03, 2008		U-179	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TRETINOIN; RENOVA</u>					
021108 001	6531141	Mar 07, 2020			
<u>TRETINOIN; RETIN-A MICRO</u>					
020475 001	4690825	Oct 04, 2005	U-134		
	5955109	Sep 21, 2016	U-134		
<u>TRETINOIN; RETIN-A MICRO</u>					
020475 002	4690825	Oct 04, 2005	U-134		
	5955109	Sep 21, 2016	U-134		
<u>TRIAMCINOLONE ACETONIDE; NASACORT</u>					
019798 001	4767612	Jan 23, 2007	U-85		
<u>TRIAMCINOLONE ACETONIDE; NASACORT AQ</u>					
020468 001	5976573	Jul 03, 2016	U-295		
	6143329	Jul 03, 2016			
<u>TRIMETHOPRIM HYDROCHLORIDE; PRIMSO</u>					
074973 001	5763449	Aug 07, 2016			
	5962461	Aug 07, 2016			
<u>TRIMETREXATE GLUCURONATE; NEUTREXIN</u>					
020326 001	4694007	May 20, 2006	U-91		
	6017922	May 18, 2018			
<u>TRIMETREXATE GLUCURONATE; NEUTREXIN</u>					
020326 002	6017922	May 18, 2018			
<u>TRIPTORELIN PAMOATE; TRELSTAR</u>					
021288 001	5225205	Jul 20, 2010			
<u>TRIPTORELIN PAMOATE; TRELSTAR DEPOT</u>					
020715 001	5225205	Jul 20, 2010			
	5776885	Jul 07, 2015			
<u>TROGLITAZONE; PRELAY</u>					
020719 001	4572912	Nov 09, 2008			
	5602133	Sep 15, 2013	U-173		
	5859037	Nov 13, 2017	U-251		
	6011049	Nov 13, 2017	U-301		
	6046202	Sep 15, 2013	U-317		
<u>TROGLITAZONE; PRELAY</u>					
020719 002	4572912	Nov 09, 2008			
	5602133	Sep 15, 2013	U-173		
	5859037	Nov 13, 2017	U-251		
	6011049	Nov 13, 2017	U-301		
	6046202	Sep 15, 2013	U-317		
<u>TROGLITAZONE; PRELAY</u>					
020719 003	4572912	Nov 09, 2008			
	5602133	Sep 15, 2013	U-173		
	5859037	Nov 13, 2017	U-251		
	6011049	Nov 13, 2017	U-301		
	6046202	Sep 15, 2013	U-317		
<u>TROGLITAZONE; REZULIN</u>					
020720 001	4572912	Nov 09, 2008			
	5602133	Sep 15, 2013	U-173		
	5859037	Nov 13, 2017	U-251		
	6011049	Nov 13, 2017	U-301		
	6046202	Sep 15, 2013	U-317		
<u>TROGLITAZONE; REZULIN</u>					
020720 002	4572912	Nov 09, 2008			
	5602133	Sep 15, 2013	U-173		
	5859037	Nov 13, 2017	U-251		
	6011049	Nov 13, 2017	U-301		
	6046202	Sep 15, 2013	U-317		
<u>TROGLITAZONE; REZULIN</u>					
020720 003	4572912	Nov 09, 2008			
	5602133	Sep 15, 2013	U-173		
	5859037	Nov 13, 2017	U-251		
	6011049	Nov 13, 2017	U-301		
	6046202	Sep 15, 2013	U-317		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TROSPIMUM CHLORIDE; SANCTURA</u>					
021595 001				NCE	May 28, 2009
<u>TROVAFLOXACIN MESYLATE; TROVAN</u>					
020759 001	5164402	Dec 18, 2011	U-282		
	5763454	Jun 15, 2015	U-282		
	6187341	Jan 20, 2019			
<u>TROVAFLOXACIN MESYLATE; TROVAN</u>					
020759 002	5164402	Dec 18, 2011	U-282		
	5763454	Jun 15, 2015	U-282		
	6187341	Jan 20, 2019			
<u>TRYPAN BLUE; VISIONBLUE</u>					
021670 001				NCE	Dec 16, 2009
<u>UNOPROSTONE ISOPROPYL; RESCULA</u>					
021214 001	5001153	Sep 19, 2008		NCE	Aug 03, 2005
	5151444	Mar 19, 2008	U-333		
	5166178	Nov 24, 2009	U-333		
	5208256	May 21, 2011	U-333		
	5212200	May 18, 2010	U-333		
	5221763	Jun 22, 2010			
<u>UREA, C-13; BREATHTEK UBT FOR H-PYLORI</u>					
020586 002	4830010	Oct 27, 2009	U-147		
	5140993	Aug 24, 2009	U-148		
<u>UREA, C-13; MERETEK UBT KIT (W/ PRANACTIN)</u>					
020586 001	4830010	Oct 27, 2009	U-147		
	5140993	Aug 24, 2009	U-148		
<u>UREA, C-14; PYTEST</u>					
020617 001	4830010	May 15, 2006	U-195		
<u>UREA, C-14; PYTEST KIT</u>					
020617 002	4830010	May 15, 2006	U-195		
<u>UROFOLLITROPIN; BRAVELLE</u>					
021289 001				I-377	Dec 19, 2005
<u>UROFOLLITROPIN; FERTINEX</u>					
019415 004	4845077	Jul 04, 2006	U-408		
	5767067	Jun 16, 2015			
<u>UROFOLLITROPIN; FERTINEX</u>					
019415 005	4845077	Jul 04, 2006	U-408		
	5767067	Jun 16, 2015			
<u>URSODIOL; URSO 250</u>					
020675 001	4859660	Nov 19, 2007		D-93	Jul 21, 2007
<u>URSODIOL; URSO FORTE</u>					
020675 002				D-93	Jul 21, 2007
<u>VALACYCLOVIR HYDROCHLORIDE; VALTREX</u>					
020487 001	4957924	Jun 23, 2009	U-530	I-403	Aug 29, 2006
	5879706	Jan 19, 2016	U-530	I-389	Apr 01, 2006
	6107302	Jan 19, 2016	U-530	I-381	Sep 09, 2005
<u>VALACYCLOVIR HYDROCHLORIDE; VALTREX</u>					
020487 002	4957924	Jun 23, 2009	U-530	I-403	Aug 29, 2006
	5879706	Jan 19, 2016	U-530	I-389	Apr 01, 2006
	6107302	Jan 19, 2016	U-530	I-381	Sep 09, 2005
<u>VALDECOXIB; BEXTRA</u>					
021341 002	5633272	Feb 13, 2015	U-462	NCE	Nov 16, 2006
<u>VALDECOXIB; BEXTRA</u>					
021341 003	5633272	Feb 13, 2015	U-462	NCE	Nov 16, 2006
<u>VALGANCICLOVIR HYDROCHLORIDE; VALCYTE</u>					
021304 001	6083953	Jul 28, 2014	U-384	I-406	Sep 12, 2006
<u>VALRUBICIN; VALSTAR PRESERVATIVE FREE</u>					
020892 001				ODE	Sep 25, 2005

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VALSARTAN; DIOVAN</u>					
020665 001	5399578	Mar 21, 2012	U-3	I-365	Aug 14, 2005
<u>VALSARTAN; DIOVAN</u>					
020665 002	5399578	Mar 21, 2012	U-3	I-365	Aug 14, 2005
<u>VALSARTAN; DIOVAN</u>					
021283 001	5399578	Mar 21, 2012		I-463	Aug 03, 2008
	5972990	Oct 26, 2016	U-692	I-365	Aug 14, 2005
	6294197	Jun 18, 2017	U-3		
<u>VALSARTAN; DIOVAN</u>					
021283 002	5399578	Mar 21, 2012		I-463	Aug 03, 2008
	5972990	Oct 26, 2016	U-692	I-365	Aug 14, 2005
	6294197	Jun 18, 2017	U-3		
<u>VALSARTAN; DIOVAN</u>					
021283 003	5399578	Mar 21, 2012		I-463	Aug 03, 2008
	5972990	Oct 26, 2016	U-692	I-365	Aug 14, 2005
	6294197	Jun 18, 2017	U-3		
<u>VALSARTAN; DIOVAN</u>					
021283 004	5399578	Mar 21, 2012		I-463	Aug 03, 2008
	5972990	Oct 26, 2016	U-692	I-365	Aug 14, 2005
	6294197	Jun 18, 2017	U-3		
<u>VARDENAFIL HYDROCHLORIDE; LEVITRA</u>					
021400 001	6362178	Oct 31, 2018	DS DP U-533	NCE	Aug 19, 2008
<u>VARDENAFIL HYDROCHLORIDE; LEVITRA</u>					
021400 002	6362178	Oct 31, 2018	DS DP U-533	NCE	Aug 19, 2008
<u>VARDENAFIL HYDROCHLORIDE; LEVITRA</u>					
021400 003	6362178	Oct 31, 2018	DS DP U-533	NCE	Aug 19, 2008
<u>VARDENAFIL HYDROCHLORIDE; LEVITRA</u>					
021400 004	6362178	Oct 31, 2018	DS DP U-533	NCE	Aug 19, 2008
<u>VENLAFAXINE HYDROCHLORIDE; EFFEXOR</u>					
020151 001	4535186	Dec 13, 2007			
	4535186*PED	Jun 13, 2008			
<u>VENLAFAXINE HYDROCHLORIDE; EFFEXOR</u>					
020151 002	4535186	Dec 13, 2007			
	4535186*PED	Jun 13, 2008			
<u>VENLAFAXINE HYDROCHLORIDE; EFFEXOR</u>					
020151 003	4535186	Dec 13, 2007			
	4535186*PED	Jun 13, 2008			
<u>VENLAFAXINE HYDROCHLORIDE; EFFEXOR</u>					
020151 004	4535186	Dec 13, 2007			
	4535186*PED	Jun 13, 2008			
<u>VENLAFAXINE HYDROCHLORIDE; EFFEXOR</u>					
020151 005	4535186	Dec 13, 2007			
	4535186*PED	Jun 13, 2008			
<u>VENLAFAXINE HYDROCHLORIDE; EFFEXOR</u>					
020151 006	4535186	Dec 13, 2007			
	4535186*PED	Jun 13, 2008			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
VENLAFAXINE HYDROCHLORIDE; EFFEXOR XR					
020699 001	4535186	Dec 13, 2007		I-261	Feb 11, 2006
	4535186*PED	Jun 13, 2008			
	5916923	Jun 28, 2013	U-398		
	5916923*PED	Dec 28, 2013	U-398		
	6274171	Mar 20, 2017			
	6274171*PED	Sep 20, 2017			
	6403120	Mar 20, 2017	U-451		
	6403120	Mar 20, 2017	U-535		
	6403120*PED	Sep 20, 2017	U-451		
	6403120*PED	Sep 20, 2017	U-535		
	6419958	Mar 20, 2017	U-535		
	6419958	Mar 20, 2017	U-459		
	6419958*PED	Sep 20, 2017	U-459		
	6419958*PED	Sep 20, 2017	U-535		
	6444708	Jun 28, 2013	U-398		
	6444708*PED	Dec 28, 2013	U-398		
VENLAFAXINE HYDROCHLORIDE; EFFEXOR XR					
020699 002	4535186	Dec 13, 2007		I-261	Feb 11, 2006
	4535186*PED	Jun 13, 2008			
	5916923	Jun 28, 2013	U-398		
	5916923*PED	Dec 28, 2013	U-398		
	6274171	Mar 20, 2017			
	6274171*PED	Sep 20, 2017			
	6403120	Mar 20, 2017	U-451		
	6403120	Mar 20, 2017	U-535		
	6403120*PED	Sep 20, 2017	U-451		
	6403120*PED	Sep 20, 2017	U-535		
	6419958	Mar 20, 2017	U-459		
	6419958	Mar 20, 2017	U-535		
	6419958*PED	Sep 20, 2017	U-535		
	6419958*PED	Sep 20, 2017	U-459		
	6444708	Jun 28, 2013	U-398		
	6444708*PED	Dec 28, 2013	U-398		
VENLAFAXINE HYDROCHLORIDE; EFFEXOR XR					
020699 003	4535186	Dec 13, 2007		I-261	Feb 11, 2006
	4535186*PED	Jun 13, 2008			
	5916923	Jun 28, 2013	U-398		
	5916923*PED	Dec 28, 2013	U-398		
	6274171	Mar 20, 2017			
	6274171*PED	Sep 20, 2017			
	6403120	Mar 20, 2017	U-451		
	6403120	Mar 20, 2017	U-535		
	6403120*PED	Sep 20, 2017	U-535		
	6403120*PED	Sep 20, 2017	U-451		
	6419958	Mar 20, 2017	U-535		
	6419958	Mar 20, 2017	U-459		
	6419958*PED	Sep 20, 2017	U-459		
	6419958*PED	Sep 20, 2017	U-535		
	6444708	Jun 28, 2013	U-398		
	6444708*PED	Dec 28, 2013	U-398		
VENLAFAXINE HYDROCHLORIDE; EFFEXOR XR					
020699 004	4535186	Dec 13, 2007		I-261	Feb 11, 2006
	4535186*PED	Jun 13, 2008			
	5916923	Jun 28, 2013	U-398		
	5916923*PED	Dec 28, 2013	U-398		
	6274171	Mar 20, 2017			
	6274171*PED	Sep 20, 2017			
	6403120	Mar 20, 2017	U-451		
	6403120	Mar 20, 2017	U-535		
	6403120*PED	Sep 20, 2017	U-451		
	6403120*PED	Sep 20, 2017	U-535		
	6419958	Mar 20, 2017	U-459		
	6419958	Mar 20, 2017	U-535		
	6419958*PED	Sep 20, 2017	U-535		
	6419958*PED	Sep 20, 2017	U-459		
	6444708	Jun 28, 2013	U-398		
	6444708*PED	Dec 28, 2013	U-398		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VERAPAMIL HYDROCHLORIDE; COVERA-HS</u>					
020552 001	4946687	Oct 02, 2007			
	5030456	Nov 07, 2008			
	5082668	Jan 21, 2009			
	5141752	Jun 27, 2006			
	5160744	Jun 27, 2011			
	5190765	Aug 14, 2007			
	5200196	Jan 22, 2008			
	5232705	Aug 31, 2010			
	5252338	Jun 27, 2011			
	5785994	Oct 22, 2009		U-315	
	6096339	Apr 04, 2017		U-365	
	6146662	Aug 14, 2007		U-366	
<u>VERAPAMIL HYDROCHLORIDE; COVERA-HS</u>					
020552 002	4946687	Oct 02, 2007			
	5030456	Nov 07, 2008			
	5082668	Jan 21, 2009			
	5141752	Jun 27, 2006			
	5160744	Jun 27, 2011			
	5190765	Aug 14, 2007			
	5200196	Jan 22, 2008			
	5232705	Aug 31, 2010			
	5252338	Jun 27, 2011			
	5785994	Oct 22, 2009		U-315	
	6096339	Apr 04, 2017		U-365	
	6146662	Aug 14, 2007		U-366	
<u>VERAPAMIL HYDROCHLORIDE; VERELAN</u>					
019614 001	4863742	Jun 19, 2007		U-3	
<u>VERAPAMIL HYDROCHLORIDE; VERELAN</u>					
019614 002	4863742	Jun 19, 2007		U-3	
<u>VERAPAMIL HYDROCHLORIDE; VERELAN</u>					
019614 003	4863742	Jun 19, 2007		U-3	
<u>VERAPAMIL HYDROCHLORIDE; VERELAN</u>					
019614 004	4863742	Jun 19, 2007			
<u>VERAPAMIL HYDROCHLORIDE; VERELAN PM</u>					
020943 001	4863742	Jun 19, 2007			
<u>VERAPAMIL HYDROCHLORIDE; VERELAN PM</u>					
020943 002	4863742	Jun 19, 2007			
<u>VERAPAMIL HYDROCHLORIDE; VERELAN PM</u>					
020943 003	4863742	Jun 19, 2007			
<u>VERTEPORFIN; VISUDYNE</u>					
021119 001	4883790	Jan 20, 2007			
	4920143	Apr 24, 2007			
	5095030	Sep 09, 2011	DS		
	5214036	May 25, 2010			
	5283255	Jan 20, 2007		U-357	
	5707608	Aug 02, 2015			
	5756541	Mar 11, 2016		U-357	
	5770619	Jan 06, 2015		U-357	
	5798349	Aug 25, 2015		U-357	
	6074666	Feb 05, 2012			
<u>VORICONAZOLE; VFEND</u>					
021266 001	5116844	Aug 11, 2009	DP	U-540	I-442 Dec 21, 2007
	5364938	Nov 15, 2011	DS		I-409 Nov 14, 2006
	5567817	May 24, 2016	DS DP	U-540	NCE May 24, 2007
	5773443	Jan 25, 2011	DS DP	U-540	
<u>VORICONAZOLE; VFEND</u>					
021266 002	5116844	Aug 11, 2009	DP	U-540	I-442 Dec 21, 2007
	5364938	Nov 15, 2011	DS		I-409 Nov 14, 2006
	5567817	May 24, 2016	DS DP	U-540	NCE May 24, 2007
	5773443	Jan 25, 2011	DS DP	U-540	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>VORICONAZOLE; VFEND</u>					
021267 001	5116844	Aug 11, 2009	DP U-540	I-442	Dec 21, 2007
	5134127	Jan 23, 2010	DP	I-409	Nov 14, 2006
	5364938	Nov 15, 2011	DS	NCE	May 24, 2007
	5376645	Jan 23, 2010	DP		
	5567817	May 24, 2016	DS DP U-540		
	5773443	Jan 25, 2011	DS DP U-540		
	6632803	Jun 02, 2018	DP		
<u>VORICONAZOLE; VFEND</u>					
021630 001	5116844	Aug 11, 2009	DP U-540	I-442	Dec 21, 2007
	5364938	Nov 15, 2011	DS	I-409	Nov 14, 2006
	5567817	May 24, 2016	DS DP U-540	NCE	May 24, 2007
	5773443	Jan 25, 2011	DS DP U-540		
<u>ZAFIRLUKAST; ACCOLATE</u>					
020547 001	4859692	Sep 26, 2010			
	5294636	Dec 11, 2011			
	5319097	Dec 11, 2011			
	5482963	Jan 09, 2013			
	5583152	Sep 26, 2010			
	5612367	Mar 18, 2014	U-189		
	6143775	Dec 11, 2011			
<u>ZAFIRLUKAST; ACCOLATE</u>					
020547 003	4859692	Sep 26, 2010			
	5294636	Dec 11, 2011			
	5319097	Dec 11, 2011			
	5482963	Jan 09, 2013			
	5583152	Sep 26, 2010			
	5612367	Mar 18, 2014	U-189		
	6143775	Dec 11, 2011			
<u>ZALCITABINE; HIVID</u>					
020199 001	4879277	Nov 07, 2006		U-65	
	5028595	Jul 02, 2008		U-65	
<u>ZALCITABINE; HIVID</u>					
020199 002	4879277	Nov 07, 2006		U-65	
	5028595	Jul 02, 2008		U-65	
<u>ZALEPLON; SONATA</u>					
020859 001	4626538	Jun 06, 2008			
<u>ZALEPLON; SONATA</u>					
020859 002	4626538	Jun 06, 2008			
<u>ZANAMIVIR; RELENZA</u>					
021036 001	5360817	Jul 26, 2013	DS DP		
	5648379	Jul 15, 2014		U-274	
	6294572	Dec 15, 2014			
<u>ZICONOTIDE; PRIALT</u>					
021060 001	5364842	Dec 30, 2011		U-55	
	5364842	Dec 30, 2011		U-48	
	5795864	Jun 27, 2015	DP		
	5859186	Dec 30, 2011		U-48	
	5859186	Dec 30, 2011		U-55	
<u>ZICONOTIDE; PRIALT</u>					
021060 002	5364842	Dec 30, 2011		U-55	
	5364842	Dec 30, 2011		U-48	
	5795864	Jun 27, 2015	DP		
	5859186	Dec 30, 2011		U-55	
	5859186	Dec 30, 2011		U-48	
<u>ZICONOTIDE; PRIALT</u>					
021060 003	5364842	Dec 30, 2011		U-55	
	5364842	Dec 30, 2011		U-48	
	5795864	Jun 27, 2015	DP		
	5859186	Dec 30, 2011		U-48	
	5859186	Dec 30, 2011		U-55	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE	
<u>ZICONOTIDE; PRIALT</u>								
021060 004	5364842	Dec	30, 2011	DP	U-55	NCE	Dec	28, 2009
	5364842	Dec	30, 2011		U-48			
	5795864	Jun	27, 2015					
	5859186	Dec	30, 2011		U-48			
	5859186	Dec	30, 2011		U-55			
<u>ZIDOVUDINE; RETROVIR</u>								
019655 001	4724232	Sep	17, 2005					
	4818538	Sep	17, 2005					
	4828838	Sep	17, 2005					
	4833130	Sep	17, 2005					
	4837208	Sep	17, 2005					
<u>ZIDOVUDINE; RETROVIR</u>								
019910 001	4724232	Sep	17, 2005					
	4818538	Sep	17, 2005					
	4833130	Sep	17, 2005					
	4837208	Sep	17, 2005					
<u>ZIDOVUDINE; RETROVIR</u>								
019951 001	4724232	Sep	17, 2005					
	4818538	Sep	17, 2005					
	4833130	Sep	17, 2005					
	4837208	Sep	17, 2005					
<u>ZIDOVUDINE; RETROVIR</u>								
020518 001	4724232	Sep	17, 2005					
	4818538	Sep	17, 2005					
	4828838	Sep	17, 2005					
	4833130	Sep	17, 2005					
	4837208	Sep	17, 2005					
<u>ZIDOVUDINE; RETROVIR</u>								
020518 002	4724232	Sep	17, 2005					
	4818538	Sep	17, 2005					
	4828838	Sep	17, 2005					
	4833130	Sep	17, 2005					
	4837208	Sep	17, 2005					
<u>ZILEUTON; ZYFLO</u>								
020471 001	4873259	Dec	10, 2010		U-168			
<u>ZILEUTON; ZYFLO</u>								
020471 003	4873259	Dec	10, 2010		U-168			
<u>ZIPRASIDONE HYDROCHLORIDE; GEODON</u>								
020825 001	4831031	Mar	02, 2007	DS DS	DP DP DP	NCE	Feb	05, 2006
	5312925	Sep	01, 2012					
	6150366	May	27, 2019					
	6245766	Dec	18, 2018					
<u>ZIPRASIDONE HYDROCHLORIDE; GEODON</u>								
020825 002	4831031	Mar	02, 2007	DS DS	DP DP DP	NCE	Feb	05, 2006
	5312925	Sep	01, 2012					
	6150366	May	27, 2019					
	6245766	Dec	18, 2018					
<u>ZIPRASIDONE HYDROCHLORIDE; GEODON</u>								
020825 003	4831031	Mar	02, 2007	DS DS	DP DP DP	NCE	Feb	05, 2006
	5312925	Sep	01, 2012					
	6150366	May	27, 2019					
	6245766	Dec	18, 2018					
<u>ZIPRASIDONE HYDROCHLORIDE; GEODON</u>								
020825 004	4831031	Mar	02, 2007	DS DS	DP DP DP	NCE	Feb	05, 2006
	5312925	Sep	01, 2012					
	6150366	May	27, 2019					
	6245766	Dec	18, 2018					
<u>ZIPRASIDONE MESYLATE; GEODON</u>								
020919 001	4831031	Mar	02, 2007			NCE	Feb	05, 2006
	6110918	Mar	26, 2017					
	6232304	Apr	01, 2017					
	6399777	Apr	01, 2017					

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnote for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE			
<u>ZOLEDRONIC ACID; ZOMETA</u>										
021223 001	4777163	Jul 24, 2007	U-53		NCE	Aug 20, 2006				
	4939130	Nov 13, 2007							ODE	Aug 20, 2008
<u>ZOLEDRONIC ACID; ZOMETA</u>										
021223 002	4777163	Jul 24, 2007	U-53		NCE	Aug 20, 2006				
	4939130	Nov 13, 2007							ODE	Aug 20, 2008
<u>ZOLMITRIPTAN; ZOMIG</u>										
020768 001	5466699	Nov 14, 2012								
	5466699*PED	May 14, 2013								
	5863935	Nov 14, 2012								
	5863935*PED	May 14, 2013								
<u>ZOLMITRIPTAN; ZOMIG</u>										
020768 002	5466699	Nov 14, 2012								
	5466699*PED	May 14, 2013								
	5863935	Nov 14, 2012								
	5863935*PED	May 14, 2013								
<u>ZOLMITRIPTAN; ZOMIG</u>										
021450 004	5466699	Nov 14, 2012	U-436	DP	NDF	Sep 26, 2006				
	5466699*PED	May 14, 2013							PED	Mar 26, 2007
	6750237	Nov 28, 2020								
	6750237*PED	May 28, 2021								
<u>ZOLMITRIPTAN; ZOMIG-ZMT</u>										
021231 001	5466699	Nov 14, 2012								
	5466699*PED	May 14, 2013								
<u>ZOLMITRIPTAN; ZOMIG-ZMT</u>										
021231 002	5466699	Nov 14, 2012								
	5466699*PED	May 14, 2013								
<u>ZOLPIDEM TARTRATE; AMBIEN</u>										
019908 001	4382938	Oct 21, 2006	U-74							
<u>ZOLPIDEM TARTRATE; AMBIEN</u>										
019908 002	4382938	Oct 21, 2006	U-74							
<u>ZOLPIDEM TARTRATE; AMBIEN CR</u>										
021774 001	4382938	Oct 21, 2006	DS	DP	U-682	NDF	Sep 02, 2008			
	6514531	Dec 01, 2019								
<u>ZOLPIDEM TARTRATE; AMBIEN CR</u>										
021774 002	4382938	Oct 21, 2006	DS	DP	U-682	NDF	Sep 02, 2008			
	6514531	Dec 01, 2019								

Footnote:

1. Patents are published upon receipt by the Orange Book Staff and may not reflect the official receipt date as described in 21 CFR 314.53(d)(5).

2. Patents submitted on FDA Form 3542 and listed after August 18, 2003 will have one to three patent codes indicating specific patent claims as submitted by the sponsor:

DS = Drug Substance claim

DP = Drug Product claim

U and number = Method of Use claim (may be multiple). Specific Method of use claims are listed at

<http://www.fda.gov/cder/orange/patex.htm>

3. Patents listed prior to August 18, 2003 are flagged with method of use claims only as applicable and submitted by the sponsor. They may not be flagged with respect to other claims which may apply.

4. *PED and PED represent pediatric exclusivity. Patents with pediatric exclusivity granted after August 18, 2003 will be indicated with *PED as was done prior to August 18, 2003. Patents with *PED added after August 18, 2003 will not contain any information relative to the patent itself other than the *PED extension. Information related specifically to the patent will be conveyed on the original patent only.

PATENT AND EXCLUSIVITY TERMS

ADB 1 of 32

PATENT & EXCLUSIVITY ABBREVIATIONS

D	NEW DOSING SCHEDULE (SEE INDIVIDUAL REFERENCES)
I	NEW INDICATION (SEE INDIVIDUAL REFERENCES)
M	MISCELLANEOUS EXCLUSIVITY CODES (SEE INDIVIDUAL REFERENCES)
NC	NEW COMBINATION
NCE	NEW CHEMICAL ENTITY
NDF	NEW DOSAGE FORM
NE	NEW ESTER OR SALT OF AN ACTIVE INGREDIENT
NP	NEW PRODUCT
NP*	NEW PRODUCT (MINT FLAVORED)
NPP	NEW PATIENT POPULATION
NR	NEW ROUTE
NS	NEW STRENGTH
ODE	ORPHAN DRUG EXCLUSIVITY
PC	PATENT CHALLENGE
PED	PEDIATRIC EXCLUSIVITY
RTO	RX TO OTC SWITCH OR OTC USE
U	PATENT USE CODE (SEE INDIVIDUAL REFERENCES)
W	EXCLUSIVITY ON THIS APPLICATION EXPIRING ON THIS DATE HAS BEEN WAIVED BY SPONSOR - SEE SECTION 1.8 OF ORANGE BOOK PREFACE WAIVED EXCLUSIVITY

EXCLUSIVITY DOSING SCHEDULE

D-1	ONCE A DAY APPLICATION
D-2	ONCE DAILY DOSING
D-3	SEVEN DAYS/SEVEN DAYS/SEVEN DAYS DOSING SCHEDULE
D-4	SEVEN DAYS/FOURTEEN DAYS DOSING SCHEDULE
D-5	TEN DAYS/ELEVEN DAYS DOSING SCHEDULE
D-6	SEVEN DAYS/NINE DAYS/FIVE DAYS DOSING SCHEDULE
D-7	BID DOSING
D-8	INTRAVENOUS, EPIDURAL AND INTRATHECAL DOSING
D-9	NARCOTIC OVERDOSE IN ADULTS
D-10	NARCOTIC OVERDOSE IN CHILDREN
D-11	POSTOPERATIVE NARCOTIC DEPRESSION IN CHILDREN
D-12	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE DUODENAL ULCER
D-13	INCREASED MAXIMUM DAILY DOSAGE RECOMMENDATION
D-14	BEDTIME DOSING OF 800MG FOR TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
D-15	SINGLE DAILY DOSE OF 25MG/37.5MG
D-16	CONTINUOUS INTRAVENOUS INFUSION
D-17	400MG EVERY 12 HOURS FOR THREE DAYS FOR UNCOMPLICATED URINARY TRACT INFECTIONS
D-18	LOWER RECOMMENDED STARTING DOSE GUIDELINES
D-19	BOLUS DOSING GUIDELINES
D-20	SINGLE 32MG DOSE
D-21	ALTERNATIVE DOSAGE OF 300MG ONCE DAILY AFTER THE EVENING MEAL
D-22	REDUCTION IN INFUSION TIME FROM 24 TO 4 HOURS FOR THE 60MG DOSE
D-23	INCREASE MAXIMUM DOSE AND VARIATIONS IN THE DOSING REGIMEN
D-24	FOR OVARIAN CANCER THE RECOMMENDED REGIMEN IS 135MG/M2 OR 175MG/M2 INTRAVENOUSLY OVER THREE HOURS EVERY THREE WEEKS
D-25	ADDITIONAL DOSAGE REGIMEN EQUAL TO HALF THE ORIGINAL DOSING REGIMEN
D-26	ONCE WEEKLY APPLICATION
D-27	BID DOSING IN PATIENTS 12 YEARS OF AGE AND OLDER FOR PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH MODERATE EMETOGENIC CANCER CHEMOTHERAPY
D-28	USE OF ISOVUE-370 IN EXCRETORY UROGRAPHY AT EQUIVALENT GRAMS OF IODINE TO THE CURRENTLY APPROVED ISOVUE-250 AND ISOVUE-300
D-29	INCREASE OF CUMULATIVE DOSE TO 0.3MMOL/KG FOR MRI OF CNS IN ADULTS
D-30	5000 IU DOSE FOR PHOPHYLAXIX AGAINST DEEP VEIN THROMBOSIS
D-31	CHANGE IN RECOMMENDED TOTAL DAILY DOSE TO 80MG (40MG BID)
D-32	REMOVAL OF THE RESTRICTIONS LIMITING TREATMENT TO TWO CONSECUTIVE WEEKS AND TO SMALL AREAS
D-33	ONCE DAILY DOSING FOR PLAQUE PSORIASIS
D-34	EVERY FOUR MONTHS DOSAGE REGIMEN
D-35	FOR A ONE WEEK DOSING OF INTERDIGITAL TINEA PEDIS
D-36	FOR A SINGLE 2MG DOSE AS AN ALTERNATIVE TO THE 1MG DOSE GIVEN TWICE DAILY

PATENT AND EXCLUSIVITY TERMS

ADB 2 of 32

EXCLUSIVITY DOSING SCHEDULE

- D-37 DOSING REGIMEN FOR ADMINISTRATION EITHER ONCE DAILY (QD) OR TWICE DAILY (BID)
- D-38 CONTINUOUS INFUSION AS AN ALTERNATE METHOD OF ADMINISTRATION
- D-39 CHANGE IN TIME TO TAKE THE DRUG PRIOR TO A MEAL TO PREVENT MEAL-INDUCED HEARTBURN SYMPTOMS FROM "...1/2 TO 1 HOUR BEFORE EATING" TO "... RIGHT BEFORE EATING OR UP TO 60MIN BEFORE CONSUMING..."
- D-40 ONCE-A-DAY DOSING REGIMEN
- D-41 DRUG MAY BE DOSED RIGHT BEFORE A MEAL OR ANY TIME UP TO 30MIN BEFORE EATING OR DRINKING FOOD AND BEVERAGES THAT WOULD BE EXPECTED TO CAUSE SYMPTOMS
- D-42 TEN DAY DOSING REGIMEN FOR TRIPLE THERAPY, PREVACID IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN, FOR THE ERADICATION OF H.PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
- D-43 INITIATION OF TREATMENT WITH 900MG/DAY BY DELETION OF THE REQUIREMENT TO TITRATE TO 900MG/DAY OVER A 3-DAY PERIOD
- D-44 IN A CLINICAL TRIAL, FEWER DISCONTINUATIONS DUE TO ADVERSE EVENTS, ESPECIALLY DIZZINESS AND VERTIGO, WERE OBSERVED WHEN TITRATING THE DOSE IN INCREMENTS OF 50MG/DAY EVERY 3 DAYS UNTIL AN EFFECTIVE DOSE (NOT EXCEEDING 400MG/DAY) WAS REACHED
- D-45 ONCE DAILY DOSING FOR MAINTENANCE ONLY
- D-46 NEW DOSING REGIMEN OF 80MG DAILY
- D-47 PREVENTION OF HEARTBURN SYMPTOMS WHEN ADMINISTERED FROM 15 MINUTES UP TO, BUT NOT INCLUDING, 1 HOUR PRIOR TO A PROVOCATIVE MEAL
- D-48 ADMINISTRATION OF CISATRICURIUM A NEUROMUSCULAR BLOCKING AGENT AT DOSES OF 3 AND 4X THE ED95 OF CISATRICURIUM FOLLOWING INDUCTION WITH THIOPENTAL
- D-49 PEDIATRIC DOSING GUIDELINES
- D-50 INFORMATION FOR USE OF CORVERT IN POST-CARDIAC SURGERY PATIENTS
- D-51 OPTIONAL STARTING DOSE OF 40MG/DAY
- D-52 ALTERNATE DOSING REGIMEN OF 1250MG TWICE DAILY
- D-53 USE IN PEDIATRIC PATIENTS FROM 1 MONTH TO 16 YEARS OF AGE
- D-54 USE OF ZYBAN FOR MAINTENANCE THERAPY. TREATMENT UP TO 6 MONTHS WAS SHOWN EFFICACIOUS
- D-55 ADDITION OF A HIGHER DOSE OF NUTROPIN FOR PUBERTAL PATIENTS (PUBERTAL DOSE LESS THAN OR EQUAL TO 0.7MG/KG/WEEK)
- D-56 ADDITION OF POSTPRANDIAL DOSING
- D-57 3-HOUR INFUSION OF TAXOL GIVEN EVERY THREE WEEKS AT A DOSE OF 175MG/M2 FOLLOWED BY CISPLATIN AT A DOSE OF 75MG/M2 FOR THE FIRST-LINE TREATMENT OF ADVANCED OVARIAN CANCER
- D-58 CHANGE IN DOSING INTERVAL TO ONCE-DAILY ADMINISTRATION
- D-59 REDUCTION OF ELEVATED LDL-C IN A NEW, HIGHER STRENGTH TABLET, 0.8MG, AND FOR EXTENSION OF THE DOSAGE RANGE TO 0.8MG DAILY
- D-60 ADDITION OF A POST-OPERATIVE DOSING REGIMEN
- D-61 ONCE WEEKLY DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-62 ONCE WEEKLY DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-63 TO ALLOW A TITRATION DOSING REGIMEN USING A 25MG DOSE
- D-64 INCREASING DOSAGE FOR NERVE BLOCK ANESTHESIA USING NAROPIN 7.5MG/ML AND FOR EXTENDING THE DURATION OF TREATMENT FOR POSTOPERATIVE ANALGESIA USING NAROPIN 2MG/ML
- D-65 CHANGE DOSING AND ADMINISTRATION TO INDICATE MAINTENANCE OF WEIGHT LOSS OVER AN 18 MONTH PERIOD THUS EXTENDING THE USE OF THIS DRUG FROM ONE TO TWO YEARS
- D-66 DOSING RECOMMENDATIONS FOR PATIENTS UNDERGOING PCI
- D-67 SHORTER TREATMENT COURSE OF THREE DAYS IN THE TREATMENT OF RECURRENT EPISODES OF GENITAL HERPES
- D-68 CHANGE OF ADMIN RATE FOR INFUSION OF AREDIA FOR TREATMENT OF MODERATE AND SEVERE HYPERCALCEMIA OF MALIGNANCY FROM 24 HOURS TO 2 HOURS UP TO BUT NOT INCLUDING 24 HOURS
- D-69 SHORTENED DOSING REGIMEN TO 5 DAYS FOR THE TREATMENT OF ACUTE EXACERBATION OF CHRONIC BRONCHITIS
- D-70 80MG ONCE DAILY DOSING REGIMEN
- D-71 EIGHT WEEK DOSING REGIMEN
- D-72 INFORMATION REGARDING INCREASED RATE OF INFUSION FOR DEPACON
- D-73 ONCE A WEEK DOSING FOR THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- D-74 ONCE A WEEK DOSING FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- D-75 INTERMITTENT DOSING REGIMEN, STARTING DAILY DOSE 14 DAYS PRIOR TO THE ANTICIPATED ONSET OF MENSTRUATION THROUGH THE FIRST FULL DAY OF MENSES AND REPEATING WITH EACH NEW CYCLE
- D-76 FOR USE ON AN "AS NEEDED" OR PRN BASIS FOR THE MANAGEMENT OF NASAL SYMPTOMS IN PATIENTS FOR WHOM THE DRUG IS INDICATED
- D-77 ADDITION OF 20MG AND 40MG DAILY AS OPTIONAL STARTING DOSES WITH 40MG INTENDED FOR PATIENTS WHO REQUIRE A LARGE REDUCTION IN LDL-C (MORE THAN 45%)
- D-78 USE OF FLEXERIL 5MG FOR THE RELIEF OF MUSCLE SPASM ASSOCIATED WITH ACUTE, PAINFUL, MUSCULOSKELETAL CONDITIONS
- D-79 NEW LOWER STARTING DOSE FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS AND/OR MODERATE TO SEVERE SYMPTOMS OF VULVAR AND VAGINAL ATROPHY ASSOCIATED W/ THE MENOPAUSE

PATENT AND EXCLUSIVITY TERMS

ADB 3 of 32

EXCLUSIVITY DOSING SCHEDULE

D-80 CHANGE OF DOSING SCHEDULE FOR LANTUS FROM ONCE DAILY AT BEDTIME TO FLEXIBLE DAILY DOSING
D-81 NEW LOWER STARTING DOSE FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
D-82 USE OF PREMARIN 0.3 MG AND 0.45 MG FOR THE PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
D-83 750 MG, ONCE DAILY FOR 5 DAYS FOR COMMUNITY ACQUIRED PNEUMONIA (CAP)
D-84 ONCE-A-DAY DOSING OF FLOXACIN OTIC FOR THE TREATMENT OF ADULTS AND PEDIATRIC PATIENTS(AGES 6 MO & OLDER) W/ OTITIS EXTERNA CAUSED BY SUSCEPTIBLE STRAINS OF E.COLI, P.AERUGINOSA AND S.AUREUS
D-85 LOWER RECOMMENDED STARTING DOSE GUIDELINES FOR TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE
D-86 FOR USE IN SELECT EXTERNAL INSULIN PUMPS
D-87 ADDITION OF ONCE-WEEKLY DOSING FOR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
D-88 NEW DOSING RANGE OF 200-400MG PER DAY IN TWO DIVIDED DOSES FOR ADULTS WITH PARTIAL SEIZURES
D-89 USE OF REYATAZ 300 MG/RITONAVIR 100 MG ONCE DAILY FOR TREATMENT IN HIV-INFECTED ANTIRETROVIRAL-EXPERIENCED PATIENTS
D-90 ADDITION OF DAYTIME ADMINISTRATION TO TREAT VULVOVAGINAL CANDIDIASIS
D-91 ALTERNATE INTERMITTENT DOSING REGIMEN
D-92 ALTERNATIVE DOSAGE OF 1000MG ONCE DAILY AT BEDTIME
D-93 ALTERNATE TWO OR THREE TIMES DAILY DOSING REGIMENS
D-94 NEW MAXIMUM DOSAGE OF 72 MG/DAY IN ADOLESCENTS 13-17 YEARS OF AGE WITH ATTENTION DEFECIT HYPERACTIVITY DISORDER (ADHD)
D-95 BROADENED INITIAL STARTING DOSE FOR HYPERTENSION FROM 50 MG TO 100 MG TO 25 MG TO 100 MG DOSE RANGE
D-96 ONCE-MONTHLY TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS WITH BONIVA (IBANDRONATE SODIUM) 150 MG TABLETS
D-97 PED CANCER PT POPULATION EXPANDED TO INCLUDE PTS 6 MOS UP TO BUT NOT INCLUDING 4 YRS AND DOSING INSTRUCTIONS TO ADMIN 30 MIN BEFPRE CHEMO WITH SECOND AND THIRD DOSES 4 & 8 HOURS AFTER FIRST DOSE
D-98 DOSING FOR PED SURGICAL PTS EXPANDED TO INCLUDE PTS 1 MONTH UP TO BUT NOT INCLUDING 2 YEARS OF AGE
D-99 ONCE DAILY ADMINISTRATION FOR THE TREATMENT OF HIV INFECTION IN THERAPY NAIVE ADULT PATIENTS
D-100 750 MG ONCE DAILY FOR FIVE DAYS FOR THE TREATMENT OF ACUTE BACTERIAL SINUSITIS
D-101 ONCE DAILY IN CHRONIC IDIOPATHIC URTICARIA FOR ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER

EXCLUSIVITY INDICATION

I-1 DYSMENORRHEA
I-2 CHOLANGIOPANCREATOGRAPHY
I-3 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
I-4 PERIPHERAL VENOGRAPHY (PHLEBOGRAPHY)
I-5 HYSTEROSALPINGOGRAPHY
I-6 TREATMENT OF JUVENILE ARTHRITIS
I-7 BIOPSY PROVEN MINIMAL CHANGE NEPHROTIC SYNDROME IN CHILDREN
I-8 ADULT INTRAVENOUS CONTRAST-ENHANCED COMPUTED TOMOGRAPHY OF THE HEAD AND BODY
I-9 PREVENTION OF POSTOPERATIVE NAUSEA AND VOMITING
I-10 PREVENTION OF POSTOPERATIVE DEEP VENOUS THROMBOSIS AND PULMONARY EMBOLISM IN TOTAL HIP REPLACEMENT SURGERY
I-11 RELIEF OF MILD TO MODERATE PAIN
I-12 TREATMENT OF CUTANEOUS CANDIDIASIS
I-13 URINARY TRACT INFECTION (UTI) PREVENTION FOR PERIODS UP TO FIVE MONTHS IN WOMEN WITH A HISTORY OF RECURRENT UTI
I-14 SEBORRHEIC DERMATITIS
I-15 PHOTOPHERESIS IN THE PALLIATIVE TREATMENT OF SKIN MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PERSONS NOT RESPONSIVE TO OTHER TREATMENT
I-16 STIMULATE THE DEVELOPMENT OF MULTIPLE FOLLICLES/OOCYTES IN OVULATORY PATIENTS PARTICIPATING IN AN IN VITRO FERTILIZATION PROGRAM
I-17 MANAGEMENT OF CONGESTIVE HEART FAILURE
I-18 ENDOSCOPIC RETROGRADE PANCREATOGRAPHY
I-19 HERNIOGRAPHY
I-20 KNEE ARTHROGRAPHY

PATENT AND EXCLUSIVITY TERMS

ADB 4 of 32

EXCLUSIVITY INDICATION

- I-21 HIGH DOSE METHOTREXATE WITH LEUCOVORIN RESCUE IN COMBINATION WITH OTHER CHEMOTHERAPEUTIC AGENTS TO DELAY RECURRENCE IN PATIENTS WITH NONMETASTATIC OSTEOSARCOMA WHO HAVE UNDERGONE SURGICAL RESECTION OR AMPUTATION FOR THE PRIMARY TUMOR
- I-22 RESCUE AFTER HIGH-DOSE METHOTREXATE THERAPY IN OSTEOSARCOMA
- I-23 SHORT-TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- I-24 TREATMENT OF RHEUMATOID ARTHRITIS
- I-25 ADULT INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY OF THE HEAD, NECK, ABDOMINAL, RENAL AND PERIPHERAL VESSELS
- I-26 TREATMENT OF LIVER FLUKES
- I-27 ADJUNCTIVE THERAPY TO DIET TO REDUCE THE RISK OF CORONARY ARTERY DISEASE
- I-28 SELECTIVE ADULT VISCERAL ARTERIOGRAPHY
- I-29 METASTATIC BREAST CANCER IN PREMENOPAUSAL WOMEN AS AN ALTERNATIVE TO OOPHORECTOMY OR OVARIAN IRRADIATION
- I-30 TREATMENT OF TINEA PEDIS
- I-31 CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS IN THE SPINE AND ASSOCIATED TISSUES
- I-32 PEDIATRIC MYELOGRAPHY
- I-33 ORAL USE OF DILUTED OMNIPAQUE INJECTION IN ADULTS FOR CONTRAST ENHANCED COMPUTED TOMOGRAPHY OF THE ABDOMEN
- I-34 ORAL USE IN ADULTS FOR PASS-THROUGH EXAMINATION OF THE GASTROINTESTINAL TRACT
- I-35 PEDIATRIC CONTRAST ENHANCEMENT OF COMPUTED TOMOGRAPHIC HEAD IMAGING
- I-36 ARTHROGRAPHY OF THE SHOULDER JOINTS IN ADULTS
- I-37 RADIOGRAPHY OF THE TEMPOROMANDIBULAR JOINT IN ADULTS
- I-38 CONTRAST ENHANCEMENT AGENT TO FACILITATE VISUALIZATION OF LESIONS OF THE CENTRAL NERVOUS SYSTEM IN CHILDREN (2 YEARS OF AGE AND OLDER)
- I-39 TREATMENT OF ACUTE MYOCARDIAL INFARCTION
- I-40 PRIMARY NOCTURNAL ENURESIS
- I-41 MIGRAINE HEADACHE PROPHYLAXIS
- I-42 HERPES ZOSTER
- I-43 HERPES SIMPLEX ENCEPHALITIS
- I-44 MAINTENANCE THERAPY IN HEALED DUODENAL ULCER PATIENTS AT DOSE OF 1 GRAM TWICE DAILY
- I-45 ACUTE TREATMENT OF VARICELLA ZOSTER VIRUS
- I-46 USE IN PEDIATRIC COMPUTED TOMOGRAPHIC HEAD AND BODY IMAGING
- I-47 TREATMENT OF PEDIATRIC PATIENTS WITH SYMPTOMATIC HUMAN IMMUNODEFICIENCY VIRUS (HIV) DISEASE
- I-48 PEDIATRIC ANGIOCARDIOGRAPHY
- I-49 TREATMENT OF TRAVELERS' DIARRHEA DUE TO SUSCEPTIBLE STRAINS OF ENTEROTOXIGENIC ESCHERICHIA COLI
- I-50 FOR USE IN WOMEN WITH AXILLARY NODE-NEGATIVE BREAST CANCER
- I-51 TREATMENT OF PRIMARY DYSMENORRHEA AND FOR THE TREATMENT OF IDIOPATHIC HEAVY MENSTRUAL BLOOD LOSS
- I-52 PEDIATRIC EXCRETORY UROGRAPHY
- I-53 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA
- I-54 RENAL CONCENTRATION CAPACITY TEST
- I-55 HYPERTENSION
- I-56 EROSIIVE GASTROESOPHAGEAL REFLUX DISEASE
- I-57 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER
- I-58 INITIAL TREATMENT OF ADVANCED OVARIAN CARCINOMA IN COMBINATION WITH OTHER APPROVED CHEMOTHERAPEUTIC AGENTS
- I-59 ENDOSCOPICALLY DIAGNOSED ESOPHAGITIS, INCLUDING EROSIIVE AND ULCERATIVE ESOPHAGITIS, AND ASSOCIATED HEARTBURN DUE TO GASTROESOPHAGEAL REFLUX DISEASE
- I-60 SINGLE APPLICATION TREATMENT OF HEAD LICE IN CHILDREN TWO MONTHS TO TWO YEARS IN AGE
- I-61 FEMALE ANDROGENETIC ALOPECIA
- I-62 PREVENTION AND TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-63 ONCE DAILY TREATMENT AS INITIAL THERAPY IN THE TREATMENT OF HYPERTENSION
- I-64 PREVENTION OF SUPRAVENTRICULAR TACHYCARDIAS
- I-65 PREVENTION OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS
- I-66 UNCOMPLICATED GONORRHEA
- I-67 TREATMENT OF ACUTE ASTHMATIC ATTACKS IN CHILDREN SIX YEARS OF AGE AND OLDER
- I-68 CENTRAL PRECOCIOUS PUBERTY
- I-69 SHORT TERM TREATMENT OF PATIENTS WITH SYMPTOMS OF GASTROESOPHAGEAL REFLUX DISEASE (GERD), AND FOR THE SHORT TERM TREATMENT OF ESOPHAGITIS DUE TO GERD INCLUDING ULCERATIVE DISEASE DIAGNOSED BY ENDOSCOPY

PATENT AND EXCLUSIVITY TERMS

ADB 5 of 32

EXCLUSIVITY INDICATION

- I-70 USE IN COMBINATION WITH 5-FLUOROURACIL TO PROLONG SURVIVAL IN THE PALLIATIVE TREATMENT OF PATIENTS WITH ADVANCED COLORECTAL CANCER
- I-71 VARICELLA INFECTIONS (CHICKENPOX)
- I-72 PREVENTION OF CMV DISEASE IN TRANSPLANT PATIENTS AT RISK FOR CMV DISEASE
- I-73 INITIATE AND MAINTAIN MONITORED ANESTHESIA CARE (MAC) SEDATION DURING DIAGNOSTIC PROCEDURES
- I-74 INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY
- I-75 TREATMENT OF ENDOSCOPICALLY DIAGNOSED EROSION ESOPHAGITIS
- I-76 PREVENTION OF OSTEOPOROSIS
- I-77 DERMAL INFECTIONS-TINEA PEDIS, TINEA CORPORIS, TINEA CRURIS DUE TO EPIDERMOPHYTON FLOCCOSUM
- I-78 CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY AND INTRAVENOUS EXCRETORY UROGRAPHY
- I-79 MANAGEMENT OF CHRONIC STABLE ANGINA AND ANGINA DUE TO CORONARY ARTERY SPASM
- I-80 DIAGNOSIS AND LOCALIZATION OF ISCHEMIA AND CORONARY HEART DISEASE
- I-81 PROPHYLAXIS IN DESIGNATED IMMUNOCOMPROMISED CONDITIONS TO REDUCE THE INCIDENCE OF OROPHARYNGEAL CANDIDIASIS
- I-82 TREATMENT OF TRAVELERS' DIARRHEA
- I-83 ANGIOCARDIOGRAPHY, CONTRAST ENHANCED COMPUTED TOMOGRAPHIC IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY IN CHILDREN
- I-84 INTRAOPERATIVE AND POSTOPERATIVE TACHYCARDIA AND/OR HYPERTENSION
- I-85 TREATMENT OF ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS
- I-86 TREATMENT OF SECONDARY CARNITINE DEFICIENCY
- I-87 RENAL IMAGING AGENT FOR USE IN CHILDREN
- I-88 MANAGEMENT OF ENDOMETRIOSIS
- I-89 EPIDURAL USE IN LABOR AND DELIVERY AS AN ANALGESIC ADJUNCT TO BUPIVACAINE
- I-90 INTENSIVE CARE UNIT SEDATION
- I-91 MONOTHERAPY USE FOR HYPERTENSION
- I-92 ADJUNCTIVE THERAPY IN THE MANAGEMENT OF HEART FAILURE
- I-93 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN CHILDREN AGES 4-11 YEARS
- I-94 USE WITH MRI IN ADULTS TO PROVIDE CONTRAST ENHANCEMENT AND FACILITATE VISUALIZATION OF LESIONS IN THE BODY [EXCLUDING THE HEART]
- I-95 TREATMENT OF LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- I-96 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA
- I-97 ORAL OR RECTAL USE IN CHILDREN FOR THE EXAMINATION OF THE GASTROINTESTINAL TRACT
- I-98 TREATMENT OF CHILDREN WHO HAVE GROWTH FAILURE ASSOCIATED WITH CHRONIC RENAL INSUFFICIENCY
- I-99 PEDIATRIC ANESTHESIA IN CHILDREN 3 YEARS AND OLDER
- I-100 TO DECREASE THE INCIDENCE OF CANDIDIASIS IN PATIENTS UNDERGOING BONE MARROW TRANSPLANTATION WHO RECEIVE CYTOTOXIC CHEMOTHERAPY AND/OR RADIATION THERAPY
- I-101 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN-DEPENDENT DIABETES MELLITUS AND RETINOPATHY
- I-102 TREATMENT OF OBSESSIVE-COMPULSIVE DISORDER
- I-103 PROPHYLAXIS AGAINST PNEUMOCYSTIS CARINII PNEUMONIA IN INDIVIDUALS WHO ARE IMMUNOCOMPROMISED AND CONSIDERED TO BE AT RISK OF DEVELOPING PNEUMOCYSTIS CARINII PNEUMONIA
- I-104 TREATMENT OF PULMONARY AND EXTRAPULMONARY ASPERGILLOSIS IN PATENTS WHO ARE INTOLERANT OF OR WHO ARE REFRACTORY TO AMPHOTERICIN B THERAPY
- I-105 TREATMENT OF METASTATIC CARCINOMA OF THE BREAST AFTER FAILURE OF FIRST-LINE OR SUBSEQUENT CHEMOTHERAPY
- I-106 TREATMENT OF ACROMEGALY
- I-107 VAGINAL CANDIDIASIS
- I-108 EXPANDED USE-FOR ICU PATIENTS UNDERGOING LONG-TERM INFUSION DURING MECHANICAL VENTILATION
- I-109 TYPHOID FEVER
- I-110 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIOTHERAPY
- I-111 TREATMENT OF PAGET'S DISEASE OF BONE
- I-112 MANAGEMENT OF MODERATE TO SEVERE PAIN
- I-113 TREATMENT OF PROSTATITIS
- I-114 USE IN CHILDREN TO VISUALIZE LESIONS WITH ABNORMAL VASCULARITY IN THE BRAIN (INTRACRANIAL LESIONS), SPINE, AND ASSOCIATED TISSUE
- I-115 USE IN MRI IN ADULTS TO VISUALIZE LESIONS IN THE HEAD AND NECK
- I-116 MAINTENANCE OF HEALING OF EROSION ESOPHAGITIS
- I-117 TO SLOW THE PROGRESSION FO CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE

PATENT AND EXCLUSIVITY TERMS

ADB 6 of 32

EXCLUSIVITY INDICATION

- I-118 PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM FOLLOWING KNEE REPLACEMENT SURGERY
- I-119 TREATMENT OF ANEMIA CAUSED BY UTERINE LEIOMYOMATA IN WOMEN WHO FAIL IRON THERAPY
- I-120 MAINTENANCE THERAPY FOR GASTRIC ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING ACUTE ULCERS
- I-121 EXPANDED PATIENT POPULATION -- USE IN ICU PATIENTS
- I-122 PSORIASIS OF THE SCALP
- I-123 RELIEF OF MILD TO MODERATE PAIN IN PATIENTS AGED 6 MONTHS AND OLDER
- I-124 LEUCOCYTE LABELED SCINTIGRAPHY AS AN ADJUNCT IN THE LOCALIZATION OF INTRA-ABDOMINAL INFECTION AND INFLAMMATORY BOWEL DISEASE
- I-125 EXPANSION OF CONSCIOUS SEDATION INDICATION TO INCLUDE SHORT THERAPEUTIC PROCEDURES
- I-126 ADJUNCT TO THALLIUM- 201 MYOCARDIAL PERFUSION IN PATIENTS UNABLE TO EXERCISE ADEQUATELY
- I-127 TREATMENT OF ACYCLOVIR-RESISTANT HERPES IN IMMUNOCOMPROMISED PATIENTS
- I-128 IN PT W/ CH DISEASE AND HYPERCHOLESTEROLEMIA: REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH; REDUCE RISK NON-FATAL MI; REDUCE RISK UNDERGOING MYOCARDIAL REVASCULARIZATION PROCEDURES; REDUCTION ELEVATED TOTAL AND LDL CHOL LEVELS...
- I-129 TREATMENT OF ALCOHOL DEPENDENCE
- I-130 MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS
- I-131 PERIPHERAL ARTERIOGRAPHY
- I-132 TREATMENT OF MANIC PHASE OF BIPOLAR DISORDER
- I-133 MANAGEMENT OF CHRONIC STABLE ANGINA
- I-134 HEART FAILURE POST MYOCARDIAL INFARCTION
- I-135 BONE METASTASES ASSOCIATED WITH MULTIPLE MYELOMA
- I-136 IDIOPATHIC CHRONIC URTICARIA
- I-137 PREVENTION OF METAL-INDUCED HEART BURN, ACID INDIGESTION, AND SOUR STOMACH WHEN TAKEN 30 MINUTES PRIOR TO CONSUMING FOOD OR BEVERAGES
- I-138 TREATMENT OF ACUTE RECURRENT GENITAL HERPES
- I-139 PALLIATIVE TREATMENT OF ADVANCED BREAST CANCER IN PRE- AND PERIMENOPAUSAL WOMEN
- I-140 PREVENTION OF CYTOMEGALOVIRUS (CMV) DISEASE IN INDIVIDUALS WITH HIV INFECTION AT RISK FOR DEVELOPING CMV DISEASE
- I-141 TREATMENT OF HEMODYNAMICALLY STABLE PATIENTS WITHIN 24 HOURS OF ACUTE MYOCARDIAL INFARCTION TO IMPROVE SURVIVAL
- I-142 LOCALIZE MYOCARDIAL ISCHEMIA(REVERSIBLE DEFECT) AND INFARCTION (NON-REVERSIBLE DEFECTS) IN EVALUATING MYOCARDIAL FUNCTION
- I-143 EPISODIC TREATMENT OF RECURRENT GENITAL HERPES IN IMMUNOCOMPETENT ADULTS
- I-144 ENHANCEMENT OF MRI OF THE ADULT BODY INTERNAL ORGANS
- I-145 0.1MMOL/KG AS A SINGLE INTRAVEOUS BOLUS FOR MRI OF THE CNS IN CHILDREN
- I-146 CONTRAST ENHANCEMENT AND FACILITATION OF VISUALIZATION OF EXTRACRANIAL HEAD AND NECK LESIONS
- I-147 PREVENTION OF GALLSTONE FORMATION IN OBESE PATIENTS EXPERIENCING RAPID WEIGHT LOSS
- I-148 TREATMENT OF ACUTE PNEUMOCYSTIC CARINI PNEUMONIA (PCP) IN HIV-INFECTED PATIENTS WHOSE ALVEOLAR-ARTERIAL OXYGEN DIFFERENCE (AaDO2) IS LESS THAN OR EQUAL TO 55 TORR
- I-149 TREATMENT OF PATIENTS WITH NON-SMALL CELL LUNG CANCER
- I-150 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER AND PANIC DISORDER
- I-151 PREVENTION OF AND PREVENTION OF FURTHER POSTOPERATIVE NAUSEA AND VOMITING IN PEDIATRIC PATIENTS RECEIVING GENERAL ANESTHESIA
- I-152 SLOWING THE PROGRESSION OF CORONARY ATHEROSCLEROSIS AND REDUCING THE RISK OF ACUTE CORONARY EVENTS
- I-153 MANAGEMENT OF SEVERE SPASTICITY [ENCOMPASES SPINAL AND CEREBRAL ORIGIN]
- I-154 PATIENT POPULATION ALTERED TO INCLUDE PEDIATRIC USE
- I-155 TREATMENT OF ONCHOMYCOSIS DUE TO DERMATOPHYTES (TINEA UNGUIUM) OF THE TOENAIL WITH OR WITHOUT FINGERNAIL INVOLVEMENT
- I-156 ADDITIONAL DATA REGARDING THE SAFE USE OF NORVASC IN PATIENTS WITH HEART FAILURE
- I-157 TREATMENT OF ACUTE UNCOMPLICATED CYSTITIS IN FEMALES
- I-158 TREATMENT OF OSTEOLYTIC BONE METASTASES OF BREAST CANCER
- I-159 FOR HYPERCHOLESTEROLEMIC PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE REDUCE THE RISK OF MYOCARDIAL INFARCTION, REVASCULARIZATION, AND DEATH DUE TO CARDIOVASCULAR CAUSES WITH NO INCREASE IN DEATH FROM NON-CARDIOVASCULAR CAUSES
- I-160 TREATMENT OF BACTERIAL CORNEAL ULCERS
- I-161 TREATMENT OF ADULT-ONSET OR CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENCY
- I-162 FOR USE IN PATIENTS 6-11 YEARS OF AGE
- I-163 TREATMENT OF PHOTOPHOBIA
- I-164 CHRONIC BACTERIAL PROSTATITIS

PATENT AND EXCLUSIVITY TERMS

ADB 7 of 32

EXCLUSIVITY INDICATION

- I-165 MANAGEMENT OF ADULTS WITH ACTIVE, CLASSIC AND DEFINITIVE RHEUMATOID ARTHRITIS WHO HAVE HAD INSUFFICIENT THERAPEUTIC RESPONSE TO OR ARE INTOLERANT OF AN ADEQUATE TRIAL OF FULL DOSES OF ONE OR MORE NON-STEROIDAL ANTI-INFLAMMATORY DRUGS
- I-166 TREATMENT OF BULIMIA
- I-167 COMPLICATED INTRA-ABDOMINAL INFECTIONS (USED IN COMBINATION WITH METRONIDAZOLE) CAUSED BY MIXED AEROBIC/ANAEROBIC PATHOGENS
- I-168 MANAGEMENT OF LOCALLY CONFINED STAGE B2-C METASTATIC CARCINOMA OF THE PROSTATE (IN COMBINATION WITH LHRH AGONISTS)
- I-169 USE IN COMBINATION WITH CORTICOSTEROIDS AS INITIAL CHEMOTHERAPY FOR THE TREATMENT OF PATIENTS WITH PAIN RELATED TO ADVANCED HORMONE-REFRACTORY PROSTATE CANCER
- I-170 PROPHYLACTIC USE DURING HEAD LICE EPIDEMICS
- I-171 RELIEF OF SYMPTOMS OF THE COMMON COLD
- I-172 TREATMENT OF INITIAL EPISODE OF GENITAL HERPES
- I-173 PREOPERATIVELY FOR THE PREVENTION OF INFECTION IN TRANSRECTAL PROSTATE BIOPSY
- I-174 PELVIC INFLAMMATORY DISEASE
- I-175 TREATMENT OF TINEA CORPORIS AND TINEA CRURIS
- I-176 TREATMENT OF POSTOPERATIVE INFLAMMATION IN PATIENTS WHO HAVE UNDERGONE CATARACT EXTRACTION
- I-177 TX OF MODERATE ACNE VULGARIS IN FEMALES, GREATER OR EQUAL TO 15YRS OF AGE, WHO HAVE NO KNOWN CONTRAINDICATIONS TO ORAL CONTRACEPTIVE THERAPY, DESIRE CONTRACEPTION, HAVE ACHIEVED MENARCHE AND ARE UNRESPONSIVE TO TOPICAL ANTI-ACNE MEDICATIONS
- I-178 TREATMENT OF ONCHOMYCOSIS OF THE FINGERNAIL WITHOUT CONCOMITANT ONCHOMYCOSIS OF THE TOENAIL WITH A PULSE DOSING REGIMEN
- I-179 NOSOCOMIAL PNEUMONIA-MILD TO MODERATE AND SEVERE CAUSED BY HAEMOPHILUS INFLUENZAE OR KLEBSIELLA PNEUMONIAE
- I-180 TREATMENT OF PLANTAR TINEA PEDIS (MOCCASIN TYPE)
- I-181 TREATMENT OF PATIENTS WITH COMPLEX PARTIAL SEIZURES WITH AND WITHOUT SECONDARY GENERALIZATION
- I-182 TREATMENT OF GROWTH FAILURE ASSOCIATED WITH TURNER SYNDROME
- I-183 MAINTENANCE THERAPY IN THE MANAGEMENT OF MILD TO MODERATE ASTHMA IN PEDIATRIC PATIENTS AGES 6-11
- I-184 TREATMENT OF PANIC DISORDER AT A RECOMMENDED DOSE RANGE OF 1 TO 2MG/DAY (MAXIMUM OF 4MG)
- I-185 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- I-186 TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR CAUSED BY OR PRESUMED TO BE CAUSED BY PITYROSPORUM ORBICULARE (ALSO KNOWN AS MALASSEZIA FURFUR OR M. ORBICULARE)
- I-187 PREVENTION OF FRACTURES IN THE TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-188 TREATMENT OF ACUTE SINUSITIS AND ACUTE EXACERBATION OF CHRONIC SINUSITIS
- I-189 TREATMENT OF ACUTE OTITIS MEDIA IN PEDIATRIC PATIENTS
- I-190 PLANAR IMAGING AS A SECOND LINE DIAGNOSTIC DRUG AFTER MAMMOGRAPHY TO ASSIST IN THE EVALUATION OF BREAST LESIONS IN PATIENTS WITH AN ABNORMAL MAMMOGRAM OR A PALPABLE BREAST MASS
- I-191 ENDOMETRIAL THINNING AGENT PRIOR TO ENDOMETRIAL ABLATION FOR DYSFUNCTIONAL UTERINE BLEEDING
- I-192 THE PREVENTION OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING ABDOMINAL SURGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS AND A NEW DOSAGE REGIMEN, 40MG ONCE DAILY, FOR THIS INDICATION
- I-193 TREATMENT OF PANIC DISORDER IN A RECOMMENDED DOSE RANGE OF 50 TO 200MG/DAY
- I-194 CONGESTIVE HEART FAILURE
- I-195 FOR USE OF LANSOPRAZOLE IN COMBINATION WITH CLARITHROMYCIN AND AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF DUODENAL ULCER
- I-196 ACUTE TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- I-197 MAINTENANCE OF HEALING OF DUODENAL ULCER
- I-198 FOR THE USE OF LANSOPRAZOLE IN COMBINATION WITH AMOXICILLIN FOR THE ERADICATION OF HELICOBACTER PYLORI IN PATIENTS WITH ACTIVE DUODENAL ULCER DISEASE OR A ONE-YEAR HISTORY OF A DUODENAL ULCER
- I-199 MONOTHERAPY AND COMBINATION THERAPY WITH SULFONYL UREAS IN THE TREATMENT OF TYPE II DIABETES
- I-200 TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR
- I-201 EMPIRICAL THERAPY FOR FEBRILE NEUTROPENIC PATIENTS
- I-202 SECOND-LINE TREATMENT OF AIDS-RELATED KAPOSI'S SARCOMA
- I-203 MAINTENANCE OF REMISSION OF ULCERATIVE COLITIS
- I-204 USE IN PEDIATRIC PATIENTS BETWEEN THE AGES OF 6 AND 11 FOR THE TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- I-205 INITIAL ANTICONSULSANT TREATMENT OF STATUS EPILEPTICUS
- I-206 TREATMENT OF EDEMA ASSOCIATED WITH CHRONIC RENAL FAILURE

PATENT AND EXCLUSIVITY TERMS

ADB 8 of 32

EXCLUSIVITY INDICATION

- I-207 FOR THE SUPPRESSION OF RECURRENT EPISODES OF GENITAL HERPES IN IMMUNOCOMPETENT ADULTS
- I-208 TREATMENT OF OBSESSIVE COMPULSIVE DISORDER IN THE PEDIATRIC POPULATION
- I-209 PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA (PSVT)
- I-210 TO SLOW THE PROGRESSION OF CORONARY ATHEROSCLEROSIS IN PATIENTS WITH CORONARY HEART DISEASE AS PART OF A TREATMENT STRATEGY TO LOWER TOTAL AND LDL CHOLESTEROL TO TARGET LEVELS
- I-211 FOR USE IN PEDIATRIC POPULATION
- I-212 TREATMENT OF SYMPTOMS OF DRY MOUTH IN PATIENTS WITH SJOGREN'S SYNDROME
- I-213 TEMPORARY RELIEF OF PAIN AND PHOTOPHOBIA IN PATIENTS UNDERGOING CORNEAL REFRACTIVE SURGERY
- I-214 TREATMENT OF OSTEOPOROSIS
- I-215 PRE-PROCEDURAL APPLICATION TO ADULT MALE GENITAL SKIN PRIOR TO SITE-SPECIFIC SUBCUTANEOUS INFILTRATION WITH LIDOCAINE FOR THE REMOVAL OF GENITAL WARTS
- I-216 FOR THE LONG-TERM TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- I-217 PREVENTION (DURING AND FOLLOWING HOSPITALIZATION) OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
- I-218 USE OF LIPITOR AS AN ADJUNCTIVE THERAPY TO DIET FOR THE TREATMENT OF PATIENTS WITH ELEVATED SERUM TRIGLYCERIDE LEVELS (FREDERICKSON TYPE IV)
- I-219 USE OF LIPITOR BY PATIENTS WITH PRIMARY DYSBETALIPOPROTEINEMIA (FREDERICKSON TYPE III) WHO DO NOT RESPOND ADEQUATELY TO DIET
- I-220 TREATMENT OF EPISODIC- HEARTBURN, ACID INDIGESTION AND SOUR STOMACH
- I-221 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA (BPH) IN MEN WITH AN ENLARGED PROSTATE TO IMPROVE SYMPTOMS, REDUCE THE RISK OF ACUTE URINARY RETENTION AND REDUCE THE RISK OF THE NEED OF SURGERY
- I-222 PREVENTION OF ISCHEMIC COMPLICATIONS OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION, WHEN CONCURRENTLY ADMINISTERED WITH ASPIRIN
- I-223 USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH ALLERGIC AND NONALLERGIC-PERENNIAL RHINITIS IN CHILDREN AGE 6-11 YEARS
- I-224 FOR THE USE IN PEDIATRIC PATIENTS 4 TO 11 YEARS OF AGE FOR THE MANAGEMENT OF THE NASAL SYMPTOMS OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS
- I-225 USE IN PATIENTS WITH PREVIOUS MI AND NORMAL CHOLESTEROL LEVELS, TO REDUCE RISK OF RECURRENT MI, MYOCARDIAL REVASCULARIZATION, AND CEREBROVASCULAR DISEASE EVENTS
- I-226 FIRST-LINE THERAPY FOR THE TREATMENT OF ADVANCED CARCINOMA OF THE OVARY IN COMBINATION WITH CISPLATIN
- I-227 SHORT-TERM TREATMENT OF SYMPTOMATIC GASTROESOPHAGEAL REFLUX DISEASE (GERD)
- I-228 PREVENTION OF MEAL INDUCED HEARTBURN AT A DOSE OF 75MG TAKEN 30-60MIN PRIOR TO A MEAL
- I-229 PRILOSEC (OMEPRazole), AMOXICILLIN, AND CLARITHROMYCIN FOR THE ERADICATION OF H. PYLORI IN PATIENTS WITH DUODENAL ULCER DISEASE
- I-230 IN COMBINATION WITH CIS-PLATIN, FOR THE FIRST LINE TREATMENT OF NON-SMALL CELL LUNG CANCER IN PATIENTS WHO ARE NOT CANDIDATES FOR POTENTIALLY CURATIVE SURGERY AND/OR RADIATION
- I-231 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR CHEMOTHERAPY
- I-232 TREATMENT OF RECURRENT MUCOCUTANEOUS HERPES SIMPLEX INFECTIONS IN HIV-AFFECTED PATIENTS AT A DOSE OF 500MG TWICE DAILY
- I-233 PROPHYLACTIC USE TO REDUCE PERIOPERATIVE BLOOD LOSS AND THE NEED FOR BLOOD TRANSFUSION IN PATIENTS UNDERGOING CARDIOPULMONARY BYPASS IN THE COURSE OF CORONARY ARTERY BYPASS GRAFT SURGERY
- I-234 FOR USE IN COMBINATION WITH CISPLATIN FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH INOPERABLE LOCALLY ADVANCED (STAGE IIIA OR IIIB) OR METASTATIC (STAGE IV) NON-SMALL CELL LUNG CANCER
- I-235 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 12 YEARS OF AGE AND OLDER
- I-236 PREVENTION OF EXERCISE-INDUCED BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-237 MAINTENANCE TREATMENT OF ASTHMA AND PREVENTION OF BRONCHOSPASM IN PATIENTS 4 YEARS OF AGE AND OLDER
- I-238 ADJUNCTIVE TREATMENT OF LENNOX-GASTAUT SYNDROME IN PEDIATRIC AND ADULT PATIENTS
- I-239 TREATMENT OF PATIENTS WITH HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA
- I-240 MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM AND RESULTANT METABOLIC BONE DISEASE IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL FAILURE (CCR 15 TO 55ML/MIN) NOT YET ON DIALYSIS
- I-241 USE IN PHOTODYNAMIC THERAPY (PDT) FOR REDUCTION OF OBSTRUCTION AND PALLIATION OF SYMPTOMS IN PATIENTS WITH COMPLETELY OR PARTIALLY OBSTRUCTING ENDOBRONCHIAL NONSMALL CELL LUNG CANCER
- I-242 TREATMENT OF MODERATE TO SEVERE VASOMOTOR SYMPTOMS ASSOCIATED WITH THE MENOPAUSE AND IN THE TREATMENT OF VULVAR AND VAGINAL ATROPHY IN WOMEN WITH AN INTACT UTERUS

PATENT AND EXCLUSIVITY TERMS

ADB 9 of 32

EXCLUSIVITY INDICATION

- I-243 USE IN THE SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH THE COMMON COLD IN CHILDREN AGE 5 TO 11 YEARS
- I-244 REDUCE THE INCIDENCE OF BREAST CANCER IN WOMEN AT HIGH RISK FOR BREAST CANCER
- I-245 TREATMENT OF ACUTE SINUSITIS
- I-246 TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS
- I-247 USE IN CONVERSION TO MONOTHERAPY IN ADULTS WITH PARTIAL SEIZURES WHO ARE RECEIVING TREATMENT WITH A SINGLE ENZYME-INDUCING ANTIEPILEPTIC DRUG
- I-248 INPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITH/WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIM AND OUTPATIENT TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMIN WITH WARFARIN SODIUM
- I-249 TREATMENT OF CHRONIC HEPATITIS C IN PATIENTS WITH COMPENSATED LIVER DISEASE PREVIOUSLY UNTREATED WITH ALPHA INTERFERON THERAPY
- I-250 PRIMARY PREVENTION OF CORONARY HEART DISEASE IN PATIENTS WITHOUT SYMPTOMATIC CARDIOVASCULAR DISEASE WHO HAVE AVERAGE TO MODERATELY ELEVATED TOTAL-C AND LDL-C AND BELOW AVERAGE HDL-C
- I-251 TREATMENT OF GENERALIZED ANXIETY DISORDER
- I-252 NEW COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS METFORMIN
- I-253 COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS INSULIN
- I-254 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS (LOSS OF BONE MASS)
- I-255 PREVENTION OF PNEUMOCYSTIS CARINII PNEUMONIA (PCP)
- I-256 USE IN TREATMENT OF SMALL CELL LUNG CANCER SENSITIVE DISEASE AFTER FAILURE OF FIRST-LINE CHEMOTHERAPY
- I-257 TREATMENT OF CHRONIC HEPATITIS B ASSOCIATED WITH EVIDENCE OF HEPATITIS B VIRAL REPLICATION AND ACTIVE LIVER INFLAMMATION
- I-258 FOR PERENNIAL NONALLERGIC RHINITIS FOR AGES 4 AND ABOVE
- I-259 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
- I-260 EXPANDED PEDIATRIC USE IN CHILDREN YOUNGER THAN ONE MONTH OF AGE TO BIRTH (WITH A GESTATIONAL AGE OF 37 WEEKS OR GREATER)
- I-261 TREATMENT OF SOCIAL ANXIETY DISORDER
- I-262 TREATMENT OR PREVENTION OF BRONCHOSPASM WITH REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE AND FOR THE PREVENTION OF EXERCISE INDUCED BRONCHOSPASM IN CHILDREN AGES 4-12
- I-263 TREATMENT OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION FOR THE PREVENTION OF ISCHEMIC COMPLICATIONS IN PATIENTS ON CONCURRENT ASPIRIN THERAPY
- I-264 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIATION, INCLUDING TOTAL BODY IRRADIATION (TBI) AND FRACTIONATED ABDOMINAL RADIATION
- I-265 TREATMENT OF ATOPIC DERMATITIS IN PEDIATRIC PATIENTS 6 YEARS AND OLDER
- I-266 USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN PEDIATRIC PATIENTS AGES 2-16 YEARS WITH PARTIAL ONSET SEIZURES
- I-267 USE IN PEDIATRIC PATIENTS 3 MONTHS OLD AND OLDER - FOR CORTICOSTEROID-RESPONSIVE DERMATOSES
- I-268 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 7-11 YEARS OF AGE
- I-269 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HIGHLY EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING CISPLATIN
- I-270 ADJUVANT TREATMENT OF NODE-POSITIVE BREAST CANCER ADMINISTERED SEQUENTIALLY TO STANDARD DOXORUBICIN-CONTAINING COMBINATION CHEMOTHERAPY
- I-271 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- I-272 TREATMENT OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN MEN AND WOMEN RECEIVING GLUCOCORTICOID IN A DAILY DOSE EQUIVALENT TO 7.5MG OR GREATER OF PREDNISONE AND WHO HAVE LOW BONE MINERAL DENSITY
- I-273 ADJUNCT TO DIET TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NON FAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
- I-274 USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES
- I-275 USE IN COMBINATION WITH METFORMIN AND SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES
- I-276 USE OF REZULIN IN COMBINATION WITH METFORMIN AND SULFONYLUREAS IN PATIENTS WITH TYPE 2 DIABETES
- I-277 TREATMENT OF TYPE III HYPERLIPOPROTEINEMIA
- I-278 TREATMENT OF PATIENTS WITH ISOLATED HYPERTRIGLYCERIDEMIA (FREDERICKSON TYPE IV)
- I-279 TREATMENT OF POST-TRAUMATIC STRESS DISORDER
- I-280 USE OF CARNITOR INJECTION FOR THE PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS
- I-281 INCREASING HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NONFAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)

PATENT AND EXCLUSIVITY TERMS

ADB 10 of 32

EXCLUSIVITY INDICATION

- I-282 TREATMENT OF PATIENTS WITH LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER AFTER FAILURE OF PRIOR PLATINUM-BASED CHEMOTHERAPY
- I-283 TO REDUCE THE INCIDENCE OF MODERATE TO SEVERE XEROSTOMIA IN PATIENTS UNDERGOING POST-OPERATIVE RADIATION TREATMENT FOR HEAD AND NECK CANCER, WHERE THE RADIATION PORT INCLUDES A SUBSTANTIAL PORTION OF THE PAROTID GLANDS
- I-284 TO REDUCE THE NUMBER OF ADENOMATOUS COLORECTAL POLYPS IN FAMILIAL ADENOMATOUS POLYPOSIS PATIENTS AS AN ADJUNCT TO USUAL CARE
- I-285 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN ADULTS AND CHILDREN 3 YEARS OF AGE AND OLDER
- I-286 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE III
- I-287 USE OF PRAVASTATIN IN PATIENTS WITH EVIDENT CORONARY HEART DISEASE TO REDUCE THE RISK OF TOTAL MORTALITY BY REDUCING CORONARY DEATH
- I-288 CHANGES IN SEVERAL SECTIONS OF THE INSERT TO INCORPORATE STATEMENTS CONCERNING THE USE OF HIGH DOSES OF LISINAPRIL TO REDUCE THE RISK OF THE COMBINED OUTCOMES OF MORTALITY AND HOSPITALIZATION IN PATIENTS WITH CONGESTIVE HEART FAILURE
- I-289 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA IN PATIENTS WITH TYPE 2 DIABETES MELLITUS WHEN DIET AND EXERCISE WITH EITHER SINGLE AGENT DOES NOT ACHIEVE ADEQUATE GLYCEMIC CONTROL
- I-290 PREVENTION OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
- I-291 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- I-292 TREATMENT OF POSTMENOPAUSAL OSTEOPOROSIS
- I-293 TREATMENT OF CORTICOSTEROID-INDUCED OSTEOPOROSIS
- I-294 TREATMENT OF UNCOMPLICATED ACUTE ILLNESS DUE TO INFLUENZA A AND B IN PEDIATRIC PATIENTS 7 YEARS AND OLDER WHO HAVE BEEN SYMPTOMATIC FOR NO MORE THAN 2 DAYS
- I-295 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR WOMEN WITH AN INTACT UTERUS
- I-296 LONG-TERM INTRAVENOUS TREATMENT OF PULMONARY HYPERTENSION ASSOCIATED WITH THE SCLERODERMA SPECTRUM OF DISEASE IN NYHA CLASS III AND CLASS IV PATIENTS WHO DO NOT RESPOND TO CONVENTIONAL THERAPY
- I-297 SHORT-TERM TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-298 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IIA AND IIB HYPERLIPOPROTEINEMIA
- I-299 USE OF CAMPTOSAR AS A COMPONENT OF FIRST-LINE THERAPY IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVARIN FOR PATIENTS WITH METASTATIC CARCINOMA OF THE COLON OR RECTUM
- I-300 PROPHYLAXIS FOR ASTHMA IN CHILDREN 2-5 YEARS OF AGE
- I-301 TREATMENT OF SIGNS AND SYMPTOMS OF ALLERGIC CONJUNCTIVITIS
- I-302 TREATMENT OF PEDIATRIC PATIENTS WITH PRADER-WILLI SYNDROME
- I-303 INCREASING HDL-CHOLESTEROL IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA AND MIXED DYSLIPIDEMIAS
- I-304 TREATMENT OF PATIENTS WITH FREDERICKSON TYPE IV
- I-305 TREATMENT OF LEVOFLOXACIN SUSCEPTIBLE STRAINS OF PENICILLIN-RESISTANT STREPTOCOCCUS PNEUMONIAE IN PATIENTS WITH COMMUNITY ACQUIRED PNEUMONIA
- I-306 INDUCTION OF SPERMATOGENESIS IN MEN WITH PRIMARY AND SECONDARY HYPOGONADOTROPIC HYPOGONADISM IN WHOM THE CAUSE OF INFERTILITY IS NOT DUE TO PRIMARY TESTICULAR FAILURE
- I-307 NEW COMBINATION USE OF METFORMIN AND INSULIN IN TYPE 2 DIABETES
- I-308 TREATMENT OF PEDIATRIC PATIENTS WITH POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS WHO RESPONDED INADEQUATELY TO SALICYLATES OR OTHER NSAIDS
- I-309 INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- I-310 REDUCTION IN RISK OF MYOCARDIAL INFARCTION, STROKE, AND DEATH FROM CARDIOVASCULAR CAUSES
- I-311 ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS AGE 3 TO 12 YEARS
- I-312 FIRST LINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER
- I-313 EXTENSION OF INDICATION TO PROVIDE FOR MAINTENANCE OF RESPONSE
- I-314 TOPICAL ANESTHETIC FOR SUPERFICIAL MINOR SURGERY OF GENITAL MUCOUS MEMBRANES AND AS AN ADJUNCT FOR LOCAL INFILTRATION ANESTHESIA IN GENITAL MUCOUS MEMBRANES
- I-315 THROMBOPROPHYLAXIS OF DEEP VEIN THROMBOSIS, WHICH MAY LEAD TO PULMONARY EMBOLISM, IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS
- I-316 TREATMENT OF NSAID-ASSOCIATED GASTRIC ULCER PATIENTS WHO CONTINUE NSAID USE AND REDUCING RISK OF NSAID-ASSOCIATED GASTRIC ULCERS IN PATIENTS WITH HISTORY OF DOCUMENTED GASTRIC ULCER WHO REQUIRE USE OF AN NSAID
- I-317 PROPHYLAXIS OF INFLUENZA IN ADULTS AND ADOLESCENTS 13 YEARS AND OLDER
- I-318 FIRSTLINE TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE OR HORMONE RECEPTOR UNKNOWN LOCALLY ADVANCED OR METASTATIC BREAST CANCER
- I-319 USE FOR SUSPECTED OR CONFIRMED METHANOL POISONING, EITHER ALONE OR IN COMBINATION WITH HEMODIALYSIS
- I-320 TREATMENT OF TYPE 2 DIABETES IN PEDIATRIC PATIENTS (AGES 10-16 YEARS)

PATENT AND EXCLUSIVITY TERMS

ADB 11 of 32

EXCLUSIVITY INDICATION

- I-321 JUVENILE RHEUMATOID ARTHRITIS
- I-322 USE OF DIPRIVAN IN PATIENTS 3 MONTHS TO 16 YEARS
- I-323 COLORECTAL CANCER
- I-324 REDUCING NEUROLOGIC DISABILITY AND/OR FREQUENCY OF CLINICAL RELAPSES IN PATIENTS WITH SECONDARY (CHRONIC) PROGRESSIVE, PROGRESSIVE RELAPSING, OR WORSENING RELAPSING-REMITTING MULTIPLE SCLEROSIS
- I-325 PREVENTION OF RELAPSE AND RECURRENCE OF DEPRESSION
- I-326 GENERALIZED ANXIETY DISORDER
- I-327 SYMPTOMATIC RELIEF OF RHINORRHEA ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS IN PATIENTS 5 YEARS AND OLDER
- I-328 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 5-6 YEARS OF AGE
- I-329 UNCOMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS
- I-330 MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND CONTROL OF DAYTIME AND NIGHTTIME HEARTBURN SYSTEMS IN PATIENTS WITH GERD
- I-331 TREATMENT OF MODERATE ACNE VULGARIS
- I-332 EMPIRIC THERAPY IN FEBRILE NEUTROPENIC PATIENTS WITH SUSPECTED FUNGAL INFECTIONS (EFTN)
- I-333 TOPICAL TREATMENT OF TINEA (PITYRIASIS) VERSICOLOR DUE TO MALASSEZIA FURFUR (FORMERLY PITYROSPORUM ORBICULARE)
- I-334 LONG-TERM TREATMENT OF GROWTH FAILURE IN CHILDREN BORN SMALL FOR GESTATIONAL AGE WHO FAIL TO MANIFEST CATCH-UP GROWTH BY TWO YEARS OF AGE
- I-335 ADJUNCTIVE THERAPY IN PATIENTS TWO YEARS AND OLDER WITH SEIZURES ASSOCIATED WITH LENNOX-GASTAUT SYNDROME
- I-336 EXPANSION OF INDICATION TO INCLUDE THE TREATMENT OF PATIENTS WITH PREDOMINATELY CLASSIC SUBFOVEAL CHOROIDAL NEOVASCULARIZATION DUE TO PATHOLOGIC MYOPIA OR PRESUMED OCULAR HISTOPLASMOSIS
- I-337 PATHOLOGICAL HYPERSECRETION ASSOCIATED WITH ZOLLINGER-ELLISON SYNDROME
- I-338 MANAGEMENT OF ACUTE PAIN IN ADULTS AND TREATMENT OF PRIMARY DYSMENORRHEA
- I-339 TREATMENT OF HEPATITIS B IN PEDIATRIC PATIENTS AGES 2-17 YEARS
- I-340 ATOPIC DERMATITIS IN PEDIATRIC PATIENTS AGES 2-5
- I-341 BREAST CANCER COMBINATION THERAPY
- I-342 USE OF FORADIL FOR LONG-TERM, TWICE DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHO-CONSTRICTION IN PATIENTS WITH COPD INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- I-343 USE OF COREG FOR SEVERE HEART FAILURE
- I-344 ACNE VULGARIS
- I-345 TREATMENT OF POSTTRAUMATIC STRESS DISORDER
- I-346 TREATMENT OF SYMPTOMATIC GASTRO ESOPHAGEAL REFLUX DISEASE (GERD)
- I-347 TREATMENT OR PREVENTION OF BRONCHOSPASM IN CHILDREN 6 YEARS OF AGE AND OLDER WITH OBSTRUCTIVE AIRWAY DISEASE
- I-348 LONG-TERM, TWICE-DAILY (MORNING AND EVENING) ADMINISTRATION IN THE MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD (INCLUDING EMPHYSEMA AND CHRONIC BRONCHITIS)
- I-349 ACUTE CORONARY SYNDROME
- I-350 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN ADOLESCENT BOYS AND GIRLS AT LEAST ONE YEAR POSTMENARCHAL, AGES 10 TO 17 YEARS, WITH A RECOMMENDED DOSING RANGE OF 10 TO 40MG ONCE DAILY
- I-351 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS FOR ALL STRENGTHS
- I-352 ANTICOAGULANT IN PATIENTS WITH OR AT RISK FOR HEPARIN-INDUCED THROMBOCYTOPENIA UNDERGOING PERCUTANEOUS CORONARY INTERVENTIONS (PCI)
- I-353 TREATMENT OF SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS
- I-354 MANAGEMENT OF POST HERPETIC NEURALGIA
- I-355 PREMENSTRUAL DYSPHORIC DISORDER
- I-356 TREATMENT OF PATHOLOGICAL HYPERSECRETORY CONDITIONS, INCLUDING ZOLLINGER-ELLISON SYNDROME
- I-357 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS
- I-358 TREATMENT OF PANIC DISORDER
- I-359 TREATMENT OF VULVAR AND VAGINAL ATROPHY ASSOCIATED WITH THE MENOPAUSE
- I-360 TREATMENT OF NASAL SYMPTOMS OF SEASONAL AND PERENNIAL RHINITIS IN CHILDREN AGES TWO UP TO AGE THREE
- I-361 TREATMENT OF MULTIPLE MYELOMA AND DOCUMENTED BONE METASTASES FROM SOLID TUMORS, IN CONJUNCTION WITH STANDARD ANTINEOPLASTIC THERAPY. PROSTATE CANCER SHOULD HAVE PROGRESSED AFTER TREATMENT WITH AT LEAST ONE HORMONAL THERAPY
- I-362 TREATMENT OF PANIC DISORDER, WITH OR WITHOUT AGORAPHOBIA
- I-363 ADJUVANT TREATMENT OF POST MENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER
- I-364 TREATMENT OF COMMUNITY-ACQUIRED PNEUMONIA IN ADULTS

PATENT AND EXCLUSIVITY TERMS

ADB 12 of 32

EXCLUSIVITY INDICATION

- I-365 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV) IN PATIENTS WHO ARE INTOLERANT TO AN ACE INHIBITOR
- I-366 PREVENTION OF RELAPSE FOLLOWING LONG-TERM TREATMENT OF MAJOR DEPRESSIVE DISORDER
- I-367 COMBINATION THERAPY WITH THIAZOLIDINEDIONE TO LOWER BLOOD GLUCOSE IN PTS WHOSE HYPERGLYCEMIA CANNOT BE CONTROLLED BY DIET/EXERCISE PLUS MONOTHERAPY WITH ANY OF THE FOLLOWING AGENTS: METFORMIN, SULFONYLUREAS, REPAGLINIDE, OR THIAZOLIDINEDIONES
- I-368 USE OF GLUCOVANCE WITH A THIAZOLIDINEDIONE WHEN GLYCEMIC CONTROL IS NOT OBTAINED WITH GLUCOVANCE ALONE
- I-369 PREVENTION AND TREATMENT OF POSTOPERATIVE NAUSEA AND VOMITING
- I-370 TREATMENT OF HETEROZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA IN CHILDREN, AGES 8-13 YEARS, WITH RECOMMENDED DOSE OF 20MG ONCE DAILY AND IN ADOLESCENTS, AGES 14-18 WITH A RECOMMENDED DOSE OF 40MG ONCE DAILY
- I-371 HELICOBACTER PYLORI ERADICATION TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
- I-372 NOSOCOMIAL PNEUMONIA
- I-373 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY
- I-374 SHORT TERM TOPICAL TREATMENT OF MILD TO MODERATE PLAQUE-TYPE PSORIASIS OF NON SCALP REGIONS
- I-375 FIRST LINE THERAPY FOR THE REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION
- I-376 TREATMENT OF NEWLY DIAGNOSED ADULT PATIENTS WITH PHILADELPHIA CHROMOSOME POSITIVE CHRONIC MYELOID LEUKEMIA (CML)
- I-377 USE OF BRAVELLE FOR MULTIPLE FOLLICULAR DEVELOPMENT (CONTROLLED OVARIAN STIMULATION) DURING ASSISTED REPRODUCTIVE TECHNOLOGY CYCLES IN PATIENTS WHO HAVE PREVIOUSLY RECEIVED PITUITARY SUPPRESSION
- I-378 RELIEF OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 2 YEARS OF AGE AND OLDER
- I-379 USE TAXOTERE IN COMBINATION WITH CISPLATIN FOR THE TREATMENT OF PATIENTS WITH UNRESECTABLE, LOCALLY ADVANCED OR METASTATIC NON-SMALL CELL LUNG CANCER WHO HAVE NOT PREVIOUSLY RECEIVED CHEMOTHERAPY FOR THIS CONDITION
- I-380 TO TREAT PATIENTS WITH SCHIZOPHRENIA OR SCHIZOAFFECTIVE DISORDER AT RISK FOR EMERGENT SUICIDAL BEHAVIOR
- I-381 TREATMENT OF COLD SORES (HERPES LABIALIS) IN ADULT AND ADOLESCENT PATIENTS 12 YEARS OF AGE AND OLDER
- I-382 FOR NEWLY-DIAGNOSED HIGH GRADE MALIGNANT GLIOMA PATIENTS AS AN ADJUNCT TO SURGERY AND RADIATION
- I-383 TREATMENT OF TYPE 2 DIABETIC NEPHROPATHY
- I-384 USE IN COMBINATION WITH INSULIN FOR THE TREATMENT OF PATIENTS WITH TYPE 2 DIABETES MELLITUS
- I-385 MODIFICATION OF THE INDICATION FOR COMMUNITY ACQUIRED PNEUMONIA TO ADD "INCLUDING PENICILLIN-RESISTANT STRAINS, MIC PENICILLIN >= 2MCG/ML TO STREPTOCOCCUS PNEUMONIAE
- I-386 RAPAMUNE (SIROLIMUS) WITHIN AN IMMUNOSUPPRESSIVE REGIMEN THAT WOULD ALLOW FOR THE WITHDRAWAL OF CYCLOSPORINE 2 TO 4 MONTHS AFTER RENAL TRANSPLANTATION IN PATIENTS CONSIDERED AT LOW TO MODERATE IMMUNOLOGIC RISK FOR RENAL TRANSPLANT REJECTION
- I-387 ADJUNCTIVE THERAPY OF PARTIAL SEIZURES IN PEDIATRIC PATIENTS GREATER THAN OR EQUAL TO 2 YEARS OF AGE
- I-388 TREATMENT OF PATIENTS WITH LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- I-389 SUPPRESSION OF RECURRENT GENITAL HERPES IN HIV-INFECTED INDIVIDUALS
- I-390 USE IN PTS AT HIGH RISK CORONARY EVENTS DUE TO EXISTING CORONARY HEART DISEASE, DIABETES, PERIPHERAL VESSEL DISEASE, STROKE HISTORY, OTHER CV DISEASE TO REDUCE RISK TOTAL MORTALITY BY REDUCING CORONARY DEATH, REDUCE NONFATAL MI & STROKE.....
- I-391 ABLATION OF HIGH-GRADE DYSPLASIA IN BARRETT'S ESOPHAGUS PATIENTS WHO DO NOT UNDERGO ESOPHAGECTOMY
- I-392 TX OF PED PATIENTS W/PH+ CHRONIC PHASE CML DISEASE RECUR AFTER STEM CELL TRNSPLT OR RESIST TO INTERFERON ALPHA THERAPY. NO CONTROLLED TRIALS DEMONSTRATING A CLINICAL BENEFIT SUCH AS IMPROVE IN DISEASE RELATED SX OR INCREASED SURVIVAL
- I-393 CHRONIC BACTERIAL PROSTATITIS
- I-394 USE IN PATIENTS WITH CORONARY HEART DISEASE TO REDUCE THE RISK OF UNDERGOING CORONARY REVASCULARIZATION PROCEDURES
- I-395 TO IMPROVE PHYSICAL FUNCTION
- I-396 EXPANDED INDICATION TO INCLUDE THE ASSESSMENT OF VENTRICULAR FUNCTION IN SUBJECTS BEING EVALUATED FOR HEART DISEASE AND/OR VENTRICULAR FUNCTION
- I-397 EXTENDED PROPHYLAXIS IN PATIENTS UNDERGOING HIP FRACTURE SURGERY
- I-398 IDIOPATHIC SHORT STATURE
- I-399 TREATMENT OF CANDIDEMIA AND THE FOLLOWING CANDIDA INFECTIONS: INTRA-ABDOMINAL ABSCESES, PERITONITIS AND PLEURAL SPACE INFECTIONS
- I-400 USE OF OLANZAPINE IN COMBINATION WITH LITHIUM OR VALPROATE FOR THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR DISORDER

PATENT AND EXCLUSIVITY TERMS

ADB 13 of 32

EXCLUSIVITY INDICATION

- I-401 LONGER-TERM EFFICACY OF ARIPIPRAZOLE IN THE TREATMENT OF SCHIZOPHRENIA
- I-402 DIABETIC FOOT INFECTIONS WITHOUT CONCOMITANT OSTEOMYELITIS
- I-403 USE OF VALTREX IN COMBINATION WITH SAFER SEX PRACTICES FOR THE REDUCTION OF THE RISK OF TRANSMISSION OF GENITAL HERPES DURING SUPPRESSIVE THERAPY OF THE SOURCE PARTNER IN A HETEROSEXUAL COUPLE
- I-404 MAINTENANCE TREATMENT OF BIPOLAR I DISORDER TO DELAY THE TIME TO OCCURRENCE OF MOOD EPISODES (DEPRESSION, MANIA, HYPOMANIA, MIXED EPISODES) IN PATIENTS TREATED FOR ACUTE MOOD EPISODES WITH STANDARD THERAPY
- I-405 TREATMENT OF PREMENSTRUAL DYSPHORIC DISORDER (PMDD) USING AN INTERMITTENT DOSING REGIMEN
- I-406 PREVENTION OF CYTOMEGALOVIRUS DISEASE IN KIDNEY, HEART, AND KIDNEY-PANCREAS TRANSPLANT PATIENTS AT HIGH RISK (DONOR CMV SEROPOSITIVE/RECIPIENT CMV SERONEGATIVE)
- I-407 IMPROVE SURVIVAL OF STABLE PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (EJECTION FRACTION<=40%) AND CLINICAL EVIDENCE OF CONGESTIVE HEART FAILURE AFTER AN ACUTE MYOCARDIAL INFARCTION
- I-408 STIMULATION OF PANCREATIC SECRETIONS TO FACILITATE THE IDENTIFICATION OF THE AMPULLA OF VATER AND ACCESSORY PAPILLA DURING ENDOSCOPIC RETROGRADE CHOLANGIO-PANCREATOGRAPHY (ERCP)
- I-409 ESOPHAGEAL CANDIDIASIS
- I-410 USE OF ADVAIR DISKUS 250/50 FOR CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD) ASSOCIATED WITH CHRONIC BRONCHITIS
- I-411 EXPANDED INDICATION FOR USE IN COMBINATION WITH ANTIDIABETIC DRUGS IN THE THIAZOLIDINEDIONE CLASS
- I-412 MONOTHERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-413 ADJUNCTIVE THERAPY FOR THE SHORT TERM TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-414 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM (PE) IN MEDICAL PATIENTS WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLICATIONS DUE TO SEVERELY RESTRICTED MOBILITY DURING ACUTE ILLNESS
- I-415 SEVERE HYPERTENSION WHEN THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY
- I-416 THE USE OF CIPRO XR FOR COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS
- I-417 USE IN THE LONG TERM TREATMENT OF BIPOLAR I DISORDER
- I-418 ADJUNCTIVE THERAPY W/ MOOD STABILIZERS (LITHIUM OR DIVALPROEX) IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDERS
- I-419 MONOTHERAPY IN THE TREATMENT OF ACUTE MANIC EPISODES ASSOCIATED WITH BIPOLAR I DISORDER
- I-420 TOPICAL TREATMENT OF CLINICALLY TYPICAL, NONHYPERKERATOTIC, NONHYPERTROPHIC ACTINIC KERATOSES ON THE FACE OR SCALP IN IMMUNOCOMPETENT ADULTS
- I-421 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND PYELONEPHRITIS DUE TO E.COLI FOR PED PATIENTS (1-17) NOT AS FIRST CHOICE
- I-422 INDICATED FOR THE IN-HOSPITAL SHORT-TERM (UP TO 4 HOURS) REDUCTION IN BLOOD PRESSURE IN PEDIATRIC PATIENTS
- I-423 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS
- I-424 MANAGEMENT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS WITH MODERATE TO SEVERE CHRONIC RENAL INSUFFICIENCY NOT YET ON DIALYSIS
- I-425 FLOXATIN IN COMBINATION WITH INFUSIONAL 5-FLUOROURACIL (5-FU) AND LEUCOVORIN (LV) FOR THE TREATMENT OF PATIENTS PREVIOUSLY UNTREATED FOR ADVANCED COLORECTAL CANCER
- I-426 TREATMENT OF ACUTE PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
- I-427 TREATMENT OF ACUTE DEEP VEIN THROMBOSIS WITHOUT PULMONARY EMBOLISM WHEN ADMINISTERED IN CONJUNCTION WITH WARFARIN SODIUM
- I-428 FOR USE IN COMBINATION WITH PACLITAXEL FOR THE FIRST-LINE TREATMENT OF PATIENTS WITH METASTATIC BREAST CANCER AFTER FAILURE OF PRIOR ANTHRACYCLINE CONTAINING ADJUVANT CHEMOTHERAPY UNLESS ANTHRACYCLINES WERE CLINICALLY CONTRAINDICATED
- I-429 FOR USE IN COMBINATION WITH PREDNISONE FOR THE TREATMENT OF PATIENTS WITH ANDROGEN INDEPENDENT (HORMONE REFRACTORY) METASTATIC PROSTATE CANCER
- I-430 FOR USE IN THE RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS
- I-431 NOSOCOMIAL PNEUMONIA AND COMMUNITY-ACQUIRED PNEUMONIA CAUSED BY STREPTOCOCCUS PNEUMONIAE INDICATION EXPANDED TO INCLUDE MULTI-DRUG RESISTANT STRAINS
- I-432 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA CAUSED BY MULTI-DRUG RESISTANT STREPTOCOCCUS PNEUMONIAE
- I-433 TREATMENT OF BIOPSY-CONFIRMED, PRIMARY SUPERFICIAL BASAL CELL CARCINOMA IN IMMUNOCOMPETENT ADULTS, WITH A MAXIMUM TUMOR DIAMETER OF 2.0CM, LOCATED ON THE TRUNK (EXCLUDING ANOGENITAL SKIN), NECK, OR EXTREMITIES (EXCLUDING HANDS AND FEET)
- I-434 PREVENTION OF CARDIOVASCULAR DISEASE IN ADULT PATIENTS WITHOUT CLINICALLY EVIDENT HEART DISEASE, BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE TO REDUCE RISK OF MI AND RISK FOR REVASCULARIZATION PROCEDURES AND ANGINA

PATENT AND EXCLUSIVITY TERMS

ADB 14 of 32

EXCLUSIVITY INDICATION

- I-435 CHRONIC IDIOPATHIC CONSTIPATION
- I-436 FOR USE IN COMBINATION WITH DOXORUBICIN AND CYCLOPHOSPHAMIDE FOR THE ADJUVANT TREATMENT OF PATIENTS WITH OPERABLE NODE-POSITIVE BREAST CANCER
- I-437 TREATMENT OF ACUTE MANIC AND MIXED EPISODES ASSOCIATED WITH BIPOLAR DISORDER
- I-438 EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS
- I-439 USED TO TREAT ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-440 FOR THE REPLACEMENT OF ENDOGENOUS GROWTH HORMONE IN ADULTS WITH GROWTH HORMONE DEFICIENCY
- I-441 USE COMBINATION WITH INFUSIONAL 5-FU/LV FOR ADJUVANT TREATMENT STAGE III COLON CANCER PTS WHO HAVE UNDERGONE COMPLETE RESECTION PRIMARY TUMOR-BASED ON IMPROVEMENT IN DISEASE FREE SURVIVAL, NO DEMONSTRATED BENEFIT OVERALL SURVIVAL AFTER 4YRS
- I-442 USED FOR CANDIDEMIA IN NONNEUTROPENIC PATIENTS AND THE FOLLOWING CANDIDA INFECTIONS: DISSEMINATED INFECTIONS IN SKIN & INFECTIONS IN ABDOMEN, KIDNEY, BLADDER WALL, AND WOUNDS
- I-443 TREATMENT OF NASAL POLYPS IN PATIENTS 18 YEARS OF AGE AND OLDER
- I-444 USE OF PROTONIX IV FOR INJECTION AS STAND ALONE THERAPY FOR THE SHORT-TERM TREATMENT OF PATIENTS HAVING GASTROESOPHAGEAL REFLUX (GERD) WITH A HISTORY OF EROSIIVE ESOPHAGITIS
- I-445 TO IMPROVE (COMPARED TO 4.25% DEXTROSE) LONG-DWELL ULTRAFILTRATION AND CLEARANCE OF CREATININE AND UREA NITROGEN IN PATIENTS WITH HIGH AVERAGE OR GREATER TRANSPORT CHARACTERISTICS, AS DEFINED USING THE PERITONEAL EQUILIBRATION TEST (PET)
- I-446 EXTENDED ADJUVANT TREATMENT OF EARLY BREAST CANCER IN POSTMENOPAUSAL WOMEN WHO HAVE RECEIVED 5 YRS ADJUVANT TAMOXIFEN THERAPY-EFFECTIVENESS BASED ON AN ANALYSIS OF DISEASE FREE SURVIVAL IN PATIENTS TREATED FOR A MEDIAN 24 MONTHS
- I-447 USE OF COPEGUS (RIBAVIRIN) FOR TREATMENT OF CHRONIC HEPATITIS C IN ADULT PATIENTS COINFECTED WITH HIV IN COMBINATION WITH PEGASYS (PEGINTERFERON ALFA-2A)
- I-448 TREATMENT OF HEART FAILURE (NYHA CLASS II-IV AND EJECTION FRACTION <=40%) TO REDUCE THE RISK OF DEATH FROM CARDIOVASCULAR CAUSES AND TO REDUCE HOSPITALIZATIONS FOR HEART FAILURE
- I-449 TO IMPROVE WAKEFULNESS IN TWO NEW PATIENT POPULATIONS WITH EXCESSIVE SLEEPINESS: THOSE WITH OBSTRUCTIVE SLEEP APNEA/HYPOPNEA SYNDROME AND THOSE WITH SHIFT WORK SLEEP DISORDER
- I-450 TREATMENT OF PATIENTS WITH NEWLY DIAGNOSED HIGH GRADE GLIOMAS CONCOMITANTLY WITH RADIOTHERAPY AND THEN AS ADJUVANT TREATMENT
- I-451 MANAGEMENT OF ENDOMETRIOSIS ASSOCIATED PAIN
- I-452 EXPANDED INDICATION TO INCLUDE TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST 1 PRIOR THERAPY
- I-453 USE IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN WHEN DIET, EXERCISE AND BOTH AGENTS DO NOT RESULT IN ADEQUATE GLYCEMIC CONTROL (TRIPLE THERAPY)
- I-454 MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CHRON'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON FOR UP TO 3 MONTHS
- I-455 MODIFIED HEART FAILURE INDICATION TO INCLUDE TREATMENT OF HEART FAILURE IN PATIENTS WITH LEFT VENTRICULAR SYSTOLIC DYSFUNCTION (NYHA CLASS II-IV; EJECTION FRACTION LESS THAN OR EQUAL TO 40%)
- I-456 TO REDUCE CARDIOVASCULAR DEATH AND TO REDUCE HEART FAILURE HOSPITALIZATIONS. INCLUDES ADDITIONAL INFORMATION ON THE ADDED EFFECT ON THESE OUTCOMES WHEN USED WITH AN ACE INHIBITOR
- I-457 TREATMENT OF PATIENTS UNDERGOING ABDOMINAL SUREGERY WHO ARE AT RISK FOR THROMBOEMBOLIC COMPLIATIONS
- I-458 USE OF BIVALIRUDIN FOR INJECTION WITH PROVISIONAL USE OF GLYCOPROTEIN IIB/IIA INHIBITOR (GPI) AS LISTED IN THE CLINICAL TRIALS REPLACE-2 SECTION FOR USE AS AN ANTICOAGULANT IN PATIENTS UNDERGOING PERCUTANEOUS CORONARY INTERVENTION (PCI)
- I-459 NON-DIALYSIS DEPENDENT CHRONIC KIDNEY DISEASE (NDD-CKD) PATIENTS RECEIVING OR NOT RECEIVING AN ERYTHROPOIETIN
- I-460 TREATMENT OF DIARRHEA CAUSED BY CRYPTOSPORIDIUM PARVUM IN NON-HIV INFECTED PATIENTS 12 YEARS OF AGE AND OLDER
- I-461 USE AS A SINGLE AGENT FOR ADJUVANT TREATMENT IN PATIENTS WITH DUKES' C COLON CANCER WHO HAVE UNDERGONE COMPLETE RESECTION OF THE PRIMARY TUMOR WHEN TREATMENT WITH FLUOROPYRIMIDINE THERAPY ALONE IS PREFERRED
- I-462 LONG TERM TREATMENT OF IDIOPATHIC SHORT STATURE
- I-463 TREATMENT OF PATIENTS POST MYOCARDIAL INFARCTION
- I-464 TREATMENT OF MODERATE TO SEVERE PRIMARY RESTLESS LEGS SYNDROME
- I-465 PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 6 MONTHS OF AGE AND OLDER
- I-466 FOR RELIEF OF THE SIGNS AND SYMPTOMS OF ANKYLOSING SPONDYLITIS
- I-467 USE OF TOPIRAMATE AS INITIAL MONOTHERAPY IN PATIENTS 10 YEARS OF AGE AND OLDER WITH PARTIAL ONSET OR PRIMARY GENERALIZED TONIC CLONIC SEIZURES
- I-468 USE IN PATIENTS WITH STABLE CORONARY ARTERY DISEASE TO REDUSE THE RISK OF CARDIOVASCULAR MORTALITY OR NON-FATAL MYOCARDIAL INFECTION
- I-469 RELIEF OF THE SIGNS AND SYMPTOMS OF PARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS IN PATIENTS 2 YEARS OF AGE AND OLDER
- I-470 DIABETIC PERIPHERAL NEUROPATHIC PAIN

PATENT AND EXCLUSIVITY TERMS

ADB 15 of 32

EXCLUSIVITY INDICATION

- I-471 INDICATED TO REDUCE THE RISK OF MYOCARDIAL INFARCTION AND STROKE IN PATIENTS WITH TYPE 2 DIABETES AND WITHOUT CLINICALLY EVIDENT CORONARY HEART DISEASE BUT WITH MULTIPLE RISK FACTORS FOR CORONARY HEART DISEASE
- I-472 USE IN PATIENTS WITH ANGIOGRAPHICALLY DOCUMENTED CORONARY ARTERY DISEASE
- I-473 USE IN COMBINATION WITH GEMCITABINE FOR THE FIRST LINE TREATMENT OF PATIENTS WITH LOCALLY ADVANCED UNRESECTABLE OR METASTATIC PANCREATIC CANCER
- I-474 TREATMENT OF IRON DEFICIENCY ANEMIA IN PERITONEAL DIALYSIS DEPENDANT CHRONIC KIDNEY DISEASE IN PATIENTS RECIEVING AN ERYTHROPOIETIN
- I-475 PREVENTION OF NAUSEA AND VOMITTING ASSOCIATED WITH INITIAL AND REPEAT COURSES OF MODERATELY EMETOGENIC CANCER CHEMOTHERAPY
- I-476 TREATMENT OF DIABETIC FOOT INFECTIONS WITHOUT OSTEOMYELITIS
- I-477 TREATMENT OF COMPLICATED SKIN AND SKIN STRUCTURE INFECTIONS CAUSED BY METHICILLIN SUSCEPTIBLE STAPHYLOCOCCUS AUREUS, ESCHERICHIA COLI, KLEBSIELLA PNEUMONIAE, OR ENTEROBACTER CLOACAE
- I-478 FOR USE AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PARTIAL SEIZURES IN CHILDREN WITH EPILEPSY AGED 2-4 YEARS
- I-479 TREATMENT OF COMPLICATED INTRA-ABDOMINAL INFECTIONS CAUSED BY E.COLI, B. FRAGILIS, S.ANGINOSUS, S.CONSTELLATUS, E. FAECALIS, P. MIRABILIS, C. PERFRINGENS, B. THETAIOAOMICRON OR PEPTOSTREPTOCOCCUS SPECIES
- I-480 PROPHYLAXIS OF INFLUENZA FOR PATIENTS BETWEEN 1-12 YEARS OF AGE
- I-481 INDICATED FOR THE ADJUVANT TREATMENT OF POSTMENOPAUSAL WOMEN WITH HORMONE RECEPTOR POSITIVE EARLY BREAST CANCER
- I-482 TREATMENT OF ACUTE MANIC OR MIXED EPISODES ASSOCIATED WITH BIPOLAR I DISORDER WITH OR WITHOUT PSYCHOTIC FEATURES
- I-483 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS

EXCLUSIVITY MISCELLANEOUS

- M-1 INFORMATION REGARDING SUPERIORITY CLAIM OVER RANITIDINE FOR DAY AND NIGHT HEARTBURN ADDED TO CLINICAL STUDIES SECTION
- M-2 APPROVAL FOR ADDITION TO CLINICAL PHARMACOLOGY SECTION OF THE LABEL REGARDING (1) IMPROVEMENT IN BONE MINERAL DENSITY IN CHILDHOOD-ONSET ADULT GROWTH HORMONE DEFICIENT PATIENTS AND (2) INCREASES IN SERUM ALKALINE PHOSPHATASE
- M-3 ADDITION OF EFFICACY AND SAFETY INFORMATION IN WHICH FOSAMAX WAS USED CONCOMITANTLY WITH ESTROGEN ALONE OR WITH ESTROGEN PLUS PROGESTIN
- M-4 CHANGES TO PEDIATRIC USE SECTION TO PROVIDE INFORMATION REGARDING SAFETY AND EFFICACY IN PEDIATRIC PATIENTS AS YOUNG AS 2 YEARS OLD
- M-5 INFORMATION REGARDING EFFECTS IN PATIENTS WITH ASTHMA ON CONCOMITANT INHALED CORTICOSTEROIDS IN CLINICAL PHARMACOLOGY SECTION
- M-6 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH GLUCOPHAGE/GLYBURIDE COMBINATION ADDED TO CLINICAL PHARMACOLOGY AND DOSING AND ADMINISTRATION
- M-7 CLINICAL PHARMACOLOGY IN PEDIATRIC PATIENTS; DOSAGE AND ADMINISTRATION INFORMATION
- M-8 ADDITIONAL INFORMATION FOR THE USE OF SONATA CAPSULES FOR UP TO 5 WEEKS (35 NIGHTS) OF TREATMENT IN A CONTROLLED TRIAL SETTING
- M-9 ADDITION TO THE CLINICAL STUDIES SECTION OF THE LABELING OF TEXT AND TWO TABLES CONTAINING INFORMATION FOR THE PRESCRIBING PHYSICIAN ON BLOOD PRESSURE, HEART RATE, AND HEART RATE VARIABILITY
- M-10 INFORMATION REGARDING MAINTENANCE OF AN ANTIDEPRESSANT EFFECT UP TO 1 YEAR OF DOSING
- M-11 USE FOR LONG-TERM TREATMENT OF POSTTRAUMATIC STRESS DISORDER
- M-12 NEW LANGUAGE FOR PEDIATRIC USE
- M-13 INFORMATION FROM PEDIATRIC STUDIES ADDED TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION
- M-14 ADDITIONAL CLINICAL TRIAL INFORMATION ADDED TO PEDIATRIC USE SUBSECTION
- M-15 LONGER TERM EFFICACY INFORMATION FOR RISPERIDONE IN THE TREATMENT OF SCHIZOPHRENIA
- M-16 CHANGE IN WORDING OF THE PEDIATRIC SECTION OF THE PACKAGE INSERT
- M-17 INFORMATION REGARDING USE OF ULTANE IN PEDIATRIC PATIENTS WITH CONGENITAL HEART DISEASE
- M-18 INFORMATION DENOTING THE EFFICACY OF REMERON IN MAINTAINING A RESPONSE IN PATIENTS WITH MAJOR DEPRESSIVE DISORDER (MDD)
- M-19 INFORMATION REGARDING USE IN PEDIATRIC PATIENTS TWO YEARS OF AGE AND OLDER
- M-20 LABELING REVISIONS RELATED TO MCCUNE ALBRIGHT SYNDROME
- M-21 COMPARISON DATA ON THE ANTIHYPERTENSIVE EFFECTS OF ATACAND AND COZAAR
- M-22 CHANGE IN TIME TO ONSET OF ACTION
- M-23 INFORMATION REGARDING ELIMINATION ADDED TO CLINICAL PHARMACOLOGY, STUDY RESULTS IN PATIENTS WITH HEPATIC AND RENAL IMPAIRMENT
- M-24 INFORMATION ON RESULTS OF A LONG TERM LONGITUDINAL GROWTH STUDY AND PEDIATRIC SAFETY INFORMATION

PATENT AND EXCLUSIVITY TERMS

ADB 16 of 32

EXCLUSIVITY MISCELLANEOUS

- M-25 ADDITIONAL SAFETY & PK INFORMATION IN CHILDREN 6 MONTHS TO LESS THAN 6 YEARS OF AGE ADDED TO PKG INSERT
- M-26 INCORPORATION OF INFORMATION CONTAINED IN THE PEG-INTRON PACKAGE INSERT INTO THE REBETOL PACKAGE INSERT AND MEDGUIDE-PEG-INTRON WAS APPROVED FOR USE IN COMBINATION WITH REBETOL FOR TREATMENT OF CHRONIC HEPATITIS C VIRUS INFECTION ON 8/7/01
- M-27 INFORMATION DESCRIBING ASPIRIN ENDOSCOPY STUDY AND THE MAXIMUM RECOMMENDED DOSE FOR PATIENTS WITH MODERATE HEPATIC INSUFFICIENCY
- M-28 INFORMATION FROM A STUDY IN PEDIATRIC PATIENTS IN ASSOCIATION WITH A NEUROLOGICAL CONDITION
- M-29 LABELING CHANGES TO PROVIDE INFORMATION IN THE MANAGEMENT OF OBESITY IN ADOLESCENTS AGED 12 TO 16 YEARS
- M-30 CHANGES TO CLINICAL PHARMACOLOGY, PRECAUTIONS, AND DOSAGE AND ADMINISTRATION SECTIONS OF LABELING CONCERNING USE OF LOTENSIN IN PEDIATRIC PATIENTS WITH HYPERTENSION
- M-31 INFORMATION FOR USE IN PEDIATRIC PATIENTS WITH CHRONIC KIDNEY DISEASE STAGE 5 (END-STAGE RENAL DISEASE)
- M-32 ADDITIONAL LANGUAGE TO CLINICAL PHARMACOLOGY AND CLINICAL STUDIES
- M-33 INFORMATION FOR USE OF ADVAIR DISKUS 100/50 IN CHILDREN 4 TO 11 YEARS OF AGE WITH ASTHMA
- M-34 EXPANDED INFORMATION TO PEDIATRIC USE SUBSECTION OF LABELING IN RESPONSE TO PEDIATRIC WRITTEN REQUEST
- M-35 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES DONE WITH ACTOS IN COMBINATION WITH METFORMIN, A SULFONYLUREA, OR INSULIN ADDED TO CLINICAL PHARMACOLOGY
- M-36 ADDITION OF INFORMATION TO CLINICAL STUDIES REGARDING PREVENTION OF CARDIOVASCULAR DISEASE
- M-37 INFORMATION ADDED TO THE LABELING THAT DETAILS INFORMATION RELATIVE TO STUDIES DONE IN PEDIATRIC POPULATIONS IN THE CLINICAL PHARMACOLOGY AND PEDIATRIC USE SUBSECTIONS
- M-38 SAFETY AND IOP-LOWERING EFFECTS OF TRUSOPT HAVE BEEN DEMONSTRATED IN PEDIATRIC PATIENTS IN A 3 MONTH, MULTI-CENTER DOUBLE MASKED ACTIVE-TREATMENT-CONTROLLED TRIAL
- M-39 FOR LABELING CHANGES BASED ON RESULTS OF THE SPD422-202 CLINICAL STUDY REPORT (CSR) SUBMITTED IN RESPONSE TO THE WRITTEN REQUEST
- M-40 ADDITIONAL INFORMATION REGARDING CLINICAL STUDIES PERFORMED IN PEDIATRIC PATIENTS WITH LEUKEMIA ADDED TO PRECAUTIONS
- M-41 REVISION TO THE PEDIATRIC USE PRECAUTIONS OF THE PRESCRIBING INFORMATION TO INCORPORATE THE RESULTS FROM THE CAPPS-169 STUDY ENTITLED "THE EFFECT OF ORTHO TRICYCLEN ON BONE MINERAL DENSITY IN PEDIATRIC SUBJECTS WITH ANOREXIA NERVOSA"
- M-42 ADDITION OF A GERIATRIC USE SUBSECTION TO THE PRECAUTIONS SECTION OF THE PACKAGE INSERT AND GERIATRIC DOSING INFORMATION
- M-43 INCLUSION OF RESULTS OF STUDY-"PLACEBO-CONTROLLED STUDY TO EVALUATE SAFETY AND PILOT EFFICACY OF ILOPROST AS ADD ON THERAPY WITH BOSENTAN IN SUBJECTS WITH PULMONARY ARTERIAL HYPERTENSION
- M-44 CLINICAL INFORMATION ADDED TO THE PEDIATRIC USE SUBSECTION OF PRECAUTIONS REGARDING THE USE OF NOVOLOG IN ADOLESCENTS WITH TYPE I DIABETES AGE 6 TO 18
- M-45 INFORMATION ADDED TO CLINICAL TRIALS SECTION OF LABELING - EFFECTS OF HUMATROPE TREATMENT IN ADULTS WITH GROWTH HORMONE DEFICIENCY
- M-46 PROVISION OF RESULTS OF STUDY AND PROPOSED REVISIONS TO PACKAGE INSERT SEE SECTION ON CARDIAC ELECTROPHYSIOLOGY
- M-47 PROVIDES FOR USE OF ANTARA WITHOUT REGARD TO MEALS
- M-48 CHANGES TO THE LABELING DESCRIBING THE RESULTS OF A STUDY OF THE USE OF NOVOLOG MIX 70/30 WITH ORAL ANTIDIABETIC AGENTS IN PATIENTS WITH TYPE 2 DIABETES
- M-49 CLINICAL DATA ADDED TO THE CLINICAL PHARMACOLOGY SECTION REGARDING EFFECT OF SINGULAIR ON GROWTH RATES IN PREPUBERTAL CHILDREN
- M-50 NEW INFO TO THE CLINICAL STUDIES, ADULT GROWTH HORMONE DEFICIENCY (GHD) SUBSECTION OF THE NUTROPIN AQ PACKAGE INSERT DESCRIBING THE EFFECTS OF SOMATROPIN ON VISCERAL ADIPOSE TISSUE IN THE ADULT GROWTH HORMONE DEFICIENT PATIENT POPULATION
- M-51 INFORMATION ADDED TO LABELING REGARDING OSTEOGENESIS IMPERFECTA STUDY

PATENT USE

- U-1 PREVENTION OF PREGNANCY
- U-2 TREATMENT OR PROPHYLAXIS OF ANGINA PECTORIS AND ARRHYTHMIA
- U-3 TREATMENT OF HYPERTENSION
- U-4 PROVIDING PREVENTION AND TREATMENT OF EMESIS AND NAUSEA IN MAMMALS
- U-5 METHOD OF PRODUCING BRONCHODILATION
- U-6 METHOD OF PRODUCING SYMPATHOMIMETIC EFFECTS
- U-7 INCREASING CARDIAC CONTRACTILITY
- U-8 ACUTE MYOCARDIAL INFARCTION
- U-9 CONTROL OF EMESIS ASSOCIATED WITH ANY CANCER CHEMOTHERAPY AGENT

PATENT AND EXCLUSIVITY TERMS

ADB 17 of 32

PATENT USE

- U-10 DIAGNOSTIC METHOD FOR DISTINGUISHING BETWEEN HYPOTHALMIC MALFUNCTIONS OR LESIONS IN HUMANS
- U-11 TREATMENT OR PROPHYLAXIS OF CARDIAC DISORDERS
- U-12 METHOD OF TREATING [A] HUMAN SUFFERING FROM DEPRESSION
- U-13 A METHOD FOR TREATING ANXIETY IN A HUMAN SUBJECT IN NEED OF SUCH TREATMENT
- U-14 ADJUNCTIVE THERAPY FOR THE PREVENTION AND TREATMENT OF HYPERAMMONEMIA IN THE CHRONIC MANAGEMENT OF PATIENTS WITH UREA CYCLE ENZYMOPATHIES
- U-15 METHOD OF LOWERING INTRAOCULAR PRESSURE
- U-16 USE IN LUNG SCANNING PROCEDURES
- U-17 TREATMENT OF VENTRICULAR AND SUPRAVENTRICULAR ARRHYTHMIAS
- U-18 METHOD FOR INHIBITING GASTRIC SECRETION IN MAMMALS
- U-19 TREATMENT OF INFLAMMATION
- U-20 A PROCESS FOR TREATING A PATIENT SUFFERING FROM PARKINSON'S SYNDROME AND IN NEED OF TREATMENT
- U-21 TREATMENT OF HUMANS SUFFERING UNDESIRE UROTOXIC SIDE EFFECTS CAUSED BY CYTOSTATICALLY ACTIVE ALKYLATING AGENTS
- U-22 METHOD OF COMBATTING PATHOLOGICALLY REDUCED CEREBRAL FUNCTIONS AND PERFORMANCE WEAKNESSES, CEREBRAL INSUFFICIENCY AND DISORDERS IN CEREBRAL CIRCULATION AND METABOLISM IN WARM-BLOODED ANIMALS
- U-23 METHOD FOR TREATING PROSTATIC CARCINOMA COMPRISING ADMINISTERING FLUTAMIDE
- U-24 METHOD FOR TREATING PROSTATE ADENOCARCINOMA COMPRISING ADMINISTERING AN ANTIANDROGEN INCLUDING FLUTAMIDE AND AN LHRH AGONIST
- U-25 REDUCING CHOLESTEROL IN CHOLELITHIASIS PATIENTS
- U-26 REDUCING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF
- U-27 DISSOLVING CHOLESTEROL GALLSTONES AND/OR FRAGMENTS THEREOF
- U-28 CEREBRAL, CORONARY, PERIPHERAL, VISCERAL AND RENAL ARTERIOGRAPHY, AORTOGRAPHY AND LEFT VENTRICULOGRAPHY
- U-29 CT IMAGING OF THE HEAD AND BODY, AND INTRAVENOUS EXCRETORY UROGRAPHY
- U-30 CEREBRAL ANGIOGRAPHY, AND VENOGRAPHY
- U-31 INTRA-ARTERIAL DIGITAL SUBTRACTION ANGIOGRAPHY
- U-32 PALLIATIVE TREATMENT OF PATIENTS WITH OVARIAN CARCINOMA RECURRENT AFTER PRIOR CHEMOTHERAPY, INCLUDING PATIENTS WHO HAVE BEEN PREVIOUSLY TREATED WITH CISPLATIN
- U-33 TREATING VIRAL INFECTIONS IN A MAMMAL
- U-34 TREATING VIRAL INFECTIONS IN A WARM-BLOODED ANIMAL
- U-35 TREATING CYTOMEGALOVIRUS IN A HUMAN WITH AN INJECTABLE COMPOSITION
- U-36 METHODS OF TREATING BACTERIAL ILLNESSES
- U-37 METHOD OF TREATING GASTROINTESTINAL DISEASE
- U-38 TREATMENT OF PAROXYSMAL SUPRAVENTRICULAR TACHYCARDIA
- U-39 ANGINA PECTORIS
- U-40 METHOD OF TREATMENT OF BURNS
- U-41 METHOD OF TREATING CARDIAC ARRHYTHMIAS
- U-42 ADJUVANT TREATMENT IN COMBINATION WITH FLUOROURACIL AFTER SURGICAL RESECTION IN PATIENTS WITH DUKES' STAGE C COLON CANCER
- U-43 MANAGEMENT OF CHRONIC PAIN IN PATIENTS REQUIRING OPIOID ANALGESIA
- U-44 RELIEF OF NAUSEA AND VOMITING
- U-45 TREATMENT OF INFLAMMATION AND ANALGESIA
- U-46 TREATMENT OF PANIC DISORDER
- U-47 STIMULATION OF THE RELEASE OF GROWTH HORMONE
- U-48 ANALGESIA
- U-49 SYMPTOMATIC CANCER-RELATED HYPERCALCEMIA
- U-50 USE IN TREATING INFLAMMATORY DERMATOSES
- U-51 BLOOD POOL IMAGING, INCLUDING CARDIAC FIRST PASS AND GATED EQUILIBRIUM IMAGING AND FOR DETECTION OF SITES OF GASTROINTESTINAL BLEEDING
- U-52 TREATMENT OF ADULT AND PEDIATRIC PATIENTS(OVER SIX MONTHS OF AGE) WITH ADVANCED HIV INFECTION
- U-53 HYPERCALCEMIA OF MALIGNANCY
- U-54 REVERSAL AGENT OR ANTAGONIST OF NONDEPOLARIZING NEUROMUSCULAR BLOCKING AGENTS
- U-55 TREATMENT OF PAIN
- U-56 AID TO SMOKING CESSATION
- U-57 OPHTHALMIC USE OF NORFLOXACIN
- U-58 METHOD OF TREATING INFLAMMATORY INTESTINAL DISEASES
- U-59 METHOD OF TREATING HYPERCHOLESTEROLEMIA
- U-60 NASAL ADMINISTRATION OF BUTORPHANOL
- U-61 CEREBRAL AND PERIPHERAL ARTERIOGRAPHY AND CT IMAGING OF THE HEAD

PATENT AND EXCLUSIVITY TERMS

ADB 18 of 32

PATENT USE

U-62 CORONARY ARTERIOGRAPHY, LEFT VENTRICULOGRAPHY, CT IMAGING OF THE BODY, INTRAVENOUS EXCRETORY UROGRAPHY, INTRAVENOUS DIGITAL SUBTRACTION ANGIOGRAPHY AND VENOGRAPHY

U-63 ISOPRENALINE ANTAGONISM ON THE HEART RATE OR BLOOD PRESSURE

U-64 TREATMENT OF VIRAL INFECTIONS

U-65 METHOD OF TREATMENT OF A PATIENT INFECTED WITH HIV

U-66 TRIPHASIC REGIMEN

U-67 METHOD OF INDUCING ANESTHESIA IN A WARM BLOODED ANIMAL

U-68 TREATMENT OF ACTINIC KERATOSIS

U-69 TREATMENT OF PNEUMOCYSTIS CARINII INFECTIONS

U-70 TREATMENT OF TRANSIENT INSOMNIA

U-71 METHOD OF TREATMENT OF HEART FAILURE

U-72 TREATMENT OF MIGRAINE

U-73 METHOD OF TREATING DISEASES OR INFECTIONS CAUSED BY MYCETES

U-74 METHOD OF PROVIDING HYPNOTIC EFFECT

U-75 RELIEF OF OCULAR ITCHING DUE TO SEASONAL ALLERGIC CONJUNCTIVITIS

U-76 USE TO IMAGE A SUBJECT WITH A MAGNETIC RESONANCE IMAGING SYSTEM

U-77 TREATMENT OF SYMPTOMS OF SEASONAL ALLERGIC RHINITIS

U-78 ULCERATIVE COLITIS

U-79 SYMPTOMATIC TREATMENT OF PATIENTS WITH NOCTURNAL HEARTBURN DUE TO GERD

U-80 METHOD OF TREATING OCULAR BACTERIAL INFECTIONS

U-81 RELIEF OF SYMPTOMS ASSOCIATED WITH SEASONAL ALLERGIC RHINITIS

U-82 TREATMENT FOR DEMENTIA IN PATIENTS WITH ALZHEIMER'S DISEASE

U-83 TREATMENT OF SEIZURES

U-84 A METHOD OF BLOCKING THE UPTAKE OF MONOAMINES BY BRAIN NEURONS IN ANIMALS

U-85 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

U-86 METHOD OF TREATING CERTAIN FORMS OF EPILEPSY

U-87 METHOD FOR NONINVASIVE ADMINISTRATION OF SEDATIVES, ANALGESICS, AND ANESTHETICS

U-88 TREATMENT OF MODERATE PLAQUE PSORIASIS

U-89 TREATMENT OR PROPHYLAXIS OF EMESIS

U-90 TREATMENT OF PSYCHOTIC DISORDERS

U-91 ALTERNATIVE THERAPY TO TRIMETHOPRIM-SULFAMETHOXAZOLE FOR TREATMENT OF MODERATE-TO-SEVERE PNEUMOCYSTIS CARINII PNEUMONIA IN IMMUNOCOMPROMISED AND AIDS PATIENTS

U-92 TREATMENT OF DIABETIC NEPHROPATHY IN PATIENTS WITH TYPE I INSULIN DEPENDENT DIABETES MELLITUS AND RETINOPATHY

U-93 USE AS AN ANTIHISTAMINE/DECONGESTANT

U-94 TREATMENT-ADULTS W/ ADVANCED HIV, INTOLERANT OF APPROVED THERAPIES, INTOLERANT OF APPROVED THERAPIES W/ PROVEN BENEFIT OR HAVE EXPERIENCED CLINICAL/IMMUNOLOGICAL DETERIORATION WHILE RECEIVING. OR FOR WHOM SUCH THERAPIES-CONTRAINDICATED

U-95 SHORT TERM MANAGEMENT OF MODERATE PRURITIS IN ADULTS WITH ATOPIC DERMATITIS AND LICHEN SIMPLEX CHRONICUS

U-96 METHOD OF TREATING VARICELLA ZOSTER (SHINGLES) INFECTIONS

U-97 A METHOD OF TREATING A PATIENT IN NEED OF MEMORY ENHANCEMENT

U-98 A METHOD OF INDUCING REGRESSION OF LEUKEMIA CELL GROWTH IN A MAMMAL

U-99 METHOD OF PROVIDING POTASSIUM TO A SUBJECT IN NEED OF POTASSIUM

U-100 METHOD OF TREATING OCULAR INFLAMMATION

U-101 ADJUNCT TO CONVENTIONAL CT OR MRI IMAGING IN THE LOCALIZATION OF STROKE IN PATIENTS IN WHOM STROKE HAS ALREADY BEEN DIAGNOSED

U-102 METHOD OF HORMONALLY TREATING MENOPAUSAL OR POST-MENOPAUSAL DISORDERS IN WOMEN

U-103 TREATMENT OF OCULAR HYPERTENSION

U-104 TREATMENT OF AQUEOUS HUMOR FORMATION AND INTRAOCULAR PRESSURE

U-105 EMESIS

U-106 TREATMENT OF EPILEPSY

U-107 TREATMENT OF HYPERTENSION AND ANGINA PECTORIS

U-108 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER, GASTROESOPHAGEAL REFLUX DISEASE (GERD), SEVERE EROSIIVE ESOPHAGITIS, POORLY RESPONSIVE SYMPTOMATIC GERD AND PATHOLOGICAL HYPERSECRETORY CONDITIONS AND MAINTENANCE HEALING OF EROSIIVE ESOPHAGITIS

U-109 ADJUNCT DIET IN THE TX OF ELEVATED TOTAL CHOLESTEROL AND LDL-C LEVELS IN PTS W/ PRIMARY HYPERCHOLESTEROLEMIA WHOSE RESPONSE TO DIETARY RESTRICTION OF SAT FAT AND CHOLESTEROL AND OTHER NONPHARMACOLOGICAL MEASURES HAS NOT BEEN ADEQUATE

U-110 USE AS A RETRIEVABLE PERSSARY

U-111 DIABETES

U-112 CONTRACEPTION

U-113 METHOD OF CONDUCTING RADIOLOGICAL EXAMINATION OF A PATIENT BY ADMINISTERING TO SAID PATIENT A RADIOPAQUE AMOUNT OF IOPROMIDE

PATENT AND EXCLUSIVITY TERMS

ADB 19 of 32

PATENT USE

- U-114 USE FOR INHIBITING BONE RESORPTION
- U-115 USE OF VASODILATORS TO EFFECT AND ENHANCE AN ERECTION (AND THUS TREAT ERECTILE DYSFUNCTION), BY INJECTION INTO THE PENIS
- U-116 METHOD OF MYOCARDIAL IMAGING
- U-117 TREATMENT OF OCULAR ALLERGIC RESPONSE IN HUMAN EYES
- U-118 METHOD OF LOWERING BLOOD SUGAR LEVEL
- U-119 TREATMENT OF NASAL HYPERSECRETION
- U-120 CONTROLLING OR PREVENTING POST-OPERATIVE INTRAOCULAR PRESSURE RISES ASSOCIATED WITH OPHTHALMIC LASER SURGICAL PROCEDURES
- U-121 METHOD OF TREATING CONDITIONS MEDIATED THROUGH HISTAMINE H2-RECEPTORS
- U-122 A THERAPEUTIC METHOD FOR CONTROLLING THROMBOSIS
- U-123 METHOD FOR CONTROLLING THROMBOSIS AND DECREASING BLOOD HYPERCOAGULATION AND HEMORRHAGING RISKS
- U-124 TREATMENT OF ACNE
- U-125 TREATMENT NEUROGENERATIVE DISEASES
- U-126 TREATMENT OF GASTRITIS
- U-127 METHOD OF PRODUCING NEUROMUSCULAR BLOCKADE
- U-128 METHOD FOR TREATMENT OF TUMORS
- U-129 METHOD TO DESTROY OR IMPAIR TARGET CELLS
- U-130 MANAGEMENT OF PATIENTS WITH MASTOCYTOSIS
- U-131 PHOTODAMAGED SKIN
- U-132 INHIBITING HIV PROTEASE
- U-133 MANAGEMENT OF OBESITY INCLUDING WEIGHT LOSS AND MAINTENANCE IN PATIENTS ON A REDUCED-CALORIE DIET
- U-134 TREATMENT OF ACNE VULGARIS
- U-135 ANTITUMOR AGENT
- U-136 PROCESS FOR WASTE NITROGEN REMOVAL
- U-137 METHOD OF TREATING BACTERIAL VAGINOSIS
- U-138 TREATMENT OF ALLERGIC RHINITIS
- U-139 TREATMENT OF ALLERGIC REACTIONS
- U-140 USE OF NORVIR TO INHIBIT HIV PROTEASE OR TO INHIBIT AN HIV INFECTION
- U-141 TREATMENT OF ULCERATIVE COLITIS
- U-142 METHOD OF TREATING ALLERGIC REACTIONS IN A MAMMAL BY USING THIS ACTIVE METABOLITE
- U-143 BIODEGRADABLE SUPERPARAMAGNETIC METAL OXIDES AS CONTRAST AGENTS FOR MR IMAGING
- U-144 BIOLOGICALLY DEGRADABLE SUPERPARAMAGNETIC MATERIALS FOR USE IN CLINICAL APPLICATIONS
- U-145 BIOLOGICALLY DEGRADABLE SUPERPARAMAGNETIC PARTICLES FOR USE AS NUCLEAR MAGNETIC RESONANCE IMAGING AGENTS
- U-146 METHOD OF TREATING SUSCEPTIBLE NEOPLASMS IN MAMMALS
- U-147 DETECTION OF GASTROINTESTINAL DISORDERS AND THE SUBSEQUENT BREATH COLLECTION AND MEASUREMENT OF $^{13}\text{CO}_2$
- U-148 DEVICE FOR COLLECTING A BREATH SAMPLE
- U-149 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS, ACUTE MANIA OR MILD ANXIETY STATES
- U-150 METHOD OF USE FOR CONTROLLING HYPERGLYCEMIA BY ADMINISTRATION OF THIS SUSTAINED RELEASE DOSAGE FORM OF GLIPIZIDE
- U-151 RELIEF OF SYMPTOMS OF THE COMMON COLD
- U-152 METHOD OF TREATING ANXIETY RELATED DISORDERS INCLUDING OBSESSIVE COMPULSIVE DISORDER
- U-153 TREATMENT OF INITIAL EPISODE GENITAL HERPES
- U-154 METHOD OF TREATING ANIMALS SUFFERING FROM AN APPETITE DISORDER
- U-155 TREATMENT OF ERECTILE DYSFUNCTION
- U-156 METHOD OF PROVIDING ANESTHESIA
- U-157 TREATMENT OF A HUMAN SUFFERING FROM VITAMIN B12 DEFICIENCY
- U-158 ANGINA
- U-159 TREATMENT OF INTERSTITIAL CYSTITIS
- U-160 TREATMENT OF BACTERIAL INFECTIOUS DISEASE
- U-161 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS IN A PATIENT
- U-162 METHOD OF USE TO INHIBIT CHOLESTEROL SYNTHESIS IN A HUMAN SUFFERING FROM HYPERCHOLESTEROLEMIA
- U-163 METHOD OF USING TROGLITAZONE TO TREAT IMPAIRED GLUCOSE TOLERANCE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS
- U-164 METHOD OF USING TROGLITAZONE TO PREVENT OR DELAY THE ONSET OF NONINSULIN-DEPENDENT DIABETES MELLITUS IN A DEFINED POPULATION OF PATIENTS
- U-165 TREATMENT OF SYMPTOMATIC BENIGN PROSTATIC HYPERPLASIA

PATENT AND EXCLUSIVITY TERMS

ADB 20 of 32

PATENT USE

U-166 TREATMENT OF H.PYLORI-ASSOCIATED DUODENAL ULCER
U-167 METHOD FOR TREATING HIV-1 INFECTION
U-168 METHOD OF INHIBITING LIPOXYGENASE ACTIVITY IN A MAMMAL WHICH IS THE MODE OF ACTION IN THE TREATMENT OF ASTHMA
U-169 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS A CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING
U-170 METHOD OF OBTAINING AN MR IMAGE USING THE COMPOSITION/DRUG PRODUCT AS A CONTRAST AGENT
U-171 METHODS OF USING THE COMPOUND/DRUG PRODUCT AS AN ORAL CONTRAST AGENT IN MAGNETIC RESONANCE IMAGING OF THE GASTROINTESTINAL TRACT
U-172 TREATMENT OF GENITAL WARTS
U-173 ADMINISTRATION TO A HOST SUFFERING FROM GESTATIONAL DIABETES
U-174 USE AS AN ANTIHISTAMINE AGENT
U-175 METHOD OF TREATING MALIGNANT TUMORS
U-176 METHOD OF TREATING A PATIENT SUFFERING FROM LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES
U-177 FUNGICIDE
U-178 FACILITATED ADHERENCE OF AGENTS TO SKIN
U-179 ENHANCED CUTANEOUS PENETRATION OF A DERMALLY-APPLIED PHARMACOLOGICALLY ACTIVE AGENT
U-180 TREATMENT OF ADULT AND PEDIATRIC PATIENTS (OVER 6 MONTHS OF AGE) WITH ADVANCED HIV INFECTION
U-181 PRODUCING ALPHA ADRENERGIC ANTAGONISTIC ACTION IN A HOST
U-182 USE OF SALMETEROL IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION
U-183 TREATMENT OF CONDITIONS CAUSED BY DISTURBANCE OF NEURONAL 5HT FUNCTION
U-184 TREATING ALLERGIC EYE DISEASES IN HUMANS
U-185 METHOD OF TREATING HYPERTENSION
U-186 METHOD FOR TREATING GI DISORDERS CAUSED BY H.RYLORI WHICH COMPRISES ADMINISTRATION OF RANITIDINE BISMUTH CITRATE AND CLARITHROMYCIN FOR A GREATER THAN ADDITIVE EFFECT
U-187 THERAPEUTIC TREATMENT OF CALCIFIC TUMORS
U-188 TREATMENT OF H.PYLORI ASSOCIATED DUODENAL ULCER
U-189 ENHANCEMENT OF THE BIOAVAILABILITY OF THE DRUG SUBSTANCE
U-190 USE OF RITONAVIR IN COMBINATION WITH ANY REVERSE TRANSCRIPTASE INHIBITOR
U-191 METHOD OF TREATMENT FOR CONTROLLING AND LOWERING INTRAOCULAR PRESSURE IN A HUMAN
U-192 USE IN TREATING ALLERGIC REACTIONS
U-193 PSORIASIS
U-194 TREATING ANGINA PECTORIS AND HIGH BLOOD PRESSURE
U-195 METHOD FOR THE DIAGNOSIS OF GASTROINTESTINAL DISORDERS BY UREA ISOTOAC OR NITROGEN LABELED CARBON
U-196 TREATMENT OF METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH ESTROGEN RECEPTOR POSITIVE TUMORS
U-197 USE IN COMBINATION WITH CERTAIN LHRH ANALOGUES FOR THE TREATMENT OF ADVANCED PROSTATE CANCER
U-198 TREATMENT METASTATIC CARCINOMA OF OVARY AFTER 1ST LINE FAILURE OR SUBSEQUENT CHEMOTHERAPY, TREATMENT OF BREAST CANCER AFTER FAILURE OF COMBINATION CHEMOTHERAPY FOR METASTATIC DISEASE AND 2ND LINE TREATMENT OF AIDS RELATED KAPOSI'S SARCOMA
U-199 METHOD OF TREATING INFECTIOUS UPPER GI TRACT DISORDERS CAUSED BY CAMPYLOBACTER PYLORIDIS INFECTION COMPRISING ADMINISTRATION OF A BISMUTH AGENT AND AN ANTIMICROBIAL AGENT
U-200 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF A BISMUTH-CONTAINING AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT
U-201 METHOD OF TREATING GI DISORDERS COMPRISING ADMINISTRATION OF CAMPYLOBACTER-INHIBITING ANTIMICROBIAL AGENT AND H2 RECEPTOR BLOCKING ANTI-SECRETORY AGENT
U-202 METHOD OF TREATING PEPTIC ULCER DISEASE CAUSED BY CAMPYLOBACTER PYLORIDIS COMPRISING ORAL ADMINISTRATION OF 50 TO 5,000MG BISMUTH DAILY FOR 3-56 DAYS
U-203 TREATMENT OF ADVANCED BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY
U-204 USE OF TAXOL IN COMBINATION WITH G-CSF FOR TREATMENT OF PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA
U-205 METHOD FOR TREATING HEARTBURN
U-206 METHOD OF USING FSH ALONE, WITHOUT THE PRESENCE OF EXOGENEOUS LH, IN IN VITRO FERTILIZATION
U-207 USE AS NASAL SPRAY
U-208 VAGINAL ADMINISTRATION USING SPECIFIED FORMULATION
U-209 VAGINAL ADMINISTRATION OF PROGESTERONE USING SPECIFIED FORMULATION
U-210 METHOD OF TREATING CONGESTIVE HEART FAILURE
U-211 USE IN PATIENTS WITH REVERSIBLE AIRWAY OBSTRUCTION
U-212 METHOD OF TREATMENT OF PARKINSON'S DISEASE

PATENT AND EXCLUSIVITY TERMS

ADB 21 of 32

PATENT USE

U-213 METHOD OF INHIBITING CHOLESTEROL BIOSYNTHESIS AND TREATING HYPERCHOLESTEROLEMIA AND METHOD FOR TREATING HYPERLIPIDEMIA
U-214 USE AS A BLOOD GLUCOSE-LOWERING AGENT
U-215 TREATMENT OF EPILEPSY TWICE DAILY. TREATING A PATIENT BY ADMINISTERING CARBAMAZEPINE IN A DOSAGE FORM CAPABLE OF MAINTAINING BLOOD CONCENTRATION FROM 4-12MCG/ML OVER 12 HOURS
U-216 TREATMENT OF ADENOCARCINOMA, INCLUDING STAGE B2-C BY ADMINISTERING AN AGONIST OF LH-RH AND FLUTAMIDE
U-217 METHOD OF PRODUCING ANESTHESIA
U-218 METHOD FOR LIMITING THE POTENTIAL FOR MICROBIAL GROWTH IN THE DRUG PRODUCT
U-219 TREATMENT OF PARKINSON'S DISEASE
U-220 METHOD OF DIAGNOSIS
U-221 SELECTIVE VASODILATION BY CONTINUOUS ADENOSINE INFUSION
U-222 METHOD OF TREATING PAGET'S DISEASE USING ACTONEL
U-223 TREATMENT OF BACTERIAL CONJUNCTIVITIS CAUSED BY SUSCEPTIBLE STRAINS OF MICROORGANISMS
U-224 CONTROLLING INTRAOCULAR PRESSURE
U-225 METHOD FOR DELIVERY
U-226 METHOD OF ENHANCING THE DISSOLUTION PROFILE OF A PHARMACEUTICAL FROM A SOLID DOSAGE FORM CONTAINING THE PHARMACEUTICAL AND SIMETHICONE
U-227 NASAL ADMINISTRATION
U-228 ASTHMA
U-229 CARDIAC INSUFFICIENCY (CONGESTIVE HEART FAILURE)
U-230 PREVENTION OF ACUTE CARDIAC ISCHEMIC EVENTS
U-231 USE IN PARKINSON'S DISEASE
U-232 METHOD OF TREATING MIGRAINE
U-233 DECREASING MORTALITY CAUSED BY CONGESTIVE HEART FAILURE
U-234 METHOD OF USING RIBAVIRIN TO TREAT VIRAL INFECTIONS IN MAMMALS
U-235 METHOD OF MODULATING TH1 AND TH2 RESPONSE IN ACTIVATED T CELLS OF A HUMAN COMPRISING ADMINISTERING RIBAVIRIN TO THE T CELLS IN A DOSAGE WHICH PROMOTES THE TH1 RESPONSE AND SUPPRESSES THE TH2 RESPONSE
U-236 TREATING MALE PATTERN BALDNESS WITH 0.05 TO 3.0MG/DAY
U-237 METHOD OF PERFORMING NMR IMAGING WITH A PATIENT COMPRISING ADMINISTERING TO THE PATIENT AN EFFECTIVE AMOUNT OF CONTRAST AGENT DISCLOSED IN THE CLAIMS
U-238 IMAGING A BODY TISSUE AND SUBJECTING TO NMR TOMOGRAPHY, ADMINISTERING AN AMOUNT OF PHARMACEUTICAL AGENT FOR AFFECTING THE RELAXATION TIMES OF ATOMS IN BODY TISSUES UNDERGOING NMR DIAGNOSIS, WHEREBY THE IMAGE CONTRAST IS ENHANCED....
U-239 TREATING OR CONTROLLING OCULAR INFLAMATION WHICH COMPRISES TOPICALLY ADMINISTERING TO AFFECTED EYE A COMPOSITION COMPRISING AN NSAID, A POLYMERIC QUATERNARY AMMONIUM COMPOUND AND BORIC ACID
U-240 TREATMENT OF ACUTE MIGRAINE ATTACKS
U-241 FOR SHORT-TERM TREATMENT ACTIVE DUODENAL ULCER, MAINTENANCE THERAPY FOR DUODENAL ULCER PATIENTS AT REDUCED DOSAGE AFTER HEALING OF ACTIVE ULCER, SHORT-TERM TREATMENT ACTIVE BENIGN GASTRIC ULCER & GERD, PATHOLOGICAL HYPERSECRETORY CONDITIONS
U-242 USE OF FOLLITROPIN ALPHA ALONE IN IN-VITRO FERTILIZATION
U-243 TOPICAL ADMINISTRATION
U-244 PLATELET AGGREGATION INHIBITORS
U-245 TREATMENT OF SEBORRHEA DERMATITIS IN HUMANS
U-246 PHOSPHATE BINDING
U-247 TREATMENT OF RHEUMATOID ARTHRITIS
U-248 TREATMENT OF HIV
U-249 METHOD OF TREATING ALLERGIC OR NON-ALLERGIC RHINITIS IN PATIENTS BY ADMINISTERING AEROSOLIZED PARTICLES OF MOMETASONE FUROATE
U-250 TREATMENT OF HEPATITIS B INFECTION
U-251 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS IN THE TREATMENT OF TYPE II DIABETES
U-252 METHOD OF TREATING A HUMAN SUBJECT HAVING GAUCHER'S DISEASE
U-253 ORAL TRANSMUCOSAL USE
U-254 USE OF AGGRASTAT IN COMBINATION WITH HEPARIN
U-255 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY
U-256 TREATMENT OF HIV INFECTION IN COMBINATION WITH ONE OR MORE ADDITIONAL HIV ANTIVIRAL AGENTS
U-257 TREATMENT OF HIV INFECTION
U-258 TREATMENT OF NEURODEGENERATIVE DISEASES
U-259 TREATMENT OF ANDROGENIC ALOPECIA BY ORAL ADMINISTRATION DRUG SUBSTANCE

PATENT AND EXCLUSIVITY TERMS

ADB 22 of 32

PATENT USE

- U-260 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA AND OCULAR HYPERTENSION WHO ARE INTOLERANT OF OTHER IOP LOWERING MEDICATIONS OR INSUFFICIENTLY RESPONSIVE TO ANOTHER IOP LOWERING MEDICATION
- U-261 TREATING BENIGN PROSTATIC HYPERPLASIA WITH A GENUS OF COMPOUNDS, INCLUDING FINASTERIDE
- U-262 TREATING BENIGN PROSTATIC HYPERTROPHY WITH FINASTERIDE
- U-263 METHOD OF TREATING A MALIGNANT CONDITION THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING LEUKEMIA OR LYMPHOMA IN A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVENOUS ADMINISTRATION OF BUSULFAN
- U-264 METHOD OF TREATING A MALIGNANT DISEASE THROUGH PARENTERAL ADMINISTRATION OF BUSULFAN. METHOD FOR TREATING A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN
- U-265 USE AS LAXATIVE
- U-266 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS; RELIEF OF THE SIGNS AND SYMPTOMS OF RHEUMATOID ARTHRITIS IN ADULTS; MANAGEMENT OF ACUTE PAIN IN ADULTS; TREATMENT OF PRIMARY DYSMENORRHEA; ACUTE TREATMENT OF MIGRAINE ATTACKS IN ADULTS
- U-267 PREVENTING HEARTBURN EPISODES FOLLOWING INGESTION OF HEARTBURN-INDUCING FOOD/BEVERAGE, COMPRISING ADMIN TO PT, 30 MIN PRIOR TO CONSUMPTION BY THE PT THE FOOD/BEVERAGE, A COMPOSITION COMPRISING 10MG FAMOTIDINE
- U-268 ACROMEGALY
- U-269 EXCESS GH-SECRETION OR GASTRO-INTESTINAL DISORDERS
- U-270 METHOD OF IMPROVING THE TIME FOR ADMINISTRATION OR THE TIME BETWEEN CHANGES OF GIVING SETS FOR THE DRUG PRODUCT
- U-271 METHOD OF TREATING TUMORS
- U-272 METHOD OF TREATING CARCINOMA
- U-273 CUTANEOUS T-CELL LYMPHOMA
- U-274 ZANAMIVIR FOR INHALATION
- U-275 METHOD OF USE OF THE DRUG SUBSTANCE
- U-276 METHOD OF USE OF LEVOBUPIVACAINE
- U-277 NEUROLOGICAL AND OTHER DISORDERS (TREATMENT OF EPILEPSY, BID ORAL DOSING)
- U-278 METHOD OF USE OF THE INDICATION OF THE DRUG PRODUCT
- U-279 METHOD OF USE OF THE APPROVED PRODUCT
- U-280 TREATING PRECIPITATED ACUTE URINARY RETENTION WITH FINASTERIDE
- U-281 ANTIMYCOTIC USES, SPECIFICALLY TREATMENT OF ONYCHOMYCOSIS
- U-282 METHOD OF TREATING BACTERIAL INFECTIONS
- U-283 METHOD FOR TREATING MENOPAUSAL SYMPTOMS IN A POSTMENOPAUSAL FEMALE
- U-284 MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE, AND VULVAR AND VAGINAL ATROPHY) AND OSTEOPOROSIS
- U-285 DEPRESSION AND SOCIAL ANXIETY DISORDER/SOCIAL PHOBIA
- U-286 DEPRESSION
- U-287 TREATMENT OR PREVENTION OF OSTEOPOROSIS
- U-288 THERAPY OF INFLUENZA
- U-289 TREATMENT OF NON-HYPERKERATOTIC ACTINIC KERATOSES OF FACE AND SCALP
- U-290 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS)
- U-291 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH CYCLOSPORIN
- U-292 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH AZATHIOPRINE
- U-293 INHIBITING TRANSPLANT REJECTION USING RAPAMYCIN (SIROLIMUS) IN COMBINATION WITH A CORTICOSTEROID
- U-294 TREATMENT OF HYPERPIGMENTARY DISORDERS
- U-295 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS
- U-296 TREATING MIGRAINE PAIN AND ONE OR MORE OF A CLUSTER OF SYMPTOMS CHARACTERISTIC OF A MIGRAINE ATTACK SYMPTOMS BEING SELECTED FROM PHOTOPHOBIA, PHONOPHOBIA NAUSEA AND FUNCTIONAL DISABILITY
- U-297 PREVENTION OR TREATMENT OF REVERSIBLE VASOCONSTRICTION BY THE INHALATION OF NITRIC OXIDE WITH AN OXYGEN CONTAINING GAS
- U-298 METHOD OF COMBATING BACTERIA IN A PATIENT
- U-299 TREATMENT OF ADENOMATOUS POLYPS
- U-300 INDICATED FOR THE REDUCTION OF ELEVATED TOTAL AND LDL CHOLESTEROL LEVELS IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA
- U-301 USE OF TROGLITAZONE IN COMBINATION WITH SULFONYLUREAS AND BIGUANIDES IN THE TREATMENT OF TYPE II DIABETES
- U-302 TO REDUCE THE RISK OF STROKE IN PATIENTS WHO HAVE HAD TRANSIENT ISCHEMIA OF THE BRAIN OR COMPLETED ISCHEMIC STROKE DUE TO THROMBOSIS

PATENT AND EXCLUSIVITY TERMS

ADB 23 of 32

PATENT USE

- U-303 METHOD OF USE PATENT-PRODUCT APPROVED FOR TREATMENT OF OSTEOPOROSIS, PAGET'S DISEASE, PREVENTION AND TREATMENT OF GLUCOCORTICOID INDUCED OSTEOPOROSIS
- U-304 A METHOD OF TREATMENT OF A CONDITION INVOLVING AN ANTIBODY ANTIGEN REACTION
- U-305 METHODS FOR USING THE DRUG PRODUCT
- U-306 TREATMENT OF POST-MENOPAUSAL UROGENITAL SYMPTOMS ASSOCIATED WITH ESTROGEN DEFICIENCY
- U-307 CLAIMS AN OLANZAPINE POLYMORPH USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATION OF THIS NDA
- U-308 CLAIMS A SOLID ORAL FORMULATION INCLUDING TABLETS AND GRANULES OF OLANZAPINE USEFUL FOR TREATING ANY NUMBER OF LISTED CONDITIONS, INCLUDING SPECIFIC PSYCHOSES, EMPLOYING OLANZAPINE AS PER THE INDICATIONS OF THIS NDA
- U-309 TREATING SJOEGREN SYNDROME
- U-310 TREATMENT OF XEROSTOMIA
- U-311 HORMONE REPLACEMENT
- U-312 PANIC DISORDER, OBSESSIVE-COMPULSIVE DISORDER, POSTTRAUMATIC STRESS DISORDER
- U-313 TREATMENT OF CONGESTIVE HEART FAILURE
- U-314 METHOD FOR TREATING HYPERPARATHYROIDISM WHICH COMPRISES SUPPRESSING PARATHYROID ACTIVITY
- U-315 METHOD FOR ADMINISTERING DRUG TO GASTROINTESTINAL TRACT
- U-316 METHOD OF TREATING A SUBJECT SUFFERING FROM PROSTATE CANCER
- U-317 METHOD OF USING TROGLITAZONE TO TREAT PATIENTS HAVING INSULIN RESISTANCE
- U-318 TREATMENT OF PATIENTS WITH AN OVERACTIVE BLADDER WITH SYMPTOMS OF URINARY FREQUENCY, URGENCY, OR URGE INCONTINENCE
- U-319 TREATMENT OF MICROBIAL INFECTIONS
- U-320 INHIBITING OR ELIMINATING ACUTE MYELOID LEUKEMIA
- U-321 REDUCTION OF ELEVATED IPTH LEVELS IN THE MGT OF SECONDARY HYPERPARATHYROIDISM IN PATIENTS UNDERGONG CHRONIC RENAL DIALYSIS
- U-322 TREATMENT OF ALZHEIMER'S DEMENTIA
- U-323 USE AS A BILE ACID SEQUESTANT
- U-324 METHOD OF TREATING AN ANIMAL, INCLUDING A HUMAN, SUFFERING FROM OR SUSCEPTIBLE TO PSYCHOSIS OR ACUTE MANIA EMPLOYING OLANZAPINE
- U-325 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS, INCLUDING "BIPOLAR DISORDER NOS" EMPLOYING OLANZAPINE
- U-326 METHOD OF TREATING SCHIZOPHRENIA AND BIPOLAR DISORDER
- U-327 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED PSYCHOTIC CONDITONS EMPLOYING OLANZAPINE
- U-328 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF LISTED CONDITIONS INCLUDING "A PSYCHOTIC CONDITION" EMPLOYING AN OLANZAPINE POLYMORPH
- U-329 USE OF AVANDIA AS MONOTHERAPY, IN COMBINATION WITH METFORMIN, AND IN COMBINATION WITH SULFONYLUREAS TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-330 TREATMENT OF NAUSEA AND VOMITING
- U-331 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-332 TREATMENT OR PREVENTION OF BRONCHOSPASM
- U-333 METHOD OF TREATING OCULAR HYPERTENSION
- U-334 TREATMENT OF EXCESSIVE FEMALE FACIAL HAIR
- U-335 USE OF PRAVASTATIN SODIUM FOR SECONDARY PREVENTION OF CORONARY EVENTS IN MEN AND WOMEN WHO HAVE HAD A MYOCARDIAL INFARCTION AND HAVE NORMAL CHOLESTEROL LEVELS
- U-336 DIAGNOSTIC RADIOIMAGING
- U-337 USE OF CARDIOLITE/MIRALUMA KIT FOR THE PREPARATION OF TC99M SESTAMIBI
- U-338 METHODS FOR TREATING DISTURBANCES OF MOOD, DISTURBANCES OF APPETITE, DEPRESSED MOOD, OR CARBOHYDRATE CRAVING ALL ASSOCIATED WITH PREMENSTRUAL SYNDROME
- U-339 PREVENTION OF CARDIO-TOXICITY CAUSED BY THE ADMINISTRATION OF DOXORUBICIN
- U-340 THE LONG TERM TREATMENT OF GROWTH FAILURE DUE TO LACK OF ADEQUATE ENDOGENOUS GROWTH HORMONE SECRETION IN CHILDREN
- U-341 METHOD FOR ENHANCING THE TREATMENT OF ... LATE LUTEAL PHASE DYSPHORIC DISORDER
- U-342 METHOD FOR TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER
- U-343 REDUCTION OF INTESTINAL GAS, CRAMPING AND ANORECTAL IRRITATION
- U-344 METHOD FOR INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH ANOTHER HIV PROTEASE INHIBITOR
- U-345 RITONAVIR AND ANOTHER HIV PROTEASE INHIBITOR FOR CONCOMITANT ADMINISTRATION FOR THE TREATMENT OF AN HIV INFECTION
- U-346 METHOD FOR INHIBITING CYTOCHROME P450 MONOOXYGENASE WITH RITONAVIR AND A METHOD FOR IMPROVING THE PHARMCOKINETICS OF A DRUG THAT IS MTABOLIZED BY CYTOCHROME P450 MONOOXYGENASE BY ADMIN THE DRUG AND RITONAVIR
- U-347 METHOD OF USE IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS
- U-348 METHOD OF USE FOR INHIBITING HIV INFECTION

PATENT AND EXCLUSIVITY TERMS

ADB 24 of 32

PATENT USE

U-349 METHOD OF USE WHICH IS SUBJECT OF THE APPLICATION
U-350 PREPARATION OF A PHARMACEUTICAL COMPOSITION FOR CONCOMITANT ADMIN WITH A REVERSE TRANSCRIPTASE INHIBITOR
U-351 INHIBITING PROTEASE WITH LOPINAVIR AND INHIBITING AN HIV INFECTION WITH LOPINAVIR
U-352 INHIBITING HIV INFECTION BY ADMINISTERING RITONAVIR IN COMBINATION WITH A REVERSE TRANSCRIPTASE INHIBITOR
U-353 PREVENTION AND TREATMENT OF OSTEOPOROSIS
U-354 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
U-355 METHOD OF ASSISTING PERSON TO QUIT SMOKING...TRANSDERMALLY ADMIN NICOTINE VIA...PATCH ADHERED TO SKIN AT DOSING RATE APPROX SAME AS ABSORBED FROM SMOKING
U-356 DELIVERING A MEDICINAL AEROSOL FORMULATION USING CFC-FREE PROPELLANT 134A.
U-357 USE OF THE DRUG PRODUCT IN PHOTODYNAMIC THERAPEUTIC PROTOCOLS FOR THE TREATMENT OF AGE-RELATED MACULAR DEGENERATION AND RELATED CONDITIONS INVOLVING UNWANTED NEOVASCULATURE IN THE EYE
U-358 DEPRESSION, OBSESSIVE COMPULSIVE DISORDER, PANIC DISORDER AND SOCIAL ANXIETY DISORDER
U-359 METHOD OF USE OF VISICOL
U-360 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS INCLUDING MENTAL DISORDERS EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011
U-361 MANAGEMENT OF ANXIETY DISORDERS AND THE SHORT-TERM RELIEF OF THE SYMPTOMS OF ANXIETY
U-362 USE OF APPROVED FORMULATIONS TO TREAT ALL APPROVED DISEASE INDICATIONS
U-363 METHOD OF TREATING A PATIENT SUFFERING FROM ANY OF A NUMBER OF PATHOLOGICAL PSYCHOLOGICAL CONDITIONS THAT RELATE TO THE USE OF A PSYCHOACTIVE SUBSTANCE EMPLOYING OLANZAPINE AS PER THE INDICATION THE SUBJECT MATTER OF SUPPLEMENT 011
U-364 TREATING A PATIENT SUFFERING FROM OR SUSCEPTIBLE TO ANY NUMBER OF LISTED CONDITIONS INCLUDING PSYCHOSIS, EMPLOYING OLANZAPINE AS PER THE INDICATION WHICH IS THE SUBJECT MATTER OF THIS SNDA-011
U-365 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN OUR EXTENDED, CONTROLLED RELEASE FORMULATION
U-366 METHOD FOR THE TREATMENT OF CARDIOVASCULAR DISEASE THROUGH THE ADMINISTRATION OF A CALCIUM BLOCKING VASODILATOR IN A DELAYED RELEASE FORMULATION
U-367 TREATMENT OF CARDIOVASCULAR DISORDERS
U-368 HEARTBURN
U-369 METHOD OF CONTROLLING AND LOWERING INTRAOCULAR PRESSURE
U-370 INTRAVAGINAL TREATMENT OF VAGINAL INFECTIONS WITH BUFFERED METRONIDAZOLE COMPOSITIONS
U-371 APPROVAL FOR MARKETING ONLY UNDER A SPECIAL RESTRICTION PROGRAM APPROVED BY FDA CALLED "SYSTEM FOR THALIDOMIDE EDUCATION AND PRESCRIBING SAFETY" (S.T.E.P.S.)
U-372 METHOD FOR ADMINISTERING A BENEFICIAL DRUG TO THE GI TRACT OF AN ANIMAL, WHICH METHOD COMPRISES ADMITTING AN OSMOTIC DEVICE ORALLY INTO THE ANIMAL...
U-373 GENERAL USE CLAIM SUBMITTED FOR 12 NEXIUM PATIENTS STATING "PERTINENT TO THE CAPSULE FORMULATION FOR NEXIUM AND ITS INDICATIONS FOR THE TREATMENT OF GERD AND ERADICATION OF H.PYLORI TO REDUCE THE RISK OF DUODENAL ULCER RECURRENCE
U-374 KIT ADAPTED AND DESIGNED TO PROVIDE BOTH DATA ON THE CURRENT REPRODUCTIVE STATUS OF A PATIENT AND CONTRACEPTION FOR THOSE WHO ARE NOT PREGNANT, BUT RECENTLY ENGAGED IN UNPROTECTED SEX
U-375 METHOD OF USING RIBAVIRIN FOR TREATING A DISEASE RESPONSIVE TO RIBAVIRIN, E.G. HEPATITIS C
U-376 TREATMENT OF INFLUENZA
U-377 METHOD OF TREATING PT WITH CHRONIC HEPATITIS C HAVING HCV GENOTYPE 1 AND VIRAL LOAD GREATER THAN 2 MILLION COPIES/ML TO ERADICATE DETECTABLE HCV-RNA BY ADMIN COMBINATION OF RIBAVIRIN AND INTERFERON ALFA-2B FOR A LEAST 24 WEEKS
U-378 METHOD FOR TREATING INCONTINENCE
U-379 METHOD OF TREATING ONYCHROMYCOSIS
U-380 COMBINATIONS OF TAXOL (PACLITAXEL) AND CISPLATIN WHICH ARE SUITABLE FOR THE TREATMENT OF OVARIAN AND NON-SMALL CELL LUNG CARCINOMAS
U-381 TREATMENT OF HYPERPHOSPHATEMIA
U-382 METHOD OF STABILIZING PROSTAGLANDIN
U-383 METHOD FOR TREATING GLAUCOMA AND OCULAR HYPERTENSION
U-384 TREATMENT OF CMV RETINITIS
U-385 TREATMENT OF PEPTIC ULCERS
U-386 TREATMENT OF PATIENTS SUFFERING FROM A LATE ASTHMATIC REACTION OR LATE PHASE ASTHMA
U-387 TREATMENT OF PATIENTS WITH RESPIRATORY DISORDERS
U-388 SMOKING CESSATION AID APPLIED TO THE SKIN

PATENT AND EXCLUSIVITY TERMS

ADB 25 of 32

PATENT USE

- U-389 SMOKING CESSATION AID APPLIED TO THE SKIN ON WAKING AND REMOVED PRIOR TO SLEEP AFTER ABOUT 16 HOURS
- U-390 METHOD OF USING THE DRUG TO TREAT NEUROIMMUNOLOGIC DISEASES (INCLUDING MULTIPLE SCLEROSIS)
- U-391 USE OF CASODEX IN COMBINATION WITH LHRH AGONISTS FOR THE TREATMENT OF PROSTATE CANCER
- U-392 TREATMENT OF PATIENTS FOR INFLAMMATION
- U-393 MANAGEMENT OF INCONTINENCE, MGT OF HORMONE REPLACEMENT THERAPY, TREATMENT OF INVOLUNTARY INCONTINENCE, MGT OVERACTIVE BLADDER AND INCREASING COMPLIANCE IN SUCH PT
- U-394 METHOD OF USE OF ALPHAGAN
- U-395 METHOD OF USE OF ALPHAGAN P
- U-396 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION
- U-397 METHOD OF TREATING PEOPLE SUFFERING FROM DEPRESSION WITHOUT AN INCREASE IN NAUSEA
- U-398 TREATMENT OF GENERALIZED ANXIETY DISORDER
- U-399 IN-THE-EYE USE OF CHLORINE DIOXIDE CONTAINING COMPOSITIONS
- U-400 USE OF RIBAVIRIN TO INCREASE TYPE 1 CYTOKINE RESPONSE AND SUPPRESS TYPE 2 CYTOKINE RESPONSE TO LYMPHOCYTES, INCLUDING METHODS THAT TAKE ADVANTAGE OF SUCH MODULATION TO TREAT INFECTIONS AND INFESTATIONS
- U-401 USE OF LOPINAVIR IN COMBINATION WITH REVERSE TRANSCRIPTASE INHIBITORS FOR TREATING HIV INFECTION AND IN COMBO WITH OTHER HIV PROTEASE INHIBITORS
- U-402 TREATMENT OF ACTINIC KERATOSES
- U-403 ANTI-ALLERGIC FOR VARIOUS ALLERGIC DISEASES
- U-404 TREATMENT OF ALLERGIC CONJUNCTIVITIS
- U-405 FOR WOMEN WITH SEVERE DIARRHEA-PREDOMINANT IRRITABLE BOWEL SYNDROME (IBS)
- U-406 METHOD OF USE OF ATOVAQUONE AND PROGUANIL
- U-407 METHOD OF TREATING OTOPATHY
- U-408 FOR INDUCING OVULATION IN CONJUNCTION WITH A GONADOTROPIN RELEASING FACTOR ANTAGONIST AND RECRUITING OOCYTES FOR IN-VITRO FERTILIZATION
- U-409 METHOD OF TREATING INFLAMMATION USING DRUG SUBSTANCE
- U-410 METHOD OF REDUCING AMOUNT OF RESPECTIVE ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (INCLUDING PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-411 METHOD OF REDUCING THE SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN PREPARATION
- U-412 TREATMENT OF TYPE 2 DIABETES
- U-413 USE OF THE ACTIVE INGREDIENT FOR INHIBITING THE BIOSYNTHESIS OF CHOLESTEROL AND TREATMENT OF ATHEROSCLEROSIS
- U-414 A METHOD OF TREATING GLYCOMETABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE
- U-415 A METHOD FOR REDUCING THE AMOUNT OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS
- U-416 A METHOD FOR REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMINISTERED TO A DIABETIC PATIENT BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE AS SAID ACTIVE COMPONENTS
- U-417 COMBINATION USE OF AD-4833 WITH A BIGUANIDE
- U-418 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING A CHEMICAL COMPOUND HAVING A PARTICULAR FORMULA (WHICH INCLUDES PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-419 A METHOD OF TREATING LIPID METABOLISM DISORDERS BY ADMINISTERING AN INSULIN SENSITIVITY ENHANCER (INCLUDING PIOGLITAZONE) IN COMBINATION WITH A BIGUANIDE
- U-420 METHOD OF TREATMENT OF TYPE II DIABETES
- U-421 USE FOR SEDATION
- U-422 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER AND ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-423 METHOD OF TREATING AT LEAST ONE OF ATTENTION DEFICIT DISORDER, ATTENTION DEFICIT HYPERACTIVITY DISORDER, OR AIDS RELATED DEMENTIA
- U-424 FOR ONCE DAILY, BOLUS ADMINISTRATION TO A PATIENT IN ORDER TO ENGENDER TREATMENT FOR A NERVOUS DISORDER FOR SUBSTANTIALLY AN ENTIRE DAY ON A CHRONIC BASIS
- U-425 METHOD OF REDUCING SIDE EFFECTS OF ACTIVE COMPONENTS ADMIN TO A DIABETIC BY ADMIN A CHEMICAL COMPOUND HAVING FORMULA (INCL PIOGLITAZONE) IN COMBINATION WITH AN INSULIN SECRETION ENHANCER
- U-426 PREVENTION OF PREMATURE LH SURGES IN WOMEN UNDERGOING CONTROLLED OVARIAN STIMULATION
- U-427 METHOD OF TREATING ALLERGIC REACTIONS IN MAMMALS
- U-428 METHOD OF TREATING ALLERGY IN A MAMMAL USING THIS ACTIVE METABOLITE
- U-429 METHOD OF USING DESLORATADINE TO TREAT ALLERGIC RHINITIS

PATENT AND EXCLUSIVITY TERMS

ADB 26 of 32

PATENT USE

- U-430 METHOD OF TREATING A DIABETIC BY ADMINISTERING AN INSULIN SENSITIZER IN COMBINATION WITH AN INSULIN SECRETION ENHANCER, AND A DRUG PRODUCT COMPRISING AN INSULIN SENSITIZER AND AN INSULIN SECRETION ENHANCER
- U-431 POSTTRAUMATIC STRESS DISORDER
- U-432 REDUCTION OF ATHEROSCLEROTIC EVENTS (MYOCARDIAL INFARCTION, STROKE, AND VASCULAR DEATH) IN PATIENTS WITH ATHEROSCLEROSIS DOCUMENTED BY RECENT STROKE, RECENT MYOCARDIAL INFARCTION OR ESTABLISHED PERIPHERAL ARTERIAL DISEASE
- U-433 USE OF LEVOCARITINE IN PREVENTION AND TREATMENT OF CARNITINE DEFICIENCY IN PATIENTS WITH END STAGE RENAL DISEASE WHO ARE UNDERGOING DIALYSIS
- U-434 CONTROLLED SYMPTOMS OF DIARRHEA, BLOATING PRESSURE AND CRAMPS, COMMONLY REFERRED TO AS GAS
- U-435 A TITRATION DOSING REGIMEN FOR THE TREATMENT OF PAIN USING AN INITIAL DOSE OF ABOUT 25MG
- U-436 ACUTE TREATMENT OF MIGRAINE ATTACKS WITH OR WITHOUT AURA IN ADULTS
- U-437 METHOD OF USE EQUAL TO PROCESS OF PREPARATION
- U-438 TREATMENT/PREVENTION OF NEURODEGENERATIVE DISEASE
- U-439 TREATMENT OF OBESITY
- U-440 METHOD FOR TRANSDERMAL ADMINISTRATION OF A DRUG THROUGH NON-SCROTAL SKIN USING A TRANSDERMAL DRUG DELIVERY DEVICE CONTAINING THE DRUG AND HAVING AN ADHESIVE SURFACE
- U-441 METHOD OF TREATING MS BY ADMINISTERING COPAXONE
- U-442 METHOD FOR DELIVERING A DRUG TO A PATIENT IN NEED OF THE DRUG, WHILE AVOIDING THE OCCURENCE OF AN ADVERSE SIDE EFFECT KNOWN OR SUSPECTED OF BEING CAUSED BY SAID DRUG
- U-443 MANAGEMENT OF MODERATE TO SEVERE PAIN WHEN A CONTINUOUS, AROUND-THE-CLOCK ANALGESIC IS NEEDED FOR AN EXTENDED PERIOD OF TIME
- U-444 TREATMENT OF MIGRAINE
- U-445 USE AS AN ANTIMYCOTIC AGENT
- U-446 TOPICAL TREATMENT OF OCULAR HYPERTENSION AND GLAUCOMA
- U-447 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-448 METHOD OF TREATING HYPERLIPIDEMIA WITH NICOTINIC ACID WITHOUT CAUSING TREATMENT-LIMITING ELEVATIONS IN URIC ACID OR GLUCOSE LEVELS OR CAUSING LIVER DAMAGE, BY DOSING ONCE PER DAY IN THE EVENING OR AT NIGHT
- U-449 USE IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLORECTAL CANCER WHERE THE DOSE OF LEUCOVORIN IS AT LEAST 200MG PER SQUARE METER
- U-450 INTERMEDIATE REL NICOTINIC ACID FORMULATIONS HAVING UNIQUE URINARY METAB PROFILES RESULTING FROM ABSORPTION PROFILES OF NICOTINIC ACID FROM THE INTERMEDIATE NICOTINIC ACID FORMULATIONS, SUITABLE FOR TX HYPERLIPIDEMIA FOLLOWING QD DOSING
- U-451 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER
- U-452 USE OF LANSOPRAZOLE FOR COMBATTING DISEASES CAUSED BY THE GENUS CAMPYLOBACTER (C.PYLORI=H.PYLORI)
- U-453 TREATMENT OF PLATELET ASSOCIATED ISCHEMIC DISORDERS
- U-454 METHOD OF TX A PT SUSPECTED OF HAVING HEPATITIS C BY ADMIN, IN COMBINATION, A CONJUGATE COMPRISING PEG 12000 & INTERFERON ALFA-2B IN AN AMT OF FROM 0.5MCG/KG TO 2MCG/KG, ONCE WEEKLY, AND RIBAVIRIN
- U-455 TREATMENT OF PULMONARY HYPERTENSION WITH UT-15
- U-456 METHOD OF DECREASING THE PRODUCTION OF A-BETA USING A COMPOSITION WHICH DECREASES BLOOD CHOLESTEROL IN PATIENTS AT RISK OF OR EXHIBITING SYMPTOMS OF ALZHEIMER'S DISEASE
- U-457 METHOD OF TREATING A VAGINAL FUNGAL INFECTION IN A FEMALE HUMAN
- U-458 METHOD OF USE OF IMAGENT
- U-459 TREATMENT OF DEPRESSION AND GENERALIZED ANXIETY DISORDER
- U-460 METHOD OF TREATING PSYCHIATRIC SYMPTOMS ASSOCIATED WITH PREMENSTRUAL DISORDERS USING SERTRALINE
- U-461 METHOD OF TREATMENT OF LATE LUTEAL PHASE DYSPHORIC DISORDER (PMDD) USING SERTRALINE
- U-462 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND ADULT RHEUMATOID ARTHRITIS AND TREATMENT OF PRIMARY DYSMENORRHEA
- U-463 VENOGRAPHY
- U-464 PERIPHERAL ARTERIOGRAPHY
- U-465 CT IMAGING OF THE HEAD
- U-466 TREATMENT OF IRRITABLE BOWEL SYNDROME
- U-467 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR FOR TREATING HYPERTENSION
- U-468 METHOD OF USING FEXOFENADINE HCL IN TREATING ALLERGIC RHINITIS
- U-469 TREATMENT OF GASTROESOPHAGEAL REFLEX DISEASE (GERD) AND ERADICATION OF H.PYLORI TO REDUCE RISK OF DUODENAL ULCER RECURRENCE
- U-470 THERAPY IN CHRONIC HEPATITIS B VIRUS INFECTION
- U-471 METHOD OF TREATING A PATIENT SUFFERING FROM DIABETES MELLITUS

PATENT AND EXCLUSIVITY TERMS

ADB 27 of 32

PATENT USE

- U-472 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING METHYLPHENIDATE BI-MODAL RELEASE PROFILE EXTENDED-RELEASE CAPSULES
- U-473 TO REDUCE PLASMA CHOLESTEROL LEVELS IN A MAMMAL
- U-474 TO REDUCE PLASMA CHOLESTEROL LEVELS BY ADMIN EZETIMIBE IN COMBO WITH CHOLESTEROL BIOSYNTHESIS INHIB SELECTED FROM GROUP CONSISTING OF HMG COA REDUCTASE INHIBITORS INCL SIMVASTATIN
- U-475 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-476 METHOD OF TREATING ANDROGEN RESPONSIVE/MEDIATED CONDITION IN MAMMAL BY ADMIN A SAFE, EFFECTIVE AMOUNT OF DUTASTERIDE OR PHARMACEUTICALLY ACCEPTABLE DERIVATIVE THEREOF..CONDITIONS INCLUDE BENIGN PROSTATIC HYPERTROPHY
- U-477 METHOD OF INHIBITING 5 ALPHA TESTOSTERONE REDUCTASE ENZYME WITH DUTASTERIDE OR ITS DERIVATIVE AND TREATING ANDROGEN RESPONSIVE/MEDIATED DISEASE INCLUDING BENIGN PROSTATIC HYPERPLASIA
- U-478 METHOD OF TREATING HEPATITIS C VIRAL INFECTION BY CONTINUOUS PARENTERAL ADMIN INTERFERON ALPHA 2-10 MILLION IU WEEKLY, SUBCUTANEOUSLY, INJECTION OF POLYMER-INTERFERON ALPHA CONJUGATE-POLYMER IS PEG-INTERFERON IS ALPHA 2B
- U-479 METHOD OF USING PEG-INTRON/REBETOL COMBINATION THERAPY AND INTRON/REBETOL COMBINATION THERAPY
- U-480 CONTRAST AGENT FOR MRI
- U-481 DISUBSTITUTED ACETYLENES BEARING HETEROAROMATIC AND HETEROBICYCLIC GROUPS HAVING RETINOID-LIKE ACTIVITY
- U-482 METHOD OF IN VITRO FERTILIZATION THERAPY INCLUDING MEANS FOR INDUCING OVULATION....
- U-483 METHOD FOR THE ADMINISTRATION OF DRUGS USING THAT COMPOUND
- U-484 METHOD OF TREATING A SKIN DISEASE WITH A CORTICOSTEROID-CONTAINING PHARMACEUTICAL COMPOSITION
- U-485 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)
- U-486 EXTERNAL PREPARATION FOR APPLICATION TO THE SKIN CONTAINING LIDOCAINE-DRUG RETAINING LAYER PLACED ON SUPPORT AND COMPRISES ADHESIVE GEL BASE 1-10% BY WEIGHT OF LIDOCAINE
- U-487 METHOD AND COMPOSITION FOR REDUCING NERVE INJURY PAIN ASSOCIATED WITH SHINGLES (HERPES ZOSTER AND POST-HERPETIC NEURALGIA)
- U-488 METHOD FOR REDUCING THE PAIN ASSOCIATED WITH HERPES-ZOSTER AND POST-HERPETIC NEURALGIA
- U-489 EXPECTORANT
- U-490 TESTOSTERONE REPLACEMENT THERAPY IN MALES FOR CONDITIONS ASSOCIATED WITH A DEFICIENCY OR ABSENCE OF ENDOGENOUS TESTOSTERONE
- U-491 METHOD OF DELIVERING A DRUG TO THE LUNG
- U-492 METHOD FOR THE TREATMENT OF SKIN, SUFFERING FROM A CONDITION SELECTED FROM A GROUP CONSISTING OF NONACNE INFLAMMATORY DERMATOSES... COMPRISING APPLYING TO AFFECTED AREA. A THERAPEUTICALLY EFFECTIVE AMT AZELAIC ACID
- U-493 TREATMENT OF TYPE 2 DIABETES MELLITUS
- U-494 TREATMENT OF ATTENTION-DEFICIT HYPERACTIVITY DISORDER
- U-495 PERITONEAL DIALYSIS SOLUTION
- U-496 METHOD FOR TREATING CHRONIC RENAL FAILURE
- U-497 RELIEF OF THE SIGNS AND SYMPTOMS OF OSTEOARTHRITIS AND RHEUMATOID ARTHRITIS
- U-498 INTRA-ARTERIAL AND INTRAVENOUS USES OF ULTRAVIST
- U-499 METHOD OF USING REBETOL CAPSULES IN COMBINATION WITH A CONJUGATE COMPRISING POLYETHYLENE GLYCOL(PEG) AND AN ALPHA INTERFERON, INCLUDING, FOR EXAMPLE, PEG-INTRON POWDER FOR INJECTION
- U-500 USE AS AN ANTIHYPERTENSIVE AGENT
- U-501 TREATMENT OF RECURRENT HERPES LABIALIS (COLD SORES) IN ADULTS
- U-502 PITYRIASIS VERSICOLOR
- U-503 GENERATOR MUST BE USED WITH INFUSION SYSTEM SPECIFICALLY LABELED FOR USE WITH GENERATOR
- U-504 TINEA PEDIS, TINEA CRURIS, TINEA CORPORIS
- U-505 ULTRASOUND CONTRAST AGENT
- U-506 PHARM PRODUCT CONTAINER 1ST CHAMBER IS DISPOSED AQUEOUS DILUENT SOL 2ND CHAMBER PHARM ACTIVE AGENT COMPRISING ACETYLCHOLINE,BUFFER IN 1ST CHAM IS SUFFICIENT TO BUFFER PH OF MIXED SOL RESULTING MIXTURE OF AQUEOUS DILUENT SOL & PHARM ACTIVE..
- U-507 ACROMEGALY IN PATIENTS W/INADEQUATE RESPONSE TO SURGERY AND/OR RADIATION THERAPY AND/OR MEDICAL THERAPIES, OR FOR WHOM THESE THERAPIES ARE NOT APPROPRIATE
- U-508 METHOD OF RELEASING 17-BETA OESTRADIOL PRECURSOR IN A SUBSTANTIALLY ZERO ORDER PATTERN FOR AT LEAST THREE WEEKS
- U-509 TREATMENT OF CUTANEOUS MANIFESTATIONS OF CUTANEOUS T-CELL LYMPHOMA IN PATIENTS WHO ARE REFRACTORY TO AT LEAST ONE PRIOR SYSTEMIC THERAPY
- U-510 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH CUTANEOUS T-CELL LYMPHOMA (STAGE IA AND IB) WHO HAVE REFRACTORY OR PERSISTENT DISEASE AFTER OTHER THERAPIES OR WHO HAVE NOT TOLERATED OTHER THERAPIES

PATENT AND EXCLUSIVITY TERMS

ADB 28 of 32

PATENT USE

U-511 USE OF QUINOLONE COMPOUNDS AGAINST ANAEROBIC PATHOGENIC BACTERIA
U-512 USE OF QUINOLONE COMPOUNDS AGAINST ATYPICAL UPPER RESPIRATORY PATHOGENIC BACTERIA
U-513 METHODS OF USE OF ANTIMICROBIAL COMPOUNDS AGAINST PATHOGENIC MYCOPLASMA BACTERIA
U-514 PREVENTION OF OVULATION IN A WOMAN
U-515 TREATMENT OF MULTIPLE MYELOMA PATIENTS WHO HAVE RECEIVED AT LEAST TWO PRIOR THERAPIES AND HAVE DEMONSTRATED DISEASE PROGRESSION ON THE LAST THERAPY
U-516 METHOD OF TREATING A PSYCHOTIC DISEASE
U-517 STABLE GEL FORMULATION FOR TOPICAL TREATMENT OF SKIN CONDITIONS
U-518 OBSESSIVE COMPULSIVE DISORDER
U-519 POST OPERATIVE NAUSEA AND VOMITING
U-520 PREMENOPAUSAL OSTEOPOROSIS
U-521 METHOD OF USING RIBAVIRIN IN COMBINATION WITH INTRON A (INTERFERON ALPHA-2 B RECOMBINANT) INJECTION TO TREAT PATIENTS WITH CHRONIC HEPATITIS C
U-522 TREATMENT OF CMV RETINITIS BY INTRAVITREAL ADMIN OF A PHOSPHOROTHIOATE OLIGONUCLEOTIDE CAPABLE OF HYBRIDIZING WITH CMV MRNA
U-523 METHOD OF TREATING INFECTION BY CRYPTOSPORIDIUM PARVUM IN AN IMMUNOCOMPROMISED MAMMAL
U-524 METHOD OF TREATING DIARRHEA
U-525 METHOD OF TREATING PARASITIC INFECTIONS
U-526 METHOD OF PROVIDING CONTROLLED RELEASE OF A TREATING AGENT USING A CONTROLLED RELEASE COMPOSITION
U-527 METHOD OF DELIVERING AN ACTIVE INGREDIENT USING A PROGRESSIVE HYDRATION BIOADHESIVE
U-528 PREVENTION OF CHEMOTHERAPY-INDUCED NAUSEA AND VOMITING
U-529 ONCE DAILY TREATMENT OF ASTHMA WITH NEBULIZED BUDESONIDE
U-530 TREATMENT OF HERPES ZOSTER, TREATMENT OF GENITAL HERPES, TREATMENT OF COLD SORES, SUPPRESSION OF GENITAL HERPES IN IMMUNOCOMPETENT AND HIV-INFECTED INDIVIDUALS, REDUCTION OF RISK OF HETEROSEXUAL TRANSMISSION OF GENITAL HERPES
U-531 TREATMENT OF PATIENTS WITH ESSENTIAL HYPERTENSION. MAY BE USED ALONE OR GIVEN WITH OTHER CLASSES OF ANTIHYPERTENSIVES, ESPECIALLY THIAZIDE DERIVATIVES
U-532 TREATMENT OF BRONCHOSPASM ASSOCIATED WITH COPD IN PATIENTS REQUIRING MORE THAN ONE BRONCHO DILATOR
U-533 ERECTILE DYSFUNCTION
U-534 HUMALOG IS AN INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS FOR THE CONTROL OF HYPERGLYCEMIA
U-535 TREATMENT OF SOCIAL ANXIETY DISORDER
U-536 CONTRAST AGENT FOR MAGNETIC RESONANCE IMAGING
U-537 TREATMENT OF CONDITIONS RELATED TO HYPERALDOSTERONISM SUCH AS HYPERTENSION AND CARDIAC INSUFFICIENCY, WITH EPLERENONE
U-538 FIRST LINE TREATMENT OF SEVERE HYPERTENSION, IN PATIENTS WITH HYPERTENSION SEVERE ENOUGH THAT THE VALUE OF ACHIEVING PROMPT BLOOD PRESSURE CONTROL EXCEEDS THE RISK OF INITIATING COMBINATION THERAPY IN THESE PATIENTS
U-539 TREATMENT OF MODERATE TO SEVERE DEMENTIA OF THE ALZHEIMER'S TYPE
U-540 TREATMENT OF FUNGAL INFECTIONS
U-541 METHOD OF TREATMENT OF ADULTS INFECTED WITH HIV-1
U-542 METHOD OF TREATING PATIENT WITH TYPE 2 DIABETES BY ONCE DAILY ADMINISTRATION
U-543 TREATMENT OF SCHIZOPHRENIA
U-544 TREATMENT OF OVERACTIVE BLADDER. TREATMENT OF URINARY INCONTINENCE.
U-545 METHOD FOR THE PREVENTION AND/OR TREATMENT OF THROMBOTIC EPISODES, SUCH AS MYOCARDIAL INFARCTION, IN A HUMAN PATIENT AND METHOD FOR THE PREVENTION OF VENOUS THROMBOSIS IN A POSTOPERATIVE HUMAN PATIENT
U-546 USE OF REPAGLINIDE IN COMBINATION WITH METFORMIN TO LOWER BLOOD GLUCOSE
U-547 MAINTENANCE MONOTHERAPY FOR BIPOLAR DISORDER
U-548 A METHOD OF REDUCING FLUSH IN AN INDIVIDUAL BEING TREATED FOR A LIPIDEMIC DISORDER AND EFFECTIVELY TREATING THE LIPIDEMIC DISORDER
U-549 USE IN THE TREATMENT OF MEN WITH ADVANCED SYMPTOMATIC PROSTATE CANCER
U-550 TREATMENT OF BIPOLAR MANIA AND SCHIZOPHRENIA
U-551 METHOD FOR REDUCING TOXICITY OF ALIMTA TREATED PATIENTS BY ADMINISTERING FOLIC ACID
U-552 TREATMENT OF HYPERTENSION AND HYPERLIPIDEMIA WITH A SINGLE COMPOSITION
U-553 MANAGEMENT OF PAIN AND DISCOMFORT ASSOCIATED WITH PERIODONTAL SCALING AND ROOT PLANNING PROCEDURES BY APPLICATION OF AN EUTECTIC MIXTURE OF LOCAL ANESTHETICS TO PERIODONTAL POCKETS
U-554 TREATING HIV INFECTION WITH INDINAVIR SULFATE IN COMBINATION WITH ANTIRETROVIRAL AGENTS
U-555 TREATMENT OF COMPLICATED URINARY TRACT INFECTIONS AND ACUTE UNCOMPLICATED PYELONEPHRITIS
U-556 USE AS ADJUNCT DIAGNOSTIC FOR SERUM THYROGLOBULIN (TG) TESTING
U-557 NASAL TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS

PATENT AND EXCLUSIVITY TERMS

ADB 29 of 32

PATENT USE

- U-558 INDICATED FOR THE RELIEF OF BRONCHOSPASM IN PATIENTS 2-12 YEARS OF AGE WITH ASTHMA (REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE)
- U-559 METHOD OF DECREASING OR REDUCING PARATHYROID HORMONE LEVEL; METHOD OF MODULATING PARATHYROID HORMONE SECRETION; METHOD OF TREATING HYPERPARATHYROIDISM; METHOD OF REDUCING SERUM IONIZED CALCIUM LEVEL
- U-560 METHOD OF DECREASING PARATHYROID HORMONE LEVEL; METHOD OF TREATING HYPERPARATHYROIDISM
- U-561 COSOPT IS INDICATED FOR THE REDUCTION OF ELEVATED INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN-ANGLE GLAUCOMA OR OCULAR HYPERTENSION WHO ARE INSUFFICIENTLY RESPONSIVE TO BETA BLOCKERS
- U-562 TOPICAL TREATMENT OF CUTANEOUS LESIONS IN PATIENTS WITH AIDS-RELATED KAPOSI'S SARCOMA
- U-563 MARINOL IS INDICATED FOR, INTER ALIA, ANOREXIA ASSOCIATED WITH WEIGHT LOSS IN PATIENTS WITH AIDS
- U-564 TREATMENT OF HIV IN CONCOMITANT THERAPY
- U-565 TREATMENT OF SEASONAL AND PERENNIAL ALLERGIC RHINITIS SYMPTOMS, AND CHRONIC URTICARIA
- U-566 FOR THE LONG-TERM, ONCE-DAILY, MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE (COPD), INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA
- U-567 METHOD OF TREATING INFERTILITY
- U-568 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION
- U-569 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THEREAFTER AN OVULATORY INDUCING AMOUNT OF HCG IS ADMINISTERED
- U-570 METHOD OF USING FSH ALONE (WITHOUT EXOGENOUS LH) IN IN VITRO FERTILIZATION AND WHEREIN THE DAILY AMOUNT OF FSH IS ABOUT 5-10 IU/KG
- U-571 TREATMENT OF AGITATION ASSOCIATED WITH SCHIZOPHRENIA AND BIPOLAR I MANIA
- U-572 INTENSIVE CARE UNIT SEDATION
- U-573 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-574 PROPHYLAXIS AND TREATMENT OF THE NASAL SYMPTOMS OF SEASONAL ALLERGIC RHINITIS AND TREATMENT OF THE NASAL SYMPTOMS OF PERENNIAL ALLERGIC RHINITIS IN ADULTS AND PEDIATRIC PATIENTS 12 YEARS OF AGE AND OLDER
- U-575 LOTEMAX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TREATMENT OF STEROID RESPONSIVE CONDITIONS OF THE PALPEBRAL BULBAR CONJUNCTIVA, CORNEA AND ANTERIOR SEGMENT OF THE GLOBE.
- U-576 ALREX OPHTHALMIC SUSPENSION IS INDICATED FOR THE TEMPORARY RELIEF OF THE SIGNS AND SYMPTOMS OF SEASONAL ALLERGIC CONJUNCTIVITIS.
- U-577 TREATMENT OF BENIGN PROSTATIC HYPERPLASIA WITH FINASTERIDE IN COMBINATION WITH DOXAZOSIN
- U-578 TREATMENT OF COMMUNITY ACQUIRED PNEUMONIA, ACUTE EXACERBATION OF CHRONIC BRONCHITIS, AND ACUTE BACTERIAL SINUSITIS CAUSED BY SUSCEPTIBLE STRAINS OF DESIGNATED MICROORGANISMS IN PATIENTS 18 YEARS AND OLDER.
- U-579 TREATMENT OF EPILEPSY AND/OR MIGRAINE.
- U-580 TREATMENT OF DISORDERS OF THE SEROTONERGIC SYSTEM SUCH AS DEPRESSION AND ANXIETY-RELATED DISORDERS
- U-581 METHOD OF TREATING A CONDITION CAPABLE OF TREATMENT BY INHALATION, E.G. ASTHMA, COMPRISING ADMINISTRATION OF A FORMULATION CLAIMED IN US PATENT NO. 6743413
- U-582 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6253762
- U-583 METHOD FOR THE TREATMENT OF A RESPIRATORY DISORDER, E.G. ASTHMA, COMPRISING ADMINISTERING TO A PATIENT BY INHALATION, A METERED AEROSOL DOSE OF A DRUG FORMULATION FROM THE METERED DOSE INHALER SYSTEM CLAIMED IN US 6546928
- U-584 SINGLE-DOSE ADMINISTRATION BY THE EPIDURAL ROUTE, AT THE LUMBAR LEVEL, FOR THE TREATMENT OF PAIN FOLLOWING MAJOR SURGERY
- U-585 TO PROMOTE WEIGHT GAIN AFTER WEIGHT LOSS IN CERTAIN TYPES OF PATIENTS
- U-586 AN INTERMEDIATE RELEASE NICOTINIC ACID FORMULATION SUITABLE FOR ORAL ADMINISTRATION ONCE-A-DAY AS A SINGLE DOSE FOR TREATING HYPERLIPIDEMIA WITHOUT CAUSING DRUG-INDUCED HEPATOTOXICITY OR ELEVATIONS IN URIC ACID OR GLUCOSE OR BOTH
- U-587 USE OF EPLERENONE IN COMBINATION WITH AN ANGIOTENSIN CONVERTING ENZYME (ACE) INHIBITOR (AND OPTIONALLY A DIURETIC) FOR TREATING CONGESTIVE HEART FAILURE AND HYPERTENSION
- U-588 SHORT-TERM TREATMENT OF ACTIVE DUODENAL ULCER; TREATMENT OF HEARTBURN AND OTHER SYMPTOMS ASSOCIATED WITH GERD; SHORT-TERM TREATMENT OF EROSIIVE ESOPHAGITIS; MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS
- U-589 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING AN EFFECTIVE AMOUNT OF AN AEROSOL COMPOSITION TO A PATIENT FROM A METERED DOSE INHALER SYSTEM AS CLAIMED IN U.S. PATENT NO. 6131966
- U-590 METHOD FOR TREATMENT OF A RESPIRATORY DISORDER, E.G., BRONCHOSPASM, COMPRISING ADMINISTERING TO A PATIENT BY ORAL OR NASAL INHALATION A DRUG FORMULATION BY USING THE METERED DOSE INHALER SYSTEM AS CLAIMED IN US PATENT NO. 6532955
- U-591 TREATMENT OF ATTENTION DEFICIT HYPERACTIVITY DISORDER USING A DOSAGE FORM WHICH PROVIDES ONCE-DAILY ORAL ADMINISTRATION OF A PHENIDATE DRUG

PATENT AND EXCLUSIVITY TERMS

ADB 30 of 32

PATENT USE

- U-592 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFA)
- U-593 TREATMENT OF PRIMARY HYPERCHOLESTEROLEMIA, MIXED HYPERLIPIDEMIA AND/OR HOMOZYGOUS FAMILIAL HYPERCHOLESTEROLEMIA (HOFH)
- U-594 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS
- U-595 35 MG ORALLY ONCE A WEEK FOR PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN; 35 MG ORALLY ONCE A WEEK FOR TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-596 TREATMENT OF HORMONE RECEPTOR POSITIVE METASTATIC BREAST CANCER IN POSTMENOPAUSAL WOMEN WITH DISEASE PROGRESSION FOLLOWING ANTIESTROGEN THERAPY
- U-597 FORTEO IS INDICATED FOR THE TREATMENT OF POST MENOPAUSAL WOMEN WITH OSTEOPOROSIS WHO ARE AT HIGH RISK FOR FRACTURE
- U-598 PROPHYLACTIC TREATMENT OF MIGRAINE
- U-599 METHOD FOR TREATING ALLERGIC CONJUNCTIVITIS
- U-600 A METHOD OF TREATING A PATIENT IN NEED OF OPHTHALMIC ANTIMICROBIAL THERAPY WITH LEVOFLOXACIN
- U-601 TREATMENT OF BIPOLAR DISORDERS
- U-602 SIGNS AND SYMPTOMS OF OSTEOARTHRITIS, RHEUMATOID ARTHRITIS IN ADULTS, AND/OR PAUCIARTICULAR OR POLYARTICULAR COURSE JUVENILE RHEUMATOID ARTHRITIS, ACUTE PAIN IN ADULTS; PRIMARY DYSMENORRHEA; AND/OR ACUTE MIGRAINE ATTACKS IN ADULTS
- U-603 METHOD OF TREATING INFECTIONS COMPRISING ORALLY ADMINISTERING AN EFFECTIVE AMOUNT OF THE FDA APPROVED ORAL SUSPENSION
- U-604 METHOD OF LOWERING BLOOD GLUCOSE BY ONCE DAILY ADMINISTRATION
- U-605 TREATMENT OF MAJOR DEPRESSIVE DISORDER(MDD);ALTHOUGH THE MECHANISM OF THE ANTIDEPRESSANT ACTION OF DULOXETINE IN HUMANS IS UNKNOWN, IT IS BELIEVED TO BE RELATED TO ITS POTENTIATION OF SERATONERGIC AND NORADRENERGIC ACTIVITY IN THE CNS
- U-606 USE OF IRINOTECAN IN COMBINATION WITH 5-FLUOROURACIL AND LEUCOVORIN FOR THE TREATMENT OF METASTATIC COLRECTAL CANCER
- U-607 CANCIDAS IS INDICATED FOR EMPIRICAL THERAPY FOR PRESUMED FUNGAL INFECTIONS IN FEBRILE, NEUTROPENIC PATIENTS.
- U-608 USE OF QUINOLONE COMPOUNDS AGAINST PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-609 USE OF QUINOLONE COMPOUNDS AGAINST QUINOLONE-RESISTANT PNEUMOCOCCAL PATHOGENIC BACTERIA
- U-610 ATROVENT HFA (IPRATROPIUM BROMIDE HFA) INHALATION AEROSOL IS INDICATED AS A BRONCHODILATOR FOR MAINTENANCE TREATMENT OF BRONCHOSPASM ASSOCIATED WITH CHRONIC OBSTRUCTIVE PULMONARY DISEASE, INCLUDING CHRONIC BRONCHITIS AND EMPHYSEMA.
- U-611 METHOD OF USING DESLORATADINE TO TREAT SEASONAL AND PERENNIAL ALLERGIC RHINITIS, PRURITIS, AND CHRONIC IDIOPATHIC URTICARIA IN PATIENTS 2 YEARS OF AGE AND OLDER
- U-612 TREATMENT OF SEASONAL ALLERGY SYMPTOMS WITH NASAL CONGESTION IN ADULTS AND CHILDREN 12 YEARS OF AGE AND OLDER
- U-613 REDUCTION OF SERUM PHOSPHATE
- U-614 TREATMENT OF SEXUAL DYSFUNCTION
- U-615 ADJUNCTIVE THERAPY TO DIET IN ADULTS TO REDUCE LDL-C, TOTAL-C, TRIGLYCERIDES AND APO B, AND INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA OR MIXED DYSLIPIDEMIA (TYPES IIA, IIB) AND TO TREAT HYPERTRIGLYCERIDEMIA (TYPES IV, V)
- U-616 MANAGEMENT OF PERSISTENT, MODERATE TO SEVERE PAIN IN PATIENTS REQUIRING CONTINUOUS, AROUND-THE-CLOCK ANALGESIA WITH A HIGH POTENCY OPIOID FOR AN EXTENDED PERIOD OF TIME GENERALLY WEEKS TO MONTHS OR LONGER
- U-617 TREATMENT OF ACUTE PROMYELOGENOUS LEUKEMIA (APL)
- U-618 USE OF ROSUVASTATIN CALCIUM TO REDUCE ELEVATED TOTAL-C, LDL-C, APOB, NONHDL-C OR TG LEVELS; TO INCREASE HDL-C; AND TO TREAT PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA
- U-619 TREATMENT OF MALIGNANT NEOPLASM
- U-620 TREATMENT OF INSOMNIA
- U-621 METHOD OF TREATING CANCER
- U-622 TREATMENT OF VEGF MEDIATED OCULAR DISEASE.
- U-623 SHORT TERM TREATMENT OF ACTIVE BENIGN GASTRIC ULCER
- U-624 REDUCTION OF RISK OF UPPER GASTROINTESTINAL BLEEDING IN CRITICALLY ILL PATIENTS
- U-625 ALLERGIC RHINITIS OR NASAL POLYPS
- U-626 CLOLAR IS INDICATED FOR THE TREATMENT OF PEDIATRIC PATIENTS 1 TO 21 YEARS OLD WITH RELAPSED OR REFRACTORY ACUTE LYMPHOBLASTIC LEUKEMIA AFTER AT LEAST TWO PRIOR REGIMENS
- U-627 TREATMENT OF PATIENTS USING EXTENDED-RELEASE CARBAMAZEPINE
- U-628 USE OF AVANDIA IN COMBINATION WITH A SULFONYLUREA, AND IN COMBINATION WITH METFORMIN AND A SULFONYLUREA TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-629 METHOD OF INDUCING A HYPNOTIC OR SEDATIVE EFFECT IN A HUMAN BY ADMINISTERING ESZOPICLONE
- U-630 TREATING URINARY INCONTINENCE BY ADMINISTERING AN EXTENDED-RELEASE FORM OF DARIFENACIN
- U-631 TREATING A DISEASE OF ALTERED MOTILITY OR TONE OF SMOOTH MUSCLE BY ADMINISTERING A MUSCARINIC RECEPTOR ANTAGONIZING AMOUNT OF DARIFENACIN

PATENT AND EXCLUSIVITY TERMS

ADB 31 of 32

PATENT USE

- U-632 METHOD OF TREATMENT OF CANCER BY ADMINISTERING PARTICLES OF PACLITAXEL THAT HAVE A PROTEIN COATING
- U-633 METHOD FOR TREATMENT OF TUMORS BY ADMINISTERING PACLITAXEL AT A DOSE IN THE RANGE OF ABOUT 30MG/METER SQUARE TO ABOUT 100MG/METER SQUARE IN A PHARMACEUTICALLY ACCEPTABLE FORMULATION THAT DOES NOT CONTAIN CREMOPHOR
- U-634 METHOD FOR DELIVERY OF A BIOLOGIC (INCLUDING ANTINEOPLASTIC AGENTS) BY ADMINISTERING TO A PATIENT AN EFFECTIVE AMOUNT OF A BIOLOGIC AS A SOLID OR LIQUID WITH A POLYMERIC BIOCOMPATIBLE MATERIAL
- U-635 TREATMENT OF GERD, MAINTENANCE OF HEALING OF EROSIIVE ESOPHAGITIS AND RISK REDUCTION OF NSAID ASSOCIATED GASTRIC ULCERS
- U-636 TREATMENT OR PREVENTION OF BRONCHOSPASM OR ASTHMATIC SYMPTOMS
- U-637 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST
- U-638 TREATMENT OF DIABETES WITH AN AMYLIN AGONIST, INCLUDING WITH INSULIN
- U-639 TREATMENT OF A MAMMAL HAVING A NEED OF OR REDUCED ABILITY TO PRODUCE INSULIN WITH AN INSULIN AND AN AMYLIN SUCH AS PRAMLINTIDE
- U-640 USE OF AN AMYLIN AGONIST TO REDUCE GASTRIC MOTILITY AND TREAT POST PRANDIAL HYPERGLYCEMIA
- U-641 USE OF AN AMYLIN AGONIST HAVING SPECIFIED BINDING ACTIVITY TO REDUCE GASTRIC MOTILITY, INCLUDING USE THROUGH PARENTERAL ADMINISTRATION
- U-642 TREATMENT AND PREVENTION OF OSTEOPOROSIS
- U-643 THE SHORT TERM TREATMENT (UP TO 10 DAYS) IN PTS HAVING GASTROESOPHAGEAL REFLUX DISEASE (GERD) AS AN ALTERNATIVE TO ORAL THERAPY IN PTS WHEN THERAPY WITH NEXIUM CAPSULES IS NOT POSSIBLE OR APPROPRIATE
- U-644 TREATMENT OF SEASONAL ALLERGIC RHINITIS
- U-645 TREATMENT OF ASTHMA
- U-646 METHOD OF TREATING OTITIS
- U-647 TREATMENT OF OSTEOPOROSIS IN POST MENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN WITH OSTEOPOROSIS
- U-648 THE TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN AND/OR THE TREATMENT TO INCREASE BONE MASS IN MEN
- U-649 A METHOD FOR TREATING A TUMOR DISEASE
- U-650 TREATMENT OF ESOPHAGEAL CANDIDIASIS AND PROPHYLAXIS OF CANDIDA INFECTIONS IN HSCT PATIENTS
- U-651 TREATMENT OF ACUTE PROMYELOCYTIC LEUKEMIA (APL)
- U-652 TREATMENT OF CARDIAC ARRHYTHMIA
- U-653 STIMULATING INSULIN RELEASE BY ADMINISTERING EXENATIDE
- U-654 LOWERING PLASMA GLUCAGON IN A SUBJECT IN NEED THEREOF, INCLUDING ONE WITH TYPE 2 DIABETES, BY ADMINISTERING AN EXEDIN OR ANALOG, SUCH AS EXENDIN-4
- U-655 TREATMENT OF MILD TO MODERATE ACTIVE CHROHN'S DISEASE INVOLVING THE ILEUM AND/OR THE ASCENDING COLON AND THE MAINTENANCE OF CLINICAL REMISSION OF MILD TO MODERATE CROHN'S DISEASE INVOLVING THE ILEUM AND/OR ASCENDING COLON FOR UP TO 3 MONTHS
- U-656 REDUCING GASTRIC MOTILITY OR DELAYING GASTRIC EMPTYING BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4
- U-657 PREVENTION OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-658 TREATMENT OF ADVANCED HORMONE-DEPENDENT BREAST CANCER
- U-659 TREATMENT OF LOCALLY ADVANCED OR METASTATIC NON SMALL-CELL LUNG CANCER (NSCLC) AFTER FAILURE OF AT LEAST ONE PRIOR CHEMOTHERAPY REGIMEN
- U-660 TREATMENT OF HYPERTENSION AND TREATMENT OF HEART FAILURE
- U-661 TREATMENT OF SEIZURE DISORDER
- U-662 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- U-663 THE TREATMENT OF UNCOMPLICATED URINARY TRACT INFECTIONS
- U-664 TREATMENT OF CONDITIONS FOR WHICH AN ALDOSTERONE RECEPTOR BLOCKER IS INDICATED, SUCH AS HYPERTENSION, HEART FAILURE, AND POST-MYOCARDIAL INFARCTION
- U-665 METHOD OF USING THE DRUG SUBSTANCE/DRUG PRODUCT FOR ULTRASOUND IMAGING
- U-666 METHOD OF TREATING ADHD
- U-667 MANAGEMENT OF INCONTINENCE; METHOD FOR TREATING INCONTINENCE
- U-668 LEVEMIR IS A LONG-ACTING BASAL INSULIN ANALOG THAT IS INDICATED IN THE TREATMENT OF PATIENTS WITH DIABETES MELLITUS
- U-669 INDICATION OF TYPE II DIABETES
- U-670 TREATMENT OF HIV-1 INFECTION BY THE CO-ADMINISTRATION OF TIPRANAVIR AND RITONAVIR.
- U-671 PREVENTION AND TREATMENT OF SECONDARY HYPERPARATHYROIDISM ASSOCIATED WITH CHRONIC KIDNEY DISEASE (CKD) STAGE 3 AND 4
- U-672 TREATMENT OF INFLAMMATION OR AN INFLAMMATION-ASSOCIATED DISORDER
- U-673 METHOD OF TREATMENT WITH ONCE-DAILY DOSES OF 625MG/5ML
- U-674 METHOD OF TREATING INSOMNIA CHARACTERIZED BY DIFFICULTY WITH SLEEP ONSET
- U-675 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA; RELIEF OF SYMPTOMS OF ALLERGIC RHINITIS

PATENT AND EXCLUSIVITY TERMS

ADB 32 of 32

PATENT USE

- U-676 METHOD OF TREATING ATTENTION DEFICIT DISORDER USING ORAL ADMINISTRATION OF A BI-MODAL OR PULSATILE RELEASE COMPOSITION
- U-677 A METHOD OF TREATING DISEASE AMENABLE TO TREATMENT WITH A PHENIDATE DRUG BY ONCE DAILY ORAL ADMINISTRATION OF AN EXTENDED RELEASE DOSAGE FORM
- U-678 METHOD OF TREATING ATTENTION DEFICIT DISORDER AND/OR ATTENTION DEFICIT HYPERACTIVITY DISORDER
- U-679 ADJUNCT TO DIET AND EXERCISE TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES WHO ARE ALREADY TREATED WITH A PIOGLITAZONE AND METFORMIN
- U-680 A METHOD OF TREATING DYSLIPIDEMIA AND DYSLIPOPROTEINEMIA USING A DOSAGE FORM THAT CAN PROVIDE AN EFFECTIVE AMOUNT OF FENOFIBRATE TO A PATIENT IN A FASTED STATE WHICH IS AT LEAST 90% OF THE AUC AMOUNT PROVIDED BY THE DOSAGE FORM
- U-681 TREATMENT OF PRIMARY IGF-1 DEFICIENCY
- U-682 NON-BENZODIAZEPINE HYPNOTIC AGENT INDICATED FOR TREATMENT OF INSOMNIA, CHARACTERIZED BY DIFFICULTIES WITH SLEEP ONSET AND/OR SLEEP MAINTENANCE
- U-683 PREVENTION OR TREATMENT OF ISCHEMIC HEART DISEASE
- U-684 TREATMENT OF UNCOMPLICATED SKIN MANIFESTATIONS OF CHRONIC IDIOPATHIC URTICARIA IN ADULTS AND CHILDREN 6 YEARS OF AGE AND OLDER
- U-685 EXPECTORANT AND COUGH SUPPRESSANT
- U-686 EXPECTORANT AND NASAL DECONGESTANT
- U-687 REDUCING FOOD INTAKE IN A SUBJECT WITH TYPE 2 DIABETES BY ADMINISTERING AN EXENDIN, SUCH AS EXENDIN-4
- U-688 TREATMENT OF HIV-INFECTION IN COMBINATION WITH OTHER ANTIRETROVIRAL AGENTS
- U-689 TREATMENT OF PATIENTS WITH T-CELL ACUTE LYMPHOBLASTIC LEUKEMIA WHOSE DISEASE HAS NOT RESPONDED TO OR HAS RELAPSED FOLLOWING TREATMENT WITH AT LEAST TWO CHEMOTHERAPY REGIMENS
- U-690 TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-691 USE AS A MONOTHERAPY, IN COMBINATION WITH A SULFONYLUREA, METFORMIN OR INSULIN OR IN COMBINATION WITH A SULFONYLUREA PLUS METFORMIN TO IMPROVE GLYCEMIC CONTROL IN PATIENTS WITH TYPE 2 DIABETES MELLITUS
- U-692 USE OF VALSARTAN TO REDUCE CARDIOVASCULAR MORTALITY IN CLINICALLY STABLE PATIENTS WITH LEFT VENTRICULAR FAILURE OR LEFT VENTRICULAR DYSFUNCTION FOLLOWING MYOCARDIAL INFARCTION
- U-693 THE RECOMMENDED INITIAL DOSE OF EQUETRO IS 400MG/DAY GIVEN IN DIVIDED DOSES, TWICE DAILY. THE DOSE SHOULD BE ADJUSTED IN 200MG DAILY INCREMENTS TO ACHIEVE OPTIMAL CLINICAL RESPONSE.