

# **APPROVED DRUG PRODUCTS**

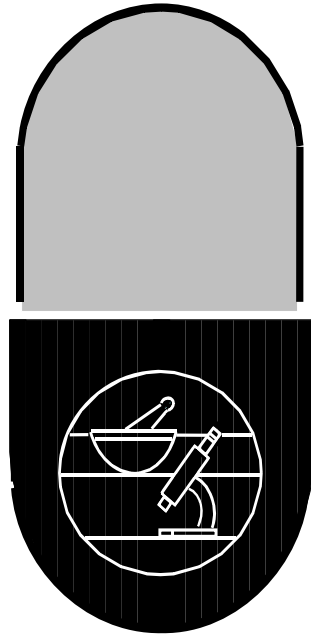
## **With Therapeutic Equivalence Evaluations**



**The "Orange Book"**

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

**CUMULATIVE  
SUPPLEMENT 09  
September 2006**



**APPROVED  
DRUG PRODUCTS**

**WITH  
THERAPEUTIC EQUIVALENCE EVALUATIONS**

**26<sup>th</sup> EDITION**

**Department of Health and Human Services**

**Food and Drug Administration  
Center for Drug Evaluation and Research  
Office of Generic Drugs**

2006

Prepared By  
Office of Generic Drugs  
Center for Drug Evaluation and Research  
Food and Drug Administration

**APPROVED DRUG PRODUCTS  
with  
THERAPEUTIC EQUIVALENCE EVALUATIONS**

**26<sup>th</sup> EDITION**

**Cumulative Supplement 09**

**September 2006**

**CONTENTS**

	<i>PAGE</i>
1.0 INTRODUCTION .....	iii
1.1 How to use the Cumulative Supplement .....	iii
1.2 Cumulative Supplement Content .....	iv
1.3 Applicant Name Changes.....	v
1.4 Availability of the Edition .....	vi
1.5 Report of Counts for the Prescription Drug Product List .....	vii
1.6 Zocor (simvastatin) patent relisting .....	viii
1.7 Levothyroxine Sodium.....	viii
1.8 Cumulative Supplement Legend .....	x
DRUG PRODUCT LISTS	
Prescription Drug Product List .....	1-1
OTC Drug Product List .....	2-1
Drug Products with Approval under Section 505 of the Act	
Administered by the Center for Biologics Evaluation and Research List.....	3-1
Orphan Product Designations and Approvals List .....	4-1
Drug Products Which Must Demonstrate in vivo Bioavailability	
Only if Product Fails to Achieve Adequate Dissolution .....	5-1
PATENT AND EXCLUSIVITY INFORMATION ADDENDUM	
A. Patent and Exclusivity Lists .....	A-1
B. Patent and Exclusivity Terms .....	B-1

**APPROVED DRUG PRODUCTS  
with  
THERAPEUTIC EQUIVALENCE EVALUATIONS**

**26<sup>th</sup> EDITION**

**CUMULATIVE SUPPLEMENT 09  
September 2006**

**1.0 INTRODUCTION**

**1.1 HOW TO USE THE CUMULATIVE SUPPLEMENT**

This Cumulative Supplement is one of a series of monthly updates to the Approved Drug Products with Therapeutic Equivalence Evaluations, 25th Edition (the List). The List is composed of four parts: approved prescription drug products with therapeutic equivalence evaluations; over-the-counter (OTC) drug products that require approved applications as a condition of marketing; drug products with approval under Section 505 of the Act administered by the Center for Biologics Evaluation and Research; and products that have never been marketed, are for exportation, are for military use, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.

The Cumulative Supplement provides, among other things, information on newly approved drugs and, if necessary, revised therapeutic equivalence evaluations and updated patent and exclusivity data. The Addendum contains appropriate drug patent and exclusivity information required of the Agency by the "Drug Price Competition and Patent Term Restoration Act of 1984" for the Prescription, OTC, and Drug Products with Approval under Section 505 of the Act Administered by the Center for Biologics Evaluation and Research Lists.

Because all parts of the publication are subject to changes, additions, or deletions, the List must be used in conjunction with the most current Cumulative Supplement. Users may wish to mark to the left of the ingredient(s) in the List to indicate that changes to that entry appear in the Cumulative Supplement. Drug product information is provided in each Cumulative Supplement for completeness to assist in locating the proper place in the List for the revision.

The presence of any therapeutic equivalence code indicates that the drug product is multisource; the deletion of a therapeutic equivalence code indicates that the drug product has become single source. (An infrequent exception exists when a therapeutic equivalence code is revised. In that case the deletion of the therapeutic equivalence code is followed immediately by the addition of the revised one.)

Products that have never been marketed, are for exportation, are for military use, or have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons, will be flagged in this Cumulative Supplement with the "@" symbol to designate their non-marketed status. All products having a "@" symbol in the 12th Cumulative Supplement of the 25th Edition List will then be added to the "Discontinued Drug Product List" appearing in the 26th Edition. The current edition

Section 2. How To Use The Drug Product Lists describes the layout and usage of the List.

New additions to the Prescription Drug Product List and OTC Drug Product List are indicated by the symbol >A>. The Patent and Exclusivity List new additions are indicated by the symbol >A> to the left of Patent Number or Exclusivity Code. The >A> symbol is then dropped in subsequent Cumulative Supplements for that item.

New deletions to the Prescription Drug Product List and OTC Drug Product List are indicated by the symbol >D> (DELETE) to the left of the line. The information line with the >D> symbol is dropped in subsequent Cumulative Supplements for that item.

The Patent and Exclusivity List is arranged in alphabetical order by active ingredient name(s) and trade name. The trade name will follow the active ingredient name separated by a dash symbol. Also shown is the application number and product number (FDA's internal file number) for reference purposes. All patents with their expiration dates are displayed for each application number. Drug substance and drug product patents are indicated as such with DS or DP in the Patent codes column. Use patents are indicated with the symbol "U" followed by a number representing a specific use. Exclusivity information for a specific drug is indicated by an abbreviation followed by the date upon which the exclusivity expires. Refer to the Exclusivity Terms, Section B, in the Patent and Exclusivity Information Addendum for an explanation of all codes and abbreviations. Refer to Section 1.3 for internet access to the most current list of Patent and Exclusivity terms.

## 1.2 CUMULATIVE SUPPLEMENT CONTENT

Since February 2005, we have been providing daily Electronic Orange Book (EOB) product information for new generic drug approvals. Daily generic updates provide the consumer with the current list of approved generic products which is important for substitution purposes. Previously, a first-time-generic product approved early in the month would not be published in the Cumulative Supplement (CS) for several weeks.

The CS monthly update publish goal is by the end of the following month's second work week (e.g., November's supplement will be updated by the end of the second full work week in December).

Currently, the monthly PDF CS includes:

- Generic product ANDA (Abbreviated New Drug Approval) approvals as of the date of publication.
- We also include all product changes received and processed as of the date of publication.
  - Refer to CS Section 1.8 Cumulative Supplement Legend for types of changes
  - Discontinued products will be processed as of the date of publication. There will be circumstances where a product is discontinued in one month, however, it will be reported in a different month's CS. For example, the Orange Book received a letter November 7 that the product has been discontinued from manufacturing and marketing. The Orange Books subsequently publishes the October CS on November 14. The product will show in the October CS that it is discontinued even though the date of discontinuance is the day that the Orange Book receives notification (November 7).

- New Drug Application (NDA) approvals (20,000 and 50,000 series) appear in the CS month they were approved.
- Patent information, also updated daily in the EOB, is current to the date of publication.
- Exclusivity information is updated monthly and current to the date of publication.

### 1.3 APPLICANT NAME CHANGES

It is not practical to identify in the Cumulative Supplement each and every product involved when an applicant transfers its entire line of approved drug products to another applicant, or when an applicant changes its name. Therefore, the cumulation of these transfers and name changes will be identified in this section only. Where only partial lines of approved products are transferred between applicants, each approved product involved will appear as an applicant name change entry in the Cumulative Supplement.

It is also not practical to identify each and every product involved when an applicant name is changed to meet internal publication standards (e.g., MSD or Zenith [Former Abbreviated Names] are changed, respectively, to Merck Sharp Dohme or Zenith Labs [New Abbreviated Names]). When this occurs, each product involved (either currently in the Cumulative Supplement or in the following year's edition) will reflect the new abbreviated name. Consequently, it will not appear as an applicant name change entry in the Cumulative Supplement nor will the cumulation of these name changes appear in this section. The Electronic Orange Book Query, updated monthly, will contain the most current applicant holder name.

<u>FORMER APPLICANT NAME</u> <u>(FORMER ABBREVIATED NAME)</u>	<u>NEW APPLICANT NAME</u> <u>(NEW ABBREVIATED NAME)</u>
AMERICAN PHARMACEUTICAL PARTNERS INC (AM PHARM PARTNERS)	ABRAXIS PHARMACEUTICAL PRODUCTS (ABRAXIS PHARM)
AMERICAN PHARMACEUTICAL CO INC SUB BURR CORP (AM PHARM)	ABRAXIS PHARMACEUTICAL PRODUCTS (ABRAXIS PHARM)
AMIDE PHARMACEUTICAL INC (AMIDE PHARM)	ACTAVIS TOTOWA LLC (ACTAVIS TOTOWA)
APOTEX CORP (APOTEX)	APOTEX INC ETOBICOKE SITE (APOTEX INC)
APOTEX CORP (APOTEX)	APOTEX INC RICHMOND HILL (APOTEX INC)
APOTEX INC (APOTEX)	APOTEX INC ETOBICOKE SITE (APOTEX INC)
APOTEX INC (APOTEX)	APOTEX INC RICHMOND HILL (APOTEX INC)
APOTEX INC (APOTEX INC)	APOTEX INC ETOBICOKE SITE (APOTEX INC)
APOTEX INC (APOTEX INC)	APOTEX INC RICHMOND HILL (APOTEX INC)
AVENTIS PHARMACEUTICALS INC (AVENTIS)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
AVENTIS PHARMACEUTICAL PRODUCTS INC (AVENTIS PHARMS)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
CLAY PARK LABORATORIES INC (CLAY PARK)	PERRIGO NEW YORK INC (PERRIGO NEW YORK)

CLAY PARK LABS INC (CLAY PARK)	PERRIGO NEW YORK INC (PERRIGO NEW YORK)
DERMIK LABORATORIES INC (DERMIK LABS)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
DERMIK LABORATORIES INC SUB RORER (DERMIK LABS)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
FIRST HORIZON PHARMACEUTICAL COMPANY (FIRST HORIZON)	SCIELE PHARMA INC (SCIELE PHARMA INC)
LOREX PHARMACEUTICALS (LOREX)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
MARTEC PHARMACEUTICALS (MARTEC)	MARTEC USA LLC (MARTEC USA LLC)
MARTEC SCIENTIFIC INC (MARTEC)	MARTEC USA LLC (MARTEC USA LLC)
MCNEIL CONSUMER AND SPECIALTY PHARMACEUTICALS DIV MCNEIL PCC IN (MCNEIL CONS SPECLT)	MCNEIL PEDIATRICS  (MCNEIL PED)
ORPHAN MEDICAL INC (ORPHAN MEDCL)	JAZZ PHARMACEUTICALS (JAZZ)
PHARMACEUTICAL FORMULATIONS INC (PHARM FORM)	LEINER HEALTH PRODUCTS INC (LEINER HLTH PRODS)
PHARMA TEK INC (PHARMA TEK)	X GEN PHARMACEUTICALS INC (X GEN)
PRIVATE FORMULATIONS INC (PRIVATE FMLTNS)	LEINER HEALTH PRODUCTS INC (LEINER HLTH PRODS)
SANKYO PHARMA INC (SANKYO)	DAIICHI SANKYO INC (DAIICHI SANKYO)
SANOFI AVENTIS US INC (SANOFI AVENTIS US)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
SANOFI-AVENTIS US INC (SANOFI AVENTIS)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
SANOFI INC (SANOFI)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
SANOFI SYNTHELABO INC (SANOFI SYNTHELABO)	SANOFI AVENTIS US LLC (SANOFI AVENTIS US)
SANOFI SYNTHELABO RESEARCH DIV SANOFI SYNTHELABO INC (SANOFI SYN RES)	SANOFI AVENTIS US LLC  (SANOFI AVENTIS US)
STERIS LABORATORIES INC (STERIS)	WATSON LABORATORIES INC (WATSON)
TRIGEN LABORATORIES INC (TRIGEN)	JUBILANT PHARMACEUTICALS INC (JUBILANT PHARMS)
UCB PHARMA INC (UCB PHARMA)	UCB INC (UCB INC)

#### 1.4 AVAILABILITY OF THE EDITION

Commencing with the 25th edition, the Annual Edition and monthly Cumulative Supplements will not be available in a published paper version.

Since 1997, the Electronic Orange Book (EOB <http://www.fda.gov/cder/ob/default.htm>), has been available on the internet and has become the updated-every-month Orange Book.

The 25th edition and current monthly supplement are available in an electronic downloadable Portable Document Format (PDF) at the EOB home page by clicking on the Annual Edition. The PDF annual and cumulative supplements will duplicate previous paper versions. Over time, there will be an archive for the annuals and each year's December Cumulative Supplement.

The Electronic Orange Book Query (EOB) is at <http://www.fda.gov/cder/ob/default.htm>. The Query provides searching of the approved drug list by active ingredient, proprietary name, applicant



holder, applicant number or patent number. Product search categories are: prescription, over-the-counter, discontinued drugs. There are links to patent and exclusivity information that may be applicable to each product. Currently, In addition to monthly updates, in the public interest, the EOB is updated on a daily basis with new generic product approval information and new patent information. Current month updates are accomplished by the third week of the following month.

There are historical lists of Orange Book cumulative supplement product monthly changes at <http://www.fda.gov/cder/rxotcdpl/pdplarchive.htm>

There are ASCII text files of the Orange Book drug product, patent, and exclusivity data at <http://www.fda.gov/cder/orange/obreadme.htm>. The drug product text files are zipped into eobzip.exe. The files are updated concurrently with the monthly cumulative supplements. Appendix A and Appendix B text files of the annual Orange Book Edition are updated quarterly.

Effective August 18, 2003, patent submissions for publication in the Orange Book and Docket \*95S-0117 need to be submitted on form FDA-3542 which may be downloaded from the FDA Forms List, <http://www.fda.gov/opacom/morechoices/fdaforms/default.html>.

The current listing of the Orphan Product Designations and Approvals is available at <http://www.fda.gov/orphan/designat/list.htm>.

## 1.5 REPORT OF COUNTS FOR THE PRESCRIPTION DRUG PRODUCT LIST

### DESCRIPTION OF REPORT

This report provides summary counts derived from the product information in the Prescription Drug Product List and the current Cumulative Supplement. Products included in the counts are domestically marketed drug products approved for both safety and effectiveness under section 505 of the Federal Food, Drug, and Cosmetic Act. Excluded are approved drug products marketed by distributors; those marketed solely abroad; and those now regarded as medical devices, biologics or foods.

The baseline column (Dec 2004) refers to the products in the Prescription Drug Product List. For each three-month period, a column of quarterly data is added which incorporates counts of product activity from the previous quarter(s) with those in the baseline count.

### DEFINITIONS

#### Drug Product

For this report, a drug product is the representation in the Prescription Drug Product List of an active moiety (molecular entity and its salts, esters and derivatives) either as a single ingredient or as a combination product provided in a specific dosage form and strength for a given route of administration with approval for marketing by a firm under a particular generic or trade name.

#### New Molecular Entity

A new molecular entity is considered an active moiety that has not previously been approved (either as the parent compound or as a salt, ester or derivative of the parent compound) in the United States for use in a drug product either as a single ingredient or as part of a

combination.

REPORT OF COUNTS FOR THE PRESCRIPTION DRUG PRODUCT LIST  
COUNTS CUMULATIVE BY QUARTER

<u>CATEGORIES COUNTED</u>	<u>DEC 2005</u>	<u>MAR 2006</u>	<u>JUN 2006</u>	<u>SEPT 2006</u>
DRUG PRODUCTS LISTED	11368	11487	11636	
SINGLE SOURCE	2428 (21.4%)	2461 (21.4%)	2447 (21.0%)	
MULTISOURCE	8851 (77.9%)	8937 (77.8%)	9000 (78.2%)	
THERAPEUTICALLY EQUIVALENT	8642 (76.04%)	8730 (76.0%)	8900 (76.5%)	
NOT THERAPEUTICALLY EQUIVALENT	209 (1.8%)	207 (1.8%)	200 (1.7%)	
EXCEPTIONS <sup>1</sup>	89 (0.8%)	89 (0.8%)	89 (0.8%)	
NEW MOLECULAR ENTITIES				
APPROVED	11	6	8	
NUMBER OF APPLICANTS	628	629	642	

<sup>1</sup>Amino acid-containing products of varying composition (see Introduction, page xx of the List).

#### 1.6 ZOCOR (SIMVASTATIN) PATENT RELISTING

U.S. Patent Nos. RE 36481 and RE 36520 are being relisted for Zocor (NDA 19-766) pursuant to the decision and related order in *Ranbaxy Labs. v. Leavitt*, No. 05-1838 (D.D.C. April 30, 2006). The '481 and '520 patents will remain listed in *Approved Drug Products with Therapeutic Equivalence Evaluations* until any applicable periods of exclusivity pursuant to section 505(j)(5)(B)(iv) of the Federal Food, Drug, and Cosmetic Act have been triggered and run, unless the agency's appeal of the decision to the U.S. Court of Appeals for the District of Columbia is decided in the agency's favor before the exclusivity periods have expired. While the patents remain listed, any new or pending ANDA referencing Zocor must contain patent certifications to these patents. For additional information on this matter, please refer to Docket Nos. 2005P-0008 and 2005P-0046.

#### 1.7 LEVOTHYROXINE SODIUM

The Description of Special Situations, Levothyroxine Sodium, published in the 26th Annual Edition of the Orange Book, has been modified in the Cumulative Supplement to include information on a supplemental approval for Genpharm ANDA 76752 approved in 2006. The full discussion as published in the 26th Annual Edition is repeated in the Cumulative Supplement and includes recent approval information on 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), Unithroid (Jerome Stevens NDA 021210) and Levothyroxine Sodium (Genpharm ANDA 76752) 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), Levothyroxine Sodium (Mylan ANDA 076187) and Levothyroxine Sodium (Genpharm ANDA 76752) 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.

Thyro-Tabs (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
LEVOTHYROXINE SODIUM	GENPHARM	0.025MG	AB2	76752	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
LEVOTHYROXINE SODIUM*	GENPHARM	0.025MG	AB3	76752	001
NOVOTHYROX	GENPHARM	0.025MG	BX	21292	001
THYRO-TABS	LLOYD	0.025MG	BX	21116	001
LEVOLET	VINTAGE	0.025MG	BX	21137	001

Trade Name	Applicant	Potency	TE Code	Appl No	Product No
	PHARMS				

\*Revised September 2006

## 1.8 CUMULATIVE SUPPLEMENT LEGEND

The List is sorted by Ingredient(s) and, within each grouping, by the Dosage Form; Route and then by trade name.

The individual product record contains the Therapeutic Equivalence Code, Reference Listed Drug symbol, applicant holder, strength(s), New Drug Application number, product number, and approval date. The last two columns describe the action. The Action Month is the CS month the action occurred. The OB Action is the type of change that has occurred.

New ingredient(s), new dosage form; route(s), new trade names, and new product additions are preceded by >A> during the action month. The change month is the current CS month; the change code for new approvals is NEWA. Following months will display the same information without the >A>.

Changes to currently listed products will list two records. The deleted product record will be preceded by >D>. The product record change addition being made will be preceded by >A>. Following months will display only the >A> record without the >A>. All changes that occur to the product through the Annual year will be listed. The change month and change code will document the change.

The change code and description:

NEWA	New drug product approval usually in the supplement month.
CAHN	Applicant holder firm name has changed.
CAIN	Change. There has been a change in the Ingredient(s) name. All products will be deleted under the old name and all products will be added under the changed ingredient(s) name.
CDFR	Change. Dosage Form; Route of Administration.
CFTG	Change. A first time generic for the innovator product. A TE Code is added.
CMFD	Change. The product is moved from the Discontinued Section due to a change in marketing status.
CMS1	Change. Miscellaneous addition to list.
CMS2	Change. Miscellaneous deletion from list.
CPOT	Change. Potency amount/unit.
CRLD	Change. Reference Listed Drug.
CTEC	Change. Therapeutic Equivalence Code.
CTNA	Change. Trade Name.
DISC	Discontinued. The Rx or OTC listed product is not being marketed and will be moved to the discontinued section in the next edition.

PRESCRIPTION DRUG PRODUCT LIST - 26TH EDITION

RX DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 9 - September 2006

1-1

>D>	<u>ABARELIX</u>								
>D>	INJECTABLE; INTRAMUSCULAR								
>D>	PLENAXIS								
>D>	+	PRAECIS	100MG/VIAL	N21320	001	Nov 25, 2003	Sep	DISC	
>A>		@	100MG/VIAL	N21320	001	Nov 25, 2003	Sep	DISC	
	<u>ACEBUTOLOL HYDROCHLORIDE</u>								
	CAPSULE; ORAL								
	SECTRAL								
AB		DR REDDYS LABS INC	EQ 200MG BASE	N18917	001	Dec 28, 1984	May	CAHN	
AB	+		EQ 400MG BASE	N18917	003	Dec 28, 1984	May	CAHN	
	<u>ACETAMINOPHEN; BUTALBITAL</u>								
	TABLET; ORAL								
	BUTAPAP								
AB	+	MIKART	650MG;50MG	N89988	001	Oct 26, 1992	Jan	CRLD	
		SEDAPAP							
		@ MAYRAND	650MG;50MG	N88944	001	Oct 17, 1985	Jan	DISC	
	<u>ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE</u>								
	TABLET; ORAL								
	ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE								
>D>	+	MIKART	712.8MG;60MG;32MG	N40316	001	Apr 28, 1999	Sep	CFTG	
>A>	AB	+	712.8MG;60MG;32MG	N40316	001	Apr 28, 1999	Sep	CFTG	
		ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITATRATE							
>A>	AB	WEST WARD	712.8MG;60MG;32MG	N40637	001	Sep 22, 2006	Sep	NEWA	
	<u>ACETAMINOPHEN; CODEINE PHOSPHATE</u>								
	SOLUTION; ORAL								
	ACETAMINOPHEN AND CODEINE PHOSPHATE								
AA	+	ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	N85861	001		Jul	CAHN	
		@ CLONMEL	120MG/5ML;12MG/5ML	N40098	001	Sep 20, 1996	Jan	DISC	
	SUSPENSION; ORAL								
	CAPITAL AND CODEINE								
AA		ACTAVIS MID ATLANTIC	120MG/5ML;12MG/5ML	N85883	001		Jul	CAHN	
	TABLET; ORAL								
	ACETAMINOPHEN AND CODEINE PHOSPHATE								
AA	+	TEVA	300MG;60MG	N88629	001	Mar 06, 1985	Apr	CRLD	
		ACETAMINOPHEN W/ CODEINE NO. 3							
		@ ROXANE	300MG;30MG	N84656	001		Jul	DISC	
	<u>ACETAMINOPHEN; HYDROCODONE BITARTRATE</u>								
	TABLET; ORAL								
	HYDROCODONE BITARTRATE AND ACETAMINOPHEN								
		@ ENDO PHARMS	500MG;7.5MG	N40280	001	Sep 30, 1998	Feb	DISC	
		@	650MG;7.5MG	N40280	002	Sep 30, 1998	Feb	DISC	
		@	650MG;10MG	N40280	003	Sep 30, 1998	Feb	DISC	
		@	750MG;7.5MG	N40281	002	Sep 30, 1998	Feb	DISC	
AA		INTERPHARM	325MG;5MG	N40736	001	Aug 25, 2006	Aug	NEWA	
AA			325MG;10MG	N40746	001	Aug 25, 2006	Aug	NEWA	
AA			500MG;5MG	N40729	001	Aug 25, 2006	Aug	NEWA	
AA			500MG;7.5MG	N40748	001	Aug 25, 2006	Aug	NEWA	
AA			650MG;7.5MG	N40754	001	Aug 25, 2006	Aug	NEWA	

## TABLET; ORAL

## HYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA	INTERPHARM	650MG;10MG	N40757	001	Aug 25, 2006	Aug	NEWA	
AA		750MG;7.5MG	N40769	001	Aug 28, 2006	Aug	NEWA	
	MIKART	300MG;5MG	N40658	001	Jan 19, 2006	Jan	NEWA	
	+	300MG;7.5MG	N40556	002	Mar 24, 2006	Mar	NEWA	
AA	VINTAGE PHARMS	325MG;5MG	N40655	001	Jan 19, 2006	Jan	NEWA	
AA		325MG;7.5MG	N40656	001	Jan 19, 2006	Jan	NEWA	
	HY-PHEN							
	@ ASCHER	500MG;5MG	N87677	001	May 03, 1982	Mar	DISC	
	ZYDONE							
	+	ENDO PHARMS	400MG;5MG	N40288	001	Nov 27, 1998	May	CTNA
	+		400MG;7.5MG	N40288	002	Nov 27, 1998	May	CTNA
	+		400MG;10MG	N40288	003	Nov 27, 1998	May	CTNA

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

## SOLUTION; ORAL

## &gt;A&gt; OXYCODONE AND ACETAMINOPHEN

>A>	AA	MALLINCKRODT	325MG/5ML;5MG/5ML	N40680	001	Sep 29, 2006	Sep	NEWA
		ROXICET						
>D>	+	ROXANE	325MG/5ML;5MG/5ML	N89351	001	Dec 03, 1986	Sep	CFTG
>A>	AA	+	325MG/5ML;5MG/5ML	N89351	001	Dec 03, 1986	Sep	CFTG

## TABLET; ORAL

## OXYCODONE AND ACETAMINOPHEN

	+	MIKART	400MG;2.5MG	N40679	001	May 16, 2006	May	NEWA
	+		400MG;5MG	N40687	001	Apr 27, 2006	Apr	NEWA
	+		400MG;7.5MG	N40698	001	Apr 27, 2006	Apr	NEWA
	+		400MG;10MG	N40692	001	Apr 27, 2006	Apr	NEWA
			500MG;10MG	N40676	001	Apr 19, 2006	Apr	NEWA

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

## TABLET; ORAL

## PROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN

AB	ACTAVIS ELIZABETH	650MG;100MG	N70910	001	Jan 02, 1987	Jun	CAHN
	CORNERSTONE	325MG;100MG	N76743	001	May 07, 2004	Jul	CAHN
AB		500MG;100MG	N76750	001	Jun 28, 2004	Jul	CAHN

ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE

## TABLET; ORAL

## TRAMADOL HYDROCHLORIDE AND ACETAMINOPHEN

AB	BARR	325MG;37.5MG	N76914	001	Jul 26, 2006	Jul	NEWA
----	------	--------------	--------	-----	--------------	-----	------

ACETAZOLAMIDE

## TABLET; ORAL

## ACETAZOLAMIDE

AB	+	TARO	250MG	N40195	002	May 28, 1997	Mar	CRLD
		DIAMOX						
		@ DURAMED PHARMS BARR	125MG	N08943	001		Mar	DISC
		@	250MG	N08943	002		Mar	DISC

ACETIC ACID, GLACIAL

## SOLUTION/DROPS; OTIC

## ACETASOL

AT	ACTAVIS MID ATLANTIC	2%	N87146	001		Jul	CAHN
----	----------------------	----	--------	-----	--	-----	------

ACETIC ACID, GLACIAL; HYDROCORTISONE

SOLUTION/DROPS; OTIC

ACETASOL HC

AT	ACTAVIS MID ATLANTIC	2%;1%	N87143 001	Jan 13, 1982	Jul	CAHN
	HYDROCORTISONE AND ACETIC ACID					
AT	VINTAGE	2%;1%	N40609 001	Feb 06, 2006	Jan	NEWA

ACETYLCHOLINE CHLORIDE

FOR SOLUTION; OPHTHALMIC

>D>	MIOCHOL					
>D>	+ NOVARTIS	20MG/VIAL	N16211 001		Sep	DISC
>A>	@	20MG/VIAL	N16211 001		Sep	DISC

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

AB	ACTAVIS ELIZABETH	200MG	N74906 001	Aug 26, 1997	Jun	CAHN
AB	CLONMEL HLTHCARE	200MG	N74833 001	Apr 22, 1997	May	CAHN

SUSPENSION; ORAL

ACYCLOVIR

AB	ACTAVIS MID ATLANTIC	200MG/5ML	N74738 001	Apr 28, 1997	Jul	CAHN
----	----------------------	-----------	------------	--------------	-----	------

TABLET; ORAL

ACYCLOVIR

AB	ACTAVIS ELIZABETH	400MG	N74870 001	Jun 05, 1997	Jun	CAHN
AB		800MG	N74870 002	Jun 05, 1997	Jun	CAHN
AB	CLONMEL HLTHCARE	400MG	N74946 001	Nov 19, 1997	May	CAHN
AB		800MG	N74946 002	Nov 19, 1997	May	CAHN

ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR SODIUM

@	HOSPIRA	EQ 500MG BASE/VIAL	N74663 001	Apr 22, 1997	Jun	DISC
@		EQ 1GM BASE/VIAL	N74663 002	Apr 22, 1997	Jun	DISC

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

@	GENPHARM	0.09MG/INH	N73045 001	Aug 19, 1997	Feb	DISC
@	PLIVA	0.09MG/INH	N74072 001	Aug 01, 1996	Feb	DISC

ALBUTEROL SULFATE

SOLUTION; INHALATION

ALBUTEROL SULFATE

AN	ACTAVIS MID ATLANTIC	EQ 0.083% BASE	N73533 001	Sep 26, 1995	Jul	CAHN
AN	RXELITE	EQ 0.083% BASE	N77569 001	Apr 04, 2006	Mar	NEWA

SYRUP; ORAL

ALBUTEROL SULFATE

AA	ACTAVIS MID ATLANTIC	EQ 2MG BASE/5ML	N74454 001	Sep 25, 1995	Jul	CAHN
AA		EQ 2MG BASE/5ML	N75262 001	Mar 30, 1999	Jul	CAHN

TABLET, EXTENDED RELEASE; ORAL

VOSPIRE ER

>A>	DAVA PHARMS INC	EQ 4MG BASE	N76130 002	Sep 26, 2002	Sep	CAHN
>A>	+	EQ 8MG BASE	N76130 003	Sep 26, 2002	Sep	CAHN
>D>	ODYSSEY PHARMS	EQ 4MG BASE	N76130 002	Sep 26, 2002	Sep	CAHN

TABLET, EXTENDED RELEASE; ORAL

VOSPIRE ER

>D> + ODYSSEY PHARMS EQ 8MG BASE N76130 003 Sep 26, 2002 Sep CAHN

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

+ SANDOZ 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 Jan CAHN

INJECTABLE; IV (INFUSION)

INFUVITE ADULT

+ SANDOZ 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 Jan CAHN

ALPRAZOLAM

TABLET; ORAL

ALPRAZOLAM

AB ACTAVIS ELIZABETH 0.25MG N74342 001 Oct 31, 1993 Jun CAHN  
AB 0.5MG N74342 002 Oct 31, 1993 Jun CAHN  
AB 1MG N74342 003 Oct 31, 1993 Jun CAHN  
AB 2MG N74342 004 Oct 31, 1993 Jun CAHN

TABLET, EXTENDED RELEASE; ORAL

ALPRAZOLAM

AB BARR 0.5MG N77725 001 Jul 31, 2006 Jul NEWA  
AB 1MG N77725 002 Jul 31, 2006 Jul NEWA  
AB 2MG N77725 004 Jul 31, 2006 Aug NEWA  
AB 3MG N77725 003 Jul 31, 2006 Jul NEWA  
AB MYLAN 0.5MG N77391 002 Jan 26, 2006 Jan NEWA  
AB 1MG N77391 003 Jan 26, 2006 Jan NEWA  
AB 2MG N77391 004 Jan 26, 2006 Jan NEWA  
AB 3MG N77391 001 Jan 26, 2006 Jan NEWA  
AB SANDOZ 0.5MG N77777 001 Jun 30, 2006 Jun NEWA  
AB 1MG N77777 002 Jun 30, 2006 Jun NEWA  
AB 2MG N77777 003 Jun 30, 2006 Jun NEWA  
AB 3MG N77777 004 Jun 30, 2006 Jun NEWA

XANAX XR

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

AMANTADINE HYDROCHLORIDE

CAPSULE; ORAL

AMANTADINE HYDROCHLORIDE

AB AMIDE PHARM 100MG N77659 001 Feb 23, 2006 Feb NEWA

SYRUP; ORAL

AMANTADINE HYDROCHLORIDE

AA + ACTAVIS MID ATLANTIC 50MG/5ML N72655 001 Oct 30, 1990 Jul CAHN



AMCINONIDE

## CREAM; TOPICAL

## AMCINONIDE

>D>	AB	ALTANA	0.1%	N76065 001	May 15, 2003	Sep	CRLD
>A>	AB	+	0.1%	N76065 001	May 15, 2003	Sep	CRLD
>D>		CYCLOCORT					
>D>	AB	+	ASTELLAS	0.1%	N18116 002	Sep	DISC
>A>		@	0.1%	N18116 002		Sep	DISC

## LOTION; TOPICAL

## AMCINONIDE

>D>	AB	ALTANA	0.1%	N76329 001	Nov 06, 2002	Sep	CRLD	
>A>		+	0.1%	N76329 001	Nov 06, 2002	Sep	CRLD	
>D>		CYCLOCORT						
>D>	AB	+	ASTELLAS	0.1%	N19729 001	Jun 13, 1988	Sep	DISC
>A>		@	0.1%	N19729 001	Jun 13, 1988	Sep	DISC	

## OINTMENT; TOPICAL

## AMCINONIDE

>D>	AB	ALTANA	0.1%	N76096 001	Nov 19, 2002	Sep	CRLD
>A>	AB	+	0.1%	N76096 001	Nov 19, 2002	Sep	CRLD
>D>		CYCLOCORT					
>D>	AB	+	ASTELLAS	0.1%	N18498 001	Sep	DISC
>A>		@	0.1%	N18498 001		Sep	DISC

AMILORIDE HYDROCHLORIDE

## TABLET; ORAL

## AMILORIDE HYDROCHLORIDE

	+	PAR PHARM	5MG	N70346 001	Jan 22, 1986	Jun	CRLD
		MIDAMOR					
		@ MERCK	5MG	N18200 001		Jun	DISC

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

## TABLET; ORAL

## AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

AB	+	MYLAN	EQ 5MG ANHYDROUS;50MG	N73209 001	Oct 31, 1991	Jun	CRLD
		MODURETIC 5-50					
		@ MERCK	EQ 5MG ANHYDROUS;50MG	N18201 001		Jun	DISC

AMINOPHYLLINE

## SOLUTION; ORAL

## AMINOPHYLLINE DYE FREE

AA	+	ACTAVIS MID ATLANTIC	105MG/5ML	N87727 001	Apr 16, 1982	Jul	CAHN
----	---	----------------------	-----------	------------	--------------	-----	------

AMIODARONE HYDROCHLORIDE

## INJECTABLE; INJECTION

## AMIODARONE HYDROCHLORIDE

AP	+	AKORN	50MG/ML	N76232 001	Jul 05, 2006	Jun	NEWA
----	---	-------	---------	------------	--------------	-----	------

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

## CAPSULE; ORAL

## LOTREL

## NOVARTIS

			EQ 5MG BASE;40MG	N20364 007	Apr 11, 2006	Apr	NEWA
			EQ 10MG BASE;20MG	N20364 005	Jun 20, 2002	Apr	CRLD
	+		EQ 10MG BASE;40MG	N20364 006	Apr 11, 2006	Apr	NEWA

AMMONIUM LACTATE

CREAM; TOPICAL

AMMONIUM LACTATE

AB	PADDOCK	EQ 12% BASE	N76829 001	Feb 07, 2006	Jan	NEWA
----	---------	-------------	------------	--------------	-----	------

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

AB	AM ANTIBIOTICS	250MG	N62058 001		Jan	CAHN
AB		500MG	N62058 002		Jan	CAHN

FOR SUSPENSION; ORAL

AMOXICILLIN

AB	AM ANTIBIOTICS	125MG/5ML	N62059 001		Jan	CAHN
AB		250MG/5ML	N62059 002		Jan	CAHN
AB	HIKMA	125MG/5ML	N65322 002	Jun 19, 2006	Jun	NEWA
AB		200MG/5ML	N65325 002	Jun 19, 2006	Jun	NEWA
AB		250MG/5ML	N65322 001	Jun 19, 2006	Jun	NEWA
AB		400MG/5ML	N65325 001	Jun 19, 2006	Jun	NEWA

TABLET; ORAL

AMOXICILLIN

AB	HIKMA	875MG	N65255 001	Mar 29, 2006	Mar	NEWA
----	-------	-------	------------	--------------	-----	------

AMPHETAMINE ASPARTATE; AMPHETAMINE SULFATE; DEXTROAMPHETAMINE SACCHARATE; DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE

AB	TEVA PHARMS	1.25MG;1.25MG;1.25MG;1.25MG	N40472 001	Sep 30, 2003	May	CAHN
AB		2.5MG;2.5MG;2.5MG;2.5MG	N40472 002	Sep 30, 2003	May	CAHN
AB		5MG;5MG;5MG;5MG	N40472 003	Sep 30, 2003	May	CAHN
AB		7.5MG;7.5MG;7.5MG;7.5MG	N40472 004	Sep 30, 2003	May	CAHN

AMPHOTERICIN B

INJECTABLE; INJECTION

AMPHOTERICIN B

AP	X GEN PHARMS	50MG/VIAL	N63206 001	Apr 29, 1992	Jun	CAHN
----	--------------	-----------	------------	--------------	-----	------

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

AP	SANDOZ	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	N65241 001	Jul 25, 2006	Jul	NEWA
AP		EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	N65310 001	Jul 25, 2006	Jul	NEWA
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	N65241 002	Jul 25, 2006	Jul	NEWA
AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	N65310 002	Jul 25, 2006	Jul	NEWA
AP		EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	N65240 001	Jul 25, 2006	Jul	NEWA

AMPICILLIN/AMPICILLIN TRIHYDRATE

CAPSULE; ORAL

AMPICILLIN TRIHYDRATE

@ AM ANTIBIOTICS

@

	EQ 250MG BASE	N61602 001		Jan	CAHN
	EQ 500MG BASE	N61602 002		Jan	CAHN

FOR SUSPENSION; ORAL

AMPICILLIN TRIHYDRATE

@ AM ANTIBIOTICS

	EQ 125MG BASE/5ML	N61601 001		Jan	CAHN
--	-------------------	------------	--	-----	------

FOR SUSPENSION; ORAL  
 AMPICILLIN TRIHYDRATE  
 @ AM ANTIBIOTICS

EQ 250MG BASE/5ML

N61601 002

Jan CAHN

ANAGRELIDE HYDROCHLORIDE

CAPSULE; ORAL  
 ANAGRELIDE HYDROCHLORIDE

AB ALPHAPHARM

EQ 0.5MG BASE

N77613 001 Jun 27, 2006 Jun NEWA

AB

EQ 1MG BASE

N77613 002 Jun 27, 2006 Jun NEWA

ANIDULAFUNGIN

INJECTABLE; IV (INFUSION)  
 ERAXIS

+ VICURON

50MG/VIAL

N21632 001 Feb 17, 2006 Feb NEWA

ANISINDIONE

TABLET; ORAL  
 MIRADON

@ SCHERING

50MG

N10909 003

Jan DISC

APREPITANT

CAPSULE; ORAL  
 EMEND

MERCK

40MG

N21549 003 Jun 30, 2006 Jun NEWA

ARIPIRAZOLE

&gt;A&gt; INJECTABLE; INTRAMUSCULAR

&gt;A&gt; ABILIFY

&gt;A&gt; + OTSUKA 9.75MG/1.3ML (7.5MG/ML)

N21866 001 Sep 20, 2006 Sep NEWA

TABLET; ORAL

ABILIFY

OTSUKA

2MG

N21436 006 Nov 15, 2002 Jul CMFD

&gt;D&gt; 10MG

N21436 001 Nov 15, 2002 Sep CRLD

&gt;A&gt; + 10MG

N21436 001 Nov 15, 2002 Sep CRLD

TABLET, ORALLY DISINTEGRATING; ORAL

ABILIFY

OTSUKA

10MG

N21729 002 Jun 07, 2006 Jun NEWA

15MG

N21729 003 Jun 07, 2006 Jun NEWA

20MG

N21729 004 Jun 07, 2006 Jun NEWA

+ 30MG

N21729 005 Jun 07, 2006 Jun NEWA

ARTICAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION  
 SEPTOCAINE

DEPROCO

4%;EQ 0.005MG BASE/ML

N22010 001 Mar 30, 2006 Mar NEWA

+ 4%; EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML)

N20971 001 Apr 03, 2000 Mar CPOT

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION  
 SEPTOCAINE

+ DEPROCO

4%;EQ 0.0085MG BASE/1.7ML(4%; EQ 0.005MG BASE/ML)

N22010 001 Mar 30, 2006 Apr CAIN

+ 4%; EQ 0.017MG BASE/1.7ML (4%; EQ 0.01MG BASE/ML)

N20971 001 Apr 03, 2000 Apr CAIN

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

INJECTABLE; IV (INFUSION)

INFUVITE PEDIATRIC

	+	SANDOZ	80MG/VIAL;0.02MG/VIAL;400 IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.1 4MG/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	Jan	CAHN
			INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE)				
	+	SANDOZ	80MG/VIAL;0.02MG/VIAL;400 IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.1 4MG/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	Jan	CAHN

ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM  
CHLORIDE; SODIUM SULFATE

FOR SOLUTION; ORAL

MOVIPREP

>D>	+	NORGINE B V	4.7GM;100GM;1.015GM;5.9GM;2.691GM ;7.5GM	N21881 001	Aug 02, 2006	Sep	CAHN
	+		4.7GM;100GM;1.015GM;5.9GM;2.691GM ;7.5GM	N21881 001	Aug 02, 2006	Aug	NEWA
>A>	+	SALIX PHARMS	4.7GM;100GM;1.015GM;5.9GM;2.691GM ;7.5GM	N21881 001	Aug 02, 2006	Sep	CAHN

ASPIRIN; BUTALBITAL; CAFFEINE

TABLET; ORAL

ASPIRIN AND CAFFEINE W/ BUTALBITAL

AB		ACTAVIS ELIZABETH	325MG;50MG;40MG	N86710 002	Aug 23, 1983	Jun	CAHN
----	--	-------------------	-----------------	------------	--------------	-----	------

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

		@ ENDO PHARMS	325MG;50MG;40MG;30MG	N75351 001	Mar 05, 1999	Feb	DISC
AB	+	WATSON PHARMS	325MG;50MG;40MG;30MG	N19429 003	Oct 26, 1990	Apr	CTNA

ASPIRIN; OXYCODONE HYDROCHLORIDE

TABLET; ORAL

ASPIRIN AND OXYCODONE

	+	ENDO PHARMS	325MG;4.8355MG	N07337 007	Aug 05, 2005	Jun	NEWA
--	---	-------------	----------------	------------	--------------	-----	------

ASPIRIN; OXYCODONE HYDROCHLORIDE; OXYCODONE TEREPHTHALATE

TABLET; ORAL

PERCODAN-DEMI

@ ENDO PHARMS

325MG;2.25MG;0.19MG

N07337 005

Feb DISC

ATENOLOL

TABLET; ORAL

ATENOLOL

AB		UNIQUE PHARM LABS	25MG	N77443 001	Sep 13, 2006	Aug	NEWA
AB			50MG	N77443 002	Sep 13, 2006	Aug	NEWA
AB			100MG	N77443 003	Sep 13, 2006	Aug	NEWA

ATOVAQUONE

TABLET; ORAL

MEPRON

@ GLAXOSMITHKLINE

250MG

N20259 001 Nov 25, 1992 May DISC

&gt;A&gt;

ATROPINE; PRALIDOXIME CHLORIDE

&gt;A&gt;

INJECTABLE; INTRAMUSCULAR

&gt;A&gt;

DUODOTE

&gt;A&gt;

+ MERIDIAN MEDCL

2.1MG/0.7ML;600MG/2ML

N21983 001 Sep 28, 2006 Sep NEWA

AZITHROMYCIN

FOR SUSPENSION; ORAL

AZITHROMYCIN

AB PLIVA

EQ 100MG BASE/5ML

N65246 002 Jul 05, 2006 Jun NEWA

AB

EQ 200MG BASE/5ML

N65246 001 Jul 05, 2006 Jun NEWA

&gt;A&gt;

AB SANDOZ

EQ 100MG BASE/5ML

N65297 001 Sep 18, 2006 Sep NEWA

&gt;A&gt;

AB

EQ 200MG BASE/5ML

N65297 002 Sep 18, 2006 Sep NEWA

ZITHROMAX

AB PFIZER

EQ 100MG BASE/5ML

N50710 001 Oct 19, 1995 Jun CFTG

AB +

EQ 200MG BASE/5ML

N50710 002 Oct 19, 1995 Jun CFTG

BACAMPICILLIN HYDROCHLORIDE

FOR SUSPENSION; ORAL

SPECTROBID

@ PFIZER

125MG/5ML

N50556 001 Mar 23, 1982 Feb DISC

TABLET; ORAL

SPECTROBID

@ PFIZER

400MG

N50520 001 Feb DISC

BACLOFEN

TABLET; ORAL

BACLOFEN

AB CARACO

10MG

N77984 001 Aug 14, 2006 Aug NEWA

AB

20MG

N77862 002 Aug 14, 2006 Aug NEWA

BENAZEPRIL HYDROCHLORIDE

TABLET; ORAL

BENAZEPRIL HYDROCHLORIDE

AB APOTEX INC

5MG

N77128 001 Mar 08, 2006 Feb NEWA

AB

10MG

N77128 002 Mar 08, 2006 Feb NEWA

AB

20MG

N77128 003 Mar 08, 2006 Feb NEWA

AB

40MG

N77128 004 Mar 08, 2006 Feb NEWA

AB BIOKEY

5MG

N76820 001 Feb 03, 2006 Jan NEWA

AB

10MG

N76820 002 Feb 03, 2006 Jan NEWA

AB

20MG

N76820 003 Feb 03, 2006 Jan NEWA

AB

40MG

N76820 004 Feb 03, 2006 Jan NEWA

BENDROFLUMETHIAZIDE

TABLET; ORAL

NATURETIN-5

@ APOTHECON

5MG

N12164 002 Jun DISC

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL  
 BENZPHETAMINE HYDROCHLORIDE  
 AA PADDOCK 50MG N40578 001 Apr 17, 2006 Apr NEWA  
 DIDREX  
 AA + PHARMACIA AND UPJOHN 50MG N12427 002 Apr CFTG

BENZQUINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION  
 EMETE-CON  
 @ PFIZER EQ 50MG BASE/VIAL N16820 001 Mar DISC

BETAINE, ANHYDROUS

FOR SOLUTION; ORAL  
 CYSTADANE  
 + JAZZ 1GM/SCOOPFUL N20576 001 Oct 25, 1996 Feb CAHN

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL  
 ALPHATREX  
 @ SAVAGE LABS EQ 0.05% BASE N19138 001 Jun 26, 1984 Jun DISC  
 BETAMETHASONE DIPROPIONATE  
 AB ACTAVIS MID ATLANTIC EQ 0.05% BASE N70885 001 Feb 03, 1987 Jun CAHN  
 AB + FOUGERA EQ 0.05% BASE N19137 001 Jun 26, 1984 Jun CRLD  
 GEL, AUGMENTED; TOPICAL  
 BETAMETHASONE DIPROPIONATE  
 AB + ALTANA EQ 0.05% BASE N75276 001 May 13, 2003 Aug CRLD  
 DIPROLENE  
 @ SCHERING EQ 0.05% BASE N19408 002 Nov 22, 1991 Aug DISC  
 LOTION; TOPICAL  
 ALPHATREX  
 @ SAVAGE LABS EQ 0.05% BASE N70273 001 Aug 12, 1985 Jun DISC  
 BETAMETHASONE DIPROPIONATE  
 AB ACTAVIS MID ATLANTIC EQ 0.05% BASE N70281 001 Jul 31, 1985 Jul CAHN  
 OINTMENT; TOPICAL  
 BETAMETHASONE DIPROPIONATE  
 AB ACTAVIS MID ATLANTIC EQ 0.05% BASE N71012 001 Feb 03, 1987 Jun CAHN  
 OINTMENT, AUGMENTED; TOPICAL  
 BETAMETHASONE DIPROPIONATE  
 AB ACTAVIS MID ATLANTIC EQ 0.05% BASE N74304 001 Aug 31, 1995 Jun CAHN

BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE

OINTMENT; TOPICAL  
 TACLONEX  
 + LEO PHARM PRODS 0.064%;0.005% N21852 001 Jan 09, 2006 Jan NEWA

BETAMETHASONE DIPROPIONATE; CLOTRIMAZOLE

CREAM; TOPICAL  
 CLOTRIMAZOLE AND BETAMETHASONE DIPROPIONATE  
 AB ACTAVIS MID ATLANTIC EQ 0.05% BASE;1% N76002 001 Aug 02, 2002 Jun CAHN

BETAMETHASONE VALERATE

CREAM; TOPICAL

VALNAC

AB	ACTAVIS MID ATLANTIC	EQ 0.1% BASE	N70050	001	Oct 10, 1984	Jun	CAHN
----	----------------------	--------------	--------	-----	--------------	-----	------

LOTION; TOPICAL

BETAMETHASONE VALERATE

AB	ACTAVIS MID ATLANTIC	EQ 0.1% BASE	N70052	001	Jul 31, 1985	Jul	CAHN
----	----------------------	--------------	--------	-----	--------------	-----	------

OINTMENT; TOPICAL

BETAMETHASONE VALERATE

AB	ACTAVIS MID ATLANTIC	EQ 0.1% BASE	N70051	001	Oct 10, 1984	Jun	CAHN
----	----------------------	--------------	--------	-----	--------------	-----	------

>A> BISKALCITRATE; METRONIDAZOLE; TETRACYCLINE

&gt;A&gt; CAPSULE; ORAL

&gt;A&gt; PYLERA

>A>	+	AXCAN SCANDIPHARM	140MG;125MG;125MG	N50786	001	Sep 28, 2006	Sep	NEWA
-----	---	-------------------	-------------------	--------	-----	--------------	-----	------

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

AB	ACTAVIS ELIZABETH	2.5MG;6.25MG	N75672	001	Sep 25, 2000	Jun	CAHN
----	-------------------	--------------	--------	-----	--------------	-----	------

AB		5MG;6.25MG	N75672	002	Sep 25, 2000	Jun	CAHN
----	--	------------	--------	-----	--------------	-----	------

AB		10MG;6.25MG	N75672	003	Sep 25, 2000	Jun	CAHN
----	--	-------------	--------	-----	--------------	-----	------

BLEOMYCIN SULFATE

INJECTABLE; INJECTION

BLEOMYCIN SULFATE

@	SICOR PHARMS	EQ 15 UNITS BASE/VIAL	N64084	001	Jun 01, 1996	Jun	DISC
---	--------------	-----------------------	--------	-----	--------------	-----	------

@		EQ 30 UNITS BASE/VIAL	N64084	002	Jun 01, 1996	Jun	DISC
---	--	-----------------------	--------	-----	--------------	-----	------

BRIMONIDINE TARTRATE

SOLUTION/DROPS; OPHTHALMIC

ALPHAGAN P

AT	+	ALLERGAN	0.15%	N21262	001	Mar 16, 2001	May	CTEC
----	---	----------	-------	--------	-----	--------------	-----	------

BRIMONIDINE TARTRATE

AT		AKORN	0.2%	N76439	001	Mar 14, 2006	Feb	NEWA
----	--	-------	------	--------	-----	--------------	-----	------

AT		ALCON RES	0.15%	N21764	001	May 22, 2006	May	NEWA
----	--	-----------	-------	--------	-----	--------------	-----	------

BRINZOLAMIDE

SUSPENSION/DROPS; OPHTHALMIC

AZOPT

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

BROMODIPHENHYDRAMINE HYDROCHLORIDE; CODEINE PHOSPHATE

SYRUP; ORAL

AMBENYL

@	FOREST LABS	12.5MG/5ML;10MG/5ML	N09319	006	Jan 10, 1984	Aug	DISC
---	-------------	---------------------	--------	-----	--------------	-----	------

MYBANIL

+	MORTON GROVE	12.5MG/5ML;10MG/5ML	N88626	001	Oct 12, 1984	Aug	CRLD
---	--------------	---------------------	--------	-----	--------------	-----	------

BUDESONIDE

POWDER, METERED; INHALATION

BUDESONIDE

	ASTRAZENECA	0.08MG/INH	N21949	001	Jul 12, 2006	Jul	NEWA
--	-------------	------------	--------	-----	--------------	-----	------

+		0.16MG/INH	N21949	002	Jul 12, 2006	Jul	NEWA
---	--	------------	--------	-----	--------------	-----	------

## SPRAY, METERED; NASAL

## RHINOCORT

	+	ASTRAZENECA	0.032MG/INH	N20746 001	Oct 01, 1999	Mar	CRLD
		@	0.064MG/INH	N20746 002	Oct 01, 1999	Mar	DISC

BUDESONIDE; FORMOTEROL FUMARATE

## SPRAY, METERED; INHALATION

## SYMBICORT

		ASTRAZENECA	0.08MG/INH;EQ 0.045MG BASE	N21929 001	Jul 21, 2006	Jul	NEWA
	+		0.016MG/INH;EQ 0.045MG BASE	N21929 002	Jul 21, 2006	Jul	NEWA

BUMETANIDE

## INJECTABLE; INJECTION

## BUMETANIDE

AP	+	BEDFORD	0.25MG/ML	N74441 001	Jan 27, 1995	Feb	CRLD
		BUMEX					
		@ ROCHE	0.25MG/ML	N18226 001	Feb 28, 1983	Feb	DISC

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

## INJECTABLE; INJECTION

## BUPIVACAINE HYDROCHLORIDE

	+	HOSPIRA	0.5%;EQ 0.009MG BASE/ML	N22046 001	Jul 13, 1983	Aug	CTNA
>A>		BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE					
>A>	AP	SEPTODONT	0.5%;0.0091MG/ML	N77250 001	Sep 27, 2006	Sep	NEWA
		BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE					
	+	HOSPIRA	0.5%;EQ 0.009MG BASE/ML	N22046 001	Jul 13, 1983	Apr	NEWA

BUPROPION HYDROCHLORIDE

## TABLET; ORAL

## BUPROPION HYDROCHLORIDE

AB		APOTEX INC	75MG	N76143 001	Jan 17, 2006	Jan	NEWA
AB			100MG	N76143 002	Jan 17, 2006	Jan	NEWA

BUSPIRONE HYDROCHLORIDE

## TABLET; ORAL

## BUSPIRONE HYDROCHLORIDE

AB		ACTAVIS TOTOWA	5MG	N75388 001	May 09, 2002	Aug	CAHN
AB			10MG	N75388 002	May 09, 2002	Aug	CAHN
AB			15MG	N75388 003	May 09, 2002	Aug	CAHN

BUSULFAN

## INJECTABLE; INJECTION

## BUSULFEX

	+	PDL BIOPHARMA INC	6MG/ML	N20954 001	Feb 04, 1999	Jan	CAHN
--	---	-------------------	--------	------------	--------------	-----	------

CAFFEINE CITRATE

## SOLUTION; INTRAVENOUS

## CAF CIT

>A>	AP	+	MEAD JOHNSON	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N20793 001	Sep 21, 1999	Sep	CFTG
-----	----	---	--------------	------------------------------------	------------	--------------	-----	------

>A>		CAFFEINE CITRATE						
>A>	AP		PHARMAFORCE	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N77233 001	Sep 21, 2006	Sep	NEWA

## SOLUTION; INTRAVENOUS, ORAL

## CAF CIT

>D>		+	MEAD JOHNSON	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N20793 001	Sep 21, 1999	Sep	CFTG
-----	--	---	--------------	------------------------------------	------------	--------------	-----	------



## SOLUTION; ORAL

## CAFCIT

>D>	+	MEAD JOHNSON	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N20793 002	Apr 12, 2000	Sep	CFTG
>A>	AA	+	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N20793 002	Apr 12, 2000	Sep	CFTG
>A>		CAFFEINE CITRATE					
>A>	AA	PHARMAFORCE	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N77304 001	Sep 21, 2006	Sep	NEWA

CAFFEINE; ERGOTAMINE TARTRATE

## SUPPOSITORY; RECTAL

## CAFERGOT

	@	NOVARTIS	100MG;2MG	N09000 002		Feb	DISC
		MIGERGOT					
	+	G AND W LABS	100MG;2MG	N86557 001	Oct 04, 1983	Feb	CRLD

CALCIPOTRIENE

## CREAM; TOPICAL

## DOVONEX

	+	LEO PHARM	0.005%	N20554 001	Jul 22, 1996	Feb	CAHN
--	---	-----------	--------	------------	--------------	-----	------

## OINTMENT; TOPICAL

## DOVONEX

	+	LEO PHARM	0.005%	N20273 001	Dec 29, 1993	Feb	CAHN
--	---	-----------	--------	------------	--------------	-----	------

## SOLUTION; TOPICAL

## DOVONEX

	+	LEO PHARM	0.005%	N20611 001	Mar 03, 1997	Feb	CAHN
--	---	-----------	--------	------------	--------------	-----	------

CALCITONIN SALMON RECOMBINANT

## SPRAY, METERED; NASAL

## FORTICAL

	+	UPSHER SMITH	200 IU/SPRAY	N21406 001	Aug 12, 2005	Jun	CAHN
--	---	--------------	--------------	------------	--------------	-----	------

CALCITONIN, SALMON

## INJECTABLE; INJECTION

## MIACALCIN

	+	NOVARTIS	200 IU/ML	N17808 002	Mar 29, 1991	Jan	CTEC
--	---	----------	-----------	------------	--------------	-----	------

CALCITRIOL

## CAPSULE; ORAL

## CALCITRIOL

AB		ROXANE	0.25UGM	N76917 001	Mar 27, 2006	Mar	NEWA
----	--	--------	---------	------------	--------------	-----	------

## INJECTABLE; INJECTION

## CALCITRIOL

AP		GENIX THERAP	0.001MG/ML	N77102 001	Feb 08, 2006	Jan	NEWA
----	--	--------------	------------	------------	--------------	-----	------

CALCIUM ACETATE

## TABLET; ORAL

## PHOSLO

## @ NABI

		EQ 169MG CALCIUM	N19976 001	Dec 10, 1990	Jun	DISC
--	--	------------------	------------	--------------	-----	------

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE

## SOLUTION; IRRIGATION

>A>		BALANCED SALT SOLUTION					
>A>	AT	AKORN	0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9M G/ML;6.4MG/ML;1.7MG/ML	N75503 001	Sep 27, 2006	Sep	NEWA

## SOLUTION; IRRIGATION

## BSS

>D>	+	ALCON	0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9M	N20742 001	Dec 10, 1997	Sep	CFTG
			G/ML;6.4MG/ML;1.7MG/ML				
>A>	AT	+	0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9M	N20742 001	Dec 10, 1997	Sep	CFTG
			G/ML;6.4MG/ML;1.7MG/ML				

CANDESARTAN CILEXETIL

## TABLET; ORAL

## ATACAND

## ASTRAZENECA

8MG

N20838 002 Jun 04, 1998 May CRLD

CAPTOPRIL

## TABLET; ORAL

## CAPTOPRIL

## @ CLONMEL HLTHCARE

12.5MG

N74423 001 Feb 13, 1996 Jan DISC

## @

25MG

N74423 002 Feb 13, 1996 Jan DISC

## @

50MG

N74423 003 Feb 13, 1996 Jan DISC

## @

100MG

N74423 004 Feb 13, 1996 Jan DISC

## @ ENDO LABS

12.5MG

N74418 001 Feb 13, 1996 Feb DISC

## @

25MG

N74418 002 Feb 13, 1996 Feb DISC

## @

50MG

N74418 003 Feb 13, 1996 Feb DISC

## @

100MG

N74418 004 Feb 13, 1996 Feb DISC

CAPTOPRIL; HYDROCHLOROTHIAZIDE

## TABLET; ORAL

## CAPTOPRIL AND HYDROCHLOROTHIAZIDE

## @ ENDO LABS

25MG;15MG

N74788 001 Dec 29, 1997 Feb DISC

## @

25MG;25MG

N74788 002 Dec 29, 1997 Feb DISC

## @

50MG;15MG

N74788 004 Dec 29, 1997 Feb DISC

## @

50MG;25MG

N74788 003 Dec 29, 1997 Feb DISC

CARBAMAZEPINE

## TABLET, CHEWABLE; ORAL

## CARBAMAZEPINE

AB

JUBILANT PHARMS

100MG

N71940 001 Feb 01, 1988 Jul CAHN

CARBIDOPA; LEVODOPA

## TABLET; ORAL

## CARBIDOPA AND LEVODOPA

AB

ACTAVIS ELIZABETH

10MG;100MG

N74260 001 Sep 03, 1993 Jun CAHN

AB

ACTAVIS ELIZABETH

25MG;100MG

N74260 002 Sep 03, 1993 Jun CAHN

AB

ACTAVIS ELIZABETH

25MG;250MG

N74260 003 Sep 03, 1993 Jun CAHN

CARBOPLATIN

## INJECTABLE; INJECTION

## CARBOPLATIN

AP

WATSON LABS

50MG/VIAL

N77383 001 Jan 27, 2006 Jan NEWA

AP

WATSON LABS

150MG/VIAL

N77383 002 Jan 27, 2006 Jan NEWA

AP

WATSON LABS

450MG/VIAL

N77383 003 Jan 27, 2006 Jan NEWA

## INJECTABLE; IV (INFUSION)

## CARBOPLATIN

## @ AM PHARM

EQ 50MG/5ML (10MG/ML)

N77247 001 Oct 21, 2004 Feb DISC

AP

## @

EQ 50MG/5ML (10MG/ML)

N77266 001 Feb 15, 2006 Jan NEWA

## @

EQ 150MG/15ML (10MG/ML)

N77247 002 Oct 21, 2004 Feb DISC

AP

## @

EQ 150MG/15ML (10MG/ML)

N77266 002 Feb 15, 2006 Jan NEWA

## INJECTABLE; IV (INFUSION)

## CARBOPLATIN

AP	AM PHARM	EQ 450MG/45ML (10MG/ML)	N77266 003	Feb 15, 2006	Jan	NEWA
AP		EQ 600MG/60ML (10MG/ML)	N77266 004	Feb 15, 2006	Jan	NEWA
AP	BEDFORD LABS	EQ 600MG/60ML (10MG/ML)	N77244 004	Jan 20, 2006	Jan	NEWA
>A>	AP	DABUR ONCOLOGY PLC	EQ 150MG/15ML (10MG/ML)	N77432 002	Sep 29, 2006	Sep NEWA
>A>	AP		EQ 450MG/45ML (10MG/ML)	N77432 003	Sep 29, 2006	Sep NEWA
>A>	AP		EQ 50MG/5ML (10MG/ML)	N77432 001	Sep 29, 2006	Sep NEWA

CASPOFUNGIN ACETATE

## INJECTABLE; IV (INFUSION)

## CANCIDAS

+	MERCK	50MG/VIAL	N21227 001	Jan 26, 2001	May	CAHN
+		70MG/VIAL	N21227 002	Jan 26, 2001	May	CAHN

CEFACTOR

## TABLET, EXTENDED RELEASE; ORAL

## CEFACTOR

AB	+	PAR PHARM	EQ 500MG BASE	N65057 001	Jan 05, 2001	May CAHN
----	---	-----------	---------------	------------	--------------	----------

CEFADROXIL/CEFADROXIL HEMIHYDRATE

## CAPSULE; ORAL

## CEFADROXIL

AB	+	IVAX PHARMS	EQ 500MG BASE	N62766 001	Mar 03, 1987	Mar CRLD
>A>	AB	ORCHID HLTHCARE	EQ 500MG BASE	N65309 001	Sep 18, 2006	Sep NEWA
AB		SANDOZ	EQ 500MG BASE	N62291 001		Aug CAHN
AB		TEVA PHARMS	EQ 500MG BASE	N65282 001	Jan 20, 2006	Jan NEWA
AB		WESTWARD	EQ 500MG BASE	N65311 001	Feb 07, 2006	Jan NEWA

## DURICEF

@	WARNER CHILCOTT	EQ 500MG BASE	N50512 001		Jan	DISC
---	-----------------	---------------	------------	--	-----	------

## FOR SUSPENSION; ORAL

## CEFADROXIL

+	RANBAXY	EQ 125MG BASE/5ML	N65115 001	Mar 26, 2003	Jul	CRLD
		EQ 125MG BASE/5ML	N65115 001	Mar 26, 2003	Feb	CTEC
AB	TEVA PHARMS	EQ 250MG BASE/5ML	N65278 001	Jan 20, 2006	Jan	NEWA
AB		EQ 500MG BASE/5ML	N65278 002	Jan 20, 2006	Jan	NEWA

## DURICEF

@	WARNER CHILCOTT	EQ 125MG BASE/5ML	N50527 002		Feb	DISC
---	-----------------	-------------------	------------	--	-----	------

## TABLET; ORAL

## CEFADROXIL

AB		HIKMA	EQ 1GM BASE	N65260 001	Mar 30, 2006	Mar NEWA
AB	+	IVAX PHARMS	EQ 1GM BASE	N62774 001	Apr 08, 1987	Feb CRLD
>A>	AB	ORCHID HLTHCARE	EQ 1GM BASE	N65301 001	Sep 18, 2006	Sep NEWA
		DURICEF				
	@	WARNER CHILCOTT	EQ 1GM BASE	N50528 001		Jan DISC

CEFAZOLIN SODIUM

## INJECTABLE; INJECTION

## CEFAZOLIN AND DEXTROSE

@	B BRAUN	EQ 500MG BASE/VIAL	N50779 001	Jul 27, 2000	May	DISC
---	---------	--------------------	------------	--------------	-----	------

CEFDINIR

## CAPSULE; ORAL

## CEFDINIR

AB	LUPIN	300MG	N65264 001	May 19, 2006	May	NEWA
----	-------	-------	------------	--------------	-----	------

CAPSULE; ORAL						
OMNICEF						
AB	+	ABBOTT	300MG	N50739	001	Dec 04, 1997 May CFTG
FOR SUSPENSION; ORAL						
CEFDIRINR						
AB		LUPIN	125MG/5ML	N65259	001	May 31, 2006 Jul CAHN
AB		LUPIN (USA)	125MG/5ML	N65259	001	May 31, 2006 May NEWA
OMNICEF						
AB		ABBOTT	125MG/5ML	N50749	001	Dec 04, 1997 May CFTG
<u>CEFOTAXIME SODIUM</u>						
INJECTABLE; INJECTION						
CEFOTAXIME						
AP		WOCKHARDT	EQ 1GM BASE/VIAL	N65197	001	Aug 29, 2006 Aug NEWA
CEFOTAXIME AND DEXTROSE 2.4% IN PLASTIC CONTAINER						
		@ B BRAUN	EQ 2GM BASE	N50792	001	Jul 29, 2004 May DISC
CEFOTAXIME AND DEXTROSE 3.9% IN PLASTIC CONTAINER						
		@ B BRAUN	EQ 1GM BASE	N50792	002	Jul 29, 2004 May DISC
CEFOTAXIME SODIUM						
AP		ORCHID HLTHCARE	EQ 500MG BASE/VIAL	N65290	001	Aug 11, 2006 Aug NEWA
AP			EQ 1GM BASE/VIAL	N65290	002	Aug 11, 2006 Aug NEWA
AP			EQ 1GM BASE/VIAL	N65293	001	Aug 10, 2006 Aug NEWA
AP			EQ 2GM BASE/VIAL	N65290	003	Aug 11, 2006 Aug NEWA
AP			EQ 2GM BASE/VIAL	N65293	002	Aug 10, 2006 Aug NEWA
AP			EQ 10GM BASE/VIAL	N65292	001	Aug 10, 2006 Aug NEWA
<u>CEFOTETAN DISODIUM</u>						
INJECTABLE; INJECTION						
CEFOTAN						
		@ ASTRAZENECA	EQ 1GM BASE/VIAL	N63293	001	Apr 29, 1993 Jun DISC
		@	EQ 2GM BASE/VIAL	N63293	002	Apr 29, 1993 Jun DISC
<u>CEFOXITIN SODIUM</u>						
INJECTABLE; INJECTION						
CEFOXITIN						
AP		ORCHID HLTHCARE	EQ 1GM BASE/VIAL	N65313	001	Jan 23, 2006 Jan NEWA
AP			EQ 2GM BASE/VIAL	N65313	002	Jan 23, 2006 Jan NEWA
AP			EQ 10GM BASE/VIAL	N65312	001	Feb 13, 2006 Jan NEWA
CEFOXITIN AND DEXTROSE IN DUPLEX CONTAINER						
AP		B BRAUN	EQ 1GM BASE/VIAL	N65214	001	Mar 10, 2006 Feb NEWA
AP			EQ 2GM BASE/VIAL	N65214	002	Mar 10, 2006 Feb NEWA
<u>CEFPROZIL</u>						
FOR SUSPENSION; ORAL						
CEFPROZIL						
AB		RANBAXY	125MG/5ML	N65202	001	Jun 30, 2006 Jun NEWA
AB			250MG/5ML	N65202	002	Jun 30, 2006 Jun NEWA
<u>CEFTAZIDIME (ARGININE FORMULATION)</u>						
INJECTABLE; INJECTION						
CEPTAZ						
		@ GLAXOSMITHKLINE	1GM/VIAL	N50646	002	Sep 27, 1990 May DISC
		@	2GM/VIAL	N50646	003	Sep 27, 1990 May DISC
		@	10GM/VIAL	N50646	004	Sep 27, 1990 May DISC

CEFTAZIDIME SODIUM

## INJECTABLE; INJECTION

## CEFTAZIDIME SODIUM IN PLASTIC CONTAINER

@	BAXTER HLTHCARE	EQ 10MG BASE/ML	N63221 001	Apr 29, 1993	Aug	DISC
@		EQ 20MG BASE/ML	N63221 002	Apr 29, 1993	Aug	DISC
@		EQ 40MG BASE/ML	N63221 003	Apr 29, 1993	Aug	DISC

## FORTAZ IN PLASTIC CONTAINER

+	GLAXOSMITHKLINE	EQ 20MG BASE/ML	N50634 002	Apr 28, 1989	Aug	CTEC
+		EQ 40MG BASE/ML	N50634 003	Apr 28, 1989	Aug	CTEC

CEFTRIAZONE SODIUM

## INJECTABLE; IM-IV

## CEFTRIAZONE

AP	AM PHARM PARTNERS	EQ 250MG BASE/VIAL	N65245 001	Feb 15, 2006	Jan	NEWA
AP		EQ 500MG BASE/VIAL	N65245 002	Feb 15, 2006	Jan	NEWA
AP		EQ 1GM BASE/VIAL	N65245 003	Feb 15, 2006	Jan	NEWA
AP		EQ 2GM BASE/VIAL	N65245 004	Feb 15, 2006	Jan	NEWA

## CEFTRIAZONE SODIUM

AP	TEVA	EQ 1GM BASE/VIAL	N65262 001	Jun 29, 2006	Jun	NEWA
AP		EQ 2GM BASE/VIAL	N65262 002	Jun 29, 2006	Jun	NEWA

## INJECTABLE; INJECTION

## CEFTRIAZONE

AP	AM PHARM	EQ 10GM BASE/VIAL	N65252 001	Feb 15, 2006	Jan	NEWA
AP	LUPIN	EQ 10GM BASE/VIAL	N65263 001	Sep 12, 2006	Aug	NEWA
AP	WOCKHARDT	EQ 1GM BASE/VIAL	N65180 001	May 12, 2006	May	NEWA

## CEFTRIAZONE SODIUM

AP	TEVA	EQ 10GM BASE/VIAL	N65274 001	May 01, 2006	Apr	NEWA
----	------	-------------------	------------	--------------	-----	------

CEFUROXIME AXETIL

## TABLET; ORAL

## CEFUROXIME AXETIL

AB	AUROBINDO PHARMA LTD	EQ 125MG BASE	N65308 001	Mar 29, 2006	Mar	NEWA
AB		EQ 250MG BASE	N65308 002	Mar 29, 2006	Mar	NEWA
AB		EQ 500MG BASE	N65308 003	Mar 29, 2006	Mar	NEWA

CEPHALEXIN

## CAPSULE; ORAL

## CEPHALEXIN

AB	HIKMA	EQ 250MG BASE	N65215 001	Jan 24, 2006	Jan	NEWA
AB		EQ 500MG BASE	N65215 002	Jan 24, 2006	Jan	NEWA

## KEFLEX

	ADVANCIS PHARM	EQ 333MG BASE	N50405 004	May 12, 2006	May	NEWA
AB		EQ 500MG BASE	N50405 003		May	CRLD
+		EQ 750MG BASE	N50405 005	May 12, 2006	May	NEWA

## FOR SUSPENSION; ORAL

## CEPHALEXIN

AB	ORCHID HLTHCARE	EQ 125MG BASE/5ML	N65326 001	Jul 10, 2006	Jun	NEWA
AB		EQ 250MG BASE/5ML	N65326 002	Jul 10, 2006	Jun	NEWA

CEPHRADINE

## CAPSULE; ORAL

## ANSPOR

@	GLAXOSMITHKLINE	250MG	N61859 001		Mar	DISC
@		500MG	N61859 002		Mar	DISC

CETYL ALCOHOL; COLFOSCERIL PALMITATE; TYLOXAPOL

FOR SUSPENSION; INTRATRACHEAL

EXOSURF NEONATAL

@	GLAXOSMITHKLINE	12MG/VIAL;108MG/VIAL;8MG/VIAL	N20044	001	Aug 02, 1990	May	DISC
---	-----------------	-------------------------------	--------	-----	--------------	-----	------

CHLORHEXIDINE GLUCONATE

SOLUTION; DENTAL

CHLORHEXIDINE GLUCONATE

AT	ACTAVIS MID ATLANTIC	0.12%	N74291	001	Dec 28, 1995	Jul	CAHN
----	----------------------	-------	--------	-----	--------------	-----	------

CHLOROPROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

NESACAINE

	+	ABRAXIS BIOSCIENCE	1%	N09435	001		Jul	CAHN
AP			2%	N09435	002		Jul	CAHN
		NESACAINE-MPF						
AP	+	ABRAXIS BIOSCIENCE	2%	N09435	006	May 02, 1996	Jul	CAHN
		@	2%	N09435	003		Jul	CAHN
		@	3%	N09435	004		Jul	CAHN
AP	+		3%	N09435	007	May 02, 1996	Jul	CAHN

CHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

DIUPRES-250

@	MERCK	250MG;0.125MG	N11635	003	Aug 26, 1987	Jun	DISC
---	-------	---------------	--------	-----	--------------	-----	------

DIUPRES-500

@	MERCK	500MG;0.125MG	N11635	006	Aug 26, 1987	Jun	DISC
---	-------	---------------	--------	-----	--------------	-----	------

CHLORPROMAZINE

SUPPOSITORY; RECTAL

THORAZINE

@	GLAXOSMITHKLINE	25MG	N09149	024		May	DISC
---	-----------------	------	--------	-----	--	-----	------

@		100MG	N09149	033		May	DISC
---	--	-------	--------	-----	--	-----	------

CHLORPROMAZINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

THORAZINE

@	GLAXOSMITHKLINE	200MG	N11120	019		May	DISC
---	-----------------	-------	--------	-----	--	-----	------

@		300MG	N11120	020		May	DISC
---	--	-------	--------	-----	--	-----	------

CONCENTRATE; ORAL

CHLORPROMAZINE HYDROCHLORIDE

AA	ACTAVIS MID ATLANTIC	100MG/ML	N86863	001		Jul	CAHN
----	----------------------	----------	--------	-----	--	-----	------

CICLOPIROX

CREAM; TOPICAL

CICLOPIROX

AB	PERRIGO	0.77%	N77364	001	Mar 03, 2006	Mar	CAHN
----	---------	-------	--------	-----	--------------	-----	------

AB	PERRIGO NEW YORK	0.77%	N77364	001	Mar 03, 2006	Feb	NEWA
----	------------------	-------	--------	-----	--------------	-----	------

CILASTATIN SODIUM; IMIPENEM

INJECTABLE; INJECTION

PRIMAXIN

@	MERCK	EQ 750MG BASE/VIAL;750MG/VIAL	N50630	002	Dec 14, 1990	Jun	DISC
---	-------	-------------------------------	--------	-----	--------------	-----	------

CILOSTAZOL

TABLET; ORAL

CILOSTAZOL

AB	MUTUAL PHARM	50MG	N77208 002	Mar 29, 2006	Mar	NEWA
AB		100MG	N77208 001	Mar 29, 2006	Mar	NEWA
AB	MYLAN	50MG	N77323 002	Apr 20, 2006	Apr	NEWA
AB		100MG	N77323 001	Apr 20, 2006	Apr	NEWA

CIMETIDINE

TABLET; ORAL

CIMETIDINE

@ ENDO PHARMS

200MG

N74281 001 May 17, 1994 Feb DISC

@

300MG

N74281 002 May 17, 1994 Feb DISC

@

400MG

N74281 003 May 17, 1994 Feb DISC

@

800MG

N74329 001 May 17, 1994 Feb DISC

CIMETIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

CIMETIDINE HYDROCHLORIDE

@ ENDO PHARMS

EQ 300MG BASE/2ML

N74005 001 Aug 31, 1994 Feb DISC

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

AA + ACTAVIS MID ATLANTIC

EQ 300MG BASE/5ML

N74176 001 Jun 01, 1994 Jul CAHN

@ ENDO PHARMS

EQ 300MG BASE/5ML

N74251 001 Dec 22, 1994 Feb DISC

CIPROFLOXACIN

INJECTABLE; INJECTION

CIPRO

AP + BAYER PHARMS

400MG/40ML (10MG/ML)

N19847 001 Dec 26, 1990 Aug CFTG

AP +

200MG/20ML (10MG/ML)

N19847 002 Dec 26, 1990 Aug CFTG

@

1200MG/120ML (10MG/ML)

N19847 003 Dec 26, 1990 Aug DISC

CIPROFLOXACIN

AP ABRAXIS PHARM

400MG/40ML (10MG/ML)

N76484 002 Aug 28, 2006 Aug NEWA

AP

200MG/20ML (10MG/ML)

N76484 001 Aug 28, 2006 Aug NEWA

AP BEDFORD LABS

400MG/40ML (10MG/ML)

N76992 002 Aug 28, 2006 Aug NEWA

AP

200MG/20ML (10MG/ML)

N76992 001 Aug 28, 2006 Aug NEWA

1200MG/120ML (10MG/ML)

N76993 001 Aug 28, 2006 Aug NEWA

AP HOSPIRA

400MG/40ML (10MG/ML)

N77245 002 Aug 28, 2006 Aug NEWA

AP

200MG/20ML (10MG/ML)

N77245 001 Aug 28, 2006 Aug NEWA

AP SICOR PHARMS

200MG/20ML (10MG/ML)

N77782 001 Aug 28, 2006 Aug NEWA

AP

400MG/40ML (10MG/ML)

N77782 002 Aug 28, 2006 Aug NEWA

CITALOPRAM HYDROBROMIDE

SOLUTION; ORAL

CITALOPRAM HYDROBROMIDE

AA AUROBINDO PHARMA LTD

EQ 10MG BASE/5ML

N77812 001 Aug 28, 2006 Aug NEWA

AA

SILARX

EQ 10MG BASE/5ML

N77629 001 Jun 15, 2006 May NEWA

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

AB ACTAVIS ELIZABETH

EQ 10MG BASE

N77033 001 Oct 28, 2004 Jun CAHN

AB

EQ 20MG BASE

N77033 002 Oct 28, 2004 Jun CAHN

AB

EQ 40MG BASE

N77033 003 Oct 28, 2004 Jun CAHN

&gt;A&gt;

AB INVAGEN PHARMS

EQ 10MG BASE

N77534 001 Oct 03, 2006 Sep NEWA

&gt;A&gt;

AB

EQ 20MG BASE

N77534 002 Oct 03, 2006 Sep NEWA

## TABLET; ORAL

## CITALOPRAM HYDROBROMIDE

>A>	AB	INVAGEN PHARMS	EQ 40MG BASE	N77534 003	Oct 03, 2006	Sep	NEWA
	AB	MUTUAL PHARM	EQ 10MG BASE	N77052 001	Jul 03, 2006	Jun	NEWA
	AB		EQ 20MG BASE	N77052 002	Jul 03, 2006	Jun	NEWA
	AB		EQ 40MG BASE	N77052 003	Jul 03, 2006	Jun	NEWA
	AB	TARO	EQ 10MG BASE	N77278 001	Mar 22, 2006	Mar	NEWA
	AB		EQ 20MG BASE	N77278 002	Mar 22, 2006	Mar	NEWA
	AB		EQ 40MG BASE	N77278 003	Mar 22, 2006	Mar	NEWA
	AB	TEVA PHARMS	EQ 10MG BASE	N77213 001	Mar 31, 2006	Mar	NEWA
	AB		EQ 20MG BASE	N77213 002	Mar 31, 2006	Mar	NEWA
	AB		EQ 40MG BASE	N77213 003	Mar 31, 2006	Mar	NEWA

CITRIC ACID; MAGNESIUM OXIDE; SODIUM CARBONATE

## SOLUTION; IRRIGATION

## IRRIGATING SOLUTION G IN PLASTIC CONTAINER

@ BAXTER HLTHCARE 3.24GM/100ML;380MG/100ML;430MG/100ML N18519 001 Jun 22, 1982 Jun DISC

## UROLOGIC G IN PLASTIC CONTAINER

+ HOSPIRA 3.24GM/100ML;380MG/100ML;430MG/100ML N18904 001 May 27, 1983 Jun CTEC

CITRIC ACID; UREA, C-13

## FOR SOLUTION, TABLET, FOR SOLUTION; ORAL

## IDKIT:HP

>A>		@ BREATHID 2006	N/A,4GM;75MG,N/A	N21314 001	Dec 17, 2002	Sep	CAHN
>D>		@ ORIDION MEDCL 1987	N/A,4GM;75MG,N/A	N21314 001	Dec 17, 2002	Sep	CAHN

CLARITHROMYCIN

## TABLET; ORAL

## CLARITHROMYCIN

	AB	WOCKHARDT	250MG	N65266 001	May 31, 2006	May	NEWA
	AB		500MG	N65266 002	May 31, 2006	May	NEWA

## TABLET, EXTENDED RELEASE; ORAL

## CLARITHROMYCIN

+ RANBAXY 1GM N65210 001 Jan 26, 2005 Apr CRLD

CLEMASTINE FUMARATE

## SYRUP; ORAL

## CLEMASTINE FUMARATE

	AA	ACTAVIS MID ATLANTIC	EQ 0.5MG BASE/5ML	N74075 001	Oct 31, 1993	Jul	CAHN
--	----	----------------------	-------------------	------------	--------------	-----	------

CLINDAMYCIN PHOSPHATE

## SOLUTION; TOPICAL

## CLINDAMYCIN PHOSPHATE

	AT	ACTAVIS MID ATLANTIC	EQ 1% BASE	N62811 001	Sep 01, 1988	Jul	CAHN
	AT	ALTANA	EQ 1% BASE	N65254 001	Feb 14, 2006	Jan	NEWA

CLOBETASOL PROPIONATE

## CREAM; TOPICAL

## CLOBETASOL PROPIONATE

	AB1	ACTAVIS MID ATLANTIC	0.05%	N74139 001	Aug 03, 1994	Jun	CAHN
--	-----	----------------------	-------	------------	--------------	-----	------

## OINTMENT; TOPICAL

## CLOBETASOL PROPIONATE

	AB	ACTAVIS MID ATLANTIC	0.05%	N74128 001	Aug 03, 1994	Jun	CAHN
--	----	----------------------	-------	------------	--------------	-----	------



## SOLUTION; TOPICAL

	CLOBETASOL PROPIONATE							
AT	ACTAVIS MID ATLANTIC	0.05%	N74331	001	Dec 15, 1995	Jul	CAHN	
	@ ALTANA	0.05%	N75391	001	Feb 08, 1999	May	DISC	

## SPRAY; TOPICAL

## CLOBEX

+	GALDERMA LABS LP	0.05%	N21835	001	Oct 27, 2005	Feb	CAHN	
---	------------------	-------	--------	-----	--------------	-----	------	--

CLOFAZIMINE

## CAPSULE; ORAL

## LAMPRENE

+	NOVARTIS	50MG	N19500	002	Dec 15, 1986	Jun	CMFD	
---	----------	------	--------	-----	--------------	-----	------	--

CLONAZEPAM

## TABLET; ORAL

## CLONAZEPAM

AB	ACTAVIS ELIZABETH	0.5MG	N74869	001	Oct 31, 1996	Jun	CAHN	
AB		1MG	N74869	002	Oct 31, 1996	Jun	CAHN	
AB		2MG	N74869	003	Oct 31, 1996	Jun	CAHN	
AB	VINTAGE PHARMS	0.5MG	N77856	001	Jun 28, 2006	Jun	NEWA	
AB		1MG	N77856	002	Jun 28, 2006	Jun	NEWA	
AB		2MG	N77856	003	Jun 28, 2006	Jun	NEWA	

CLONIDINE HYDROCHLORIDE

## TABLET; ORAL

## CLONIDINE HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	0.1MG	N70974	001	Dec 16, 1986	Jun	CAHN	
AB		0.2MG	N70975	001	Dec 16, 1986	Jun	CAHN	
AB		0.3MG	N70976	001	Dec 16, 1986	Jun	CAHN	

CLOPIDOGREL BISULFATE

## TABLET; ORAL

## CLOPIDOGREL BISULFATE

AB	APOTEX	EQ 75MG BASE	N76274	001	Jan 20, 2006	Jan	NEWA	
AB	PLAVIX							
AB	+ SANOFI SYNTHELABO	EQ 75MG BASE	N20839	001	Nov 17, 1997	Jan	CFTG	

CLOZAPINE

## TABLET, ORALLY DISINTEGRATING; ORAL

## FAZACLO ODT

	AVANIR PHARMS	25MG	N21590	001	Feb 10, 2004	Jul	CAHN	
		50MG	N21590	003	Jun 03, 2005	Jul	CAHN	
+		100MG	N21590	002	Feb 10, 2004	Jul	CAHN	

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		Jun	DISC	
--	----------	-----------------	--------	-----	--	-----	------	--

CODEINE PHOSPHATE; PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

## SYRUP; ORAL

## PROMETH VC W/ CODEINE

+	ACTAVIS MID ATLANTIC	10MG/5ML;5MG/5ML;6.25MG/5ML	N88764	001	Oct 31, 1984	Jul	CAHN	
+	ALPHARMA US PHARMS	10MG/5ML;5MG/5ML;6.25MG/5ML	N88764	001	Oct 31, 1984	Jan	CTEC	

## SYRUP; ORAL

PROMETHAZINE VC W/ CODEINE

@ MORTON GROVE	10MG/5ML;5MG/5ML;6.25MG/5ML	N88896 001	Jan 04, 1985	Jan	DISC
----------------	-----------------------------	------------	--------------	-----	------

CODEINE PHOSPHATE; PROMETHAZINE HYDROCHLORIDE

## SYRUP; ORAL

PROMETH W/ CODEINE

AA +	ACTAVIS MID ATLANTIC	10MG/5ML;6.25MG/5ML	N88763 001	Oct 31, 1984	Jul	CAHN
------	----------------------	---------------------	------------	--------------	-----	------

PROMETHAZINE WITH CODEINE SYRUP

AA	VINTAGE	10MG/5ML;6.25MG/5ML	N40650 001	Jan 31, 2006	Jan	NEWA
----	---------	---------------------	------------	--------------	-----	------

CODEINE PHOSPHATE; PSEUDOEPHEDRINE HYDROCHLORIDE; TRIPROLIDINE HYDROCHLORIDE

## SYRUP; ORAL

TRIACIN-C

AA	ACTAVIS MID ATLANTIC	10MG/5ML;30MG/5ML;1.25MG/5ML	N88704 001	Mar 22, 1985	Jul	CAHN
----	----------------------	------------------------------	------------	--------------	-----	------

COLESTIPOL HYDROCHLORIDE

## GRANULE; ORAL

COLESTID

AB	PHARMACIA AND UPJOHN	5GM/SCOOPFUL	N17563 003	Sep 22, 1995	Apr	CFTG
----	----------------------	--------------	------------	--------------	-----	------

AB +		5GM/PACKET	N17563 004	Sep 22, 1995	Apr	CFTG
------	--	------------	------------	--------------	-----	------

COLESTIPOL HYDROCHLORIDE

AB	IMPAX LABS	5GM/SCOOPFUL	N77277 001	May 02, 2006	Apr	NEWA
----	------------	--------------	------------	--------------	-----	------

AB		5GM/PACKET	N77277 002	May 02, 2006	Apr	NEWA
----	--	------------	------------	--------------	-----	------

FLAVORED COLESTID

PHARMACIA AND UPJOHN 5GM/PACKET

N17563 001		Apr	CFTG
------------	--	-----	------

CROMOLYN SODIUM

## CONCENTRATE; ORAL

GASTROCROM

+ AZUR PHARMA	100MG/5ML	N20479 001	Feb 29, 1996	May	CDFR
---------------	-----------	------------	--------------	-----	------

## SOLUTION; INHALATION

CROMOLYN SODIUM

AN	ACTAVIS MID ATLANTIC	10MG/ML	N75067 001	Jul 19, 1999	Jul	CAHN
----	----------------------	---------	------------	--------------	-----	------

## SOLUTION, CONCENTRATE; ORAL

GASTROCROM

+ AZUR PHARMA	100MG/5ML	N20479 001	Feb 29, 1996	Feb	CAHN
---------------	-----------	------------	--------------	-----	------

CYANOCOBALAMIN

## GEL, METERED; NASAL

NASCOBAL

@ QOL MEDCL	0.5MG/INH	N19722 001	Nov 05, 1996	Mar	DISC
-------------	-----------	------------	--------------	-----	------

CYCLOBENZAPRINE HYDROCHLORIDE

## TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

AB	JUBILANT PHARMS	5MG	N77563 001	Apr 19, 2006	Apr	NEWA
----	-----------------	-----	------------	--------------	-----	------

AB		10MG	N77563 002	Apr 19, 2006	Apr	NEWA
----	--	------	------------	--------------	-----	------

CYCLOBENZAPRINE HYDROCHLORIDE

AB	AMIDE PHARM	5MG	N77291 001	Feb 03, 2006	Jan	NEWA
----	-------------	-----	------------	--------------	-----	------

AB	MUTUAL PHARM	5MG	N73541 002	Apr 06, 2006	Mar	NEWA
----	--------------	-----	------------	--------------	-----	------

AB	MYLAN	5MG	N73144 002	Feb 03, 2006	Jan	NEWA
----	-------	-----	------------	--------------	-----	------

AB	SANDOZ	5MG	N72854 002	Feb 03, 2006	Jan	NEWA
----	--------	-----	------------	--------------	-----	------

AB	WATSON LABS	5MG	N71611 002	Feb 03, 2006	Jan	NEWA
----	-------------	-----	------------	--------------	-----	------

		7.5MG	N71611 003	Feb 03, 2006	Jan	NEWA
--	--	-------	------------	--------------	-----	------

TABLET; ORAL					
FLEXERIL					
AB	MCNEIL CONS SPECLT	5MG	N17821 001	Jan	CFTG
<u>CYPROHEPTADINE HYDROCHLORIDE</u>					
SYRUP; ORAL					
CYPROHEPTADINE HYDROCHLORIDE					
AA	+ ACTAVIS MID ATLANTIC	2MG/5ML	N86833 001	Jul	CAHN
AA	+ ALPHARMA US PHARMS	2MG/5ML	N86833 001	Jun	CTEC
AA	LYNE	2MG/5ML	N40668 001	Jun 28, 2006	Jun NEWA
TABLET; ORAL					
CYPROHEPTADINE HYDROCHLORIDE					
	@ TG UNITED LABS	4MG	N88212 001	May 26, 1983	May CAHN
CYPROHEPTADINE HYDROCHLORIDE					
AA	STASON PHARMS	4MG	N40644 001	May 30, 2006	May NEWA
<u>DAPTOMYCIN</u>					
INJECTABLE; IV (INFUSION)					
CUBICIN					
	@ CUBIST	250MG/VIAL	N21572 001	Sep 12, 2003	Jun DISC
<u>DARUNAVIR ETHANOLATE</u>					
TABLET; ORAL					
PREZISTA					
+	TIBOTEC	EQ 300MG BASE	N21976 001	Jun 23, 2006	Jun NEWA
<u>DASATINIB</u>					
TABLET; ORAL					
SPRYCEL					
	BRISTOL MYERS SQUIBB	20MG	N21986 001	Jun 28, 2006	Jun NEWA
		50MG	N21986 002	Jun 28, 2006	Jun NEWA
+		70MG	N21986 003	Jun 28, 2006	Jun NEWA
<u>DECITABINE</u>					
INJECTABLE; INTRAVENOUS					
DACOGEN					
+	MGI PHARMA INC	50MG/VIAL	N21790 001	May 02, 2006	May NEWA
<u>DEFEROXAMINE MESYLATE</u>					
INJECTABLE; INJECTION					
DEFEROXAMINE MESYLATE					
AP	SICOR PHARMS	500MG/VIAL	N76806 001	Mar 31, 2006	Mar NEWA
AP		2GM/VIAL	N76806 002	Mar 31, 2006	Mar NEWA
<u>DEMECLOCYCLINE HYDROCHLORIDE</u>					
TABLET; ORAL					
DECLOMYCIN					
	@ GLADES PHARMS LLC	75MG	N50261 001	Mar	CAHN
AB		150MG	N50261 002	Mar	CAHN
AB	+	300MG	N50261 003	Mar	CAHN
	@ PROTEIN DESIGN LABS	75MG	N50261 001	Feb	CAHN
AB		150MG	N50261 002	Feb	CAHN
AB	+	300MG	N50261 003	Feb	CAHN

DESLORATADINE; PSEUDOEPHEDRINE SULFATE

TABLET, EXTENDED RELEASE; ORAL

CLARINEX-D 12 HOUR

+	SCHERING	2.5MG;120MG	N21313 001	Feb 01, 2006	Feb	NEWA
---	----------	-------------	------------	--------------	-----	------

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DESMOPRESSIN ACETATE

@	BEDFORD	0.004MG/ML	N74575 001	Feb 18, 2000	Jan	DISC
---	---------	------------	------------	--------------	-----	------

DESMOPRESSIN ACETATE PRESERVATIVE FREE

@	BEDFORD	0.004MG/ML	N74574 001	Feb 18, 2000	Jan	DISC
---	---------	------------	------------	--------------	-----	------

TABLET; ORAL

DESMOPRESSIN ACETATE

AB	APOTEX	0.1MG	N77414 001	Mar 07, 2006	Feb	NEWA
----	--------	-------	------------	--------------	-----	------

AB		0.2MG	N77414 002	Mar 07, 2006	Feb	NEWA
----	--	-------	------------	--------------	-----	------

AB	TEVA PHARMS	0.1MG	N77122 001	Jan 25, 2006	Jan	NEWA
----	-------------	-------	------------	--------------	-----	------

AB		0.2MG	N77122 002	Jan 25, 2006	Jan	NEWA
----	--	-------	------------	--------------	-----	------

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-28

MIRCETTE

AB	+	DURAMED	0.15MG,N/A;0.02MG,0.01MG	N20713 001	Apr 22, 1998	Feb	CAHN
----	---	---------	--------------------------	------------	--------------	-----	------

AB	+		0.15MG,N/A;0.02MG,0.01MG	N20713 001	Apr 22, 1998	Feb	CAHN
----	---	--	--------------------------	------------	--------------	-----	------

DESONIDE

&gt;A&gt; AEROSOL, FOAM; TOPICAL

&gt;A&gt; VERDESO

>A>	+	CONNETICS	0.05%	N21978 001	Sep 19, 2006	Sep	NEWA
-----	---	-----------	-------	------------	--------------	-----	------

DEXAMETHASONE

ELIXIR; ORAL

DEXAMETHASONE

AA	+	ACTAVIS MID ATLANTIC	0.5MG/5ML	N84754 001		Jul	CAHN
----	---	----------------------	-----------	------------	--	-----	------

AA	+	ALPHARMA US PHARMS	0.5MG/5ML	N84754 001		Jun	CRLD
----	---	--------------------	-----------	------------	--	-----	------

HEXADROL

@	ORGANON USA INC	0.5MG/5ML	N12674 001		Jun	DISC
---	-----------------	-----------	------------	--	-----	------

MYMETHASONE

AA	+	MORTON GROVE	0.5MG/5ML	N88254 001	Jul 27, 1983	Jun	CRLD
----	---	--------------	-----------	------------	--------------	-----	------

TABLET; ORAL

DECADRON

@	MERCK	0.5MG	N11664 001		Aug	DISC
---	-------	-------	------------	--	-----	------

@		0.75MG	N11664 002		Aug	DISC
---	--	--------	------------	--	-----	------

DEXAMETHASONE

BP	ROXANE	0.5MG	N84611 001		Aug	CTEC
----	--------	-------	------------	--	-----	------

BP		0.75MG	N84613 001		Aug	CTEC
----	--	--------	------------	--	-----	------

HEXADROL

@	ORGANON USA INC	4MG	N12675 010		Jun	DISC
---	-----------------	-----	------------	--	-----	------

DEXAMETHASONE SODIUM PHOSPHATE

INJECTABLE; INJECTION

DEXAMETHASONE

AP	BAXTER HLTHCARE	EQ 10MG PHOSPHATE/ML	N87702 001	Sep 07, 1982	Mar	CAHN
----	-----------------	----------------------	------------	--------------	-----	------

DEXMETHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL  
 FOCALIN XR  
 NOVARTIS 15MG

N21802 004 Aug 01, 2006 Aug NEWA

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL  
 DEXTROAMPHETAMINE SULFATE  
 @ ENDO PHARMS 5MG

N40299 001 May 13, 1999 Feb DISC

DEXTROMETHORPHAN HYDROBROMIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL  
 PROMETH W/ DEXTROMETHORPHAN  
 AA + ACTAVIS MID ATLANTIC 15MG/5ML; 6.25MG/5ML  
 PROMETHAZINE DM  
 AA VINTAGE 15MG/5ML; 6.25MG/5ML

N88762 001 Oct 31, 1984 Jul CAHN

N40649 001 Feb 14, 2006 Jan NEWA

DIAZEPAM

TABLET; ORAL  
 DIAZEPAM  
 AB ACTAVIS ELIZABETH 2MG  
 AB 5MG  
 AB 10MG  
 AB VINTAGE PHARMS 2MG  
 AB 5MG  
 AB 10MG

N70781 001 Mar 19, 1986 Jun CAHN

N70706 001 Mar 19, 1986 Jun CAHN

N70707 001 Mar 19, 1986 Jun CAHN

N77749 001 Mar 31, 2006 Mar NEWA

N77749 002 Mar 31, 2006 Mar NEWA

N77749 003 Mar 31, 2006 Mar NEWA

DICLOFENAC SODIUM

TABLET, DELAYED RELEASE; ORAL  
 DICLOFENAC SODIUM  
 AB ACTAVIS ELIZABETH 50MG  
 AB 75MG  
 AB SANDOZ 75MG  
 VOLTAREN  
 AB + NOVARTIS 75MG  
 TABLET, EXTENDED RELEASE; ORAL  
 DICLOFENAC SODIUM  
 AB ACTAVIS ELIZABETH 100MG

N74514 001 Mar 26, 1996 Jun CAHN

N74514 002 Mar 26, 1996 Jun CAHN

N74394 001 Nov 30, 1995 May CRLD

N19201 003 Jul 28, 1988 May CMFD

N75910 001 Jan 07, 2002 Jun CAHN

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL  
 DIETHYLPROPION HYDROCHLORIDE  
 @ TG UNITED LABS 25MG  
 @ 25MG

N88267 001 Aug 25, 1983 May CAHN

N88268 001 Aug 25, 1983 May CAHN

DIFLORASONE DIACETATE

CREAM; TOPICAL  
 DIFLORASONE DIACETATE  
 BX + ALTANA 0.05%  
 FLORONE  
 @ PHARMACIA AND UPJOHN 0.05%  
 FLORONE E  
 @ PHARMACIA AND UPJOHN 0.05%

N76263 001 Dec 20, 2002 Jan CRLD

N17741 001 Jan DISC

N19259 001 Aug 28, 1985 Jan DISC

## OINTMENT; TOPICAL

	DIFLORASONE DIACETATE							
AB	+ TARO	0.05%	N75331	001	May 14, 1999	Jan	CRLD	
	FLORONE							
	@ PHARMACIA AND UPJOHN	0.05%	N17994	001		Jan	DISC	
	PSORCON							
	@ PHARMACIA AND UPJOHN	0.05%	N19260	001	Aug 28, 1985	Jan	DISC	

DIFLUNISAL

## TABLET; ORAL

	DIFLUNISAL							
AB	+ TEVA	500MG	N73673	001	Jul 31, 1992	Jun	CRLD	
	DOLOBID							
	@ MERCK	250MG	N18445	001	Apr 19, 1982	Jun	DISC	
	@	500MG	N18445	002	Apr 19, 1982	Jun	DISC	

DIGOXIN

## INJECTABLE; INJECTION

	DIGOXIN							
AP	SANDOZ	0.25MG/ML	N40481	001	Aug 21, 2003	Jan	CAHN	

DILTIAZEM HYDROCHLORIDE

## CAPSULE, EXTENDED RELEASE; ORAL

	DILTIAZEM HYDROCHLORIDE							
AB3	ACTAVIS ELIZABETH	120MG	N74984	001	Dec 20, 1999	Jun	CAHN	
AB3		180MG	N74984	002	Dec 20, 1999	Jun	CAHN	
AB3		240MG	N74984	003	Dec 20, 1999	Jun	CAHN	
AB3		300MG	N74984	004	Dec 20, 1999	Jun	CAHN	
AB4	KV PHARM	120MG	N76563	002	Sep 12, 2006	Aug	NEWA	
AB4		180MG	N76563	003	Sep 12, 2006	Aug	NEWA	
AB4		240MG	N76563	004	Sep 12, 2006	Aug	NEWA	
AB4		300MG	N76563	005	Sep 12, 2006	Aug	NEWA	
AB4		360MG	N76563	006	Sep 12, 2006	Aug	NEWA	
AB4		420MG	N76563	001	Sep 12, 2006	Aug	NEWA	
	DILTZAC							
AB4	APOTEX INC	120MG	N76395	001	Feb 01, 2006	Jan	NEWA	
AB4		180MG	N76395	002	Feb 01, 2006	Jan	NEWA	
AB4		240MG	N76395	003	Feb 01, 2006	Jan	NEWA	
AB4		300MG	N76395	004	Feb 01, 2006	Jan	NEWA	
AB4		360MG	N76395	005	Feb 01, 2006	Jan	NEWA	
	TIAZAC							
AB4	+ BIOVAIL	420MG	N20401	006	Oct 16, 1998	Aug	CFTG	

DIMETHYL SULFOXIDE

## SOLUTION; INTRAVESICAL

	RIMSO-50							
AT	+ BIONICHE PHARMA	50%	N17788	001		Jul	CAHN	

DIPYRIDAMOLE

## TABLET; ORAL

	DIPYRIDAMOLE							
AB	AMIDE PHARM	25MG	N40542	001	Apr 21, 2006	Apr	NEWA	
AB		50MG	N40542	002	Apr 21, 2006	Apr	NEWA	
AB		75MG	N40542	003	Apr 21, 2006	Apr	NEWA	
	@ GLENMARK PHARMA	25MG	N88999	001	Feb 05, 1991	Jul	CAHN	

## TABLET; ORAL

	DIPYRIDAMOLE							
AB	GLENMARK PHARMA	50MG	N89000	001	Feb 05, 1991	Jul	CAHN	
	@	75MG	N89001	001	Feb 05, 1991	Jul	CAHN	

DISOPYRAMIDE PHOSPHATE

## CAPSULE; ORAL

## DISOPYRAMIDE PHOSPHATE

	@ IVAX PHARMS	EQ 100MG BASE	N70186	001	Nov 18, 1985	Jan	DISC	
	@	EQ 150MG BASE	N70187	001	Nov 18, 1985	Jan	DISC	
	@ SANDOZ	EQ 100MG BASE	N70470	001	Dec 10, 1985	Jan	DISC	
	@	EQ 150MG BASE	N70471	001	Dec 10, 1985	Jan	DISC	

DOLASETRON MESYLATE

## INJECTABLE; INJECTION

## ANZEMET

+	SANOFI AVENTIS US	EQ 12.5MG BASE/0.625ML (EQ 20MG BASE/ML)	N20624	002	Sep 11, 1997	Mar	CAIN	
+		EQ 100MG BASE/5ML (EQ 20MG BASE/ML)	N20624	001	Sep 11, 1997	Mar	CAIN	
+		EQ 500MG BASE/25ML (EQ 20MG BASE/ML)	N20624	003	Dec 11, 2001	Mar	CAIN	

## TABLET; ORAL

## ANZEMET

	SANOFI AVENTIS US	EQ 50MG BASE	N20623	001	Sep 11, 1997	Mar	CAIN	
+		EQ 100MG BASE	N20623	002	Sep 11, 1997	Mar	CAIN	

DOPAMINE HYDROCHLORIDE

## INJECTABLE; INJECTION

## DOPAMINE HYDROCHLORIDE

	@ HOSPIRA	40MG/ML	N74403	001	May 23, 1996	Apr	DISC	
	@ INTL MEDICATION	40MG/ML	N18014	001		Apr	DISC	
	@ SICOR PHARMS	40MG/ML	N72999	001	Oct 23, 1991	Apr	DISC	
	@	80MG/ML	N73000	001	Oct 23, 1991	Apr	DISC	

DOXAZOSIN MESYLATE

## TABLET; ORAL

## DOXAZOSIN MESYLATE

AB	ACTAVIS ELIZABETH	EQ 1MG BASE	N75574	001	Oct 18, 2000	Jun	CAHN	
AB		EQ 2MG BASE	N75574	002	Oct 18, 2000	Jun	CAHN	
AB		EQ 4MG BASE	N75574	003	Oct 18, 2000	Jun	CAHN	
AB		EQ 8MG BASE	N75574	004	Oct 18, 2000	Jun	CAHN	

DOXYCYCLINE

## CAPSULE; ORAL

## DOXYCYCLINE

	@ PAR PHARM	EQ 75MG BASE	N65055	004	Apr 18, 2005	Mar	DISC	
	@	EQ 150MG BASE	N65055	003	Jul 15, 2005	Mar	DISC	
	RANBAXY	EQ 75MG BASE	N65053	003	Sep 10, 2003	Mar	CTEC	

## CAPSULE, DELAYED RELEASE; ORAL

## ORACEA

+	COLLAGENEX PHARMS	40MG	N50805	001	May 26, 2006	May	NEWA	
---	-------------------	------	--------	-----	--------------	-----	------	--

## TABLET; ORAL

## DOXYCYCLINE

AB	PAR PHARM	EQ 75MG BASE	N65070	003	Dec 30, 2002	May	CFTG	
		EQ 75MG BASE	N65070	003	Dec 30, 2002	Mar	CMFD	
AB	RANBAXY	EQ 50MG BASE	N65356	001	May 31, 2006	May	NEWA	

## TABLET; ORAL

## DOXYCYCLINE

AB	RANBAXY	EQ 75MG BASE	N65356 002	May 31, 2006	May	NEWA
AB		EQ 100MG BASE	N65356 003	May 31, 2006	May	NEWA

DOXYCYCLINE HYCLATE

## CAPSULE, COATED PELLETS; ORAL

## DORYX

>D>	@ FH FAULDING CO LTD	EQ 75MG BASE	N50582 002	Aug 13, 2001	Sep	CAHN
>D>	@	EQ 100MG BASE	N50582 001	Jul 22, 1985	Sep	CAHN

## CAPSULE, DELAYED RELEASE; ORAL

## DORYX

>D>	@ FH FAULDING CO LTD	EQ 75MG BASE	N50582 002	Aug 13, 2001	Sep	CAHN
	@	EQ 75MG BASE	N50582 002	Aug 13, 2001	Apr	DISC
>D>	@	EQ 100MG BASE	N50582 001	Jul 22, 1985	Sep	CAHN
	@	EQ 100MG BASE	N50582 001	Jul 22, 1985	Apr	DISC
>A>	@ MAYNE PHARMA INTL	EQ 75MG BASE	N50582 002	Aug 13, 2001	Sep	CAHN
>A>	@	EQ 100MG BASE	N50582 001	Jul 22, 1985	Sep	CAHN
	@ WARNER CHILCOTT	EQ 100MG BASE	N62653 001	Oct 30, 1985	Apr	DISC

## DOXYCYCLINE HYCLATE

## SANDOZ

		EQ 75MG BASE	N65281 001	Dec 21, 2005	Apr	CTEC
--	--	--------------	------------	--------------	-----	------

+		EQ 100MG BASE	N65281 002	Dec 21, 2005	Apr	CRLD
---	--	---------------	------------	--------------	-----	------

## TABLET; ORAL

## DOXYCYCLINE HYCLATE

AB	PAR PHARM	EQ 20MG BASE	N65287 001	Feb 28, 2006	Feb	NEWA
----	-----------	--------------	------------	--------------	-----	------

DROSPIRENONE; ETHINYL ESTRADIOL

## TABLET; ORAL

## YAZ

+	BERLEX	3MG;0.02MG	N21676 001	Mar 16, 2006	Jun	CAHN
+	BERLEX LABS	3MG;0.02MG	N21676 001	Mar 16, 2006	Mar	NEWA

DYPHYLLINE

## TABLET; ORAL

## DILOR

	@ SAVAGE LABS	200MG	N84514 001		Jun	DISC
--	---------------	-------	------------	--	-----	------

## DILOR-400

	@ SAVAGE LABS	400MG	N84751 001		Jun	DISC
--	---------------	-------	------------	--	-----	------

## LUFYLLIN

## MEDPOINTE PHARM HLC

		200MG	N84566 001		Jun	CTEC
--	--	-------	------------	--	-----	------

+		400MG	N84566 002		Jun	CRLD
---	--	-------	------------	--	-----	------

EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE

## TABLET; ORAL

## ATRIPLA

+	GILEAD	600MG;200MG;300MG	N21937 001	Jul 12, 2006	Jul	NEWA
---	--------	-------------------	------------	--------------	-----	------

EPINEPHRINE; LIDOCAINE HYDROCHLORIDE

## INJECTABLE; INJECTION

## XYLOCAINE W/ EPINEPHRINE

AP	+	ABRAXIS BIOSCIENCE	0.005MG/ML;0.5%	N06488 012		Jul	CAHN
AP	+		0.005MG/ML;1%	N06488 018	Nov 13, 1986	Jul	CAHN
AP	+		0.005MG/ML;1.5%	N06488 017		Jul	CAHN
AP	+		0.005MG/ML;2%	N06488 019	Nov 13, 1986	Jul	CAHN
AP	+		0.01MG/ML;1%	N06488 004		Jul	CAHN



## INJECTABLE; INJECTION

## XYLOCAINE W/ EPINEPHRINE

@	ABRAXIS BIOSCIENCE	0.01MG/ML;2%	N06488 003		Jul	CAHN
@		0.02MG/ML;2%	N06488 005		Jul	CAHN
+	DENTSPLY PHARM	0.02MG/ML;2%	N21381 002		Mar	CRLD

## PATCH; IONTOPHORESIS, TOPICAL

## LIDOSITE TOPICAL SYSTEM KIT

+	VYTERIS	1.05MG/PATCH;100MG/PATCH	N21504 001	May 06, 2004	May	CDFR
---	---------	--------------------------	------------	--------------	-----	------

## SOLUTION; IONTOPHORESIS, TOPICAL

## LIDOCAINE HYDROCHLORIDE AND EPINEPHRINE

+	EMPI	0.01MG/ML;2%	N21486 001	Oct 26, 2004	May	CDFR
---	------	--------------	------------	--------------	-----	------

EPIRUBICIN HYDROCHLORIDE

&gt;A&gt; INJECTABLE; IV (INFUSION)

&gt;A&gt; EPIRUBICIN HYDROCHLORIDE

>A>	+	MAYNE PHARMA USA	50MG/VIAL	N50807 001	Sep 15, 2006	Sep	NEWA
>A>	+		200MG/VIAL	N50807 002	Sep 15, 2006	Sep	NEWA

ERYTHROMYCIN

## SOLUTION; TOPICAL

## A/T/S

AT	TARO PHARMS NORTH	2%	N62405 001	Nov 18, 1982	Feb	CAHN
----	-------------------	----	------------	--------------	-----	------

## SWAB; TOPICAL

## ERYTHROMYCIN

AT	ALTANA	2%	N65320 001	Jul 25, 2006	Jul	NEWA
----	--------	----	------------	--------------	-----	------

ERYTHROMYCIN ETHYLSUCCINATE

## TABLET, CHEWABLE; ORAL

## E.E.S.

@	ABBOTT	EQ 200MG BASE	N50297 002		Aug	DISC
---	--------	---------------	------------	--	-----	------

## ERYPED

@	ABBOTT	EQ 200MG BASE	N50297 003	Jul 05, 1988	Aug	DISC
---	--------	---------------	------------	--------------	-----	------

ESCITALOPRAM OXALATE

## TABLET; ORAL

## ESCITALOPRAM OXALATE

AB	IVAX PHARMS	5MG	N76765 001	May 22, 2006	May	NEWA
AB		10MG	N76765 002	May 22, 2006	May	NEWA
AB		20MG	N76765 003	May 22, 2006	May	NEWA

## LEXAPRO

AB	FOREST LABS	5MG	N21323 001	Aug 14, 2002	May	CFTG
AB		10MG	N21323 002	Aug 14, 2002	May	CFTG
AB	+	20MG	N21323 003	Aug 14, 2002	May	CFTG

ESTAZOLAM

## TABLET; ORAL

## ESTAZOLAM

AB	PAR PHARM	1MG	N74826 001	Jul 03, 1997	Apr	CAHN
AB		2MG	N74826 002	Jul 03, 1997	Apr	CAHN

ESTRADIOL

## FILM, EXTENDED RELEASE; TRANSDERMAL

## CLIMARA

AB2	BERLEX	0.025MG/24HR	N20375 004	Mar 05, 1999	Aug	CRLD
AB		0.0375MG/24HR	N20375 005	May 27, 2003	Aug	CRLD

## FILM, EXTENDED RELEASE; TRANSDERMAL

## CLIMARA

AB	+	BERLEX	0.0375MG/24HR	N20375 005	May 27, 2003	Jul	CFTG
AB2			0.05MG/24HR	N20375 001	Dec 22, 1994	Aug	CRLD
AB			0.06MG/24HR	N20375 006	May 27, 2003	Aug	CRLD
AB	+		0.06MG/24HR	N20375 006	May 27, 2003	Jul	CFTG
AB2			0.075MG/24HR	N20375 003	Mar 23, 1998	Aug	CRLD

## ESTRADERM

BX		NOVARTIS	0.05MG/24HR	N19081 002	Sep 10, 1986	Aug	CRLD
----	--	----------	-------------	------------	--------------	-----	------

## ESTRADIOL

AB		MYLAN TECHNOLOGIES	0.0375MG/24HR	N75182 004	Jul 20, 2006	Jul	NEWA
AB			0.06MG/24HR	N75182 005	Jul 20, 2006	Jul	NEWA

## MENOSTAR

+		BERLEX	0.014MG/24HR	N21674 001	Jun 08, 2004	Jun	CAHN
---	--	--------	--------------	------------	--------------	-----	------

## VIVELLE-DOT

BX		NOVARTIS	0.025MG/24HR	N20538 009	May 03, 2002	Aug	CRLD
BX			0.0375MG/24HR	N20538 005	Jan 08, 1999	Aug	CRLD
AB1			0.05MG/24HR	N20538 006	Jan 08, 1999	Aug	CRLD
BX			0.075MG/24HR	N20538 007	Jan 08, 1999	Aug	CRLD

## GEL; TOPICAL

## ESTROGEL

@		ASCEND	0.06%	N21166 001	Feb 09, 2004	Jan	CAHN
---	--	--------	-------	------------	--------------	-----	------

## GEL, METERED; TOPICAL

## ESTROGEL

+		ASCEND	0.06%	N21166 002	Feb 09, 2004	Jan	CAHN
---	--	--------	-------	------------	--------------	-----	------

## TABLET; ORAL

## ESTRADIOL

@		RADIUS PHARMS	0.5MG	N40275 001	Dec 29, 1998	Aug	CAHN
@			1MG	N40275 002	Dec 29, 1998	Aug	CAHN
@			2MG	N40275 003	Dec 29, 1998	Aug	CAHN

ESTRADIOL HEMIHYDRATE

## EMULSION; TOPICAL

## ESTRASORB

+		ESPRIT PHARMA	0.25%	N21371 001	Oct 09, 2003	Feb	CAHN
---	--	---------------	-------	------------	--------------	-----	------

ESTROGENS, CONJUGATED SYNTHETIC B

## TABLET; ORAL

## ENJUVIA

		DURAMED	0.3MG	N21443 001	Dec 20, 2004	Mar	CMFD
			0.45MG	N21443 002	Dec 20, 2004	Mar	CMFD
			0.625MG	N21443 003	May 10, 2004	Mar	CMFD
+			1.25MG	N21443 004	May 10, 2004	Mar	CMFD

ETHACRYNATE SODIUM

## INJECTABLE; INJECTION

## EDECIN

+		ATON	EQ 50MG BASE/VIAL	N16093 001		Jul	CAHN
---	--	------	-------------------	------------	--	-----	------

ETHINYL ESTRADIOL; LEVONORGESTREL

## TABLET; ORAL

## QUASENSE

AB		WATSON LABS	0.03MG;0.15MG	N77101 001	Sep 06, 2006	Aug	NEWA
AB	+	DURAMED	0.03MG;0.15MG	N21544 001	Sep 05, 2003	Aug	CFTG

## TABLET; ORAL

## SEASONIQUE

	+	DURAMED RES	0.03MG,0.01MG;0.15MG,N/A	N21840 001	May 25, 2006	May	NEWA
--	---	-------------	--------------------------	------------	--------------	-----	------

## TABLET; ORAL-28

## LEVONORGESTREL AND ETHINYL ESTRADIOL

AB2		WATSON LABS	0.02MG;0.1MG	N77681 001	May 31, 2006	May	NEWA
-----	--	-------------	--------------	------------	--------------	-----	------

ETHINYL ESTRADIOL; NORETHINDRONE

## TABLET; ORAL-21

## BALZIVA-21

## BARR

0.035MG;0.4MG

N76198 001	Apr 22, 2004	Mar	CRLD
------------	--------------	-----	------

## TABLET; ORAL-28

## OVCON-35

AB	+	WARNER CHILCOTT	0.035MG;0.4MG	N17716 001		Mar	CRLD
----	---	-----------------	---------------	------------	--	-----	------

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

## TABLET; ORAL

## LOESTRIN 24 FE

	+	WARNER CHILCOTT	0.02MG;1MG	N21871 001	Feb 17, 2006	Feb	NEWA
--	---	-----------------	------------	------------	--------------	-----	------

ETHINYL ESTRADIOL; NORGESTIMATE

## TABLET; ORAL-28

## NORGESTIMATE AND ETHINYL ESTRADIOL

AB		WATSON LABS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	N76626 001	Aug 17, 2006	Aug	NEWA
----	--	-------------	---	------------	--------------	-----	------

AB			0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	N76626 001	Aug 17, 2006	Aug	NEWA
----	--	--	---	------------	--------------	-----	------

AB			0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	N76626 001	Aug 17, 2006	Aug	NEWA
----	--	--	---	------------	--------------	-----	------

AB			0.035MG;0.25MG	N76627 001	Aug 17, 2006	Aug	NEWA
----	--	--	----------------	------------	--------------	-----	------

ETODOLAC

## CAPSULE; ORAL

## ETODOLAC

## @ ENDO PHARMS

200MG

N74842 001	Jul 17, 1997	Feb	DISC
------------	--------------	-----	------

## @

300MG

N74842 002	Jul 17, 1997	Feb	DISC
------------	--------------	-----	------

AB	+	TARO	300MG	N75078 002	Apr 30, 1998	Mar	CRLD
----	---	------	-------	------------	--------------	-----	------

## LODINE

## @ WYETH PHARMS INC

300MG

N18922 003	Jan 31, 1991	Mar	DISC
------------	--------------	-----	------

## TABLET; ORAL

## ETODOLAC

AB		ACTAVIS ELIZABETH	400MG	N74819 001	Feb 28, 1997	Jun	CAHN
----	--	-------------------	-------	------------	--------------	-----	------

## @ ENDO PHARMS

400MG

N74841 001	Jun 27, 1997	Feb	DISC
------------	--------------	-----	------

ETONOGESTREL

## IMPLANT; IMPLANTATION

## IMPLANON

	+	ORGANON USA INC	68MG/IMPLANT	N21529 001	Jul 17, 2006	Jul	NEWA
--	---	-----------------	--------------	------------	--------------	-----	------

ETOPOSIDE

## INJECTABLE; INJECTION

## TOPOSAR

## @ SICOR PHARMS

20MG/ML

N74166 001	Feb 27, 1995	Jun	DISC
------------	--------------	-----	------

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

AB	ACTAVIS ELIZABETH	20MG	N75650 001	Sep 14, 2001	Jun	CAHN
AB		40MG	N75650 002	Sep 14, 2001	Jun	CAHN

FENOFIBRATE

CAPSULE; ORAL

ANTARA (MICRONIZED)

	OSCIENT	43MG	N21695 001	Nov 30, 2004	Aug	CAHN
	@	87MG	N21695 002	Nov 30, 2004	Aug	CAHN
+		130MG	N21695 003	Nov 30, 2004	Aug	CAHN
	LIPOFEN					
	CIPHER	50MG	N21612 001	Jan 11, 2006	Jan	NEWA
		100MG	N21612 002	Jan 11, 2006	Jan	NEWA
+		150MG	N21612 003	Jan 11, 2006	Jan	NEWA

TABLET; ORAL

FENOFIBRATE

AB	+ TEVA	160MG	N76433 002	May 13, 2005	Jan	CRLD
	TRICOR					
	@ ABBOTT	54MG	N21203 001	Sep 04, 2001	Jan	DISC
	@	160MG	N21203 003	Sep 04, 2001	Jan	DISC

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

FENOLDOPAM MESYLATE

AP	SANDOZ	EQ 10MG BASE/ML	N77155 001	Feb 15, 2005	Jan	CAHN
----	--------	-----------------	------------	--------------	-----	------

FENOPROFEN CALCIUM

TABLET; ORAL

FENOPROFEN CALCIUM

AB	ACTAVIS ELIZABETH	EQ 600MG BASE	N72274 001	May 02, 1988	Jun	CAHN
	@ CLONMEL HLTHCARE	EQ 600MG BASE	N72326 001	Aug 17, 1988	Jan	DISC

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

FENTANYL

AB	LAVIPHARM LABS	25UGM/HR	N77051 001	Aug 04, 2006	Jul	NEWA
AB		50UGM/HR	N77051 002	Aug 04, 2006	Jul	NEWA
AB		75UGM/HR	N77051 003	Aug 04, 2006	Jul	NEWA
AB		100UGM/HR	N77051 004	Aug 04, 2006	Jul	NEWA

FENTANYL CITRATE

>A>	TABLET; BUCCAL					
>A>	FENTORA					
>A>	CEPHALON	EQ 0.1MG BASE	N21947 001	Sep 25, 2006	Sep	NEWA
>A>		EQ 0.2MG BASE	N21947 002	Sep 25, 2006	Sep	NEWA
>A>	+	EQ 0.4MG BASE	N21947 003	Sep 25, 2006	Sep	NEWA
>A>		EQ 0.6MG BASE	N21947 004	Sep 25, 2006	Sep	NEWA
>A>		EQ 0.8MG BASE	N21947 005	Sep 25, 2006	Sep	NEWA
	TROCHE/LOZENGE; ORAL					
	FENTANYL					
	@ CEPHALON	EQ 0.1MG BASE	N20195 007	Oct 30, 1995	Jul	CAHN
	@	EQ 0.2MG BASE	N20195 001	Oct 04, 1993	Jul	CAHN

## TROCHE/LOZENGE; ORAL

## FENTANYL

@	CEPHALON	EQ 0.3MG BASE	N20195 002	Oct 04, 1993	Jul	CAHN
@		EQ 0.4MG BASE	N20195 003	Oct 04, 1993	Jul	CAHN

## TROCHE/LOZENGE; TRANSMUCOSAL

## ACTIQ (SUGAR-FREE)

	CEPHALON	EQ 0.2MG BASE	N20747 001	Nov 04, 1998	Mar	CTNA
+		EQ 0.4MG BASE	N20747 002	Nov 04, 1998	Mar	CTNA
		EQ 0.6MG BASE	N20747 003	Nov 04, 1998	Mar	CTNA
		EQ 0.8MG BASE	N20747 004	Nov 04, 1998	Mar	CTNA
		EQ 1.2MG BASE	N20747 005	Nov 04, 1998	Mar	CTNA
		EQ 1.6MG BASE	N20747 006	Nov 04, 1998	Mar	CTNA

FEXOFENADINE HYDROCHLORIDE

## TABLET; ORAL

## FEXOFENADINE HYDROCHLORIDE

AB	DR REDDYS LABS LTD	30MG	N76502 001	Apr 11, 2006	Mar	NEWA
AB		60MG	N76502 002	Apr 11, 2006	Mar	NEWA
AB		180MG	N76502 003	Apr 11, 2006	Mar	NEWA

FINASTERIDE

## TABLET; ORAL

## FINASTERIDE

AB	DR REDDYS LABS INC	1MG	N76436 001	Jul 28, 2006	Jul	NEWA
AB	IVAX PHARMS	5MG	N76340 001	Jun 19, 2006	Jun	NEWA
AB	PROPECIA					
AB	+ MERCK	1MG	N20788 001	Dec 19, 1997	Jul	CFTG
AB	PROSCAR					
AB	+ MERCK	5MG	N20180 001	Jun 19, 1992	Jun	CFTG

FLUCONAZOLE

## TABLET; ORAL

## FLUCONAZOLE

AB	GLENMARK PHARMA	50MG	N77253 001	Jan 25, 2006	Jan	NEWA
AB		100MG	N77253 002	Jan 25, 2006	Jan	NEWA
AB		150MG	N77253 003	Jan 25, 2006	Jan	NEWA
AB		200MG	N77253 004	Jan 25, 2006	Jan	NEWA

FLUMAZENIL

## INJECTABLE; INJECTION

## FLUMAZENIL

AP	SANDOZ	1MG/10ML (0.1MG/ML)	N77071 002	May 03, 2005	Jan	CAHN
AP		0.5MG/5ML (0.1MG/ML)	N77071 001	May 03, 2005	Jan	CAHN

FLUNISOLIDE

## AEROSOL, METERED; INHALATION

## AEROSPAN HFA

+	FOREST LABS	EQ 78UGM BASE/INH	N21247 001	Jan 27, 2006	Jan	NEWA
---	-------------	-------------------	------------	--------------	-----	------

## SPRAY, METERED; NASAL

## FLUNISOLIDE

AB	+ BAUSCH AND LOMB	0.025MG/SPRAY	N74805 001	Feb 20, 2002	Jul	CTEC
AB	QPHARMA	0.025MG/SPRAY	N77704 001	Aug 03, 2006	Jul	NEWA

FLUOCINONIDE

CREAM; TOPICAL

FLUOCINONIDE

AB1	ACTAVIS MID ATLANTIC	0.05%	N73085	001	Feb 14, 1992	Jun	CAHN
	FLUOCINONIDE EMULSIFIED BASE						
AB2	ACTAVIS MID ATLANTIC	0.05%	N74204	001	Jun 13, 1995	Jun	CAHN
	SOLUTION; TOPICAL						
	FLUOCINONIDE						
AT	ACTAVIS MID ATLANTIC	0.05%	N71535	001	Dec 02, 1988	Jul	CAHN

FLUORESCEIN SODIUM

INJECTABLE; INTRAVENOUS

FLUORESCITE

+	ALCON	EQ 500MG BASE/5ML (EQ 100MG BASE/ML)	N21980	001	Mar 28, 2006	May	CAHN
+	ALCON RES	EQ 500MG BASE/5ML (EQ 100MG BASE/ML)	N21980	001	Mar 28, 2006	Mar	NEWA

FLUOROURACIL

INJECTABLE; INJECTION

ADRUCIL

	@ SICOR PHARMS	50MG/ML	N40023	001	Oct 18, 1991	Jun	DISC
AP	+	50MG/ML	N40023	001	Oct 18, 1991	Mar	CRLD
	@	50MG/ML	N81225	001	Aug 28, 1991	Jun	DISC

FLUOROURACIL

AP	+	AM PHARM PARTNERS	50MG/ML	N40278	001	Sep 30, 1998	Mar	CRLD
AP	+		50MG/ML	N40279	001	Sep 30, 1998	Mar	CRLD
AP	+		50MG/ML	N40291	001	Mar 24, 1999	Mar	CRLD
AP	+		50MG/ML	N40379	001	Nov 15, 2000	Mar	CRLD
	@ BEDFORD	50MG/ML	N89508	001	Jan 26, 1988	Apr	DISC	
AP	+		50MG/ML	N89508	001	Jan 26, 1988	Mar	CRLD
AP	+	SICOR PHARMS	50MG/ML	N40333	001	Jan 27, 2000	Mar	CRLD
AP	+		50MG/ML	N40334	001	Feb 25, 2000	Mar	CRLD
AP	+	STERIS	50MG/ML	N87792	001	Oct 13, 1982	Mar	CRLD
	@ WATSON LABS	50MG/ML	N87792	001	Oct 13, 1982	Apr	DISC	

FLUOXETINE HYDROCHLORIDE

SOLUTION; ORAL

FLUOXETINE

AA	ACTAVIS MID ATLANTIC	EQ 20MG BASE/5ML	N75690	001	Jan 31, 2002	Jul	CAHN
	TABLET; ORAL						
	FLUOXETINE HYDROCHLORIDE						
+	IVAX PHARMS	EQ 40MG BASE	N75865	003	Aug 30, 2004	Aug	CRLD
	SARAFEM						
	WARNER CHILCOTT	EQ 10MG BASE	N21860	001	May 19, 2006	May	NEWA
		EQ 15MG BASE	N21860	002	May 19, 2006	May	NEWA
+		EQ 20MG BASE	N21860	003	May 19, 2006	May	NEWA

FLUPHENAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

FLUPHENAZINE HYDROCHLORIDE

+	PHARM ASSOC	5MG/ML	N74725	001	Sep 16, 1996	Aug	CRLD
	@ TEVA PHARMS	5MG/ML	N73058	001	Aug 30, 1991	Aug	DISC
	PROLIXIN						
	@ APOTHECON	5MG/ML	N70533	001	Nov 07, 1985	Jul	DISC

## ELIXIR; ORAL

## FLUPHENAZINE HYDROCHLORIDE

+	PHARM ASSOC	2.5MG/5ML	N40146 001	Aug 21, 1996	Apr	CRLD
	@ TEVA PHARMS	2.5MG/5ML	N81310 001	Apr 29, 1993	Apr	DISC
	PROLIXIN					
	@ APOTHECON	2.5MG/5ML	N12145 003		Apr	DISC

FLURBIPROFEN SODIUM

## SOLUTION/DROPS; OPHTHALMIC

## FLURBIPROFEN SODIUM

AT	BAUSCH AND LOMB	0.03%	N74447 001	Jan 04, 1995	Aug	CAHN
----	-----------------	-------	------------	--------------	-----	------

FLUTAMIDE

## CAPSULE; ORAL

## EULEXIN

## @ SCHERING

125MG

N18554 001 Jan 27, 1989 May DISC

## FLUTAMIDE

AB	PAR PHARM	125MG	N75298 001	Sep 18, 2001	Apr	CAHN
AB	+ SANDOZ	125MG	N75818 001	Sep 18, 2001	May	CRLD

FLUTICASONE PROPIONATE

## CREAM; TOPICAL

## FLUTICASONE PROPIONATE

AB	G AND W LABS	0.05%	N77055 001	Jun 30, 2006	Jun	NEWA
----	--------------	-------	------------	--------------	-----	------

## OINTMENT; TOPICAL

## FLUTICASONE PROPIONATE

AB	G AND W LABS	0.005%	N77168 001	Mar 03, 2006	Feb	NEWA
----	--------------	--------	------------	--------------	-----	------

## SPRAY, METERED; NASAL

## FLONASE

AB	+ GLAXOSMITHKLINE	0.05MG/SPRAY	N20121 001	Oct 19, 1994	Feb	CFTG
----	-------------------	--------------	------------	--------------	-----	------

## FLUTICASONE PROPIONATE

AB	ROXANE	0.05MG/SPRAY	N76504 001	Feb 22, 2006	Feb	NEWA
----	--------	--------------	------------	--------------	-----	------

FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE

## AEROSOL, METERED; INHALATION

## ADVAIR HFA

+	GLAXOSMITHKLINE	0.045MG/INH;EQ 0.021MG BASE/INH	N21254 001	Jun 08, 2006	Jun	NEWA
+		0.115MG/INH;EQ 0.021MG BASE/INH	N21254 002	Jun 08, 2006	Jun	NEWA
+		0.23MG/INH;EQ 0.021MG BASE/INH	N21254 003	Jun 08, 2006	Jun	NEWA

FLUVOXAMINE MALEATE

## TABLET; ORAL

## FLUVOXAMINE MALEATE

AB	ACTAVIS ELIZABETH	25MG	N75901 001	Dec 28, 2000	Jun	CAHN
AB		50MG	N75901 002	Dec 28, 2000	Jun	CAHN
AB		100MG	N75901 003	Dec 28, 2000	Jun	CAHN
AB	CARACO	25MG	N75900 001	Feb 23, 2006	Feb	NEWA
AB		50MG	N75900 002	Feb 23, 2006	Feb	NEWA
AB		100MG	N75900 003	Feb 23, 2006	Feb	NEWA

FOLLITROPIN ALFA/BETA

## INJECTABLE; SUBCUTANEOUS

## FOLLISTIM AQ

## @ ORGANON USA INC

150 IU/0.18ML

N21211 003 Feb 11, 2004 Jul DISC

+ 300 IU/0.36ML

N21211 001 Mar 23, 2004 Jul CMFD

## INJECTABLE; SUBCUTANEOUS

FOLLISTIM AQ

@ ORGANON USA INC

300 IU/0.36ML

N21211 001 Mar 23, 2004 Jun DISC

FOSCARNET SODIUM

INJECTABLE; INJECTION

FOSCARNET SODIUM

AP HOSPIRA

2.4GM/100ML

N77174 001 May 31, 2005 Feb CAHN

FOSINOPRIL SODIUM

TABLET; ORAL

FOSINOPRIL SODIUM

AB COBALT

10MG

N77531 001 Aug 31, 2006 Aug NEWA

AB

20MG

N77531 002 Aug 31, 2006 Aug NEWA

AB

40MG

N77531 003 Aug 31, 2006 Aug NEWA

FOSINOPRIL SODIUM; HYDROCHLOROTHIAZIDE

TABLET; ORAL

FOSINOPRIL SODIUM AND HYDROCHLOROTHIAZIDE

AB GENPHARM

10MG;12.5MG

N77705 001 Aug 14, 2006 Aug NEWA

AB

20MG;12.5MG

N77705 002 Aug 14, 2006 Aug NEWA

AB TEVA

10MG;12.5MG

N76945 001 Jul 05, 2006 Jun NEWA

AB

20MG;12.5MG

N76945 002 Jul 05, 2006 Jun NEWA

FUROSEMIDE

TABLET; ORAL

FUROSEMIDE

&gt;A&gt; AB OHM LABS

20MG

N78010 001 Sep 18, 2006 Sep NEWA

&gt;A&gt; AB

40MG

N78010 002 Sep 18, 2006 Sep NEWA

&gt;A&gt; AB

80MG

N78010 003 Sep 18, 2006 Sep NEWA

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

AB ACTAVIS ELIZABETH

100MG

N75350 001 Sep 12, 2003 Jun CAHN

AB

300MG

N75350 002 Sep 12, 2003 Jun CAHN

AB

400MG

N75350 003 Sep 12, 2003 Jun CAHN

AB SANDOZ

100MG

N75428 001 Jan 24, 2006 Jan NEWA

AB

300MG

N75428 002 Jan 24, 2006 Jan NEWA

AB

400MG

N75428 003 Jan 24, 2006 Jan NEWA

AB SUN PHARM INDS LTD

100MG

N77242 001 Aug 24, 2006 Aug NEWA

AB

300MG

N77242 002 Aug 24, 2006 Aug NEWA

AB

400MG

N77242 003 Aug 24, 2006 Aug NEWA

TABLET; ORAL

GABAPENTIN

AB ACTAVIS ELIZABETH

600MG

N75694 001 Oct 21, 2004 Jun CAHN

AB

800MG

N75694 002 Oct 21, 2004 Jun CAHN

&gt;A&gt; AB APOTEX INC

100MG

N77894 001 Oct 10, 2006 Sep NEWA

&gt;A&gt; AB

300MG

N77894 002 Oct 10, 2006 Sep NEWA

&gt;A&gt; AB

400MG

N77894 003 Oct 10, 2006 Sep NEWA

AB

600MG

N77661 004 Sep 13, 2006 Aug NEWA

AB

800MG

N77661 005 Sep 13, 2006 Aug NEWA

AB GLENMARK PHARMS

600MG

N77662 001 Aug 18, 2006 Aug NEWA

AB

800MG

N77662 002 Aug 18, 2006 Aug NEWA

&gt;D&gt; IVAX PHARMS

100MG

N76017 001 Apr 28, 2004 Sep CFTG



## TABLET; ORAL

## GABAPENTIN

>A>	AB	IVAX PHARMS	100MG	N76017 001	Apr 28, 2004	Sep	CFTG
>D>			300MG	N76017 002	Apr 28, 2004	Sep	CFTG
>A>	AB		300MG	N76017 002	Apr 28, 2004	Sep	CFTG
>D>			400MG	N76017 003	Apr 28, 2004	Sep	CFTG
>A>	AB		400MG	N76017 003	Apr 28, 2004	Sep	CFTG
	AB	SANDOZ	600MG	N76120 001	Jan 27, 2006	Jan	NEWA
	AB		600MG	N76877 001	Jul 06, 2006	Jun	NEWA
	AB		800MG	N76120 002	Jan 27, 2006	Jan	NEWA
	AB		800MG	N76877 002	Jul 06, 2006	Jun	NEWA
	AB	SUN PHARM INDS LTD	600MG	N77525 001	Aug 24, 2006	Aug	NEWA
	AB		800MG	N77525 002	Aug 24, 2006	Aug	NEWA

GADOVERSETAMIDE

## INJECTABLE; INJECTION

## OPTIMARK

+		MALLINCKRODT	1654.5MG/5ML (330.9MG/ML)	N20937 001	Dec 08, 1999	Jan	CPOT
+			3309MG/10ML (330.9MG/ML)	N20937 002	Dec 08, 1999	Jan	NEWA
+			4963.5MG/15ML (330.9MG/ML)	N20937 003	Dec 08, 1999	Jan	NEWA
+			6618MG/20ML (330.9MG/ML)	N20937 004	Dec 08, 1999	Jan	NEWA
+			16.545GM/50ML (330.9MG/ML)	N20975 001	Dec 08, 1999	Jan	CPOT

## OPTIMARK IN PLASTIC CONTAINER

+		MALLINCKRODT	1654.5MG/5ML (330.9MG/ML)	N20976 001	Dec 08, 1999	Jan	CPOT
+			3309MG/10ML (330.9MG/ML)	N20976 002	Dec 08, 1999	Jan	NEWA
+			4963.5MG/15ML (330.9MG/ML)	N20976 003	Dec 08, 1999	Jan	NEWA
+			6618MG/20ML (330.9MG/ML)	N20976 004	Dec 08, 1999	Jan	NEWA

GALANTAMINE HYDROBROMIDE

## CAPSULE, EXTENDED RELEASE; ORAL

## RAZADYNE ER

+		JANSSEN	EQ 8MG BASE	N21615 001	Apr 01, 2005	May	CMS2
			EQ 16MG BASE	N21615 002	Apr 01, 2005	May	CMS2
			EQ 24MG BASE	N21615 003	Apr 01, 2005	May	CMS2

GANCICLOVIR

## CAPSULE; ORAL

## CYTOVENE

		@ ROCHE PALO	250MG	N20460 001	Dec 22, 1994	Jun	DISC
		@	500MG	N20460 002	Dec 12, 1997	Jun	DISC

## GANCICLOVIR

		RANBAXY	250MG	N76457 001	Jun 27, 2003	Jun	CTEC
+			500MG	N76457 002	Jun 27, 2003	Jun	CRLD

GATIFLOXACIN

## INJECTABLE; INJECTION

## TEQUIN

		@ BRISTOL MYERS SQUIBB	400MG/40ML(10MG/ML)	N21062 004	Dec 17, 1999	May	DISC
		TEQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER					
		@ BRISTOL MYERS SQUIBB	200MG/100ML(2MG/ML)	N21062 001	Dec 17, 1999	May	DISC
		@	400MG/200ML(2MG/ML)	N21062 002	Dec 17, 1999	May	DISC

## SUSPENSION; ORAL

## TEQUIN

		@ BRISTOL MYERS SQUIBB	200MG/5ML	N21678 001	Aug 27, 2004	Jun	DISC
--	--	------------------------	-----------	------------	--------------	-----	------

## TABLET; ORAL

## TEQUIN

@ BRISTOL MYERS SQUIBB 200MG

N21061 001 Dec 17, 1999 May DISC

@ 400MG

N21061 002 Dec 17, 1999 May DISC

GEMFIBROZIL

## TABLET; ORAL

## GEMFIBROZIL

AB INVAGEN PHARMS 600MG

N77836 001 Jul 27, 2006 Jul NEWA

GENTAMICIN SULFATE

## SOLUTION/DROPS; OPHTHALMIC

## GENTACIDIN

@ NOVARTIS EQ 0.3% BASE

N62480 001 Mar 30, 1984 Jun DISC

GLIMEPIRIDE

## TABLET; ORAL

## GLIMEPIRIDE

AB COBALT 1MG

N77280 001 Feb 03, 2006 Jan NEWA

AB 2MG

N77280 002 Feb 03, 2006 Jan NEWA

AB 4MG

N77280 003 Feb 03, 2006 Jan NEWA

AB GENPHARM 1MG

N77486 001 Feb 10, 2006 Jan NEWA

AB 2MG

N77486 002 Feb 10, 2006 Jan NEWA

AB 4MG

N77486 003 Feb 10, 2006 Jan NEWA

GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE

## TABLET; ORAL

## DUETACT

TAKEDA GLOBAL 2MG;30MG

N21925 001 Jul 28, 2006 Jul NEWA

&gt;D&gt; 4MG;30MG

N21925 002 Jul 28, 2006 Sep CRLD

&gt;A&gt; + 4MG;30MG

N21925 002 Jul 28, 2006 Sep CRLD

4MG;30MG

N21925 002 Jul 28, 2006 Jul NEWA

GLIPIZIDE

## TABLET; ORAL

## GLIPIZIDE

AB CARACO 5MG

N77820 001 Jul 11, 2006 Jun NEWA

AB 10MG

N77820 002 Jul 11, 2006 Jun NEWA

@ ENDO PHARMS 5MG

N74378 001 Nov 28, 1994 Feb DISC

@ 10MG

N74378 002 Nov 28, 1994 Feb DISC

## TABLET, EXTENDED RELEASE; ORAL

## GLIPIZIDE

AB WATSON LABS 2.5MG

N76467 003 Mar 27, 2006 Mar NEWA

## GLUCOTROL XL

AB PFIZER 2.5MG

N20329 003 Aug 10, 1999 Mar CFTG

GLYBURIDE

## TABLET; ORAL

## DIABETA

BX + SANOFI AVENTIS US 5MG

N17532 003 May 01, 1984 Feb CRLD

GLYBURIDE; METFORMIN HYDROCHLORIDE

## TABLET; ORAL

## GLYBURIDE AND METFORMIN HYDROCHLORIDE

AB ACTAVIS ELIZABETH 1.25MG;250MG

N76716 001 Jun 28, 2005 Jun CAHN

## TABLET; ORAL

## GLYBURIDE AND METFORMIN HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	2.5MG;500MG	N76716 002	Jun 28, 2005	Jun	CAHN
AB		5MG;500MG	N76716 003	Jun 28, 2005	Jun	CAHN

GLYCOPYRROLATE

## TABLET; ORAL

## GLYCOPYRROLATE

AA	KALI LABS	1MG	N40653 001	Aug 31, 2006	Aug	NEWA
AA		2MG	N40653 002	Aug 31, 2006	Aug	NEWA
ROBINUL						
AA	+ SCIELE PHARMA INC	1MG	N12827 001		Jun	CAHN
ROBINUL FORTE						
AA	+ SCIELE PHARMA INC	2MG	N12827 002		Jun	CAHN

GUANFACINE HYDROCHLORIDE

## TABLET; ORAL

## TENEX

AB	DR REDDYS LABS INC	EQ 1MG BASE	N19032 001	Oct 27, 1986	May	CAHN
AB	+ @	EQ 2MG BASE	N19032 002	Nov 07, 1988	May	CAHN
		EQ 3MG BASE	N19032 003	Nov 07, 1988	May	CAHN

HALOBETASOL PROPIONATE

## CREAM; TOPICAL

## HALOBETASOL PROPIONATE

AB	PERRIGO ISRAEL	0.05%	N77123 001	Dec 16, 2004	Apr	CAHN
----	----------------	-------	------------	--------------	-----	------

## OINTMENT; TOPICAL

## HALOBETASOL PROPIONATE

AB	ACTAVIS MID ATLANTIC	0.05%	N77109 001	Jun 14, 2005	Jun	CAHN
AB	G AND W LABS	0.05%	N77721 001	Sep 07, 2006	Aug	NEWA
AB	PERRIGO	0.05%	N76872 001	Dec 16, 2004	Mar	CAHN

HALOPERIDOL DECANOATE

## INJECTABLE; INJECTION

## HALOPERIDOL DECANOATE

AO	SANDOZ	EQ 50MG BASE/ML	N76463 001	Jun 24, 2005	Jan	CAHN
AO		EQ 100MG BASE/ML	N76463 002	Jun 24, 2005	Jan	CAHN

HALOPERIDOL LACTATE

## INJECTABLE; INJECTION

## HALOPERIDOL

AP	SANDOZ	EQ 5MG BASE/ML	N76464 001	Sep 29, 2004	Jan	CAHN
----	--------	----------------	------------	--------------	-----	------

## SOLUTION; ORAL

## HALOPERIDOL LACTATE

	ACTAVIS MID ATLANTIC	EQ 1MG BASE/ML	N74536 001	Nov 02, 1995	Jul	CAHN
--	----------------------	----------------	------------	--------------	-----	------

HALOTHANE

## LIQUID; INHALATION

## HALOTHANE

	@ HOSPIRA	99.99%	N83254 001		Aug	DISC
--	-----------	--------	------------	--	-----	------

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

## SYRUP; ORAL

## HYDROCODONE COMPOUND

AA	ACTAVIS MID ATLANTIC	1.5MG/5ML;5MG/5ML	N88017 001	Jul 05, 1983	Jul	CAHN
----	----------------------	-------------------	------------	--------------	-----	------

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDRALAZINE HYDROCHLORIDE

@ SICOR PHARMS	20MG/ML	N40373 001	Feb 23, 2000	Aug	DISC
----------------	---------	------------	--------------	-----	------

TABLET; ORAL

APRESOLINE

@ NOVARTIS	10MG	N08303 004		Feb	DISC
@	25MG	N08303 001		Feb	DISC
@	50MG	N08303 002		Feb	DISC
@	100MG	N08303 005		Feb	DISC

HYDRALAZINE HYDROCHLORIDE

AA +	PLIVA	10MG	N89097 001	Dec 18, 1985	Feb	CRLD
AA +		25MG	N88467 001	May 01, 1984	Feb	CRLD
AA +		50MG	N88468 001	May 01, 1984	Feb	CRLD
AA +		100MG	N89098 001	Dec 18, 1985	Feb	CRLD
	@ RADIUS PHARMS	25MG	N86243 001		Aug	CAHN
	@	50MG	N86242 002		Aug	CAHN
	@ TG UNITED LABS	10MG	N88846 001	Feb 26, 1985	May	CAHN
	@	25MG	N88847 001	Feb 26, 1985	May	CAHN
	@	50MG	N88848 001	Feb 26, 1985	May	CAHN
	@	100MG	N88849 001	Feb 26, 1985	May	CAHN

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

CAM-AP-ES

@ TG UNITED LABS	25MG;15MG;0.1MG	N84897 001		May	CAHN
------------------	-----------------	------------	--	-----	------

HYDROCHLOROTHIAZIDE

TABLET; ORAL

HYDROCHLOROTHIAZIDE

AB	ACTAVIS ELIZABETH	25MG	N85054 002		Jun	CAHN
AB		50MG	N85208 001		Jun	CAHN
	@ TG UNITED LABS	25MG	N85683 001		May	CAHN
	@	50MG	N83965 001		May	CAHN
AB	WEST WARD	25MG	N84878 002	Jul 12, 2006	Jun	NEWA

HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

AB	ACTAVIS ELIZABETH	12.5MG;10MG	N76230 001	Jul 01, 2002	Jun	CAHN
AB		12.5MG;20MG	N76230 002	Jul 01, 2002	Jun	CAHN
AB		25MG;20MG	N76230 003	Jul 01, 2002	Jun	CAHN
AB	AUROBINDO	12.5MG;10MG	N77606 001	Mar 14, 2006	Feb	NEWA
AB		12.5MG;20MG	N77606 002	Mar 14, 2006	Feb	NEWA
AB		25MG;20MG	N77606 003	Mar 14, 2006	Feb	NEWA
>A>	AB	LUPIN	N77912 001	Sep 27, 2006	Sep	NEWA
>A>	AB		N77912 002	Sep 27, 2006	Sep	NEWA
>A>	AB		N77912 003	Sep 27, 2006	Sep	NEWA

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

ALDORIL 15

@ MERCK	15MG;250MG	N13402 001		Jun	DISC
---------	------------	------------	--	-----	------

## TABLET; ORAL

ALDORIL 25									
@ MERCK	25MG;250MG		N13402	002			Jun	DISC	
ALDORIL D30									
@ MERCK	30MG;500MG		N13402	003			Jun	DISC	
ALDORIL D50									
@ MERCK	50MG;500MG		N13402	004			Jun	DISC	
METHYLDOPA AND HYDROCHLOROTHIAZIDE									
MYLAN	15MG;250MG		N70264	001	Jan 23, 1986	Jun	CTEC		
+	25MG;250MG		N70265	001	Jan 23, 1986	Jun	CRLD		
@ PAR PHARM	15MG;250MG		N70616	001	Feb 02, 1987	Jun	DISC		
@	25MG;250MG		N70612	001	Feb 02, 1987	Jun	DISC		
@	30MG;500MG		N70613	001	Feb 02, 1987	Jun	DISC		
@	50MG;500MG		N70614	001	Feb 02, 1987	Jun	DISC		
@ SANDOZ	15MG;250MG		N70182	001	Jan 15, 1986	Jun	DISC		
@	25MG;250MG		N70183	001	Jan 15, 1986	Jun	DISC		
@	30MG;500MG		N70543	001	Jan 15, 1986	Jun	DISC		
@	50MG;500MG		N70544	001	Jan 15, 1986	Jun	DISC		
@ WATSON LABS	15MG;250MG		N70958	001	Feb 06, 1989	Jun	DISC		
@	25MG;250MG		N70959	001	Jan 19, 1989	Jun	DISC		
@	30MG;500MG		N71069	001	Jan 19, 1989	Jun	DISC		
@	50MG;500MG		N70960	001	Feb 06, 1989	Jun	DISC		

HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE

## TABLET, EXTENDED RELEASE; ORAL

DUTOPROL									
ASTRAZENECA	12.5MG;EQ 25MG TARTRATE		N21956	001	Aug 28, 2006	Aug	NEWA		
	12.5MG;EQ 50MG TARTRATE		N21956	002	Aug 28, 2006	Aug	NEWA		
+	12.5MG;EQ 100MG TARTRATE		N21956	003	Aug 28, 2006	Aug	NEWA		

HYDROCHLOROTHIAZIDE; PROPRANOLOL HYDROCHLORIDE

## TABLET; ORAL

PROPRANOLOL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE									
AB ACTAVIS ELIZABETH	25MG;40MG		N70851	001	May 15, 1986	Jun	CAHN		
AB	25MG;80MG		N70852	001	May 15, 1986	Jun	CAHN		

HYDROCHLOROTHIAZIDE; TIMOLOL MALEATE

## TABLET; ORAL

TIMOLIDE 10-25									
@ MERCK	25MG;10MG		N18061	001		Jun	DISC		

HYDROCHLOROTHIAZIDE; VALSARTAN

## TABLET; ORAL

DIOVAN HCT									
NOVARTIS	12.5MG;320MG		N20818	004	Apr 28, 2006	Apr	NEWA		
	25MG;160MG		N20818	003	Jan 17, 2002	Apr	CRLD		
+	25MG;320MG		N20818	005	Apr 28, 2006	Apr	NEWA		

HYDROCORTISONE

## CREAM; TOPICAL

HYDROCORTISONE									
AT ACTAVIS MID ATLANTIC	1%		N87795	001	May 03, 1983	Jun	CAHN		
AT	2.5%		N89682	001	Mar 10, 1988	Jun	CAHN		

OINTMENT; TOPICAL						
HYDROCORTISONE						
AT	ACTAVIS MID ATLANTIC	1%	N87796	001	Oct 13, 1982	Jun CAHN
POWDER; FOR RX COMPOUNDING						
HYDRO-RX						
+	X GEN PHARMS	100%	N85982	001		Jun CTNA
TABLET; ORAL						
CORTEF						
	@ PHARMACIA AND UPJOHN	10MG	N08697	001		Jun CTEC
HYDROCORTONE						
	@ MERCK	10MG	N08506	007		Jun DISC
	@	20MG	N08506	011		Jun DISC
<u>HYDROCORTISONE SODIUM SUCCINATE</u>						
INJECTABLE; INJECTION						
A-HYDROCORT						
AP	HOSPIRA	EQ 100MG BASE/VIAL	N40666	001	Apr 06, 2006	Mar NEWA
<u>HYDROCORTISONE; NEOMYCIN; POLYMYXIN B SULFATE</u>						
SUSPENSION/DROPS; OTIC						
NEOMYCIN AND POLYMYXIN B SULFATES AND HYDROCORTISONE						
AT	PHARMAFORCE	1%;EQ 3.5MG BASE;10,000 UNITS	N65219	001	May 01, 2006	Apr NEWA
<u>HYDROXYZINE HYDROCHLORIDE</u>						
SYRUP; ORAL						
ATARAX						
	@ ROERIG	10MG/5ML	N10485	001		Jun DISC
HYDROXYZINE HYDROCHLORIDE						
AA	+	ACTAVIS MID ATLANTIC	10MG/5ML	N86880	001	Jul CAHN
AA	+	ALPHARMA US PHARMS	10MG/5ML	N86880	001	Jun CRLD
AA	+	HI TECH PHARMA	10MG/5ML	N40010	001	Oct 28, 1994 Jun CRLD
AA	+	MORTON GROVE	10MG/5ML	N87294	001	Apr 12, 1982 Jun CRLD
AA	+	VINTAGE PHARMS	10MG/5ML	N40391	001	Apr 10, 2002 Jun CRLD
<u>HYDROXYZINE PAMOATE</u>						
CAPSULE; ORAL						
HYDROXYZINE PAMOATE						
	BARR	EQ 100MG HCL	N88488	001	Jun 15, 1984	Mar CTEC
VISTARIL						
	@ PFIZER	EQ 100MG HCL	N11459	006		Mar DISC
<u>IBANDRONATE SODIUM</u>						
INJECTABLE; INTRAVENOUS						
BONIVA						
+	ROCHE	EQ 3MG BASE/3ML	N21858	001	Jan 06, 2006	Jan NEWA
<u>IBUPROFEN</u>						
SUSPENSION; ORAL						
IBUPROFEN						
AB	ACTAVIS MID ATLANTIC	100MG/5ML	N74978	001	Mar 25, 1998	Jul CAHN
TABLET; ORAL						
IBU						
	@ BASF	400MG	N18197	001		Jun DISC
	@	400MG	N70083	001	Feb 22, 1985	Jun DISC
	@	600MG	N70099	001	Mar 29, 1985	Jun DISC

## TABLET; ORAL

IBU

@ BASF	800MG	N70745 001	Jul 23, 1986	Jun	DISC
--------	-------	------------	--------------	-----	------

IBUPROFEN LYSINE

## INJECTABLE; INTRAVENOUS

NEOPROFEN

+ FARMACON IL	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N21903 001	Apr 13, 2006	Apr	NEWA
---------------	------------------------------------	------------	--------------	-----	------

INDAPAMIDE

## TABLET; ORAL

INDAPAMIDE

AB	ACTAVIS ELIZABETH	1.25MG	N74722 001	Jun 17, 1996	Jun	CAHN
AB		2.5MG	N74722 002	Jun 17, 1996	Jun	CAHN

INDIUM IN-111 OXYQUINOLINE

## INJECTABLE; INJECTION

INDIUM IN-111 OXYQUINOLINE

+ GE HEALTHCARE	1mCi/ML	N19044 001	Dec 24, 1985	Aug	CRLD
-----------------	---------	------------	--------------	-----	------

INDIUM IN-111 PENTETATE DISODIUM

## INJECTABLE; INTRATHECAL

MPI INDIUM DTPA IN 111

+ GE HEALTHCARE	1mCi/ML	N17707 001	Feb 18, 1982	Aug	CRLD
-----------------	---------	------------	--------------	-----	------

INDIUM IN-111 PENTETREOTIDE KIT

## INJECTABLE; INJECTION

OCTREOSCAN

+ MALLINCKRODT	3mCi/ML	N20314 001	Jun 02, 1994	Aug	CRLD
----------------	---------	------------	--------------	-----	------

INDOMETHACIN

## CAPSULE; ORAL

INDOCIN

@ MERCK	25MG	N16059 001		Jun	DISC
---------	------	------------	--	-----	------

@	50MG	N16059 002		Jun	DISC
---	------	------------	--	-----	------

INDO-LEMMON

@ TEVA	25MG	N70266 001	Nov 07, 1985	Jun	DISC
--------	------	------------	--------------	-----	------

@	50MG	N70267 001	Nov 07, 1985	Jun	DISC
---	------	------------	--------------	-----	------

INDOMETHACIN

@ CLONMEL HLTHCARE	25MG	N18851 001	May 18, 1984	Jun	DISC
--------------------	------	------------	--------------	-----	------

@	50MG	N18851 002	May 18, 1984	Jun	DISC
---	------	------------	--------------	-----	------

@ IVAX PHARMS	25MG	N70719 001	Feb 12, 1986	Jun	DISC
---------------	------	------------	--------------	-----	------

@	50MG	N70756 001	Feb 12, 1986	Jun	DISC
---	------	------------	--------------	-----	------

@ MUTUAL PHARM	25MG	N70899 001	Feb 09, 1987	Jun	DISC
----------------	------	------------	--------------	-----	------

@	50MG	N70900 001	Feb 09, 1987	Jun	DISC
---	------	------------	--------------	-----	------

MYLAN	25MG	N18858 001	Apr 20, 1984	Jun	CTEC
-------	------	------------	--------------	-----	------

AB	+	50MG	N18858 002	Apr 20, 1984	Jun	CRLD
----	---	------	------------	--------------	-----	------

@ PAR PHARM	25MG	N18829 002	Aug 06, 1984	Jun	DISC
-------------	------	------------	--------------	-----	------

@	50MG	N18829 001	Aug 06, 1984	Jun	DISC
---	------	------------	--------------	-----	------

@	50MG	N70651 001	Mar 05, 1986	Jun	DISC
---	------	------------	--------------	-----	------

@ PLIVA	25MG	N71148 001	Mar 18, 1987	Jun	DISC
---------	------	------------	--------------	-----	------

@	50MG	N71149 001	Mar 18, 1987	Jun	DISC
---	------	------------	--------------	-----	------

@ RADIUS PHARMS	25MG	N18851 001	May 18, 1984	Aug	CAHN
-----------------	------	------------	--------------	-----	------

@	50MG	N18851 002	May 18, 1984	Aug	CAHN
---	------	------------	--------------	-----	------

@ SANDOZ	25MG	N70673 001	Apr 29, 1987	Jun	DISC
----------	------	------------	--------------	-----	------

## CAPSULE; ORAL

## INDOMETHACIN

@ SANDOZ	50MG	N70674 001	Apr 29, 1987	Jun	DISC
@ TEVA	25MG	N71342 001	Apr 18, 1988	Jun	DISC
@	50MG	N71343 001	Apr 18, 1988	Jun	DISC

## CAPSULE, EXTENDED RELEASE; ORAL

## INDOCIN SR

+ SANDOZ	75MG	N74464 001	May 28, 1998	Jun	CTEC
@ INWOOD LABS	75MG	N72410 001	Mar 15, 1989	Jun	DISC

## SUPPOSITORY; RECTAL

## INDOMETHACIN

+ G AND W LABS	50MG	N73314 001	Aug 31, 1992	Apr	CTNA
----------------	------	------------	--------------	-----	------

INSULIN GLULISINE RECOMBINANT

## INJECTABLE; SUBCUTANEOUS

## APIDRA

+ SANOFI AVENTIS US	1000 UNITS/10ML (100 UNITS/ML)	N21629 001	Apr 16, 2004	Mar	CMFD
+	300 UNITS/3ML (100 UNITS/ML)	N21629 002	Dec 20, 2005	Mar	CMFD

INSULIN PURIFIED PORK

## INJECTABLE; INJECTION

## ILETIN II

@ LILLY	500 UNITS/ML	N18344 002		Jun	DISC
---------	--------------	------------	--	-----	------

INSULIN RECOMBINANT HUMAN

## POWDER; INHALATION

## EXUBERA

PFIZER	1MG/INH	N21868 001	Jan 27, 2006	Jan	NEWA
+	3MG/INH	N21868 002	Jan 27, 2006	Jan	NEWA

IPRATROPIUM BROMIDE

## AEROSOL, METERED; INHALATION

## ATROVENT

@ BOEHRINGER INGELHEIM	0.018MG/INH	N19085 001	Dec 29, 1986	May	DISC
------------------------	-------------	------------	--------------	-----	------

## SOLUTION; INHALATION

## ATROVENT

@ BOEHRINGER INGELHEIM	0.02%	N20228 001	Sep 29, 1993	May	DISC
------------------------	-------	------------	--------------	-----	------

## IPRATROPIUM BROMIDE

AN	ACTAVIS MID ATLANTIC	0.02%	N75111 001	Apr 22, 1999	Jul	CAHN
AN	+ DEY	0.02%	N74755 001	Jan 10, 1997	May	CRLD
	@ ROXANE	0.02%	N75867 001	Jul 22, 2002	May	DISC

ISONIAZID

## INJECTABLE; INJECTION

## ISONIAZID

AP	SANDOZ	100MG/ML	N40648 001	Jul 05, 2005	Jan	CAHN
----	--------	----------	------------	--------------	-----	------

ISOSORBIDE MONONITRATE

## TABLET; ORAL

## ISMO

AB	DR REDDYS LABS INC	20MG	N19091 001	Dec 30, 1991	May	CAHN
	ISOSORBIDE MONONITRATE					
AB	ACTAVIS ELIZABETH	10MG	N75037 002	Oct 30, 1998	Jun	CAHN
AB		20MG	N75037 001	Oct 30, 1998	Jun	CAHN



TABLET, EXTENDED RELEASE; ORAL						
ISOSORBIDE MONONITRATE						
AB	ACTAVIS ELIZABETH	30MG	N75306 001	Dec 31, 1998	Jun	CAHN
AB		60MG	N75306 002	Dec 31, 1998	Jun	CAHN
AB	WEST WARD	30MG	N76813 002	Mar 30, 2006	Mar	NEWA
<u>ISOTRETINOIN</u>						
CAPSULE; ORAL						
CLARAVIS						
AB	BARR	30MG	N76135 003	May 11, 2006	May	NEWA
SOTRET						
AB	RANBAXY	30MG	N76503 001	Jun 20, 2003	May	CTEC
<u>ISRADIPINE</u>						
CAPSULE; ORAL						
DYNACIRC						
	@ RELIANT PHARMS	2.5MG	N19546 001	Dec 20, 1990	May	DISC
ISRADIPINE						
AB	+ ABRIKA PHARMS	5MG	N77317 002	Jan 05, 2006	Jun	CRLD
AB		5MG	N77317 002	Jan 05, 2006	Apr	CTEC
AB	AMIDE PHARM	2.5MG	N77169 001	Apr 24, 2006	Apr	NEWA
AB		5MG	N77169 002	Apr 24, 2006	Apr	NEWA
<u>IVERMECTIN</u>						
TABLET; ORAL						
STROMECTOL						
	+ MERCK	3MG	N50742 002	Oct 08, 1998	Jun	CRLD
	@	6MG	N50742 001	Nov 22, 1996	Jun	DISC
<u>KETOCONAZOLE</u>						
GEL; TOPICAL						
XOLEGEL						
	+ BARRIER THERAP	2%	N21946 001	Jul 28, 2006	Jul	NEWA
<u>KETOPROFEN</u>						
CAPSULE; ORAL						
KETOPROFEN						
AB	RADIUS PHARMS	25MG	N74014 001	Jan 29, 1993	Jul	CAHN
AB		50MG	N74014 002	Jan 29, 1993	Jul	CAHN
AB		75MG	N74014 003	Jan 29, 1993	Jul	CAHN
<u>KETOROLAC TROMETHAMINE</u>						
INJECTABLE; INJECTION						
KETOROLAC TROMETHAMINE						
AP	SANDOZ	15MG/ML	N76271 001	Oct 06, 2004	Jan	CAHN
AP		30MG/ML	N76271 002	Oct 06, 2004	Jan	CAHN
<u>KETOTIFEN FUMARATE</u>						
SOLUTION/DROPS; OPHTHALMIC						
KETOTIFEN FUMARATE						
AT	APOTEX	EQ 0.025% BASE	N77354 001	May 09, 2006	Apr	NEWA
ZADITOR						
AT	+ NOVARTIS	EQ 0.025% BASE	N21066 001	Jul 02, 1999	Apr	CFTG

LACTULOSE

SOLUTION; ORAL

CONSTULOSE

AA + ACTAVIS MID ATLANTIC 10GM/15ML N70288 001 Aug 15, 1988 Jul CAHN

SOLUTION; ORAL, RECTAL

ENULOSE

AA + ACTAVIS MID ATLANTIC 10GM/15ML N71548 001 Aug 15, 1988 Jul CAHN

LAMOTRIGINE

TABLET; ORAL

LAMICTAL

AB + GLAXOSMITHKLINE 25MG N20241 005 Dec 27, 1994 Aug CFTG

AB 100MG N20241 001 Dec 27, 1994 Aug CFTG

AB 150MG N20241 002 Dec 27, 1994 Aug CFTG

AB 200MG N20241 003 Dec 27, 1994 Aug CFTG

LAMOTRIGINE

AB TEVA 25MG N76388 001 Aug 30, 2006 Aug NEWA

AB 100MG N76388 002 Aug 30, 2006 Aug NEWA

AB 150MG N76388 003 Aug 30, 2006 Aug NEWA

AB 200MG N76388 004 Aug 30, 2006 Aug NEWA

TABLET, CHEWABLE; ORAL

LAMICTAL CD

AB GLAXOSMITHKLINE 5MG N20764 001 Aug 24, 1998 Jun CFTG

AB + 25MG N20764 002 Aug 24, 1998 Jun CFTG

LAMOTRIGINE

AB TEVA 5MG N76420 001 Jun 21, 2006 Jun NEWA

AB 25MG N76420 002 Jun 21, 2006 Jun NEWA

LANSOPRAZOLE; NAPROXEN

CAPSULE, DELAYED REL PELLETS, TABLET; ORAL

PREVACID NAPRAPAC 250 (COPACKAGED)

TAP PHARM 15MG,N/A;N/A,250MG N21507 002 Nov 14, 2003 Feb CTNA

PREVACID NAPRAPAC 375 (COPACKAGED)

TAP PHARM 15MG,N/A;N/A,375MG N21507 003 Nov 14, 2003 Feb CTNA

PREVACID NAPRAPAC 500 (COPACKAGED)

+ TAP PHARM 15MG,N/A;N/A,500MG N21507 004 Nov 14, 2003 Feb CTNA

LENALIDOMIDE

CAPSULE; ORAL

REVLIMID

CELGENE 10MG N21880 002 Dec 27, 2005 Jul CRLD

15MG N21880 003 Jun 29, 2006 Jul NEWA

+ 25MG N21880 004 Jun 29, 2006 Jul NEWA

LEVETIRACETAM

INJECTABLE; IV (INFUSION)

KEPPRA

+ UCB INC 500MG/5ML (100MG/ML) N21872 001 Jul 31, 2006 Jul NEWA

TABLET; ORAL

KEPPRA

UCB INC 750MG N21035 003 Nov 30, 1999 Mar CRLD

+ 1GM N21035 004 Jan 06, 2006 Mar NEWA

LEVOBETAXOLOL HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

BETAXON

@ ALCON

EQ 0.5% BASE

N21114 001 Feb 23, 2000 Feb CAHN

LEVOCABASTINE HYDROCHLORIDE

SUSPENSION/DROPS; OPHTHALMIC

LIVOSTIN

@ NOVARTIS

EQ 0.05% BASE

N20219 001 Nov 10, 1993 May DISC

LEVONORGESTREL

TABLET; ORAL

PLAN B

+ DURAMED

0.75MG

N21045 002 Aug 24, 2006 Aug NEWA

LEVOTHYROXINE SODIUM\*\*

\*\*Refer to Preface Section 1.7 Levothyroxine Sodium for amplifying information

TABLET; ORAL

LEVOTHYROXINE SODIUM

>D>	AB2	GENPHARM	0.025MG	N76752 001	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.025MG	N76752 001	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.05MG	N76752 002	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.05MG	N76752 002	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.075MG	N76752 003	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.075MG	N76752 003	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.088MG	N76752 004	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.088MG	N76752 004	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.1MG	N76752 005	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.1MG	N76752 005	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.112MG	N76752 006	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.112MG	N76752 006	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.125MG	N76752 007	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.125MG	N76752 007	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.15MG	N76752 008	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.15MG	N76752 008	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.175MG	N76752 009	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.175MG	N76752 009	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.2MG	N76752 010	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.2MG	N76752 010	Jun 16, 2005	Sep	CTEC
	AB3						
>D>	AB2		0.3MG	N76752 011	Jun 16, 2005	Sep	CTEC
>A>	AB2,		0.3MG	N76752 011	Jun 16, 2005	Sep	CTEC
	AB3						

LIDOCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

XYLOCAINE

AP + ABRAXIS BIOSCIENCE

0.5%

N06488 008

Jul CAHN

AP +

1%

N06488 007

Jul CAHN

AP +

1.5%

N06488 010

Jul CAHN

## INJECTABLE; INJECTION

		XYLOCAINE						
		@ ABRAXIS BIOSCIENCE	2%		N06488	002		Jul CAHN
		XYLOCAINE 4% PRESERVATIVE FREE						
AP	+	ABRAXIS BIOSCIENCE	4%		N10417	001		Jul CAHN
		XYLOCAINE PRESERVATIVE FREE						
AP		ABRAXIS BIOSCIENCE	1%		N16801	005	Jan 19, 1988	Jul CAHN
AP	+		2%		N16801	001		Jul CAHN
AP			4%		N16801	002		Jul CAHN
AP	+		10%		N16801	003		Jul CAHN
AP	+		20%		N16801	004		Jul CAHN

## INJECTABLE; SPINAL

		XYLOCAINE 1.5% W/ DEXTROSE 7.5%						
		@ ABRAXIS BIOSCIENCE	1.5%		N16297	001		Jul CAHN

## JELLY; TOPICAL

		ANESTACON						
AT	+	POLYMEDICA	2%		N80429	001		Jan CDFR
		XYLOCAINE						
AT	+	ABRAXIS BIOSCIENCE	2%		N08816	001		Jul CAHN

## SOLUTION; ORAL

		LIDOCAINE HYDROCHLORIDE VISCOUS						
AT		ACTAVIS MID ATLANTIC	2%		N86578	001		Jul CAHN
		XYLOCAINE VISCOUS						
AT	+	ABRAXIS BIOSCIENCE	2%		N09470	001		Jul CAHN

## SOLUTION; TOPICAL

		XYLOCAINE 4% PRESERVATIVE FREE						
AT	+	ABRAXIS BIOSCIENCE	4%		N10417	002		Jul CAHN

LIDOCAINE; PRILOCAINE

## CREAM; TOPICAL

		EMLA						
AB	+	ABRAXIS BIOSCIENCE	2.5%;2.5%		N19941	001	Dec 30, 1992	Jul CAHN

LIDOCAINE; TETRACAINE

## CREAM; TOPICAL

		LIDOCAINE AND TETRACAINE						
	+	ZARS	7%;7%		N21717	001	Jun 29, 2006	Jun NEWA

## PATCH; TOPICAL

		SYNERA						
	+	ENDO PHARMS	70MG;70MG		N21623	001	Jun 23, 2005	Feb CAHN

LINDANE

## LOTION; TOPICAL

		LINDANE						
AT	+	ACTAVIS MID ATLANTIC	1%		N87313	001		Jul CAHN

## SHAMPOO; TOPICAL

		LINDANE						
AT	+	ACTAVIS MID ATLANTIC	1%		N87266	001		Jul CAHN

LIOTHYRONINE SODIUM

## INJECTABLE; INJECTION

## LIOTHYRONINE SODIUM

>D>	AP	PHARMAFORCE	EQ 0.01MG BASE/ML		N76923	001	Aug 17, 2005	Sep CAHN
>A>	AP	X GEN PHARMS	EQ 0.01MG BASE/ML		N76923	001	Aug 17, 2005	Sep CAHN

LISINOPRIL

TABLET; ORAL

LISINOPRIL

AB	ACTAVIS ELIZABETH	2.5MG	N76180 001	Jul 01, 2002	Jun	CAHN
AB		5MG	N76180 002	Jul 01, 2002	Jun	CAHN
AB		10MG	N76180 003	Jul 01, 2002	Jun	CAHN
AB		20MG	N76164 001	Jul 01, 2002	Jun	CAHN
AB		30MG	N76164 002	Jul 01, 2002	Jun	CAHN
AB		40MG	N76164 003	Jul 01, 2002	Jun	CAHN
AB	AUROBINDO	2.5MG	N77622 001	Feb 22, 2006	Feb	NEWA
AB		5MG	N77622 002	Feb 22, 2006	Feb	NEWA
AB		10MG	N77622 003	Feb 22, 2006	Feb	NEWA
AB		20MG	N77622 004	Feb 22, 2006	Feb	NEWA
AB		30MG	N77622 005	Feb 22, 2006	Feb	NEWA
AB		40MG	N77622 006	Feb 22, 2006	Feb	NEWA
	PRINIVIL					
	@ MERCK	2.5MG	N19558 006	Jan 28, 1994	Jun	DISC

LITHIUM CARBONATE

TABLET; ORAL

LITHIUM CARBONATE

	@ PFIZER	300MG	N16834 001		Aug	DISC
+	ROXANE	300MG	N18558 001	Jan 29, 1982	Aug	CRLD
	TABLET, EXTENDED RELEASE; ORAL					
	ESKALITH CR					
	@ GLAXOSMITHKLINE	450MG	N18152 001	Mar 29, 1982	Jul	DISC
	LITHIUM CARBONATE					
	@ BARR	450MG	N76366 001	Aug 21, 2003	Jul	DISC
AB	+ ROXANE	450MG	N76691 001	Jan 05, 2004	Jul	CRLD

LORAZEPAM

TABLET; ORAL

LORAZEPAM

AB	ACTAVIS ELIZABETH	0.5MG	N71403 001	Apr 21, 1987	Jun	CAHN
AB		1MG	N71404 001	Apr 21, 1987	Jun	CAHN
AB		2MG	N71141 001	Apr 21, 1987	Jun	CAHN
AB	MYLAN	0.5MG	N77657 001	Mar 16, 2006	Mar	NEWA
AB		1MG	N77657 002	Mar 16, 2006	Mar	NEWA
AB		2MG	N77657 003	Mar 16, 2006	Mar	NEWA
AB	VINTAGE PHARMS	0.5MG	N77754 001	May 10, 2006	Apr	NEWA
AB		1MG	N77754 002	May 10, 2006	Apr	NEWA
AB		2MG	N77754 003	May 10, 2006	Apr	NEWA

LOVASTATIN

TABLET; ORAL

LOVASTATIN

AB	ACTAVIS ELIZABETH	10MG	N75828 001	Dec 17, 2001	Jun	CAHN
AB		20MG	N75828 002	Dec 17, 2001	Jun	CAHN
AB		40MG	N75828 003	Dec 17, 2001	Jun	CAHN
AB	MUTUAL PHARM	10MG	N77520 001	Apr 14, 2006	Apr	NEWA
AB		20MG	N77520 002	Apr 14, 2006	Apr	NEWA
AB		40MG	N77520 003	Apr 14, 2006	Apr	NEWA
	MEVACOR					
	@ MERCK	10MG	N19643 002	Mar 28, 1991	Aug	DISC

LOVASTATIN; NIACIN

TABLET, EXTENDED RELEASE; ORAL  
ADVICOR

+	KOS LIFE	20MG;750MG	N21249 002	Dec 17, 2001	Feb	CMFD
+		40MG;1GM	N21249 004	Apr 27, 2006	Jul	NEWA

LUBIPROSTONE

CAPSULE; ORAL  
AMITIZA

+	SUCAMPO PHARMS	24UGM	N21908 001	Jan 31, 2006	Jan	NEWA
---	----------------	-------	------------	--------------	-----	------

MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

SOLUTION; INJECTION  
NORMOCARB HF 25

+	DIALYSIS SUPS	0.21GM/100ML;2.8GM/100ML;9.07GM/100ML	N21910 001	Jul 26, 2006	Jul	NEWA
---	---------------	---------------------------------------	------------	--------------	-----	------

NORMOCARB HF 35

+	DIALYSIS SUPS	0.21GM/100ML;3.97GM/100ML;8.3GM/100ML	N21910 002	Jul 26, 2006	Jul	NEWA
---	---------------	---------------------------------------	------------	--------------	-----	------

MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE

TABLET, CHEWABLE; ORAL  
ZEGERID

	SANTARUS	700MG;20MG;600MG	N21850 001	Mar 24, 2006	Mar	NEWA
+		700MG;40MG;600MG	N21850 002	Mar 24, 2006	Mar	NEWA

MEDROXYPROGESTERONE ACETATE

INJECTABLE; SUBCUTANEOUS  
DEPO-SUBQ PROVERA 104

+	PHARMACIA AND UPJOHN	104MG/0.65ML	N21583 001	Dec 17, 2004	Jan	CAHN
---	----------------------	--------------	------------	--------------	-----	------

MEGESTROL ACETATE

SUSPENSION; ORAL  
MEGESTROL ACETATE

AB	APOTEX	40MG/ML	N77404 001	Feb 16, 2006	Jan	NEWA
----	--------	---------	------------	--------------	-----	------

MELOXICAM

TABLET; ORAL  
MELOXICAM

AB	ACTAVIS TOTOWA	7.5MG	N77938 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77938 002	Jul 19, 2006	Jul	NEWA
AB	APOTEX INC	7.5MG	N77882 001	Jul 20, 2006	Jul	NEWA
AB		15MG	N77882 002	Jul 20, 2006	Jul	NEWA
>A>	AUROBINDO PHARMA	7.5MG	N78008 001	Oct 02, 2006	Sep	NEWA
>A>		15MG	N78008 002	Oct 02, 2006	Sep	NEWA
AB	BRECKENRIDGE PHARM	7.5MG	N77920 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77920 002	Jul 19, 2006	Jul	NEWA
AB	CARACO	7.5MG	N77937 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77937 002	Jul 19, 2006	Jul	NEWA
AB	COREPHARMA	7.5MG	N77930 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77930 002	Jul 19, 2006	Jul	NEWA
AB	DR REDDYS LABS INC	7.5MG	N77931 001	Jul 25, 2006	Jul	NEWA
AB		15MG	N77931 002	Jul 25, 2006	Jul	NEWA
AB	GENPHARM	7.5MG	N77934 001	Jul 20, 2006	Jul	NEWA
AB		15MG	N77934 002	Jul 20, 2006	Jul	NEWA

## TABLET; ORAL

## MELOXICAM

AB	GLENMARK PHARMS LTD	7.5MG	N77932 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77932 002	Jul 19, 2006	Jul	NEWA
AB	LUPIN PHARMS	7.5MG	N77944 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77944 002	Jul 19, 2006	Jul	NEWA
AB	MUTUAL PHARM	7.5MG	N77935 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77935 002	Jul 19, 2006	Jul	NEWA
AB	MYLAN	7.5MG	N77923 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77923 002	Jul 19, 2006	Jul	NEWA
AB	PAR PHARM	7.5MG	N77933 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77933 002	Jul 19, 2006	Jul	NEWA
AB	ROXANE	7.5MG	N77925 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77925 002	Jul 19, 2006	Jul	NEWA
AB	TEVA PHARMS	7.5MG	N77936 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77936 002	Jul 19, 2006	Jul	NEWA
AB	WATSON LABS	7.5MG	N77929 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77929 002	Jul 19, 2006	Jul	NEWA
AB	ZYDUS PHARMS USA	7.5MG	N77921 001	Jul 19, 2006	Jul	NEWA
AB		15MG	N77921 002	Jul 19, 2006	Jul	NEWA
	MOBIC					
AB	BOEHRINGER INGELHEIM	7.5MG	N20938 001	Apr 13, 2000	Jul	CFTG
AB	+	15MG	N20938 002	Aug 23, 2000	Jul	CFTG

MEPROBAMATE

## TABLET; ORAL

## MEPROBAMATE

	@ ROXANE	600MG	N84332 001		Jan	DISC
	@ SANDOZ	200MG	N14547 002		Jan	DISC
	@	400MG	N14547 001		Jan	DISC
	@	400MG	N80655 001		Jan	DISC
	@ SCHERER LABS	400MG	N83343 001		Jan	DISC
	@ TABLICAPS	400MG	N83494 001		Jan	DISC
AA	+ WATSON LABS	200MG	N83304 001		Jan	CRLD
	@	200MG	N85720 001		Jan	DISC
	+	400MG	N83308 001		Jan	CRLD
	@	400MG	N85721 001		Jan	DISC
	MILTOWN					
	@ MEDPOINTE PHARM HLC	200MG	N09698 004		Jan	DISC
	@	400MG	N09698 002		Jan	DISC
	TRANMEP					
	@ SOLVAY	400MG	N16249 001		Jan	DISC

MESALAMINE

## ENEMA; RECTAL

## ROWASA

AB	+ ALAVEN PHARM	4GM/60ML	N19618 001	Dec 24, 1987	Apr	CAHN
----	----------------	----------	------------	--------------	-----	------

## SUPPOSITORY; RECTAL

## CANASA

## @ AXCAN SCANDIPHARM

## ROWASA

## @ ALAVEN PHARM

		500MG	N21252 001	Jan 05, 2001	May	DISC
		500MG	N19919 001	Dec 18, 1990	Jul	CAHN

METARAMINOL BITARTRATE

INJECTABLE; INJECTION

ARAMINE

@	MERCK	EQ 10MG BASE/ML	N09509 002	Dec 22, 1987	Jun	DISC
+	ABRAXIS PHARM	EQ 10MG BASE/ML	N80722 001		Jun	CRLD

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	500MG	N76033 001	Jan 24, 2002	Jun	CAHN
AB		850MG	N76033 002	Jan 24, 2002	Jun	CAHN
AB		1GM	N76033 003	Jan 24, 2002	Jun	CAHN
AB	AMNEAL PHARM	500MG	N77853 001	Jul 28, 2006	Jul	NEWA
AB		850MG	N77853 002	Jul 28, 2006	Jul	NEWA
AB		1GM	N77853 003	Jul 28, 2006	Jul	NEWA
AB	DR REDDYS LABS INC	500MG	N77787 001	Aug 23, 2006	Aug	NEWA
AB		850MG	N77787 002	Aug 23, 2006	Aug	NEWA
AB		1GM	N77787 003	Aug 23, 2006	Aug	NEWA
AB	INTERPHARM	500MG	N77880 001	Jun 05, 2006	May	NEWA
AB		850MG	N77880 002	Jun 05, 2006	May	NEWA
AB		1GM	N77880 003	Jun 05, 2006	May	NEWA

TABLET, EXTENDED RELEASE; ORAL

GLUMETZA

BX	DEPOMED INC	500MG	N21748 001	Jun 03, 2005	Jan	CAHN
BX	+	1GM	N21748 002	Jun 03, 2005	Aug	CRLD
BX		1GM	N21748 002	Jun 03, 2005	Jan	CAHN

METFORMIN HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	500MG	N76450 001	Oct 01, 2004	Jun	CAHN
AB		750MG	N76878 001	Apr 13, 2005	Jun	CAHN
AB	NOSTRUM	500MG	N76756 001	Jul 26, 2006	Jul	NEWA
AB	SUN PHARM INDS (IN)	500MG	N77336 001	Feb 09, 2006	Jan	NEWA
AB		750MG	N77336 002	Feb 09, 2006	Jan	NEWA

METHIMAZOLE

TABLET; ORAL

METHIMAZOLE

+	CEDAR PHARMS	20MG	N40547 004	Feb 18, 2005	May	CRLD
AB	JONES PHARMA	5MG	N40320 001	Mar 31, 2000	May	CTNA
AB		10MG	N40320 002	Mar 31, 2000	May	CTNA

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE

AP	+	ABRAXIS PHARM	EQ 50MG BASE/2ML (25MG/ML)	N40263 001	Feb 26, 1999	Jun	CMFD
AP	+		EQ 250MG BASE/10ML (25MG/ML)	N40263 002	Feb 26, 1999	Jul	CRLD
AP			EQ 250MG BASE/10ML (25MG/ML)	N40263 002	Feb 26, 1999	Jun	NEWA

METHOTREXATE PRESERVATIVE FREE

+	BEDFORD	EQ 1GM BASE/VIAL	N40632 001	Aug 12, 2005	Apr	CAIN
+	MAYNE PHARMA USA	EQ 1GM BASE/40ML (25MG/ML)	N11719 012	Apr 13, 2005	Jun	CPOT
+		EQ 1GM BASE/40ML (25 MG/ML)	N11719 012	Apr 13, 2005	Apr	CPOT

METHOTREXATE SODIUM

AP	+	BEDFORD	EQ 50MG BASE/2ML (25MG/ML)	N89340 001	Sep 16, 1986	Apr	CPOT
+			EQ 100MG BASE/4ML (25MG/ML)	N89341 001	Sep 16, 1986	Apr	CTEC



## INJECTABLE; INJECTION

## METHOTREXATE SODIUM

+	BEDFORD	EQ 200MG BASE/8ML (25MG/ML)	N89342 001	Sep 16, 1986	Apr	CTEC
+		EQ 250MG BASE/10ML (25 MG/ML)	N89343 001	Sep 16, 1986	Apr	CTEC

## TABLET; ORAL

## METHOTREXATE SODIUM

AB	+	STADA PHARMS	EQ 2.5MG BASE	N08085 002		Aug	CAHN
----	---	--------------	---------------	------------	--	-----	------

METHYLPHENIDATE

## FILM, EXTENDED RELEASE; TRANSDERMAL

## DAYTRANA

+	SHIRE	10MG/9HR (1.1MG/HR)	N21514 001	Apr 06, 2006	Apr	NEWA
+		15MG/9HR (1.6MG/HR)	N21514 002	Apr 06, 2006	Apr	NEWA
+		20MG/9HR (2.2MG/HR)	N21514 003	Apr 06, 2006	Apr	NEWA
+		30MG/9HR (3.3MG/HR)	N21514 004	Apr 06, 2006	Apr	NEWA

METHYLPHENIDATE HYDROCHLORIDE

## CAPSULE, EXTENDED RELEASE; ORAL

## METADATE CD

BX		UCB INC	40MG	N21259 004	Feb 19, 2006	Feb	NEWA
			50MG	N21259 005	Feb 19, 2006	Feb	NEWA
	+		60MG	N21259 006	Feb 19, 2006	Feb	NEWA

## RITALIN LA

BX	+	NOVARTIS	40MG	N21284 003	Jun 05, 2002	Feb	CTEC
----	---	----------	------	------------	--------------	-----	------

## TABLET; ORAL

## METHYLPHENIDATE HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	5MG	N40321 001	Feb 05, 2002	Jun	CAHN
AB			10MG	N40321 002	Feb 05, 2002	Jun	CAHN
AB			20MG	N40321 003	Feb 05, 2002	Jun	CAHN

## TABLET, EXTENDED RELEASE; ORAL

## METHYLPHENIDATE HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	20MG	N75450 001	Dec 21, 2001	Jun	CAHN
----	--	-------------------	------	------------	--------------	-----	------

METOCLOPRAMIDE HYDROCHLORIDE

## SOLUTION; ORAL

## METOCLOPRAMIDE

AA	+	JVL	EQ 5MG BASE/5ML	N74703 001	Oct 31, 1997	Aug	CRLD
		@ VISTAPHARM	EQ 5MG BASE/5ML	N75051 001	Jan 26, 2001	Aug	DISC

## METOCLOPRAMIDE HYDROCHLORIDE

AA		ACTAVIS MID ATLANTIC	EQ 5MG BASE/5ML	N71340 001	Aug 18, 1988	Jul	CAHN
		@ ROXANE	EQ 5MG BASE/5ML	N72038 001	Dec 05, 1988	Aug	DISC
		@ TEVA	EQ 5MG BASE/5ML	N70819 001	Jul 10, 1987	Aug	DISC
		@	EQ 5MG BASE/5ML	N71315 001	Jun 30, 1993	Aug	DISC

## TABLET; ORAL

## METOCLOPRAMIDE

AB		VINTAGE PHARMS	EQ 5MG BASE	N77878 001	Aug 28, 2006	Aug	NEWA
AB			EQ 10MG BASE	N77878 002	Aug 28, 2006	Aug	NEWA

## METOCLOPRAMIDE HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	EQ 10MG BASE	N70581 001	Oct 17, 1985	Jun	CAHN
AB		MUTUAL PHARM	EQ 5MG BASE	N71536 002	Jan 16, 1997	Apr	CMFD
AB			EQ 10MG BASE	N71536 001	Apr 28, 1993	Apr	CMFD

## TABLET, ORALLY DISINTEGRATING; ORAL

## REGLAN ODT

## @ SCHWARZ PHARMA

## @

EQ 5MG BASE	N21793 001	Jun 10, 2005	May	DISC
EQ 10MG BASE	N21793 002	Jun 10, 2005	May	DISC

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

AB	SANDOZ	EQ 25MG TARTRATE	N76969 001	Jul 31, 2006	Jul	NEWA
	TOPROL-XL					
AB	ASTRAZENECA	EQ 25MG TARTRATE	N19962 004	Feb 05, 2001	Jul	CFTG

METRONIDAZOLE

GEL; TOPICAL

METROGEL

AB	+ GALDERMA LABS LP	0.75%	N19737 001	Nov 22, 1988	May	CFTG
	METRONIDAZOLE					
AB	ALTANA	0.75%	N77018 001	Jun 06, 2006	May	NEWA
AB	QLT USA	0.75%	N77547 001	Jul 13, 2006	Jun	NEWA
AB	TARO	0.75%	N77819 001	Jul 18, 2006	Jul	NEWA

LOTION; TOPICAL

METROLOTION

AB	+ GALDERMA LABS LP	0.75%	N20901 001	Nov 24, 1998	May	CFTG
	METRONIDAZOLE					
AB	ALTANA	0.75%	N77197 001	May 24, 2006	May	NEWA

MICAFUNGIN SODIUM

INJECTABLE; IV (INFUSION)

MYCAMINE

ASTELLAS

100MG/VIAL

N21506 003 Jun 27, 2006 Jun NEWA

MICONAZOLE NITRATE

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

AB	ACTAVIS MID ATLANTIC	200MG	N73508 001	Nov 19, 1993	Jun	CAHN
	MONISTAT 3					
AB	+ PERSONAL PRODS	200MG	N18888 001	Aug 15, 1984	Aug	CAHN

MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE

OINTMENT; TOPICAL

VUSION

+ BARRIER

0.25%;81.35%;15%

N21026 001 Feb 16, 2006 Feb NEWA

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

AP	+ HOSPIRA	EQ 5MG BASE/ML	N75293 002	Jun 20, 2000	Mar	CRLD
----	-----------	----------------	------------	--------------	-----	------

MIDODRINE HYDROCHLORIDE

TABLET; ORAL

MIDODRINE HYDROCHLORIDE

AB	APOTEX INC	2.5MG	N77746 001	Sep 12, 2006	Aug	NEWA
AB		5MG	N77746 002	Sep 12, 2006	Aug	NEWA
AB		10MG	N77746 003	Sep 12, 2006	Aug	NEWA

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

MINOCIN

AB	TRIAx PHARMS	EQ 50MG BASE	N50649 001	May 31, 1990	Feb	CAHN
----	--------------	--------------	------------	--------------	-----	------

## CAPSULE; ORAL

## MINOCIN

	@ TRIAX PHARMS	EQ 75MG BASE	N50649 003	Feb 12, 2001	Feb	CAHN
AB	+	EQ 100MG BASE	N50649 002	May 31, 1990	Feb	CAHN

## TABLET, EXTENDED RELEASE; ORAL

## SOLODYN

	MEDICIS	EQ 45MG BASE	N50808 001	May 08, 2006	May	NEWA
		EQ 90MG BASE	N50808 002	May 08, 2006	May	NEWA
	+	EQ 135MG BASE	N50808 003	May 08, 2006	May	NEWA

MIRTAZAPINE

## TABLET; ORAL

## MIRTAZAPINE

AB	ACTAVIS ELIZABETH	15MG	N76308 001	Jun 20, 2003	Jun	CAHN
AB		30MG	N76308 002	Jun 20, 2003	Jun	CAHN
AB		45MG	N76308 003	Jun 20, 2003	Jun	CAHN

## TABLET, ORALLY DISINTEGRATING; ORAL

## MIRTAZAPINE

AB	AUROBINDO PHARMA LTD	45MG	N77376 004	Feb 28, 2006	Feb	NEWA
AB	BARR	45MG	N76307 003	Feb 28, 2006	Feb	NEWA

MITOXANTRONE HYDROCHLORIDE

## INJECTABLE; INJECTION

## MITOXANTRONE

AP	AM PHARM	EQ 20MG BASE/10ML (2MG/ML)	N77496 001	Apr 11, 2006	Mar	NEWA	
AP		EQ 25MG BASE/12.5ML (2MG/ML)	N77496 002	Apr 11, 2006	Mar	NEWA	
AP		EQ 30MG BASE/15ML (2MG/ML)	N77496 003	Apr 11, 2006	Mar	NEWA	
AP	BEDFORD	EQ 20MG BASE/10ML (2MG/ML)	N76611 001	Apr 11, 2006	Mar	NEWA	
AP		EQ 25MG BASE/12.5ML (2MG/ML)	N76611 002	Apr 11, 2006	Mar	NEWA	
AP		EQ 30MG BASE/15ML (2MG/ML)	N76611 003	Apr 11, 2006	Mar	NEWA	
AP	MAYNE PHARMA USA	EQ 20MG BASE/10ML (2MG/ML)	N76871 001	Apr 11, 2006	Mar	NEWA	
AP		EQ 25MG BASE/12.5ML (2MG/ML)	N76871 002	Apr 11, 2006	Mar	NEWA	
AP		EQ 30MG BASE/15ML (2MG/ML)	N76871 003	Apr 11, 2006	Mar	NEWA	
AP	SICOR PHARMS	EQ 20MG BASE/10ML (2MG/ML)	N77356 001	Apr 11, 2006	Mar	NEWA	
AP		EQ 25MG BASE/12.5ML (2MG/ML)	N77356 002	Apr 11, 2006	Mar	NEWA	
AP		EQ 30MG BASE/15ML (2MG/ML)	N77356 003	Apr 11, 2006	Mar	NEWA	
	NOVANTRONE						
AP	+	SERONO INC	EQ 20MG BASE/10ML (2MG/ML)	N19297 001	Dec 23, 1987	Mar	CFTG
AP	+		EQ 25MG BASE/12.5ML (2MG/ML)	N19297 002	Dec 23, 1987	Mar	CFTG
AP	+		EQ 30MG BASE/15ML (2MG/ML)	N19297 003	Dec 23, 1987	Mar	CFTG

>D> MIVACURIUM CHLORIDE

## &gt;D&gt; INJECTABLE; INJECTION

## &gt;D&gt; MIVACRON

>D>	+	ABBOTT	EQ 2MG BASE/ML	N20098 001	Jan 22, 1992	Sep	DISC
>A>		@	EQ 2MG BASE/ML	N20098 001	Jan 22, 1992	Sep	DISC
>D>		MIVACRON IN DEXTROSE 5%	IN PLASTIC CONTAINER				
>D>	+	ABBOTT	EQ 50MG BASE/100ML	N20098 003	Jan 22, 1992	Sep	DISC
>A>		@	EQ 50MG BASE/100ML	N20098 003	Jan 22, 1992	Sep	DISC

MOMETASONE FUROATE

## CREAM; TOPICAL

## MOMETASONE FUROATE

AB	G AND W LABS	0.1%	N77447 001	May 22, 2006	May	NEWA
----	--------------	------	------------	--------------	-----	------

## LOTION; TOPICAL

## MOMETASONE FUROATE

AB	PERRIGO	0.1%	N77180 001	Apr 06, 2005	Mar	CAHN
AB	TARO	0.1%	N76788 001	Mar 15, 2006	Feb	NEWA

## OINTMENT; TOPICAL

## MOMETASONE FUROATE

AB	G AND W LABS	0.1%	N77401 001	Jun 20, 2006	Jun	NEWA
----	--------------	------	------------	--------------	-----	------

MORPHINE SULFATE

## INJECTABLE; INJECTION

## MORPHINE SULFATE

## HOSPIRA

5MG/ML

N19916 002	Mar 30, 2006	Mar	NEWA
------------	--------------	-----	------

NABILONE

## CAPSULE; ORAL

## CESAMET

## + VALEANT

1MG

N18677 001	Dec 26, 1985	May	CMFD
------------	--------------	-----	------

NABUMETONE

## TABLET; ORAL

## NABUMETONE

## @ COPLEY PHARM

750MG

N75179 001	Jun 06, 2000	Jun	DISC
------------	--------------	-----	------

AB	PAR PHARM	500MG	N76009 001	Jan 24, 2003	Apr	CAHN
----	-----------	-------	------------	--------------	-----	------

AB		750MG	N76009 002	Jan 24, 2003	Apr	CAHN
----	--	-------	------------	--------------	-----	------

AB	+ TEVA	750MG	N75189 002	Sep 24, 2001	Jul	CRLD
----	--------	-------	------------	--------------	-----	------

## RELAFEN

## @ SMITHKLINE BEECHAM

500MG

N19583 001	Dec 24, 1991	Jul	DISC
------------	--------------	-----	------

## @

750MG

N19583 002	Dec 24, 1991	Jul	DISC
------------	--------------	-----	------

NADOLOL

## TABLET; ORAL

## CORGARD

AB	KING PHARMS	20MG	N18063 005	Oct 28, 1986	Jun	CAHN
----	-------------	------	------------	--------------	-----	------

AB		40MG	N18063 001		Jun	CAHN
----	--	------	------------	--	-----	------

AB		80MG	N18063 002		Jun	CAHN
----	--	------	------------	--	-----	------

AB		120MG	N18063 003		Jun	CAHN
----	--	-------	------------	--	-----	------

AB	+	160MG	N18063 004		Jun	CAHN
----	---	-------	------------	--	-----	------

NAFCILLIN SODIUM

## INJECTABLE; INJECTION

## NAFCILLIN SODIUM

AP	+ SANDOZ	EQ 1GM BASE/VIAL	N62527 002	Aug 02, 1984	Apr	CRLD
----	----------	------------------	------------	--------------	-----	------

AP	+	EQ 1GM BASE/VIAL	N62732 001	Dec 23, 1986	Apr	CRLD
----	---	------------------	------------	--------------	-----	------

AP	+	EQ 2GM BASE/VIAL	N62527 003	Aug 02, 1984	Apr	CRLD
----	---	------------------	------------	--------------	-----	------

AP	+	EQ 2GM BASE/VIAL	N62732 002	Dec 23, 1986	Apr	CRLD
----	---	------------------	------------	--------------	-----	------

AP	+	EQ 10GM BASE/VIAL	N62527 004	Aug 02, 1984	Apr	CRLD
----	---	-------------------	------------	--------------	-----	------

NAFTIFINE HYDROCHLORIDE

## CREAM; TOPICAL

## NAFTIN

>D>	MERZ PHARMS	1%	N19599 001	Feb 29, 1988	Sep	CRLD
-----	-------------	----	------------	--------------	-----	------

>A>	+	1%	N19599 001	Feb 29, 1988	Sep	CRLD
-----	---	----	------------	--------------	-----	------

NALTREXONEFOR SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR  
VIVITROL

+ ALKERMES 380MG/VIAL N21897 001 Apr 13, 2006 Apr NEWA

NAPROXEN

TABLET, DELAYED RELEASE; ORAL

NAPROXEN

AB ACTAVIS ELIZABETH 375MG N74936 001 Feb 24, 1998 Jun CAHN  
AB 500MG N74936 002 Feb 24, 1998 Jun CAHNNEOMYCIN SULFATE

TABLET; ORAL

NEOMYCIN SULFATE

@ BRISTOL MYERS SQUIBB 500MG N60365 001 Jul CPOT  
@ LANNETT 500MG N60607 001 Jul CPOT  
@ LILLY 500MG N60385 001 Jul CPOT  
@ ROXANE 500MG N62173 001 Jul CPOT  
@ SANDOZ 500MG N61586 001 Jul CPOT  
AB + TEVA 500MG N60304 001 Jul CFTG  
AB X GEN PHARMS 500MG N65220 001 Jul 28, 2006 Jul NEWANEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION; IRRIGATION

NEOMYCIN AND POLYMYXIN B SULFATE

AT WATSON LABS EQ 40MG BASE/ML;200,000 UNITS/ML N62664 001 Apr 08, 1986 Jul CMFD  
AT X GEN PHARMS EQ 40MG BASE/ML;200,000 UNITS/ML N65106 001 Jan 31, 2006 Jun CTNA  
AT EQ 800MG BASE/20ML;4,000,000 UNITS/20ML N65108 001 Jan 31, 2006 Jun CTNA  
NEOSPORIN AND POLYMYXIN B SULFATE  
AT X GEN PHARMS EQ 40MG BASE/ML;200,000 UNITS/ML N65106 001 Jan 31, 2006 Jan NEWA  
AT EQ 800MG BASE/20ML;4,000,000 UNITS/20ML (EQ 40MG BASE/ML;200,000 UNITS/ML) N65108 001 Jan 31, 2006 Jan NEWA  
NEOSPORIN G.U. IRRIGANT  
AT + MONARCH PHARMS EQ 40MG BASE/ML;200,000 UNITS/ML N60707 001 Jan CTEC  
AT + EQ 800MG BASE/20ML;4,000,000 UNITS/20ML (EQ 40MG BASE/ML;200,000 UNITS/ML) N60707 002 Jan NEWANIACIN

TABLET, EXTENDED RELEASE; ORAL

NIACIN

>D> NIACIN  
>D> AB BARR 500MG N76378 001 Apr 26, 2005 Sep DISC  
>A> @ 500MG N76378 001 Apr 26, 2005 Sep DISC  
>D> AB 750MG N76378 002 Apr 26, 2005 Sep DISC  
>A> @ 750MG N76378 002 Apr 26, 2005 Sep DISC  
>D> AB 1GM N76250 001 Apr 14, 2005 Sep DISC  
>A> @ 1GM N76250 001 Apr 14, 2005 Sep DISC  
NIASPAN  
>D> AB + KOS LIFE 500MG N20381 002 Jul 28, 1997 Sep CTEC  
>A> + 500MG N20381 002 Jul 28, 1997 Sep CTEC  
>D> AB + 750MG N20381 003 Jul 28, 1997 Sep CTEC  
>A> + 750MG N20381 003 Jul 28, 1997 Sep CTEC  
>D> AB + 1GM N20381 004 Jul 28, 1997 Sep CTEC  
>A> + 1GM N20381 004 Jul 28, 1997 Sep CTEC

NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL

NICARDIPINE HYDROCHLORIDE

AB	BARR	20MG	N74439 001	Dec 10, 1996	Apr	CAHN
AB		30MG	N74439 002	Dec 10, 1996	Apr	CAHN

INJECTABLE; INJECTION

CARDENE

+	PDL BIOPHARMA INC	2.5MG/ML	N19734 001	Jan 30, 1992	Jan	CAHN
---	-------------------	----------	------------	--------------	-----	------

NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

AB	ACTAVIS ELIZABETH	10MG	N72579 001	Jan 08, 1991	Jun	CAHN
AB		20MG	N72556 001	Sep 20, 1990	Jun	CAHN

TABLET, EXTENDED RELEASE; ORAL

AFEDITAB CR

AB1	WATSON LABS	30MG	N75128 001	Mar 10, 2000	Jan	CAHN
AB1		60MG	N75659 001	Oct 26, 2001	Jan	CAHN

NISOLDIPINE

TABLET, EXTENDED RELEASE; ORAL

SULAR

+	SCIELE PHARMA INC	10MG	N20356 001	Feb 02, 1995	Jun	CAHN
		20MG	N20356 002	Feb 02, 1995	Jun	CAHN
+		30MG	N20356 003	Feb 02, 1995	Jun	CAHN
+		40MG	N20356 004	Feb 02, 1995	Jun	CAHN

NITROGLYCERIN

AEROSOL; SUBLINGUAL

NITROLINGUAL

@	POHL BOSKAMP	0.4MG/SPRAY	N18705 001	Oct 31, 1985	Jun	CAHN
---	--------------	-------------	------------	--------------	-----	------

SPRAY, METERED; SUBLINGUAL

NITROLINGUAL PUMPSPRAY

+	POHL BOSKAMP	0.4MG/SPRAY	N18705 002	Jan 10, 1997	Jun	CAHN
---	--------------	-------------	------------	--------------	-----	------

NOREPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

NOREPINEPHRINE BITARTRATE

AP	METRICS PHARM	EQ 1MG BASE/ML	N40522 001	Sep 30, 2004	May	CAHN
----	---------------	----------------	------------	--------------	-----	------

NORTRIPTYLINE HYDROCHLORIDE

SOLUTION; ORAL

NORTRIPTYLINE HYDROCHLORIDE

AA	TARO	EQ 10MG BASE/5ML	N77965 001	Jun 20, 2006	Jun	NEWA
----	------	------------------	------------	--------------	-----	------

NYSTATIN

CREAM; TOPICAL

NYSTATIN

AT	ACTAVIS MID ATLANTIC	100,000 UNITS/GM	N62949 001	Jun 13, 1988	Jun	CAHN
	@ TARO	100,000 UNITS/GM	N62457 001	Jul 28, 1983	May	DISC
AT	VINTAGE	100,000 UNITS/GM	N65315 001	May 31, 2006	May	NEWA

OINTMENT; TOPICAL

NYSTATIN

AT	ACTAVIS MID ATLANTIC	100,000 UNITS/GM	N62840 001	Nov 13, 1987	Jun	CAHN
----	----------------------	------------------	------------	--------------	-----	------

## POWDER; ORAL

## NILSTAT

>D>	@ CLONMEL HLTHCARE	100%	N50576 001	Dec 22, 1983	Sep	CAHN
>A>	@ STADA PHARMS	100%	N50576 001	Dec 22, 1983	Sep	CAHN

## POWDER; TOPICAL

## NYSTATIN

AT	KV PHARM	100,000 UNITS/GM	N65321 001	Aug 18, 2006	Aug	NEWA
----	----------	------------------	------------	--------------	-----	------

## SUSPENSION; ORAL

## NYSTATIN

AA	ACTAVIS MID ATLANTIC	100,000 UNITS/ML	N62349 001	Jul 14, 1982	Jul	CAHN
AA	TARO	100,000 UNITS/ML	N62876 001	Feb 29, 1988	May	CMFD

NYSTATIN; TRIAMCINOLONE ACETONIDE

## CREAM; TOPICAL

## MYKACET

AT	ACTAVIS MID ATLANTIC	100,000 UNITS/GM;0.1%	N62367 001	May 28, 1985	Jun	CAHN
----	----------------------	-----------------------	------------	--------------	-----	------

## OINTMENT; TOPICAL

## MYKACET

AT	ACTAVIS MID ATLANTIC	100,000 UNITS/GM;0.1%	N62733 001	Mar 09, 1987	Jun	CAHN
----	----------------------	-----------------------	------------	--------------	-----	------

OCTREOTIDE ACETATE

## INJECTABLE; INJECTION

## OCTREOTIDE ACETATE

AP	AM PHARM	EQ 0.2MG BASE/ML	N77450 001	Feb 10, 2006	Jan	NEWA
AP		EQ 1MG BASE/ML	N77450 002	Feb 10, 2006	Jan	NEWA

## OCTREOTIDE ACETATE (PRESERVATIVE FREE)

AP	AM PHARM	EQ 0.05MG BASE/ML	N77457 001	Feb 10, 2006	Jan	NEWA
AP		EQ 0.1MG BASE/ML	N77457 002	Feb 10, 2006	Jan	NEWA
AP		EQ 0.5MG BASE/ML	N77457 003	Feb 10, 2006	Jan	NEWA

OFLOXACIN

## SOLUTION/DROPS; OTIC

## OFLOXACIN

AT	APOTEX INC	0.3%	N76527 001	Nov 18, 2005	Aug	CAHN
----	------------	------	------------	--------------	-----	------

## TABLET; ORAL

## OFLOXACIN

AB	DR REDDYS LABS LTD	200MG	N77098 001	Feb 10, 2006	Jan	NEWA
AB		300MG	N77098 002	Feb 10, 2006	Jan	NEWA
AB		400MG	N77098 003	Feb 10, 2006	Jan	NEWA

OMEPRAZOLE; SODIUM BICARBONATE

## CAPSULE; ORAL

## ZEGERID

	SANTARUS	20MG;1.1GM	N21849 001	Feb 27, 2006	Feb	NEWA
+		40MG;1.1GM	N21849 002	Feb 27, 2006	Feb	NEWA

## FOR SUSPENSION; ORAL

## ZEGERID

	SANTARUS	20MG/PACKET;1.68GM/PACKET	N21636 001	Jun 15, 2004	Feb	CAIN
+		40MG/PACKET;1.68GM/PACKET	N21706 001	Dec 21, 2004	Feb	CAIN

ORPHENADRINE CITRATE

## INJECTABLE; INJECTION

## ORPHENADRINE CITRATE

AP	AKORN	30MG/ML	N40484 001	May 24, 2006	May	NEWA
----	-------	---------	------------	--------------	-----	------

OXALIPLATIN

INJECTABLE; IV (INFUSION)

ELOXATIN

@ SANOFI AVENTIS US 50MG/VIAL  
 @ 100MG/VIAL

N21492 001 Aug 09, 2002 Jun DISC  
 N21492 002 Aug 09, 2002 Jun DISC

OXAPROZIN

TABLET; ORAL

OXAPROZIN

AB ACTAVIS ELIZABETH 600MG

N75843 001 Oct 03, 2001 Jun CAHN

OXAZEPAM

CAPSULE; ORAL

OXAZEPAM

AB ACTAVIS ELIZABETH 10MG  
 AB 15MG  
 AB 30MG

N72251 001 Apr 14, 1988 Jun CAHN  
 N72252 001 Apr 14, 1988 Jun CAHN  
 N72253 001 Apr 14, 1988 Jun CAHN

OXYCODONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

OXYCONTIN

>A> PURDUE PHARMA LP 15MG  
 >A> 30MG  
 >A> 60MG

N20553 006 Sep 18, 2006 Sep NEWA  
 N20553 007 Sep 18, 2006 Sep NEWA  
 N20553 008 Sep 18, 2006 Sep NEWA

OXYMETHOLONE

TABLET; ORAL

ANADROL-50

+ ALAVEN PHARM 50MG

N16848 001 Apr CAHN

OXYMORPHONE HYDROCHLORIDE

TABLET; ORAL

OPANA

ENDO PHARMS 5MG  
 + 10MG

N21611 001 Jun 22, 2006 Jun NEWA  
 N21611 002 Jun 22, 2006 Jun NEWA

TABLET, EXTENDED RELEASE; ORAL

OPANA ER

ENDO PHARMS 5MG  
 10MG  
 20MG  
 + 40MG

N21610 001 Jun 22, 2006 Jun NEWA  
 N21610 002 Jun 22, 2006 Jun NEWA  
 N21610 003 Jun 22, 2006 Jun NEWA  
 N21610 004 Jun 22, 2006 Jun NEWA

PACLITAXEL

FOR SUSPENSION; IV (INFUSION)

ABRAXANE

+ ABRAXIS BIOSCIENCE 100MG/VIAL

N21660 001 Jan 07, 2005 Apr CAHN

PARICALCITOL

INJECTABLE; INJECTION

ZEMPLAR

+ ABBOTT 0.002MG/ML

N20819 002 Feb 01, 2000 Aug CRLD



PAROXETINE MESYLATE

TABLET; ORAL

PEXEVA

JDS PHARMS	EQ 10MG BASE	N21299 001	Jul 03, 2003	Apr	CAHN
	EQ 20MG BASE	N21299 002	Jul 03, 2003	Apr	CAHN
	EQ 30MG BASE	N21299 003	Jul 03, 2003	Apr	CAHN
+	EQ 40MG BASE	N21299 004	Jul 03, 2003	Apr	CAHN

PEGAPTANIB SODIUM

INJECTABLE; INTRAVITREAL

MACUGEN

+	OSI EYETECH	EQ 0.3MG ACID/0.09ML	N21756 001	Dec 17, 2004	May	CAHN
---	-------------	----------------------	------------	--------------	-----	------

PENICILLAMINE

CAPSULE; ORAL

CUPRIMINE

@	ATON	125MG	N19853 002		Aug	CAHN
+		250MG	N19853 001		Aug	CAHN
@	MERCK	125MG	N19853 002		Jun	DISC

PENICILLIN V POTASSIUM

FOR SOLUTION; ORAL

PENICILLIN V POTASSIUM

AA	AM ANTIBIOTICS	EQ 125MG BASE/5ML	N61529 001		Jan	CAHN
AA		EQ 250MG BASE/5ML	N61529 002		Jan	CAHN

TABLET; ORAL

PENICILLIN V POTASSIUM

@	AM ANTIBIOTICS	EQ 250MG BASE	N61528 001		Jan	CAHN
@		EQ 500MG BASE	N61528 002		Jan	CAHN

PENTOSTATIN

INJECTABLE; INJECTION

NIPENT

+	MAYNE PHARMA USA	10MG/VIAL	N20122 001	Oct 11, 1991	Aug	CAHN
---	------------------	-----------	------------	--------------	-----	------

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

AB	ACTAVIS ELIZABETH	400MG	N74878 001	Jul 09, 1997	Jun	CAHN
AB	RADIUS PHARMS	400MG	N74877 001	Jul 08, 1997	Jul	CAHN

PERGOLIDE MESYLATE

TABLET; ORAL

PERGOLIDE MESYLATE

AB	PAR PHARM	EQ 0.05MG BASE	N76061 001	Nov 27, 2002	Apr	CAHN
AB		EQ 0.25MG BASE	N76061 002	Nov 27, 2002	Apr	CAHN
AB		EQ 1MG BASE	N76061 003	Nov 27, 2002	Apr	CAHN
	PERMAX					
AB	VALEANT	EQ 0.05MG BASE	N19385 001	Dec 30, 1988	Jan	CRLD
AB	+	EQ 0.25MG BASE	N19385 002	Dec 30, 1988	Jan	CRLD

PERMETHRIN

CREAM; TOPICAL

PERMETHRIN

AB	ACTAVIS MID ATLANTIC	5%	N74806	001	Jan 23, 1998	Jun	CAHN
----	----------------------	----	--------	-----	--------------	-----	------

PHENDIMETRAZINE TARTRATE

TABLET; ORAL

CAM-METRAZINE

	@ TG UNITED LABS	35MG	N83922	001		May	CAHN
	@	35MG	N85318	001		May	CAHN
	@	35MG	N85320	001		May	CAHN
	@	35MG	N85321	001		May	CAHN

PHENDIMETRAZINE TARTRATE

	@ TG UNITED LABS	35MG	N85761	001		May	CAHN
	@	35MG	N85941	001	Jun 27, 1983	May	CAHN

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHENTERMINE HYDROCHLORIDE

	@ TG UNITED LABS	18.75MG	N88576	001	May 23, 1984	May	CAHN
	@	30MG	N85417	001		May	CAHN
	@	30MG	N86732	002		May	CAHN
	@	30MG	N87215	001		May	CAHN
	@	37.5MG	N87915	001	Dec 22, 1983	May	CAHN
	@	37.5MG	N87918	001	Dec 22, 1983	May	CAHN
	@	37.5MG	N87930	001	Oct 14, 1983	May	CAHN
	@	37.5MG	N88610	001	Jun 04, 1984	May	CAHN
	@	37.5MG	N88611	001	Jun 04, 1984	May	CAHN
	@	37.5MG	N88625	001	Aug 23, 1984	May	CAHN

TABLET; ORAL

PHENTERMINE HYDROCHLORIDE

AA	ACTAVIS ELIZABETH	37.5MG	N40276	001	Nov 25, 1998	Jun	CAHN
	@ TG UNITED LABS	8MG	N83923	001		May	CAHN
	@	8MG	N85319	001		May	CAHN
	@	37.5MG	N87805	001	Dec 06, 1982	May	CAHN
	@	37.5MG	N88596	001	Apr 04, 1984	May	CAHN

PHENYLEPHRINE HYDROCHLORIDE; PROMETHAZINE HYDROCHLORIDE

SYRUP; ORAL

PROMETH VC PLAIN

+	ACTAVIS MID ATLANTIC	5MG/5ML; 6.25MG/5ML	N88761	001	Nov 08, 1984	Jul	CAHN
+	ALPHARMA US PHARMS	5MG/5ML; 6.25MG/5ML	N88761	001	Nov 08, 1984	Jan	CTEC
	PROMETHAZINE VC PLAIN						
	@ MORTON GROVE	5MG/5ML; 6.25MG/5ML	N88897	001	Jan 04, 1985	Jan	DISC

PHENYTOIN

SUSPENSION; ORAL

PHENYTOIN

AB	ACTAVIS MID ATLANTIC	125MG/5ML	N89892	001	Sep 25, 1992	Jul	CAHN
----	----------------------	-----------	--------	-----	--------------	-----	------

PHENYTOIN SODIUM

CAPSULE; ORAL

EXTENDED PHENYTOIN SODIUM

AB	TARO	100MG EXTENDED	N40684	001	Sep 05, 2006	Aug	NEWA
----	------	----------------	--------	-----	--------------	-----	------

## INJECTABLE; INJECTION

## PHENYTOIN SODIUM

AP	HIKMA FARMACEUTICA	50MG/ML	N40573 001	Sep 13, 2006	Aug	NEWA
----	--------------------	---------	------------	--------------	-----	------

PHYTONADIONE

## INJECTABLE; INJECTION

## AQUAMEPHYTON

@ MERCK

1MG/0.5ML

N12223 002

Feb DISC

@

10MG/ML

N12223 001

Feb DISC

## VITAMIN K1

BP	+	HOSPIRA	1MG/0.5ML	N87954 001	Jul 25, 1983	Feb	CRLD
----	---	---------	-----------	------------	--------------	-----	------

	+		10MG/ML	N87955 001	Jul 25, 1983	Feb	CRLD
--	---	--	---------	------------	--------------	-----	------

PILOCARPINE HYDROCHLORIDE

## TABLET; ORAL

## PILOCARPINE HYDROCHLORIDE

AB	IMPAX LABS	5MG	N77248 001	Mar 31, 2006	Mar	NEWA
----	------------	-----	------------	--------------	-----	------

AB		7.5MG	N77248 002	Mar 31, 2006	Mar	NEWA
----	--	-------	------------	--------------	-----	------

## SALAGEN

AB	+	MGI PHARMA INC	7.5MG	N20237 002	Apr 18, 2003	Mar	CFTG
----	---	----------------	-------	------------	--------------	-----	------

POLYETHYLENE GLYCOL 3350

## FOR SOLUTION; ORAL

## POLYETHYLENE GLYCOL 3350

AA	COASTAL PHARMS	17GM/SCOOPFUL	N77893 001	May 26, 2006	May	NEWA
----	----------------	---------------	------------	--------------	-----	------

AA	KALI LABS	17GM/SCOOPFUL	N77736 001	May 26, 2006	May	NEWA
----	-----------	---------------	------------	--------------	-----	------

AA	TEVA PHARMS	17GM/SCOOPFUL	N77445 001	May 04, 2006	Apr	NEWA
----	-------------	---------------	------------	--------------	-----	------

>A>	AA	YVR THERAP	17GM/SCOOPFUL	N77706 001	Sep 27, 2006	Sep	NEWA
-----	----	------------	---------------	------------	--------------	-----	------

>A> POSACONAZOLE

## &gt;A&gt; SUSPENSION; ORAL

## &gt;A&gt; NOXAFIL

>A>	+	SCHERING	40MG/ML	N22003 001	Sep 15, 2006	Sep	NEWA
-----	---	----------	---------	------------	--------------	-----	------

POTASSIUM CHLORIDE

## INJECTABLE; INJECTION

## POTASSIUM CHLORIDE 10MEQ IN PLASTIC CONTAINER

AP	+	BAXTER HLTHCARE	14.9MG/ML	N19904 001	Dec 26, 1989	Jun	CTEC
----	---	-----------------	-----------	------------	--------------	-----	------

AP	+		746MG/100ML	N19904 005	Dec 17, 1990	Jun	CTEC
----	---	--	-------------	------------	--------------	-----	------

## POTASSIUM CHLORIDE 20MEQ IN PLASTIC CONTAINER

AP	+	BAXTER HLTHCARE	1.49GM/100ML	N19904 006	Dec 17, 1990	Jun	CTEC
----	---	-----------------	--------------	------------	--------------	-----	------

## TABLET, EXTENDED RELEASE; ORAL

## KLOR-CON

AB	UPSHER SMITH	8MEQ	N19123 001	Apr 17, 1986	Jan	CRLD
----	--------------	------	------------	--------------	-----	------

## POTASSIUM CHLORIDE

AB	+	COPELY PHARM	8MEQ	N70618 001	Sep 09, 1987	Jan	CRLD
----	---	--------------	------	------------	--------------	-----	------

POTASSIUM CITRATE

## TABLET, EXTENDED RELEASE; ORAL

## POTASSIUM CITRATE

AB	COREPHARMA	5MEQ	N77440 001	Jun 09, 2006	May	NEWA
----	------------	------	------------	--------------	-----	------

AB		10MEQ	N77440 002	Jun 09, 2006	May	NEWA
----	--	-------	------------	--------------	-----	------

## UROCIT-K

AB	MISSION PHARMA	5MEQ	N19071 001	Aug 30, 1985	May	CFTG
----	----------------	------	------------	--------------	-----	------

AB	+		10MEQ	N19071 002	Aug 31, 1992	May	CFTG
----	---	--	-------	------------	--------------	-----	------

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOL

AB	BRISTOL MYERS SQUIBB	10MG	N19898 002	Oct 31, 1991	Apr	CFTG
AB		20MG	N19898 003	Oct 31, 1991	Apr	CFTG
AB		40MG	N19898 004	Mar 22, 1993	Apr	CFTG
PRAVASTATIN SODIUM						
AB	TEVA	10MG	N76056 001	Apr 24, 2006	Apr	NEWA
AB		20MG	N76056 002	Apr 24, 2006	Apr	NEWA
AB		40MG	N76056 003	Apr 24, 2006	Apr	NEWA

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

@ CLONMEL HLTHCARE

EQ 1MG BASE

N72705 001 May 16, 1989 Jan DISC

@

EQ 5MG BASE

N72707 001 May 16, 1989 Jan DISC

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLLIENT

>D>	+	SANOFI AVENTIS US	0.1%	N20279 001	Oct 29, 1993	Sep	CFTG
>A>	AB	+	0.1%	N20279 001	Oct 29, 1993	Sep	CFTG
>A>		PREDNICARBATE					
>A>	AB	ALTANA	0.1%	N77287 001	Sep 19, 2006	Sep	NEWA

PREDNISOLONE ACETATE

INJECTABLE; INJECTION

PREDNISOLONE ACETATE

@ STERIS

25MG/ML

N83398 001

Mar DISC

@

50MG/ML

N83764 001

Mar DISC

SUSPENSION/DROPS; OPHTHALMIC

OMNIPRED

AB	ALCON	1%	N17469 001		May	CTNA
----	-------	----	------------	--	-----	------

PREDNISOLONE SODIUM PHOSPHATE

TABLET, ORALLY DISINTEGRATING; ORAL

ORAPRED ODT

BIOMARIN PHARM

EQ 10MG BASE

N21959 001 Jun 01, 2006 Jun NEWA

EQ 15MG BASE

N21959 002 Jun 01, 2006 Jun NEWA

+

EQ 30MG BASE

N21959 003 Jun 01, 2006 Jun NEWA

PRIMIDONE

TABLET; ORAL

PRIMIDONE

AB	WEST WARD	50MG	N40667 001	Jul 27, 2006	Jul	NEWA
AB		250MG	N40667 002	Jul 27, 2006	Jul	NEWA

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPAZINE

@ GLAXOSMITHKLINE

2.5MG

N11127 003

May DISC

@

5MG

N11127 001

May DISC

PROCHLORPERAZINE EDISYLATE

SYRUP; ORAL

COMPAZINE

@ GLAXOSMITHKLINE	EQ 5MG BASE/5ML	N11188 001		May	DISC
-------------------	-----------------	------------	--	-----	------

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

COMPAZINE

@ GLAXOSMITHKLINE	EQ 10MG BASE	N21019 001	Oct 06, 1999	May	DISC
-------------------	--------------	------------	--------------	-----	------

PROGESTERONE

GEL; VAGINAL

CRINONE

COLUMBIA LABS	4%	N20701 001	Jul 31, 1997	May	CAHN
+	8%	N20701 002	Jul 31, 1997	May	CAHN

PROMETHAZINE HYDROCHLORIDE

SUPPOSITORY; RECTAL

PHENERGAN

@ WYETH PHARMS INC	12.5MG	N10926 002		Mar	DISC
--------------------	--------	------------	--	-----	------

@	25MG	N10926 001		Mar	DISC
---	------	------------	--	-----	------

PROMETHAZINE HYDROCHLORIDE

AB	+	G AND W LABS	25MG	N40428 001	Feb 05, 2002	Mar	CRLD
----	---	--------------	------	------------	--------------	-----	------

PROMETHEGAN

+	G AND W LABS	50MG	N87165 001	Aug 14, 1987	Jan	CRLD
---	--------------	------	------------	--------------	-----	------

SYRUP; ORAL

PROMETH PLAIN

@ ACTAVIS MID ATLANTIC	6.25MG/5ML	N85953 001		Jul	CAHN
------------------------	------------	------------	--	-----	------

PROMETHAZINE HYDROCHLORIDE

AA	VINTAGE	6.25MG/5ML	N40643 001	Apr 26, 2006	Apr	NEWA
----	---------	------------	------------	--------------	-----	------

TABLET; ORAL

PROMETHAZINE HYDROCHLORIDE

AB	KVK-TECH INC	25MG	N40712 001	Jul 31, 2006	Jul	NEWA
----	--------------	------	------------	--------------	-----	------

AB		50MG	N40713 001	Jul 31, 2006	Jul	NEWA
----	--	------	------------	--------------	-----	------

AB	VINTAGE PHARMS	12.5MG	N40622 001	Jul 18, 2006	Jul	NEWA
----	----------------	--------	------------	--------------	-----	------

AB		25MG	N40622 002	Jul 18, 2006	Jul	NEWA
----	--	------	------------	--------------	-----	------

AB		50MG	N40622 003	Jul 18, 2006	Jul	NEWA
----	--	------	------------	--------------	-----	------

AB		50MG	N40622 003	Jul 18, 2006	Jun	NEWA
----	--	------	------------	--------------	-----	------

AB	ZYDUS PHARMS USA	12.5MG	N40596 001	Nov 18, 2005	Jul	CTEC
----	------------------	--------	------------	--------------	-----	------

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

AB	+	ABRAXIS BIOSCIENCE	10MG/ML	N19627 002	Jun 11, 1996	Jul	CAHN
----	---	--------------------	---------	------------	--------------	-----	------

@	10MG/ML	N19627 001	Oct 02, 1989	Jul	CAHN
---	---------	------------	--------------	-----	------

PROPOFOL

AB	HOSPIRA	10MG/ML	N77908 001	Mar 17, 2006	Mar	NEWA
----	---------	---------	------------	--------------	-----	------

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DOLENE

@ RADIUS PHARMS	65MG	N80530 001		Aug	CAHN
-----------------	------	------------	--	-----	------

PROPOXYPHENE HYDROCHLORIDE

AA	PAR PHARM	65MG	N80269 001		Mar	CAHN
----	-----------	------	------------	--	-----	------

PROPRANOLOL HYDROCHLORIDE

INJECTABLE; INJECTION  
PROPRANOLOL HYDROCHLORIDE

AP		SANDOZ	1MG/ML	N76400 001	Feb 26, 2003	Jan	CAHN
----	--	--------	--------	------------	--------------	-----	------

PROPYLTHIOURACIL

TABLET; ORAL  
PROPYLTHIOURACIL

BD		ACTAVIS ELIZABETH	50MG	N80172 001		Jun	CAHN
>D>	BD	+ CLONMEL HLTHCARE	50MG	N06188 001		Sep	CAHN
>A>	BD	+ STADA PHARMS	50MG	N06188 001		Sep	CAHN

PYRIDOSTIGMINE BROMIDE

INJECTABLE; INJECTION  
REGONOL

AP		SANDOZ	5MG/ML	N17398 001		Jan	CAHN
----	--	--------	--------	------------	--	-----	------

QUAZEPAM

TABLET; ORAL  
DORAL

@ QUESTCOR PHARMS  
+

7.5MG  
15MG

N18708 003	Feb 26, 1987	May	CAHN
N18708 001	Dec 27, 1985	May	CAHN

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL  
QUINAPRIL

AB		LUPIN	EQ 5MG BASE	N77690 001	Jun 20, 2006	Jun	NEWA
AB			EQ 10MG BASE	N77690 002	Jun 20, 2006	Jun	NEWA
AB			EQ 20MG BASE	N77690 003	Jun 20, 2006	Jun	NEWA
AB			EQ 40MG BASE	N77690 004	Jun 20, 2006	Jun	NEWA

QUINAPRIL HYDROCHLORIDE

AB		TORPHARM	EQ 5MG BASE	N76240 001	Jan 26, 2006	Jan	NEWA
AB			EQ 10MG BASE	N76240 002	Jan 26, 2006	Jan	NEWA
AB			EQ 20MG BASE	N76240 003	Jan 26, 2006	Jan	NEWA
AB			EQ 40MG BASE	N76240 004	Jan 26, 2006	Jan	NEWA

QUINIDINE GLUCONATE

TABLET, EXTENDED RELEASE; ORAL  
QUINIDINE GLUCONATE

BX	+	MUTUAL PHARM	324MG	N89338 001	Feb 11, 1987	Jan	CTEC
BX		WATSON LABS	324MG	N87810 001	Sep 29, 1982	Jan	CMFD

QUINIDINE SULFATE

TABLET; ORAL

QUINIDINE SULFATE  
@ CLONMEL HLTHCARE  
@ LANNETT  
@ MUTUAL PHARM

			200MG	N87011 001		Jan	DISC
			200MG	N83743 001		Jan	DISC
			100MG	N81029 001	Apr 14, 1989	Jan	DISC
AB			300MG	N81031 001	Apr 14, 1989	Jul	CMFD
		@ PHARM FORM	200MG	N83808 001		Jan	DISC
		@ SANDOZ	200MG	N84631 001		Jan	DISC
		@	200MG	N84914 001		Jan	DISC
AB			200MG	N88072 002		Jan	NEWA
		@	300MG	N89839 001	Sep 29, 1988	Jan	DISC

## TABLET; ORAL

## QUINIDINE SULFATE

AB	WATSON LABS	200MG	N83288 001	Jul	CMFD
	@	200MG	N83288 001	Jan	DISC
	@	200MG	N85140 002	Jan	DISC

QUININE SULFATE

## CAPSULE; ORAL

## QUININE SULFATE

>A>	+	AR HOLDING CO INC	324MG	N21799 001	Aug 12, 2005	Sep	CAHN
>D>	+	MUTUAL PHARM	324MG	N21799 001	Aug 12, 2005	Sep	CAHN

RANITIDINE

## INJECTABLE; INJECTION

## RANITIDINE

AP	BEDFORD	EQ 25MG BASE/ML	N77458 001	Feb 16, 2006	Feb	NEWA
----	---------	-----------------	------------	--------------	-----	------

RANOLAZINE

## TABLET, EXTENDED RELEASE; ORAL

## RANEXA

	+	CV THERAP	500MG	N21526 002	Jan 27, 2006	Jan	NEWA
--	---	-----------	-------	------------	--------------	-----	------

RASAGILINE MESYLATE

## TABLET; ORAL

## AZILECT

	+	TEVA	EQ 0.5MG BASE	N21641 001	May 16, 2006	May	NEWA
			EQ 1MG BASE	N21641 002	May 16, 2006	May	NEWA

RIBAVIRIN

## TABLET; ORAL

## RIBAVIRIN

>A>	AB	SANDOZ	200MG	N77743 001	Oct 03, 2006	Sep	NEWA
-----	----	--------	-------	------------	--------------	-----	------

RISPERIDONE

## TABLET, ORALLY DISINTEGRATING; ORAL

## RISPERDAL

		JANSSEN PHARMA	3MG	N21444 004	Dec 23, 2004	Mar	CMFD
			4MG	N21444 005	Dec 23, 2004	Mar	CMFD

ROPIVACAINE HYDROCHLORIDE MONOHYDRATE

## INJECTABLE; INJECTION

## NAROPIN

		ABRAXIS BIOSCIENCE	2MG/ML	N20533 001	Sep 24, 1996	Jul	CAHN
			5MG/ML	N20533 003	Sep 24, 1996	Jul	CAHN
			7.5MG/ML	N20533 004	Sep 24, 1996	Jul	CAHN
	+		10MG/ML	N20533 005	Sep 24, 1996	Jul	CAHN

SECRETIN SYNTHETIC PORCINE

## FOR SOLUTION; INTRAVENOUS

## SECFLO

	+	CHIRHOCLIN	16UGM/VIAL	N21136 001	Apr 04, 2002	May	CTNA
--	---	------------	------------	------------	--------------	-----	------

SELEGILINE

FILM, EXTENDED RELEASE; TRANSDERMAL  
EMSAM

	SOMERSET	6MG/24HR	N21336 001	Feb 27, 2006	Mar	CAIN
		9MG/24HR	N21336 002	Feb 27, 2006	Mar	CAIN
+		12MG/24HR	N21336 003	Feb 27, 2006	Mar	CAIN

SELEGILINE HYDROCHLORIDE

CAPSULE; ORAL

>D>	SELEGILINE HYDROCHLORIDE					
>D>	AB	AAIPHARMA LLC	5MG	N75145 001	Sep 15, 2003	Sep DISC
>A>		@	5MG	N75145 001	Sep 15, 2003	Sep DISC

FILM, EXTENDED RELEASE; TRANSDERMAL  
EMSAM

	SOMERSET	6MG/24HR	N21336 001	Feb 27, 2006	Feb	NEWA
		9MG/24HR	N21336 002	Feb 27, 2006	Feb	NEWA
+		12MG/24HR	N21336 003	Feb 27, 2006	Feb	NEWA

TABLET, ORALLY DISINTEGRATING; ORAL  
ZELAPAR

+	VALEANT PHARM INTL	1.25MG	N21479 001	Jun 14, 2006	Jun	NEWA
---	--------------------	--------	------------	--------------	-----	------

SELENIUM SULFIDE

LOTION/SHAMPOO; TOPICAL

SELENIUM SULFIDE

AT	ACTAVIS MID ATLANTIC	2.5%	N84394 001		Jul	CAHN
----	----------------------	------	------------	--	-----	------

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

AB	ROXANE	EQ 20MG BASE/ML	N76934 001	Jun 30, 2006	Jun	NEWA
----	--------	-----------------	------------	--------------	-----	------

ZOLOFT

AB	+	PFIZER	EQ 20MG BASE/ML	N20990 001	Dec 07, 1999	Jun CFTG
----	---	--------	-----------------	------------	--------------	----------

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

AB	IVAX PHARMS	EQ 25MG BASE	N75719 003	Jun 30, 2006	Jun	NEWA
AB		EQ 50MG BASE	N75719 001	Jun 30, 2006	Jun	NEWA
AB		EQ 100MG BASE	N75719 002	Jun 30, 2006	Jun	NEWA
AB	TEVA	EQ 25MG BASE	N76465 001	Aug 11, 2006	Aug	NEWA
AB		EQ 50MG BASE	N76465 002	Aug 11, 2006	Aug	NEWA
AB		EQ 100MG BASE	N76465 003	Aug 11, 2006	Aug	NEWA

ZOLOFT

AB	PFIZER	EQ 25MG BASE	N19839 005	Mar 06, 1996	Jun	CFTG
AB		EQ 50MG BASE	N19839 001	Dec 30, 1991	Jun	CFTG
AB	+	EQ 100MG BASE	N19839 002	Dec 30, 1991	Jun	CFTG

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

AB	IVAX PHARMS	5MG	N76052 001	Jun 23, 2006	Jun	NEWA
AB		10MG	N76052 002	Jun 23, 2006	Jun	NEWA
AB		20MG	N76052 003	Jun 23, 2006	Jun	NEWA
AB		40MG	N76052 004	Jun 23, 2006	Jun	NEWA
AB	RANBAXY	80MG	N76285 005	Jun 23, 2006	Jun	NEWA



## TABLET; ORAL

## ZOCOR

AB		MERCK	5MG	N19766 001	Dec 23, 1991	Jun	CFTG
AB			10MG	N19766 002	Dec 23, 1991	Jun	CFTG
AB			20MG	N19766 003	Dec 23, 1991	Jun	CFTG
AB			40MG	N19766 004	Dec 23, 1991	Jun	CFTG
AB	+		80MG	N19766 005	Jul 10, 1998	Jun	CFTG

SODIUM CHLORIDE

## INJECTABLE; INJECTION

## SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP	+	B BRAUN	900MG/100ML	N17464 001		May	CRLD
AP	+		900MG/100ML	N19635 002	Mar 09, 1988	May	CRLD
AP	+	BAXTER HLTHCARE	900MG/100ML	N16677 001		May	CRLD
AP	+		900MG/100ML	N20178 001	Dec 07, 1992	May	CRLD
AP	+	HOSPIRA	900MG/100ML	N16366 001		May	CRLD
AP	+		900MG/100ML	N19465 001	Jul 15, 1985	May	CRLD
AP	+		900MG/100ML	N19480 001	Sep 17, 1985	May	CRLD
	+	MALLINCKRODT	45MG/50ML (9MG/ML)	N21569 001	Jul 27, 2006	Aug	CDFR
			112.5MG/125ML (9MG/ML)	N21569 002	Jul 27, 2006	Aug	CDFR
AP		TARO PHARMS IRELAND	9MG/ML	N77407 001	Aug 11, 2006	Aug	NEWA

## INJECTABLE; INTRAVASCULAR

## SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

	+	MALLINCKRODT	45MG/50ML (0.9MG/ML)	N21569 001	Jul 27, 2006	Jul	NEWA
			112.5MG/125ML (0.9MG/ML)	N21569 002	Jul 27, 2006	Jul	NEWA

SODIUM IODIDE, I-123

## CAPSULE; ORAL

## SODIUM IODIDE I 123

>D>	AA	GE HEALTHCARE	100uCi	N17630 001		Sep	CRLD
>A>	AA	+	100uCi	N17630 001		Sep	CRLD
>D>	AA	SYNCOR PHARMS	100uCi	N18671 001	May 27, 1982	Sep	CRLD
>A>	AA	+	100uCi	N18671 001	May 27, 1982	Sep	CRLD
>D>	AA		200uCi	N18671 002	May 27, 1982	Sep	CRLD
>A>	AA	+	200uCi	N18671 002	May 27, 1982	Sep	CRLD

## SOLUTION; ORAL

## SODIUM IODIDE I 123

>D>		GE HEALTHCARE	2mCi/ML	N17630 002		Sep	CRLD
>A>		+	2mCi/ML	N17630 002		Sep	CRLD

SODIUM IODIDE, I-131

## CAPSULE; ORAL

## SODIUM IODIDE I 131

		DRAXIMAGE	100mCi	N21305 004	Nov 18, 2004	Jun	NEWA
--	--	-----------	--------	------------	--------------	-----	------

SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE

## TABLET; ORAL

## OSMOPREP

	+	SALIX PHARMS	0.398GM;1.102GM	N21892 001	Mar 16, 2006	Mar	NEWA
--	---	--------------	-----------------	------------	--------------	-----	------

SOMATROPIN RECOMBINANT

## INJECTABLE; INJECTION

## GENOTROPIN

BX	+	PHARMACIA AND UPJOHN	5.8MG/VIAL	N20280 006	Aug 24, 1995	May	CTEC
----	---	----------------------	------------	------------	--------------	-----	------

## INJECTABLE; INJECTION

GENOTROPIN PRESERVATIVE FREE									
BX	PHARMACIA AND UPJOHN	1.5MG/VIAL	N20280	004	Aug 24, 1995	May	CTEC		
NUTROPIN DEPOT									
	@ GENENTECH	13.5MG/VIAL	N21075	001	Dec 22, 1999	Jul	DISC		
	@	18MG/VIAL	N21075	002	Dec 22, 1999	Jul	DISC		
	@	22.5MG/VIAL	N21075	003	Dec 22, 1999	Jul	DISC		
OMNITROPE									
BX	SANDOZ	1.5MG/VIAL	N21426	002	May 30, 2006	May	NEWA		
BX		5.8MG/VIAL	N21426	001	May 30, 2006	May	NEWA		

SPIRONOLACTONE

## TABLET; ORAL

SPIRONOLACTONE									
AB	ACTAVIS ELIZABETH	25MG	N40353	003	Mar 15, 2006	Jun	CAHN		
AB		50MG	N40353	001	Jul 29, 1999	Jun	CAHN		
AB		100MG	N40353	002	Jul 29, 1999	Jun	CAHN		
AB	PUREPAC PHARM	25MG	N40353	003	Mar 15, 2006	Feb	NEWA		
AB	VINTAGE	25MG	N40750	001	Aug 29, 2006	Aug	NEWA		
AB		50MG	N40750	002	Aug 29, 2006	Aug	NEWA		
AB		100MG	N40750	003	Aug 29, 2006	Aug	NEWA		

SUCCINYLCHOLINE CHLORIDE

## INJECTABLE; INJECTION

ANECTINE									
AP	+ SANDOZ	20MG/ML	N08453	002		Jan	CAHN		
	@	50MG/ML	N08453	003		Jan	CAHN		
	@	500MG/VIAL	N08453	001		Jan	CAHN		
	@	1GM/VIAL	N08453	004		Jan	CAHN		
SUCCINYLCHOLINE CHLORIDE									
	@ ORGANON USA INC	20MG/ML	N80997	001		Jun	DISC		

SULCONAZOLE NITRATE

## SOLUTION; TOPICAL

EXELDERM									
	+ WESTWOOD SQUIBB	1%	N18738	001	Aug 30, 1985	Jun	CMFD		

SULFACETAMIDE SODIUM

## SOLUTION/DROPS; OPHTHALMIC

SULFACETAMIDE SODIUM									
AT	ALCON	30%	N89068	001	May 05, 1987	Apr	CAHN		

SULFAMETHOXAZOLE; TRIMETHOPRIM

## SUSPENSION; ORAL

BACTRIM PEDIATRIC									
AB	MUTUAL PHARM	200MG/5ML; 40MG/5ML	N17560	002		Apr	CMFD		
SULFATRIM									
AB	ACTAVIS MID ATLANTIC	200MG/5ML; 40MG/5ML	N18615	002	Jan 07, 1983	Jul	CAHN		
SULFATRIM PEDIATRIC									
AB	ACTAVIS MID ATLANTIC	200MG/5ML; 40MG/5ML	N18615	001	Jan 07, 1983	Jul	CAHN		

SULFASALAZINE

## TABLET; ORAL

SULFASALAZINE									
	@ RADIUS PHARMS	500MG	N80197	001		Aug	CAHN		

SULINDAC

TABLET; ORAL

CLINORIL

@ MERCK 150MG N17911 001 Jun DISC

SULINDAC

@ RADIUS PHARMS 150MG N73261 001 Sep 06, 1991 Jul CAHN

@ 200MG N73262 001 Sep 06, 1991 Jul CAHN

SUMATRIPTAN SUCCINATE

INJECTABLE; SUBCUTANEOUS

IMITREX

+ GLAXOSMITHKLINE EQ 6MG BASE/0.5ML (12MG/ML) N20080 001 Dec 28, 1992 Feb CDFR

IMITREX STATDOSE

+ GLAXOSMITHKLINE EQ 4MG BASE/0.5ML (8MG/ML) N20080 002 Feb 01, 2006 Feb NEWA

+ EQ 6MG BASE/0.5ML (12MG/ML) N20080 003 Dec 23, 1996 Feb NEWA

SUNITINIB MALATE

CAPSULE; ORAL

SUTENT

CPPI CV

EQ 12.5MG BASE N21938 001 Jan 26, 2006 Jul CAHN

EQ 25MG BASE N21938 002 Jan 26, 2006 Jul CAHN

+ EQ 50MG BASE N21938 003 Jan 26, 2006 Jul CAHN

PFIZER

EQ 12.5MG BASE N21938 001 Jan 26, 2006 Jun CPOT

12.5MG N21938 001 Jan 26, 2006 Jan NEWA

EQ 25MG BASE N21938 002 Jan 26, 2006 Jun CPOT

25MG N21938 002 Jan 26, 2006 Jan NEWA

+ EQ 50MG BASE N21938 003 Jan 26, 2006 Jun CPOT

+ 50MG N21938 003 Jan 26, 2006 Jan NEWA

TAMOXIFEN CITRATE

SOLUTION; ORAL

SOLTAMOX

+ SAVIENT PHARMA EQ 10MG BASE/5ML N21807 001 Oct 29, 2005 Jun CAHN

+ SAVIENT PHARMS EQ 10MG BASE/5ML N21807 001 Oct 29, 2005 Jul CAHN

TECHNETIUM TC-99M APCITIDE

INJECTABLE; INJECTION

ACUTECT

@ CIS BIO INTL SA N/A N20887 001 Sep 14, 1998 May DISC

N/A N20887 001 Sep 14, 1998 Apr CAHN

TECHNETIUM TC-99M DEPREOTIDE

INJECTABLE; INJECTION

NEO TECT KIT

@ CIS BIO INTL SA N/A N21012 001 Aug 03, 1999 May DISC

+ N/A N21012 001 Aug 03, 1999 Apr CAHN

TECHNETIUM TC-99M MEBROFENIN KIT

INJECTABLE; INJECTION

CHOLETEC

+ BRACCO N/A N18963 001 Jan 21, 1987 Aug CRLD

TECHNETIUM TC-99M TETROFOSMIN KIT

INJECTABLE; INJECTION

MYOVIEW 30ML

@ GE HEALTHCARE

N/A

N20372 002 Jul 07, 2005 May DISC

TEMAZEPAM

CAPSULE; ORAL

TEMAZEPAM

AB ACTAVIS ELIZABETH

15MG

N71638 001 Aug 07, 1987 Jun CAHN

AB 30MG

N71620 001 Aug 07, 1987 Jun CAHN

TERBUTALINE SULFATE

INJECTABLE; INJECTION

BRETHINE

@ AAIPHARMA LLC

1MG/ML

N18571 001 Aug DISC

TERBUTALINE SULFATE

AP + BEDFORD

1MG/ML

N76770 001 Apr 23, 2004 Aug CRLD

TERCONAZOLE

CREAM; VAGINAL

TERAZOL 3

&gt;D&gt; @ ORTHO MCNEIL PHARM

0.8%

N19964 001 Feb 21, 1991 Sep CMFD

&gt;A&gt; AB + 0.8%

N19964 001 Feb 21, 1991 Sep CMFD

@ 0.8%

N19964 001 Feb 21, 1991 Jul DISC

TERAZOL 7

&gt;D&gt; @ ORTHO MCNEIL PHARM

0.4%

N19579 001 Dec 31, 1987 Sep CMFD

&gt;A&gt; AB + 0.4%

N19579 001 Dec 31, 1987 Sep CMFD

@ 0.4%

N19579 001 Dec 31, 1987 Jul DISC

TERCONAZOLE

&gt;D&gt; AB + TARO

0.4%

N76043 001 Jan 19, 2005 Sep CRLD

&gt;A&gt; AB 0.4%

N76043 001 Jan 19, 2005 Sep CRLD

AB + 0.4%

N76043 001 Jan 19, 2005 Jul CRLD

&gt;D&gt; BX + 0.8%

N75953 001 Apr 06, 2004 Sep CRLD

&gt;A&gt; AB 0.8%

N75953 001 Apr 06, 2004 Sep CRLD

BX + 0.8%

N75953 001 Apr 06, 2004 Jul CRLD

SUPPOSITORY; VAGINAL

TERAZOL 3

AB + ORTHO MCNEIL PHARM

80MG

N19641 001 May 24, 1988 Mar CFTG

TERCONAZOLE

AB ALTANA

80MG

N76850 001 Jul 12, 2006 Jun NEWA

AB PERRIGO NEW YORK

80MG

N77149 001 Mar 17, 2006 Mar NEWA

TESTOSTERONE

GEL; TRANSDERMAL

ANDROGEL

AB + UNIMED PHARMS

1%

N21015 001 Feb 28, 2000 Jan CTEC

TESTOSTERONE

AB WATSON LABS

1%

N76737 001 Jan 27, 2006 Jan NEWA

TESTOSTERONE CYPIONATE

INJECTABLE; INJECTION

TESTOSTERONE CYPIONATE

AO SANDOZ

100MG/ML

N40615 001 Aug 10, 2006 Aug NEWA

AO 200MG/ML

N40615 002 Aug 10, 2006 Aug NEWA

TESTOSTERONE ENANTHATE

	INJECTABLE; INJECTION				
	DELATESTRYL				
	@ INDEVUS PHARMS	200MG/ML	N09165 001	Jan	CAHN
AO	+	200MG/ML	N09165 003	Jan	CAHN
	TESTOSTERONE ETHANATE				
AO	PADDOCK	200MG/ML	N40575 001	Jun 14, 2006	May NEWA

TETRACYCLINE HYDROCHLORIDE

	CAPSULE; ORAL				
	ACHROMYCIN V				
	@ RADIUS PHARMS	250MG	N50278 003	May	CAHN
	@	500MG	N50278 001	May	CAHN
	@ SCIREG INTL INC	250MG	N50278 003	Apr	CAHN
	@	500MG	N50278 001	Apr	CAHN
	SUSPENSION; ORAL				
	SUMYCIN				
	+ PAR PHARM	125MG/5ML	N60400 001	Aug	CAHN

THALLOUS CHLORIDE, TL-201

	INJECTABLE; INJECTION				
	THALLOUS CHLORIDE TL 201				
AP	TRACE RADIOCHEMICALS	1mCi/ML	N75569 001	Nov 21, 2001	Feb CAHN

THEOPHYLLINE

	ELIXIR; ORAL				
	THEOPHYLLINE				
AA	ACTAVIS MID ATLANTIC	80MG/15ML	N85863 001	Jul	CAHN
	SOLUTION; ORAL				
	THEOLAIR				
	@ 3M	80MG/15ML	N86107 001	Aug	DISC
	THEOPHYLLINE				
	+ ROXANE	80MG/15ML	N87449 001	Sep 15, 1983	Aug CRLD
	TABLET, EXTENDED RELEASE; ORAL				
	THEOPHYLLINE				
AB	NOSTRUM	400MG	N40595 001	Apr 21, 2006	Apr NEWA
AB		600MG	N40560 002	Apr 21, 2006	Apr NEWA
	T-PHYL				
	@ PURDUE FREDERICK	200MG	N88253 001	Aug 17, 1983	Jun DISC
	UNIPHYL				
AB	+ PURDUE FREDERICK	400MG	N87571 001	Sep 01, 1982	Apr CTEC
AB	+	600MG	N40086 001	Apr 15, 1996	Apr CTEC

THIABENDAZOLE

	SUSPENSION; ORAL				
	MINTEZOL				
	@ MERCK	500MG/5ML	N16097 001	Jun	DISC

THIORIDAZINE HYDROCHLORIDE

	CONCENTRATE; ORAL				
	THIORIDAZINE HYDROCHLORIDE				
AA	ACTAVIS MID ATLANTIC	100MG/ML	N88229 001	Aug 23, 1983	Jul CAHN

TICARCILLIN DISODIUM

INJECTABLE; INJECTION

TICAR

@	GLAXOSMITHKLINE	EQ 1GM BASE/VIAL	N50497 001		May	DISC
@		EQ 3GM BASE/VIAL	N50497 002		May	DISC
@		EQ 20GM BASE/VIAL	N50497 004		May	DISC
@		EQ 30GM BASE/VIAL	N50497 005	Apr 04, 1984	May	DISC

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	250MG	N75253 001	Aug 20, 1999	Jun	CAHN
----	-------------------	-------	------------	--------------	-----	------

TIMOLOL MALEATE

TABLET; ORAL

BLOCADREN

@	MERCK	5MG	N18017 001		Jun	DISC
@		10MG	N18017 002		Jun	DISC
@		20MG	N18017 004		Jun	DISC

TIMOLOL MALEATE

AB	+	MYLAN	20MG	N72668 001	Jun 08, 1990	Jun	CRLD
----	---	-------	------	------------	--------------	-----	------

TINIDAZOLE

TABLET; ORAL

TINDAMAX

	MISSION PHARMA	250MG	N21618 001	May 17, 2004	Jan	CAHN
+		500MG	N21618 002	May 17, 2004	Jan	CAHN

TIROFIBAN HYDROCHLORIDE

INJECTABLE; INJECTION

AGGRASTAT

+	MEDICURE	EQ 0.05MG BASE/ML	N20913 001	May 14, 1998	Aug	CAHN
+		EQ 0.25MG BASE/ML	N20912 001	May 14, 1998	Aug	CAHN

TIZANIDINE HYDROCHLORIDE

TABLET; ORAL

TIZANIDINE HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	EQ 2MG BASE	N76283 001	Jul 12, 2002	Jun	CAHN
AB		EQ 4MG BASE	N76283 002	Jul 12, 2002	Jun	CAHN

TOBRAMYCIN

SOLUTION; INHALATION

TOBI

+	NOVARTIS PHARMS	300MG/5ML	N50753 001	Dec 22, 1997	Jul	CAHN
---	-----------------	-----------	------------	--------------	-----	------

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

TOBRAMYCIN

AP	+	ABRAXIS PHARM	EQ 40MG BASE/ML	N65122 002	Nov 29, 2002	Jul	CRLD
----	---	---------------	-----------------	------------	--------------	-----	------

TOLMETIN SODIUM

CAPSULE; ORAL

TOLMETIN SODIUM

AB	ACTAVIS ELIZABETH	EQ 400MG BASE	N73308 001	Jan 24, 1992	Jun	CAHN
----	-------------------	---------------	------------	--------------	-----	------

## TABLET; ORAL

## TOLMETIN SODIUM

AB	ACTAVIS ELIZABETH	EQ 600MG BASE	N73527 001	Jun 30, 1992	Jun	CAHN
----	-------------------	---------------	------------	--------------	-----	------

TOPIRAMATE

## TABLET; ORAL

## TOPAMAX

AB	+	ORTHO MCNEIL PHARM	25MG	N20505 004	Dec 24, 1996	Aug	CFTG
AB			100MG	N20505 001	Dec 24, 1996	Aug	CFTG
AB			200MG	N20505 002	Dec 24, 1996	Aug	CFTG

## TOPIRAMATE

AB	MYLAN	25MG	N76314 001	Sep 11, 2006	Aug	NEWA
AB		100MG	N76314 002	Sep 11, 2006	Aug	NEWA
AB		200MG	N76314 003	Sep 11, 2006	Aug	NEWA

TRAMADOL HYDROCHLORIDE

## TABLET; ORAL

## TRAMADOL HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	50MG	N75960 001	Jun 19, 2002	Jun	CAHN
	@ IVAX PHARMS	50MG	N75963 001	Jul 03, 2002	Jan	DISC

TRANLYCYPROMINE SULFATE

## TABLET; ORAL

## PARNATE

AB	+	GLAXOSMITHKLINE	EQ 10MG BASE	N12342 003	Aug 16, 1985	Jun	CFTG
----	---	-----------------	--------------	------------	--------------	-----	------

## TRANLYCYPROMINE SULFATE

AB	KALI LABS	EQ 10MG BASE	N40640 001	Jun 29, 2006	Jun	NEWA
----	-----------	--------------	------------	--------------	-----	------

TRAVOPROST

## SOLUTION/DROPS; OPHTHALMIC

>A>	TRAVATAN Z						
>A>	+	ALCON	0.004%	N21994 001	Sep 21, 2006	Sep	NEWA

TRAZODONE HYDROCHLORIDE

## TABLET; ORAL

## DESYREL

## @ APOTHECON

## @

## @

## @

## TRAZODONE HYDROCHLORIDE

		50MG	N18207 001		Aug	DISC	
		100MG	N18207 002		Aug	DISC	
		150MG	N18207 003	Mar 25, 1985	Aug	DISC	
		300MG	N18207 004	Nov 07, 1988	Aug	DISC	
AB	ACTAVIS ELIZABETH	50MG	N71636 001	Apr 18, 1988	Jun	CAHN	
AB		100MG	N71514 001	Apr 18, 1988	Jun	CAHN	
>D>	AB	BARR	300MG	N71196 003	Apr 26, 1999	Sep	CRLD
>A>	AB	+	300MG	N71196 003	Apr 26, 1999	Sep	CRLD

TRIAMCINOLONE

## TABLET; ORAL

## ARISTOCORT

>D>	BP	+	ASTELLAS	4MG	N11161 007	Sep	DISC
>A>			@	4MG	N11161 007	Sep	DISC

## KENACORT

>D>	BP	BRISTOL MYERS SQUIBB	4MG	N11283 006	Sep	CRLD
>A>		+	4MG	N11283 006	Sep	CRLD

TRIAMCINOLONE ACETONIDE

## CREAM; TOPICAL

>D>		ARISTOCORT A						
>D>	AT	ASTELLAS	0.025%	N88818	001	Oct 16, 1984	Sep	DISC
>A>		@	0.025%	N88818	001	Oct 16, 1984	Sep	DISC
>D>	AT		0.1%	N88819	001	Oct 16, 1984	Sep	DISC
>A>		@	0.1%	N88819	001	Oct 16, 1984	Sep	DISC
>D>	AT		0.5%	N88820	001	Oct 16, 1984	Sep	DISC
>A>		@	0.5%	N88820	001	Oct 16, 1984	Sep	DISC

## TRIAMCINOLONE ACETONIDE

AT		ACTAVIS MID ATLANTIC	0.1%	N87798	001	Jun 04, 1982	Jun	CAHN
AT		VINTAGE	0.025%	N40671	001	Jun 09, 2006	May	NEWA
AT			0.1%	N40671	002	Jun 09, 2006	May	NEWA

## OINTMENT; TOPICAL

## TRIAMCINOLONE ACETONIDE

AT		ACTAVIS MID ATLANTIC	0.1%	N87799	001	Jun 07, 1982	Jun	CAHN
----	--	----------------------	------	--------	-----	--------------	-----	------

TRIAMCINOLONE DIACETATE

## INJECTABLE; INJECTION

## ARISTOCORT

@	SANDOZ	25MG/ML		N11685	003		Jan	CAHN
@		40MG/ML		N12802	001		Jan	CAHN

TRIAMCINOLONE HEXACETONIDE

## INJECTABLE; INJECTION

## ARISTOSPAN

+	SANDOZ	5MG/ML		N16466	001		Jan	CAHN
+		20MG/ML		N16466	002		Jan	CAHN

TRICHLORMETHIAZIDE

## TABLET; ORAL

## TRICHLORMETHIAZIDE

@	TG UNITED LABS	4MG		N85568	001		May	CAHN
---	----------------	-----	--	--------	-----	--	-----	------

TRIFLUOPERAZINE HYDROCHLORIDE

## INJECTABLE; INJECTION

## STELAZINE

@	GLAXOSMITHKLINE	EQ 2MG BASE/ML		N11552	005		May	DISC
---	-----------------	----------------	--	--------	-----	--	-----	------

TRIMIPRAMINE MALEATE

## CAPSULE; ORAL

## SURMONTIL

AB		ODYSSEY PHARMS	EQ 25MG BASE	N16792	001		Jul	CFTG
AB			EQ 50MG BASE	N16792	002		Jul	CFTG
AB	+		EQ 100MG BASE	N16792	003	Sep 15, 1982	Jul	CFTG

## TRIMIPRAMINE MALEATE

AB		ACTAVIS TOTOWA	EQ 25MG BASE	N77361	001	Aug 02, 2006	Jul	NEWA
AB			EQ 50MG BASE	N77361	002	Aug 02, 2006	Jul	NEWA
AB			EQ 100MG BASE	N77361	003	Aug 02, 2006	Jul	NEWA

TRIPLENNAMINE HYDROCHLORIDE

## TABLET; ORAL

## PBZ

@	NOVARTIS	50MG		N05914	002		Jan	DISC
---	----------	------	--	--------	-----	--	-----	------



TROLEANDOMYCIN

CAPSULE; ORAL

TAO

@ PFIZER

EQ 250MG BASE

N50336 002

Mar DISC

UNOPROSTONE ISOPROPYL

SOLUTION/DROPS; OPHTHALMIC

RESCULA

@ R TECH UENO LTD

0.15%

N21214 001 Aug 03, 2000 Jun DISC

+

0.15%

N21214 001 Aug 03, 2000 Feb CAHN

UREA

INJECTABLE; INJECTION

UREAPHIL

@ HOSPIRA

40GM/VIAL

N12154 001

Jun DISC

UROKINASE

INJECTABLE; INJECTION

ABBOKINASE

@ IMARX THERAP

5,000 IU/VIAL

N21846 003

Apr CAHN

@

9,000 IU/VIAL

N21846 002

Apr CAHN

+

250,000 IU/VIAL

N21846 001

Apr CAHN

URSODIOL

CAPSULE; ORAL

URSODIOL

AB

COREPHARMA

300MG

N77895 001 Jul 27, 2006 Jul NEWA

VALRUBICIN

SOLUTION; INTRAVESICAL

VALSTAR PRESERVATIVE FREE

+ VALERA

40MG/ML

N20892 001 Sep 25, 1998 Jul CAHN

VARENICLINE TARTRATE

TABLET; ORAL

CHANTIX

&gt;D&gt;

CPPI CV

EQ 0.5MG BASE

N21928 001 May 10, 2006 Sep CAHN

EQ 0.5MG BASE

N21928 001 May 10, 2006 Jul CAHN

&gt;D&gt;

+

EQ 1MG BASE

N21928 002 May 10, 2006 Sep CAHN

+

EQ 1MG BASE

N21928 002 May 10, 2006 Jul CAHN

PFIZER

EQ 0.5MG BASE

N21928 001 May 10, 2006 May NEWA

+

EQ 1MG BASE

N21928 002 May 10, 2006 May NEWA

&gt;A&gt;

PFIZER INC

EQ 0.5MG BASE

N21928 001 May 10, 2006 Sep CAHN

&gt;A&gt;

+

EQ 1MG BASE

N21928 002 May 10, 2006 Sep CAHN

VECURONIUM BROMIDE

INJECTABLE; INJECTION

VECURONIUM BROMIDE

AP

+

BEDFORD

20MG/VIAL

N75549 002 Jun 13, 2000 Jan CRLD

VENLAFAXINE HYDROCHLORIDE

TABLET; ORAL

EFFEXOR

AB

WYETH PHARMS INC

EQ 25MG BASE

N20151 002 Dec 28, 1993 Jul CFTG

## TABLET; ORAL

## EFFEXOR

AB	WYETH PHARMS INC	EQ 37.5MG BASE	N20151 006	Dec 28, 1993	Jul	CFTG
AB	+	EQ 50MG BASE	N20151 003	Dec 28, 1993	Jul	CFTG
AB		EQ 75MG BASE	N20151 004	Dec 28, 1993	Jul	CFTG
AB		EQ 100MG BASE	N20151 005	Dec 28, 1993	Jul	CFTG

## VENLAFAXINE HYDROCHLORIDE

AB	TEVA	EQ 25MG BASE	N76690 001	Aug 03, 2006	Jul	CTNA
AB		EQ 37.5MG BASE	N76690 002	Aug 03, 2006	Jul	NEWA
AB		EQ 50MG BASE	N76690 003	Aug 03, 2006	Jul	NEWA
AB		EQ 75MG BASE	N76690 004	Aug 03, 2006	Jul	CTNA
AB		EQ 100MG BASE	N76690 005	Aug 03, 2006	Jul	NEWA

VERAPAMIL HYDROCHLORIDE

## TABLET; ORAL

## VERAPAMIL HYDROCHLORIDE

AB	ACTAVIS ELIZABETH	80MG	N71019 001	Sep 24, 1986	Jun	CAHN
AB		120MG	N70468 001	Sep 24, 1986	Jun	CAHN
	@ RADIUS PHARMS	80MG	N71880 001	Apr 05, 1988	Jul	CAHN
	@	120MG	N71881 001	Apr 05, 1988	Jul	CAHN

WARFARIN SODIUM

## TABLET; ORAL

## WARFARIN SODIUM

AB	PLIVA	1MG	N40616 009	Jul 05, 2006	Jun	NEWA
AB		2MG	N40616 001	Jul 05, 2006	Jun	NEWA
AB		2.5MG	N40616 002	Jul 05, 2006	Jun	NEWA
AB		3MG	N40616 003	Jul 05, 2006	Jun	NEWA
AB		4MG	N40616 004	Jul 05, 2006	Jun	NEWA
AB		5MG	N40616 005	Jul 05, 2006	Jun	NEWA
AB		6MG	N40616 006	Jul 05, 2006	Jun	NEWA
AB		7.5MG	N40616 007	Jul 05, 2006	Jun	NEWA
AB		10MG	N40616 008	Jul 05, 2006	Jun	NEWA
AB	ZYDUS PHARMS USA	1MG	N40663 001	May 30, 2006	May	NEWA
AB		2MG	N40663 002	May 30, 2006	May	NEWA
AB		2.5MG	N40663 003	May 30, 2006	May	NEWA
AB		3MG	N40663 004	May 30, 2006	May	NEWA
AB		4MG	N40663 005	May 30, 2006	May	NEWA
AB		5MG	N40663 006	May 30, 2006	May	NEWA
AB		6MG	N40663 007	May 30, 2006	May	NEWA
AB		7.5MG	N40663 008	May 30, 2006	May	NEWA
AB		10MG	N40663 009	May 30, 2006	May	NEWA

WATER FOR INJECTION, STERILE

## LIQUID; N/A

## STERILE WATER FOR INJECTION IN PLASTIC CONTAINER

AP	TARO PHARMS IRELAND	100%	N77393 001	Aug 11, 2006	Aug	NEWA
----	---------------------	------	------------	--------------	-----	------

ZIDOVUDINE

## CAPSULE; ORAL

## RETROVIR

AB	+	GLAXOSMITHKLINE	100MG	N19655 001	Mar 19, 1987	Mar	CFTG
		ZIDOVUDINE					
AB		AUROBINDO PHARMA LTD	100MG	N78128 001	Mar 27, 2006	Mar	NEWA

ZIPRASIDONE HYDROCHLORIDESUSPENSION; ORAL  
GEODON

+ PFIZER INC EQ 10MG BASE/ML N21483 001 Mar 29, 2006 Mar NEWA

ZONISAMIDECAPSULE; ORAL  
ZONISAMIDE

AB	BANNER PHARMACAPS	25MG	N77813 001	Aug 16, 2006	Aug	NEWA
AB		50MG	N77813 002	Aug 16, 2006	Aug	NEWA
AB		100MG	N77813 003	Aug 16, 2006	Aug	NEWA
>A>	DR REDDYS LABS LTD	25MG	N77891 001	Sep 29, 2006	Sep	NEWA
>A>		50MG	N77891 002	Sep 29, 2006	Sep	NEWA
AB	GLENMARK PHARMS	25MG	N77651 001	Jan 30, 2006	Jan	NEWA
AB		50MG	N77651 002	Jan 30, 2006	Jan	NEWA
AB		100MG	N77651 003	Jan 30, 2006	Jan	NEWA
AB	INVAGEN PHARMS	25MG	N77869 001	May 31, 2006	May	NEWA
AB		50MG	N77869 002	May 31, 2006	May	NEWA
AB		100MG	N77869 003	May 31, 2006	May	NEWA
AB	SUN PHARM INDS (IN)	25MG	N77634 001	Mar 17, 2006	Mar	NEWA
AB		50MG	N77634 002	Mar 17, 2006	Mar	NEWA
AB		100MG	N77634 003	Mar 17, 2006	Mar	NEWA
AB	WATSON LABS	25MG	N77650 001	Apr 20, 2006	Apr	NEWA
AB		50MG	N77650 002	Apr 20, 2006	Apr	NEWA
AB		100MG	N77650 003	Apr 20, 2006	Apr	NEWA
AB	WOCKHARDT	25MG	N77636 003	Jul 27, 2006	Jul	NEWA
AB		50MG	N77636 002	Jul 27, 2006	Jul	NEWA

OTC DRUG PRODUCT LIST - 26TH EDITION

OTC DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 9 - September 2006

2-1

ACETAMINOPHEN

SUPPOSITORY; RECTAL

ACETAMINOPHEN

ACTAVIS MID ATLANTIC	120MG	N18337 003	Sep 12, 1983	Jun	CAHN
	325MG	N18337 002		Jun	CAHN
+	650MG	N18337 001		Jun	CAHN
INFANTS' FEVERALL					
ACTAVIS MID ATLANTIC	80MG	N18337 004	Aug 26, 1992	Jun	CAHN
TYLENOL					
@ MCNEIL CONS	120MG	N17756 002		Jun	CAHN
@	650MG	N17756 001		Jun	CAHN
TABLET, EXTENDED RELEASE; ORAL					
TYLENOL (CAPLET)					
+	MCNEIL CONS	650MG	N19872 001	Jun 08, 1994	Jun CAHN
TYLENOL (GELTAB)					
+	MCNEIL CONS	650MG	N19872 002	Jan 11, 2001	Jun CAHN

AVOBENZONE; ECAMSULE; OCTOCRYLENE

CREAM; TOPICAL

ANTHELIOS SX

+	LOREAL USA	2%;2%;10%	N21502 001	Jul 21, 2006	Jul NEWA
---	------------	-----------	------------	--------------	----------

CHLORHEXIDINE GLUCONATE

CLOTH; TOPICAL

HALO

+	SAGE PRODS	2%	N21669 001	Apr 25, 2005	Aug CTNA
---	------------	----	------------	--------------	----------

SPONGE; TOPICAL

CHLORHEXIDINE GLUCONATE

	BECTON DICKINSON	4%	N72525 001	Oct 24, 1989	Aug CAHN
	@ KENDALL IL	4%	N19490 001	Mar 27, 1987	Aug DISC

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SPONGE; TOPICAL

CHLORAPREP WITH TINT

+	MEDI FLEX INC	2%;70% (10.5ML)	N20832 005	Apr 03, 2006	Apr NEWA
---	---------------	-----------------	------------	--------------	----------

CLOTRIMAZOLE

CREAM; VAGINAL

CLOTRIMAZOLE

	ACTAVIS MID ATLANTIC	1%	N74165 001	Jul 16, 1993	Jun CAHN
--	----------------------	----	------------	--------------	----------

CROMOLYN SODIUM

SPRAY, METERED; NASAL

CROMOLYN SODIUM

	ACTAVIS MID ATLANTIC	5.2MG/SPRAY	N74800 001	Jul 26, 2001	Jul CAHN
	ALPHARMA US PHARMS	5.2MG/SPRAY	N74800 001	Jul 26, 2001	Jan CPOT
+	BAUSCH AND LOMB	5.2MG/SPRAY	N75702 001	Jul 03, 2001	Jan CRLD
NASALCROM					
	@ PHARMACIA UPJOHN	5.2MG/SPRAY	N20463 001	Jan 03, 1997	Jan DISC

DEXTROMETHORPHAN HYDROBROMIDE; GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

MUCINEX DM

	ADAMS RESP THERAP	30MG;600MG	N21620 002	Apr 29, 2004	May	CAHN
+		60MG;1.2GM	N21620 001	Apr 29, 2004	May	CAHN

DEXTROMETHORPHAN POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

DELSYM

+	ADAMS RESP THERAP	EQ 30MG HBR/5ML	N18658 001	Oct 08, 1982	Jun	CAHN
---	-------------------	-----------------	------------	--------------	-----	------

FAMOTIDINE

TABLET; ORAL

FAMOTIDINE

>A>	DR REDDYS LABS LTD	20MG	N77367 001	Sep 25, 2006	Sep	NEWA
>A>	PERRIGO	20MG	N77351 001	Sep 25, 2006	Sep	NEWA

GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

HUMABID

+	ADAMS RESP THERAP	1.2GM	N21282 002	Dec 18, 2002	Aug	CTNA
---	-------------------	-------	------------	--------------	-----	------

MUCINEX

	ADAMS RESP THERAP	600MG	N21282 001	Jul 12, 2002	May	CAHN
+		1.2GM	N21282 002	Dec 18, 2002	May	CAHN

GUAIFENESIN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

MUCINEX D

	ADAMS RESP THERAP	600MG;60MG	N21585 001	Jun 22, 2004	May	CAHN
+		1.2GM;120MG	N21585 002	Jun 22, 2004	May	CAHN

IBUPROFEN

SUSPENSION/DROPS; ORAL

CHILDREN'S MOTRIN

+	MCNEIL CONS	40MG/ML	N20603 001	Jun 10, 1996	Jun	CAHN
---	-------------	---------	------------	--------------	-----	------

SUSPENSION; ORAL

CHILDREN'S ELIXSURE

	ALTERNA-TCHP LLC	100MG/5ML	N21604 001	Jan 07, 2004	Jul	CAHN
--	------------------	-----------	------------	--------------	-----	------

IBUPROFEN

	ACTAVIS MID ATLANTIC	100MG/5ML	N74916 001	Apr 30, 1999	Jul	CAHN
--	----------------------	-----------	------------	--------------	-----	------

TABLET, CHEWABLE; ORAL

CHILDREN'S MOTRIN

	MCNEIL CONS	50MG	N20601 001	Nov 15, 1996	Jun	CAHN
--	-------------	------	------------	--------------	-----	------

JUNIOR STRENGTH MOTRIN

+	MCNEIL CONS	100MG	N20601 003	Nov 15, 1996	Jun	CAHN
---	-------------	-------	------------	--------------	-----	------

TABLET; ORAL

JUNIOR STRENGTH MOTRIN

	MCNEIL CONS	100MG	N20602 001	Jun 10, 1996	Jun	CAHN
--	-------------	-------	------------	--------------	-----	------

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

SUSPENSION; ORAL

CHILDREN'S MOTRIN COLD

+	MCNEIL CONS	100MG/5ML;15MG/5ML	N21128 001	Aug 01, 2000	Jun	CAHN
---	-------------	--------------------	------------	--------------	-----	------

TABLET; ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

DR REDDYS LABS LTD 200MG;30MG

N77628 001 Aug 14, 2006 Aug NEWA

SINE-AID IB

MCNEIL CONS 200MG;30MG

N19899 001 Dec 31, 1992 Jun CAHN

INSULIN PURIFIED PORK

INJECTABLE; INJECTION

REGULAR ILETIN II (PORK)

@ LILLY 100 UNITS/ML

N18344 001 Jun DISC

INSULIN SUSP ISOPHANE BEEF/PORK

INJECTABLE; INJECTION

NPH ILETIN I (BEEF-PORK)

@ LILLY 100 UNITS/ML

N17936 002 Jun DISC

INSULIN SUSP ISOPHANE PURIFIED PORK

INJECTABLE; INJECTION

NPH ILETIN II (PORK)

@ LILLY 100 UNITS/ML

N18345 001 Jun DISC

INSULIN ZINC SUSP EXTENDED RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN U

@ LILLY 100 UNITS/ML

N19571 002 Jun 10, 1987 Jun DISC

INSULIN ZINC SUSP PURIFIED PORK

INJECTABLE; INJECTION

LENTE ILETIN II (PORK)

@ LILLY 100 UNITS/ML

N18347 001 Jun DISC

>A> IODINE POVACRYLEX; ISOPROPYL ALCOHOL

&gt;A&gt; SPONGE; TOPICAL

&gt;A&gt; DURAPREP

&gt;A&gt; 3M EQ 0.7% IODINE;74% (26ML)

N21586 002 Sep 29, 2006 Sep NEWA

&gt;A&gt; + EQ 0.7% IODINE; 74% (6ML)

N21586 001 Sep 29, 2006 Sep NEWA

KETOCONAZOLE

SHAMPOO; TOPICAL

NIZORAL A-D

+ MCNEIL CONS 1%

N20310 001 Oct 10, 1997 Jun CAHN

KETOPROFEN

TABLET; ORAL

ACTRON

@ BAYER 12.5MG

N20499 001 Oct 06, 1995 Feb DISC

ORUDIS KT

@ WYETH CONS 12.5MG

N20429 001 Oct 06, 1995 Feb DISC

LEVONORGESTREL

TABLET; ORAL

PLAN B

+ DURAMED 0.75MG

N21045 002 Aug 24, 2006 Aug NEWA

LOPERAMIDE HYDROCHLORIDE

SOLUTION; ORAL

IMODIUM A-D

+ MCNEIL CONS 1MG/5ML N19487 001 Mar 01, 1988 Jun CAHN

SUSPENSION; ORAL

IMODIUM A-D

+ MCNEIL 1MG/7.5ML N19487 002 Jul 08, 2004 Mar CDFR

+ MCNEIL CONS 1MG/7.5ML N19487 002 Jul 08, 2004 Jun CAHN

TABLET; ORAL

IMODIUM A-D

+ MCNEIL CONS 2MG N19860 001 Nov 22, 1989 Jun CAHN

LOPERAMIDE HYDROCHLORIDE; SIMETHICONE

TABLET; ORAL

IMODIUM ADVANCED

+ MCNEIL CONS 2MG;125MG N21140 001 Nov 30, 2000 Jun CAHN

LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE

RANBAXY 2MG;125MG N77500 001 Sep 06, 2006 Aug NEWA

LORATADINE

SYRUP; ORAL

LORATADINE

SILARX 1MG/ML N77421 001 Jun 29, 2006 Jun NEWA

TABLET, CHEWABLE; ORAL

CHILDREN'S CLARITIN

+ SCHERING PLOUGH 5MG N21891 001 Aug 23, 2006 Aug NEWA

TABLET; ORAL

LORATADINE

APOTEX 10MG N76471 001 Feb 14, 2006 Jan NEWA

MICONAZOLE NITRATE

CREAM, SUPPOSITORY; TOPICAL, VAGINAL

M-ZOLE 3 COMBINATION PACK

ACTAVIS MID ATLANTIC 2%,200MG N74926 001 Apr 16, 1999 Jun CAHN

2%,200MG N74926 001 Apr 16, 1999 Jun CAHN

2%,200MG N74926 001 Apr 16, 1999 Jun CAHN

2%,200MG N74926 001 Apr 16, 1999 Jun CAHN

2%,200MG N74926 001 Apr 16, 1999 Jun CAHN

2%,200MG N74926 001 Apr 16, 1999 Jun CAHN

2%,200MG N74926 001 Apr 16, 1999 Jun CAHN

2%,200MG N74926 001 Apr 16, 1999 Jun CAHN

M-ZOLE 7 DUAL PACK

ACTAVIS MID ATLANTIC 2%,100MG N74586 001 Jul 17, 1997 Jun CAHN

CREAM; VAGINAL

MICONAZOLE 7

ACTAVIS MID ATLANTIC 2% N74164 001 Mar 29, 1996 Jun CAHN

SUPPOSITORY; VAGINAL

MICONAZOLE NITRATE

ACTAVIS MID ATLANTIC 100MG N73507 001 Nov 19, 1993 Jun CAHN

MINOXIDIL

AEROSOL, FOAM; TOPICAL

MEN'S ROGAINE

+ PHARMACIA AND UPJOHN 5% N21812 001 Jan 20, 2006 Jan NEWA

## SOLUTION; TOPICAL

MINOXIDIL (FOR MEN)

ACTAVIS MID ATLANTIC 2%

N74588 001 Apr 05, 1996 Jul CAHN

MINOXIDIL EXTRA STRENGTH (FOR MEN)

ACTAVIS MID ATLANTIC 5%

N75518 001 Nov 17, 2000 Jul CAHN

NAPROXEN SODIUM

CAPSULE; ORAL

NAPROXEN SODIUM

>D>	@ BANNER PHARMACAPS	EQ 200MG BASE	N21920 001	Feb 17, 2006	Sep	CMFD
>D>	@	EQ 200MG BASE	N21920 001	Feb 17, 2006	Sep	CMFD
>A>		EQ 200MG BASE	N21920 001	Feb 17, 2006	Sep	CMFD
	@	EQ 200MG BASE	N21920 001	Feb 17, 2006	Jul	DISC
	+	EQ 200MG BASE	N21920 001	Feb 17, 2006	Feb	NEWA

NAPROXEN SODIUM; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

NAPROXEN SODIUM AND PSEUDOEPHEDRINE HYDROCHLORIDE

>A>	DR REDDYS LABS INC	EQ 220MG BASE;120MG	N77381 001	Sep 27, 2006	Sep	NEWA
-----	--------------------	---------------------	------------	--------------	-----	------

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICORETTE

GLAXOSMITHKLINE

EQ 2MG BASE

N18612 003 Dec 23, 1998 Mar CRLD

EQ 2MG BASE

N18612 004 Sep 25, 2000 Mar CRLD

EQ 4MG BASE

N20066 003 Dec 23, 1998 Mar CRLD

EQ 4MG BASE

N20066 004 Sep 25, 2000 Mar CRLD

NICORETTE (MINT)

GLAXOSMITHKLINE

EQ 2MG BASE

N18612 003 Dec 23, 1998 Apr CTNA

EQ 4MG BASE

N20066 003 Dec 23, 1998 Apr CTNA

NICOTINE POLACRILEX

PERRIGO

EQ 2MG BASE

N76776 001 Sep 16, 2004 Apr CTNA

EQ 2MG BASE

N76777 001 Sep 16, 2004 Apr CTNA

EQ 4MG BASE

N76778 001 Sep 16, 2004 Apr CTNA

EQ 4MG BASE

N76779 001 Sep 16, 2004 Apr CTNA

WATSON LABS

EQ 2MG BASE

N76569 001 Jul 29, 2004 Apr CTNA

EQ 4MG BASE

N76568 002 Jul 29, 2004 Apr CTNA

TROCHE/LOZENGE; ORAL

NICOTINE POLACRILEX

PERRIGO R AND D

EQ 2MG BASE

N77007 001 Jan 31, 2006 Jan NEWA

EQ 4MG BASE

N77007 002 Jan 31, 2006 Jan NEWA

PERMETHRIN

LOTION; TOPICAL

PERMETHRIN

ACTAVIS MID ATLANTIC 1%

N75014 001 Mar 28, 2000 Jun CAHN

POVIDONE-IODINE

SOLUTION; TOPICAL

E-Z PREP

@ CLINIPAD

10%

N19382 001 Jul 25, 1989 Aug DISC

SPONGE; TOPICAL

E-Z PREP

@ CLINIPAD

5%

N19382 002 Jul 25, 1989 Aug DISC

E-Z PREP 220

@ CLINIPAD

5%

N19382 003 Jul 25, 1989 Aug DISC



RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE

WOCKHARDT

EQ 75MG BASE

N76760 001 Feb 24, 2006 Feb NEWA

TERBINAFINE

GEL; TOPICAL

LAMISIL AT

+ NOVARTIS

1%

N21958 001 Jul 24, 2006 Jul NEWA

**DRUG PRODUCTS WITH APPROVAL UNDER SECTION 505 OF THE ACT  
ADMINISTERED BY THE CENTER FOR BIOLOGICS EVALUATION AND RESEARCH LIST**

**CUMULATIVE SUPPLEMENT NUMBER 09 SEPTEMBER 2006**

NO SEPTEMBER 2006 APPROVALS

## 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**

NO SEPTEMBER 2006 ADDITIONS

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ABACAVIR SULFATE; LAMIVUDINE - EPZICOM</u>					
021652 001	5034394	Dec 18, 2011	DS DP	D-40	Aug 02, 2007
	5034394*PED	Jun 18, 2012			
	5047407	Nov 17, 2009	DS DP	U-257	
	5047407*PED	May 17, 2010			
	5089500	Jun 26, 2009		U-257	
	5089500*PED	Dec 26, 2009			
	5905082	May 18, 2016	DS DP		
	5905082*PED	Nov 18, 2016			
	6294540	May 14, 2018	DS DP	U-257	
	6294540*PED	Nov 14, 2018			
<u>ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE - TRIZIVIR</u>					
021205 001	>A> 5034394	Dec 18, 2011	DS DP		
	>A> 5034394*PED	Jun 18, 2012			
<u>ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE - ULTRACET</u>					
021123 001	RE39221	Aug 09, 2011	DS DP	U-55	
<u>ALBUTEROL SULFATE - PROAIR HFA</u>					
021457 001				I-235	Feb 03, 2009
<u>ALBUTEROL SULFATE - VENTOLIN HFA</u>					
020983 001	6131566	Apr 14, 2015	DP	U-716	
	6131566	Apr 14, 2015	DP	U-589	
	6532955	Apr 14, 2015	DP	U-716	
	6532955	Apr 14, 2015	DP	U-590	
<u>ALENDRONATE SODIUM; CHOLECALCIFEROL - FOSAMAX PLUS D</u>					
021762 001				NC	Apr 07, 2008
<u>ALFUZOSIN HYDROCHLORIDE - UROXATRAL</u>					
021287 001	4661491	May 27, 2007		U-706	
<u>ALITRETINOIN - PANRETIN</u>					
020886 001	7056954	Aug 02, 2012	DP		
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u>					
020364 006	4879303	Mar 25, 2007	DS DP	NS	Apr 11, 2009
	6162802	Dec 19, 2017	DS DP	U-185	
<u>AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL</u>					
020364 007	4879303	Mar 25, 2007	DS DP	NS	Apr 11, 2009
	6162802	Dec 19, 2017	DS DP	U-185	
<u>ANIDULAFUNGIN - ERAXIS</u>					
021632 001	5965525	Oct 12, 2016	DS DP	U-540	NCE
	6384013	Mar 19, 2012	DS		
	6743777	Mar 19, 2012	DP	U-540	
	6960564	Apr 12, 2021	DP	U-540	
<u>APREPITANT - EMEND</u>					
021549 001	5145684	Jan 25, 2011	DP		
	5719147	Jun 29, 2012	DS DP		
	6096742	Jul 01, 2018	DS DP	U-745	
	6235735	Jun 29, 2012		U-746	
	6235735	Jun 29, 2012		U-747	
<u>APREPITANT - EMEND</u>					
021549 002	5145684	Jan 25, 2011	DP		
	5719147	Jun 29, 2012	DS DP		
	6096742	Jul 01, 2018	DS DP	U-745	
	6235735	Jun 29, 2012		U-747	
	6235735	Jun 29, 2012		U-746	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>APREPITANT - EMEND</u>					
021549 003	5145684	Jan 25, 2011	DP	I-498	Jun 30, 2009
	5719147	Jun 29, 2012	DS DP	NS	Jun 30, 2009
	6096742	Jul 01, 2018	DS DP U-745	NCE	Mar 26, 2008
	6235735	Jun 29, 2012	U-747		
	6235735	Jun 29, 2012	U-746		
<u>ARIPIPRAZOLE - ABILIFY</u>					
021436 001				I-488	Mar 01, 2008
<u>ARIPIPRAZOLE - ABILIFY</u>					
021436 002				I-488	Mar 01, 2008
<u>ARIPIPRAZOLE - ABILIFY</u>					
021436 003				I-488	Mar 01, 2008
<u>ARIPIPRAZOLE - ABILIFY</u>					
021436 004				I-488	Mar 01, 2008
<u>ARIPIPRAZOLE - ABILIFY</u>					
021436 005				I-488	Mar 01, 2008
<u>ARIPIPRAZOLE - ABILIFY</u>					
021436 006				I-488	Mar 01, 2008
<u>ARIPIPRAZOLE - ABILIFY</u>					
021713 001	6977257	Apr 24, 2022	DS DP	I-488	Mar 01, 2008
<u>ARIPIPRAZOLE - ABILIFY</u>					
021729 002				>A> I-488 >A> I-437 >A> NCE	Mar 01, 2008 Sep 29, 2007 Nov 15, 2007
<u>ARIPIPRAZOLE - ABILIFY</u>					
021729 003				>A> I-488 >A> I-437 >A> NCE	Mar 01, 2008 Sep 29, 2007 Nov 15, 2007
<u>ARIPIPRAZOLE - ABILIFY</u>					
021729 004				>A> I-488 >A> I-437 >A> NCE	Mar 01, 2008 Sep 29, 2007 Nov 15, 2007
<u>ARIPIPRAZOLE - ABILIFY</u>					
021729 005				>A> I-488 >A> I-437 >A> NCE	Mar 01, 2008 Sep 29, 2007 Nov 15, 2007
<u>ARIPIPRAZOLE - ABILIFY</u>					
021866 001				>A> NDF >A> NCE	Sep 20, 2009 Nov 15, 2007
<u>ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE - SEPTOCAINE</u>					
022010 001				NP	Mar 30, 2009
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u>					
021881 001				NP	Aug 02, 2009
<u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u>					
021411 001	5658590	Nov 26, 2016		U-494	
	5658590*PED	May 26, 2017		U-494	
<u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u>					
021411 002	5658590	Nov 26, 2016		U-494	
	5658590*PED	May 26, 2017		U-494	
<u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u>					
021411 003	5658590	Nov 26, 2016		U-494	
	5658590*PED	May 26, 2017		U-494	
<u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u>					
021411 004	5658590	Nov 26, 2016		U-494	
	5658590*PED	May 26, 2017		U-494	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u>					
021411 005	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
<u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u>					
021411 006	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
<u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u>					
021411 007	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
<u>ATOMOXETINE HYDROCHLORIDE - STRATTERA</u>					
021411 008	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
<u>AVOBENZONE; ECAMSULE; OCTOCRYLENE - ANTHELIOS SX</u>					
021502 001	4585597	Jun 16, 2007	DS DP	U-752	NC Jul 21, 2009
	5587150	Dec 24, 2013	DP	U-752	
<u>AZELASTINE HYDROCHLORIDE - ASTELIN</u>					
020114 001				D-102	Feb 17, 2009
<u>BALSALAZIDE DISODIUM - COLAZAL</u>					
020610 001	4412992*PED	Jan 08, 2007			
<u>BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX</u>					
021852 001	4866048	Dec 29, 2007	DS DP	U-88	NC Jan 09, 2009
	4866048	Dec 29, 2007	DS DP	U-193	
	5763426	Jun 09, 2015	DS DP		
	6753013	Jan 27, 2020	DP	U-88	
	6753013	Jan 27, 2020	DP	U-193	
<u>BETAMETHASONE VALERATE - LUXIQ</u>					
020934 001	7078058	Mar 01, 2016	DS DP		
<u>BETAXOLOL HYDROCHLORIDE - BETOPTIC S</u>					
019845 001	4911920	Mar 27, 2007			
	4911920*PED	Sep 27, 2007			
<u>BIMATOPROST - LUMIGAN</u>					
021275 001	5688819	Aug 19, 2014		U-446	
	6403649	Sep 21, 2012	DS DP	U-446	
<u>BIVALIRUDIN - ANGIOMAX</u>					
020873 001				I-486	Nov 30, 2008
<u>BORTEZOMIB - VELCADE</u>					
021602 001				ODE	Mar 25, 2012
<u>BRIMONIDINE TARTRATE - ALPHAGAN P</u>					
021770 001	5424078	Jun 13, 2012	DP		
	5424078*PED	Dec 13, 2012			
	6562873	Jul 10, 2021	DP		
	6562873*PED	Jan 10, 2022			
	6627210	Jul 18, 2021	DP		
	6627210*PED	Jan 18, 2022			
	6641834	Jul 28, 2021	DP		
	6641834*PED	Jan 28, 2022			
	6673337	Jul 26, 2021	DP		
	6673337*PED	Jan 26, 2022			
<u>BRINZOLAMIDE - AZOPT</u>					
020816 001	5240923	Aug 31, 2010		U-224	
	5240923*PED	Mar 01, 2011			
	5378703	Apr 01, 2012		U-224	
	5378703*PED	Oct 01, 2012			
	5461081	Oct 24, 2012		U-225	
	5461081*PED	Apr 24, 2013			

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>BROMFENAC SODIUM - XIBROM</u>					
021664 001				I-485	Jan 27, 2009
<u>BUDESONIDE - BUDESONIDE</u>					
021949 001				NP	Jul 12, 2009
<u>BUDESONIDE - BUDESONIDE</u>					
021949 002				NP	Jul 12, 2009
<u>BUDESONIDE - PULMICORT RESPULES</u>					
020929 001	6598603*PED	Jun 23, 2019			
	6899099	Dec 23, 2018	U-751		
	6899099*PED	Jun 23, 2019			
<u>BUDESONIDE - PULMICORT RESPULES</u>					
020929 002	6899099	Dec 23, 2018	U-751		
	6899099*PED	Jun 23, 2019			
<u>BUDESONIDE - RHINOCORT</u>					
020746 001	6986904	Apr 29, 2017	DP U-699		
<u>BUDESONIDE - RHINOCORT</u>					
020746 002	6986904	Apr 29, 2017	DP U-699		
<u>BUDESONIDE; FORMOTEROL FUMARATE - SYMBICORT</u>					
021929 001	5674860	Oct 07, 2014	DP U-754	NC	Jul 21, 2009
	5972919	Dec 17, 2012	DP U-754		
	6123924	Sep 26, 2017	DP		
	6641800	Sep 26, 2017	DP		
<u>BUDESONIDE; FORMOTEROL FUMARATE - SYMBICORT</u>					
021929 002	5674860	Oct 07, 2014	DP U-754	NC	Jul 21, 2009
	5972919	Dec 17, 2012	DP U-754		
	6123924	Sep 26, 2017	DP		
	6641800	Sep 26, 2017	DP		
<u>BUPROPION HYDROCHLORIDE - WELLBUTRIN XL</u>					
021515 001				I-497	Jun 12, 2009
<u>BUPROPION HYDROCHLORIDE - WELLBUTRIN XL</u>					
021515 002				I-497	Jun 12, 2009
<u>CALCIUM CARBONATE; RISEDRONATE SODIUM - ACTONEL WITH CALCIUM (COPACKAGED)</u>					
021823 001				M-52	Jan 24, 2009
<u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 100</u>					
021485 002	4963590	Nov 27, 2007	DP U-219		
	5112861	May 12, 2009	U-219		
	5135950	Oct 31, 2010	DS DP U-219		
	6500867	Jun 29, 2020	DP U-219		
<u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 150</u>					
021485 003	4963590	Nov 27, 2007	DP U-219		
	5112861	May 12, 2009	U-219		
	5135950	Oct 31, 2010	DS DP U-219		
	6500867	Jun 29, 2020	DP U-219		
<u>CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 50</u>					
021485 001	4963590	Nov 27, 2007	DP U-219		
	5112861	May 12, 2009	U-219		
	5135950	Oct 31, 2010	DS DP U-219		
	6500867	Jun 29, 2020	DP U-219		
<u>CARVEDILOL - COREG</u>					
020297 001				M-56	Aug 28, 2009
<u>CARVEDILOL - COREG</u>					
020297 002				M-56	Aug 28, 2009
<u>CARVEDILOL - COREG</u>					
020297 003				M-56	Aug 28, 2009
<u>CARVEDILOL - COREG</u>					
020297 004				M-56	Aug 28, 2009



**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CEFDITOREN PIVOXIL - SPECTRACEF</u>					
021222 001	4839350	Apr 14, 2009	DS DP		
<u>CELECOXIB - CELEBREX</u>					
020998 001	5466823	Nov 30, 2013	DS	I-466	Jul 29, 2008
	5466823*PED	May 30, 2014		PED	Jan 29, 2009
	5563165	Nov 30, 2013	DP		
	5563165*PED	May 30, 2014			
	5760068	Jun 02, 2015		U-672	
	5760068	Jun 02, 2015		U-299	
	5760068*PED	Dec 02, 2015			
	5972986	Oct 14, 2017		U-299	
	5972986*PED	Apr 14, 2018			
<u>CELECOXIB - CELEBREX</u>					
020998 002	5466823	Nov 30, 2013	DS	I-466	Jul 29, 2008
	5466823*PED	May 30, 2014		PED	Jan 29, 2009
	5563165	Nov 30, 2013	DP		
	5563165*PED	May 30, 2014			
	5760068	Jun 02, 2015		U-672	
	5760068	Jun 02, 2015		U-299	
	5760068*PED	Dec 02, 2015			
	5972986	Oct 14, 2017		U-299	
	5972986*PED	Apr 14, 2018			
<u>CELECOXIB - CELEBREX</u>					
020998 003	5466823	Nov 30, 2013	DS	I-466	Jul 29, 2008
	5466823*PED	May 30, 2014		PED	Jan 29, 2009
	5563165	Nov 30, 2013	DP		
	5563165*PED	May 30, 2014			
	5760068	Jun 02, 2015		U-672	
	5760068	Jun 02, 2015		U-299	
	5760068*PED	Dec 02, 2015			
	5972986	Oct 14, 2017		U-299	
	5972986*PED	Apr 14, 2018			
<u>CETIRIZINE HYDROCHLORIDE - ZYRTEC</u>					
019835 001	4525358	Jun 25, 2007	DS DP	U-565	
	4525358*PED	Dec 25, 2007			
<u>CETIRIZINE HYDROCHLORIDE - ZYRTEC</u>					
019835 002	4525358	Jun 25, 2007	DS DP	U-565	
	4525358*PED	Dec 25, 2007			
<u>CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ZYRTEC-D 12 HOUR</u>					
021150 001	7014867	Jun 10, 2022		DP	
<u>CHLORHEXIDINE GLUCONATE - HALO</u>					
021669 001	7066916	Feb 17, 2024		U-737	
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>					
020832 004	5690958	Sep 30, 2016		DP	
	6536975	Nov 10, 2020		DP	
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP FREPP</u>					
020832 003	5538353	Aug 25, 2015		DP	
	5690958	Sep 30, 2016		DP	
	5752363	Apr 22, 2017		DP	
	5772346	Apr 22, 2017		DP	
	D386849	Nov 25, 2011		DP	
	D396911	Aug 11, 2012		DP	
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP SINGLE SWABSTICK</u>					
021555 002	5690958	Sep 30, 2016		DP	
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>					
020832 002	6991393	Mar 14, 2023		DP	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>					
020832 005	5690958	Sep 30, 2016	DP		
	6536975	Nov 10, 2020	DP		
	6729786	Mar 14, 2023	DP		
	6991393	Jan 31, 2024	DP		
<u>CICLOPIROX - LOPROX</u>					
020519 001	7018656	Sep 05, 2018	DP		
	7026337	Apr 02, 2018		U-714	
<u>CIPROFLOXACIN - CIPRO</u>					
019847 002				>A> I-421	Mar 25, 2007
				>A> PED	Sep 25, 2007
<u>CIPROFLOXACIN - CIPRO</u>					
019847 003				>A> I-421	Mar 25, 2007
				>A> PED	Sep 25, 2007
<u>CLOBETASOL PROPIONATE - CLOBEX</u>					
021835 001	5972920	Feb 12, 2018	DP	NDF	Oct 27, 2008
	5990100	Mar 24, 2018	DP	U-742	
<u>CLOPIDOGREL BISULFATE - CLOPIDOGREL BISULFATE</u>					
076274 001				>A> PC	Feb 04, 2007
<u>CLOPIDOGREL BISULFATE - PLAVIX</u>					
020839 001				I-502	Aug 17, 2009
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>					
021176 001	>A> 5693675	Dec 02, 2014	DS		
	>A> 5917007	Apr 29, 2014	DS	U-323	
	>A> 5919832	Jun 10, 2014	DS		
	>A> 6066678	Jun 10, 2014	DS	U-323	
	>A> 6433026	Jun 10, 2014	DS		
	>A> 6784254	Jun 10, 2014	DS	DP	
	>A> 7101960	Apr 29, 2014	DS	DP	U-757
<u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL</u>					
021697 001	5723606	Mar 03, 2015	DS	DP	U-698
<u>DAPSONE - ACZONE</u>					
021794 001	5863560	Sep 11, 2016	DP		
	6620435	Sep 11, 2016	DP		
<u>DAPTOMYCIN - CUBICIN</u>					
021572 001	6468967	Sep 24, 2019		U-282	
	RE39071	Jun 15, 2016	DS	DP	U-728
<u>DAPTOMYCIN - CUBICIN</u>					
021572 002	6468967	Sep 24, 2019		U-282	
	RE39071	Jun 15, 2016	DS	DP	U-728
<u>DARUNAVIR ETHANOLATE - PREZISTA</u>					
021976 001	5843946	Dec 01, 2015	DP	U-744	NCE Jun 23, 2011
	6248775	Aug 25, 2012	DS		
	6335460	Aug 25, 2012	DS	DP	U-744
<u>DASATINIB - SPRYCEL</u>					
021986 001	6596746	Apr 13, 2020	DS	DP	U-748 NCE Jun 28, 2011
<u>DASATINIB - SPRYCEL</u>					
021986 002	6596746	Apr 13, 2020	DS	DP	U-748 NCE Jun 28, 2011
<u>DASATINIB - SPRYCEL</u>					
021986 003	6596746	Apr 13, 2020	DS	DP	U-748 NCE Jun 28, 2011
<u>DECITABINE - DACOGEN</u>					
021790 001				NCE	May 02, 2011
<u>DEFERASIROX - EXJADE</u>					
021882 001	6465504	Jun 24, 2017	DS	DP	
	6596750	Jun 24, 2017	DS		U-735

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DEFERASIROX - EXJADE</u>					
021882 002	6465504	Jun 24, 2017	DS DP		
	6596750	Jun 24, 2017	DS	U-735	
<u>DEFERASIROX - EXJADE</u>					
021882 003	6465504	Jun 24, 2017	DS DP		
	6596750	Jun 24, 2017	DS	U-735	
<u>DESFLURANE - SUPRANE</u>					
020118 001	>A> 4762856	Feb 02, 2007		U-67	
	>A> 4762856*PED	Aug 02, 2007			
	>A> 5617906	Apr 08, 2014	DP		
	>A> 5617906*PED	Oct 08, 2014			
<u>DES Loratadine - CLARINEX</u>					
021165 001	4659716	Mar 31, 2007	DP	U-725	
	4659716	Mar 31, 2007	DP	U-427	
	4659716*PED	Oct 01, 2007		U-427	
<u>DES Loratadine - CLARINEX</u>					
021300 001	4659716	Mar 31, 2007	DP	U-725	
	4659716	Mar 31, 2007	DP	U-611	
	4659716*PED	Oct 01, 2007			
<u>DES Loratadine - CLARINEX</u>					
021312 001	4659716	Mar 31, 2007	DP	U-725	
	4659716	Mar 31, 2007	DP	U-427	
	4659716*PED	Oct 01, 2007		U-427	
	5178878	Jan 12, 2010	DP		
	5607697	Jun 07, 2015	DP		
<u>DES Loratadine - CLARINEX</u>					
021312 002	4659716	Mar 31, 2007	DP	U-725	
	4659716	Mar 31, 2007	DP	U-427	
	4659716*PED	Oct 01, 2007	DP		
	5178878	Jan 12, 2010	DP		
	5607697	Jun 07, 2015	DP		
<u>DES Loratadine; Pseudoephedrine Sulfate - CLARINEX D 24 HOUR</u>					
021605 001	4659716	Mar 31, 2007	DP	U-726	
	4659716	Mar 31, 2007	DP	U-644	
	4659716*PED	Oct 01, 2007			
	6979463	Mar 28, 2022	DP		
<u>DES Loratadine; Pseudoephedrine Sulfate - CLARINEX-D 12 HOUR</u>					
021313 001	4659716	Mar 31, 2007	DP	U-707	>A> NP
	4659716*PED	Oct 01, 2007			NCE
	6100274	Jul 07, 2019	DP		NC
	6100274*PED	Jan 07, 2020			PED
	6709676	Feb 18, 2021	DP	U-707	
<u>DES MOPRESSIN ACETATE - DDAVP</u>					
019955 001	7022340	Apr 30, 2023	DP		
<u>DES MOPRESSIN ACETATE - DDAVP</u>					
019955 002	7022340	Apr 30, 2023	DP		
<u>DESONIDE - VERDESO</u>					
021978 001				>A> NDF	Sep 19, 2009
<u>DEXMEDETOMIDINE - PRECEDEX</u>					
021038 001	>A> 4910214	Jul 15, 2013	DS DP	U-421	
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u>					
021802 004				>A> NDF	May 26, 2008
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>					
021392 001	>A> 7108866	Dec 17, 2019	DP	U-107	
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>					
021392 002	>A> 7108866	Dec 17, 2019	DP	U-107	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>					
021392 003	>A> 7108866	Dec 17, 2019	DP U-107		
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>					
021392 004	>A> 7108866	Dec 17, 2019	DP U-107		
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>					
021392 005	>A> 7108866	Dec 17, 2019	DP U-107		
<u>DILTIAZEM HYDROCHLORIDE - CARDIZEM LA</u>					
021392 006	>A> 7108866	Dec 17, 2019	DP U-107		
<u>DOCETAXEL - TAXOTERE</u>					
020449 001	5750561	Jul 03, 2012	DP	I-490	Mar 22, 2009
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT ODT</u>					
021720 001	4895841	Nov 25, 2010	DS DP U-713		
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT ODT</u>					
021720 002	4895841	Nov 25, 2010	DS DP U-713		
<u>DROSPIRENONE; ETHINYL ESTRADIOL - YAZ</u>					
021676 001	5569652	Oct 29, 2013		U-1 NP	Mar 16, 2009
	5798338	Jul 10, 2015	DP		
	6787531	Aug 31, 2020	DP		
	6933395	Aug 11, 2017	DP		
	6958326	Dec 20, 2021	DP		
	RE37564	Jun 30, 2014	DP		
	RE37838	Jun 30, 2014	DP		
	RE38253	Jun 30, 2014	DP		
<u>EFAVIRENZ - SUSTIVA</u>					
020972 001	5519021	May 21, 2013	DS DP		
	5663169	Sep 02, 2014		U-257	
	5811423	Aug 07, 2012	DS DP	U-256	
	6238695	Apr 06, 2019	DP		
<u>EFAVIRENZ - SUSTIVA</u>					
020972 002	5519021	May 21, 2013	DS DP		
	5663169	Sep 02, 2014		U-257	
	5811423	Aug 07, 2012	DS DP	U-256	
	6238695	Apr 06, 2019	DP		
<u>EFAVIRENZ - SUSTIVA</u>					
020972 003	5519021	May 21, 2013	DS DP		
	5663169	Sep 02, 2014		U-257	
	5811423	Aug 07, 2012	DS DP	U-256	
	6238695	Apr 06, 2019	DP		
<u>EFAVIRENZ - SUSTIVA</u>					
021360 002	5519021	May 21, 2013	DS DP		

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u>						
021937 001	5210085	May 11, 2010			U-750	
	5210085*PED	Nov 11, 2010				
	5519021	May 21, 2013	DS	DP		
	5663169	Sep 02, 2014			U-750	
	5811423	Aug 07, 2012			U-750	
	5814639	Sep 29, 2015	DS	DP		
	5814639*PED	Mar 29, 2016				
	5914331	Sep 29, 2015	DS			
	5914331*PED	Mar 29, 2016				
	5922695	Jul 25, 2017	DS		U-750	
	5935946	Jul 25, 2017	DS	DP	U-750	
	5977089	Jul 25, 2017	DS	DP	U-750	
	6043230	Jul 25, 2017			U-750	
	6238695	Apr 06, 2019		DP		
	6555133	Apr 06, 2019			U-750	
	6639071	Nov 13, 2021	DS			
	6642245	Nov 04, 2020			U-750	
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS	DP		
	6703396*PED	Sep 09, 2021				
	6939964	Jan 20, 2018	DS			
<u>EMTRICITABINE - EMTRIVA</u>						
021500 001	5210085	May 11, 2010			NCE	Jul 02, 2008
	5210085*PED	Nov 11, 2010			PED	Jan 02, 2009
	5814639	Sep 29, 2015				
	5814639*PED	Mar 29, 2016				
	5914331	Sep 29, 2015				
	5914331*PED	Mar 29, 2016				
	6642245	Nov 04, 2020			U-541	
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS	DP		
	6703396*PED	Sep 09, 2021				
<u>EMTRICITABINE - EMTRIVA</u>						
021896 001	5210085	May 11, 2010			U-257	
	5210085*PED	Nov 11, 2010			NDF	Sep 27, 2008
	5814639	Sep 29, 2015	DS	DP	NCE	Jul 02, 2008
	5814639*PED	Mar 29, 2016			PED	Mar 27, 2009
	5914331	Sep 29, 2015	DS		PED	Jan 02, 2009
	5914331*PED	Mar 29, 2016				
	6642245	Nov 04, 2020			U-257	
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS	DP		
	6703396*PED	Sep 09, 2021				
<u>EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA</u>						
021752 001	5210085	May 11, 2010			U-248	
	5210085	May 11, 2010			U-541	
	5210085*PED	Nov 11, 2010			NCE	Jul 02, 2008
	5814639	Sep 29, 2015	DS	DP	PED	Jan 02, 2009
	5814639*PED	Mar 29, 2016				
	5914331	Sep 29, 2015	DS	DP	U-248	
	5914331*PED	Mar 29, 2016				
	6642245	Nov 04, 2020			U-248	
	6642245	Nov 04, 2020			U-541	
	6642245*PED	May 04, 2021				
	6703396	Mar 09, 2021	DS	DP		
	6703396*PED	Sep 09, 2021				

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ENTACAPONE - COMTAN</u>					
020796 001	4963590	Nov 27, 2007	DP U-219		
	5112861	May 12, 2009	U-219		
	5135950	Oct 31, 2010	DS DP U-219		
	6599530	Sep 14, 2018	DP U-219		
<u>EPLERENONE - INSPRA</u>					
021437 001	4559332	Apr 09, 2007	DS DP U-537		
<u>EPLERENONE - INSPRA</u>					
021437 002	4559332	Apr 09, 2007	DS DP U-537		
<u>EPLERENONE - INSPRA</u>					
021437 003	4559332	Apr 09, 2007	DS DP U-537		
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>					
021153 001	4738974	Apr 19, 2007	DS DP U-635	I-484	Nov 24, 2007
	4738974	Apr 19, 2007	DS DP U-373	NPP	Apr 28, 2009
	4738974*PED	Oct 19, 2007	U-373		
	4786505	Apr 20, 2007	DP U-373		
	4786505	Apr 20, 2007	DP U-729		
	4853230	Apr 20, 2007	DP U-729		
	4853230	Apr 20, 2007	DP U-373		
	5690960	Nov 25, 2014	DP U-729		
	5690960	Nov 25, 2014	DP U-373		
	5714504	Feb 03, 2015	DP U-729		
	5714504	Feb 03, 2015	DP U-373		
	5877192	May 27, 2014	DP U-373		
	5877192	May 27, 2014	DP U-729		
	5900424	May 04, 2016	DS U-729		
	5900424	May 04, 2016	DS U-373		
	6369085	May 25, 2018	DS DP U-729		
	6428810	Nov 03, 2019	DP U-469		
	6428810	Nov 03, 2019	DP U-729		
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>					
021153 002	4738974	Apr 19, 2007	DS DP U-635	I-484	Nov 24, 2007
	4738974	Apr 19, 2007	DS DP U-373	NPP	Apr 28, 2009
	4738974*PED	Oct 19, 2007	U-373		
	4786505	Apr 20, 2007	DP U-373		
	4786505	Apr 20, 2007	DP U-729		
	4853230	Apr 20, 2007	DP U-729		
	4853230	Apr 20, 2007	DP U-373		
	5690960	Nov 25, 2014	DP U-729		
	5690960	Nov 25, 2014	DP U-373		
	5714504	Feb 03, 2015	DP U-729		
	5714504	Feb 03, 2015	DP U-373		
	5877192	May 27, 2014	DP U-373		
	5877192	May 27, 2014	DP U-729		
	5900424	May 04, 2016	DS U-729		
	5900424	May 04, 2016	DS U-373		
	6369085	May 25, 2018	DS DP U-729		
	6428810	Nov 03, 2019	DP U-469		
	6428810	Nov 03, 2019	DP U-729		
<u>ESTRADIOL - ESTRADIOL</u>					
075182 004				PC	Feb 06, 2007
<u>ESTRADIOL - ESTRADIOL</u>					
075182 005				PC	Feb 06, 2007
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u>					
021840 001				NP	May 25, 2009
<u>ETHINYL ESTRADIOL; NORELGESTROMIN - ORTHO EVRA</u>					
021180 001	5876746	Nov 20, 2015	DP U-514		

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ETHINYL ESTRADIOL; NORETHINDRONE - OVCON-35</u>					
021490 001	6667050	Apr 06, 2019	DP U-1		
<u>ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LOESTRIN 24 FE</u>					
021871 001	5552394	Jul 22, 2014	U-1	NP	Feb 17, 2009
<u>ETONOGESTREL - IMPLANON</u>					
021529 001	4957119	Aug 05, 2008	DP	NDF	Jul 17, 2009
	5150718	Sep 29, 2009	U-749		
<u>EXEMESTANE - AROMASIN</u>					
020753 001				I-495	Oct 05, 2008
<u>EZETIMIBE - ZETIA</u>					
021445 001	7030106	Jan 25, 2022	DP	I-493	May 23, 2009
	>A> RE37721	Oct 25, 2016	DS DP	U-473	
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>					
021687 001	>A> RE37721	Oct 25, 2016	DS DP	U-473	
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>					
021687 002	>A> RE37721	Oct 25, 2016	DS DP	U-473	
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>					
021687 003	>A> RE37721	Oct 25, 2016	DS DP	U-473	
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>					
021687 004	>A> RE37721	Oct 25, 2016	DS DP	U-473	
<u>FAMCICLOVIR - FAMVIR</u>					
020363 001				D-103 I-501	Jul 28, 2009 Jul 28, 2009
<u>FAMCICLOVIR - FAMVIR</u>					
020363 002				D-103 I-501	Jul 28, 2009 Jul 28, 2009
<u>FAMCICLOVIR - FAMVIR</u>					
020363 003				D-103 I-501	Jul 28, 2009 Jul 28, 2009
<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>					
021695 001	7101574	Aug 20, 2020	DS DP	M-47	Oct 21, 2008
<u>FENOFIBRATE - ANTARA (MICRONIZED)</u>					
021695 003	7101574	Aug 20, 2020	DS DP	M-47	Oct 21, 2008
<u>FENOFIBRATE - FENOFIBRATE</u>					
076433 001				PC	May 22, 2006
<u>FENOFIBRATE - FENOFIBRATE</u>					
076433 002				PC	May 22, 2006
<u>FENOFIBRATE - LIPOFEN</u>					
021612 001	5545628	Jan 10, 2015		U-701	
<u>FENOFIBRATE - LIPOFEN</u>					
021612 002	5545628	Jan 10, 2015		U-701	
<u>FENOFIBRATE - LIPOFEN</u>					
021612 003	5545628	Jan 10, 2015		U-701	
<u>FENOFIBRATE - TRICOR</u>					
021656 001	7037529	Jan 09, 2018	DP		
	7041319	Jan 09, 2018	DP		
<u>FENOFIBRATE - TRICOR</u>					
021656 002	7037529	Jan 09, 2018	DP		
	7041319	Jan 09, 2018	DP		
<u>FENTANYL CITRATE - FENTORA</u>					
021947 001				>A> NDF	Sep 25, 2009
<u>FENTANYL CITRATE - FENTORA</u>					
021947 002				>A> NDF	Sep 25, 2009
<u>FENTANYL CITRATE - FENTORA</u>					
021947 003				>A> NDF	Sep 25, 2009

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FENTANYL CITRATE - FENTORA</u>					
021947 004				>A> NDF	Sep 25, 2009
<u>FENTANYL CITRATE - FENTORA</u>					
021947 005				>A> NDF	Sep 25, 2009
<u>FENTANYL HYDROCHLORIDE - IONSYS</u>					
021338 001	5232438	Oct 03, 2008	DP U-736	NDF	May 22, 2009
	5445606	Dec 11, 2011	DP		
	5697896	Dec 16, 2014	DP		
	5843014	Dec 01, 2015	DP		
	6169920	Jan 02, 2018	DP		
	6171294	Jun 05, 2015		U-736	
	6181963	Nov 02, 2019	DP		
	6195582	Jan 28, 2019	DP	U-736	
	6216033	Jun 05, 2015	DP		
	6317629	Jun 02, 2012	DP		
	6425892	Jun 05, 2015		U-736	
	6842640	Jun 02, 2015	DP		
	6881208	Apr 19, 2022		U-736	
	6975902	Apr 01, 2024	DP		
	7018370	Jun 05, 2015		U-736	
	7027859	Sep 26, 2014	DP		
<u>FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA D 24 HOUR</u>					
021704 001	RE39069	May 29, 2018	DP		
<u>FINASTERIDE - FINASTERIDE</u>					
076340 001				PC	Dec 16, 2006
<u>FLUNISOLIDE - AEROSPAN HFA</u>					
021247 001				NP	Jan 27, 2009
<u>FLUOCINOLONE ACETONIDE - RETISERT</u>					
021737 001	6217895	Mar 22, 2019	DP U-708		
	6548078	Mar 22, 2019	DP U-708		
<u>FLUOCINONIDE - VANOS</u>					
021758 001				I-487	Mar 02, 2009
<u>FLUOXETINE HYDROCHLORIDE - PROZAC WEEKLY</u>					
021235 001	RE39030	May 29, 2017	DP U-397		
	RE39030	May 29, 2017	DP U-396		
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>					
021433 001	5658549	Sep 19, 2014	DP U-710	NPP	Feb 28, 2009
	5674472	Oct 07, 2014	DP		
	6251368	Dec 04, 2012	DP		
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>					
021433 002	5658549	Sep 19, 2014	DP U-710	NPP	Feb 28, 2009
	5674472	Oct 07, 2014	DP		
	6251368	Dec 04, 2012	DP		
<u>FLUTICASONE PROPIONATE - FLOVENT HFA</u>					
021433 003	5658549	Sep 19, 2014	DP U-710	NPP	Feb 28, 2009
	5674472	Oct 07, 2014	DP		
	6251368	Dec 04, 2012	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 100/50</u>					
021077 001	4992474	Feb 12, 2008		U-211	
	4992474*PED	Aug 12, 2008		U-211	
	5126375	Feb 12, 2008			
	5126375*PED	Aug 12, 2008			
	5225445	Feb 12, 2008		U-211	
	5225445*PED	Aug 12, 2008		U-211	
	6536427	Mar 01, 2011	DP		
	6536427*PED	Sep 01, 2011			



**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 250/50</u>					
021077 002	4992474	Feb 12, 2008		U-211	
	4992474*PED	Aug 12, 2008		U-211	
	5126375	Feb 12, 2008			
	5126375*PED	Aug 12, 2008			
	5225445	Feb 12, 2008		U-211	
	5225445*PED	Aug 12, 2008		U-211	
	6536427	Mar 01, 2011	DP		
	6536427*PED	Sep 01, 2011			
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 500/50</u>					
021077 003	4992474	Feb 12, 2008		U-211	
	4992474*PED	Aug 12, 2008		U-211	
	5126375	Feb 12, 2008			
	5126375*PED	Aug 12, 2008			
	5225445	Feb 12, 2008		U-211	
	5225445*PED	Aug 12, 2008		U-211	
	6536427	Mar 01, 2011	DP		
	6536427*PED	Sep 01, 2011			
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>					
021254 001	4992474	Feb 12, 2008	DS DP	U-738	NP Jun 08, 2009
	4992474*PED	Aug 12, 2008			
	5126375	Feb 12, 2008	DS DP		
	5126375*PED	Aug 12, 2008			
	5225445	Feb 12, 2008		U-738	
	5225445*PED	Aug 12, 2008			
	5270305	Sep 07, 2010		U-738	
	5658549	Aug 19, 2014	DP	U-738	
	5674472	Oct 07, 2014	DP		
	6143277	Apr 14, 2015	DP	U-738	
	6251368	Dec 04, 2012	DP		
	6253762	Apr 14, 2015	DP	U-738	
	6315173	Dec 23, 2017	DP		
	6510969	Dec 23, 2017	DP		
	6524555	Apr 14, 2015	DP		
	6546928	Apr 14, 2015	DP		
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>					
021254 002	4992474	Feb 12, 2008	DS DP	U-738	NP Jun 08, 2009
	4992474*PED	Aug 12, 2008			
	5126375	Feb 12, 2008	DS DP		
	5126375*PED	Aug 12, 2008			
	5225445	Feb 12, 2008		U-738	
	5225445*PED	Aug 12, 2008			
	5270305	Sep 07, 2010		U-738	
	5658549	Aug 19, 2014	DP	U-738	
	5674472	Oct 07, 2014	DP		
	6143277	Apr 14, 2015	DP	U-738	
	6251368	Dec 04, 2012	DP		
	6253762	Apr 14, 2015	DP	U-738	
	6315173	Dec 23, 2017	DP		
	6510969	Dec 23, 2017	DP		
	6524555	Apr 14, 2015	DP		
	6546928	Apr 14, 2015	DP		

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA</u>					
021254 003	4992474	Feb 12, 2008	DS DP	U-738	NP
	4992474*PED	Aug 12, 2008			
	5126375	Feb 12, 2008	DS DP		
	5126375*PED	Aug 12, 2008			
	5225445	Feb 12, 2008		U-738	
	5225445*PED	Aug 12, 2008			
	5270305	Sep 07, 2010		U-738	
	5658549	Aug 19, 2014	DP	U-738	
	5674472	Oct 07, 2014	DP		
	6143277	Apr 14, 2015	DP	U-738	
	6251368	Dec 04, 2012	DP		
	6253762	Apr 14, 2015	DP	U-738	
	6315173	Dec 23, 2017	DP		
	6510969	Dec 23, 2017	DP		
	6524555	Apr 14, 2015	DP		
	6546928	Apr 14, 2015	DP		
<u>FROVATRIPTAN SUCCINATE - FROVA</u>					
021006 001	5464864	Nov 07, 2015		U-436	
<u>FULVESTRANT - FASLODEX</u>					
021344 001	>A> 4659516	Dec 11, 2007	DS DP	U-596	
<u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u>					
021615 001				NDF	Apr 01, 2008
<u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u>					
021615 002				NDF	Apr 01, 2008
<u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u>					
021615 003				NDF	Apr 01, 2008
<u>GANIRELIX ACETATE - GANIRELIX ACETATE INJECTION</u>					
021057 001	4801577	Feb 05, 2012	DS DP		
<u>GEMCITABINE HYDROCHLORIDE - GEMZAR</u>					
020509 001				I-499	Jul 14, 2009
<u>GEMCITABINE HYDROCHLORIDE - GEMZAR</u>					
020509 002				I-499	Jul 14, 2009
<u>GEMIFLOXACIN MESYLATE - FACTIVE</u>					
021158 001	5776944	Apr 04, 2017	DS DP		
<u>GLIMEPIRIDE - AMARYL</u>					
020496 001				M-54 PED	Nov 28, 2008 May 28, 2009
<u>GLIMEPIRIDE - AMARYL</u>					
020496 002				M-54 PED	Nov 28, 2008 May 28, 2009
<u>GLIMEPIRIDE - AMARYL</u>					
020496 003				M-54 PED	Nov 28, 2008 May 28, 2009
<u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u>					
021925 001	4687777	Jan 17, 2011	DS		
	6150383	Jun 19, 2016		U-753	
	6211205	Jun 19, 2016		U-753	
	6303640	Aug 09, 2016		U-753	
	6329404	Jun 19, 2016	DP	U-753	
<u>GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT</u>					
021925 002	4687777	Jan 17, 2011	DS		
	6150383	Jun 19, 2016		U-753	
	6211205	Jun 19, 2016		U-753	
	6303640	Aug 09, 2016		U-753	
	6329404	Jun 19, 2016	DP	U-753	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>					
021700 001	5002953	Sep 17, 2011	DS DP	U-690	
	5002953*PED	Mar 17, 2012			
	5741803	Apr 21, 2015	DS DP	U-690	
	5741803*PED	Oct 21, 2015			
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>					
021700 002	5002953	Sep 17, 2011	DS DP	U-690	
	5002953*PED	Mar 17, 2012			
	5741803	Apr 21, 2015	DS DP	U-690	
	5741803*PED	Oct 21, 2015			
<u>GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL</u>					
021700 003	5002953	Sep 17, 2011	DS DP	U-690	
	5002953*PED	Mar 17, 2012			
	5741803	Apr 21, 2015	DS DP	U-690	
	5741803*PED	Oct 21, 2015			
<u>HYALURONIDASE RECOMBINANT HUMAN - HYLENEX RECOMBINANT</u>					
021859 001				NCE	Dec 02, 2010
<u>HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL</u>					
021956 001				NC	Aug 28, 2009
<u>HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL</u>					
021956 002				NC	Aug 28, 2009
<u>HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL</u>					
021956 003				NC	Aug 28, 2009
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>					
020818 004				NS	Apr 28, 2009
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>					
020818 005				NS	Apr 28, 2009
<u>IBANDRONATE SODIUM - BONIVA</u>					
021455 001	4927814	Jul 09, 2007	DS DP	U-642	
	6143326	Apr 21, 2017		U-642	
<u>IBANDRONATE SODIUM - BONIVA</u>					
021858 001	4927814	Jul 09, 2007	DS DP	U-700	NDF Jan 06, 2009
	5662918	Sep 02, 2014	DP		NCE May 16, 2008
<u>IBUPROFEN LYSINE - NEOPROFEN</u>					
021903 001				NE	Apr 13, 2009
				ODE	Apr 13, 2013
<u>IMATINIB MESYLATE - GLEEVEC</u>					
021335 001	5521184	Jan 04, 2015	DS DP	I-392	May 20, 2006
	5521184*PED	Jul 04, 2015		I-376	Dec 20, 2005
	6894051	May 23, 2019	DS DP	U-649	NCE May 10, 2006
	6894051*PED	Nov 23, 2019			ODE Feb 01, 2009
	6958335	Dec 19, 2021	DS DP		ODE May 10, 2008
	6958335*PED	Jun 19, 2022			PED Aug 01, 2009
					PED Nov 10, 2008
					PED Nov 20, 2006
					PED Nov 10, 2006
					PED Jun 20, 2006
<u>IMATINIB MESYLATE - GLEEVEC</u>					
021335 002	5521184	Jan 04, 2015		I-392	May 20, 2006
	5521184*PED	Jul 04, 2015		I-376	Dec 20, 2005
	6894051	May 23, 2019	DS DP	U-649	NCE May 10, 2006
	6894051*PED	Nov 23, 2019			ODE Feb 01, 2009
	6958335	Dec 19, 2021	DS DP		ODE May 10, 2008
	6958335*PED	Jun 19, 2022			PED Aug 01, 2009
					PED Nov 10, 2008
					PED Nov 20, 2006
					PED Nov 10, 2006
					PED Nov 10, 2006
					PED Jun 20, 2006

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IMATINIB MESYLATE - GLEEVEC</u>					
021588 001	5521184	Jan 04, 2015	DS DP	I-392	May 20, 2006
	5521184*PED	Jul 04, 2015		I-376	Dec 20, 2005
	6894051	May 23, 2019	DS DP U-649	NCE	May 10, 2006
	6894051*PED	Nov 23, 2019		ODE	Feb 01, 2009
	6958335	Dec 19, 2021	DS DP	ODE	May 10, 2008
	6958335*PED	Jun 19, 2022		PED	Aug 01, 2009
				PED	Nov 10, 2008
				PED	Nov 20, 2006
				PED	Nov 10, 2006
				PED	Jun 20, 2006
<u>IMATINIB MESYLATE - GLEEVEC</u>					
021588 002	5521184	Jan 04, 2015		I-392	May 20, 2006
	5521184*PED	Jul 04, 2015		I-376	Dec 20, 2005
	6894051	May 23, 2019	DS DP U-649	NCE	May 10, 2006
	6894051*PED	Nov 23, 2019		ODE	Feb 01, 2009
	6958335	Dec 19, 2021	DS DP	ODE	May 10, 2008
	6958335*PED	Jun 19, 2022		PED	Aug 01, 2009
				PED	Nov 10, 2008
				PED	Nov 20, 2006
				PED	Nov 10, 2006
				PED	Jun 20, 2006
<u>INSULIN DETEMIR RECOMBINANT - LEVEMIR</u>					
021536 001				I-489	Oct 19, 2008
<u>INSULIN RECOMBINANT HUMAN - EXUBERA</u>					
021868 001	5740794	Apr 21, 2015	DP	NP	Jan 27, 2009
	5997848	Mar 07, 2014		U-704	
	6051256	Mar 07, 2014	DP		
	6257233	May 14, 2019		U-704	
	6423344	Mar 07, 2014	DP		
	6543448	Sep 21, 2014	DP		
	6546929	May 14, 2019		U-704	
	6582728	Jun 24, 2020	DP		
	6592904	Mar 07, 2014	DP		
	6685967	Sep 11, 2018	DP		
	6737045	Mar 07, 2014		U-704	
	RE37872	Feb 12, 2010	DP		
	RE38385	Feb 12, 2010	DP		
<u>INSULIN RECOMBINANT HUMAN - EXUBERA</u>					
021868 002	5740794	Apr 21, 2015	DP	NP	Jan 27, 2009
	5997848	Mar 07, 2014		U-704	
	6051256	Mar 07, 2014	DP		
	6257233	May 14, 2019		U-704	
	6423344	Mar 07, 2014	DP		
	6543448	Sep 21, 2014	DP		
	6546929	May 14, 2019		U-704	
	6582728	Jun 24, 2020	DP		
	6592904	Mar 07, 2014	DP		
	6685967	Sep 11, 2018	DP		
	6737045	Mar 07, 2014		U-704	
	RE37872	Feb 12, 2010	DP		
	RE38385	Feb 12, 2010	DP		
<u>IPRATROPIUM BROMIDE - ATROVENT HFA</u>					
021527 001	6983743	May 26, 2020	DP		
<u>LAMOTRIGINE - LAMOTRIGINE</u>					
076420 001				>A> PC	Feb 25, 2007
<u>LAMOTRIGINE - LAMOTRIGINE</u>					
076420 002				PC	Dec 25, 2006
<u>LANSOPRAZOLE - PREVACID</u>					
020406 001	6749864	Feb 13, 2007	DP		

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes 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 002	6749864	Feb 13, 2007	DP		
<u>LANSOPRAZOLE - PREVACID</u>					
021281 001	6749864	Feb 13, 2007	DP		
<u>LANSOPRAZOLE - PREVACID</u>					
021281 002	6749864	Feb 13, 2007	DP		
<u>LANSOPRAZOLE - PREVACID</u>					
021428 001	6749864	Feb 13, 2007	DP		
<u>LANSOPRAZOLE - PREVACID</u>					
021428 002	6749864	Feb 13, 2007	DP		
<u>LANTHANUM CARBONATE - FOSRENOL</u>					
021468 003	5968976	Mar 19, 2016	DP	U-613	
<u>LANTHANUM CARBONATE - FOSRENOL</u>					
021468 004	5968976	Mar 19, 2016	DP	U-613	
<u>LENALIDOMIDE - REVLIMID</u>					
021880 001	5635517	Jul 24, 2016	DS		Dec 27, 2012
	6045501	Aug 28, 2018		U-694	
	6315720	Oct 23, 2020		U-694	
	6555554	Jul 24, 2016	DP		
	6561976	Aug 28, 2018		U-694	
	6561977	Oct 23, 2020		U-694	
	6755784	Oct 23, 2020		U-694	
	6908432	Aug 28, 2018		U-694	
<u>LENALIDOMIDE - REVLIMID</u>					
021880 002	5635517	Jul 24, 2016	DS		Dec 27, 2012
	6045501	Aug 28, 2018		U-694	
	6315720	Oct 23, 2020		U-694	
	6555554	Jul 24, 2016	DP		
	6561976	Aug 28, 2018		U-694	
	6561977	Oct 23, 2020		U-694	
	6755784	Oct 23, 2020		U-694	
	6908432	Aug 28, 2018		U-694	
<u>LENALIDOMIDE - REVLIMID</u>					
021880 003	5635517	Jul 24, 2016	DS		Jun 29, 2009
	6045501	Aug 28, 2018		U-694	Dec 27, 2010
	6315720	Oct 23, 2020		U-694	
	6555554	Jul 24, 2016	DP		
	6561976	Aug 28, 2018		U-694	
	6561977	Oct 23, 2020		U-694	
	6755784	Oct 23, 2020		U-694	
	6908432	Aug 28, 2018		U-694	
<u>LENALIDOMIDE - REVLIMID</u>					
021880 004	5635517	Jul 24, 2016	DS		Jun 29, 2009
	6045501	Aug 28, 2018		U-694	Dec 27, 2010
	6315720	Oct 23, 2020		U-694	
	6555554	Jul 24, 2016	DP		
	6561976	Aug 28, 2018		U-694	
	6561977	Oct 23, 2020		U-694	
	6755784	Oct 23, 2020		U-694	
	6908432	Aug 28, 2018		U-694	
<u>LEVALBUTEROL TARTRATE - XOPENEX HFA</u>					
021730 001	>A> 5362755	Mar 25, 2013		U-636	
<u>LEVETIRACETAM - KEPPRA</u>					
021035 004	4943639	Jul 14, 2008	DS	NPP	Jun 21, 2008
<u>LEVETIRACETAM - KEPPRA</u>					
021872 001	4943639	Jul 14, 2008	DS	NDF	Jul 31, 2009

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOBETAXOLOL HYDROCHLORIDE - BETAXON</u>					
021114 001	4911920	Mar 27, 2007		U-369	
	4911920*PED	Sep 27, 2007			
	5540918	Jul 30, 2013			
	5540918*PED	Jan 30, 2014			
<u>LEVONORGESTREL - PLAN B</u>					
021045 002				NP	Aug 24, 2009
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 001	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 002	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 003	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 004	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 005	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 006	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 007	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 008	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 009	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 010	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 011	7067148	Feb 15, 2022	DP		
<u>LEVOTHYROXINE SODIUM - LEVOXYL</u>					
021301 012	7067148	Feb 15, 2022	DP		
<u>LIDOCAINE; TETRACAINE - LIDOCAINE AND TETRACAINE</u>					
021717 001	5919479	Jul 28, 2015	DP	NP	Jun 29, 2009
	6528086	Sep 28, 2019	DP		
<u>LIDOCAINE; TETRACAINE - SYNERA</u>					
021623 001				NC	Jun 23, 2008
<u>LOPINAVIR; RITONAVIR - KALETRA</u>					
021906 001	5541206	Jul 30, 2013	DS DP	U-688	
	5541206*PED	Jan 30, 2014			
	5635523	Jun 03, 2014		U-688	
	5635523*PED	Dec 03, 2014			
	5648497	Jul 15, 2014	DS DP		
	5648497*PED	Jan 15, 2015			
	5674882	Oct 07, 2014		U-688	
	5674882*PED	Apr 07, 2015			
	5846987	Dec 29, 2012		U-688	
	5846987*PED	Jun 29, 2013			
	5886036	Dec 29, 2012	DP		
	5886036*PED	Jun 29, 2013			
	6037157	Jun 26, 2016		U-688	
	6037157*PED	Dec 26, 2016			
	6703403	Jun 26, 2016		U-688	
	6703403*PED	Dec 26, 2016			
<u>LOVASTATIN; NIACIN - ADVICOR</u>					
021249 001	7011848	Sep 20, 2013		U-712	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LOVASTATIN; NIACIN - ADVICOR</u>					
021249 002	7011848	Sep 20, 2013		U-712	
<u>LOVASTATIN; NIACIN - ADVICOR</u>					
021249 003	7011848	Sep 20, 2013		U-712	
<u>LUBIPROSTONE - AMITIZA</u>					
021908 001	5284858	Feb 08, 2011	DS DP	NCE	Jan 31, 2011
	5317032	May 31, 2011	DS DP	U-717	
	6414016	Sep 05, 2020	DS DP	U-717	
	6583174	Oct 16, 2020	DS DP		
	7064148	Aug 30, 2022	DS DP	U-739	
<u>MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>					
021850 001	6489346	Jul 16, 2016	DS DP	U-623	
	6489346	Jul 16, 2016	DS DP	U-588	
	6645988	Jul 16, 2016	DS DP	U-623	
	6645988	Jul 16, 2016	DS DP	U-588	
	6699885	Jul 16, 2016		U-623	
	6699885	Jul 16, 2016		U-588	
<u>MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>					
021850 002	6489346	Jul 16, 2016	DS DP	U-623	
	6489346	Jul 16, 2016	DS DP	U-588	
	6645988	Jul 16, 2016	DS DP	U-623	
	6645988	Jul 16, 2016	DS DP	U-588	
	6699885	Jul 16, 2016		U-623	
	6699885	Jul 16, 2016		U-588	
<u>MECASERMIN RINFABATE RECOMBINANT - IPLEX</u>					
021884 001	5200509	Apr 06, 2010	DS		
	5681818	Oct 28, 2014		U-697	
<u>MEGESTROL ACETATE - MEGACE ES</u>					
021778 001	7101576	Apr 22, 2024		U-755	
<u>MELOXICAM - MOBIC</u>					
020938 001				ODE PED	Aug 11, 2012 Feb 11, 2013
<u>MELOXICAM - MOBIC</u>					
020938 002				ODE PED	Aug 11, 2012 Feb 11, 2013
<u>MELOXICAM - MOBIC</u>					
021530 001				I-469 ODE PED PED	Aug 11, 2008 Aug 11, 2012 Feb 11, 2013 Feb 11, 2009
<u>MEQUINOL; TRETINOIN - SOLAGE</u>					
020922 001	5194247	Dec 10, 2013	DP	U-294	
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 001	5002953	Sep 17, 2011	DS DP	U-690	I-494
	5002953	Sep 17, 2011	DS DP	U-734	May 19, 2009
	5002953	Sep 17, 2011	DS DP	U-691	
	5002953	Sep 17, 2011	DS DP	U-493	
	5741803	Apr 21, 2015	DS DP	U-734	
	5741803	Apr 21, 2015	DS DP	U-493	
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 002	5002953	Sep 17, 2011	DS DP	U-690	I-494
	5002953	Sep 17, 2011	DS DP	U-734	May 19, 2009
	5002953	Sep 17, 2011	DS DP	U-691	
	5002953	Sep 17, 2011	DS DP	U-493	
	5741803	Apr 21, 2015	DS DP	U-734	
	5741803	Apr 21, 2015	DS DP	U-493	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes 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 003	5002953	Sep 17, 2011	DS DP	U-691	I-494 May 19, 2009
	5002953	Sep 17, 2011	DS DP	U-734	
	5002953	Sep 17, 2011	DS DP	U-690	
	5002953	Sep 17, 2011	DS DP	U-493	
	5741803	Apr 21, 2015	DS DP	U-734	
	5741803	Apr 21, 2015	DS DP	U-493	
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 004	5002953	Sep 17, 2011	DS DP	U-691	I-494 May 19, 2009
	5002953	Sep 17, 2011	DS DP	U-734	
	5002953	Sep 17, 2011	DS DP	U-690	
	5002953	Sep 17, 2011	DS DP	U-493	
	5741803	Apr 21, 2015	DS DP	U-734	
	5741803	Apr 21, 2015	DS DP	U-493	
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 005	5002953	Sep 17, 2011	DS DP	U-690	I-494 May 19, 2009
	5002953	Sep 17, 2011	DS DP	U-734	
	5002953	Sep 17, 2011	DS DP	U-691	
	5002953	Sep 17, 2011	DS DP	U-493	
	5741803	Apr 21, 2015	DS DP	U-734	
	5741803	Apr 21, 2015	DS DP	U-493	
	5741803*PED	Oct 21, 2015			
<u>METHYLPHENIDATE - DAYTRANA</u>					
021514 001	5958446	Dec 12, 2012	DP		NDF Apr 06, 2009
	6210705	Sep 30, 2018	DP	U-727	
	6348211	Sep 30, 2018	DP	U-727	
<u>METHYLPHENIDATE - DAYTRANA</u>					
021514 002	5958446	Dec 12, 2012	DP		NDF Apr 06, 2009
	6210705	Sep 30, 2018	DP	U-727	
	6348211	Sep 30, 2018	DP	U-727	
<u>METHYLPHENIDATE - DAYTRANA</u>					
021514 003	5958446	Dec 12, 2012	DP		NDF Apr 06, 2009
	6210705	Sep 30, 2018	DP	U-727	
	6348211	Sep 30, 2018	DP	U-727	
<u>METHYLPHENIDATE - DAYTRANA</u>					
021514 004	5958446	Dec 12, 2012	DP		NDF Apr 06, 2009
	6210705	Sep 30, 2018	DP	U-727	
	6348211	Sep 30, 2018	DP	U-727	
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>					
021259 001	6344215	Oct 27, 2020	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>					
021259 002	6344215	Oct 27, 2020	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>					
021259 003	6344215	Oct 27, 2020	DP		
<u>METHYLPHENIDATE HYDROCHLORIDE - METADATE CD</u>					
021259 004	6344215	Oct 27, 2020	DP		
<u>METOPROLOL SUCCINATE - TOPROL-XL</u>					
019962 001	4927640	May 22, 2007	DP		D-95 Feb 15, 2008
	4927640*PED	Nov 22, 2007			PED Aug 15, 2008
	4957745	Sep 18, 2007	DP	U-107	
	4957745*PED	Mar 18, 2008			
	5001161	Sep 18, 2007	DP		
	5001161*PED	Mar 18, 2008			
	5081154	Sep 18, 2007	DS		
	5081154*PED	Mar 18, 2008			



**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>METOPROLOL SUCCINATE - TOPROL-XL</u>					
019962 002	4927640	May 22, 2007	DP	D-95	Feb 15, 2008
	4927640*PED	Nov 22, 2007		PED	Aug 15, 2008
	4957745	Sep 18, 2007	DP	U-107	
	4957745*PED	Mar 18, 2008			
	5001161	Sep 18, 2007	DP		
	5001161*PED	Mar 18, 2008			
	5081154	Sep 18, 2007	DS		
	5081154*PED	Mar 18, 2008			
<u>METOPROLOL SUCCINATE - TOPROL-XL</u>					
019962 003	4927640	May 22, 2007	DP	D-95	Feb 15, 2008
	4927640*PED	Nov 22, 2007		PED	Aug 15, 2008
	4957745	Sep 18, 2007	DP	U-107	
	4957745*PED	Mar 18, 2008			
	5001161	Sep 18, 2007	DP		
	5001161*PED	Mar 18, 2008			
	5081154	Sep 18, 2007	DS		
	5081154*PED	Mar 18, 2008			
<u>METOPROLOL SUCCINATE - TOPROL-XL</u>					
019962 004	4927640	May 22, 2007	DP	D-95	Feb 15, 2008
	4927640*PED	Nov 22, 2007		PED	Aug 15, 2008
	4957745	Sep 18, 2007	DP	U-107	
	4957745*PED	Mar 18, 2008			
	5001161	Sep 18, 2007	DP	U-107	
	5001161*PED	Mar 18, 2008			
	5081154	Sep 18, 2007	DS	U-107	
	5081154*PED	Mar 18, 2008			
<u>METRONIDAZOLE - METROGEL</u>					
021789 001	6881726	Feb 21, 2022	DP	U-743	
<u>MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE - VUSION</u>					
021026 001	4911932	Mar 27, 2007	DP	U-718	NP Feb 16, 2009
<u>MINOXIDIL - MEN'S ROGAINE</u>					
021812 001	6946120	Apr 20, 2019	DP	U-702	NDF Jan 20, 2009
<u>MODAFINIL - PROVIGIL</u>					
020717 001	4927855	May 22, 2007		U-255	I-449 Jan 23, 2007
	4927855*PED	Nov 22, 2007			ODE Dec 24, 2005
	RE37516	Oct 06, 2014		U-255	PED Jul 23, 2007
	RE37516*PED	Apr 06, 2015			PED Jun 24, 2006
<u>MODAFINIL - PROVIGIL</u>					
020717 002	4927855	May 22, 2007		U-255	I-449 Jan 23, 2007
	4927855*PED	Nov 22, 2007			ODE Dec 24, 2005
	RE37516	Oct 06, 2014		U-255	PED Jul 23, 2007
	RE37516*PED	Apr 06, 2015			PED Jun 24, 2006
<u>MORPHINE SULFATE - KADIAN</u>					
020616 004	5378474	Mar 23, 2010			
<u>MORPHINE SULFATE - KADIAN</u>					
020616 005	5202128	Apr 13, 2010			
	5378474	Mar 23, 2010			
<u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX</u>					
021085 001	4990517	Dec 08, 2011	DS DP	U-298	
	6610327	Oct 29, 2019	DP	U-298	
<u>MOXIFLOXACIN HYDROCHLORIDE - AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER</u>					
021277 001	4990517	Dec 08, 2011	DS DP	U-298	
	6548079	Jul 25, 2020	DP	U-298	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MOXIFLOXACIN HYDROCHLORIDE - VIGAMOX</u>					
021598 001	4990517	Dec 08, 2011	DS DP	U-709	
	4990517*PED	Jun 08, 2012			
<u>NALTREXONE - VIVITROL</u>					
021897 001	5792477	May 02, 2017	DP	NDF	Apr 13, 2009
	5916598	May 02, 2017	DP		
	6110503	May 02, 2017	DP		
	6194006	Dec 30, 2018	DP		
	6264987	May 19, 2020	DP		
	6331317	Nov 12, 2019	DP		
	6379703	Dec 30, 2018	DP		
	6379704	May 19, 2020	DP		
	6395304	Nov 12, 2019	DP		
	6403114	May 02, 2017	DP		
	6495164	May 25, 2020	DP		
	6495166	Nov 12, 2019	DP		
	6534092	May 19, 2020	DP		
	6537586	Nov 12, 2019	DP		
	6596316	Dec 30, 2018	DP		
	6667061	May 25, 2020	DP		
	6713090	Nov 12, 2019	DP		
	6939033	Nov 12, 2019	DP		
<u>NELARABINE - ARRANON</u>					
021877 001	5747472	Feb 20, 2013		U-696	
	5747472	Feb 20, 2013		U-695	
	5747472	Feb 20, 2013		U-689	
	5821236	Feb 20, 2013		U-695	
<u>NIACIN - NIASPAN</u>					
020381 001	7011848	Sep 20, 2013		U-712	
<u>NIACIN - NIASPAN</u>					
020381 002	7011848	Sep 20, 2013		U-712	
<u>NIACIN - NIASPAN</u>					
020381 003	7011848	Sep 20, 2013		U-712	
<u>NIACIN - NIASPAN</u>					
020381 004	7011848	Sep 20, 2013		U-712	
<u>NIACIN - NIASPAN TITRATION STARTER PACK</u>					
020381 005	7011848	Sep 20, 2013		U-712	
<u>NICOTINE POLACRILEX - NICOTINE POLACRILEX</u>					
077007 001				PC	Aug 21, 2006
<u>NICOTINE POLACRILEX - NICOTINE POLACRILEX</u>					
077007 002				PC	Aug 21, 2006
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>					
019667 001	5753618	May 19, 2015			
	5753618*PED	Nov 19, 2015			
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>					
019667 002	5753618	May 19, 2015			
	5753618*PED	Nov 19, 2015			
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>					
019667 003	5753618	May 19, 2015			
	5753618*PED	Nov 19, 2015			
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>					
019667 004	5753618	May 19, 2015			
	5753618*PED	Nov 19, 2015			
<u>OCTREOTIDE ACETATE - SANDOSTATIN</u>					
019667 005	5753618	May 19, 2015			
	5753618*PED	Nov 19, 2015			

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes 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 LAR</u>					
021008 001	5538739	Jul 23, 2013	DP	M-55	May 10, 2009
	5538739*PED	Jan 23, 2014		ODE	Nov 25, 2005
	5639480	Jun 17, 2014	DP	PED	Nov 10, 2009
	5639480*PED	Dec 17, 2014		PED	May 25, 2006
	5688530	Nov 18, 2014		U-268	
	5688530*PED	May 18, 2015			
	5922338	Jul 13, 2016	DP		
	5922338*PED	Jan 13, 2017			
	5922682	Jul 13, 2016	DP		
	5922682*PED	Jan 13, 2017			
<u>OCTREOTIDE ACETATE - SANDOSTATIN LAR</u>					
021008 002	5538739	Jul 23, 2013		M-55	May 10, 2009
	5538739*PED	Jan 23, 2014		ODE	Nov 25, 2005
	5639480	Jun 17, 2014	DP	PED	Nov 10, 2009
	5639480*PED	Dec 17, 2014		PED	May 25, 2006
	5688530	Nov 18, 2014		U-268	
	5688530*PED	May 18, 2015			
	5922338	Jul 13, 2016	DP		
	5922338*PED	Jan 13, 2017			
	5922682	Jul 13, 2016	DP		
	5922682*PED	Jan 13, 2017			
<u>OCTREOTIDE ACETATE - SANDOSTATIN LAR</u>					
021008 003	5538739	Jul 23, 2013		M-55	May 25, 2009
	5538739*PED	Jan 23, 2014		ODE	Nov 25, 2005
	5639480	Jun 17, 2014	DP	PED	Nov 25, 2009
	5639480*PED	Dec 17, 2014		PED	May 25, 2006
	5688530	Nov 18, 2014		U-268	
	5688530*PED	May 18, 2015			
	5922338	Jul 13, 2016	DP		
	5922338*PED	Jan 13, 2017			
	5922682	Jul 13, 2016	DP		
	5922682*PED	Jan 13, 2017			
<u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>					
021849 001	6489346	Jul 16, 2016	DS DP	U-588	
	6645988	Jul 16, 2016	DS DP		
	6699885	Jul 16, 2016		U-588	
<u>OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID</u>					
021849 002	6489346	Jul 16, 2016	DS DP	U-623	
	6489346	Jul 16, 2016	DS DP	U-588	
	6645988	Jul 16, 2016	DS DP		
	6699885	Jul 16, 2016		U-623	
	6699885	Jul 16, 2016		U-588	
<u>OXALIPLATIN - ELOXATIN</u>					
021492 001	>A> 5290961	Jan 12, 2013		>A> I-441	Nov 04, 2007
	>A> 5290961*PED	Jul 12, 2013		>A> I-425	Jan 09, 2007
	>A> 5338874	Apr 07, 2013	DS	>A> NCE	Aug 09, 2007
	>A> 5338874*PED	Oct 07, 2013		>A> PED	May 04, 2008
	>A> 5420319	Aug 09, 2016	DS	>A> PED	Feb 09, 2008
	>A> 5420319*PED	Aug 09, 2016		>A> PED	Jul 09, 2007
<u>OXALIPLATIN - ELOXATIN</u>					
021492 002	>A> 5290961	Jan 12, 2013		>A> I-441	Nov 04, 2007
	>A> 5290961*PED	Jul 12, 2013		>A> I-425	Jan 09, 2007
	>A> 5338874	Apr 07, 2013	DS	>A> NCE	Aug 09, 2007
	>A> 5338874*PED	Oct 07, 2013		>A> PED	May 04, 2008
	>A> 5420319	Aug 09, 2016	DS	>A> PED	Feb 09, 2008
	>A> 5420319*PED	Aug 09, 2016		>A> PED	Jul 09, 2007

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OXALIPLATIN - ELOXATIN</u>					
021759 001	>A> 5290961	Jan 12, 2013	DS	>A> I-441	Nov 04, 2007
	>A> 5290961*PED	Jul 12, 2013		>A> NCE	Aug 09, 2007
	>A> 5338874	Apr 07, 2013	DS	>A> PED	May 04, 2008
	>A> 5338874*PED	Oct 07, 2013		>A> PED	Feb 09, 2008
	>A> 5420319	Aug 08, 2016	DS		
	>A> 5420319*PED	Feb 08, 2017			
	>A> 5716988	Aug 07, 2015	DP		
	>A> 5716988*PED	Feb 07, 2016			
<u>OXALIPLATIN - ELOXATIN</u>					
021759 002	>A> 5290961	Jan 12, 2013	DS	>A> I-441	Nov 04, 2007
	>A> 5290961*PED	Jul 12, 2013		>A> NCE	Aug 09, 2007
	>A> 5338874	Apr 07, 2013	DS	>A> PED	May 04, 2008
	>A> 5338874*PED	Oct 07, 2013		>A> PED	Feb 09, 2008
	>A> 5420319	Aug 08, 2016	DS		
	>A> 5420319*PED	Feb 08, 2017			
	>A> 5716988	Aug 07, 2015	DP		
	>A> 5716988*PED	Feb 07, 2016			
<u>OXCARBAZEPINE - TRILEPTAL</u>					
021014 001	7037525	Feb 12, 2018		U-724	
<u>OXCARBAZEPINE - TRILEPTAL</u>					
021014 002	7037525	Feb 12, 2018		U-724	
<u>OXCARBAZEPINE - TRILEPTAL</u>					
021014 003	7037525	Feb 12, 2018		U-724	
<u>OXCARBAZEPINE - TRILEPTAL</u>					
021285 001	7037525	Feb 12, 2018		U-724	
<u>OXYMORPHONE HYDROCHLORIDE - OPANA</u>					
021611 001				NDF	Jun 22, 2009
<u>OXYMORPHONE HYDROCHLORIDE - OPANA</u>					
021611 002				NDF	Jun 22, 2009
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>					
021610 001	5128143	Sep 19, 2008	DP	NDF	Jun 22, 2009
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>					
021610 002	5128143	Sep 19, 2008	DP	NDF	Jun 22, 2009
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>					
021610 003	5128143	Sep 19, 2008	DP	NDF	Jun 22, 2009
<u>OXYMORPHONE HYDROCHLORIDE - OPANA ER</u>					
021610 004	5128143	Sep 19, 2008	DP	NDF	Jun 22, 2009
<u>PAROXETINE HYDROCHLORIDE - PAXIL</u>					
020031 004	6133289	May 19, 2015		U-358	
<u>PAROXETINE HYDROCHLORIDE - PAXIL</u>					
020031 005	6133289	May 19, 2015		U-358	
<u>POSACONAZOLE - NOXAFIL</u>					
022003 001				>A> NCE	Sep 15, 2011
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>					
020667 001	4843086	Jun 27, 2006		U-231	
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>					
020667 002	4843086	Jun 27, 2006		U-231	
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>					
020667 003	4843086	Jun 27, 2006		U-231	
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>					
020667 004	4843086	Jun 27, 2006		U-231	
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>					
020667 005	4843086	Jun 27, 2006		U-231	

# PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX</u>					
020667 006	4843086	Jun 27, 2006		U-231	
<u>PRAVASTATIN SODIUM - PRAVASTATIN SODIUM</u>					
076056 001				PC	Oct 21, 2006
<u>PRAVASTATIN SODIUM - PRAVASTATIN SODIUM</u>					
076056 002				PC	Oct 21, 2006
<u>PRAVASTATIN SODIUM - PRAVASTATIN SODIUM</u>					
076056 003				PC	Oct 21, 2006
<u>QUETIAPINE FUMARATE - SEROQUEL</u>					
020639 006	4879288	Sep 26, 2011	DS DP	U-550	
<u>QUETIAPINE FUMARATE - SEROQUEL</u>					
020639 007	4879288	Sep 26, 2011	DS DP	U-550	
<u>RALOXIFENE HYDROCHLORIDE - EVISTA</u>					
020815 001	RE38968	Jul 28, 2012		U-662	
	RE38968	Jul 28, 2012		U-657	
	RE39049	Jul 28, 2012		U-662	
	RE39049	Jul 28, 2012		U-657	
	RE39050	Mar 02, 2014		U-662	
	RE39050	Mar 02, 2014		U-657	
<u>RAMIPRIL - ALTACE</u>					
019901 001	5061722	Oct 19, 2008			
<u>RANOLAZINE - RANEXA</u>					
021526 002	4567264	May 18, 2007	DS		Jan 27, 2011
	6303607	May 27, 2019		U-705	
	6369062	May 27, 2019		DP	
	6479496	May 27, 2019		U-705	
	6503911	May 27, 2019		DP	
	6525057	May 27, 2019		U-705	
	6562826	May 27, 2019		U-705	
	6617328	May 27, 2019		DP	
	6620814	May 27, 2019		U-705	
	6852724	May 27, 2019		U-705	
	6864258	May 27, 2019		U-705	
<u>RASAGILINE MESYLATE - AZILECT</u>					
021641 001	5387612	Feb 07, 2012		U-219	May 16, 2011
	5453446	Feb 07, 2012		U-219	
	5457133	Feb 07, 2012	DS DP		
	5532415	Jul 02, 2013	DS		
	5786390	Feb 07, 2012		DP	
	6126968	Sep 18, 2016		DP	
<u>RASAGILINE MESYLATE - AZILECT</u>					
021641 002	5387612	Feb 07, 2012		U-219	May 16, 2011
	5453446	Feb 07, 2012		U-219	
	5457133	Feb 07, 2012	DS DP		
	5532415	Jul 02, 2013	DS		
	5786390	Feb 07, 2012		DP	
	6126968	Sep 18, 2016		DP	
<u>RIFAXIMIN - XIFAXAN</u>					
021361 001	7045620	May 22, 2024	DS		
<u>RISEDRONATE SODIUM - ACTONEL</u>					
020835 001				M-52	Jan 24, 2009
<u>RISEDRONATE SODIUM - ACTONEL</u>					
020835 002				M-52	Jan 24, 2009
<u>RISEDRONATE SODIUM - ACTONEL</u>					
020835 003	>A> 5583122	Dec 10, 2013	DS DP	U-756	Aug 11, 2009
	>A> 5583122	Dec 10, 2013	DS DP	U-222	Jan 24, 2009
	>A> 6096342	Nov 22, 2011		DP	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes 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>					
020588 001	RE39181	Jul 11, 2014	DP		
<u>RIVASTIGMINE TARTRATE - EXELON</u>					
020823 003	4948807	Aug 14, 2012	DS	U-322	
<u>RIVASTIGMINE TARTRATE - EXELON</u>					
020823 004	4948807	Aug 14, 2012	DS	U-322	
<u>RIVASTIGMINE TARTRATE - EXELON</u>					
020823 005	4948807	Aug 14, 2012	DS	U-322	
<u>RIVASTIGMINE TARTRATE - EXELON</u>					
020823 006	4948807	Aug 14, 2012	DS	U-322	
<u>RIVASTIGMINE TARTRATE - EXELON</u>					
021025 001	4948807	Aug 14, 2012	DS	U-322	
<u>ROCURONIUM BROMIDE - ZEMURON</u>					
020214 003	4894369	Apr 13, 2008			
<u>SALMETEROL XINAFOATE - SEREVENT</u>					
020236 001	4992474	Feb 12, 2008			
	4992474*PED	Aug 12, 2008			
	5126375	Feb 12, 2008			
	5126375*PED	Aug 12, 2008			
	5225445	Feb 12, 2008		U-182	
	5225445*PED	Aug 12, 2008			
<u>SALMETEROL XINAFOATE - SEREVENT</u>					
020692 001	4992474	Feb 12, 2008			
	4992474*PED	Aug 12, 2008			
	5126375	Feb 12, 2008			
	5126375*PED	Aug 12, 2008			
	5225445	Feb 12, 2008		U-211	
	5225445*PED	Aug 12, 2008			
<u>SELEGILINE - EMSAM</u>					
021336 001	7070808	May 10, 2018	DS DP	NDF	Feb 27, 2009
	RE34579	Aug 18, 2007	DS DP	U-711	
<u>SELEGILINE - EMSAM</u>					
021336 002	RE34579	Aug 18, 2007	DS DP	U-711	NDF Feb 27, 2009
<u>SELEGILINE - EMSAM</u>					
021336 003	RE34579	Aug 18, 2007	DS DP	U-711	NDF Feb 27, 2009
<u>SELEGILINE HYDROCHLORIDE - ZELAPAR</u>					
021479 001	5648093	Jul 15, 2014	DP		
	6423342	Mar 01, 2016	DP		
<u>SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE</u>					
075719 001				PC	Feb 06, 2007
<u>SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE</u>					
075719 002				PC	Feb 06, 2007
<u>SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE</u>					
075719 003				PC	Feb 06, 2007
<u>SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE</u>					
076934 001				PC	Feb 03, 2007
<u>SERTRALINE HYDROCHLORIDE - ZOLOFT</u>					
020990 001	6727283	Oct 11, 2019	DP	U-580	
	6727283*PED	Apr 11, 2020			
	7067555	Nov 10, 2019	DP		
	7067555*PED	May 10, 2020			
<u>SEVELAMER HYDROCHLORIDE - RENAGEL</u>					
021179 001	7014846	Aug 11, 2013	DP	U-246	
<u>SEVELAMER HYDROCHLORIDE - RENAGEL</u>					
021179 002	7014846	Aug 11, 2013	DP	U-246	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SIMVASTATIN - SIMVASTATIN</u>					
076052 001				PC	Dec 20, 2006
<u>SIMVASTATIN - SIMVASTATIN</u>					
076052 002				PC	Dec 20, 2006
<u>SIMVASTATIN - SIMVASTATIN</u>					
076052 003				PC	Dec 20, 2006
<u>SIMVASTATIN - SIMVASTATIN</u>					
076052 004				PC	Dec 20, 2006
<u>SIMVASTATIN - SIMVASTATIN</u>					
076285 005				PC	Dec 20, 2006
<u>SIMVASTATIN - ZOCOR</u>					
019766 001	RE36481 ***	Jul 10, 2007		U-300	
	RE36481*PED	Jan 10, 2008		U-300	
	RE36520 ***	May 26, 2009		U-300	
	RE36520*PED	Nov 26, 2009		U-300	
<u>SIMVASTATIN - ZOCOR</u>					
019766 002	RE36481 ***	Jul 10, 2007		U-300	
	RE36481*PED	Jan 10, 2008		U-300	
	RE36520 ***	May 26, 2009		U-300	
	RE36520*PED	Nov 26, 2009		U-300	
<u>SIMVASTATIN - ZOCOR</u>					
019766 003	RE36481 ***	Jul 10, 2007		U-300	
	RE36481*PED	Jan 10, 2008		U-300	
	RE36520 ***	May 26, 2009		U-300	
	RE36520*PED	Nov 26, 2009		U-300	
<u>SIMVASTATIN - ZOCOR</u>					
019766 004	RE36481 ***	Jul 10, 2007		U-300	
	RE36481*PED	Jan 10, 2008		U-300	
	RE36520 ***	May 26, 2009		U-300	
	RE36520*PED	Nov 26, 2009		U-300	
<u>SIMVASTATIN - ZOCOR</u>					
019766 005	RE36481 ***	Jul 10, 2007		U-300	
	RE36481*PED	Jan 10, 2008		U-300	
	RE36520 ***	May 26, 2009		U-300	
	RE36520*PED	Nov 26, 2009		U-300	
<u>SIROLIMUS - RAPAMUNE</u>					
021083 001	>A> 5100899	Jul 07, 2013		U-290	
	>A> 5100899*PED	Jan 07, 2014			
<u>SIROLIMUS - RAPAMUNE</u>					
021110 001	>A> 5100899	Jul 07, 2013		U-290	
	>A> 5100899*PED	Jan 07, 2014			
<u>SIROLIMUS - RAPAMUNE</u>					
021110 002	>A> 5100899	Jul 07, 2013		U-290	
	>A> 5100899*PED	Jan 07, 2014			
<u>SIROLIMUS - RAPAMUNE</u>					
021110 003	>A> 5100899	Jul 07, 2013		U-290	
	>A> 5100899*PED	Jan 07, 2014			
<u>SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP</u>					
021892 001	5616346	May 18, 2013	DP	U-715 NP	Mar 16, 2009
<u>SOMATROPIN RECOMBINANT - GENOTROPIN</u>					
020280 006	4968299	Jun 28, 2008	DP	I-496	Apr 27, 2009
<u>SOMATROPIN RECOMBINANT - GENOTROPIN</u>					
020280 007	4968299	Jun 28, 2008	DP	I-496	Apr 27, 2009
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 001	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		

# PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 002	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 003	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 004				I-496	Apr 27, 2009
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 005	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 008	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 009	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 010	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 011	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 012	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE</u>					
020280 013	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
<u>SOMATROPIN RECOMBINANT - OMNITROPE</u>					
021426 001				NP	May 30, 2009
<u>SOMATROPIN RECOMBINANT - OMNITROPE</u>					
021426 002				NP	May 30, 2009
<u>SORAFENIB TOSYLATE - NEXAVAR</u>					
021923 001				ODE	Dec 20, 2012
<u>SUMATRIPTAN SUCCINATE - IMITREX STATDOSE</u>					
020080 003	4816470	Dec 28, 2006		U-72	
	4816470*PED	Jun 28, 2007			
	5037845	Aug 06, 2008		U-72	
	5037845*PED	Feb 06, 2009			
<u>SUNITINIB MALATE - SUTENT</u>					
021938 001	6573293	Feb 15, 2021	DS DP	U-703	Jan 26, 2011
<u>SUNITINIB MALATE - SUTENT</u>					
021938 002	6573293	Feb 15, 2021	DS DP	U-703	Jan 26, 2011
<u>SUNITINIB MALATE - SUTENT</u>					
021938 003	6573293	Feb 15, 2021	DS DP	U-703	Jan 26, 2011
<u>TACROLIMUS - PROGRAF</u>					
050708 001				ODE	Mar 29, 2013
<u>TACROLIMUS - PROGRAF</u>					
050708 002				ODE	Mar 29, 2013
<u>TACROLIMUS - PROGRAF</u>					
050708 003				ODE	Mar 29, 2013
<u>TACROLIMUS - PROGRAF</u>					
050709 001				ODE	Mar 29, 2013



**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u>					
021318 001	6977077	Aug 19, 2019		U-597	
<u>THALIDOMIDE - THALOMID</u>					
020785 001	5629327	Mar 01, 2013		U-731	
	6045501	Aug 28, 2018		U-731	
	6045501	Aug 28, 2018		U-371	
	6235756	Mar 01, 2013		U-731	
	6315720	Oct 23, 2020		U-442	
	6315720	Oct 23, 2020		U-731	
	6561976	Aug 28, 2018		U-731	
	6561976	Aug 28, 2018		U-371	
	6561977	Oct 23, 2020		U-371	
	6561977	Oct 23, 2020		U-731	
	6755784	Sep 23, 2020		U-731	
	6755784	Sep 23, 2020		U-371	
	6869399	Oct 23, 2020		U-371	
	6869399	Oct 23, 2020		U-732	
	6869399	Oct 23, 2020		U-733	
	6869399	Oct 23, 2020		U-731	
	6908432	Aug 28, 2018		U-371	
	6908432	Aug 28, 2018		U-731	
<u>THALIDOMIDE - THALOMID</u>					
020785 002	5629327	Mar 01, 2013		U-731	
	6045501	Aug 28, 2018		U-731	
	6045501	Aug 28, 2018		U-371	
	6235756	Mar 01, 2013		U-731	
	6315720	Oct 23, 2020		U-442	
	6315720	Oct 23, 2020		U-731	
	6561976	Aug 28, 2018		U-731	
	6561976	Aug 28, 2018		U-371	
	6561977	Oct 23, 2020		U-371	
	6561977	Oct 23, 2020		U-731	
	6755784	Sep 23, 2020		U-731	
	6755784	Sep 23, 2020		U-371	
	6869399	Oct 23, 2020		U-371	
	6869399	Oct 23, 2020		U-733	
	6869399	Oct 23, 2020		U-732	
	6869399	Oct 23, 2020		U-731	
	6908432	Aug 28, 2018		U-371	
	6908432	Aug 28, 2018		U-731	
<u>THALIDOMIDE - THALOMID</u>					
020785 003	5629327	Mar 01, 2013		U-731	
	6045501	Aug 28, 2018		U-731	
	6045501	Aug 28, 2018		U-371	
	6235756	Mar 01, 2013		U-731	
	6315720	Oct 23, 2020		U-442	
	6315720	Oct 23, 2020		U-731	
	6561976	Aug 28, 2018		U-731	
	6561976	Aug 28, 2018		U-371	
	6561977	Oct 23, 2020		U-371	
	6561977	Oct 23, 2020		U-731	
	6755784	Sep 23, 2020		U-731	
	6755784	Sep 23, 2020		U-371	
	6869399	Oct 23, 2020		U-371	
	6869399	Oct 23, 2020		U-732	
	6869399	Oct 23, 2020		U-733	
	6869399	Oct 23, 2020		U-731	
	6908432	Aug 28, 2018		U-371	
	6908432	Aug 28, 2018		U-731	

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes 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					M-53	Jan 23, 2009
<u>TIOTROPIUM BROMIDE MONOHYDRATE - SPIRIVA</u>						
021395 001	7070800	Jan 22, 2022	DP	U-566		
	RE38912	Oct 11, 2021	DP			
<u>TOPIRAMATE - TOPAMAX</u>						
020505 001	7018983	Oct 13, 2015		U-723		
<u>TOPIRAMATE - TOPAMAX</u>						
020505 002	7018983	Oct 13, 2015		U-723		
<u>TOPIRAMATE - TOPAMAX</u>						
020505 003	7018983	Oct 13, 2015		U-723		
<u>TOPIRAMATE - TOPAMAX</u>						
020505 004	7018983	Oct 13, 2015		U-723		
<u>TOPIRAMATE - TOPAMAX</u>						
020505 005	7018983	Oct 13, 2015		U-723		
<u>TOPIRAMATE - TOPAMAX</u>						
020505 006	7018983	Oct 13, 2015		U-723		
<u>TOPIRAMATE - TOPAMAX SPRINKLE</u>						
020844 001	7018983	Oct 13, 2015		U-723		
<u>TOPIRAMATE - TOPAMAX SPRINKLE</u>						
020844 002	7018983	Oct 13, 2015		U-723		
<u>TOPIRAMATE - TOPAMAX SPRINKLE</u>						
020844 003	7018983	Oct 13, 2015		U-723		
<u>TOPOTECAN HYDROCHLORIDE - Hycamtin</u>						
020671 001	5004758	May 28, 2010	DS DP	U-741		
<u>TREPROSTINIL SODIUM - REMODULIN</u>						
021272 001	5153222	Oct 06, 2014		U-455		
<u>TREPROSTINIL SODIUM - REMODULIN</u>						
021272 002	5153222	Oct 06, 2014		U-455		
<u>TREPROSTINIL SODIUM - REMODULIN</u>						
021272 003	5153222	Oct 06, 2014		U-455		
<u>TREPROSTINIL SODIUM - REMODULIN</u>						
021272 004	5153222	Oct 06, 2014		U-455		
<u>URSODIOL - URSO FORTE</u>						
020675 002	4859660	Nov 19, 2007		U-740		
<u>VARENICLINE TARTRATE - CHANTIX</u>						
021928 001	6410550	Nov 13, 2018	DS DP	U-56	NCE	May 10, 2011
	6890927	May 06, 2022	DS DP	U-56		
<u>VARENICLINE TARTRATE - CHANTIX</u>						
021928 002	6410550	Nov 13, 2018	DS DP	U-56	NCE	May 10, 2011
	6890927	May 06, 2022	DS DP	U-56		
<u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
076690 001					PC	Jan 30, 2007
<u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
076690 002					PC	Jan 30, 2007
<u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
076690 003					PC	Jan 30, 2007
<u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
076690 004					PC	Jan 30, 2007
<u>VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE</u>						
076690 005					PC	Jan 30, 2007

**PRESCRIPTION AND OTC DRUG PRODUCT  
PATENT AND EXCLUSIVITY LIST**

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ZANAMIVIR - RELENZA</u>					
021036 001	5648379	Jul 15, 2014	U-722	I-491	Mar 29, 2009
	5648379	Jul 15, 2014	U-721		
	5648379	Jul 15, 2014	U-274		
	6294572	Dec 15, 2014	DS DP		
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>					
020825 001	4831031	Mar 02, 2012	DS DP U-720	I-492	Aug 19, 2007
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>					
020825 002	4831031	Mar 02, 2012	DS DP U-720	I-492	Aug 19, 2007
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>					
020825 003	4831031	Mar 02, 2012	DS DP U-720	I-492	Aug 19, 2007
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>					
020825 004	4831031	Mar 02, 2012	DS DP U-720	I-492	Aug 19, 2007
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>					
021483 001	4831031	Mar 02, 2012	DS DP U-720	I-492	Aug 19, 2007
	5312925	Sep 01, 2012	DS DP U-720		
	6150366	May 27, 2019	DP U-719		
	6245766	Dec 18, 2018	U-601		
<u>ZIPRASIDONE MESYLATE - GEODON</u>					
020919 001	4831031	Mar 02, 2012	DS DP U-720		
<u>ZOLEDRONIC ACID - ZOMETA</u>					
021223 001	4939130	Sep 02, 2012	DS DP U-53		
<u>ZOLEDRONIC ACID - ZOMETA</u>					
021223 002	4939130	Sep 02, 2012	DS DP U-53		

## Footnotes:

- 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).
- 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>
- 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.
- \*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.
- \*\*\* U.S. Patent Nos. RE 36481 and RE 36520 are being relisted for Zocor (NDA 19-766) pursuant to the decision and related order in Ranbaxy Labs. v. Leavitt, No. 05-1838 (D.D.C. April 30, 2006). The '481 and '520 patents will remain listed in Approved Drug Products with Therapeutic Equivalence Evaluations until any applicable periods of exclusivity pursuant to section 505(j)(5)(B)(iv) of the Federal Food, Drug, and Cosmetic Act have been triggered and run, unless the agency's appeal of the decision to the U.S. Court of Appeals for the District of Columbia is decided in the agency's favor before the exclusivity periods have expired. While the patents remain listed, any new or pending ANDA referencing Zocor must contain patent certifications to these patents. For additional information on this matter, please refer to Docket Nos. 2005P-0008 and 2005P-0046.

## PATENT AND EXCLUSIVITY TERMS

Due to space limitations in the patent and exclusivity columns, abbreviations and references have been developed. Refer to the APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 25<sup>th</sup> Edition for a full listing of patent and exclusivity terms (Abbreviations, Dosing Schedule, Indications, and Patent Use Codes).

The current complete list of patent terms is available at <http://www.accessdata.fda.gov/scripts/cder/ob/docs/pattermsall.cfm>

The current complete list of exclusivity terms is available at <http://www.accessdata.fda.gov/scripts/cder/ob/docs/excltermsall.cfm>