

APPROVED DRUG PRODUCTS

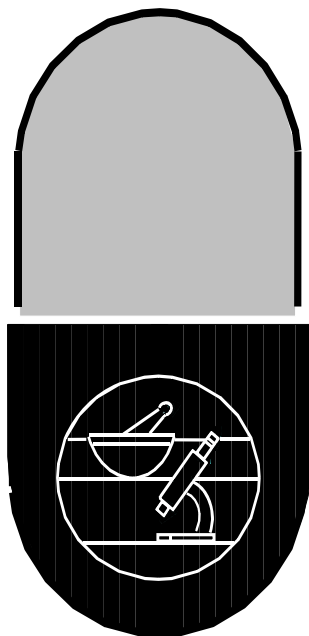
With Therapeutic Equivalence Evaluations



The "Orange Book"

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**CUMULATIVE
SUPPLEMENT 02
February 2010**



**APPROVED
DRUG PRODUCTS**

**WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS**

30th EDITION

**Department of Health and Human Services
Food and Drug Administration
Center for Drug Evaluation and Research
Office of Generic Drugs**

2010

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

**APPROVED DRUG PRODUCTS
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THERAPEUTIC EQUIVALENCE EVALUATIONS**

30th EDITION

Cumulative Supplement 02

February 2010

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30th EDITION

**CUMULATIVE SUPPLEMENT 02
February 2010**

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, 30th 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are for exportation, are for military use, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.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 30th Edition List will then be added to the "Discontinued Drug Product List" appearing in the 30th 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.
- 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 Staff received a letter November 7 that the product has been discontinued from manufacturing and marketing. The Orange Book 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 Staff 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.

Every effort is made to ensure the Cumulative Supplement is current and accurate. Applicant holders are requested to inform the FDA Orange Book Staff (OBS) of any changes or corrections. The OBS can be contacted by email at drugproducts@fda.hhs.gov. Send Changes by FAX: 240-276-8974; mail to:

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

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>
GOLDLINE LABORATORIES INC (GOLDLINE)	IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA (IVAX SUB TEVA PHARMS)
HLR TECHNOLOGY (HLR)	HOFFMANN LA ROCHE INC (HOFFMANN LA ROCHE)
IVAX PHARMACEUTICALS INC (IVAX PHARMS)	IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA (IVAX SUB TEVA PHARMS)
PROCTER AND GAMBLE CO (PROCTER AND GAMBLE)	WARNER CHILCOTT CO LLC (WARNER CHILCOTT)
PROCTER AND GAMBLE CO PHARMACEUTICALS INC SUB PROCTER AND GAMBLE CO (PROCTER AND GAMBLE)	WARNER CHILCOTT CO LLC (WARNER CHILCOTT)
TEVA PHARMACEUTICALS USA (TEVA PHARMS)	IVAX PHARMACEUTICALS INC SUB TEVA PHARMACEUTICALS USA (IVAX SUB TEVA PHARMS)

ZENITH GOLDLINE LABORATORIES INC

(ZENITH GOLDLINE)

ZENITH GOLDLINE PHARMACEUTICALS

(ZENITH GOLDLINE)

ZENITH GOLDLINE PHARMACEUTICALS INC

(ZENITH GOLDLINE)

IVAX PHARMACEUTICALS INC SUB

TEVA PHARMACEUTICALS USA

(IVAX SUB TEVA PHARMS)

IVAX PHARMACEUTICALS INC SUB

TEVA PHARMACEUTICALS USA

(IVAX SUB TEVA PHARMS)

IVAX PHARMACEUTICALS INC SUB

TEVA PHARMACEUTICALS USA

(IVAX SUB TEVA PHARMS)

1.4 AVAILABILITY OF THE EDITION

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

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

The downloaded Annual Edition and Cumulative Supplements are also available in a paper version (Approved Drug Products with Therapeutic Equivalence Evaluations, ADP) from the U.S. Government Printing Office: <http://bookstore.gpo.gov>; toll free 866-512-1800.

There are historical lists of Orange Book cumulative supplement product monthly changes at <http://www.fda.gov/Drugs/InformationOnDrugs/ucm086229.htm>. There are ASCII text files of the Orange Book drug product, patent, and exclusivity data at <http://www.fda.gov/Drugs/InformationOnDrugs/ucm129689.htm>. The drug product text files are provided in eobzip.zip format. The files are updated concurrently with the monthly cumulative supplements. The annual Orange Book Edition Appendices A, B, and C in PDF format 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 2008) 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 2009</u>	<u>MAR 2010</u>	<u>JUN 2010</u>	<u>SEPT 2010</u>	<u>DEC 2010</u>
DRUG PRODUCTS LISTED	13065				
SINGLE SOURCE	2460				
	(18.8%)				
MULTISOURCE	10516				
	(80.5%)				
THERAPEUTICALLY	10367				
EQUIVALENT	(79.3%)				
NOT THERAPEUTICALLY	149				
EQUIVALENT	(1.1%)				
EXCEPTIONS ¹	89				
	(0.7%)				
NEW MOLECULAR ENTITIES					
APPROVED	3				
NUMBER OF APPLICANTS	718				

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

1.6 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 application number preceded by "N" is a New Drug Application (NDA or innovator). The application number preceded by an "A" is an Abbreviated New Drug Application (ANDA or generic). 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 - 30TH EDITION

RX DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 2 - February 2010

1-1

ACETAMINOPHEN; BUTALBITAL

TABLET; ORAL

BUTAPAP

>D>	AB	MIKART	325MG;50MG	A089987	001	Oct 26, 1992	Feb	CTEC
>A>	AA		325MG;50MG	A089987	001	Oct 26, 1992	Feb	CTEC
PHRENILIN								
>D>	AB	+ VALEANT	325MG;50MG	A087811	001	Jun 19, 1985	Feb	CTEC
>A>	AA	+	325MG;50MG	A087811	001	Jun 19, 1985	Feb	CTEC

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

>D>		NEXGEN PHARMA	300MG;50MG;40MG	A040885	001	Nov 16, 2009	Feb	CRLD
>A>		+	300MG;50MG;40MG	A040885	001	Nov 16, 2009	Feb	CRLD
>D>	AB	WEST WARD	500MG;50MG;40MG	A040261	001	Oct 28, 1998	Feb	CTEC
>A>	AA		500MG;50MG;40MG	A040261	001	Oct 28, 1998	Feb	CTEC
ESGIC-PLUS								
>D>	AB	+ MIKART	500MG;50MG;40MG	A040085	001	Mar 28, 1996	Feb	CTEC
>A>	AA	+	500MG;50MG;40MG	A040085	001	Mar 28, 1996	Feb	CTEC

TABLET; ORAL

BUTALBITAL, ACETAMINOPHEN AND CAFFEINE

>D>	AB	CONCORD LABS NJ	325MG;50MG;40MG	A040864	001	Dec 01, 2008	Feb	CTEC
>A>	AA		325MG;50MG;40MG	A040864	001	Dec 01, 2008	Feb	CTEC
>D>	AB		500MG;50MG;40MG	A040883	001	Dec 23, 2008	Feb	CTEC
>A>	AA		500MG;50MG;40MG	A040883	001	Dec 23, 2008	Feb	CTEC
>D>	AB	MALLINCKRODT	325MG;50MG;40MG	A087804	001	Jan 24, 1985	Feb	CTEC
>A>	AA		325MG;50MG;40MG	A087804	001	Jan 24, 1985	Feb	CTEC
>D>	AB	MIKART	325MG;50MG;40MG	A089175	001	Jan 21, 1987	Feb	CTEC
>A>	AA		325MG;50MG;40MG	A089175	001	Jan 21, 1987	Feb	CTEC
>D>	AB	VINTAGE PHARMS	325MG;50MG;40MG	A040511	001	Aug 27, 2003	Feb	CTEC
>A>	AA		325MG;50MG;40MG	A040511	001	Aug 27, 2003	Feb	CTEC
>D>	AB		500MG;50MG;40MG	A040513	001	Aug 25, 2003	Feb	CTEC
>A>	AA		500MG;50MG;40MG	A040513	001	Aug 25, 2003	Feb	CTEC
>D>	AB	WATSON LABS	500MG;50MG;40MG	A040267	001	Jul 30, 1998	Feb	CTEC
>A>	AA		500MG;50MG;40MG	A040267	001	Jul 30, 1998	Feb	CTEC
>D>	AB	WEST WARD	325MG;50MG;40MG	A089718	001	Jun 12, 1995	Feb	CTEC
>A>	AA		325MG;50MG;40MG	A089718	001	Jun 12, 1995	Feb	CTEC
>D>	AB		500MG;50MG;40MG	A040336	001	Aug 18, 1999	Feb	CTEC
>A>	AA		500MG;50MG;40MG	A040336	001	Aug 18, 1999	Feb	CTEC
ESGIC-PLUS								
>D>	AB	+ MIKART	500MG;50MG;40MG	A089451	001	May 23, 1988	Feb	CTEC
>A>	AA	+	500MG;50MG;40MG	A089451	001	May 23, 1988	Feb	CTEC
FIORICET								
>D>	AB	+ WATSON PHARMS	325MG;50MG;40MG	A088616	001	Nov 09, 1984	Feb	CTEC
>A>	AA	+	325MG;50MG;40MG	A088616	001	Nov 09, 1984	Feb	CTEC

ACETAMINOPHEN; HYDROCODONE BITARTRATE

SOLUTION; ORAL

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

>A>		+ MIKART	300MG/15ML;10MG/15ML	A040881	001	Feb 25, 2010	Feb	NEWA
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ACETAMINOPHEN; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

ACETAMINOPHEN AND PENTAZOCINE HYDROCHLORIDE

>D>	AB	WATSON LABS	650MG;EQ 25MG BASE	A074699 001	Mar 24, 2000	Feb	CRLD
>A>	AB	+	650MG;EQ 25MG BASE	A074699 001	Mar 24, 2000	Feb	CRLD
>D>		TALACEN					
>D>	AB	+	SANOFI AVENTIS US	650MG;EQ 25MG BASE	N018458 001	Sep 23, 1982	Feb DISC
>A>		@	650MG;EQ 25MG BASE	N018458 001	Sep 23, 1982	Feb	DISC

ACITRETIN

CAPSULE; ORAL

SORIATANE

STIEFEL LABS INC

17.5MG

N019821 003 Aug 06, 2009 Jan NEWA

22.5MG

N019821 004 Aug 06, 2009 Jan NEWA

ALENDRONATE SODIUM

TABLET; ORAL

ALENDRONATE SODIUM

AB		CADISTA PHARMS	EQ 5MG BASE	A090557 001	Feb 18, 2010	Jan	NEWA
AB			EQ 10MG BASE	A090557 002	Feb 18, 2010	Jan	NEWA
AB			EQ 35MG BASE	A090557 003	Feb 18, 2010	Jan	NEWA
AB			EQ 70MG BASE	A090557 004	Feb 18, 2010	Jan	NEWA

ALPRAZOLAM

TABLET, ORALLY DISINTEGRATING; ORAL

ALPRAZOLAM

>A>	AB	ACTAVIS ELIZABETH	0.25MG	A078561 001	Mar 16, 2010	Feb	NEWA
>A>	AB		0.5MG	A078561 002	Mar 16, 2010	Feb	NEWA
>A>	AB		1MG	A078561 003	Mar 16, 2010	Feb	NEWA
>A>	AB		2MG	A078561 004	Mar 16, 2010	Feb	NEWA

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

AB		LUPIN PHARMS	EQ 2.5MG BASE;10MG	A078466 001	Feb 05, 2010	Jan	NEWA
AB			EQ 5MG BASE;10MG	A078466 002	Feb 05, 2010	Jan	NEWA
AB			EQ 5MG BASE;20MG	A078466 003	Feb 05, 2010	Jan	NEWA
AB			EQ 10MG BASE;20MG	A078466 004	Feb 05, 2010	Jan	NEWA

AMOXICILLIN

FOR SUSPENSION; ORAL

TRIMOX

>D>		APOTHECON	50MG/ML	A061886 001		Feb	DISC
>A>		@	50MG/ML	A061886 001		Feb	DISC
>D>	AB		125MG/5ML	A061886 002		Feb	DISC
>A>		@	125MG/5ML	A061886 002		Feb	DISC
>D>	AB		125MG/5ML	A062885 001	Mar 08, 1988	Feb	DISC
>A>		@	125MG/5ML	A062885 001	Mar 08, 1988	Feb	DISC
>D>	AB		250MG/5ML	A061886 003		Feb	DISC
>A>		@	250MG/5ML	A061886 003		Feb	DISC
>D>	AB		250MG/5ML	A062885 002	Mar 08, 1988	Feb	DISC
>A>		@	250MG/5ML	A062885 002	Mar 08, 1988	Feb	DISC

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL

AUGMENTIN '125'

>D>	@ GLAXOSMITHKLINE	125MG/5ML;EQ 31.25MG BASE/5ML	N050575 001	Aug 06, 1984	Feb	CMFD
>A>	AB	125MG/5ML;EQ 31.25MG BASE/5ML	N050575 001	Aug 06, 1984	Feb	CMFD

AUGMENTIN '250'

>D>	@ GLAXOSMITHKLINE	250MG/5ML;EQ 62.5MG BASE/5ML	N050575 002	Aug 06, 1984	Feb	CMFD
>A>	AB	250MG/5ML;EQ 62.5MG BASE/5ML	N050575 002	Aug 06, 1984	Feb	CMFD

TABLET; ORAL

AUGMENTIN '250'

>D>	@ GLAXOSMITHKLINE	250MG;EQ 125MG BASE	N050564 001	Aug 06, 1984	Feb	CMFD
>A>	AB	250MG;EQ 125MG BASE	N050564 001	Aug 06, 1984	Feb	CMFD

AUGMENTIN '500'

>D>	@ GLAXOSMITHKLINE	500MG;EQ 125MG BASE	N050564 002	Aug 06, 1984	Feb	CMFD
>A>	AB	500MG;EQ 125MG BASE	N050564 002	Aug 06, 1984	Feb	CMFD

AUGMENTIN '875'

>D>	@ GLAXOSMITHKLINE	875MG;EQ 125MG BASE	N050720 001	Feb 13, 1996	Feb	CMFD
>A>	AB	875MG;EQ 125MG BASE	N050720 001	Feb 13, 1996	Feb	CMFD

TABLET, EXTENDED RELEASE; ORAL

AUGMENTIN XR

>D>	@ GLAXOSMITHKLINE	1GM;EQ 62.5MG BASE	N050785 001	Sep 25, 2002	Feb	CMFD
>A>	AB	1GM;EQ 62.5MG BASE	N050785 001	Sep 25, 2002	Feb	CMFD

AMPHOTERICIN B

INJECTABLE, LIPID COMPLEX; INJECTION

ABELCET

>D>	+ ENZON	5MG/ML	N050724 001	Nov 20, 1995	Feb	CAHN
>A>	+ SIGMA TAU	5MG/ML	N050724 001	Nov 20, 1995	Feb	CAHN

ARMODAFINIL

TABLET; ORAL

NUVIGIL

>D>	CEPHALON	100MG	N021875 002	Mar 26, 2009	Feb	DISC
>A>	@	100MG	N021875 002	Mar 26, 2009	Feb	DISC
>D>		200MG	N021875 005	Mar 26, 2009	Feb	DISC
>A>	@	200MG	N021875 005	Mar 26, 2009	Feb	DISC

ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

ARTICAINE HYDROCHLORIDE WITH EPINEPHRINE

>A>	PIERREL	4%;EQ 0.009MG BASE/1.8ML (EQ 0.005MG BASE/ML)	N022466 001	Feb 26, 2010	Feb	NEWA
>A>	+	4%;EQ 0.018MG BASE/1.8ML (EQ 0.01MG BASE/ML)	N022466 002	Feb 26, 2010	Feb	NEWA

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

FIORINAL

>D>	AB	+ WATSON PHARMS	325MG;50MG;40MG	N017534 005	Apr 16, 1986	Feb	CTEC
>A>	AA	+	325MG;50MG;40MG	N017534 005	Apr 16, 1986	Feb	CTEC

LANORINAL

>D>	AB	LANNETT	325MG;50MG;40MG	A086996 002	Oct 11, 1985	Feb	CTEC
>A>	AA		325MG;50MG;40MG	A086996 002	Oct 11, 1985	Feb	CTEC

TABLET; ORAL

BUTALBITAL, ASPIRIN AND CAFFEINE

>D>	AB	ACTAVIS ELIZABETH	325MG;50MG;40MG	A086710 002	Aug 23, 1983	Feb	CTEC
>A>	AA		325MG;50MG;40MG	A086710 002	Aug 23, 1983	Feb	CTEC
>D>	AB	+ WEST WARD	325MG;50MG;40MG	A086162 002	Feb 16, 1984	Feb	CTEC
>A>	AA	+	325MG;50MG;40MG	A086162 002	Feb 16, 1984	Feb	CTEC

AZELASTINE HYDROCHLORIDE

SPRAY, METERED; NASAL

ASTELIN

AB	+	MEDA PHARMS	EQ 0.125MG BASE/SPRAY	N020114 001	Nov 01, 1996	Jan	CTEC
AB		APOTEX INC	EQ 0.125MG BASE/SPRAY	A077954 001	Apr 30, 2009	Jan	CMFD

AZITHROMYCIN

INJECTABLE; INJECTION

AZITHROMYCIN

>D>	AP	GENERAMEDIX	EQ 500MG BASE/VIAL	A065501 001	Nov 09, 2009	Feb	CAHN
>A>	AP	GLAND PHARMA LTD	EQ 500MG BASE/VIAL	A065501 001	Nov 09, 2009	Feb	CAHN

AZTREONAM

FOR SOLUTION; INHALATION

>A>		CAYSTON					
>A>	+	GILEAD	75MG/VIAL	N050814 001	Feb 22, 2010	Feb	NEWA

BACLOFEN

TABLET; ORAL

BACLOFEN

AB		MATRIX LABS LTD	10MG	A090334 001	Feb 18, 2010	Jan	NEWA
AB			20MG	A090334 002	Feb 18, 2010	Jan	NEWA

TABLET, ORALLY DISINTEGRATING; ORAL

KEMSTRO

@ SCHWARZ PHARMA

@

10MG	N021589 001	Oct 30, 2003	Jan	DISC
20MG	N021589 002	Oct 30, 2003	Jan	DISC

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

QVAR 40

>A>	+	TEVA BRANDED PHARM	0.04MG/INH	N020911 002	Sep 15, 2000	Feb	CAHN
>D>	+	TEVA GLOBAL	0.04MG/INH	N020911 002	Sep 15, 2000	Feb	CAHN

QVAR 80

>A>	+	TEVA BRANDED PHARM	0.08MG/INH	N020911 001	Sep 15, 2000	Feb	CAHN
>D>	+	TEVA GLOBAL	0.08MG/INH	N020911 001	Sep 15, 2000	Feb	CAHN

BENZTROPINE MESYLATE

TABLET; ORAL

BENZTROPINE MESYLATE

>A>	AA	+ LANNETT	0.5MG	A088877 001	Apr 11, 1985	Feb	CAHN
>A>	AA	+	1MG	A088894 001	Apr 11, 1985	Feb	CAHN
>A>	AA	+	2MG	A088895 001	Apr 11, 1985	Feb	CAHN
>D>	AA	+ PAR PHARM	0.5MG	A088877 001	Apr 11, 1985	Feb	CAHN
>D>	AA	+	1MG	A088894 001	Apr 11, 1985	Feb	CAHN
>D>	AA	+	2MG	A088895 001	Apr 11, 1985	Feb	CAHN

BETAXOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

BETAXOLOL HYDROCHLORIDE

AT	AKORN	EQ 0.5% BASE	A075386 001	Jun 30, 2000	Jan	CTNA
AT	NOVEX	EQ 0.5% BASE	A075446 001	Sep 28, 2000	Jan	CTNA
AT	WOCKHARDT	EQ 0.5% BASE	A078694 001	Nov 16, 2009	Jan	CAIN

TABLET; ORAL

BETAXOLOL HYDROCHLORIDE

>D>	AB	ACTAVIS TOTOWA	10MG	A075541 001	Oct 22, 1999	Feb	CAHN
>D>	AB	+	20MG	A075541 002	Oct 22, 1999	Feb	CAHN
>A>	AB	MIKAH PHARMA	10MG	A075541 001	Oct 22, 1999	Feb	CAHN
>A>	AB	+	20MG	A075541 002	Oct 22, 1999	Feb	CAHN

BUSPIRONE HYDROCHLORIDE

TABLET; ORAL

BUSPAR

>D>	AB	+	BRISTOL MYERS SQUIBB	15MG	N018731 003	Apr 22, 1996	Feb	DISC
>A>		@		15MG	N018731 003	Apr 22, 1996	Feb	DISC

BUSPIRONE HYDROCHLORIDE

>D>	AB	TEVA	30MG	A075022 004	Mar 25, 2004	Feb	CRLD
>A>	AB	+	30MG	A075022 004	Mar 25, 2004	Feb	CRLD

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

SOLUTION; IRRIGATION

BALANCED SALT

AT	B BRAUN	0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9MG/ML;6.4MG/ML;1.7MG/ML	A091387 001	Feb 03, 2010	Jan	NEWA
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CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

COREG CR

>D>		SB PHARMCO	40MG	N022012 003	Oct 20, 2006	Feb	CRLD
>A>		+	40MG	N022012 003	Oct 20, 2006	Feb	CRLD
>D>		+	80MG	N022012 004	Oct 20, 2006	Feb	CRLD
>A>			80MG	N022012 004	Oct 20, 2006	Feb	CRLD

CEFADROXIL/CEFADROXIL HEMIHYDRATE

FOR SUSPENSION; ORAL

CEFADROXIL

AB	+	LUPIN	EQ 500MG BASE/5ML	A065396 002	Feb 21, 2008	Jan	CRLD
		DURICEF					
		@ WARNER CHILCOTT	EQ 250MG BASE/5ML	N050527 003		Jan	DISC
		@	EQ 500MG BASE/5ML	N050527 001		Jan	DISC

CEFOTAXIME SODIUM

INJECTABLE; INJECTION

CEFOTAXIME SODIUM

AP		CEPHAZONE PHARMA	EQ 10GM BASE/VIAL	A065348 001	Jan 25, 2010	Jan	NEWA
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CEFOXITIN SODIUM

INJECTABLE; INJECTION

CEFOXITIN

>A>	AP	HIKMA FARMACEUTICA	EQ 1GM BASE/VIAL	A065238 001	Mar 12, 2010	Feb	NEWA
>A>	AP		EQ 2GM BASE/VIAL	A065238 002	Mar 12, 2010	Feb	NEWA

INJECTABLE; INJECTION

CEFOXITIN

>A>	AP	HIKMA FARMACEUTICA	EQ 10GM BASE/VIAL	A065239 001	Mar 02, 2010	Feb	NEWA
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CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

>D>	AB	LUPIN	250MG/5ML	A065261 002	Dec 19, 2005	Feb	CTEC
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>A>	AB	+	250MG/5ML	A065261 002	Dec 19, 2005	Feb	CTEC
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TABLET; ORAL

CEFPROZIL

>D>	AB	LUPIN	500MG	A065276 002	Dec 08, 2005	Feb	CRLD
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>A>	AB	+	500MG	A065276 002	Dec 08, 2005	Feb	CRLD
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CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CETIRIZINE HYDROCHLORIDE

AA		ACTAVIS MID ATLANTIC	5MG/5ML	A078617 001	Feb 02, 2010	Jan	NEWA
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CICLOPIROX

SHAMPOO; TOPICAL

CICLOPIROX

AT		PERRIGO	1%	A078594 001	Feb 16, 2010	Jan	NEWA
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SOLUTION; TOPICAL

CICLOPIROX

AT		VERSAPHARM	8%	A078975 001	Feb 17, 2010	Jan	NEWA
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CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPRO

>D>	AB	BAYER HLTHCARE	EQ 500MG BASE	N019537 003	Oct 22, 1987	Feb	CRLD
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>A>	AB	+	EQ 500MG BASE	N019537 003	Oct 22, 1987	Feb	CRLD
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>D>	AB	+	EQ 750MG BASE	N019537 004	Oct 22, 1987	Feb	CRLD
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>A>	AB		EQ 750MG BASE	N019537 004	Oct 22, 1987	Feb	CRLD
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CYANOCOBALAMIN

SPRAY, METERED; NASAL

CALOMIST

@ FLEMING

25MCG/SPRAY

N022102 001	Jul 27, 2007	Jan	DISC
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CYCLOPHOSPHAMIDE

INJECTABLE; INJECTION

CYTOXAN

>D>		@ BAXTER HLTHCARE	500MG/VIAL	N012142 003		Feb	CMFD
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>A>	AP	+	500MG/VIAL	N012142 003		Feb	CMFD
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>D>		@	1GM/VIAL	N012142 004	Aug 30, 1982	Feb	CMFD
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>A>	AP	+	1GM/VIAL	N012142 004	Aug 30, 1982	Feb	CMFD
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>D>		@	2GM/VIAL	N012142 005	Aug 30, 1982	Feb	CMFD
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>A>	AP	+	2GM/VIAL	N012142 005	Aug 30, 1982	Feb	CMFD
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>D>		LYOPHILIZED CYTOXAN					
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>D>		+	BAXTER HLTHCARE	100MG/VIAL	Dec 05, 1985	Feb	DISC
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>A>		@	100MG/VIAL	N012142 006	Dec 05, 1985	Feb	DISC
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>D>		+	200MG/VIAL	N012142 007	Dec 10, 1985	Feb	DISC
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>A>		@	200MG/VIAL	N012142 007	Dec 10, 1985	Feb	DISC
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>D>	AP	+	500MG/VIAL	N012142 008	Jan 04, 1984	Feb	DISC
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>A>		@	500MG/VIAL	N012142 008	Jan 04, 1984	Feb	DISC
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INJECTABLE; INJECTION

>D>		LYOPHILIZED CYTOXAN						
>D>	AP	+	BAXTER HLTHCARE	1GM/VIAL	N012142	010	Sep 24, 1985	Feb DISC
>A>		@		1GM/VIAL	N012142	010	Sep 24, 1985	Feb DISC
>D>	AP	+		2GM/VIAL	N012142	009	Dec 10, 1984	Feb DISC
>A>		@		2GM/VIAL	N012142	009	Dec 10, 1984	Feb DISC

DALFAMPRIDINE

TABLET, EXTENDED RELEASE; ORAL
AMPYRA

+	ACORDA	10MG	N022250	001	Jan 22, 2010	Jan	NEWA
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DES Loratadine

TABLET; ORAL

CLARINEX

AB	+	SCHERING PLOUGH	5MG	N021165	001	Dec 21, 2001	Jan CFTG
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DES Loratadine

AB		ORCHID HLTHCARE	5MG	A078357	001	Feb 19, 2010	Jan NEWA
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DESMOPRESSIN ACETATE

SPRAY, METERED; NASAL

STIMATE (NEEDS NO REFRIGERATION)

@	CSL BEHRING	1.5MG/SPRAY	N020355	002	Oct 24, 2007	Jan	DISC
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DEXMEDETOMIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

PRECEDEX

+	HOSPIRA	EQ 100MCG BASE/ML (EQ100MCG BASE/ML)	N021038	001	Dec 17, 1999	Jan	CAIN
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DIAZOXIDE

CAPSULE; ORAL

PROGLYCEM

>A>		@	TEVA BRANDED PHARM	50MG	N017425	001	Feb	CAHN
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>A>		@		100MG	N017425	002	Feb	CAHN
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>D>		@	TEVA GLOBAL	50MG	N017425	001	Feb	CAHN
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>D>		@		100MG	N017425	002	Feb	CAHN
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DICLOFENAC POTASSIUM

TABLET; ORAL

DICLOFENAC POTASSIUM

@	SANDOZ	50MG	A075582	001	Feb 23, 2001	Jan	DISC
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DIFLORASONE DIACETATE

CREAM; TOPICAL

PSORCON

>D>	AB1	+	SANOFI AVENTIS US	0.05%	N020205	001	Nov 20, 1992	Feb DISC
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>A>		@		0.05%	N020205	001	Nov 20, 1992	Feb DISC
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DILTIAZEM HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CARDIZEM LA

>D>			BIOVAIL LABS INTL	120MG	N021392	001	Feb 06, 2003	Feb CFTG
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>A>	AB			120MG	N021392	001	Feb 06, 2003	Feb CFTG
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>D>				180MG	N021392	002	Feb 06, 2003	Feb CFTG
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>A>	AB			180MG	N021392	002	Feb 06, 2003	Feb CFTG
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TABLET, EXTENDED RELEASE; ORAL

CARDIZEM LA

>D>		BIOVAIL LABS INTL	240MG	N021392 003	Feb 06, 2003	Feb	CFTG
>A>	AB		240MG	N021392 003	Feb 06, 2003	Feb	CFTG
>D>			300MG	N021392 004	Feb 06, 2003	Feb	CFTG
>A>	AB		300MG	N021392 004	Feb 06, 2003	Feb	CFTG
>D>			360MG	N021392 005	Feb 06, 2003	Feb	CFTG
>A>	AB		360MG	N021392 005	Feb 06, 2003	Feb	CFTG
>D>		+	420MG	N021392 006	Feb 06, 2003	Feb	CFTG
>A>	AB	+	420MG	N021392 006	Feb 06, 2003	Feb	CFTG
>A>		DILTIAZEM HYDROCHLORIDE					
>A>	AB	WATSON LABS FLORIDA	120MG	A077686 006	Mar 15, 2010	Feb	NEWA
>A>	AB		180MG	A077686 005	Mar 15, 2010	Feb	NEWA
>A>	AB		240MG	A077686 004	Mar 15, 2010	Feb	NEWA
>A>	AB		300MG	A077686 003	Mar 15, 2010	Feb	NEWA
>A>	AB		360MG	A077686 002	Mar 15, 2010	Feb	NEWA
>A>	AB		420MG	A077686 001	Mar 15, 2010	Feb	NEWA

EPINEPHRINE

INJECTABLE; IM-SC

TWINJECT 0.15

+		SHIONOGI PHARMA	EQ 0.15MG /DELIVERY	N020800 002	May 28, 2004	Jan	CAHN
		TWINJECT 0.3					
+		SHIONOGI PHARMA	EQ 0.3MG /DELIVERY	N020800 001	May 30, 2003	Jan	CAHN

ESTRADIOL VALERATE

INJECTABLE; INJECTION

ESTRADIOL VALERATE

AO		PHARMAFORCE	20MG/ML	A090920 001	Jan 19, 2010	Jan	NEWA
AO			40MG/ML	A090920 002	Jan 19, 2010	Jan	NEWA

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

NORETHINDRONE AND ETHINYL ESTRADIOL

>A>		WATSON LABS	0.035MG;0.4MG	A078379 001	Feb 23, 2010	Feb	NEWA
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TABLET; ORAL-28

NORETHINDRONE AND ETHINYL ESTRADIOL

AB		WATSON LABS	0.035MG,0.035MG,0.035MG;0.5MG,0.75MG,1MG	A076393 001	Feb 04, 2010	Jan	NEWA
AB			0.035MG;0.4MG	A078323 001	Feb 04, 2010	Jan	NEWA

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

AB		WATSON LABS	0.02MG;1MG	A078267 001	Sep 01, 2009	Jan	CDFR
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TABLET; ORAL-28

NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE

>D>	AB	TEVA PHARMS	0.02MG;1MG	A077077 001	May 20, 2005	Feb	CAHN
>D>	AB		0.03MG;1.5MG	A077075 001	Apr 28, 2005	Feb	CAHN
>A>	AB	VINTAGE	0.02MG;1MG	A077077 001	May 20, 2005	Feb	CAHN
>A>	AB		0.03MG;1.5MG	A077075 001	Apr 28, 2005	Feb	CAHN

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28

PREVIFEM

>D>	AB	TEVA PHARMS	0.035MG;0.25MG	A076334 001	Jan 09, 2004	Feb	CAHN
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TABLET; ORAL-28

PREVIFEM

>A>	AB	VINTAGE	0.035MG;0.25MG	A076334 001	Jan 09, 2004	Feb	CAHN
TRI-PREVIFEM							
>D>	AB	TEVA PHARMS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A076335 001	Mar 26, 2004	Feb	CAHN
>A>	AB	VINTAGE	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	A076335 001	Mar 26, 2004	Feb	CAHN

FAMOTIDINE

INJECTABLE; INJECTION

FAMOTIDINE

>D>	AP	BAXTER HLTHCARE	10MG/ML	A075488 001	Apr 16, 2001	Feb	CRLD
>A>	AP	+	10MG/ML	A075488 001	Apr 16, 2001	Feb	CRLD
>D>	AP		10MG/ML	A075799 001	Apr 30, 2002	Feb	CRLD
>A>	AP	+	10MG/ML	A075799 001	Apr 30, 2002	Feb	CRLD
FAMOTIDINE PRESERVATIVE FREE							
>D>	AP	BAXTER HLTHCARE	10MG/ML	A075486 001	Apr 16, 2001	Feb	CRLD
>A>	AP	+	10MG/ML	A075486 001	Apr 16, 2001	Feb	CRLD
>D>	AP		10MG/ML	A075789 001	Apr 30, 2002	Feb	CRLD
>A>	AP	+	10MG/ML	A075789 001	Apr 30, 2002	Feb	CRLD
FAMOTIDINE PRESERVATIVE FREE IN PLASTIC CONTAINER							
>D>	AP	BAXTER HLTHCARE	0.4MG/ML	A075591 001	May 10, 2001	Feb	CRLD
>A>	AP	+	0.4MG/ML	A075591 001	May 10, 2001	Feb	CRLD

TABLET, ORALLY DISINTEGRATING; ORAL

FLUXID

@ SCHWARZ PHARMA

20MG

N021712 001 Sep 24, 2004 Jan DISC

@

40MG

N021712 002 Sep 24, 2004 Jan DISC

FENOFIBRATE

CAPSULE; ORAL

ANTARA (MICRONIZED)

LUPIN ATLANTIS

43MG

N021695 001 Nov 30, 2004 Jan CAHN

@

87MG

N021695 002 Nov 30, 2004 Jan CAHN

+

130MG

N021695 003 Nov 30, 2004 Jan CAHN

TABLET; ORAL

FENOGLIDE

SHIONOGI PHARMA

40MG

N022118 001 Aug 10, 2007 Jan CAHN

+

120MG

N022118 002 Aug 10, 2007 Jan CAHN

FINASTERIDE

TABLET; ORAL

FINASTERIDE

>A>	AB	ACCORD HLTHCARE INC	5MG	A090121 001	Feb 23, 2010	Feb	NEWA
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FLUCONAZOLE

INJECTABLE; INJECTION

FLUCONAZOLE IN SODIUM CHLORIDE 0.9%

BEDFORD

100MG/50ML (2MG/ML)

A076087 002 Sep 26, 2008 Jan CTNA

FLUCONAZOLE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

>D>	AP	HOSPIRA	200MG/100ML (2MG/ML)	A076617 001	Jul 29, 2004	Feb	DISC
>A>		@	200MG/100ML (2MG/ML)	A076617 001	Jul 29, 2004	Feb	DISC
>D>	AP		400MG/200ML (2MG/ML)	A076617 002	Jul 29, 2004	Feb	DISC
>A>		@	400MG/200ML (2MG/ML)	A076617 002	Jul 29, 2004	Feb	DISC

FLUNISOLIDE

AEROSOL, METERED; INHALATION

AEROSPAN HFA

>A>	+	ACTON PHARMS	EQ 78MCG BASE/INH	N021247 001	Jan 27, 2006	Feb	CAHN
>D>	+	FOREST LABS	EQ 78MCG BASE/INH	N021247 001	Jan 27, 2006	Feb	CAHN

SPRAY, METERED; NASAL

NASAREL

>A>	AB	+	TEVA BRANDED PHARM	0.029MG/SPRAY	N020409 001	Mar 08, 1995	Feb	CAHN
>D>	AB	+	TEVA GLOBAL	0.029MG/SPRAY	N020409 001	Mar 08, 1995	Feb	CAHN

FLUOROURACIL

CREAM; TOPICAL

FLUOROURACIL

>A>	AB		TARO	5%	A090368 001	Mar 05, 2010	Feb	NEWA
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FLUOXETINE HYDROCHLORIDE

SOLUTION; ORAL

FLUOXETINE HYDROCHLORIDE

>A>	AA		LANNETT	EQ 20MG BASE/5ML	A076458 001	May 14, 2004	Feb	CAHN
>D>	AA		PAR PHARM	EQ 20MG BASE/5ML	A076458 001	May 14, 2004	Feb	CAHN

FOLIC ACID

TABLET; ORAL

FOLIC ACID

AA	+	PHARMAX	1MG	A040625 001	Jul 21, 2005	Jan	CRLD
AA	+	WATSON LABS	1MG	A080680 001		Jan	CMFD

FOSPHENYTOIN SODIUM

INJECTABLE; INJECTION

FOSPHENYTOIN SODIUM

>D>	AP		IV THERAP	EQ 50MG PHENYTOIN NA/ML	A078277 001	Aug 06, 2007	Feb	CAHN
>A>	AP		PHARMAFORCE	EQ 50MG PHENYTOIN NA/ML	A078277 001	Aug 06, 2007	Feb	CAHN

GLYBURIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLYBURIDE AND METFORMIN HYDROCHLORIDE

@	TEVA	1.25MG;250MG	A076821 001	Jan 27, 2005	Jan	DISC
@		2.5MG;500MG	A076821 002	Jan 27, 2005	Jan	DISC
@		5MG;500MG	A076821 003	Jan 27, 2005	Jan	DISC

GLYCOPYRROLATE

TABLET; ORAL

ROBINUL

AA	+	SHIONOGI PHARMA	1MG	N012827 001		Jan	CAHN
AA	+	SHIONOGI PHARMA	2MG	N012827 002		Jan	CAHN

GRISEOFULVIN, MICROCRYSTALLINE

TABLET; ORAL

FULVICIN-U/F

>A>	@	ELORAC	250MG	A060569 002		Feb	CAHN
>A>	@		500MG	A060569 001		Feb	CAHN
>D>	@	SCHERING	250MG	A060569 002		Feb	CAHN
>D>	@		500MG	A060569 001		Feb	CAHN

GUANFACINE HYDROCHLORIDE

TABLET; ORAL

GUANFACINE HYDROCHLORIDE

>D>	AB	ACTAVIS TOTOWA	EQ 1MG BASE	A074673 001	Feb 28, 1997	Feb	CAHN
>D>	AB		EQ 2MG BASE	A074673 002	Feb 28, 1997	Feb	CAHN
>A>	AB	MIKAH PHARMA	EQ 1MG BASE	A074673 001	Feb 28, 1997	Feb	CAHN
>A>	AB		EQ 2MG BASE	A074673 002	Feb 28, 1997	Feb	CAHN

HALOPERIDOL LACTATE

INJECTABLE; INJECTION

HALDOL

>D>	AP	+	ORTHO MCNEIL JANSSEN	EQ 5MG BASE/ML	N015923 001		Feb	DISC
>A>			@	EQ 5MG BASE/ML	N015923 001		Feb	DISC
			HALOPERIDOL					
>D>	AP		TEVA PARENTERAL	EQ 5MG BASE/ML	A076035 001	Aug 29, 2001	Feb	CRLD
>A>	AP	+		EQ 5MG BASE/ML	A076035 001	Aug 29, 2001	Feb	CRLD

HYDRALAZINE HYDROCHLORIDE

TABLET; ORAL

HYDRALAZINE HYDROCHLORIDE

	AA		HERITAGE PHARMS INC	10MG	A086242 001	Feb 04, 2010	Jan	NEWA
	AA			100MG	A086242 004	Feb 04, 2010	Jan	NEWA
>A>	AA		ZYDUS PHARMS USA	10MG	A040858 001	Feb 26, 2010	Feb	NEWA
>A>	AA			25MG	A040858 002	Feb 26, 2010	Feb	NEWA
>A>	AA			50MG	A040858 003	Feb 26, 2010	Feb	NEWA
>A>	AA			100MG	A040858 004	Feb 26, 2010	Feb	NEWA

HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

HYDROCHLOROTHIAZIDE

	AB		UNICHEM	12.5MG	A090510 001	Jan 19, 2010	Jan	NEWA
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HYDROCHLOROTHIAZIDE; LISINAPRIL

TABLET; ORAL

LISINAPRIL AND HYDROCHLOROTHIAZIDE

			@ TEVA	12.5MG;10MG	A075869 001	Jul 01, 2002	Jan	DISC
			@	12.5MG;20MG	A075869 002	Jul 01, 2002	Jan	DISC
			@	25MG;20MG	A075869 003	Jul 01, 2002	Jan	DISC

HYDROCHLOROTHIAZIDE; METOPROLOL TARTRATE

TABLET; ORAL

METOPROLOL TARTRATE AND HYDROCHLOROTHIAZIDE

	AB	+	MYLAN	25MG;100MG	A076792 002	Aug 20, 2004	Jan	CRLD
				50MG;100MG	A076792 003	Aug 20, 2004	Jan	CTEC

HYDROCORTISONE

CREAM; TOPICAL

HYDROCORTISONE

>D>	AT		FOUGERA	1%	A080693 003		Feb	CRLD
>A>	AT	+		1%	A080693 003		Feb	CRLD
>D>	AT			2.5%	A089414 001	Dec 16, 1986	Feb	CRLD
>A>	AT	+		2.5%	A089414 001	Dec 16, 1986	Feb	CRLD
>D>			HYTONE					
>D>	AT	+	SANOFI AVENTIS US	1%	A080472 003		Feb	DISC
>A>			@	1%	A080472 003		Feb	DISC

CREAM; TOPICAL

>D>		HYTONE						
>D>	AT	+	SANOFI AVENTIS US	2.5%	A080472	004		Feb DISC
>A>		@		2.5%	A080472	004		Feb DISC

LOTION; TOPICAL

>D>		HYTONE						
>D>	AT	+	SANOFI AVENTIS US	1%	A080473	003		Feb DISC
>A>		@		1%	A080473	003		Feb DISC
>D>	AT	+		2.5%	A080473	004	Nov 30, 1982	Feb DISC
>A>		@		2.5%	A080473	004	Nov 30, 1982	Feb DISC

IBUPROFEN

TABLET; ORAL

IBUPROFEN

AB		CONTRACT PHARMACAL	400MG	A071267	001	Oct 15, 1986	Jan	CAHN
AB			600MG	A071268	001	Oct 15, 1986	Jan	CAHN
AB			800MG	A072300	001	Jul 01, 1988	Jan	CAHN

IFOSFAMIDE

INJECTABLE; INJECTION

IFEX

>D>		@ BAXTER HLTHCARE	1GM/VIAL	N019763	001	Dec 30, 1988	Feb	CMFD
>A>	AP		1GM/VIAL	N019763	001	Dec 30, 1988	Feb	CMFD
>D>		@	3GM/VIAL	N019763	002	Dec 30, 1988	Feb	CMFD
>A>	AP		3GM/VIAL	N019763	002	Dec 30, 1988	Feb	CMFD

IFOSFAMIDE

>D>		+ APP PHARMS	1GM/VIAL	A076078	001	May 28, 2002	Feb	CTEC	
>A>	AP	+	1GM/VIAL	A076078	001	May 28, 2002	Feb	CTEC	
		+	1GM/VIAL	A076078	001	May 28, 2002	Jan	CTEC	
	AP		1GM/20ML (50MG/ML)	A090181	001	Sep 22, 2009	Jan	CPOT	
>D>		+	3GM/VIAL	A076078	002	May 28, 2002	Feb	CTEC	
>A>	AP	+	3GM/VIAL	A076078	002	May 28, 2002	Feb	CTEC	
		+	3GM/VIAL	A076078	002	May 28, 2002	Jan	CTEC	
	AP		3GM/60ML (50MG/ML)	A090181	002	Sep 22, 2009	Jan	CPOT	
	AP	+	TEVA PARENTERAL	1GM/20ML (50MG/ML)	A076657	001	Apr 04, 2007	Jan	CTEC
	AP	+		3GM/60ML (50MG/ML)	A076657	002	Apr 04, 2007	Jan	CTEC

IFOSFAMIDE; MESNA

INJECTABLE; INJECTION

IFEX/MESNEX KIT

>D>		+ BAXTER HLTHCARE	1GM/VIAL;100MG/ML	N019763	003	Oct 10, 1992	Feb	DISC
>A>		@	1GM/VIAL;100MG/ML	N019763	003	Oct 10, 1992	Feb	DISC
>D>		+	3GM/VIAL;100MG/ML	N019763	004	Oct 10, 1992	Feb	DISC
>A>		@	3GM/VIAL;100MG/ML	N019763	004	Oct 10, 1992	Feb	DISC

IMIPRAMINE HYDROCHLORIDE

TABLET; ORAL

IMIPRAMINE HYDROCHLORIDE

>A>	AB		LUPIN LTD	10MG	A090443	001	Mar 11, 2010	Feb	NEWA
>A>	AB			25MG	A090442	001	Mar 11, 2010	Feb	NEWA
>A>	AB			50MG	A090441	001	Mar 11, 2010	Feb	NEWA

IMIQUIMOD

CREAM; TOPICAL

ALDARA

>D>		+ GRACEWAY	5%	N020723	001	Feb 27, 1997	Feb	CFTG
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CREAM; TOPICALALDARA

>A>	AB	+	GRACEWAY	5%	N020723	001	Feb 27, 1997	Feb	CFTG
>A>			IMIQUIMOD						
>A>	AB		NYCOMED US	5%	A078548	001	Feb 25, 2010	Feb	NEWA

INAMRINONE LACTATEINJECTABLE; INJECTIONAMRINONE LACTATE

>D>	AP		BAXTER HLTHCARE CORP	EQ 5MG BASE/ML	A075542	001	May 10, 2000	Feb	DISC
>A>			@	EQ 5MG BASE/ML	A075542	001	May 10, 2000	Feb	DISC
>D>	AP	+	BEDFORD	EQ 5MG BASE/ML	A075513	001	May 09, 2000	Feb	CTEC
>A>		+		EQ 5MG BASE/ML	A075513	001	May 09, 2000	Feb	CTEC

LABETALOL HYDROCHLORIDEINJECTABLE; INJECTIONLABETALOL HYDROCHLORIDE

	AP		SAGENT STRIDES	5MG/ML	A079134	001	Feb 03, 2010	Jan	NEWA
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TABLET; ORALTRANDATE

>D>	AB		PROMETHEUS LABS	200MG	N018716	002	Aug 01, 1984	Feb	CRLD
>A>	AB	+		200MG	N018716	002	Aug 01, 1984	Feb	CRLD
>D>	AB	+		300MG	N018716	003	Aug 01, 1984	Feb	CRLD
>A>	AB			300MG	N018716	003	Aug 01, 1984	Feb	CRLD

LENALIDOMIDECAPSULE; ORALREVLIMID

		+	CELGENE	5MG	N021880	001	Dec 27, 2005	Jan	CRLD
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LEVETIRACETAMSOLUTION; ORALLEVETIRACETAM

>A>	AA		WOCKHARDT	100MG/ML	A090028	001	Mar 03, 2010	Feb	NEWA
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TABLET; ORALLEVETIRACETAM

	AB		TARO	250MG	A078960	004	Feb 01, 2010	Jan	NEWA
	AB			500MG	A078960	003	Feb 01, 2010	Jan	NEWA
	AB			750MG	A078960	002	Feb 01, 2010	Jan	NEWA
	AB			1GM	A078960	001	Feb 01, 2010	Jan	NEWA

LEVOTHYROXINE SODIUMCAPSULE; ORALTIROSINT

>D>		@	INST BIOCHIMIQUE	0.013MG	N022121	001	Aug 01, 2007	Feb	CMFD
>A>				0.013MG	N022121	001	Aug 01, 2007	Feb	CMFD
>D>		@		0.025MG	N021924	002	Oct 13, 2006	Feb	CMFD
>A>				0.025MG	N021924	002	Oct 13, 2006	Feb	CMFD
>D>		@		0.05MG	N021924	003	Oct 13, 2006	Feb	CMFD
>A>				0.05MG	N021924	003	Oct 13, 2006	Feb	CMFD
>D>		@		0.075MG	N021924	004	Oct 13, 2006	Feb	CMFD
>A>				0.075MG	N021924	004	Oct 13, 2006	Feb	CMFD
>A>				0.088MG	N021924	010	Oct 02, 2009	Feb	NEWA
>D>		@		0.1MG	N021924	005	Oct 13, 2006	Feb	CMFD
>A>				0.1MG	N021924	005	Oct 13, 2006	Feb	CMFD
>A>				0.112MG	N021924	008	Oct 02, 2009	Feb	NEWA

CAPSULE; ORAL

TIROSINT

>D>	@ INST BIOCHIMIQUE	0.125MG	N021924 006	Oct 13, 2006	Feb	CMFD
>A>		0.125MG	N021924 006	Oct 13, 2006	Feb	CMFD
>A>		0.137MG	N021924 009	Oct 02, 2009	Feb	NEWA
>D>	@	0.15MG	N021924 007	Oct 13, 2006	Feb	CMFD
>A>		0.15MG	N021924 007	Oct 13, 2006	Feb	CMFD

LIRAGLUTIDE RECOMBINANT

SOLUTION; SUBCUTANEOUS

VICTOZA

+	NOVO NORDISK INC	18MG/3ML (6MG/ML)	N022341 001	Jan 25, 2010	Jan	NEWA
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LISINOPRIL

TABLET; ORAL

LISINOPRIL

@	TEVA	2.5MG	A075783 001	Jul 01, 2002	Jan	DISC
@		5MG	A075783 002	Jul 01, 2002	Jan	DISC
@		10MG	A075783 003	Jul 01, 2002	Jan	DISC
@		20MG	A075783 004	Jul 01, 2002	Jan	DISC
@		30MG	A075783 005	Jul 01, 2002	Jan	DISC
@		40MG	A075783 006	Jul 01, 2002	Jan	DISC

LITHIUM CARBONATE

CAPSULE; ORAL

ESKALITH

@	NOVEN THERAP	300MG	N016860 001		Jan	DISC
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MECLIZINE HYDROCHLORIDE

TABLET; ORAL

ANTIVERT

AA	+	PFIZER	50MG	N010721 001	Jan 20, 1982	Jan	CMFD
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TABLET, CHEWABLE; ORAL

ANTIVERT

@	PFIZER	25MG	N010721 005		Jan	DISC
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MEPERIDINE HYDROCHLORIDE

SYRUP; ORAL

MEPERIDINE HYDROCHLORIDE

>D>	AA	ROXANE	50MG/5ML	A088744 001	Jan 30, 1985	Feb	CRLD
>A>	+		50MG/5ML	A088744 001	Jan 30, 1985	Feb	CRLD

TABLET; ORAL

DEMEROL

>D>	AA	+	SANOFI AVENTIS US	50MG	N005010 001	Feb	DISC
>A>	@			50MG	N005010 001	Feb	DISC

MEPERIDINE HYDROCHLORIDE

>D>	AA	ACTAVIS TOTOWA	50MG	A040331 001	May 28, 1999	Feb	CAHN
>D>	AA		100MG	A040331 002	May 28, 1999	Feb	CAHN
>A>	AA	MIKAH PHARMA	50MG	A040331 001	May 28, 1999	Feb	CAHN
>A>	AA		100MG	A040331 002	May 28, 1999	Feb	CAHN

MESALAMINE

TABLET, DELAYED RELEASE; ORAL

ASACOL

>D>	+	PROCTER AND GAMBLE	400MG	N019651 001	Jan 31, 1992	Feb	CAHN
>A>	+	WARNER CHILCOTT INC	400MG	N019651 001	Jan 31, 1992	Feb	CAHN

TABLET, DELAYED RELEASE; ORAL

ASACOL HD

>D>	+	PROCTER AND GAMBLE	800MG	N021830 001	May 29, 2008	Feb	CAHN
>A>	+	WARNER CHILCOTT INC	800MG	N021830 001	May 29, 2008	Feb	CAHN

METFORMIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

METFORMIN HYDROCHLORIDE

AB		TORRENT PHARMS	750MG	A079226 001	Feb 18, 2010	Jan	NEWA
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METHENAMINE HIPPURATE

TABLET; ORAL

UREX

>A>	AB	CNTY LINE PHARMS	1GM	N016151 001		Feb	CAHN
>D>	AB	VATRING PHARMS	1GM	N016151 001		Feb	CAHN

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE HYDROCHLORIDE

>D>	AP	HOSPIRA	EQ 5MG BASE/ML	A074147 001	Aug 02, 1996	Feb	DISC
>A>		@	EQ 5MG BASE/ML	A074147 001	Aug 02, 1996	Feb	DISC

TABLET; ORAL

METOCLOPRAMIDE HYDROCHLORIDE

		@ SANDOZ	EQ 10MG BASE	A074478 002	Oct 05, 1995	Jan	DISC
		@ WATSON LABS	EQ 10MG BASE	A070511 001	Jan 22, 1986	Jan	DISC

MILRINONE LACTATE

INJECTABLE; INJECTION

MILRINONE LACTATE IN DEXTROSE 5% IN PLASTIC CONTAINER

>D>	AP	BAXTER HLTHCARE	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075834 001	May 28, 2002	Feb	CTEC	
>A>	AP	+	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A075834 001	May 28, 2002	Feb	CTEC	
>D>	AP		EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A075834 002	May 28, 2002	Feb	CRLD	
>A>	AP	+	EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A075834 002	May 28, 2002	Feb	CRLD	
			MILRINONE LACTATE IN PLASTIC CONTAINER					
	AP	HIKMA FARMACEUTICA	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	A090038 001	Jan 21, 2010	Jan	NEWA	
	AP		EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	A090038 002	Jan 21, 2010	Jan	NEWA	
>D>			PRIMACOR IN DEXTROSE 5% IN PLASTIC CONTAINER					
>D>	AP	+	SANOFI AVENTIS US	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	N020343 003	Aug 09, 1994	Feb	DISC
>A>		@	EQ 20MG BASE/100ML (EQ 0.2MG BASE/ML)	N020343 003	Aug 09, 1994	Feb	DISC	
>D>	AP	+	EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	N020343 004	Aug 09, 1994	Feb	DISC	
>A>		@	EQ 40MG BASE/200ML (EQ 0.2MG BASE/ML)	N020343 004	Aug 09, 1994	Feb	DISC	

MORPHINE SULFATE

SOLUTION; ORAL

MORPHINE SULFATE

		ROXANE	20MG/5ML	N022195 002	Mar 17, 2008	Jan	CRLD
		+	100MG/5ML	N022195 003	Jan 25, 2010	Jan	NEWA

TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

>D>		@ AB GENERICS	15MG	A074862 001	Jul 07, 1998	Feb	CAHN
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TABLET, EXTENDED RELEASE; ORAL

MORPHINE SULFATE

>D>	@ AB GENERICS	30MG	A074862 002	Jul 07, 1998	Feb	CAHN
>D>	@	60MG	A074862 003	Jul 07, 1998	Feb	CAHN
>D>	@	100MG	A074769 001	Jul 02, 1998	Feb	CAHN
>D>	@	200MG	A074769 002	Jul 02, 1998	Feb	CAHN
>A>	@ PURDUE PHARMA LP	15MG	A074862 001	Jul 07, 1998	Feb	CAHN
>A>	@	30MG	A074862 002	Jul 07, 1998	Feb	CAHN
>A>	@	60MG	A074862 003	Jul 07, 1998	Feb	CAHN
>A>	@	100MG	A074769 001	Jul 02, 1998	Feb	CAHN
>A>	@	200MG	A074769 002	Jul 02, 1998	Feb	CAHN

NALIDIXIC ACID

TABLET; ORAL

NEGGRAM

>D>						
>D>	SANOFI AVENTIS US	250MG	N014214 002		Feb	DISC
>A>	@	250MG	N014214 002		Feb	DISC
>D>		500MG	N014214 004		Feb	DISC
>A>	@	500MG	N014214 004		Feb	DISC
>D>	+	1GM	N014214 005		Feb	DISC
>A>	@	1GM	N014214 005		Feb	DISC

NALOXONE HYDROCHLORIDE; PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

PENTAZOCINE AND NALOXONE HYDROCHLORIDES

>D>	AB	WATSON LABS	EQ 0.5MG BASE;EQ 50MG BASE	A074736 001	Jan 21, 1997	Feb	CRLD	
>A>	AB	+	EQ 0.5MG BASE;EQ 50MG BASE	A074736 001	Jan 21, 1997	Feb	CRLD	
>D>		TALWIN NX						
>D>	AB	+	SANOFI AVENTIS US	EQ 0.5MG BASE;EQ 50MG BASE	N018733 001	Dec 16, 1982	Feb	DISC
>A>		@		EQ 0.5MG BASE;EQ 50MG BASE	N018733 001	Dec 16, 1982	Feb	DISC

NISOLDIPINE

TABLET, EXTENDED RELEASE; ORAL

SULAR

+	SHIONOGI PHARMA	8.5MG	N020356 008	Jan 02, 2008	Jan	CAHN
	@	10MG	N020356 001	Feb 02, 1995	Jan	CAHN
+		17MG	N020356 007	Jan 02, 2008	Jan	CAHN
	@	20MG	N020356 002	Feb 02, 1995	Jan	CAHN
		25.5MG	N020356 006	Jan 02, 2008	Jan	CAHN
	@	30MG	N020356 003	Feb 02, 1995	Jan	CAHN
+		34MG	N020356 005	Jan 02, 2008	Jan	CAHN
	@	40MG	N020356 004	Feb 02, 1995	Jan	CAHN

NITROFURANTOIN

SUSPENSION; ORAL

FURADANTIN

+	SHIONOGI PHARMA	25MG/5ML	N009175 001		Jan	CAHN
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NITROGLYCERIN

AEROSOL; SUBLINGUAL

NITROLINGUAL

@	SHIONOGI PHARMA	0.4MG/SPRAY	N018705 001	Oct 31, 1985	Jan	CAHN
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SPRAY, METERED; SUBLINGUAL

NITROLINGUAL PUMPSPRAY

+	SHIONOGI PHARMA	0.4MG/SPRAY	N018705 002	Jan 10, 1997	Jan	CAHN
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OFLOXACIN

TABLET; ORAL

OFLOXACIN

@	LARKEN LABS	200MG	A076093 001	Sep 02, 2003	Jan	CAHN
@		300MG	A076093 002	Sep 02, 2003	Jan	CAHN
@		400MG	A076093 003	Sep 02, 2003	Jan	CAHN

ORPHENADRINE CITRATE

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATE

>D>	AB	ACTAVIS TOTOWA	100MG	A040284 001	Jun 19, 1998	Feb	CAHN
>A>	AB	GAVIS PHARMS	100MG	A040284 001	Jun 19, 1998	Feb	CAHN

OXALIPLATIN

INJECTABLE; INJECTION

OXALIPLATIN

AP	+	HOSPIRA INC	50MG/VIAL	A078815 001	Sep 30, 2009	Jan	CRLD
AP	+		100MG/VIAL	A078815 002	Sep 30, 2009	Jan	CRLD

OXCARBAZEPINE

TABLET; ORAL

OXCARBAZEPINE

AB		CADISTA PHARMS	150MG	A090239 001	Jan 25, 2010	Jan	NEWA
AB			300MG	A090239 002	Jan 25, 2010	Jan	NEWA
AB			600MG	A090239 003	Jan 25, 2010	Jan	NEWA

PALIPERIDONE PALMITATE

SUSPENSION, EXTENDED RELEASE; INTRAMUSCULAR

INVEGA SUSTENNA

+	JOHNSON AND JOHNSON	234MG/1.5ML (156MG/ML)	N022264 005	Jul 31, 2009	Jan	CRLD
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PAMIDRONATE DISODIUM

INJECTABLE; INJECTION

PAMIDRONATE DISODIUM

>D>	AP	GENERAMEDIX	30MG/VIAL	A078300 001	Mar 10, 2009	Feb	CAHN
>D>	AP		90MG/VIAL	A078300 002	Mar 10, 2009	Feb	CAHN
>A>	AP	MN PHARMS	30MG/VIAL	A078300 001	Mar 10, 2009	Feb	CAHN
>A>	AP		90MG/VIAL	A078300 002	Mar 10, 2009	Feb	CAHN

PEGADEMASE BOVINE

INJECTABLE; INJECTION

ADAGEN

>D>	+	ENZON PHARMS	250 UNITS/ML	N019818 001	Mar 21, 1990	Feb	CAHN
>A>	+	SIGMA TAU	250 UNITS/ML	N019818 001	Mar 21, 1990	Feb	CAHN

PERINDOPRIL ERBUMINE

TABLET; ORAL

PERINDOPRIL ERBUMINE

AB		LUPIN LTD	2MG	A078263 001	Jan 27, 2010	Jan	NEWA
AB			4MG	A078263 002	Jan 27, 2010	Jan	NEWA
AB			8MG	A078263 003	Jan 27, 2010	Jan	NEWA

PHENDIMETRAZINE TARTRATE

CAPSULE, EXTENDED RELEASE; ORAL

BONTRIL

>D>	BC	MALLINCKRODT	105MG	A088021 001	Sep 21, 1982	Feb	CAHN
>A>	BC	VALEANT	105MG	A088021 001	Sep 21, 1982	Feb	CAHN

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

FOR SOLUTION; ORAL

LAX-LYTE WITH FLAVOR PACKS

>A>	AA	PADDOCK	420GM/BOT;1.48GM/BOT;5.72GM/BOT;1.2GM/BOT	A079232 001	Feb 25, 2010	Feb	NEWA
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POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE ANHYDROUS

FOR SOLUTION; ORAL

PEG 3350 AND ELECTROLYTES

AA	MYLAN		236GM;2.97GM;6.74GM;5.86GM;22.74GM	A090928 001	Jan 28, 2010	Jan	NEWA
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POLYETHYLENE GLYCOL 3350 AND ELECTROLYTES

>A>	AA	PADDOCK LABS	240GM/BOT;2.98GM/BOT;6.72GM/BOT;5.84GM/BOT;22.72GM/BOT	A090712 001	Feb 25, 2010	Feb	NEWA
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POTASSIUM CITRATE

TABLET, EXTENDED RELEASE; ORAL

UROCIT-K

>A>		MISSION PHARMA	15MEQ	N019071 003	Dec 30, 2009	Feb	NEWA
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PRAMIPEXOLE DIHYDROCHLORIDE

>A> TABLET, EXTENDED RELEASE; ORAL

>A> MIRAPEX ER

>A>		BOEHRINGER INGELHEIM	0.375MG	N022421 001	Feb 19, 2010	Feb	NEWA
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>A>			0.75MG	N022421 002	Feb 19, 2010	Feb	NEWA
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>A>			1.5MG	N022421 003	Feb 19, 2010	Feb	NEWA
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>A>			3MG	N022421 004	Feb 19, 2010	Feb	NEWA
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>A>	+		4.5MG	N022421 005	Feb 19, 2010	Feb	NEWA
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PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

PRAZOSIN HYDROCHLORIDE

@ WATSON LABS

EQ 1MG BASE

A072352 001 May 16, 1989 Jan DISC

@

EQ 2MG BASE

A072333 001 May 16, 1989 Jan DISC

PREDNISONE

TABLET; ORAL

PREDNISONE

AB		CONTRACT PHARMACAL	5MG	A080209 001		Jan	CAHN
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PREGABALIN

SOLUTION; ORAL

LYRICA

+ PFIZER 20MG/ML

N022488 001 Jan 04, 2010 Jan NEWA

PROCAINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINE HYDROCHLORIDE

@ WATSON LABS

1%

A080658 001 Jan DISC

PROCHLORPERAZINE MALEATE

TABLET; ORAL

PROCHLORPERAZINE MALEATE

	@ DURAMED PHARMS BARR	EQ 5MG BASE	A040207 001	May 01, 1997	Jan	DISC
	@	EQ 10MG BASE	A040207 002	May 01, 1997	Jan	DISC
	PROCOMP					
AB	CADISTA PHARMS	EQ 5MG BASE	A040268 001	Feb 27, 1998	Jan	CTNA
AB		EQ 10MG BASE	A040268 002	Feb 27, 1998	Jan	CTNA

PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

PROPOXYPHENE HYDROCHLORIDE

	@ PAR PHARM	65MG	A080269 001		Jan	DISC
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QUINIDINE SULFATE

TABLET; ORAL

QUINIDINE SULFATE

	@ CONTRACT PHARMACAL	200MG	A083808 001		Jan	CAHN
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RISPERIDONE

INJECTABLE; INTRAMUSCULAR

RISPERDAL CONSTA

+	ORTHO MCNEIL JANSSEN	25MG/VIAL	N021346 001	Oct 29, 2003	Jan	CRLD
		50MG/VIAL	N021346 003	Oct 29, 2003	Jan	CRLD

TABLET; ORAL

RISPERIDONE

>D>	AB	SYNTHON PHARMS	0.25MG	A078187 001	Oct 22, 2009	Feb	DISC
>A>		@	0.25MG	A078187 001	Oct 22, 2009	Feb	DISC
>D>	AB		0.5MG	A078187 002	Oct 22, 2009	Feb	DISC
>A>		@	0.5MG	A078187 002	Oct 22, 2009	Feb	DISC
>D>	AB		1MG	A078187 003	Oct 22, 2009	Feb	DISC
>A>		@	1MG	A078187 003	Oct 22, 2009	Feb	DISC
>D>	AB		2MG	A078187 004	Oct 22, 2009	Feb	DISC
>A>		@	2MG	A078187 004	Oct 22, 2009	Feb	DISC
>D>	AB		3MG	A078187 005	Oct 22, 2009	Feb	DISC
>A>		@	3MG	A078187 005	Oct 22, 2009	Feb	DISC
>D>	AB		4MG	A078187 006	Oct 22, 2009	Feb	DISC
>A>		@	4MG	A078187 006	Oct 22, 2009	Feb	DISC

RITONAVIR

>A>	TABLET; ORAL						
>A>	RITONAVIR						
>A>	+	ABBOTT LABS	100MG	N022417 001	Feb 10, 2010	Feb	NEWA

RIVASTIGMINE TARTRATE

CAPSULE; ORAL

EXELON

>D>	AB	+	NOVARTIS	EQ 6MG BASE	N020823 006	Apr 21, 2000	Feb	CRLD
>A>	AB			EQ 6MG BASE	N020823 006	Apr 21, 2000	Feb	CRLD

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

ROPINIROLE HYDROCHLORIDE

>A>	AB	GLENMARK GENERICS	EQ 0.25MG BASE	A090135 001	Feb 25, 2010	Feb	NEWA
>A>	AB		EQ 0.5MG BASE	A090135 002	Feb 25, 2010	Feb	NEWA

TABLET; ORALROPINIROLE HYDROCHLORIDE

>A>	AB	GLENMARK GENERICS	EQ 1MG BASE	A090135 003	Feb 25, 2010	Feb	NEWA
>A>	AB		EQ 2MG BASE	A090135 004	Feb 25, 2010	Feb	NEWA
>A>	AB		EQ 3MG BASE	A090135 005	Feb 25, 2010	Feb	NEWA
>A>	AB		EQ 4MG BASE	A090135 006	Feb 25, 2010	Feb	NEWA
>A>	AB		EQ 5MG BASE	A090135 007	Feb 25, 2010	Feb	NEWA

SODIUM BICARBONATEINJECTABLE; INJECTIONSODIUM BICARBONATE

>D>		HOSPIRA	0.9MEQ/ML	A077394 001	Nov 09, 2005	Feb	CRLD
>A>	+		0.9MEQ/ML	A077394 001	Nov 09, 2005	Feb	CRLD

STAVUDINECAPSULE; ORALSTAVUDINE

>D>	AB	MATRIX LABS INC	15MG	A079069 001	Dec 29, 2008	Feb	CAHN
>D>	AB		20MG	A079069 002	Dec 29, 2008	Feb	CAHN
>D>	AB		30MG	A079069 003	Dec 29, 2008	Feb	CAHN
>D>	AB		40MG	A079069 004	Dec 29, 2008	Feb	CAHN
>A>	AB	MYLAN	15MG	A079069 001	Dec 29, 2008	Feb	CAHN
>A>	AB		20MG	A079069 002	Dec 29, 2008	Feb	CAHN
>A>	AB		30MG	A079069 003	Dec 29, 2008	Feb	CAHN
>A>	AB		40MG	A079069 004	Dec 29, 2008	Feb	CAHN

SULFAMETHOXAZOLE; TRIMETHOPRIMTABLET; ORALSULFAMETHOXAZOLE AND TRIMETHOPRIM

AB		AUROBINDO PHARMA	400MG;80MG	A090624 001	Feb 16, 2010	Jan	NEWA
AB			800MG;160MG	A090624 002	Feb 16, 2010	Jan	NEWA

TAMSULOSIN HYDROCHLORIDECAPSULE; ORALFLOMAX

>D>	+	BOEHRINGER INGELHEIM	0.4MG	N020579 001	Apr 15, 1997	Feb	CFTG
>A>	AB	+	0.4MG	N020579 001	Apr 15, 1997	Feb	CFTG
>A>		TAMSULOSIN HYDROCHLORIDE					
>A>	AB	IMPAX LABS	0.4MG	A090377 001	Mar 02, 2010	Feb	NEWA

TECHNETIUM TC-99M SODIUM PERTECHNETATE GENERATORSOLUTION; INJECTION, ORALTECHNELITE

>D>		LANTHEUS MEDCL	0.0083-2.7 CI/GENERATOR	N017771 001		Feb	CRLD
>A>	+		0.0083-2.7 CI/GENERATOR	N017771 001		Feb	CRLD
>D>		ULTRA-TECHNEKOW FM					
>D>		MALLINCKRODT	0.25-3 CI/GENERATOR	N017243 002		Feb	CRLD
>A>	+		0.25-3 CI/GENERATOR	N017243 002		Feb	CRLD

TEMOZOLOMIDECAPSULE; ORALTEMODAR

>D>		SCHERING	5MG	N021029 001	Aug 11, 1999	Feb	CFTG
>A>	AB		5MG	N021029 001	Aug 11, 1999	Feb	CFTG
>D>			20MG	N021029 002	Aug 11, 1999	Feb	CFTG
>A>	AB		20MG	N021029 002	Aug 11, 1999	Feb	CFTG

CAPSULE; ORALTEMODAR

>D>		SCHERING	100MG	N021029 003	Aug 11, 1999	Feb	CFTG
>A>	AB		100MG	N021029 003	Aug 11, 1999	Feb	CFTG
>D>			140MG	N021029 005	Oct 19, 2006	Feb	CFTG
>A>	AB		140MG	N021029 005	Oct 19, 2006	Feb	CFTG
>D>			180MG	N021029 006	Oct 19, 2006	Feb	CFTG
>A>	AB		180MG	N021029 006	Oct 19, 2006	Feb	CFTG
>D>		+	250MG	N021029 004	Aug 11, 1999	Feb	CFTG
>A>	AB	+	250MG	N021029 004	Aug 11, 1999	Feb	CFTG

TEMOZOLOMIDE

>A>	AB	BARR	5MG	A078879 001	Mar 01, 2010	Feb	NEWA
>A>	AB		20MG	A078879 002	Mar 01, 2010	Feb	NEWA
>A>	AB		100MG	A078879 003	Mar 01, 2010	Feb	NEWA
>A>	AB		140MG	A078879 005	Mar 01, 2010	Feb	NEWA
>A>	AB		180MG	A078879 006	Mar 01, 2010	Feb	NEWA
>A>	AB		250MG	A078879 004	Mar 01, 2010	Feb	NEWA

TERCONAZOLECREAM; VAGINALTERCONAZOLE

BX	+	NYCOMED US	0.8%	N021735 001	Oct 01, 2004	Jan	CAHN
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TINZAPARIN SODIUMINJECTABLE; INJECTIONINNOHEP

>D>		+	CELGENE	20,000 IU/ML	N020484 001	Jul 14, 2000	Feb	CAHN
>A>		+	LEO PHARMA AS	20,000 IU/ML	N020484 001	Jul 14, 2000	Feb	CAHN

TOPIRAMATETABLET; ORALTOPIRAMATE

		@	PLIVA HRVATSKA DOO	25MG	A077905 001	Mar 30, 2009	Jan	DISC
		@		50MG	A077905 002	Mar 30, 2009	Jan	DISC
		@		100MG	A077905 003	Mar 30, 2009	Jan	DISC
		@		200MG	A077905 004	Mar 30, 2009	Jan	DISC

TRANDOLAPRILTABLET; ORALTRANDOLAPRIL

>A>	AB	EPIC PHARMA	1MG	A078508 003	Jun 18, 2008	Feb	CAHN
>A>	AB		2MG	A078508 001	Jun 18, 2008	Feb	CAHN
>A>	AB		4MG	A078508 002	Jun 18, 2008	Feb	CAHN
>D>	AB	GLENMARK GENERICS	1MG	A078508 003	Jun 18, 2008	Feb	CAHN
>D>	AB		2MG	A078508 001	Jun 18, 2008	Feb	CAHN
>D>	AB		4MG	A078508 002	Jun 18, 2008	Feb	CAHN

TRAZODONE HYDROCHLORIDETABLET; ORALTRAZODONE HYDROCHLORIDE

>D>	AB	TEVA	50MG	A072192 001	Feb 02, 1989	Feb	CAHN
>D>	AB		100MG	A072193 001	Feb 02, 1989	Feb	CAHN
>A>	AB	VINTAGE	50MG	A072192 001	Feb 02, 1989	Feb	CAHN
>A>	AB		100MG	A072193 001	Feb 02, 1989	Feb	CAHN

>A>	TABLET, EXTENDED RELEASE; ORAL						
>A>	OLEPTRO						
>A>	LABOPHARM	150MG	N022411	001	Feb 02, 2010	Feb	NEWA
>A>	+	300MG	N022411	002	Feb 02, 2010	Feb	NEWA

TRETINOIN

	CAPSULE; ORAL						
	TRETINOIN						
>D>	AB	BARR	10MG	A077684	001	Jun 22, 2007	Feb CRLD
>A>	+		10MG	A077684	001	Jun 22, 2007	Feb CRLD
>D>	VESANOID						
>D>	AB	+ ROCHE	10MG	N020438	001	Nov 22, 1995	Feb DISC
>A>	@		10MG	N020438	001	Nov 22, 1995	Feb DISC

UNOPROSTONE ISOPROPYL

SOLUTION/DROPS; OPHTHALMIC

RESCULA

@ SUCAMPO PHARMS

0.15%

N021214 001 Aug 03, 2000 Jan DISC

UREA, C-14

CAPSULE; ORAL

PYTEST

>A>	+	AVENT	1uCi	N020617	001	May 09, 1997	Feb CAHN
>D>	+	BALLARD MEDCL	1uCi	N020617	001	May 09, 1997	Feb CAHN
		PYTEST KIT					
>A>	+	AVENT	1uCi	N020617	002	May 09, 1997	Feb CAHN
>D>	+	BALLARD MEDCL	1uCi	N020617	002	May 09, 1997	Feb CAHN

URSODIOL

CAPSULE; ORAL

URSODIOL

>D>	AB	ACTAVIS TOTOWA	300MG	A075517	001	Mar 14, 2000	Feb CAHN
>A>	AB	MIKAH PHARMA	300MG	A075517	001	Mar 14, 2000	Feb CAHN
	AB	MYLAN	300MG	A090530	001	Feb 17, 2010	Jan NEWA

VALPROATE SODIUM

INJECTABLE; INJECTION

VALPROATE SODIUM

AP HIKMA FARMACEUTICA

EQ 100MG BASE/ML

A078523 001 Feb 17, 2010 Jan NEWA

VELAGLUCERASE ALFA

>A>	INJECTABLE; IV (INFUSION)						
>A>	VPRIV						
>A>	SHIRE HUMAN GENETIC	400 UNITS/VIAL	N022575	001	Feb 26, 2010	Feb	NEWA
>A>	POWDER; IV (INFUSION)						
>A>	VPRIV						
>A>	+	SHIRE HUMAN GENETIC	200 UNITS/VIAL	N022575	002	Feb 26, 2010	Feb NEWA

OTC DRUG PRODUCT LIST - 30TH EDITION

OTC DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 2 - February 2010

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CETIRIZINE HYDROCHLORIDE

SYRUP; ORAL

CHILDREN'S CETIRIZINE HYDROCHLORIDE ALLERGY

AUROBINDO PHARMA 5MG/5ML A090750 002 Feb 02, 2010 Jan NEWA

CHILDREN'S CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AUROBINDO PHARMA 5MG/5ML A090750 001 Feb 02, 2010 Jan NEWA

TABLET; ORAL

CETIRIZINE HYDROCHLORIDE ALLERGY

AMNEAL PHARMS NY 5MG A078780 001 Jan 21, 2010 Jan NEWA

10MG A078780 004 Jan 21, 2010 Jan NEWA

CETIRIZINE HYDROCHLORIDE HIVES RELIEF

AMNEAL PHARMS NY 5MG A078780 003 Jan 21, 2010 Jan NEWA

10MG A078780 002 Jan 21, 2010 Jan NEWA

CIMETIDINE

TABLET; ORAL

CIMETIDINE

CONTRACT PHARMACAL 200MG A074961 001 Jun 19, 1998 Jan CAHN

200MG A074963 001 Jun 19, 1998 Jan CAHN

IBUPROFEN

CAPSULE; ORAL

IBUPROFEN

+ CONTRACT PHARMACAL 200MG A074782 001 Jul 06, 1998 Jan CAHN

TABLET; ORAL

IBUPROFEN

CONTRACT PHARMACAL 200MG A071732 001 Sep 10, 1987 Jan CAHN

200MG A071735 001 Sep 10, 1987 Jan CAHN

200MG A072299 001 Jul 01, 1988 Jan CAHN

200MG A073691 001 Feb 25, 1994 Jan CAHN

PROFEN

CONTRACT PHARMACAL 200MG A071265 001 Oct 15, 1986 Jan CAHN

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET; ORAL

IBUPROFEN AND PSEUDOEPHEDRINE HYDROCHLORIDE

CONTRACT PHARMACAL 200MG;30MG A075588 001 Apr 08, 2002 Jan CAHN

LOPERAMIDE HYDROCHLORIDE

TABLET; ORAL

LOPERAMIDE HYDROCHLORIDE

CONTRACT PHARMACAL 2MG A073254 001 Jul 30, 1993 Jan CAHN

MICONAZOLE NITRATE

CREAM; VAGINAL

MICONAZOLE NITRATE

PERRIGO R AND D 4% A091366 001 Jan 15, 2010 Jan NEWA

NAPHAZOLINE HYDROCHLORIDE; PHENIRAMINE MALEATE

SOLUTION/DROPS; OPHTHALMIC

VISINE-A

+ JOHNSON AND JOHNSON 0.025%;0.3% N020485 001 Jan 31, 1996 Jan CRLD

PSEUDOEPHEDRINE HYDROCHLORIDETABLET, EXTENDED RELEASE; ORAL
SUDAFED 24 HOUR

>D>	+	ALZA	240MG	N020021	002	Dec 15, 1992	Feb	CAHN
>A>	+	MCNEIL CONS	240MG	N020021	002	Dec 15, 1992	Feb	CAHN

RANITIDINE HYDROCHLORIDE

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

CONTRACT PHARMACAL EQ 75MG BASE

A075094 001 Jun 21, 1999 Jan CAHN

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

CUMULATIVE SUPPLEMENT NUMBER 02 FEBRUARY 2010

NO FEBRUARY 2010 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 FEBRUARY 2010 ADDITIONS

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 2 - February 2010

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N022217 001					>A> NCE	Mar 05, 2012
<u>ALISKIREN HEMIFUMARATE; VALSARTAN - VALTURNA</u>						
N022217 002					>A> NCE	Mar 05, 2012
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u>						
N021881 001	7658914	Sep 01, 2024	DS DP			
<u>AZTREONAM - CAYSTON</u>						
N050814 001	>A> 7208141	Dec 20, 2021	DP U-1031			
	>A> 7214364	Dec 20, 2021	DP			
	>A> 7427633	Dec 20, 2021	DP U-1031			
<u>BALSALAZIDE DISODIUM - COLAZAL</u>						
N020610 001	7452872	Aug 24, 2026	U-141			
	7452872*PED	Feb 24, 2027				
	7625884	Aug 24, 2026	U-141			
	7625884*PED	Feb 24, 2027				
<u>BEXAROTENE - TARGRETIN</u>						
N021055 001	>A> 7655699	Apr 22, 2012	DS DP U-509			
<u>BEXAROTENE - TARGRETIN</u>						
N021056 001	>A> 7655699	Apr 22, 2012	DS DP U-510			
<u>BUDESONIDE - RHINOCORT</u>						
N020746 001	6686346	Apr 29, 2017	DP U-557	Y		
	6686346*PED	Oct 29, 2017				
	6986904	Apr 29, 2017	DP U-699	Y		
	6986904*PED	Oct 29, 2017				
<u>BUDESONIDE - RHINOCORT</u>						
N020746 002	6686346	Apr 29, 2017	DP U-557			
	6686346*PED	Oct 29, 2017				
	6986904	Apr 29, 2017	DP U-699			
	6986904*PED	Oct 29, 2017				
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N022108 001	7645802	Jun 27, 2026	DP			
	7649019	Jun 27, 2026	DP			
	>A> 7662407	Jun 27, 2026	DP			
	>A> 7671094	Jun 27, 2026	DP			
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N022108 002	7645802	Jun 27, 2026	DP			
	7649019	Jun 27, 2026	DP			
	>A> 7662407	Jun 27, 2026	DP			
	>A> 7671094	Jun 27, 2026	DP			
<u>BUPROPION HYDROBROMIDE - APLENZIN</u>						
N022108 003	7645802	Jun 27, 2026	DP			
	7649019	Jun 27, 2026	DP			
	>A> 7662407	Jun 27, 2026	DP			
	>A> 7671094	Jun 27, 2026	DP			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 2 - February 2010

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CAPSAICIN - QUTENZA</u>						
N022395	001				ODE	Nov 16, 2016
<u>CEFTIBUTEN DIHYDRATE - CEDAX</u>						
N050686	001	5599557	Feb 04, 2014	DP U-578		
		5599557	Feb 04, 2014	DP U-282		
<u>CEFTIBUTEN DIHYDRATE - CEDAX</u>						
N050686	002	5599557	Feb 04, 2014	DP U-578		
		5599557	Feb 04, 2014	DP U-282		
<u>CLINDAMYCIN PHOSPHATE; TRETINOIN - ZIANA</u>						
N050802	001	>A> RE41134	Feb 24, 2015	DP U-1033		
<u>CLONIDINE HYDROCHLORIDE - JENLOGA</u>						
N022331	001	5869100	Oct 13, 2013	DP		
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N022362	001	>A> 5607669	Jun 10, 2014	U-757		
		>A> 5607669*PED	Dec 10, 2014			
		>A> 5679717	Apr 29, 2014	U-757		
		>A> 5679717*PED	Oct 29, 2014			
		>A> 5693675	Dec 02, 2014	DS		
		>A> 5693675*PED	Jun 02, 2015			
		>A> 5917007	Apr 29, 2014	DS	U-757	
		>A> 5917007*PED	Oct 29, 2014			
		>A> 5919832	Apr 29, 2014	DS		
		>A> 5919832*PED	Oct 29, 2014			
		>A> 6066678	Apr 29, 2014	DS	U-757	
		>A> 6066678*PED	Oct 29, 2014			
		>A> 6433026	Apr 29, 2014	DS		
		>A> 6433026*PED	Oct 29, 2014			
		>A> 6784254	Apr 29, 2014	DS DP		
		>A> 6784254*PED	Oct 29, 2014			
		>A> 7101960	Apr 29, 2014	DS DP	U-757	
		>A> 7101960*PED	Oct 29, 2014			
		>A> 7229613	Apr 17, 2022	U-493		
		>A> 7229613*PED	Oct 17, 2022			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 2 - February 2010

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE (S)	EXCLUSIVITY EXPIRATION DATE
<u>COLESEVELAM HYDROCHLORIDE - WELCHOL</u>						
N022362 002	>A> 5607669	Jun 10, 2014	U-757			
	>A> 5607669*PED	Dec 10, 2014				
	>A> 5679717	Apr 29, 2014	U-757			
	>A> 5679717*PED	Oct 29, 2014				
	>A> 5693675	Dec 02, 2014	DS			
	>A> 5693675*PED	Jun 02, 2015				
	>A> 5917007	Apr 29, 2014	DS U-757			
	>A> 5917007*PED	Oct 29, 2014				
	>A> 5919832	Apr 29, 2014	DS			
	>A> 5919832*PED	Oct 29, 2014				
	>A> 6066678	Apr 29, 2014	DS U-757			
	>A> 6066678*PED	Oct 29, 2014				
	>A> 6433026	Apr 29, 2014	DS			
	>A> 6433026*PED	Oct 29, 2014				
	>A> 6784254	Apr 29, 2014	DS DP			
	>A> 6784254*PED	Oct 29, 2014				
	>A> 7101960	Apr 29, 2014	DS DP U-757			
	>A> 7101960*PED	Oct 29, 2014				
	>A> 7229613	Apr 17, 2022	U-493			
	>A> 7229613*PED	Oct 17, 2022				
<u>DALFAMPRIDINE - AMPYRA</u>						
N022250 001	>A> 5370879	Dec 06, 2011	DP		NCE	Jan 22, 2015
	>A> 5540938	Jul 30, 2013	U-1030		>A> ODE	Jan 22, 2017
<u>DICLOFENAC POTASSIUM - CAMBIA</u>						
N022165 001					>A> NDF	Jun 17, 2012
<u>DICLOFENAC POTASSIUM - ZIPSOR</u>						
N022202 001	6365180	Jul 15, 2019	DP U-980			
	>A> 7662858	Feb 24, 2029	U-1035			
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>						
N021427 003	5023269	Jun 11, 2013	DS DP U-797			
	5508276	Jul 18, 2014	DP			
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE</u>						
N021840 001	>A> 7615545	Jun 15, 2023	U-1			
<u>EZETIMIBE - ZETIA</u>						
N021445 001	7612058	Jan 25, 2022	U-1027			
	7612058*PED	Jul 25, 2022				
<u>FENOFIBRATE - FENOGLIDE</u>						
N022118 001	>A> 7658944	Dec 09, 2024	DP			
<u>FENOFIBRATE - FENOGLIDE</u>						
N022118 002	>A> 7658944	Dec 09, 2024	DP			
<u>HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM - HYZAAR</u>						
N020387 001	>A> 5608075****	Mar 04, 2009			Y	
	>A> 5608075*PED	Sep 04, 2009				

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 2 - February 2010

See List footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	PATENT DELIST REQUESTED	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>HYDROCHLOROTHIAZIDE; LOSARTAN POTASSIUM - HYZAAR</u>						
N020387 002	>A> 5608075****	Mar 04, 2009		Y		
	>A> 5608075*PED	Sep 04, 2009				
<u>HYDROCHLOROTHIAZIDE; OLMESARTAN MEDOXOMIL - BENICAR HCT</u>						
N021532 002	>A> 6878703	Nov 19, 2021	U-3	Y		
<u>IMIQUIMOD - IMIQUIMOD</u>						
A078548 001					>A> PC	Aug 24, 2010
<u>INSULIN GLULISINE RECOMBINANT - APIDRA SOLOSTAR</u>						
N021629 003	6221633	Jun 18, 2018	DS DP U-471			
	6960561	Jan 25, 2023	DP U-471			
	7452860	Mar 22, 2022	DP			
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N022115 001					>A> I-622 NDF PED	Jan 29, 2013 May 29, 2012 Nov 29, 2012
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N022115 002					>A> I-622 NDF PED	Jan 29, 2013 May 29, 2012 Nov 29, 2012
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N022115 003					>A> I-622 NDF PED	Jan 29, 2013 May 29, 2012 Nov 29, 2012
<u>LAMOTRIGINE - LAMICTAL XR</u>						
N022115 004					>A> I-622 NDF PED	Jan 29, 2013 May 29, 2012 Nov 29, 2012
<u>LAPATINIB DITOSYLATE - TYKERB</u>						
N022059 001					I-620	Jan 29, 2013
<u>LEVETIRACETAM - KEPPRA</u>						
N021872 001					I-563 PED	Mar 19, 2010 Sep 19, 2010
<u>LIRAGLUTIDE RECOMBINANT - VICTOZA</u>						
N022341 001	6268343	Aug 22, 2017	DS DP U-968		NCE	Jan 25, 2015
	6458924	Aug 22, 2017	DS DP U-968			
	7235627	Aug 22, 2017	DS DP			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N021977 001	7655630	Feb 24, 2023	DS			
	>A> 7659253	Feb 24, 2023	DS DP U-727			
	>A> 7659254	Feb 24, 2023	U-1034			
	>A> 7662787	Feb 24, 2023	DS			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N021977 002	7655630	Feb 24, 2023	DS			
	>A> 7659253	Feb 24, 2023	DS DP U-727			
	>A> 7659254	Feb 24, 2023	U-1034			
	>A> 7662787	Feb 24, 2023	DS			

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<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N021977 003	7655630	Feb 24, 2023	DS			
	>A> 7659253	Feb 24, 2023	DS DP U-727			
	>A> 7659254	Feb 24, 2023	U-1034			
	>A> 7662787	Feb 24, 2023	DS			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N021977 004	7655630	Feb 24, 2023	DS			
	>A> 7659253	Feb 24, 2023	DS DP U-727			
	>A> 7659254	Feb 24, 2023	U-1034			
	>A> 7662787	Feb 24, 2023	DS			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N021977 005	7655630	Feb 24, 2023	DS			
	>A> 7659253	Feb 24, 2023	DS DP U-727			
	>A> 7659254	Feb 24, 2023	U-1034			
	>A> 7662787	Feb 24, 2023	DS			
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>						
N021977 006	7655630	Feb 24, 2023	DS			
	>A> 7659253	Feb 24, 2023	DS DP U-727			
	>A> 7659254	Feb 24, 2023	U-1034			
	>A> 7662787	Feb 24, 2023	DS			
<u>LOSARTAN POTASSIUM - COZAAR</u>						
N020386 001	>A> 5608075****	Mar 04, 2009			Y	
	>A> 5608075*PED	Sep 04, 2009				
<u>LOSARTAN POTASSIUM - COZAAR</u>						
N020386 002	>A> 5608075****	Mar 04, 2009			Y	
	>A> 5608075*PED	Sep 04, 2009				
<u>LOSARTAN POTASSIUM - COZAAR</u>						
N020386 003	>A> 5608075****	Mar 04, 2009			Y	
	>A> 5608075*PED	Sep 04, 2009				
<u>MESALAMINE - SFROWASA</u>						
N019618 002	7645801	Jul 24, 2027	DS DP			
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.83% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N019734 004	>A> 7659291	Apr 18, 2027			U-1029	
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 0.86% SODIUM CHLORIDE IN PLASTIC CONTAINER</u>						
N019734 003	>A> 7659291	Apr 18, 2027			U-1029	
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 4.8% DEXTROSE IN PLASTIC CONTAINER</u>						
N019734 002	>A> 7659291	Apr 18, 2027			U-1029	
<u>NICARDIPINE HYDROCHLORIDE - CARDENE IN 5.0% DEXTROSE IN PLASTIC CONTAINER</u>						
N019734 005	>A> 7659291	Apr 18, 2027			U-1029	
<u>OLMESARTAN MEDOXOMIL - BENICAR</u>						
N021286 001					>A> NPP	Feb 04, 2013
					>A> PED	Aug 04, 2010
<u>OLMESARTAN MEDOXOMIL - BENICAR</u>						
N021286 003					>A> NPP	Feb 04, 2013
					>A> PED	Aug 04, 2010

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<u>OLMESARTAN MEDOXOMIL - BENICAR</u>						
N021286	004				>A> NPP >A> PED	Feb 04, 2013 Aug 04, 2010
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N021087	001				>A> M-90	Feb 22, 2013
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N021087	002				>A> M-90	Feb 22, 2013
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N021087	003				>A> M-90	Feb 22, 2010
<u>OSELTAMIVIR PHOSPHATE - TAMIFLU</u>						
N021246	001				>A> M-90	Feb 22, 2013
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N022210	001	>A> 7658918	Feb 20, 2028	DP		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N022210	002	>A> 7658918	Feb 20, 2028	DP		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N022210	003	>A> 7658918	Feb 20, 2028	DP		
<u>PANCRELIPASE (AMYLASE;LIPASE;PROTEASE) - ZENPEP</u>						
N022210	004	>A> 7658918	Feb 20, 2028	DP		
<u>PANTOPRAZOLE SODIUM - PROTONIX</u>						
N022020	001	7544370	Jun 07, 2026	DP		
		7544370*PED	Dec 07, 2026			
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421	001	>A> 4886812	Oct 08, 2010	DS DP		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421	002	>A> 4886812	Oct 08, 2010	DS DP		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421	003	>A> 4886812	Oct 08, 2010	DS DP		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421	004	>A> 4886812	Oct 08, 2010	DS DP		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX ER</u>						
N022421	005	>A> 4886812	Oct 08, 2010	DS DP		
<u>PRAMIPEXOLE DIHYDROCHLORIDE - PRAMIPEXOLE DIHYDROCHLORIDE</u>						
A077724	001				PC	Jul 03, 2010
<u>PRAMIPEXOLE DIHYDROCHLORIDE - PRAMIPEXOLE DIHYDROCHLORIDE</u>						
A077724	002				PC	Jul 03, 2010
<u>PRAMIPEXOLE DIHYDROCHLORIDE - PRAMIPEXOLE DIHYDROCHLORIDE</u>						
A077724	003				PC	Jul 03, 2010
<u>PRAMIPEXOLE DIHYDROCHLORIDE - PRAMIPEXOLE DIHYDROCHLORIDE</u>						
A077724	004				PC	Jul 03, 2010
<u>PRAMIPEXOLE DIHYDROCHLORIDE - PRAMIPEXOLE DIHYDROCHLORIDE</u>						
A077724	005				PC	Jul 03, 2010

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<u>PREGABALIN - LYRICA</u>						
N022488 001	5563175	Oct 08, 2013	U-661		I-535	Jun 21, 2010
	6001876	Dec 30, 2018	U-819			
	6001876	Dec 30, 2018	U-55			
	6197819	Dec 30, 2018	DS DP			
<u>REGADENOSON - LEXISCAN</u>						
N022161 001	7655636	Jun 22, 2019	U-869			
	7655637	Jun 22, 2019	DS DP U-869			
<u>RITONAVIR - RITONAVIR</u>						
N022417 001	>A> 5541206	Jul 30, 2013	DS DP U-688			
	>A> 5541206*PED	Jan 30, 2014				
	>A> 5635523	Jun 03, 2014	U-688			
	>A> 5635523*PED	Dec 03, 2014				
	>A> 5648497	Jul 15, 2014	DS			
	>A> 5648497*PED	Jan 15, 2015				
	>A> 5674882	Oct 07, 2014	U-688			
	>A> 5674882*PED	Apr 07, 2015				
	>A> 6037157	Jun 26, 2016	U-688			
	>A> 6037157*PED	Dec 26, 2016				
	>A> 6703403	Jun 26, 2016	U-688			
	>A> 6703403*PED	Dec 26, 2016				
	>A> 7148359	Jul 19, 2019	DP			
	>A> 7148359*PED	Jan 19, 2020				
	>A> 7364752	Nov 10, 2020	DP U-688			
	>A> 7364752*PED	May 10, 2021				
<u>ROMIDEPSIN - ISTODAX</u>						
N022393 001					ODE	Nov 05, 2016
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N021366 002	>A> 7030152	Apr 02, 2018	U-1032		>A> I-621	Feb 08, 2013
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N021366 003	>A> 7030152	Apr 02, 2018	U-1032		>A> I-621	Feb 08, 2013
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N021366 004	>A> 7030152	Apr 02, 2018	U-1032		>A> I-621	Feb 08, 2013
<u>ROSUVASTATIN CALCIUM - CRESTOR</u>						
N021366 005	>A> 7030152	Apr 02, 2018	U-1032		>A> I-621	Feb 08, 2013
<u>SODIUM OXYBATE - XYREM</u>						
N021196 001	>A> 7668730	Mar 07, 2024	U-1028			
<u>TIOTROPIUM BROMIDE MONOHYDRATE - SPIRIVA</u>						
N021395 001	7642268	Sep 24, 2021	DS DP			
<u>TOLTERODINE TARTRATE - DETROL</u>						
N020771 001	5559269	Nov 05, 2013	U-318	Y		
	5559269*PED	May 05, 2014				
<u>TOLTERODINE TARTRATE - DETROL</u>						
N020771 002	5559269	Nov 05, 2013	U-318	Y		
	5559269*PED	May 05, 2014				

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<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N022411 001	>A> 6607748	Jun 29, 2020	DP		>A> NDF	Feb 02, 2013
<u>TRAZODONE HYDROCHLORIDE - OLEPTRO</u>						
N022411 002	>A> 6607748	Jun 29, 2020	DP		>A> NDF	Feb 02, 2013
<u>VELAGLUCERASE ALFA - VPRIV</u>						
N022575 001					>A> NCE	Feb 26, 2015
<u>VELAGLUCERASE ALFA - VPRIV</u>						
N022575 002					>A> NCE	Feb 26, 2015
<u>VORINOSTAT - ZOLINZA</u>						
N021991 001	7652069	Mar 04, 2023	DS			

Footnote:

1. Patents are published upon receipt by the Orange Book Staff and may not reflect the official receipt date as described in 21 CFR 314.53(d)(5).
2. Patents 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.
3. **** The expiration date for U.S. Patent No. 5,608,075 is March 4, 2009.

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, 29th 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>