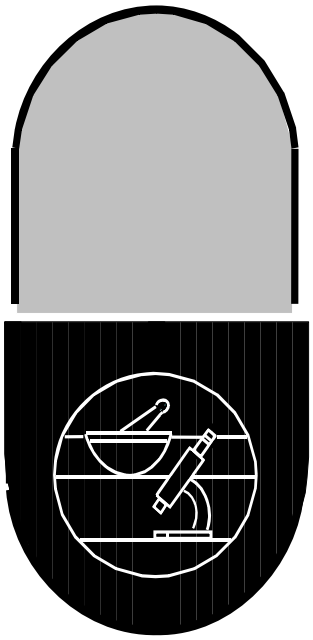
CUMULATIVE SUPPLEMENT 12
December 2006



APPROVED DRUG PRODUCTS

WITH THERAPEUTIC EQUIVALENCE EVALUATIONS

26th 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

26th EDITION

Cumulative Supplement 12

December 2006

CONTENTS

PAGE

APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS

26th EDITION

CUMULATIVE SUPPLEMENT 12
December 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are for exportation, are for military use, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.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.
- All product changes received and processed as of the date of publication.
  - o Refer to CS Section 1.8 Cumulative Supplement Legend for types of changes
  - o 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.

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.

FORMER APPLICANT NAME (FORMER ABBREVIATED NAME)	NEW APPLICANT NAME (NEW ABBREVIATED NAME)
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	PERRIGO NEW YORK INC

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

1.4 AVAILABILITY OF THE EDITION

Since 1997, the Electronic Orange Book Query (EOBQ) http://www.fda.gov/cder/ob/default.htm, 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 Annual Edition. 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://www.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/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 provided in eobzip.exe and 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 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

CATEGORIES COUNTED	MAR 2006	JUN 2006	SEPT 2006	DEC 2006
DRUG PRODUCTS LISTED	11487	11636	11741	11896
SINGLE SOURCE	2461	2447	2456	2471
	(21.4%)	(21.0%)	(20.9%)	(20.8%)
MULTISOURCE	8937	9000	9196	9336
	(77.8%)	(78.2%)	(78.3%)	(78.5%)
THERAPEUTICALLY EQUIVALENT	8730	8900	8997	9139
	(76.0%)	(76.5%)	(76.6%)	(76.8%)
NOT THERAPEUTICALLY EQUIVALENT	207	200	199	197
	(1.8%)	(1.7%)	(1.7%)	(1.7%)
EXCEPTIONS[1]	89	89	89	89
	(0.8%)	(0.8%)	(0.8%)	(0.7%)
NEW MOLECULAR ENTITIES APPROVED	6	8	1	10
NUMBER OF APPLICANTS	629	642	649	666

[1]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 were 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 remained 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 were triggered and run. For additional information on this matter, please refer to Docket Nos. 2005P-0008 and 2005P-0046. Patents were subsequently delisted in the December 2006 Orange Book update as the exclusivity periods have triggered and run to expiration.

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.

Levothyroxine Sodium (Mylan ANDA 76187) tablets have been determined to be therapeutically equivalent to corresponding strengths of Levothroid (Lloyd NDA 021116) 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.

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
LEVOTHROID	LLOYD	0.025MG	AB4	21116	001
LEVOTHYROXINE SODIUM**	MYLAN	0.025MG	AB4	76187	001
NOVOTHYROX	GENPHARM	0.025MG	BX	21292	001
LEVOLET	VINTAGE PHARMS	0.025MG	BX	21137	001

*Revised September 2006 **Revised December 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 proceeded 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.

ABARELIX

INJECTABLE; INTRAMUSCULAR

		PLENAXIS						
	@	PRAECIS	100MG/VIAL	N21320	001	Nov 25, 2003	Sep	DISC

ACEBUTOLOL HYDROCHLORIDE

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

ACETAMINOPHEN; BUTALBITAL

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

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

TABLET; ORAL

		ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE						
AB	+	MIKART	712.8MG;60MG;32MG	N40316	001	Apr 28, 1999	Sep	CFTG
		ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITATRATE						
AB		WEST WARD	712.8MG;60MG;32MG	N40637	001	Sep 22, 2006	Sep	NEWA

ACETAMINOPHEN; CODEINE PHOSPHATE

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

ACETAMINOPHEN; HYDROCODONE BITARTRATE

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
AA			650MG;10MG	N40757	001	Aug 25, 2006	Aug	NEWA
AA			750MG;7.5MG	N40769	001	Aug 28, 2006	Aug	NEWA

	TABLET; ORAL							
	HYDROCODONE BITARTRATE AND ACETAMINOPHEN							
	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							
	OXYCODONE AND ACETAMINOPHEN							
AA	MALLINCKRODT	325MG/5ML;5MG/5ML	N40680	001	Sep 29, 2006	Sep	NEWA	
	ROXICET							
AA +	ROXANE	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

MIOCHOL

@	NOVARTIS	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
>A> AA	VINTAGE	EQ 2MG BASE/5ML	N78105	001	Dec 27, 2006	Dec	NEWA

TABLET, EXTENDED RELEASE; ORAL

VOSPIRE ER

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

ALBUTEROL SULFATE; IPRATROPIUM BROMIDE

SOLUTION; INHALATION

>A>			ALBUTEROL SULFATE AND IPRATROPIUM BROMIDE						
>A>	AN		SANDOZ	EQ 0.083% BASE;0.017%	N76867	001	Dec 21, 2006	Dec	NEWA
			DUONEB						
>D>		+	DEY	EQ 0.083% BASE;0.017%	N20950	001	Mar 21, 2001	Dec	CFTG
>A>	AN	+		EQ 0.083% BASE;0.017%	N20950	001	Mar 21, 2001	Dec	CFTG

ALITRETINOIN

GEL; TOPICAL

			PANRETIN						
		+	EISAI MEDCL RES	EQ 0.1% BASE	N20886	001	Feb 02, 1999	Oct	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
AA	+	VINTAGE	50MG/5ML	N77992	001	Dec 12, 2006	Nov	NEWA

AMCINONIDE

CREAM; TOPICAL
AMCINONIDE

AB	+	ALTANA	0.1%	N76065	001	May 15, 2003	Sep	CRLD

CYCLOCORT

		@ ASTELLAS	0.1%	N18116	002		Sep	DISC

LOTION; TOPICAL
AMCINONIDE

	+	ALTANA	0.1%	N76329	001	Nov 06, 2002	Sep	CRLD

CYCLOCORT

		@ ASTELLAS	0.1%	N19729	001	Jun 13, 1988	Sep	DISC

OINTMENT; TOPICAL
AMCINONIDE

AB	+	ALTANA	0.1%	N76096	001	Nov 19, 2002	Sep	CRLD

CYCLOCORT

		@ ASTELLAS	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
>D> AP	+	APOTEX INC	50MG/ML	N77161	001	Apr 20, 2005	Dec	CAHN
>A> AP	+	GLAND PHARMA LTD	50MG/ML	N77161	001	Apr 20, 2005	Dec	CAHN

AMITRIPTYLINE HYDROCHLORIDE; PERPHENAZINE

TABLET; ORAL
PERPHENAZINE AND AMITRIPTYLINE HYDROCHLORIDE

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

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
>A>	AB	AUROBINDO	200MG/5ML	N65334	001	Dec 28, 2006	Dec	NEWA
>A>	AB		400MG/5ML	N65334	002	Dec 28, 2006	Dec	NEWA
	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

		TABLET; ORAL							
		DEXTROAMP SACCHARATE, AMP ASPARTATE, DEXTROAMP SULFATE AND AMP SULFATE							
	AB	TEVA PHARMS	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
		LOTION; TOPICAL							
		FUNGIZONE							
		@ APOTHECON	3%	N60570	001		Nov	DISC
		AMPICILLIN SODIUM; SULBACTAM SODIUM							
		INJECTABLE; INJECTION							
		AMPICILLIN AND SULBACTAM							
	AP	INSTITUTO BIOCHIMICO	EQ 10GM BASE/VIAL;EQ 5GM BASE/VIAL	N65314	001	Nov 27, 2006	Nov	NEWA
	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
>D>	AB	CLONMEL	EQ 250MG BASE	N62883	001	Feb 25, 1988	Dec	CTEC
>A>			EQ 250MG BASE	N62883	001	Feb 25, 1988	Dec	CTEC
>D>	AB		EQ 500MG BASE	N62882	001	Feb 25, 1988	Dec	CRLD
>A>		+	EQ 500MG BASE	N62882	001	Feb 25, 1988	Dec	CRLD
>D>		PRINCIPEN						
>D>	AB	APOTHECON	EQ 250MG BASE	N61392	001		Dec	DISC
>A>		@	EQ 250MG BASE	N61392	001		Dec	DISC
		@	EQ 250MG BASE	N62888	001	Mar 04, 1988	Nov	DISC
>D>	AB	+	EQ 500MG BASE	N61392	002		Dec	DISC
>A>		@	EQ 500MG BASE	N61392	002		Dec	DISC
	AB	+	EQ 500MG BASE	N61392	002		Nov	CRLD
		@	EQ 500MG BASE	N62888	002	Mar 04, 1988	Nov	DISC
		FOR SUSPENSION; ORAL							
		AMPICILLIN TRIHYDRATE							
		@ AM ANTIBIOTICS	EQ 125MG BASE/5ML	N61601	001		Jan	CAHN
		@	EQ 250MG BASE/5ML	N61601	002		Jan	CAHN
		PRINCIPEN							
>D>		APOTHECON	EQ 100MG BASE/ML	N61394	001		Dec	DISC
>A>		@	EQ 100MG BASE/ML	N61394	001		Dec	DISC
		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	
+	100MG/VIAL	N21632	002	Nov 14, 2006	Nov	NEWA	

ANISINDIONE

TABLET; ORAL						
MIRADON						
@ SCHERING	50MG	N10909	003		Jan	DISC

APOMORPHINE HYDROCHLORIDE

INJECTABLE; SUBCUTANEOUS						
APOKYN						
@ VERNALIS	20MG/2ML (10MG/ML)	N21264	001	Apr 20, 2004	Oct	DISC

APREPITANT

CAPSULE; ORAL						
EMEND						
MERCK	40MG	N21549	003	Jun 30, 2006	Jun	NEWA

ARFORMOTEROL TARTRATE

SOLUTION; INHALATION						
BROVANA						
+ SEPRACOR	EQ 0.015MG BASE/2ML	N21912	001	Oct 06, 2006	Oct	NEWA

ARIPIPRAZOLE

INJECTABLE; INTRAMUSCULAR						
ABILIFY						
+ 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
+	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;0.14MG/VIAL;17MG/VIAL;1MG/VIAL;1.4MG/VIAL;1.2MG/VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL	N21265	001	Feb 21, 2001	Jan	CAHN

INFUVITE PEDIATRIC (PHARMACY BULK PACKAGE)

+	SANDOZ	80MG/VIAL;0.02MG/VIAL;400 IU/VIAL;0.001MG/VIAL;5MG/VIAL;0.14MG/VIAL;17MG/VIAL;1MG/VIAL;1.4MG/VIAL;1.2MG/VIAL;7 IU/VIAL;2,300 IU/VIAL;0.2MG/VIAL	N21646	001	Jan 29, 2004	Jan	CAHN

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

FOR SOLUTION; ORAL

MOVIPREP

+	NORGINE B V	4.7GM;100GM;1.015GM;5.9GM;2.691GM;7.5GM	N21881	001	Aug 02, 2006	Aug	NEWA
+	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

FIORINAL W/CODEINE

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

ATAZANAVIR SULFATE

CAPSULE; ORAL

REYATAZ

	BRISTOL MYERS SQUIBB	EQ 200MG BASE	N21567	003	Jun 20, 2003	Oct	CRLD
+		EQ 300MG BASE	N21567	004	Oct 16, 2006	Oct	NEWA

ATENOLOL

TABLET; ORAL

ATENOLOL

>A>	AB	OHM LABS	25MG	N77877	001	Dec 27, 2006	Dec	NEWA
>A>	AB		50MG	N77877	002	Dec 27, 2006	Dec	NEWA
>A>	AB		100MG	N77877	003	Dec 27, 2006	Dec	NEWA

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

ATROPINE; PRALIDOXIME CHLORIDE

INJECTABLE; INTRAMUSCULAR
DUODOTE

+	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
AB	SANDOZ	EQ 100MG BASE/5ML	N65297	001	Sep 18, 2006	Sep	NEWA
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

TABLET; ORAL
AZITHROMYCIN

>A> AB	MYLAN	EQ 600MG BASE	N65360	001	Jan 08, 2007	Dec	NEWA

AZITHROMYCIN HYDROGENCITRATE

INJECTABLE; INJECTION
AZITHROMYCIN

>A>	+ SICOR PHARMS	EQ 500MG BASE/VIAL	N50809	001	Dec 19, 2006	Dec	NEWA
>A>	+	EQ 2.5GM BASE/VIAL	N50809	002	Dec 19, 2006	Dec	NEWA

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

TABLET; ORAL
BENAZEPRIL HYDROCHLORIDE

AB		APOTEX INC	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

BETHANECHOL CHLORIDE

TABLET; ORAL
BETHANECHOL CHLORIDE

AA	IMPAX LABS	5MG		N40739	001	Nov 01, 2006	Oct	NEWA
AA		10MG		N40741	001	Nov 01, 2006	Oct	NEWA
AA		25MG		N40740	001	Nov 01, 2006	Oct	NEWA
AA		50MG		N40721	004	Nov 01, 2006	Oct	NEWA

BEXAROTENE

CAPSULE; ORAL
TARGRETIN

+	EISAI MEDCL RES	75MG		N21055	001	Dec 29, 1999	Oct	CAHN

GEL; TOPICAL
TARGRETIN

+	EISAI MEDCL RES	1%		N21056	001	Jun 28, 2000	Oct	CAHN

BISKALCITRATE; METRONIDAZOLE; TETRACYCLINE

CAPSULE; ORAL
PYLERA

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

BISOPROLOL FUMARATE

TABLET; ORAL
BISOPROLOL FUMARATE

>A> AB	AUROBINDO PHARMA	5MG		N77910	001	Dec 27, 2006	Dec	NEWA
>A> AB		10MG		N77910	002	Dec 27, 2006	Dec	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

		INJECTABLE; INJECTION							
		BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE							
	AP	SEPTODONT	0.5%;0.0091MG/ML	N77250	001	Sep 27, 2006	Sep	NEWA	
		BUPIVACAINE HYDROCHLORIDE W/EPINEPHRINE							
	AP +	HOSPIRA	0.5%;0.0091MG/ML	N22046	001	Jul 13, 1983	Oct	CTEC	
	+		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	
		TABLET, EXTENDED RELEASE; ORAL							
		BUPROPION HYDROCHLORIDE							
>A>	AB3	ANCHEN PHARMS	150MG	N77284	001	Dec 14, 2006	Dec	NEWA	
>A>	AB3		300MG	N77284	002	Dec 14, 2006	Dec	NEWA	
>A>	AB3	IMPAX PHARMS	300MG	N77415	002	Dec 15, 2006	Dec	NEWA	
		WELLBUTRIN XL							
>D>	+	SMITHKLINE BEECHAM	150MG	N21515	001	Aug 28, 2003	Dec	CFTG	
>A>	AB3 +		150MG	N21515	001	Aug 28, 2003	Dec	CFTG	
>D>			300MG	N21515	002	Aug 28, 2003	Dec	CFTG	
>A>	AB3		300MG	N21515	002	Aug 28, 2003	Dec	CFTG	
		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							
		CAFCIT							
	AP +	MEAD JOHNSON	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N20793	001	Sep 21, 1999	Sep	CFTG	
		CAFFEINE CITRATE							
	AP	PHARMAFORCE	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N77233	001	Sep 21, 2006	Sep	NEWA	
		SOLUTION; ORAL							
		CAFCIT							
	AA +	MEAD JOHNSON	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N20793	002	Apr 12, 2000	Sep	CFTG	
		CAFFEINE CITRATE							
	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 FRESENIUS MEDCL 0.001MG/ML N75766 001 Feb 20, 2003 Nov CAHN
AP 0.002MG/ML N75766 002 Feb 20, 2003 Nov CAHN
AP GENIX THERAP 0.001MG/ML N77102 001 Feb 08, 2006 Jan NEWA

CALCIUM ACETATE

CAPSULE; ORAL
PHOSLO
@ FRESENIUS MEDCL EQ 84.5MG CALCIUM N21160 001 Apr 02, 2001 Nov CAHN
@ EQ 169MG CALCIUM N21160 002 Apr 02, 2001 Nov CAHN
PHOSLO GELCAPS
\+ FRESENIUS MEDCL EQ 169MG CALCIUM N21160 003 Apr 02, 2001 Nov CAHN

TABLET; ORAL
PHOSLO
@ FRESENIUS MEDCL EQ 169MG CALCIUM N19976 001 Dec 10, 1990 Nov CAHN
@ NABI EQ 169MG CALCIUM N19976 001 Dec 10, 1990 Jun DISC

CALCIUM CHLORIDE; DEXTROSE; LACTIC ACID; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE

INJECTABLE; INJECTION
PRISMASOL BGK 0/2.5 IN PLASTIC CONTAINER
\+ GAMBRO RENAL PRODS 3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML;3.05GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46GM/1000ML N21703 006 Oct 25, 2006 Oct NEWA
PRISMASOL BGK 2/0 IN PLASTIC CONTAINER
\+ GAMBRO RENAL PRODS N/A/1000ML;20GM/1000ML;5.4GM/1000ML;2.03GM/1000ML;0.157GM/1000ML;3.09GM/1000ML;6.46GM/1000ML N21703 002 Oct 25, 2006 Oct NEWA

INJECTABLE; INJECTION

PRISMASOL BGK 2/3.5 IN PLASTIC CONTAINER

+	GAMBRO RENAL PRODS	5.15GM/1000ML;20GM/1000ML;5.4GM/1000ML;2.03GM/1000ML;0.157GM/1000ML;3.09GM/1000ML;6.46GM/1000ML	N21703	003	Oct 25, 2006	Oct	NEWA

PRISMASOL BGK 4/0 IN PLASTIC CONTAINER

+	GAMBRO RENAL PRODS	N/A/1000ML;20GM/1000ML;5.4GM/1000ML;3.05GM/1000ML;0.314GM/1000ML;3.09GM/1000ML;6.46GM/1000ML	N21703	005	Oct 25, 2006	Oct	NEWA

PRISMASOL BGK 4/2.5 IN PLASTIC CONTAINER

+	GAMBRO RENAL PRODS	3.68GM/1000ML;20GM/1000ML;5.4GM/1000ML;3.05GM/1000ML;0.314GM/1000ML;3.09GM/1000ML;6.46GM/1000ML	N21703	004	Oct 25, 2006	Oct	NEWA

PRISMASOL BK 0/0 IN PLASTIC CONTAINER

+	GAMBRO RENAL PRODS	N/A/1000ML;N/A/1000ML;5.4GM/1000ML;3.05GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46GM/1000ML	N21703	007	Oct 25, 2006	Oct	NEWA

PRISMASOL BK 0/3.5 IN PLASTIC CONTAINER

+	GAMBRO RENAL PRODS	5.15GM/1000ML;N/A/1000ML;5.4GM/1000ML;2.03GM/1000ML;N/A/1000ML;3.09GM/1000ML;6.46GM/1000ML	N21703	001	Oct 25, 2006	Oct	NEWA

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE

SOLUTION; INTRAPERITONEAL

DIANEAL LOW CALCIUM W/ DEXTROSE 2.5% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	18.3MG/100ML;2.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML	N20183	002	Dec 04, 1992	Nov	CTEC

DIANEAL LOW CALCIUM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	18.3MG/100ML;3.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML	N20183	003	Dec 04, 1992	Nov	CTEC

DIANEAL LOW CALCIUM W/ DEXTROSE 4.25% IN PLASTIC CONTAINER

	BAXTER HLTHCARE	18.3MG/100ML;4.25GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML	N20183	004	Dec 04, 1992	Nov	CTEC

INPERSOL-LC/LM W/ DEXTROSE 3.5% IN PLASTIC CONTAINER

	FRESENIUS	18.4MG/100ML;3.5GM/100ML;5.08MG/100ML;538MG/100ML;448MG/100ML	N20374	003	Jun 13, 1994	Nov	CTEC

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

SOLUTION; IRRIGATION

BALANCED SALT SOLUTION

AT		AKORN	0.48MG/ML;0.3MG/ML;0.75MG/ML;3.9MG/ML;6.4MG/ML;1.7MG/ML	N75503	001	Sep 27, 2006	Sep	NEWA

BSS

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

CANDESARTAN CILEXETIL

TABLET; ORAL

ATACAND

	ASTRAZENECA	8MG	N20838	002	Jun 04, 1998	May	CRLD

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

@	APOTHECON	12.5MG	N74472	001	Mar 31, 1995	Nov	DISC
@		25MG	N74472	002	Mar 31, 1995	Nov	DISC
@		50MG	N74472	003	Mar 31, 1995	Nov	DISC
@		100MG	N74472	004	Mar 31, 1995	Nov	DISC
@	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

TABLET; ORAL
CAPTOPRIL

@	CLONMEL HLTHCARE	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		25MG;100MG	N74260	002	Sep 03, 1993	Jun	CAHN
AB		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		150MG/VIAL	N77383	002	Jan 27, 2006	Jan	NEWA
AP		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
AP		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
AP	DABUR ONCOLOGY PLC	EQ 450MG/45ML (10MG/ML)	N77432	003	Sep 29, 2006	Sep	NEWA
AP		EQ 50MG/5ML (10MG/ML)	N77432	001	Sep 29, 2006	Sep	NEWA
AP		EQ 150MG/15ML (10MG/ML)	N77432	002	Sep 29, 2006	Sep	NEWA

CARVEDILOL

TABLET; ORAL
COREG

	SMITHKLINE BEECHAM	3.125MG	N20297	004	May 29, 1997	Nov	CAHN
		6.25MG	N20297	003	Sep 14, 1995	Nov	CAHN
+		12.5MG	N20297	002	Sep 14, 1995	Nov	CAHN
		25MG	N20297	001	Sep 14, 1995	Nov	CAHN

CARVEDILOL PHOSPHATE

CAPSULE, EXTENDED RELEASE; ORAL

COREG CR

	SB PHARMCO	10MG	N22012	001	Oct 20, 2006	Oct	NEWA
		20MG	N22012	002	Oct 20, 2006	Oct	NEWA
		40MG	N22012	003	Oct 20, 2006	Oct	NEWA
+		80MG	N22012	004	Oct 20, 2006	Oct	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

CEFACLOR

TABLET, EXTENDED RELEASE; ORAL

CEFACLOR

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
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

AB	ORCHID HLTHCARE	EQ 250MG BASE/5ML	N65307	002	Oct 16, 2006	Oct	NEWA
AB		EQ 500MG BASE/5ML	N65307	003	Oct 16, 2006	Oct	NEWA
	+ 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
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

AP	ORCHID HLTHCARE	EQ 1GM BASE/VIAL	N65244	001	Aug 12, 2005	Nov	CPOT
AP		EQ 10GM BASE/VIAL	N65247	001	Aug 12, 2005	Nov	CPOT

CEFAZOLIN AND DEXTROSE

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

		INJECTABLE; INJECTION						
		CEFAZOLIN SODIUM						
	+	SAMSON MEDCL	EQ 100GM BASE/VIAL	N65141	001	Nov 29, 2006	Nov	NEWA
	+		EQ 300GM BASE/VIAL	N65141	002	Nov 29, 2006	Nov	NEWA
		CEFDINIR						
		CAPSULE; ORAL						
		CEFDINIR						
AB		LUPIN	300MG	N65264	001	May 19, 2006	May	NEWA
		OMNICEF						
AB	+	ABBOTT	300MG	N50739	001	Dec 04, 1997	May	CFTG
		FOR SUSPENSION; ORAL						
		CEFDINIR						
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
		CEFDITOREN PIVOXIL						
		TABLET; ORAL						
		SPECTRACEF						
>A>	+	CORNERSTONE	200MG	N21222	001	Aug 29, 2001	Dec	CAHN
>D>	+	PURDUE PHARMA LP	200MG	N21222	001	Aug 29, 2001	Dec	CAHN
		CEFOTAXIME SODIUM						
		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
		CEFOTETAN DISODIUM						
		INJECTABLE; INJECTION						
		CEFOTAN						
		@ ASTRAZENECA	EQ 1GM BASE/VIAL	N50588	001	Dec 27, 1985	Oct	DISC
		@	EQ 1GM BASE/VIAL	N63293	001	Apr 29, 1993	Jun	DISC
		@	EQ 2GM BASE/VIAL	N50588	002	Dec 27, 1985	Oct	DISC
		@	EQ 2GM BASE/VIAL	N63293	002	Apr 29, 1993	Jun	DISC
		@	EQ 10GM BASE/VIAL	N50588	003	Apr 25, 1988	Oct	DISC
		CEFOTAN IN PLASTIC CONTAINER						
		@ ASTRAZENECA	EQ 20MG BASE/ML	N50694	002	Jul 30, 1993	Oct	DISC
		@	EQ 40MG BASE/ML	N50694	001	Jul 30, 1993	Oct	DISC

CEFOXITIN SODIUM

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

CEFPROZIL

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

TABLET; ORAL

CEFPROZIL

AB	RANBAXY	250MG	N65198	001	Dec 13, 2006	Nov	NEWA
AB		500MG	N65198	002	Dec 13, 2006	Nov	NEWA

CEFTAZIDIME (ARGININE FORMULATION)

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

CEFTRIAXONE SODIUM

INJECTABLE; IM-IV

CEFTRIAXONE

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

CEFTRIAXONE 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

CEFTRIAXONE

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

CEFTRIAXONE 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

CEPHRADINE

>D> AB	TEVA	250MG	N62683	001	Jan 09, 1987	Dec	CTEC
>A>		250MG	N62683	001	Jan 09, 1987	Dec	CTEC
>D> AB		500MG	N62683	002	Jan 09, 1987	Dec	CRLD
>A> +		500MG	N62683	002	Jan 09, 1987	Dec	CRLD

>D> VELOSEF

>D> AB +	APOTHECON	500MG	N61764	002		Dec	DISC
>A> @		500MG	N61764	002		Dec	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

INJECTABLE; INJECTION
NESACAINE-MPF

AP	+	ABRAXIS BIOSCIENCE	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

CHLORPHENIRAMINE POLISTIREX; HYDROCODONE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL
TUSSIONEX PENNKINETIC

	+	UCB INC	EQ 8MG MALEATE/5ML;EQ 10MG BITARTRATE/5ML	N19111	001	Dec 31, 1987	Nov	CTNA

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

CICLESONIDE

SPRAY, METERED; NASAL
OMNARIS

	+	ALTANA PHARMA	0.05MG/INH	N22004	001	Oct 20, 2006	Oct	NEWA

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

SUSPENSION; TOPICAL
CICLOPIROX

>A>	AB		PERRIGO NEW YORK	0.77%	N77676	001	Dec 15, 2006	Dec	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

TABLET; ORAL
CILOSTAZOL

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
	@		1200MG/120ML (10MG/ML)	N19847	003	Dec 26, 1990	Aug	DISC
AP	+		200MG/20ML (10MG/ML)	N19847	002	Dec 26, 1990	Aug	CFTG

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	200MG/20ML (10MG/ML)	N76992	001	Aug 28, 2006	Aug	NEWA
AP			400MG/40ML (10MG/ML)	N76992	002	Aug 28, 2006	Aug	NEWA
			1200MG/120ML (10MG/ML)	N76993	001	Aug 28, 2006	Aug	NEWA
AP		HOSPIRA	200MG/20ML (10MG/ML)	N77245	001	Aug 28, 2006	Aug	NEWA
AP			400MG/40ML (10MG/ML)	N77245	002	Aug 28, 2006	Aug	NEWA
AP		SICOR PHARMS	400MG/40ML (10MG/ML)	N77782	002	Aug 28, 2006	Aug	NEWA
AP			200MG/20ML (10MG/ML)	N77782	001	Aug 28, 2006	Aug	NEWA

SOLUTION/DROPS; OPHTHALMIC
CIPROFLOXACIN

AT		NEXUS PHARMS	EQ 0.3% BASE	N77689	001	Dec 13, 2006	Nov	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
AB		INTERPHARM	EQ 10MG BASE	N77289	001	Nov 30, 2006	Nov	NEWA
AB			EQ 20MG BASE	N77289	002	Nov 30, 2006	Nov	NEWA
AB			EQ 40MG BASE	N77289	003	Nov 30, 2006	Nov	NEWA
AB		INVAGEN PHARMS	EQ 10MG BASE	N77534	001	Oct 03, 2006	Sep	NEWA

TABLET; ORAL
CITALOPRAM HYDROBROMIDE

AB	INVAGEN PHARMS	EQ 20MG BASE	N77534	002	Oct 03, 2006	Sep	NEWA
AB		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

TABLET, ORALLY DISINTEGRATING; ORAL
CITALOPRAM HYDROBROMIDE

@	BIOVAIL LABS INTL	EQ 10MG BASE	N21763	001	Dec 20, 2005	Oct	DISC
@		EQ 20MG BASE	N21763	002	Dec 20, 2005	Oct	DISC
@		EQ 40MG BASE	N21763	003	Dec 20, 2005	Oct	DISC

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

@	BREATHID 2006	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

CLINDAMYCIN PHOSPHATE; TRETINOIN

GEL; TOPICAL
ZIANA

+	MEDICIS	1.2%;0.025%	N50802	001	Nov 07, 2006	Nov	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

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

TABLET, ORALLY DISINTEGRATING; ORAL
FAZACLO ODT

+	AVANIR PHARMS	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 HYDROCHLORIDE,PHENYLEPHRINE HYDROCHLORIDE W/CODEINE PHOSPHATE

AA		VINTAGE	10MG/5ML;5MG/5ML;6.25MG/5ML	N40660	001	Dec 07, 2006	Nov	NEWA

PROMETH VC W/ CODEINE

AA	+	ACTAVIS MID ATLANTIC	10MG/5ML;5MG/5ML;6.25MG/5ML	N88764	001	Oct 31, 1984	Nov	CFTG
	+		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

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/PACKET	N17563	004	Sep 22, 1995	Apr	CFTG
AB			5GM/SCOOPFUL	N17563	003	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

TABLET; ORAL
COLESTID

AB	+	PHARMACIA AND UPJOHN	1GM	N20222	001	Jul 19, 1994	Oct	CFTG

COLESTIPOL HYDROCHLORIDE

AB		IMPAX LABS	1GM	N77510	001	Oct 24, 2006	Oct	NEWA

COPPER

INTRAUTERINE DEVICE; INTRAUTERINE
PARAGARD T 380A

	+	DURAMED PHARMS BARR	309MG/COPPER	N18680	001	Nov 15, 1984	Oct	CTNA

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

INJECTABLE; INJECTION

CYANOCOBALAMIN

@	ABRAXIS PHARM	0.1MG/ML	N80557	002		Nov	DISC
AP +	BIONICHE PHARMA	1MG/ML	N40451	001	Sep 23, 2003	Nov	CRLD
AP +	LUITPOLD	1MG/ML	N80737	001		Nov	CRLD
@	WATSON LABS	0.1MG/ML	N80573	002		Nov	DISC
@		1MG/ML	N80573	001		Nov	DISC

RUBRAMIN PC

@	BRISTOL MYERS SQUIBB	1MG/ML	N06799	010	Apr 28, 1988	Nov	DISC

VIBISONE

AP +	ABRAXIS PHARM	1MG/ML	N80557	003		Nov	CRLD

CYCLOBENZAPRINE HYDROCHLORIDE

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

AB	AMIDE PHARM	5MG	N77291	001	Feb 03, 2006	Jan	NEWA
AB	JUBILANT PHARMS	5MG	N77563	001	Apr 19, 2006	Apr	NEWA
AB		10MG	N77563	002	Apr 19, 2006	Apr	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

FLEXERIL

AB	MCNEIL CONS SPECLT	5MG	N17821	001		Jan	CFTG

CYPROHEPTADINE HYDROCHLORIDE

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 HYDROCLORIDE

AA	STASON PHARMS	4MG	N40644	001	May 30, 2006	May	NEWA

DAPTOMYCIN

INJECTABLE; IV (INFUSION)
CUBICIN

	@ CUBIST	250MG/VIAL	N21572	001	Sep 12, 2003	Jun		DISC

DARUNAVIR ETHANOLATE

TABLET; ORAL
PREZISTA

+	TIBOTEC	EQ 300MG BASE	N21976	001	Jun 23, 2006	Jun		NEWA

DASATINIB

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

DECITABINE

INJECTABLE; INTRAVENOUS
DACOGEN

+	MGI PHARMA INC	50MG/VIAL	N21790	001	May 02, 2006	May		NEWA

DEFEROXAMINE MESYLATE

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

DEMECLOCYCLINE HYDROCHLORIDE

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

TABLET; ORAL
DESMOPRESSIN ACETATE

AB		TEVA PHARMS	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

AEROSOL, FOAM; TOPICAL
VERDESO

	+	CONNETICS	0.05%	N21978	001	Sep 19, 2006	Sep	NEWA

GEL; TOPICAL
DESONATE

	+	SKINMEDICA	0.05%	N21844	001	Oct 20, 2006	Oct	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	Oct	CRLD
AP			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	N88762	001	Oct 31, 1984	Jul	CAHN

PROMETHAZINE DM

AA		VINTAGE	15MG/5ML;6.25MG/5ML	N40649	001	Feb 14, 2006	Jan	NEWA

DIAZEPAM

TABLET; ORAL

DIAZEPAM

AB		ACTAVIS ELIZABETH	2MG	N70781	001	Mar 19, 1986	Jun	CAHN
AB			5MG	N70706	001	Mar 19, 1986	Jun	CAHN
AB			10MG	N70707	001	Mar 19, 1986	Jun	CAHN
AB		VINTAGE PHARMS	2MG	N77749	001	Mar 31, 2006	Mar	NEWA
AB			5MG	N77749	002	Mar 31, 2006	Mar	NEWA
AB			10MG	N77749	003	Mar 31, 2006	Mar	NEWA

DIAZOXIDE

INJECTABLE; INJECTION

HYPERSTAT

		@ SCHERING	15MG/ML	N16996	001		Oct	DISC

DICLOFENAC SODIUM

TABLET, DELAYED RELEASE; ORAL

DICLOFENAC SODIUM

AB		ACTAVIS ELIZABETH	50MG	N74514	001	Mar 26, 1996	Jun	CAHN
AB			75MG	N74514	002	Mar 26, 1996	Jun	CAHN
AB		SANDOZ	75MG	N74394	001	Nov 30, 1995	May	CRLD

VOLTAREN

AB	+	NOVARTIS	75MG	N19201	003	Jul 28, 1988	May	CMFD

TABLET, EXTENDED RELEASE; ORAL

DICLOFENAC SODIUM

AB		ACTAVIS ELIZABETH	100MG	N75910	001	Jan 07, 2002	Jun	CAHN

DIETHYLPROPION HYDROCHLORIDE

TABLET; ORAL

DIETHYLPROPION HYDROCHLORIDE

		@ TG UNITED LABS	25MG	N88267	001	Aug 25, 1983	May	CAHN
		@	25MG	N88268	001	Aug 25, 1983	May	CAHN

TENUATE

>D>	+	SANOFI AVENTIS US	25MG	N11722	002		Dec	CAHN
>A>	+	WATSON PHARMS	25MG	N11722	002		Dec	CAHN

TABLET, EXTENDED RELEASE; ORAL

TENUATE DOSPAN

>D>	+	SANOFI AVENTIS US	75MG	N12546	001		Dec	CAHN
>A>	+	WATSON PHARMS	75MG	N12546	001		Dec	CAHN

DIFLORASONE DIACETATE

CREAM; TOPICAL

DIFLORASONE DIACETATE

BX	+	ALTANA	0.05%	N76263	001	Dec 20, 2002	Jan	CRLD

FLORONE

		@ PHARMACIA AND UPJOHN	0.05%	N17741	001		Jan	DISC

CREAM; TOPICAL
FLORONE E
@ PHARMACIA AND UPJOHN 0.05% 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

DILTIAZEM MALATE; ENALAPRIL MALEATE

TABLET, EXTENDED RELEASE; ORAL
TECZEM
@ BIOVAIL EQ 180MG HCL;5MG N20507 001 Oct 04, 1996 Oct DISC

DIMETHYL SULFOXIDE

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

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL
BENADRYL

>A>		@ MCNEIL CONS	25MG	N05845	007		Dec	CAHN
>A>		@	50MG	N05845	001		Dec	CAHN
>D>		@ PARKE DAVIS	25MG	N05845	007		Dec	CAHN
>D>		@	50MG	N05845	001		Dec	CAHN

ELIXIR; ORAL
BENADRYL

>A>		@ MCNEIL CONS	12.5MG/5ML	N05845	004		Dec	CAHN
>D>		@ PARKE DAVIS	12.5MG/5ML	N05845	004		Dec	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
	AB	GLENMARK PHARMA	25MG	N88999	001	Feb 05, 1991	Oct	CMFD
		@	25MG	N88999	001	Feb 05, 1991	Jul	CAHN
	AB		50MG	N89000	001	Feb 05, 1991	Jul	CAHN
	AB		75MG	N89001	001	Feb 05, 1991	Oct	CMFD
		@	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

DISULFIRAM

TABLET; ORAL
ANTABUSE

		ODYSSEY PHARMS	250MG	N88482	001	Dec 08, 1983	Oct	CRLD
	+		500MG	N88483	001	Dec 08, 1983	Oct	CMFD

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

INJECTABLE; INJECTION
DOPAMINE HYDROCHLORIDE

	@	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		MYLAN	EQ 50MG BASE	N65377	001	Nov 07, 2006	Oct	NEWA
AB			EQ 75MG BASE	N65377	002	Nov 07, 2006	Oct	NEWA
AB			EQ 100MG BASE	N65377	003	Nov 07, 2006	Oct	NEWA
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
AB			EQ 75MG BASE	N65356	002	May 31, 2006	May	NEWA
AB			EQ 100MG BASE	N65356	003	May 31, 2006	May	NEWA
AB		SANDOZ	EQ 50MG BASE	N65353	001	Nov 27, 2006	Nov	NEWA
AB			EQ 75MG BASE	N65353	002	Nov 27, 2006	Nov	NEWA
AB			EQ 100MG BASE	N65353	003	Nov 27, 2006	Nov	NEWA

DOXYCYCLINE HYCLATE

CAPSULE, DELAYED RELEASE; ORAL
DORYX

	@	FH FAULDING CO LTD	EQ 75MG BASE	N50582	002	Aug 13, 2001	Apr	DISC
	@		EQ 100MG BASE	N50582	001	Jul 22, 1985	Apr	DISC
	@	MAYNE PHARMA INTL	EQ 75MG BASE	N50582	002	Aug 13, 2001	Sep	CAHN
	@		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

INJECTABLE; INJECTION
DOXY 100

AP	+	ABRAXIS PHARM	EQ 100MG BASE/VIAL	N62475	001	Dec 09, 1983	Oct	CTEC

DOXYCYCLINE

AP	+	BEDFORD	EQ 100MG BASE/VIAL	N62569	001	Mar 09, 1988	Oct	CMFD

TABLET; ORAL
DOXYCYCLINE HYCLATE

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

TABLET, DELAYED RELEASE; ORAL
DORYX

	MAYNE PHARMA INTL	EQ 75MG BASE	N50795	001	May 06, 2005	Oct	CAHN
+		EQ 100MG BASE	N50795	002	May 06, 2005	Oct	CAHN

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
	@		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

INJECTABLE; IV (INFUSION)
EPIRUBICIN HYDROCHLORIDE

	+	MAYNE PHARMA USA	50MG/VIAL	N50807	001	Sep 15, 2006	Sep	NEWA
	+		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	EQ 5MG BASE	N76765	001	May 22, 2006	Oct	CPOT
AB			5MG	N76765	001	May 22, 2006	May	NEWA
AB			EQ 10MG BASE	N76765	002	May 22, 2006	Oct	CPOT
AB			10MG	N76765	002	May 22, 2006	May	NEWA
AB			EQ 20MG BASE	N76765	003	May 22, 2006	Oct	CPOT
AB			20MG	N76765	003	May 22, 2006	May	NEWA

LEXAPRO

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

ESOMEPRAZOLE MAGNESIUM

FOR SUSPENSION, DELAYED RELEASE; ORAL
NEXIUM

		ASTRAZENECA	EQ 20MG BASE/PACKET	N21957	001	Oct 20, 2006	Oct	NEWA
	+		EQ 40MG BASE/PACKET	N21957	002	Oct 20, 2006	Oct	NEWA

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
AB	+		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

		FILM, EXTENDED RELEASE; TRANSDERMAL							
		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	
>D>		GEL, METERED; TOPICAL							
>D>		ESTROGEL							
>D>	+	ASCEND	0.06%	N21166	002	Feb 09, 2004	Dec	CDFR	
	+		0.06%	N21166	002	Feb 09, 2004	Jan	CAHN	
		GEL, METERED; TRANSDERMAL							
>A>		ELESTRIN							
>A>	BX +	BIOSANTE	0.06%	N21813	001	Dec 15, 2006	Dec	NEWA	
>A>		ESTROGEL							
>A>	BX +	ASCEND	0.06%	N21166	002	Feb 09, 2004	Dec	CDFR	
		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	
		ESTRADIOL; NORETHINDRONE ACETATE							
		TABLET; ORAL							
		ACTIVELLA							
>A>		NOVO NORDISK INC	0.5MG;0.1MG	N20907	002	Dec 28, 2006	Dec	NEWA	
		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							
		EDECRIN							
	+	ATON	EQ 50MG BASE/VIAL	N16093	001		Jul	CAHN	
		ETHACRYNIC ACID							
		TABLET; ORAL							
		EDECRIN							
	+	ATON	25MG	N16092	001		Oct	CAHN	
		@	50MG	N16092	002		Oct	CAHN	

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL

QUASENSE

AB		WATSON LABS	0.03MG;0.15MG	N77101	001	Sep 06, 2006	Aug	NEWA

SEASONALE

AB	+	DURAMED	0.03MG;0.15MG	N21544	001	Sep 05, 2003	Aug	CFTG

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

TABLET; BUCCAL
FENTORA
CEPHALON EQ 0.1MG BASE N21947 001 Sep 25, 2006 Sep NEWA
EQ 0.2MG BASE N21947 002 Sep 25, 2006 Sep NEWA
+ EQ 0.4MG BASE N21947 003 Sep 25, 2006 Sep NEWA
EQ 0.6MG BASE N21947 004 Sep 25, 2006 Sep NEWA

TABLET; BUCCAL
FENTORA
CEPHALON 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
@ 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
CEPHALON EQ 0.2MG BASE N20747 001 Nov 04, 1998 Oct CTNA
+ EQ 0.4MG BASE N20747 002 Nov 04, 1998 Oct CTNA
EQ 0.6MG BASE N20747 003 Nov 04, 1998 Oct CTNA
EQ 0.8MG BASE N20747 004 Nov 04, 1998 Oct CTNA
EQ 1.2MG BASE N20747 005 Nov 04, 1998 Oct CTNA
EQ 1.6MG BASE N20747 006 Nov 04, 1998 Oct CTNA
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

SUSPENSION; ORAL
ALLEGRA
+ SANOFI AVENTIS US 30MG/5ML N21963 001 Oct 16, 2006 Oct NEWA

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
>A> AB GEDEON RICHTER USA 5MG N77251 001 Dec 22, 2006 Dec NEWA
AB IVAX PHARMS 5MG N76340 001 Jun 19, 2006 Jun NEWA
>A> AB MYLAN 5MG N77578 001 Dec 18, 2006 Dec NEWA
>A> AB TEVA 5MG N76511 001 Dec 15, 2006 Dec NEWA
PROPECIA
AB + MERCK 1MG N20788 001 Dec 19, 1997 Jul CFTG
PROSCAR
AB + MERCK 5MG N20180 001 Jun 19, 1992 Jun CFTG

FLUCONAZOLE

FOR SUSPENSION; ORAL
FLUCONAZOLE
>A> AB TARO PHARM INDS 50MG/5ML N76918 001 Dec 18, 2006 Dec NEWA
>A> AB 200MG/5ML N76918 002 Dec 18, 2006 Dec NEWA

TABLET; ORAL
FLUCONAZOLE
@ GEDEON RICHTER USA 50MG N76432 001 Jul 29, 2004 Nov DISC

TABLET; ORAL
FLUCONAZOLE

	@	GEDEON RICHTER USA	100MG	N76432	002	Jul 29, 2004	Nov	DISC
	@		150MG	N76432	003	Jul 29, 2004	Nov	DISC
	@		200MG	N76432	004	Jul 29, 2004	Nov	DISC
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

INJECTABLE; INJECTION
FLUOROURACIL

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
>D>		PROZAC						
>D>	AB	+ LILLY	EQ 10MG BASE	N20974	001	Mar 09, 1999	Dec	DISC
>A>		@	EQ 10MG BASE	N20974	001	Mar 09, 1999	Dec	DISC
		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

AEROSOL, METERED; INHALATION
FLOVENT

	@	GLAXOSMITHKLINE	0.044MG/INH	N20548	001	Mar 27, 1996	Oct	DISC
	@		0.11MG/INH	N20548	002	Mar 27, 1996	Oct	DISC
	@		0.22MG/INH	N20548	003	Mar 27, 1996	Oct	DISC

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
	@		300 IU/0.36ML	N21211	001	Mar 23, 2004	Jun	DISC

FORMOTEROL FUMARATE

POWDER; INHALATION
FORADIL CERTIHALER

>A>		NOVARTIS	EQ 0.0085MG BASE/INH	N21592	001	Dec 15, 2006	Dec	NEWA

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

AB	OHM LABS	20MG	N78010	001	Sep 18, 2006	Sep	NEWA
AB		40MG	N78010	002	Sep 18, 2006	Sep	NEWA
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
AB	APOTEX INC	100MG	N77894	001	Oct 10, 2006	Sep	NEWA
AB		300MG	N77894	002	Oct 10, 2006	Sep	NEWA
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
AB	IVAX PHARMS	100MG	N76017	001	Apr 28, 2004	Sep	CFTG
AB		300MG	N76017	002	Apr 28, 2004	Sep	CFTG
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

		INJECTABLE; INJECTION						
		OPTIMARK						
	+	MALLINCKRODT	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						
		INJECTABLE; INJECTION						
		GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER						
AP	+	B BRAUN	EQ 0.8MG BASE/ML	N62814	001	Aug 28, 1987	Nov	CRLD
AP	+		EQ 1.2MG BASE/ML	N62814	002	Aug 28, 1987	Nov	CRLD
AP	+		EQ 1.4MG BASE/ML	N62814	003	Aug 28, 1987	Nov	CRLD
AP	+		EQ 1.6MG BASE/ML	N62814	004	Aug 28, 1987	Nov	CRLD
AP	+		EQ 1.8MG BASE/ML	N62814	005	Aug 28, 1987	Nov	CRLD
AP	+		EQ 2MG BASE/ML	N62814	006	Aug 28, 1987	Nov	CRLD
AP	+		EQ 2.4MG BASE/ML	N62814	007	Aug 28, 1987	Nov	CRLD

INJECTABLE; INJECTION

GENTAMICIN SULFATE IN SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP	+	B BRAUN	EQ 40MG BASE/100ML	N62814	008	Aug 28, 1987	Nov	CRLD
AP	+		EQ 60MG BASE/100ML	N62814	009	Aug 28, 1987	Nov	CRLD
AP	+		EQ 70MG BASE/100ML	N62814	010	Aug 28, 1987	Nov	CRLD
AP	+		EQ 80MG BASE/100ML	N62814	011	Aug 28, 1987	Nov	CRLD
AP	+		EQ 90MG BASE/100ML	N62814	012	Aug 28, 1987	Nov	CRLD
AP	+		EQ 100MG BASE/100ML	N62814	013	Aug 28, 1987	Nov	CRLD
AP	+		EQ 120MG BASE/100ML	N62814	014	Aug 28, 1987	Nov	CRLD

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
	+		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
AB			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

>D>	AP	APOTEX INC	EQ 5MG BASE/ML	N76774	001	Aug 25, 2004	Dec	CAHN
>A>	AP	GLAND PHARMA LTD	EQ 5MG BASE/ML	N76774	001	Aug 25, 2004	Dec	CAHN
	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

>D> HALOTHANE

>D> LIQUID; INHALATION

>D> HALOTHANE

>D>	AN	HALOCARBON	99.99%	N80810	001		Dec	DISC
>A>		@	99.99%	N80810	001		Dec	DISC
		@ 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

TABLET; ORAL
TUSSIGON

>D>	AA		JONES PHARMA	1.5MG;5MG	N88508	001	Jul 30, 1985	Dec	CAHN
>A>	AA		KING PHARMS	1.5MG;5MG	N88508	001	Jul 30, 1985	Dec	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; LISINOPRIL

TABLET; ORAL
LISINOPRIL 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
AB			LUPIN	12.5MG;10MG	N77912	001	Sep 27, 2006	Sep	NEWA
AB				12.5MG;20MG	N77912	002	Sep 27, 2006	Sep	NEWA
AB				25MG;20MG	N77912	003	Sep 27, 2006	Sep	NEWA

HYDROCHLOROTHIAZIDE; METHYLDOPA

TABLET; ORAL

	ALDORIL 15							
	@ MERCK	15MG;250MG	N13402	001		Jun	DISC	
	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; TRIAMTERENE

TABLET; ORAL

	TRIAMTERENE AND HYDROCHLOROTHIAZIDE							
AB	APOTEX INC	25MG;37.5MG	N71251	002	May 05, 1998	Nov	CAHN	
AB		50MG;75MG	N71251	001	Apr 17, 1988	Nov	CAHN	

HYDROCHLOROTHIAZIDE; VALSARTAN

TABLET; ORAL

	DIOVAN HCT							
	NOVARTIS	12.5MG;320MG	N20818	004	Apr 28, 2006	Apr	NEWA	

TABLET; ORAL
DIOVAN HCT

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

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL
HYDROCODONE BITARTRATE AND IBUPROFEN

AB	+	INTERPHARM	5MG;200MG	N76642	002	Mar 18, 2004	Oct	CTEC
AB		VINTAGE PHARMS	5MG;200MG	N77727	001	Nov 06, 2006	Oct	NEWA
AB			7.5MG;200MG	N77723	001	Nov 06, 2006	Oct	NEWA
			10MG;200MG	N77723	002	Nov 06, 2006	Oct	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

PENECORT

		@ ALLERGAN HERBERT	1%	N88216	001	Jun 06, 1984	Oct	DISC

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

SOLUTION; TOPICAL
PENECORT

		@ ALLERGAN HERBERT	1%	N88214	001	Jun 06, 1984	Oct	DISC

TABLET; ORAL
CORTEF

		@ PHARMACIA AND UPJOHN	10MG	N08697	001		Jun	CTEC

HYDROCORTONE

		@ MERCK	10MG	N08506	007		Jun	DISC
		@	20MG	N08506	011		Jun	DISC

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION
A-HYDROCORT

AP		HOSPIRA	EQ 100MG BASE/VIAL	N40666	001	Apr 06, 2006	Mar	NEWA

HYDROCORTISONE; NEOMYCIN; POLYMYXIN B SULFATE

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

HYDROXOCOBALAMIN

INJECTABLE; INJECTION
CYANOKIT

>A>								
>A>	+	EMD PHARMS	2.5GM/VIAL (5GM/KIT)	N22041	002	Dec 15, 2006	Dec	NEWA

HYDROXYPROPYL CELLULOSE

INSERT; OPHTHALMIC
LACRISERT

	+	ATON	5MG	N18771	001		Oct	CAHN

HYDROXYZINE HYDROCHLORIDE

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

HYDROXYZINE PAMOATE

CAPSULE; ORAL

HYDROXYZINE PAMOATE

	BARR	EQ 100MG HCL	N88488	001	Jun 15, 1984	Mar	CTEC

VISTARIL

	@ PFIZER	EQ 100MG HCL	N11459	006		Mar	DISC

IBANDRONATE SODIUM

INJECTABLE; INTRAVENOUS

BONIVA

	+ ROCHE	EQ 3MG BASE/3ML	N21858	001	Jan 06, 2006	Jan	NEWA

IBUPROFEN

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
	@	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

IDARUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

IDARUBICIN HYDROCHLORIDE

>A>	AP BEDFORD LABS	1MG/ML	N65275	001	Dec 14, 2006	Dec	NEWA

IMATINIB MESYLATE

CAPSULE; ORAL

GLEEVEC

	@ NOVARTIS	EQ 50MG BASE	N21335	001	May 10, 2001	Nov	CPOT
	@	EQ 100MG BASE	N21335	002	May 10, 2001	Nov	CPOT

TABLET; ORAL

GLEEVEC

	NOVARTIS	EQ 100MG BASE	N21588	001	Apr 18, 2003	Nov	CPOT
	+	EQ 400MG BASE	N21588	002	Apr 18, 2003	Nov	CPOT

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
@		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

INDOMETHACIN

@	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

ISOTRETINOIN

CAPSULE; ORAL
CLARAVIS

AB		BARR	30MG	N76135	003	May 11, 2006	May	NEWA

CAPSULE; ORAL
SOTRET

AB		RANBAXY	30MG	N76503	001	Jun 20, 2003	May	CTEC

ISRADIPINE

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

IVERMECTIN

TABLET; ORAL
STROMECTOL

	+	MERCK	3MG	N50742	002	Oct 08, 1998	Jun	CRLD
	@		6MG	N50742	001	Nov 22, 1996	Jun	DISC

KETOCONAZOLE

GEL; TOPICAL
XOLEGEL

	+	BARRIER THERAP	2%	N21946	001	Jul 28, 2006	Jul	NEWA

KETOPROFEN

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

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION
KETOROLAC TROMETHAMINE

>D>	AP	APOTEX INC	15MG/ML	N76722	001	Jul 27, 2004	Dec	CAHN
>D>	AP		30MG/ML	N76722	002	Jul 27, 2004	Dec	CAHN
>A>	AP	GLAND PHARMA LTD	15MG/ML	N76722	001	Jul 27, 2004	Dec	CAHN
>A>	AP		30MG/ML	N76722	002	Jul 27, 2004	Dec	CAHN
	AP	SANDOZ	15MG/ML	N76271	001	Oct 06, 2004	Jan	CAHN
	AP		30MG/ML	N76271	002	Oct 06, 2004	Jan	CAHN

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC
KETOTIFEN FUMARATE

AT		APOTEX	EQ 0.025% BASE	N77354	001	May 09, 2006	Apr	NEWA
		APOTEX INC	EQ 0.025% BASE	N77354	001	May 09, 2006	Oct	CTEC

KUNECATECHINS

OINTMENT; TOPICAL
VEREGEN

	+	MEDIGENE	15%	N21902	001	Oct 31, 2006	Oct	NEWA

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 375 (COPACKAGED)

>D>								
>A>		@ TAP PHARM	15MG,N/A;N/A,375MG	N21507	003	Nov 14, 2003	Dec	DISC

CAPSULE, DELAYED REL PELLETS, TABLET; ORAL
PREVACID NAPRAPAC 250 (COPACKAGED)

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

PREVACID NAPRAPAC 375 (COPACKAGED)

>D>								
>D>		TAP PHARM	15MG,N/A;N/A,375MG	N21507	003	Nov 14, 2003	Dec	DISC
			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

CAPSULE; ORAL
TIROSINT

	INST BIOCHIMIQUE	0.025MG	N21924	002	Oct 13, 2006	Oct	NEWA
		0.05MG	N21924	003	Oct 13, 2006	Oct	NEWA
		0.075MG	N21924	004	Oct 13, 2006	Oct	NEWA
		0.1MG	N21924	005	Oct 13, 2006	Oct	NEWA
		0.125MG	N21924	006	Oct 13, 2006	Oct	NEWA
+		0.15MG	N21924	007	Oct 13, 2006	Oct	NEWA

TABLET; ORAL
LEVOTHROID

>D>	BX	LLOYD	0.025MG	N21116	001	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.025MG	N21116	001	Oct 24, 2002	Dec	CTEC
>D>	BX		0.05MG	N21116	002	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.05MG	N21116	002	Oct 24, 2002	Dec	CTEC
>D>	BX		0.075MG	N21116	003	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.075MG	N21116	003	Oct 24, 2002	Dec	CTEC
>D>	BX		0.088MG	N21116	010	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.088MG	N21116	010	Oct 24, 2002	Dec	CTEC
>D>	BX		0.1MG	N21116	004	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.1MG	N21116	004	Oct 24, 2002	Dec	CTEC
>D>	BX		0.112MG	N21116	011	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.112MG	N21116	011	Oct 24, 2002	Dec	CTEC
>D>	BX		0.125MG	N21116	005	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.125MG	N21116	005	Oct 24, 2002	Dec	CTEC
>D>	BX		0.137MG	N21116	012	Dec 07, 2004	Dec	CTEC
>A>	AB4		0.137MG	N21116	012	Dec 07, 2004	Dec	CTEC
>D>	BX		0.15MG	N21116	006	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.15MG	N21116	006	Oct 24, 2002	Dec	CTEC
>D>	BX		0.175MG	N21116	007	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.175MG	N21116	007	Oct 24, 2002	Dec	CTEC
>D>	BX		0.2MG	N21116	008	Oct 24, 2002	Dec	CTEC
>A>	AB4		0.2MG	N21116	008	Oct 24, 2002	Dec	CTEC
>D>	BX	+	0.3MG	N21116	009	Oct 24, 2002	Dec	CTEC
>A>	AB4	+	0.3MG	N21116	009	Oct 24, 2002	Dec	CTEC

TABLET; ORAL

LEVOTHYROXINE SODIUM

	AB2, AB3	GENPHARM	0.025MG	N76752	001	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.05MG	N76752	002	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.075MG	N76752	003	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.088MG	N76752	004	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.1MG	N76752	005	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.112MG	N76752	006	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.125MG	N76752	007	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.15MG	N76752	008	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.175MG	N76752	009	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.2MG	N76752	010	Jun 16, 2005	Sep	CTEC
	AB2, AB3		0.3MG	N76752	011	Jun 16, 2005	Sep	CTEC
>D>	AB1, AB2, AB3	MYLAN	0.025MG	N76187	001	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.025MG	N76187	001	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.05MG	N76187	002	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.05MG	N76187	002	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.075MG	N76187	003	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.075MG	N76187	003	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.088MG	N76187	004	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.088MG	N76187	004	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.1MG	N76187	005	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.1MG	N76187	005	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.112MG	N76187	006	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.112MG	N76187	006	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.125MG	N76187	007	Jun 05, 2002	Dec	CTEC

TABLET; ORAL

LEVOTHYROXINE SODIUM

>A>	AB1, AB2, AB3, AB4	MYLAN	0.125MG		N76187	007	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.137MG		N76187	012	Dec 13, 2006	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.137MG		N76187	012	Dec 13, 2006	Dec	CTEC
	AB1, AB2, AB3		0.137MG		N76187	012	Dec 13, 2006	Nov	NEWA
>D>	AB1, AB2, AB3		0.15MG		N76187	008	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.15MG		N76187	008	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.175MG		N76187	009	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.175MG		N76187	009	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.2MG		N76187	010	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.2MG		N76187	010	Jun 05, 2002	Dec	CTEC
>D>	AB1, AB2, AB3		0.3MG		N76187	011	Jun 05, 2002	Dec	CTEC
>A>	AB1, AB2, AB3, AB4		0.3MG		N76187	011	Jun 05, 2002	Dec	CTEC
		LEVOXYL							
	AB1, AB3	JONES PHARMA	0.137MG		N21301	008	May 25, 2001	Nov	CTEC
		SYNTHROID							
	AB1, AB2	ABBOTT	0.137MG		N21402	008	Jul 24, 2002	Nov	CTEC

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
	@		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						
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

TABLET; ORAL
LISINOPRIL
AB AUROBINDO 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

>D> LORACARBEF
>D> CAPSULE; ORAL
>D> LORABID
>D> KING PHARMS 200MG N50668 001 Dec 31, 1991 Dec DISC
>A> @ 200MG N50668 001 Dec 31, 1991 Dec DISC
>D> \+ 400MG N50668 002 Apr 05, 1996 Dec DISC
>A> @ 400MG N50668 002 Apr 05, 1996 Dec DISC
>D> FOR SUSPENSION; ORAL
>D> LORABID
>D> KING PHARMS 100MG/5ML N50667 001 Dec 31, 1991 Dec DISC
>A> @ 100MG/5ML N50667 001 Dec 31, 1991 Dec DISC
>D> \+ 200MG/5ML N50667 002 Dec 31, 1991 Dec DISC
>A> @ 200MG/5ML N50667 002 Dec 31, 1991 Dec DISC

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 IVAX PHARMS 0.5MG N77396 001 Dec 13, 2006 Nov NEWA
AB 1MG N77396 002 Dec 13, 2006 Nov NEWA
AB 2MG N77396 003 Dec 13, 2006 Nov NEWA
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

TABLET; ORAL
LOVASTATIN

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
AB	AUROBINDO PHARMA	7.5MG	N78008	001	Oct 02, 2006	Sep	NEWA
AB		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

TABLET; ORAL
MELOXICAM

	AB		CARACO	15MG	N77937	002	Jul 19, 2006	Jul	NEWA
	AB		CARLSBAD	7.5MG	N77918	001	Dec 07, 2006	Nov	NEWA
	AB			15MG	N77918	002	Dec 07, 2006	Nov	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
	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		RANBAXY	7.5MG	N78039	001	Dec 14, 2006	Nov	NEWA
	AB			15MG	N78039	002	Dec 14, 2006	Nov	NEWA
	AB		ROXANE	7.5MG	N77925	001	Jul 19, 2006	Jul	NEWA
	AB			15MG	N77925	002	Jul 19, 2006	Jul	NEWA
	AB		TARO	7.5MG	N78102	001	Nov 07, 2006	Oct	NEWA
	AB			15MG	N78102	002	Nov 07, 2006	Oct	NEWA
	AB		TEVA PHARMS	7.5MG	N77936	001	Jul 19, 2006	Jul	NEWA
	AB			15MG	N77936	002	Jul 19, 2006	Jul	NEWA
>A>	AB		UNICHEM	7.5MG	N77927	001	Dec 20, 2006	Dec	NEWA
>A>	AB			15MG	N77927	002	Dec 20, 2006	Dec	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

TABLET; ORAL
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 500MG N21252 001 Jan 05, 2001 May DISC
ROWASA
@ ALAVEN PHARM 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
METARAMINOL BITARTRATE
+ ABRAXIS PHARM EQ 10MG BASE/ML N80722 001 Jun CRLD

METAXALONE

TABLET; ORAL
SKELAXIN
>D> @ JONES PHARMA INC 400MG N13217 001 Dec CAHN
>D> + 800MG N13217 003 Aug 30, 2002 Dec CAHN
>A> @ KING PHARMS 400MG N13217 001 Dec CAHN
>A> + 800MG N13217 003 Aug 30, 2002 Dec CAHN

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

TAPAZOLE

>D>	AB		JONES PHARMA	5MG	N40320	001	Mar 31, 2000	Dec	CAHN
	AB			5MG	N40320	001	Mar 31, 2000	May	CTNA
>D>	AB			10MG	N40320	002	Mar 31, 2000	Dec	CAHN
	AB			10MG	N40320	002	Mar 31, 2000	May	CTNA
>A>	AB		KING PHARMS	5MG	N40320	001	Mar 31, 2000	Dec	CAHN
>A>	AB			10MG	N40320	002	Mar 31, 2000	Dec	CAHN

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
		+		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

METHSCOPOLAMINE BROMIDE

TABLET; ORAL

METHSCOPOLAMINE BROMIDE

>A>									
>A>	AA		BOCA PHARMA	2.5MG	N40624	001	Dec 28, 2006	Dec	NEWA
>A>	AA			5MG	N40624	002	Dec 28, 2006	Dec	NEWA

PAMINE

>D>		+	BRADLEY PHARMS	2.5MG	N08848	001		Dec	CTEC
>A>	AA	+		2.5MG	N08848	001		Dec	CTEC

PAMINE FORTE

>D>		+	BRADLEY PHARMS	5MG	N08848	002	Mar 25, 2003	Dec	CFTG
>A>	AA	+		5MG	N08848	002	Mar 25, 2003	Dec	CFTG

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

METHYLPREDNISOLONE ACETATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE ACETATE

AB		SICOR PHARMS	40MG/ML	N40620	001	Oct 27, 2006	Oct	NEWA
AB			80MG/ML	N40620	002	Oct 27, 2006	Oct	NEWA

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

METOPROLOL TARTRATE

TABLET; ORAL
METOPROLOL TARTRATE

	@	APOTHECON	50MG	N74258	001	Jan 27, 1994	Nov	DISC
	@		100MG	N74258	002	Jan 27, 1994	Nov	DISC

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

GEL; VAGINAL
METROGEL-VAGINAL

AB	+	3M	0.75%	N20208	001	Aug 17, 1992	Oct	CFTG

METRONIDAZOLE

AB		QLT USA	0.75%	N77264	001	Oct 31, 2006	Oct	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

METYROSINE

CAPSULE; ORAL
DEMSER

	+	ATON	250MG	N17871	001		Oct	CAHN

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

MILRINONE LACTATE

INJECTABLE; INJECTION
MILRINONE LACTATE

AP	GLAND PHARMA LTD	EQ 1MG BASE/ML	N77190	001	Oct 31, 2006	Oct	NEWA

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL
MINOCIN

AB	TRIAX PHARMS	EQ 50MG BASE	N50649	001	May 31, 1990	Feb	CAHN
	@	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

MIVACURIUM CHLORIDE

INJECTABLE; INJECTION
MIVACRON
@ ABBOTT EQ 2MG BASE/ML N20098 001 Jan 22, 1992 Sep DISC
MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER
@ ABBOTT EQ 50MG BASE/100ML N20098 003 Jan 22, 1992 Sep DISC

MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL
MOEXIPRIL HYDROCHLORIDE
AB PADDOCK 7.5MG N77536 001 Nov 30, 2006 Nov NEWA
AB 15MG N77536 002 Nov 30, 2006 Nov NEWA

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

CAPSULE, EXTENDED RELEASE; ORAL
KADIAN
ALPHARMA US PHARMS 80MG N20616 006 Oct 27, 2006 Oct NEWA
INJECTABLE; INJECTION
MORPHINE SULFATE
\+ HOSPIRA 5MG/ML N19916 002 Oct 27, 2006 Oct 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

TABLET; ORAL
CORGARD

AB		KING PHARMS	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

	+	MERZ PHARMS	1%	N19599	001	Feb 29, 1988	Sep	CRLD

NALTREXONE

FOR 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	CAHN

NEOMYCIN 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	NEWA

NEOMYCIN 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

SOLUTION; IRRIGATION
NEOSPORIN G.U. IRRIGANT

AT	+	MONARCH PHARMS	EQ 800MG BASE/20ML;4,000,000 UNITS/20ML (EQ 40MG BASE/ML;200,000 UNITS/ML)	N60707	002			Jan	NEWA

NIACIN

TABLET, EXTENDED RELEASE; ORAL
NIACIN

	@	BARR	500MG	N76378	001	Apr 26, 2005		Sep	DISC
	@		750MG	N76378	002	Apr 26, 2005		Sep	DISC
	@		1GM	N76250	001	Apr 14, 2005		Sep	DISC

NIASPAN

	+	KOS LIFE	500MG	N20381	002	Jul 28, 1997		Sep	CTEC
	+		750MG	N20381	003	Jul 28, 1997		Sep	CTEC
	+		1GM	N20381	004	Jul 28, 1997		Sep	CTEC

NICARDIPINE HYDROCHLORIDE

CAPSULE; ORAL
CARDENE

>A>	AB	PDL BIOPHARMA INC	20MG	N19488	001	Dec 21, 1988		Dec	CAHN
>A>	AB	+	30MG	N19488	002	Dec 21, 1988		Dec	CAHN
>D>	AB	ROCHE PALO	20MG	N19488	001	Dec 21, 1988		Dec	CAHN
>D>	AB	+	30MG	N19488	002	Dec 21, 1988		Dec	CAHN

NICARDIPINE HYDROCHLORIDE

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

CAPSULE, EXTENDED RELEASE; ORAL
CARDENE SR

>A>	+	PDL BIOPHARMA INC	30MG	N20005	001	Feb 21, 1992		Dec	CAHN
>A>	+		45MG	N20005	002	Feb 21, 1992		Dec	CAHN
>A>	+		60MG	N20005	003	Feb 21, 1992		Dec	CAHN
>D>	+	ROCHE PALO	30MG	N20005	001	Feb 21, 1992		Dec	CAHN
>D>	+		45MG	N20005	002	Feb 21, 1992		Dec	CAHN
>D>	+		60MG	N20005	003	Feb 21, 1992		Dec	CAHN

INJECTABLE; INJECTION
CARDENE

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

NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL
NICOTINE

		AVEVA	7MG/24HR	N74645	001	Oct 20, 1997		Oct	CAHN
			14MG/24HR	N74611	001	Oct 20, 1997		Oct	CAHN
			21MG/24HR	N74612	001	Oct 20, 1997		Oct	CAHN

SPRAY, METERED; NASAL
NICOTROL

	+	PFIZER INC	0.5MG/SPRAY	N20385	001	Mar 22, 1996		Nov	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

NIFEDIPINE

AB1	ABRIKA PHARMS	30MG	N77899	001	Dec 13, 2006	Nov	NEWA
AB1		60MG	N77899	002	Dec 13, 2006	Nov	NEWA

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

AEROSOL, METERED; SUBLINGUAL
NITROMIST

+	NOVADEL	0.4MG/SPRAY	N21780	001	Nov 02, 2006	Nov	NEWA

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

@	STADA PHARMS	100%	N50576	001	Dec 22, 1983	Sep	CAHN

NYSTATIN

@	PADDOCK	100%	N62613	001	Nov 26, 1985	Oct	DISC

POWDER; TOPICAL
NYSTATIN

AT	KV PHARM	100,000 UNITS/GM	N65321	001	Aug 18, 2006	Aug	NEWA
AT	PHARMAFORCE	100,000 UNITS/GM	N65203	001	Jul 15, 2004	Nov	CAHN

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
FLOXIN OTIC

+	DAIICHI	0.3%	N20799	001	Dec 16, 1997	Oct	CTEC

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

OLOPATADINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC
PATADAY

+	ALCON	0.2%	N21545	001	Dec 22, 2004	Oct	CTNA

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

ONDANSETRON

TABLET, ORALLY DISINTEGRATING; ORAL
ONDANSETRON

>A>								
>A>	AB	KALI LABS	4MG	N76506	001	Dec 26, 2006	Dec	NEWA
>A>	AB		8MG	N76506	002	Dec 26, 2006	Dec	NEWA
>A>			16MG	N77406	001	Dec 26, 2006	Dec	NEWA
>A>			24MG	N77406	002	Dec 26, 2006	Dec	NEWA

TABLET, ORALLY DISINTEGRATING; ORAL

ZOFRAN ODT

>D>			GLAXOSMITHKLINE	4MG	N20781	001	Jan 27, 1999	Dec	CFTG
>A>	AB			4MG	N20781	001	Jan 27, 1999	Dec	CFTG
				4MG	N20781	001	Jan 27, 1999	Nov	CPOT
>D>		+		8MG	N20781	002	Jan 27, 1999	Dec	CFTG
>A>	AB	+		8MG	N20781	002	Jan 27, 1999	Dec	CFTG
		+		8MG	N20781	002	Jan 27, 1999	Nov	CPOT

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON

	AP		TEVA	EQ 2MG BASE/ML	N76876	001	Nov 22, 2006	Nov	NEWA

ONDANSETRON AND DEXTROSE IN PLASTIC CONTAINER

>D>	AP		SICOR PHARMS	EQ 0.64MG BASE/ML	N77480	001	Nov 22, 2006	Dec	CTNA
	AP			EQ 0.64MG BASE/ML	N77480	001	Nov 22, 2006	Nov	NEWA

ONDANSETRON HYDROCHLORIDE

>A>	AP		ABRAXIS PHARM	EQ 2MG BASE/ML	N76974	001	Dec 26, 2006	Dec	NEWA
>A>	AP		APOTEX	EQ 2MG BASE/ML	N77368	001	Dec 26, 2006	Dec	NEWA
>A>	AP		BAXTER HLTHCARE	EQ 2MG BASE/ML	N77365	001	Dec 26, 2006	Dec	NEWA
>A>	AP		BEDFORD	EQ 2MG BASE/ML	N76967	001	Dec 26, 2006	Dec	NEWA
>A>	AP		HIKMA FARMACEUTICA	EQ 2MG BASE/ML	N76781	001	Dec 26, 2006	Dec	NEWA
>A>	AP		HOSPIRA	EQ 2MG BASE/ML	N77473	001	Dec 26, 2006	Dec	NEWA
>A>	AP		MAYNE PHARMA USA	EQ 2MG BASE/ML	N76695	001	Dec 26, 2006	Dec	NEWA
>A>	AP		PHARMAFORCE	EQ 2MG BASE/ML	N77582	001	Dec 26, 2006	Dec	NEWA
>A>	AP		PLIVA HRVATSKA DOO	EQ 2MG BASE/ML	N77544	001	Dec 26, 2006	Dec	NEWA
>A>	AP		SUN PHARM INDS (IN)	EQ 2MG BASE/ML	N77172	001	Dec 26, 2006	Dec	NEWA
>A>	AP		TEVA	EQ 2MG BASE/ML	N76876	001	Nov 22, 2006	Dec	NEWA
>A>	AP		WOCKHARDT	EQ 2MG BASE/ML	N77577	001	Dec 26, 2006	Dec	NEWA

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

>A>	AP		SICOR PHARMS	EQ 0.64MG BASE/ML	N77480	001	Nov 22, 2006	Dec	CTNA

ONDANSETRON HYDROCHLORIDE AND SODIUM CHLORIDE IN PLASTIC CONTAINER

>A>	AP	+	BAXTER HLTHCARE	EQ 0.64MG BASE/ML	N21915	002	Dec 27, 2006	Dec	NEWA

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

>A>	AP		ABRAXIS PHARM	EQ 2MG BASE/ML	N76972	001	Dec 26, 2006	Dec	NEWA
>A>	AP		APOTEX INC	EQ 2MG BASE/ML	N77343	001	Dec 26, 2006	Dec	NEWA
>A>	AP		BAXTER HLTHCARE	EQ 2MG BASE/ML	N77541	001	Dec 26, 2006	Dec	NEWA
>A>	AP		BEDFORD LABS	EQ 2MG BASE/ML	N77011	001	Dec 26, 2006	Dec	NEWA
>A>	AP		HIKMA FARMACEUTICA	EQ 2MG BASE/ML	N76780	001	Dec 26, 2006	Dec	NEWA
>A>	AP		HOSPIRA	EQ 2MG BASE/ML	N77548	001	Dec 26, 2006	Dec	NEWA
>A>	AP		MAYNE PHARMA USA	EQ 2MG BASE/ML	N76696	001	Dec 26, 2006	Dec	NEWA
>A>	AP		PHARMAFORCE	EQ 2MG BASE/ML	N77387	001	Dec 26, 2006	Dec	NEWA
>A>	AP		SUN PHARM INDS LTD	EQ 2MG BASE/ML	N77173	001	Dec 26, 2006	Dec	NEWA
>A>	AP		TEVA	EQ 2MG BASE/ML	N76759	001	Nov 22, 2006	Dec	CTNA
>A>	AP		WOCKHARDT	EQ 2MG BASE/ML	N77716	001	Dec 26, 2006	Dec	NEWA

ONDANSETRON PRESERVATIVE FREE

>D>	AP		TEVA	EQ 2MG BASE/ML	N76759	001	Nov 22, 2006	Dec	CTNA
	AP			EQ 2MG BASE/ML	N76759	001	Nov 22, 2006	Nov	NEWA

ZOFRAN

	AP	+	GLAXOSMITHKLINE	EQ 2MG BASE/ML	N20007	001	Jan 04, 1991	Nov	CFTG

ZOFRAN IN PLASTIC CONTAINER

	AP	+	GLAXOSMITHKLINE	EQ 0.64MG BASE/ML	N20403	001	Jan 31, 1995	Nov	CFTG

ZOFRAN PRESERVATIVE FREE

	AP	+	GLAXOSMITHKLINE	EQ 2MG BASE/ML	N20007	003	Dec 10, 1993	Nov	CFTG

		SOLUTION; ORAL							
>A>		ONDANSETRON HYDROCHLORIDE							
>A>	AA	ROXANE	EQ 4MG BASE/5ML	N76960	001	Dec 26, 2006	Dec	NEWA	
		ZOFRAN							
>D>	+	GLAXOSMITHKLINE	EQ 4MG BASE/5ML	N20605	001	Jan 24, 1997	Dec	CFTG	
>A>	AA	+	EQ 4MG BASE/5ML	N20605	001	Jan 24, 1997	Dec	CFTG	
		TABLET; ORAL							
>A>		ONDANSETRON HYDROCHLORIDE							
>A>	AB	DR REDDYS LABS LTD	EQ 4MG BASE	N76183	003	Dec 26, 2006	Dec	NEWA	
>A>	AB		EQ 8MG BASE	N76183	002	Dec 26, 2006	Dec	NEWA	
>A>			EQ 16MG BASE	N76559	001	Dec 26, 2006	Dec	NEWA	
>A>	AB		EQ 24MG BASE	N76183	001	Dec 26, 2006	Dec	NEWA	
		ZOFRAN							
>D>		GLAXOSMITHKLINE	EQ 4MG BASE	N20103	001	Dec 31, 1992	Dec	CFTG	
>A>	AB		EQ 4MG BASE	N20103	001	Dec 31, 1992	Dec	CFTG	
>D>			EQ 8MG BASE	N20103	002	Dec 31, 1992	Dec	CFTG	
>A>	AB		EQ 8MG BASE	N20103	002	Dec 31, 1992	Dec	CFTG	
>D>	+		EQ 24MG BASE	N20103	003	Aug 27, 1999	Dec	CFTG	
>A>	AB	+	EQ 24MG BASE	N20103	003	Aug 27, 1999	Dec	CFTG	
		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	N21492	001	Aug 09, 2002	Jun	DISC	
		@	100MG/VIAL	N21492	002	Aug 09, 2002	Jun	DISC	
		+	200MG/40ML (5MG/ML)	N21759	003	Nov 17, 2006	Nov	NEWA	
		OXANDROLONE							
		TABLET; ORAL							
		OXANDRIN							
	AB	SAVIENT PHARMS	2.5MG	N13718	001		Nov	CFTG	
	AB	+	10MG	N13718	002	Nov 05, 2001	Nov	CFTG	
		OXANDROLONE							
	AB	SANDOZ	2.5MG	N76897	001	Dec 01, 2006	Nov	NEWA	
	AB		10MG	N76897	002	Dec 01, 2006	Nov	NEWA	
	AB	UPSHER SMITH	2.5MG	N76761	001	Dec 01, 2006	Nov	NEWA	
		OXAPROZIN							
		TABLET; ORAL							
		OXAPROZIN							
	AB	ACTAVIS ELIZABETH	600MG	N75843	001	Oct 03, 2001	Jun	CAHN	
		OXAZEPAM							
		CAPSULE; ORAL							
		OXAZEPAM							
	AB	ACTAVIS ELIZABETH	10MG	N72251	001	Apr 14, 1988	Jun	CAHN	
	AB		15MG	N72252	001	Apr 14, 1988	Jun	CAHN	
	AB		30MG	N72253	001	Apr 14, 1988	Jun	CAHN	

OXYBUTYNIN CHLORIDE

TABLET, EXTENDED RELEASE; ORAL

DITROPAN XL

AB		ALZA	5MG	N20897	001	Dec 16, 1998	Oct	CFTG
AB			10MG	N20897	002	Dec 16, 1998	Oct	CFTG
AB	+		15MG	N20897	003	Jun 22, 1999	Oct	CFTG

OXYBUTYNIN CHLORIDE

AB		IMPAX PHARMS	15MG	N76745	001	Nov 09, 2006	Oct	NEWA
AB		MYLAN	5MG	N76702	001	Nov 09, 2006	Oct	NEWA
AB			10MG	N76644	001	Nov 09, 2006	Oct	NEWA

OXYCODONE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

OXYCONTIN

		PURDUE PHARMA LP	15MG	N20553	006	Sep 18, 2006	Sep	NEWA
			30MG	N20553	007	Sep 18, 2006	Sep	NEWA
			60MG	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	N21611	001	Jun 22, 2006	Jun	NEWA
	+		10MG	N21611	002	Jun 22, 2006	Jun	NEWA

TABLET, EXTENDED RELEASE; ORAL

OPANA ER

		ENDO PHARMS	5MG	N21610	001	Jun 22, 2006	Jun	NEWA
			10MG	N21610	002	Jun 22, 2006	Jun	NEWA
			20MG	N21610	003	Jun 22, 2006	Jun	NEWA
	+		40MG	N21610	004	Jun 22, 2006	Jun	NEWA

PACLITAXEL

FOR SUSPENSION; IV (INFUSION)

ABRAXANE

	+	ABRAXIS BIOSCIENCE	100MG/VIAL	N21660	001	Jan 07, 2005	Apr	CAHN

INJECTABLE; INJECTION

PACLITAXEL

AP		DABUR ONCOLOGY PLC	6MG/ML	N77574	001	Nov 27, 2006	Nov	NEWA
AP		SICOR PHARMS	6MG/ML	N75184	001	Jan 25, 2002	Nov	CAHN
AP			6MG/ML	N75297	001	Jan 25, 2002	Nov	CAHN

>A> PALIPERIDONE

>A> TABLET, EXTENDED RELEASE; ORAL

>A> INVEGA

>A>		JANSSEN LP	3MG	N21999	001	Dec 19, 2006	Dec	NEWA
>A>			6MG	N21999	002	Dec 19, 2006	Dec	NEWA
>A>	+		9MG	N21999	003	Dec 19, 2006	Dec	NEWA
>A>	@		12MG	N21999	004	Dec 19, 2006	Dec	DISC

PARICALCITOL

INJECTABLE; INJECTION

ZEMPLAR

+	ABBOTT	0.002MG/ML	N20819	002	Feb 01, 2000	Aug	CRLD

PAROXETINE HYDROCHLORIDE

SUSPENSION; ORAL

PAROXETINE HYDROCHLORIDE

AB	APOTEX INC	EQ 10MG BASE/5ML	N77395	001	Dec 05, 2006	Nov	NEWA

PAXIL

AB	+ GLAXOSMITHKLINE	EQ 10MG BASE/5ML	N20710	001	Jun 25, 1997	Nov	CFTG

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

VEETIDS

	@ APOTHECON	EQ 250MG BASE	N61411	001		Nov	DISC
	@	EQ 500MG BASE	N61411	002		Nov	DISC

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

TABLET, EXTENDED RELEASE; ORAL
PENTOXIFYLLINE

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

AA	+	SANDOZ	15MG	N87190	002		Nov	CMS1
		@	15MG	N87301	001		Nov	DISC
		@	30MG	N87208	001		Nov	DISC
		@	30MG	N87223	001		Nov	DISC
		@ 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

PHENYLEPHRINE HYDROCHLORIDE AND PROMETHAZINE HYDROCHLORIDE

AA		VINTAGE	5MG/5ML;6.25MG/5ML	N40654	001	Dec 07, 2006	Nov	NEWA

PROMETH VC PLAIN

AA	+	ACTAVIS MID ATLANTIC	5MG/5ML;6.25MG/5ML	N88761	001	Nov 08, 1984	Nov	CFTG
	+		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		SUN PHARM INDS (IN)	100MG EXTENDED	N40621	001	Dec 11, 2006	Nov	NEWA
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

		@ ATON	1MG/0.5ML	N12223	002		Oct	CAHN
		@	10MG/ML	N12223	001		Oct	CAHN
		@ 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

TABLET; ORAL

MEPHYTON

	+	ATON	5MG	N10104	003		Oct	CAHN

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

PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

ACTOS

		TAKEDA GLOBAL	EQ 15MG BASE	N21073	001	Jul 15, 1999	Oct	CAHN
			EQ 30MG BASE	N21073	002	Jul 15, 1999	Oct	CAHN
	+		EQ 45MG BASE	N21073	003	Jul 15, 1999	Oct	CAHN

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

POSACONAZOLE

SUSPENSION; ORAL

NOXAFIL

	+	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	+	COPLEY PHARM	8MEQ	N70618	001	Sep 09, 1987	Jan	CRLD

POTASSIUM CHLORIDE; SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.45% AND POTASSIUM CHLORIDE 0.15% IN PLASTIC CONTAINER

	+	BAXTER HLTHCARE	450MG/100ML;150MG/100ML	N17648	005	Nov 26, 2002	Oct	NEWA

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		APOTEX	10MG	N76341	001	Oct 23, 2006	Oct	NEWA
AB			20MG	N76341	002	Oct 23, 2006	Oct	NEWA
AB			40MG	N76341	003	Oct 23, 2006	Oct	NEWA
AB		COBALT	10MG	N76939	004	Oct 23, 2006	Oct	NEWA
AB			20MG	N76939	003	Oct 23, 2006	Oct	NEWA
AB			40MG	N76939	002	Oct 23, 2006	Oct	NEWA
AB		DR REDDYS LABS INC	10MG	N76714	001	Oct 23, 2006	Oct	NEWA

TABLET; ORAL
PRAVASTATIN SODIUM

AB	DR REDDYS LABS INC	20MG	N76714	002	Oct 23, 2006	Oct	NEWA
AB		40MG	N76714	003	Oct 23, 2006	Oct	NEWA
AB	GENPHARM	10MG	N77013	001	Oct 23, 2006	Oct	NEWA
AB		20MG	N77013	002	Oct 23, 2006	Oct	NEWA
AB		40MG	N77013	003	Oct 23, 2006	Oct	NEWA
AB	LEK PHARMS DD	10MG	N76397	003	Oct 23, 2006	Oct	NEWA
AB		20MG	N76397	002	Oct 23, 2006	Oct	NEWA
AB		40MG	N76397	001	Oct 23, 2006	Oct	NEWA
AB	PLIVA HRVATSKA DOO	10MG	N77730	001	Nov 21, 2006	Nov	NEWA
AB		20MG	N77730	002	Nov 21, 2006	Nov	NEWA
		30MG	N77730	003	Nov 21, 2006	Nov	NEWA
AB		40MG	N77730	005	Nov 21, 2006	Nov	NEWA
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
AB	WATSON LABS	10MG	N77904	001	Oct 23, 2006	Oct	NEWA
AB		20MG	N77904	002	Oct 23, 2006	Oct	NEWA
AB		40MG	N77904	003	Oct 23, 2006	Oct	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

AB	+ SANOFI AVENTIS US	0.1%	N20279	001	Oct 29, 1993	Sep	CFTG

PREDNICARBATE

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
	MEDICIS	EQ 10MG BASE	N21959	001	Jun 01, 2006	Oct	CAHN
		EQ 15MG BASE	N21959	002	Jun 01, 2006	Oct	CAHN
+		EQ 30MG BASE	N21959	003	Jun 01, 2006	Oct	CAHN

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

INJECTABLE; INJECTION
PROMETHAZINE HYDROCHLORIDE

AP		SANDOZ	25MG/ML	N40593	001	Nov 08, 2006	Oct	NEWA
AP			50MG/ML	N40593	002	Nov 08, 2006	Oct	NEWA

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
AB		TARO	12.5MG	N40603	001	Oct 26, 2006	Oct	NEWA
AB			25MG	N40603	002	Oct 26, 2006	Oct	NEWA

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

TABLET; ORAL
PROMETHAZINE HYDROCHLORIDE

AB	VINTAGE PHARMS	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
	@ IMPAX LABS	50MG	N80159	001		Nov	DISC
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	N18708	003	Feb 26, 1987	May	CAHN
	+	15MG	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

TABLET; ORAL
QUINAPRIL HYDROCHLORIDE

AB	TORPHARM	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	200MG	N87011	001		Jan	DISC
	@ LANNETT	200MG	N83743	001		Jan	DISC
	@ MUTUAL PHARM	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
AB	WATSON LABS	200MG	N83288	001		Jul	CMFD
	@	200MG	N83288	001		Jan	DISC
	@	200MG	N85140	002		Jan	DISC

QUININE SULFATE

CAPSULE; ORAL
QUININE SULFATE

	+ AR HOLDING CO INC	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

RANITIDINE HYDROCHLORIDE

TABLET; ORAL
RANITIDINE HYDROCHLORIDE

AB	INTERPHARM	EQ 150MG BASE	N77824	001	Oct 13, 2006	Oct	NEWA
AB		EQ 300MG BASE	N77824	002	Oct 13, 2006	Oct	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

	COPEGUS							
AB	ROCHE	200MG	N21511	001	Dec 03, 2002	Nov	CRLD	
	RIBAVIRIN							
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	

SALMETEROL XINAFOATE

AEROSOL, METERED; INHALATION

	SEREVENT							
	@ GLAXOSMITHKLINE	EQ 0.021MG BASE/INH	N20236	001	Feb 04, 1994	Oct	DISC	

SAQUINAVIR

CAPSULE; ORAL

	FORTOVASE							
	@ HLR	200MG	N20828	001	Nov 07, 1997	Nov	DISC	

SECRETIN SYNTHETIC PORCINE

FOR SOLUTION; INTRAVENOUS

	SECREFLO							
+	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

	SELEGILINE HYDROCHLORIDE							
	@ AAIPHARMA LLC	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
>A> AB AUROBINDO PHARMA 5MG N77691 001 Dec 20, 2006 Dec NEWA
>A> AB 10MG N77691 002 Dec 20, 2006 Dec NEWA
>A> AB 20MG N77691 003 Dec 20, 2006 Dec NEWA
>A> AB 40MG N77691 004 Dec 20, 2006 Dec NEWA
>A> AB 80MG N77691 005 Dec 20, 2006 Dec NEWA
>A> AB COBALT 5MG N76685 001 Dec 20, 2006 Dec NEWA
>A> AB 10MG N76685 002 Dec 20, 2006 Dec NEWA
>A> AB 20MG N76685 003 Dec 20, 2006 Dec NEWA
>A> AB 40MG N76685 004 Dec 20, 2006 Dec NEWA
>A> AB 80MG N76685 005 Dec 20, 2006 Dec NEWA
>A> AB DR REDDYS LABS INC 10MG N77752 001 Dec 20, 2006 Dec NEWA
>A> AB 20MG N77752 002 Dec 20, 2006 Dec NEWA
>A> AB 40MG N77752 003 Dec 20, 2006 Dec NEWA
>A> AB 80MG N77752 004 Dec 20, 2006 Dec NEWA
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
>A> AB 80MG N76052 005 Dec 20, 2006 Dec NEWA
>A> AB PERRIGO R AND D 5MG N78034 001 Dec 20, 2006 Dec NEWA
>A> AB 10MG N78034 002 Dec 20, 2006 Dec NEWA
>A> AB 20MG N78034 003 Dec 20, 2006 Dec NEWA
>A> AB 40MG N78034 004 Dec 20, 2006 Dec NEWA
>A> AB 80MG N78034 005 Dec 20, 2006 Dec NEWA
>A> AB RANBAXY 5MG N76285 001 Dec 20, 2006 Dec NEWA

TABLET; ORAL
SIMVASTATIN

>A>	AB		RANBAXY	10MG	N76285	002	Dec 20, 2006	Dec	NEWA
>A>	AB			20MG	N76285	003	Dec 20, 2006	Dec	NEWA
>A>	AB			40MG	N76285	004	Dec 20, 2006	Dec	NEWA
	AB			80MG	N76285	005	Jun 23, 2006	Jun	NEWA
>A>	AB		SANDOZ	5MG	N77766	001	Dec 20, 2006	Dec	NEWA
>A>	AB			10MG	N77766	002	Dec 20, 2006	Dec	NEWA
>A>	AB			20MG	N77766	003	Dec 20, 2006	Dec	NEWA
>A>	AB			40MG	N77766	004	Dec 20, 2006	Dec	NEWA
>A>	AB			80MG	N77766	005	Dec 20, 2006	Dec	NEWA
>A>	AB		ZYDUS PHARMS USA	5MG	N77837	001	Dec 20, 2006	Dec	NEWA
>A>	AB			10MG	N77837	002	Dec 20, 2006	Dec	NEWA
>A>	AB			20MG	N77837	003	Dec 20, 2006	Dec	NEWA
>A>	AB			40MG	N77837	004	Dec 20, 2006	Dec	NEWA
>A>	AB			80MG	N77837	005	Dec 20, 2006	Dec	NEWA

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

SITAGLIPTIN PHOSPHATE

TABLET; ORAL
JANUVIA

			MERCK CO INC	EQ 25MG BASE	N21995	001	Oct 16, 2006	Oct	NEWA
				EQ 50MG BASE	N21995	002	Oct 16, 2006	Oct	NEWA
		+		EQ 100MG BASE	N21995	003	Oct 16, 2006	Oct	NEWA

SODIUM CHLORIDE

INJECTABLE; INJECTION
SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

	AP	+	ABRAXIS PHARM	9MG/ML	N88912	001	Jan 10, 1985	Oct	CRLD
	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	9MG/ML	N18803	001	Oct 29, 1982	Oct	CRLD
	AP	+		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	Oct	CRLD
	AP			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

	AA	+	GE HEALTHCARE	100uCi	N17630	001		Sep	CRLD
	AA	+	SYNCOR PHARMS	100uCi	N18671	001	May 27, 1982	Sep	CRLD

		CAPSULE; ORAL							
		SODIUM IODIDE I 123							
AA	+	SYNCOR PHARMS	200uCi	N18671	002	May 27, 1982	Sep	CRLD	
		SOLUTION; ORAL							
		SODIUM IODIDE I 123							
	+	GE HEALTHCARE	2mCi/ML	N17630	002		Sep	CRLD	
		SODIUM IODIDE, I-131							
		CAPSULE; ORAL							
		SODIUM IODIDE I 131							
>D>		DRAXIMAGE	100mCi	N21305	004	Nov 18, 2004	Dec	CPOT	
>A>			100uCi	N21305	004	Nov 18, 2004	Dec	CPOT	
			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	
		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	
		SOYBEAN OIL							
		INJECTABLE; INJECTION							
		SOYACAL 10%							
		@ ALPHA THERA	10%	N18465	001	Jun 29, 1983	Nov	DISC	
		SOYACAL 20%							
		@ ALPHA THERA	20%	N18786	001	Jun 29, 1983	Nov	DISC	
		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	

INJECTABLE; INJECTION

ANECTINE

@	SANDOZ	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

LOTION; TOPICAL

KLARON

AB	+	SANOFI AVENTIS US	10%	N19931	001	Dec 23, 1996	Nov	CFTG

SULFACETAMIDE SODIUM

AB		ALTANA	10%	N77015	001	Nov 17, 2006	Nov	NEWA

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

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB		TEVA PHARMS	200MG/5ML;40MG/5ML	N77612	001	Nov 13, 2006	Oct	NEWA

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

SULFANILAMIDE

CREAM; VAGINAL

AVC

+	PHARMELLE	15%	N06530	003	Jan 27, 1987	Oct	CMFD

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							
+ ROSEMONT	EQ 10MG BASE/5ML	N21807	001	Oct 29, 2005	Nov	CAHN
+ 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

TELBIVUDINE

TABLET; ORAL							
TYZEKA							
+ IDENIX	600MG	N22011	001	Oct 25, 2006	Oct	NEWA

TEMAZEPAM

CAPSULE; ORAL

		TEMAZEPAM						
AB		ACTAVIS ELIZABETH	15MG	N71638	001	Aug 07, 1987	Jun	CAHN
AB			30MG	N71620	001	Aug 07, 1987	Jun	CAHN

TEMOZOLOMIDE

CAPSULE; ORAL

		TEMODAR						
		SCHERING	140MG	N21029	005	Oct 19, 2006	Oct	NEWA
			180MG	N21029	006	Oct 19, 2006	Oct	NEWA

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						
AB	+	ORTHO MCNEIL PHARM	0.8%	N19964	001	Feb 21, 1991	Sep	CMFD
		@	0.8%	N19964	001	Feb 21, 1991	Jul	DISC
		TERAZOL 7						
AB	+	ORTHO MCNEIL PHARM	0.4%	N19579	001	Dec 31, 1987	Sep	CMFD
		@	0.4%	N19579	001	Dec 31, 1987	Jul	DISC
		TERCONAZOLE						
AB		TARO	0.4%	N76043	001	Jan 19, 2005	Sep	CRLD
AB	+		0.4%	N76043	001	Jan 19, 2005	Jul	CRLD
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
AO		SYNERX PHARMA	200MG/ML	N40652	001	Dec 11, 2006	Nov	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

INJECTABLE; INTRAVENOUS

		THALLOUS CHLORIDE TL 201						
AP	+	BRISTOL MYERS SQUIBB	2mCi/ML	N17806	002	Oct 09, 1998	Oct	CFTG
AP		TCI MEDCL	2mCi/ML	N77698	001	Nov 09, 2006	Oct	NEWA

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	Nov	CRLD
AB	+		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

TRANDOLAPRIL

TABLET; ORAL

MAVIK

AB		ABBOTT	1MG	N20528	001	Apr 26, 1996	Nov	CFTG
AB			2MG	N20528	002	Apr 26, 1996	Nov	CFTG
AB	+		4MG	N20528	003	Apr 26, 1996	Nov	CFTG

TRANDOLAPRIL

AB		TEVA PHARMS	1MG	N77489	001	Dec 12, 2006	Nov	NEWA
AB			2MG	N77489	002	Dec 12, 2006	Nov	NEWA
AB			4MG	N77489	003	Dec 12, 2006	Nov	NEWA

TRANYLCYPROMINE SULFATE

TABLET; ORAL

PARNATE

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

TRANYLCYPROMINE SULFATE

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

TRAVOPROST

SOLUTION/DROPS; OPHTHALMIC

TRAVATAN Z

	+	ALCON	0.004%	N21994	001	Sep 21, 2006	Sep	NEWA

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

DESYREL

		@ APOTHECON	50MG	N18207	001		Aug	DISC
		@	100MG	N18207	002		Aug	DISC
		@	150MG	N18207	003	Mar 25, 1985	Aug	DISC

TABLET; ORAL
DESYREL
@ APOTHECON 300MG N18207 004 Nov 07, 1988 Aug DISC
TRAZODONE HYDROCHLORIDE
AB ACTAVIS ELIZABETH 50MG N71636 001 Apr 18, 1988 Jun CAHN
AB 100MG N71514 001 Apr 18, 1988 Jun CAHN
AB APOTEX INC 100MG N71196 001 Mar 25, 1987 Nov CAHN
AB 150MG N71196 002 Apr 26, 1999 Nov CAHN
AB + 300MG N71196 003 Apr 26, 1999 Nov CAHN
AB + BARR 300MG N71196 003 Apr 26, 1999 Sep CRLD

TRETINOIN

CREAM; TOPICAL
TRETINOIN
AB TRIAX PHARMS LLC 0.025% N75264 001 Dec 24, 1998 Nov CAHN
AB1 0.05% N75265 001 Dec 24, 1998 Nov CAHN
AB 0.1% N75213 001 Dec 24, 1998 Nov CAHN

TRIAMCINOLONE

TABLET; ORAL
ARISTOCORT
@ ASTELLAS 4MG N11161 007 Sep DISC
>D> KENACORT
>D> + BRISTOL MYERS SQUIBB 4MG N11283 006 Dec DISC
>A> @ 4MG N11283 006 Dec DISC
+ 4MG N11283 006 Sep CRLD

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL
ARISTOCORT A
@ ASTELLAS 0.025% N88818 001 Oct 16, 1984 Sep DISC
@ 0.1% N88819 001 Oct 16, 1984 Sep DISC
@ 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

LOTION; TOPICAL
TRIAMCINOLONE ACETONIDE
AT VINTAGE 0.1% N40672 002 Dec 13, 2006 Nov NEWA

OINTMENT; TOPICAL
TRIAMCINOLONE ACETONIDE
AT ACTAVIS MID ATLANTIC 0.1% N87799 001 Jun 07, 1982 Jun CAHN

SPRAY, METERED; NASAL
NASACORT HFA
@ SANOFI AVENTIS US 0.055MG/SPRAY N20784 001 Apr 07, 2004 Nov DISC

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

TRIENTINE HYDROCHLORIDE

CAPSULE; ORAL
SYPRINE

+	ATON	250MG	N19194	001	Nov 08, 1985	Oct	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

TRIPELENNAMINE 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

VALPROIC ACID

SOLUTION; ORAL

	VALPROIC ACID							
AA	VINTAGE	250MG/5ML	N77960	001	Oct 13, 2006		Oct	NEWA

VALRUBICIN

SOLUTION; INTRAVESICAL

	VALSTAR PRESERVATIVE FREE							
+	VALERA	40MG/ML	N20892	001	Sep 25, 1998		Jul	CAHN

VARENICLINE TARTRATE

TABLET; ORAL

	CHANTIX							
	CPPI CV	EQ 0.5MG BASE	N21928	001	May 10, 2006		Jul	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
	PFIZER INC	EQ 0.5MG BASE	N21928	001	May 10, 2006		Sep	CAHN
+		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
AB		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

TABLET, EXTENDED RELEASE; ORAL
ISOPTIN SR

AB	+	FSC	120MG	N19152	003	Mar 06, 1991	Nov	CAHN
AB	+		180MG	N19152	002	Dec 15, 1989	Nov	CAHN
AB	+		240MG	N19152	001	Dec 16, 1986	Nov	CAHN

VORINOSTAT

CAPSULE; ORAL
ZOLINZA

	+	MERCK	100MG	N21991	001	Oct 06, 2006	Oct	NEWA

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

ZALEPLON

CAPSULE; ORAL
SONATA

>D>		JONES PHARMA	5MG	N20859	001	Aug 13, 1999	Dec	CAHN
>D>	+		10MG	N20859	002	Aug 13, 1999	Dec	CAHN
>A>		KING PHARMS	5MG	N20859	001	Aug 13, 1999	Dec	CAHN
>A>	+		10MG	N20859	002	Aug 13, 1999	Dec	CAHN

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 HYDROCHLORIDE

SUSPENSION; ORAL

GEODON

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

ZONISAMIDE

CAPSULE; 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
AB		DR REDDYS LABS LTD	25MG	N77891	001	Sep 29, 2006	Sep	NEWA
AB			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
AB		ZYDUS PHARMS USA	25MG	N77625	001	Oct 16, 2006	Oct	NEWA
AB			50MG	N77625	002	Oct 16, 2006	Oct	NEWA
AB			100MG	N77625	003	Oct 16, 2006	Oct	NEWA

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
CAPITAL SOLEIL 15							
+ LOREAL USA	2%;3%;10%	N21501	001	Oct 02, 2006	Oct		NEWA

AVOBENZONE; ECAMSULE; OCTOCRYLENE; TITANIUM DIOXIDE

CREAM; TOPICAL							
ANTHELIOS 20							
+ LOREAL USA	2%;2%;10%;2%	N21471	001	Oct 05, 2006	Oct		NEWA

CHLORHEXIDINE GLUCONATE

CLOTH; TOPICAL							
CHLORHEXIDINE GLUCONATE							
+ SAGE PRODS	2%	N21669	001	Apr 25, 2005	Oct		CTNA
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 ONE-STEP							
+ MEDI FLEX INC	2%;70% (26ML)	N20832	006	Nov 21, 2006	Nov		NEWA
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						
>A>	@ MCNEIL CONS	5.2MG/SPRAY	N20463	001	Jan 03, 1997	Dec	CAHN
>D>	@ PHARMACIA UPJOHN	5.2MG/SPRAY	N20463	001	Jan 03, 1997	Dec	CAHN
	@	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							
	DR REDDYS LABS LTD	20MG	N77367	001	Sep 25, 2006	Sep	NEWA
	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

TABLET, CHEWABLE; ORAL
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 RECOMBINANT HUMAN

INJECTABLE; INJECTION
VELOSULIN BR
@ NOVO NORDISK INC 100 UNITS/ML N21028 001 Jul 19, 1999 Nov 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

INSULIN ZINC SUSP RECOMBINANT HUMAN

INJECTABLE; INJECTION
NOVOLIN L
@ NOVO NORDISK INC 100 UNITS/ML N19965 001 Jun 25, 1991 Nov DISC

IODINE POVACRYLEX; ISOPROPYL ALCOHOL

SPONGE; TOPICAL
DURAPREP
+ 3M EQ 0.7% IODINE;74% (26ML) N21586 002 Sep 29, 2006 Oct CRLD

SPONGE; TOPICAL
DURAPREP
\+ 3M EQ 0.7% IODINE; 74% (6ML) N21586 001 Sep 29, 2006 Sep NEWA
EQ 0.7% IODINE;74% (26ML) N21586 002 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

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC
>A> ALAWAY
>A> \+ ALIMERA SCIENCES INC EQ 0.025% BASE N21996 001 Dec 01, 2006 Dec NEWA
ZADITOR
\+ NOVARTIS EQ 0.025% BASE N21066 002 Oct 19, 2006 Oct NEWA

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, ORALLY DISINTEGRATING; ORAL
>A> CLARITIN REDITABS
>A> + SCHERING PLOUGH 5MG N21993 001 Dec 12, 2006 Dec 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
>A> + JOHNSON AND JOHNSON 5% N21812 001 Jan 20, 2006 Dec CAHN
>D> + PHARMACIA AND UPJOHN 5% N21812 001 Jan 20, 2006 Dec CAHN
+ 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
ROGAINE (FOR MEN)
>A> + JOHNSON AND JOHNSON 2% N19501 002 Feb 09, 1996 Dec CAHN
>D> + PHARMACIA AND UPJOHN 2% N19501 002 Feb 09, 1996 Dec CAHN
ROGAINE (FOR WOMEN)
>A> + JOHNSON AND JOHNSON 2% N19501 003 Feb 09, 1996 Dec CAHN
>D> + PHARMACIA AND UPJOHN 2% N19501 003 Feb 09, 1996 Dec CAHN
ROGAINE EXTRA STRENGTH (FOR MEN)
>A> + JOHNSON AND JOHNSON 5% N20834 001 Nov 14, 1997 Dec CAHN
>D> + PHARMACIA AND UPJOHN 5% N20834 001 Nov 14, 1997 Dec CAHN

NAPHAZOLINE HYDROCHLORIDE; PHENIRAMINE MALEATE

SOLUTION/DROPS; OPHTHALMIC
VISINE-A
>A> JOHNSON AND JOHNSON 0.025%;0.3% N20485 001 Jan 31, 1996 Dec CAHN
>D> PFIZER 0.025%;0.3% N20485 001 Jan 31, 1996 Dec CAHN

NAPROXEN SODIUM

CAPSULE; ORAL

NAPROXEN SODIUM

+	BANNER PHARMACAPS	EQ 200MG BASE	N21920	001	Feb 17, 2006	Oct	CRLD
		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

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

NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

NICOTROL

>A>	@ MCNEIL CONS	15MG/16HR	N20536	001	Jul 03, 1996	Dec	CAHN
>D>	@ PHARMACIA AND UPJOHN	15MG/16HR	N20536	001	Jul 03, 1996	Dec	CAHN

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICORETTE

GLAXOSMITHKLINE	EQ 2MG BASE	N18612	004	Sep 25, 2000	Mar	CRLD
	EQ 2MG BASE	N18612	003	Dec 23, 1998	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
PERRIGO R AND D	EQ 2MG BASE	N78325	001	Oct 30, 2006	Oct	NEWA
	EQ 4MG BASE	N78326	001	Oct 30, 2006	Oct	NEWA
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

OXYMETAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

VISINE L.R.

>A>	+ JOHNSON AND JOHNSON	0.025%	N19407	001	Mar 31, 1989	Dec	CAHN
>D>	+ PFIZER	0.025%	N19407	001	Mar 31, 1989	Dec	CAHN

PERMETHRIN

LOTION; TOPICAL

PERMETHRIN

ACTAVIS MID ATLANTIC	1%	N75014	001	Mar 28, 2000	Jun	CAHN

POLYETHYLENE GLYCOL 3350

FOR SOLUTION; ORAL
MIRALAX

+	BRAINTREE	17GM/SCOOPFUL	N22015	001	Oct 06, 2006	Oct	NEWA
+	SCHERING PLOUGH	17GM/SCOOPFUL	N22015	001	Oct 06, 2006	Nov	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, EFFERVESCENT; ORAL
ZANTAC 75

>A>	@	MCNEIL CONS	EQ 75MG BASE	N20745	001	Feb 26, 1998	Dec	CAHN
>D>	@	WARNER LAMBERT	EQ 75MG BASE	N20745	001	Feb 26, 1998	Dec	CAHN

TABLET; ORAL
RANITIDINE

		WOCKHARDT	EQ 75MG BASE	N76760	001	Feb 24, 2006	Feb	NEWA

ZANTAC 150

>A>	+	MCNEIL CONS	EQ 150MG BASE	N21698	001	Aug 31, 2004	Dec	CAHN
>D>	+	PFIZER CONS HLTHCARE	EQ 150MG BASE	N21698	001	Aug 31, 2004	Dec	CAHN

ZANTAC 75

>A>		MCNEIL CONS	EQ 75MG BASE	N20520	001	Dec 19, 1995	Dec	CAHN
>D>		WARNER LAMBERT	EQ 75MG BASE	N20520	001	Dec 19, 1995	Dec	CAHN

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 12 DECEMBER 2006

NO DECEMBER 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 DECEMBER 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 EXPIRATON DATE	PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ABACAVIR SULFATE; LAMIVUDINE - EPZICOM						
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				
ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE - TRIZIVIR						
021205 001	5034394	Dec 18, 2011	DS	DP		
	5034394*PED	Jun 18, 2012				
ACETAMINOPHEN; TRAMADOL HYDROCHLORIDE - ULTRACET						
021123 001	RE39221	Aug 09, 2011	DS	DP U-55		
ALBUTEROL SULFATE - PROAIR HFA						
021457 001					I-235	Feb 03, 2009
ALBUTEROL SULFATE - VENTOLIN HFA						
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		
ALENDRONATE SODIUM; CHOLECALCIFEROL - FOSAMAX PLUS D						
021762 001					NC	Apr 07, 2008
ALFUZOSIN HYDROCHLORIDE - UROXATRAL						
021287 001	4661491	May 27, 2007		U-706		
ALITRETINOIN - PANRETIN						
020886 001	7056954	Aug 02, 2012		DP		
AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL						
020364 006	4879303	Mar 25, 2007	DS	DP	NS	Apr 11, 2009
	6162802	Dec 19, 2017	DS	DP U-185		
AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE - LOTREL						
020364 007	4879303	Mar 25, 2007	DS	DP	NS	Apr 11, 2009
	6162802	Dec 19, 2017	DS	DP U-185		
ANIDULAFUNGIN - ERAXIS						
021632 001	5965525	Oct 12, 2016	DS	DP U-540	NCE	Feb 17, 2011
	6384013	Mar 19, 2012	DS			
	6743777	Mar 19, 2012		DP U-540		
	6960564	Apr 12, 2021		DP U-540		
ANIDULAFUNGIN - ERAXIS						
021632 002					**>A>** NCE	Feb 17, 2011
APREPITANT - EMEND						
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		
APREPITANT - EMEND						
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 EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
APREPITANT - EMEND							
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		
ARFORMOTEROL TARTRATE - BROVANA							
021912 001						NP	Oct 06, 2009
ARIPIPRAZOLE - ABILIFY							
021436 001	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
ARIPIPRAZOLE - ABILIFY							
021436 002	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
ARIPIPRAZOLE - ABILIFY							
021436 003	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
ARIPIPRAZOLE - ABILIFY							
021436 004	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
ARIPIPRAZOLE - ABILIFY							
021436 005	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
ARIPIPRAZOLE - ABILIFY							
021436 006	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
ARIPIPRAZOLE - ABILIFY							
021713 001	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
	6977257	Apr 24, 2022	DS	DP			
ARIPIPRAZOLE - ABILIFY							
021729 002	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
						I-437	Sep 29, 2007
						NCE	Nov 15, 2007
ARIPIPRAZOLE - ABILIFY							
021729 003	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
						I-437	Sep 29, 2007
						NCE	Nov 15, 2007
ARIPIPRAZOLE - ABILIFY							
021729 004	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
						I-437	Sep 29, 2007
						NCE	Nov 15, 2007
ARIPIPRAZOLE - ABILIFY							
021729 005	5006528	Oct 20, 2014	DS	DP	U-761	I-488	Mar 01, 2008
						I-437	Sep 29, 2007
						NCE	Nov 15, 2007
ARIPIPRAZOLE - ABILIFY							
021866 001	5006528	Oct 20, 2014	DS	DP	U-763	NCE	Nov 15, 2007
	7115587	Jul 21, 2024	DS	DP	U-764	NDF	Sep 20, 2009
ARTICAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE - SEPTOCAINE							
022010 001						NP	Mar 30, 2009
ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP							
021881 001						NP	Aug 02, 2009
ATAZANAVIR SULFATE - REYATAZ							
021567 001	5849911	Apr 09, 2017	DS	DP	U-167		
	6087383	Dec 21, 2018	DS	DP			
ATAZANAVIR SULFATE - REYATAZ							
021567 002	5849911	Apr 09, 2017	DS	DP	U-167		
	6087383	Dec 21, 2018	DS	DP			
ATAZANAVIR SULFATE - REYATAZ							
021567 003	5849911	Apr 09, 2017	DS	DP	U-167		
	6087383	Dec 21, 2018	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 EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ATAZANAVIR SULFATE - REYATAZ					
021567 004	5849911	Apr 09, 2017	DS DP U-167	D-89	Jul 06, 2007
	6087383	Dec 21, 2018	DS DP	NCE	Jun 20, 2008
ATOMOXETINE HYDROCHLORIDE - STRATTERA					
021411 001	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
ATOMOXETINE HYDROCHLORIDE - STRATTERA					
021411 002	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
ATOMOXETINE HYDROCHLORIDE - STRATTERA					
021411 003	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
ATOMOXETINE HYDROCHLORIDE - STRATTERA					
021411 004	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
ATOMOXETINE HYDROCHLORIDE - STRATTERA					
021411 005	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
ATOMOXETINE HYDROCHLORIDE - STRATTERA					
021411 006	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
ATOMOXETINE HYDROCHLORIDE - STRATTERA					
021411 007	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
ATOMOXETINE HYDROCHLORIDE - STRATTERA					
021411 008	5658590	Nov 26, 2016	U-494		
	5658590*PED	May 26, 2017	U-494		
ATROPINE; PRALIDOXIME CHLORIDE - DUODOTE					
021983 001	5092843	Apr 12, 2010	DP		
AVOBENZONE; ECAMSULE; OCTOCRYLENE - ANTHELIOS SX					
021502 001	4585597	Jun 16, 2007	DS DP U-752	NC	Jul 21, 2009
	5587150	Dec 24, 2013	DP U-752		
AZELASTINE HYDROCHLORIDE - ASTELIN					
020114 001				D-102	Feb 17, 2009
BALSALAZIDE DISODIUM - COLAZAL					
020610 001	4412992*PED	Jan 08, 2007		>A> NPP	Dec 20, 2009
				>A> PED	Jun 20, 2010
BETAMETHASONE DIPROPIONATE; CALCIPOTRIENE HYDRATE - TACLONEX					
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		
BETAMETHASONE VALERATE - LUXIQ					
020934 001	7078058	Mar 01, 2016	DS DP		
BETAXOLOL HYDROCHLORIDE - BETOPTIC S					
019845 001	4911920	Mar 27, 2007			
	4911920*PED	Sep 27, 2007			
BIMATOPROST - LUMIGAN					
021275 001	5688819	Aug 19, 2014	U-446		
	6403649	Sep 21, 2012	DS DP U-446		
BIVALIRUDIN - ANGIOMAX					
020873 001				I-486	Nov 30, 2008
BORTEZOMIB - VELCADE					
021602 001				>A> I-521	Dec 08, 2009
				ODE	Mar 25, 2012

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
BRIMONIDINE TARTRATE - ALPHAGAN P							
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					
BRINZOLAMIDE - AZOPT							
020816 001	5240923	Aug 31, 2010	DS	DP	U-224	M-54	Sep 28, 2009
	5240923*PED	Mar 01, 2011				PED	Mar 28, 2010
	5378703	Apr 01, 2012	DS	DP	U-224		
	5378703*PED	Oct 01, 2012					
	5461081	Oct 24, 2012		DP	U-225		
	5461081*PED	Apr 24, 2013					
BROMFENAC SODIUM - XIBROM							
021664 001						I-485	Jan 27, 2009
BUDESONIDE - BUDESONIDE							
021949 001	**>A>** 4907583	Mar 13, 2007		DP		NP	Jul 12, 2009
	>A> 6027714	Jan 09, 2018		DP	U-787		
	>A> 6142145	May 08, 2018		DP			
	>A> 6287540	Jan 09, 2018		DP			
	>A> 7143764	Mar 13, 2018		DP			
BUDESONIDE - BUDESONIDE							
021949 002	**>A>** 4907583	Mar 13, 2007		DP		NP	Jul 12, 2009
	>A> 6027714	Jan 09, 2018		DP	U-787		
	>A> 6142145	May 08, 2018		DP			
	>A> 6287540	Jan 09, 2018		DP			
	>A> 7143764	Mar 13, 2018		DP			
BUDESONIDE - PULMICORT RESPULES							
020929 001	6598603*PED	Jun 23, 2019					
	6899099	Dec 23, 2018			U-751		
	6899099*PED	Jun 23, 2019					
BUDESONIDE - PULMICORT RESPULES							
020929 002	6899099	Dec 23, 2018			U-751		
	6899099*PED	Jun 23, 2019					
BUDESONIDE - RHINOCORT							
020746 001	6986904	Apr 29, 2017		DP	U-699		
BUDESONIDE - RHINOCORT							
020746 002	6986904	Apr 29, 2017		DP	U-699		
BUDESONIDE; FORMOTEROL FUMARATE - SYMBICORT							
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			
BUDESONIDE; FORMOTEROL FUMARATE - SYMBICORT							
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			
BUPROPION HYDROCHLORIDE - WELLBUTRIN XL							
021515 001						I-497	Jun 12, 2009
BUPROPION HYDROCHLORIDE - WELLBUTRIN XL							
021515 002						I-497	Jun 12, 2009

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
BUSULFAN - BUSULFEX							
020954 001	5559148	Sep 30, 2013			U-264		
	5559148*PED	Mar 30, 2014			U-264		
CALCIUM CARBONATE; RISEDRONATE SODIUM - ACTONEL WITH CALCIUM (COPACKAGED)							
021823 001						M-52	Jan 24, 2009
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 100							
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		
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 150							
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		
CARBIDOPA; ENTACAPONE; LEVODOPA - STALEVO 50							
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		
CARVEDILOL - COREG							
020297 001	4503067	Mar 05, 2007			U-3	M-56	Aug 28, 2009
	4503067*PED	Sep 05, 2007				PED	Feb 28, 2010
	5760069	Jun 07, 2015			U-233		
	5760069*PED	Dec 07, 2015					
	5902821	Feb 07, 2016			U-313		
	5902821*PED	Aug 07, 2016					
CARVEDILOL - COREG							
020297 002	4503067	Mar 05, 2007			U-3	M-56	Aug 28, 2009
	4503067*PED	Sep 05, 2007				PED	Feb 28, 2010
	5760069	Jun 07, 2015			U-233		
	5760069*PED	Dec 07, 2015					
	5902821	Feb 07, 2016			U-313		
	5902821*PED	Aug 07, 2016					
CARVEDILOL - COREG							
020297 003	4503067	Mar 05, 2007			U-3	M-56	Aug 28, 2009
	4503067*PED	Sep 05, 2007				PED	Feb 28, 2010
	5760069	Jun 07, 2015			U-233		
	5760069*PED	Dec 07, 2015					
	5902821	Feb 07, 2016			U-313		
	5902821*PED	Aug 07, 2016					
CARVEDILOL - COREG							
020297 004	4503067	Mar 05, 2007			U-3	M-56	Aug 28, 2009
	4503067*PED	Sep 05, 2007				PED	Feb 28, 2010
	5760069	Jun 07, 2015			U-233		
	5760069*PED	Dec 07, 2015					
	5902821	Feb 07, 2016			U-313		
	5902821*PED	Aug 07, 2016					

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
CARVEDILOL PHOSPHATE - COREG CR					
022012 001	>A> 4503067	Mar 05, 2007	DS DP U-3	>A> M-56	Aug 28, 2009
	>A> 4503067*PED	Sep 05, 2007		>A> NDF	Oct 20, 2009
	>A> 5760069	Jun 07, 2015	U-777	>A> PED	Feb 28, 2010
	>A> 5760069*PED	Dec 07, 2015		>A> PED	Apr 20, 2010
	>A> 5902821	Feb 07, 2016	U-313		
	>A> 5902821	Feb 07, 2016	U-777		
	>A> 5902821*PED	Aug 07, 2016			
	>A> 6022562	Oct 17, 2015	DP		
	>A> 6022562*PED	Apr 17, 2016			
CARVEDILOL PHOSPHATE - COREG CR					
022012 002	>A> 4503067	Mar 05, 2007	DS DP U-3	>A> M-56	Aug 28, 2009
	>A> 4503067*PED	Sep 05, 2007		>A> NDF	Oct 20, 2009
	>A> 5760069	Jun 07, 2015	U-777	>A> PED	Feb 28, 2010
	>A> 5760069*PED	Dec 07, 2015		>A> PED	Apr 20, 2010
	>A> 5902821	Feb 07, 2016	U-777		
	>A> 5902821	Feb 07, 2016	U-313		
	>A> 5902821*PED	Aug 07, 2016			
	>A> 6022562	Oct 17, 2015	DP		
	>A> 6022562*PED	Apr 17, 2016			
CARVEDILOL PHOSPHATE - COREG CR					
022012 003	>A> 4503067	Mar 05, 2007	DS DP U-3	>A> M-56	Aug 28, 2009
	>A> 4503067*PED	Sep 05, 2007		>A> NDF	Oct 20, 2009
	>A> 5760069	Jun 07, 2015	U-777	>A> PED	Feb 28, 2010
	>A> 5760069*PED	Dec 07, 2015		>A> PED	Apr 20, 2010
	>A> 5902821	Feb 07, 2016	U-777		
	>A> 5902821	Feb 07, 2016	U-313		
	>A> 5902821*PED	Aug 07, 2016			
	>A> 6022562	Oct 17, 2015	DP		
	>A> 6022562*PED	Apr 17, 2016			
CARVEDILOL PHOSPHATE - COREG CR					
022012 004	>A> 4503067	Mar 05, 2007	DS DP U-3	>A> M-56	Aug 28, 2009
	>A> 4503067*PED	Sep 05, 2007		>A> NDF	Oct 20, 2009
	>A> 5760069	Jun 07, 2015	U-777	>A> PED	Feb 28, 2010
	>A> 5760069*PED	Dec 07, 2015		>A> PED	Apr 20, 2010
	>A> 5902821	Feb 07, 2016	U-777		
	>A> 5902821	Feb 07, 2016	U-313		
	>A> 5902821*PED	Aug 07, 2016			
	>A> 6022562	Oct 17, 2015	DP		
	>A> 6022562*PED	Apr 17, 2016			
CASPOFUNGIN ACETATE - CANCIDAS					
021227 001	5514650	Jan 26, 2015	DP U-607		
CASPOFUNGIN ACETATE - CANCIDAS					
021227 002	5514650	Jan 26, 2015	DP U-607		
CEFDITOREN PIVOXIL - SPECTRACEF					
021222 001	4839350	Apr 14, 2009	DS DP		
CELECOXIB - CELEBREX					
020998 001	5466823	Nov 30, 2013	DS	I-466	Jul 29, 2008
	5466823*PED	May 30, 2014		>A> NPP	Dec 15, 2009
	5563165	Nov 30, 2013	DP	>A> PED	Jun 15, 2010
	5563165*PED	May 30, 2014		PED	Jan 29, 2009
	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			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
CELECOXIB - CELEBREX					
020998 002	5466823	Nov 30, 2013	DS	I-466	Jul 29, 2008
	5466823*PED	May 30, 2014		>A> NPP	Dec 15, 2009
	5563165	Nov 30, 2013	DP	>A> PED	Jun 15, 2010
	5563165*PED	May 30, 2014		PED	Jan 29, 2009
	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			
CELECOXIB - CELEBREX					
020998 003	5466823	Nov 30, 2013	DS	I-466	Jul 29, 2008
	5466823*PED	May 30, 2014		>A> NPP	Dec 15, 2009
	5563165	Nov 30, 2013	DP	>A> PED	Jun 15, 2010
	5563165*PED	May 30, 2014		PED	Jan 29, 2009
	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			
CETIRIZINE HYDROCHLORIDE - ZYRTEC					
019835 001	4525358	Jun 25, 2007	DS DP U-565		
	4525358*PED	Dec 25, 2007			
CETIRIZINE HYDROCHLORIDE - ZYRTEC					
019835 002	4525358	Jun 25, 2007	DS DP U-565		
	4525358*PED	Dec 25, 2007			
CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ZYRTEC-D 12 HOUR					
021150 001	7014867	Jun 10, 2022	DP		
CHLORHEXIDINE GLUCONATE - CHLORHEXIDINE GLUCONATE					
021669 001	7066916	Feb 17, 2024	U-737		
CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP					
020832 004	5690958	Sep 30, 2016	DP		
	6536975	Nov 10, 2020	DP		
CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP FREPP					
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		
CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP SINGLE SWABSTICK					
021555 002	5690958	Sep 30, 2016	DP		
CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT					
020832 002	6991393	Mar 14, 2023	DP		
CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT					
020832 005	5690958	Sep 30, 2016	DP		
	6536975	Nov 10, 2020	DP		
	6729786	Mar 14, 2023	DP		
	6991393	Jan 31, 2024	DP		
CICLESONIDE - OMNARIS					
022004 001				NCE	Oct 20, 2011
CICLOPIROX - LOPROX					
020519 001	7018656	Sep 05, 2018	DP		
	7026337	Apr 02, 2018	U-714		
CIPROFLOXACIN - CIPRO					
019847 002				I-421	Mar 25, 2007
				PED	Sep 25, 2007

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
CIPROFLOXACIN - CIPRO					
019847 003				I-421	Mar 25, 2007
				PED	Sep 25, 2007
CLOBETASOL PROPIONATE - CLOBEX					
021835 001	5972920	Feb 12, 2018	DP	NDF	Oct 27, 2008
	5990100	Mar 24, 2018	DP U-742		
CLOPIDOGREL BISULFATE - CLOPIDOGREL BISULFATE					
076274 001				PC	Feb 04, 2007
CLOPIDOGREL BISULFATE - PLAVIX					
020839 001				I-502	Aug 17, 2009
COLESEVELAM HYDROCHLORIDE - WELCHOL					
021176 001	5693675	Dec 02, 2014	DS		
	5917007	Apr 29, 2014	DS U-323		
	5919832	Jun 10, 2014	DS		
	6066678	Jun 10, 2014	DS U-323		
	6433026	Jun 10, 2014	DS		
	6784254	Jun 10, 2014	DS DP		
	7101960	Apr 29, 2014	DS DP U-757		
CONIVAPTAN HYDROCHLORIDE - VAPRISOL					
021697 001	5723606	Mar 03, 2015	DS DP U-698		
DAPSONE - ACZONE					
021794 001	5863560	Sep 11, 2016	DP		
	6620435	Sep 11, 2016	DP		
DAPTOMYCIN - CUBICIN					
021572 001	6468967	Sep 24, 2019	U-282		
	RE39071	Jun 15, 2016	DS DP U-728		
DAPTOMYCIN - CUBICIN					
021572 002	6468967	Sep 24, 2019	U-282	I-505	May 25, 2009
	RE39071	Jun 15, 2016	DS DP U-728		
DARUNAVIR ETHANOLATE - PREZISTA					
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		
DASATINIB - SPRYCEL					
021986 001	6596746	Apr 13, 2020	DS DP U-748	NCE	Jun 28, 2011
	7125875	Apr 13, 2020	U-780	ODE	Jun 28, 2013
	7125875	Apr 13, 2020	U-779	ODE	Jun 28, 2013
DASATINIB - SPRYCEL					
021986 002	6596746	Apr 13, 2020	DS DP U-748	NCE	Jun 28, 2011
	7125875	Apr 13, 2020	U-780	ODE	Jun 28, 2013
	7125875	Apr 13, 2020	U-779	ODE	Jun 28, 2013
DASATINIB - SPRYCEL					
021986 003	6596746	Apr 13, 2020	DS DP U-748	NCE	Jun 28, 2011
	7125875	Apr 13, 2020	U-780	ODE	Jun 28, 2013
	7125875	Apr 13, 2020	U-779	ODE	Jun 28, 2013
DECITABINE - DACOGEN					
021790 001				NCE	May 02, 2011
DEFERASIROX - EXJADE					
021882 001	6465504	Jun 24, 2017	DS DP		
	6596750	Jun 24, 2017	DS U-735		
DEFERASIROX - EXJADE					
021882 002	6465504	Jun 24, 2017	DS DP		
	6596750	Jun 24, 2017	DS U-735		
DEFERASIROX - EXJADE					
021882 003	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 EXPIRATON DATE		PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
DESFLURANE - SUPRANE						
020118 001	4762856	Feb 02, 2007		U-67		
	4762856*PED	Aug 02, 2007				
	5617906	Apr 08, 2014		DP		
	5617906*PED	Oct 08, 2014				
DESIRUDIN RECOMBINANT - IPRIVASK						
021271 001	4745177	May 17, 2010	DS	DP		
DESLORATADINE - CLARINEX						
021165 001	4659716	Mar 31, 2007		DP U-725		
	4659716	Mar 31, 2007		DP U-427		
	4659716*PED	Oct 01, 2007		U-427		
DESLORATADINE - CLARINEX						
021300 001	4659716	Mar 31, 2007		DP U-725		
	4659716	Mar 31, 2007		DP U-611		
	4659716*PED	Oct 01, 2007				
DESLORATADINE - CLARINEX						
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		
DESLORATADINE - CLARINEX						
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		
DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX D 24 HOUR						
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		
DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX-D 12 HOUR						
021313 001	4659716	Mar 31, 2007		DP U-707	NP	Feb 01, 2009
	4659716*PED	Oct 01, 2007			NC	Mar 03, 2008
	6100274	Jul 07, 2019		DP	NCE	Dec 21, 2006
	6100274*PED	Jan 07, 2020			PED	Jun 21, 2007
	6709676	Feb 18, 2021		DP U-707		
DESMOPRESSIN ACETATE - DDAVP						
019955 001	7022340	Apr 30, 2023		DP		
DESMOPRESSIN ACETATE - DDAVP						
019955 002	7022340	Apr 30, 2023		DP		
DESONIDE - DESONATE						
021844 001	6387383	Aug 03, 2020	DS	DP U-783		
DESONIDE - VERDESO						
021978 001	6730288	Sep 08, 2019		DP	NDF	Sep 19, 2009
	7029659	Sep 08, 2019		DP		
DEXMEDETOMIDINE - PRECEDEX						
021038 001	4910214	Jul 15, 2013	DS	DP U-421		
DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR						
021802 004					NDF	May 26, 2008
DIAZEPAM - DIASTAT						
020648 001	5462740	Sep 17, 2013		DP		
DIAZEPAM - DIASTAT						
020648 002	5462740	Sep 17, 2013		DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
DIAZEPAM - DIASTAT						
020648 003	5462740	Sep 17, 2013		DP		
DIAZEPAM - DIASTAT						
020648 004	5462740	Sep 17, 2013		DP		
DIAZEPAM - DIASTAT						
020648 005	5462740	Sep 17, 2013		DP		
DIAZEPAM - DIASTAT ACUDIAL						
020648 006	5462740	Sep 17, 2013		DP		
DIAZEPAM - DIASTAT ACUDIAL						
020648 007	5462740	Sep 17, 2013		DP		
DILTIAZEM HYDROCHLORIDE - CARDIZEM LA						
021392 001	7108866	Dec 17, 2019		DP U-107		
DILTIAZEM HYDROCHLORIDE - CARDIZEM LA						
021392 002	7108866	Dec 17, 2019		DP U-107		
DILTIAZEM HYDROCHLORIDE - CARDIZEM LA						
021392 003	7108866	Dec 17, 2019		DP U-107		
DILTIAZEM HYDROCHLORIDE - CARDIZEM LA						
021392 004	7108866	Dec 17, 2019		DP U-107		
DILTIAZEM HYDROCHLORIDE - CARDIZEM LA						
021392 005	7108866	Dec 17, 2019		DP U-107		
DILTIAZEM HYDROCHLORIDE - CARDIZEM LA						
021392 006	7108866	Dec 17, 2019		DP U-107		
DOCETAXEL - TAXOTERE						
020449 001	4814470	May 14, 2010	DS	DP	I-519	Oct 17, 2009
	5438072	Nov 22, 2013		DP	I-490	Mar 22, 2009
	5698582	Jul 03, 2012		DP		
	5714512	Jul 03, 2012		DP		
	5750561	Jul 03, 2012		DP		
DONEPEZIL HYDROCHLORIDE - ARICEPT ODT						
021720 001	4895841	Nov 25, 2010	DS	DP U-713		
DONEPEZIL HYDROCHLORIDE - ARICEPT ODT						
021720 002	4895841	Nov 25, 2010	DS	DP U-713		
DROSPIRENONE; ETHINYL ESTRADIOL - YAZ						
021676 001	5569652	Oct 29, 2013		U-758	I-508	Oct 04, 2009
	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		
	6987101	Dec 22, 2017		U-758		
	6987101	Dec 22, 2017		U-1		
	RE37564	Jun 30, 2014		DP		
	RE37838	Jun 30, 2014		DP		
	RE38253	Jun 30, 2014		DP		
EFAVIRENZ - SUSTIVA						
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		
EFAVIRENZ - SUSTIVA						
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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
EFAVIRENZ - SUSTIVA					
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		
EFAVIRENZ - SUSTIVA					
021360 002	5519021	May 21, 2013	DS DP		
EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA					
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		
EMTRICITABINE - EMTRIVA					
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			
EMTRICITABINE - EMTRIVA					
021896 001	5210085	May 11, 2010	U-257	**>A>** NPP	Dec 22, 2009
	5210085*PED	Nov 11, 2010		NDF	Sep 27, 2008
	5814639	Sep 29, 2015	DS DP	NCE	Jul 02, 2008
	5814639*PED	Mar 29, 2016		**>A>** PED	Jun 22, 2010
	5914331	Sep 29, 2015	DS	PED	Mar 27, 2009
	5914331*PED	Mar 29, 2016		PED	Jan 02, 2009
	6642245	Nov 04, 2020	U-257		
	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 EXPIRATON DATE			PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - TRUVADA							
021752 001	5210085	May 11, 2010			U-248	NCE	Jul 02, 2008
	5210085	May 11, 2010			U-541	PED	Jan 02, 2009
	5210085*PED	Nov 11, 2010					
	5814639	Sep 29, 2015	DS	DP			
	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					
ENFUVIRTIDE - FUZEON							
021481 001						M-59	Sep 29, 2009
ENTACAPONE - COMTAN							
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		
EPLERENONE - INSPRA							
021437 001	4559332	Apr 09, 2007	DS	DP	U-537		
EPLERENONE - INSPRA							
021437 002	4559332	Apr 09, 2007	DS	DP	U-537		
EPLERENONE - INSPRA							
021437 003	4559332	Apr 09, 2007	DS	DP	U-537		
ERTAPENEM SODIUM - INVANZ							
021337 001						I-515	Aug 10, 2009
ESCITALOPRAM OXALATE - LEXAPRO							
021323 001	RE34712	Sep 14, 2011	DS	DP			
	RE34712*PED	Mar 14, 2012					
ESCITALOPRAM OXALATE - LEXAPRO							
021323 002	RE34712	Sep 14, 2011	DS	DP			
	RE34712*PED	Mar 14, 2012					
ESCITALOPRAM OXALATE - LEXAPRO							
021323 003	RE34712	Sep 14, 2011	DS	DP			
	RE34712*PED	Mar 14, 2012					
ESCITALOPRAM OXALATE - LEXAPRO							
021365 001	RE34712	Sep 14, 2011	DS	DP			
	RE34712*PED	Mar 14, 2012					

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ESOMEPRAZOLE MAGNESIUM - NEXIUM							
021153 001	4738974	Apr 19, 2007	DS	DP	U-770	I-504	Oct 11, 2009
	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		
	4786505	Apr 20, 2007		DP	U-770		
	4853230	Apr 20, 2007		DP	U-373		
	4853230	Apr 20, 2007		DP	U-770		
	4853230	Apr 20, 2007		DP	U-729		
	5690960	Nov 25, 2014		DP	U-770		
	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-770		
	5714504	Feb 03, 2015		DP	U-373		
	5877192	May 27, 2014		DP	U-373		
	5877192	May 27, 2014		DP	U-770		
	5877192	May 27, 2014		DP	U-729		
	5900424	May 04, 2016	DS		U-770		
	5900424	May 04, 2016	DS		U-729		
	5900424	May 04, 2016	DS		U-373		
	6369085	May 25, 2018	DS	DP	U-729		
	6369085	May 25, 2018	DS	DP	U-770		
	6428810	Nov 03, 2019		DP	U-469		
	6428810	Nov 03, 2019		DP	U-770		
	6428810	Nov 03, 2019		DP	U-729		
ESOMEPRAZOLE MAGNESIUM - NEXIUM							
021153 002	4738974	Apr 19, 2007	DS	DP	U-635	I-504	Oct 11, 2009
	4738974	Apr 19, 2007	DS	DP	U-770	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-770		
	4786505	Apr 20, 2007		DP	U-729		
	4853230	Apr 20, 2007		DP	U-373		
	4853230	Apr 20, 2007		DP	U-729		
	4853230	Apr 20, 2007		DP	U-770		
	5690960	Nov 25, 2014		DP	U-770		
	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-770		
	5714504	Feb 03, 2015		DP	U-373		
	5877192	May 27, 2014		DP	U-373		
	5877192	May 27, 2014		DP	U-770		
	5877192	May 27, 2014		DP	U-729		
	5900424	May 04, 2016	DS		U-729		
	5900424	May 04, 2016	DS		U-770		
	5900424	May 04, 2016	DS		U-373		
	6369085	May 25, 2018	DS	DP	U-729		
	6369085	May 25, 2018	DS	DP	U-770		
	6428810	Nov 03, 2019		DP	U-469		
	6428810	Nov 03, 2019		DP	U-770		
	6428810	Nov 03, 2019		DP	U-729		

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ESOMEPRAZOLE MAGNESIUM - NEXIUM					
021957 001	4738974	Apr 19, 2007	DS DP U-773	I-504	Oct 11, 2009
	4738974	Apr 19, 2007	DS DP U-729	I-484	Nov 24, 2007
	4783974*PED	Oct 19, 2007		NPP	Apr 28, 2009
	4786505	Apr 20, 2007	DP U-729		
	4786505	Apr 20, 2007	DP U-773		
	4786505*PED	Oct 20, 2007			
	4853230	Apr 20, 2007	DP U-729		
	4853230	Apr 20, 2007	DP U-773		
	4853230*PED	Oct 20, 2007			
	5690960	Nov 25, 2014	DP U-729		
	5690960	Nov 25, 2014	DP U-773		
	5690960*PED	May 25, 2015			
	5714504	Feb 03, 2015	DP U-773		
	5714504	Feb 03, 2015	DP U-729		
	5714504*PED	Aug 03, 2015			
	5877192	May 27, 2014	U-773		
	5877192	May 27, 2014	U-729		
	5877192*PED	Nov 27, 2014			
	5900424	May 04, 2016	DS U-729		
	5900424	May 04, 2016	DS U-773		
	5900424*PED	Nov 04, 2016			
	6369085	May 25, 2018	DS DP U-729		
	6369085	May 25, 2018	DS DP U-773		
	6369085*PED	Nov 25, 2018			
	6428810	Nov 03, 2019	DP U-729		
	6428810	Nov 03, 2019	DP U-773		
	6428810*PED	May 03, 2020			
	6875872	May 27, 2014	DS		
	6875872*PED	Nov 27, 2014			
ESOMEPRAZOLE MAGNESIUM - NEXIUM					
021957 002	4738974	Apr 19, 2007	DS DP U-729	I-504	Oct 11, 2009
	4738974	Apr 19, 2007	DS DP U-773	I-484	Nov 24, 2007
	4783974*PED	Oct 19, 2007		NPP	Apr 28, 2009
	4786505	Apr 20, 2007	DP U-773		
	4786505	Apr 20, 2007	DP U-729		
	4786505*PED	Oct 20, 2007			
	4853230	Apr 20, 2007	DP U-773		
	4853230	Apr 20, 2007	DP U-729		
	4853230*PED	Oct 20, 2007			
	5690960	Nov 25, 2014	DP U-773		
	5690960	Nov 25, 2014	DP U-729		
	5690960*PED	May 25, 2015			
	5714504	Feb 03, 2015	DP U-773		
	5714504	Feb 03, 2015	DP U-729		
	5714504*PED	Aug 03, 2015			
	5877192	May 27, 2014	U-729		
	5877192	May 27, 2014	U-773		
	5877192*PED	Nov 27, 2014			
	5900424	May 04, 2016	DS U-773		
	5900424	May 04, 2016	DS U-729		
	5900424*PED	Nov 04, 2016			
	6369085	May 25, 2018	DS DP U-773		
	6369085	May 25, 2018	DS DP U-729		
	6369085*PED	Nov 25, 2018			
	6428810	Nov 03, 2019	DP U-773		
	6428810	Nov 03, 2019	DP U-729		
	6428810*PED	May 03, 2020			
	6875872	May 27, 2014	DS		
	6875872*PED	Nov 27, 2014			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ESTRADIOL - ELESTRIN							
021813 001						**>A>** NP	Dec 15, 2009
ESTRADIOL - ESTRADIOL							
075182 004						PC	Feb 06, 2007
ESTRADIOL - ESTRADIOL							
075182 005						PC	Feb 06, 2007
ESTRADIOL; NORETHINDRONE ACETATE - ACTIVELLA							
020907 002						**>A>** D-104	Dec 29, 2009
						>A> NS	Dec 28, 2009
ETHINYL ESTRADIOL; LEVONORGESTREL - SEASONIQUE							
021840 001						NP	May 25, 2009
ETHINYL ESTRADIOL; NORELGESTROMIN - ORTHO EVRA							
021180 001	5876746	Nov 20, 2015		DP	U-514		
ETHINYL ESTRADIOL; NORETHINDRONE - OVCON-35							
021490 001	6667050	Apr 06, 2019		DP	U-1		
ETHINYL ESTRADIOL; NORETHINDRONE ACETATE - LOESTRIN 24 FE							
021871 001	5552394	Jul 22, 2014			U-1	NP	Feb 17, 2009
ETONOGESTREL - IMPLANON							
021529 001	4957119	Aug 05, 2008		DP		NDF	Jul 17, 2009
	5150718	Sep 29, 2009			U-749		
EXEMESTANE - AROMASIN							
020753 001						I-495	Oct 05, 2008
EXENATIDE SYNTHETIC - BYETTA							
021773 001						**>A>** I-520	Dec 22, 2009
EXENATIDE SYNTHETIC - BYETTA							
021773 002						**>A>** I-520	Dec 22, 2009
EZETIMIBE - ZETIA							
021445 001	7030106	Jan 25, 2022		DP		I-493	May 23, 2009
	RE37721	Oct 25, 2016	DS	DP	U-473	M-57	Mar 16, 2009
EZETIMIBE; SIMVASTATIN - VYTORIN							
021687 001	RE37721	Oct 25, 2016	DS	DP	U-473	M-60	Oct 03, 2009
						M-58	Mar 16, 2009
EZETIMIBE; SIMVASTATIN - VYTORIN							
021687 002	RE37721	Oct 25, 2016	DS	DP	U-473	M-60	Oct 03, 2009
						M-58	Mar 16, 2009
EZETIMIBE; SIMVASTATIN - VYTORIN							
021687 003	RE37721	Oct 25, 2016	DS	DP	U-473	M-60	Oct 03, 2009
						M-58	Mar 16, 2009
EZETIMIBE; SIMVASTATIN - VYTORIN							
021687 004	RE37721	Oct 25, 2016	DS	DP	U-473	M-60	Oct 03, 2009
						M-58	Mar 16, 2009
FAMCICLOVIR - FAMVIR							
020363 001						D-103	Jul 28, 2009
						I-501	Jul 28, 2009
FAMCICLOVIR - FAMVIR							
020363 002						D-103	Jul 28, 2009
						I-501	Jul 28, 2009
FAMCICLOVIR - FAMVIR							
020363 003						D-103	Jul 28, 2009
						I-501	Jul 28, 2009
FENOFIBRATE - ANTARA (MICRONIZED)							
021695 001	7101574	Aug 20, 2020	DS	DP		M-47	Oct 21, 2008
FENOFIBRATE - ANTARA (MICRONIZED)							
021695 003	7101574	Aug 20, 2020	DS	DP		M-47	Oct 21, 2008
FENOFIBRATE - FENOFIBRATE							
076433 001						PC	May 22, 2006

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FENOFIBRATE - FENOFIBRATE					
076433 002				PC	May 22, 2006
FENOFIBRATE - LIPOFEN					
021612 001	5545628	Jan 10, 2015	U-701		
FENOFIBRATE - LIPOFEN					
021612 002	5545628	Jan 10, 2015	U-701		
FENOFIBRATE - LIPOFEN					
021612 003	5545628	Jan 10, 2015	U-701		
FENOFIBRATE - TRICOR					
021203 001	**>A>** 7037529	Jan 09, 2018	DP		
	>A> 7041319	Jan 09, 2018	DP		
FENOFIBRATE - TRICOR					
021203 003	**>A>** 7037529	Jan 09, 2018	DP		
	>A> 7041319	Jan 09, 2018	DP		
FENOFIBRATE - TRICOR					
021656 001	7037529	Jan 09, 2018	DP		
	7041319	Jan 09, 2018	DP		
FENOFIBRATE - TRICOR					
021656 002	7037529	Jan 09, 2018	DP		
	7041319	Jan 09, 2018	DP		
FENTANYL CITRATE - FENTORA					
021947 001	6200604	Mar 26, 2019	U-767	NDF	Sep 25, 2009
	6974590	Mar 26, 2019	U-767		
FENTANYL CITRATE - FENTORA					
021947 002	6200604	Mar 26, 2019	U-767	NDF	Sep 25, 2009
	6974590	Mar 26, 2019	U-767		
FENTANYL CITRATE - FENTORA					
021947 003	6200604	Mar 26, 2019	U-767	NDF	Sep 25, 2009
	6974590	Mar 26, 2019	U-767		
FENTANYL CITRATE - FENTORA					
021947 004	6200604	Mar 26, 2019	U-767	NDF	Sep 25, 2009
	6974590	Mar 26, 2019	U-767		
FENTANYL CITRATE - FENTORA					
021947 005	6200604	Mar 26, 2019	U-767	NDF	Sep 25, 2009
	6974590	Mar 26, 2019	U-767		
FENTANYL HYDROCHLORIDE - IONSYS					
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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FEXOFENADINE HYDROCHLORIDE - ALLEGRA					
020625 001	**>A>** 7135571	May 18, 2014	DS		
	>A> 7135571*PED	Nov 18, 2014			
	>A> 7138524	May 18, 2014	DS		
	>A> 7138524*PED	Nov 18, 2014			
FEXOFENADINE HYDROCHLORIDE - ALLEGRA					
020872 001	**>A>** 7135571	May 18, 2014	DS		
	>A> 7135571*PED	Nov 18, 2014			
	>A> 7138524	May 18, 2014	DS		
	>A> 7138524*PED	Nov 18, 2014			
FEXOFENADINE HYDROCHLORIDE - ALLEGRA					
020872 002	**>A>** 7135571	May 18, 2014	DS		
	>A> 7135571*PED	Nov 18, 2014			
	>A> 7138524	May 18, 2014	DS		
	>A> 7138524*PED	Nov 18, 2014			
FEXOFENADINE HYDROCHLORIDE - ALLEGRA					
020872 004	**>A>** 7135571	May 18, 2014	DS		
	>A> 7135571*PED	Nov 18, 2014			
	>A> 7138524	May 18, 2014	DS		
	>A> 7138524*PED	Nov 18, 2014			
FEXOFENADINE HYDROCHLORIDE - ALLEGRA					
021963 001	5578610	Nov 26, 2013	DS DP U-772		
	6037353	Mar 14, 2017	U-772		
	6187791	May 11, 2012	U-772		
	6399632	May 11, 2012	U-772		
	>A> 7138524	May 18, 2014	DS		
	>A> 7138524*PED	Nov 18, 2014			
FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA D 24 HOUR					
021704 001	**>A>** 7138524	May 18, 2014	DS		
	>A> 7138524*PED	Nov 18, 2014			
	RE39069	May 29, 2018	DP		
FEXOFENADINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ALLEGRA-D 12 HOUR					
020786 001	**>A>** 7135571	May 18, 2014	DS		
	>A> 7135571*PED	Nov 18, 2014			
	>A> 7138524	May 18, 2014	DS		
	>A> 7138524*PED	Nov 18, 2014			
FINASTERIDE - FINASTERIDE					
076340 001				PC	Dec 16, 2006
FLUNISOLIDE - AEROSPAN HFA					
021247 001				NP	Jan 27, 2009
FLUOCINOLONE ACETONIDE - RETISERT					
021737 001	6217895	Mar 22, 2019	DP U-708		
	6548078	Mar 22, 2019	DP U-708		
FLUOCINONIDE - VANOS					
021758 001				I-487	Mar 02, 2009
FLUOXETINE HYDROCHLORIDE - PROZAC WEEKLY					
021235 001	RE39030	May 29, 2017	DP U-397		
	RE39030	May 29, 2017	DP U-396		
FLUTICASONE PROPIONATE - FLOVENT HFA					
021433 001	5658549	Sep 19, 2014	DP U-710	NPP	Feb 28, 2009
	5674472	Oct 07, 2014	DP		
	6251368	Dec 04, 2012	DP		
FLUTICASONE PROPIONATE - FLOVENT HFA					
021433 002	5658549	Sep 19, 2014	DP U-710	NPP	Feb 28, 2009
	5674472	Oct 07, 2014	DP		
	6251368	Dec 04, 2012	DP		

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FLUTICASONE PROPIONATE - FLOVENT HFA							
021433 003	5658549	Sep 19, 2014		DP	U-710	NPP	Feb 28, 2009
	5674472	Oct 07, 2014		DP			
	6251368	Dec 04, 2012		DP			
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 100/50							
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					
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 250/50							
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					
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR DISKUS 500/50							
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					
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA							
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			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA					
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		
FLUTICASONE PROPIONATE; SALMETEROL XINAFOATE - ADVAIR HFA					
021254 003	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		
FLUVASTATIN SODIUM - LESCOL					
020261 001				I-507	Apr 10, 2009
				PED	Oct 10, 2009
FLUVASTATIN SODIUM - LESCOL					
020261 002				I-507	Apr 10, 2009
				PED	Oct 10, 2009
FLUVASTATIN SODIUM - LESCOL XL					
021192 001				I-507	Apr 10, 2009
				PED	Oct 10, 2009
FORMOTEROL FUMARATE - FORADIL					
020831 001	6887459	Nov 28, 2020	U-762		
FROVATRIPTAN SUCCINATE - FROVA					
021006 001	5464864	Nov 07, 2015	U-436		
FULVESTRANT - FASLODEX					
021344 001	4659516	Dec 11, 2007	DS DP U-596		
GALANTAMINE HYDROBROMIDE - RAZADYNE ER					
021615 001				NDF	Apr 01, 2008
GALANTAMINE HYDROBROMIDE - RAZADYNE ER					
021615 002				NDF	Apr 01, 2008
GALANTAMINE HYDROBROMIDE - RAZADYNE ER					
021615 003				NDF	Apr 01, 2008
GANIRELIX ACETATE - GANIRELIX ACETATE INJECTION					
021057 001	4801577	Feb 05, 2012	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 EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
GEMCITABINE HYDROCHLORIDE - GEMZAR					
020509 001				I-499	Jul 14, 2009
GEMCITABINE HYDROCHLORIDE - GEMZAR					
020509 002				I-499	Jul 14, 2009
GEMIFLOXACIN MESYLATE - FACTIVE					
021158 001	5776944	Apr 04, 2017	DS DP		
GLIMEPIRIDE - AMARYL					
020496 001				M-54	Nov 28, 2008
				PED	May 28, 2009
GLIMEPIRIDE - AMARYL					
020496 002				M-54	Nov 28, 2008
				PED	May 28, 2009
GLIMEPIRIDE - AMARYL					
020496 003				M-54	Nov 28, 2008
				PED	May 28, 2009
GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT					
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		
GLIMEPIRIDE; PIOGLITAZONE HYDROCHLORIDE - DUETACT					
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		
GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL					
021700 001	5002953	Sep 17, 2011	DS DP U-781		
	5002953	Sep 17, 2011	DS DP U-690		
	5002953*PED	Mar 17, 2012	U-781		
	5741803	Apr 21, 2015	DS DP U-781		
	5741803	Apr 21, 2015	DS DP U-690		
	5741803*PED	Oct 21, 2015			
GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL					
021700 002	5002953	Sep 17, 2011	DS DP U-781		
	5002953	Sep 17, 2011	DS DP U-690		
	5002953*PED	Mar 17, 2012			
	5741803	Apr 21, 2015	DS DP U-781		
	5741803	Apr 21, 2015	DS DP U-690		
	5741803*PED	Oct 21, 2015			
GLIMEPIRIDE; ROSIGLITAZONE MALEATE - AVANDARYL					
021700 003	5002953	Sep 17, 2011	DS DP U-781		
	5002953	Sep 17, 2011	DS DP U-690		
	5002953*PED	Mar 17, 2012			
	5741803	Apr 21, 2015	DS DP U-781		
	5741803	Apr 21, 2015	DS DP U-690		
	5741803*PED	Oct 21, 2015			
HYALURONIDASE RECOMBINANT HUMAN - HYLENEX RECOMBINANT					
021859 001				NCE	Dec 02, 2010
HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL					
021956 001				NC	Aug 28, 2009
HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL					
021956 002				NC	Aug 28, 2009
HYDROCHLOROTHIAZIDE; METOPROLOL SUCCINATE - DUTOPROL					
021956 003				NC	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 EXPIRATON DATE	PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT							
020818 004						NS	Apr 28, 2009
HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT							
020818 005						NS	Apr 28, 2009
HYDROXOCOBALAMIN - CYANOKIT							
022041 002						**>A>** NP	Dec 15, 2009
IBANDRONATE SODIUM - BONIVA							
021455 001	4927814	Jul 09, 2007	DS	DP	U-642		
	6143326	Apr 21, 2017			U-642		
IBANDRONATE SODIUM - BONIVA							
021858 001	4927814	Jul 09, 2007	DS	DP	U-700	NDF	Jan 06, 2009
	5662918	Sep 02, 2014		DP		NCE	May 16, 2008
IBUPROFEN LYSINE - NEOPROFEN							
021903 001						NE	Apr 13, 2009
						ODE	Apr 13, 2013
ICODEXTRIN - EXTRANEAL							
021321 001	4761237	Aug 09, 2009			U-495		
IMATINIB MESYLATE - GLEEVEC							
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	Nov 10, 2008
						PED	Nov 20, 2006
						PED	Jun 20, 2006
						PED	Aug 01, 2009
						PED	Nov 10, 2006
IMATINIB MESYLATE - GLEEVEC							
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	Nov 10, 2008
						PED	Nov 20, 2006
						PED	Jun 20, 2006
						PED	Aug 01, 2009
						PED	Nov 10, 2006
IMATINIB MESYLATE - GLEEVEC							
021588 001	5521184	Jan 04, 2015	DS	DP		I-514	Oct 19, 2009
	5521184*PED	Jul 04, 2015				I-510	Oct 19, 2009
	6894051	May 23, 2019	DS	DP	U-649	I-511	Oct 19, 2009
	6894051*PED	Nov 23, 2019				I-513	Oct 19, 2009
	6958335	Dec 19, 2021	DS	DP		I-512	Oct 19, 2009
	6958335*PED	Jun 19, 2022				I-376	Dec 20, 2005
						I-392	May 20, 2006
						NCE	May 10, 2006
						ODE	Oct 19, 2013
						ODE	Oct 19, 2013
						ODE	Oct 19, 2013
						ODE	Oct 19, 2013
						ODE	Oct 19, 2013
						ODE	Feb 01, 2009
						ODE	May 10, 2008
						PED	Aug 01, 2009
						PED	Nov 10, 2008
						PED	Nov 20, 2006
						PED	Jun 20, 2006
						PED	Nov 10, 2006

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
IMATINIB MESYLATE - GLEEVEC					
021588 002	5521184	Jan 04, 2015		I-514	Oct 19, 2009
	5521184*PED	Jul 04, 2015		I-512	Oct 19, 2009
	6894051	May 23, 2019	DS DP U-649	I-513	Oct 19, 2009
	6894051*PED	Nov 23, 2019		I-511	Oct 19, 2009
	6958335	Dec 19, 2021	DS DP	I-510	Oct 19, 2009
	6958335*PED	Jun 19, 2022		I-376	Dec 20, 2005
				I-392	May 20, 2006
				NCE	May 10, 2006
				ODE	Oct 19, 2013
				ODE	Oct 19, 2013
				ODE	Oct 19, 2013
				ODE	Oct 19, 2013
				ODE	Oct 19, 2013
				ODE	Feb 01, 2009
				ODE	May 10, 2008
				PED	Nov 10, 2006
				PED	Aug 01, 2009
				PED	Jun 20, 2006
				PED	Nov 10, 2008
				PED	Nov 20, 2006
IMIQUIMOD - ALDARA					
020723 001	4689338	Aug 25, 2009	U-172	I-433	Jul 14, 2007
	4689338*PED	Feb 25, 2010		I-420	Mar 02, 2007
	5238944	Aug 24, 2010		PED	Sep 02, 2007
	5238944*PED	Feb 24, 2011		PED	Jan 14, 2008
INSULIN DETEMIR RECOMBINANT - LEVEMIR					
021536 001				I-489	Oct 19, 2008
INSULIN GLARGINE RECOMBINANT - LANTUS					
021081 001	6100376	Nov 06, 2009	DS DP U-771		
	6100376*PED	May 06, 2010			
INSULIN RECOMBINANT HUMAN - EXUBERA					
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		
INSULIN RECOMBINANT HUMAN - EXUBERA					
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		
IODINE POVACRYLEX; ISOPROPYL ALCOHOL - DURAPREP					
021586 001				NC	Sep 29, 2009

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
IODINE POVACRYLEX; ISOPROPYL ALCOHOL - DURAPREP					
021586 002				NC	Sep 29, 2009
IPRATROPIUM BROMIDE - ATROVENT HFA					
021527 001	6983743	May 26, 2020	DP		
KETOCONAZOLE - XOLEGEL					
021946 001				NDF	Jul 28, 2009
KETOTIFEN FUMARATE - KETOTIFEN FUMARATE					
077354 001				PC	Dec 30, 2006
KUNECATECHINS - VEREGEN					
021902 001	5795911	Apr 10, 2017	U-172	NP	Oct 31, 2009
	5968973	Apr 10, 2017	U-172		
LAMOTRIGINE - LAMICTAL					
020241 001				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL					
020241 002				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL					
020241 003				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL					
020241 004				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL					
020241 005				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL					
020241 006				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL CD					
020764 001				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL CD					
020764 002				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL CD					
020764 003				I-516	Sep 22, 2009
LAMOTRIGINE - LAMICTAL CD					
020764 004				I-516	Sep 22, 2009
LAMOTRIGINE - LAMOTRIGINE					
076420 001				PC	Feb 25, 2007
LAMOTRIGINE - LAMOTRIGINE					
076420 002				PC	Dec 25, 2006
LANSOPRAZOLE - PREVACID					
020406 001	6749864	Feb 13, 2007	DP		
LANSOPRAZOLE - PREVACID					
020406 002	6749864	Feb 13, 2007	DP		
LANSOPRAZOLE - PREVACID					
021281 001	6749864	Feb 13, 2007	DP		
LANSOPRAZOLE - PREVACID					
021281 002	6749864	Feb 13, 2007	DP		
LANSOPRAZOLE - PREVACID					
021428 001	6749864	Feb 13, 2007	DP		
LANSOPRAZOLE - PREVACID					
021428 002	6749864	Feb 13, 2007	DP		
LANTHANUM CARBONATE - FOSRENOL					
021468 003	5968976	Mar 19, 2016	DP	U-613	
LANTHANUM CARBONATE - FOSRENOL					
021468 004	5968976	Mar 19, 2016	DP	U-613	
LATANOPROST - XALATAN					
020597 001	5296504	Mar 22, 2011	DP	U-778	
	5422368	Mar 22, 2011	DP	U-778	
	6429226	Sep 06, 2009	DP	U-778	

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
LENALIDOMIDE - REVLIMID					
021880 001	5635517	Jul 24, 2016	DS	ODE	Jun 29, 2013
	6045501	Aug 28, 2018	U-694	ODE	Dec 27, 2012
	6281230	Jul 24, 2016	U-769		
	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		
	7119106	Jul 24, 2016	DP		
LENALIDOMIDE - REVLIMID					
021880 002	5635517	Jul 24, 2016	DS	ODE	Jun 29, 2013
	6045501	Aug 28, 2018	U-694	ODE	Dec 27, 2012
	6281230	Jul 24, 2016	U-769		
	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		
	7119106	Jul 24, 2016	DP		
LENALIDOMIDE - REVLIMID					
021880 003	5635517	Jul 24, 2016	DS	I-500	Jun 29, 2009
	6045501	Aug 28, 2018	U-694	NCE	Dec 27, 2010
	6281230	Jul 24, 2016	U-769	ODE	Jun 29, 2013
	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		
	7119106	Jul 24, 2016	DP		
LENALIDOMIDE - REVLIMID					
021880 004	5635517	Jul 24, 2016	DS	I-500	Jun 29, 2009
	6045501	Aug 28, 2018	U-694	NCE	Dec 27, 2010
	6281230	Jul 24, 2016	U-769	ODE	Jun 29, 2013
	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		
	7119106	Jul 24, 2016	DP		
LEVALBUTEROL TARTRATE - XOPENEX HFA					
021730 001	5362755	Mar 25, 2013	U-636		
LEVETIRACETAM - KEPPRA					
021035 001				I-506	Aug 15, 2009
LEVETIRACETAM - KEPPRA					
021035 002				I-506	Aug 15, 2009
LEVETIRACETAM - KEPPRA					
021035 003				I-506	Aug 15, 2009
LEVETIRACETAM - KEPPRA					
021035 004	4943639	Jul 14, 2008	DS	I-506	Aug 15, 2009
				NPP	Jun 21, 2008
LEVETIRACETAM - KEPPRA					
021505 001				I-506	Aug 15, 2009

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
LEVETIRACETAM - KEPPRA							
021872 001	4943639	Jul 14, 2008	DS			NDF	Jul 31, 2009
LEVOBETAXOLOL HYDROCHLORIDE - BETAXON							
021114 001	4911920	Mar 27, 2007	DS	DP	U-191	M-54	Sep 28, 2009
	4911920	Mar 27, 2007	DS	DP	U-369	PED	Mar 28, 2010
	4911920*PED	Sep 27, 2007					
	5540918	Jul 30, 2013		DP			
	5540918*PED	Jan 30, 2014					
LEVONORGESTREL - PLAN B							
021045 002						NP	Aug 24, 2009
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 001	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 002	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 003	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 004	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 005	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 006	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 007	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 008	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 009	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 010	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 011	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LEVOTHYROXINE SODIUM - LEVOXYL							
021301 012	7067148	Feb 15, 2022		DP			
	7101569	Aug 14, 2022			U-759		
LIDOCAINE; TETRACAINE - LIDOCAINE AND TETRACAINE							
021717 001	5919479	Jul 28, 2015		DP		NP	Jun 29, 2009
	6528086	Sep 28, 2019		DP			
LIDOCAINE; TETRACAINE - SYNERA							
021623 001						NC	Jun 23, 2008
LOPERAMIDE HYDROCHLORIDE; SIMETHICONE - LOPERAMIDE HYDROCHLORIDE AND SIMETHICONE							
077500 001						PC	May 12, 2007

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO		PATENT NO	PATENT EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
LOPINAVIR; RITONAVIR - KALETRA								
021226 001	>A>	5886036	Nov 19, 2013	DS	DP			
	>A>	5886036*PED	May 19, 2014					
	>A>	5948436	Sep 13, 2013		DP			
	>A>	5948436*PED	Mar 13, 2014					
	>A>	7141593	May 22, 2020		DP			
	>A>	7141593*PED	Nov 22, 2020					
LOPINAVIR; RITONAVIR - KALETRA								
021251 001	>A>	5886036	Nov 19, 2013	DS	DP			
	>A>	5886036*PED	May 19, 2014					
	>A>	5948436	Sep 13, 2013		DP			
	>A>	5948436*PED	Mar 13, 2014					
LOPINAVIR; RITONAVIR - KALETRA								
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					
	>A>	5886036	Nov 19, 2013	DS	DP			
	>A>	5886036*PED	May 19, 2014					
		6037157	Jun 26, 2016			U-688		
		6037157*PED	Dec 26, 2016					
		6703403	Jun 26, 2016			U-688		
		6703403*PED	Dec 26, 2016					
LOVASTATIN; NIACIN - ADVICOR								
021249 001		6469035	Mar 15, 2018			U-768		
		7011848	Sep 20, 2013			U-712		
LOVASTATIN; NIACIN - ADVICOR								
021249 002		6469035	Mar 15, 2018			U-768		
		7011848	Sep 20, 2013			U-712		
LOVASTATIN; NIACIN - ADVICOR								
021249 003		6469035	Mar 15, 2018			U-768		
		7011848	Sep 20, 2013			U-712		
LOVASTATIN; NIACIN - ADVICOR								
021249 004		6469035	Mar 15, 2018			U-768		
LUBIPROSTONE - AMITIZA								
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		
MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - NORMOCARB HF 25								
021910 001	>A>	5945449	Oct 31, 2017		DP	U-785	ODE	Jul 26, 2013
MAGNESIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE - NORMOCARB HF 35								
021910 002	>A>	5945449	Oct 31, 2017		DP	U-785	ODE	Jul 26, 2013
MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID								
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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
MAGNESIUM HYDROXIDE; OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID					
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		
MECASERMIN RINFABATE RECOMBINANT - IPLEX					
021884 001	5200509	Apr 06, 2010	DS		
	5681818	Oct 28, 2014	U-697		
MEGESTROL ACETATE - MEGACE ES					
021778 001	7101576	Apr 22, 2024	U-755		
MELOXICAM - MOBIC					
020938 001				ODE	Aug 11, 2012
				PED	Feb 11, 2013
MELOXICAM - MOBIC					
020938 002				ODE	Aug 11, 2012
				PED	Feb 11, 2013
MELOXICAM - MOBIC					
021530 001				I-469	Aug 11, 2008
				ODE	Aug 11, 2012
				PED	Feb 11, 2013
				PED	Feb 11, 2009
MEQUINOL; TRETINOIN - SOLAGE					
020922 001	5194247	Dec 10, 2013	DP U-294		
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET					
021410 001	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		
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET					
021410 002	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		
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET					
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		
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET					
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		

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET							
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					
METHYLPHENIDATE - DAYTRANA							
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		
METHYLPHENIDATE - DAYTRANA							
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		
METHYLPHENIDATE - DAYTRANA							
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		
METHYLPHENIDATE - DAYTRANA							
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		
METHYLPHENIDATE HYDROCHLORIDE - METADATE CD							
021259 001	6344215	Oct 27, 2020		DP			
METHYLPHENIDATE HYDROCHLORIDE - METADATE CD							
021259 002	6344215	Oct 27, 2020		DP			
METHYLPHENIDATE HYDROCHLORIDE - METADATE CD							
021259 003	6344215	Oct 27, 2020		DP			
METHYLPHENIDATE HYDROCHLORIDE - METADATE CD							
021259 004	6344215	Oct 27, 2020		DP			
METOPROLOL SUCCINATE - TOPROL-XL							
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					
	>A> 5246714	Sep 21, 2010					
	>A> 5246714*PED	Mar 21, 2011					
METOPROLOL SUCCINATE - TOPROL-XL							
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					
	>A> 5246714	Sep 21, 2010					
	>A> 5246714*PED	Mar 21, 2011					

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
METOPROLOL SUCCINATE - TOPROL-XL						
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				
	>A> 5246714	Sep 21, 2010				
	>A> 5246714*PED	Mar 21, 2011				
METOPROLOL SUCCINATE - TOPROL-XL						
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				
	>A> 5246714	Sep 21, 2010				
	>A> 5246714*PED	Mar 21, 2011				
METRONIDAZOLE - METROGEL						
021789 001	6881726	Feb 21, 2022	DP	U-743		
MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE - VUSION						
021026 001	4911932	Mar 27, 2007	DP	U-718	NP	Feb 16, 2009
MINOXIDIL - MEN'S ROGAINE						
021812 001	6946120	Apr 20, 2019	DP	U-702	NDF	Jan 20, 2009
MODAFINIL - PROVIGIL						
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
MODAFINIL - PROVIGIL						
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
MORPHINE SULFATE - KADIAN						
020616 004	5378474	Mar 23, 2010				
MORPHINE SULFATE - KADIAN						
020616 005	5202128	Apr 13, 2010				
	5378474	Mar 23, 2010				
MOXIFLOXACIN HYDROCHLORIDE - AVELOX						
021085 001	4990517	Dec 08, 2011	DS	DP U-298		
	6610327	Oct 29, 2019		DP U-298		
MOXIFLOXACIN HYDROCHLORIDE - AVELOX IN SODIUM CHLORIDE 0.8% IN PLASTIC CONTAINER						
021277 001	4990517	Dec 08, 2011	DS	DP U-298		
	6548079	Jul 25, 2020		DP U-298		
MOXIFLOXACIN HYDROCHLORIDE - VIGAMOX						
021598 001	4990517	Dec 08, 2011	DS	DP U-709		
	4990517*PED	Jun 08, 2012				

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
NALTREXONE - VIVITROL					
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		
NELARABINE - ARRANON					
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		
NIACIN - NIASPAN					
020381 001	7011848	Sep 20, 2013	U-712		
NIACIN - NIASPAN					
020381 002	6469035	Mar 15, 2018	U-768		
	7011848	Sep 20, 2013	U-712		
NIACIN - NIASPAN					
020381 003	6469035	Mar 15, 2018	U-768		
	7011848	Sep 20, 2013	U-712		
NIACIN - NIASPAN					
020381 004	6469035	Mar 15, 2018	U-768		
	7011848	Sep 20, 2013	U-712		
NIACIN - NIASPAN TITRATION STARTER PACK					
020381 005	7011848	Sep 20, 2013	U-712		
NICOTINE POLACRILEX - NICOTINE POLACRILEX					
077007 001				PC	Aug 21, 2006
NICOTINE POLACRILEX - NICOTINE POLACRILEX					
077007 002				PC	Aug 21, 2006
NITROGLYCERIN - NITROMIST					
021780 001				NP	Nov 02, 2009
OCTREOTIDE ACETATE - SANDOSTATIN					
019667 001	5753618	May 19, 2015			
	5753618*PED	Nov 19, 2015			
OCTREOTIDE ACETATE - SANDOSTATIN					
019667 002	5753618	May 19, 2015			
	5753618*PED	Nov 19, 2015			
OCTREOTIDE ACETATE - SANDOSTATIN					
019667 003	5753618	May 19, 2015			
	5753618*PED	Nov 19, 2015			
OCTREOTIDE ACETATE - SANDOSTATIN					
019667 004	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 EXPIRATON DATE			PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
OCTREOTIDE ACETATE - SANDOSTATIN								
019667 005	5753618	May	19,	2015				
	5753618*PED	Nov	19,	2015				
OCTREOTIDE ACETATE - SANDOSTATIN LAR								
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				
OCTREOTIDE ACETATE - SANDOSTATIN LAR								
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				
OCTREOTIDE ACETATE - SANDOSTATIN LAR								
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				
OLOPATADINE HYDROCHLORIDE - PATADAY								
021545 001	5116863	Dec	18,	2010	DS	DP		
	5641805	Jun	06,	2015		U-765		
	6995186	Nov	12,	2023		DP U-765		
OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID								
021849 001	6489346	Jul	16,	2016	DS	DP U-588		
	6645988	Jul	16,	2016	DS	DP		
	6699885	Jul	16,	2016		U-588		
OMEPRAZOLE; SODIUM BICARBONATE - ZEGERID								
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		
ONDANSETRON HYDROCHLORIDE - ONDANSETRON HYDROCHLORIDE								
076960 001							**>A>** PC	Jun 25, 2007

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
OXALIPLATIN - ELOXATIN					
021492 001	5290961	Jan 12, 2013		I-441	Nov 04, 2007
	5290961*PED	Jul 12, 2013		I-425	Jan 09, 2007
	5338874	Apr 07, 2013	DS	NCE	Aug 09, 2007
	5338874*PED	Oct 07, 2013		PED	Jul 09, 2007
	5420319	Aug 09, 2016	DS	PED	Feb 09, 2008
	5420319*PED	Aug 09, 2016		PED	May 04, 2008
OXALIPLATIN - ELOXATIN					
021492 002	5290961	Jan 12, 2013		I-441	Nov 04, 2007
	5290961*PED	Jul 12, 2013		I-425	Jan 09, 2007
	5338874	Apr 07, 2013	DS	NCE	Aug 09, 2007
	5338874*PED	Oct 07, 2013		PED	Jul 09, 2007
	5420319	Aug 09, 2016	DS	PED	Feb 09, 2008
	5420319*PED	Aug 09, 2016		PED	May 04, 2008
OXALIPLATIN - ELOXATIN					
021759 001	5290961	Jan 12, 2013	DS	I-441	Nov 04, 2007
	5290961*PED	Jul 12, 2013		NCE	Aug 09, 2007
	5338874	Apr 07, 2013	DS	PED	Feb 09, 2008
	5338874*PED	Oct 07, 2013		PED	May 04, 2008
	5420319	Aug 08, 2016	DS		
	5420319*PED	Feb 08, 2017			
	5716988	Aug 07, 2015	DP		
	5716988*PED	Feb 07, 2016			
OXALIPLATIN - ELOXATIN					
021759 002	5290961	Jan 12, 2013	DS	I-441	Nov 04, 2007
	5290961*PED	Jul 12, 2013		NCE	Aug 09, 2007
	5338874	Apr 07, 2013	DS	PED	Feb 09, 2008
	5338874*PED	Oct 07, 2013		PED	May 04, 2008
	5420319	Aug 08, 2016	DS		
	5420319*PED	Feb 08, 2017			
	5716988	Aug 07, 2015	DP		
	5716988*PED	Feb 07, 2016			
OXALIPLATIN - ELOXATIN					
021759 003				**>A>** I-441	Nov 04, 2007
				>A> NCE	Aug 09, 2007
				>A> PED	Feb 09, 2008
				>A> PED	May 04, 2008
OXCARBAZEPINE - TRILEPTAL					
021014 001	7037525	Feb 12, 2018	U-724		
OXCARBAZEPINE - TRILEPTAL					
021014 002	7037525	Feb 12, 2018	U-724		
OXCARBAZEPINE - TRILEPTAL					
021014 003	7037525	Feb 12, 2018	U-724		
OXCARBAZEPINE - TRILEPTAL					
021285 001	7037525	Feb 12, 2018	U-724		
OXYBUTYNIN CHLORIDE - OXYBUTYNIN CHLORIDE					
076644 001				PC	May 09, 2007
OXYBUTYNIN CHLORIDE - OXYBUTYNIN CHLORIDE					
076702 001				PC	May 09, 2007
OXYBUTYNIN CHLORIDE - OXYBUTYNIN CHLORIDE					
076745 001				PC	May 09, 2007
OXYCODONE HYDROCHLORIDE - OXYCONTIN					
020553 001	5266331	Oct 26, 2007	DP		
	5549912	Oct 26, 2007	DP		
OXYCODONE HYDROCHLORIDE - OXYCONTIN					
020553 002	5266331	Oct 26, 2007	DP		
	5549912	Oct 26, 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 EXPIRATON DATE			PATENT CODES		EXCLUSIVITY CODE(S)		EXCLUSIVITY EXPIRATION DATE
OXYCODONE HYDROCHLORIDE - OXYCONTIN									
020553 003	5266331	Oct	26,	2007	DP				
	5549912	Oct	26,	2007	DP				
OXYCODONE HYDROCHLORIDE - OXYCONTIN									
020553 004	5266331	Oct	26,	2007	DP				
	5549912	Oct	26,	2007	DP				
OXYCODONE HYDROCHLORIDE - OXYCONTIN									
020553 005	5266331	Oct	26,	2007	DP				
	5549912	Oct	26,	2007	DP				
OXYCODONE HYDROCHLORIDE - OXYCONTIN									
020553 006	5266331	Oct	26,	2007	DP				
	5508042	Apr	16,	2013		U-443			
	5549912	Oct	26,	2007	DP				
	5656295	Oct	26,	2007	DP	U-443			
OXYCODONE HYDROCHLORIDE - OXYCONTIN									
020553 007	5266331	Oct	26,	2007	DP				
	5508042	Apr	16,	2013		U-443			
	5549912	Oct	26,	2007	DP				
	5656295	Oct	26,	2007	DP	U-443			
OXYCODONE HYDROCHLORIDE - OXYCONTIN									
020553 008	5266331	Oct	26,	2007	DP				
	5508042	Apr	16,	2013		U-443			
	5549912	Oct	26,	2007	DP				
	5656295	Oct	26,	2007	DP	U-443			
OXYMORPHONE HYDROCHLORIDE - OPANA									
021611 001							NDF	Jun	22, 2009
OXYMORPHONE HYDROCHLORIDE - OPANA									
021611 002							NDF	Jun	22, 2009
OXYMORPHONE HYDROCHLORIDE - OPANA ER									
021610 001	5128143	Sep	19,	2008	DP		NDF	Jun	22, 2009
OXYMORPHONE HYDROCHLORIDE - OPANA ER									
021610 002	5128143	Sep	19,	2008	DP		NDF	Jun	22, 2009
OXYMORPHONE HYDROCHLORIDE - OPANA ER									
021610 003	5128143	Sep	19,	2008	DP		NDF	Jun	22, 2009
OXYMORPHONE HYDROCHLORIDE - OPANA ER									
021610 004	5128143	Sep	19,	2008	DP		NDF	Jun	22, 2009
PALIPERIDONE - INVEGA									
021999 001							**>A>** NCE	Dec	19, 2011
PALIPERIDONE - INVEGA									
021999 002							**>A>** NCE	Dec	19, 2011
PALIPERIDONE - INVEGA									
021999 003							**>A>** NCE	Dec	19, 2011
PALIPERIDONE - INVEGA									
021999 004							**>A>** NCE	Dec	19, 2011
PAROXETINE HYDROCHLORIDE - PAXIL									
020031 004	6133289	May	19,	2015		U-358			
PAROXETINE HYDROCHLORIDE - PAXIL									
020031 005	6133289	May	19,	2015		U-358			
POLYETHYLENE GLYCOL 3350 - MIRALAX									
022015 001							NP	Oct	06, 2009
POSACONAZOLE - NOXAFIL									
022003 001	5661151	Aug	26,	2014	DS	DP U-760	NCE	Sep	15, 2011
	5703079	Dec	30,	2014	DS	DP U-760			
	6958337	Sep	25,	2020	DS	DP U-760			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX						
020667 001	4843086	Jun 27, 2006		U-231	I-517	Nov 07, 2009
	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX						
020667 002	4843086	Jun 27, 2006		U-231	I-517	Nov 07, 2009
	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX						
020667 003	4843086	Jun 27, 2006		U-231	I-517	Nov 07, 2007
	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX						
020667 004	4843086	Jun 27, 2006		U-231		
PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX						
020667 005	4843086	Jun 27, 2006		U-231	I-517	Nov 07, 2009
	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
PRAMIPEXOLE DIHYDROCHLORIDE - MIRAPEX						
020667 006	4843086	Jun 27, 2006		U-231	I-517	Nov 07, 2009
	6001861	Jan 16, 2018		U-784		
	6194445	Jan 16, 2018		U-784		
PRAVASTATIN SODIUM - PRAVASTATIN SODIUM						
076056 001					PC	Oct 21, 2006
PRAVASTATIN SODIUM - PRAVASTATIN SODIUM						
076056 002					PC	Oct 21, 2006
PRAVASTATIN SODIUM - PRAVASTATIN SODIUM						
076056 003					PC	Oct 21, 2006
PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT						
021959 001	5178878	Jan 12, 2010		DP		
PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT						
021959 002	5178878	Jan 12, 2010		DP		
PREDNISOLONE SODIUM PHOSPHATE - ORAPRED ODT						
021959 003	5178878	Jan 12, 2010		DP		
QUETIAPINE FUMARATE - SEROQUEL						
020639 001					I-503	Oct 20, 2009
QUETIAPINE FUMARATE - SEROQUEL						
020639 002					I-503	Oct 20, 2009
QUETIAPINE FUMARATE - SEROQUEL						
020639 003					I-503	Oct 20, 2009
QUETIAPINE FUMARATE - SEROQUEL						
020639 004					I-503	Oct 20, 2009
QUETIAPINE FUMARATE - SEROQUEL						
020639 005					I-503	Oct 20, 2009
QUETIAPINE FUMARATE - SEROQUEL						
020639 006	4879288	Sep 26, 2011	DS	DP U-550	I-503	Oct 20, 2009
QUETIAPINE FUMARATE - SEROQUEL						
020639 007	4879288	Sep 26, 2011	DS	DP U-550	I-503	Oct 20, 2009
RALOXIFENE HYDROCHLORIDE - EVISTA						
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		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
RAMIPRIL - ALTACE					
019901 001	5061722	Oct 19, 2008			
RANOLAZINE - RANEXA					
021526 002	4567264	May 18, 2007	DS	NCE	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		
RASAGILINE MESYLATE - AZILECT					
021641 001	5387612	Feb 07, 2012	U-219	NCE	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		
RASAGILINE MESYLATE - AZILECT					
021641 002	5387612	Feb 07, 2012	U-219	NCE	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		
RIFAXIMIN - XIFAXAN					
021361 001	7045620	May 22, 2024	DS		
RISEDRONATE SODIUM - ACTONEL					
020835 001				M-52	Jan 24, 2009
RISEDRONATE SODIUM - ACTONEL					
020835 002				M-52	Jan 24, 2009
RISEDRONATE SODIUM - ACTONEL					
020835 003	5583122	Dec 10, 2013	DS DP U-756	I-309	Aug 11, 2009
	5583122	Dec 10, 2013	DS DP U-222	M-52	Jan 24, 2009
	6096342	Nov 22, 2011	DP		
RISPERIDONE - RISPERDAL					
020272 001				I-509	Oct 06, 2009
RISPERIDONE - RISPERDAL					
020272 002				I-509	Oct 06, 2009
RISPERIDONE - RISPERDAL					
020272 003				I-509	Oct 06, 2009
RISPERIDONE - RISPERDAL					
020272 004				I-509	Oct 06, 2009
RISPERIDONE - RISPERDAL					
020272 007				I-509	Oct 06, 2009
RISPERIDONE - RISPERDAL					
020272 008				I-509	Oct 06, 2009
RISPERIDONE - RISPERDAL					
020588 001	RE39181	Jul 11, 2014	DP	I-509	Oct 06, 2009
RISPERIDONE - RISPERDAL					
021444 001				I-509	Oct 06, 2009
RISPERIDONE - RISPERDAL					
021444 002				I-509	Oct 06, 2009

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
RISPERIDONE - RISPERDAL					
021444 003				I-509	Oct 06, 2006
RISPERIDONE - RISPERDAL					
021444 004				I-509	Oct 06, 2006
RISPERIDONE - RISPERDAL					
021444 005				I-509	Oct 06, 2009
RITONAVIR - NORVIR					
020659 001	**>A>** 5948436	Sep 13, 2013	DP		
	>A> 5948436*PED	Mar 13, 2014			
RITONAVIR - NORVIR					
020945 001	**>A>** 5948436	Sep 13, 2013	DP		
	>A> 5948436*PED	Mar 13, 2014			
	>A> 7141593	May 22, 2020	DP		
	>A> 7141593*PED	Nov 22, 2020			
RIVASTIGMINE TARTRATE - EXELON					
020823 003	4948807	Aug 14, 2012	DS U-322		
RIVASTIGMINE TARTRATE - EXELON					
020823 004	4948807	Aug 14, 2012	DS U-322		
RIVASTIGMINE TARTRATE - EXELON					
020823 005	4948807	Aug 14, 2012	DS U-322		
RIVASTIGMINE TARTRATE - EXELON					
020823 006	4948807	Aug 14, 2012	DS U-322		
RIVASTIGMINE TARTRATE - EXELON					
021025 001	4948807	Aug 14, 2012	DS U-322		
ROCURONIUM BROMIDE - ZEMURON					
020214 003	4894369	Apr 13, 2008			
SALMETEROL XINAFOATE - SEREVENT					
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			
SALMETEROL XINAFOATE - SEREVENT					
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			
	6536427	Mar 01, 2011	DP		
SELEGILINE - EMSAM					
021336 001	7070808	May 10, 2018	DS DP	NDF	Feb 27, 2009
	RE34579	Aug 18, 2007	DS DP U-711		
SELEGILINE - EMSAM					
021336 002	RE34579	Aug 18, 2007	DS DP U-711	NDF	Feb 27, 2009
SELEGILINE - EMSAM					
021336 003	RE34579	Aug 18, 2007	DS DP U-711	NDF	Feb 27, 2009
SELEGILINE HYDROCHLORIDE - ZELAPAR					
021479 001	5648093	Jul 15, 2014	DP	NDF	Jun 14, 2009
	6423342	Mar 01, 2016	DP		
SERTACONAZOLE NITRATE - ERTACZO					
021385 001	**>A>** 5135943	May 31, 2014	DS DP U-786		
SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE					
075719 001				PC	Feb 06, 2007

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE						
075719 002					PC	Feb 06, 2007
SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE						
075719 003					PC	Feb 06, 2007
SERTRALINE HYDROCHLORIDE - SERTRALINE HYDROCHLORIDE						
076934 001					PC	Feb 03, 2007
SERTRALINE HYDROCHLORIDE - ZOLOFT						
020990 001	6727283	Oct 11, 2019	DP	U-580		
	6727283*PED	Apr 11, 2020				
	7067555	Nov 10, 2019	DP			
	7067555*PED	May 10, 2020				
SEVELAMER HYDROCHLORIDE - RENAGEL						
021179 001	7014846	Aug 11, 2013	DP	U-246		
SEVELAMER HYDROCHLORIDE - RENAGEL						
021179 002	7014846	Aug 11, 2013	DP	U-246		
SIMVASTATIN - SIMVASTATIN						
076052 001					PC	Dec 20, 2006
SIMVASTATIN - SIMVASTATIN						
076052 002					PC	Dec 20, 2006
SIMVASTATIN - SIMVASTATIN						
076052 003					PC	Dec 20, 2006
SIMVASTATIN - SIMVASTATIN						
076052 004					PC	Dec 20, 2006
SIMVASTATIN - SIMVASTATIN						
076285 005					PC	Dec 20, 2006
SIROLIMUS - RAPAMUNE						
021083 001	5100899	Jul 07, 2013		U-290		
	5100899*PED	Jan 07, 2014				
SIROLIMUS - RAPAMUNE						
021110 001	5100899	Jul 07, 2013		U-290		
	5100899*PED	Jan 07, 2014				
SIROLIMUS - RAPAMUNE						
021110 002	5100899	Jul 07, 2013		U-290		
	5100899*PED	Jan 07, 2014				
SIROLIMUS - RAPAMUNE						
021110 003	5100899	Jul 07, 2013		U-290		
	5100899*PED	Jan 07, 2014				
SITAGLIPTIN PHOSPHATE - JANUVIA						
021995 001	6303661	Apr 24, 2017		U-774	NCE	Oct 16, 2011
	6699871	Jul 26, 2022	DS	DP U-774		
	6890898	Feb 02, 2019		U-775		
	7078381	Feb 02, 2019		U-775		
	7125873	Jul 26, 2022		U-775		
SITAGLIPTIN PHOSPHATE - JANUVIA						
021995 002	6303661	Apr 24, 2017		U-774	NCE	Oct 16, 2011
	6699871	Jul 26, 2022	DS	DP U-774		
	6890898	Feb 02, 2019		U-775		
	7078381	Feb 02, 2019		U-775		
	7125873	Jul 26, 2022		U-775		
SITAGLIPTIN PHOSPHATE - JANUVIA						
021995 003	6303661	Apr 24, 2017		U-774	NCE	Oct 16, 2011
	6699871	Jul 26, 2022	DS	DP U-774		
	6890898	Feb 02, 2019		U-775		
	7078381	Feb 02, 2019		U-775		
	7125873	Jul 26, 2022		U-775		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
SODIUM PHOSPHATE, DIBASIC, ANHYDROUS; SODIUM PHOSPHATE, MONOBASIC, MONOHYDRATE - OSMOPREP					
021892 001	5616346	May 18, 2013	DP U-715	NP	Mar 16, 2009
SOMATROPIN RECOMBINANT - GENOTROPIN					
020280 006	4968299	Jun 28, 2008	DP	I-496	Apr 27, 2009
SOMATROPIN RECOMBINANT - GENOTROPIN					
020280 007	4968299	Jun 28, 2008	DP	I-496	Apr 27, 2009
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 001	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 002	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 003	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 004				I-496	Apr 27, 2009
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 005	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 008	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 009	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 010	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 011	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 012	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - GENOTROPIN PRESERVATIVE FREE					
020280 013	5435076	Apr 16, 2013	DP	I-496	Apr 27, 2009
	5716338	Feb 10, 2015	DP		
SOMATROPIN RECOMBINANT - HUMATROPE					
019640 004				I-518	Nov 01, 2009
SOMATROPIN RECOMBINANT - HUMATROPE					
019640 005				I-518	Nov 01, 2009
SOMATROPIN RECOMBINANT - HUMATROPE					
019640 006				I-518	Nov 01, 2009
SOMATROPIN RECOMBINANT - HUMATROPE					
019640 007				I-518	Nov 01, 2009
SOMATROPIN RECOMBINANT - OMNITROPE					
021426 001				NP	May 30, 2009
SOMATROPIN RECOMBINANT - OMNITROPE					
021426 002				NP	May 30, 2009
SORAFENIB TOSYLATE - NEXAVAR					
021923 001				ODE	Dec 20, 2012
STAVUDINE - ZERIT XR					
021453 001	7135465	Feb 18, 2023	DP U-167		
	7135465*PED	Aug 18, 2023			

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE		PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
STAVUDINE - ZERIT XR							
021453 002	7135465	Feb 18, 2023		DP	U-167		
	7135465*PED	Aug 18, 2023					
STAVUDINE - ZERIT XR							
021453 003	7135465	Feb 18, 2023		DP	U-167		
	7135465*PED	Aug 18, 2023					
STAVUDINE - ZERIT XR							
021453 004	7135465	Feb 18, 2023		DP	U-167		
	7135465*PED	Aug 18, 2023					
SUMATRIPTAN SUCCINATE - IMITREX STATDOSE							
020080 003	4816470	Dec 28, 2006			U-72		
	4816470*PED	Jun 28, 2007					
	5037845	Aug 06, 2008			U-72		
	5037845*PED	Feb 06, 2009					
SUNITINIB MALATE - SUTENT							
021938 001	6573293	Feb 15, 2021	DS	DP	U-703	NCE	Jan 26, 2011
	7125905	Feb 15, 2021	DS	DP			
SUNITINIB MALATE - SUTENT							
021938 002	6573293	Feb 15, 2021	DS	DP	U-703	NCE	Jan 26, 2011
	7125905	Feb 15, 2021	DS	DP			
SUNITINIB MALATE - SUTENT							
021938 003	6573293	Feb 15, 2021	DS	DP	U-703	NCE	Jan 26, 2011
	7125905	Feb 15, 2021	DS	DP			
TACROLIMUS - PROGRAF							
050708 001						ODE	Mar 29, 2013
TACROLIMUS - PROGRAF							
050708 002						ODE	Mar 29, 2013
TACROLIMUS - PROGRAF							
050708 003						ODE	Mar 29, 2013
TACROLIMUS - PROGRAF							
050709 001						ODE	Mar 29, 2013
TELBIVUDINE - TYZEKA							
022011 001	6395716	Aug 10, 2019			U-782	NCE	Oct 25, 2011
	6444652	Aug 10, 2019			U-782		
	6566344	Aug 10, 2019			U-782		
	6569837	Aug 10, 2019			U-782		
TEMOZOLOMIDE - TEMODAR							
021029 005						I-450	Mar 15, 2008
						ODE	Mar 15, 2012
						ODE	Aug 11, 2006
						PED	Feb 11, 2007
TEMOZOLOMIDE - TEMODAR							
021029 006						I-450	Mar 15, 2008
						ODE	Mar 15, 2012
						ODE	Aug 11, 2006
						PED	Feb 11, 2007

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
TERBINAFINE - LAMISIL					
020846 001	4755534	Dec 30, 2006		U-445	
	4755534*PED	Jun 30, 2007			
	5681849	Oct 28, 2014	DP		
	5681849*PED	Apr 28, 2015			
	5856355	May 18, 2012	DP	U-502	
	5856355	May 18, 2012	DP	U-504	
	5856355	May 18, 2012	DP	U-540	
	5856355*PED	Nov 18, 2012			
	6005001	May 18, 2012	DP	U-502	
	6005001	May 18, 2012	DP	U-504	
	6005001	May 18, 2012	DP	U-540	
	6005001*PED	Nov 18, 2012			
	6121314	May 18, 2012	DP	U-504	
	6121314	May 18, 2012	DP	U-540	
	6121314	May 18, 2012	DP	U-502	
	6121314*PED	Nov 18, 2012			
TERBINAFINE HYDROCHLORIDE - LAMISIL					
020192 001	4755534	Dec 30, 2006		U-73	
	4755534*PED	Jun 30, 2007			
TERBINAFINE HYDROCHLORIDE - LAMISIL					
020539 001	4755534	Dec 30, 2006		U-73	
	4755534*PED	Jun 30, 2007			
TERBINAFINE HYDROCHLORIDE - LAMISIL					
020749 001	4755534	Dec 30, 2006			
	4755534*PED	Jun 30, 2007			
	6121314	May 18, 2012		U-502	
	6121314*PED	Nov 18, 2012			
TERBINAFINE HYDROCHLORIDE - LAMISIL					
020980 001	4755534	Dec 30, 2006		U-73	
	4755534*PED	Jun 30, 2007			
TERBINAFINE HYDROCHLORIDE - LAMISIL AT					
021124 001	4755534	Dec 30, 2006		U-73	
	4755534*PED	Jun 30, 2007			
	5681849	Oct 28, 2014			
	5681849*PED	Apr 28, 2015			
	6121314	May 18, 2012		U-504	
	6121314*PED	Nov 18, 2012			
TERBINAFINE HYDROCHLORIDE - LAMISIL AT					
021124 002	4755534	Dec 30, 2006		U-73	
	4755534*PED	Jun 30, 2007			
	5681849	Oct 28, 2014			
	5681849*PED	Apr 28, 2015			
	6121314	May 18, 2012		U-504	
	6121314*PED	Nov 18, 2012			
TERIPARATIDE RECOMBINANT HUMAN - FORTEO					
021318 001	6977077	Aug 19, 2019		U-597	
	7144861	Dec 08, 2018	DP		

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO		PATENT NO	PATENT EXPIRATON DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
THALIDOMIDE - THALOMID						
020785 001	**>A>**	5629327	May 13, 2014	U-731	ODE	May 25, 2013
		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-371		
		6561976	Aug 28, 2018	U-731		
		6561977	Oct 23, 2020	U-371		
		6561977	Oct 23, 2020	U-731		
		6755784	Oct 23, 2020	U-731		
		6755784	Oct 23, 2020	U-371		
		6869399	Oct 23, 2020	U-732		
		6869399	Oct 23, 2020	U-733		
		6869399	Oct 23, 2020	U-731		
		6869399	Oct 23, 2020	U-371		
		6908432	Aug 28, 2018	U-371		
		6908432	Aug 28, 2018	U-731		
	>A>	7141018	Oct 23, 2020	U-732		
	>A>	7141018	Oct 23, 2020	U-371		
	>A>	7141018	Oct 23, 2020	U-731		
	>A>	7141018	Oct 23, 2020	U-733		
THALIDOMIDE - THALOMID						
020785 002	**>A>**	5629327	May 13, 2014	U-731	ODE	May 25, 2013
		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-371		
		6561976	Aug 28, 2018	U-731		
		6561977	Oct 23, 2020	U-371		
		6561977	Oct 23, 2020	U-731		
		6755784	Oct 23, 2020	U-731		
		6755784	Oct 23, 2020	U-371		
		6869399	Oct 23, 2020	U-733		
		6869399	Oct 23, 2020	U-732		
		6869399	Oct 23, 2020	U-731		
		6869399	Oct 23, 2020	U-371		
		6908432	Aug 28, 2018	U-371		
		6908432	Aug 28, 2018	U-731		
	>A>	7141018	Oct 23, 2020	U-733		
	>A>	7141018	Oct 23, 2020	U-371		
	>A>	7141018	Oct 23, 2020	U-731		
	>A>	7141018	Oct 23, 2020	U-732		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO		PATENT NO	PATENT EXPIRATON DATE		PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
THALIDOMIDE - THALOMID							
020785 003	**>A>**	5629327	May 13, 2014		U-731	ODE	May 25, 2013
		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-371		
		6561976	Aug 28, 2018		U-731		
		6561977	Oct 23, 2020		U-371		
		6561977	Oct 23, 2020		U-731		
		6755784	Oct 23, 2020		U-731		
		6755784	Oct 23, 2020		U-371		
		6869399	Oct 23, 2020		U-732		
		6869399	Oct 23, 2020		U-733		
		6869399	Oct 23, 2020		U-731		
		6869399	Oct 23, 2020		U-371		
		6908432	Aug 28, 2018		U-371		
		6908432	Aug 28, 2018		U-731		
	>A>	7141018	Oct 23, 2020		U-733		
	>A>	7141018	Oct 23, 2020		U-371		
	>A>	7141018	Oct 23, 2020		U-731		
	>A>	7141018	Oct 23, 2020		U-732		
THYROTROPIN ALFA - THYROGEN							
020898 001						M-53	Jan 23, 2009
TIOTROPIUM BROMIDE MONOHYDRATE - SPIRIVA							
021395 001		7070800	Jan 22, 2022		DP U-566		
		RE38912	Oct 11, 2021		DP		
TOPIRAMATE - TOPAMAX							
020505 001		7018983	Oct 13, 2015		U-723		
TOPIRAMATE - TOPAMAX							
020505 002		7018983	Oct 13, 2015		U-723		
TOPIRAMATE - TOPAMAX							
020505 003		7018983	Oct 13, 2015		U-723		
TOPIRAMATE - TOPAMAX							
020505 004		7018983	Oct 13, 2015		U-723		
TOPIRAMATE - TOPAMAX							
020505 005		7018983	Oct 13, 2015		U-723		
TOPIRAMATE - TOPAMAX							
020505 006		7018983	Oct 13, 2015		U-723		
TOPIRAMATE - TOPAMAX SPRINKLE							
020844 001		7018983	Oct 13, 2015		U-723		
		7125560	Mar 01, 2019		U-766		
TOPIRAMATE - TOPAMAX SPRINKLE							
020844 002		7018983	Oct 13, 2015		U-723		
		7125560	Mar 01, 2019		U-766		
TOPIRAMATE - TOPAMAX SPRINKLE							
020844 003		7018983	Oct 13, 2015		U-723		
		7125560	Mar 01, 2019		U-766		
TOPOTECAN HYDROCHLORIDE - HYCAMTIN							
020671 001		5004758	May 28, 2010	DS	DP U-741		
TRAVOPROST - TRAVATAN Z							
021994 001		5510383	Aug 03, 2013		DP U-383		
		5889052	Dec 02, 2014		DP U-383		
		6503497	May 06, 2012		DP		
		6849253	May 06, 2012		DP		

PRESCRIPTION AND OTC DRUG PRODUCT PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE			PATENT CODES		EXCLUSIVITY CODE(S)		EXCLUSIVITY EXPIRATION DATE	
TREPROSTINIL SODIUM - REMODULIN										
021272 001	5153222	Oct	06,	2014		U-455				
TREPROSTINIL SODIUM - REMODULIN										
021272 002	5153222	Oct	06,	2014		U-455				
TREPROSTINIL SODIUM - REMODULIN										
021272 003	5153222	Oct	06,	2014		U-455				
TREPROSTINIL SODIUM - REMODULIN										
021272 004	5153222	Oct	06,	2014		U-455				
URSODIOL - URSO FORTE										
020675 002	4859660	Nov	19,	2007		U-740				
VALDECOXIB - BEXTRA										
021341 002	7135489	Aug	12,	2017	DS	DP				
VALDECOXIB - BEXTRA										
021341 003	7135489	Aug	12,	2017	DS	DP				
VARENICLINE TARTRATE - CHANTIX										
021928 001	6410550	Nov	13,	2018	DS	DP	U-56	NCE	May	10, 2011
	6890927	May	06,	2022	DS	DP	U-56			
VARENICLINE TARTRATE - CHANTIX										
021928 002	6410550	Nov	13,	2018	DS	DP	U-56	NCE	May	10, 2011
	6890927	May	06,	2022	DS	DP	U-56			
VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE										
076690 001								PC	Jan	30, 2007
VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE										
076690 002								PC	Jan	30, 2007
VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE										
076690 003								PC	Jan	30, 2007
VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE										
076690 004								PC	Jan	30, 2007
VENLAFAXINE HYDROCHLORIDE - VENLAFAXINE HYDROCHLORIDE										
076690 005								PC	Jan	30, 2007
VORINOSTAT - ZOLINZA										
021991 001	6087367	Oct	04,	2011		U-776		NCE	Oct	06, 2011
	RE38506	Nov	29,	2011	DS	DP		>A> ODE	Oct	06, 2013
ZANAMIVIR - RELENZA										
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				
ZIPRASIDONE HYDROCHLORIDE - GEODON										
020825 001	4831031	Mar	02,	2012	DS	DP	U-720	I-492	Aug	19, 2007
ZIPRASIDONE HYDROCHLORIDE - GEODON										
020825 002	4831031	Mar	02,	2012	DS	DP	U-720	I-492	Aug	19, 2007
ZIPRASIDONE HYDROCHLORIDE - GEODON										
020825 003	4831031	Mar	02,	2012	DS	DP	U-720	I-492	Aug	19, 2007
ZIPRASIDONE HYDROCHLORIDE - GEODON										
020825 004	4831031	Mar	02,	2012	DS	DP	U-720	I-492	Aug	19, 2007
ZIPRASIDONE HYDROCHLORIDE - GEODON										
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			
ZIPRASIDONE MESYLATE - GEODON										
020919 001	4831031	Mar	02,	2012	DS	DP	U-720			

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY LIST

See report footnotes for information regarding report content

APPL/PROD NO	PATENT NO	PATENT EXPIRATON DATE	PATENT CODES		EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
ZOLEDRONIC ACID - ZOMETA						
021223 001	4939130	Sep 02, 2012	DS	DP U-53		
ZOLEDRONIC ACID - ZOMETA						
021223 002	4939130	Sep 02, 2012	DS	DP U-53		
ZOLPIDEM TARTRATE - AMBIEN						
019908 001	4382938	Oct 21, 2006		U-74		
	4382938*PED	Apr 21, 2007				
ZOLPIDEM TARTRATE - AMBIEN						
019908 002	4382938	Oct 21, 2006		U-74		
	4382938*PED	Apr 21, 2007				
ZOLPIDEM TARTRATE - AMBIEN CR						
021774 001	4382938	Oct 21, 2006	DS	U-682	NDF	Sep 02, 2008
	4382938*PED	Apr 21, 2007			PED	Mar 02, 2009
	6514531	Dec 01, 2019		DP		
	6514531*PED	Jun 01, 2020				
ZOLPIDEM TARTRATE - AMBIEN CR						
021774 002	4382938	Oct 21, 2006	DS	U-682	NDF	Sep 02, 2008
	4382938*PED	Apr 21, 2007			PED	Mar 02, 2009
	6514531	Dec 01, 2019		DP		
	6514531*PED	Jun 01, 2020				

Footnotes:

1. Patents are published upon receipt by the Orange Book Staff and may not reflect the official receipt date as described in 21 CFR 314.53(d)(5).
2. Patents submitted on FDA Form 3542 and listed after August 18, 2003 will have one to three patent codes indicating specific patent claims as submitted by the sponsor:
   DS = Drug Substance claim
   DP = Drug Product claim
   U and number = Method of Use claim (may be multiple). Specific Method of use claims are listed at http://www.fda.gov/cder/orange/patex.htm
3. Patents listed prior to August 18, 2003 are flagged with method of use claims only as applicable and submitted by the sponsor. They may not be flagged with respect to other claims which may apply.
4. *PED and PED represent pediatric exclusivity. Patents with pediatric exclusivity granted after August 18, 2003 will be indicated with *PED as was done prior to August 18, 2003. Patents with *PED added after August 18, 2003 will not contain any information relative to the patent itself other than the *PED extension. Information related specifically to the patent will be conveyed on the original patent only.

5. *** U.S. Patent Nos. RE 36481 and RE 36520 were 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 remained 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 were triggered and run. For additional information on this matter, please refer to Docket Nos. 2005P-0008 and 2005P-0046. Patents were subsequently delisted in the December 2006 Orange Book update as the exclusivity periods have triggered and run to expiration.

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, 25th 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