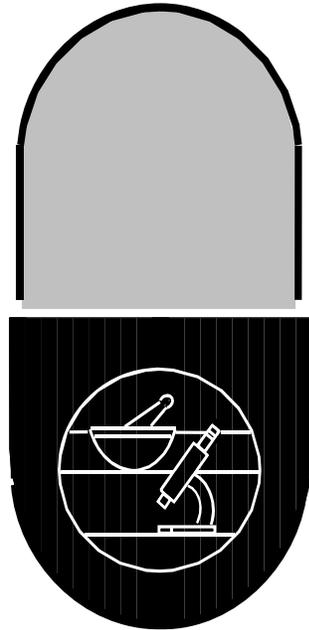


**CUMULATIVE
SUPPLEMENT 06
June 2007**



**APPROVED
DRUG PRODUCTS**

**WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS**

27th EDITION

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

2007

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

**APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS**

27th EDITION

Cumulative Supplement 06

June 2007

CONTENTS

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

**APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS**

27th EDITION

**CUMULATIVE SUPPLEMENT 06
June 2007**

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, 27th 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are for exportation, are for military use, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.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 26th Edition List will then be added to the "Discontinued Drug Product List" appearing in the 27th Edition. The current edition

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

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

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

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

1.2 CUMULATIVE SUPPLEMENT CONTENT

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

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

Currently, the monthly PDF CS includes:

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

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

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

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

1.3 APPLICANT NAME CHANGES

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

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

<u>FORMER APPLICANT NAME</u> <u>(FORMER ABBREVIATED NAME)</u>	<u>NEW APPLICANT NAME</u> <u>(NEW ABBREVIATED NAME)</u>
ALPHARMA BRANDED PRODUCTS DIVISION INC (ALPHARMA BRANDED)	ALPHARMA PHARMACEUTICALS LLC (ALPHARMA PHARMS)
ALPHARMA US PHARMACEUTICALS DIVISION (ALPHARMA US PHARMS)	ALPHARMA PHARMACEUTICALS LLC (ALPHARMA PHARMS)
ALPHARMA USPD INC (ALPHARMA)	ALPHARMA PHARMACEUTICALS LLC (ALPHARMA PHARMS)
BAKER NORTON PHARMACEUTICALS INC (BAKER NORTON)	TEVA PARENTERAL MEDICINES INC (TEVA PARENTERAL)
BERLEX INC (BERLEX)	BAYER HEALTHCARE PHARMACEUTICALS INC (BAYER HLTHC)
BIONICHE PHARMA (BIONICHE PHARMA)	BIONICHE PHARMA USA LLC (BIONICHE PHARMA)

BIONICHE PHARMA (CANADA) LTD (BIONICHE (CANADA))	BIONICHE PHARMA USA LLC (BIONICHE PHARMA)
BIONICHE PHARMA USA INC (BIONICHE PHARM USA)	BIONICHE PHARMA USA LLC (BIONICHE PHARMA)
FAULDING INC (FAULDING)	ALPHARMA PHARMACEUTICALS LLC (ALPHARMA PHARMS)
FAULDING PHARMACEUTICALS INC (FAULDING PHARMS)	ALPHARMA PHARMACEUTICALS LLC (ALPHARMA PHARMS)
GENSIA LABORATORIES LTD (GENSIA)	TEVA PARENTERAL MEDICINES INC (TEVA PARENTERAL)
GENSIA SICOR PHARMACEUTICALS INC (GENSIA SICOR PHARMS)	TEVA PARENTERAL MEDICINES INC (TEVA PARENTERAL)
IVAX PHARMACEUTICALS INC (IVAX PHARMS)	TEVA PARENTERAL MEDICINES INC (TEVA PARENTERAL)
KOS LIFE SCIENCES INC (KOS LIFE)	ABBOTT LABORATORIES (ABBOTT)
MAYNE PHARMA USA INC (MAYNE PHARMA USA)	HOSPIRA INC (HOSPIRA)
PHARMACIA AND UPJOHN CO (PHARMACIA AND	TEVA PARENTERAL MEDICINES INC (TEVA PARENTERAL)
QLT USA INC (QLT USA)	TOLMAR INC (TOLMAR)
SICOR PHARMACEUTICALS INC (SICOR PHARMS)	TEVA PARENTERAL MEDICINES INC (TEVA PARENTERAL)
ZENITH GOLDLINE PHARMACEUTICALS (ZENITH GOLDLINE)	TEVA PARENTERAL MEDICINES INC (TEVA PARENTERAL)

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

<u>CATEGORIES COUNTED</u>	<u>DEC 2006</u>	<u>MAR 2007</u>	<u>JUN 2007</u>	<u>SEPT 2007</u>
DRUG PRODUCTS LISTED	11896	12063	11900	
SINGLE SOURCE	2471 (20.8%)	2471 (20.5%)	2483 (20.9%)	

MULTISOURCE	9336 (78.5%)	9503 (78.8%)	9328 (78.4%)
THERAPEUTICALLY EQUIVALENT	9139 (76.8%)	9320 (77.3%)	9148 (76.9%)
NOT THERAPEUTICALLY EQUIVALENT EXCEPTIONS ¹	197 (1.7%) 89 (0.7%)	183 (1.5%) 89 (0.7%)	180 (1.5%) 89 (0.7%)
NEW MOLECULAR ENTITIES APPROVED	10	4	7
NUMBER OF APPLICANTS	666	675	679

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

1.6 CUMULATIVE SUPPLEMENT LEGEND

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

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

New ingredient(s), new dosage form; route(s), new trade names, and new product additions are preceded by >A> during the action month.

The change month is the current CS month; the change code for new approvals is NEWA. Following months will display the same information without the >A>.

Changes to currently listed products will list two records. The deleted product record will be preceded by >D>. The product record change addition being made will be preceded by >A>. Following months will display only the >A> record without the >A>.

All changes that occur to the product through the Annual year will be listed. The change month and change code will document the change.

The change code and description:

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

PRESCRIPTION DRUG PRODUCT LIST - 27TH EDITION

RX DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 6 - June 2007

1-1

ABARELIX

INJECTABLE; INTRAMUSCULAR

PLENAXIS

@ SPECIALITY EUROPEAN 100MG/VIAL N21320 001 Nov 25, 2003 Feb CAHN

ACETAMINOPHEN; BUTALBITAL

CAPSULE; ORAL

>D> BUCET

>D> AB MALLINCKRODT 650MG;50MG N88991 001 Jun 28, 1985 Jun DISC

>A> @ 650MG;50MG N88991 001 Jun 28, 1985 Jun DISC

PHRENILIN FORTE

>D> AB + VALEANT 650MG;50MG N88831 001 Jun 19, 1985 Jun CTEC

>A> + 650MG;50MG N88831 001 Jun 19, 1985 Jun CTEC

>D> TENCON

>D> AB MALLINCKRODT 650MG;50MG N89405 001 May 15, 1990 Jun DISC

>A> @ 650MG;50MG N89405 001 May 15, 1990 Jun DISC

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

TABLET; ORAL

>D> BUTALBITAL, APAP, AND CAFFEINE

>D> AB WATSON LABS 325MG;50MG;40MG N89536 001 Feb 16, 1988 Jun DISC

>A> @ 325MG;50MG;40MG N89536 001 Feb 16, 1988 Jun DISC

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

CAPSULE; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

>D> AA BOCA PHARMA 356.4MG;30MG;16MG N40688 001 Apr 03, 2007 Jun CAHN

AA 356.4MG;30MG;16MG N40688 001 Apr 03, 2007 Mar NEWA

>A> AA E5 PHARMA INC 356.4MG;30MG;16MG N40688 001 Apr 03, 2007 Jun CAHN

AA + MIKART 356.4MG;30MG;16MG N40109 001 Aug 26, 1997 Mar CTEC

TABLET; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE

AA BOCA PHARMA 712.8MG;60MG;32MG N40701 001 Apr 03, 2007 Mar NEWA

AA + MIKART 712.8MG;60MG;32MG N40316 001 Apr 28, 1999 Feb CTEC

AA WEST WARD 712.8MG;60MG;32MG N40637 001 Sep 22, 2006 Feb CTEC

ACETAMINOPHEN; CODEINE PHOSPHATE

TABLET; ORAL

ACETAMINOPHEN AND CODEINE PHOSPHATE

>D> AA MUTUAL PHARM 300MG;30MG N89672 001 Feb 10, 1988 Jun DISC

>A> @ 300MG;30MG N89672 001 Feb 10, 1988 Jun DISC

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

>D> ALLAY

>D> AA IVAX PHARMS 500MG;5MG N89907 001 Jan 13, 1989 Jun DISC

>A> @ 500MG;5MG N89907 001 Jan 13, 1989 Jun DISC

>D> HYDROCET

>D> AA MALLINCKRODT 500MG;5MG N89006 001 Aug 09, 1985 Jun DISC

>A> @ 500MG;5MG N89006 001 Aug 09, 1985 Jun DISC

HYDROCODONE BITARTRATE AND ACETAMINOPHEN

>D> AA MALLINCKRODT 500MG;5MG N88956 001 Jul 19, 1985 Jun DISC

>A> @ 500MG;5MG N88956 001 Jul 19, 1985 Jun DISC

TABLET; ORALHYDROCODONE BITARTRATE AND ACETAMINOPHEN

AA	INTERPHARM	500MG;10MG	N40813	001	Feb 23, 2007	Feb	NEWA
>D>	AA	IVAX PHARMS	N89696	001	Apr 21, 1988	Jun	DISC
>A>	@	500MG;5MG	N89696	001	Apr 21, 1988	Jun	DISC
>D>	AA	MIKART	N89271	001	Jul 16, 1986	Jun	DISC
>A>	@	500MG;5MG	N89271	001	Jul 16, 1986	Jun	DISC
>D>	AA	500MG;5MG	N89697	001	Jan 28, 1992	Jun	DISC
>A>	@	500MG;5MG	N89697	001	Jan 28, 1992	Jun	DISC
	@ UCB INC	500MG;10MG	N40210	001	Aug 13, 1997	May	DISC
	@	650MG;7.5MG	N40134	001	Nov 21, 1996	May	DISC
>D>	AA	VINTAGE PHARMS	N89831	001	Sep 07, 1988	Jun	DISC
>A>	@	500MG;5MG	N89831	001	Sep 07, 1988	Jun	DISC

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATETABLET; ORALPROPOXYPHENE NAPSYLATE AND ACETAMINOPHEN

>D>	AB	MUTUAL PHARM	N70771	001	Mar 21, 1986	Jun	DISC
>A>	@	650MG;100MG	N70771	001	Mar 21, 1986	Jun	DISC
>D>	AB	650MG;100MG	N70775	001	Mar 21, 1986	Jun	DISC
>A>	@	650MG;100MG	N70775	001	Mar 21, 1986	Jun	DISC
>D>	AB	MYLAN	N72195	001	Feb 16, 1988	Jun	DISC
>A>	@	650MG;100MG	N72195	001	Feb 16, 1988	Jun	DISC
	AB	WOCKHARDT	N77677	001	Mar 16, 2007	Mar	NEWA
	AB	650MG;100MG	N77677	002	Mar 16, 2007	Mar	NEWA

ACETAZOLAMIDECAPSULE, EXTENDED RELEASE; ORALDIAMOX

	+	DURAMED PHARMS BARR	500MG	N12945	001		May	CMFD
--	---	---------------------	-------	--------	-----	--	-----	------

ACETIC ACID, GLACIALSOLUTION/DROPS; OTICVOSOL

>A>	@	HI TECH PHARMA	2%	N12179	001		Jun	CAHN
>D>	@	MEDPOINTE PHARM HLC	2%	N12179	001		Jun	CAHN

ACETIC ACID, GLACIAL; HYDROCORTISONESOLUTION/DROPS; OTICVOSOL HC

>A>	AT	+	HI TECH PHARMA	2%;1%	N12770	001		Jun	CAHN
>D>	AT	+	MEDPOINTE PHARM HLC	2%;1%	N12770	001		Jun	CAHN

ACYCLOVIRCAPSULE; ORALACYCLOVIR

>D>	AB	CLONMEL HLTHCARE	200MG	N74872	001	Apr 22, 1997	Jun	DISC
>A>	@	200MG	N74872	001	Apr 22, 1997	Jun	DISC	
>D>	AB	MYLAN	200MG	N74727	001	Apr 22, 1997	Jun	DISC
>A>	@	200MG	N74727	001	Apr 22, 1997	Jun	DISC	
>D>	AB	TEVA	200MG	N74828	001	Apr 22, 1997	Jun	DISC
>A>	@	200MG	N74828	001	Apr 22, 1997	Jun	DISC	
>D>	AB	TEVA PHARMS	200MG	N74914	001	Nov 26, 1997	Jun	DISC
>A>	@	200MG	N74914	001	Nov 26, 1997	Jun	DISC	

TABLET; ORAL

ACYCLOVIR

>D>	AB	MYLAN	400MG	N75211 001	Sep 28, 1998	Jun	DISC
>A>		@	400MG	N75211 001	Sep 28, 1998	Jun	DISC
>D>	AB		800MG	N75211 002	Sep 28, 1998	Jun	DISC
>A>		@	800MG	N75211 002	Sep 28, 1998	Jun	DISC
>D>	AB	TEVA PHARMS	400MG	N75021 001	Mar 18, 1998	Jun	DISC
>A>		@	400MG	N75021 001	Mar 18, 1998	Jun	DISC
>D>	AB		800MG	N75021 002	Mar 18, 1998	Jun	DISC
>A>		@	800MG	N75021 002	Mar 18, 1998	Jun	DISC

ADAPALENE

GEL; TOPICAL

DIFFERIN

>A>	+	GALDERMA LABS LP	0.3%	N21753 001	Jun 19, 2007	Jun	NEWA
-----	---	------------------	------	------------	--------------	-----	------

ADENOSINE

INJECTABLE; INJECTION

ADENOSINE

>A>	AP	GLAND PHARMA LTD	3MG/ML	N77283 001	Jun 14, 2007	Jun	NEWA
-----	----	------------------	--------	------------	--------------	-----	------

ALBUTEROL

AEROSOL, METERED; INHALATION

ALBUTEROL

+		ARMSTRONG PHARMS	0.09MG/INH	N72273 001	Aug 14, 1996	Mar	CRLD
		@ IVAX PHARMS	0.09MG/INH	N73272 001	Dec 28, 1995	Mar	DISC
		VENTOLIN					
		@ GLAXOSMITHKLINE	0.09MG/INH	N18473 001		Mar	DISC

ALBUTEROL SULFATE

SOLUTION; INHALATION

ALBUTEROL SULFATE

AN		APOTEX INC	EQ 0.083% BASE	N75717 001	Feb 02, 2007	Jan	NEWA
		@ ROXANE	EQ 0.083% BASE	N75129 001	Feb 13, 2001	May	DISC

SYRUP; ORAL

ALBUTEROL SULFATE

>A>	AA	VISTAPHARM	EQ 2MG BASE/5ML	N77788 001	Jun 26, 2007	Jun	NEWA
-----	----	------------	-----------------	------------	--------------	-----	------

TABLET; ORAL

ALBUTEROL SULFATE

>D>	AB	TEVA	EQ 2MG BASE	N72619 001	Dec 05, 1989	Jun	DISC
>A>		@	EQ 2MG BASE	N72619 001	Dec 05, 1989	Jun	DISC
>D>	AB		EQ 2MG BASE	N72779 001	Jun 25, 1993	Jun	DISC
>A>		@	EQ 2MG BASE	N72779 001	Jun 25, 1993	Jun	DISC
>D>	AB		EQ 2MG BASE	N72938 001	Mar 30, 1990	Jun	DISC
>A>		@	EQ 2MG BASE	N72938 001	Mar 30, 1990	Jun	DISC
>D>	AB		EQ 4MG BASE	N72620 001	Dec 05, 1989	Jun	DISC
>A>		@	EQ 4MG BASE	N72620 001	Dec 05, 1989	Jun	DISC
>D>	AB		EQ 4MG BASE	N72939 001	Mar 30, 1990	Jun	DISC
>A>		@	EQ 4MG BASE	N72939 001	Mar 30, 1990	Jun	DISC
>D>	AB	WATSON LABS	EQ 2MG BASE	N72629 001	Jan 31, 1991	Jun	DISC
>A>		@	EQ 2MG BASE	N72629 001	Jan 31, 1991	Jun	DISC
>D>	AB		EQ 4MG BASE	N72765 001	Aug 28, 1991	Jun	DISC
>A>		@	EQ 4MG BASE	N72765 001	Aug 28, 1991	Jun	DISC

TABLET, EXTENDED RELEASE; ORAL

ALBUTEROL SULFATE

AB		MYLAN	EQ 4MG BASE	N78092 002	Jan 29, 2007	Jan	NEWA
----	--	-------	-------------	------------	--------------	-----	------

TABLET, EXTENDED RELEASE; ORAL

ALBUTEROL SULFATE

AB	MYLAN	EQ 8MG BASE	N78092 001	Jan 29, 2007	Jan	NEWA
	VOSPIRE ER					
AB	DAVA PHARMS INC	EQ 4MG BASE	N76130 002	Sep 26, 2002	Jan	CTEC
AB	+	EQ 8MG BASE	N76130 003	Sep 26, 2002	Jan	CTEC

ALENDRONATE SODIUM; CHOLECALCIFEROL

TABLET; ORAL

FOSAMAX PLUS D

	MERCK	EQ 70MG BASE;2,800 IU	N21762 001	Apr 07, 2005	Apr	CRLD
+		EQ 70MG BASE;5,600 IU	N21762 002	Apr 26, 2007	Apr	NEWA

ALISKIREN HEMIFUMARATE

TABLET; ORAL

TEKTURNA

	NOVARTIS	EQ 150MG BASE	N21985 001	Mar 05, 2007	Mar	NEWA
+		EQ 300MG BASE	N21985 002	Mar 05, 2007	Mar	NEWA

ALLOPURINOL SODIUM

INJECTABLE; INJECTION

ALOPRIM

AP	+	BIONICHE PHARMA	EQ 500MG BASE/VIAL	N20298 001	May 17, 1996	May	CAHN
----	---	-----------------	--------------------	------------	--------------	-----	------

ALPRAZOLAM

TABLET; ORAL

ALPRAZOLAM

AB	APOTEX INC	0.25MG	N77741 001	Jan 19, 2007	Jan	NEWA	
AB		0.5MG	N77741 002	Jan 19, 2007	Jan	NEWA	
AB		1MG	N77741 003	Jan 19, 2007	Jan	NEWA	
AB		2MG	N77741 004	Jan 19, 2007	Jan	NEWA	
AB	DAVA INTL INC	0.25MG	N74174 001	Oct 19, 1993	Mar	CMFD	
	@	0.25MG	N74174 001	Oct 19, 1993	Feb	CAHN	
AB		0.5MG	N74174 002	Oct 19, 1993	Mar	CMFD	
	@	0.5MG	N74174 002	Oct 19, 1993	Feb	CAHN	
AB		1MG	N74174 003	Oct 19, 1993	Mar	CMFD	
	@	1MG	N74174 003	Oct 19, 1993	Feb	CAHN	
AB		2MG	N74174 004	Oct 19, 1993	Mar	CMFD	
	@	2MG	N74174 004	Oct 19, 1993	Feb	CAHN	
>D>	AB	IVAX PHARMS	0.25MG	N74294 001	Jul 29, 1994	Jun	DISC
>A>		@	0.25MG	N74294 001	Jul 29, 1994	Jun	DISC
>D>	AB		0.5MG	N74294 002	Jul 29, 1994	Jun	DISC
>A>		@	0.5MG	N74294 002	Jul 29, 1994	Jun	DISC
>D>	AB		1MG	N74294 003	Jul 29, 1994	Jun	DISC
>A>		@	1MG	N74294 003	Jul 29, 1994	Jun	DISC
>D>	AB		2MG	N74294 004	Jul 29, 1994	Jun	DISC
>A>		@	2MG	N74294 004	Jul 29, 1994	Jun	DISC
>D>	AB	WATSON LABS	0.25MG	N74479 001	Jan 21, 1997	Jun	DISC
>A>		@	0.25MG	N74479 001	Jan 21, 1997	Jun	DISC
>D>	AB		0.5MG	N74479 002	Jan 21, 1997	Jun	DISC
>A>		@	0.5MG	N74479 002	Jan 21, 1997	Jun	DISC
>D>	AB		1MG	N74479 003	Jan 21, 1997	Jun	DISC
>A>		@	1MG	N74479 003	Jan 21, 1997	Jun	DISC

TABLET, EXTENDED RELEASE; ORAL

ALPRAZOLAM

AB	ACTAVIS ELIZABETH	0.5MG	N78056 001	Feb 13, 2007	Jan	NEWA
----	-------------------	-------	------------	--------------	-----	------

TABLET, EXTENDED RELEASE; ORAL

ALPRAZOLAM

AB	ACTAVIS ELIZABETH	1MG	N78056 002	Feb 13, 2007	Jan	NEWA
AB		2MG	N78056 003	Feb 13, 2007	Jan	NEWA
AB		3MG	N78056 004	Feb 13, 2007	Jan	NEWA
AB	COREPHARMA	0.5MG	N77996 001	Jan 31, 2007	Jan	NEWA
AB		1MG	N77996 002	Jan 31, 2007	Jan	NEWA
AB		2MG	N77996 003	Jan 31, 2007	Jan	NEWA
AB		3MG	N77996 004	Jan 31, 2007	Jan	NEWA
AB	IMPAX LABS	0.5MG	N77968 004	May 24, 2007	May	NEWA
AB		1MG	N77968 003	May 24, 2007	May	NEWA
AB		2MG	N77968 002	May 24, 2007	May	NEWA
AB		3MG	N77968 001	May 24, 2007	May	NEWA
AB	TEVA PHARMS	0.5MG	N77979 001	Feb 28, 2007	Feb	NEWA
AB		1MG	N77979 002	Feb 28, 2007	Feb	NEWA
AB		2MG	N77979 003	Feb 28, 2007	Feb	NEWA
AB		3MG	N77979 004	Feb 28, 2007	Feb	NEWA

TABLET, ORALLY DISINTEGRATING; ORAL

NIRAVAM

+	SCHWARZ PHARMA	1MG	N21726 003	Jan 19, 2005	Jan	CRLD
		2MG	N21726 004	Jan 19, 2005	Jan	CRLD

ALPROSTADIL

INJECTABLE; INJECTION

CAVERJECT IMPULSE

	PHARMACIA AND UPJOHN	0.01MG/VIAL	N21212 001	Jun 11, 2002	Apr	CTNA
		0.02MG/VIAL	N21212 002	Jun 11, 2002	Apr	CTNA

>A> AMBRISENTAN

>A> TABLET; ORAL

>A> LETAIRIS

>A>	GILEAD	5MG	N22081 001	Jun 15, 2007	Jun	NEWA
>A>	+	10MG	N22081 002	Jun 15, 2007	Jun	NEWA

AMILORIDE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

AMILORIDE HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

>D>	AB	TEVA	EQ 5MG ANHYDROUS;50MG	N70795 001	Apr 17, 1988	Jun	DISC
>A>		@	EQ 5MG ANHYDROUS;50MG	N70795 001	Apr 17, 1988	Jun	DISC

AMINO ACIDS

INJECTABLE; INJECTION

NOVAMINE 11.4%

+	HOSPIRA	11.4% (11.4GM/100ML)	N17957 003	Aug 09, 1982	Jan	CRLD
---	---------	----------------------	------------	--------------	-----	------

NOVAMINE 15%

+	HOSPIRA	15% (15GM/100ML)	N17957 004	Nov 28, 1986	Jan	CRLD
---	---------	------------------	------------	--------------	-----	------

>D> AMINOSALICYLATE SODIUM

>D> TABLET; ORAL

>D> SODIUM P.A.S.

>D>	+	LANNETT	500MG	N80138 002		Jun	DISC
>A>		@	500MG	N80138 002		Jun	DISC

AMIODARONE HYDROCHLORIDE

INJECTABLE; INJECTION

CORDARONE

@	WYETH PHARMS INC	50MG/ML	N20377	001	Aug 03, 1995	May	DISC
---	------------------	---------	--------	-----	--------------	-----	------

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AMITRIPTYLINE HYDROCHLORIDE

>D>	AB	TEVA	10MG	N84910	003		Jun	DISC
>A>		@	10MG	N84910	003		Jun	DISC
>D>	AB		25MG	N85031	001		Jun	DISC
>A>		@	25MG	N85031	001		Jun	DISC
>D>	AB		100MG	N85836	001		Jun	DISC
>A>		@	100MG	N85836	001		Jun	DISC

AMITRIPTYLINE HYDROCHLORIDE; CHLORDIAZEPOXIDE

TABLET; ORAL

CHLORDIAZEPOXIDE AND AMITRIPTYLINE HYDROCHLORIDE

>D>	AB	WATSON LABS	EQ 12.5MG BASE;5MG	N72052	001	Dec 16, 1988	Jun	DISC
>A>		@	EQ 12.5MG BASE;5MG	N72052	001	Dec 16, 1988	Jun	DISC
>D>	AB		EQ 25MG BASE;10MG	N72053	001	Dec 16, 1988	Jun	DISC
>A>		@	EQ 25MG BASE;10MG	N72053	001	Dec 16, 1988	Jun	DISC

AMLODIPINE BESYLATE

TABLET; ORAL

AMLODIPINE BESYLATE

	AB	APOTEX	EQ 2.5MG BASE	N76719	001	May 23, 2007	May	NEWA
	AB		EQ 5MG BASE	N76719	002	May 23, 2007	May	NEWA
	AB		EQ 10MG BASE	N76719	003	May 23, 2007	May	NEWA
>A>	AB	GENPHARM	EQ 2.5MG BASE	N77362	001	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 5MG BASE	N77362	002	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 10MG BASE	N77362	003	Jul 09, 2007	Jun	NEWA
>A>	AB	RANBAXY	EQ 2.5MG BASE	N77974	001	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 5MG BASE	N77974	002	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 10MG BASE	N77974	003	Jul 09, 2007	Jun	NEWA
>A>	AB	ROXANE	EQ 2.5MG BASE	N77262	001	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 5MG BASE	N77262	002	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 10MG BASE	N77262	003	Jul 09, 2007	Jun	NEWA
>A>	AB	SYNTHON LAB	EQ 2.5MG BASE	N77080	001	Jun 27, 2007	Jun	NEWA
>A>	AB		EQ 5MG BASE	N77080	002	Jun 27, 2007	Jun	NEWA
>A>	AB		EQ 10MG BASE	N77080	003	Jun 27, 2007	Jun	NEWA
>A>	AB	TEVA	EQ 2.5MG BASE	N76846	001	Jun 28, 2007	Jun	NEWA
>A>	AB		EQ 5MG BASE	N76846	002	Jun 28, 2007	Jun	NEWA
>A>	AB		EQ 10MG BASE	N76846	003	Jun 28, 2007	Jun	NEWA
>A>	AB	UPSHER SMITH	EQ 2.5MG BASE	N77759	001	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 5MG BASE	N77759	002	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 10MG BASE	N77759	003	Jul 09, 2007	Jun	NEWA
>A>	AB	ZYDUS PHARMS USA	EQ 2.5MG BASE	N78226	001	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 5MG BASE	N78226	002	Jul 09, 2007	Jun	NEWA
>A>	AB		EQ 10MG BASE	N78226	003	Jul 09, 2007	Jun	NEWA

AMLODIPINE BESYLATE; BENAZEPRIL HYDROCHLORIDE

CAPSULE; ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

AB	TEVA PHARMS	EQ 2.5MG BASE;10MG	N77179	001	May 18, 2007	May	NEWA
----	-------------	--------------------	--------	-----	--------------	-----	------

CAPSULE; ORAL

AMLODIPINE BESYLATE AND BENAZEPRIL HYDROCHLORIDE

AB	TEVA PHARMS	EQ 5MG BASE;10MG	N77179 002	May 18, 2007	May	NEWA
AB		EQ 5MG BASE;20MG	N77179 003	May 18, 2007	May	NEWA
AB		EQ 10MG BASE;20MG	N77179 004	May 18, 2007	May	NEWA

LOTREL

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

>A> AMLODIPINE BESYLATE; VALSARTAN

>A> TABLET; ORAL

>A> EXFORGE

>A>	NOVARTIS	EQ 5MG BASE;160MG	N21990 002	Jun 20, 2007	Jun	NEWA
>A>		EQ 5MG BASE;320MG	N21990 004	Jun 20, 2007	Jun	NEWA
>A>		EQ 10MG BASE;160MG	N21990 003	Jun 20, 2007	Jun	NEWA
>A>	+	EQ 10MG BASE;320MG	N21990 005	Jun 20, 2007	Jun	NEWA

AMMONIUM LACTATE

CREAM; TOPICAL

LAC-HYDRIN

>A>	AB	+	RANBAXY	EQ 12% BASE	N20508 001	Aug 29, 1996	Jun	CAHN
>D>	AB	+	WESTWOOD SQUIBB	EQ 12% BASE	N20508 001	Aug 29, 1996	Jun	CAHN

LOTION; TOPICAL

LAC-HYDRIN

>A>	AB	+	RANBAXY	EQ 12% BASE	N19155 001	Apr 24, 1985	Jun	CAHN
>D>	AB	+	WESTWOOD SQUIBB	EQ 12% BASE	N19155 001	Apr 24, 1985	Jun	CAHN

AMOXAPINE

TABLET; ORAL

AMOXAPINE

>D>	AB		SANDOZ	25MG	N72943 001	Jun 28, 1991	Jun	DISC
>A>			@	25MG	N72943 001	Jun 28, 1991	Jun	DISC
>D>	AB			50MG	N72944 001	Jun 28, 1991	Jun	DISC
>A>			@	50MG	N72944 001	Jun 28, 1991	Jun	DISC
>D>	AB			100MG	N72878 001	Jun 28, 1991	Jun	DISC
>A>			@	100MG	N72878 001	Jun 28, 1991	Jun	DISC
>D>	AB			150MG	N72879 001	Jun 28, 1991	Jun	DISC
>A>			@	150MG	N72879 001	Jun 28, 1991	Jun	DISC
>D>	AB		WATSON LABS	25MG	N72418 001	May 11, 1989	Jun	DISC
>A>			@	25MG	N72418 001	May 11, 1989	Jun	DISC
>D>	AB			50MG	N72419 001	May 11, 1989	Jun	DISC
>A>			@	50MG	N72419 001	May 11, 1989	Jun	DISC
>D>	AB			100MG	N72420 001	May 11, 1989	Jun	DISC
>A>			@	100MG	N72420 001	May 11, 1989	Jun	DISC
>D>	AB	+		150MG	N72421 001	May 11, 1989	Jun	DISC
>A>			@	150MG	N72421 001	May 11, 1989	Jun	DISC
>D>	AB			150MG	N72691 001	Aug 28, 1992	Jun	CRLD
>A>	AB	+		150MG	N72691 001	Aug 28, 1992	Jun	CRLD

AMOXICILLIN

CAPSULE; ORAL

AMOXICILLIN

AB	HIKMA PHARMS	250MG	N65291 001	Feb 05, 2007	Jan	NEWA
AB		500MG	N65291 002	Feb 05, 2007	Jan	NEWA

CAPSULE; ORAL

AMOXICILLIN

>D>	AB	TEVA	250MG	N62853 001	Dec 22, 1987	Jun	DISC
>A>		@	250MG	N62853 001	Dec 22, 1987	Jun	DISC
>D>	AB		500MG	N62854 001	Dec 22, 1987	Jun	DISC
>A>		@	500MG	N62854 001	Dec 22, 1987	Jun	DISC
		TRIMOX					
		@ APOTHECON	250MG	N61885 001		May	DISC
		@	500MG	N61885 002		May	DISC

FOR SOLUTION; ORAL

AMOXICILLIN

>A>	AB	MORTON GROVE	400MG/5ML	N65319 002	Jun 18, 2007	Jun	NEWA
-----	----	--------------	-----------	------------	--------------	-----	------

FOR SUSPENSION; ORAL

AMOXICILLIN

	AB	SANDOZ	125MG/5ML	N65387 001	Mar 26, 2007	Mar	NEWA
	AB		200MG/5ML	N65378 001	Mar 26, 2007	Mar	NEWA
	AB		250MG/5ML	N65387 002	Mar 26, 2007	Mar	NEWA
	AB		400MG/5ML	N65378 002	Mar 26, 2007	Mar	NEWA
>D>	AB	TEVA	125MG/5ML	N62946 001	Nov 01, 1988	Jun	DISC
>A>		@	125MG/5ML	N62946 001	Nov 01, 1988	Jun	DISC
>D>	AB		250MG/5ML	N63001 001	Jan 06, 1989	Jun	DISC
>A>		@	250MG/5ML	N63001 001	Jan 06, 1989	Jun	DISC

TABLET, CHEWABLE; ORAL

AMOXICILLIN

>D>	AB	TEVA	125MG	N64031 001	Dec 19, 1996	Jun	DISC
>A>		@	125MG	N64031 001	Dec 19, 1996	Jun	DISC
>D>	AB		250MG	N64031 002	Dec 19, 1996	Jun	DISC
>A>		@	250MG	N64031 002	Dec 19, 1996	Jun	DISC

TABLET, FOR SUSPENSION; ORAL

AMOXICILLIN

	AB	AUROBINDO PHARMA	200MG	N65324 001	Jan 17, 2007	Jan	NEWA
	AB		400MG	N65324 002	Jan 17, 2007	Jan	NEWA
		DISPERMOX					
	AB	RANBAXY	200MG	N65080 002	Aug 11, 2003	Jan	CTEC
	AB	+	400MG	N65080 001	Aug 11, 2003	Jan	CTEC

AMOXICILLIN; CLAVULANATE POTASSIUM

FOR SUSPENSION; ORAL

AMOXICILLIN AND CLAVULANATE POTASSIUM

	AB	RANBAXY	600MG/5ML;EQ 42.9MG BASE/5ML	N65207 002	Jan 30, 2007	Jan	NEWA	
		AUGMENTIN ES-600						
	AB	+	SMITHKLINE BEECHAM	600MG/5ML;EQ 42.9MG BASE/5ML	N50755 001	Jun 22, 2001	Mar	CAHN

AMPHOTERICIN B

>D>		CREAM; TOPICAL					
>D>		FUNGIZONE					
>D>	+	APOTHECON	3%	N50314 001		Jun	DISC
>A>		@	3%	N50314 001		Jun	DISC

AMPICILLIN SODIUM; SULBACTAM SODIUM

INJECTABLE; INJECTION

AMPICILLIN AND SULBACTAM

>A>	AP	GENERAMEDIX	EQ 1GM BASE/VIAL;EQ 500MG BASE/VIAL	N65316 001	Jun 29, 2007	Jun	NEWA
>A>	AP		EQ 2GM BASE/VIAL;EQ 1GM BASE/VIAL	N65316 002	Jun 29, 2007	Jun	NEWA

ARIPIPIRAZOLE

SOLUTION; ORAL

ABILIFY

+ OTSUKA PHARM 1MG/ML N21713 001 Dec 10, 2004 Apr CAHN

TABLET; ORAL

ABILIFY

OTSUKA 15MG N21436 002 Nov 15, 2002 Feb CRLD

30MG N21436 004 Nov 15, 2002 Feb CRLD

TABLET, ORALLY DISINTEGRATING; ORAL

ABILIFY

+ OTSUKA 10MG N21729 002 Jun 07, 2006 Feb CRLD

30MG N21729 005 Jun 07, 2006 Feb CRLD

>A> ARMODAFINIL

>A> TABLET; ORAL

>A> NUVIGIL

>A> CEPHALON 50MG N21875 001 Jun 15, 2007 Jun NEWA

>A> @ 100MG N21875 002 Jun 15, 2007 Jun DISC

>A> 150MG N21875 003 Jun 15, 2007 Jun NEWA

>A> + 250MG N21875 004 Jun 15, 2007 Jun NEWA

ASPIRIN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, AND CAFFEINE

>A> AB MUTUAL PHARM 325MG;50MG;40MG N78149 001 Jun 13, 2007 Jun NEWA

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

NORGESIC

AB GRACEWAY 385MG;30MG;25MG N13416 003 Oct 27, 1982 Jan CAHN

NORGESIC FORTE

AB + GRACEWAY 770MG;60MG;50MG N13416 004 Oct 27, 1982 Jan CAHN

ASPIRIN; CAFFEINE; PROPOXYPHENE HYDROCHLORIDE

CAPSULE; ORAL

DARVON COMPOUND-65

>D> AA + XANODYNE PHARM 389MG;32.4MG;65MG N10996 007 Mar 08, 1983 Jun CTEC

>A> + 389MG;32.4MG;65MG N10996 007 Mar 08, 1983 Jun CTEC

>D> PROPOXYPHENE COMPOUND 65

>D> AA TEVA 389MG;32.4MG;65MG N89025 001 Mar 29, 1985 Jun DISC

>A> @ 389MG;32.4MG;65MG N89025 001 Mar 29, 1985 Jun DISC

ATENOLOL

TABLET; ORAL

ATENOLOL

>A> AB CARACO 25MG N78210 001 Jul 10, 2007 Jun NEWA

>A> AB 50MG N78210 002 Jul 10, 2007 Jun NEWA

>A> AB 100MG N78210 003 Jul 10, 2007 Jun NEWA

>D> AB TEVA 50MG N73315 001 May 28, 1993 Jun DISC

>A> @ 50MG N73315 001 May 28, 1993 Jun DISC

>D> AB 100MG N73316 001 May 28, 1993 Jun DISC

>A> @ 100MG N73316 001 May 28, 1993 Jun DISC

>D> AB TEVA PHARMS 50MG N74120 001 Feb 24, 1995 Jun DISC

>A> @ 50MG N74120 001 Feb 24, 1995 Jun DISC

TABLET; ORAL

ATENOLOL

>D>	AB	TEVA PHARMS	100MG	N74120 002	Feb 24, 1995	Jun	DISC
>A>	@		100MG	N74120 002	Feb 24, 1995	Jun	DISC

ATRACURIUM BESYLATE

INJECTABLE; INJECTION

ATRACURIUM BESYLATE

>D>	AP	BAXTER HLTHCARE	10MG/ML	N74824 001	Sep 30, 1997	Jun	DISC
>A>	@		10MG/ML	N74824 001	Sep 30, 1997	Jun	DISC
>D>	AP	HOSPIRA	10MG/ML	N74632 001	Dec 23, 1996	Jun	DISC
>A>	@		10MG/ML	N74632 001	Dec 23, 1996	Jun	DISC
ATRACURIUM BESYLATE PRESERVATIVE FREE							
>D>	AP	HOSPIRA	10MG/ML	N74633 001	Dec 23, 1996	Jun	DISC
>A>	@		10MG/ML	N74633 001	Dec 23, 1996	Jun	DISC
>D>	AP		10MG/ML	N74639 001	Mar 25, 1997	Jun	DISC
>A>	@		10MG/ML	N74639 001	Mar 25, 1997	Jun	DISC

AZACITIDINE

INJECTABLE; IV-SC

VIDAZA

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

AZATHIOPRINE

TABLET; ORAL

AZATHIOPRINE

AB	ZYDUS PHARMS USA	50MG	N77621 001	Mar 15, 2007	Mar	NEWA
----	------------------	------	------------	--------------	-----	------

AZITHROMYCIN

INJECTABLE; INJECTION

AZITHROMYCIN

AP	PLIVA HRVATSKA DOO	EQ 500MG BASE/VIAL	N65265 001	Jan 18, 2007	Jan	NEWA	
AP	+	TEVA PARENTERAL	EQ 500MG BASE/VIAL	N50809 001	Dec 19, 2006	May	CAHN
	+		EQ 2.5GM BASE/VIAL	N50809 002	Dec 19, 2006	May	CAHN

SOLUTION/DROPS; OPHTHALMIC

AZASITE

+	INSITE VISION	1%	N50810 001	Apr 27, 2007	Apr	NEWA
+	INSPIRE	1%	N50810 001	Apr 27, 2007	May	CAHN

TABLET; ORAL

AZITHROMYCIN

AB	MYLAN	EQ 250MG BASE	N65365 001	May 30, 2007	May	NEWA
AB		EQ 500MG BASE	N65366 001	May 30, 2007	May	NEWA

BACITRACIN ZINC; HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

CORTISPORIN

@	MONARCH PHARMS	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	N50416 002		Feb	DISC	
NEOMYCIN AND POLYMYXIN B SULFATES, BACITRACIN ZINC AND HYDROCORTISONE							
+	BAUSCH AND LOMB	400 UNITS/GM;1%;EQ 3.5MG BASE/GM;10,000 UNITS/GM	N64068 001	Oct 30, 1995	Feb	CTEC	

BACITRACIN ZINC; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

NEOSPORIN

@	MONARCH PHARMS	400 UNITS/GM;EQ 3.5MG BASE/GM;10,000 UNITS/GM	N50417 001		Feb	DISC
---	----------------	--	------------	--	-----	------

BACITRACIN ZINC; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

POLYSPORIN

@	MONARCH PHARMS	500 UNITS/GM;10,000 UNITS/GM	N61229 001			Feb	DISC
---	----------------	------------------------------	------------	--	--	-----	------

BACLOFEN

TABLET; ORAL

BACLOFEN

>A>	AB	LANNETT	10MG	N78220 001	Jul 06, 2007	Jun	NEWA
>D>	AB	WATSON LABS	10MG	N73092 001	Jan 28, 1994	Jun	DISC
>A>		@	10MG	N73092 001	Jan 28, 1994	Jun	DISC

BECLOMETHASONE DIPROPIONATE

AEROSOL, METERED; INHALATION

QVAR 40

+	IVAX RES	0.04MG/INH	N20911 002	Sep 15, 2000	Mar	CAHN
---	----------	------------	------------	--------------	-----	------

QVAR 80

+	IVAX RES	0.08MG/INH	N20911 001	Sep 15, 2000	Mar	CAHN
---	----------	------------	------------	--------------	-----	------

BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL

CORZIDE

AB		KING PHARMS	5MG;40MG	N18647 001	May 25, 1983	Mar	CFTG
----	--	-------------	----------	------------	--------------	-----	------

AB	+		5MG;80MG	N18647 002	May 25, 1983	Mar	CFTG
----	---	--	----------	------------	--------------	-----	------

NADOLOL AND BENDROFLUMETHIAZIDE

AB		IMPAX LABS	5MG;40MG	N77833 001	Mar 30, 2007	Mar	NEWA
----	--	------------	----------	------------	--------------	-----	------

AB			5MG;80MG	N77833 002	Mar 30, 2007	Mar	NEWA
----	--	--	----------	------------	--------------	-----	------

BENZONATATE

CAPSULE; ORAL

BENZONATATE

AA		THE PHARMA NETWORK	100MG	N40627 001	Mar 30, 2007	Mar	NEWA
----	--	--------------------	-------	------------	--------------	-----	------

>A>	AA	ZYDUS PHARMS USA	100MG	N40597 001	Jun 08, 2007	Jun	NEWA
-----	----	------------------	-------	------------	--------------	-----	------

>A>	AA		200MG	N40597 002	Jun 08, 2007	Jun	NEWA
-----	----	--	-------	------------	--------------	-----	------

TESSALON

>D>		+	FOREST LABS	200MG	N11210 003	Jun 25, 1999	Jun	CFTG
-----	--	---	-------------	-------	------------	--------------	-----	------

>A>	AA	+		200MG	N11210 003	Jun 25, 1999	Jun	CFTG
-----	----	---	--	-------	------------	--------------	-----	------

BENZOYL PEROXIDE; CLINDAMYCIN PHOSPHATE

GEL; TOPICAL

BENZACLIN

@	SANOFI AVENTIS US	5%;EQ 1% BASE	N50756 002	Apr 20, 2007	Apr	DISC
---	-------------------	---------------	------------	--------------	-----	------

BENZPHETAMINE HYDROCHLORIDE

TABLET; ORAL

BENZPHETAMINE HYDROCHLORIDE

AA		TEDOR PHARM	50MG	N40747 001	Mar 30, 2007	Mar	NEWA
----	--	-------------	------	------------	--------------	-----	------

AA		TYCO HLTHCARE	50MG	N40773 001	Apr 25, 2007	Apr	NEWA
----	--	---------------	------	------------	--------------	-----	------

BETAINE HYDROCHLORIDE

FOR SOLUTION; ORAL

CYSTADANE

+	RARE DIS	1GM/SCOOPFUL	N20576 001	Oct 25, 1996	Apr	CAHN
---	----------	--------------	------------	--------------	-----	------

BETAMETHASONE DIPROPIONATE

CREAM; TOPICAL

BETAMETHASONE DIPROPIONATE

>D>	AB	TARO	EQ 0.05% BASE	N71143	001	Jun 17, 1987	Jun	DISC
>A>		@	EQ 0.05% BASE	N71143	001	Jun 17, 1987	Jun	DISC
>D>	AB	TEVA	EQ 0.05% BASE	N71476	001	Aug 10, 1987	Jun	DISC
>A>		@	EQ 0.05% BASE	N71476	001	Aug 10, 1987	Jun	DISC

LOTION; TOPICAL

BETAMETHASONE DIPROPIONATE

>D>	AB	TARO	EQ 0.05% BASE	N72276	001	Aug 24, 1988	Jun	DISC
>A>		@	EQ 0.05% BASE	N72276	001	Aug 24, 1988	Jun	DISC
>D>	AB		EQ 0.05% BASE	N74272	001	Sep 30, 1994	Jun	DISC
>A>		@	EQ 0.05% BASE	N74272	001	Sep 30, 1994	Jun	DISC
>D>	AB	TEVA PHARMS	EQ 0.05% BASE	N71882	001	Jun 06, 1988	Jun	DISC
>A>		@	EQ 0.05% BASE	N71882	001	Jun 06, 1988	Jun	DISC

LOTION, AUGMENTED; TOPICAL

BETAMETHASONE DIPROPIONATE

	AB	ALTANA	EQ 0.05% BASE	N77111	001	May 21, 2007	May	NEWA
	AB	TARO	EQ 0.05% BASE	N77477	001	May 21, 2007	May	NEWA

DIPROLENE

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

OINTMENT; TOPICAL

BETAMETHASONE DIPROPIONATE

>D>	AB	TEVA	EQ 0.05% BASE	N71477	001	Aug 10, 1987	Jun	DISC
>A>		@	EQ 0.05% BASE	N71477	001	Aug 10, 1987	Jun	DISC

BETAMETHASONE VALERATE

LOTION; TOPICAL

BETAMETHASONE VALERATE

>D>	AB	TEVA PHARMS	EQ 0.1% BASE	N71883	001	Apr 22, 1988	Jun	DISC
>A>		@	EQ 0.1% BASE	N71883	001	Apr 22, 1988	Jun	DISC

OINTMENT; TOPICAL

BETA-VAL

>D>	AB	TEVA	EQ 0.1% BASE	N70069	001	Dec 19, 1985	Jun	DISC
>A>		@	EQ 0.1% BASE	N70069	001	Dec 19, 1985	Jun	DISC

BETHANECHOL CHLORIDE

TABLET; ORAL

BETHANECHOL CHLORIDE

>D>	AA	ACTAVIS TOTOWA	50MG	N40551	001	Oct 28, 2004	Jun	DISC
>A>		@	50MG	N40551	001	Oct 28, 2004	Jun	DISC

BISMUTH SUBCITRATE POTASSIUM; METRONIDAZOLE; TETRACYCLINE

CAPSULE; ORAL

PYLERA

		+ AXCAN SCANDIPHARM	140MG;125MG;125MG	N50786	001	Sep 28, 2006	May	CAIN
--	--	---------------------	-------------------	--------	-----	--------------	-----	------

BISOPROLOL FUMARATE; HYDROCHLOROTHIAZIDE

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

>D>	AB	TEVA	2.5MG;6.25MG	N75686	001	Jan 19, 2001	Jun	DISC
>A>		@	2.5MG;6.25MG	N75686	001	Jan 19, 2001	Jun	DISC
>D>	AB		5MG;6.25MG	N75686	002	Jan 19, 2001	Jun	DISC
>A>		@	5MG;6.25MG	N75686	002	Jan 19, 2001	Jun	DISC

TABLET; ORAL

BISOPROLOL FUMARATE AND HYDROCHLOROTHIAZIDE

>D>	AB	TEVA	10MG;6.25MG	N75686 003	Jan 19, 2001	Jun	DISC
>A>		@	10MG;6.25MG	N75686 003	Jan 19, 2001	Jun	DISC

BUDESONIDE

POWDER, METERED; INHALATION

PULMICORT FLEXHALER

		ASTRAZENECA	0.08MG/INH	N21949 001	Jul 12, 2006	Feb	CTNA
	+		0.16MG/INH	N21949 002	Jul 12, 2006	Feb	CTNA

BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE

SPRAY, METERED; INHALATION

SYMBICORT

	+	ASTRAZENECA	0.08MG/INH;0.045MG/INH	N21929 001	Jul 21, 2006	Jan	CAIN
	+		0.16MG/INH;0.045MG/INH	N21929 002	Jul 21, 2006	Jan	CAIN

BUMETANIDE

INJECTABLE; INJECTION

BUMETANIDE

>D>	AP	HOSPIRA	0.25MG/ML	N74160 001	Oct 30, 1997	Jun	DISC
>A>		@	0.25MG/ML	N74160 001	Oct 30, 1997	Jun	DISC

BUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

SENSORCAINE

AP		ABRAXIS BIOSCIENCE	0.25%	N18304 001		May	CAHN
AP			0.25%	N70552 001	May 21, 1986	May	CAHN
AP			0.5%	N18304 002		May	CAHN
AP			0.5%	N70553 001	May 21, 1986	May	CAHN
AP			0.75%	N18304 003		May	CAHN
AP			0.75%	N70554 001	May 21, 1986	May	CAHN

INJECTABLE; SPINAL

SENSORCAINE

AP		ABRAXIS BIOSCIENCE	0.75%	N71202 001	Apr 15, 1987	May	CAHN
----	--	--------------------	-------	------------	--------------	-----	------

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE

INJECTABLE; INJECTION

BUPIVACAINE HYDROCHLORIDE AND EPINEPHRINE

>D>		HOSPIRA	0.25%;0.005MG/ML	N71166 001	Jun 16, 1988	Jun	DISC
>A>		@	0.25%;0.005MG/ML	N71166 001	Jun 16, 1988	Jun	DISC

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE

INJECTABLE; INJECTION

SENSORCAINE

AP		ABRAXIS BIOSCIENCE	0.25%;0.0091MG/ML	N70966 001	Oct 13, 1987	May	CAHN
AP			0.25%;0.0091MG/ML	N70967 001	Oct 13, 1987	May	CAHN
AP			0.5%;0.0091MG/ML	N18304 004	Sep 02, 1983	May	CAHN
AP			0.5%;0.0091MG/ML	N70968 001	Oct 13, 1987	May	CAHN
AP			0.75%;0.0091MG/ML	N18304 005	Sep 02, 1983	May	CAHN

BUPRENORPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

BUPRENORPHINE HYDROCHLORIDE

AP		PHARMAFORCE	EQ 0.3MG BASE/ML	N78331 001	Mar 27, 2007	Mar	NEWA
----	--	-------------	------------------	------------	--------------	-----	------

BUPROPION HYDROCHLORIDE

TABLET; ORAL

BUPROPION HYDROCHLORIDE

>D>	AB	SANDOZ	75MG	N75613 002	Oct 10, 2000	Jun	DISC
>A>		@	75MG	N75613 002	Oct 10, 2000	Jun	DISC
>D>	AB		100MG	N75613 001	Oct 10, 2000	Jun	DISC
>A>		@	100MG	N75613 001	Oct 10, 2000	Jun	DISC

TABLET, EXTENDED RELEASE; ORAL

BUPROPION HYDROCHLORIDE

>D>	AB1	SANDOZ	100MG	N76845 001	Jul 14, 2005	Jun	DISC
>A>		@	100MG	N76845 001	Jul 14, 2005	Jun	DISC
>D>	AB1		150MG	N76845 002	Jul 14, 2005	Jun	DISC
>A>		@	150MG	N76845 002	Jul 14, 2005	Jun	DISC
>A>	AB3	WATSON LABS	300MG	N77715 002	Jun 13, 2007	Jun	NEWA

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

>D>	AP	HOSPIRA	1MG/ML	N75342 001	Nov 04, 1999	Jun	DISC
>A>		@	1MG/ML	N75342 001	Nov 04, 1999	Jun	DISC
>D>	AP		1MG/ML	N75559 001	Mar 20, 2000	Jun	DISC
>A>		@	1MG/ML	N75559 001	Mar 20, 2000	Jun	DISC
>D>	AP		2MG/ML	N75342 002	Nov 04, 1999	Jun	DISC
>A>		@	2MG/ML	N75342 002	Nov 04, 1999	Jun	DISC
>D>	AP		2MG/ML	N75559 002	Mar 20, 2000	Jun	DISC
>A>		@	2MG/ML	N75559 002	Mar 20, 2000	Jun	DISC

CABERGOLINE

TABLET; ORAL

CABERGOLINE

>A>	AB	BARR	0.5MG	N77843 001	Jul 03, 2007	Jun	NEWA
	AB	IVAX PHARMS INC	0.5MG	N77750 001	Mar 07, 2007	Feb	NEWA

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS

CAFFEINE CITRATE

AP		LUITPOLD	EQ 30MG BASE/3ML (EQ 10MG BASE/ML)	N77906 001	May 15, 2007	May	NEWA
----	--	----------	------------------------------------	------------	--------------	-----	------

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

>D>	AB	IVAX PHARMS	12.5MG	N74590 004	Aug 30, 1996	Jun	DISC
>A>		@	12.5MG	N74590 004	Aug 30, 1996	Jun	DISC
>D>	AB		25MG	N74590 002	Aug 30, 1996	Jun	DISC
>A>		@	25MG	N74590 002	Aug 30, 1996	Jun	DISC
>D>	AB		50MG	N74590 001	Aug 30, 1996	Jun	DISC
>A>		@	50MG	N74590 001	Aug 30, 1996	Jun	DISC
>D>	AB		100MG	N74590 003	Aug 30, 1996	Jun	DISC
>A>		@	100MG	N74590 003	Aug 30, 1996	Jun	DISC
>D>	AB	TEVA	12.5MG	N74433 001	Feb 13, 1996	Jun	DISC
>A>		@	12.5MG	N74433 001	Feb 13, 1996	Jun	DISC
>D>	AB		25MG	N74433 002	Feb 13, 1996	Jun	DISC
>A>		@	25MG	N74433 002	Feb 13, 1996	Jun	DISC
>D>	AB		50MG	N74433 003	Feb 13, 1996	Jun	DISC

TABLET; ORALCAPTOPRIL

>A>	@	TEVA	50MG	N74433 003	Feb 13, 1996	Jun	DISC
>D>	AB		100MG	N74433 004	Feb 13, 1996	Jun	DISC
>A>	@		100MG	N74433 004	Feb 13, 1996	Jun	DISC
>D>	AB	TEVA PHARMS	12.5MG	N74462 001	Feb 13, 1996	Jun	DISC
>A>	@		12.5MG	N74462 001	Feb 13, 1996	Jun	DISC
>D>	AB		25MG	N74462 002	Feb 13, 1996	Jun	DISC
>A>	@		25MG	N74462 002	Feb 13, 1996	Jun	DISC
>D>	AB		50MG	N74462 003	Feb 13, 1996	Jun	DISC
>A>	@		50MG	N74462 003	Feb 13, 1996	Jun	DISC
>D>	AB		100MG	N74462 004	Feb 13, 1996	Jun	DISC
>A>	@		100MG	N74462 004	Feb 13, 1996	Jun	DISC

CARBACHOLSOLUTION; INTRAOCULAR

>D>		CARBASTAT					
>D>	AT	NOVARTIS	0.01%	N73677 001	Apr 28, 1995	Jun	DISC
>A>	@		0.01%	N73677 001	Apr 28, 1995	Jun	DISC
		MIOSTAT					
>D>	AT	+ ALCON	0.01%	N16968 001		Jun	CTEC
>A>	+		0.01%	N16968 001		Jun	CTEC

CARBIDOPA; ENTACAPONE; LEVODOPATABLET; ORALSTALEVO 50

	+	ORION PHARMA INC	12.5MG;200MG;50MG	N21485 001	Jun 11, 2003	Apr	CRLD
--	---	------------------	-------------------	------------	--------------	-----	------

CARBOPLATININJECTABLE; IV (INFUSION)CARBOPLATIN

AP		GENERAMEDIX	EQ 50MG/5ML (10MG/ML)	N77998 001	Apr 24, 2007	Apr	NEWA
AP			EQ 150MG/ML (10MG/ML)	N77998 002	Apr 24, 2007	Apr	NEWA
AP			EQ 450MG/45ML (10MG/ML)	N77998 003	Apr 24, 2007	Apr	NEWA
AP		SICOR PHARMS	EQ 50MG/5ML (10MG/ML)	N77389 001	Mar 30, 2007	Mar	NEWA
AP			EQ 150MG/15ML (10MG/ML)	N77389 002	Mar 30, 2007	Mar	NEWA
AP			EQ 450MG/45ML (10MG/ML)	N77389 003	Mar 30, 2007	Mar	NEWA
AP		WATSON LABS	EQ 50MG/5ML (10MG/ML)	N77861 001	Jan 18, 2007	Jan	NEWA
AP			EQ 150MG/15ML (10MG/ML)	N77861 002	Jan 18, 2007	Jan	NEWA
AP			EQ 450MG/45ML (10MG/ML)	N77861 003	Jan 18, 2007	Jan	NEWA
AP			EQ 600MG/60ML (10MG/ML)	N77861 004	Jan 18, 2007	Jan	NEWA

CARISOPRODOLTABLET; ORALCARISOPRODOL

AA		SUN PHARM INDS LTD	350MG	N40755 001	Feb 27, 2007	Feb	NEWA
----	--	--------------------	-------	------------	--------------	-----	------

CEFACLORCAPSULE; ORALCEFACLOR

	@	CEPH INTL	EQ 250MG BASE	N62205 001		Apr	DISC
	@		EQ 500MG BASE	N62205 002		Apr	DISC
AB		HIKMA	EQ 250MG BASE	N65350 001	Apr 03, 2007	Mar	NEWA
AB			EQ 500MG BASE	N65350 002	Apr 03, 2007	Mar	NEWA
>D>	AB	IVAX PHARMS	EQ 250MG BASE	N64061 001	Apr 27, 1995	Jun	DISC

CAPSULE; ORAL

CEFACTOR

>A>	@	IVAX PHARMS	EQ 250MG BASE	N64061 001	Apr 27, 1995	Jun	DISC
>D>	AB		EQ 500MG BASE	N64061 002	Apr 27, 1995	Jun	DISC
>A>	@		EQ 500MG BASE	N64061 002	Apr 27, 1995	Jun	DISC
	@	TEVA	EQ 250MG BASE	N64145 001	Jun 24, 1996	May	DISC
	@		EQ 500MG BASE	N64145 002	Jun 24, 1996	May	DISC

FOR SUSPENSION; ORAL

CEFACTOR

>D>	AB	IVAX PHARMS	EQ 375MG BASE/5ML	N64070 001	Apr 28, 1995	Jun	DISC
>A>	@		EQ 375MG BASE/5ML	N64070 001	Apr 28, 1995	Jun	DISC

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL

CEFADROXIL

AB	AUROBINDO PHARMA	EQ 500MG BASE	N65352 001	Jan 25, 2007	May	CPOT
AB		500MG	N65352 001	Jan 25, 2007	Jan	NEWA
AB	LUPIN	EQ 500MG BASE	N65392 001	May 29, 2007	May	NEWA

CEFAZOLIN SODIUM

INJECTABLE; INJECTION

CEFAZOLIN SODIUM

AP	SANDOZ	EQ 1GM BASE/VIAL	N65345 001	May 09, 2007	May	NEWA
----	--------	------------------	------------	--------------	-----	------

CEFDINIR

CAPSULE; ORAL

CEFDINIR

AB	SANDOZ	300MG	N65330 001	Apr 06, 2007	Mar	NEWA
AB	TEVA PHARMS	300MG	N65368 001	May 09, 2007	May	NEWA

FOR SUSPENSION; ORAL

CEFDINIR

AB	LUPIN	250MG/5ML	N65259 002	May 07, 2007	Apr	NEWA
AB	SANDOZ	125MG/5ML	N65337 001	Apr 06, 2007	Mar	NEWA
AB		250MG/5ML	N65337 002	Apr 06, 2007	Mar	NEWA
AB	TEVA PHARMS	125MG/5ML	N65332 001	May 04, 2007	Apr	NEWA
AB		250MG/5ML	N65332 002	May 04, 2007	Apr	NEWA
	OMNICEF					
AB	+ ABBOTT	250MG/5ML	N50749 002	Jul 29, 2004	Mar	CFTG

CEFEPIME HYDROCHLORIDE (ARGININE FORMULATION)

INJECTABLE; INJECTION

CEFEPIME HYDROCHLORIDE

>A>	AP	ORCHID HLTHCARE	EQ 500MG BASE/VIAL	N65369 001	Jun 18, 2007	Jun	NEWA
>A>	AP		EQ 1GM BASE/VIAL	N65369 002	Jun 18, 2007	Jun	NEWA
>A>	AP		EQ 2GM BASE/VIAL	N65369 003	Jun 18, 2007	Jun	NEWA
		MAXIPIME					
>D>	+	BRISTOL MYERS SQUIBB	EQ 500MG BASE/VIAL	N50679 001	Jan 18, 1996	Jun	CFTG
>A>	AP		EQ 500MG BASE/VIAL	N50679 001	Jan 18, 1996	Jun	CFTG
>D>	+		EQ 1GM BASE/VIAL	N50679 002	Jan 18, 1996	Jun	CFTG
>A>	AP		EQ 1GM BASE/VIAL	N50679 002	Jan 18, 1996	Jun	CFTG
>D>	+		EQ 2GM BASE/VIAL	N50679 003	Jan 18, 1996	Jun	CFTG
>A>	AP		EQ 2GM BASE/VIAL	N50679 003	Jan 18, 1996	Jun	CFTG

CEFIXIME

SUSPENSION; ORAL

CEFIXIME

	LUPIN PHARMS	200MG/5ML	N65355 001	Apr 10, 2007	Mar	NEWA
	SUPRAX					
	LUPIN PHARMS	100MG/5ML	N65129 001	Feb 23, 2004	Apr	CRLD
+		200MG/5ML	N65355 001	Apr 10, 2007	Apr	CRLD

CEFOXITIN SODIUM

INJECTABLE; INJECTION

>D>	MEFOXIN					
>D>	AP	MERCK	EQ 1GM BASE/VIAL	N62757 001	Jan 08, 1987	Jun DISC
>A>		@	EQ 1GM BASE/VIAL	N62757 001	Jan 08, 1987	Jun DISC
>D>	AP		EQ 2GM BASE/VIAL	N62757 002	Jan 08, 1987	Jun DISC
>A>		@	EQ 2GM BASE/VIAL	N62757 002	Jan 08, 1987	Jun DISC

CEFPODOXIME PROXETIL

FOR SUSPENSION; ORAL

CEFPODOXIME PROXETIL

>A>	AB	AUROBINDO PHARMA	EQ 50MG BASE/5ML	N65409 001	Jun 08, 2007	Jun NEWA
>A>	AB		EQ 100MG BASE/5ML	N65409 002	Jun 08, 2007	Jun NEWA

TABLET; ORAL

CEFPODOXIME PROXETIL

>A>	AB	AUROBINDO PHARMA	EQ 100MG BASE	N65370 001	Jun 11, 2007	Jun NEWA
>A>	AB		EQ 200MG BASE	N65370 002	Jun 11, 2007	Jun NEWA

CEFPROZIL

FOR SUSPENSION; ORAL

CEFPROZIL

AB	AUROBINDO PHARMA	125MG/5ML	N65381 001	Jan 30, 2007	Jan	NEWA
AB		250MG/5ML	N65381 002	Jan 30, 2007	Jan	NEWA

TABLET; ORAL

CEFPROZIL

AB	AUROBINDO PHARMA LTD	250MG	N65340 001	May 24, 2007	May	NEWA
AB		500MG	N65340 002	May 24, 2007	May	NEWA
>A>	AB	WOCKHARDT	250MG	N65428 001	Jun 14, 2007	Jun NEWA
>A>	AB		500MG	N65428 002	Jun 14, 2007	Jun NEWA

CEFTIZOXIME SODIUM

INJECTABLE; INJECTION

CEFIZOX

	@ ASTELLAS	EQ 1GM BASE/VIAL	N50560 002	Sep 15, 1983	Feb	DISC
+		EQ 1GM BASE/VIAL	N63294 002	Mar 31, 1994	Feb	CRLD

CEFTRIAZONE SODIUM

INJECTABLE; IM-IV

CEFTRIAZONE

AP	CEPHAZONE PHARMA	EQ 250MG BASE/VIAL	N65294 001	Mar 26, 2007	Mar	NEWA
AP		EQ 500MG BASE/VIAL	N65294 002	Mar 26, 2007	Mar	NEWA
AP		EQ 1GM BASE/VIAL	N65294 003	Mar 26, 2007	Mar	NEWA
AP		EQ 2GM BASE/VIAL	N65294 004	Mar 26, 2007	Mar	NEWA
AP	HANFORD GC	EQ 1GM BASE/VIAL	N65268 001	Feb 28, 2007	Feb	NEWA
AP		EQ 2GM BASE/VIAL	N65268 002	Feb 28, 2007	Feb	NEWA
AP	SICOR PHARMS	EQ 250MG BASE/VIAL	N65227 001	Mar 15, 2007	Mar	NEWA

INJECTABLE; IM-IV

CEFTRIAZONE

AP	SICOR PHARMS	EQ 500MG BASE/VIAL	N65227 002	Mar 15, 2007	Mar	NEWA
AP		EQ 1GM BASE/VIAL	N65227 003	Mar 15, 2007	Mar	NEWA
AP		EQ 2GM BASE/VIAL	N65227 004	Mar 15, 2007	Mar	NEWA
AP	TEVA	EQ 1GM BASE/VIAL	N65262 001	Jun 29, 2006	Apr	CTNA
AP		EQ 2GM BASE/VIAL	N65262 002	Jun 29, 2006	Apr	CTNA
AP	WOCKHARDT	EQ 250MG BASE/VIAL	N65391 001	Apr 12, 2007	Apr	NEWA
AP		EQ 500MG BASE/VIAL	N65391 002	Apr 12, 2007	Apr	NEWA
AP		EQ 2GM BASE/VIAL	N65391 003	Apr 12, 2007	Apr	NEWA

INJECTABLE; INJECTION

CEFTRIAZONE

AP	HANFORD GC	EQ 10GM BASE/VIAL	N65269 001	Feb 28, 2007	Feb	NEWA
----	------------	-------------------	------------	--------------	-----	------

CELECOXIB

CAPSULE; ORAL

CELEBREX

GD SEARLE

50MG

N20998 004 Dec 15, 2006 Jan NEWA

CEPHALEXIN

CAPSULE; ORAL

CEPHALEXIN

>D>	AB	TEVA	EQ 250MG BASE	N62760 001	Apr 24, 1987	Jun	DISC
>A>		@	EQ 250MG BASE	N62760 001	Apr 24, 1987	Jun	DISC
>D>	AB		EQ 500MG BASE	N62761 001	Apr 24, 1987	Jun	DISC
>A>		@	EQ 500MG BASE	N62761 001	Apr 24, 1987	Jun	DISC

FOR SUSPENSION; ORAL

CEPHALEXIN

@ CEPH INTL

EQ 100MG BASE/ML

N62117 001 May DISC

@ TEVA

EQ 125MG BASE/5ML

N62767 001 Jun 16, 1987 May DISC

@

EQ 250MG BASE/5ML

N62768 001 Jun 16, 1987 May DISC

CEPHRADINE

CAPSULE; ORAL

CEPHRADINE

@ TEVA

250MG

N62683 001 Jan 09, 1987 May DISC

@

500MG

N62683 002 Jan 09, 1987 May DISC

FOR SUSPENSION; ORAL

ANSPOR

@ GLAXOSMITHKLINE

125MG/5ML

N61866 001 May DISC

@

250MG/5ML

N61866 002 May DISC

CEPHRADINE

@ TEVA

125MG/5ML

N62693 001 Jan 09, 1987 May DISC

@

250MG/5ML

N62693 002 Jan 09, 1987 May DISC

VELOSEF '125'

@ APOTHECON

125MG/5ML

N61763 001 May DISC

VELOSEF '250'

@ APOTHECON

250MG/5ML

N61763 002 May DISC

CETRORELIX

INJECTABLE; INJECTION

CETROTIDE

>A>	+	EMD SERONO	EQ 0.25MG BASE/ML	N21197 001	Aug 11, 2000	Jun	CAHN
>A>	+		EQ 3MG BASE/ML	N21197 002	Aug 11, 2000	Jun	CAHN
>D>	+	SERONO INC	EQ 0.25MG BASE/ML	N21197 001	Aug 11, 2000	Jun	CAHN
>D>	+		EQ 3MG BASE/ML	N21197 002	Aug 11, 2000	Jun	CAHN

CHLORAMPHENICOL SODIUM SUCCINATE

INJECTABLE; INJECTION

CHLORAMPHENICOL SODIUM SUCCINATE

+	ABRAXIS PHARM	EQ 1GM BASE/VIAL	N62365 001	Aug 25, 1982	Feb	CRLD
	CHLOROMYCETIN					
	@ PARKEDALE	EQ 1GM BASE/VIAL	N50155 001		Feb	DISC

CHLOROTHIAZIDE

SUSPENSION; ORAL

DIURIL

+	SALIX PHARMS	250MG/5ML	N11870 001		Feb	CAHN
---	--------------	-----------	------------	--	-----	------

CHLORPHENIRAMINE POLISTIREX; CODEINE POLISTIREX

SUSPENSION, EXTENDED RELEASE; ORAL

CODEPREX

	@ UCB INC	EQ 4MG MALEATE/5ML;EQ 20MG BASE/5ML	N21369 001	Jun 21, 2004	Mar	DISC
--	-----------	--	------------	--------------	-----	------

CHOLESTYRAMINE

POWDER; ORAL

CHOLESTYRAMINE

>D>	AB	TEVA	EQ 4GM RESIN/PACKET	N74347 001	May 28, 1998	Jun	DISC
>D>	AB		EQ 4GM RESIN/SCOOPFUL	N74347 002	May 28, 1998	Jun	DISC
>A>		@	EQ 4GM RESIN/SCOOPFUL	N74347 002	May 28, 1998	Jun	DISC
>A>		@	EQ 4GM RESIN/PACKET	N74347 001	May 28, 1998	Jun	DISC
>D>	AB	TEVA PHARMS	EQ 4GM RESIN/SCOOPFUL	N74554 002	Oct 02, 1996	Jun	DISC
>D>	AB		EQ 4GM RESIN/PACKET	N74554 001	Oct 02, 1996	Jun	DISC
>A>		@	EQ 4GM RESIN/PACKET	N74554 001	Oct 02, 1996	Jun	DISC
>A>		@	EQ 4GM RESIN/SCOOPFUL	N74554 002	Oct 02, 1996	Jun	DISC

CHOLESTYRAMINE LIGHT

>D>	AB	TEVA	EQ 4GM RESIN/PACKET	N74348 001	May 28, 1998	Jun	DISC
>D>	AB		EQ 4GM RESIN/SCOOPFUL	N74348 002	May 28, 1998	Jun	DISC
>A>		@	EQ 4GM RESIN/PACKET	N74348 001	May 28, 1998	Jun	DISC
>A>		@	EQ 4GM RESIN/SCOOPFUL	N74348 002	May 28, 1998	Jun	DISC
>D>	AB	TEVA PHARMS	EQ 4GM RESIN/PACKET	N74555 001	Sep 30, 1998	Jun	DISC
>D>	AB		EQ 4GM RESIN/SCOOPFUL	N74555 002	Sep 30, 1998	Jun	DISC
>A>		@	EQ 4GM RESIN/SCOOPFUL	N74555 002	Sep 30, 1998	Jun	DISC
>A>		@	EQ 4GM RESIN/PACKET	N74555 001	Sep 30, 1998	Jun	DISC

CHORIOGONADOTROPIN ALFA

INJECTABLE; INJECTION

OVIDREL

>A>		@ EMD SERONO	0.25MG/VIAL	N21149 001	Sep 20, 2000	Jun	CAHN
>D>		@ SERONO INC	0.25MG/VIAL	N21149 001	Sep 20, 2000	Jun	CAHN

INJECTABLE; SUBCUTANEOUS

OVIDREL

>A>	+	EMD SERONO	EQ 0.25MG /0.5ML	N21149 002	Oct 06, 2003	Jun	CAHN
>D>	+	SERONO INC	EQ 0.25MG /0.5ML	N21149 002	Oct 06, 2003	Jun	CAHN

CICLOPIROX

>D>		SOLUTION; TOPICAL					
>D>		PENLAC					
>D>	+	SANOFI AVENTIS US	8%	N21022 001	Dec 17, 1999	Jun	DISC
>A>		@	8%	N21022 001	Dec 17, 1999	Jun	DISC

CIMETIDINE

TABLET; ORAL

CIMETIDINE

>D>	AB	IVAX PHARMS	800MG	N74402 001	May 30, 1995	Jun	DISC
>A>		@	800MG	N74402 001	May 30, 1995	Jun	DISC
>D>	AB	TEVA	200MG	N74365 001	Feb 28, 1995	Jun	DISC
>A>		@	200MG	N74365 001	Feb 28, 1995	Jun	DISC
>D>	AB		300MG	N74365 002	Feb 28, 1995	Jun	DISC
>A>		@	300MG	N74365 002	Feb 28, 1995	Jun	DISC
>D>	AB		400MG	N74365 003	Feb 28, 1995	Jun	DISC
>A>		@	400MG	N74365 003	Feb 28, 1995	Jun	DISC
>D>	AB		800MG	N74365 004	Feb 28, 1995	Jun	DISC
>A>		@	800MG	N74365 004	Feb 28, 1995	Jun	DISC

CIMETIDINE HYDROCHLORIDE

SOLUTION; ORAL

CIMETIDINE HYDROCHLORIDE

>D>	AA	TEVA PHARMS	EQ 300MG BASE/5ML	N74859 001	Jul 09, 1998	Jun	DISC
>A>		@	EQ 300MG BASE/5ML	N74859 001	Jul 09, 1998	Jun	DISC

CIPROFLOXACIN HYDROCHLORIDE

TABLET; ORAL

CIPROFLOXACIN

	AB	AUROBINDO PHARMA	EQ 250MG BASE	N77859 001	Apr 26, 2007	Apr	NEWA
	AB		EQ 500MG BASE	N77859 002	Apr 26, 2007	Apr	NEWA
	AB		EQ 750MG BASE	N77859 003	Apr 26, 2007	Apr	NEWA
>D>	AB	MYLAN	EQ 250MG BASE	N75685 002	Jun 09, 2004	Jun	DISC
>A>		@	EQ 250MG BASE	N75685 002	Jun 09, 2004	Jun	DISC
>D>	AB		EQ 500MG BASE	N75685 003	Jun 09, 2004	Jun	DISC
>A>		@	EQ 500MG BASE	N75685 003	Jun 09, 2004	Jun	DISC
>D>	AB		EQ 750MG BASE	N75685 001	Jun 09, 2004	Jun	DISC
>A>		@	EQ 750MG BASE	N75685 001	Jun 09, 2004	Jun	DISC
			CIPROFLOXACIN HYDROCHLORIDE				
>A>	AB	GENPHARM	EQ 100MG BASE	N75817 001	Jun 25, 2007	Jun	NEWA

CIPROFLOXACIN; CIPROFLOXACIN HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

CIPRO XR

	AB	+	BAYER PHARMS	212.6MG;EQ 287.5MG BASE	N21473 001	Dec 13, 2002	Mar	CFTG
	AB	+		425.2MG;EQ 574.9MG BASE	N21473 002	Aug 28, 2003	Mar	CFTG

CIPROFLOXACIN EXTENDED RELEASE

	AB		DR REDDYS LABS LTD	425.2MG;EQ 574.9MG BASE	N77701 001	Mar 26, 2007	Mar	NEWA
	AB		MYLAN	212.6MG;EQ 287.5MG BASE	N78183 001	Mar 22, 2007	Mar	NEWA
	AB			425.2MG;EQ 574.9MG BASE	N78183 002	Mar 22, 2007	Mar	NEWA

CISPLATIN

INJECTABLE; INJECTION

CISPLATIN

		@	BEDFORD	10MG/VIAL	N74713 001	Nov 14, 2000	Mar	DISC
		@		50MG/VIAL	N74713 002	Nov 14, 2000	Mar	DISC

PLATINOL

	+		BRISTOL MYERS	10MG/VIAL	N18057 001		Mar	CTEC
	+			50MG/VIAL	N18057 002		Mar	CTEC

CITALOPRAM HYDROBROMIDE

CAPSULE; ORAL

CITALOPRAM HYDROBROMIDE

		ALPHAPHARM	EQ 10MG BASE	N77668 001	Feb 28, 2007	Feb	NEWA
			EQ 20MG BASE	N77668 002	Feb 28, 2007	Feb	NEWA
		+	EQ 40MG BASE	N77668 003	Feb 28, 2007	Feb	NEWA

TABLET; ORAL

CITALOPRAM HYDROBROMIDE

		@ ROXANE	EQ 10MG BASE	N77041 001	Nov 23, 2004	May	DISC
		@	EQ 20MG BASE	N77041 002	Nov 23, 2004	May	DISC
>D>	AB	SANDOZ	EQ 10MG BASE	N77040 001	Aug 17, 2005	Jun	DISC
>A>		@	EQ 10MG BASE	N77040 001	Aug 17, 2005	Jun	DISC
>D>	AB		EQ 20MG BASE	N77040 002	Aug 17, 2005	Jun	DISC
>A>		@	EQ 20MG BASE	N77040 002	Aug 17, 2005	Jun	DISC
>D>	AB		EQ 40MG BASE	N77040 003	Aug 17, 2005	Jun	DISC
>A>		@	EQ 40MG BASE	N77040 003	Aug 17, 2005	Jun	DISC
	AB	TORRENT PHARMS	EQ 10MG BASE	N78216 001	Mar 27, 2007	Mar	NEWA
	AB		EQ 20MG BASE	N78216 002	Mar 27, 2007	Mar	NEWA
	AB		EQ 40MG BASE	N78216 003	Mar 27, 2007	Mar	NEWA

CLEMASTINE FUMARATE

SYRUP; ORAL

CLEMASTINE FUMARATE

>D>	AA	TEVA PHARMS	EQ 0.5MG BASE/5ML	N73095 001	Apr 21, 1992	Jun	DISC
>A>		@	EQ 0.5MG BASE/5ML	N73095 001	Apr 21, 1992	Jun	DISC

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLINDAMYCIN HYDROCHLORIDE

>D>	AB	TEVA	EQ 75MG BASE	N63027 001	Sep 20, 1989	Jun	DISC
>A>		@	EQ 75MG BASE	N63027 001	Sep 20, 1989	Jun	DISC

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE

	AP	ABRAXIS PHARM	EQ 150MG BASE/ML	N65346 001	Mar 29, 2007	Mar	NEWA
	AP		EQ 150MG BASE/ML	N65347 001	May 09, 2007	May	NEWA

SWAB; TOPICAL

CLINDETS

>A>	AT	PERRIGO	EQ 1% BASE	N64136 001	Sep 30, 1996	Jun	CAHN
>D>	AT	STIEFEL	EQ 1% BASE	N64136 001	Sep 30, 1996	Jun	CAHN

CLOBETASOL PROPIONATE

AEROSOL, FOAM; TOPICAL

OLUX E

		+ CONNETICS	0.05%	N22013 001	Jan 12, 2007	Jan	NEWA
--	--	-------------	-------	------------	--------------	-----	------

GEL; TOPICAL

CLOBETASOL PROPIONATE

>A>	AB	PERRIGO	0.05%	N75027 001	Oct 31, 1997	Jun	CAHN
>D>	AB	STIEFEL	0.05%	N75027 001	Oct 31, 1997	Jun	CAHN

SOLUTION; TOPICAL

CLOBETASOL PROPIONATE

	AT	ALTANA	0.05%	N75391 001	Feb 08, 1999	Feb	CMFD
--	----	--------	-------	------------	--------------	-----	------

CLOCORTOLONE PIVALATE

CREAM; TOPICAL

CLODERM

+ CORIA	0.1%	N17765 001	May	CAHN
---------	------	------------	-----	------

CLOFIBRATE

CAPSULE; ORAL

CLOFIBRATE

>D> AB	TEVA	500MG	N72600 001	Jul 25, 1991	Jun	DISC
>A>	@	500MG	N72600 001	Jul 25, 1991	Jun	DISC

CLOMIPHENE CITRATE

TABLET; ORAL

SEROPHENE

>A> AB	EMD SERONO	50MG	N18361 001	Mar 22, 1982	Jun	CAHN
>D> AB	SERONO	50MG	N18361 001	Mar 22, 1982	Jun	CAHN

CLOMIPRAMINE HYDROCHLORIDE

CAPSULE; ORAL

CLOMIPRAMINE HYDROCHLORIDE

>D> AB	TEVA	25MG	N74849 001	Apr 04, 1997	Jun	DISC
>A>	@	25MG	N74849 001	Apr 04, 1997	Jun	DISC
>D> AB		25MG	N74958 001	Aug 26, 1997	Jun	DISC
>A>	@	25MG	N74958 001	Aug 26, 1997	Jun	DISC
>D> AB		50MG	N74849 002	Apr 04, 1997	Jun	DISC
>A>	@	50MG	N74849 002	Apr 04, 1997	Jun	DISC
>D> AB		50MG	N74958 002	Aug 26, 1997	Jun	DISC
>A>	@	50MG	N74958 002	Aug 26, 1997	Jun	DISC
>D> AB		75MG	N74849 003	Apr 04, 1997	Jun	DISC
>A>	@	75MG	N74849 003	Apr 04, 1997	Jun	DISC
>D> AB		75MG	N74958 003	Aug 26, 1997	Jun	DISC
>A>	@	75MG	N74958 003	Aug 26, 1997	Jun	DISC
>D> AB	WATSON LABS	25MG	N74600 001	Nov 27, 1996	Jun	DISC
>A>	@	25MG	N74600 001	Nov 27, 1996	Jun	DISC
>D> AB		25MG	N74751 001	Sep 30, 1998	Jun	DISC
>A>	@	25MG	N74751 001	Sep 30, 1998	Jun	DISC
>D> AB		50MG	N74600 002	Nov 27, 1996	Jun	DISC
>A>	@	50MG	N74600 002	Nov 27, 1996	Jun	DISC
>D> AB		50MG	N74751 002	Sep 30, 1998	Jun	DISC
>A>	@	50MG	N74751 002	Sep 30, 1998	Jun	DISC
>D> AB		75MG	N74600 003	Nov 27, 1996	Jun	DISC
>A>	@	75MG	N74600 003	Nov 27, 1996	Jun	DISC
>D> AB		75MG	N74751 003	Sep 30, 1998	Jun	DISC
>A>	@	75MG	N74751 003	Sep 30, 1998	Jun	DISC

CLONAZEPAM

TABLET; ORAL

CLONAZEPAM

>D> AB	SANDOZ	0.5MG	N74925 001	Sep 30, 1997	Jun	DISC
>A>	@	0.5MG	N74925 001	Sep 30, 1997	Jun	DISC
>D> AB		1MG	N74925 002	Sep 30, 1997	Jun	DISC
>A>	@	1MG	N74925 002	Sep 30, 1997	Jun	DISC
>D> AB		2MG	N74925 003	Sep 30, 1997	Jun	DISC
>A>	@	2MG	N74925 003	Sep 30, 1997	Jun	DISC
>D> AB	TEVA	0.5MG	N74920 001	Aug 04, 1998	Jun	DISC

TABLET; ORAL

CLONAZEPAM

>A>	@	TEVA	0.5MG	N74920 001	Aug 04, 1998	Jun	DISC
>D>	AB		1MG	N74920 002	Aug 04, 1998	Jun	DISC
>A>	@		1MG	N74920 002	Aug 04, 1998	Jun	DISC
>D>	AB		2MG	N74920 003	Aug 04, 1998	Jun	DISC
>A>	@		2MG	N74920 003	Aug 04, 1998	Jun	DISC

CLONIDINE HYDROCHLORIDE

TABLET; ORAL

CLONIDINE HYDROCHLORIDE

AB		VINTAGE	0.1MG	N77901 001	Mar 09, 2007	Feb	NEWA
AB			0.2MG	N77901 002	Mar 09, 2007	Feb	NEWA
AB			0.3MG	N77901 003	Mar 09, 2007	Feb	NEWA

CLORAZEPATE DIPOTASSIUM

CAPSULE; ORAL

CLORAZEPATE DIPOTASSIUM

>D>	AB	MYLAN	3.75MG	N71509 001	Oct 19, 1987	Jun	DISC
>A>	@		3.75MG	N71509 001	Oct 19, 1987	Jun	DISC
>D>	AB		7.5MG	N71510 001	Oct 19, 1987	Jun	DISC
>A>	@		7.5MG	N71510 001	Oct 19, 1987	Jun	DISC
>D>	AB	+	15MG	N71511 001	Oct 19, 1987	Jun	DISC
>A>	@		15MG	N71511 001	Oct 19, 1987	Jun	DISC
>D>	AB	SANDOZ	3.75MG	N72219 001	Aug 26, 1988	Jun	CTEC
>A>			3.75MG	N72219 001	Aug 26, 1988	Jun	CTEC
>D>	AB		15MG	N72112 001	Aug 26, 1988	Jun	CRLD
>A>	+		15MG	N72112 001	Aug 26, 1988	Jun	CRLD

CLOXACILLIN SODIUM

CAPSULE; ORAL

CLOXACILLIN SODIUM

>D>	AB	TEVA	EQ 250MG BASE	N62240 001		Jun	DISC
>A>	@		EQ 250MG BASE	N62240 001		Jun	DISC
>D>	AB		EQ 500MG BASE	N62240 002		Jun	DISC
>A>	@		EQ 500MG BASE	N62240 002		Jun	DISC
>D>	AB	GLAXOSMITHKLINE	EQ 250MG BASE	N61806 001		Jun	CTEC
>A>			EQ 250MG BASE	N61806 001		Jun	CTEC
>D>	AB	+	EQ 500MG BASE	N61806 002		Jun	CTEC
>A>	+		EQ 500MG BASE	N61806 002		Jun	CTEC
>D>		FOR SOLUTION; ORAL					
>D>		CLOXACILLIN SODIUM					
>D>	+	TEVA	EQ 125MG BASE/5ML	N62268 001		Jun	DISC
>A>	@		EQ 125MG BASE/5ML	N62268 001		Jun	DISC

CLOZAPINE

TABLET, ORALLY DISINTEGRATING; ORAL

FAZACLO ODT

		AVANIR PHARMS	12.5MG	N21590 004	May 30, 2007	May	NEWA
--	--	---------------	--------	------------	--------------	-----	------

CORTISONE ACETATE

INJECTABLE; INJECTION

CORTISONE ACETATE

	@	PHARMACIA AND UPJOHN	25MG/ML	N08126 002		Apr	DISC
--	---	----------------------	---------	------------	--	-----	------

TABLET; ORAL

CORTISONE ACETATE

	@ PHARMACIA AND UPJOHN	5MG	N08126 003	Apr	DISC
	@	10MG	N08126 004	Apr	DISC
	@	25MG	N08126 001	Apr	DISC
+	WEST WARD	25MG	N80776 002	Apr	CRLD

CROMOLYN SODIUM

SOLUTION; INHALATION

CROMOLYN SODIUM

	@ ROXANE	10MG/ML	N75175 001	Sep 30, 1999	May	DISC
--	----------	---------	------------	--------------	-----	------

CROTAMITON

CREAM; TOPICAL

EURAX

>D>	+	RANBAXY	10%	N06927 001	Jun	CAHN
>A>	+		10%	N06927 001	Jun	CAHN

LOTION; TOPICAL

EURAX

>A>	AT	+	RANBAXY	10%	N09112 003	Jun	CAHN
>D>	AT	+	WESTWOOD SQUIBB	10%	N09112 003	Jun	CAHN

CYANOCOBALAMIN, CO-57

>D>			CAPSULE; ORAL			
>D>			RUBRATOPE-57			
>D>	+	BRACCO	0.5-1uCi	N16089 002	Jun	DISC
>A>	@		0.5-1uCi	N16089 002	Jun	DISC

CYCLOBENZAPRINE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

AMRIX

		ECR	15MG	N21777 001	Feb 01, 2007	Feb	NEWA
	+		30MG	N21777 002	Feb 01, 2007	Feb	NEWA

TABLET; ORAL

CYCLOBENZAPRINE HYDROCHLORIDE

AB		VINTAGE PHARMS	5MG	N77797 001	Feb 28, 2007	Feb	NEWA
AB			10MG	N77797 002	Feb 28, 2007	Feb	NEWA

CYCLOPENTOLATE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

CYCLOPENTOLATE HYDROCHLORIDE

>D>	AT		ALCON UNIVERSAL	1%	N89162 001	Jan 24, 1991	Jun	DISC
>A>	@			1%	N89162 001	Jan 24, 1991	Jun	DISC

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE

>D>	@	SICOR PHARMS	100MG/VIAL	N16793 001	Jun	CAHN	
>D>	@		500MG/VIAL	N16793 002	Jun	CAHN	
>D>	@		1GM/VIAL	N16793 003	Dec 21, 1987	Jun	CAHN
>D>	@		2GM/VIAL	N16793 004	Dec 21, 1987	Jun	CAHN
>A>	@	TEVA PARENTERAL	100MG/VIAL	N16793 001	Jun	CAHN	
>A>	@		500MG/VIAL	N16793 002	Jun	CAHN	
>A>	@		1GM/VIAL	N16793 003	Dec 21, 1987	Jun	CAHN
>A>	@		2GM/VIAL	N16793 004	Dec 21, 1987	Jun	CAHN

DALTEPARIN SODIUM

INJECTABLE; SUBCUTANEOUS

FRAGMIN

+	PHARMACIA AND UPJOHN	2,500IU/0.2ML (12,500IU/ML)	N20287 001	Dec 22, 1994	May	CDFR
+		5,000IU/0.2ML (25,000IU/ML)	N20287 003	Mar 18, 1996	May	CDFR
+		7,500IU/0.3ML (25,000IU/ML)	N20287 005	Apr 04, 2002	May	CDFR
+		10,000IU/ML (10,000IU/ML)	N20287 004	Jan 30, 1998	May	CDFR
+		10,000IU/0.4ML (25,000IU/ML)	N20287 002	May 01, 2007	May	NEWA
+		12,500IU/0.5ML (25,000IU/ML)	N20287 009	May 01, 2007	May	NEWA
+		15,000IU/0.6ML (25,000IU/ML)	N20287 010	May 01, 2007	May	NEWA
+		18,000IU/0.72ML (25,000IU/ML)	N20287 011	May 01, 2007	May	NEWA
+		95,000IU/9.5ML (10,000IU/ML)	N20287 007	Apr 04, 2002	May	CDFR
+		95,000IU/3.8ML (25,000IU/ML)	N20287 006	Apr 04, 2002	May	CDFR

DANAZOL

CAPSULE; ORAL

DANAZOL

AB	BARR	50MG	N74582 003	May 29, 1998	Apr	CTEC
AB		100MG	N74582 002	May 29, 1998	Apr	CTEC
AB	LANNETT	50MG	N78214 001	Apr 19, 2007	Apr	NEWA
AB		100MG	N78214 002	Apr 19, 2007	Apr	NEWA

>D> DAPIPRAZOLE HYDROCHLORIDE

>D> SOLUTION/DROPS; OPHTHALMIC

>D> REV-EYES

>D>	+	ANGELINI	0.5%	N19849 001	Dec 31, 1990	Jun	DISC
>A>		@	0.5%	N19849 001	Dec 31, 1990	Jun	DISC

DAPSONE

GEL; TOPICAL

ACZONE

@ QLT USA

5%

N21794 001 Jul 07, 2005 May CAHN

DEFEROXAMINE MESYLATE

INJECTABLE; INJECTION

DEFEROXAMINE MESYLATE

AP	BEDFORD	500MG/VIAL	N78086 001	May 30, 2007	May	NEWA
AP		2GM/VIAL	N78086 002	May 30, 2007	May	NEWA

DEMECLOCYCLINE HYDROCHLORIDE

TABLET; ORAL

DECLOMYCIN

@ STIEFEL

75MG

N50261 001 Mar CAHN

AB 150MG N50261 002 Mar CAHN

AB + 300MG N50261 003 Mar CAHN

DESIPRAMINE HYDROCHLORIDE

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

AB	ACTAVIS TOTOWA	10MG	N74430 001	Feb 09, 1996	Mar	CAHN
AB		25MG	N71601 001	Jun 05, 1987	Mar	CAHN
AB		75MG	N71602 001	Oct 05, 1987	Mar	CAHN
AB		100MG	N71766 001	Oct 05, 1987	Mar	CAHN
AB		150MG	N74430 002	Feb 09, 1996	Mar	CAHN

TABLET; ORAL

DESIPRAMINE HYDROCHLORIDE

AB	AMIDE PHARM	50MG	N71588 001	Jun 05, 1987	Mar	CAHN
----	-------------	------	------------	--------------	-----	------

DESMOPRESSIN ACETATE

INJECTABLE; INJECTION

DDAVP

>D>	+	SANOFI AVENTIS US	0.015MG/ML	N18938 002	Apr 25, 1995	Jun	DISC
-----	---	-------------------	------------	------------	--------------	-----	------

>A>	@		0.015MG/ML	N18938 002	Apr 25, 1995	Jun	DISC
-----	---	--	------------	------------	--------------	-----	------

>D>		SOLUTION; NASAL					
-----	--	-----------------	--	--	--	--	--

>D>		DDAVP					
-----	--	-------	--	--	--	--	--

>D>	+	SANOFI AVENTIS US	0.01%	N17922 001		Jun	DISC
-----	---	-------------------	-------	------------	--	-----	------

>A>	@		0.01%	N17922 001		Jun	DISC
-----	---	--	-------	------------	--	-----	------

SPRAY, METERED; NASAL

STIMATE

+	CSL BEHRING	0.15MG/SPRAY	N20355 001	Mar 07, 1994	Apr	CAHN
---	-------------	--------------	------------	--------------	-----	------

DESONIDE

CREAM; TOPICAL

DESONIDE

AB	+	PERRIGO NEW YORK	0.05%	N17010 001		May	CTNA
----	---	------------------	-------	------------	--	-----	------

>D>	AB	TEVA PHARMS	0.05%	N74027 001	Sep 28, 1992	Jun	DISC
-----	----	-------------	-------	------------	--------------	-----	------

>A>	@		0.05%	N74027 001	Sep 28, 1992	Jun	DISC
-----	---	--	-------	------------	--------------	-----	------

OINTMENT; TOPICAL

DESONIDE

AB	+	PERRIGO NEW YORK	0.05%	N17426 001		May	CTNA
----	---	------------------	-------	------------	--	-----	------

DEXMETHYLPHENIDATE HYDROCHLORIDE

TABLET; ORAL

DEXMETHYLPHENIDATE HYDROCHLORIDE

AB		TEVA PHARMS	2.5MG	N77107 003	Jan 29, 2007	Jan	NEWA
----	--	-------------	-------	------------	--------------	-----	------

AB			5MG	N77107 001	Jan 29, 2007	Jan	NEWA
----	--	--	-----	------------	--------------	-----	------

AB			10MG	N77107 002	Jan 29, 2007	Jan	NEWA
----	--	--	------	------------	--------------	-----	------

FOCALIN

AB		NOVARTIS	2.5MG	N21278 001	Nov 13, 2001	Jan	CFTG
----	--	----------	-------	------------	--------------	-----	------

AB			5MG	N21278 002	Nov 13, 2001	Jan	CFTG
----	--	--	-----	------------	--------------	-----	------

AB	+		10MG	N21278 003	Nov 13, 2001	Jan	CFTG
----	---	--	------	------------	--------------	-----	------

DEXTROAMPHETAMINE SULFATE

TABLET; ORAL

DEXEDRINE

@	GLAXOSMITHKLINE	5MG	N84935 001		Feb	DISC
---	-----------------	-----	------------	--	-----	------

DEXTROSTAT

AA	+	SHIRE	5MG	N84051 001		Mar	CAHN
----	---	-------	-----	------------	--	-----	------

AA	+		10MG	N84051 002		Mar	CAHN
----	---	--	------	------------	--	-----	------

DIAZOXIDE

SUSPENSION; ORAL

PROGLYCEM

+	IVAX RES	50MG/ML	N17453 001		May	CAHN
---	----------	---------	------------	--	-----	------

DICLOFENAC EPOLAMINE

PATCH; TOPICAL

FLECTOR

+	INST BIOCHEM	1.3%	N21234 001	Jan 31, 2007	Jan	NEWA
---	--------------	------	------------	--------------	-----	------

DICLOFENAC SODIUM

TABLET, DELAYED RELEASE; ORAL
DICLOFENAC SODIUM

>D>	AB	TEVA	50MG	N74723 001	Mar 30, 1999	Jun	DISC
>A>		@	50MG	N74723 001	Mar 30, 1999	Jun	DISC
>D>	AB		75MG	N74390 001	Aug 15, 1996	Jun	DISC
>A>		@	75MG	N74390 001	Aug 15, 1996	Jun	DISC
>D>	AB	TEVA PHARMS	25MG	N74459 001	Jun 25, 1997	Jun	DISC
>A>		@	25MG	N74459 001	Jun 25, 1997	Jun	DISC
>D>	AB		50MG	N74459 002	Jun 25, 1997	Jun	DISC
>A>		@	50MG	N74459 002	Jun 25, 1997	Jun	DISC
>D>	AB		75MG	N74459 003	Jun 25, 1997	Jun	DISC
>A>		@	75MG	N74459 003	Jun 25, 1997	Jun	DISC
>A>	AB	UNIQUE PHARM LABS	75MG	N77863 003	Jun 08, 2007	Jun	NEWA

DIDANOSINE

FOR SOLUTION; ORAL
DIDANOSINE

AA		AUROBINDO PHARMA	10MG/ML	N78112 001	Mar 08, 2007	Feb	NEWA
AA	+	BRISTOL MYERS SQUIBB	10MG/ML	N20156 001	Oct 09, 1991	Feb	CFTG

DIFLORASONE DIACETATE

OINTMENT; TOPICAL
PSORCON E

@	PHARMACIA AND UPJOHN	0.05%	N17994 001			May	CTNA
---	----------------------	-------	------------	--	--	-----	------

DIFLUNISAL

TABLET; ORAL
DIFLUNISAL

>D>	AB	TEVA	250MG	N73679 001	Jul 31, 1992	Jun	DISC
>A>		@	250MG	N73679 001	Jul 31, 1992	Jun	DISC

DILTIAZEM HYDROCHLORIDE

TABLET; ORAL
DILTIAZEM HYDROCHLORIDE

>D>	AB	IVAX PHARMS	30MG	N74168 001	Mar 03, 1995	Jun	DISC
>A>		@	30MG	N74168 001	Mar 03, 1995	Jun	DISC
>D>	AB		60MG	N74168 002	Mar 03, 1995	Jun	DISC
>A>		@	60MG	N74168 002	Mar 03, 1995	Jun	DISC
>D>	AB		90MG	N74168 003	Mar 03, 1995	Jun	DISC
>A>		@	90MG	N74168 003	Mar 03, 1995	Jun	DISC
>D>	AB		120MG	N74168 004	Mar 03, 1995	Jun	DISC
>A>		@	120MG	N74168 004	Mar 03, 1995	Jun	DISC

DIMYRISTOYL LECITHIN; PERFLEXANE

INJECTABLE; INTRAVENOUS
IMAGENT

@	IMCOR PHARMS CO	0.92MG/VIAL;0.092MG/VIAL	N21191 001	May 31, 2002	Feb	CAHN
---	-----------------	--------------------------	------------	--------------	-----	------

DIPHENHYDRAMINE HYDROCHLORIDE

INJECTABLE; INJECTION
BENADRYL

@	MCNEIL CONS	10MG/ML	N06146 001		Mar	CAHN
---	-------------	---------	------------	--	-----	------

INJECTABLE; INJECTION

BENADRYL

AP	+	MCNEIL CONS	50MG/ML	N06146 002			Mar	CAHN
BENADRYL PRESERVATIVE FREE								
AP	+	MCNEIL CONS	50MG/ML	N09486 001			Mar	CAHN

DIPYRIDAMOLE

TABLET; ORAL

DIPYRIDAMOLE

AB		MURTY PHARMS	25MG	N40733 001	Feb 13, 2007	Jan	NEWA
AB			50MG	N40733 002	Feb 13, 2007	Jan	NEWA
AB			75MG	N40733 003	Feb 13, 2007	Jan	NEWA

DOXAZOSIN MESYLATE

TABLET; ORAL

DOXAZOSIN MESYLATE

>D>	AB	TEVA	EQ 1MG BASE	N75353 001	Jan 12, 2001	Jun	DISC
>A>		@	EQ 1MG BASE	N75353 001	Jan 12, 2001	Jun	DISC
>D>	AB		EQ 2MG BASE	N75353 002	Jan 12, 2001	Jun	DISC
>A>		@	EQ 2MG BASE	N75353 002	Jan 12, 2001	Jun	DISC
>D>	AB		EQ 4MG BASE	N75353 003	Jan 12, 2001	Jun	DISC
>A>		@	EQ 4MG BASE	N75353 003	Jan 12, 2001	Jun	DISC
>D>	AB		EQ 8MG BASE	N75353 004	Jan 12, 2001	Jun	DISC
>A>		@	EQ 8MG BASE	N75353 004	Jan 12, 2001	Jun	DISC

DOXEPIN HYDROCHLORIDE

CAPSULE; ORAL

DOXEPIN HYDROCHLORIDE

>D>	AB	SANDOZ	EQ 10MG BASE	N71487 001	Mar 02, 1987	Jun	DISC
>A>		@	EQ 10MG BASE	N71487 001	Mar 02, 1987	Jun	DISC
>D>	AB	WATSON LABS	EQ 10MG BASE	N72985 001	Mar 29, 1991	Jun	DISC
>A>		@	EQ 10MG BASE	N72985 001	Mar 29, 1991	Jun	DISC
>D>	AB		EQ 25MG BASE	N72986 001	Mar 29, 1991	Jun	DISC
>A>		@	EQ 25MG BASE	N72986 001	Mar 29, 1991	Jun	DISC
>D>	AB		EQ 50MG BASE	N72987 001	Mar 29, 1991	Jun	DISC
>A>		@	EQ 50MG BASE	N72987 001	Mar 29, 1991	Jun	DISC

DOXORUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

DOXORUBICIN HYDROCHLORIDE

AP	+	BEDFORD	10MG/VIAL	N62921 001	Mar 17, 1989	Apr	CRLD	
AP	+		20MG/VIAL	N62921 002	Mar 17, 1989	Apr	CRLD	
AP	+		50MG/VIAL	N62921 003	Mar 17, 1989	Apr	CRLD	
RUBEX								
		@ BRISTOL MYERS SQUIBB	10MG/VIAL	N62926 001	Apr 13, 1989	Mar	DISC	
		@	50MG/VIAL	N62926 002	Apr 13, 1989	Mar	DISC	
		@	100MG/VIAL	N62926 003	Apr 13, 1989	Mar	DISC	

DOXYCYCLINE

CAPSULE; ORAL

DOXYCYCLINE

AB		RANBAXY	EQ 75MG BASE	N65053 003	Sep 10, 2003	Jan	CTEC	
MONODOX								
AB		OCLASSEN	EQ 75MG BASE	N50641 003	Oct 18, 2006	Jan	NEWA	

TABLET; ORAL

DOXYCYCLINE

>A>	AB	MYLAN	EQ 150MG BASE	N65427 001	Jun 07, 2007	Jun	NEWA
>D>	+	PAR PHARM	EQ 150MG BASE	N65070 004	Jul 14, 2005	Jun	CTEC
>A>	AB	+	EQ 150MG BASE	N65070 004	Jul 14, 2005	Jun	CTEC

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

DOXYCYCLINE HYCLATE

>D>	AB	MYLAN	EQ 50MG BASE	N62337 001	Mar 29, 1982	Jun	DISC
>A>		@	EQ 50MG BASE	N62337 001	Mar 29, 1982	Jun	DISC
>D>	AB		EQ 100MG BASE	N62337 002	Mar 29, 1982	Jun	DISC
>A>		@	EQ 100MG BASE	N62337 002	Mar 29, 1982	Jun	DISC

TABLET; ORAL

DOXYCYCLINE HYCLATE

>D>	AB	MYLAN	EQ 100MG BASE	N62432 001	Feb 15, 1983	Jun	DISC
>A>		@	EQ 100MG BASE	N62432 001	Feb 15, 1983	Jun	DISC

ECONAZOLE NITRATE

CREAM; TOPICAL

ECONAZOLE NITRATE

>A>	AB	DPT LABS	1%	N76574 001	Dec 17, 2004	Jun	CAHN
>D>	AB	PRASCO	1%	N76574 001	Dec 17, 2004	Jun	CAHN
	AB		1%	N76574 001	Dec 17, 2004	May	CAHN

EDETATE CALCIUM DISODIUM

INJECTABLE; INJECTION

CALCIUM DISODIUM VERSENATE

	+	GRACEWAY	200MG/ML	N08922 001		Jan	CAHN
--	---	----------	----------	------------	--	-----	------

TABLET; ORAL

CALCIUM DISODIUM VERSENATE

	@	GRACEWAY	500MG	N08922 002		Jan	CAHN
--	---	----------	-------	------------	--	-----	------

ENALAPRILAT

INJECTABLE; INJECTION

ENALAPRILAT

AP	+	BEDFORD	1.25MG/ML	N75634 001	Aug 22, 2000	Mar	CRLD
----	---	---------	-----------	------------	--------------	-----	------

VASOTEC

AP	+	BIOVAIL LABS INTL	1.25MG/ML	N19309 001	Feb 09, 1988	May	CMFD
----	---	-------------------	-----------	------------	--------------	-----	------

	@		1.25MG/ML	N19309 001	Feb 09, 1988	Mar	DISC
--	---	--	-----------	------------	--------------	-----	------

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ELLENC

AP		PFIZER INC	2MG/ML (50MG/25ML)	N50778 002	Sep 15, 1999	Apr	CFTG
----	--	------------	--------------------	------------	--------------	-----	------

AP	+		2MG/ML (200MG/100ML)	N50778 001	Sep 15, 1999	Apr	CFTG
----	---	--	----------------------	------------	--------------	-----	------

EPIRUBICIN HYDROCHLORIDE

>A>	AP	BEDFORD	2MG/ML (50MG/25ML)	N65289 001	Jun 27, 2007	Jun	NEWA
-----	----	---------	--------------------	------------	--------------	-----	------

>A>	AP		2MG/ML (200MG/100ML)	N65289 002	Jun 27, 2007	Jun	NEWA
-----	----	--	----------------------	------------	--------------	-----	------

AP		HOSPIRA	2MG/ML (50MG/25ML)	N65343 002	Apr 19, 2007	Apr	NEWA
----	--	---------	--------------------	------------	--------------	-----	------

			2MG/ML (10MG/5ML)	N65343 001	Apr 19, 2007	Apr	NEWA
--	--	--	-------------------	------------	--------------	-----	------

			2MG/ML (150MG/75ML)	N65343 003	Apr 19, 2007	Apr	NEWA
--	--	--	---------------------	------------	--------------	-----	------

AP			2MG/ML (200MG/100ML)	N65343 004	Apr 19, 2007	Apr	NEWA
----	--	--	----------------------	------------	--------------	-----	------

ERGOTAMINE TARTRATE

TABLET; SUBLINGUAL

ERGOMAR

	+	ROSEDALE THERAPEUTIC	2MG	N87693	001	Feb 24, 1983	Jan	CAHN
--	---	----------------------	-----	--------	-----	--------------	-----	------

ERYTHROMYCIN

GEL; TOPICAL

ERYTHROMYCIN

>A>	AT	PERRIGO	2%	N63211	001	Jan 29, 1993	Jun	CAHN
>D>	AT	STIEFEL	2%	N63211	001	Jan 29, 1993	Jun	CAHN

OINTMENT; TOPICAL

AKNE-MYCIN

	+	CORIA	2%	N50584	001	Jan 10, 1985	May	CAHN
--	---	-------	----	--------	-----	--------------	-----	------

SWAB; TOPICAL

C-SOLVE-2

	@	IVAX PHARMS	2%	N62751	001	Jul 30, 1993	May	DISC
--	---	-------------	----	--------	-----	--------------	-----	------

ERYCETTE

	@	J AND J	2%	N50594	001	Feb 15, 1985	May	DISC
--	---	---------	----	--------	-----	--------------	-----	------

ERYTHROMYCIN

	AT	+	ALTANA	2%	N65320	001	Jul 25, 2006	May	CRLD
>A>	AT	+	PERRIGO	2%	N64126	001	Jul 03, 1996	Jun	CAHN
>D>	AT	+	STIEFEL	2%	N64126	001	Jul 03, 1996	Jun	CAHN
	AT	+		2%	N64126	001	Jul 03, 1996	May	CRLD
		@		2%	N64128	001	Jul 03, 1996	May	DISC

ERYTHROMYCIN ETHYLSUCCINATE

TABLET; ORAL

E.E.S. 400

>D>	AB	+	ABBOTT	EQ 400MG BASE	N61905	002	Aug 12, 1982	Jun	CTEC
>A>		+		EQ 400MG BASE	N61905	002	Aug 12, 1982	Jun	CTEC
>D>			ERYTHROMYCIN ETHYLSUCCINATE						
>D>	AB		MYLAN	EQ 400MG BASE	N62847	001	Sep 14, 1988	Jun	DISC
>A>		@		EQ 400MG BASE	N62847	001	Sep 14, 1988	Jun	DISC

ERYTHROMYCIN STEARATE

TABLET; ORAL

ERYTHROCIN STEARATE

>D>	AB	+	ABBOTT	EQ 500MG BASE	N60359	003		Jun	CTEC
>A>		+		EQ 500MG BASE	N60359	003		Jun	CTEC
>D>	AB		MYLAN	EQ 250MG BASE	N61505	001		Jun	DISC
>A>		@		EQ 250MG BASE	N61505	001		Jun	DISC
>D>	AB			EQ 500MG BASE	N61505	002		Jun	DISC
>A>		@		EQ 500MG BASE	N61505	002		Jun	DISC

ESMOLOL HYDROCHLORIDE

INJECTABLE; INJECTION

ESMOLOL HYDROCHLORIDE

>A>	AP		BIONICHE PHARMA	10MG/ML	N76474	001	May 02, 2005	Jun	CAHN
>D>	AP		PHARMAFORCE	10MG/ML	N76474	001	May 02, 2005	Jun	CAHN

ESTAZOLAM

	TABLET; ORAL								
	ESTAZOLAM								
AB	+ TEVA	2MG		N74921	002	Jul 10, 1997	May	CRLD	
	PROSOM								
	@ ABBOTT	1MG		N19080	001	Dec 26, 1990	Apr	DISC	
	@	2MG		N19080	002	Dec 26, 1990	Apr	DISC	

ESTRADIOL

>A>	GEL; TOPICAL								
>A>	DIVIGEL								
>A>	+ UPSHER SMITH	0.1%		N22038	003	Jun 04, 2007	Jun	NEWA	
	GEL, METERED; TRANSDERMAL								
	ELESTRIN								
BX	+ BRADLEY PHARMS	0.06%		N21813	001	Dec 15, 2006	Jan	CAHN	
	TABLET; ORAL								
	ESTRADIOL								
>A>	@ AAI PHARMA INC	0.5MG		N40138	001	Jan 30, 1998	Jun	DISC	
>A>	@	1MG		N40138	002	Jan 30, 1998	Jun	DISC	
>A>	@	2MG		N40138	003	Jan 30, 1998	Jun	DISC	
>D>	AB APPLIED ANAL	0.5MG		N40138	001	Jan 30, 1998	Jun	DISC	
>D>	AB	1MG		N40138	002	Jan 30, 1998	Jun	DISC	
>D>	AB	2MG		N40138	003	Jan 30, 1998	Jun	DISC	
	@ HERITAGE PHARMS INC	0.5MG		N40275	001	Dec 29, 1998	Feb	CAHN	
	@	1MG		N40275	002	Dec 29, 1998	Feb	CAHN	
	@	2MG		N40275	003	Dec 29, 1998	Feb	CAHN	

ESTROGENS, CONJUGATED SYNTHETIC B

	TABLET; ORAL								
	ENJUVIA								
	DURAMED	0.9MG		N21443	005	Apr 27, 2007	Apr	NEWA	

ESTROPIPATE

	TABLET; ORAL								
	ESTROPIPATE								
>D>	AB MYLAN	0.75MG		N40359	001	Aug 26, 1999	Jun	DISC	
>A>	@	0.75MG		N40359	001	Aug 26, 1999	Jun	DISC	
>D>	AB	1.5MG		N40359	002	Aug 26, 1999	Jun	DISC	
>A>	@	1.5MG		N40359	002	Aug 26, 1999	Jun	DISC	
>D>	AB	3MG		N40359	003	Aug 26, 1999	Jun	DISC	
>A>	@	3MG		N40359	003	Aug 26, 1999	Jun	DISC	

ETHINYL ESTRADIOL; LEVONORGESTREL

	TABLET; ORAL								
	LYBREL								
	+ WYETH PHARMS INC	0.02MG;0.09MG		N21864	001	May 22, 2007	May	NEWA	
	TABLET; ORAL-28								
	LEVONORGESTREL AND ETHINYL ESTRADIOL								
AB1	+ WATSON LABS	0.02MG;0.1MG		N76625	001	Nov 18, 2004	Apr	CRLD	

ETHINYL ESTRADIOL; NORETHINDRONE

	TABLET, CHEWABLE; ORAL-28								
	OVCON-35 FE								
	+ WARNER CHILCOTT	0.035MG;0.4MG		N21490	001	Nov 14, 2003	Jan	CTNA	

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL-21		MICROGESTIN 1.5/30	
AB	WATSON LABS	0.03MG;1.5MG	N75548 002 Jul 30, 2003 Jan NEWA
TABLET; ORAL-28		NORETHINDRONE ACETATE AND ETHINYL ESTRADIOL AND FERROUS FUMARATE	
AB	TEVA PHARMS	0.02MG;1MG	N77077 001 May 20, 2005 Apr CAHN
AB		0.03MG;1.5MG	N77075 001 Apr 28, 2005 Apr CAHN

ETHINYL ESTRADIOL; NORGESTIMATE

TABLET; ORAL-28		PREVIFEM	
AB	TEVA PHARMS	0.035MG;0.25MG	N76334 001 Jan 09, 2004 Apr CAHN
TABLET; ORAL-28		TRI-PREVIFEM	
AB	TEVA PHARMS	0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	N76335 001 Mar 26, 2004 Apr CAHN
AB		0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	N76335 001 Mar 26, 2004 Apr CAHN
AB		0.035MG,0.035MG,0.035MG;0.18MG,0.215MG,0.25MG	N76335 001 Mar 26, 2004 Apr CAHN

ETODOLAC

CAPSULE; ORAL

ETODOLAC

@	AAIPHARMA LLC	300MG	N74929 001 Jan 30, 1998 Apr DISC
@	GENPHARM	200MG	N75071 001 Sep 30, 1998 Apr DISC
@		300MG	N75071 002 Sep 30, 1998 Apr DISC
@	MYLAN	200MG	N74932 001 May 16, 1997 Apr DISC
@		300MG	N74932 002 May 16, 1997 Apr DISC
@	SANDOZ	200MG	N74840 001 Aug 29, 1997 Apr DISC
@		200MG	N74942 001 Sep 30, 1997 Apr DISC
@		300MG	N74840 002 Aug 29, 1997 Apr DISC
@		300MG	N74942 002 Sep 30, 1997 Apr DISC
@	TEVA	200MG	N75126 001 Sep 16, 1999 Apr DISC
@	WATSON LABS	200MG	N74844 001 Dec 23, 1997 Apr DISC
@		300MG	N74844 002 Dec 23, 1997 Apr DISC

TABLET; ORAL

ETODOLAC

@	AAIPHARMA LLC	400MG	N74927 001 Oct 30, 1997 Apr DISC
@	GENPHARM	400MG	N75012 001 Sep 30, 1998 Apr DISC
@		500MG	N75012 002 Sep 30, 1998 Apr DISC
@	IVAX PHARMS	400MG	N74883 001 Feb 28, 1997 Apr DISC
@		500MG	N74883 002 Nov 20, 1998 Apr DISC
@	RANBAXY	400MG	N75226 001 Nov 24, 1998 Apr DISC
@		500MG	N75226 002 Nov 24, 1998 Apr DISC
@	SANDOZ	400MG	N74839 001 Jul 11, 1997 Apr DISC
@		400MG	N74846 001 Feb 28, 1997 Apr DISC
@	TEVA	400MG	N74847 001 Apr 23, 1999 Apr DISC
@		500MG	N74847 002 Apr 23, 1999 Apr DISC
@	WATSON LABS	400MG	N74892 001 Apr 16, 1997 Apr DISC
@		400MG	N75069 001 Apr 16, 1998 Apr DISC
@		500MG	N74892 002 Oct 29, 1998 Apr DISC

TABLET, EXTENDED RELEASE; ORAL

ETODOLAC

@	ANDRX PHARMS	400MG	N75829 001	Nov 30, 2001	Apr	DISC
@		500MG	N75829 002	Nov 30, 2001	Apr	DISC
@	SANDOZ	400MG	N75943 001	Jul 26, 2002	Apr	DISC
@		500MG	N75943 002	Jul 26, 2002	Apr	DISC
@		600MG	N75943 003	Jul 26, 2002	Apr	DISC

FAMOTIDINE

FOR SUSPENSION; ORAL

PEPCID

+	SALIX PHARMS	40MG/5ML	N19527 001	Feb 02, 1987	Feb	CAHN
---	--------------	----------	------------	--------------	-----	------

FENOFIBRATE

TABLET; ORAL

TRIGLIDE

	SKYEPHARMA AG	50MG	N21350 001	May 07, 2005	Jan	CAHN
BX		160MG	N21350 002	May 07, 2005	Jan	CAHN

FENOLDOPAM MESYLATE

INJECTABLE; INJECTION

FENOLDOPAM MESYLATE

AP	SICOR PHARMS	EQ 10MG BASE/ML	N77826 001	Mar 07, 2007	Feb	NEWA
----	--------------	-----------------	------------	--------------	-----	------

FENTANYL

FILM, EXTENDED RELEASE; TRANSDERMAL

DURAGESIC-12

AB	ALZA	12.5UGM/HR	N19813 005	Feb 04, 2005	Jan	CFTG
----	------	------------	------------	--------------	-----	------

FENTANYL-100

AB	LAVIPHARM LABS	100UGM/HR	N77051 004	Aug 04, 2006	Jan	CTNA
----	----------------	-----------	------------	--------------	-----	------

AB	MYLAN TECHNOLOGIES	100UGM/HR	N76258 004	Jan 28, 2005	Jan	CTNA
----	--------------------	-----------	------------	--------------	-----	------

FENTANYL-12

AB	MYLAN TECHNOLOGIES	12.5UGM/HR	N76258 005	Jan 23, 2007	Jan	NEWA
----	--------------------	------------	------------	--------------	-----	------

FENTANYL-25

AB	LAVIPHARM LABS	25UGM/HR	N77051 001	Aug 04, 2006	Jan	CTNA
----	----------------	----------	------------	--------------	-----	------

AB	MYLAN TECHNOLOGIES	25UGM/HR	N76258 001	Jan 28, 2005	Jan	CTNA
----	--------------------	----------	------------	--------------	-----	------

FENTANYL-50

AB	LAVIPHARM LABS	50UGM/HR	N77051 002	Aug 04, 2006	Jan	CTNA
----	----------------	----------	------------	--------------	-----	------

AB	MYLAN TECHNOLOGIES	50UGM/HR	N76258 002	Jan 28, 2005	Jan	CTNA
----	--------------------	----------	------------	--------------	-----	------

FENTANYL-75

AB	LAVIPHARM LABS	75UGM/HR	N77051 003	Aug 04, 2006	Jan	CTNA
----	----------------	----------	------------	--------------	-----	------

AB	MYLAN TECHNOLOGIES	75UGM/HR	N76258 003	Jan 28, 2005	Jan	CTNA
----	--------------------	----------	------------	--------------	-----	------

FENTANYL CITRATE

TABLET; BUCCAL

FENTORA

CEPHALON

EQ 0.3MG BASE

N21947 006	Mar 02, 2007	Mar	NEWA
------------	--------------	-----	------

FEXOFENADINE HYDROCHLORIDE

TABLET; ORAL

FEXOFENADINE HYDROCHLORIDE

AB	MYLAN	180MG	N77081 001	Apr 16, 2007	Apr	NEWA
----	-------	-------	------------	--------------	-----	------

FINASTERIDE

TABLET; ORAL

FINASTERIDE

AB	ACTAVIS TOTOWA	5MG	N77914 001	Mar 28, 2007	Mar	NEWA
AB	DR REDDYS LABS LTD	5MG	N76437 001	Feb 28, 2007	Feb	NEWA

FLECAINIDE ACETATE

TABLET; ORAL

TAMBOCOR

AB	GRACEWAY	50MG	N18830 004	Aug 23, 1988	Jan	CAHN
AB		100MG	N18830 001	Oct 31, 1985	Jan	CAHN
AB	+	150MG	N18830 003	Jun 03, 1988	Jan	CAHN
	@	200MG	N18830 002	Oct 31, 1985	Jan	CAHN

FLUDARABINE PHOSPHATE

INJECTABLE; INJECTION

FLUDARABINE PHOSPHATE

AP	MAYNE PHARMA USA	50MG/VIAL	N77790 001	Apr 06, 2007	Mar	NEWA
----	------------------	-----------	------------	--------------	-----	------

FLUDROCORTISONE ACETATE

TABLET; ORAL

FLORINEF

>D>						
>D>	AB	+	KING PHARMS	0.1MG		Jun DISC
>A>			@	0.1MG		Jun DISC

FLUDROCORTISONE ACETATE

>D>	AB		BARR	0.1MG	N40425 001	Jan 21, 2003 Jun CRLD
>A>	AB	+		0.1MG	N40425 001	Jan 21, 2003 Jun CRLD

FLUOCINONIDE

SOLUTION; TOPICAL

FLUOCINONIDE

>D>	AT		TEVA PHARMS	0.05%	N72522 001	Sep 28, 1990 Jun DISC
>A>			@	0.05%	N72522 001	Sep 28, 1990 Jun DISC

FLUOROMETHOLONE

SUSPENSION/DROPS; OPHTHALMIC

FLUOR-OP

>D>	AB		NOVARTIS	0.1%	N70185 001	Feb 27, 1986 Jun DISC
>A>			@	0.1%	N70185 001	Feb 27, 1986 Jun DISC

FML

>D>	AB	+	ALLERGAN	0.1%	N16851 002	Jul 28, 1982 Jun CTEC
>A>			+	0.1%	N16851 002	Jul 28, 1982 Jun CTEC

FLUOROMETHOLONE; SULFACETAMIDE SODIUM

SUSPENSION/DROPS; OPHTHALMIC

FML-S

>D>						
>D>		+	ALLERGAN	0.1%;10%	N19525 001	Sep 29, 1989 Jun DISC
>A>			@	0.1%;10%	N19525 001	Sep 29, 1989 Jun DISC

FLUOROURACIL

INJECTABLE; INJECTION

FLUOROURACIL

@ ABRAXIS PHARM

50MG/ML

N40291 001 Mar 24, 1999 Apr DISC

@

50MG/ML

N40379 001 Nov 15, 2000 Apr DISC

INJECTABLE; INJECTION

FLUOROURACIL

AP	+	ABRAXIS PHARM	500MG/10ML (50MG/ML)	N40279 002	Sep 30, 1998	Apr	NEWA
AP	+		1GM/20ML (50MG/ML)	N40279 001	Sep 30, 1998	Apr	CPOT
AP	+		2.5GM/50ML (50MG/ML)	N40278 001	Sep 30, 1998	Apr	CPOT
AP	+		5GM/100ML (50MG/ML)	N40278 002	Sep 30, 1998	Apr	NEWA
AP	+	GENERAMEDIX	500MG/10ML (50MG/ML)	N40743 002	Apr 26, 2007	Apr	NEWA
AP	+		1GM/20ML (50MG/ML)	N40743 001	Apr 26, 2007	Apr	NEWA
AP	+		2.5GM/50ML (50MG/ML)	N40798 002	Apr 26, 2007	Apr	NEWA
AP	+		5GM/100ML (50MG/ML)	N40798 001	Apr 26, 2007	Apr	NEWA
AP	+	SICOR PHARMS	500MG/10ML (50MG/ML)	N40333 001	Jan 27, 2000	Apr	CPOT
AP	+		2.5GM/50ML (50MG/ML)	N40334 001	Feb 25, 2000	Apr	CPOT
AP	+		5GM/100ML (50MG/ML)	N40334 002	Feb 25, 2000	Apr	NEWA
AP	+	VALEANT	500MG/10ML (50MG/ML)	N12209 001		Apr	CPOT

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

FLUOXETINE

AB		MYLAN	EQ 40MG BASE	N75207 003	May 25, 2007	May	NEWA
>D>	AB	WATSON LABS	EQ 10MG BASE	N75662 001	Jan 29, 2002	Jun	DISC
>A>		@	EQ 10MG BASE	N75662 001	Jan 29, 2002	Jun	DISC
>D>	AB		EQ 20MG BASE	N75662 002	Jan 29, 2002	Jun	DISC
>A>		@	EQ 20MG BASE	N75662 002	Jan 29, 2002	Jun	DISC

SOLUTION; ORAL

FLUOXETINE HYDROCHLORIDE

AA		SILARX	EQ 20MG BASE/5ML	N77849 001	Feb 09, 2007	Jan	NEWA
----	--	--------	------------------	------------	--------------	-----	------

FLUOXETINE HYDROCHLORIDE; OLANZAPINE

CAPSULE; ORAL

SYMBYAX

LILLY

EQ 25MG BASE;EQ 3MG BASE

N21520 001 Apr 09, 2007 Apr NEWA

FLUPHENAZINE DECANOATE

INJECTABLE; INJECTION

FLUPHENAZINE DECANOATE

AO	+	BEDFORD	25MG/ML	N74531 001	Aug 30, 1996	Jan	CRLD
		PROLIXIN DECANOATE					
		@ BRISTOL MYERS SQUIBB	25MG/ML	N16727 001		Jan	DISC

FLUPHENAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

FLUPHENAZINE HYDROCHLORIDE

+ ABRAXIS PHARM 2.5MG/ML

N89556 001 Apr 16, 1987 Jan CRLD

PROLIXIN

@ APOTHECON 2.5MG/ML

N11751 005 Jan DISC

TABLET; ORAL

FLUPHENAZINE HYDROCHLORIDE

AB	+	MYLAN	10MG	N89804 001	Aug 12, 1988	Feb	CRLD
		PROLIXIN					
		@ APOTHECON	1MG	N11751 004		Jan	DISC
		@	2.5MG	N11751 001		Jan	DISC
		@	5MG	N11751 003		Jan	DISC
		@	10MG	N11751 002		Jan	DISC

FLURBIPROFEN

TABLET; ORAL

FLURBIPROFEN

>D>	AB	TEVA	50MG	N74405 002	May 24, 1995	Jun	DISC
>A>		@	50MG	N74405 002	May 24, 1995	Jun	DISC
>D>	AB		100MG	N74405 001	May 24, 1995	Jun	DISC
>A>		@	100MG	N74405 001	May 24, 1995	Jun	DISC
	AB	THERAGEN	100MG	N74560 002	May 16, 1997	Apr	CAHN

FLUTICASONE FUROATE

SPRAY, METERED; NASAL

VERAMYST

	+	GLAXOSMITHKLINE	0.0275MG/INH	N22051 001	Apr 27, 2007	Apr	NEWA
--	---	-----------------	--------------	------------	--------------	-----	------

FLUTICASONE PROPIONATE

POWDER; INHALATION

FLOVENT

>D>							
>D>	+	GLAXOSMITHKLINE	0.044MG/INH	N20549 001	Nov 07, 1997	Jun	DISC
>A>		@	0.044MG/INH	N20549 001	Nov 07, 1997	Jun	DISC
>D>	+		0.088MG/INH	N20549 002	Nov 07, 1997	Jun	DISC
>A>		@	0.088MG/INH	N20549 002	Nov 07, 1997	Jun	DISC
>D>	+		0.22MG/INH	N20549 003	Nov 07, 1997	Jun	DISC
>A>		@	0.22MG/INH	N20549 003	Nov 07, 1997	Jun	DISC

FLUVOXAMINE MALEATE

TABLET; ORAL

FLUVOXAMINE MALEATE

>D>	AB	MUTUAL PHARM	25MG	N76125 001	Apr 29, 2002	Jun	DISC
>A>		@	25MG	N76125 001	Apr 29, 2002	Jun	DISC
>D>	AB		50MG	N76125 002	Apr 29, 2002	Jun	DISC
>A>		@	50MG	N76125 002	Apr 29, 2002	Jun	DISC
>D>	AB		100MG	N76125 003	Apr 29, 2002	Jun	DISC
>A>		@	100MG	N76125 003	Apr 29, 2002	Jun	DISC
>D>	AB	SANDOZ	25MG	N75887 001	Jan 05, 2001	Jun	DISC
>A>		@	25MG	N75887 001	Jan 05, 2001	Jun	DISC
>D>	AB		50MG	N75887 002	Jan 05, 2001	Jun	DISC
>A>		@	50MG	N75887 002	Jan 05, 2001	Jun	DISC
>D>	AB		100MG	N75887 003	Jan 05, 2001	Jun	DISC
>A>		@	100MG	N75887 003	Jan 05, 2001	Jun	DISC
>D>	AB	WATSON LABS	25MG	N75894 001	Apr 18, 2001	Jun	DISC
>A>		@	25MG	N75894 001	Apr 18, 2001	Jun	DISC
>D>	AB		50MG	N75894 002	Apr 18, 2001	Jun	DISC
>A>		@	50MG	N75894 002	Apr 18, 2001	Jun	DISC
>D>	AB		100MG	N75894 003	Apr 18, 2001	Jun	DISC
>A>		@	100MG	N75894 003	Apr 18, 2001	Jun	DISC

FOLIC ACID

INJECTABLE; INJECTION

FOLIC ACID

	+	ABRAXIS PHARM	5MG/ML	N89202 001	Feb 18, 1986	Jan	CTEC
		@ BEN VENUE	5MG/ML	N81066 001	Dec 29, 1993	Jan	DISC

FOLLITROPIN ALFA/BETA

INJECTABLE; SUBCUTANEOUS

GONAL-F

@	EMD SERONO	37.5 IU/VIAL	N20378 003	May 25, 2000	Mar	CAHN
@		37.5 IU/VIAL	N21765 001	Mar 25, 2004	Mar	CAHN
@		75 IU/VIAL	N20378 001	Sep 29, 1997	Mar	CAHN
@		150 IU/VIAL	N20378 002	Sep 29, 1997	Mar	CAHN
@		150 IU/VIAL	N21765 003	Mar 25, 2004	Mar	CAHN
+		450 IU/VIAL	N20378 005	Mar 26, 2004	Mar	CAHN
@		1,050 IU/VIAL	N20378 004	Feb 28, 2001	Mar	CAHN

GONAL-F RFF

+	EMD SERONO	75 IU/VIAL	N21765 002	Mar 25, 2004	Mar	CAHN
---	------------	------------	------------	--------------	-----	------

GONAL-F RFF PEN

>A>	+	EMD SERONO	300 IU/0.5ML	N21684 001	May 25, 2004	Jun	CAHN
>A>	+		450 IU/0.75ML	N21684 002	May 25, 2004	Jun	CAHN
>A>	+		900 IU/1.5ML	N21684 003	May 25, 2004	Jun	CAHN
>D>	+	SERONO INC	300 IU/0.5ML	N21684 001	May 25, 2004	Jun	CAHN
>D>	+		450 IU/0.75ML	N21684 002	May 25, 2004	Jun	CAHN
>D>	+		900 IU/1.5ML	N21684 003	May 25, 2004	Jun	CAHN

FORMOTEROL FUMARATE

POWDER; INHALATION

FORADIL CERTIHALER

+	NOVARTIS	0.0085MG/INH	N21592 001	Dec 15, 2006	Jan	CRLD
---	----------	--------------	------------	--------------	-----	------

SOLUTION; INHALATION

PERFOROMIST

+	DEY LP	0.02MG/2ML	N22007 001	May 11, 2007	May	NEWA
---	--------	------------	------------	--------------	-----	------

FOSAMPRENAVIR CALCIUM

>A>		SUSPENSION; ORAL					
>A>		LEXIVA					
>A>	+	GLAXO	50MG BASE/ML	N22116 001	Jun 14, 2007	Jun	NEWA

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

AP	WOCKHARDT	10MG/ML	N77941 001	Mar 22, 2007	Mar	NEWA
----	-----------	---------	------------	--------------	-----	------

GABAPENTIN

CAPSULE; ORAL

GABAPENTIN

AB	WATSON LABS	100MG	N75485 003	May 11, 2007	May	NEWA
AB		300MG	N75485 002	May 11, 2007	May	NEWA
AB		400MG	N75485 001	May 11, 2007	May	NEWA

TABLET; ORAL

GABAPENTIN

AB	+	IVAX PHARMS	800MG	N76017 005	Apr 29, 2005	Feb	CRLD
AB		PFIZER PHARMS	800MG	N20882 002	Oct 09, 1998	Feb	CRLD

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

AB	IMPAX PHARMS	600MG	N78207 001	Jun 01, 2007	May	NEWA
----	--------------	-------	------------	--------------	-----	------

TABLET; ORAL

GEMFIBROZIL

>D>	AB	MYLAN	600MG	N74452 001	Feb 16, 1995	Jun	DISC
>A>		@	600MG	N74452 001	Feb 16, 1995	Jun	DISC
	AB	PERRIGO R AND D	600MG	N78012 001	Mar 26, 2007	Mar	NEWA

GLIPIZIDE

TABLET; ORAL

GLIPIZIDE

>D>	AB	TEVA	5MG	N74387 001	Mar 04, 1996	Jun	DISC
>A>		@	5MG	N74387 001	Mar 04, 1996	Jun	DISC
>D>	AB		10MG	N74387 002	Mar 04, 1996	Jun	DISC
>A>		@	10MG	N74387 002	Mar 04, 1996	Jun	DISC

GLIPIZIDE; METFORMIN HYDROCHLORIDE

TABLET; ORAL

GLIPIZIDE AND METFORMIN HYDROCHLORIDE

	AB	MYLAN	2.5MG;250MG	N78083 001	Apr 12, 2007	Apr	NEWA
	AB		2.5MG;500MG	N78083 002	Apr 12, 2007	Apr	NEWA
	AB		5MG;500MG	N78083 003	Apr 12, 2007	Apr	NEWA

GRISEOFULVIN, MICROCRYSTALLINE

SUSPENSION; ORAL

GRISEOFULVIN

>A>	AB	ACTAVIS MID ATLANTIC	125MG/5ML	N65394 001	Jul 06, 2007	Jun	NEWA
-----	----	----------------------	-----------	------------	--------------	-----	------

GUANABENZ ACETATE

TABLET; ORAL

GUANABENZ ACETATE

>D>	AB	TEVA PHARMS	EQ 4MG BASE	N74267 001	Jun 01, 1994	Jun	DISC
>A>		@	EQ 4MG BASE	N74267 001	Jun 01, 1994	Jun	DISC
>D>	AB		EQ 8MG BASE	N74267 002	Jun 01, 1994	Jun	DISC
>A>		@	EQ 8MG BASE	N74267 002	Jun 01, 1994	Jun	DISC

HALCINONIDE

CREAM; TOPICAL

HALOG

>A>	+	RANBAXY	0.1%	N17556 001		Jun	CAHN
>D>	+	WESTWOOD SQUIBB	0.1%	N17556 001		Jun	CAHN

HALOG-E

>A>	@	RANBAXY	0.1%	N18234 001		Jun	CAHN
>D>	@	WESTWOOD SQUIBB	0.1%	N18234 001		Jun	CAHN

OINTMENT; TOPICAL

HALOG

>A>	+	RANBAXY	0.1%	N17824 001		Jun	CAHN
>D>	+	WESTWOOD SQUIBB	0.1%	N17824 001		Jun	CAHN

SOLUTION; TOPICAL

HALOG

>A>		RANBAXY	0.1%	N17823 001		Jun	CAHN
>D>		WESTWOOD SQUIBB	0.1%	N17823 001		Jun	CAHN

HALOBETASOL PROPIONATE

CREAM; TOPICAL

HALOBETASOL PROPIONATE

	AB	G AND W LABS	0.05%	N78162 001	Apr 24, 2007	Apr	NEWA
--	----	--------------	-------	------------	--------------	-----	------

CREAM; TOPICALULTRAVATE

>A>	AB	+	RANBAXY	0.05%	N19967	001	Dec 27, 1990	Jun	CAHN
>D>	AB	+	WESTWOOD SQUIBB	0.05%	N19967	001	Dec 27, 1990	Jun	CAHN

OINTMENT; TOPICALULTRAVATE

>A>	AB	+	RANBAXY	0.05%	N19968	001	Dec 17, 1990	Jun	CAHN
>D>	AB	+	WESTWOOD SQUIBB	0.05%	N19968	001	Dec 17, 1990	Jun	CAHN

HEPARIN SODIUMINJECTABLE; INJECTIONHEPARIN SODIUM

@	ABRAXIS PHARM		1,000 UNITS/ML	N17979	001			Mar	DISC
@			10,000 UNITS/ML	N17979	002			Mar	DISC
@	WATSON LABS		1,000 UNITS/ML	N17064	002			Feb	CAHN
@			2,500 UNITS/ML	N17064	015			Feb	CAHN
@			3,000 UNITS/ML	N17064	016			Feb	CAHN
@			4,000 UNITS/ML	N17064	017			Feb	CAHN
@			5,000 UNITS/ML	N17064	003			Feb	CAHN
@			6,000 UNITS/ML	N17064	018			Feb	CAHN
@			7,500 UNITS/ML	N17064	019			Feb	CAHN
@			10,000 UNITS/ML	N17064	004			Feb	CAHN
@			20,000 UNITS/ML	N17064	005			Feb	CAHN
@			40,000 UNITS/ML	N17064	006			Feb	CAHN

HISTRELIN ACETATEIMPLANT; SUBCUTANEOUSSUPPRELIN LA

+	INDEVUS		50MG	N22058	001	May 03, 2007	May	NEWA
+	INDEVUS		50MG	N21732	001	Oct 12, 2004	Apr	CAHN

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATESYRUP; ORALHYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE

>D>	AA		IVAX PHARMS	1.5MG/5ML; 5MG/5ML	N40285	001	Jul 19, 1999	Jun	DISC
>A>		@		1.5MG/5ML; 5MG/5ML	N40285	001	Jul 19, 1999	Jun	DISC

HYDRALAZINE HYDROCHLORIDETABLET; ORALHYDRALAZINE HYDROCHLORIDE

@	HERITAGE PHARMS INC		25MG	N86243	001			Feb	CAHN
@			50MG	N86242	002			Feb	CAHN

HYDROCHLOROTHIAZIDETABLET; ORALHYDROCHLOROTHIAZIDE

AB	ACTAVIS ELIZABETH		12.5MG	N40707	001	Feb 27, 2007	Feb	NEWA
AB	CARACO		25MG	N40810	001	Mar 27, 2007	Mar	NEWA
AB			50MG	N40810	002	Mar 27, 2007	Mar	NEWA
AB	EXCELLIUM		25MG	N40702	001	Mar 16, 2007	Mar	NEWA
AB			50MG	N40702	002	Mar 16, 2007	Mar	NEWA
AB	HERITAGE PHARMS INC		25MG	N85181	001		Feb	CAHN
AB			50MG	N85182	001		Feb	CAHN
AB	MYLAN		12.5MG	N40770	001	Jan 23, 2007	Feb	CTEC

TABLET; ORAL

HYDROCHLOROTHIAZIDE

	MYLAN	12.5MG	N40770 001	Jan 23, 2007	Jan	NEWA
AB		25MG	N40735 002	Jan 23, 2007	Jan	NEWA
AB		50MG	N40735 003	Jan 23, 2007	Jan	NEWA

HYDROCHLOROTHIAZIDE; MOEXIPRIL HYDROCHLORIDE

TABLET; ORAL

MOEXIPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

AB	TEVA	12.5MG;7.5MG	N76980 001	Mar 07, 2007	Feb	NEWA
AB		12.5MG;15MG	N76980 003	Mar 07, 2007	Feb	NEWA
AB		25MG;15MG	N76980 002	Mar 07, 2007	Feb	NEWA

UNIRETIC

AB	SCHWARZ PHARMA	12.5MG;7.5MG	N20729 001	Jun 27, 1997	Feb	CFTG
AB		12.5MG;15MG	N20729 003	Feb 14, 2002	Feb	CFTG
AB	+	25MG;15MG	N20729 002	Jun 27, 1997	Feb	CFTG

HYDROCHLOROTHIAZIDE; QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE AND HYDROCHLOROTHIAZIDE

>D>	MYLAN	12.5MG;EQ 10MG BASE	N77093 001	Mar 28, 2005	Jun	DISC
>A>	@	12.5MG;EQ 10MG BASE	N77093 001	Mar 28, 2005	Jun	DISC
>D>	AB	12.5MG;EQ 20MG BASE	N77093 002	Mar 28, 2005	Jun	DISC
>A>	@	12.5MG;EQ 20MG BASE	N77093 002	Mar 28, 2005	Jun	DISC
>D>	AB	25MG;EQ 20MG BASE	N77093 003	Mar 28, 2005	Jun	DISC
>A>	@	25MG;EQ 20MG BASE	N77093 003	Mar 28, 2005	Jun	DISC

HYDROCODONE BITARTRATE; IBUPROFEN

TABLET; ORAL

HYDROCODONE BITARTRATE AND IBUPROFEN

AB	INTERPHARM	5MG;200MG	N76642 002	Mar 18, 2004	Apr	CRLD
----	------------	-----------	------------	--------------	-----	------

HYDROCORTISONE

LOTION; TOPICAL

STIE-CORT

>A>	AT	PERRIGO	1%	N89066 001	Nov 25, 1985	Jun	CAHN
>A>	AT		2.5%	N89074 001	Nov 26, 1985	Jun	CAHN
>D>	AT	STIEFEL	1%	N89066 001	Nov 25, 1985	Jun	CAHN
>D>	AT		2.5%	N89074 001	Nov 26, 1985	Jun	CAHN

TABLET; ORAL

CORTEF

AB	PHARMACIA AND UPJOHN	5MG	N08697 003		Mar	CFTG
AB		10MG	N08697 001		Mar	CFTG
		10MG	N08697 001		Feb	CMFD
AB	+	20MG	N08697 002		Mar	CTEC

HYDROCORTISONE

AB	STIEFEL	5MG	N40646 001	Mar 30, 2007	Mar	NEWA
AB		10MG	N40646 002	Mar 30, 2007	Mar	NEWA
AB		20MG	N40646 003	Mar 30, 2007	Mar	NEWA

HYDROCORTISONE BUTYRATE

LOTION; TOPICAL

LOCROID

+	FERNDALE LABS	0.1%	N22076 001	May 18, 2007	May	NEWA
---	---------------	------	------------	--------------	-----	------

HYDROCORTISONE VALERATE

CREAM; TOPICAL

HYDROCORTISONE VALERATE

>D>	AB	TEVA PHARMS	0.2%	N74489 001	Aug 12, 1998	Jun	DISC
>A>		@	0.2%	N74489 001	Aug 12, 1998	Jun	DISC
WESTCORT							
>A>	AB	+ RANBAXY	0.2%	N17950 001		Jun	CAHN
>D>	AB	+ WESTWOOD SQUIBB	0.2%	N17950 001		Jun	CAHN

OINTMENT; TOPICAL

WESTCORT

>A>	AB	+ RANBAXY	0.2%	N18726 001	Aug 08, 1983	Jun	CAHN
>D>	AB	+ WESTWOOD SQUIBB	0.2%	N18726 001	Aug 08, 1983	Jun	CAHN

HYDROMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROMORPHONE HYDROCHLORIDE

AP		BARR	10MG/ML	N76444 001	Apr 25, 2003	Apr	CAHN
		@ WATSON LABS	10MG/ML	N74317 001	Aug 23, 1995	Feb	DISC

HYDROXYCHLOROQUINE SULFATE

TABLET; ORAL

HYDROXYCHLOROQUINE SULFATE

>A>	AB	IPCA	200MG	N40766 001	Jun 14, 2007	Jun	NEWA
-----	----	------	-------	------------	--------------	-----	------

HYDROXYUREA

CAPSULE; ORAL

HYDROXYUREA

>D>	AB	ROXANE	500MG	N74476 001	Aug 18, 1995	Jun	DISC
>A>		@	500MG	N74476 001	Aug 18, 1995	Jun	DISC

HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HYDROCHLORIDE

		@ HOSPIRA	25MG/ML	N87416 001		Feb	DISC
		@	50MG/ML	N87546 001		Feb	DISC
		@ WATSON LABS	50MG/ML	N85779 001		Feb	DISC

TABLET; ORAL

HYDROXYZINE HYDROCHLORIDE

AB		KVK-TECH INC	10MG	N40786 001	Mar 20, 2007	Mar	NEWA
AB			25MG	N40787 001	Mar 20, 2007	Mar	NEWA
AB			50MG	N40788 001	Mar 20, 2007	Mar	NEWA

IBUPROFEN

TABLET; ORAL

IBUPROFEN

>A>	AB	INTERPHARM	400MG	N78558 001	Jun 18, 2007	Jun	NEWA
>A>	AB		600MG	N78558 002	Jun 18, 2007	Jun	NEWA
>A>	AB		800MG	N78558 003	Jun 18, 2007	Jun	NEWA
>D>	AB	MYLAN	400MG	N70045 001	Sep 24, 1985	Jun	DISC
>A>		@	400MG	N70045 001	Sep 24, 1985	Jun	DISC
>D>	AB		600MG	N70057 001	Sep 24, 1985	Jun	DISC
>A>		@	600MG	N70057 001	Sep 24, 1985	Jun	DISC
>D>	AB		800MG	N71999 001	Dec 03, 1987	Jun	DISC
>A>		@	800MG	N71999 001	Dec 03, 1987	Jun	DISC

IBUPROFEN LYSINEINJECTABLE; INTRAVENOUS
NEOPROFEN

>D>	+	FARMACON IL	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N21903 001	Apr 13, 2006	Jun	CAHN
>A>	+	OVATION PHARMS	EQ 20MG BASE/2ML (EQ 10MG BASE/ML)	N21903 001	Apr 13, 2006	Jun	CAHN

IDARUBICIN HYDROCHLORIDEINJECTABLE; INJECTION
IDARUBICIN HYDROCHLORIDE

AP		BEDFORD LABS	1MG/ML	N65288 001	May 15, 2007	May	NEWA
----	--	--------------	--------	------------	--------------	-----	------

IFOSFAMIDEINJECTABLE; INJECTION
IFSOFAMIDE

+	SICOR PHARMS	1GM/20ML (50MG/ML)	N76657 001	Apr 04, 2007	Mar	NEWA
+		3GM/60ML (50MG/ML)	N76657 002	Apr 04, 2007	Mar	NEWA

ILOPROSTSOLUTION; INHALATION
VENTAVIS

+	ACTELION	10UGM/ML (10UGM/ML)	N21779 002	Dec 08, 2005	Jan	CAHN
+		20UGM/2ML (10UGM/ML)	N21779 001	Dec 29, 2004	Jan	CAHN
+	ACTELION PHARM	10UGM/ML (10UGM/ML)	N21779 002	Dec 08, 2005	Feb	CAHN
+		20UGM/2ML (10UGM/ML)	N21779 001	Dec 29, 2004	Feb	CAHN

IMIQUIMODCREAM; TOPICAL
ALDARA

+	GRACEWAY	5%	N20723 001	Feb 27, 1997	Jan	CAHN
---	----------	----	------------	--------------	-----	------

INDAPAMIDETABLET; ORAL
INDAPAMIDE

>D>	AB	TEVA	1.25MG	N74498 002	Feb 12, 1998	Jun	DISC
>A>		@	1.25MG	N74498 002	Feb 12, 1998	Jun	DISC
>D>	AB		2.5MG	N74498 001	Oct 31, 1996	Jun	DISC
>A>		@	2.5MG	N74498 001	Oct 31, 1996	Jun	DISC

INDIUM IN-111 CHLORIDEINJECTABLE; INJECTION
INDICLOR

+	GE HEALTHCARE	2mCi/0.2ML	N19862 001	Dec 29, 1992	May	CAHN
---	---------------	------------	------------	--------------	-----	------

INDOMETHACINCAPSULE; ORAL
INDOMETHACIN

		@ HERITAGE PHARMS INC	25MG	N18851 001	May 18, 1984	Feb	CAHN
		@	50MG	N18851 002	May 18, 1984	Feb	CAHN
AB		MYLAN	25MG	N18858 001	Apr 20, 1984	Feb	CTEC
		@	50MG	N18858 002	Apr 20, 1984	Mar	DISC
AB	+		50MG	N70624 001	Sep 04, 1985	Mar	CRLD
AB		SANDOZ	25MG	N70673 001	Apr 29, 1987	Feb	CMFD
AB			50MG	N70674 001	Apr 29, 1987	Feb	CMFD

INSULIN GLULISINE RECOMBINANT

INJECTABLE; IV (INFUSION)-SC

APIDRA

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

IRON SUCROSE

INJECTABLE; INTRAVENOUS

VENOFER

@ LUITPOLD

EQ 50MG BASE/2.5ML(EQ 20MG
BASE/ML)

N21135 002 Mar 20, 2005 Feb DISC

@

EQ 75MG BASE/3.75ML(EQ 20MG
BASE/ML)

N21135 003 Mar 29, 2005 Feb DISC

EQ 200MG BASE/10ML(EQ 20MG
BASE/ML)

N21135 004 Feb 09, 2007 Feb NEWA

ISONIAZID

TABLET; ORAL

>D>		LANIAZID				
>D>		LANNETT	50MG	N80140 001		Jun DISC
>A>		@	50MG	N80140 001		Jun DISC
>D>	AA		100MG	N80140 002		Jun DISC
>A>		@	100MG	N80140 002		Jun DISC
>D>	AA		300MG	N89776 001	Jun 13, 1988	Jun DISC
>A>		@	300MG	N89776 001	Jun 13, 1988	Jun DISC

ISOSORBIDE DINITRATE

TABLET; ORAL

ISOSORBIDE DINITRATE

AB		WEST WARD	30MG	N40591 001	Jan 10, 2007	Jan NEWA
----	--	-----------	------	------------	--------------	----------

ISRADIPINE

CAPSULE; ORAL

ISRADIPINE

>D>	AB		ABRIKA PHARMS	2.5MG	N77317 001	Jan 05, 2006	Jun CAHN
>D>	AB	+		5MG	N77317 002	Jan 05, 2006	Jun CAHN
>A>	AB		COBALT LABS INC	2.5MG	N77317 001	Jan 05, 2006	Jun CAHN
>A>	AB	+		5MG	N77317 002	Jan 05, 2006	Jun CAHN

KETOCONAZOLE

>A>			AEROSOL, FOAM; TOPICAL			
>A>			EXTINA			
>A>		+	STIEFEL LABS INC	2%	N21738 001	Jun 12, 2007 Jun NEWA

TABLET; ORAL

KETOCONAZOLE

>D>	AB		AAIPHARMA LLC	200MG	N75341 001	Jul 27, 1999	Jun DISC
>A>		@		200MG	N75341 001	Jul 27, 1999	Jun DISC
>D>	AB		TEVA	200MG	N74971 001	Jun 15, 1999	Jun DISC
>A>		@		200MG	N74971 001	Jun 15, 1999	Jun DISC

KETOPROFEN

CAPSULE; ORAL

KETOPROFEN

>D>	AB		HERITAGE PHARMS INC	25MG	N74014 001	Jan 29, 1993	Jun CTEC
>A>				25MG	N74014 001	Jan 29, 1993	Jun CTEC
	AB			25MG	N74014 001	Jan 29, 1993	Feb CAHN

CAPSULE; ORALKETOPROFEN

AB	HERITAGE PHARMS INC	50MG	N74014 002	Jan 29, 1993	Feb	CAHN
AB		75MG	N74014 003	Jan 29, 1993	Feb	CAHN
>D>	AB	TEVA	25MG	N73515 001	Dec 22, 1992	Jun DISC
>A>	@	25MG	N73515 001	Dec 22, 1992	Jun	DISC

KETOROLAC TROMETHAMINEINJECTABLE; INJECTIONKETOROLAC TROMETHAMINE

	@ AMPHASTAR PHARM	15MG/ML	N76209 001	Jul 21, 2004	Mar	DISC
	@	30MG/ML	N76209 002	Jul 21, 2004	Mar	DISC
	@ APOTEX INC	30MG/ML	N75626 001	Jul 24, 2001	Mar	DISC
	@	30MG/ML	N77201 001	Oct 14, 2005	Mar	DISC
	@ GLAND PHARMA LTD	15MG/ML	N76722 001	Jul 27, 2004	Mar	DISC
	@	30MG/ML	N76722 002	Jul 27, 2004	Mar	DISC
AP	WOCKHARDT	15MG/ML	N77942 001	Mar 27, 2007	Mar	NEWA
AP		30MG/ML	N77942 002	Mar 27, 2007	Mar	NEWA
AP		30MG/ML	N77943 001	Mar 27, 2007	Mar	NEWA

>D> KUNECATECHINS>D> OINTMENT; TOPICAL>D> VEREGEN

>D>	+	MEDIGENE	15%	N21902 001	Oct 31, 2006	Jun	CAIN
-----	---	----------	-----	------------	--------------	-----	------

LABETALOL HYDROCHLORIDETABLET; ORALLABETALOL HYDROCHLORIDE

>D>	AB	TEVA	100MG	N74989 001	Sep 30, 1998	Jun	DISC
>A>	@		100MG	N74989 001	Sep 30, 1998	Jun	DISC
>D>	AB		200MG	N74989 002	Sep 30, 1998	Jun	DISC
>A>	@		200MG	N74989 002	Sep 30, 1998	Jun	DISC
>D>	AB		300MG	N74989 003	Sep 30, 1998	Jun	DISC
>A>	@		300MG	N74989 003	Sep 30, 1998	Jun	DISC

LACTULOSESOLUTION; ORALEVALOSE

>D>	AA	TEVA PHARMS	10GM/15ML	N73497 001	May 28, 1993	Jun	DISC
>A>	@		10GM/15ML	N73497 001	May 28, 1993	Jun	DISC

SOLUTION; ORAL, RECTALHEPTALAC

>D>	AA	TEVA PHARMS	10GM/15ML	N73504 001	May 28, 1993	Jun	DISC
>A>	@		10GM/15ML	N73504 001	May 28, 1993	Jun	DISC

LAPATINIB DITOSYLATETABLET; ORALTYKERB

+	GLAXOSMITHKLINE	EQ 250MG BASE	N22059 001	Mar 13, 2007	Mar	NEWA
+	SMITHKLINE BEECHAM	EQ 250MG BASE	N22059 001	Mar 13, 2007	Apr	CAHN

LEUCOVORIN CALCIUMINJECTABLE; INJECTIONLEUCOVORIN CALCIUM

@	PHARMACHEMIE	EQ 350MG BASE/VIAL	N40262 001	Dec 15, 1999	May	DISC
---	--------------	--------------------	------------	--------------	-----	------

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

@ PHARMACHEMIE USA

EQ 50MG BASE/VIAL

N89628 001 Apr 17, 1997 May DISC

TABLET; ORAL

LEUCOVORIN CALCIUM

@ PHARMACHEMIE

EQ 5MG BASE

N73099 001 Mar 28, 1997 Jan DISC

@

EQ 25MG BASE

N73101 001 Mar 28, 1997 Jan DISC

LEUPROLIDE ACETATE

INJECTABLE; SUBCUTANEOUS

ELIGARD

>A>	+	QLT USA	7.5MG/VIAL	N21343 001	Jan 23, 2002	Jun	CAHN
>A>	+		22.5MG/VIAL	N21379 001	Jul 24, 2002	Jun	CAHN
>A>	+		30MG/VIAL	N21488 001	Feb 13, 2003	Jun	CAHN
>A>	+		45MG/VIAL	N21731 001	Dec 14, 2004	Jun	CAHN
>D>	+	SANOFI AVENTIS US	7.5MG/VIAL	N21343 001	Jan 23, 2002	Jun	CAHN
	+		7.5MG/VIAL	N21343 001	Jan 23, 2002	Jan	CAHN
>D>	+		22.5MG/VIAL	N21379 001	Jul 24, 2002	Jun	CAHN
	+		22.5MG/VIAL	N21379 001	Jul 24, 2002	Jan	CAHN
>D>	+		30MG/VIAL	N21488 001	Feb 13, 2003	Jun	CAHN
	+		30MG/VIAL	N21488 001	Feb 13, 2003	Jan	CAHN
>D>	+		45MG/VIAL	N21731 001	Dec 14, 2004	Jun	CAHN
	+		45MG/VIAL	N21731 001	Dec 14, 2004	Jan	CAHN

LEVOCARNITINE

SOLUTION; ORAL

CARNITOR SF

AA

SIGMA TAU

1GM/10ML

N19257 002 Mar 28, 2007 Mar NEWA

LEVOCETIRIZINE DIHYDROCHLORIDE

TABLET; ORAL

XYZAL

+ UCB INC

5MG

N22064 001 May 25, 2007 May NEWA

LIDOCAINE

PATCH; TOPICAL

LIDOCAINE

+ NOVEN

46.1MG/PATCH

N20575 002 May 21, 1996 Jan CDFR

LIDOCAINE HYDROCHLORIDE

SOLUTION; ORAL

LIDOCAINE HYDROCHLORIDE VISCOUS

AT

VINTAGE

2%

N40708 001 Feb 27, 2007 Feb NEWA

SOLUTION; TOPICAL

LIDOCAINE HYDROCHLORIDE

AT

VINTAGE

4%

N40710 001 Feb 27, 2007 Feb NEWA

LIDOCAINE; TETRACAINE

CREAM; TOPICAL

LIDOCAINE AND TETRACAINE

+ GALDERMA LABS LP

7%;7%

N21717 001 Jun 29, 2006 Apr CAHN

LIOTHYRONINE SODIUMINJECTABLE; INJECTION
TRIOSTAT

AP	+	KING PHARMS	EQ 0.01MG BASE/ML	N20105 001	Dec 31, 1991	Apr	CAHN
----	---	-------------	-------------------	------------	--------------	-----	------

LISDEXAMFETAMINE DIMESYLATE

CAPSULE; ORAL

VYVANSE

		NEW RIVER	30MG	N21977 001	Feb 23, 2007	Feb	NEWA
			50MG	N21977 002	Feb 23, 2007	Feb	NEWA
	+		70MG	N21977 003	Feb 23, 2007	Feb	NEWA
		NEW RIVER PHARMS	30MG	N21977 001	Feb 23, 2007	May	CAHN
			50MG	N21977 002	Feb 23, 2007	May	CAHN
	+		70MG	N21977 003	Feb 23, 2007	May	CAHN

LISINAPRIL

TABLET; ORAL

LISINAPRIL

AB		WOCKHARDT	2.5MG	N78402 001	Apr 19, 2007	Apr	NEWA
AB			5MG	N78402 002	Apr 19, 2007	Apr	NEWA
AB			10MG	N78402 003	Apr 19, 2007	Apr	NEWA
AB			20MG	N78402 004	Apr 19, 2007	Apr	NEWA
AB			30MG	N78402 005	Apr 19, 2007	Apr	NEWA
AB			40MG	N78402 006	Apr 19, 2007	Apr	NEWA

LOPERAMIDE HYDROCHLORIDE

CAPSULE; ORAL

LOPERAMIDE HYDROCHLORIDE

>D>	AB	TEVA	2MG	N73122 001	Aug 30, 1991	Jun	DISC
>A>		@	2MG	N73122 001	Aug 30, 1991	Jun	DISC

LOVASTATIN

TABLET; ORAL

LOVASTATIN

AB		APOTEX INC	10MG	N77748 001	Feb 28, 2007	Feb	NEWA
AB			20MG	N77748 002	Feb 28, 2007	Feb	NEWA
AB			40MG	N77748 003	Feb 28, 2007	Feb	NEWA

TABLET, EXTENDED RELEASE; ORAL

ALTOPREV

		@ ANDRX LABS LLC	10MG	N21316 001	Jun 26, 2002	Feb	DISC
--	--	------------------	------	------------	--------------	-----	------

LOVASTATIN; NIACIN

TABLET, EXTENDED RELEASE; ORAL

ADVICOR

	+	ABBOTT	20MG;500MG	N21249 001	Dec 17, 2001	Mar	CAHN
	+		20MG;750MG	N21249 002	Dec 17, 2001	Mar	CAHN
	+		20MG;1GM	N21249 003	Dec 17, 2001	Mar	CAHN
	+		40MG;1GM	N21249 004	Apr 27, 2006	Mar	CAHN

LUTROPIN ALFA

INJECTABLE; SUBCUTANEOUS

LUVERIS

>A>	+	EMD SERONO	75 IU/VIAL	N21322 001	Oct 08, 2004	Jun	CAHN
>D>	+	SERONO INC	75 IU/VIAL	N21322 001	Oct 08, 2004	Jun	CAHN

MAFENIDE ACETATE

CREAM; TOPICAL

SULFAMYLON

+	UDL LABS	EQ 85MG BASE/GM	N16763 001		May	CAHN
---	----------	-----------------	------------	--	-----	------

FOR SOLUTION; TOPICAL

SULFAMYLON

+	UDL LABS	5%	N19832 003	Jun 05, 1998	May	CAHN
---	----------	----	------------	--------------	-----	------

MAGNESIUM SULFATE

INJECTABLE; INJECTION

MAGNESIUM SULFATE IN PLASTIC CONTAINER

HOSPIRA

2GM/50ML (40MG/ML)

N20309 003	Jan 26, 2007	Jan	NEWA
------------	--------------	-----	------

+		4GM/100ML (40MG/ML)	N20309 001	Jun 24, 1994	Jan	CPOT
---	--	---------------------	------------	--------------	-----	------

+		4GM/50ML (80MG/ML)	N20309 002	Jun 24, 1994	Jan	CPOT
---	--	--------------------	------------	--------------	-----	------

MAPROTILINE HYDROCHLORIDE

TABLET; ORAL

MAPROTILINE HYDROCHLORIDE

>D>	AB	MYLAN	25MG	N72284 001	Oct 03, 1988	Jun	CTEC
>A>			25MG	N72284 001	Oct 03, 1988	Jun	CTEC
>D>	AB	+	50MG	N72285 001	Oct 03, 1988	Jun	CTEC
>A>		+	50MG	N72285 001	Oct 03, 1988	Jun	CTEC
>D>	AB		75MG	N72286 001	Oct 03, 1988	Jun	CTEC
>A>			75MG	N72286 001	Oct 03, 1988	Jun	CTEC
>D>	AB	WATSON LABS	25MG	N72162 001	Jun 01, 1988	Jun	DISC
>A>		@	25MG	N72162 001	Jun 01, 1988	Jun	DISC
>D>	AB		50MG	N72163 001	Jun 01, 1988	Jun	DISC
>A>		@	50MG	N72163 001	Jun 01, 1988	Jun	DISC
>D>	AB		75MG	N72164 001	Jun 01, 1988	Jun	DISC
>A>		@	75MG	N72164 001	Jun 01, 1988	Jun	DISC

MECLIZINE HYDROCHLORIDE

TABLET; ORAL

MECLIZINE HYDROCHLORIDE

>D>	AA	IVAX PHARMS	12.5MG	N84975 001		Jun	DISC
>A>		@	12.5MG	N84975 001		Jun	DISC
>D>	AA		25MG	N84657 001		Jun	DISC
>A>		@	25MG	N84657 001		Jun	DISC

MEDRYSONE

>D> SUSPENSION; OPHTHALMIC

>D> HMS

>D>	+	ALLERGAN	1%	N16624 003		Jun	DISC
-----	---	----------	----	------------	--	-----	------

>A>		@	1%	N16624 003		Jun	DISC
-----	--	---	----	------------	--	-----	------

MEGESTROL ACETATE

TABLET; ORAL

MEGESTROL ACETATE

>D>	AB	TEVA	40MG	N74745 001	Feb 27, 1998	Jun	DISC
-----	----	------	------	------------	--------------	-----	------

>A>		@	40MG	N74745 001	Feb 27, 1998	Jun	DISC
-----	--	---	------	------------	--------------	-----	------

MEPERIDINE HYDROCHLORIDE

INJECTABLE; INJECTION

MEPERIDINE HYDROCHLORIDE PRESERVATIVE FREE

>D>	AP	MALLINCKRODT	10MG/ML	N40163 001	May 12, 1997	Jun	DISC
>A>		@	10MG/ML	N40163 001	May 12, 1997	Jun	DISC

TABLET; ORAL

MEPERIDINE HYDROCHLORIDE

>D>	AA	MUTUAL PHARM	50MG	N80448 001		Jun	DISC
>A>		@	50MG	N80448 001		Jun	DISC
>D>	AA		100MG	N80448 002		Jun	DISC
>A>		@	100MG	N80448 002		Jun	DISC

MEPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION

POLOCAINE

AP		ABRAXIS BIOSCIENCE	1%	N89407 001	Dec 01, 1986	May	CAHN
AP			2%	N89410 001	Dec 01, 1986	May	CAHN

POLOCAINE-MPF

AP		ABRAXIS BIOSCIENCE	1%	N89406 001	Dec 01, 1986	May	CAHN
AP			1.5%	N89408 001	Dec 01, 1986	May	CAHN
AP			2%	N89409 001	Dec 01, 1986	May	CAHN

MERCAPTOPYRINE

TABLET; ORAL

MERCAPTOPYRINE

AB		MYLAN	50MG	N40594 001	Jul 01, 2005	May	CAIN
----	--	-------	------	------------	--------------	-----	------

MESALAMINE

TABLET, DELAYED RELEASE; ORAL

LIALDA

+		SHIRE	1.2GM	N22000 001	Jan 16, 2007	Jan	NEWA
---	--	-------	-------	------------	--------------	-----	------

METAPROTERENOL SULFATE

TABLET; ORAL

METAPROTERENOL SULFATE

>D>	AB	TEVA	10MG	N72519 001	Mar 30, 1990	Jun	DISC
>A>		@	10MG	N72519 001	Mar 30, 1990	Jun	DISC
>D>	AB		20MG	N72520 001	Mar 30, 1990	Jun	DISC
>A>		@	20MG	N72520 001	Mar 30, 1990	Jun	DISC

METFORMIN HYDROCHLORIDE

TABLET; ORAL

METFORMIN HYDROCHLORIDE

>D>	AB	TEVA	500MG	N76328 001	Dec 16, 2002	Jun	DISC
>A>		@	500MG	N76328 001	Dec 16, 2002	Jun	DISC
>D>	AB		850MG	N76328 002	Dec 16, 2002	Jun	DISC
>A>		@	850MG	N76328 002	Dec 16, 2002	Jun	DISC
>D>	AB		1GM	N76328 003	Dec 16, 2002	Jun	DISC
>A>		@	1GM	N76328 003	Dec 16, 2002	Jun	DISC
	AB	TORRENT PHARMS	500MG	N77711 001	Jan 24, 2007	Jan	NEWA
	AB		850MG	N77711 002	Jan 24, 2007	Jan	NEWA
	AB		1GM	N77711 003	Jan 24, 2007	Jan	NEWA

METFORMIN HYDROCHLORIDE; PIOGLITAZONE HYDROCHLORIDE

TABLET; ORAL

ACTOPLUS MET

	TAKEDA GLOBAL	500MG;EQ 15MG BASE	N21842 001	Aug 29, 2005	May	CAIN
+		850MG;EQ 15MG BASE	N21842 002	Aug 29, 2005	May	CAIN

METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE

TABLET; ORAL

JANUMET

	MERCK	500MG;EQ 50MG BASE	N22044 001	Mar 30, 2007	Mar	NEWA
+		1GM;EQ 50MG BASE	N22044 002	Mar 30, 2007	Mar	NEWA

METHADONE HYDROCHLORIDE

CONCENTRATE; ORAL

METHADONE HYDROCHLORIDE

>D>	AA	UDL	10MG/ML	N40088 001	Nov 30, 1994	Jun	CAHN
>A>	AA	VISTAPHARM	10MG/ML	N40088 001	Nov 30, 1994	Jun	CAHN

TABLET; ORAL

METHADONE HYDROCHLORIDE

	@	ROXANE	5MG	N88108 001	Mar 08, 1983	May	DISC
	@		10MG	N88109 001	Mar 08, 1983	May	DISC
	@		40MG	N74081 001	Apr 28, 1995	May	DISC

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE LPF

	@	HOSPIRA	EQ 25MG BASE/ML	N11719 007	Mar 31, 1982	Apr	CAHN
--	---	---------	-----------------	------------	--------------	-----	------

METHOTREXATE PRESERVATIVE FREE

	@	HOSPIRA	EQ 20MG BASE/2ML (10 MG/ML)	N11719 014	Apr 13, 2005	Apr	CAHN
	@		EQ 500MG BASE/20ML (25 MG/ML)	N11719 013	Apr 13, 2005	Apr	CAHN
	@		EQ 2.5GM BASE/100ML (25 MG/ML)	N11719 011	Apr 13, 2005	Apr	CAHN

METHOTREXATE SODIUM

AP	+	ABRAXIS PHARM	EQ 50MG BASE/2ML (25MG/ML)	N40263 001	Feb 26, 1999	Apr	CTNA
AP	+		EQ 250MG BASE/10ML (25MG/ML)	N40263 002	Feb 26, 1999	Apr	CTNA
	@	HOSPIRA	EQ 2.5MG BASE/ML	N11719 004		Apr	CAHN
	@		EQ 20MG BASE/VIAL	N11719 001		Apr	CAHN
	@		EQ 25MG BASE/ML	N11719 005		Apr	CAHN
	@		EQ 50MG BASE/VIAL	N11719 003		Apr	CAHN
AP	+		EQ 50MG BASE/2ML (25 MG/ML)	N11719 010	Dec 15, 2004	Apr	CTNA
	@		EQ 100MG BASE/VIAL	N11719 006		Apr	CAHN

METHOTREXATE SODIUM PRESERVATIVE FREE

AP	+	BEDFORD	EQ 50MG BASE/2ML (25MG/ML)	N89340 001	Sep 16, 1986	Apr	CTNA
AP	+		EQ 250MG BASE/10ML (25MG/ML)	N89343 001	Sep 16, 1986	Apr	CTEC
AP	+	GENERAMEDIX	EQ 50MG BASE/2ML (25MG/ML)	N40767 001	Apr 30, 2007	Apr	NEWA
AP	+		EQ 250MG BASE/10ML (25MG/ML)	N40768 001	Apr 30, 2007	Apr	NEWA
AP	+		EQ 1GM BASE/40ML (25MG/ML)	N40716 001	Apr 30, 2007	Apr	NEWA
	@	HOSPIRA	EQ 1GM BASE/VIAL	N11719 009	Apr 07, 1988	Apr	CAHN
AP	+		EQ 1GM BASE/40ML (25MG/ML)	N11719 012	Apr 13, 2005	Apr	CAHN

METHYCLOTHIAZIDE

TABLET; ORAL

METHYCLOTHIAZIDE

>D>	AB	IVAX PHARMS	2.5MG	N87913 001	Jun 03, 1982	Jun	DISC
>A>	@		2.5MG	N87913 001	Jun 03, 1982	Jun	DISC

METHYLPHENIDATE HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

METADATE CD

BX	UCB INC	30MG	N21259 002	Jun 19, 2003	May	CRLD
----	---------	------	------------	--------------	-----	------

TABLET, EXTENDED RELEASE; ORAL

CONCERTA

	ALZA	18MG	N21121 001	Aug 01, 2000	Apr	CAHN
		27MG	N21121 004	Apr 01, 2002	Apr	CAHN
		36MG	N21121 002	Aug 01, 2000	Apr	CAHN
+		54MG	N21121 003	Dec 08, 2000	Apr	CAHN
	JOHNSON AND JOHNSON	18MG	N21121 001	Aug 01, 2000	Feb	CAHN
		27MG	N21121 004	Apr 01, 2002	Feb	CAHN
		36MG	N21121 002	Aug 01, 2000	Feb	CAHN
+		54MG	N21121 003	Dec 08, 2000	Feb	CAHN

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

AP	BEDFORD LABS	EQ 40MG BASE/VIAL	N40662 001	Feb 21, 2007	Feb	NEWA
AP		EQ 125MG BASE/VIAL	N40641 002	Feb 21, 2007	Feb	NEWA
AP		EQ 500MG BASE/VIAL	N40641 003	Feb 21, 2007	Feb	NEWA
AP		EQ 500MG BASE/VIAL	N40709 001	Feb 21, 2007	Feb	NEWA
AP		EQ 1GM BASE/VIAL	N40641 004	Feb 21, 2007	Feb	NEWA
AP		EQ 1GM BASE/VIAL	N40709 002	Feb 21, 2007	Feb	NEWA

METOLAZONE

TABLET; ORAL

METOLAZONE

	@ ROXANE	10MG	N76482 002	Apr 29, 2004	May	DISC
--	----------	------	------------	--------------	-----	------

METOPROLOL SUCCINATE

TABLET, EXTENDED RELEASE; ORAL

METOPROLOL SUCCINATE

AB	KV PHARM	EQ 100MG TARTRATE	N76640 002	May 18, 2007	May	NEWA
AB		EQ 200MG TARTRATE	N76640 001	May 18, 2007	May	NEWA
AB	SANDOZ	EQ 50MG TARTRATE	N76969 002	May 18, 2007	May	NEWA
	TOPROL-XL					
AB	+ ASTRAZENECA	EQ 50MG TARTRATE	N19962 001	Jan 10, 1992	May	CFTG
AB		EQ 100MG TARTRATE	N19962 002	Jan 10, 1992	May	CFTG
AB	+	EQ 200MG TARTRATE	N19962 003	Jan 10, 1992	May	CFTG

METOPROLOL TARTRATE

INJECTABLE; INJECTION

METOPROLOL TARTRATE

AP	HIKMA FARMACEUTICA	1MG/ML	N77761 001	May 30, 2007	May	NEWA
----	--------------------	--------	------------	--------------	-----	------

TABLET; ORAL

METOPROLOL TARTRATE

>D>	AB	MYLAN	50MG	N73666 001	Dec 21, 1993	Jun	DISC
>A>		@	50MG	N73666 001	Dec 21, 1993	Jun	DISC
>D>	AB		100MG	N73666 002	Dec 21, 1993	Jun	DISC
>A>		@	100MG	N73666 002	Dec 21, 1993	Jun	DISC
>D>	AB	TEVA	50MG	N74143 001	Sep 30, 1994	Jun	DISC
>A>		@	50MG	N74143 001	Sep 30, 1994	Jun	DISC
>D>	AB		100MG	N74143 002	Sep 30, 1994	Jun	DISC

TABLET; ORAL

METOPROLOL TARTRATE

>A>	@	TEVA	100MG	N74143 002	Sep 30, 1994	Jun	DISC
>D>	AB	TEVA PHARMS	50MG	N74333 001	Jan 27, 1994	Jun	DISC
>A>	@		50MG	N74333 001	Jan 27, 1994	Jun	DISC
>D>	AB		100MG	N74333 002	Jan 27, 1994	Jun	DISC
>A>	@		100MG	N74333 002	Jan 27, 1994	Jun	DISC

METRONIDAZOLE

GEL; VAGINAL

METROGEL-VAGINAL

AB	+	GRACEWAY	0.75%	N20208 001	Aug 17, 1992	Jan	CAHN
----	---	----------	-------	------------	--------------	-----	------

MEXILETINE HYDROCHLORIDE

CAPSULE; ORAL

MEXILETINE HYDROCHLORIDE

AB	+	TEVA	250MG	N74377 003	May 16, 1995	Apr	CRLD
		MEXITIL					
	@	BOEHRINGER INGELHEIM	150MG	N18873 002	Dec 30, 1985	Apr	DISC
	@		200MG	N18873 003	Dec 30, 1985	Apr	DISC
	@		250MG	N18873 004	Dec 30, 1985	Apr	DISC

MIDAZOLAM HYDROCHLORIDE

INJECTABLE; INJECTION

MIDAZOLAM HYDROCHLORIDE

AP	+	HOSPIRA	EQ 1MG BASE/ML	N75293 001	Jun 20, 2000	Apr	CRLD
AP		TAYLOR	EQ 1MG BASE/ML	N75494 001	Jun 30, 2000	Feb	CAHN
AP			EQ 5MG BASE/ML	N75494 002	Jun 30, 2000	Feb	CAHN
		MIDAZOLAM HYDROCHLORIDE PRESERVATIVE FREE					
AP	+	HOSPIRA	EQ 1MG BASE/ML	N75857 001	Jul 22, 2002	Feb	CTNA
AP	+		EQ 5MG BASE/ML	N75857 002	Jul 22, 2002	Feb	CTNA

MIRTAZAPINE

TABLET; ORAL

MIRTAZAPINE

>D>	AB	SANDOZ	15MG	N76189 001	Jun 19, 2003	Jun	DISC
>A>	@		15MG	N76189 001	Jun 19, 2003	Jun	DISC
>D>	AB		30MG	N76189 002	Jun 19, 2003	Jun	DISC
>A>	@		30MG	N76189 002	Jun 19, 2003	Jun	DISC
>D>	AB		45MG	N76189 003	Jun 19, 2003	Jun	DISC
>A>	@		45MG	N76189 003	Jun 19, 2003	Jun	DISC

MITOMYCIN

INJECTABLE; INJECTION

MITOZYTREX

	+	SUPERGEN	5MG/VIAL	N50763 001	Nov 14, 2002	Jan	CTNA
--	---	----------	----------	------------	--------------	-----	------

MITOXANTRONE HYDROCHLORIDE

INJECTABLE; INJECTION

NOVANTRONE

AP	+	EMD SERONO	EQ 20MG BASE/10ML (2MG/ML)	N19297 001	Dec 23, 1987	Mar	CAHN
AP	+		EQ 25MG BASE/12.5ML (2MG/ML)	N19297 002	Dec 23, 1987	Mar	CAHN
AP	+		EQ 30MG BASE/15ML (2MG/ML)	N19297 003	Dec 23, 1987	Mar	CAHN

MOMETASONE FUROATE

CREAM; TOPICAL

MOMETASONE FUROATE

AB	TOLMAR	0.1%	N76591 001	Apr 18, 2007	Apr	NEWA
----	--------	------	------------	--------------	-----	------

MORPHINE SULFATE

CAPSULE, EXTENDED RELEASE; ORAL

AVINZA

BX	KING PHARMS	30MG	N21260 001	Mar 20, 2002	Mar	CAHN
----	-------------	------	------------	--------------	-----	------

BX		60MG	N21260 002	Mar 20, 2002	Mar	CAHN
----	--	------	------------	--------------	-----	------

		90MG	N21260 003	Mar 20, 2002	Mar	CAHN
--	--	------	------------	--------------	-----	------

+		120MG	N21260 004	Mar 20, 2002	Mar	CAHN
---	--	-------	------------	--------------	-----	------

KADIAN

ALPHARMA BRANDED

		10MG	N20616 008	Apr 20, 2007	Apr	NEWA
--	--	------	------------	--------------	-----	------

		20MG	N20616 001	Jul 03, 1996	Apr	CAHN
--	--	------	------------	--------------	-----	------

BX		30MG	N20616 004	Mar 09, 2001	Apr	CAHN
----	--	------	------------	--------------	-----	------

		50MG	N20616 002	Jul 03, 1996	Apr	CAHN
--	--	------	------------	--------------	-----	------

BX		60MG	N20616 005	Mar 09, 2001	Apr	CAHN
----	--	------	------------	--------------	-----	------

		80MG	N20616 006	Oct 27, 2006	Apr	CAHN
--	--	------	------------	--------------	-----	------

+		100MG	N20616 003	Jul 03, 1996	Apr	CAHN
---	--	-------	------------	--------------	-----	------

		200MG	N20616 007	Feb 27, 2007	Apr	CAHN
--	--	-------	------------	--------------	-----	------

ALPHARMA US PHARMS

		200MG	N20616 007	Feb 27, 2007	Feb	NEWA
--	--	-------	------------	--------------	-----	------

INJECTABLE; INJECTION

ASTRAMORPH PF

AP	ABRAXIS BIOSCIENCE	0.5MG/ML	N71050 001	Oct 07, 1986	May	CAHN
----	--------------------	----------	------------	--------------	-----	------

AP		0.5MG/ML	N71051 001	Oct 07, 1986	May	CAHN
----	--	----------	------------	--------------	-----	------

AP		1MG/ML	N71052 001	Oct 07, 1986	May	CAHN
----	--	--------	------------	--------------	-----	------

AP		1MG/ML	N71053 001	Oct 07, 1986	May	CAHN
----	--	--------	------------	--------------	-----	------

NADOLOL

TABLET; ORAL

NADOLOL

>D>	AB	TEVA PHARMS	80MG	N74368 001	Aug 31, 1994	Jun	DISC
-----	----	-------------	------	------------	--------------	-----	------

>A>		@	80MG	N74368 001	Aug 31, 1994	Jun	DISC
-----	--	---	------	------------	--------------	-----	------

>D>	AB		120MG	N74368 002	Aug 31, 1994	Jun	DISC
-----	----	--	-------	------------	--------------	-----	------

>A>		@	120MG	N74368 002	Aug 31, 1994	Jun	DISC
-----	--	---	-------	------------	--------------	-----	------

>D>	AB		160MG	N74368 003	Aug 31, 1994	Jun	DISC
-----	----	--	-------	------------	--------------	-----	------

>A>		@	160MG	N74368 003	Aug 31, 1994	Jun	DISC
-----	--	---	-------	------------	--------------	-----	------

NALBUPHINE HYDROCHLORIDE

INJECTABLE; INJECTION

NALBUPHINE HYDROCHLORIDE

AP	BARR	10MG/ML	N74471 001	Mar 19, 1998	Apr	CAHN
----	------	---------	------------	--------------	-----	------

AP		20MG/ML	N74471 002	Mar 19, 1998	Apr	CAHN
----	--	---------	------------	--------------	-----	------

AP	+	HOSPIRA	10MG/ML	N70914 001	Feb 03, 1989	Feb	CRLD
----	---	---------	---------	------------	--------------	-----	------

AP	+		10MG/ML	N70915 001	Feb 03, 1989	Feb	CRLD
----	---	--	---------	------------	--------------	-----	------

AP	+		20MG/ML	N70916 001	Feb 03, 1989	Feb	CRLD
----	---	--	---------	------------	--------------	-----	------

AP	+		20MG/ML	N70918 001	Feb 03, 1989	Feb	CRLD
----	---	--	---------	------------	--------------	-----	------

NUBAIN

@ ENDO PHARMS

		10MG/ML	N18024 001		Feb	DISC
--	--	---------	------------	--	-----	------

@

		20MG/ML	N18024 002	May 27, 1982	Feb	DISC
--	--	---------	------------	--------------	-----	------

NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

>D>		VASOCON							
>D>	AT	NOVARTIS	0.1%	N80235	002	Mar 24,	1983	Jun	DISC
>A>		@	0.1%	N80235	002	Mar 24,	1983	Jun	DISC

NAPROXEN

TABLET; ORAL

NAPROXEN

	AB	GLENMARK PHARMS INC	250MG	N78250	001	Mar 28,	2007	Mar	NEWA
	AB		375MG	N78250	002	Mar 28,	2007	Mar	NEWA
	AB		500MG	N78250	003	Mar 28,	2007	Mar	NEWA
>D>	AB	TEVA	250MG	N74129	001	Dec 21,	1993	Jun	DISC
>A>		@	250MG	N74129	001	Dec 21,	1993	Jun	DISC
>D>	AB		250MG	N74216	001	Apr 11,	1996	Jun	DISC
>A>		@	250MG	N74216	001	Apr 11,	1996	Jun	DISC
>D>	AB		375MG	N74129	002	Dec 21,	1993	Jun	DISC
>A>		@	375MG	N74129	002	Dec 21,	1993	Jun	DISC
>D>	AB		375MG	N74216	002	Apr 11,	1996	Jun	DISC
>A>		@	375MG	N74216	002	Apr 11,	1996	Jun	DISC
>D>	AB		500MG	N74129	003	Dec 21,	1993	Jun	DISC
>A>		@	500MG	N74129	003	Dec 21,	1993	Jun	DISC
>D>	AB		500MG	N74216	003	Apr 11,	1996	Jun	DISC
>A>		@	500MG	N74216	003	Apr 11,	1996	Jun	DISC
>D>	AB	TEVA PHARMS	250MG	N74207	001	Dec 21,	1993	Jun	DISC
>A>		@	250MG	N74207	001	Dec 21,	1993	Jun	DISC
>D>	AB		375MG	N74207	002	Dec 21,	1993	Jun	DISC
>A>		@	375MG	N74207	002	Dec 21,	1993	Jun	DISC
>D>	AB		500MG	N74207	003	Dec 21,	1993	Jun	DISC
>A>		@	500MG	N74207	003	Dec 21,	1993	Jun	DISC
>A>	AB	ZYDUS PHARMS USA	250MG	N78620	001	Jun 07,	2007	Jun	NEWA
>A>	AB		375MG	N78620	002	Jun 07,	2007	Jun	NEWA
>A>	AB		500MG	N78620	003	Jun 07,	2007	Jun	NEWA

NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

	AB	GLENMARK PHARMS	EQ 250MG BASE	N78314	001	Apr 27,	2007	Apr	NEWA
	AB		EQ 500MG BASE	N78314	002	Apr 27,	2007	Apr	NEWA
	AB	INTERPHARM	EQ 250MG BASE	N78432	001	Apr 25,	2007	Apr	NEWA
	AB		EQ 500MG BASE	N78432	002	Apr 25,	2007	Apr	NEWA
>D>	AB	MYLAN	EQ 250MG BASE	N74367	001	Aug 31,	1994	Jun	DISC
>A>		@	EQ 250MG BASE	N74367	001	Aug 31,	1994	Jun	DISC
>D>	AB		EQ 500MG BASE	N74367	002	Aug 31,	1994	Jun	DISC
>A>		@	EQ 500MG BASE	N74367	002	Aug 31,	1994	Jun	DISC
>D>	AB	TEVA	EQ 250MG BASE	N74142	001	Dec 21,	1993	Jun	DISC
>A>		@	EQ 250MG BASE	N74142	001	Dec 21,	1993	Jun	DISC
>D>	AB		EQ 500MG BASE	N74142	002	Dec 21,	1993	Jun	DISC
>A>		@	EQ 500MG BASE	N74142	002	Dec 21,	1993	Jun	DISC
>D>	AB	TEVA PHARMS	EQ 250MG BASE	N74289	001	Jan 27,	1994	Jun	DISC
>A>		@	EQ 250MG BASE	N74289	001	Jan 27,	1994	Jun	DISC
>D>	AB		EQ 500MG BASE	N74289	002	Jan 27,	1994	Jun	DISC
>A>		@	EQ 500MG BASE	N74289	002	Jan 27,	1994	Jun	DISC

NEFAZODONE HYDROCHLORIDE

TABLET; ORAL

NEFAZODONE HYDROCHLORIDE

>D>	AB	MYLAN	100MG	N76129 002	Sep 16, 2003	Jun	DISC
>A>		@	100MG	N76129 002	Sep 16, 2003	Jun	DISC
>D>	AB		150MG	N76129 003	Sep 16, 2003	Jun	DISC
>A>		@	150MG	N76129 003	Sep 16, 2003	Jun	DISC
>D>	AB		200MG	N76129 004	Sep 16, 2003	Jun	DISC
>A>		@	200MG	N76129 004	Sep 16, 2003	Jun	DISC
>D>	AB		250MG	N76129 005	Sep 16, 2003	Jun	DISC
>A>		@	250MG	N76129 005	Sep 16, 2003	Jun	DISC
>D>	AB	SANDOZ	50MG	N76072 001	Sep 16, 2003	Jun	DISC
>A>		@	50MG	N76072 001	Sep 16, 2003	Jun	DISC
>D>	AB		100MG	N76072 002	Sep 16, 2003	Jun	DISC
>A>		@	100MG	N76072 002	Sep 16, 2003	Jun	DISC
>D>	AB		150MG	N76072 003	Sep 16, 2003	Jun	DISC
>A>		@	150MG	N76072 003	Sep 16, 2003	Jun	DISC
>D>	AB		200MG	N76072 004	Sep 16, 2003	Jun	DISC
>A>		@	200MG	N76072 004	Sep 16, 2003	Jun	DISC
>D>	AB		250MG	N76072 005	Sep 16, 2003	Jun	DISC
>A>		@	250MG	N76072 005	Sep 16, 2003	Jun	DISC
>D>	AB	WATSON LABS	100MG	N76073 002	Sep 16, 2003	Jun	DISC
>A>		@	100MG	N76073 002	Sep 16, 2003	Jun	DISC
>D>	AB		150MG	N76073 003	Sep 16, 2003	Jun	DISC
>A>		@	150MG	N76073 003	Sep 16, 2003	Jun	DISC
>D>	AB		200MG	N76073 004	Sep 16, 2003	Jun	DISC
>A>		@	200MG	N76073 004	Sep 16, 2003	Jun	DISC
>D>	AB		250MG	N76073 005	Sep 16, 2003	Jun	DISC
>A>		@	250MG	N76073 005	Sep 16, 2003	Jun	DISC

NELFINAVIR MESYLATE

POWDER; ORAL

VIRACEPT

+	AGOURON	EQ 50MG BASE/SCOOPFUL	N20778 001	Mar 14, 1997	Mar	CDFR
---	---------	-----------------------	------------	--------------	-----	------

NEOMYCIN SULFATE

TABLET; ORAL

NEOMYCIN SULFATE

AA	+	TEVA	500MG	N60304 001		Apr	CTEC
AA		X GEN PHARMS	500MG	N65220 001	Jul 28, 2006	Apr	CTEC

NESIRITIDE RECOMBINANT

FOR SOLUTION; INTRAVENOUS

NATRECOR

+	ALZA CORP	1.5MG/VIAL	N20920 001	Aug 10, 2001	Jan	CAHN
+	SCIOS	1.5MG/VIAL	N20920 001	Aug 10, 2001	Feb	CAHN

NIACIN

TABLET, EXTENDED RELEASE; ORAL

NIASPAN

	@	ABBOTT	375MG	N20381 001	Jul 28, 1997	Mar	CAHN
			500MG	N20381 002	Jul 28, 1997	Apr	CRLD
+			500MG	N20381 002	Jul 28, 1997	Mar	CAHN
+			750MG	N20381 003	Jul 28, 1997	Mar	CAHN
+			1GM	N20381 004	Jul 28, 1997	Mar	CAHN

TABLET, EXTENDED RELEASE; ORAL

NIASPAN TITRATION STARTER PACK

@	ABBOTT	375MG;500MG;750MG	N20381	005	Jul 28, 1997	Mar	CAHN
---	--------	-------------------	--------	-----	--------------	-----	------

NIFEDIPINE

CAPSULE; ORAL

NIFEDIPINE

>D>	AB	TEVA	10MG	N72651	001	Feb 19, 1992	Jun	DISC
>A>		@	10MG	N72651	001	Feb 19, 1992	Jun	DISC

NIMODIPINE

CAPSULE; ORAL

NIMODIPINE

AB	BARR	30MG	N77811	001	May 02, 2007	Apr	NEWA
AB	SUN PHARM INDS INC	30MG	N77067	001	Apr 17, 2007	Apr	NEWA
AB	+ BAYER PHARMS	30MG	N18869	001	Dec 28, 1988	Apr	CFTG

NITROGLYCERIN

FILM, EXTENDED RELEASE; TRANSDERMAL

MINITRAN

AB1	GRACEWAY	0.1MG/HR	N89771	001	Aug 30, 1996	Jan	CAHN
AB1		0.2MG/HR	N89772	001	Aug 30, 1996	Jan	CAHN
AB1		0.4MG/HR	N89773	001	Aug 30, 1996	Jan	CAHN
AB1		0.6MG/HR	N89774	001	Aug 30, 1996	Jan	CAHN

NITROGLYCERIN

>D>	AB1	MYLAN TECHNOLOGIES	0.1MG/HR	N74992	004	Nov 12, 1999	Jun	DISC
>A>		@	0.1MG/HR	N74992	004	Nov 12, 1999	Jun	DISC
>D>	AB1		0.2MG/HR	N74992	003	Nov 12, 1999	Jun	DISC
>A>		@	0.2MG/HR	N74992	003	Nov 12, 1999	Jun	DISC
>D>	AB1		0.4MG/HR	N74992	002	Nov 12, 1999	Jun	DISC
>A>		@	0.4MG/HR	N74992	002	Nov 12, 1999	Jun	DISC
>D>	AB1		0.6MG/HR	N74992	001	Nov 12, 1999	Jun	DISC
>A>		@	0.6MG/HR	N74992	001	Nov 12, 1999	Jun	DISC

NORTRIPTYLINE HYDROCHLORIDE

CAPSULE; ORAL

NORTRIPTYLINE HYDROCHLORIDE

>D>	AB	SANDOZ	EQ 10MG BASE	N74054	001	Dec 31, 1992	Jun	DISC
>A>		@	EQ 10MG BASE	N74054	001	Dec 31, 1992	Jun	DISC
>D>	AB		EQ 25MG BASE	N74054	002	Dec 31, 1992	Jun	DISC
>A>		@	EQ 25MG BASE	N74054	002	Dec 31, 1992	Jun	DISC
>D>	AB		EQ 50MG BASE	N74054	003	Dec 31, 1992	Jun	DISC
>A>		@	EQ 50MG BASE	N74054	003	Dec 31, 1992	Jun	DISC
>D>	AB		EQ 75MG BASE	N74054	004	Dec 31, 1992	Jun	DISC
>A>		@	EQ 75MG BASE	N74054	004	Dec 31, 1992	Jun	DISC
>D>	AB	TEVA	EQ 10MG BASE	N73667	001	Apr 11, 1996	Jun	DISC
>A>		@	EQ 10MG BASE	N73667	001	Apr 11, 1996	Jun	DISC
>D>	AB		EQ 25MG BASE	N73667	002	Apr 11, 1996	Jun	DISC
>A>		@	EQ 25MG BASE	N73667	002	Apr 11, 1996	Jun	DISC
>D>	AB		EQ 50MG BASE	N73667	003	Apr 11, 1996	Jun	DISC
>A>		@	EQ 50MG BASE	N73667	003	Apr 11, 1996	Jun	DISC
>D>	AB		EQ 75MG BASE	N73667	004	Apr 11, 1996	Jun	DISC
>A>		@	EQ 75MG BASE	N73667	004	Apr 11, 1996	Jun	DISC

NYSTATIN

CREAM; TOPICAL

MYCOSTATIN

>A>	AT	+	RANBAXY	100,000 UNITS/GM	N60575 001		Jun	CAHN
>D>	AT	+	WESTWOOD SQUIBB	100,000 UNITS/GM	N60575 001		Jun	CAHN

POWDER; TOPICAL

MYCOSTATIN

>A>	AT	+	RANBAXY	100,000 UNITS/GM	N60578 001		Jun	CAHN
>D>	AT	+	WESTWOOD SQUIBB	100,000 UNITS/GM	N60578 001		Jun	CAHN

NYSTATIN

>A>	AT		COASTAL PHARMS	100,000 UNITS/GM	N65203 001	Jul 15, 2004	Jun	CAHN
>D>	AT		PHARMAFORCE	100,000 UNITS/GM	N65203 001	Jul 15, 2004	Jun	CAHN

ONDANSETRON

TABLET, ORALLY DISINTEGRATING; ORAL

ONDANSETRON

>A>	AB		BARR	4MG	N76693 001	Jun 25, 2007	Jun	NEWA
>A>	AB			8MG	N76693 002	Jun 25, 2007	Jun	NEWA
>A>	AB		GLENMARK PHARMS INC	4MG	N78152 001	Jun 27, 2007	Jun	NEWA
>A>	AB			8MG	N78152 002	Jun 27, 2007	Jun	NEWA
>A>	AB		KV PHARM	4MG	N77717 001	Jun 25, 2007	Jun	NEWA
>A>	AB			8MG	N77717 002	Jun 25, 2007	Jun	NEWA
>A>	AB		MYLAN	4MG	N78139 001	Jun 25, 2007	Jun	NEWA
>A>	AB			8MG	N78139 002	Jun 25, 2007	Jun	NEWA
>A>	AB		TEVA	4MG	N76810 001	Jun 25, 2007	Jun	NEWA
>A>	AB			8MG	N76810 002	Jun 25, 2007	Jun	NEWA

ONDANSETRON HYDROCHLORIDE

INJECTABLE; INJECTION

ONDANSETRON HYDROCHLORIDE

AP			HOSPIRA	EQ 2MG BASE/ML	N77840 001	Jan 19, 2007	Jan	NEWA
>A>	AP		PHARMAFORCE	EQ 2MG BASE/ML	N77582 001	Dec 26, 2006	Jun	CAHN
>D>	AP		PLIVA	EQ 2MG BASE/ML	N77582 001	Dec 26, 2006	Jun	CAHN
	AP			EQ 2MG BASE/ML	N77582 001	Dec 26, 2006	Mar	CAHN
>A>	AP		SANDOZ	EQ 2MG BASE/ML	N77430 001	Jun 27, 2007	Jun	NEWA
	AP		SPECTRUM PHARMS	EQ 2MG BASE/ML	N78180 001	Mar 26, 2007	Mar	NEWA

ONDANSETRON HYDROCHLORIDE AND DEXTROSE IN PLASTIC CONTAINER

AP			HOSPIRA	EQ 0.64MG BASE/ML	N77348 001	Feb 01, 2007	Jan	NEWA
AP			MAYNE PHARMA USA	EQ 0.64MG BASE/ML	N76978 001	Feb 26, 2007	Feb	NEWA

ONDANSETRON HYDROCHLORIDE PRESERVATIVE FREE

>A>	AP		PHARMAFORCE	EQ 2MG BASE/ML	N77387 001	Dec 26, 2006	Jun	CAHN
>D>	AP		PLIVA	EQ 2MG BASE/ML	N77387 001	Dec 26, 2006	Jun	CAHN
	AP			EQ 2MG BASE/ML	N77387 001	Dec 26, 2006	Mar	CAHN
>A>	AP		SANDOZ	EQ 2MG BASE/ML	N77551 001	Jun 27, 2007	Jun	NEWA

ZOFRAN AND DEXTROSE IN PLASTIC CONTAINER

AP	+		GLAXOSMITHKLINE	EQ 0.64MG BASE/ML	N20403 001	Jan 31, 1995	Jan	CTNA
----	---	--	-----------------	-------------------	------------	--------------	-----	------

SOLUTION; ORAL

ONDANSETRON HYDROCHLORIDE

>A>	AA		APOTEX	EQ 4MG BASE/5ML	N78127 001	Jun 25, 2007	Jun	NEWA
-----	----	--	--------	-----------------	------------	--------------	-----	------

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

>A>	AB		APOTEX	EQ 4MG BASE	N77306 001	Jun 25, 2007	Jun	NEWA
>A>	AB			EQ 8MG BASE	N77306 002	Jun 25, 2007	Jun	NEWA
>A>	AB		GLENMARK PHARMS	EQ 4MG BASE	N77535 001	Jun 25, 2007	Jun	NEWA
>A>	AB			EQ 8MG BASE	N77535 002	Jun 25, 2007	Jun	NEWA

TABLET; ORAL

ONDANSETRON HYDROCHLORIDE

>A>	AB	GLENMARK PHARMS	EQ 24MG BASE	N77535 003	Jun 25, 2007	Jun	NEWA
>A>	AB	KALI LABS	EQ 4MG BASE	N77303 001	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 8MG BASE	N77303 002	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 24MG BASE	N77303 004	Jun 25, 2007	Jun	NEWA
>A>	AB	MYLAN	EQ 4MG BASE	N76930 001	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 8MG BASE	N76930 002	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 24MG BASE	N76930 004	Jun 25, 2007	Jun	NEWA
>A>	AB	NATCO PHARMA LTD	EQ 4MG BASE	N77851 001	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 8MG BASE	N77851 002	Jun 25, 2007	Jun	NEWA
>A>	AB	PLIVA HRVATSKA DOO	EQ 4MG BASE	N77112 001	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 8MG BASE	N77112 002	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 24MG BASE	N77112 003	Jun 25, 2007	Jun	NEWA
>A>	AB	SANDOZ	EQ 4MG BASE	N77517 001	Jun 25, 2007	Jun	NEWA
>A>	AA		EQ 8MG BASE	N77517 002	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 24MG BASE	N77517 003	Jun 25, 2007	Jun	NEWA
>A>	AB	SUN PHARM INDS (IN)	EQ 4MG BASE	N77050 001	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 8MG BASE	N77050 002	Jun 25, 2007	Jun	NEWA
>A>	AB	TEVA	EQ 4MG BASE	N76252 001	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 8MG BASE	N76252 002	Jun 25, 2007	Jun	NEWA
>A>	AB		EQ 24MG BASE	N76252 003	Jun 25, 2007	Jun	NEWA

ORPHENADRINE CITRATE

INJECTABLE; INJECTION

NORFLEX

AP	+	GRACEWAY	30MG/ML	N13055 001		Jan	CAHN
----	---	----------	---------	------------	--	-----	------

TABLET, EXTENDED RELEASE; ORAL

NORFLEX

@ GRACEWAY

100MG

N12157 001

Jan CAHN

ORPHENADRINE CITRATE

AB		ACTAVIS TOTOWA	100MG	N40284 001	Jun 19, 1998	Mar	CAHN
----	--	----------------	-------	------------	--------------	-----	------

OXACILLIN SODIUM

CAPSULE; ORAL

BACTOCILL

>D>	AB	GLAXOSMITHKLINE	EQ 250MG BASE	N61336 001		Jun	CTEC
>A>			EQ 250MG BASE	N61336 001		Jun	CTEC
>D>	AB	+	EQ 500MG BASE	N61336 002		Jun	CTEC
>A>		+	EQ 500MG BASE	N61336 002		Jun	CTEC
>D>		OXACILLIN SODIUM					
>D>	AB	TEVA	EQ 250MG BASE	N62222 001		Jun	DISC
>A>		@	EQ 250MG BASE	N62222 001		Jun	DISC
>D>	AB		EQ 500MG BASE	N62222 002		Jun	DISC
>A>		@	EQ 500MG BASE	N62222 002		Jun	DISC

FOR SOLUTION; ORAL

BACTOCILL

>D>	AA	GLAXOSMITHKLINE	EQ 250MG BASE/5ML	N62321 001		Jun	CRLD
>A>		+	EQ 250MG BASE/5ML	N62321 001		Jun	CRLD
>D>		OXACILLIN SODIUM					
>D>	AA	TEVA	EQ 250MG BASE/5ML	N62252 001		Jun	DISC
>A>		@	EQ 250MG BASE/5ML	N62252 001		Jun	DISC

OXANDROLONE

TABLET; ORAL

OXANDROLONE

>A>	AB	KALI LABS	2.5MG	N77827 001	Jun 22, 2007	Jun	NEWA
>A>	AB		10MG	N77827 002	Jun 22, 2007	Jun	NEWA
>A>	AB	ROXANE	2.5MG	N77249 001	Jul 10, 2007	Jun	NEWA
>A>	AB		10MG	N77249 002	Jul 10, 2007	Jun	NEWA
	AB	UPSHER SMITH	10MG	N78033 001	Mar 22, 2007	Mar	NEWA

OXAPROZIN

TABLET; ORAL

OXAPROZIN

>D>	AB	MYLAN	600MG	N75851 001	Aug 17, 2001	Jun	DISC
>A>		@	600MG	N75851 001	Aug 17, 2001	Jun	DISC

OXYBUTYNIN CHLORIDE

TABLET, EXTENDED RELEASE; ORAL

OXYBUTYNIN CHLORIDE

	AB	IMPAX PHARMS	5MG	N76745 002	May 09, 2007	May	NEWA
	AB		10MG	N76745 003	May 09, 2007	May	NEWA
	AB	MYLAN	15MG	N78293 001	May 10, 2007	May	NEWA

OXYCODONE HYDROCHLORIDE

TABLET; ORAL

OXYCODONE HYDROCHLORIDE

	AB	KV PHARM	5MG	N77290 001	Dec 08, 2005	Mar	CTEC
	AB	TYCO HLTHCARE	5MG	N78206 001	Mar 19, 2007	Mar	NEWA
	AB	VINTAGE PHARMS	15MG	N77712 001	Jan 31, 2007	Jan	NEWA
	AB		30MG	N77712 002	Jan 31, 2007	Jan	NEWA

ROXICODONE

	AB	+ XANODYNE PHARMS INC	15MG	N21011 001	Aug 31, 2000	Feb	CAHN
	AB		30MG	N21011 002	Aug 31, 2000	Feb	CAHN

TABLET, EXTENDED RELEASE; ORAL

OXYCONTIN

@ PURDUE PHARMA LP

@

@

			15MG	N20553 006	Sep 18, 2006	May	DISC
			30MG	N20553 007	Sep 18, 2006	May	DISC
			60MG	N20553 008	Sep 18, 2006	May	DISC

OXYMORPHONE HYDROCHLORIDE

INJECTABLE; INJECTION

NUMORPHAN

>D>							
>D>		+ ENDO PHARMS	1MG/ML	N11707 002		Jun	CTNA
>D>		@	1.5MG/ML	N11707 001		Jun	CTNA
>A>		OPANA					
>A>		+ ENDO PHARMS	1MG/ML	N11707 002		Jun	CTNA
>A>		@	1.5MG/ML	N11707 001		Jun	CTNA

PAROXETINE HYDROCHLORIDE

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

>A>	AB	CARACO	EQ 10MG BASE	N78194 001	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 20MG BASE	N78194 002	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 30MG BASE	N78194 003	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 40MG BASE	N78194 004	Jun 29, 2007	Jun	NEWA

TABLET; ORAL

PAROXETINE HYDROCHLORIDE

>A>	AB	ROXANE	EQ 10MG BASE	N78026 001	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 20MG BASE	N78026 002	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 30MG BASE	N78026 003	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 40MG BASE	N78026 004	Jun 29, 2007	Jun	NEWA
>D>	AB	SANDOZ	EQ 10MG BASE	N75566 001	Mar 08, 2004	Jun	DISC
>A>	@		EQ 10MG BASE	N75566 001	Mar 08, 2004	Jun	DISC
>D>	AB		EQ 20MG BASE	N75566 002	Mar 08, 2004	Jun	DISC
>A>	@		EQ 20MG BASE	N75566 002	Mar 08, 2004	Jun	DISC
>D>	AB		EQ 30MG BASE	N75566 003	Mar 08, 2004	Jun	DISC
>A>	@		EQ 30MG BASE	N75566 003	Mar 08, 2004	Jun	DISC
>D>	AB		EQ 40MG BASE	N75566 004	Mar 08, 2004	Jun	DISC
>A>	@		EQ 40MG BASE	N75566 004	Mar 08, 2004	Jun	DISC
>A>	AB	TEVA PHARMS	EQ 10MG BASE	N77082 001	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 20MG BASE	N77082 002	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 30MG BASE	N77082 003	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 40MG BASE	N77082 004	Jun 29, 2007	Jun	NEWA
	AB	ZYDUS PHARMS USA	EQ 10MG BASE	N77584 001	Mar 07, 2007	Feb	NEWA
	AB		EQ 20MG BASE	N77584 002	Mar 07, 2007	Feb	NEWA
	AB		EQ 30MG BASE	N77584 003	Mar 07, 2007	Feb	NEWA
	AB		EQ 40MG BASE	N77584 004	Mar 07, 2007	Feb	NEWA

TABLET, EXTENDED RELEASE; ORAL

PAROXETINE HYDROCHLORIDE

>A>	AB	MYLAN	EQ 12.5MG BASE	N77873 001	Jun 29, 2007	Jun	NEWA
>A>	AB		EQ 25MG BASE	N77873 002	Jun 29, 2007	Jun	NEWA
		PAXIL CR					
>D>		GLAXOSMITHKLINE	EQ 12.5MG BASE	N20936 001	Feb 16, 1999	Jun	CFTG
>A>	AB		EQ 12.5MG BASE	N20936 001	Feb 16, 1999	Jun	CFTG
>D>			EQ 25MG BASE	N20936 002	Feb 16, 1999	Jun	CFTG
>A>	AB		EQ 25MG BASE	N20936 002	Feb 16, 1999	Jun	CFTG

PEGADEMASE BOVINE

INJECTABLE; INJECTION

ADAGEN

>D>	+	ENZON	250 UNITS/ML	N19818 001	Mar 21, 1990	Jun	CAHN
>A>	+	ENZON PHARMS	250 UNITS/ML	N19818 001	Mar 21, 1990	Jun	CAHN

PENTOSTATIN

INJECTABLE; INJECTION

NIPENT

	+	HOSPIRA INC	10MG/VIAL	N20122 001	Oct 11, 1991	Mar	CAHN
--	---	-------------	-----------	------------	--------------	-----	------

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE

	AB	HERITAGE PHARMS INC	400MG	N74877 001	Jul 08, 1997	Feb	CAHN
>D>	AB	TEVA	400MG	N75199 001	Sep 03, 1999	Jun	DISC
>A>	@		400MG	N75199 001	Sep 03, 1999	Jun	DISC

PERGOLIDE MESYLATE

TABLET; ORAL

PERGOLIDE MESYLATE

	@	IVAX PHARMS	EQ 0.05MG BASE	N76094 001	Sep 04, 2003	Mar	DISC
	@		EQ 0.25MG BASE	N76094 002	Sep 04, 2003	Mar	DISC
	@		EQ 1MG BASE	N76094 003	Sep 04, 2003	Mar	DISC

TABLET; ORAL

PERGOLIDE MESYLATE

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

PERPHENAZINE

TABLET; ORAL

PERPHENAZINE

>D>	AB	SANDOZ	2MG	N89683 001	Dec 08, 1988	Jun	DISC
>A>		@	2MG	N89683 001	Dec 08, 1988	Jun	DISC
>D>	AB		4MG	N89684 001	Dec 08, 1988	Jun	DISC
>A>		@	4MG	N89684 001	Dec 08, 1988	Jun	DISC
>D>	AB		8MG	N89685 001	Dec 08, 1988	Jun	DISC
>A>		@	8MG	N89685 001	Dec 08, 1988	Jun	DISC
>D>	AB		16MG	N89686 001	Dec 08, 1988	Jun	DISC
>A>		@	16MG	N89686 001	Dec 08, 1988	Jun	DISC

PHEENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

PHEENTERMINE HYDROCHLORIDE

>D>	AA	MUTUAL PHARM	37.5MG	N40527 001	Oct 23, 2003	Jun	DISC	
>A>		@	37.5MG	N40527 001	Oct 23, 2003	Jun	DISC	
	AA	+	SANDOZ	30MG	N87190 001	Feb	CRLD	
	AA		TG UNITED INC	30MG	N40083 001	Mar 07, 1997	Feb	CAHN

>D> PHEENTERMINE RESIN COMPLEX

>D> CAPSULE, EXTENDED RELEASE; ORAL

>D> IONAMIN

>D>		UCB INC	EQ 15MG BASE	N11613 004		Jun	DISC
>A>		@	EQ 15MG BASE	N11613 004		Jun	DISC
>D>		@	EQ 30MG BASE	N11613 002		Jun	DISC
>A>		@	EQ 30MG BASE	N11613 002		Jun	DISC

PILOCARPINE HYDROCHLORIDE

TABLET; ORAL

PILOCARPINE HYDROCHLORIDE

AB		ROXANE	7.5MG	N76963 002	Feb 27, 2007	Feb	NEWA
----	--	--------	-------	------------	--------------	-----	------

PINDOLOL

TABLET; ORAL

PINDOLOL

>D>	AB	GENPHARM	5MG	N74013 001	Sep 24, 1992	Jun	DISC
>A>		@	5MG	N74013 001	Sep 24, 1992	Jun	DISC
>D>	AB		10MG	N74018 001	Sep 24, 1992	Jun	DISC
>A>		@	10MG	N74018 001	Sep 24, 1992	Jun	DISC
>D>	AB	MUTUAL PHARM	5MG	N74063 001	Jan 27, 1994	Jun	DISC
>A>		@	5MG	N74063 001	Jan 27, 1994	Jun	DISC

TABLET; ORALPINDOLOL

>D>	AB	MUTUAL PHARM	10MG	N74063 002	Jan 27, 1994	Jun	DISC
>A>		@	10MG	N74063 002	Jan 27, 1994	Jun	DISC
>D>	AB	TEVA	5MG	N73661 001	Oct 31, 1993	Jun	DISC
>A>		@	5MG	N73661 001	Oct 31, 1993	Jun	DISC
>D>	AB		5MG	N74123 001	Apr 17, 1997	Jun	DISC
>A>		@	5MG	N74123 001	Apr 17, 1997	Jun	DISC
>D>	AB		10MG	N73661 002	Oct 31, 1993	Jun	DISC
>A>		@	10MG	N73661 002	Oct 31, 1993	Jun	DISC
>D>	AB		10MG	N74123 002	Apr 17, 1997	Jun	DISC
>A>		@	10MG	N74123 002	Apr 17, 1997	Jun	DISC

PIPERACILLIN SODIUMINJECTABLE; INJECTIONPIPERACILLIN

+	ISTITUTO BIOCHIMICO	EQ 2GM BASE/VIAL	N65114 001	Nov 14, 2003	Feb	CAHN
+		EQ 3GM BASE/VIAL	N65114 002	Nov 14, 2003	Feb	CAHN
+		EQ 4GM BASE/VIAL	N65114 003	Nov 14, 2003	Feb	CAHN
+		EQ 40GM BASE/VIAL	N65157 001	Jul 12, 2004	Feb	CAHN

PIRBUTEROL ACETATEAEROSOL, METERED; INHALATIONMAXAIR

	@ GRACEWAY	EQ 0.2MG BASE/INH	N19009 001	Dec 30, 1986	Jan	CAHN
+		EQ 0.2MG BASE/INH	N20014 001	Nov 30, 1992	Jan	CAHN

PIROXICAMCAPSULE; ORALPIROXICAM

>D>	AB	GENPHARM	10MG	N74043 001	Sep 22, 1992	Jun	DISC
>A>		@	10MG	N74043 001	Sep 22, 1992	Jun	DISC
>D>	AB		20MG	N74043 002	Sep 22, 1992	Jun	DISC
>A>		@	20MG	N74043 002	Sep 22, 1992	Jun	DISC
>D>	AB	TEVA	10MG	N73637 001	Jan 28, 1994	Jun	DISC
>A>		@	10MG	N73637 001	Jan 28, 1994	Jun	DISC
>D>	AB		20MG	N73638 001	Jan 28, 1994	Jun	DISC
>A>		@	20MG	N73638 001	Jan 28, 1994	Jun	DISC
>D>	AB	TEVA PHARMS	10MG	N74103 001	Aug 28, 1992	Jun	DISC
>A>		@	10MG	N74103 001	Aug 28, 1992	Jun	DISC
>D>	AB		20MG	N74103 002	Aug 28, 1992	Jun	DISC
>A>		@	20MG	N74103 002	Aug 28, 1992	Jun	DISC

POLYETHYLENE GLYCOL 3350FOR SOLUTION; ORALPOLYETHYLENE GLYCOL 3350

AA	ANABOLIC LABS	17GM/SCOOPFUL	N77706 001	Sep 27, 2006	Jan	CAHN
AA	PADDOCK	17GM/SCOOPFUL	N77893 001	May 26, 2006	May	CAHN

POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM SULFATE, ANHYDROUSFOR SUSPENSION; ORALGO-EVAC

AA	BOCA PHARMA	236GM/BOT; 2.97GM/BOT; 6.74GM/BOT; 5.86GM/BOT; 22.74GM/BOT	N73433 001	Apr 28, 1992	May	CAHN
----	-------------	--	------------	--------------	-----	------

>D> POTASSIUM AMINOSALICYLATE

>D> CAPSULE; ORAL

>D> PASKALIUM

>D> + GLENWOOD 500MG N09395 004 Jun DISC

>A> @ 500MG N09395 004 Jun DISC

>D> TABLET; ORAL

>D> PASKALIUM

>D> + GLENWOOD 1GM N09395 003 Jun DISC

>A> @ 1GM N09395 003 Jun DISC

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MICRO-K

>D> AB KV PHARM 8MEQ N18238 001 Jun CTEC

>A> 8MEQ N18238 001 Jun CTEC

POTASSIUM CHLORIDE

>D> AB TEVA 8MEQ N73531 001 Apr 26, 1996 Jun DISC

>A> @ 8MEQ N73531 001 Apr 26, 1996 Jun DISC

>D> AB 10MEQ N73532 001 Apr 26, 1996 Jun DISC

>A> @ 10MEQ N73532 001 Apr 26, 1996 Jun DISC

PRAVASTATIN SODIUM

TABLET; ORAL

PRAVACHOL

AB + BRISTOL MYERS SQUIBB 80MG N19898 008 Dec 18, 2001 Apr CFTG

PRAVASTATIN SODIUM

AB GLENMARK PHARMS 10MG N77987 001 May 11, 2007 May NEWA

AB 20MG N77987 002 May 11, 2007 May NEWA

AB 40MG N77987 003 May 11, 2007 May NEWA

AB RANBAXY 10MG N76445 001 Apr 23, 2007 Apr NEWA

AB 20MG N76445 002 Apr 23, 2007 Apr NEWA

AB 40MG N76445 003 Apr 23, 2007 Apr NEWA

AB 80MG N76445 004 Apr 23, 2007 Apr NEWA

PREDNICARBATE

CREAM; TOPICAL

DERMATOP E EMOLLIENT

>D> AB + SANOFI AVENTIS US 0.1% N20279 001 Oct 29, 1993 Jun DISC

>A> @ 0.1% N20279 001 Oct 29, 1993 Jun DISC

PREDNICARBATE

>D> AB ALTANA 0.1% N77287 001 Sep 19, 2006 Jun CRLD

>A> + 0.1% N77287 001 Sep 19, 2006 Jun CRLD

OINTMENT; TOPICAL

DERMATOP

>D> AB + SANOFI AVENTIS US 0.1% N19568 001 Sep 23, 1991 Jun DISC

>A> @ 0.1% N19568 001 Sep 23, 1991 Jun DISC

AB + 0.1% N19568 001 Sep 23, 1991 Feb CFTG

PREDNICARBATE

>D> AB ALTANA 0.1% N77236 001 Mar 09, 2007 Jun CRLD

>A> + 0.1% N77236 001 Mar 09, 2007 Jun CRLD

AB 0.1% N77236 001 Mar 09, 2007 Feb NEWA

PREDNISOLONE

SYRUP; ORAL

PREDNISOLONE

>D>	AA	TEVA PHARMS	15MG/5ML	N40322 001	Jan 19, 2000	Jun	DISC
>A>		@	15MG/5ML	N40322 001	Jan 19, 2000	Jun	DISC

TABLET; ORAL

PREDNISOLONE

>D>	BX	LANNETT	5MG	N80531 002		Jun	DISC
>A>		@	5MG	N80531 002		Jun	DISC
>D>	BX	+ WATSON LABS	5MG	N80354 001		Jun	CTEC
>A>		+	5MG	N80354 001		Jun	CTEC

PREDNISOLONE SODIUM PHOSPHATE

SOLUTION/DROPS; OPHTHALMIC

INFLAMASE FORTE

>D>	AT	+ NOVARTIS	EQ 0.9% PHOSPHATE	N80751 002		Jun	DISC
>A>		@	EQ 0.9% PHOSPHATE	N80751 002		Jun	DISC
>D>		INFLAMASE MILD					
>D>	AT	+ NOVARTIS	EQ 0.11% PHOSPHATE	N80751 001		Jun	DISC
>A>		@	EQ 0.11% PHOSPHATE	N80751 001		Jun	DISC

PREDNISOLONE SODIUM PHOSPHATE

>D>	AT	BAUSCH AND LOMB	EQ 0.11% PHOSPHATE	N40065 001	Jul 29, 1994	Jun	CTEC
>A>		+	EQ 0.11% PHOSPHATE	N40065 001	Jul 29, 1994	Jun	CTEC
>D>	AT		EQ 0.9% PHOSPHATE	N40070 001	Jul 29, 1994	Jun	CRLD
>A>		+	EQ 0.9% PHOSPHATE	N40070 001	Jul 29, 1994	Jun	CRLD

PROCAINAMIDE HYDROCHLORIDE

CAPSULE; ORAL

PROCAINAMIDE HYDROCHLORIDE

		@ IVAX PHARMS	250MG	N84604 001		Jan	DISC
		@	375MG	N84595 001		Jan	DISC
		@	500MG	N84606 001		Jan	DISC
		@ WATSON LABS	250MG	N83287 001		Jan	DISC
		@	375MG	N84403 001		Jan	DISC
		@	500MG	N84280 001		Jan	DISC
		PRONESTYL					
		@ APOTHECON	250MG	N07335 001		Jan	DISC
		@	375MG	N07335 004		Jan	DISC
		@	500MG	N07335 003		Jan	DISC

TABLET, EXTENDED RELEASE; ORAL

PRONESTYL-SR

		@ APOTHECON	500MG	N87361 001		Jan	DISC
--	--	-------------	-------	------------	--	-----	------

PROCHLORPERAZINE

SUPPOSITORY; RECTAL

COMPAZINE

		@ GLAXOSMITHKLINE	25MG	N11127 002		Feb	DISC
AB	+	G AND W LABS	25MG	N40058 001	Nov 24, 1993	Feb	CRLD

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

COMPAZINE

		@ GLAXOSMITHKLINE	EQ 5MG BASE/ML	N10742 002		Feb	DISC
--	--	-------------------	----------------	------------	--	-----	------

INJECTABLE; INJECTION

	PROCHLORPERAZINE								
	@ BAXTER HLTHCARE	EQ 5MG BASE/ML	N87759	001	Oct 01, 1982	Feb	DISC		
	PROCHLORPERAZINE EDISYLATE								
AP	+ BAXTER HLTHCARE	EQ 5MG BASE/ML	N89903	001	Aug 29, 1989	Feb	CRLD		
	@ HOSPIRA	EQ 5MG BASE/ML	N89703	001	Apr 07, 1988	Feb	DISC		
	@ WATSON LABS	EQ 5MG BASE/ML	N89530	001	Jul 08, 1987	Feb	DISC		

PROCHLORPERAZINE MALEATE

	CAPSULE, EXTENDED RELEASE; ORAL								
	COMPAZINE								
	@ GLAXOSMITHKLINE	EQ 15MG BASE	N21019	002	Oct 06, 1999	Jan	DISC		
	TABLET; ORAL								
	COMPAZINE								
	@ GLAXOSMITHKLINE	EQ 5MG BASE	N10571	001		Feb	DISC		
	@	EQ 10MG BASE	N10571	002		Feb	DISC		
	@	EQ 25MG BASE	N10571	003		Feb	DISC		
	PROCHLORPERAZINE MALEATE								
AB	+ SANDOZ	EQ 10MG BASE	N40101	002	Jul 19, 1996	Feb	CRLD		
	@	EQ 25MG BASE	N40101	003	Jul 19, 1996	Feb	DISC		

PROGESTERONE

>A>	INSERT; VAGINAL								
>A>	ENDOMETRIN								
>A>	+ FERRING	100MG	N22057	001	Jun 21, 2007	Jun	NEWA		

PROMETHAZINE HYDROCHLORIDE

	SYRUP; ORAL								
	PROMETHAZINE HYDROCHLORIDE								
AA	TARO	6.25MG/5ML	N40718	001	Apr 04, 2007	Mar	NEWA		
	TABLET; ORAL								
	PROMETHAZINE HYDROCHLORIDE								
AB	KVK-TECH INC	12.5MG	N40712	002	May 04, 2007	Apr	NEWA		

PROPOXYPHENE HYDROCHLORIDE

	CAPSULE; ORAL								
	DOLENE								
	@ HERITAGE PHARMS INC	65MG	N80530	001		Feb	CAHN		

PROPRANOLOL HYDROCHLORIDE

	CAPSULE, EXTENDED RELEASE; ORAL								
	INDERAL LA								
AB	WYETH PHARMS INC	60MG	N18553	004	Mar 18, 1987	Jan	CTEC		
AB		80MG	N18553	002	Apr 19, 1983	Jan	CTEC		
AB		120MG	N18553	003	Apr 19, 1983	Jan	CTEC		
AB	+	160MG	N18553	001	Apr 19, 1983	Jan	CTEC		
	PROPRANOLOL HYDROCHLORIDE								
AB	MYLAN	60MG	N78022	001	Feb 15, 2007	Feb	NEWA		
AB		80MG	N78022	002	Feb 15, 2007	Feb	NEWA		
AB		120MG	N78022	003	Feb 15, 2007	Feb	NEWA		
AB		160MG	N78022	004	Feb 15, 2007	Feb	NEWA		
AB	PAR PHARM	60MG	N78065	001	Jan 26, 2007	Jan	NEWA		
AB		80MG	N78065	002	Jan 26, 2007	Jan	NEWA		
AB		120MG	N78065	003	Jan 26, 2007	Jan	NEWA		
AB		160MG	N78065	004	Jan 26, 2007	Jan	NEWA		

CONCENTRATE; ORAL

PROPRANOLOL HYDROCHLORIDE INTENSOL

@ ROXANE 80MG/ML

N71388 001 May 15, 1987 May DISC

TABLET; ORAL

INDERAL

@ WYETH PHARMS INC 10MG

N16418 001 Jan DISC

@ 20MG

N16418 003 Jan DISC

PROPRANOLOL HYDROCHLORIDE

>D> AB TEVA 40MG

N70234 001 Jun 23, 1986 Jun DISC

>A> @ 40MG

N70234 001 Jun 23, 1986 Jun DISC

PYRIDOXINE HYDROCHLORIDE

INJECTABLE; INJECTION

PYRIDOXINE HYDROCHLORIDE

+ ABRAXIS PHARM 100MG/ML

N80618 001 Jan CRLD

@ WATSON LABS 100MG/ML

N80572 001 Jan DISC

QUETIAPINE FUMARATE

TABLET, EXTENDED RELEASE; ORAL

SEROQUEL XR

ASTRAZENECA 50MG

N22047 001 May 17, 2007 May NEWA

200MG

N22047 002 May 17, 2007 May NEWA

300MG

N22047 003 May 17, 2007 May NEWA

+ 400MG

N22047 004 May 17, 2007 May NEWA

QUINAPRIL HYDROCHLORIDE

TABLET; ORAL

QUINAPRIL HYDROCHLORIDE

>D> AB MYLAN EQ 5MG BASE

N76694 001 Dec 23, 2004 Jun DISC

>A> @ EQ 5MG BASE

N76694 001 Dec 23, 2004 Jun DISC

>D> AB EQ 10MG BASE

N76694 002 Dec 23, 2004 Jun DISC

>A> @ EQ 10MG BASE

N76694 002 Dec 23, 2004 Jun DISC

>D> AB EQ 20MG BASE

N76694 003 Dec 23, 2004 Jun DISC

>A> @ EQ 20MG BASE

N76694 003 Dec 23, 2004 Jun DISC

>D> AB EQ 40MG BASE

N76694 004 Dec 23, 2004 Jun DISC

>A> @ EQ 40MG BASE

N76694 004 Dec 23, 2004 Jun DISC

QUINIDINE SULFATE

TABLET; ORAL

QUINIDINE SULFATE

@ WATSON LABS 200MG

N83288 001 Mar DISC

@ 300MG

N85583 001 Apr DISC

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE; ORAL

ACIPHEX

AB + EISAI MEDCL RES 20MG

N20973 002 Aug 19, 1999 Feb CFTG

RABEPRAZOLE SODIUM

AB TEVA 20MG

N76822 001 Feb 21, 2007 Feb NEWA

RAMIPRIL

CAPSULE; ORAL

ALTACE

KING PHARMS 1.25MG

N19901 001 Jan 28, 1991 Jan CTEC

2.5MG

N19901 002 Jan 28, 1991 Jan CTEC

CAPSULE; ORAL

ALTACE

	KING PHARMS	5MG	N19901 003	Jan 28, 1991	Jan	CTEC
+		10MG	N19901 004	Jan 28, 1991	Jan	CTEC
	RAMIPRIL					
	@ COBALT	1.25MG	N76549 001	Oct 24, 2005	Jan	DISC
	@	2.5MG	N76549 002	Oct 24, 2005	Jan	DISC
	@	5MG	N76549 003	Oct 24, 2005	Jan	DISC
	@	10MG	N76549 004	Oct 24, 2005	Jan	DISC

TABLET; ORAL

ALTACE

	COBALT	1.25MG	N22021 001	Feb 27, 2007	Feb	NEWA
		2.5MG	N22021 002	Feb 27, 2007	Feb	NEWA
		5MG	N22021 003	Feb 27, 2007	Feb	NEWA
+		10MG	N22021 004	Feb 27, 2007	Feb	NEWA
	KING PHARMS	1.25MG	N22021 001	Feb 27, 2007	Apr	CAHN
		2.5MG	N22021 002	Feb 27, 2007	Apr	CAHN
		5MG	N22021 003	Feb 27, 2007	Apr	CAHN
+		10MG	N22021 004	Feb 27, 2007	Apr	CAHN

RANITIDINE HYDROCHLORIDE

CAPSULE; ORAL

RANITIDINE HYDROCHLORIDE

>D>	AB	GENPHARM	EQ 150MG BASE	N75564 001	Oct 27, 2000	Jun	DISC
>A>		@	EQ 150MG BASE	N75564 001	Oct 27, 2000	Jun	DISC
>D>	AB		EQ 300MG BASE	N75564 002	Oct 27, 2000	Jun	DISC
>A>		@	EQ 300MG BASE	N75564 002	Oct 27, 2000	Jun	DISC
>D>	AB	TEVA	EQ 150MG BASE	N75557 001	Oct 31, 2003	Jun	DISC
>A>		@	EQ 150MG BASE	N75557 001	Oct 31, 2003	Jun	DISC
>D>	AB		EQ 300MG BASE	N75557 002	Oct 31, 2003	Jun	DISC
>A>		@	EQ 300MG BASE	N75557 002	Oct 31, 2003	Jun	DISC

SYRUP; ORAL

RANITIDINE HYDROCHLORIDE

AA		ALPHARMA US PHARMS	EQ 15MG BASE/ML	N76124 001	Feb 21, 2007	Feb	NEWA
		ZANTAC					
AA	+	GLAXOSMITHKLINE	EQ 15MG BASE/ML	N19675 001	Dec 30, 1988	Feb	CFTG

TABLET; ORAL

RANITIDINE HYDROCHLORIDE

>D>	AB	MYLAN	EQ 150MG BASE	N74552 001	Jul 30, 1998	Jun	DISC
>A>		@	EQ 150MG BASE	N74552 001	Jul 30, 1998	Jun	DISC
>D>	AB		EQ 300MG BASE	N74552 002	Jul 30, 1998	Jun	DISC
>A>		@	EQ 300MG BASE	N74552 002	Jul 30, 1998	Jun	DISC

RANOLAZINE

TABLET, EXTENDED RELEASE; ORAL

RANEXA

>D>	+	CV THERAP	500MG	N21526 002	Jan 27, 2006	Jun	CRLD
>A>			500MG	N21526 002	Jan 27, 2006	Jun	CRLD
>A>	+		1GM	N21526 001	Feb 12, 2007	Jun	NEWA

RETAPAMULIN

OINTMENT; TOPICAL

ALTABAX

+	GLAXO GRP LTD	1%	N22055 001	Apr 12, 2007	Apr	NEWA
---	---------------	----	------------	--------------	-----	------

RIBAVIRIN

TABLET; ORAL

RIBAVIRIN

AB	THREE RIVERS PHARMS	400MG	N77456 002	Dec 05, 2005	Mar	CTEC
AB	+	600MG	N77456 003	Dec 05, 2005	Mar	CTEC
AB	ZYDUS PHARMS USA	400MG	N77094 002	Mar 16, 2007	Mar	NEWA
AB		600MG	N77094 003	Mar 16, 2007	Mar	NEWA

RISEDRONATE SODIUM

TABLET; ORAL

ACTONEL

+	PROCTER AND GAMBLE	75MG	N20835 004	Apr 16, 2007	Apr	NEWA
---	--------------------	------	------------	--------------	-----	------

RISPERIDONE

INJECTABLE; INTRAMUSCULAR

RISPERDAL CONSTA

	JANSSEN PHARMA	12.5MG/VIAL	N21346 004	Apr 12, 2007	Apr	NEWA
--	----------------	-------------	------------	--------------	-----	------

ROTIGOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL

NEUPRO

	SCHWARZ BIOSCIENCES	2MG/24HR	N21829 001	May 09, 2007	May	NEWA
		4MG/24HR	N21829 002	May 09, 2007	May	NEWA
+		6MG/24HR	N21829 003	May 09, 2007	May	NEWA

SECRETIN SYNTHETIC HUMAN

FOR SOLUTION; INTRAVENOUS

>A>	CHIRHOSTIM 40					
>A>	CHIRHOCLIN	40UGM/VIAL	N21256 002	Jun 21, 2007	Jun	NEWA

SELEGILINE HYDROCHLORIDE

TABLET; ORAL

SELEGILINE HYDROCHLORIDE

>D>	AB	IVAX PHARMS	5MG	N74756 001	Nov 25, 1998	Jun	DISC
>A>		@	5MG	N74756 001	Nov 25, 1998	Jun	DISC
>D>	AB	TEVA	5MG	N74537 001	Aug 02, 1996	Jun	DISC
>A>		@	5MG	N74537 001	Aug 02, 1996	Jun	DISC
>D>	AB		5MG	N74744 001	Jan 27, 1997	Jun	DISC
>A>		@	5MG	N74744 001	Jan 27, 1997	Jun	DISC

SERMORELIN ACETATE

INJECTABLE; INJECTION

GEREF

+	EMD SERONO	EQ 0.05MG BASE/AMP	N19863 001	Dec 28, 1990	Feb	CAHN
	@	EQ 0.5MG BASE/VIAL	N20443 001	Sep 26, 1997	Feb	CAHN
	@	EQ 1MG BASE/VIAL	N20443 002	Sep 26, 1997	Feb	CAHN

SERTRALINE HYDROCHLORIDE

CONCENTRATE; ORAL

SERTRALINE HYDROCHLORIDE

AA	RANBAXY	EQ 20MG BASE/ML	N78053 001	Feb 05, 2007	Feb	CTEC
AB		EQ 20MG BASE/ML	N78053 001	Feb 05, 2007	Jan	NEWA
AA	ROXANE	EQ 20MG BASE/ML	N76934 001	Jun 30, 2006	Feb	CTEC

CONCENTRATE; ORAL

ZOLOFT

AA	+	PFIZER	EQ 20MG BASE/ML	N20990 001	Dec 07, 1999	Feb	CTEC
----	---	--------	-----------------	------------	--------------	-----	------

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

AB		ACTAVIS ELIZABETH	EQ 25MG BASE	N77345 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77345 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77345 003	Feb 06, 2007	Jan	NEWA
AB		APOTEX INC	EQ 25MG BASE	N76882 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N76882 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N76882 003	Feb 06, 2007	Jan	NEWA
AB		AUROBINDO PHARMA	EQ 25MG BASE	N77206 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77206 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77206 003	Feb 06, 2007	Jan	NEWA
AB		COBALT	EQ 25MG BASE	N77663 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77663 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77663 003	Feb 06, 2007	Jan	NEWA
AB		DR REDDYS LABS LTD	EQ 25MG BASE	N76442 001	Apr 30, 2007	Apr	NEWA
AB			EQ 50MG BASE	N76442 002	Apr 30, 2007	Apr	NEWA
AB			EQ 100MG BASE	N76442 003	Apr 30, 2007	Apr	NEWA
AB		GENPHARM	EQ 25MG BASE	N76540 001	Mar 20, 2007	Mar	NEWA
AB			EQ 50MG BASE	N76540 002	Mar 20, 2007	Mar	NEWA
AB			EQ 100MG BASE	N76540 003	Mar 20, 2007	Mar	NEWA
AB		INVAGEN PHARMS	EQ 25MG BASE	N77397 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77397 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77397 003	Feb 06, 2007	Jan	NEWA
AB		LUPIN	EQ 25MG BASE	N77670 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77670 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77670 003	Feb 06, 2007	Jan	NEWA
AB		MUTUAL PHARM	EQ 25MG BASE	N77818 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77818 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77818 003	Feb 06, 2007	Jan	NEWA
AB		MYLAN	EQ 25MG BASE	N76671 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N76671 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N76671 003	Feb 06, 2007	Jan	NEWA
AB		PLIVA HRVATSKA DOO	EQ 25MG BASE	N77299 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77299 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77299 003	Feb 06, 2007	Jan	NEWA
AB		RANBAXY	EQ 25MG BASE	N77977 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77977 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77977 003	Feb 06, 2007	Jan	NEWA
			EQ 150MG BASE	N77977 004	Feb 06, 2007	Jan	NEWA
			EQ 200MG BASE	N77977 005	Feb 06, 2007	Jan	NEWA
AB		ROXANE	EQ 25MG BASE	N76881 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N76881 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N76881 003	Feb 06, 2007	Jan	NEWA
AB		SANDOZ	EQ 25MG BASE	N77713 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77713 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77713 003	Feb 06, 2007	Jan	NEWA
AB		SUN PHARM INDS (IN)	EQ 25MG BASE	N78108 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N78108 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N78108 003	Feb 06, 2007	Jan	NEWA
AB		TORRENT PHARMS	EQ 25MG BASE	N77765 001	Feb 06, 2007	Jan	NEWA
AB			EQ 50MG BASE	N77765 002	Feb 06, 2007	Jan	NEWA
AB			EQ 100MG BASE	N77765 003	Feb 06, 2007	Jan	NEWA
AB		WATSON LABS	EQ 25MG BASE	N77162 001	Feb 06, 2007	Jan	NEWA

TABLET; ORAL

SERTRALINE HYDROCHLORIDE

AB	WATSON LABS	EQ 50MG BASE	N77162 002	Feb 06, 2007	Jan	NEWA
AB		EQ 100MG BASE	N77162 003	Feb 06, 2007	Jan	NEWA
AB	ZYDUS PHARMS USA	EQ 25MG BASE	N77106 001	Feb 06, 2007	Jan	NEWA
AB		EQ 50MG BASE	N77106 002	Feb 06, 2007	Jan	NEWA
AB		EQ 100MG BASE	N77106 003	Feb 06, 2007	Jan	NEWA

SEVOFLURANE

LIQUID; INHALATION

SOJOURN

AN	MINRAD	100%	N77867 001	May 02, 2007	Apr	NEWA
----	--------	------	------------	--------------	-----	------

SIMVASTATIN

TABLET; ORAL

SIMVASTATIN

AB	LUPIN	10MG	N78103 001	May 11, 2007	May	NEWA
AB		20MG	N78103 002	May 11, 2007	May	NEWA
AB		40MG	N78103 003	May 11, 2007	May	NEWA
AB		80MG	N78103 004	May 11, 2007	May	NEWA

>A> SINECATECHINS

>A> OINTMENT; TOPICAL

>A> VEREGEN

>A>	+ MEDIGENE	15%	N21902 001	Oct 31, 2006	Jun	CAIN
-----	------------	-----	------------	--------------	-----	------

SODIUM CHLORIDE

INJECTABLE; INJECTION

SODIUM CHLORIDE 0.9% IN PLASTIC CONTAINER

AP	FRESENIUS MEDCL	900MG/100ML	N78177 001	Apr 12, 2007	Apr	NEWA
----	-----------------	-------------	------------	--------------	-----	------

SODIUM LACTATE

INJECTABLE; INJECTION

SODIUM LACTATE IN PLASTIC CONTAINER

	+ HOSPIRA	5MEQ/ML	N18947 001	Sep 05, 1984	Feb	CRLD
--	-----------	---------	------------	--------------	-----	------

SOMATROPIN RECOMBINANT

INJECTABLE; INJECTION

SAIZEN

BX	EMD SERONO	4MG/VIAL	N19764 005	Jan 16, 2007	Feb	CAHN
BX		5MG/VIAL	N19764 002	Oct 08, 1996	Feb	CAHN
	@	6MG/VIAL	N19764 001	Oct 08, 1996	Feb	CAHN
	+ SERONO	8.8MG/VIAL	N19764 003	Aug 29, 2000	Feb	CAHN
BX	SERONO	4MG/VIAL	N19764 005	Jan 16, 2007	Jan	NEWA

SEROSTIM

BX	EMD SERONO	4MG/VIAL	N20604 003	Jul 25, 1997	Feb	CAHN
BX		5MG/VIAL	N20604 002	Aug 23, 1996	Feb	CAHN
BX		6MG/VIAL	N20604 001	Aug 23, 1996	Feb	CAHN
	@	8.8MG/VIAL	N20604 004	Sep 06, 2001	Feb	CAHN

VALTROPIN

BX	LG LIFE	5MG/VIAL	N21905 001	Apr 19, 2007	Apr	NEWA
----	---------	----------	------------	--------------	-----	------

ZORBTIVE

>A>	@ EMD SERONO	4MG/VIAL	N21597 001	Dec 01, 2003	Jun	CAHN
>A>	@	5MG/VIAL	N21597 002	Dec 01, 2003	Jun	CAHN
>A>	@	6MG/VIAL	N21597 003	Dec 01, 2003	Jun	CAHN

INJECTABLE; INJECTION

ZORBTIVE

>A>	+	EMD SERONO	8.8MG/VIAL	N21597 004	Dec 01, 2003	Jun	CAHN
>D>		@ SERONO INC	4MG/VIAL	N21597 001	Dec 01, 2003	Jun	CAHN
>D>		@	5MG/VIAL	N21597 002	Dec 01, 2003	Jun	CAHN
>D>		@	6MG/VIAL	N21597 003	Dec 01, 2003	Jun	CAHN
>D>	+		8.8MG/VIAL	N21597 004	Dec 01, 2003	Jun	CAHN

INJECTABLE; SUBCUTANEOUS

SEROSTIM LQ

@ EMD SERONO

6MG/0.5ML

N20604 005 Feb 11, 2005 Feb CAHN

SOTALOL HYDROCHLORIDE

TABLET; ORAL

SOTALOL HYDROCHLORIDE

AB2		MYLAN	80MG	N77616 001	Feb 07, 2007	Jan	NEWA
AB2			120MG	N77616 002	Feb 07, 2007	Jan	NEWA
AB2			160MG	N77616 003	Feb 07, 2007	Jan	NEWA

SPIRONOLACTONE

TABLET; ORAL

SPIRONOLACTONE

>D>	AB	MYLAN	25MG	N87086 001		Jun	DISC
>A>		@	25MG	N87086 001		Jun	DISC

STREPTOZOCIN

INJECTABLE; INJECTION

ZANOSAR

>D>	+	SICOR PHARMS	1GM/VIAL	N50577 001	May 07, 1982	Jun	CAHN
>A>	+	TEVA PARENTERAL	1GM/VIAL	N50577 001	May 07, 1982	Jun	CAHN

SUFENTANIL CITRATE

INJECTABLE; INJECTION

SUFENTA

>D>	AP	+	AKORN	EQ 0.05MG BASE/ML	N19050 001	May 04, 1984	Jun	CTNA
>A>			SUFENTA PRESERVATIVE FREE					
>A>	AP	+	AKORN	EQ 0.05MG BASE/ML	N19050 001	May 04, 1984	Jun	CTNA

SULCONAZOLE NITRATE

CREAM; TOPICAL

EXELDERM

>A>	+	RANBAXY	1%	N18737 001	Feb 28, 1989	Jun	CAHN
>D>	+	WESTWOOD SQUIBB	1%	N18737 001	Feb 28, 1989	Jun	CAHN

SOLUTION; TOPICAL

EXELDERM

>A>	+	RANBAXY	1%	N18738 001	Aug 30, 1985	Jun	CAHN
>D>	+	WESTWOOD SQUIBB	1%	N18738 001	Aug 30, 1985	Jun	CAHN

SULFACETAMIDE SODIUM

LOTION; TOPICAL

KLARON

>D>								
>D>	AB	+	SANOFI AVENTIS US	10%	N19931 001	Dec 23, 1996	Jun	DISC
>A>		@		10%	N19931 001	Dec 23, 1996	Jun	DISC

SULFACETAMIDE SODIUM

>D>	AB		ALTANA	10%	N77015 001	Nov 17, 2006	Jun	CRLD
>A>		+		10%	N77015 001	Nov 17, 2006	Jun	CRLD

SULFAMETHOXAZOLE; TRIMETHOPRIM

INJECTABLE; INJECTION

BACTRIM

@	MUTUAL PHARM	80MG/ML;16MG/ML	N18374 001		Feb	DISC
---	--------------	-----------------	------------	--	-----	------

SUSPENSION; ORAL

BACTRIM PEDIATRIC

@	MUTUAL PHARM	200MG/5ML;40MG/5ML	N17560 002		Feb	DISC
---	--------------	--------------------	------------	--	-----	------

SEPTRA

@	MONARCH PHARMS	200MG/5ML;40MG/5ML	N17598 001		Feb	DISC
---	----------------	--------------------	------------	--	-----	------

SEPTRA GRAPE

@	MONARCH PHARMS	200MG/5ML;40MG/5ML	N17598 002	Feb 12, 1986	Feb	DISC
---	----------------	--------------------	------------	--------------	-----	------

SULFAMETHOXAZOLE AND TRIMETHOPRIM

@	TEVA	200MG/5ML;40MG/5ML	N18812 001	Jan 28, 1983	Feb	DISC
---	------	--------------------	------------	--------------	-----	------

@		200MG/5ML;40MG/5ML	N18812 002	Jun 10, 1983	Feb	DISC
---	--	--------------------	------------	--------------	-----	------

AB	+	TEVA PHARMS	200MG/5ML;40MG/5ML	N77612 001	Nov 13, 2006	Feb	CRLD
----	---	-------------	--------------------	------------	--------------	-----	------

AB		VINTAGE	200MG/5ML;40MG/5ML	N77785 001	Jan 24, 2007	Jan	NEWA
----	--	---------	--------------------	------------	--------------	-----	------

SULFATRIM

@	ACTAVIS MID ATLANTIC	200MG/5ML;40MG/5ML	N18615 002	Jan 07, 1983	Mar	DISC
---	----------------------	--------------------	------------	--------------	-----	------

TABLET; ORAL

>D> COTRIM

>D>	AB	TEVA	400MG;80MG	N70034 001	May 16, 1985	Jun	DISC
-----	----	------	------------	------------	--------------	-----	------

>A>		@	400MG;80MG	N70034 001	May 16, 1985	Jun	DISC
-----	--	---	------------	------------	--------------	-----	------

>D> COTRIM D.S.

>D>	AB	TEVA	800MG;160MG	N70048 001	Mar 18, 1985	Jun	DISC
-----	----	------	-------------	------------	--------------	-----	------

>A>		@	800MG;160MG	N70048 001	Mar 18, 1985	Jun	DISC
-----	--	---	-------------	------------	--------------	-----	------

SULFAMETHOXAZOLE AND TRIMETHOPRIM

AB		VINTAGE	400MG;80MG	N78060 002	Jan 25, 2007	Jan	NEWA
----	--	---------	------------	------------	--------------	-----	------

AB			800MG;160MG	N78060 001	Jan 25, 2007	Jan	NEWA
----	--	--	-------------	------------	--------------	-----	------

SULFANILAMIDE

CREAM; VAGINAL

AVC

+	AZUR PHARMA	15%	N06530 003	Jan 27, 1987	Apr	CAHN
---	-------------	-----	------------	--------------	-----	------

SUPPOSITORY; VAGINAL

AVC

@	AZUR PHARMA	1.05GM	N06530 004	Jan 27, 1987	Apr	CAHN
---	-------------	--------	------------	--------------	-----	------

SULFASALAZINE

TABLET; ORAL

SULFASALAZINE

@	HERITAGE PHARMS INC	500MG	N80197 001		Feb	CAHN
---	---------------------	-------	------------	--	-----	------

SULFINPYRAZONE

CAPSULE; ORAL

SULFINPYRAZONE

>D>	AB	IVAX PHARMS	200MG	N87770 001	Nov 19, 1982	Jun	DISC
-----	----	-------------	-------	------------	--------------	-----	------

>A>		@	200MG	N87770 001	Nov 19, 1982	Jun	DISC
-----	--	---	-------	------------	--------------	-----	------

SULINDAC

TABLET; ORAL

SULINDAC

@	HERITAGE PHARMS INC	150MG	N73262 002	Sep 06, 1991	Feb	CAHN
---	---------------------	-------	------------	--------------	-----	------

@		200MG	N73262 001	Sep 06, 1991	Feb	CAHN
---	--	-------	------------	--------------	-----	------

>D>	AB	TEVA	150MG	N72972 001	Feb 28, 1992	Jun	DISC
-----	----	------	-------	------------	--------------	-----	------

TABLET; ORAL

SULINDAC

>A>	@	TEVA	150MG	N72972 001	Feb 28, 1992	Jun	DISC
>D>	AB		200MG	N72973 001	Feb 28, 1992	Jun	DISC
>A>	@		200MG	N72973 001	Feb 28, 1992	Jun	DISC

TADALAFIL

TABLET; ORAL

CIALIS

LILLY

			5MG	N21368 001	Nov 21, 2003	Apr	CAHN
			10MG	N21368 002	Nov 21, 2003	Apr	CAHN
	+		20MG	N21368 003	Nov 21, 2003	Apr	CAHN

TAMOXIFEN CITRATE

TABLET; ORAL

NOLVADEX

@ ASTRAZENECA

EQ 10MG BASE

N17970 001

Mar DISC

@

EQ 20MG BASE

N17970 002

Mar 21, 1994 Mar DISC

TAMOXIFEN CITRATE

>D>	AB	TEVA	EQ 10MG BASE	N74504 001	Apr 28, 2003	Jun	DISC
>A>	@		EQ 10MG BASE	N74504 001	Apr 28, 2003	Jun	DISC
>D>	AB		EQ 20MG BASE	N74504 002	Apr 28, 2003	Jun	DISC
>A>	@		EQ 20MG BASE	N74504 002	Apr 28, 2003	Jun	DISC
AB	+	TEVA PHARMS	EQ 20MG BASE	N74858 001	Feb 20, 2003	May	CRLD

TECHNETIUM TC-99M MEDRONATE KIT

INJECTABLE; INJECTION

DRAXIMAGE MDP-10

AP	+	DRAXIMAGE	N/A	N18035 001		Jan	CTNA
----	---	-----------	-----	------------	--	-----	------

DRAXIMAGE MDP-25

	+	DRAXIMAGE	N/A	N18035 002	Feb 27, 2004	Jan	NEWA
--	---	-----------	-----	------------	--------------	-----	------

TECHNETIUM TC-99M SESTAMIBI KIT

INJECTABLE; INJECTION

MIRALUMA

@ BRISTOL MYERS SQUIBB N/A

N19785 003

May 23, 1997 May DISC

TEGASEROD MALEATE

TABLET; ORAL

ZELNORM

@ NOVARTIS

EQ 2MG BASE

N21200 001

Jul 24, 2002 Mar DISC

@

EQ 6MG BASE

N21200 002

Jul 24, 2002 Mar DISC

TEMSIROLIMUS

SOLUTION; INTRAVENOUS

TORISEL

	+	WYETH PHARMS INC	25MG/ML (25MG/ML)	N22088 001	May 30, 2007	May	NEWA
--	---	------------------	-------------------	------------	--------------	-----	------

TERBINAFINE HYDROCHLORIDE

TABLET; ORAL

LAMISIL

>D>	+	NOVARTIS	EQ 250MG BASE	N20539 001	May 10, 1996	Jun	CFTG
>A>	AB	+	EQ 250MG BASE	N20539 001	May 10, 1996	Jun	CFTG
>A>		TERBINAFINE HYDROCHLORIDE					
>A>	AB	AMNEAL PHARM	EQ 250MG BASE	N77919 001	Jul 02, 2007	Jun	NEWA

TABLET; ORALTERBINAFINE HYDROCHLORIDE

>A>	AB	APOTEX	EQ 250MG BASE	N78199 001	Jul 02, 2007	Jun	NEWA
>A>	AB	AUROBINDO PHARMA	EQ 250MG BASE	N78297 001	Jul 02, 2007	Jun	NEWA
>A>	AB	DR REDDYS LABS INC	EQ 250MG BASE	N76390 001	Jul 02, 2007	Jun	NEWA
>A>	AB	GEDEON RICHTER USA	EQ 250MG BASE	N77065 001	Jul 02, 2007	Jun	NEWA
>A>	AB	GENPHARM	EQ 250MG BASE	N77136 001	Jul 02, 2007	Jun	NEWA
>A>	AB	GLENMARK PHARMS LTD	EQ 250MG BASE	N78157 001	Jul 02, 2007	Jun	NEWA
>A>	AB	INVAGEN PHARMS	EQ 250MG BASE	N77533 001	Jul 02, 2007	Jun	NEWA
>A>	AB	MYLAN	EQ 250MG BASE	N77195 001	Jul 02, 2007	Jun	NEWA
>A>	AB	ORCHID HLTHCARE	EQ 250MG BASE	N78163 001	Jul 02, 2007	Jun	NEWA
>A>	AB	ROXANE	EQ 250MG BASE	N77223 001	Jul 02, 2007	Jun	NEWA
>A>	AB	TEVA	EQ 250MG BASE	N76377 001	Jul 02, 2007	Jun	NEWA
>A>	AB	WATSON LABS	EQ 250MG BASE	N77137 001	Jul 02, 2007	Jun	NEWA
>A>	AB	WOCKHARDT	EQ 250MG BASE	N78229 001	Jul 02, 2007	Jun	NEWA

TERBUTALINE SULFATETABLET; ORALBRETHINE

@	AAIPHARMA LLC	2.5MG	N17849 001			Apr	DISC
@		5MG	N17849 002			Apr	DISC

TERCONAZOLESUPPOSITORY; VAGINALTERCONAZOLE

AB	TARO	80MG	N77553 001	Mar 09, 2007	Feb	NEWA
----	------	------	------------	--------------	-----	------

TESTOSTERONEGEL; TRANSDERMALTESTOSTERONE

AB	PAR PHARM	1%	N76744 001	May 23, 2007	May	NEWA
----	-----------	----	------------	--------------	-----	------

TETRACYCLINE HYDROCHLORIDECAPSULE; ORALACHROMYCIN V

@	HERITAGE PHARMS INC	250MG	N50278 003			Apr	CAHN
@		500MG	N50278 001			Apr	CAHN

TETRACYCLINE HYDROCHLORIDE

>D>	AB	MYLAN	250MG	N60783 001		Jun	DISC
>A>	@		250MG	N60783 001		Jun	DISC
>D>	AB		500MG	N60783 002		Jun	DISC
>A>	@		500MG	N60783 002		Jun	DISC

FIBER, EXTENDED RELEASE; PERIODONTALACTISITE

@	ON SITE	12.7MG/FIBER	N50653 001	Mar 25, 1994	May	CAHN
---	---------	--------------	------------	--------------	-----	------

THALIDOMIDECAPSULE; ORALTHALOMID

	CELGENE	150MG	N20785 004	Jan 10, 2007	Jan	NEWA
--	---------	-------	------------	--------------	-----	------

THEOPHYLLINETABLET; ORALQUIBRON-T

@	MONARCH PHARMS	300MG	N88656 001	Aug 22, 1985	Feb	DISC
---	----------------	-------	------------	--------------	-----	------

TABLET; ORAL

THEOLAIR

+ GRACEWAY	125MG	N86399 001	Mar	CAHN
+	250MG	N86399 002	Mar	CAHN

TABLET, EXTENDED RELEASE; ORAL

QUIBRON-T/SR

@ MONARCH PHARMS	300MG	N87563 001	Jun 21, 1983	Feb	DISC
------------------	-------	------------	--------------	-----	------

>D> THIETHYLPERAZINE MALEATE

>D> TABLET; ORAL

>D> TORECAN

>D> + NOVARTIS	10MG	N12753 001	Jun	DISC
>A> @	10MG	N12753 001	Jun	DISC

THIORIDAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

THIORIDAZINE HYDROCHLORIDE

>D> AA PHARM ASSOC	30MG/ML	N40187 001	Aug 28, 1997	Jun	CRLD
>A> +	30MG/ML	N40187 001	Aug 28, 1997	Jun	CRLD
>D> AA	100MG/ML	N40213 001	May 29, 1998	Jun	CRLD
>A> AA +	100MG/ML	N40213 001	May 29, 1998	Jun	CRLD
>D> AA + TEVA PHARMS	30MG/ML	N89602 001	Nov 09, 1987	Jun	DISC
>A> @	30MG/ML	N89602 001	Nov 09, 1987	Jun	DISC
>D> AA +	100MG/ML	N89603 001	Nov 09, 1987	Jun	DISC
>A> @	100MG/ML	N89603 001	Nov 09, 1987	Jun	DISC

THIORIDAZINE HYDROCHLORIDE INTENSOL

@ ROXANE

@	30MG/ML	N88941 001	Dec 16, 1985	May	DISC
---	---------	------------	--------------	-----	------

@

@	100MG/ML	N88942 001	Dec 16, 1985	May	DISC
---	----------	------------	--------------	-----	------

TABLET; ORAL

THIORIDAZINE HYDROCHLORIDE

>D> AB IVAX PHARMS	10MG	N88270 001	Apr 14, 1983	Jun	DISC
>A> @	10MG	N88270 001	Apr 14, 1983	Jun	DISC
>D> AB	15MG	N88271 001	Apr 14, 1983	Jun	DISC
>A> @	15MG	N88271 001	Apr 14, 1983	Jun	DISC
>D> AB	25MG	N88272 001	Apr 14, 1983	Jun	DISC
>A> @	25MG	N88272 001	Apr 14, 1983	Jun	DISC
>D> AB TEVA	100MG	N88456 001	May 17, 1985	Jun	DISC
>A> @	100MG	N88456 001	May 17, 1985	Jun	DISC

THIOTHIXENE HYDROCHLORIDE

CONCENTRATE; ORAL

NAVANE

>D> AA + PFIZER	EQ 5MG BASE/ML	N16758 001	Jun	CTEC
>A> +	EQ 5MG BASE/ML	N16758 001	Jun	CTEC

>D> THIOTHIXENE HYDROCHLORIDE

>D> AA TEVA PHARMS	EQ 5MG BASE/ML	N71554 001	Oct 16, 1987	Jun	DISC
>A> @	EQ 5MG BASE/ML	N71554 001	Oct 16, 1987	Jun	DISC

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HYDROCHLORIDE

>D> AB MYLAN	250MG	N75316 001	Nov 02, 1999	Jun	DISC
>A> @	250MG	N75316 001	Nov 02, 1999	Jun	DISC

TOLBUTAMIDE

TABLET; ORAL

TOLBUTAMIDE

>D>	AB	IVAX PHARMS	500MG	N87093 001		Jun	DISC
>A>		@	500MG	N87093 001		Jun	DISC

TOLCAPONE

TABLET; ORAL

TASMAR

		VALEANT PHARM INTL	100MG	N20697 001	Jan 29, 1998	Jan	CAHN
		+	200MG	N20697 002	Jan 29, 1998	Jan	CAHN

TOLMETIN SODIUM

CAPSULE; ORAL

TOLMETIN SODIUM

>D>	AB	IVAX PHARMS	EQ 400MG BASE	N73392 001	Jan 24, 1992	Jun	DISC
>A>		@	EQ 400MG BASE	N73392 001	Jan 24, 1992	Jun	DISC

TRAMADOL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

ULTRAM ER

		BIOVAIL LABS INTL	100MG	N21692 001	Sep 08, 2005	Apr	CRLD
			100MG	N21692 001	Sep 08, 2005	Feb	CTNA
			200MG	N21692 002	Sep 08, 2005	Feb	CTNA
			300MG	N21692 003	Sep 08, 2005	Apr	CRLD
		+	300MG	N21692 003	Sep 08, 2005	Feb	CTNA

TRANDOLAPRIL

TABLET; ORAL

TRANDOLAPRIL

>A>	AB	AUROBINDO PHARMA	1MG	N78438 001	Jun 12, 2007	Jun	NEWA
>A>	AB		2MG	N78438 002	Jun 12, 2007	Jun	NEWA
>A>	AB		4MG	N78438 003	Jun 12, 2007	Jun	NEWA
>A>	AB	CIPLA	1MG	N77307 002	Jun 12, 2007	Jun	NEWA
>A>	AB		2MG	N77307 001	Jun 12, 2007	Jun	NEWA
>A>	AB		4MG	N77307 003	Jun 12, 2007	Jun	NEWA
>A>	AB	COBALT	1MG	N77805 001	Jun 12, 2007	Jun	NEWA
>A>	AB		2MG	N77805 002	Jun 12, 2007	Jun	NEWA
>A>	AB		4MG	N77805 003	Jun 12, 2007	Jun	NEWA
>A>	AB	COREPHARMA	1MG	N77256 001	Jun 12, 2007	Jun	NEWA
>A>	AB		2MG	N77256 002	Jun 12, 2007	Jun	NEWA
>A>	AB		4MG	N77256 003	Jun 12, 2007	Jun	NEWA
>A>	AB	INVAGEN PHARMS	1MG	N78320 001	Jun 12, 2007	Jun	NEWA
>A>	AB		2MG	N78320 002	Jun 12, 2007	Jun	NEWA
>A>	AB		4MG	N78320 003	Jun 12, 2007	Jun	NEWA
>A>	AB	LUPIN	1MG	N77522 001	Jun 12, 2007	Jun	NEWA
>A>	AB		2MG	N77522 002	Jun 12, 2007	Jun	NEWA
>A>	AB		4MG	N77522 003	Jun 12, 2007	Jun	NEWA

TRANDOLAPRIL; VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

TARKA

		ABBOTT	1MG; 240MG	N20591 003	Oct 22, 1996	May	CRLD
			2MG; 180MG	N20591 001	Oct 22, 1996	May	CRLD

TABLET, EXTENDED RELEASE; ORAL

TARKA

ABBOTT	2MG;240MG	N20591 004	Oct 22, 1996	May	CRLD
--------	-----------	------------	--------------	-----	------

TRAZODONE HYDROCHLORIDE

TABLET; ORAL

TRAZODONE HYDROCHLORIDE

AB	APOTEX	50MG	N71258 001	Mar 25, 1987	Mar	CAHN	
>D>	AB	MYLAN	50MG	N71405 001	Feb 27, 1991	Jun	DISC
>A>	@	50MG	N71405 001	Feb 27, 1991	Jun	DISC	
>D>	AB	TEVA	150MG	N74357 001	Apr 30, 1997	Jun	DISC
>A>	@	150MG	N74357 001	Apr 30, 1997	Jun	DISC	

TRETINOIN

CAPSULE; ORAL

TRETINOIN

>A>	AB	BARR	10MG	N77684 001	Jun 22, 2007	Jun	NEWA
		VESANOID					
>D>	+	ROCHE	10MG	N20438 001	Nov 22, 1995	Jun	CFTG
>A>	AB	+	10MG	N20438 001	Nov 22, 1995	Jun	CFTG

SOLUTION; TOPICAL

TRETINOIN

>D>	AT	TEVA PHARMS	0.05%	N74873 001	Jun 19, 1998	Jun	DISC
>A>	@		0.05%	N74873 001	Jun 19, 1998	Jun	DISC

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

KENALOG

@ APOTHECON

0.5%

N83943 001

Mar

DISC

TRIACET

>D>	AT	TEVA	0.1%	N84908 002		Jun	DISC
>A>	@		0.1%	N84908 002		Jun	DISC

TRIAMCINOLONE ACETONIDE

AT + ALTANA

0.5%

N85692 002

Apr

CRLD

LOTION; TOPICAL

KENALOG

>D>	AT	APOTHECON	0.025%	N84343 001		Jun	DISC
>A>	@		0.025%	N84343 001		Jun	DISC
>D>	AT	+	0.1%	N84343 002		Jun	DISC
>A>	@		0.1%	N84343 002		Jun	DISC

TRIAMCINOLONE ACETONIDE

>D> AT MORTON GROVE

0.1%

N88451 001

Apr 03, 1985

Jun

CRLD

>A>	AT	+	0.1%	N88451 001	Apr 03, 1985	Jun	CRLD
-----	----	---	------	------------	--------------	-----	------

SPRAY; TOPICAL

KENALOG

>D>	+	APOTHECON	0.147MG/GM	N12104 001		Jun	CAHN
>A>	+	RANBAXY	0.147MG/GM	N12104 001		Jun	CAHN

TRIMETREXATE GLUCURONATE

>D> INJECTABLE; INJECTION

>D> NEUTREXIN

>D>	+	MEDIMMUNE ONCOLOGY	EQ 25MG BASE/VIAL	N20326 001	Dec 17, 1993	Jun	DISC
>A>	@		EQ 25MG BASE/VIAL	N20326 001	Dec 17, 1993	Jun	DISC
>D>	+		EQ 200MG BASE/VIAL	N20326 002	Jul 31, 1998	Jun	DISC
>A>	@		EQ 200MG BASE/VIAL	N20326 002	Jul 31, 1998	Jun	DISC

VALACYCLOVIR HYDROCHLORIDE

TABLET; ORAL

VALACYCLOVIR HYDROCHLORIDE

AB	RANBAXY	EQ 500MG BASE	N76588 001	Jan 31, 2007	Jan	NEWA
AB		EQ 1GM BASE	N76588 002	Jan 31, 2007	Jan	NEWA
	VALTREX					
AB	GLAXOSMITHKLINE	EQ 500MG BASE	N20487 001	Jun 23, 1995	Jan	CFTG
AB	+	EQ 1GM BASE	N20487 002	Jun 23, 1995	Jan	CFTG

>D> VALDECOXIB

>D> TABLET; ORAL

>D> BEXTRA

>D>	GD SEARLE	10MG	N21341 002	Nov 16, 2001	Jun	DISC
>A>	@	10MG	N21341 002	Nov 16, 2001	Jun	DISC
>D>	+	20MG	N21341 003	Nov 16, 2001	Jun	DISC
>A>	@	20MG	N21341 003	Nov 16, 2001	Jun	DISC

VALRUBICIN

SOLUTION; INTRAVESICAL

VALSTAR PRESERVATIVE FREE

+	INDEVUS PHARMS	40MG/ML	N20892 001	Sep 25, 1998	Apr	CAHN
---	----------------	---------	------------	--------------	-----	------

VERAPAMIL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

VERELAN

AB	ELAN DRUG	120MG	N19614 001	May 29, 1990	May	CRLD
AB		180MG	N19614 003	Jan 09, 1992	May	CRLD
AB		240MG	N19614 002	May 29, 1990	May	CRLD

INJECTABLE; INJECTION

ISOPTIN

@	FSC	2.5MG/ML	N18485 001		Feb	DISC
---	-----	----------	------------	--	-----	------

VERAPAMIL HYDROCHLORIDE

@	LUITPOLD	2.5MG/ML	N70225 001	Nov 12, 1985	Feb	DISC
---	----------	----------	------------	--------------	-----	------

AP	+	2.5MG/ML	N70617 001	Nov 12, 1985	Feb	CRLD
----	---	----------	------------	--------------	-----	------

TABLET; ORAL

CALAN

AB	+	GD SEARLE LLC	120MG	N18817 002	Sep 10, 1984	Feb	CRLD
----	---	---------------	-------	------------	--------------	-----	------

ISOPTIN

@	FSC	40MG	N18593 003	Nov 23, 1987	Feb	DISC
---	-----	------	------------	--------------	-----	------

@		80MG	N18593 001	Mar 08, 1982	Feb	DISC
---	--	------	------------	--------------	-----	------

@		120MG	N18593 002	Mar 08, 1982	Feb	DISC
---	--	-------	------------	--------------	-----	------

VERAPAMIL HYDROCHLORIDE

@	HERITAGE PHARMS INC	80MG	N71880 001	Apr 05, 1988	Feb	CAHN
---	---------------------	------	------------	--------------	-----	------

@		120MG	N71881 001	Apr 05, 1988	Feb	CAHN
---	--	-------	------------	--------------	-----	------

>D>	AB	MUTUAL PHARM	80MG	N71488 001	Jan 13, 1988	Jun	DISC
-----	----	--------------	------	------------	--------------	-----	------

>A>	@		80MG	N71488 001	Jan 13, 1988	Jun	DISC
-----	---	--	------	------------	--------------	-----	------

>D>	AB		120MG	N71489 001	Jan 13, 1988	Jun	DISC
-----	----	--	-------	------------	--------------	-----	------

>A>	@		120MG	N71489 001	Jan 13, 1988	Jun	DISC
-----	---	--	-------	------------	--------------	-----	------

ZICONOTIDE

INJECTABLE; INTRATHECAL

PRIALT

+	ELAN PHARMS	100UGM/1ML(100UGM/ML)	N21060 002	Dec 28, 2004	May	CPOT
---	-------------	-----------------------	------------	--------------	-----	------

+		500UGM/20ML(25UGM/ML)	N21060 001	Dec 28, 2004	May	CPOT
---	--	-----------------------	------------	--------------	-----	------

INJECTABLE; INTRATHECAL

PRIALT

+	ELAN PHARMS	500UGM/5ML(100UGM/ML)	N21060 004	Dec 28, 2004	May	CPOT
---	-------------	-----------------------	------------	--------------	-----	------

ZIDOVUDINE

CAPSULE; ORAL

ZIDOVUDINE

AB	CIPLA LTD	100MG	N78349 001	May 23, 2007	May	NEWA
----	-----------	-------	------------	--------------	-----	------

ZILEUTON

TABLET, EXTENDED RELEASE; ORAL

ZYFLO CR

+	CRITICAL	600MG	N22052 001	May 30, 2007	May	NEWA
---	----------	-------	------------	--------------	-----	------

ZOLEDRONIC ACID

INJECTABLE; IV (INFUSION)

RECLAST

+	NOVARTIS	EQ 5MG BASE/100ML	N21817 001	Apr 16, 2007	Apr	NEWA
---	----------	-------------------	------------	--------------	-----	------

ZOLPIDEM TARTRATE

TABLET; ORAL

AMBIEN

AB	SANOFI AVENTIS US	5MG	N19908 001	Dec 16, 1992	Apr	CFTG
----	-------------------	-----	------------	--------------	-----	------

AB	+	10MG	N19908 002	Dec 16, 1992	Apr	CFTG
----	---	------	------------	--------------	-----	------

ZOLPIDEM TARTRATE

AB	APOTEX INC	5MG	N77884 001	Apr 23, 2007	Apr	NEWA
----	------------	-----	------------	--------------	-----	------

AB		10MG	N77884 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	AUROBINDO PHARMA	5MG	N78413 001	May 04, 2007	Apr	NEWA
----	------------------	-----	------------	--------------	-----	------

AB		10MG	N78413 002	May 04, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	CARACO	5MG	N77359 001	Apr 23, 2007	Apr	NEWA
----	--------	-----	------------	--------------	-----	------

AB		10MG	N77359 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	CARLSBAD	5MG	N77990 001	Apr 23, 2007	Apr	NEWA
----	----------	-----	------------	--------------	-----	------

AB		10MG	N77990 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	DR REDDYS LABS LTD	5MG	N77985 001	Apr 23, 2007	Apr	NEWA
----	--------------------	-----	------------	--------------	-----	------

AB		10MG	N77985 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	GENPHARM	5MG	N78016 001	Apr 23, 2007	Apr	NEWA
----	----------	-----	------------	--------------	-----	------

AB		10MG	N78016 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	LEK PHARMS DD	5MG	N77322 001	Apr 23, 2007	Apr	NEWA
----	---------------	-----	------------	--------------	-----	------

AB		10MG	N77322 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	MUTUAL PHARMA	5MG	N77288 001	Apr 23, 2007	Apr	NEWA
----	---------------	-----	------------	--------------	-----	------

AB		10MG	N77288 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	MYLAN	5MG	N76578 001	Apr 23, 2007	Apr	NEWA
----	-------	-----	------------	--------------	-----	------

AB		10MG	N76578 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

>D>	AB	PAR PHARM	5MG	N76062 001	Apr 23, 2007	Jun	DISC
-----	----	-----------	-----	------------	--------------	-----	------

>A>		@	5MG	N76062 001	Apr 23, 2007	Jun	DISC
-----	--	---	-----	------------	--------------	-----	------

AB		5MG	N76062 001	Apr 23, 2007	Apr	NEWA
----	--	-----	------------	--------------	-----	------

>D>	AB		10MG	N76062 002	Apr 23, 2007	Jun	DISC
-----	----	--	------	------------	--------------	-----	------

>A>		@	10MG	N76062 002	Apr 23, 2007	Jun	DISC
-----	--	---	------	------------	--------------	-----	------

AB		10MG	N76062 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	RANBAXY	5MG	N78055 001	Apr 23, 2007	Apr	NEWA
----	---------	-----	------------	--------------	-----	------

AB		10MG	N78055 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	ROXANE	5MG	N77214 001	Apr 23, 2007	Apr	NEWA
----	--------	-----	------------	--------------	-----	------

AB		10MG	N77214 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	SYNTHON PHARMS	5MG	N77540 001	Apr 23, 2007	Apr	NEWA
----	----------------	-----	------------	--------------	-----	------

AB		10MG	N77540 002	Apr 23, 2007	Apr	NEWA
----	--	------	------------	--------------	-----	------

AB	TEVA	5MG	N76410 001	Apr 23, 2007	Apr	NEWA
----	------	-----	------------	--------------	-----	------

TABLET; ORAL

ZOLPIDEM TARTRATE

AB	TEVA	10MG	N76410 002	Apr 23, 2007	Apr	NEWA
AB	WATSON LABS	5MG	N77773 001	Apr 23, 2007	Apr	NEWA
AB		10MG	N77773 002	Apr 23, 2007	Apr	NEWA
AB	WOCKHARDT	5MG	N78426 001	May 15, 2007	May	NEWA
AB		10MG	N78426 002	May 15, 2007	May	NEWA

>D> TABLET, ORALLY DISINTEGRATING; ORAL

>D> TOVALT ODT

>D>	BIOVAIL LABS INTL	5MG	N21412 001	Apr 25, 2007	Jun	DISC
>A>	@	5MG	N21412 001	Apr 25, 2007	Jun	DISC
		5MG	N21412 001	Apr 25, 2007	Apr	NEWA
>D>	+	10MG	N21412 002	Apr 25, 2007	Jun	DISC
>A>	@	10MG	N21412 002	Apr 25, 2007	Jun	DISC
	+	10MG	N21412 002	Apr 25, 2007	Apr	NEWA

ZONISAMIDE

CAPSULE; ORAL

ZONISAMIDE

AB	COREPHARMA	25MG	N77876 001	Feb 21, 2007	Feb	NEWA
AB		50MG	N77876 002	Feb 21, 2007	Feb	NEWA
AB		100MG	N77876 003	Feb 21, 2007	Feb	NEWA

OTC DRUG PRODUCT LIST - 27TH EDITION

OTC DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 6 - June 2007

2-1

BROMPHENIRAMINE MALEATE; PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

EFIDAC 24 PSEUDOEPHEDRINE HYDROCHLORIDE/BROMPHENIRAMINE MALEATE

@ ALZA 16MG;240MG N19672 001 Mar 29, 1996 May DISC

CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL

SPONGE; TOPICAL

CHLORAPREP ONE-STEP

+ ENTURIA INC 2%;70% (3ML) N20832 001 Jul 14, 2000 Jan CAHN

+ 2%;70% (10.5ML) N20832 004 Aug 20, 2003 Jan CAHN

+ 2%;70% (26ML) N20832 006 Nov 21, 2006 Jan CAHN

CHLORAPREP ONE-STEP FREPP

+ ENTURIA INC 2%;70% (1.5ML) N20832 003 Apr 26, 2002 Jan CAHN

CHLORAPREP WITH TINT

+ ENTURIA INC 2%;70% (3ML) N20832 007 Oct 10, 2006 Feb NEWA

+ 2%;70% (26ML) N20832 002 May 03, 2005 Jan CAHN

+ 2%;70% (10.5ML) N20832 005 Apr 03, 2006 Jan CAHN

SWAB; TOPICAL

CHLORAPREP ONE-STEP SEPP

+ ENTURIA INC 2%;70% (0.67ML) N21555 001 Oct 07, 2002 Jan CAHN

CHLORAPREP SINGLE SWABSTICK

+ ENTURIA INC 2%;70% (1.75ML) N21555 002 May 10, 2005 Jan CAHN

CLOTRIMAZOLE

TABLET; VAGINAL

GYNIX

>D>

TEVA PHARMS 100MG N73249 001 Feb 13, 1998 Jun DISC

>D>

@ 100MG N73249 001 Feb 13, 1998 Jun DISC

>A>

DIPHENHYDRAMINE HYDROCHLORIDE; IBUPROFEN

CAPSULE; ORAL

ADVIL PM

+ WYETH CONS 25MG;EQ 200MG FREE ACID AND POTASSIUM SALT N21393 001 Dec 21, 2005 Apr CAIN

DOXYLAMINE SUCCINATE

TABLET; ORAL

DOXYLAMINE SUCCINATE

>D> COPLEY PHARM 25MG N88900 002 Feb 12, 1988 Jun DISC

>A> @ 25MG N88900 002 Feb 12, 1988 Jun DISC

UNISOM

+ CHATTEM 25MG N18066 001 Feb CAHN

+ MCNEIL CONS 25MG N18066 001 Jan CAHN

FAMOTIDINE

TABLET, CHEWABLE; ORAL

FAMOTIDINE

+ PERRIGO 10MG N75715 001 Aug 22, 2003 Mar CRLD

PEPCID AC

@ MERCK 10MG N20801 001 Sep 24, 1998 Mar DISC

GUAIFENESIN

TABLET, EXTENDED RELEASE; ORAL

HUMIBID

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

IBUPROFEN

CAPSULE; ORAL

ADVIL LIQUI-GELS

+	WYETH CONS	EQ 200MG FREE ACID AND POTASSIUM SALT	N20402 001	Apr 20, 1995	Jan	CAIN
---	------------	---------------------------------------	------------	--------------	-----	------

ADVIL MIGRAINE LIQUI-GELS

+	WYETH CONS	EQ 200MG FREE ACID AND POTASSIUM SALT	N20402 002	Mar 16, 2000	Jan	CAIN
---	------------	---------------------------------------	------------	--------------	-----	------

TABLET; ORAL

IBUPROFEN

	IVAX PHARMS	200MG	N71144 001	Jan 20, 1987	Jun	DISC
>D>	@	200MG	N71144 001	Jan 20, 1987	Jun	DISC
>A>	MYLAN	200MG	N71870 001	May 05, 1988	Jun	DISC
>D>	@	200MG	N71870 001	May 05, 1988	Jun	DISC

IBUPROFEN; PSEUDOEPHEDRINE HYDROCHLORIDE

CAPSULE; ORAL

ADVIL COLD AND SINUS

+	WYETH CONS	EQ 200MG FREE ACID AND POTASSIUM SALT; 30MG	N21374 001	May 30, 2002	Apr	CAIN
---	------------	---	------------	--------------	-----	------

INSULIN ZINC SUSP RECOMBINANT HUMAN

INJECTABLE; INJECTION

HUMULIN L

	LILLY	100 UNITS/ML	N19377 002	Sep 30, 1985	Jun	DISC
>D>	@	100 UNITS/ML	N19377 002	Sep 30, 1985	Jun	DISC

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC

ALAWAY

+	BAUSCH AND LOMB	EQ 0.025% BASE	N21996 001	Dec 01, 2006	Jan	CAHN
---	-----------------	----------------	------------	--------------	-----	------

LOPERAMIDE HYDROCHLORIDE

SOLUTION; ORAL

LOPERAMIDE HYDROCHLORIDE

	TEVA	1MG/5ML	N73478 001	Jun 23, 1995	Jun	DISC
>D>	@	1MG/5ML	N73478 001	Jun 23, 1995	Jun	DISC

TABLET, CHEWABLE; ORAL

IMODIUM A-D EZ CHEWS

+	MCNEIL	2MG	N20448 001	Jul 24, 1997	Jan	CTNA
---	--------	-----	------------	--------------	-----	------

LORATADINE

TABLET, ORALLY DISINTEGRATING; ORAL

LORATADINE REDIDOSE

	RANBAXY	10MG	N77153 001	Apr 11, 2007	Apr	NEWA
--	---------	------	------------	--------------	-----	------

MICONAZOLE NITRATE

CREAM; VAGINAL

MICONAZOLE NITRATE

	TEVA	2%	N74136 001	Jan 04, 1995	Jun	DISC
>D>	@	2%	N74136 001	Jan 04, 1995	Jun	DISC

CREAM; VAGINAL

MICONAZOLE NITRATE

>D>	TEVA PHARMS	2%	N74030	001	Oct 30, 1992	Jun	DISC
>A>	@	2%	N74030	001	Oct 30, 1992	Jun	DISC

MINOXIDIL

SOLUTION; TOPICAL

MINOXIDIL (FOR MEN)

>D>	COPLEY PHARM	2%	N74500	001	May 23, 1996	Jun	DISC
>A>	@	2%	N74500	001	May 23, 1996	Jun	DISC
>D>	TEVA	2%	N74589	001	Apr 05, 1996	Jun	DISC
>A>	@	2%	N74589	001	Apr 05, 1996	Jun	DISC

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL

NICOTINE POLACRILEX

	PERRIGO R AND D	EQ 2MG BASE	N78547	001	May 24, 2007	May	NEWA
		EQ 4MG BASE	N78546	001	May 24, 2007	May	NEWA
>A>	THRIVE						
>A>	NOVARTIS	EQ 2MG BASE	N77658	001	Jun 19, 2007	Jun	NEWA
>A>		EQ 4MG BASE	N77656	001	Jun 19, 2007	Jun	NEWA

NONOXYNOL-9

SPONGE; VAGINAL

TODAY

>D>	+ ALLENDALE PHARMS	1GM	N18683	001	Apr 01, 1983	Jun	CAHN
		1GM	N18683	001	Apr 01, 1983	Feb	CMFD
>A>	+ ALLENDALE PHARMS INC	1GM	N18683	001	Apr 01, 1983	Jun	CAHN

ORLISTAT

CAPSULE; ORAL

ALLI

	+ GLAXOSMITHKLINE CONS	60MG	N21887	001	Feb 07, 2007	Feb	NEWA
--	------------------------	------	--------	-----	--------------	-----	------

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

SUDAFED 12 HOUR

	+ MCNEIL CONS	120MG	N73585	001	Oct 31, 1991	Mar	CAHN
--	---------------	-------	--------	-----	--------------	-----	------

SUDAFED 24 HOUR

	+ ALZA	240MG	N20021	002	Dec 15, 1992	May	CTNA
--	--------	-------	--------	-----	--------------	-----	------

RANITIDINE HYDROCHLORIDE

TABLET, EFFERVESCENT; ORAL

ZANTAC 75

	@ BOEHRINGER INGELHEIM	EQ 75MG BASE	N20745	001	Feb 26, 1998	Feb	CAHN
--	------------------------	--------------	--------	-----	--------------	-----	------

TABLET; ORAL

ZANTAC 150

>A>	BOEHRINGER INGELHEIM	EQ 150MG BASE	N21698	002	Mar 13, 2007	Jun	NEWA
-----	----------------------	---------------	--------	-----	--------------	-----	------

	+	EQ 150MG BASE	N21698	001	Aug 31, 2004	Jan	CAHN
--	---	---------------	--------	-----	--------------	-----	------

ZANTAC 75

	BOEHRINGER INGELHEIM	EQ 75MG BASE	N20520	001	Dec 19, 1995	Jan	CAHN
--	----------------------	--------------	--------	-----	--------------	-----	------

SODIUM MONOFLUOROPHOSPHATE

>D>	GEL; DENTAL						
>D>	EXTRA-STRENGTH AIM						
>D>	+ CHESEBROUGH PONDS	1.2%	N19518	002	Aug 06, 1986	Jun	DISC

>D>	GEL; DENTAL								
>D>	EXTRA-STRENGTH AIM								
>A>	@ CHESEBROUGH PONDS	1.2%		N19518	002	Aug 06, 1986	Jun	DISC	
>D>	PASTE; DENTAL								
>D>	EXTRA-STRENGTH AIM								
>D>	+ CHESEBROUGH PONDS	1.2%		N19518	001	Jun 03, 1987	Jun	DISC	
>A>	@	1.2%		N19518	001	Jun 03, 1987	Jun	DISC	

TERBINAFINE HYDROCHLORIDE

	CREAM; TOPICAL								
	TERBINAFINE HYDROCHLORIDE								
>A>	TARO	1%		N77511	001	Jul 02, 2007	Jun	NEWA	

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

CUMULATIVE SUPPLEMENT NUMBER 06 JUNE 2007

NO JUNE 2007 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 JUNE 2007 ADDITIONS

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ABACAVIR SULFATE - ZIAGEN</u>					
020978 001	>A> 6641843	Feb 04, 2020	DP		
<u>ABACAVIR SULFATE; LAMIVUDINE - EPZICOM</u>					
021652 001	7119202	Feb 08, 2009	DS		
	7119202*PED	Aug 08, 2009			
<u>ABACAVIR SULFATE; LAMIVUDINE; ZIDOVUDINE - TRIZIVIR</u>					
021205 001	7119202	Feb 08, 2009	DS		
	7119202*PED	Aug 08, 2009			
<u>ACARBOSE - PRECOSE</u>					
020482 001	4904769	Sep 06, 2009			
<u>ACARBOSE - PRECOSE</u>					
020482 002	4904769	Sep 06, 2009			
<u>ACARBOSE - PRECOSE</u>					
020482 004	4904769	Sep 06, 2009			
<u>ADAPALENE - DIFFERIN</u>					
021753 001	>A> 4717720	May 31, 2010	DS DP	>A> NP	Jun 19, 2010
	>A> RE34440	May 31, 2010		U-818	
<u>ALBUTEROL SULFATE - VENTOLIN HFA</u>					
020983 001	6558651	Dec 19, 2016	DP		
	6743413	Jun 01, 2021	DP	U-716	
<u>ALENDRONATE SODIUM; CHOLECALCIFEROL - FOSAMAX PLUS D</u>					
021762 001				M-51	Dec 21, 2008
<u>ALENDRONATE SODIUM; CHOLECALCIFEROL - FOSAMAX PLUS D</u>					
021762 002				D-107 M-51 NC	Apr 26, 2010 Dec 21, 2008 Apr 07, 2008
<u>ALFUZOSIN HYDROCHLORIDE - UROXATRAL</u>					
021287 001	4661491	Jan 18, 2011		U-706	
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>					
021985 001	5559111	Apr 04, 2015	DS DP	U-3	NCE
<u>ALISKIREN HEMIFUMARATE - TEKTURNA</u>					
021985 002	5559111	Apr 04, 2015	DS DP	U-3	NCE
<u>AMBRISENTAN - LETAIRIS</u>					
022081 001				>A> NCE	Jun 15, 2012
<u>AMBRISENTAN - LETAIRIS</u>					
022081 002				>A> NCE	Jun 15, 2012
<u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u>					
021990 002				>A> NC	Jun 20, 2010
<u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u>					
021990 003				>A> NC	Jun 20, 2010
<u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u>					
021990 004				>A> NC	Jun 20, 2010
<u>AMLODIPINE BESYLATE; VALSARTAN - EXFORGE</u>					
021990 005				>A> NC	Jun 20, 2010
<u>ANIDULAFUNGIN - ERAXIS</u>					
021632 001	7198796	Jun 13, 2022	DP		
<u>ANIDULAFUNGIN - ERAXIS</u>					
021632 002	5965525	Oct 12, 2016	DS DP	U-540	
	6384013	Mar 19, 2012	DS		
	6743777	Mar 19, 2012		DP	U-540
	6960564	Apr 12, 2021		DP	U-540
	7198796	Jun 13, 2022		DP	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ARFORMOTEROL TARTRATE - BROVANA</u>					
021912 001	5795564	Apr 03, 2012	U-793		
	6068833	Apr 03, 2012	U-793		
	6589508	Apr 03, 2012	U-793		
	6866839	Apr 03, 2012	U-793		
<u>ASCORBIC ACID; POLYETHYLENE GLYCOL 3350; POTASSIUM CHLORIDE; SODIUM ASCORBATE; SODIUM CHLORIDE; SODIUM SULFATE - MOVIPREP</u>					
021881 001	7169381	Sep 01, 2024	DS DP		
<u>ATORVASTATIN CALCIUM - LIPITOR</u>					
020702 001				I-523	Mar 02, 2010
<u>ATORVASTATIN CALCIUM - LIPITOR</u>					
020702 002				I-523	Mar 02, 2010
<u>ATORVASTATIN CALCIUM - LIPITOR</u>					
020702 003				I-523	Mar 02, 2010
<u>ATORVASTATIN CALCIUM - LIPITOR</u>					
020702 004				I-523	Mar 02, 2010
<u>AVOBENZONE; ECAMSULE; OCTOCRYLENE - CAPITAL SOLEIL 15</u>					
021501 001				NC NP	Jul 21, 2009 Oct 02, 2009
<u>AVOBENZONE; ECAMSULE; OCTOCRYLENE; TITANIUM DIOXIDE - ANTHELIOS 20</u>					
021471 001				NC	Oct 05, 2009
<u>BALSALAZIDE DISODIUM - COLAZAL</u>					
020610 001				ODE PED	Dec 20, 2013 Jun 20, 2014
<u>BORTEZOMIB - VELCADE</u>					
021602 001	5780454	May 03, 2017	DP		
<u>BOSENTAN - TRACLEER</u>					
021290 001				M-64	Feb 15, 2010
<u>BOSENTAN - TRACLEER</u>					
021290 002				M-64	Feb 15, 2010
<u>BROMFENAC SODIUM - XIBROM</u>					
021664 001	>A> 4910225	Jan 24, 2009	DP	U-810	
<u>BUDESONIDE - PULMICORT RESPULES</u>					
020929 001	6899099	Dec 23, 2018		U-529	
<u>BUDESONIDE - PULMICORT RESPULES</u>					
020929 002	6899099	Dec 23, 2018		U-529	
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>					
021929 001	6641800	Sep 23, 2012	DP		
<u>BUDESONIDE; FORMOTEROL FUMARATE DIHYDRATE - SYMBICORT</u>					
021929 002	6641800	Sep 23, 2012	DP		
<u>BUPROPION HYDROCHLORIDE - BUPROPION HYDROCHLORIDE</u>					
077284 002				PC	Jun 12, 2007
<u>BUPROPION HYDROCHLORIDE - BUPROPION HYDROCHLORIDE</u>					
077415 002				PC	Jun 12, 2007
<u>CARVEDILOL - COREG</u>					
020297 001				M-61 PED	Feb 23, 2010 Aug 23, 2010
<u>CARVEDILOL - COREG</u>					
020297 002				M-61 PED	Feb 23, 2010 Aug 23, 2010
<u>CARVEDILOL - COREG</u>					
020297 003				M-61 PED	Feb 23, 2010 Aug 23, 2010

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLEMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CARVEDILOL - COREG</u>					
020297 004				M-61 PED	Feb 23, 2010 Aug 23, 2010
<u>CELECOXIB - CELEBREX</u>					
020998 004	5466823	Nov 30, 2013	DS	I-466	Jul 29, 2008
	5466823*PED	May 30, 2014		NPP	Dec 15, 2009
	5563165	Nov 30, 2013	DP	PED	Jun 15, 2010
	5563165*PED	May 30, 2014		PED	Jan 29, 2009
	5760068	Jun 02, 2015		U-672	
	5760068*PED	Dec 02, 2015			
<u>CETIRIZINE HYDROCHLORIDE; PSEUDOEPHEDRINE HYDROCHLORIDE - ZYRTEC-D 12 HOUR</u>					
021150 001	>A> 7226614	Jun 10, 2022		U-295	
<u>CETRORELIX - CETROTIDE</u>					
021197 001	5198533	Oct 24, 2010	DS DP		
<u>CETRORELIX - CETROTIDE</u>					
021197 002	5198533	Oct 24, 2010	DS DP		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP ONE-STEP</u>					
020832 006	6991394	Jan 31, 2024	DP		
<u>CHLORHEXIDINE GLUCONATE; ISOPROPYL ALCOHOL - CHLORAPREP WITH TINT</u>					
020832 002	6991394	Jan 31, 2024	DP		
<u>CICLESONIDE - OMNARIS</u>					
022004 001	5482934	Jan 09, 2013	DS DP	U-557	
	6767901	Oct 21, 2020		DP	
	6939559	Apr 21, 2019		DP	
<u>CLOBETASOL PROPIONATE - OLUX E</u>					
022013 001	6730288	Sep 08, 2019	DP	NP	Jan 12, 2010
	7029659	Sep 08, 2019	DP		
<u>CLOZAPINE - FAZACLO ODT</u>					
021590 001	5178878	Jan 12, 2010	DP		
	>A> 6024981	Apr 09, 2018	DP		
	>A> 6221392	Apr 09, 2018	DP		
<u>CLOZAPINE - FAZACLO ODT</u>					
021590 002	5178878	Jan 12, 2010	DP		
	>A> 6024981	Apr 09, 2018	DP		
	>A> 6221392	Apr 09, 2018	DP		
<u>CLOZAPINE - FAZACLO ODT</u>					
021590 003	5178878	Jan 12, 2010	DP		
	>A> 6024981	Apr 09, 2018	DP		
	>A> 6221392	Apr 09, 2018	DP		
<u>CLOZAPINE - FAZACLO ODT</u>					
021590 004	5178878	Jan 12, 2010	DP		
	>A> 6024981	Apr 09, 2018	DP		
	>A> 6221392	Apr 09, 2018	DP		
<u>COLESTIPOL HYDROCHLORIDE - COLESTIPOL HYDROCHLORIDE</u>					
077510 001				PC	Jun 12, 2007
<u>CONIVAPTAN HYDROCHLORIDE - VAPRISOL</u>					
021697 001				I-526	Feb 28, 2010
<u>CYANOCOBALAMIN - NASCOBAL</u>					
021642 001	>A> 7229636	Jun 11, 2024	DP	U-817	
<u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u>					
021777 001				NDF	Feb 01, 2010
<u>CYCLOBENZAPRINE HYDROCHLORIDE - AMRIX</u>					
021777 002				NDF	Feb 01, 2010

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>CYTARABINE - DEPOCYT</u>					
021041 001	5455044	May 14, 2013		U-806	
	5723147	Mar 03, 2015	DP	U-806	
	5962016	Jan 31, 2017	DP	U-806	
	6071534	Feb 18, 2008	DP		
	7198801	Jun 25, 2022	DP		
<u>DALTEPARIN SODIUM - FRAGMIN</u>					
020287 001				I-534	May 01, 2010
<u>DALTEPARIN SODIUM - FRAGMIN</u>					
020287 002				I-534	May 01, 2010
<u>DALTEPARIN SODIUM - FRAGMIN</u>					
020287 003				I-534	May 01, 2010
<u>DALTEPARIN SODIUM - FRAGMIN</u>					
020287 004				I-534	May 01, 2010
<u>DALTEPARIN SODIUM - FRAGMIN</u>					
020287 005				I-534	May 01, 2010
<u>DALTEPARIN SODIUM - FRAGMIN</u>					
020287 006				I-534	May 01, 2010
<u>DALTEPARIN SODIUM - FRAGMIN</u>					
020287 007				I-534	May 01, 2010
<u>DAPSONE - ACZONE</u>					
021794 001	>A> 6060085	Sep 11, 2016		U-124	
<u>DASATINIB - SPRYCEL</u>					
021986 001	7153856	Apr 28, 2020		U-780	
<u>DASATINIB - SPRYCEL</u>					
021986 002	7153856	Apr 28, 2020		U-780	
<u>DASATINIB - SPRYCEL</u>					
021986 003	7153856	Apr 28, 2020		U-780	
<u>DECITABINE - DACOGEN</u>					
021790 001				ODE	May 02, 2013
<u>DESLORATADINE - CLARINEX</u>					
021165 001	7211582	Dec 30, 2014		U-809	
	7211582*PED	Jun 30, 2015			
	>A> 7214683	Dec 30, 2014	DP		
	>A> 7214684	Dec 30, 2014		U-138	
<u>DESLORATADINE - CLARINEX</u>					
021300 001	7211582	Dec 30, 2014		U-809	
	7211582*PED	Jun 30, 2015			
	>A> 7214683	Dec 30, 2014	DP		
	>A> 7214684	Dec 30, 2014		U-138	
<u>DESLORATADINE - CLARINEX</u>					
021312 001	7211582	Dec 30, 2014		U-809	
	7211582*PED	Jun 30, 2015			
	>A> 7214683	Dec 30, 2014	DP		
	>A> 7214684	Dec 30, 2014		U-138	
<u>DESLORATADINE - CLARINEX</u>					
021312 002	7211582	Dec 30, 2014		U-809	
	7211582*PED	Jun 30, 2015			
	>A> 7214683	Dec 30, 2014	DP		
	>A> 7214684	Dec 30, 2014		U-138	
<u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX D 24 HOUR</u>					
021605 001	>A> 7214683	Dec 30, 2014	DP		
	>A> 7214684	Dec 30, 2014		U-138	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DESLORATADINE; PSEUDOEPHEDRINE SULFATE - CLARINEX-D 12 HOUR</u>					
021313 001	>A> 7214683	Dec 30, 2014	DP		
	>A> 7214684	Dec 30, 2014		U-138	
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u>					
021802 001	>A> 6355656	Dec 04, 2015	DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u>					
021802 002	>A> 6355656	Dec 04, 2015	DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u>					
021802 003	>A> 6355656	Dec 04, 2015	DP		
<u>DEXMETHYLPHENIDATE HYDROCHLORIDE - FOCALIN XR</u>					
021802 004	5837284	Dec 04, 2015	DP		
	5908850	Dec 04, 2015		U-678	
	6228398	Nov 01, 2019	DP	U-676	
	>A> 6355656	Dec 04, 2015	DP		
	6528530	Dec 04, 2015	DP		
	6635284	Dec 04, 2015	DP	U-677	
	6730325	Nov 01, 2019	DP	U-676	
<u>DICLOFENAC EPOLAMINE - FLECTOR</u>					
021234 001	4948805	Nov 09, 2007	DS	NE	Jan 31, 2010
	5607690	Apr 13, 2014	DP	NDF	Jan 31, 2010
<u>DOCOSANOL - ABREVA</u>					
020941 001	>A> 4874794	Apr 28, 2014		U-815	
	>A> 5534554	Dec 13, 2013	DP	U-815	
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u>					
020690 001				I-529	Oct 13, 2009
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u>					
020690 002				I-529	Oct 13, 2009
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT</u>					
021719 001				I-529	Oct 13, 2009
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT ODT</u>					
021720 001				I-529	Oct 13, 2009
<u>DONEPEZIL HYDROCHLORIDE - ARICEPT ODT</u>					
021720 002				I-529	Oct 13, 2009
<u>DOXORUBICIN HYDROCHLORIDE - DOXIL</u>					
050718 001				>A> ODE	May 17, 2014
<u>DROSPIRENONE; ETHINYL ESTRADIOL - YAZ</u>					
021676 001	7163931	Dec 20, 2021		U-1	
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>					
021427 001	5023269	Jun 11, 2013	DS DP	U-795	I-524
	5023269	Jun 11, 2013	DS DP	U-799	Feb 23, 2010
	5023269	Jun 11, 2013	DS DP	U-797	
	5023269	Jun 11, 2013	DS DP	U-796	
	5023269	Jun 11, 2013	DS DP	U-605	
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>					
021427 002	5023269	Jun 11, 2013	DS DP	U-797	I-524
	5023269	Jun 11, 2013	DS DP	U-799	Feb 23, 2010
	5023269	Jun 11, 2013	DS DP	U-796	
	5023269	Jun 11, 2013	DS DP	U-795	
	5023269	Jun 11, 2013	DS DP	U-605	
<u>DULOXETINE HYDROCHLORIDE - CYMBALTA</u>					
021427 004	5023269	Jun 11, 2013	DS DP	U-795	I-524
	5023269	Jun 11, 2013	DS DP	U-797	Feb 23, 2010
	5023269	Jun 11, 2013	DS DP	U-799	
	5023269	Jun 11, 2013	DS DP	U-796	
	5023269	Jun 11, 2013	DS DP	U-605	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>DUTASTERIDE - AVODART</u>					
021319 001	5565467	Nov 20, 2015	DS DP		
<u>EFAVIRENZ; EMTRICITABINE; TENOFOVIR DISOPROXIL FUMARATE - ATRIPLA</u>					
021937 001	6639071	Feb 14, 2018	DS	NCE PED	Jul 02, 2008 Jan 02, 2009
<u>EMTRICITABINE - EMTRIVA</u>					
021500 001	5210085	May 11, 2010		U-257	
	5814639	Sep 29, 2015	DS DP		
	5914331	Sep 29, 2015	DS		
<u>ENOXAPARIN SODIUM - LOVENOX</u>					
020164 009				I-533	May 16, 2010
<u>ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)</u>					
020164 001				I-533	May 16, 2010
<u>ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)</u>					
020164 002				I-533	May 16, 2010
<u>ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)</u>					
020164 003				I-533	May 16, 2010
<u>ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)</u>					
020164 004				I-533	May 16, 2010
<u>ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)</u>					
020164 005				I-533	May 16, 2010
<u>ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)</u>					
020164 006				I-533	May 16, 2010
<u>ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)</u>					
020164 007				I-533	May 16, 2010
<u>ENOXAPARIN SODIUM - LOVENOX (PRESERVATIVE FREE)</u>					
020164 008				I-533	May 16, 2010
<u>ENTECAVIR - BARACLUDE</u>					
021798 001	5908638	Jul 26, 2015	DP		
<u>EPLERENONE - INSPRA</u>					
021437 001	4559332	Aug 11, 2007	DS DP	U-537	
	7157101	Dec 08, 2019	DP	U-664	
<u>EPLERENONE - INSPRA</u>					
021437 002	4559332	Aug 11, 2007	DS DP	U-537	
	7157101	Dec 08, 2019	DP	U-664	
<u>EPLERENONE - INSPRA</u>					
021437 003	4559332	Aug 11, 2007	DS DP	U-537	
	7157101	Dec 08, 2019	DP	U-664	
<u>ERTAPENEM SODIUM - INVANZ</u>					
021337 001	5478820	Nov 21, 2015	DS DP	U-160	
	5478820*PED	May 21, 2016			
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>					
021153 001	4738974	Sep 01, 2007	DS DP	U-635	
	4738974	Sep 01, 2007	DS DP	U-770	
	4738974	Sep 01, 2007	DS DP	U-373	
	4738974*PED	Mar 01, 2008		U-373	
	4853230	Sep 01, 2007	DP	U-729	
	4853230	Sep 01, 2007	DP	U-770	
	4853230	Sep 01, 2007	DP	U-373	
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>					
021153 002	4738974	Sep 01, 2007	DS DP	U-770	
	4738974	Sep 01, 2007	DS DP	U-635	
	4738974	Sep 01, 2007	DS DP	U-373	
	4738974*PED	Mar 01, 2008		U-373	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>					
021957 001	4738974	Sep 01, 2007	DS DP U-773		
	4738974	Sep 01, 2007	DS DP U-729		
	4783974*PED	Mar 01, 2008			
<u>ESOMEPRAZOLE MAGNESIUM - NEXIUM</u>					
021957 002	4738974	Sep 01, 2007	DS DP U-773		
	4738974	Sep 01, 2007	DS DP U-729		
	4783974*PED	Mar 01, 2008			
<u>ESTRADIOL - DIVIGEL</u>					
022038 003				>A> NP	Jun 04, 2010
<u>ESTRADIOL - ELESTRIN</u>					
021813 001	7198801	Jun 25, 2022	DP		
<u>ESTRADIOL; NORETHINDRONE ACETATE - ACTIVELLA</u>					
020907 002				D-104 I-525	Dec 28, 2009 Dec 29, 2009
<u>ESTROGENS, CONJUGATED SYNTHETIC A - CENESTIN</u>					
020992 001	5908638	Jul 26, 2015	DP		
<u>ESTROGENS, CONJUGATED SYNTHETIC B - ENJUVA</u>					
021443 001				I-528	Apr 23, 2010
<u>ETHINYL ESTRADIOL; LEVONORGESTREL - LYBREL</u>					
021864 001	>A> 6500814	Sep 03, 2018	U-1	NP	May 22, 2010
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>					
021687 001	>A> 7229982	Dec 11, 2023	DP U-592		
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>					
021687 002	>A> 7229982	Dec 11, 2023	DP U-592		
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>					
021687 003	>A> 7229982	Dec 11, 2023	DP U-592		
<u>EZETIMIBE; SIMVASTATIN - VYTORIN</u>					
021687 004	>A> 7229982	Dec 11, 2023	DP U-592		
<u>FENTANYL CITRATE - ACTIQ</u>					
020747 001				M-63	Feb 06, 2010
<u>FENTANYL CITRATE - ACTIQ</u>					
020747 002				M-63	Feb 06, 2010
<u>FENTANYL CITRATE - ACTIQ</u>					
020747 003				M-63	Feb 06, 2010
<u>FENTANYL CITRATE - ACTIQ</u>					
020747 004				M-63	Feb 06, 2010
<u>FENTANYL CITRATE - ACTIQ</u>					
020747 005				M-63	Feb 06, 2010
<u>FENTANYL CITRATE - ACTIQ</u>					
020747 006				M-63	Feb 06, 2010
<u>FENTANYL CITRATE - FENTORA</u>					
021947 006				NDF	Sep 25, 2009
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>					
021520 001	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5229382*PED	Oct 23, 2011		PED	Jun 24, 2007
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>					
021520 002	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5229382*PED	Oct 23, 2011		PED	Jun 24, 2007
	5945416	Mar 24, 2017	DS DP		
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>					
021520 003	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5229382*PED	Oct 23, 2011		PED	Jun 24, 2007
	5945416	Mar 24, 2017	DS DP		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>					
021520 004	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5229382*PED	Oct 23, 2011		PED	Jun 24, 2007
	5945416	Mar 24, 2017	DS DP		
<u>FLUOXETINE HYDROCHLORIDE; OLANZAPINE - SYMBYAX</u>					
021520 005	5229382	Apr 23, 2011	DS DP	NC	Dec 24, 2006
	5229382*PED	Oct 23, 2011		PED	Jun 24, 2007
	5945416	Mar 24, 2017	DS DP		
<u>FLUTICASON FUROATE - VERAMYST</u>					
022051 001	6858596	Aug 03, 2021	DP	U-808	
	7101866	Aug 03, 2021	DS DP	U-808	
<u>FORMOTEROL FUMARATE - FORADIL CERTIHALER</u>					
021592 001	6182655	Dec 05, 2016	DP	NP	Dec 15, 2009
	6645466	Nov 10, 2019	DP		
<u>FORMOTEROL FUMARATE - PERFOROMIST</u>					
022007 001	>A> 6667344	Jun 22, 2021	DP	NP	May 11, 2010
	>A> 6814953	Jun 22, 2021	DP	U-813	
<u>FOSAMPRENAVIR CALCIUM - LEXIVA</u>					
022116 001				>A> NDF	Jun 14, 2010
<u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u>					
021615 001	7160559	Dec 20, 2019	DP		
<u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u>					
021615 002	7160559	Dec 20, 2019	DP		
<u>GALANTAMINE HYDROBROMIDE - RAZADYNE ER</u>					
021615 003	7160559	Dec 20, 2019	DP		
<u>GEMIFLOXACIN MESYLATE - FACTIVE</u>					
021158 001				D-106	May 01, 2010
<u>GOSERELIN ACETATE - ZOLADEX</u>					
019726 001	>A> 7118552	Apr 13, 2022	DP		
	>A> 7220247	Apr 09, 2022	DP		
<u>GOSERELIN ACETATE - ZOLADEX</u>					
020578 001	>A> 7118552	Apr 13, 2022	DP		
	>A> 7220247	Apr 09, 2022	DP		
<u>HISTRELIN ACETATE - SUPPRELIN LA</u>					
022058 001	>A> 5266325	Nov 30, 2010	DP	NP	May 03, 2010
	>A> 5292515	Mar 08, 2011	DP		
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>					
020818 004	5399578	Mar 21, 2012	DS DP	U-3	
	6294197	Jun 18, 2017	DP	U-3	
<u>HYDROCHLOROTHIAZIDE; VALSARTAN - DIOVAN HCT</u>					
020818 005	5399578	Mar 21, 2012	DS DP	U-3	
	6294197	Jun 18, 2017	DP	U-3	
<u>HYDROCORTISONE BUTYRATE - LOCOID</u>					
022076 001				NDF	May 18, 2010
<u>HYDROXOCOBALAMIN - CYANOKIT</u>					
022041 002	5834448	Nov 14, 2016	DP	U-789	Dec 15, 2013
<u>IBANDRONATE SODIUM - BONIVA</u>					
021455 001	4927814	Jul 09, 2008	DS DP	U-700	
	4927814	Jul 09, 2008	DS DP	U-642	
<u>IBANDRONATE SODIUM - BONIVA</u>					
021455 002	4927814	Jul 09, 2008	DS DP	U-700	
	4927814	Jul 09, 2008	DS DP	U-642	
	7192938	May 06, 2023		U-798	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>IBUPROFEN LYSINE - NEOPROFEN</u>					
021903 001	6342530	Nov 14, 2020	DS DP U-794		
	6344479	Mar 20, 2021	DS DP U-794		
<u>IMATINIB MESYLATE - GLEEVEC</u>					
021335 001	6958335	Dec 19, 2021		U-791	
	6958335*PED	Jun 19, 2022			
<u>IMATINIB MESYLATE - GLEEVEC</u>					
021335 002	6958335	Dec 19, 2021		U-791	
	6958335*PED	Jun 19, 2022			
<u>IMATINIB MESYLATE - GLEEVEC</u>					
021588 001	6958335	Dec 19, 2021		U-791	
	6958335*PED	Jun 19, 2022			
<u>IMATINIB MESYLATE - GLEEVEC</u>					
021588 002	6958335	Dec 19, 2021		U-791	
	6958335*PED	Jun 19, 2022			
<u>IRON SUCROSE - VENOFER</u>					
021135 004				I-474 I-459	Oct 17, 2008 Jun 17, 2008
<u>KETOCONAZOLE - XOLEGEL</u>					
021946 001	7179475	Dec 04, 2018	DP U-792		
<u>LAMIVUDINE - EPIVIR</u>					
020564 001	7119202	Feb 08, 2009	DS		
	7119202*PED	Aug 08, 2009			
<u>LAMIVUDINE - EPIVIR</u>					
020564 003	7119202	Feb 08, 2009	DS		
	7119202*PED	Aug 08, 2009			
<u>LAMIVUDINE - EPIVIR</u>					
020596 001	7119202	Feb 08, 2009	DS		
	7119202*PED	Aug 08, 2009			
<u>LAMIVUDINE - EPIVIR-HBV</u>					
021003 001	7119202	Feb 08, 2009	DS		
	7119202*PED	Aug 08, 2008			
<u>LAMIVUDINE - EPIVIR-HBV</u>					
021004 001	7119202	Feb 08, 2009	DS		
	7119202*PED	Aug 08, 2009			
<u>LAMIVUDINE; ZIDOVUDINE - COMBIVIR</u>					
020857 001	7119202	Feb 08, 2009	DS		
	7119202*PED	Aug 08, 2009			
<u>LAMOTRIGINE - LAMICTAL</u>					
020241 001	4602017	Jul 22, 2008		U-106	
	4602017*PED	Jan 22, 2009		I-516 PED	Sep 22, 2009 Mar 22, 2010
<u>LAMOTRIGINE - LAMICTAL</u>					
020241 002	4602017	Jul 22, 2008		U-106	
	4602017*PED	Jan 22, 2009		I-516 PED	Sep 22, 2009 Mar 22, 2010
<u>LAMOTRIGINE - LAMICTAL</u>					
020241 003	4602017	Jul 22, 2008		U-106	
	4602017*PED	Jan 22, 2009		I-516 PED	Sep 22, 2009 Mar 22, 2010
<u>LAMOTRIGINE - LAMICTAL</u>					
020241 004	4602017	Jul 22, 2008		U-106	
	4602017*PED	Jan 22, 2009		I-516 PED	Sep 22, 2009 Mar 22, 2010
<u>LAMOTRIGINE - LAMICTAL</u>					
020241 005	4602017	Jul 22, 2008		U-106	
	4602017*PED	Jan 22, 2009		I-516 PED	Sep 22, 2009 Mar 22, 2010

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LAMOTRIGINE - LAMICTAL</u>					
020241 006	4602017	Jul 22, 2008	U-106	I-516	Sep 22, 2009
	4602017*PED	Jan 22, 2009		PED	Mar 22, 2010
<u>LAMOTRIGINE - LAMICTAL CD</u>					
020764 001	4602017	Jul 22, 2008	U-106	I-516	Sep 22, 2009
	4602017*PED	Jan 22, 2009		PED	Mar 22, 2010
	5698226	Jan 29, 2012			
	5698226*PED	Jul 29, 2012			
<u>LAMOTRIGINE - LAMICTAL CD</u>					
020764 002	4602017	Jul 22, 2008	U-106	I-516	Sep 22, 2009
	4602017*PED	Jan 22, 2009		PED	Mar 22, 2010
	5698226	Jan 29, 2012			
	5698226*PED	Jul 29, 2012			
<u>LAMOTRIGINE - LAMICTAL CD</u>					
020764 003	4602017	Jul 22, 2008	U-106	I-516	Sep 22, 2009
	4602017*PED	Jan 22, 2009		PED	Mar 22, 2010
	5698226	Jan 29, 2012			
	5698226*PED	Jul 29, 2012			
<u>LAMOTRIGINE - LAMICTAL CD</u>					
020764 004	4602017	Jul 22, 2008	U-106	I-516	Sep 22, 2009
	4602017*PED	Jan 22, 2009		PED	Mar 22, 2010
	5698226	Jan 29, 2012			
	5698226*PED	Jul 29, 2012			
<u>LAPATINIB DITOSYLATE - TYKERB</u>					
022059 001	6391874	Jul 11, 2017	DS DP	U-800	NCE
	6713485	Jan 08, 2019	DS DP	U-800	Mar 13, 2012
	6727256	Jan 08, 2019	DS DP	U-800	
	6828320	Jul 11, 2017		U-800	
	7157466	Nov 19, 2021	DS DP		
<u>LATANOPROST - XALATAN</u>					
020597 001	7163959	Jun 19, 2010	DS		
<u>LENALIDOMIDE - REVLIMID</u>					
021880 001	7189740	Apr 11, 2023		U-769	
<u>LENALIDOMIDE - REVLIMID</u>					
021880 002	7189740	Apr 11, 2023		U-769	
<u>LENALIDOMIDE - REVLIMID</u>					
021880 003	7189740	Apr 11, 2023		U-769	
<u>LENALIDOMIDE - REVLIMID</u>					
021880 004	7189740	Apr 11, 2023		U-769	
<u>LEUPROLIDE ACETATE - ELIGARD</u>					
021343 001	6395293	Sep 28, 2013	DP	U-801	
	6565874	Oct 28, 2018	DP	U-801	
	6626870	Mar 27, 2020	DP		
	6773714	Oct 28, 2018	DP	U-801	
	RE37950	Oct 03, 2008	DP	U-801	
<u>LEUPROLIDE ACETATE - ELIGARD</u>					
021379 001	6395293	Sep 28, 2013	DP		
	6565874	Oct 28, 2018	DP	U-801	
	6626870	Mar 27, 2020	DP		
	6773714	Oct 28, 2018	DP	U-801	
	RE37950	Oct 03, 2008	DP	U-801	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEUPROLIDE ACETATE - ELIGARD</u>					
021488 001	6395293	Sep 28, 2013	DP		
	6565874	Oct 28, 2018	DP	U-801	
	6626870	Mar 27, 2020	DP		
	6773714	Oct 28, 2018	DP	U-801	
	RE37950	Oct 03, 2008	DP	U-801	
<u>LEVETIRACETAM - KEPRA</u>					
021035 001				I-527	Mar 19, 2010
<u>LEVETIRACETAM - KEPRA</u>					
021035 002				I-527	Mar 19, 2010
<u>LEVETIRACETAM - KEPRA</u>					
021035 003				I-527	Mar 19, 2010
<u>LEVETIRACETAM - KEPRA</u>					
021035 004				I-527	Mar 19, 2010
<u>LEVETIRACETAM - KEPRA</u>					
021505 001				I-527	Mar 19, 2010
<u>LEVOCETIRIZINE DIHYDROCHLORIDE - XYZAL</u>					
022064 001	>A> 4525358	Jun 25, 2007	DS DP	U-811	May 25, 2010
	>A> 5698558	Sep 24, 2012		U-812	
<u>LEVOFLOXACIN - LEVAQUIN</u>					
020634 001	5053407	Dec 20, 2010		D-100	Aug 04, 2008
	5053407*PED	Jun 20, 2011		D-83	Oct 23, 2006
				PED	Feb 04, 2009
				PED	Apr 23, 2007
<u>LEVOFLOXACIN - LEVAQUIN</u>					
020634 002	5053407	Dec 20, 2010		D-100	Aug 04, 2008
	5053407*PED	Jun 20, 2011		D-83	Oct 23, 2006
				PED	Feb 04, 2009
				PED	Apr 23, 2007
<u>LEVOFLOXACIN - LEVAQUIN</u>					
020634 003	5053407	Dec 20, 2010		D-100	Aug 04, 2008
	5053407*PED	Jun 20, 2011		D-83	Oct 23, 2006
				PED	Apr 23, 2007
				PED	Feb 04, 2009
<u>LEVOFLOXACIN - LEVAQUIN</u>					
020635 001	5053407	Dec 20, 2010		D-100	Aug 04, 2008
	5053407*PED	Jun 20, 2011		D-83	Oct 23, 2006
				PED	Feb 04, 2009
				PED	Apr 23, 2007
<u>LEVOFLOXACIN - LEVAQUIN</u>					
020635 004	5053407	Dec 20, 2010		D-100	Aug 04, 2008
	5053407*PED	Jun 20, 2011		D-83	Oct 23, 2006
				PED	Apr 23, 2007
				PED	Feb 04, 2009
<u>LEVOFLOXACIN - LEVAQUIN</u>					
021721 001	5053407	Dec 20, 2010	DS	U-36	Aug 04, 2008
	5053407*PED	Jun 20, 2011			Feb 04, 2009
	6806256	Feb 26, 2022	DP		
	6806256*PED	Aug 26, 2022			
<u>LEVOFLOXACIN - LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
020635 002	5053407	Dec 20, 2010		D-100	Aug 04, 2008
	5053407*PED	Jun 20, 2011		D-83	Oct 23, 2006
				PED	Feb 04, 2009
				PED	Apr 23, 2007
<u>LEVOFLOXACIN - LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
020635 003	5053407	Dec 20, 2010		D-100	Aug 04, 2008
	5053407*PED	Jun 20, 2011		D-83	Oct 23, 2006
				PED	Feb 04, 2009
				PED	Apr 23, 2007

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>LEVOFLOXACIN - LEVAQUIN IN DEXTROSE 5% IN PLASTIC CONTAINER</u>					
020635 005	5053407	Dec 20, 2010		D-100	Aug 04, 2008
	5053407*PED	Jun 20, 2011		D-83	Oct 23, 2006
				PED	Apr 23, 2007
				PED	Feb 04, 2009
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>					
021977 001	7105486	Jun 29, 2023	U-727	NCE	Feb 23, 2012
	>A> 7223735	Jun 29, 2023	DP		
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>					
021977 002	7105486	Jun 29, 2023	U-727	NCE	Feb 23, 2012
	>A> 7223735	Jun 29, 2023	DP		
<u>LISDEXAMFETAMINE DIMESYLATE - VYVANSE</u>					
021977 003	7105486	Jun 29, 2023	U-727	NCE	Feb 23, 2012
	>A> 7223735	Jun 29, 2023	DP		
<u>LOPERAMIDE HYDROCHLORIDE - IMODIUM A-D EZ CHEWS</u>					
020448 001	5489436	Feb 06, 2013	DP		
	6814978	Aug 26, 2021	DP		
<u>LOPINAVIR; RITONAVIR - KALETRA</u>					
021251 001	5914332	Dec 13, 2015		U-351	
<u>LOPINAVIR; RITONAVIR - KALETRA</u>					
021906 001	7148359	Jul 19, 2019	DP		
<u>MEMANTINE HYDROCHLORIDE - NAMENDA</u>					
021627 001	>A> 5061703	Apr 11, 2010		U-539	
	>A> 5614560	Mar 25, 2014		U-539	
<u>MESALAMINE - LIALDA</u>					
022000 001	6773720	Jun 08, 2020	DP	NP	Jan 16, 2010
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 001				M-62	Jan 31, 2010
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 002				M-62	Jan 31, 2010
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 003				M-62	Jan 31, 2010
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 004				M-62	Jan 31, 2010
<u>METFORMIN HYDROCHLORIDE; ROSIGLITAZONE MALEATE - AVANDAMET</u>					
021410 005				M-62	Jan 31, 2010
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>					
022044 001	6303661	Apr 24, 2017		U-802	Mar 30, 2010
	6699871	Jul 26, 2022	DS DP	U-802	Oct 16, 2011
	6890898	Feb 02, 2019		U-803	
	7078381	Feb 02, 2019		U-803	
	7125873	Jul 26, 2022	DP	U-803	
<u>METFORMIN HYDROCHLORIDE; SITAGLIPTIN PHOSPHATE - JANUMET</u>					
022044 002	6303661	Apr 24, 2017		U-802	Mar 30, 2010
	6699871	Jul 26, 2022	DS DP	U-802	Oct 16, 2011
	6890898	Feb 02, 2019		U-803	
	7078381	Feb 02, 2019		U-803	
	7125873	Jul 26, 2022	DP	U-803	
<u>METHYL AMINOLEVULINATE HYDROCHLORIDE - METVIXIA</u>					
021415 001	6034267	Mar 08, 2016		U-804	
<u>MICONAZOLE NITRATE; PETROLATUM, WHITE; ZINC OXIDE - VUSION</u>					
021026 001	4911932	Mar 27, 2008	DP	U-718	

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>MONTELUKAST SODIUM - SINGULAIR</u>					
020829 002	5565473	Feb 03, 2012	DS DP U-675	I-530	Apr 13, 2010
	5565473	Feb 03, 2012	DS DP U-807		
	5565473	Feb 03, 2012	DS DP U-228		
<u>MONTELUKAST SODIUM - SINGULAIR</u>					
020830 001	5565473	Feb 03, 2012	DS DP U-675	I-530	Apr 13, 2010
	5565473	Feb 03, 2012	DS DP U-807		
	5565473	Feb 03, 2012	DS DP U-228		
<u>MONTELUKAST SODIUM - SINGULAIR</u>					
020830 002	5565473	Feb 03, 2012	DS DP U-807	I-530	Apr 13, 2010
	5565473	Feb 03, 2012	DS DP U-675		
	5565473	Feb 03, 2012	DS DP U-228		
<u>MONTELUKAST SODIUM - SINGULAIR</u>					
021409 001	5565473	Feb 03, 2012	DS DP U-807	I-530	Apr 13, 2010
	5565473	Feb 03, 2012	DS DP U-675		
<u>MORPHINE SULFATE - KADIAN</u>					
020616 006	5202128	Apr 13, 2010	DP		
	5378474	Mar 23, 2010	DP		
<u>MORPHINE SULFATE - KADIAN</u>					
020616 007	5202128	Apr 13, 2010	DP		
	5378474	Mar 23, 2010	DP		
<u>MORPHINE SULFATE - KADIAN</u>					
020616 008	5202128	Apr 13, 2010			
<u>MOXIFLOXACIN HYDROCHLORIDE - VIGAMOX</u>					
021598 001	6716830	Sep 29, 2019	DP		
<u>OLANZAPINE - ZYPREXA</u>					
020592 001	5229382	Apr 23, 2011	DS DP U-149	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS DP U-547	PED	Jul 14, 2007
	5229382*PED	Oct 23, 2011			
	5605897	Feb 25, 2014		U-176	
	5605897*PED	Aug 25, 2014			
	5627178	Apr 23, 2011		U-364	
	5627178*PED	Oct 23, 2011			
	5736541	Mar 24, 2015		U-307	
	5736541*PED	Sep 24, 2015			
	5817655	Apr 23, 2011		U-364	
	5817655*PED	Oct 23, 2011			
	5817656	Apr 23, 2011		U-360	
	5817656*PED	Oct 23, 2011			
	5817657	Apr 23, 2011		U-363	
	5817657*PED	Oct 23, 2011			
	5919485	Mar 24, 2015		U-308	
	5919485*PED	Sep 24, 2015			
	6251895	Sep 23, 2017			
	6251895*PED	Mar 23, 2018			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLANZAPINE - ZYPREXA</u>							
020592 002	5229382	Apr 23, 2011	DS	DP	U-547	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS	DP	U-149	PED	Jul 14, 2007
	5229382*PED	Oct 23, 2011					
	5605897	Feb 25, 2014			U-176		
	5605897*PED	Aug 25, 2014					
	5627178	Apr 23, 2011			U-364		
	5627178*PED	Oct 23, 2011					
	5736541	Mar 24, 2015			U-307		
	5736541*PED	Sep 24, 2015					
	5817655	Apr 23, 2011			U-364		
	5817655*PED	Oct 23, 2011					
	5817656	Apr 23, 2011			U-360		
	5817656*PED	Oct 23, 2011					
	5817657	Apr 23, 2011			U-363		
	5817657*PED	Oct 23, 2011					
	5919485	Mar 24, 2015			U-308		
	5919485*PED	Sep 24, 2015					
	6251895	Sep 23, 2017					
	6251895*PED	Mar 23, 2018					
	<u>OLANZAPINE - ZYPREXA</u>						
020592 003	5229382	Apr 23, 2011	DS	DP	U-547	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS	DP	U-149	PED	Jul 14, 2007
	5229382*PED	Oct 23, 2011					
	5605897	Feb 25, 2014			U-176		
	5605897*PED	Aug 25, 2014					
	5627178	Apr 23, 2011			U-364		
	5627178*PED	Oct 23, 2011					
	5736541	Mar 24, 2015			U-307		
	5736541*PED	Sep 24, 2015					
	5817655	Apr 23, 2011			U-364		
	5817655*PED	Oct 23, 2011					
	5817656	Apr 23, 2011			U-360		
	5817656*PED	Oct 23, 2011					
	5817657	Apr 23, 2011			U-363		
	5817657*PED	Oct 23, 2011					
	5919485	Mar 24, 2015			U-308		
	5919485*PED	Sep 24, 2015					
	6251895	Sep 23, 2017					
	6251895*PED	Mar 23, 2018					
	<u>OLANZAPINE - ZYPREXA</u>						
020592 004	5229382	Apr 23, 2011	DS	DP	U-547	I-417	Jan 14, 2007
	5229382	Apr 23, 2011	DS	DP	U-149	PED	Jul 14, 2007
	5229382*PED	Oct 23, 2011					
	5605897	Feb 25, 2014			U-176		
	5605897*PED	Aug 25, 2014					
	5627178	Apr 23, 2011			U-364		
	5627178*PED	Oct 23, 2011					
	5736541	Mar 24, 2015			U-307		
	5736541*PED	Sep 24, 2015					
	5817655	Apr 23, 2011			U-364		
	5817655*PED	Oct 23, 2011					
	5817656	Apr 23, 2011			U-360		
	5817656*PED	Oct 23, 2011					
	5817657	Apr 23, 2011			U-363		
	5817657*PED	Oct 23, 2011					
	5919485	Mar 24, 2015			U-308		
	5919485*PED	Sep 24, 2015					
	6251895	Sep 23, 2017					
	6251895*PED	Mar 23, 2018					

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLANZAPINE - ZYPREXA</u>					
020592 005	5229382	Apr 23, 2011	DS DP	U-149	I-417 Jan 14, 2007
	5229382	Apr 23, 2011	DS DP	U-547	PED Jul 14, 2007
	5229382*PED	Oct 23, 2011			
	5605897	Feb 25, 2014		U-176	
	5605897*PED	Aug 25, 2014			
	5627178	Apr 23, 2011		U-364	
	5627178*PED	Oct 23, 2011			
	5736541	Mar 24, 2015		U-307	
	5736541*PED	Sep 24, 2015			
	5817655	Apr 23, 2011		U-364	
	5817655*PED	Oct 23, 2011			
	5817656	Apr 23, 2011		U-360	
	5817656*PED	Oct 23, 2011			
	5817657	Apr 23, 2011		U-363	
	5817657*PED	Oct 23, 2011			
	5919485	Mar 24, 2015		U-308	
	5919485*PED	Sep 24, 2015			
	6251895	Sep 23, 2017			
6251895*PED	Mar 23, 2018				
<u>OLANZAPINE - ZYPREXA</u>					
020592 006	5229382	Apr 23, 2011	DS DP	U-547	I-417 Jan 14, 2007
	5229382	Apr 23, 2011	DS DP	U-149	PED Jul 14, 2007
	5229382*PED	Oct 23, 2011			
	5605897	Feb 25, 2014		U-176	
	5605897*PED	Aug 25, 2014			
	5627178	Apr 23, 2011		U-364	
	5627178*PED	Oct 23, 2011			
	5736541	Mar 24, 2015		U-307	
	5736541*PED	Sep 24, 2015			
	5817655	Apr 23, 2011		U-364	
	5817655*PED	Oct 23, 2011			
	5817656	Apr 23, 2011		U-360	
	5817656*PED	Oct 23, 2011			
	5817657	Apr 23, 2011		U-363	
	5817657*PED	Oct 23, 2011			
	5919485	Mar 24, 2015		U-308	
	5919485*PED	Sep 24, 2015			
	6251895	Sep 23, 2017			
6251895*PED	Mar 23, 2018				
<u>OLANZAPINE - ZYPREXA</u>					
021253 001	5229382	Apr 23, 2011	DS DP	U-571	NP Mar 29, 2007
	5229382*PED	Oct 23, 2011			NDF Mar 29, 2007
					PED Sep 29, 2007
					PED Sep 29, 2007
<u>OLANZAPINE - ZYPREXA ZYDIS</u>					
021086 001	5229382	Apr 23, 2011		U-324	I-400 Jul 10, 2006
	5229382*PED	Oct 23, 2011			I-417 Jan 14, 2007
					PED Jan 10, 2007
					PED Jul 14, 2007
<u>OLANZAPINE - ZYPREXA ZYDIS</u>					
021086 002	5229382	Apr 23, 2011		U-324	I-400 Jul 10, 2006
	5229382*PED	Oct 23, 2011			I-417 Jan 14, 2007
					PED Jan 10, 2007
					PED Jul 14, 2007
<u>OLANZAPINE - ZYPREXA ZYDIS</u>					
021086 003	5229382	Apr 23, 2011		U-324	I-400 Jul 10, 2006
	5229382*PED	Oct 23, 2011			I-417 Jan 14, 2007
					PED Jan 10, 2007
					PED Jul 14, 2007

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>OLANZAPINE - ZYPREXA ZYDIS</u>					
021086 004	5229382	Apr 23, 2011	U-324	I-400	Jul 10, 2006
	5229382*PED	Oct 23, 2011		I-417	Jan 14, 2007
				PED	Jan 10, 2007
				PED	Jul 14, 2007
<u>ONDANSETRON - ONDANSETRON</u>					
076506 001				PC	Jun 24, 2007
<u>ONDANSETRON - ONDANSETRON</u>					
076506 002				PC	Jun 24, 2007
<u>ONDANSETRON HYDROCHLORIDE - ONDANSETRON HYDROCHLORIDE</u>					
076183 001				PC	Jun 24, 2007
<u>ONDANSETRON HYDROCHLORIDE - ONDANSETRON HYDROCHLORIDE</u>					
076183 002				PC	Jun 24, 2007
<u>ONDANSETRON HYDROCHLORIDE - ONDANSETRON HYDROCHLORIDE</u>					
076183 003				PC	Jun 24, 2007
<u>ORLISTAT - ALLI</u>					
021887 001	4598089	Jun 18, 2009	DS DP	NP	Feb 07, 2010
	6004996	Jan 06, 2018	DP		
<u>OXALIPLATIN - ELOXATIN</u>					
021492 001				M-61	Jan 10, 2010
				PED	Jul 10, 2010
<u>OXALIPLATIN - ELOXATIN</u>					
021492 002				M-61	Jan 10, 2010
				PED	Jul 10, 2010
<u>OXALIPLATIN - ELOXATIN</u>					
021759 001	5420319	Aug 09, 2016	DS		
	5420319*PED	Feb 09, 2017			
<u>OXALIPLATIN - ELOXATIN</u>					
021759 002	5420319	Aug 09, 2016	DS		
	5420319*PED	Feb 09, 2017			
<u>OXALIPLATIN - ELOXATIN</u>					
021759 003	5290961	Jan 12, 2013	DS		
	5290961*PED	Jul 12, 2013			
	5338874	Apr 07, 2013	DS		
	5338874*PED	Oct 07, 2013			
	5420319	Aug 09, 2016	DS		
	5420319*PED	Feb 09, 2017			
	5716988	Aug 07, 2015		DP	
	5716988*PED	Feb 07, 2016			
<u>OXCARBAZEPINE - TRILEPTAL</u>					
021014 001	7037525	Feb 12, 2018		U-724	
	7037525*PED	Aug 12, 2018			
<u>OXCARBAZEPINE - TRILEPTAL</u>					
021014 002	7037525	Feb 12, 2018		U-724	
	7037525*PED	Aug 12, 2018			
<u>OXCARBAZEPINE - TRILEPTAL</u>					
021014 003	7037525	Feb 12, 2018		U-724	
	7037525*PED	Aug 12, 2018			
<u>OXCARBAZEPINE - TRILEPTAL</u>					
021285 001	7037525*PED	Aug 12, 2018			
<u>OXYBUTYNIN - OXYTROL</u>					
021351 002	7179483	Apr 26, 2020	DS DP	U-318	
<u>PALIPERIDONE - INVEGA</u>					
021999 001	5158952	Oct 27, 2009	DP	U-90	Apr 27, 2010

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>PALIPERIDONE - INVEGA</u>					
021999 002	5158952	Oct 27, 2009	DP U-90	I-531	Apr 27, 2010
<u>PALIPERIDONE - INVEGA</u>					
021999 003	5158952	Oct 27, 2009	DP U-90	I-531	Apr 27, 2010
<u>PALIPERIDONE - INVEGA</u>					
021999 004	5158952	Oct 27, 2009	DP U-90	I-531	Apr 27, 2010
<u>PALONOSETRON HYDROCHLORIDE - ALOXI</u>					
021372 001	5202333	Apr 13, 2015 DS	DP U-528		
<u>PAROXETINE HYDROCHLORIDE - PAROXETINE HYDROCHLORIDE</u>					
077395 001				PC	Jun 10, 2007
<u>PAROXETINE HYDROCHLORIDE - PAXIL CR</u>					
020936 001	5789449	Jan 06, 2009		U-788	
	5789449*PED	Jul 06, 2009			
	>A> 7229640	Jul 19, 2016	DP U-816		
<u>PAROXETINE HYDROCHLORIDE - PAXIL CR</u>					
020936 002	5789449	Jan 06, 2009		U-788	
	5789449*PED	Jul 06, 2009			
	>A> 7229640	Jul 19, 2016	DP U-816		
<u>PAROXETINE HYDROCHLORIDE - PAXIL CR</u>					
020936 003	5789449	Jan 06, 2009		U-788	
	5789449*PED	Jul 06, 2009			
	>A> 7229640	Jul 19, 2016	DP U-816		
<u>PEMETREXED DISODIUM - ALIMTA</u>					
021462 001	5344932	Jul 24, 2016 DS	DP		
<u>PREGABALIN - LYRICA</u>					
021446 001				>A> I-535	Jun 21, 2010
<u>PREGABALIN - LYRICA</u>					
021446 002				>A> I-535	Jun 21, 2010
<u>PREGABALIN - LYRICA</u>					
021446 003				>A> I-535	Jun 21, 2010
<u>PREGABALIN - LYRICA</u>					
021446 004				>A> I-535	Jun 21, 2010
<u>PREGABALIN - LYRICA</u>					
021446 005				>A> I-535	Jun 21, 2010
<u>PREGABALIN - LYRICA</u>					
021446 006				>A> I-535	Jun 21, 2010
<u>PREGABALIN - LYRICA</u>					
021446 007				>A> I-535	Jun 21, 2010
<u>PREGABALIN - LYRICA</u>					
021446 008				>A> I-535	Jun 21, 2010
<u>PROGESTERONE - ENDOMETRIN</u>					
022057 001				>A> NP	Jun 21, 2010
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>					
022047 001	>A> 4879288	Sep 26, 2011 DS	DP U-814	NDF	May 17, 2010
	>A> 5948437	May 28, 2017	DP U-814		
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>					
022047 002	>A> 4879288	Sep 26, 2011 DS	DP U-814	NDF	May 17, 2010
	>A> 5948437	May 28, 2017	DP U-814		
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>					
022047 003	>A> 4879288	Sep 26, 2011 DS	DP U-814	NDF	May 17, 2010
	>A> 5948437	May 28, 2017	DP U-814		
<u>QUETIAPINE FUMARATE - SEROQUEL XR</u>					
022047 004	>A> 4879288	Sep 26, 2011 DS	DP U-814	NDF	May 17, 2010
	>A> 5948437	May 28, 2017	DP U-814		

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES			EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RAMIPRIL - ALTACE</u>							
019901 001	5061722	Oct 29, 2008					
<u>RAMIPRIL - ALTACE</u>							
019901 002	5061722	Oct 29, 2008					
<u>RAMIPRIL - ALTACE</u>							
019901 003	5061722	Oct 29, 2008					
<u>RAMIPRIL - ALTACE</u>							
019901 004	5061722	Oct 29, 2008					
<u>RAMIPRIL - ALTACE</u>							
022021 001	5061722	Oct 29, 2008	DS	DP	U-185		
	5403856	Apr 04, 2012			U-71		
<u>RAMIPRIL - ALTACE</u>							
022021 002	5061722	Oct 29, 2008	DS	DP	U-185		
	5403856	Apr 04, 2012			U-71		
<u>RAMIPRIL - ALTACE</u>							
022021 003	5061722	Oct 29, 2008	DS	DP	U-185		
	5403856	Apr 04, 2012			U-71		
<u>RAMIPRIL - ALTACE</u>							
022021 004	5061722	Oct 29, 2008	DS	DP	U-185		
	5403856	Apr 04, 2012			U-71		
<u>RANITIDINE HYDROCHLORIDE - RANITIDINE HYDROCHLORIDE</u>							
076124 001						PC	Sep 15, 2007
<u>RANITIDINE HYDROCHLORIDE - ZANTAC 150</u>							
021698 001	>A> 5098715	Dec 20, 2010		DP			
<u>RANITIDINE HYDROCHLORIDE - ZANTAC 150</u>							
021698 002						>A> NP	Aug 31, 2007
<u>RETAPAMULIN - ALTABAX</u>							
022055 001	RE39128	Oct 27, 2018	DS	DP	U-805	NCE	Apr 12, 2012
<u>RISEDRONATE SODIUM - ACTONEL</u>							
020835 004	5583122	Dec 10, 2013	DS	DP	U-353	D-105	Apr 16, 2010
	6096342	Nov 22, 2011		DP	U-353	M-52	Jan 24, 2009
	6165513	Jun 10, 2018		DP			
<u>RISPERIDONE - RISPERDAL</u>							
020272 001	4804663	Dec 29, 2007			U-90	I-509	Oct 06, 2009
	4804663*PED	Jun 29, 2008				I-413	Dec 04, 2006
						I-412	Dec 04, 2006
						PED	Jun 04, 2007
						PED	Jun 04, 2007
						PED	Apr 06, 2010
<u>RISPERIDONE - RISPERDAL</u>							
020272 002	4804663	Dec 29, 2007			U-90	I-509	Oct 06, 2009
	4804663*PED	Jun 29, 2008				I-413	Dec 04, 2006
						I-412	Dec 04, 2006
						PED	Jun 04, 2007
						PED	Jun 04, 2007
						PED	Apr 06, 2010
<u>RISPERIDONE - RISPERDAL</u>							
020272 003	4804663	Dec 29, 2007			U-90	I-509	Oct 06, 2009
	4804663*PED	Jun 29, 2008				I-413	Dec 04, 2006
						I-412	Dec 04, 2006
						PED	Jun 04, 2007
						PED	Jun 04, 2007
						PED	Apr 06, 2010

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE		
<u>RISPERIDONE - RISPERDAL</u>							
020272 004	4804663	Dec 29, 2007	U-90	I-509	Oct 06, 2009		
	4804663*PED	Jun 29, 2008		I-413	Dec 04, 2006		
				I-412	Dec 04, 2006		
				PED	Jun 04, 2007		
				PED	Jun 04, 2007		
				PED	Apr 06, 2010		
<u>RISPERIDONE - RISPERDAL</u>							
020272 005	4804663	Dec 29, 2007	U-90	I-413	Dec 04, 2006		
	4804663*PED	Jun 29, 2008		I-412	Dec 04, 2006		
				PED	Jun 04, 2007		
				PED	Jun 04, 2007		
<u>RISPERIDONE - RISPERDAL</u>							
020272 007	4804663	Dec 29, 2007	U-90	I-509	Oct 06, 2009		
	4804663*PED	Jun 29, 2008		I-413	Dec 04, 2006		
				I-412	Dec 04, 2006		
				PED	Jun 04, 2007		
				PED	Jun 04, 2007		
				PED	Apr 06, 2010		
<u>RISPERIDONE - RISPERDAL</u>							
020272 008	4804663	Dec 29, 2007	U-90	I-509	Oct 06, 2009		
	4804663*PED	Jun 29, 2008		I-413	Dec 04, 2006		
				I-412	Dec 04, 2006		
				PED	Jun 04, 2007		
				PED	Jun 04, 2007		
				PED	Apr 06, 2010		
<u>RISPERIDONE - RISPERDAL</u>							
020588 001	4804663	Dec 29, 2007	U-90	I-509	Oct 06, 2009		
	4804663*PED	Jun 29, 2008		I-413	Dec 04, 2006		
	5453425	Jul 11, 2014		I-412	Dec 04, 2006		
	5453425*PED	Jan 11, 2015		PED	Apr 06, 2010		
	5616587	Jul 11, 2014		PED	Jun 04, 2007		
	5616587*PED	Jan 11, 2015		PED	Jun 04, 2007		
	RE39181	Jul 11, 2014	DP				
	RE39181*PED	Jan 11, 2015					
<u>RISPERIDONE - RISPERDAL</u>							
021444 001	4804663	Dec 29, 2007	DS	DP	U-516	I-509	Oct 06, 2009
	4804663*PED	Jun 29, 2008				I-413	Dec 04, 2006
	5648093	Jul 15, 2014		DP		I-412	Dec 04, 2006
	5648093*PED	Jan 15, 2015				PED	Jun 04, 2007
	6224905	Jun 10, 2017		DP		PED	Jun 04, 2007
	6244905*PED	Dec 10, 2017				PED	Apr 06, 2010
<u>RISPERIDONE - RISPERDAL</u>							
021444 002	4804663	Dec 29, 2007	DS	DP	U-516	I-509	Oct 06, 2009
	4804663*PED	Jun 29, 2008				I-413	Dec 04, 2006
	5648093	Jul 15, 2014		DP		I-412	Dec 04, 2006
	5648093*PED	Jan 15, 2015				PED	Jun 04, 2007
	6224905	Jun 10, 2017		DP		PED	Jun 04, 2007
	6244905*PED	Dec 10, 2017				PED	Apr 06, 2010
<u>RISPERIDONE - RISPERDAL</u>							
021444 003	4804663	Dec 29, 2007	DS	DP	U-516	I-509	Oct 06, 2009
	4804663*PED	Jun 29, 2008				I-413	Dec 04, 2006
	5648093	Jul 15, 2014		DP		I-412	Dec 04, 2006
	5648093*PED	Jan 15, 2015				PED	Apr 06, 2010
	6224905	Jun 10, 2017		DP		PED	Jun 04, 2007
	6244905*PED	Dec 10, 2017				PED	Jun 04, 2007

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RISPERIDONE - RISPERDAL</u>					
021444 004	4804663	Dec 29, 2007	DS DP	U-516	I-509 Oct 06, 2009
	4804663	Dec 29, 2007	DS DP	U-543	I-413 Dec 04, 2006
	4804663*PED	Jun 29, 2008			I-412 Dec 04, 2006
	5648093	Jul 15, 2014	DP		PED Jun 04, 2007
	5648093*PED	Jan 15, 2015			PED Jun 04, 2007
	6224905	Jun 10, 2017	DP		PED Apr 06, 2010
	6244905*PED	Dec 10, 2017			
<u>RISPERIDONE - RISPERDAL</u>					
021444 005	4804663	Dec 29, 2007	DS DP	U-516	I-509 Oct 06, 2009
	4804663	Dec 29, 2007	DS DP	U-543	I-413 Dec 04, 2006
	4804663*PED	Jun 29, 2008			I-412 Dec 04, 2006
	5648093	Jul 15, 2014	DP		PED Jun 04, 2007
	5648093*PED	Jan 15, 2015			PED Jun 04, 2007
	6224905	Jun 10, 2017	DP		PED Apr 06, 2010
	6244905*PED	Dec 10, 2017			
<u>RISPERIDONE - RISPERDAL CONSTA</u>					
021346 001	4804663	Dec 29, 2007			NDF Oct 29, 2006
	4804663*PED	Jun 29, 2008			PED Apr 29, 2007
	5688801	Nov 18, 2014			
	5688801*PED	May 18, 2015			
	5770231	Nov 19, 2013			
	5770231*PED	May 19, 2014			
	5792477	May 02, 2017			
	5792477*PED	Nov 02, 2017			
	5916598	May 02, 2017			
	5916598*PED	Nov 02, 2017			
	5965168	Nov 19, 2013			
	5965168*PED	May 19, 2014			
	6110503	May 02, 2017			
	6110503*PED	Nov 02, 2017			
	6110921	Nov 19, 2013			
	6110921*PED	May 19, 2014			
	6194006	Dec 30, 2018			
	6194006*PED	Jun 30, 2019			
	6264987	May 19, 2020			
	6264987*PED	Nov 19, 2020			
	6368632	Nov 19, 2013		U-543	
	6368632*PED	May 19, 2014			
	6379703	Dec 30, 2018	DP		
	6379703*PED	Jun 30, 2019			
	6379704	May 19, 2020	DP		
	6379704*PED	Nov 19, 2020			
	6403114	May 02, 2017			
	6403114*PED	Nov 02, 2017			
	6534092	May 19, 2020	DP		
	6534092*PED	Nov 19, 2020			
	6596316	Dec 30, 2018	DP		
	6596316*PED	Jun 30, 2019			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RISPERIDONE - RISPERDAL CONSTA</u>					
021346 002	4804663	Dec 29, 2007		NDF	Oct 29, 2006
	4804663*PED	Jun 29, 2008		PED	Apr 29, 2007
	5688801	Nov 18, 2014			
	5688801*PED	May 18, 2015			
	5770231	Nov 19, 2013			
	5770231*PED	May 19, 2014			
	5792477	May 02, 2017			
	5792477*PED	Nov 02, 2017			
	5916598	May 02, 2017			
	5916598*PED	Nov 02, 2017			
	5965168	Nov 19, 2013			
	5965168*PED	May 19, 2014			
	6110503	May 02, 2017			
	6110503*PED	Nov 02, 2017			
	6110921	Nov 19, 2013			
	6110921*PED	May 19, 2014			
	6194006	Dec 30, 2018			
	6194006*PED	Jun 30, 2019			
	6264987	May 19, 2020			
	6264987*PED	Nov 19, 2020			
	6368632	Nov 19, 2013		U-543	
	6368632*PED	May 19, 2014			
	6379703	Dec 30, 2018	DP		
	6379703*PED	Jun 30, 2019			
	6379704	May 19, 2020	DP		
	6379704*PED	Nov 19, 2020			
	6403114	May 02, 2017			
	6403114*PED	Nov 02, 2017			
	6534092	May 19, 2020	DP		
	6534092*PED	Nov 19, 2020			
	6596316	Dec 30, 2018	DP		
	6596316*PED	Jun 30, 2019			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>RISPERIDONE - RISPERDAL CONSTA</u>					
021346 003	4804663	Dec 29, 2007		NDF	Oct 29, 2006
	4804663*PED	Jun 29, 2008		PED	Apr 29, 2007
	5688801	Nov 18, 2014			
	5688801*PED	May 18, 2015			
	5770231	Nov 19, 2013			
	5770231*PED	May 19, 2014			
	5792477	May 02, 2017			
	5792477*PED	Nov 02, 2017			
	5916598	May 02, 2017			
	5916598*PED	Nov 02, 2017			
	5965168	Nov 19, 2013			
	5965168*PED	May 19, 2014			
	6110503	May 02, 2017			
	6110503*PED	Nov 02, 2017			
	6110921	Nov 19, 2013			
	6110921*PED	May 19, 2014			
	6194006	Dec 30, 2018			
	6194006*PED	Jun 30, 2019			
	6264987	May 19, 2020			
	6264987*PED	Nov 19, 2020			
	6368632	Nov 19, 2013		U-543	
	6368632*PED	May 19, 2014			
	6379703	Dec 30, 2018	DP		
	6379703*PED	Jun 30, 2019			
	6379704	May 19, 2020		DP	
	6379704*PED	Nov 19, 2020			
	6403114	May 02, 2017			
	6403114*PED	Nov 02, 2017			
	6534092	May 19, 2020		DP	
	6534092*PED	Nov 19, 2020			
	6596316	Dec 30, 2018		DP	
	6596316*PED	Jun 30, 2019			
<u>RISPERIDONE - RISPERDAL CONSTA</u>					
021346 004	4804663	Dec 29, 2007	DS DP	U-543	
	4804663*PED	Jun 29, 2008			
	5688801	Nov 18, 2014		U-543	
	5688801*PED	May 18, 2015			
	5770231	Nov 19, 2013		DP	
	5770231*PED	May 19, 2014			
	5792477	May 02, 2017		DP	
	5792477*PED	Nov 02, 2017			
	5916598	May 02, 2017		DP	
	5916598*PED	Nov 02, 2017			
	5965168	Nov 19, 2013		DP	
	5965168*PED	May 19, 2014			
	6110503	May 02, 2017		DP	
	6110503*PED	Nov 02, 2017			
	6110921	Nov 19, 2013		U-543	
	6110921*PED	May 19, 2014			
	6194006	Dec 30, 2018		DP	
	6194006*PED	Jun 30, 2019			
	6368632	Nov 19, 2013		U-543	
	6368632*PED	May 19, 2014			
	6379703	Dec 30, 2018		DP	
	6379703*PED	Jun 30, 2019			
	6403114	May 02, 2017		DP	
	6403114*PED	Nov 02, 2017			
	6596316	Dec 30, 2018		DP	
	6596316*PED	Jun 30, 2019			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>ROTIGOTINE - NEUPRO</u>					
021829 001				NCE	May 09, 2012
<u>ROTIGOTINE - NEUPRO</u>					
021829 002				NCE	May 09, 2012
<u>ROTIGOTINE - NEUPRO</u>					
021829 003				NCE	May 09, 2012
<u>SELEGILINE - EMSAM</u>					
021336 001	7150881	Jun 12, 2018	DS DP		
<u>SELEGILINE - EMSAM</u>					
021336 002	7150881	Jun 12, 2018	DS DP		
<u>SELEGILINE - EMSAM</u>					
021336 003	7150881	Jun 12, 2018	DS DP		
<u>SOMATROPIN RECOMBINANT - SAIZEN</u>					
019764 005				I-440	Aug 26, 2007
<u>SOMATROPIN RECOMBINANT - VALTROPIN</u>					
021905 001				NP	Apr 19, 2010
<u>TADALAFIL - CIALIS</u>					
021368 001	5859006	Nov 21, 2017	DS DP		
	7182958	Apr 26, 2020	DP	U-155	
<u>TADALAFIL - CIALIS</u>					
021368 002	5859006	Nov 21, 2017	DS DP		
	7182958	Apr 26, 2020	DP	U-155	
<u>TADALAFIL - CIALIS</u>					
021368 003	5859006	Nov 21, 2017	DS DP		
	7182958	Apr 26, 2020	DP	U-155	
<u>TEMSIROLIMUS - TORISEL</u>					
022088 001	>A> 5362718	Apr 18, 2014	DS DP	NCE	May 30, 2012
<u>TERIPARATIDE RECOMBINANT HUMAN - FORTEO</u>					
021318 001	7163684	Aug 19, 2019		U-790	
<u>THALIDOMIDE - THALOMID</u>					
020785 001	>A> 7230012	Dec 09, 2023	DP		
<u>THALIDOMIDE - THALOMID</u>					
020785 002	>A> 7230012	Dec 09, 2023	DP		
<u>THALIDOMIDE - THALOMID</u>					
020785 003	>A> 7230012	Dec 09, 2023	DP		
<u>THALIDOMIDE - THALOMID</u>					
020785 004	5629327	May 13, 2014		U-731	May 23, 2013
	6045501	Aug 28, 2018		U-731	
	6235756	Mar 01, 2013		U-731	
	6315720	Oct 23, 2020		U-731	
	6561976	Aug 28, 2018		U-731	
	6561977	Oct 23, 2020		U-731	
	6755784	Oct 23, 2020		U-731	
	6869399	Oct 23, 2020		U-731	
	6908432	Aug 28, 2018		U-731	
	7141018	Oct 23, 2020		U-731	
<u>TIMOLOL MALEATE - ISTALOL</u>					
021516 001	>A> 6335335	Nov 02, 2018	DP		
<u>TIMOLOL MALEATE - TIMOLOL MALEATE</u>					
020963 001	6174524	Mar 26, 2019			
	6174524*PED	Sep 26, 2019			
<u>TIMOLOL MALEATE - TIMOLOL MALEATE</u>					
020963 002	6174524	Mar 26, 2019			
	6174524*PED	Sep 26, 2019			

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
<u>TINIDAZOLE - TINDAMAX</u>					
021618 001				I-532	May 21, 2010
<u>TINIDAZOLE - TINDAMAX</u>					
021618 002				I-532	May 21, 2010
<u>ZICONOTIDE - PRIALT</u>					
021060 001	>A> 5364842	Dec 30, 2016	U-55		
	>A> 5364842	Dec 30, 2016	U-48		
<u>ZICONOTIDE - PRIALT</u>					
021060 002	>A> 5364842	Dec 30, 2016	U-55		
	>A> 5364842	Dec 30, 2016	U-48		
<u>ZICONOTIDE - PRIALT</u>					
021060 003	>A> 5364842	Dec 30, 2016	U-55		
	>A> 5364842	Dec 30, 2016	U-48		
<u>ZICONOTIDE - PRIALT</u>					
021060 004	>A> 5364842	Dec 30, 2016	U-55		
	>A> 5364842	Dec 30, 2016	U-48		
<u>ZILEUTON - ZYFLO CR</u>					
022052 001	>A> 4873259	Dec 09, 2010	DS		
	>A> 5422123	Jun 06, 2012	DP		
<u>ZIPRASIDONE HYDROCHLORIDE - GEODON</u>					
021483 001	7175855	May 18, 2020	DP		
<u>ZOLEDRONIC ACID - RECLAST</u>					
021817 001				NP	Apr 16, 2010
<u>ZOLMITRIPTAN - ZOMIG</u>					
021450 004	>A> 7220767	Nov 28, 2020	DP		
	>A> 7220767*PED	May 28, 2021			
<u>ZOLPIDEM TARTRATE - AMBIEN</u>					
019908 001				M-54 PED	Mar 29, 2010 Sep 29, 2010
<u>ZOLPIDEM TARTRATE - AMBIEN</u>					
019908 002				M-54 PED	Mar 29, 2010 Sep 29, 2010

PATENT & EXCLUSIVITY DRUG PRODUCT LIST - CUMULATIVE SUPPLMENT 6 - June 2007

See report footnote for information regarding List content

APPL/PROD NO	PATENT NO	PATENT EXPIRATION DATE	PATENT CODES	EXCLUSIVITY CODE(S)	EXCLUSIVITY EXPIRATION DATE
-----------------	-----------	---------------------------	-----------------	------------------------	--------------------------------

Footnote:

1. Patents are published upon receipt by the Orange Book Staff and may not reflect the official receipt date as described in 21 CFR 314.53(d)(5).
2. Patents submitted on FDA Form 3542 and listed after August 18, 2003 will have one to three patent codes indicating specific patent claims as submitted by the sponsor:
 - DS = Drug Substance claim
 - DP = Drug Product claim
 - U and number = Method of Use claim (may be multiple). Specific Method of use claims are listed at <http://www.fda.gov/cder/orange/patex.htm>
3. Patents listed prior to August 18, 2003 are flagged with method of use claims only as applicable and submitted by the sponsor. They may not be flagged with respect to other claims which may apply.
4. *PED and PED represent pediatric exclusivity. Patents with pediatric exclusivity granted after August 18, 2003 will be indicated with *PED as was done prior to August 18, 2003. Patents with *PED added after August 18, 2003 will not contain any information relative to the patent itself other than the *PED extension. Information related specifically to the patent will be conveyed on the original patent only.
5. *** U.S. Patent Nos. RE 36481 and RE 36520 are being relisted for Zocor (NDA 19-766) pursuant to the decision and related order in *Ranbaxy Labs. v. Leavitt*, No. 05-1838 (D.D.C. April 30, 2006). The '481 and '520 patents will remain listed in Approved Drug Products with Therapeutic Equivalence Evaluations until any applicable periods of exclusivity pursuant to section 505(j)(5)(B)(iv) of the Federal Food, Drug, and Cosmetic Act have been triggered and run, unless the agency's appeal of the decision to the U.S. Court of Appeals for the District of Columbia is decided in the agency's favor before the exclusivity periods have expired. While the patents remain listed, any new or pending ANDA referencing Zocor must contain patent certifications to these patents. For additional information on this matter, please refer to docket Nos. 2005P-0008 and 2005P-0046.

PATENT AND EXCLUSIVITY TERMS

Due to space limitations in the patent and exclusivity columns, abbreviations and references have been developed. Refer to the APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 27th 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>