

CUMULATIVE
SUPPLEMENT 11
NOV'99

APPROVED DRUG PRODUCTS

WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS

19TH EDITION

U.S. DEPARTMENT OF HEALTH AND HUMAN SERVICES
PUBLIC HEALTH SERVICE
FOOD AND DRUG ADMINISTRATION
CENTER FOR DRUG EVALUATION AND RESEARCH
OFFICE OF INFORMATION TECHNOLOGY
DIVISION OF DATA MANAGEMENT AND SERVICES

1999

RM
301.45
.A66
1999
Nov
Suppl



RM301.45 .A66 1999 Nov Suppl

Approved drug products with
therapeutic equivalence

C:355661 M:174736 O:12937927

Prepared By
Division of Data Management and Services
Office of Information Technology
Center for Drug Evaluation and Research
Food and Drug Administration

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New 20th Edition



**APPROVED
DRUG PRODUCTS**

WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS

**20TH EDITION
2000**

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- Discontinued Drug Product List
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- Drug Products Which Must Demonstrate in vivo Bioavailability Only if Product Fails to Achieve Adequate Dissolution
- Patent and Exclusivity Information

See Subscription Form Inside Back Cover

APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS

19TH EDITION

Cumulative Supplement 11

NOVEMBER 1999

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APPROVED DRUG PRODUCTS
with
THERAPEUTIC EQUIVALENCE EVALUATIONS

19TH EDITION

CUMULATIVE SUPPLEMENT 11
NOVEMBER 1999

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, 19th 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and products that have never been marketed, have been discontinued from marketing or that have had their approvals withdrawn for other than safety or efficacy reasons.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.

The Patent and Exclusivity Lists are arranged in alphabetical order by active ingredient name. For those products with multiple active ingredients, only the first active ingredient (in alphabetical sort) will appear. In addition, the trade name will be displayed to the right of the active ingredient name for each product. 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. Use patents are indicated with the symbol "U" followed by a number representing a specific use. Patent and Exclusivity information for a specific drug is indicated by an abbreviation followed by the date upon which the patent and/or exclusivity expires. Refer to the Patent and Exclusivity Terms section in the Patent and Exclusivity Information Addendum for an explanation of all codes and abbreviations. Information regarding drug patents and exclusivity for an approved product will appear in this addendum as it is received and may not correspond with the month the drug product is approved and published in the Rx/OTC sections of the Cumulative Supplement.

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 place an asterisk (*) 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. [Strength(s) which already exist in the List will not be repeated for context.]

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.)

New additions to the Prescription Drug Product List, OTC Drug Product List, and the Patent and Exclusivity Data are indicated by the symbol >**ADD**> to the left of the line on which new information exists. The >**ADD**> 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 >**DLT**> (DELETE) to the left of the line. The >**DLT**> symbol is dropped in subsequent Cumulative Supplements for that item. The shaded print remains in the Prescription Drug Product List and OTC Drug Product Lists in all Cumulative Supplements (hard copy only) for this edition.

Products that have never been marketed, 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 19th Edition List will then be added to the "Discontinued Drug Product List" appearing in the 20th Edition.

1.2 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

APPLICANT NAME CHANGES

FORMER APPLICANT NAME (FORMER ABBREVIATED NAME)

NEW APPLICANT NAME (NEW ABBREVIATED NAME)

ASTRA PHARMACEUTICALS LP
(ASTRA PHARMS)

ASTRA ZENECA LP
(ASTRA ZENECA)

PENEDERM INC
(PENEDERM)

BERTEK PHARMACEUTICALS
(BERTEK PHARMS)

SANOFI PHARMACEUTICALS INC
(SANOFI)

SANOFI SYNTHELABO INC
(SANOFI SYNTHELABO)

1.3 DICLOFENAC SODIUM OPHTHALMIC SOLUTION 0.1%

Two NDAs have been approved for diclofenac sodium ophthalmic solution 0.1% (DSOS), (1) Ciba's NDA 20-037 for Voltaren and (2) Falcon Pharms' (Alcon) NDA 20-809 for DSOS. Alcon was required to do a study comparing their DSOS to Voltaren and to a placebo control in post cataract surgical inflammation. This study was necessary to demonstrate that the different formulation of the Alcon drug product did not affect the safety and/or effectiveness of the proposed drug product for this indication. Prior to the approval of Alcon's DSOS Ciba did clinical studies and was approved for two additional indications for the temporary relief of pain and photophobia in patients undergoing corneal refractive surgery. Three years of Waxman-Hatch marketing exclusivity was granted to Ciba for these two new uses.

Since the treatment of pain has a different site of action than the anti-inflammatory or photophobia indications the Agency did not have information to support a recommendation that the Alcon and Ciba DSOS are therapeutically equivalent for the treatment of pain. The designation of therapeutic equivalence at this time applies only to the anti-inflammatory indication. The therapeutic equivalence designation will apply to the photophobia indication upon expiration of Ciba's marketing exclusivity.

1.4 AVAILABILITY OF THE EDITION

The 19th Edition of the Orange Book and its monthly cumulative supplements are available by subscription from the Government Printing Office:

Superintendent of Documents
Government Printing Office
P.O. Box 371954
Pittsburgh, PA 15250-7954

The telephone number to charge your subscription is 202-512-1800. The cost is \$90.00 annually.

The Approved Drug Products with Therapeutic Equivalence Evaluation (Orange Book) and related drug information is also available on the Internet at the Food and Drug Administration, Center for Drug Evaluation and Research, Drug Info page.

There is an Electronic Orange Book Query (EOB) at <http://www.fda.gov/cder/ob>. The Query provides searching of the approved drug list by active ingredient, proprietary name, applicant holder or applicant 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. The data is updated concurrently with the publication of the annual edition or monthly cumulative supplements.

The Internet version of the hard copy Orange Book annual edition is at <http://www.fda.gov/cder/orange/adp.htm>.

The Internet version of the hard copy monthly supplement is at <http://www.fda.gov/cder/orange/supplement/cspreface.htm>. Changes to the annual edition are listed separately by month.

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

The 19th annual edition of the 1998 Orange Book Patent and Exclusivity List is at <http://www.fda.gov/cder/orange/19bookpub.pdf>.

The current year Patent and Exclusivity cumulative supplement list that denotes the current month additions is at <http://www.fda.gov/cder/orange/supplement/patents.pdf>.

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 1998) 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 1998</u>	<u>MAR 1999</u>	<u>JUN 1999</u>	<u>SEP 1999</u>
DRUG PRODUCTS LISTED	9923	9975	10009	9939
SINGLE SOURCE	2504 (25.2%)	2520 (25.3%)	2523 (25.2%)	2557 (25.7%)
MULTISOURCE	7308 (73.6%)	7344 (73.6%)	7375 (73.7%)	7271 (73.2%)
THERAPEUTICALLY EQUIVALENT	6934 (69.9%)	6969 (69.9%)	7012 (70.1%)	6923 (69.7%)
NOT THERAPEUTICALLY EQUIVALENT	374 (3.8%)	375 (3.8%)	363 (3.6%)	348 (3.5%)
EXCEPTIONS ¹	111 (1.1%)	111 (1.1%)	111 (1.1%)	111 (1.1%)
NEW MOLECULAR ENTITIES APPROVED	10	3	5	13
NUMBER OF APPLICANTS	563	570	568	572

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

PRESCRIPTION DRUG PRODUCT LIST
19TH EDITION

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN'99 - NOV'99

1

ACETAMINOPHEN; BUTALBITAL; CAFFEINE

CAPSULE; ORAL

AB BUTALBITAL, ACETAMINOPHEN, CAFFEINE
GRAHAM DM 325MG; 50MG; 40MG

N88758 001
MAR 27, 1985
N88758 001
MAR 27, 1985

@ MALLINCKRODT 325MG; 50MG; 40MG

AB FEMCET
MALLINCKRODT 325MG; 50MG; 40MG

N89102 001
JUN 19, 1985
N89102 001
JUN 19, 1985

@ 325MG; 50MG; 40MG

AB TRIAD
MALLINCKRODT 325MG; 50MG; 40MG

N89023 001
JUN 19, 1985
N89023 001
JUN 19, 1985

@ 325MG; 50MG; 40MG

TABLET; ORAL
BUTALBITAL, ACETAMINOPHEN, AND CAFFEINE
WEST WARD 500MG; 50MG; 40MG

N40336 001
AUG 18, 1999

ACETAMINOPHEN; CAFFEINE; DIHYDROCODEINE BITARTRATE

TABLET; ORAL

ACETAMINOPHEN, CAFFEINE, AND DIHYDROCODEINE BITARTRATE
+ MIKART 712.8MG; 60MG; 32MG

N40316 001
APR 28, 1999

ACETAMINOPHEN; HYDROCODONE BITARTRATE

CAPSULE; ORAL

AA ALLAY
NORTON HN 500MG; 5MG

N89907 001
JAN 13, 1989

AA ZENITH GOLDLINE 500MG; 5MG

N89907 001
JAN 13, 1989

AA HYDROCODONE BITARTRATE AND ACETAMINOPHEN
MALLINCKRODT 500MG; 5MG

N88956 001
JUL 19, 1985

AA ZYDONE
MALLINCKRODT 500MG; 5MG

N88956 001
JUL 19, 1985

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE

CAPSULE; ORAL

AA OXYCODONE AND ACETAMINOPHEN
DURAMED 500MG; 5MG

N40289 001
MAR 16, 1999

TABLET; ORAL

AA OXYCODONE AND ACETAMINOPHEN
AMIDE PHARM 325MG; 5MG

N40203 001
MAR 15, 1999

PERCOCET
ENDO PHARMS 325MG; 2.5MG

N40330 001
JUN 25, 1999

AA ENDO PHARMS 325MG; 5MG

N40330 002
JUN 25, 1999

+ 325MG; 2.5MG

+ 500MG; 7.5MG

+ 650MG; 10MG

N40341 001
JUL 26, 1999

N40341 002
JUL 26, 1999

ACETAMINOPHEN; PROPOXYPHENE NAPSYLATE

TABLET; ORAL

AB PROPACET 100
TEVA 650MG; 100MG

N70107 001
JUN 12, 1985

@ 650MG; 100MG

N70107 001
JUN 12, 1985

ACITRETIN

CAPSULE; ORAL

SORIATANE
HLR 10MG

N19821 001
OCT 28, 1996

+ 25MG

+ 10MG

+ 25MG

N19821 001
OCT 28, 1996

N19821 002
OCT 28, 1996

ACYCLOVIR

CAPSULE; ORAL

ACYCLOVIR

STASON

200MGN75090 001
JAN 26, 1999
> ADD >
> ADDAN

SOLUTION; INHALATION

ALBUTEROL SULFATE

STERIPAK

EQ 0.083% BASEN75343 001
NOV 09, 1999

TABLET; ORAL

ACYCLOVIR

CARLSEAD

400MG

N75382 001

APR 30, 1999

N75382 002

APR 30, 1999

ALBUTEROL SULFATE

UDL

EQ 2MG BASE/5MLN75262 001
MAR 30, 1999

VENTOLIN

AA + GLAXO WELLCOME

EQ 2MG BASE/5MLN13621 001
JUN 10, 1987

©

EQ 2MG BASE/5MLN13621 001
JUN 10, 1987ACYCLOVIR SODIUM

INJECTABLE; INJECTION

ACYCLOVIR

ABBOTT

EQ 50MG BASE/MLN75114 001
JUL 26, 1999ACYCLOVIR SODIUM
AP + AM PHARM PARTNERSEQ 50MG BASE/ML

N74930 001

MAY 13, 1998

N74930 001

MAY 13, 1998

N75065 001

FEB 25, 1999

EQ 50MG BASE/ML

AP MERIDIAN MEDCL TECHN

ALCOHOL; DEXTROSE

INJECTABLE; INJECTION

ALCOHOL 5% IN DEXTROSE 5% IN WATER

BAXTER HILTCARE

5ML/100ML; 5GM/100ML5ML/100ML; 5GM/100MLN83256 001
N83256 001ALFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ALFENTA

AP + AKORN

EQ 0.5MG BASE/MLN19353 001
DEC 29, 1986

AEROSOL, METERED; INHALATION

ALBUTEROL

MEDEVA

0.09MG/INH

N72273 001

AUG 14, 1996

N72273 001

AUG 14, 1996

0.09MG/INH

AB MEDEVA PHARMS MA

ALBUTEROLALBUTEROL SULFATE

SOLUTION; INHALATION

ALBUTEROL SULFATE

HI TECH PHARMA

EQ 0.083% BASE

N75063 001

FEB 09, 1999

N75394 001

NOV 22, 1999

AB MORTON GROVE

EQ 0.083% BASEEQ 0.1% BASEN20886 001
FEB 02, 1999ALITRETINOLIN

GEL; TOPICAL

PANRETIN

+ LIGAND

> ADD >
> ADD >

ALLOPURINOLTABLET; ORAL

AB ZYLOPRIM
AB FARO PHARMS
AB +
AB GLAXO WELLCOME
AB *

100MG
300MG
100MG
300MG

N16084 001
N16084 002
N16084 001
N16084 002

ALLOPURINOL SODIUMINJECTABLE; INJECTION

ALOPRIM
+ CATALYTICA PHARMS

N20298 001
MAY 17, 1996

EQ 500MG BASE/VIAL

ZYLOPRIM
* CATALYTICA PHARMS

N20298 001
MAY 17, 1996

EQ 500MG BASE/VIAL

ALPHA-TOCOPHEROL; ASCORBIC ACID; BIOTIN, D-; CHOLECALCIFEROL;
CYANOCOBALAMIN; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID;
PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A

INJECTABLE; INJECTION
CERNEVIT-12
+ BAXTER HLTHCARE

11.2 IU/VIAL; 125MG/VIAL; 60 UGM/VIAL;
200 IU/VIAL; 5.5MG/VIAL; 414 UGM/VIAL;
46MG/VIAL; 17.25MG/VIAL; 4.53MG/VIAL;
4.14MG/VIAL; 3.51MG/VIAL;
3,500 IU/VIAL

N20924 001
APR 06, 1999

ALPRAZOLAM

CONCENTRATE; ORAL
ALPRAZOLAM
ROXANE

1MG/ML
1MG/ML
N74312 001
OCT 31, 1993
N74312 001
OCT 31, 1993

SOLUTION; ORAL
ALPRAZOLAM
ROXANE

N74314 001
OCT 31, 1993

0.5MG/5ML

ALPRAZOLAM

SOLUTION; ORAL
ALPRAZOLAM
+ ROXANE

0.5MG/5ML

N74314 001
OCT 31, 1993

TABLET; ORAL
ALPRAZOLAM
ALPHAPHARM

2MG

N74046 004
MAY 07, 1997

ALPROSTADILINJECTABLE; INJECTION

ALPROSTADIL
AP GENSLA SICOR PHARMS

0.5MG/ML

N75196 001
APR 30, 1999

AMCINONIDE

CREAM; TOPICAL
CYCLOCORT
@ FUJISAWA HLTHCARE

0.025%
0.1%
0.025%
0.1%

N18116 001
N18116 002
N18116 001
N18116 002

LOTION; TOPICAL
CYCLOCORT

0.1%

N19729 001
JUN 13, 1988
N19729 001
JUN 13, 1988

0.1%

OINTMENT; TOPICAL
CYCLOCORT

0.1%
0.1%

N18498 001
N18498 001

AMIFOSTINE

INJECTABLE; INJECTION
ETHYOL
US BIOSCIENCE

375MG/VIAL

N20221 002
SEP 10, 1999

AMINO ACIDSINJECTABLE; INJECTION

AMINESS 5.2% ESSENTIAL AMINO ACIDS W/ HISTADINE

FRESENIUS KABI 5.2%

N18901 001

APR 06, 1984

N18901 001

APR 06, 1984

NEOPHAM 6.4%

@ FRESENIUS KABI

N18792 001

JAN 17, 1984

N18792 001

JAN 17, 1984

NOVAMINE 11.4%

FRESENIUS KABI

N17957 003

AUG 09, 1982

N17957 003

AUG 09, 1982

NOVAMINE 15%

FRESENIUS KABI

N17957 004

NOV 28, 1986

N17957 004

NOV 28, 1986

NOVAMINE 8.5%

@ FRESENIUS KABI

N17957 002

AUG 09, 1982

N17957 002

AUG 09, 1982

@ PHARMACIA AND UPJOHN 8.5%

AMINO ACIDS; MAGNESIUM CHLORIDE; POTASSIUM ACETATE; POTASSIUM CHLORIDE; SODIUM ACETATEINJECTABLE; INJECTION

VEINAMINE 8%

@ FRESENIUS KABI

8%; 61MG/100ML; 211MG/100ML;

56MG/100ML; 388MG/100ML

8%; 61MG/100ML; 211MG/100ML;

56MG/100ML; 388MG/100ML

N17957 001

AMINOPHYLLINEINJECTABLE; INJECTION

AMINOPHYLLINE

SMITH AND NEPHEW

25MG/ML

N88749 001

MAY 30, 1985

N88749 001

MAY 30, 1985

25MG/ML

AMIODARONE HYDROCHLORIDETABLET; ORAL

AMIODARONE HCL

ALPHAPHARM

200MG

N75188 001

FEB 24, 1999

N74895 001

APR 16, 1999

200MG

AMITRIPTYLINE HYDROCHLORIDEINJECTABLE; INJECTION

AMITRIPTYLINE HCL

@ STERIS

10MG/ML

10MG/ML

N85594 001

N85594 001

N12704 001

N12704 001

10MG/ML

10MG/ML

10MG/ML

10MG/ML

ELAVIL

@ ZENECA

+

TABLET; ORAL

AMITRIPTYLINE HCL

COFLEY PHARM

10MG

25MG

50MG

75MG

100MG

150MG

10MG

25MG

50MG

75MG

100MG

150MG

10MG

25MG

50MG

75MG

100MG

150MG

10MG

25MG

50MG

75MG

100MG

150MG

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

N88421 001

AMITRIPTYLINE HYDROCHLORIDE

TABLET; ORAL

AB ENDER
AB ROCHE
AB

50MG
75MG
100MG
10MG
25MG
50MG
75MG
100MG

@
@
@
@
@

N83639 003
N83639 004
N83639 005
N83639 001
N83639 002
N83639 003
N83639 004
N83639 005

INJECTABLE, LIPID COMPLEX; INJECTION
AMPHOTEC
+ ALZA

50MG/VIAL
100MG/VIAL
50MG/VIAL
100MG/VIAL
N50729 001
NOV 22, 1996
N50729 002
NOV 22, 1996
N50729 001
NOV 22, 1996
N50729 002
NOV 22, 1996

AMOXICILLIN

CAPSULE; ORAL

AB AMOXICILLIN
AB RANBAXY

250MG
500MG

N65016 001
APR 08, 1999
N65016 002
APR 08, 1999

CAPSULE; ORAL
AGENERASE
GLAXO WELLCOME

50MG
150MG
N21007 001
APR 15, 1999
N21007 002
APR 15, 1999

POWDER FOR RECONSTITUTION; ORAL
AMOXIL

SMITHKLINE BEECHAM
+
200MG/5ML
400MG/5ML

N50760 001
APR 15, 1999
N50760 002
APR 15, 1999

SOLUTION; ORAL
AGENERASE
+ GLAXO WELLCOME

15MG/ML
N21039 001
APR 15, 1999

TABLET; ORAL

AB AMOXIL

500MG
500MG

N50754 002
JUL 10, 1998
N50754 002
JUL 10, 1998

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC
ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE;
RIBOFLAVIN; THIAMINE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION
KABIVITE PED F + W KIT
+ FRESENIUS KABI

N/A, 80MG/VIAL; N/A, 0.02MG/VIAL; N/A,
0.001MG/VIAL; 400 IU/10ML, N/A, N/A,
0.14MG/VIAL; N/A, 17MG/VIAL; N/A,
5MG/VIAL; 0.2MG/10ML, N/A; N/A,
1MG/VIAL; N/A, 1.4MG/VIAL; N/A,
1.2MG/VIAL; EQ 2,300 UNITS BASE/10ML,
N/A; 7 IU/10ML, N/A
N20176 001
DEC 29, 1993

TABLET, CHEWABLE; ORAL

AB AMOXIL
AB SMITHKLINE BEECHAM

200MG
400MG

N50761 001
APR 15, 1999
N50761 002
APR 15, 1999

> DLT >
> DLT >
> ADD >
> ADD >

ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PANTOTHENIC ACID; PHYTONADIONE; PYRIDOXINE; RIBOFLAVIN; THIAMINE; VITAMIN A PALMITATE; VITAMIN E

INJECTABLE; INJECTION

KABIVITE PED F + W KIT

* PHARMACIA AND UPJOHN

N/A, 80MG/VIAL, N/A, 0.02MG/VIAL, N/A, 0.001MG/VIAL; 400 IU/10ML, N/A, N/A, 0.14MG/VIAL, N/A, 1.7MG/VIAL, N/A, 5MG/VIAL, 0.2MG/10ML, N/A, N/A, 1MG/VIAL, N/A, 1.4MG/VIAL, N/A, 1.2MG/VIAL; EQ 2.300 UNITS BASE/10ML, N/A, 7 IU/10ML, N/A
N20176 001
DEC 29, 1993

ASPIRIN; BUTALBITAL; CAFFEINE; CODEINE PHOSPHATE

CAPSULE; ORAL

BUTALBITAL, ASPIRIN, CAFFEINE, AND CODEINE PHOSPHATE

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

ASPIRIN; CAFFEINE; ORPHENADRINE CITRATE

TABLET; ORAL

ORPHENADRINE CITRATE, ASPIRIN, AND CAFFEINE

AB STEVENS J 385MG; 30MG; 25MG N74988 001
APR 30, 1999
AB 770MG; 60MG; 50MG N74988 002
APR 30, 1999

ASPIRIN; DIPYRIDAMOLE

CAPSULE, EXTENDED RELEASE; ORAL

AGGRENOL + BOEHRINGER INGELHEIM 25MG; 200MG

> ADD >
> ADD >
> ADD >
> ADD >
> ADD >

ASTEMIZOLE

TABLET; ORAL

HISMANAL

* JANSSEN

10MG
N19402 001
DEC 29, 1988

ATENOLOL

TABLET; ORAL

ATENOLOL

APOTHECON

AB 50MG N73317 001
MAR 20, 1992
AB 100MG N73318 001
MAR 20, 1992
@ 50MG N73317 001
MAR 20, 1992
@ 100MG N73318 001
MAR 20, 1992
MYLAN 25MG N73457 002
APR 26, 1999

ATACURIUM BESYLATE

INJECTABLE; INJECTION

TRACRIUM

+ ABBOTT

10MG/ML

N18831 002
JUN 20, 1985
N18831 002
JUN 20, 1985

AB GLAXO WELLCOME

10MG/ML

TRACRIUM PRESERVATIVE FREE

+ ABBOTT

10MG/ML

N18831 001
NOV 23, 1983
N18831 001
NOV 23, 1983

AB GLAXO WELLCOME

10MG/ML

ATROPINE SULFATE; DIPHENOXYLATE HYDROCHLORIDE

TABLET; ORAL

DIPHENOXYLATE HCL AND ATROPINE SULFATE

PHARMERXAL 0.025MG; 2.5MG

R AND S PHARMA 0.025MG; 2.5MG

DIPHENOXYLATE HCL W/ ATROPINE SULFATE

ZENITH GOLDLINE 0.025MG; 2.5MG

0.025MG; 2.5MG

N85035 001
N85035 001
N86727 001
N86727 001

AZATHIOPRINE

TABLET; ORAL

AZATHIOPRINE

APPLIED ANAL

50MG

N75252 001
JUN 07, 1999

IMURAN

+ FARO PHARMS

50MG

N16324 001

AZATHIOPRINE

TABLET; ORAL

IMURAN
@ FARO PHARMS
25MG
AB * GLAXO WELLCOME
50MG
25MG

N16324 002
N16324 001
N16324 002

TABLET; ORAL
CORZIDE
+ APOTHECON

5MG;80MG

N18647 002
MAY 25, 1983
N18647 001
MAY 25, 1983

BRISTOL MYERS SQUIBB
5MG;40MG

5MG;80MG

N18647 002
MAY 25, 1983

AZATHIOPRINE SODIUM

INJECTABLE; INJECTION

IMURAN
+ FARO PHARMS
EQ 100MG BASE/VIAL
AB * GLAXO WELLCOME
EQ 100MG BASE/VIAL

N17391 001
N17391 001

BENZYL PENICILLOYL-POLYLYSINE

INJECTABLE; INJECTION

PRE-PEN
* BAYER
+ HOLLISTER STIER LABS 60 UMOLAR

N50114 001
N50114 001

AZITHROMYCIN DIHYDRATE; TROVAFLOXACIN MESYLATE

POWDER FOR RECONSTITUTION, TABLET; ORAL

TRIOVAN/ZITHROMAX COMPLIANCE PAK
* PFIZER
EQ 1GM BASE, N/A; N/A;
EQ 100MG BASE

N50762 001
DEC 18, 1998

@
EQ 1GM BASE, N/A; N/A;
EQ 100MG BASE

N50762 001
DEC 18, 1998

BETAMETHASONE DIPROPIONATE

ointment, augmented; TOPICAL

BETAMETHASONE DIPROPIONATE
AB ALTANA
EQ 0.05% BASE

N75373 001
JUN 22, 1999

BACITRACIN; HYDROCORTISONE ACETATE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

ointment; OPHTHALMIC

BACITRACIN-NEOMYCIN-POLYMYXIN W/ HYDROCORTISONE ACETATE
AT * ALTANA
400 UNITS/GM; 1%; EQ 3.5MG BASE/GM;
10,000 UNITS/GM
N60731 002

@
400 UNITS/GM; 1%; EQ 3.5MG BASE/GM;
10,000 UNITS/GM

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

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

EQ 0.12% BASE

N20934 001
FEB 28, 1999

BENDROFLUMETHIAZIDE; NADOLOL

TABLET; ORAL

CORZIDE
+ APOTHECON

5MG;40MG
N18647 001
MAY 25, 1983

N75541 001
OCT 22, 1999
N75541 002
OCT 22, 1999

N19507 001
OCT 27, 1989
N19507 002
OCT 27, 1989

BETAXOLOL HYDROCHLORIDE

TABLET; ORAL
KERLONE
LOREX

10MG
20MG

N19507 001
OCT 27, 1989
N19507 002
OCT 27, 1989

BRETYLIUM TOSYLATE

INJECTABLE; INJECTION
BRETYLIUM TOSYLATE
ASTRA PHARMS

50MG/ML
50MG/ML

N71153 001
AUG 10, 1987
N71153 001
AUG 10, 1987

BUDESONIDE

SPRAY, METERED; NASAL
RHINOCORT
ASTRAZENECA

0.032MG/INH
0.064MG/INH

N20746 001
OCT 01, 1999
N20746 002
OCT 01, 1999

BUPROPION HYDROCHLORIDE

TABLET; ORAL
BUPROPION HCL
TEVA

75MG
100MG

N75310 001
NOV 29, 1999
N75310 002
NOV 29, 1999

WELLBUTRIN
GLAXO WELLCOME

75MG
100MG

N18644 002
DEC 30, 1985
N18644 003
DEC 30, 1985
N18644 002
DEC 30, 1985
N18644 003
DEC 30, 1985

BUPROPION HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
WELLBUTRIN
+ GLAXO WELLCOME

50MG
100MG
150MG

N20358 001
OCT 04, 1996
N20358 002
OCT 04, 1996
N20358 003
OCT 04, 1996

WELLBUTRIN SR
+ GLAXO WELLCOME

50MG
100MG
150MG

N20358 001
OCT 04, 1996
N20358 002
OCT 04, 1996
N20358 003
OCT 04, 1996

BUSULFAN

INJECTABLE; INJECTION
BUSULFEX
+ ORPHAN MEDCL

6MG/ML

N20954 001
FEB 04, 1999

BUTOCONAZOLE NITRATE

CREAM; VAGINAL
FENSTAT
+ SYNTEX

2%
2%

N19215 001
NOV 25, 1985
N19215 001
NOV 25, 1985

FEMSTAT ONE
+ KV PHARM

2%

N19881 001
FEB 07, 1997
N19881 001
FEB 07, 1997

+ SYNTEX

2%

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION
BUTORPHANOL TARTRATE
MERIDIAN MEDCL TECHN

1MG/ML

N75342 001
NOV 04, 1999

> ADD >
> ADD >

> ADD >
> ADD >
> ADD >
> ADD >
> DLT >
> DLT >
> DLT >

BUTORPHANOL TARTRATE

INJECTABLE; INJECTION

BUTORPHANOL TARTRATE

> ADD > AP MERIDIAN MEDCL TECHN 2MG/ML

> ADD

N75342 002
NOV 04, 1999

CAFFEINE CITRATE

SOLUTION; INTRAVENOUS, ORAL

CAFCIT

+ OPR DEVELOP LP

EQ 10MG BASE/ML

N20793 001
SEP 21, 1999

CALCIUM CHLORIDE; DEXTROSE; GLUTATHIONE DISULFIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM BICARBONATE; SODIUM CHLORIDE; SODIUM PHOSPHATE

SOLUTION; IRRIGATION

ENDOSOL EXTRA

AKORN

0.154MG/ML; 0.92MG/ML; 0.184MG/ML;
0.2MG/ML; 0.38MG/ML; 2.1MG/ML;

7.14MG/ML; 0.42MG/ML

N20079 001
NOV 27, 1991

AT ALLERGAN

0.154MG/ML; 0.92MG/ML; 0.184MG/ML;
0.2MG/ML; 0.38MG/ML; 2.1MG/ML;

7.14MG/ML; 0.42MG/ML

N20079 001
NOV 27, 1991

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE

INJECTABLE; INJECTION

ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN

37MG/100ML; 5GM/100ML; 31MG/100ML;
120MG/100ML; 330MG/100ML;

88MG/100ML

37MG/100ML; 5GM/100ML; 31MG/100ML;

120MG/100ML; 330MG/100ML;

88MG/100ML

N18271 001

@

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

INJECTABLE; INJECTION

ISOLYTE R W/ DEXTROSE 5% IN PLASTIC CONTAINER

B BRAUN

35MG/100ML; 5GM/100ML; 30MG/100ML;
74MG/100ML; 640MG/100ML; 500MG/100ML;

74MG/100ML

N18269 002
JAN 17, 1983

@

35MG/100ML; 5GM/100ML; 30MG/100ML;

74MG/100ML; 640MG/100ML; 500MG/100ML;

74MG/100ML

N18269 002
JAN 17, 1983

CALFACTANT

INJECTABLE; INJECTION

INFASURF PRESERVATIVE FREE

+ ONY

35MG/ML

N20521 001
JUL 01, 1998

SUSPENSION; INTRATRACHEAL

INFASURF PRESERVATIVE FREE

+ ONY

35MG/ML

N20521 001
JUL 01, 1998

CAPECITABINE

TABLET; ORAL

XELODA

HLR

ROCHE

150MG

150MG

N20896 001
APR 30, 1998
N20896 001
APR 30, 1998

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL

BAKER NORTON

12.5MG

25MG

50MG

N74590 004
AUG 30, 1996
N74590 002
AUG 30, 1996
N74590 001
AUG 30, 1996

CAPTOPRIL

TABLET; ORAL

CAPTOPRIL
BAKER NORTON

100MG

N74590 003
AUG 30, 1996

AB ZENITH GOLDLINE

12.5MG

N74590 004
AUG 30, 1996

25MG

N74590 002
AUG 30, 1996

50MG

N74590 001
AUG 30, 1996

100MG

N74590 003
AUG 30, 1996

CARBENICILLIN DISODIUM

INJECTABLE; INJECTION

GEOPEN

+ ROERIG

EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 5GM BASE/VIAL
EQ 10GM BASE/VIAL
EQ 30GM BASE/VIAL
EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 5GM BASE/VIAL
EQ 10GM BASE/VIAL
EQ 30GM BASE/VIAL

N50306 001
N50306 004
N50306 002
N50306 006
N50306 007
N50306 001
N50306 004
N50306 002
N50306 006
N50306 007

CARBIDOPA; LEVODOPA

TABLET, EXTENDED RELEASE; ORAL

CARBIDOPA AND LEVODOPA

MYLAN

50MG; 200MG

N75091 001
SEP 30, 1999

SINEMET CR

AB + DUPONT PHARMS

50MG; 200MG

50MG; 200MG

N19856 001
MAY 30, 1991
N19856 001
MAY 30, 1991

CARTEOLOL HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

OCUPRESS

+ CIBA

1%

* OTSUKA

1%

N19972 001
MAY 23, 1990
N19972 001
MAY 23, 1990

CEFACTOR

CAPSULE; ORAL

CEFACTOR

LEDERLE

EQ 250MG BASE

N64107 001
APR 27, 1995

EQ 500MG BASE

N64107 002
APR 27, 1995

EQ 250MG BASE

N64107 001
APR 27, 1995

EQ 500MG BASE

N64081 001
SEP 16, 1996

EQ 500MG BASE

N64081 002
SEP 16, 1996

EQ 250MG BASE

N64081 001
SEP 16, 1996

EQ 500MG BASE

N64081 002
SEP 16, 1996

POWDER FOR RECONSTITUTION; ORAL

CEFACTOR

LEDERLE

EQ 125MG BASE/5ML

N64114 001
APR 28, 1995

EQ 187MG BASE/5ML

N64115 001
APR 28, 1995

EQ 250MG BASE/5ML

N64116 001
APR 28, 1995

EQ 375MG BASE/5ML

N64110 001
APR 28, 1995

EQ 125MG BASE/5ML

N64114 001
APR 28, 1995

EQ 187MG BASE/5ML

N64115 001
APR 28, 1995

EQ 250MG BASE/5ML

N64116 001
APR 28, 1995

EQ 375MG BASE/5ML

N64110 001
APR 28, 1995

CEFADROXIL/CEFADROXIL HEMIHYDRATE

CAPSULE; ORAL
CEFADROXIL
 BARR

AB EQ 500MG BASE

N65015 001
 JUN 22, 1999

POWDER FOR RECONSTITUTION; ORAL

CEFADROXIL
 APOTHECON

AB EQ 125MG BASE/5ML
 AB EQ 250MG BASE/5ML
 AB EQ 500MG BASE/5ML
 @ EQ 125MG BASE/5ML
 @ EQ 250MG BASE/5ML
 @ EQ 500MG BASE/5ML

N62334 001
 N62334 002
 N62334 003
 N62334 001
 N62334 002
 N62334 003

DURICEF

AB * EQ 125MG BASE/5ML
 AB * EQ 250MG BASE/5ML
 AB * EQ 500MG BASE/5ML
 + EQ 125MG BASE/5ML
 + EQ 250MG BASE/5ML
 + EQ 500MG BASE/5ML

N50527 002
 N50527 003
 N50527 001
 N50527 002
 N50527 003
 N50527 001

TABLET; ORAL

CEFADROXIL
 RANBAXY

AB EQ 1GM BASE

N65018 001
 APR 23, 1999

AB ULTRACEF
 APOTHECON

AB EQ 1GM BASE

N62390 001
 JUN 10, 1982
 N62390 001
 JUN 10, 1982

CEFTAZIDIME (ARGININE FORMULATION)

INJECTABLE; INJECTION

CEPTAZ

AB * GLAXO WELLCOME

AB EQ 1GM/VIAL

N50646 002
 SEP 27, 1990

AB *

AB EQ 2GM/VIAL

N50646 003
 SEP 27, 1990

AB *

AB EQ 10GM/VIAL

N50646 004
 SEP 27, 1990

+

AB EQ 1GM/VIAL

N50646 002
 SEP 27, 1990

+

AB EQ 2GM/VIAL

N50646 003
 SEP 27, 1990

CEFTAZIDIME (ARGININE FORMULATION)

INJECTABLE; INJECTION

CEPTAZ

+ GLAXO WELLCOME

AB EQ 10GM/VIAL

N50646 004
 SEP 27, 1990

PENTACEF

SMITHKLINE BEECHAM

AB EQ 1GM/VIAL

N63322 001
 NOV 07, 1995

AB

AB EQ 1GM/VIAL

N64006 001
 MAR 31, 1992

AB

AB EQ 2GM/VIAL

N63322 002
 NOV 07, 1995

AB

AB EQ 2GM/VIAL

N64006 002
 MAR 31, 1992

AB

AB EQ 10GM/VIAL

N64008 002
 MAR 31, 1992

AB

AB EQ 6GM/VIAL

N64008 001
 MAR 31, 1992

@

AB EQ 1GM/VIAL

N63322 001
 NOV 07, 1995

@

AB EQ 1GM/VIAL

N64006 001
 MAR 31, 1992

@

AB EQ 2GM/VIAL

N63322 002
 NOV 07, 1995

@

AB EQ 2GM/VIAL

N64006 002
 MAR 31, 1992

@

AB EQ 6GM/VIAL

N64008 001
 MAR 31, 1992

@

AB EQ 10GM/VIAL

N64008 002
 MAR 31, 1992

CEFUROXIME SODIUM

INJECTABLE; INJECTION

CEFUROXIME

AB ASTRA PHARMS

AB EQ 750MG BASE/VIAL

N64192 002
 APR 16, 1998

AB

AB EQ 1.5GM BASE/VIAL

N64192 001
 APR 16, 1998

AB

AB EQ 7.5GM BASE/VIAL

N64191 001
 APR 16, 1998

AB

AB EQ 750MG BASE/VIAL

N64192 002
 APR 16, 1998

AB

AB EQ 1.5GM BASE/VIAL

N64192 001
 APR 16, 1998

AB

AB EQ 7.5GM BASE/VIAL

N64191 001
 APR 16, 1998

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN '99 - NOV '99

CEPHALEXIN

CAPSULE; ORAL
CEPHALEXIN
RANBAXY

AB EQ 250MG BASE
SEP 16, 1999
AB EQ 500MG BASE
SEP 16, 1999

N65007 001
SEP 16, 1999
N65007 002
SEP 16, 1999

N85517 001
N85517 001
N86217 001
N86217 001

CEPHALOTHIN SODIUM

INJECTABLE; INJECTION
CEPHALOTHIN SODIUM
ABBOTT

AP EQ 1GM BASE/VIAL
SEP 11, 1985
AP EQ 2GM BASE/VIAL
SEP 11, 1985
EQ 1GM BASE/VIAL
SEP 11, 1985
EQ 2GM BASE/VIAL
SEP 11, 1985

N62547 001
SEP 11, 1985
N62547 002
SEP 11, 1985
N62547 001
SEP 11, 1985
N62547 002
SEP 11, 1985

AA EQ 1GM BASE/VIAL
SEP 11, 1985
AA EQ 2GM BASE/VIAL
SEP 11, 1985
EQ 1GM BASE/VIAL
SEP 11, 1985
EQ 2GM BASE/VIAL
SEP 11, 1985

N87228 001
N87228 001
N83082 002
SEP 17, 1999

CERIVASTATIN SODIUM

TABLET; ORAL
BAYCOL
* BAYER

0.3MG
JUN 26, 1997
0.3MG
JUN 26, 1997
0.4MG
MAY 24, 1999

N20740 004
JUN 26, 1997
N20740 004
JUN 26, 1997
N20740 005
MAY 24, 1999

N84652 001
N84652 001
N88102 004
N88102 004

CHLORAMPHENICOL

SOLUTION/DROPS; OPHTHALMIC
CHLORAMPHENICOL
ALCON

AT 0.5%
SEP 25, 1985
AT 0.5%
SEP 25, 1985

N62628 001
SEP 25, 1985
N62628 001
SEP 25, 1985

CHLOROTHALIDONE

TABLET; ORAL

CHLOROTHALIDONE
ABBOTT

AB 25MG
SEP 16, 1999
AB 25MG
SEP 16, 1999
AB 25MG
SEP 16, 1999
AB 25MG
SEP 16, 1999

N87364 001
N87364 001
N87296 001
N87296 001
N87706 001
N87706 001
N87521 001
N87521 001

CHLORDIAZEPOXIDE HYDROCHLORIDE

CAPSULE; ORAL
A-POXIDE
ABBOTT

AB 10MG
SEP 16, 1999
AB 10MG
SEP 16, 1999

N85517 001
N85517 001
N86217 001
N86217 001

CHLOROQUINE PHOSPHATE

TABLET; ORAL

CHLOROQUINE PHOSPHATE
MD PHARM

AA EQ 150MG BASE
SEP 17, 1999
AA EQ 150MG BASE
SEP 17, 1999
AA EQ 300MG BASE
SEP 17, 1999

N87228 001
N87228 001
N83082 002
SEP 17, 1999

CHLOROTRIANISENE

CAPSULE; ORAL

CHLOROTRIANISENE
BANNER PHARMACAPS

AA 12MG
SEP 16, 1999
AA 12MG
SEP 16, 1999
AA 12MG
SEP 16, 1999

N84652 001
N84652 001
N88102 004
N88102 004

CHLORPROMAZINE HYDROCHLORIDE

CONCENTRATE; ORAL

CHLORPROMAZINE HCL
PHARM ASSOC

AA 100MG/ML
SEP 16, 1999

N40224 001
JAN 26, 1999

CLINDAMYCIN HYDROCHLORIDE

CAPSULE; ORAL

CLEOCIN HCL

AB

AB

AB

AB

* PHARMACIA AND UPJOHN

EQ 75MG BASE

EQ 75MG BASE

EQ 150MG BASE

EQ 150MG BASE

EQ 300MG BASE

EQ 300MG BASE

APR 14, 1988

APR 14, 1988

CLINDAMYCIN PHOSPHATE

INJECTABLE; INJECTION

CLINDAMYCIN PHOSPHATE

SOLOPAK

AP

AP

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

EQ 150MG BASE/ML

SUPPOSITORY; VAGINAL

CLEOCIN

+ PHARMACIA AND UPJOHN 100MG

CLOBETASOL PROPIONATE

CREAM; TOPICAL

CLOBETASOL PROPIONATE

ALPHARMA US PHARM

0.05%

0.05%

AB

AB

N74139 001

AUG 03, 1994

N74139 001

AUG 03, 1994

CLOBETASOL PROPIONATE

CREAM; TOPICAL

CLOBETASOL PROPIONATE (EMOLLIENT)

ALTANA

AB2

> ADD >

> ADD >

> DLT >

> DLT >

N50162 001

N50162 001

N50162 002

N50162 002

N50162 003

APR 14, 1988

APR 14, 1988

GEL; TOPICAL

CLOBETASOL PROPIONATE

TARO

AB

0.05%

N75430 001

MAY 26, 1999

N75430 001

MAY 26, 1999

N75279 001

MAY 28, 1999

SOLUTION; TOPICAL

CLOBETASOL PROPIONATE

ALTANA

AT

0.05%

N75391 001

FEB 08, 1999

CLOMIPHENE CITRATE

TABLET; ORAL

CLOMIPHENE CITRATE

PAR PHARM

AB

50MG

N75528 001

AUG 30, 1999

CLOXACILLIN SODIUM

CAPSULE; ORAL

CLOXAPEN

SMITHKLINE BEECHAM

AB

AB

EQ 250MG BASE

EQ 500MG BASE

EQ 250MG BASE

EQ 500MG BASE

N62233 001

N62233 002

N62233 001

N62233 002

POWDER FOR RECONSTITUTION; ORAL

CLOXACILLIN SODIUM

TEVA

AA

EQ 125MG BASE/5ML

EQ 125MG BASE/5ML

N62268 001

N62268 001

+ TEGOPEN

* APOTHECON

AA

EQ 125MG BASE/5ML

EQ 125MG BASE/5ML

N61453 001

N61453 001

CLOZAPINE

TABLET; ORAL

AB CLOZAPINE
CREIGHTON

25MG

N74546 001
AUG 30, 1996

100MG

N74546 002
AUG 30, 1996

AB GENEVA PHARMS

25MG

N74546 001
AUG 30, 1996

100MG

N74546 002
AUG 30, 1996

AB MYLAN

25MG

N75417 001
MAY 27, 1999

100MG

N75417 002
MAY 27, 1999

COLISTIMETHATE SODIUM

INJECTABLE; INJECTION

AP COLISTIMETHATE
PHARMA TEK

EQ 150MG BASE/VIAL

N64216 001
FEB 26, 1999

AP COLY-MYCIN M

AP + PARKEDALE
*

EQ 150MG BASE/VIAL
EQ 150MG BASE/VIAL

N50108 002
N50108 002

CORTICOTROPIN

INJECTABLE; INJECTION

AP CORTICOTROPIN
STERIS

40 UNITS/VIAL

N88772 001
NOV 21, 1984

40 UNITS/VIAL

N88772 001
NOV 21, 1984

CROMOLYN SODIUM

CAPSULE; ORAL

AB GASTROCROM
MEDEVA
*

100MG

N19188 001
DEC 22, 1989

100MG

N19188 001
DEC 22, 1989

CROMOLYN SODIUM

SOLUTION; INHALATION

AN CROMOLYN SODIUM
ALPHARMA

10MG/ML

N75067 001
JUL 19, 1999

AN MORTON GROVE

10MG/ML

N75346 001
OCT 25, 1999

AN ROXANE

10MG/ML

N75175 001
SEP 30, 1999

SOLUTION/DROPS; OPHTHALMIC

AT CROMOLYN SODIUM
ALCON

4%

N75282 001
JUN 16, 1999

AT CROMOPTIC
KING PHARMS

4%

N75088 001
APR 27, 1999

CROTAMITON

LOTION; TOPICAL

AT CROTAN
CAIDSENA

10%

N87204 001

10%

N87204 001

CYANOCOBALAMIN

INJECTABLE; INJECTION

AP RUVITE
SAVAGE LABS
*

1MG/ML

N80570 002

1MG/ML

N80570 002

CYCLOPHOSPHAMIDE

TABLET; ORAL

AB CYCLOPHOSPHAMIDE
ROXANE

25MG

N40032 001

50MG

N40032 002

AB CYTOSAN

25MG

N12141 002

50MG

N12141 001

50MG

N12141 002

50MG

N12141 001

50MG

N12141 001

CYCLOSPORINE

INJECTABLE; INJECTION
CYCLOSPORINE

50MG/ML

N65004 001
OCT 29, 1999

AP DACARBAZINE
AM PHARM PARTNERS

100MG/VIAL

N75371 001
AUG 27, 1999

AP SANDIMMUNE
+ NOVARTIS

50MG/ML

N50573 001
NOV 14, 1983

AP DTIC-DOME
+ BAYER

200MG/VIAL

N75371 002
AUG 27, 1999

50MG/ML

N50573 001
NOV 14, 1983

AP DTIC-DOME
+ BAYER

100MG/VIAL
100MG/VIAL

N17575 001
N17575 001

CYPROHEPTADINE HYDROCHLORIDE

TABLET; ORAL

CYPROHEPTADINE HCL

4MG

N87566 001
NOV 10, 1982

INJECTABLE; INJECTION
SYNERCID

350MG/VIAL;
EQ 150MG BASE/VIAL

N87566 001
NOV 10, 1982

350MG/VIAL;
EQ 150MG BASE/VIAL

N50747 001
SEP 21, 1999

CYSTEINE HYDROCHLORIDE

INJECTABLE; INJECTION
CYSTEINE HCL

7.25%

N19523 001
OCT 22, 1986

INJECTABLE; INJECTION
CERUBIDINE

EQ 20MG BASE/VIAL

N64103 001
FEB 03, 1995

PHARMACIA AND UPJOHN

EQ 20MG BASE/VIAL

N64103 001
FEB 03, 1995

CYTARABINE

INJECTABLE; INJECTION

CYTARABINE
FAULDING

2GM/VIAL

N75383 001
NOV 22, 1999

INJECTABLE; INJECTION
CERUBIDINE

EQ 20MG BASE/VIAL

N50731 001
JAN 30, 1998

INJECTABLE, LIPOSOMAL; INJECTION
DEPOCYT

10MG/ML

N21041 001
APR 01, 1999

INJECTABLE; INJECTION
CERUBIDINE

EQ 20MG BASE/VIAL

N50731 001
JAN 30, 1998

+ SKYEPHARMA

EQ 20MG BASE/VIAL

N50731 001
JAN 30, 1998

DESMOPRESSIN ACETATE

SPRAY, METERED; NASAL

AB DDAVE
AB + RHONE POULENC RORER 0.01MG/SPRAY

AB *
AB 0.01MG/SPRAY

AB *
AB 0.01MG/SPRAY

AB + RHONE POULENC RORER 0.01MG/SPRAY

AB + RHONE POULENC RORER 0.01MG/SPRAY

AB DESMOPRESSIN ACETATE

AB BAUSCH AND LOMB 0.01MG/SPRAY

DESOGESTREL; ETHINYL ESTRADIOL

TABLET; ORAL-21

AB DESOGESTREL AND ETHINYL ESTRADIOL
 DURAMED 0.15MG;0.03MG

AB ORTHO-CEPT
AB + JOHNSON RW 0.15MG;0.03MG

AB *
AB 0.15MG;0.03MG

TABLET; ORAL-28

AB DESOGESTREL AND ETHINYL ESTRADIOL
 DURAMED 0.15MG;0.03MG

DESONIDE

CREAM; TOPICAL

AB TRIDESILON
AB * BAYER 0.05%
AB + CLAY PARK 0.05%

OINTMENT; TOPICAL

AB TRIDESILON
AB * BAYER 0.05%
AB + CLAY PARK 0.05%

DEXAMETHASONE

SUSPENSION/DROPS; OPHTHALMIC

AT DEXAMETHASONE
AT STERIS 0.1%

AT *
AT 0.1%

AT MAXIDEX
AT * ALCON 0.1%

AT +
AT 0.1%

TABLET; ORAL

BP DEXONE 0.5
BP SOLVAY 0.5MG

BP *
BP 0.5MG

BP DEXONE 0.75
BP SOLVAY 0.75MG

BP *
BP 0.75MG

BP DEXONE 4
BP SOLVAY 4MG

DEXAMETHASONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

OINTMENT; OPHTHALMIC

AT MAXITROL
AT * ALCON 0.1%;EQ 3.5MG BASE/GM;
 10,000 UNITS/GM

AT + FALCON PHARMS 0.1%;EQ 3.5MG BASE/GM;
 10,000 UNITS/GM

SUSPENSION/DROPS; OPHTHALMIC

AT MAXITROL
AT * ALCON 0.1%;EQ 3.5MG BASE/ML;
 10,000 UNITS/ML

AT + FALCON PHARMS 0.1%;EQ 3.5MG BASE/ML;
 10,000 UNITS/ML

DEXAMETHASONE; TOBRAMYCIN

SUSPENSION; OPHTHALMIC

AB TOBRADEX
AB * ALCON 0.1%;0.3%

N89170 001
 MAY 09, 1989
 N89170 001
 MAY 09, 1989

N13422 001
 N13422 001

N84991 001
 N84991 001

N84993 001
 N84993 001

N84992 001
 N84992 001

N50065 002
 N50065 002

N50023 002
 N50023 002

N50592 001
 AUG 18, 1988

DEXAMETHASONE; TOBRAMYCINSUSPENSION/DROPS; OPHTHALMIC

AB + TOBREADX ALCON 0.1%; 0.3% N50592 001
AUG 18, 1988

AB TOBRAMYCIN AND DEXAMETHASONE N64134 001
OCT 27, 1999

DEXAMETHASONE SODIUM PHOSPHATEINJECTABLE; INJECTION

AP * DECADRON MERCK EQ 24MG PHOSPHATE/ML N12071 004
+ EQ 24MG PHOSPHATE/ML N12071 004

AP DEXAMETHASONE SODIUM PHOSPHATE EQ 4MG PHOSPHATE/ML N83702 001
AP STERIS EQ 24MG PHOSPHATE/ML N85606 001
AP EQ 4MG PHOSPHATE/ML N83702 001
EQ 24MG PHOSPHATE/ML N85606 001

DEXTROAMPHETAMINE SULFATETABLET; ORAL

AA DEXTROAMPHETAMINE SULFATE ENDO PHARMS 5MG N40299 001
MAY 13, 1999

DEXTROSEINJECTABLE; INJECTION

DEXTROSE 25% 250MG/ML N19445 002
ABBOTT NOV 23, 1998

DEXTROSE; SODIUM CHLORIDEINJECTABLE; INJECTION

DEXTROSE 2.5% AND SODIUM CHLORIDE 0.45% IN PLASTIC
CONTAINER N18030 001
AP B BRAUN 2.5GM/100ML; 450MG/100ML N18030 001
+ DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER N18030 001
B BRAUN 5GM/100ML; 110MG/100ML N18030 005

DEXTROSE; SODIUM CHLORIDEINJECTABLE; INJECTION

DEXTROSE 5% AND SODIUM CHLORIDE 0.11% IN PLASTIC CONTAINER N18030 005
AP B BRAUN 5GM/100ML; 110MG/100ML N18030 005
+ DEXTROSE 5% AND SODIUM CHLORIDE 0.2% IN PLASTIC CONTAINER N18030 004
B BRAUN 5GM/100ML; 200MG/100ML N18030 004
+ DEXTROSE 5% AND SODIUM CHLORIDE 0.33% IN PLASTIC CONTAINER N18030 003
B BRAUN 5GM/100ML; 330MG/100ML N18030 003
+ DEXTROSE 5% AND SODIUM CHLORIDE 0.45% IN PLASTIC CONTAINER N18030 002
AP B BRAUN 5GM/100ML; 450MG/100ML N18030 002
+ 5GM/100ML; 450MG/100ML N18030 002

DIATRIZOATE MEGLUMINE; DIATRIZOATE SODIUMINJECTABLE; INJECTION

AP MD-76 55%; 10% N87073 001
+ MALLINCKRODT 66%; 10% N87073 001

DIAZEPAMGEL; RECTAL

DIATAT 2.5MG/0.5ML N20648 001
* ATHENA 5MG/ML N20648 002
10MG/2ML N20648 003
15MG/3ML N20648 004
20MG/4ML N20648 005
+ ELAN PHARMS 2.5MG/0.5ML N20648 001
5MG/ML N20648 002
10MG/2ML N20648 003
15MG/3ML N20648 004
20MG/4ML N20648 005
+ JUL 29, 1997

DIAZEPAMINJECTABLE; INJECTION

AP DIAZEPAM
STERIS

5MG/ML

N70911 001

AUG 28, 1986

5MG/ML

N70930 001

DEC 01, 1986

5MG/ML

N70911 001

AUG 28, 1986

5MG/ML

N70930 001

DEC 01, 1986

DICLOFENAC POTASSIUMTABLET; ORAL

AB DICLOFENAC POTASSIUM
MYLAN

50MG

N75463 001

JUL 26, 1999

DICLOFENAC SODIUMSOLUTION/DROPS; OPHTHALMIC

AB DICLOFENAC SODIUM
ALCON

0.1%

AB FALCON PHARMS

0.1%

N20809 001

MAY 04, 1998

N20809 001

MAY 04, 1998

TABLET, DELAYED RELEASE; ORAL

AB DICLOFENAC SODIUM
MARTEC

50MG

N74986 001

FEB 26, 1999

N74986 002

FEB 26, 1999

N74723 001

MAR 30, 1999

N74432 002

JUL 29, 1999

N74432 003

JUL 29, 1999

DICYCLOMINE HYDROCHLORIDECAPSULE; ORAL

AB DICYCLOMINE HCL
MYLAN

10MG

N40319 001
SEP 07, 1999

TABLET; ORAL

AB DICYCLOMINE HCL
LANNETT

20MG

N40230 001
FEB 26, 1999
N40317 001
SEP 07, 1999

AB MYLAN

20MG

DIDANOSINE

TABLET, CHEWABLE; ORAL
VIDEX

25MG

N20154 002
OCT 09, 1991

N20154 005

OCT 09, 1991

N20154 002

OCT 09, 1991

N20154 005

OCT 09, 1991

N20154 006

OCT 28, 1999

DIFLORASONE DIACETATECREAM; TOPICAL

AB DIFLORASONE DIACETATE
TARO

0.05%

N75331 001
MAY 14, 1999

ointment; TOPICAL

AB DIFLORASONE DIACETATE
ALTANA

0.05%

N75374 001
APR 27, 1999

N75331 001

MAY 14, 1999

PSORCON

AB PHARMACIA AND UPJOHN

0.05%

N19260 001
AUG 28, 1985
N19280 001
AUG 28, 1985

[†] SEE SECTION 1.3 OF INTRODUCTION

DIFLUNISAL

TABLET; ORAL

DIFLUNISAL
PUREPAC PHARM

AB 250MG

AB 500MG

AB 250MG

AB 500MG

N74285 001

MAY 07, 1996

N74285 002

MAY 07, 1996

N74285 001

MAY 07, 1996

N74285 002

MAY 07, 1996

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CARDIZEM CD

AB3 + CARDERM

AB3 + 120MG

AB3 + 180MG

AB3 + 240MG

AB3 + 300MG

BC * 120MG

BC * 180MG

BC * 240MG

BC * 300MG

BC * 120MG

BC * 180MG

BC * 240MG

BC * 300MG

CARTIA XT
ANDRX PHARMS

AB3 120MG

AB3 180MG

AB3 240MG

AB3 300MG

AB3 120MG

AB3 180MG

AB3 240MG

DILTIAZEM HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

CARTIA XT

ANDRX PHARMS

AB1 300MG

AB1 60MG

AB1 90MG

AB1 120MG

N74752 004

JUL 09, 1998

N74845 001

SEP 15, 1999

N74845 002

SEP 15, 1999

N74845 003

SEP 15, 1999

INJECTABLE; INJECTION

DILTIAZEM HCL

APOTEX

AP 5MG/ML

AP 5MG/ML

N75375 001

SEP 30, 1999

N75106 001

APR 29, 1999

DIPHENHYDRAMINE HYDROCHLORIDE

CAPSULE; ORAL

DIPHENHYDRAMINE HCL

PUREPAC PHARM

AA 25MG

AA 50MG

AA 25MG

AA 50MG

N85156 001

N85150 001

N85156 001

N85150 001

ELIXIR; ORAL

HYDRAMINE

ALPHA

AA 12.5MG/5ML

AA 12.5MG/5ML

N80763 002

N80763 002

DIPIVEFRIN HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

DIPIVEFRIN HCL

ALCON

AT 0.1%

AT 0.1%

N73636 001

JUN 30, 1994

N73636 001

JUN 30, 1994

DIPYRIDAMOLE

TABLET; ORAL

AB DIPYRIDAMOLE
AB PUREPAC PHARM

25MG

50MG

25MG

50MG

N89425 001

JUL 12, 1990

N89426 001

JUL 12, 1990

N89425 001

JUL 12, 1990

N89426 001

JUL 12, 1990

> ADD >

> ADD >

> DLT >

> DLT >

EQ 1MG BASE/ML

EQ 1MG BASE/ML

N19946 001

MAR 07, 1991

N19946 001

MAR 07, 1991

DIRITHROMYCIN

TABLET, DELAYED RELEASE; ORAL

AB DYNABAC

AB LILLY

250MG

250MG

+ LILLY RES LABS

N20862 001

JUN 09, 1999

2.5 UGM

DOFETILIDE

CAPSULE; ORAL

AB TIKOSYN

AB PFIZER

0.125MG

0.25MG

0.5MG

N50718 001

NOV 17, 1995

N50718 001

NOV 17, 1995

2MG/ML

2MG/ML

+ ALZA

+ SEQUUS

DOXORUBICIN HYDROCHLORIDE

INJECTABLE, LIPOSOMAL; INJECTION

AB DOXIL

AB ALZA

N50678 001

JUN 19, 1995

N50678 001

JUN 19, 1995

DOXYCYCLINE HYCLATE

CAPSULE; ORAL

AB DOXYCYCLINE HYCLATE

AB PUREPAC PHARM

EQ 50MG BASE

EQ 100MG BASE

+ RANBAXY

EQ 50MG BASE

EQ 100MG BASE

N62479 001

DEC 23, 1983

N62479 002

DEC 23, 1983

N62479 001

DEC 23, 1983

N62479 002

DEC 23, 1983

DOPAMINE HYDROCHLORIDE

INJECTABLE; INJECTION

AB DOPAMINE HCL

AB SMITH AND NEPHEW

40MG/ML

80MG/ML

40MG/ML

80MG/ML

N70046 001

AUG 29, 1985

N70047 001

AUG 29, 1985

N70046 001

AUG 29, 1985

N70047 001

AUG 29, 1985

DOXYLAMINE SUCCINATE; PYRIDOXINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

AB BENDECTIN

AB HOECHST MARION RSSL

10MG;10MG

N10598 002

DROPERIDOL

INJECTABLE; INJECTION

DROPERIDOL
SOLOPAK

2.5MG/ML

N71754 001

SEP 06, 1988

2.5MG/ML

N71755 001

SEP 06, 1988

2.5MG/ML

N71754 001

SEP 06, 1988

2.5MG/ML

N73520 001

SEP 06, 1988

2.5MG/ML

N73521 001

NOV 27, 1991

2.5MG/ML

N73523 001

NOV 27, 1991

2.5MG/ML

N73520 001

NOV 27, 1991

2.5MG/ML

N73521 001

NOV 27, 1991

2.5MG/ML

N73523 001

NOV 27, 1991

ENALAPRIL MALEATE; FELODIPINE

TABLET, EXTENDED RELEASE; ORAL

LEXCEL

+ ASTRAZENECA 5MG; 2.5MG

ENTACAPONE

TABLET; ORAL

COMTAN

+ ORION

200MG

EPINEPHRINE

INJECTABLE; INJECTION

SUS-PHRINE

* FOREST LABS

SUS-PHRINE SULFITE-FREE

FOREST LABS 1.5MG/AMP

5MG/ML

N67942 001

FEB 05, 1999

N07942 003

FEB 05, 1999

EPINEPHRINE

INJECTABLE; INJECTION

SUS-PHRINE SULFITE-FREE

+ FOREST LABS 5MG/ML

N07942 001

EPIRUBICIN HYDROCHLORIDE

INJECTABLE; INJECTION

ELLENC

+ PHARMACIA AND UPJOHN 2MG/ML

N50778 001

SEP 15, 1999

EPROSARTAN MESYLATE

TABLET; ORAL

TEVETEN

SMITHKLINE BEECHAM

EQ 300MG BASE

N20738 004

DEC 22, 1997

EQ 400MG BASE

N20738 005

DEC 22, 1997

EQ 600MG BASE

N20738 006

MAY 27, 1999

UNIMED PHARMS

N20738 004

DEC 22, 1997

EQ 400MG BASE

N20738 005

DEC 22, 1997

EQ 600MG BASE

N20738 006

MAY 27, 1999

ERGOTAMINE TARTRATE

AEROSOL, METERED; INHALATION

MEDIHALER ERGOTAMINE

* 3M

@

0.36MG/INH

0.36MG/INH

N12102 001

N12102 001

ERYTHROMYCIN

CAPSULE, DELAYED REL PELLETS; ORAL

ERYC

@ FARKE DAVIS

@ WARNER CHILCOTT

250MG

250MG

N62338 001

N62338 001

ERYTHROMYCIN

SOLUTION; TOPICAL

C-SOLVE-2

AT BIOGLAN PHAR 2%

AT ZENITH GOLDLINE 2%

N62468 001
JUL 03, 1985
N62468 001
JUL 03, 1985

ESTRADIOL

FILM, EXTENDED RELEASE; TRANSDERMAL

ALORA

PROCTER AND GAMBLE

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

WATSON LABS

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

CLIMARA

+ BERLEX LABS

0.025MG/24HR

ESCLIM

FOURNIER

0.025MG/24HR

0.0375MG/24HR

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

WOMEN FIRST HLTHCARE

0.025MG/24HR

0.0375MG/24HR

0.05MG/24HR

0.075MG/24HR

ESTRADIOL

FILM, EXTENDED RELEASE; TRANSDERMAL

ESCLIM

WOMEN FIRST HLTHCARE

0.1MG/24HR

ESTRADIOL

CYGNUS CA

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

0.0375MG/24HR

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

MENOREST

0.0375MG/24HR

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

MYETH AYERST

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

VIVELLE-DOT

NOVARTIS

0.0375MG/24HR

0.05MG/24HR

0.075MG/24HR

0.1MG/24HR

TABLET; ORAL

ESTRADIOL

MYLAN

0.5MG

1MG

2MG

INNOFEM

NOVO NORDISK

0.5MG

N20847 005
AUG 04, 1998

N21048 001
SEP 20, 1999

N21048 002
SEP 20, 1999

N21048 003
SEP 20, 1999

N20538 001
JUL 31, 1996

N20538 003
JUL 31, 1996

N20538 002
JUL 31, 1996

N20538 004
JUL 31, 1996

N21048 001
SEP 20, 1999

N21048 002
SEP 20, 1999

N21048 003
SEP 20, 1999

N20538 001
JUL 31, 1996

N20538 003
JUL 31, 1996

N20538 002
JUL 31, 1996

N20538 004
JUL 31, 1996

N21048 001
SEP 20, 1999

N21048 002
SEP 20, 1999

N21048 003
SEP 20, 1999

N20538 001
JUL 31, 1996

N20538 003
JUL 31, 1996

N20538 002
JUL 31, 1996

N20538 004
JUL 31, 1996

N40326 001
APR 21, 1999

N40326 002
APR 21, 1999

N40326 003
APR 21, 1999

N40312 001
NOV 19, 1999

ESTRADIOL

TABLET; ORAL
INNOFEM
NOVO NORDISK

> ADD >
> ADD >
> ADD >
> ADD >
> ADD >

1MG
2MG

N40312 002
NOV 19, 1999
N40312 003
NOV 19, 1999

N84710 001

TABLET; VAGINAL
VAGIFEM
+ NOVO NORDISK

25 UGM

N20908 001
MAR 26, 1999

ESTRADIOL; NORGESTIMATE

TABLET; ORAL
ORTHO-PREFEST
+ JOHNSON RW

1MG, 1MG; 0.09MG, N/A

N21040 001
OCT 22, 1999

ESTROGENS, CONJUGATED SYNTHETIC A

TABLET; ORAL
CENESTIN
DURAMED

0.9MG
0.625MG
0.9MG

N20992 003
MAR 24, 1999
N20992 002
MAR 24, 1999
N20992 003
MAR 24, 1999

N83220 004

ESTRONE

INJECTABLE; INJECTION
NATURAL ESTROGENIC SUBSTANCE-ESTRONE
+ STERIS

2MG/ML
2MG/ML

N85237 001
NOV 23, 1982
N85237 001
NOV 23, 1982

N75095 001

NOV 30, 1999
N75095 002
NOV 30, 1999

ESTROPIPATE

CREAM; VAGINAL
OGEN
+ ABBOTT

1.5MG/GM

N84710 001

ESTROPIPATE

CREAM; VAGINAL
OGEN
+ PHARMACIA AND UPJOHN

1.5MG/GM

TABLET; ORAL
ESTROPIPATE
DURAMED

> ADD >
> ADD >
> ADD >
> ADD >
> ADD >
> ADD >

0.75MG

N40296 001

NOV 01, 1999

N40296 002

NOV 01, 1999

N40296 003

NOV 01, 1999

N40359 001

AUG 26, 1999

N40359 002

AUG 26, 1999

N40359 003

AUG 26, 1999

MYLAN

0.75MG

N40359 001

AUG 26, 1999

N40359 002

AUG 26, 1999

N40359 003

AUG 26, 1999

OGEN .625

ABBOTT

PHARMACIA AND UPJOHN

0.75MG

N83220 001

N83220 001

OGEN 1.25

ABBOTT

PHARMACIA AND UPJOHN

1.5MG

N83220 002

N83220 002

OGEN 2.5

ABBOTT

PHARMACIA AND UPJOHN

3MG

N83220 003

N83220 003

OGEN 5

ABBOTT

PHARMACIA AND UPJOHN

5MG

N83220 004

N83220 004

ETHAMBUTOL HYDROCHLORIDE

TABLET; ORAL
ETHAMBUTOL HCL
WEST WARD

> ADD >
> ADD >
> ADD >
> ADD >
> ADD >

100MG

N75095 001

NOV 30, 1999

N75095 002

NOV 30, 1999

MYAMBUTOL

LEDERLE

> ADD >
> ADD >
> DLT >
> DLT >

100MG

N16320 001

N16320 003

N16320 001

N16320 003

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN'99 - NOV'99

ETHCHLORVYNOL

CAPSULE; ORAL
ETHCHLORVYNOL
BANNER PHARMACAPS

AA 200MG
AA 500MG
AA 750MG
AA 100MG
@ 100MG
@ 200MG
@ 500MG
@ 750MG

PLACIDYL
+ ABBOTT

AA 200MG
AA 500MG
AA 750MG
+ 200MG
+ 500MG
+ 750MG

N84463 002
N84463 003
N84463 004
N84463 001
N84463 001
N84463 002
N84463 003
N84463 004
N10021 007
N10021 002
N10021 010
N10021 007
N10021 002
N10021 010

TABLET; ORAL-28

BREVICON 28-DAY
WATSON LABS

AB 0.035MG; 0.5MG

NORINYL 1+35 28-DAY

AB 0.035MG; 1MG

SEARLE 0.035MG; 1MG

WATSON LABS

TRI-NORINYL 28-DAY

AB 0.035MG; 0.035MG; 0.5MG; 1MG

SEARLE 0.035MG; 0.035MG; 0.5MG; 1MG

WATSON LABS

0.035MG; 0.035MG; 0.5MG; 1MG

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

FEMHRT

+ PARKE DAVIS

0.005MG; 1MG

N21065 002

OCT 15, 1999

ETHINYL ESTRADIOL; LEVONORGESTREL

TABLET; ORAL-21

LEVITE

BERLEX LABS

+ 0.02MG; 0.1MG

0.02MG; 0.1MG

JUL 13, 1998

JUL 13, 1998

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-21

BREVICON 21-DAY

SEARLE

WATSON LABS

NORINYL 1+35 21-DAY

AB 0.035MG; 0.5MG

AB 0.035MG; 0.5MG

SEARLE 0.035MG; 1MG

WATSON LABS

TRI-NORINYL 21-DAY

AB 0.035MG; 0.035MG; 0.5MG; 1MG

SEARLE 0.035MG; 0.035MG; 0.5MG; 1MG

WATSON LABS

0.035MG; 0.035MG; 0.5MG; 1MG

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

BREVICON 28-DAY

SEARLE

AB 0.035MG; 0.5MG

N17743 001

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

BREVICON 28-DAY

WATSON LABS

AB 0.035MG; 0.5MG

NORINYL 1+35 28-DAY

AB 0.035MG; 1MG

SEARLE 0.035MG; 1MG

WATSON LABS

TRI-NORINYL 28-DAY

AB 0.035MG; 0.035MG; 0.5MG; 1MG

SEARLE 0.035MG; 0.035MG; 0.5MG; 1MG

WATSON LABS

0.035MG; 0.035MG; 0.5MG; 1MG

ETHINYL ESTRADIOL; NORETHINDRONE ACETATE

TABLET; ORAL

FEMHRT

+ PARKE DAVIS

0.005MG; 1MG

N21065 002

OCT 15, 1999

ETHINYL ESTRADIOL; NORGESTREL

TABLET; ORAL-21

LO/OVRAL

AB 0.03MG; 0.3MG

+ WYETH AYERST

0.03MG; 0.3MG

JUL 28, 1999

JUL 28, 1999

LOW-OGESTREL-21

AB 0.03MG; 0.3MG

SCS

WATSON LABS

0.03MG; 0.3MG

TABLET; ORAL-28

LO/OVRAL-28

AB 0.03MG; 0.3MG

WYETH AYERST

0.03MG; 0.3MG

JUL 28, 1999

JUL 28, 1999

LOW-OGESTREL-28

AB 0.03MG; 0.3MG

SCS

WATSON LABS

0.03MG; 0.3MG

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

BREVICON 28-DAY

SEARLE

AB 0.035MG; 0.5MG

N17743 001

JUL 28, 1999

JUL 28, 1999

LOW-OGESTREL-28

AB 0.03MG; 0.3MG

SCS

WATSON LABS

0.03MG; 0.3MG

ETHINYL ESTRADIOL; NORETHINDRONE

TABLET; ORAL-28

BREVICON 28-DAY

SEARLE

AB 0.035MG; 0.5MG

N17743 001

JUL 28, 1999

JUL 28, 1999

LOW-OGESTREL-28

AB 0.03MG; 0.3MG

SCS

WATSON LABS

0.03MG; 0.3MG

ETODOLAC

CAPSULE; ORAL
ETODOLAC
NOVOPHARM NC

AB 200MG
AB 300MG

N75126 001
SEP 16, 1999
N75126 002
SEP 16, 1999

25MG

N19369 002
SEP 30, 1986

TABLET; ORAL
ETODOLAC
EON

AB 500MG
AB 400MG
AB 500MG

N74903 002
APR 19, 1999
N74847 001
APR 23, 1999
N74847 002
APR 23, 1999

TABLET; ORAL
AROMASIN
+ PHARMACIA AND UPJOHN 25MG

N20753 001
OCT 21, 1999

ETOPOSIDE

INJECTABLE; INJECTION
ETOPOSIDE
AM PHARM PARTNERS

AP 20MG/ML
AP 20MG/ML
AP 20MG/ML
20MG/ML

N74983 001
SEP 30, 1998
N74983 001
SEP 30, 1998
N74228 001
OCT 15, 1996
N74228 001
OCT 15, 1996

67MG
67MG
134MG
200MG

N19304 002
FEB 09, 1998
N19304 002
FEB 09, 1998
N19304 003
JUN 30, 1999
N19304 004
JUN 30, 1999

VEPESID
AP + BRISTOL

AP 20MG/ML
AP 20MG/ML

N18768 001
NOV 10, 1983
N18768 001
NOV 10, 1983

FERRIC SODIUM GLUCONATE
INJECTABLE; INJECTION
FERRLECIT
+ R AND D LABS

62.5MG/5ML

N20955 001
FEB 18, 1999

ETRETINATE

CAPSULE; ORAL
TEGISON
ROCHE

10MG
25MG
10MG

N19369 001
SEP 30, 1986
N19369 002
SEP 30, 1986
N19369 001
SEP 30, 1986

FLUOCINONIDE

AB
FLUOCINONIDE
TARO

0.05%

N75008 001
JUN 30, 1999

FLUOROURACIL

INJECTABLE; INJECTION

FLUOROURACIL

AP BIGNAR

50MG/ML

N40291 001

MAR 24, 1999

AP SMITH AND NEPHEW

50MG/ML

N88767 001

DEC 28, 1984

AP

50MG/ML

N89434 001

MAR 26, 1987

AP

50MG/ML

N88767 001

DEC 28, 1984

AP

50MG/ML

N89434 001

MAR 26, 1987

FLUOXETINE HYDROCHLORIDE

CAPSULE; ORAL

PROZAC

+ LILLY

EQ 20MG BASE

N18936 001

DEC 29, 1987

EQ 20MG BASE

N18936 001

DEC 29, 1987

EQ 40MG BASE

N18936 003

JUN 15, 1999

EQ 60MG BASE

N18936 004

JUN 15, 1999

TABLET; ORAL

PROZAC

LILLY

EQ 10MG BASE

N20974 001

MAR 09, 1999

EQ 20MG BASE

N20974 002

MAR 09, 1999

EQ 10MG BASE

N20974 001

MAR 09, 1999

EQ 20MG BASE

N20974 002

MAR 09, 1999

FOLIC ACID

TABLET; ORAL

FOLIC ACID

VINTAGE PHARMS

1MG

N86296 001

N86296 001

FUROSEMIDE

INJECTABLE; INJECTION

FUROSEMIDE

ABBOTT

10MG/ML

N75241 001

MAY 28, 1999

AP SMITH AND NEPHEW

10MG/ML

N70078 001

FEB 05, 1986

AP

10MG/ML

N70078 001

FEB 05, 1986

AP

10MG/ML

N70019 001

SEP 22, 1986

AP

10MG/ML

N70604 001

JAN 02, 1987

AP

10MG/ML

N70019 001

SEP 22, 1986

AP

10MG/ML

N70604 001

JAN 02, 1987

GALLAMINE TRIETHIOIDE

INJECTABLE; INJECTION

FLAXEDIL

+ DAVIS AND GECK

+

+

+

20MG/ML

100MG/ML

20MG/ML

100MG/ML

N07842 001

N07842 002

N07842 001

N07842 002

GALLIUM NITRATE

INJECTABLE; INJECTION

GANITE

+ SOLOPAK

+

+

25MG/ML

25MG/ML

N19961 002

JAN 17, 1991

N19961 002

JAN 17, 1991

GANIRELIX ACETATE

INJECTABLE; INJECTION

ANTAGON

+ ORGANON INC

EQ 250 UGM BASE/0.5ML

N21057 001

JUL 29, 1999

GEMFIBROZIL

TABLET; ORAL

GEMFIBROZIL

AB PUREPAC PHARM

600MG

N74360 001

AUG 31, 1994

N74360 001

AUG 31, 1994

600MG

@

GENTAMICIN SULFATE

INJECTABLE; INJECTION

AFOGEN

AB KING PHARMS

AP

AP

AP

EQ 10MG BASE/ML

EQ 40MG BASE/ML

EQ 10MG BASE/ML

EQ 40MG BASE/ML

N62289 001

N62289 002

N62289 001

N62289 002

BRISTAGEN

AB BRISTOL

@

GENTAMICIN SULFATE

AB SOLOPAK

AP

AP

AP

EQ 10MG BASE/ML

EQ 40MG BASE/ML

EQ 10MG BASE/ML

EQ 40MG BASE/ML

N62507 001

N62507 002

N62507 001

N62507 002

JUN 06, 1985

N62507 002

JUN 06, 1985

SOLUTION/DROPS; OPHTHALMIC

GENTAMICIN SULFATE

AT ALCON

AT

AT

EQ 0.3% BASE

EQ 0.3% BASE

N62196 001

N62196 001

GLIPIZIDE

TABLET; ORAL

GLUCOTROL

PFIZER

@

TABLET, EXTENDED RELEASE; ORAL

GLUCOTROL XL

+ PFIZER

2.5MG

N20329 003

AUG 10, 1999

GLYBURIDE

TABLET; ORAL

GLYBURIDE (MICRONIZED)

AB MOVA

4.5MG

N74591 003

DEC 22, 1997

N74591 003

DEC 22, 1997

N74792 003

AUG 17, 1999

N74686 001

APR 20, 1999

N74686 002

APR 20, 1999

N74686 003

APR 20, 1999

N74686 004

APR 20, 1999

AB MYLAN

AB NOVOPHARM

AB

AB

AB

AB

AB

AB

AB

AB

AB

AB

GLYCOPYRROLATE

INJECTABLE; INJECTION

GLYCOPYRROLATE

AP STERIS

AP

AP

AP

9.2MG/ML

9.2MG/ML

9.2MG/ML

9.2MG/ML

N86947 001

JUN 24, 1983

N86947 001

JUN 24, 1983

TABLET; ORAL

ROBINUL

AB HORIZON PHARM

AB ROBIN PHARM

AB ROBINUL FORTE

AB HORIZON PHARM

AB ROBIN PHARM

AB

1MG

1MG

2MG

2MG

N12827 001

N12827 001

N12827 002

N12827 002

GRAMICIDIN; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OPHTHALMIC

NEOMYCIN AND POLYMYXIN B SULFATES AND GRAMICIDIN

AT STERIS

AT

AT

AT

AT

AT

AT

AT

0.025MG/ML; EQ 1.75MG BASE/ML;

10,000 UNITS/ML

N62788 001

JUN 11, 1987

0.025MG/ML; EQ 1.75MG BASE/ML;

10,000 UNITS/ML

N62788 001

JUN 11, 1987

HALOPERIDOLTABLET, ORALHALOPERIDOL
SCS

<u>AB</u>	<u>0.5MG</u>	N70720 001
		JUN 10, 1986
<u>AB</u>	<u>1MG</u>	N70721 001
		JUN 10, 1986
<u>AB</u>	<u>2MG</u>	N70722 001
		JUN 10, 1986
<u>AB</u>	<u>5MG</u>	N70723 001
		JUN 10, 1986
<u>AB</u>	<u>10MG</u>	N70724 001
		JUN 10, 1986
<u>AB</u>	<u>20MG</u>	N70725 001
		SEP 24, 1986
<u>AB</u>	<u>0.5MG</u>	N70720 001
		JUN 10, 1986
<u>AB</u>	<u>1MG</u>	N70721 001
		JUN 10, 1986
<u>AB</u>	<u>2MG</u>	N70722 001
		JUN 10, 1986
<u>AB</u>	<u>5MG</u>	N70723 001
		JUN 10, 1986
<u>AB</u>	<u>10MG</u>	N70724 001
		JUN 10, 1986
<u>AB</u>	<u>20MG</u>	N70725 001
		SEP 24, 1986

HALOPERIDOL DECANOATEINJECTABLE, INJECTIONHALOPERIDOL DECANOATE
GENSIA SICOR PHARMS

<u>AO</u>	<u>EQ 50MG BASE/ML</u>	N75393 001
		MAY 11, 1999
<u>AO</u>	<u>EQ 100MG BASE/ML</u>	N75393 002
		MAY 11, 1999

HALOPERIDOL LACTATECONCENTRATE, ORALHALOPERIDOL
SCS

<u>AA</u>	<u>EQ 2MG BASE/ML</u>	N70726 001
		JUN 10, 1986
<u>AA</u>	<u>EQ 2MG BASE/ML</u>	N70726 001
		JUN 10, 1986

HALOPERIDOL LACTATEINJECTABLE, INJECTIONHALOPERIDOL
SOLOPAK

<u>AP</u>	<u>EQ 5MG BASE/ML</u>	N70801 001
		DEC 14, 1987
<u>AP</u>	<u>EQ 5MG BASE/ML</u>	N70864 001
		DEC 14, 1987
<u>AP</u>	<u>EQ 5MG BASE/ML</u>	N70801 001
		DEC 14, 1987
<u>AP</u>	<u>EQ 5MG BASE/ML</u>	N70864 001
		DEC 14, 1987
<u>AP</u>	<u>EQ 5MG BASE/ML</u>	N70713 001
		MAY 17, 1988
<u>AP</u>	<u>EQ 5MG BASE/ML</u>	N70744 001
		MAY 17, 1988
<u>AP</u>	<u>EQ 5MG BASE/ML</u>	N70713 001
		MAY 17, 1988
<u>AP</u>	<u>EQ 5MG BASE/ML</u>	N70744 001
		MAY 17, 1988

HEPARIN SODIUMINJECTABLE, INJECTIONHEPARIN LOCK FLUSH
SMITH AND NEPHEW

<u>AP</u>	<u>10 UNITS/ML</u>	N88458 001
		JUL 26, 1984
<u>AP</u>	<u>10 UNITS/ML</u>	N88580 001
		OCT 25, 1984
<u>AP</u>	<u>100 UNITS/ML</u>	N88460 001
		JUL 26, 1984
<u>AP</u>	<u>100 UNITS/ML</u>	N88581 001
		OCT 25, 1984
<u>AP</u>	<u>10 UNITS/ML</u>	N88458 001
		JUL 26, 1984
<u>AP</u>	<u>10 UNITS/ML</u>	N88580 001
		OCT 25, 1984
<u>AP</u>	<u>100 UNITS/ML</u>	N88460 001
		JUL 26, 1984
<u>AP</u>	<u>100 UNITS/ML</u>	N88581 001
		OCT 25, 1984
<u>AP</u>	<u>100 UNITS/ML</u>	N88459 001
		JUL 26, 1984
<u>AP</u>	<u>100 UNITS/ML</u>	N88459 001
		JUL 26, 1984
<u>AP</u>	<u>1,000 UNITS/ML</u>	N88239 001
		JUL 26, 1984

HEPARIN SODIUM

INJECTABLE; INJECTION
HEPARIN SODIUM
@ SMITH AND NEPHEW

1,000 UNITS/ML

N88239 001
JUL 26, 1984

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL
RESERPINE, HYDRALAZINE HCL AND HYDROCHLOROTHIAZIDE
@ SOLVAY

25MG;15MG;0.1MG
N88376 001
OCT 28, 1983

HOMATROPINE METHYLBROMIDE; HYDROCODONE BITARTRATE

SYRUP; ORAL

AA HYDROCODONE BITARTRATE AND HOMATROPINE METHYLBROMIDE
HALSEY 1.5MG/5ML; 5MG/5ML

N40285 001
JUL 19, 1999

HYDRALAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

AP * HYDRALAZINE HCL

AP * LUITFOLD

20MG/ML

N40136 001
JUN 30, 1997
N40136 001
JUN 30, 1997
N88517 001
AUG 22, 1985
N88517 001
AUG 22, 1985

20MG/ML

AP SOLOPAK

20MG/ML

@

20MG/ML

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE

CAPSULE; ORAL

AB HYDRALAZINE HCL AND HYDROCHLOROTHIAZIDE
SOLVAY

25MG; 25MG

N87608 001

FEB 08, 1982

50MG; 50MG

N87213 001

FEB 08, 1982

25MG; 25MG

N87608 001

FEB 08, 1982

50MG; 50MG

N87213 001

FEB 08, 1982

HYDRALAZINE HYDROCHLORIDE; HYDROCHLOROTHIAZIDE; RESERPINE

TABLET; ORAL

BP RESERPINE, HYDRALAZINE HCL AND HYDROCHLOROTHIAZIDE
SOLVAY

25MG;15MG;0.1MG

N88376 001

OCT 28, 1983

HYDROCHLOROTHIAZIDE

TABLET; ORAL

AB HYDROCHLOROTHIAZIDE
MAST MM

25MG

50MG

25MG

50MG

N86192 001

N86192 002

N86192 001

N86192 002

HYDROCHLOROTHIAZIDE; IRBESARTAN

TABLET; ORAL

AVALIDE

@ SANOFI SYNTHELABO

12.5MG; 75MG

12.5MG; 150MG

12.5MG; 300MG

+

AVAPRO HCT

@ SANOFI

12.5MG; 75MG

12.5MG; 150MG

N20758 001

SEP 30, 1997

N20758 002

SEP 30, 1997

N20758 003

AUG 31, 1998

N20758 001

SEP 30, 1997

N20758 002

SEP 30, 1997

HYDROCHLOROTHIAZIDE; TRIAMTERENE

CAPSULE; ORAL

AB TRIAMTERENE AND HYDROCHLOROTHIAZIDE
DURAMED

25MG; 37.5MG

N75052 001

JUN 18, 1999

HYDROCORTISONE

CREAM; TOPICAL

HYDROCORTISONE

@ G AND W LABS

1%

1%

N84059 001

N84059 001

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN'99 - NOV'99

HYDROCORTISONE

ENEMA; RECTAL

CORTENEMA

+ SOLVAY

AB
BR100MG/60ML
100MG/60MLN16199 001
N16199 001HYDROCORTISONE

+ COPLEY PHARM

AB

BR

100MG/60ML
100MG/60MLN74171 001
MAY 27, 1994
N74171 001
MAY 27, 1994

LOTION; TOPICAL

DERMACORT

+ SOLVAY

AT
AT

0.5%

1%

0.5%

1%

N84573 002
N84573 002
N84573 002
N84573 002HYDROCORTISONE

+ THAMES

AT

2.5%

N40247 001
JUL 23, 1999

SOLUTION; TOPICAL

TEXACORT

+ BIOGLAN PHAR

AT

1%

2.5%

N80425 001
N81271 001
APR 17, 1992
N81271 001
APR 17, 1992

+ GENDERM

AT

2.5%

N80425 001

+ MEDICIS

AT

1%

HYDROCORTISONE; NEOMYCIN SULFATE; POLYMYXIN B SULFATE

SOLUTION/DROPS; OTIC

NEO-OTOSOL-HC

+ ALCON

AT

1%; EQ 3.5MG BASE/ML;
10,000 UNITS/MLN62423 001
AUG 25, 19831%; EQ 3.5MG BASE/ML;
10,000 UNITS/MLN62423 001
AUG 25, 1983

+ STERIS

AT

> ADD >
> ADD >
> ADD >
> DLT >
> DLT >
> DLT >HYDROCORTISONE ACETATE

CREAM; TOPICAL

HYDROCORTISONE ACETATE

+ FERNDAL LABS

AT

N40259 001

JUL 29, 1999

2.5%

HYDROCORTISONE SODIUM SUCCINATE

INJECTABLE; INJECTION

HYDROCORTISONE SODIUM SUCCINATE

+ STERIS

AP

AP

AP

AP

AP

AP

AP

AP

EQ 100MG BASE/VIAL
EQ 100MG BASE/VIAL
EQ 100MG BASE/VIAL
EQ 250MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 1GM BASE/VIAL
EQ 100MG BASE/VIAL
EQ 100MG BASE/VIAL
EQ 100MG BASE/VIAL
EQ 250MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 1GM BASE/VIALN84737 002
N84738 001
N84737 001
N84737 001
N84747 001
N84748 001
N84737 002
N84738 001
N84737 001
N84747 001
N84748 001HYDROXYAMPHETAMINE HYDROBROMIDE

SOLUTION/DROPS; OPHTHALMIC

+ PAREDRINE

+ AKORN

+ PHARMICS

1%

1%

N00004 004
N00004 004HYDROXYUREA

CAPSULE; ORAL

HYDROXYUREA

+ PAR PHARM

AB

500MG

N75340 001
FEB 24, 1999HYDROXYZINE HYDROCHLORIDE

INJECTABLE; INJECTION

HYDROXYZINE HCL

+ SOLOPAK

AP

AP

AP

25MG/ML

50MG/ML

50MG/ML

25MG/ML

50MG/ML

50MG/ML

N87591 001
N87593 001
N87595 001
N87591 001
N87591 001
N87593 001
N87595 001IBUPROFEN

SUSPENSION; ORAL

MOTRIN

+ MCNEIL

AB

100MG/5ML

N13842 001
SEP 19, 1989

IBUPROFEN

SUSPENSION; ORAL
MOTRIN

AB + MCNEIL CONS

100MG/5ML

N19842 001
SEP 19, 1989

TABLET; ORAL
IBUPROFEN
NORTON HN

AB 400MG

AB 600MG

AB 800MG

AB ZENITH GOLDLINE

AB 400MG

AB 600MG

AB 800MG

MOTRIN
MCNEIL

AB 300MG

AB 400MG

AB 600MG

AB 800MG

AB 100MG

AB MCNEIL CONS

AB 300MG

AB 400MG

AB 600MG

AB 800MG

AB 100MG

TABLET, CHEWABLE; ORAL
MOTRIN
MCNEIL

AB 50MG

AB 100MG

AB 50MG

AB 100MG

AB MCNEIL CONS

AB 100MG

IMIGLUCERASE

INJECTABLE; INJECTION

CEREZYME
* GENZYME

200 UNITS/VIAL

N20367 001
MAY 23, 1994

200 UNITS/VIAL

N20367 001
MAY 23, 1994

400 UNITS/VIAL

N20367 002
SEP 22, 1999

INDOMETHACIN

CAPSULE, EXTENDED RELEASE; ORAL

INDOCIN SR
EON

AB 75MG

AB 75MG

AB 75MG

AB 75MG

AB 75MG

N74464 001
MAY 28, 1998

N74464 001
MAY 28, 1998

N18185 001
FEB 23, 1982

N18185 001
FEB 23, 1982

N18185 001
FEB 23, 1982

N18185 001
FEB 23, 1982

INSULIN HUMAN

INJECTABLE; INJECTION

VELOSULIN BR
+ NOVO NORDISK

100 UNITS/ML

N21028 001
JUL 19, 1999

IODAMIDE MEGLUMINE

INJECTABLE; INJECTION
RENOVUE-65

* BRACCO
@

65%

N17902 001
N17902 001

IPRATROPIUM BROMIDE

SOLUTION; INHALATION
IPRATROPIUM BROMIDE

AN ALPHARMA

0.02%

N75111 001
APR 22, 1999

ISOETHARINE HYDROCHLORIDESOLUTION; INHALATIONISOETHARINE HCLINTL MEDICATION

AN	0.08%
AN	0.1%
AN	0.1%
AN	0.167%
AN	0.167%
AN	0.25%
AN	0.25%
AN	0.08%

ISOETHARINE HCL S/FDEX

AN	0.08%
AN	0.1%
AN	0.17%
AN	0.25%
AN	0.08%
AN	0.1%
AN	0.17%
AN	0.25%

ISOFLURANELIQUID; INHALATIONFORANE

AN	99.9%
AN	99.9%
AN	99.9%

ISOFLURANEHALOCARBON PRODS

99.9%

ISONIAZID; RIFAMPINCAPSULE; ORALRIFAMATE

AN	150MG; 300MG
AN	150MG; 300MG

DOV PHARMHOECHST MARION RSSLISOSORBIDE DINITRATETABLET; ORALSORBITRATEZENECA

30MG

@

30MG

ISOSORBIDE MONONITRATETABLET, EXTENDED RELEASE; ORALISOSORBIDE MONONITRATEBRIGHTSTONE

60MG

AB

N75166 001
OCT 07, 1999ISOTRETINOINCAPSULE; ORALACCUTANEHLR

10MG

N18662 002

MAY 07, 1982

N18662 004

MAR 28, 1983

N18662 003

MAY 07, 1982

N18662 002

MAY 07, 1982

N18662 004

MAR 28, 1983

N18662 003

MAY 07, 1982

N18662 002

MAY 07, 1982

N18662 004

MAR 28, 1983

N18662 003

MAY 07, 1982

ITRACONAZOLEINJECTABLE; INJECTIONSPORANOX+ JANSSEN

10MG/ML

N20966 001

MAR 30, 1999

KANAMYCIN SULFATEINJECTABLE; INJECTIONKANAMYCIN SULFATESOLOPAX

EQ 75MG BASE/2ML

N62605 003

FEB 26, 1986

KANAMYCIN SULFATE

INJECTABLE; INJECTION

KANAMYCIN SULFATE

SOLOPAK

<u>AP</u>	<u>EQ 500MG BASE/2ML</u>	<u>N62605 001</u>	<u>AB</u>	<u>ORUVAIL</u>	<u>150MG</u>	<u>N19816 002</u>
		FEB 26, 1986		WYETH AYERST		FEB 08, 1995
<u>AP</u>	<u>EQ 1GM BASE/3ML</u>	<u>N62605 002</u>			<u>100MG</u>	<u>N19816 003</u>
		FEB 26, 1986				FEB 08, 1995
	<u>EQ 75MG BASE/2ML</u>	<u>N62605 003</u>			<u>150MG</u>	<u>N19816 002</u>
		FEB 26, 1986				FEB 08, 1995
	<u>EQ 500MG BASE/2ML</u>	<u>N62605 001</u>				
		FEB 26, 1986				
	<u>EQ 1GM BASE/3ML</u>	<u>N62605 002</u>				
		FEB 26, 1986				

KETOCONAZOLE

TABLET; ORAL

KETOCONAZOLE

APPLIED ANAL

<u>AB</u>	<u>200MG</u>	<u>N75341 001</u>	<u>AP</u>	<u>ABBOTT</u>	<u>15MG/ML</u>	<u>N74993 001</u>
		JUL 27, 1999				JAN 27, 1999
<u>AB</u>	<u>200MG</u>	<u>N75314 001</u>			<u>30MG/ML</u>	<u>N74993 002</u>
		JUN 15, 1999				JAN 27, 1999
<u>AB</u>	<u>200MG</u>	<u>N74971 001</u>	<u>AP</u>	<u>BAXTER PHARM PROD</u>	<u>15MG/ML</u>	<u>N75299 001</u>
		JUN 15, 1999				NOV 03, 1999
<u>AB</u>	<u>200MG</u>	<u>N75362 001</u>	<u>AP</u>	<u>BEDFORD</u>	<u>30MG/ML</u>	<u>N75299 002</u>
		JUN 15, 1999			<u>15MG/ML</u>	NOV 03, 1999
<u>AB</u>	<u>200MG</u>	<u>N75319 001</u>	<u>AP</u>		<u>15MG/ML</u>	<u>N75222 001</u>
		JUN 15, 1999				APR 26, 1999
<u>AB</u>	<u>200MG</u>	<u>N75273 001</u>	<u>AP</u>		<u>30MG/ML</u>	<u>N75230 002</u>
		JUN 15, 1999				OCT 25, 1999
<u>AB</u>	<u>200MG</u>	<u>N18533 001</u>	<u>AP</u>		<u>30MG/ML</u>	<u>N75222 002</u>
		<u>N18533 001</u>				APR 26, 1999
<u>AB</u>	<u>200MG</u>		<u>AP</u>		<u>30MG/ML</u>	<u>N75228 001</u>
						APR 26, 1999
<u>AB</u>	<u>200MG</u>		<u>AP</u>		<u>30MG/ML</u>	<u>N75230 001</u>
						APR 26, 1999
<u>AB</u>	<u>200MG</u>		<u>AP</u>		<u>30MG/ML</u>	<u>N75230 001</u>
						OCT 25, 1999

KETOPROFEN

CAPSULE, EXTENDED RELEASE; ORAL

KETOPROFEN

ANDRX PHARMS

<u>AB</u>	<u>100MG</u>	<u>N75270 002</u>	<u>AB</u>	<u>ORUVAIL</u>	<u>10MG</u>	<u>N75284 001</u>
		MAR 24, 1999		WYETH AYERST		JUN 23, 1999
<u>AB</u>	<u>150MG</u>	<u>N75270 003</u>				
		MAR 24, 1999				
<u>AB</u>	<u>200MG</u>	<u>N75270 001</u>				
		MAR 24, 1999				
<u>AB</u>	<u>100MG</u>	<u>N19816 003</u>				
		FEB 08, 1995				

KETOPROFEN

CAPSULE, EXTENDED RELEASE; ORAL

ORUVAIL

WYETH AYERST

<u>AB</u>	<u>150MG</u>	<u>N19816 002</u>
		FEB 08, 1995
<u>AB</u>	<u>100MG</u>	<u>N19816 003</u>
		FEB 08, 1995
<u>AB</u>	<u>150MG</u>	<u>N19816 002</u>
		FEB 08, 1995

KETOROLAC TROMETHAMINE

INJECTABLE; INJECTION

KETOROLAC TROMETHAMINE

ABBOTT

<u>AP</u>	<u>15MG/ML</u>	<u>N74993 001</u>
		JAN 27, 1999
<u>AP</u>	<u>30MG/ML</u>	<u>N74993 002</u>
		JAN 27, 1999
<u>AP</u>	<u>15MG/ML</u>	<u>N75299 001</u>
		NOV 03, 1999
<u>AP</u>	<u>30MG/ML</u>	<u>N75299 002</u>
		NOV 03, 1999
<u>AP</u>	<u>15MG/ML</u>	<u>N75222 001</u>
		APR 26, 1999
<u>AP</u>	<u>15MG/ML</u>	<u>N75230 002</u>
		OCT 25, 1999
<u>AP</u>	<u>30MG/ML</u>	<u>N75222 002</u>
		APR 26, 1999
<u>AP</u>	<u>30MG/ML</u>	<u>N75228 001</u>
		APR 26, 1999
<u>AP</u>	<u>30MG/ML</u>	<u>N75230 001</u>
		OCT 25, 1999

SOLUTION/DROPS; OPHTHALMIC

ACULAR

+ ALLERGAN

0.5%

* SYNTAX

0.5%

		<u>N19700 001</u>
		NOV 09, 1992
		<u>N19700 001</u>
		NOV 09, 1992

TABLET; ORAL

KETOROLAC TROMETHAMINE

SIDMAK LABS CA

10MG

<u>AB</u>	<u>10MG</u>	<u>N75284 001</u>
		JUN 23, 1999

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN'99 - NOV'99

KETOTIFEN FUMARATE

SOLUTION/DROPS; OPHTHALMIC
ZADITOR
+ CIBA

EQ 0.025% BASE

N21066 001
JUL 02, 1999

N18716 003
AUG 01, 1984
N18716 004
AUG 01, 1984

300MG
400MG

LABETALOL HYDROCHLORIDE

TABLET; ORAL

AB TRANDATE
GLAXO WELLCOME

⊕

LABETALOL HYDROCHLORIDE

INJECTABLE; INJECTION

LABETALOL HCL

ABBOTT

5MG/ML

N75239 001

NOV 29, 1999

5MG/ML

N75240 001

NOV 29, 1999

5MG/ML

N75355 001

NOV 29, 1999

5MG/ML

N75303 001

MAY 28, 1999

5MG/ML

N75242 001

SEP 30, 1999

5MG/ML

N75431 001

NOV 29, 1999

5MG/ML

N75524 001

NOV 29, 1999

SOLUTION; ORAL

LACTULOSEAA PACO10GM/15ML

N73160 001

AUG 25, 1992

10GM/15ML

N73160 001

AUG 25, 1992

SOLUTION; ORAL, RECTAL

LACTULOSEAA PACO10GM/15ML

N72029 001

AUG 25, 1992

10GM/15ML

N72029 001

AUG 25, 1992

TABLET; ORAL

LABETALOL HCL
MUTUAL PHARM

100MG

N75215 001

JUL 29, 1999

200MG

N75215 002

JUL 29, 1999

300MG

N75215 003

JUL 29, 1999

100MG

N18716 001

MAY 24, 1985

200MG

N18716 002

AUG 01, 1984

300MG

N18716 003

AUG 01, 1984

400MG

N18716 004

AUG 01, 1984

100MG

N18716 001

MAY 24, 1985

200MG

N18716 002

AUG 01, 1984

SOLUTION; ORAL

EPIVIR-HBV

* GLAXO WELLCOME5MG/ML

N20596 002

DEC 08, 1998

5MG/ML

N20596 002

DEC 08, 1998

5MG/ML

N21004 001

DEC 08, 1998

+

TABLET; ORAL

EPIVIR-HBV

* GLAXO WELLCOME100MG

N20564 002

DEC 08, 1998

100MG

N20564 002

DEC 08, 1998

100MG

N21003 001

DEC 08, 1998

+

LEUCOVORIN CALCIUM

INJECTABLE; INJECTION

LEUCOVORIN CALCIUM

AP * ABBOTT EQ 10MG BASE/ML
AP * BEDFORD EQ 200MG BASE/VIAL

LEUCOVORIN CALCIUM PRESERVATIVE FREE
+ ABBOTT EQ 10MG BASE/ML

AP + BEDFORD EQ 200MG BASE/VIAL
AP BIGNAR EQ 10MG BASE/ML
AP EQ 200MG BASE/VIAL
+ EQ 500MG BASE/VIAL
AP GENSLA SICOR PHARMS EQ 10MG BASE/ML

N40147 001
JUN 25, 1997
N40056 001
MAY 23, 1995
N40286 001
FEB 26, 1999
N40258 001
FEB 26, 1999
N40286 001
FEB 26, 1999
N40332 001
JUN 28, 1999

> ADD >
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TABLET; ORAL
KEPPRA
UCB

N21035 001
NOV 30, 1999
N21035 002
NOV 30, 1999
N21035 003
NOV 30, 1999

LEVOBUPIVACAINE HYDROCHLORIDE

INJECTABLE; INJECTION
CHIROCAINE
DARWIN DISCOVERY

EQ 2.5MG BASE/ML
EQ 5MG BASE/ML
EQ 7.5MG BASE/ML

N20997 001
AUG 05, 1999
N20997 002
AUG 05, 1999
N20997 003
AUG 05, 1999

TABLET; ORAL

LEUCOVORIN CALCIUM

AB INVAMED EQ 15MG BASE

WELLCOVORIN
GLAXO WELLCOME

AB EQ 5MG BASE
AB EQ 25MG BASE
EQ 5MG BASE
EQ 25MG BASE

N75327 001
MAR 24, 1999
N18342 001
JUL 08, 1983
N18342 002
JUL 08, 1983
N18342 001
JUL 08, 1983
N18342 002
JUL 08, 1983

LEVOMETHADYL ACETATE HYDROCHLORIDE

CONCENTRATE; ORAL
ORLAAM
* BIODEVELOPMENT

10MG/ML
10MG/ML

N20315 001
JUL 09, 1993
N20315 001
JUL 09, 1993

LEVONORGESTREL

TABLET; ORAL

PLAN B
+ WOMENS CAPITAL

0.75MG

N21045 001
JUL 28, 1999

LEVALBUTEROL HYDROCHLORIDE

SOLUTION; INHALATION

XOPENEX

+ SEPRACOR

EQ 0.021% BASE

EQ 0.042% BASE

N20837 001
MAR 25, 1999
N20837 002
MAR 25, 1999

LIDOCAINE

FILM, EXTENDED RELEASE; TOPICAL

LIDODERM

+ TEIKOKU PHARMA USA 700MG/12HR

N20612 001
MAR 19, 1999

LORACARBEE

POWDER FOR RECONSTITUTION; ORAL

LORABID
+ KING PHARMS 200MG/5ML

LILLY 100MG/5ML

+ 200MG/5ML

N50667 002
DEC 31, 1991
N50667 001
DEC 31, 1991
N50667 002
DEC 31, 1991

LOTEPREDNOL ETABONATE

SUSPENSION/DROPS; OPHTHALMIC

LOTEMAX
+ PHARMOS 0.5%

@ 0.5%

N20841 001
MAR 09, 1998
N20841 001
MAR 09, 1998

MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; POTASSIUM PHOSPHATE,
MONOBASIC; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM GLUCONATE,
SODIUM PHOSPHATE, DIBASIC

INJECTABLE; INJECTION

ISOLYTE S PH 7.4 IN PLASTIC CONTAINER
B. BRAUN

30MG/100ML; 37MG/100ML; 0.82MG/100ML;
370MG/100ML; 530MG/100ML; 500MG/100ML;
12MG/100ML N19006 001

APR 04, 1984
30MG/100ML; 37MG/100ML; 0.82MG/100ML;
370MG/100ML; 530MG/100ML; 500MG/100ML;
12MG/100ML N19006 001

APR 04, 1984

MALATHION

LOTION; TOPICAL
OVIDE

@ MEDICIS

0.5%

0.5%

N18613 001
AUG 02, 1982
N18613 001
AUG 02, 1982

MANNITOL

INJECTABLE; INJECTION

MANNITOL 25%

AP STERIS

12.5GM/50ML

@ 12.5GM/50ML

N87460 001
JUN 27, 1983
N87460 001
JUN 27, 1983

MEBENDAZOLE

TABLET, CHEWABLE; ORAL

VERMOX

AB + JENSEN

AB + MCNEIL CONS

100MG

100MG

N17481 001
N17481 001

MECLOCYCLINE SULFOSALICYLATE

CREAM; TOPICAL

MECLAN

+ JOHNSON AND JOHNSON 1%

@ 1%

N50518 001
N50518 001

MEDROXYPROGESTERONE ACETATE

TABLET; ORAL

CYCRIN

ESI

2.5MG

5MG

10MG

2.5MG

5MG

10MG

N81239 001
OCT 30, 1992
N81240 001
OCT 30, 1992
N89386 001
SEP 09, 1987
N81239 001
OCT 30, 1992
N81240 001
OCT 30, 1992
N89386 001
SEP 09, 1987

MENOTROPINS (FSH; LH)

INJECTABLE; INJECTION

REPRONEX

FERRING

75 IU/VIAL; 75 IU/VIAL

N21047 001
AUG 27, 1999

INJECTABLE; INJECTION
REPRONEX
FERRING

150 IU/VIAL;150 IU/VIAL N21047 002
AUG 27, 1999

LECTURE: INJECTION

AP	DEMEROL	25MG/ML
AP	ABBOTT	50MG/ML
AP	+	75MG/ML
AP	+	100MG/ML
AP	+	25MG/ML
AP	+	50MG/ML
AP	+	75MG/ML
AP	+	100MG/ML
AP	SANOFI	SYNTHELABO

MEPERIDINE HCL	10MG/ML	N88432 001
ABBOTT		AUG 16, 1984
ASTRA PHARMS	10MG/ML	N81002 001
		JUL 30, 1993
INTL MEDICATION	10MG/ML	N81309 001
		AUG 30, 1993
WALLINCKRODT	10MG/ML	N40163 001
		MAY 12, 1997
STERIS	10MG/ML	N73443 001
		MAR 17, 1992

FREE PRESERVATIVE HCL. FREE

<u>AP</u>	<u>MEPERIDINE HCL INJECTION</u>	<u>10MG/ML</u>	<u>N88432 001</u>
<u>AP</u>	<u>ABBOTT</u>	<u>10MG/ML</u>	<u>AUG 16, 1984</u>
<u>AP</u>	<u>ASTRA PHARMS</u>	<u>10MG/ML</u>	<u>N81002 001</u>
<u>AP</u>	<u>FAULDING</u>	<u>10MG/ML</u>	<u>JUL 30, 1993</u>
<u>AP</u>	<u>INTL MEDICATION</u>	<u>10MG/ML</u>	<u>N40305 001</u>
<u>AP</u>	<u>MALLINCKRODT</u>	<u>10MG/ML</u>	<u>MAR 10, 1999</u>
<u>AP</u>	<u>STERIS</u>	<u>10MG/ML</u>	<u>N81309 001</u>
<u>AP</u>			<u>AUG 30, 1993</u>
<u>AP</u>			<u>N40163 001</u>
<u>AP</u>			<u>MAY 12, 1997</u>
<u>AP</u>			<u>N73443 001</u>
<u>AP</u>			<u>MAR 17, 1992</u>

SYRUP: ORAL

DEMEROL	50MG/5ML	N05010 005
+ ABBOTT	50MG/5ML	N05010 005

TABLET: ORAL

AA	DEMEROL	50MG	N05010 001
+	ABBOTT		
AA		100MG	N05010 004
+		50MG	N05010 001
AA	SANOFI SYNTHELABO	100MG	N05010 004
+			
AA	MEPERIDINE HCL	50MG	N40331 001
AA	AMIDE PHARM		MAY 28, 1999
AA		100MG	N40331 002
AA			MAY 28, 1999
AA		50MG	N40318 001
AA	DURAMED		OCT 05, 1999
AA		100MG	N40318 002
AA			OCT 05, 1999

MEPROBAMATE

AA **TABLET; ORAL**
MEPROBAMATE
PHARMAVITE (4)

THEOPHYLLINE. SODIUM. THEOPHYLLINE

INJECTABLE: INJECTION
MERSALYL-THEOPHYLLINE
* STERIS

N84875 001
N84875 001

METFORMIN HYDROCHLORIDE

TABLET; ORAL
GLUCOPHAGE
+ BRISTOL MYERS SQUIBB 500MG

N20357 001
MAR 03, 1995
N20357 003
NOV 05, 1998
N20357 004
NOV 05, 1998
N20357 001
MAR 03, 1995
N20357 003
NOV 05, 1998

2

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN'99 - NOV'99

40

METFORMIN HYDROCHLORIDE

TABLET; ORAL
GLUCOPHAGE
@ BRISTOL MYERS SQUIBB 750MG

N20357 004
NOV 05, 1998

METHOTREXATE SODIUM

INJECTABLE; INJECTION

METHOTREXATE

BIGMAR EQ 25MG BASE/ML

METHOTREXATE PRESERVATIVE FREE

BIGMAR EQ 25MG BASE/ML

EQ 1GM BASE/VIAL

METHOTREXATE SODIUM

* LEDERLE

EQ 1GM BASE/VIAL

METHOTREXATE SODIUM PRESERVATIVE FREE

EQ 1GM BASE/VIAL

TABLET; ORAL

METHOTREXATE SODIUM

DURAMED EQ 2.5MG BASE

METHOTRIMEPAZINE

INJECTABLE; INJECTION

LEVOPROME

+ IMMUNEX

20MG/ML

20MG/ML

METHOXSALEN

INJECTABLE; INJECTION

UVADEX

+ THERAKOS

0.02MG/ML

METHYLDOPATE HYDROCHLORIDE

INJECTABLE; INJECTION

METHYLDOPATE HCL

SMITH AND NEPHEW

50MG/ML

@

50MG/ML

N70841 001
JAN 02, 1987
N70841 001
JAN 02, 1987

METHYLPHENIDATE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

METADATE ER

MEDEVA

10MG

N40306 001
OCT 20, 1999

METHYLPREDNISOLONE SODIUM SUCCINATE

INJECTABLE; INJECTION

METHYLPREDNISOLONE SODIUM SUCCINATE

STERIS

EQ 40MG BASE/VIAL

EQ 125MG BASE/VIAL

EQ 500MG BASE/VIAL

EQ 1GM BASE/VIAL

EQ 40MG BASE/VIAL

EQ 125MG BASE/VIAL

EQ 500MG BASE/VIAL

EQ 1GM BASE/VIAL

N86953 001
JUL 22, 1982
N87030 001
JUL 22, 1982
N88523 001
JUL 24, 1984
N88524 001
JUL 24, 1984
N86953 001
JUL 22, 1982
N87030 001
JUL 22, 1982
N88523 001
JUL 24, 1984
N88524 001
JUL 24, 1984

METOCLOPRAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

METOCLOPRAMIDE HCL

SMITH AND NEPHEW

EQ 5MG BASE/ML

@

EQ 5MG BASE/ML

N20969 001
FEB 25, 1999

N70623 001
MAR 02, 1987
N70623 001
MAR 02, 1987

METOLAZONE

TABLET; ORAL
ZAROXOLYN
MEDVEA
+

2.5MG
2.5MG

N17386 001
N17386 001

N18592 001
OCT 27, 1989
N18592 001
OCT 27, 1989

METOPROLOL TARTRATE

TABLET; ORAL
METOPROLOL TARTRATE
PUREPAC PHARM

50MG

N74380 001

N20682 001

100MG

N74380 002

N20682 002

50MG

N74380 001

DEC 18, 1986

100MG

N74380 002

DEC 18, 1986

METRONIDAZOLE

INJECTABLE; INJECTION

FLAGYL I.V. RTU IN PLASTIC CONTAINER

+ BAXTER HLTHCARE

500MG/100ML

500MG/100ML

500MG/100ML

500MG/100ML

500MG/100ML

500MG/100ML

500MG/100ML

MICONAZOLE NITRATE

INSERT, CREAM; VAGINAL, TOPICAL
MONISTAT DUAL- PAK
+ ADVANCED CARE PRODS

1.2GM, 2%

N63065 002
JUN 10, 1999
N65005 001
MAR 23, 1999
N65005 002
MAR 23, 1999
N63066 001
AUG 14, 1990
N63067 001
JUL 31, 1990

SUPPOSITORY; VAGINAL
MICONAZOLE NITRATE
ALPHARMA US PHARM

200MG

N73508 001

N63066 001

200MG

N73508 001

N63067 002

200MG

N73508 001

SEP 15, 1999

200MG

N73508 001

N63067 001

200MG

N73508 001

JUL 31, 1990

MICONAZOLE NITRATE

TAMPON; VAGINAL
MONISTAT 5

100MG

* ADVANCED CARE PRODS

100MG

@

MIGLITOL

TABLET; ORAL
GLYSET
BAYER

25MG

25MG

50MG

50MG

100MG

100MG

100MG

100MG

25MG

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50MG

100MG

100MG

25MG

25MG

50MG

MINOCYCLINE HYDROCHLORIDE

CAPSULE; ORAL

VECTRIN

WARNER CHILCOTT

EQ 75MG BASE

N63067 002
SEP 15, 1999

0.02MG/ML

SMITH AND NEPHEW

AP

N71671 001
NOV 17, 1987

0.4MG/ML

SMITH AND NEPHEW

AP

N71681 001
NOV 17, 1987MIVACURIUM CHLORIDE

INJECTABLE; INJECTION

MIVACRON

+ ABBOTT

EQ 2MG BASE/ML

N20098 001

NOV 17, 1987

* GLAXO WELLCOME

EQ 2MG BASE/ML

N20098 001

NOV 17, 1987

MIVACRON IN DEXTROSE 5% IN PLASTIC CONTAINER

+ ABBOTT

EQ 0.5MG BASE/ML

N20098 002

NOV 17, 1987

+ EQ 50MG BASE/100ML

N20098 003

NOV 17, 1987

* GLAXO WELLCOME

EQ 0.5MG BASE/ML

N20098 002

NOV 17, 1987

* EQ 50MG BASE/100ML

N20098 003

NOV 17, 1987

NADOLOL

TABLET; ORAL

CORGARD

APOTHECON

20MG

N18063 005

FEB 10, 1986

40MG

N18063 001

FEB 10, 1986

80MG

N18063 002

FEB 10, 1986

120MG

N18063 003

FEB 10, 1986

160MG

N18063 004

FEB 10, 1986

20MG

N18063 005

FEB 10, 1986

+ BRISTOL MYERS SQUIBB

40MG

N18063 001

FEB 10, 1986

80MG

N18063 002

FEB 10, 1986

120MG

N18063 003

FEB 10, 1986

160MG

N18063 004

FEB 10, 1986

NALOXONE HYDROCHLORIDE

INJECTABLE; INJECTION

NALOXONE HCL

SMITH AND NEPHEW

0.02MG/ML

N71671 001
NOV 17, 1987

0.4MG/ML

SMITH AND NEPHEW

N71681 001
NOV 17, 1987

0.02MG/ML

SMITH AND NEPHEW

N71671 001
NOV 17, 1987

0.4MG/ML

SMITH AND NEPHEW

N71681 001
NOV 17, 1987

0.4MG/ML

SMITH AND NEPHEW

N71339 001
NOV 18, 1987

0.4MG/ML

SMITH AND NEPHEW

N71339 001
NOV 18, 1987NALTREXONE HYDROCHLORIDE

TABLET; ORAL

NALTREXONE HCL

AMIDE PHARM

50MG

N75274 001
MAY 26, 1999NANDROLONE DECANOATE

INJECTABLE; INJECTION

NANDROLONE DECANOATE

STERIS

50MG/ML

N88554 001
FEB 10, 1986

50MG/ML

N88554 001
FEB 10, 1986NANDROLONE PHENPROPIONATE

INJECTABLE; INJECTION

DURABOLIN

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

* ORGANON

25MG/ML

N11891 001
JUN 17, 1983

50MG/ML

N11891 002
JUN 17, 1983

25MG/ML

N11891 001
JUN 17, 1983

50MG/ML

N11891 002
JUN 17, 1983

25MG/ML

N11891 001
JUN 17, 1983

50MG/ML

N11891 002
JUN 17, 1983

25MG/ML

N11891 001
JUN 17, 1983

50MG/ML

N11891 002
JUN 17, 1983

25MG/ML

N11891 001
JUN 17, 1983

50MG/ML

N11891 002
JUN 17, 1983

25MG/ML

N11891 001
JUN 17, 1983

50MG/ML

N11891 002
JUN 17, 1983

25MG/ML

N11891 001
JUN 17, 1983

50MG/ML

N11891 002
JUN 17, 1983

25MG/ML

N11891 001
JUN 17, 1983

50MG/ML

N11891 002
JUN 17, 1983

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN'99 - NOV'99

NANDROLONE PHENPROPIONATE

INJECTABLE; INJECTION

NANDROLONE PHENPROPIONATE

* STERIS 25MG/ML

* 50MG/ML

N86386 001
JUN 17, 1983
N87488 001
JUN 17, 1983N20750 001
OCT 01, 1997NAPHAZOLINE HYDROCHLORIDE

SOLUTION/DROPS; OPHTHALMIC

NAPHAZOLINE HCL* AKORN 0.1%
TAYLOR 0.1%N83590 001
N83590 00115MG/5ML; 3.75MG/5ML;
600MG/5ML
15MG/5ML; 3.75MG/5ML;
600MG/5MLN08378 003
N08378 003NAPROXEN

TABLET, DELAYED RELEASE; ORAL

NAPROXEN

* SIDMAK LABS CA

375MG

500MG

N75337 001
MAY 26, 1999
N75337 002
MAY 26, 1999

FILM, EXTENDED RELEASE; TRANSDERMAL

* PROSTEP

* ELAN PHARM

11MG/24HR

22MG/24HR

11MG/24HR

22MG/24HR

N19983 001
JAN 28, 1992
N1983 002
JAN 28, 1992
N19983 001
JAN 28, 1992
N19983 002
JAN 28, 1992NAPROXEN SODIUM

TABLET; ORAL

NAPROXEN SODIUM

* PUREPAC PHARM

EQ 250MG BASE

EQ 500MG BASE

EQ 250MG BASE

EQ 500MG BASE

N74319 001
MAR 20, 1995
N74319 002
MAR 20, 1995
N74319 001
MAR 20, 1995
N74319 002
MAR 20, 1995NITROGLYCERIN

FILM, EXTENDED RELEASE; TRANSDERMAL

NITROGLYCERIN

* MYLAN TECHNOLOGIES

0.1MG/HR

0.2MG/HR

0.4MG/HR

0.6MG/HR

N75076 001
NOV 12, 1999
N75073 001
NOV 12, 1999
N75075 001
NOV 12, 1999
N74992 001
NOV 12, 1999NEDOCROMIL SODIUM

SOLUTION; INHALATION

TILADE

* RHONE POULENC RORER 0.5%

N20750 001
OCT 01, 1997N70633 001
JUN 19, 1986

NITROGLYCERIN

INJECTABLE; INJECTION

NITROGLYCERIN

@ SMITH AND NEPHEW

5MG/ML

N70633 001
JUN 19, 1986

OINTMENT; TRANSFERMAL

NITROGLYCERIN

ALTANA

2%

N87355 001
JUL 08, 1988
N87355 001
JUL 08, 1988

+

2%

SPRAY, METERED; SUBLINGUAL

NITROLINGUAL PUMPSPRAY

+ POHL BOSKAMP

0.4MG/SPRAY

N18705 002
JAN 10, 1997OLANZAPINE

TABLET; ORAL

ZYPREXA

+ LILLY

2.5MG

N20592 001
SEP 30, 1996
N20592 001
SEP 30, 1996

2.5MG

OMEPRAZOLE

CAPSULE, DELAYED REL PELLETS; ORAL

PRILOSEC

+ ASTRA PHARMS

10MG

N19810 003
OCT 05, 1995
N19810 003
OCT 05, 1995

ASTRAZENECA

10MG

ONDANSETRON

TABLET, ORALLY DISINTEGRATING; ORAL

ZOFTRAN ODT

GLAXO WELLCOME

EQ 4MG BASE

N20781 001
JAN 27, 1999

EQ 8MG BASE

N20781 002
JAN 27, 1999ONDANSETRON HYDROCHLORIDE

TABLET; ORAL

ZOFTRAN

+ GLAXO WELLCOME

EQ 8MG BASE

N20103 002
DEC 31, 1992

EQ 8MG BASE

N20103 002
DEC 31, 1992

EQ 24MG BASE

N20103 003
AUG 27, 1999ORLISTAT

CAPSULE; ORAL

XENICAL

+ ROCHE

120MG

N20766 001
APR 23, 1999ORPHENADRINE CITRATE

TABLET, EXTENDED RELEASE; ORAL

ORPHENADRINE CITRATE

KIEL

100MG

N40249 001
JAN 29, 1999OSELTAMIVIR PHOSPHATE

CAPSULE; ORAL

TAMIFLU

+ ROCHE

EQ 75MG BASE

N21087 001
OCT 27, 1999OXYBUTYNIN CHLORIDE

SYRUP; ORAL

OXYBUTYNIN CHLORIDE

AA MIKART

5MG/5ML

N75039 001
JAN 29, 1999

TABLET, EXTENDED RELEASE; ORAL

DITROPAN XL

+ ALZA

15MG

N20897 003
JUN 22, 1999

OXYTETRACYCLINE CALCIUM

SYRUP; ORAL
TERRAMYCIN
* PFIZER
@

EQ 125MG BASE/5ML
EQ 125MG BASE/5ML

N60595 001
N60595 001

OXYTETRACYCLINE HYDROCHLORIDE

INJECTABLE; INJECTION
TERRAMYCIN
* PFIZER
*
@
@

EQ 250MG BASE/VIAL
EQ 500MG BASE/VIAL
EQ 250MG BASE/VIAL
EQ 500MG BASE/VIAL

N60586 001
N60586 002
N60586 001
N60586 002

OXYTETRACYCLINE HYDROCHLORIDE; POLYMYXIN B SULFATE

OINTMENT; OTIC
TERRAMYCIN W/ POLYMYXIN
* PFIZER
@

EQ 5MG BASE/GM;
10,000 UNITS/GM
EQ 5MG BASE/GM;
10,000 UNITS/GM

N61841 001
N61841 001

TABLET; VAGINAL
TERRAMYCIN-POLYMYXIN
* PFIZER
@

EQ 100MG BASE;
100,000 UNITS
EQ 100MG BASE;
100,000 UNITS

N61009 001
N61009 001

PAROMOMYCIN SULFATE

CAPSULE; ORAL
HUMATIN
* FARKEDALE
AA
AA
AA

EQ 250MG BASE
EQ 250MG BASE
EQ 250MG BASE
EQ 250MG BASE

N60521 001
N62310 001
N62310 001
N60521 001

PAROXETINE HYDROCHLORIDE

CAPSULE; ORAL
PAXIL
SMITHKLINE BEECHAM

EQ 10MG BASE
EQ 20MG BASE
EQ 30MG BASE
EQ 40MG BASE
EQ 10MG BASE
EQ 20MG BASE
EQ 30MG BASE
EQ 40MG BASE

N20885 001
OCT 09, 1998
N20885 002
OCT 09, 1998
N20885 003
OCT 09, 1998
N20885 004
OCT 09, 1998
N20885 001
OCT 09, 1998
N20885 002
OCT 09, 1998
N20885 003
OCT 09, 1998
N20885 004
OCT 09, 1998

TABLET, EXTENDED RELEASE; ORAL
PAXIL CR
+ SMITHKLINE BEECHAM

EQ 12.5MG BASE
EQ 25MG BASE

N20936 001
FEB 16, 1999
N20936 002
FEB 16, 1999

PEMIROLAST POTASSIUM

SOLUTION/DROPS; OPHTHALMIC
ALAMAST
+ SANTEN

0.1%

N21079 001
SEP 24, 1999

PEMOLINE

TABLET; ORAL
CYLERT
ABBOTT

37.5MG
75MG
18.75MG
37.5MG
75MG
18.75MG

N16832 002
N16832 003
N16832 001
N16832 002
N16832 003
N16832 001

AB
AB
+
PEMOLINE
COPLEY PHARM

37.5MG

N75030 001
JAN 29, 1999

PEMOLINE

TABLET; ORAL

PEMOLINE
@ COPLEY PHARM

75MG

N75030 002
JAN 29, 1999

37.5MG

N75286 002
JUN 30, 1999

75MG

N75286 003
JUN 30, 1999

PENTAZOCINE HYDROCHLORIDE

TABLET; ORAL

TALWIN 50
@ SANOFI SYNTHELABO
@ STERLING WINTHROP

EQ 50MG BASE
EQ 50MG BASE

N16732 001
N16732 001

PENTOXIFYLLINE

TABLET, EXTENDED RELEASE; ORAL

PENTOXIFYLLINE
IMPAX PHARM

400MG

N75093 001
AUG 10, 1999

400MG

N74874 001
MAY 25, 1999

400MG

N75199 001
SEP 03, 1999

400MG

N75191 001
JUN 09, 1999

PENTOXIL
@ UPSHER SMITH

400MG

N74962 001
MAR 31, 1999

PERINDOPRIL ERBUMINE

TABLET; ORAL

ACEON
@ HUESCHEN

2MG

N20184 001
DEC 30, 1993

4MG

N20184 002
DEC 30, 1993

8MG

N20184 003
DEC 30, 1993

2MG

N20184 001
DEC 30, 1993

@ SOLVAY

PERINDOPRIL ERBUMINE

TABLET; ORAL

ACEON
@ SOLVAY

4MG

N20184 002
DEC 30, 1993

8MG

N20184 003
DEC 30, 1993

PHENDIMETRAZINE TARTRATE

CAPSULE; ORAL

PHENAZINE
MAST MM

35MG

N86523 001

35MG

N86524 001

35MG

N86525 001

35MG

N86523 001

35MG

N86524 001

35MG

N86525 001

TABLET; ORAL

PHENAZINE
MAST MM

35MG

N87305 001

35MG

N87305 001

35MG

N85914 001

35MG

N85914 001

35MG

N85697 001

35MG

N85697 001

PHENTERMINE HYDROCHLORIDE

CAPSULE; ORAL

FASTIN
@ SMITHKLINE BEECHAM

30MG

N17352 001

30MG

N17352 001

30MG

N86945 001

30MG

N86945 001

30MG

N86945 001

PHENYTOIN SODIUM

INJECTABLE; INJECTION

PHENYTOIN SODIUM
STERIS

50MG/ML

N85434 001

PHENYTOIN SODIUM

INJECTABLE; INJECTION
PHENYTOIN SODIUM
@ STERIS

50MG/ML

N85434 001

PIGLITAZONE HYDROCHLORIDE

TABLET; ORAL
ACTOS
TAKEDA

EQ 15MG BASE

N21073 001

EQ 30MG BASE

N21073 002

EQ 45MG BASE

N21073 003

+

POLYETHYLENE GLYCOL 3350

POWDER FOR RECONSTITUTION; ORAL
MIRALAX
+ BRAINTREE

17GM/SCOOPFUL

N20698 001
FEB 18, 1999

POLYMYXIN B SULFATE; TRIMETHOPRIM SULFATE

SOLUTION/DROPS; OPHTHALMIC
TRIMETHOPRIM SULFATE AND POLYMYXIN B SULFATE
AT ALCON

10,000 UNITS/ML;
EQ 1MG BASE/ML

N64211 001
APR 13, 1998

10,000 UNITS/ML;
EQ 1MG BASE/ML

N64211 001
APR 13, 1998

PORACTANT ALPHA

SUSPENSION; INTRATRACHEAL
CUROSURF
+ DEY

80MG/ML

N20744 001
NOV 18, 1999

POTASSIUM CHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL

MICRO-K
KV PHARM
@ ROBINS AH

8MEQ
8MEQ

N18238 001
N18238 001

MICRO-K 10
KV PHARM

10MEQ

N18238 002
MAY 14, 1984
N18238 002
MAY 14, 1984

* ROBINS AH

10MEQ

GRANULE, FOR RECONSTITUTION ER; ORAL

MICRO-K LS
@ KV PHARM

20MEQ/PACKET

N19561 003
AUG 26, 1988

@ ROBINS AH

20MEQ/PACKET

N19561 003
AUG 26, 1988

INJECTABLE; INJECTION

POTASSIUM CHLORIDE
AM PHARM PARTNERS

3MEQ/ML

N80225 003

3MEQ/ML

N80225 003

4MEQ/ML

N80195 004

4MEQ/ML

N80195 004

2MEQ/ML

N86208 001

2MEQ/ML

N89163 001

2MEQ/ML

N89163 001

2MEQ/ML

N89421 001

3MEQ/ML

N86210 001

2MEQ/ML

N86208 001

2MEQ/ML

N89163 001

2MEQ/ML

N89421 001

3MEQ/ML

N86210 001

POTASSIUM CHLORIDE; SODIUM CHLORIDE; TROMETHAMINE

INJECTABLE; INJECTION

THAM-E
+ ABBOTT

370MG/VIAL; 1.75GM/VIAL;

36GM/VIAL

N13025 001

370MG/VIAL; 1.75GM/VIAL;

36GM/VIAL

N13025 001

POTASSIUM CITRATE

POWDER FOR RECONSTITUTION; ORAL
 POTASSIUM CITRATE
 @ MISSION PHARMA 10MEQ/PACKET
 @ 20MEQ/PACKET
 @ UNIV TX 10MEQ/PACKET
 @ 20MEQ/PACKET

N19647 002
 OCT 13, 1988
 N19647 001
 OCT 13, 1988
 N19647 002
 OCT 13, 1988
 N19647 001
 OCT 13, 1988

PREDNISOLONE ACETATE

INJECTABLE; INJECTION
 PREDNISOLONE ACETATE

* STERIS 40MG/ML
 @ 40MG/ML

N83767 001
 N83767 001

SUSPENSION/DROPS; OPHTHALMIC
 ECONOPRED PLUS

AB ALCON 1%
 AB FALCON PHARMS 1%

N17469 001
 N17469 001

POVIDONE-IODINE

SOLUTION/DROPS; OPHTHALMIC
 BETADINE
 + ALCON 5%

* PURDUE FREDERICK 5%

N18634 001
 DEC 17, 1986
 N18634 001
 DEC 17, 1986

INJECTABLE; INJECTION
 HYDELTRASOL

* MERCK EQ 20MG PHOSPHATE/ML
 @ EQ 20MG PHOSPHATE/ML

N11583 002
 N11583 002

PRAZOSIN HYDROCHLORIDE

CAPSULE; ORAL

AB PRAZOSIN HCL
 ZENITH GOLDLINE

EQ 2MG BASE

AB EQ 2MG BASE

AB EQ 5MG BASE

AB EQ 5MG BASE

N71995 001
 MAY 16, 1989
 N71995 001
 SEP 12, 1988
 N71745 001
 MAY 16, 1989
 N71745 001
 SEP 12, 1988

INJECTABLE; INJECTION

BP HYDELTRA-TBA

* MERCK 20MG/ML
 + 20MG/ML

N10562 001
 N10562 001

BP PREDNISOLONE TEBUTATE

* STERIS 20MG/ML

N83362 001
 FEB 17, 1984
 N83362 001
 FEB 17, 1984

PREDNISONE

TABLET; ORAL

AB ORASONE

* SOLVAY 50MG
 @ 50MG

N85999 001
 N85999 001

EX PREDNISONE

* PHARMAVITE 5MG
 @ 5MG

N84662 002
 N84662 002

PROBENECID

TABLET; ORAL

AB BENEMID

* MERCK 500MG

N07898 004

PREDNISOLONE

SYRUP; ORAL

AA HALSEY

15MG/5ML

AA 15MG/5ML

N40287 001
 MAY 28, 1999
 N40323 001
 MAY 13, 1999

AA UDL

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN'99 - NOV'99

PROBENECID

TABLET; ORAL

BENEMID

@ MERCK

PROBENECID

NYLAN

AB

AB +

500MG

500MG500MG

N07898 004

N84211 002

N84211 002

N21019 002
OCT 06, 1999PROCAINAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

PROCAINAMIDE HCL

SMITH AND NEPHEW

100MG/ML500MG/ML

100MG/ML

500MG/ML

100MG/ML500MG/ML

100MG/ML

500MG/ML

STERIS

AP

AP

AP

AP

PROCHLORPERAZINE EDISYLATE

INJECTABLE; INJECTION

PROCHLORPERAZINE EDISYLATE

SMITH AND NEPHEW

EQ 5MG BASE/ML

EQ 5MG BASE/ML

EQ 5MG BASE/ML

EQ 5MG BASE/ML

EQ 5MG BASE/ML

EQ 5MG BASE/ML

STERIS

AP

AP

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

COMPAZINE

SMITHKLINE BEECHAM EQ 10MG BASE

N21019 001

OCT 06, 1999

PROCHLORPERAZINE MALEATE

CAPSULE, EXTENDED RELEASE; ORAL

COMPAZINE

SMITHKLINE BEECHAM EQ 15MG BASE

CAPSULE; ORAL

PROMETRIUM

* SCHERING PLOUGH

100MG

100MG

200MG

300MG

N19781 001
MAY 14, 1998N19781 001
MAY 14, 1998N19781 002
MAY 14, 1998N19781 002
OCT 15, 1999N19781 003
OCT 15, 1999N19781 003
OCT 15, 1999PROMAZINE HYDROCHLORIDE

INJECTABLE; INJECTION

PROMAZINE HCL

STERIS

50MG/ML

25MG/ML

25MG/ML

50MG/ML

50MG/ML

50MG/ML

SPARINE

* WYETH AYERST

+

N84517 001

N84510 001

N84510 001

N84517 001

N10349 006

N10349 006

PROPOFOL

INJECTABLE; INJECTION

DIPRIVAN

+ ZENECA

*

10MG/ML

10MG/ML

N19627 002

JUN 11, 1996

N19627 002

JUN 11, 1996

PROPOFOL

GENSIA SICOR PHARMS

10MG/ML

N75102 001

JAN 04, 1999

PROPRANOLOL HYDROCHLORIDE

CAPSULE, EXTENDED RELEASE; ORAL
INDERAL LA

<u>AB</u>	<u>* WYETH AYERST</u>	<u>60MG</u>	<u>N18553 004</u>
<u>AB</u>	<u>*</u>	<u>80MG</u>	<u>MAR 18, 1987</u>
<u>AB</u>	<u>*</u>	<u>120MG</u>	<u>N18553 002</u>
<u>AB</u>	<u>*</u>	<u>160MG</u>	<u>APR 19, 1983</u>
<u>AB</u>	<u>*</u>	<u>60MG</u>	<u>N18553 003</u>
<u>AB</u>	<u>*</u>	<u>80MG</u>	<u>APR 19, 1983</u>
<u>AB</u>	<u>*</u>	<u>120MG</u>	<u>N18553 001</u>
<u>AB</u>	<u>*</u>	<u>160MG</u>	<u>APR 19, 1983</u>
<u>AB</u>	<u>*</u>	<u>60MG</u>	<u>N18553 004</u>
<u>AB</u>	<u>*</u>	<u>80MG</u>	<u>MAR 18, 1987</u>
<u>AB</u>	<u>*</u>	<u>120MG</u>	<u>N18553 002</u>
<u>AB</u>	<u>*</u>	<u>160MG</u>	<u>APR 19, 1983</u>
<u>AB</u>	<u>*</u>	<u>60MG</u>	<u>N18553 003</u>
<u>AB</u>	<u>*</u>	<u>80MG</u>	<u>APR 19, 1983</u>
<u>AB</u>	<u>*</u>	<u>120MG</u>	<u>N18553 001</u>
<u>AB</u>	<u>*</u>	<u>160MG</u>	<u>APR 19, 1983</u>

PROPRANOLOL HCL
INWOOD LABS

<u>AB</u>	<u>60MG</u>	<u>N72499 001</u>
<u>AB</u>	<u>80MG</u>	<u>APR 11, 1989</u>
<u>AB</u>	<u>120MG</u>	<u>N72500 001</u>
<u>AB</u>	<u>160MG</u>	<u>APR 11, 1989</u>
<u>AB</u>	<u>60MG</u>	<u>N72501 001</u>
<u>AB</u>	<u>80MG</u>	<u>APR 11, 1989</u>
<u>AB</u>	<u>120MG</u>	<u>N72502 001</u>
<u>AB</u>	<u>160MG</u>	<u>APR 11, 1989</u>
<u>AB</u>	<u>60MG</u>	<u>N72499 001</u>
<u>AB</u>	<u>80MG</u>	<u>APR 11, 1989</u>
<u>AB</u>	<u>120MG</u>	<u>N72500 001</u>
<u>AB</u>	<u>160MG</u>	<u>APR 11, 1989</u>
<u>AB</u>	<u>60MG</u>	<u>N72501 001</u>
<u>AB</u>	<u>80MG</u>	<u>APR 11, 1989</u>
<u>AB</u>	<u>120MG</u>	<u>N72502 001</u>
<u>AB</u>	<u>160MG</u>	<u>APR 11, 1989</u>

INJECTABLE; INJECTION
INDERAL

<u>AP</u>	<u>* WYETH AYERST</u>	<u>1MG/ML</u>	<u>N16419 001</u>
<u>AP</u>	<u>* WYETH AYERST</u>	<u>1MG/ML</u>	<u>N16419 001</u>
<u>AP</u>	<u>PROPRANOLOL HCL</u>	<u>1MG/ML</u>	<u>N70137 001</u>
<u>AP</u>	<u>SMITH AND NEEHEW</u>	<u>1MG/ML</u>	<u>APR 15, 1986</u>
<u>AP</u>	<u>SMITH AND NEEHEW</u>	<u>1MG/ML</u>	<u>N70137 001</u>
<u>AP</u>	<u>SMITH AND NEEHEW</u>	<u>1MG/ML</u>	<u>APR 15, 1986</u>

RABEPRAZOLE SODIUM

TABLET, DELAYED RELEASE; ORAL
ACIPHEX
+ EISAI

<u>N20973 002</u>
<u>AUG 19, 1999</u>

RANITIDINE BISMUTH CITRATE

TABLET; ORAL
TRITEC
* GLAXO WELLCOME

<u>400MG</u>
<u>400MG</u>

<u>N20559 001</u>
<u>AUG 08, 1996</u>
<u>N20559 001</u>
<u>AUG 08, 1996</u>

RANITIDINE HYDROCHLORIDE

TABLET; ORAL
RANITIDINE HCL
GRANULET PHARMS

<u>AB</u>	<u>EQ 150MG BASE</u>	<u>N74488 001</u>
<u>AB</u>	<u>EQ 300MG BASE</u>	<u>JUL 31, 1997</u>
<u>AB</u>	<u>EQ 150MG BASE</u>	<u>N74488 002</u>
<u>AB</u>	<u>EQ 300MG BASE</u>	<u>JUL 31, 1997</u>
<u>AB</u>	<u>EQ 150MG BASE</u>	<u>N74488 001</u>
<u>AB</u>	<u>EQ 300MG BASE</u>	<u>JUL 31, 1997</u>
<u>AB</u>	<u>EQ 150MG BASE</u>	<u>N74488 002</u>
<u>AB</u>	<u>EQ 300MG BASE</u>	<u>JUL 31, 1997</u>
<u>AB</u>	<u>EQ 150MG BASE</u>	<u>N75180 001</u>
<u>AB</u>	<u>EQ 300MG BASE</u>	<u>JAN 28, 1999</u>
<u>AB</u>	<u>EQ 150MG BASE</u>	<u>N75180 002</u>
<u>AB</u>	<u>EQ 300MG BASE</u>	<u>JAN 28, 1999</u>

RAPACURONIUM BROMIDE

INJECTABLE; INJECTION
RAPLON
ORGANON

<u>100MG/VIAL</u>
<u>200MG/VIAL</u>

<u>N20984 001</u>
<u>AUG 18, 1999</u>
<u>N20984 002</u>
<u>AUG 18, 1999</u>

REMIFENTANIL HYDROCHLORIDE

INJECTABLE; INJECTION

ULTIVA
ABBOTT

> ADD >
> ADD >
> ADD >
> ADD >
> ADD >
> ADD >
> DLT >
> DLT >
> DLT >
> DLT >
> DLT >

EQ 1MG BASE/VIAL
EQ 2MG BASE/VIAL
EQ 5MG BASE/VIAL
EQ 1MG BASE/VIAL
EQ 2MG BASE/VIAL
EQ 5MG BASE/VIAL

N20630 001
JUL 12, 1996
N20630 002
JUL 12, 1996
N20630 003
JUL 12, 1996
N20630 001
JUL 12, 1996
N20630 002
JUL 12, 1996
N20630 003
JUL 12, 1996

GLAXO WELLCOME

RITONAVIR

SOLUTION; ORAL

NORVIR
ABBOTT

+

80MG/ML
80MG/ML

N20659 001
MAR 01, 1996
N20659 001
MAR 01, 1996

ROFECOXIB

SUSPENSION; ORAL

VIOXX
MERCK

+

12.5MG/5ML
25MG/5ML

N21052 001
MAY 20, 1999
N21052 002
MAY 20, 1999

RIFAMPIN

INJECTABLE; INJECTION

RIFADIN

AP + HOECHST MARION RSSL

+

600MG/VIAL
600MG/VIAL

N50627 001
MAY 25, 1989
N50627 001
MAY 25, 1989

RIFAMPIN
BEDFORD

AP

600MG/VIAL

N64217 001
OCT 29, 1999

TABLET; ORAL

VIOXX
MERCK

+

12.5MG
25MG

N21042 001
MAY 20, 1999
N21042 002
MAY 20, 1999

ROPINIROLE HYDROCHLORIDE

TABLET; ORAL

REQUIP
SMITHKLINE BEECHAM

+

EQ 3MG BASE
EQ 4MG BASE

N20658 006
JAN 27, 1999
N20658 007
JAN 27, 1999

TABLET; ORAL

RISPERDAL

JANSSEN

+

0.25MG
0.5MG

N20272 008
MAY 10, 1999
N20272 007
JAN 27, 1999

ROSIGLITAZONE MALEATE

TABLET; ORAL

AVANDIA
SMITHKLINE BEECHAM

+

EQ 2MG BASE
EQ 4MG BASE
EQ 8MG BASE

N21071 002
MAY 25, 1999
N21071 003
MAY 25, 1999
N21071 004
MAY 25, 1999

RITONAVIR

CAPSULE; ORAL

NORVIR
ABBOTT

100MG

N20945 001
JUN 29, 1999

RX DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN '99 - NOV '99

52

SAQUINAVIR

POWDER; ORAL
BUPHENYL
+ MEDICIS
* UICYCLYD

200MG
200MG

N20828 001
NOV 07, 1997
N20828 001
NOV 07, 1997

3GM/TEASPOONFUL
3GM/TEASPOONFUL

N20573 001
APR 30, 1996
N20573 001
APR 30, 1996

SAQUINAVIR MESYLATE

POWDER; ORAL
BUPHENYL
+ MEDICIS
* UICYCLYD

EQ 200MG BASE
EQ 200MG BASE

N20628 001
DEC 06, 1995
N20628 001
DEC 06, 1995

500MG
500MG

N20572 001
MAY 13, 1996
N20572 001
MAY 13, 1996

SELEGILINE HYDROCHLORIDE

TABLET; ORAL
ELDEPRYL
* SOMERSET

5MG

N19334 001
JUN 05, 1989

AB SELEGILINE HCL
+ SOMERSET

5MG

N19334 001
JUN 05, 1989

SILVER SULFADIAZINE

CREAM; TOPICAL
THERMAZENE
KENDALL LP

1%

N18810 001

DEC 23, 1985

AB SHERWOOD MEDCL

1%

N18810 001
DEC 23, 1985

SIROLIMUS

SOLUTION; ORAL
RAPAMUNE
+ WYETH AYERST

1MG/ML

N21083 001
SEP 15, 1999

+ HUMATROPE
LILLY

BX

6MG/VIAL

N19640 005
FEB 04, 1999

INJECTABLE; INJECTION
GENOTROPIN PRESERVATIVE FREE
* PHARMACIA AND UPJOHN

1.5MG/VIAL

N20280 004
AUG 24, 1995
N20280 001
JAN 27, 1998

0.2MG/VIAL

N20280 002
JAN 27, 1998

0.4MG/VIAL

N20280 003
JAN 27, 1998

0.6MG/VIAL

N20280 005
JAN 27, 1998

0.8MG/VIAL

N20280 008
JAN 27, 1998

1MG/VIAL

N20280 009
JAN 27, 1998

1.2MG/VIAL

N20280 010
JAN 27, 1998

1.4MG/VIAL

N20280 011
JAN 27, 1998

1.5MG/VIAL

N20280 012
JAN 27, 1998

1.6MG/VIAL

N20280 013
JAN 27, 1998

1.8MG/VIAL

N20280 014
JAN 27, 1998

2MG/VIAL

N20280 015
JAN 27, 1998

6MG/VIAL

N20280 016
JAN 27, 1998

SOMATROPIN, BIOSYNTHETICINJECTABLE; INJECTION

HUMATROPE

+ LILLY

12MG/VIAL

+

24MG/VIAL

N19640 006
FEB 04, 1999
N19640 007
FEB 04, 199950MG
100MG
25MGN40353 001
JUL 29, 1999
N40353 002
JUL 29, 1999
N87998 001
OCT 14, 1983SOYBEAN OILINJECTABLE; INJECTION

INTRALIPID 10%

AP + FRESenius KABI 10%

AP + PHARMACIA AND UPJOHN 10%

AP + INTRALIPID 20%

AP + FRESenius KABI 20%

AP + FRESenius KABI 20%

AP + PHARMACIA AND UPJOHN 20%

AP + PHARMACIA AND UPJOHN 20%

AP + INTRALIPID 30%

AP + FRESenius KABI 30%

AP + PHARMACIA AND UPJOHN 30%

AP + PHARMACIA AND UPJOHN 30%

N17643 001
N17643 001
N18449 001
N20248 001
AUG 07, 1996
N18449 001
N20248 001
AUG 07, 1996
N19942 001
DEC 30, 1993
N19942 001
DEC 30, 199310%
15%
30%
10%
15%
30%
10%
30%N83021 001
N83021 002
N83021 003
N83021 001
N83021 002
N83021 003
N40215 001
MAY 25, 1999
N40216 001
MAY 25, 1999SPIRONOLACTONETABLET; ORAL

ALDACTONE

SEARLE

AB 50MG

AB 100MG

AB 50MG

AB 100MG

AB 50MG

AB 100MG

AB 25MG

AB 25MG

N12151 008
DEC 30, 1982
N12151 010
DEC 30, 1983
N12151 008
DEC 30, 1982
N12151 010
DEC 30, 1983
N89424 002
AUG 11, 1999
N89424 003
AUG 11, 1999
N87998 001
OCT 14, 1983500MG
500MG
500MG
500MG
500MGN40091 001
JUL 29, 1994
N40091 001
JUL 29, 1994
N80081 001
N80081 001
N80081 001
N80081 001
N80081 001SPIRONOLACTONE

MUTUAL PHARM

+

+

+

+

+

+

N89424 002
AUG 11, 1999
N89424 003
AUG 11, 1999
N87998 001
OCT 14, 198350MG
100MG
25MGN71556 001
DEC 29, 1987SPIRONOLACTONETABLET; ORAL

SPIRONOLACTONE

PUREPAC PHARM

50MG

100MG

25MG

N40353 001
JUL 29, 1999
N40353 002
JUL 29, 1999
N87998 001
OCT 14, 1983SULFACETAMIDE SODIUMSOLUTION/DROPS; OPHTHALMIC

SODIUM SULFACETAMIDE

AKORN

AT 10%

AT 15%

AT 30%

AT 10%

AT 15%

AT 30%

AT 10%

AT 30%

AT 10%

AT 30%

N83021 001
N83021 002
N83021 003
N83021 001
N83021 002
N83021 003
N40215 001
MAY 25, 1999
N40216 001
MAY 25, 1999SULFADIAZINETABLET; ORAL

SULFADIAZINE

EON

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

AT 500MG

N40091 001
JUL 29, 1994
N40091 001
JUL 29, 1994
N80081 001
N80081 001
N80081 001
N80081 001
N80081 001SULFAMETHOXAZOLE; TRIMETHOPRIMINJECTABLE; INJECTION

SULFAMETHOXAZOLE AND TRIMETHOPRIM

STERIS

80MG/ML; 16MG/ML

AP 80MG/ML; 16MG/ML

N71556 001
DEC 29, 1987

SULFAMETHOXAZOLE, TRIMETHOPRIM

INJECTABLE; INJECTION
SULFAMETHOXAZOLE AND TRIMETHOPRIM
 @ STERIS 80MG/ML; 16MG/ML

N71556 001
 DEC 29, 1987

INJECTABLE; INJECTION
 HEPATOLITE

N/A

N18467 001
 MAR 16, 1982
 N18467 001
 MAR 16, 1982

DUPONT PHARMS

N/A

TACROLIMUS

CAPSULE; ORAL

PROGRAF

* FUJISAWA HLTHCARE

EQ 1MG BASE

N50708 001

APR 08, 1994

INJECTABLE; INJECTION

TECHNETIUM TC-99M LIDOPENIN KIT

N/A

EQ 0.5MG BASE

N50708 003

AUG 24, 1998

DRAXIMAGE

N/A

EQ 1MG BASE

N50708 001

APR 08, 1994

@

N/A

N18489 001
 OCT 31, 1986
 N18489 001
 OCT 31, 1986

TECHNETIUM TC-99M ALBUMIN AGGREGATED KIT

INJECTABLE; INJECTION

A-N STANNOUS AGGREGATED ALBUMIN

@ NORTH AM CHEMS

N/A

N17916 001

APR 08, 1994

INJECTABLE; INJECTION

OSTEOLITE

N/A

N17972 001
 N17972 001

@ SYNCOR PHARMS

N/A

N17916 001

APR 08, 1994

DUPONT PHARMS

N/A

BS

CIS

DUPONT PHARMS

N/A

N17776 001

APR 08, 1994

TECHNETIUM TC-99M PYRO/TRIMETA PHOSPHATES KIT

INJECTABLE; INJECTION
 PYROLITE

CIS

N/A

N17684 001

APR 08, 1994

DUPONT PHARMS

N/A

N17884 001
 N17884 001

DUPONT PHARMS

N/A

N18263 001

MAR 25, 1983

TEMZOLOMIDE

N/A

DUPONT PHARMS

N/A

N18263 001

MAR 25, 1983

CAPSULE; ORAL

5MG

TEMODAR

SCHERING

20MG

N21029 001
 AUG 11, 1999
 N21029 002
 AUG 11, 1999
 N21029 003
 AUG 11, 1999
 N21029 004
 AUG 11, 1999

TECHNETIUM TC-99M DEPREOTIDE

INJECTABLE; INJECTION

NEO TECT KIT

+ DIATIDE

N/A

N21012 001

AUG 03, 1999

+

250MG

TERBUTALINE SULFATEINJECTABLE; INJECTIONBRETHINE

AP * NOVARTIS

1MG/ML
1MG/MLN18571 001
N18571 001

+ BRICANYL

AP * HOECHST MARION RSSL

1MG/ML
1MG/MLN17466 001
N17466 001TABLET; ORALBRETHINE

BP * NOVARTIS

2.5MG

N17849 001

5MG

N17849 002

2.5MG

N17618 001

5MG

N17618 002

2.5MG

N17618 001

5MG

N17618 002

TERIPARATIDE ACETATEINJECTABLE; INJECTIONPARATHAR

* RHONE POULENC RORER

200 UNITS/VIAL

N19498 001

200 UNITS/VIAL

N19498 001

DEC 23, 1987

TESTOSTERONE ENANTHATEINJECTABLE; INJECTIONTESTOSTERONE ENANTHATE

STERIS

AP *

100MG/ML

100MG/ML

N85599 001

N85599 001

TETRACYCLINE HYDROCHLORIDECAPSULE; ORALTETRACYCLINE HCL

PUREPAC PHARM

250MG

N60290 001

500MG

N60290 002

250MG

N60290 001

500MG

N60290 002

TETRACYCLINE HYDROCHLORIDECAPSULE; ORALTETRACYN

AP * PFIZER

250MG

N60082 003

500MG

N60082 004

250MG

N60082 003

500MG

N60082 004

THEOPHYLLINECAPSULE; ORALBRONKODYL

AP * SANOFI SYNTHELABO

100MG

N85264 001

200MG

N85264 002

100MG

N85264 001

200MG

N85264 002

CAPSULE, EXTENDED RELEASE; ORALTHEOCLEAR L.A.-130

130MG

N86569 001

130MG

N86569 001

130MG

N86569 001

130MG

N86569 001

130MG

N86569 001

130MG

N86569 001

130MG

N86569 001

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130MG

N86569 001

130MG

N86569 001

130MG

N86569 001

130MG

N86569 001

THEOPHYLLINE

SYRUP; ORAL

AQUAPHYLLIN

FERDALE LABS

80MG/15ML

80MG/15ML

TABLET, EXTENDED RELEASE; ORAL

SUSTAIRE

ROERIG

100MG

300MG

100MG

300MG

THIOXIXENE HYDROCHLORIDE

CONCENTRATE; ORAL

THIOXIXENE HCL

ALPHARMA

EQ 5MG BASE/ML

EQ 5MG BASE/ML

TIAGABINE HYDROCHLORIDE

TABLET; ORAL

GABITRIL

+ ABBOTT

2MG

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLID

ROCHE

125MG

AB + SYNTEX

250MG

125MG

SYNTEX USA

250MG

TICLOPIDINE HCL

EON

250MG

TICLOPIDINE HYDROCHLORIDE

TABLET; ORAL

TICLOPIDINE HCL

GENPHARM

250MG

AB

N87917 001

JAN 18, 1983

AB

N87917 001

JAN 18, 1983

AB

N85665 001

N85665 002

N85665 001

N85665 002

AB

N85665 002

AB

N85665 002

AB

N85665 002

TIMOLOL MALEATE

SOLUTION, GEL FORMING/DROPS; OPHTHALMIC

TIMOLOL MALEATE

ALCON

N70969 001

OCT 16, 1987

N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

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N70969 001

OCT 16, 1987

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N70969 001

OCT 16, 1987

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N70969 001

OCT 16, 1987

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N70969 001

OCT 16, 1987

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N70969 001

OCT 16, 1987

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N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

AB

N70969 001

OCT 16, 1987

N75161 001
SEP 13, 1999
N75318 001
AUG 20, 1999
N75316 001
NOV 02, 1999
N75253 001
AUG 20, 1999
N75149 001
AUG 20, 1999
N75089 001
JUL 01, 1999

N20963 001
OCT 21, 1998
N20963 002
OCT 21, 1998
N20963 001
OCT 21, 1998
N20963 002
OCT 21, 1998

N74466 001
MAR 25, 1997
N74466 001
MAR 25, 1997
N74261 001
APR 28, 1995
N74262 001
APR 28, 1995
N74261 001
APR 28, 1995
N74262 001
APR 28, 1995

TOBRAMYCIN

SOLUTION/DROPS; OPHTHALMIC

AT * TOBREX 0.3%
 AT * ALCON 0.3%
 AT + FALCON PHARMS

N50541 001
 N50541 001

TOBRAMYCIN SULFATE

INJECTABLE; INJECTION

AP * TOBRAMYCIN SULFATE
 AP * APOTHECON

EQ 40MG BASE/ML

N64026 001
 MAY 31, 1994

EQ 40MG BASE/ML

N64026 001
 MAY 31, 1994

TOPOTECAN HYDROCHLORIDE

INJECTABLE; INJECTION

HYCAMTIN

EQ 4MG BASE/VIAL

N20671 001
 MAY 28, 1996

EQ 4MG BASE/VIAL

N20671 001
 MAY 28, 1996

TRETINOLIN

GEL; TOPICAL

RETIN-A MICRO

* ADV POLYMER

0.1%

+ JOHNSON AND JOHNSON

0.1%

N20475 001
 FEB 07, 1997

N20475 001
 FEB 07, 1997

SOLUTION; TOPICAL

TRETINOLIN

MORTON GROVE

0.05%

N75260 001
 JAN 25, 1999

TRIAMCINOLONE

TABLET; ORAL

ARISTOCORT

BP FUJISAWA HLTHCARE

4MG

N11161 007

N11161 011

TRIAMCINOLONE

TABLET; ORAL

ARISTOCORT

FUJISAWA HLTHCARE

1MG

N11161 009

LEDERLE

2MG

N11161 004

BP

2MG

N11161 007

BP

4MG

N11161 011

BP

8MG

N11161 009

* TRIAMCINOLONE

ROXANE

2MG

N84708 001

4MG

N84709 001

8MG

N84707 001

2MG

N84708 001

4MG

N84709 001

8MG

N84707 001

4MG

N84775 001

4MG

N84775 001

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL

ARISTOCORT

FUJISAWA HLTHCARE

0.025%

N83017 003

0.1%

N83016 004

0.5%

N83015 002

0.025%

N83017 003

0.1%

N83016 004

0.5%

N83015 002

ARISTOCORT A

FUJISAWA HLTHCARE

0.025%

N83017 004

0.025%

N88818 001

OCT 16, 1984

N83016 005

0.1%

N88819 001

OCT 16, 1984

N83015 003

0.5%

N88820 001

OCT 16, 1984

N83017 004

0.025%

N88818 001

OCT 16, 1984

N88819 001

0.1%

N83016 005

OCT 16, 1984

N88819 001

0.5%

N83015 003

OCT 16, 1984

N83017 004

0.025%

N88818 001

OCT 16, 1984

N88819 001

0.1%

N83016 005

OCT 16, 1984

N88819 001

0.5%

N83015 003

TRIAMCINOLONE ACETONIDE

CREAM; TOPICAL
ARISTOCORT A
LEDERLE

0.5%

N88820 001
OCT 16, 1984

TRIAMCINOLONE DIACETATE

SYRUP; ORAL
ARISTOCORT
LEDERLE

2MG/5ML

N11960 004

GEL; TOPICAL

ARISTOGEL
@ FUJISAWA HLTHCARE
LEDERLE

0.1%
0.1%

N83380 001
N83380 001

LOTION; TOPICAL

KENALOG
@ APOTHECON

0.025%
0.1%
0.025%
0.1%

N11602 003
N11602 001
N11602 003
N11602 001

@ BRISTOL MYERS SQUIBB
@

> DLT >
> DLT >
> ADD >
> ADD >

SMITHKLINE BEECHAM

50MG
100MG
50MG
100MG

N13174 001
N13174 002
N13174 001
N13174 002

OINTMENT; TOPICAL

ARISTOCORT
FUJISAWA HLTHCARE

0.1%
0.5%
0.1%
0.5%

N80750 004
N80745 002
N80750 004
N80745 002

ARISTOCORT A
FUJISAWA HLTHCARE

0.1%
0.1%

N80750 003
N88780 001
OCT 01, 1984

0.5%
0.5%

N80745 003
N88781 001
OCT 05, 1984

0.1%
0.1%

N80750 003
N88780 001
OCT 01, 1984

0.5%
0.5%

N80745 003
N88781 001
OCT 05, 1984

TRIAMCINOLONE ACETONIDE
ALPHARMA

0.5%

N89913 001
DEC 23, 1988

0.5%

N89913 001
DEC 23, 1988

TRIAMCINOLONE DIACETATE

SYRUP; ORAL
ARISTOCORT
FUJISAWA HLTHCARE

2MG/5ML

INJECTABLE; INJECTION

TIGAN

+ KING PHARMS
+ ROBERTS LABS

100MG/ML
100MG/ML

N17530 001
N17530 001

TABLET; ORAL

TRICHLORMAS
MAST NM

4MG
4MG

N86259 001
N86259 001

TRIFLURIDINE

SOLUTION/DROPS; OPHTHALMIC

TRIFLURIDINE

1%

N74311 001
OCT 06, 1995
N74311 001
OCT 06, 1995

ALCON

1%

STERIS

1%

TRIHENYPHENIDYL HYDROCHLORIDE

ELIXIR; ORAL

TRIHENYPHENIDYL HCL
MIKART

2MG/5ML

N40251 001
SEP 27, 1999

TRIMETHOENZAMIDE HYDROCHLORIDE

INJECTABLE; INJECTION

TIGAN

+ KING PHARMS
+ ROBERTS LABS

100MG/ML
100MG/ML

N17530 001
N17530 001

VERAPAMIL HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL

VERAPAMIL HCL

DURAMED

120MG

AB

240MG

AB

N75072 001

MAY 25, 1999

N75072 003

MAY 25, 1999

6MG

AB

7.5MG

AB

10MG

AB

N40301 008

JUL 15, 1999

N40301 009

JUL 15, 1999

N40301 001

JUL 15, 1999

VINCRISTINE SULFATE

INJECTABLE; INJECTION

VINCRISTINE SULFATE PFS

AP GENSLA SICOR PHARMS 1MG/ML

N75493 001

SEP 01, 1999

VITAMIN A PALMITATE

CAPSULE; ORAL

DEL-VI-A

DEL RAY LABS

EQ 50,000 UNITS BASE

AA

EQ 50,000 UNITS BASE

N80830 001

N80830 001

25GM/BOT

25GM/BOT

N17605 001

N17605 001

25GM/BOT

25GM/BOT

N18856 001

MAR 26, 1987

N18856 001

MAR 26, 1987

WARFARIN SODIUM

TABLET; ORAL

COMADIN

* DUPONT MERCK

2MG

2MG

2.5MG

2.5MG

5MG

5MG

1MG

2MG

2.5MG

3MG

4MG

5MG

AB

AB

AB

AB

AB

AB

AB

AB

AB

AB

AB

AB

AB

5MG

10MG

N20859 001

AUG 13, 1999

N20859 002

AUG 13, 1999

ZANAMIVIR

CAPSULE; INHALATION

RELENZA

+ GLAXO WELLCOME

5MG

N21036 001

JUL 26, 1999

IBUPROFEN

SUSPENSION; ORAL
IBUPROFEN
ALPHARMA

100MG/5ML

N74916 001
APR 30, 1999

TABLET; ORAL
IBUPROFEN
LNK

200MG

N75010 001
MAR 01, 1999

200MG

N75139 001
MAR 01, 1999

200MG

N71144 001
JAN 20, 1987

200MG

N72901 001
DEC 19, 1991

200MG

N72903 001
DEC 19, 1991

200MG

N71144 001
JAN 20, 1987

200MG

N72901 001
DEC 19, 1991

200MG

N72903 001
DEC 19, 1991

JUNIOR STRENGTH IBUPROFEN
PERRIGO 100MG

N75367 001
APR 22, 1999

IBUPROFEN POTASSIUM

CAPSULE; ORAL
PROVEL

200MG

N20402 001
APR 20, 1995

200MG

N20402 001
APR 20, 1995

200MG

N20402 001
APR 20, 1995

INSULIN ZINC SUSP PURIFIED BEEF

INJECTABLE; INJECTION
LENTE Iletin II

100 UNITS/ML
100 UNITS/ML

N18477 001
N18477 001

MICONAZOLE NITRATE

CREAM, SUPPOSITORY; TOPICAL, VAGINAL
M-ZOLE 3 COMBINATION PACK
ALPHARMA US PHARM 2%,200MG

N74926 001
APR 16, 1999

MICONAZOLE NITRATE COMBINATION PACK
PERRIGO 2%,200MG

N75329 001
APR 20, 1999

SUPPOSITORY; VAGINAL
MICONAZOLE NITRATE
ALPHARMA US PHARM 100MG

N73507 001
NOV 19, 1993

100MG

N73507 001
NOV 19, 1993

100MG

N73507 001
NOV 19, 1993

MINOXIDIL

SOLUTION; TOPICAL
MINOXIDIL (FOR MEN)
PERRIGO 2%

N75357 001
JUL 30, 1999

2%

N75357 002
JUL 30, 1999

2%

N75357 002
JUL 30, 1999

NAPROXEN SODIUM

TABLET; ORAL
NAPROXEN SODIUM
GRANTEC

EQ 200MG BASE

N74635 001
JAN 13, 1997

EQ 200MG BASE

N74635 001
JAN 13, 1997

EQ 200MG BASE

NAPROXEN SODIUM, PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
ALEVE COLD AND SINUS
+ BAYER

N21076 001
NOV 29, 1999

EQ 200MG BASE,120MG

OTC DRUG PRODUCT LIST / CUMULATIVE SUPPLEMENT NUMBER 11 / JAN'99 - NOV'99

NICOTINE

FILM, EXTENDED RELEASE; TRANSDERMAL
PROSTEP
+ ELAN PHARM
11MG/24HR
22MG/24HR
+

N19983 003
DEC 23, 1998
N19983 004
DEC 23, 1998

TABLET; ORAL
ZANTAC 75
+ WARNER LAMBERT
EQ 75MG BASE

N20520 001
DEC 19, 1995

NICOTINE POLACRILEX

GUM, CHEWING; BUCCAL
NICOTINE POLACRILEX
WATSON LAB

EQ 2MG BASE
EQ 4MG BASE

N74507 001
MAR 15, 1999
N74707 001
MAR 19, 1999

CREAM; TOPICAL
LAMISIL
+ NOVARTIS

N20980 001
MAR 09, 1999

TERBINAFINE HYDROCHLORIDE

1%

NONOXYNOL-9

SPONGE; VAGINAL

TODAY
@ ALLENDALE PHARMS
1GM
@ WHITEHALL ROBINS
1GM

N18683 001
APR 01, 1983
N18683 001
APR 01, 1983

PSEUDOEPHEDRINE HYDROCHLORIDE

TABLET, EXTENDED RELEASE; ORAL
PSEUDOEPHEDRINE HCL
PERRIGO
120MG

N75153 001
FEB 26, 1999

RANITIDINE HYDROCHLORIDE

TABLET; ORAL
RANITIDINE HCL
NOVOPHARM
RANBAXY
ZANTAC 75
+ GLAXO WELLCOME

EQ 75MG BASE
EQ 75MG BASE
EQ 75MG BASE

N75094 001
JUN 21, 1999
N75132 001
N20520 001
DEC 19, 1995

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

CUMULATIVE SUPPLEMENT NUMBER 11 NOV '99

This data is provided to the Division of Data Management and Services from the Office of Orphan Products Development and it is not edited prior to publication.

Orphan Product Designations and Approvals List November 1999

Name	Indication Designated	Sponsor & Address
Generic Name		DD= Date Designated
TN=Trade Name		MA=Marketing Approval
111Indium pentetreotide TN= SomatoTher	Treatment of somatostatin receptor positive neuroendocrine tumors.	Louisiana State University Medical Center Foundation 1600 Canal St. 10th Floor New Orleans, LA 70112 DD=06/10/1999
166Ho-DOTMP TN=	Treatment of multiple myeloma.	NeoRx Corporation 410 W. Harrison Seattle, WA 98119 DD=02/10/1999
506U78 TN=	Treatment of chronic lymphocytic leukemia.	Glaxo Wellcome, Inc. Five Moore Dr. P.O. Box 13398 Durham, NC 27709 DD=09/02/1999
6-hydroxymethyla cylfulvene TN=	Treatment of histologically confirmed advanced or metastatic pancreatic cancer.	MGI Pharma, Inc. 6300 West Old Shakoppe Rd. Suite 110 Bloomington, MN 55438 DD=04/06/1999
6-hydroxymethyla cylfulvene TN=	Treatment of ovarian cancer.	MGI Pharma, Inc. 6300 West Old Shakoppe Rd. Suite 110 Bloomington, MN 55438 DD=07/06/1999

Orphan Product Designations and Approvals List November 1999

Name	Generic Name	TN=Trade Name	Indication Designated	Sponsor & Address	DD= Date Designated	MA=Marketing Approval
6-hydroxymethyla cylfulvene TN=	Treatment of renal cell carcinoma.			MGI Pharma, Inc. 6300 West Old Shakoppe Rd. Suite 110 Bloomington, MN 55438 DD=07/27/1999		
AP1903 TN=	Treatment of acute graft-versus-host disease in patients undergoing bone marrow transplantation.			Ariad Gene Therapeutics, Inc. 26 Landsdowne St. Cambridge, MA 02139 DD=11/24/1999		
Alitretinoin TN= Panretin	Topical treatment of cutaneous lesions in patients with AIDS-related Kaposi's sarcoma.			Ligand Pharmaceuticals Inc. 10275 Science Center Drive San Diego, CA 92121 DD=03/24/1998 MA=02/02/1999		
Alpha1-proteinase inhibitor (human) TN=	For slowing the progression of emphysema in alpha1-antitrypsin deficient patients.			Centeon LLC 1020 First Ave. King of Prussia, PA 19406 DD=11/24/1999		
Amifostine TN= Ethyol	Reduction of the incidence of moderate to severe xerostomia in patients undergoing post-operative radiation treatment for head and neck cancer.			U.S. Bioscience, Inc. One Tower Bridge 100 Front Street, Suite 400 Conshohocken, PA 19428 DD=05/12/1998 MA=06/24/1999		

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Amifostine TN= Ethyol	Treatment of myelodysplastic syndromes.	US Bioscience, Inc. One Tower Bridge 100 Front St., Suite 400 West Conshohocken, PA 19428 DD=10/04/1999
Antihemophilic factor/von Willebrand factor complex (human), dried, pasteurized TN= Humate-P	Treatment and prevention of bleeding in hemophilia A (classical hemophilia) in adult patients; and treatment of spontaneous and trauma-induced bleeding episodes in severe von Willebrand disease, and in mild and moderate von Willebrand disease where use of desmopressin is known or suspected to be inadequate in adult and pediatric patients.	Centeon Pharma GmbH Emil-von-Behring-Strasse 76 35041 Marburg Germany DD=10/16/1992 MA=04/01/1999
Artesunate TN=	Treatment of malaria.	World Health Organisation Special Programme for Research and Training in Tropical Diseases Via Appia Geneva 27, Switzerland DD=07/19/1999
Atovaquone TN= Mepron	Prevention of Pneumocystis carinii pneumonia (PCP) in high-risk, HIV-infected patients defined by a history of one or more episodes of PCP and/or a peripheral CD4+ (T4 helper/inducer) lymphocyte count less than or equal to 200/mm ³ .	Glaxo Wellcome Research and Development 5 Moore Drive PO Box 13398 Research Triangle Park, NC 27709 DD=08/14/1991 MA=01/05/1999

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Autologous DNP-conjugated tumor vaccine TN= M-Vax	For adjuvant therapy in melanoma patients with surgically resectable lymph node metastasis (Stage III and limited Stage IV disease).	Avax Technologies, Inc. 4520 Main St. Suite 930 Kansas City, MO 64111 DD=02/23/1999
Azathioprine TN= Imuran	Treatment of oral manifestations of graft-versus-host disease.	Oral Solutions, Inc. 787 Seventh Ave., 48th Floor New York, NY 10019 DD=09/14/1999
Beraprost TN=	Treatment of pulmonary arterial hypertension associated with any New York Heart Association classification (Class I, II, III, or IV).	United Therapeutics Corporation 68 T.W. Alexander Drive, PO Box 14186 Research Triangle Park, NC 27709 DD=04/29/1999
Bexarotene TN= Targretin	Treatment of cutaneous T-cell lymphoma.	Ligand Pharmaceuticals, Inc. 10275 Science Center Dr. San Diego, CA 92121 DD=06/18/1999
Bleomycin TN= Blenoxane	Treatment of pancreatic cancer.	Genetronics, Inc. 11199 Sorrento Valley Rd. San Diego, CA 92121 DD=02/09/1999
Botulinum toxin type A TN= Dysport	Treatment of dynamic muscle contractures in pediatric cerebral palsy patients.	Ipsen Limited 1 Bath Rd., Maidenhead Berkshire SL6 4UH England, UK DD=10/20/1999

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name		Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Busulfan TN= Busulfex		As preparative therapy in the treatment of malignancies with bone marrow transplantation.	Orphan Medical, Inc. 13911 Ridgedale Drive Suite 475 Minnetonka, MN 55305 DD=07/28/1994 MA=02/04/1999
CT-2584 mesylate TN=		Treatment of adult soft tissue sarcoma.	Cell Therapeutics, Inc. 201 Elliott Ave. West Suite 400 Seattle, WA 98119 DD=04/16/1999
CT-2584 mesylate TN=		Treatment of malignant mesothelioma.	Cell Therapeutics, Inc. 201 Elliott Ave. West Seattle, WA 98119 DD=04/16/1999
Caffeine TN= Cafcit		Treatment of apnea of prematurity.	O.P.R. Development, L.P. 1501 Wakarusa Drive Lawrence, KS 66047 DD=09/20/1988 MA=09/21/1999
Coagulation factor VIIa (recombinant) TN= NovoSeven		Treatment of bleeding episodes in hemophilia A or B patients with inhibitors to Factor VIII or Factor IX.	Novo Nordisk Pharmaceuticals, Inc. 100 Overlook Center Suite 200 Princeton, NJ 08540 DD=06/06/1988 MA=03/25/1999
Cytarabine liposomal TN= DepoCyt		Treatment of neoplastic meningitis.	DepoTech Corporation 10450 Science Center Drive San Diego, CA 92121 DD=06/02/1993 MA=04/01/1999

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Decitabine TN=	Treatment of myelodysplastic syndromes.	Pharmachemie B.V. Swensweg 5 2031 GA Haarlem, The Netherlands DD=03/08/1999
Decitabine TN=	Treatment of chronic myelogenous leukemia.	Pharmachemie B.V. Swensweg 5 2031 GA Haarlem, The Netherlands DD=03/08/1999
Denileukin diftitox TN= Ontak	Treatment of patients with persistent or recurrent cutaneous T-cell lymphoma whose malignant cells express the CD25 component of the IL-2 receptor.	Seragen, Inc. 97 South Street Hopkinton, MA 01748 DD=08/21/1996 MA=02/05/1999
Doxorubicin liposome TN= Doxil	Treatment of ovarian cancer.	Alza Corporation 1550 Plymouth St. PO Box 7210 Mountain View, CA 94039 DD=11/04/1998 MA=06/28/1999
Epirubicin TN= Ellence	Treatment of breast cancer.	Pharmacia & Upjohn Company 0634-298-113 7000 Portage Rd. Kalamazoo, MI 49001 DD=09/14/1999 MA=09/15/1999
Epoprostenol TN= Flolan	Treatment of secondary pulmonary hypertension due to intrinsic precapillary pulmonary vascular disease.	Glaxo Wellcome Inc. Five Moore Dr. PO Box 13398 Research Triangle Park, NC 27709 DD=03/22/1999

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Etanercept TN= Enbrel	Reduction in signs and symptoms of moderately to severely active polyarticular-course juvenile rheumatoid arthritis in patients who have had an inadequate response to one or more disease-modifying anti-rheumatic drugs.	Immunex Corporation 51 University St. Seattle, WA 98101 DD=10/27/1998 MA=05/27/1999
Etanercept TN= Enbrel	Treatment of Wegener's granulomatosis.	Immunex Corporation 51 University Street Seattle, WA 98101 DD=04/06/1999
Exemestane TN= Aromasin	Treatment of advanced breast cancer in postmenopausal women whose disease has progressed following tamoxifen therapy.	Pharmacia & Upjohn 7000 Portage Road Mail Stop: 0636-298-113 Kalamazoo, MI 49001 DD=09/19/1991 MA=10/21/1999
Fluoxetine TN= Prozac	Treatment of autism.	Hollander, MD, Eric Mt. Sinai School of Medicine, Dept. of Psychiatry Box 1230, One Gustave L. Levy Place New York, NY 10029 DD=04/30/1999
Gemtuzumab zogamicin TN=	Treatment of CD33-positive acute myeloid leukemia.	Wyeth-Ayerst Laboratories PO Box 8299 Philadelphia, PA 19101 DD=11/24/1999

Orphan Product Designations and Approvals List November 1999

Name	Indication Designated	Sponsor & Address
Generic Name		DD= Date Designated
TN=Trade Name		MA=Marketing Approval
Guanfacine TN= Tenex	Treatment of fragile X syndrome.	Watson Laboratories, Inc. 311 Bonnie Circle PO Box 1900 Corona, CA 91718 DD=08/05/1999
HLA-B7/Beta2M DNA Lipid (DMRIE/DOPE) Complex TN= Allovectin-7	Treatment of invasive and metastatic melanoma (Stages II, III, and IV).	Vical Incorporated 9373 Towne Centre Dr., Suite 100 San Diego, CA 92121 DD=09/30/1999
Humanized MAb (IDEC-131) to CD40L TN=	Treatment of systemic lupus erythematosus.	Idex Pharmaceuticals Corporation 3030 Callan Rd. San Diego, CA 92121 DD=02/09/1999
Interferon beta-1a (recombinant human) TN= Avonex	Treatment of pulmonary fibrosis.	Biogen, Inc. 14 Cambridge Center Cambridge, MA 02142 DD=01/07/1999
Iodine I-131 radiolabeled chimeric MAb tumor necrosis treatment (TNT-1B) TN= 131IohTNT-1	Treatment of glioblastoma multiforme and anaplastic astrocytoma.	Techniclone Corporation 14282 Franklin Ave. Tustin, CA 92780 DD=02/12/1999
Japanese encephalitis vaccine (live, attenuated) TN=	Prevention of Japanese encephalitis.	Boran Pharmaceuticals 3F, Koryo Academytel, 437-3 Ahyun-Dong, Mapo-Gu, Seoul 121-010 South Korea, DD=05/19/1999

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
L-5-hydroxytryptophan TN=	Treatment of tetrahydrobiopterin deficiency.	Watson Laboratories, Inc. 311 Bonnie Circle P.O. Box 1900 Corona, CA 91718 DD=01/20/1999
L-glutamyl-L-tryptophan TN=	Treatment of AIDS-related Kaposi's sarcoma.	Cytran Incorporated 10230 N.E. Points Dr. Suite 530 Kirkland, WA 98033 DD=10/20/1999
Lactic acid TN= Aphthaid	Treatment of severe aphthous stomatitis in severely, terminally immunocompromised patients.	Frontier Pharmaceutical, Inc. SUNY Farmingdale Conklin Hall Farmingdale, NY 11735 DD=06/29/1999
Lidocaine patch 5% TN= Lidoderm Patch	For relief of allodynia (painful hypersensitivity), and chronic pain in post-herpetic neuralgia.	Hind Health Care, Inc. 3707 Williams Rd., Suite 101 San Jose, CA 95117 DD=10/24/1995 MA=03/19/1999
Liposomal-cis-bis-nitrosyl-L-arginine TN=	Treatment of malignant mesothelioma.	Aronex Pharmaceuticals, Inc. 8707 Technology Forest Place The Woodlands, TX 77381 DD=09/01/1999
Lisofylline TN=	Treatment of patients undergoing induction therapy for acute myeloid leukemia.	Cell Therapeutics, Inc. 201 Elliot Ave. W., Suite 400 Seattle, WA 98119 DD=06/10/1999

Orphan Product Designations and Approvals List November 1999

Name	Indication Designated	Sponsor & Address
Generic Name		DD= Date Designated
TN=Trade Name		MA=Marketing Approval
Marijuana TN=	Treatment of HIV-associated wasting syndrome.	Multidisciplinary Association for Psychedelic Studies, Inc. 3 Francis St. Belmont, MA 02478 DD=05/25/1999
Mitoxantrone TN= Novantrone	Treatment of secondary-progressive multiple sclerosis.	Immunex Corporation 51 University St. Seattle, WA 98101 DD=08/13/1999
Mitoxantrone TN= Novantrone	Treatment of progressive-relapsing multiple sclerosis.	Immunex Corporation 51 University St. Seattle, WA 98101 DD=08/13/1999
Murine MAb to polymorphic epithelial mucin, human milk fat globule 1 TN= Theragyn	Adjuvant treatment of ovarian cancer.	Antisoma West Africa House Hanger Lane London W5 3QR, UK DD=03/22/1999
N-acetylgalactos amine-4-sulfatas e, recombinant human TN=	Treatment of mucopolysaccharidosis Type VI (Maroteaux-Lamy syndrome).	BioMarin Pharmaceutical, Inc. 11 Pimental Court Novato, CA 94949 DD=02/17/1999
Parovirus B19 (recombinant VP1 and VP2; S.frugiperda cells) vaccine TN= MEDI-491	Prevention of transient aplastic crisis in patients with sickle cell anemia.	MedImmune, Inc. 35 West Watkins Mill Rd. Gaithersburg, MD 20878 DD=05/07/1999

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Peginterferon alfa-2a TN= PEGASYS	Treatment of chronic myelogenous leukemia.	Hoffman-La Roche Inc. 340 Kingsland St. Nutley, NJ 07110 DD=09/30/1999
Pegylated arginine deiminase TN= Hepacid	Treatment of hepatocellular carcinoma.	Phoenix Pharmacologics, Inc. 115 John Robert Thomas Dr. Exton, PA 19341 DD=03/26/1999
Pegylated arginine deiminase TN= Melanocid	Treatment of invasive malignant melanoma.	Phoenix Pharmacologics, Inc. 115 John Robert Thomas Dr. Exton, PA 19341 DD=04/12/1999
Pentostatin TN= Nipent	Treatment of peripheral T-cell lymphomas.	SuperGen, Inc. Two Annabel Lane, Suite 220 San Ramon, CA 94583 DD=11/24/1999
Polyethylene glycol-modified uricase TN= Zurase	Prophylaxis of hyperuricemia in cancer patients prone to develop tumor lysis syndrome during chemotherapy.	Phoenix Pharmacologics, Inc. 115 John Robert Thomas Dr. Exton, PA 19341 DD=09/14/1999
Recombinant human C1-esterase inhibitor TN=	Prophylactic treatment of angioedema caused by hereditary or acquired C1-esterase inhibitor deficiency.	Pharming N.V. Cipalstreet 3 B-2440 Geel, Belgium DD=02/23/1999

Orphan Product Designations and Approvals List November 1999

Name	Indication Designated	Sponsor & Address
Generic Name		DD= Date Designated
TN=Trade Name		MA=Marketing Approval
Recombinant human C1-esterase inhibitor TN=	Treatment of (acute attacks of) angioedema caused by hereditary or acquired C1-esterase inhibitor deficiency.	Pharming N.V. Cipalstreet 3 B-2440 Geel, Belgium DD=02/23/1999
Recombinant human insulin-like growth factor-I/insulin -like growth factor binding protein-3 TN=	Treatment of major burns that require hospitalization.	Celtrix Pharmaceuticals, Inc. 3055 Patrick Henry Dr. Santa Clara, CA 95054 DD=06/15/1999
Recombinant human nerve growth factor TN=	Treatment of HIV-associated sensory neuropathy.	Genentech, Inc. 1 DNA Way South San Francisco, CA 94080 DD=04/16/1999
Recombinant humanized MAb 5c8 TN=	Prevention of rejection of solid organ transplants.	Biogen, Inc. 14 Cambridge Center Cambridge, MA 02142 DD=03/22/1999
Recombinant humanized MAb 5c8 TN=	Prevention of rejection of pancreatic islet cell transplants.	Biogen, Inc. 14 Cambridge Center Cambridge, MA 02142 DD=03/22/1999
Rifalazil TN=	Treatment of pulmonary tuberculosis.	PathoGenesis Corporation 201 Elliott Avenue West Suite 150 Seattle, WA 98119 DD=04/13/1999

Orphan Product Designations and Approvals List November 1999

Name		Sponsor & Address
Generic Name		DD= Date Designated
TN=Trade Name	Indication Designated	MA=Marketing Approval
SCH 58500 TN=	Treatment of primary ovarian cancer.	Schering Corporation 2000 Galloping Hill Rd. Kenilworth, NJ 07033 DD=04/12/1999
Silver sulfadiazine and cerium nitrate TN= Flammacerium	Prevention of mortality in severely burned patients.	Synthes (USA) 1690 Russell Road PO Box 1766 Paoli, PA 19301 DD=11/17/1999
Sodium 1,3-propanedisul fonate TN=	Treatment of secondary amyloidosis.	Neurochem, Inc. 7220 Frederick Banting, Suite 100 Saint-Laurent, Quebec Canada H4S 2A1 DD=04/06/1999
Sodium dichloroacetate TN= Ceresine	Treatment of severe head injury.	Cypros Pharmaceutical Corporation 2714 Loker Avenue West Carlsbad, CA 92008 DD=06/14/1999
Somatropin (rDNA origin) TN= Neutropin Depot	Long-term treatment of children who have growth failure due to a lack of adequate endogenous growth hormone secretion.	Genentech, Inc. 1 DNA Way South San Francisco, CA 94080 DD=10/28/1999
Somatropin [rDNA] TN= Genotropin	Treatment of short stature in patients with Prader-Willi syndrome.	Pharmacia & Upjohn 7000 Portage Rd. 0633-298-113 Kalamazoo, MI 49001 DD=07/06/1999

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Synthetic human secretin TN=	For use in the evaluation of exocrine pancreas function.	ChiRhoClin, Inc. 1550 Gallaudet Ave. Silver Spring, MD 20905 DD=06/16/1999
Synthetic human secretin TN=	For use in obtaining desquamated pancreatic cells for cytopathologic examination in pancreatic carcinoma.	ChiRhoClin, Inc. 15500 Gallaudet Ave. Silver Spring, MD 20905 DD=06/16/1999
Synthetic human secretin TN=	For use in the diagnosis of gastrinoma associated with Zollinger-Ellison syndrome.	ChiRhoClin, Inc. 15500 Gallaudet Ave. Silver Spring, MD 20905 DD=06/16/1999
Synthetic porcine secretin TN=	For use in the evaluation of exocrine pancreas function.	ChiRhoClin, Inc. 15500 Gallaudet Ave. Silver Spring, MD 20905 DD=06/18/1999
Synthetic porcine secretin TN=	For use in obtaining desquamated pancreatic cells for cytopathologic examination in pancreatic carcinoma.	ChiRhoClin, Inc. 15500 Gallaudet Ave. Silver Spring, MD 20905 DD=06/18/1999
Synthetic porcine secretin TN=	For use in the diagnosis of gastrinoma associated with Zollinger-Ellison syndrome.	ChiRhoClin, Inc. 15500 Gallaudet Ave. Silver Spring, MD 20905 DD=06/18/1999

Orphan Product Designations and Approvals List November 1999

Name Generic Name TN=Trade Name	Indication Designated	Sponsor & Address DD= Date Designated MA=Marketing Approval
Temoporfin TN= Foscan	Palliative treatment of recurrent, refractory or second primary squamous cell carcinomas of the head and neck in patients considered to be incurable with surgery or radiotherapy.	Scotia Pharmaceuticals Ltd. Scotia House, Castle Business Park Stirling Scotland FK9 4TZ DD=10/28/1999
Temozolomide TN= Temodar	Treatment of recurrent malignant glioma.	Schering-Plough Research Institute 2000 Galloping Hill Rd. Kenilworth, NJ 07033 DD=10/05/1998 MA=08/11/1999
Teriparatide TN= Parathar	Treatment of idiopathic osteoporosis.	Henri Beaufour Institute USA, Inc. 27 Maple St. Milford, MA 01757 DD=10/28/1999
Thalidomide TN= Thalomid	Treatment of Crohn's disease.	Celgene Corporation 7 Powder Horn Dr. Warren, NJ 07059 DD=04/06/1999
Tobramycin TN= Tobi	Treatment of bronchiectasis patients infected with <i>Pseudomonas aeruginosa</i> .	PathoGenesis Corporation 201 Elliott Avenue West Suite 150 Seattle, WA 98119 DD=06/18/1999
Transgenic human alpha 1 antitrypsin TN=	Treatment of emphysema secondary to alpha 1 antitrypsin deficiency.	PPL Therapeutics (Scotland) Limited Roslin, Edinburgh EH25 9PP Scotland U.K. DD=05/19/1999

Orphan Product Designations and Approvals List November 1999

Name	Indication Designated	Sponsor & Address
Generic Name		DD= Date Designated
TN=Trade Name		MA=Marketing Approval
Yttrium-90 radiolabeled humanized monoclonal anti-carcinoembr yonic antigen IgG antibody TN= Cea-Cide	Treatment of ovarian carcinoma.	Immunomedics, Inc. 300 American Rd. Morris Plains, NJ 07950 DD=08/03/1999
pVGI.1 (VEGF2) TN=	Treatment of thromboangiitis obliterans.	Vascular Genetics, Inc. 200 Westpark Corporate Center 4364 South Alston Ave. Durham, NC 27713 DD=11/09/1999

DRUG PRODUCTS WHICH MUST DEMONSTRATE *IN VIVO* BIOAVAILABILITY ONLY
IF PRODUCT FAILS TO ACHIEVE ADEQUATE DISSOLUTION

NO NOVEMBER 1999 ADDITIONS

PATENT AND EXCLUSIVITY TERMS

DUE TO SPACE LIMITATIONS IN THE PATENT AND EXCLUSIVITY COLUMNS, ABBREVIATIONS AND REFERENCES HAVE BEEN DEVELOPED. PLEASE REFER BACK TO THE APPROVED DRUG PRODUCTS WITH THERAPEUTIC EQUIVALENCE EVALUATIONS, 19TH EDITION FOR A FULL LISTING OF PATENT AND EXCLUSIVITY TERMS (ABBREVIATIONS, NEW DOSING SCHEDULE, NEW INDICATIONS AND PATENT USE CODES). THE CUMULATIVE SUPPLEMENT WILL LIST NEW CODES ADDED SINCE THE LAST ANNUAL EDITION.

ABBREVIATIONS

NP[®] NEW PRODUCT (MINT FLAVORED)

REFERENCES

NEW DOSING SCHEDULE

D-50 INFORMATION FOR USE OF CORVERT IN POST-CARDIAC SURGERY PATIENTS

NEW INDICATION

- I-250 PRIMARY PREVENTION OF CORONARY HEART DISEASE IN PATIENTS WITHOUT SYMPTOMATIC CARDIOVASCULAR DISEASE WHO HAVE AVERAGE TO MODERATELY ELEVATED TOTAL-C AND LDL-C AND BELOW AVERAGE HDL-C
- I-251 TREATMENT OF GENERALIZED ANXIETY DISORDER
- I-252 NEW COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS METFORMIN
- I-253 COMBINATION USE OF PRECOSE FOR PATIENTS WITH TYPE 2 DIABETES TREATED WITH DIET PLUS INSULIN
- I-254 PREVENTION OF POSTMENOPAUSAL OSTEOPOROSIS (LOSS OF BONE MASS)
- I-255 PREVENTION OF PNEUMOCYSTIS CARINII PNEUMONIA (PCP)
- I-256 USE IN TREATMENT OF SMALL CELL LUNG CANCER SENSITIVE DISEASE AFTER FAILURE OF FIRST-LINE CHEMOTHERAPY
- I-257 TREATMENT OF CHRONIC HEPATITIS B ASSOCIATED WITH EVIDENCE OF HEPATITIS B VIRAL REPLICATION AND ACTIVE LIVER INFLAMMATION
- I-258 FOR PERENNIAL NONALLERGIC RHINITIS FOR AGES FOUR AND ABOVE
- I-259 PROPHYLAXIS OF DEEP VEIN THROMBOSIS (DVT), WHICH MAY LEAD TO PULMONARY EMBOLISM, IN PATIENTS UNDERGOING HIP REPLACEMENT SURGERY
- I-260 EXPANDED PEDIATRIC USE IN CHILDREN YOUNGER THAN ONE MONTH OF AGE TO BIRTH (WITH A GESTATIONAL AGE OF 37 WEEKS OR GREATER)
- I-261 TREATMENT OF SOCIAL ANXIETY DISORDER
- I-262 TREATMENT OR PREVENTION OF BRONCHOSPASM WITH REVERSIBLE OBSTRUCTIVE AIRWAY DISEASE AND FOR THE PREVENTION OF EXERCISE INDUCED BRONCHOSPASM IN CHILDREN AGES 4-12
- I-263 TREATMENT OF UNSTABLE ANGINA AND NON-Q-WAVE MYOCARDIAL INFARCTION FOR THE PREVENTION OF ISCHEMIC COMPLICATIONS IN PATIENTS ON CONCURRENT ASPIRIN THERAPY
- I-264 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH RADIATION, INCLUDING TOTAL BODY IRRADIATION (TBI) AND FRACTIONATED ABDOMINAL RADIATION
- I-265 TREATMENT OF ATOPIC DERMATITIS IN PEDIATRIC PATIENTS 6 YEARS AND OLDER
- I-266 USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN PEDIATRIC PATIENTS AGES 2-16 YEARS WITH PARTIAL ONSET SEIZURES
- I-267 USE IN PEDIATRIC PATIENTS 3 MONTHS OLD AND OLDER FOR CORTICOSTEROID-RESPONSIVE DERMATOSES
- I-268 PROPHYLAXIS AND CHRONIC TREATMENT OF ASTHMA IN PATIENTS 7-11 YEARS OF AGE

PATENT AND EXCLUSIVITY TERMS

NEW INDICATION

- I-269 PREVENTION OF NAUSEA AND VOMITING ASSOCIATED WITH HIGHLY EMETOGENIC CANCER CHEMOTHERAPY, INCLUDING CISPLATIN
- I-270 ADJUVANT TREATMENT OF NODE-POSITIVE BREAST CANCER ADMINISTERED SEQUENTIALLY TO STANDARD DOXORUBICIN-CONTAINING COMBINATION CHEMOTHERAPY
- I-271 TREATMENT OF OSTEOPOROSIS IN POSTMENOPAUSAL WOMEN
- I-272 TREATMENT OF GLUCOCORTICOID-INDUCED OSTEOPOROSIS IN MEN AND WOMEN RECEIVING GLUCOCORTICOID IN A DAILY DOSE EQUIVALENT TO 7.5MG OR GREATER OF PREDNISONE AND WHO HAVE LOW BONE MINERAL DENSITY
- I-273 ADJUNCT TO DIET TO INCREASE HDL-C IN PATIENTS WITH PRIMARY HYPERCHOLESTEROLEMIA (HETEROZYGOUS FAMILIAL AND NONFAMILIAL) AND MIXED DYSLIPIDEMIA (FREDERICKSON TYPES IIA AND IIB)
- I-274 USE OF TOPAMAX AS ADJUNCTIVE THERAPY IN THE TREATMENT OF PRIMARY GENERALIZED TONIC-CLONIC SEIZURES
- I-275 USE IN COMBINATION WITH METFORMIN AND SULFONYLUREAS IN PATIENTS WITH TYPE 2 DIABETES
- I-276 USE OF REZULIN IN COMBINATION WITH METFORMIN AND SULFONYLUREAS IN PATIENTS WITH TYPE 2 DIABETES

REFERENCES

MISCELLANEOUS EXCLUSIVITY CODES

- M-1 INFORMATION REGARDING SUPERIORITY CLAIM OVER RANITIDINE FOR DAY AND NIGHT HEARTBURN ADDED TO CLINICAL STUDIES SECTION

PATENT USE CODE

- U-254 USE OF AGGRASTAT IN COMBINATION WITH HEPARIN
- U-255 IMPROVED WAKEFULNESS IN PATIENTS WITH EXCESSIVE DAYTIME SLEEPINESS ASSOCIATED WITH NARCOLEPSY
- U-256 TREATMENT OF HIV INFECTION IN COMBINATION WITH ONE OR MORE ADDITIONAL HIV ANTIVIRAL AGENTS
- U-257 TREATMENT OF HIV INFECTION
- U-258 TREATMENT OF NEURODEGENERATIVE DISEASES
- U-259 TREATMENT OF ANDROGENIC ALOPECIA BY ORAL ADMINISTRATION OF DRUG SUBSTANCE
- U-260 REDUCTION OF INTRAOCULAR PRESSURE IN PATIENTS WITH OPEN ANGLE GLAUCOMA AND OCULAR HYPERTENSION WHO ARE INTOLERANT OF OTHER IOP LOWERING MEDICATIONS OR INSUFFICIENTLY RESPONSIVE TO ANOTHER IOP LOWERING MEDICATION
- U-261 TREATING BENIGN PROSTATIC HYPERPLASIA WITH A GENUS OF COMPOUNDS, INCLUDING FINASTERIDE
- U-262 TREATING BENIGN PROSTATIC HYPERTROPHY WITH FINASTERIDE
- U-263 METHOD OF TREATING A MALIGNANT CONDITION THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN, METHOD FOR TREATING LEUKEMIA OR LYMPHOMA IN A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVENOUS ADMINISTRATION OF BUSULFAN.
- U-264 METHOD OF TREATING A MALIGNANT DISEASE THROUGH PARENTERAL ADMINISTRATION OF BUSULFAN, METHOD FOR TREATING A PATIENT UNDERGOING A BONE MARROW TRANSPLANT THROUGH INTRAVASCULAR ADMINISTRATION OF BUSULFAN.
- U-265 USE AS A LAXATIVE
- U-266 OSTEOARTHRITIS
- U-267 METHOD FOR PREVENTING HEARTBURN
- U-268 ACROMEGALY
- U-269 EXCESS GH-SECRETION OR GASTRO-INTESTINAL DISORDERS

PATENT AND EXCLUSIVITY TERMS

PATENT USE CODE

U-270	METHOD FOR IMPROVING THE TIME FOR ADMINISTRATION OR THE TIME BETWEEN CHANGES OF GIVING SETS FOR THE DRUG PRODUCT
U-271	METHOD OF TREATING TUMORS
U-272	METHOD OF TREATING CARCINOMA
U-273	CUTANEOUS T-CELL LYMPHOMA
U-274	ZANAMIVIR FOR INHALATION
U-275	METHOD OF USE OF THE DRUG SUBSTANCE
U-276	METHOD OF USE OF LEVOBUPIVACAINE
U-277	NEUROLOGICAL AND OTHER DISORDERS (TREATMENT OF EPILEPSY, BID ORAL DOSING)
U-278	METHOD OF USE OF THE INDICATION OF THE DRUG PRODUCT
U-279	METHOD OF USE OF THE APPROVED PRODUCT
U-280	TREATING PRECIPITATED ACUTE URINARY RETENTION WITH FINASTERIDE
U-281	ANTIMYCOTIC USES, SPECIFICALLY, TREATMENT OF ONYCHOMYCOSIS
U-282	METHOD OF TREATING BACTERIAL INFECTIONS
U-283	METHOD FOR TREATING MENOPAUSAL SYMPTOMS IN A POSTMENOPAUSAL FEMALE
U-284	MENOPAUSAL AND POSTMENOPAUSAL DISORDERS (INCLUDING VASOMOTOR SYMPTOMS ASSOCIATED WITH MENOPAUSE, AND VULVAR AND VAGINAL ATROPHY) AND OSTEOPOROSIS
U-285	DEPRESSION AND SOCIAL ANXIETY DISORDER/SOCIAL PHOBIA
U-286	DEPRESSION
U-287	TREATMENT OR PREVENTION OF OSTEOPOROSIS
U-288	THERAPY OF INFLUENZA

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020482 004	ACARBOSE; PRECOSE	4717720	MAY 31, 2010	U-134	I-252	SEP 29, 2001
020338 001	ADAPALENE; DIFFERIN	RE34440	MAR 31, 2010	U-275	I-253	SEP 29, 2001
020380 001	ADAPALENE; DIFFERIN	4717720	MAY 31, 2010	U-134		
020059 001	ADENOSINE; ADENOSCAN	RE34440	MAY 31, 2010	U-275		
020760 001	ALATROFLOXACIN MESYLATE; TROVAN PRESERVATIVE FREE	5070877	MAY 18, 2009	U-116		
020760 002	ALATROFLOXACIN MESYLATE; TROVAN PRESERVATIVE FREE	5164402	NOV 17, 2009	U-282		
020503 001	ALBUTEROL SULFATE; PROVENTIL-HFA	5763454	JUN 15, 2015	U-282		
019621 001	ALBUTEROL SULFATE; VENTOLIN	5164402	NOV 17, 2009	U-282		
020560 001	ALENDRONATE SODIUM; FOSAMAX	5763454	JUN 15, 2015	U-282		
020560 002	ALENDRONATE SODIUM; FOSAMAX	4594359	JUN 10, 2003			
020560 003	ALENDRONATE SODIUM; FOSAMAX					
020886 001	ALITRETINOIN; PANRETIN					
020221 001	AMIFOSTINE; ETHYOL					
020965 001	AMINOLEVULINIC ACID HYDROCHLORIDE; LEVULAN KERASTICK	5079262	JUL 28, 2009	U-289	I-262	JUN 02, 2002
		5211938	MAY 18, 2010	U-289	I-272	JUN 16, 2002
		5422093	JUL 28, 2009	U-289	I-272	JUN 16, 2002
020364 002	AMLODIPINE BESYLATE; LOTREL	4572909	JUL 31, 2006		ODE	FEB 02, 2006
020364 003	AMLODIPINE BESYLATE; LOTREL	4572909	JUL 31, 2006		NCE	FEB 02, 2004
020364 004	AMLODIPINE BESYLATE; LOTREL	4572909	JUL 31, 2006		ODE	FEB 02, 2006
020508 001	AMMONIUM LACTATE; LAC-HYDRIN				NCE	FEB 02, 2004
021007 001	AMPRENAVIR; AGENERASE				ODE	JUN 24, 2006
021007 002	AMPRENAVIR; AGENERASE				NCE	JUN 24, 2006
021039 001	AMPRENAVIR; AGENERASE				NCE	DEC 03, 2004
020884 001	ASPIRIN; AGGRENOL	5585397	DEC 17, 2013			
020702 001	ATORVASTATIN CALCIUM; LIPITOR	5969156	JUL 08, 2016		PED	FEB 29, 2000
020702 002	ATORVASTATIN CALCIUM; LIPITOR	5969156	JUL 08, 2016		NCE	APR 15, 2004
020702 003	ATORVASTATIN CALCIUM; LIPITOR	5969156	JUL 08, 2016		NCE	APR 15, 2004
020500 001	ATOVAQUONE; MEPRON				NCE	APR 15, 2004
					NC	NOV 22, 2002
020746 001	BUDESONIDE; RHINOCORT					
020746 002	BUDESONIDE; RHINOCORT				ODE	JAN 05, 2006
020711 002	BUPROPION HYDROCHLORIDE; ZYBAN	5763493	AUG 12, 2013		I-255	JAN 05, 2002
020711 003	BUPROPION HYDROCHLORIDE; ZYBAN	5430057	SEP 30, 2013		NDF	OCT 01, 2002
020954 001	BUSULFAN; BUSULFEX	5559148	MAY 24, 2015		NDF	OCT 01, 2002
020793 001	CAFFEINE CITRATE; CAFECIT					
020313 002	CALCITONIN, SALMON; MIALCALCIN	5759565	JUN 02, 2015			
020896 001	CAPECITABINE; XELODA	5472949	DEC 14, 2013			
020896 002	CAPECITABINE; XELODA	4966891	NOV 08, 2008		U-271	SEP 21, 2002
020712 001	CARBAMAZEPINE; CARBATROL	5472949	DEC 14, 2013		U-271	
020712 002	CARBAMAZEPINE; CARBATROL	4966891	NOV 08, 2008		U-272	
		5912013	JUN 15, 2016		U-277	
		5912013	JUN 15, 2016		U-277	

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020998 001	CELECOXIB; CELEBREX	5760068	JUN 02, 2015	U-19		
		5466823	NOV 30, 2013			
020998 002	CELECOXIB; CELEBREX	5563165	NOV 30, 2013	U-19		
		5760068	JUN 02, 2015			
		5466823	NOV 30, 2013			
020740 005	CERIVASTATIN SODIUM; BAYCOL	5563165	NOV 30, 2013			
		5006530	JAN 17, 2009		NS	MAY 24, 2002
		5177080	NOV 26, 2011			
020638 001	CIDOFOVIR; VISTIDE	5142051	JUN 26, 2010			
020863 001	CILOSTAZOL; PLETAL	4277479	AUG 29, 2000		NCE	JAN 15, 2004
020863 002	CILOSTAZOL; PLETAL	4277479	AUG 29, 2000		NCE	JAN 15, 2004
019537 004	CIPROFLOXACIN HYDROCHLORIDE; CIPRO	4670444	DEC 09, 2003	U-36		
		5286754	FEB 15, 2011			
		5648093	JUL 15, 2014			
020767 001	CISAPRIDE MONOHYDRATE; PROPULSID QUICKSOLV					
020463 001	CROMOLYN SODIUM; NASALCROM					
021041 001	CYTARABINE; DEPOCYT					
020287 001	DALTEPARIN SODIUM; FRAGMIN					
020287 003	DALTEPARIN SODIUM; FRAGMIN					
020287 004	DALTEPARIN SODIUM; FRAGMIN					
017922 001	DESMOPRESSIN ACETATE; DDAVP	5763407	JUN 29, 2013			
017922 002	DESMOPRESSIN ACETATE; DDAVP	5763407	JUN 29, 2013			
018938 001	DESMOPRESSIN ACETATE; DDAVP	5763407	JUN 29, 2013			
018938 002	DESMOPRESSIN ACETATE; DDAVP	5763407	JUN 29, 2013			
019955 001	DESMOPRESSIN ACETATE; DDAVP	5763407	JUN 29, 2013			
019955 002	DESMOPRESSIN ACETATE; DDAVP	5763407	JUN 29, 2013			
017922 003	DESMOPRESSIN ACETATE; DDAVP (NEEDS NO REFRIGERATION)	5763407	JUN 29, 2013			
021038 001	DEXMETOMIDINE; PRECEDEX					
074752 001	DILTIAZEM HYDROCHLORIDE; CARTIA XT					
074752 003	DILTIAZEM HYDROCHLORIDE; CARTIA XT					
074752 004	DILTIAZEM HYDROCHLORIDE; CARTIA XT					
020449 001	DOCETAXEL; TAXOTERE					
		4814470	MAY 14, 2010			
		5438072	NOV 22, 2013			
		5698582	JUL 03, 2012			
		5714512	JUL 03, 2012			
020931 001	DOFETILIDE; TIKOSYN					
020931 002	DOFETILIDE; TIKOSYN					
020931 003	DOFETILIDE; TIKOSYN					
020869 001	DORZOLAMIDE HYDROCHLORIDE; COSOPT					
020862 001	DOXERCALCIFEROL; HECTOROL	4797413	APR 28, 2008			
		4619939	OCT 28, 2003			
		5602116	APR 03, 2115	U-278	NCE	JUN 09, 2004
		5707980	FEB 11, 2117	U-278		
050718 001	DOXORUBICIN HYDROCHLORIDE; DOXIL					
020972 001	EFAVIRENZ; SUSTIVA					
		5811423	AUG 07, 2012	U-256	ODE	JUN 28, 2006
		5519021	MAY 21, 2013			
		5663169	SEP 02, 2014			

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020972 002	EFAVIRENZ; SUSTIVA	5811423 5519021 5663169 5519021 5663169 5811423	AUG 07, 2012 MAY 21, 2013 SEP 02, 2014 MAY 21, 2013 SEP 02, 2014 AUG 07, 2012	U-256 U-257 U-257 U-256		
020972 003	EFAVIRENZ; SUSTIVA	5223261	JUN 29, 2010			
020796 001	ENTACAPONE; COMTAN				NCE	OCT 19, 2004
050778 001	EPIDUBICIN HYDROCHLORIDE; ELLENCE				ODE	SEP 15, 2006
020375 001	ESTRADIOL; CLIMARA				I-254	MAR 05, 2002
020375 002	ESTRADIOL; CLIMARA				I-254	MAR 05, 2002
020375 003	ESTRADIOL; CLIMARA				I-254	MAR 05, 2002
020375 004	ESTRADIOL; CLIMARA				I-254	MAR 05, 2002
020908 001	ESTRADIOL; VAGIFEM				NP	MAR 26, 2002
020992 002	ESTROGENS, CONJUGATED SYNTHETIC A; CENESTIN				NP	MAR 24, 2002
020992 003	ESTROGENS, CONJUGATED SYNTHETIC A; CENESTIN				NP	MAR 24, 2002
010971 005	ESTROGENS, CONJUGATED; PMB 200					
010971 003	ESTROGENS, CONJUGATED; PMB 400					
004782 001	ESTROGENS, CONJUGATED; PREMARIN					
004782 002	ESTROGENS, CONJUGATED; PREMARIN					
004782 003	ESTROGENS, CONJUGATED; PREMARIN					
004782 004	ESTROGENS, CONJUGATED; PREMARIN					
004782 005	ESTROGENS, CONJUGATED; PREMARIN					
010402 001	ESTROGENS, CONJUGATED; PREMARIN					
020216 001	ESTROGENS, CONJUGATED; PREMPHASE 14/14					
020303 002	ESTROGENS, CONJUGATED; PREMPHASE 14/14					
020527 002	ESTROGENS, CONJUGATED; PREMPRO 14/14					
020303 001	ESTROGENS, CONJUGATED; PREMPRO 14/14					
020527 001	ESTROGENS, CONJUGATED; PREMPRO 14/14					
020527 003	ESTROGENS, CONJUGATED; PREMPRO 14/14					
021065 001	ETHINYL ESTRADIOL; FEMHRT					
021065 002	ETHINYL ESTRADIOL; FEMHRT					
021065 003	ETHINYL ESTRADIOL; FEMHRT					
020584 001	ETODOLAC; LODINE XL					
020584 002	ETODOLAC; LODINE XL					
020584 003	ETODOLAC; LODINE XL					
020753 001	EXEMESTANE; AROMASIN					
020363 001	FAMCICLOVIR; FAMVIR					
020363 003	FAMCICLOVIR; FAMVIR					
020325 001	FAMOTIDINE; PEPCID AC					
020801 001	FAMOTIDINE; PEPCID AC					

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
019304 002	FENOFIBRATE;TRICOR (MICRONIZED)	4895726	JAN 19, 2009			
019304 003	FENOFIBRATE;TRICOR (MICRONIZED)	4895726	JAN 19, 2009			
019304 004	FENOFIBRATE;TRICOR (MICRONIZED)	4895726	JAN 19, 2009			
020747 001	FENTANYL CITRATE;ACTIQ				NP	NOV 04, 2001
020747 002	FENTANYL CITRATE;ACTIQ				NP	NOV 04, 2001
020747 003	FENTANYL CITRATE;ACTIQ				NP	NOV 04, 2001
020747 004	FENTANYL CITRATE;ACTIQ				NP	NOV 04, 2001
020747 005	FENTANYL CITRATE;ACTIQ				NP	NOV 04, 2001
020747 006	FENTANYL CITRATE;ACTIQ				NP	NOV 04, 2001
020955 001	FERRIC SODIUM GLUCONATE;FERRLECIT				NCE	FEB 18, 2004
020625 001	FEXOFENADINE HYDROCHLORIDE;ALLEGRA	5855912	FEB 28, 2015			
		5738872	FEB 28, 2015			
020788 001	FINASTERIDE; PROPECIA	5571817	NOV 05, 2013	U-259		
		5886184	NOV 19, 2012			
020180 001	FINASTERIDE; PROSCAR	4760071	JUN 19, 2006	U-262		
		5886184	NOV 19, 2012			
		4377584	MAR 22, 2000	U-261		
		5942519	OCT 23, 2018	U-280		
019452 001	FLUCINOLONE ACETONIDE; DERMA-SMOOTHIE/FS				I-265	AUG 18, 2002
018936 003	FLUOXETINE HYDROCHLORIDE;PROZAC	4314081	FEB 02, 2001	U-154		
		4626549	DEC 02, 2003	U-84		
		4626549	DEC 02, 2003			
018936 004	FLUOXETINE HYDROCHLORIDE;PROZAC	4314081	FEB 02, 2001	U-154		
		4626549	DEC 02, 2003	U-84		
		4626549	DEC 02, 2003	U-84		
020101 001	FLUOXETINE HYDROCHLORIDE;PROZAC	4626549	DEC 02, 2003	U-84		
020974 001	FLUOXETINE HYDROCHLORIDE;PROZAC	4626549	DEC 02, 2003	U-154		
		4314081	FEB 02, 2001			
020974 002	FLUOXETINE HYDROCHLORIDE;PROZAC	4626549	DEC 02, 2003	U-84		
		4626549	DEC 02, 2003	U-154		
		4314081	FEB 02, 2001			
019958 001	FLUTICASON PROPIONATE;CUTIVATE					
020121 001	FLUTICASON PROPIONATE;FLONASE	5767251	JUN 16, 2015			
020582 001	FOLLITROPIN ALFA/BETA; FOLLISTIM	5767251	JUN 16, 2015			
020582 002	FOLLITROPIN ALFA/BETA; FOLLISTIM	4087544	JAN 16, 2000	U-106		
020882 001	GABAPENTIN;NEURONTIN	5084479	JAN 02, 2010	U-258		
		4087544	JAN 16, 2000	U-106		
020882 002	GABAPENTIN;NEURONTIN	5084479	JAN 02, 2010	U-258		
					NCE	DEC 08, 2004
					NCE	DEC 08, 2004
					NCE	DEC 08, 2004
					NCE	JUL 29, 2004
020937 001	GADOVERSETAMIDE;OPTIMARK					
020975 001	GADOVERSETAMIDE;OPTIMARK					
020976 001	GADOVERSETAMIDE;OPTIMARK					
021057 001	GANIRELIX ACETATE;ANTAGON	4801577	FEB 05, 2007			
		5767082	JUN 16, 2015			
021061 001	GATIFLOXACIN;TEQUIN	4980470	DEC 25, 2007			
021061 002	GATIFLOXACIN;TEQUIN	4980470	DEC 25, 2007			
021062 001	GATIFLOXACIN;TEQUIN	4980470	DEC 25, 2007			
					NCE	DEC 17, 2004
					NCE	DEC 17, 2004
					NCE	DEC 17, 2004

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APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
>ADD>	CATIFLOXACIN; TEQUIN	4980470	DEC 25, 2007		NCE	DEC 17, 2004
021062 002	GEMCITABINE HYDROCHLORIDE; GEMZAR	4808614	MAY 15, 2010			
020509 001	GEMCITABINE HYDROCHLORIDE; GEMZAR	4808614	MAY 15, 2010			
020509 002	GEMCITABINE HYDROCHLORIDE; GEMZAR	4379785	APR 06, 2005	U-118		
020496 001	GLIMEPIRIDE; AMARYL	4379785	APR 06, 2005	U-118		
020496 002	GLIMEPIRIDE; AMARYL	4379785	APR 06, 2005	U-118		
020496 003	GLIMEPIRIDE; AMARYL	4379785	APR 06, 2005			
020305 001	GRANISETRON HYDROCHLORIDE; KYTRIL	5994348	JUN 07, 2015		I-264	JUL 27, 2002
020305 002	GRANISETRON HYDROCHLORIDE; KYTRIL	5994348	JUN 07, 2015		I-264	JUL 27, 2002
020758 001	HYDROCHLOROTHIAZIDE; AVALIDE	5994348	JUN 07, 2015		NCE	SEP 30, 2002
020758 002	HYDROCHLOROTHIAZIDE; AVALIDE	5994348	JUN 07, 2015			
020758 003	HYDROCHLOROTHIAZIDE; AVALIDE	5270317	MAR 20, 2011		D-50	MAY 13, 2002
020491 001	IBUTILIDE FUMARATE; CORVERT	4689338	AUG 25, 2009	U-172		
020723 001	IMIQUIMOD; ALDARA	4791111	DEC 23, 2005		NP	JUL 19, 2002
021028 001	INSULIN HUMAN; VELOSULIN BR	4791111	DEC 23, 2005			
020083 001	ITRACONAZOLE; SPORANOX	4791111	DEC 23, 2005			
020657 001	ITRACONAZOLE; SPORANOX	4791111	DEC 23, 2005			
020966 001	ITRACONAZOLE; SPORANOX	4267179	JUN 23, 2000		NCE	JUL 02, 2004
021066 001	KETOTIFEN FUMARATE; ZADITOR	5905082	MAY 18, 2016			
020564 001	LAMIVUDINE; EPIVIR	5047407	FEB 08, 2009		I-257	DEC 08, 2001
021003 001	LAMIVUDINE; EPIVIR-HBV	5532246	JUL 02, 2013	U-250		
021004 001	LAMIVUDINE; EPIVIR-HBV	5905082	MAY 18, 2016			
020764 001	LAMOTRIGINE; LAMICTAL CD	5047407	FEB 08, 2009			
020764 002	LAMOTRIGINE; LAMICTAL CD	5532246	JUL 02, 2013	U-250		
020764 003	LAMOTRIGINE; LAMICTAL CD	5905082	MAY 18, 2016			
020406 001	LANSOPRAZOLE; PREVACID	5047407	FEB 08, 2009		I-257	DEC 08, 2001
020406 002	LANSOPRAZOLE; PREVACID	5532246	JUL 02, 2013	U-250		
020597 001	LATANOPROST; XALATAN	5296504	MAR 22, 2011			
020517 002	LEUPROLIDE ACETATE; LUPRON DEPOT-4	5422368	MAR 22, 2011		U-260	
020837 001	LEVALBUTEROL HYDROCHLORIDE; XOPENEX	4593353	JUL 28, 2006		U-260	
020837 002	LEVALBUTEROL HYDROCHLORIDE; XOPENEX	4652441	NOV 01, 2004			
020035 001	LEVAMISOLE HYDROCHLORIDE; ERGAMISOL	4584305	JUN 18, 2004			
021035 001	LEVETIRACETAM; KEPPRA	4943639	JUN 06, 2006	U-42		
021035 002	LEVETIRACETAM; KEPPRA	4837223	JUN 06, 2006		NCE	NOV 30, 2004
021035 003	LEVETIRACETAM; KEPPRA	4943639	JUN 06, 2006		NCE	NOV 30, 2004
020997 001	LEVOBUPIVACAINE HYDROCHLORIDE; CHIROCAINE	4837223	JUN 06, 2006		NCE	NOV 30, 2004
020997 002	LEVOBUPIVACAINE HYDROCHLORIDE; CHIROCAINE	4943639	JUN 06, 2006			
020997 003	LEVOBUPIVACAINE HYDROCHLORIDE; CHIROCAINE	4837223	JUN 06, 2006			
021045 001	LEVONORGESTREL; PLAN B	4943639	JUN 06, 2006			
019941 001	LIDOCAINE; EMLA	4837223	JUN 06, 2006			
>ADD>	LEVOBUPIVACAINE HYDROCHLORIDE; CHIROCAINE	5708011	OCT 13, 2014	U-276		
>ADD>	LEVOBUPIVACAINE HYDROCHLORIDE; CHIROCAINE	5708011	OCT 13, 2014	U-276		
>ADD>	LEVONORGESTREL; PLAN B	5708011	OCT 13, 2014	U-276		
>ADD>	LIDOCAINE; EMLA	5708011	OCT 13, 2014		NP	JUL 28, 2002
>ADD>					I-260	MAR 11, 2002

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APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020612 001	LIDOCAINE; LIDODERM	4374829	DEC 30, 2001		ODE	MAR 19, 2006
019777 006	LISINAPRIL; ZESTRIL	4231938	JUN 15, 2001		I-250	MAR 11, 2002
019643 002	LOVASTATIN; MEVACOR	4231938	JUN 15, 2001		I-250	MAR 11, 2002
019643 003	LOVASTATIN; MEVACOR	4231938	JUN 15, 2001		I-250	MAR 11, 2002
019643 004	LOVASTATIN; MEVACOR	4231938	JUN 15, 2001		I-250	MAR 11, 2002
021047 001	LUTEINIZING HORMONE; REPRONEX				NP	AUG 27, 2002
021047 002	LUTEINIZING HORMONE; REPRONEX				NP	AUG 27, 2002
020922 001	MEQUINOL; SOLAGE	5194247	MAR 16, 2010	U-294		
		5470567	MAR 19, 2010	U-294		
020357 003	METFORMIN HYDROCHLORIDE; GLUCOPHAGE				NCE	MAR 03, 2000
020357 004	METFORMIN HYDROCHLORIDE; GLUCOPHAGE				NCE	MAR 03, 2000
020357 005	METFORMIN HYDROCHLORIDE; GLUCOPHAGE				NCE	MAR 03, 2000
020969 001	METHOXSALEN; UVADEX				NP	FEB 25, 2002
020968 001	MICONAZOLE NITRATE; MONISTAT DUAL- PAK	4845075	APR 11, 2009	U-273		
020682 001	MIGLITOL; GLYSET	5036102	JUL 30, 2008			
020682 002	MIGLITOL; GLYSET	4999375	MAR 12, 2008			
020682 003	MIGLITOL; GLYSET	5514698	MAR 21, 2014		NP	JUN 30, 2002
019436 001	MILIRINONE LACTATE; PRIMACOR	4639436	JAN 27, 2009	U-111		
020343 001	MILIRINONE LACTATE; PRIMACOR	4639436	JAN 27, 2009	U-111		
020343 002	MILIRINONE LACTATE; PRIMACOR	4639436	JAN 27, 2009	U-111		
020343 003	MILIRINONE LACTATE; PRIMACOR	4639436	JAN 27, 2009	U-111		
020717 001	MODAFINIL; PROVIGIL	4313951	NOV 26, 2001			
		4313951	NOV 26, 2001			
		4313951	NOV 26, 2001			
		4313951	NOV 26, 2001			
		4177290	NOV 26, 2001			
		5618845	MAR 09, 1999			
020717 002	MODAFINIL; PROVIGIL	4927855	OCT 06, 2014	U-255		
		4927855	MAY 22, 2007	U-255		
		4177290	MAR 09, 1999			
		5618845	OCT 06, 2014	U-255		
		4927855	MAY 22, 2007	U-255		
021085 001	MOXIFLOXACIN HYDROCHLORIDE; AVELOX	4474787	OCT 02, 2006		NCE	DEC 10, 2004
021009 001	NEDOCROMIL SODIUM; ABREVIA	4760072	JUL 26, 2005		NP	DEC 08, 2002
		5366972	NOV 22, 2011			
020933 001	NEVIRAPINE; VIRAMUNE					
018612 003	NICOTINE POLACRILEX; NICORETTE (MINT)	5656255	AUG 12, 2014			
020666 003	NICOTINE POLACRILEX; NICORETTE (MINT)	5400808	JUN 08, 2010			
020385 001	NICOTINE; NICOTROL	4917120	APR 17, 2007			
020714 001	NICOTINE; NICOTROL	4800903	JAN 31, 2006		NP*	DEC 23, 2001
		4793366	DEC 27, 2005		NP*	DEC 23, 2001
		5167242	JUN 08, 2010			
		5501236	JUN 08, 2010			
		5186925	FEB 16, 2010			
018705 002	NITROGLYCERIN; NITROLINGUAL PUMPSPRAY	5688530	NOV 18, 2014	U-268	ODE	NOV 25, 2005
021008 001	OCITREOTIDE ACETATE; SANDOSTATIN LAR	4395403	NOV 21, 2002	U-269		
		5538739	JUL 23, 2013			
		5639480	JUN 17, 2014			
		5922338	JUL 13, 2016			
		5922682	JUL 13, 2016			

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021008 002	OCTREOTIDE ACETATE; SANDOSTATIN LAR	5688530 4395403 5538739 5639480 5922338 5922682 5688530 4395403 5538739 5639480 5922338 5922682	NOV 18, 2014 NOV 21, 2002 JUL 23, 2013 JUN 17, 2014 JUL 13, 2016 JUL 13, 2016 NOV 18, 2014 NOV 21, 2002 JUL 23, 2013 JUN 17, 2014 JUL 13, 2016 JUL 13, 2016	U-268 ODE U-269		NOV 25, 2005
021008 003	OCTREOTIDE ACETATE; SANDOSTATIN LAR	5688530 4395403 5538739 5639480 5922338 5922682	NOV 18, 2014 NOV 21, 2002 JUL 23, 2013 JUN 17, 2014 JUL 13, 2016 JUL 13, 2016	U-268 ODE U-269		NOV 25, 2005
020103 001	ONDANSETRON HYDROCHLORIDE; ZOFTRAN	4598089	JUN 18, 2004		I-269	AUG 27, 2002
020103 002	ONDANSETRON HYDROCHLORIDE; ZOFTRAN	5763483	DEC 27, 2016		I-269	AUG 27, 2002
020766 001	ORLISTAT; XENICAL	5866601	FEB 02, 2016		NCE	APR 23, 2004
021087 001	OSELTAMIVIR PHOSPHATE; TAMIFLU	5952375	FEB 02, 2016		NCE	OCT 27, 2004
>ADD> >ADD>	OXAPROZIN; DAYPRO				PED	APR 29, 2000
020897 001	OXYBUTYNYN CHLORIDE; DITROPAN XL	4783337 5674895 5082668 5840754 4612008 4519801 5912268 5840754 5674895 5082668 4783337 4612008 4519801	SEP 16, 2003 MAY 22, 2015 SEP 16, 2003 MAY 22, 2015 SEP 16, 2003 JUL 12, 2002 MAY 22, 2015 MAY 22, 2015 MAY 22, 2015 SEP 16, 2003 SEP 16, 2003 SEP 16, 2003 JUL 12, 2002		NCE NP	OCT 29, 1999 DEC 16, 2001
020897 002	OXYBUTYNYN CHLORIDE; DITROPAN XL	5912268 5912268 4612008 4519801 5912268 5840754 5674895 5082668 4783337 4612008 4519801	MAY 22, 2015 MAY 22, 2015 SEP 16, 2003 JUL 12, 2002 MAY 22, 2015 MAY 22, 2015 MAY 22, 2015 SEP 16, 2003 SEP 16, 2003 SEP 16, 2003 JUL 12, 2002		NP	DEC 16, 2001
020897 003	OXYBUTYNYN CHLORIDE; DITROPAN XL	5912268 5912268 4612008 4519801 5912268 5840754 5674895 5082668 4783337 4612008 4519801	MAY 22, 2015 MAY 22, 2015 SEP 16, 2003 JUL 12, 2002 MAY 22, 2015 MAY 22, 2015 MAY 22, 2015 SEP 16, 2003 SEP 16, 2003 SEP 16, 2003 JUL 12, 2002		NP	DEC 16, 2001
020553 001	OXYCODONE HYDROCHLORIDE; OXYCONTIN	5508042 5656295 5508042 5656295 5508042 5656295 5508042 5656295 5508042 5656295	APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008			
020553 002	OXYCODONE HYDROCHLORIDE; OXYCONTIN	5508042 5656295 5508042 5656295 5508042 5656295 5508042 5656295 5508042 5656295	APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008			
020553 003	OXYCODONE HYDROCHLORIDE; OXYCONTIN	5508042 5656295 5508042 5656295 5508042 5656295 5508042 5656295 5508042 5656295	APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008			
020553 004	OXYCODONE HYDROCHLORIDE; OXYCONTIN	5508042 5656295 5508042 5656295 5508042 5656295 5508042 5656295 5508042 5656295	APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008 APR 16, 2013 FEB 05, 2008			

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PATENT AND EXCLUSIVITY DATA
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020262 001	PACLITAXEL; TAXOL	5872132	MAY 19, 2015		I-270	OCT 25, 2002
020031 001	PAROXETINE HYDROCHLORIDE; PAXIL	5900423	MAY 19, 2015		I-261	MAY 17, 2002
020031 002	PAROXETINE HYDROCHLORIDE; PAXIL	5789449	JAN 06, 2009	U-285		
020031 003	PAROXETINE HYDROCHLORIDE; PAXIL	5872132	MAY 19, 2015		I-261	MAY 17, 2002
020031 004	PAROXETINE HYDROCHLORIDE; PAXIL	5900423	MAY 19, 2015			
020031 005	PAROXETINE HYDROCHLORIDE; PAXIL	5789449	JAN 06, 2009	U-285		
020710 001	PAROXETINE HYDROCHLORIDE; PAXIL	5872132	MAY 19, 2015		I-261	MAY 17, 2002
020885 001	PAROXETINE HYDROCHLORIDE; PAXIL	5900423	MAY 19, 2015			
020885 002	PAROXETINE HYDROCHLORIDE; PAXIL	5789449	JAN 06, 2009	U-285		
020885 003	PAROXETINE HYDROCHLORIDE; PAXIL	5872132	MAY 19, 2015		I-261	MAY 17, 2002
020885 004	PAROXETINE HYDROCHLORIDE; PAXIL	5900423	MAY 19, 2015			
020936 001	PAROXETINE HYDROCHLORIDE; PAXIL CR	5789449	JAN 06, 2009	U-285		
020936 002	PAROXETINE HYDROCHLORIDE; PAXIL CR	5872132	MAY 19, 2015		I-261	MAY 17, 2002
021079 001	PEMITROLAST POTASSIUM; ALAMAST	5034230	DEC 23, 2008	U-184	PED	MAR 24, 2005
		5034230	JUN 23, 2009	U-184	NCE	SEP 24, 2004

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APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
021073 001	PIOGLITAZONE HYDROCHLORIDE; ACTOS	4444779	JUL 27, 1999		NCE	JUL 15, 2004
021073 002	PIOGLITAZONE HYDROCHLORIDE; ACTOS	4687777	JAN 17, 2006			
021073 003	PIOGLITAZONE HYDROCHLORIDE; ACTOS	4444779	JUL 27, 1999		NCE	JUL 15, 2004
020698 001	POLYETHYLENE GLYCOL 3350; MIRALAX	4687777	JAN 17, 2006		NCE	JUL 15, 2004
020744 001	PORACTANT ALPHA; CUROSURF	5710183	JUL 14, 2015	U-265	NP	FEB 18, 2002
020667 001	PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4886812	MAR 25, 2011		NCE	NOV 18, 2004
020667 002	PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4886812	MAR 25, 2011			
020667 003	PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4886812	MAR 25, 2011			
020667 004	PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4886812	MAR 25, 2011			
020667 005	PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4886812	MAR 25, 2011			
020667 006	PRAMIPEXOLE DIHYDROCHLORIDE; MIRAPEX	4886812	MAR 25, 2011			
019627 002	PROPOFOL; DIPRIVAN	5714520	MAR 22, 2015			
075102 001	PROPOFOL; PROPOFOL	5714520*	PED SEP 22, 2015	U-217		
020639 004	QUETIAPINE FUMARATE; SEROQUEL	5731355	MAR 22, 2015	U-217		
020973 002	RABEPRAZOLE SODIUM; ACIPHEX	5731355*	PED SEP 22, 2015	U-217		
020815 001	RALOXIFENE HYDROCHLORIDE; EVISTA	5731356	MAR 22, 2015	U-218		
		5731356*	PED SEP 22, 2015	U-218		
		5908869	MAR 22, 2015	U-270		
		5908869*	PED SEP 22, 2015	U-270		
075102 001	PROPOFOL; PROPOFOL	4879288	MAR 20, 2007		PC	OCT 17, 1999
020639 004	QUETIAPINE FUMARATE; SEROQUEL				NCE	SEP 26, 2002
020973 002	RABEPRAZOLE SODIUM; ACIPHEX				NCE	AUG 19, 2004
020815 001	RALOXIFENE HYDROCHLORIDE; EVISTA				I-271	SEP 30, 2002
075094 001	RANITIDINE HYDROCHLORIDE; RANITIDINE HCL	5466810	JUN 10, 2014			
020984 001	RAPACURONIUM BROMIDE; RAPLON	5514826	JUN 07, 2015			
020984 002	RAPACURONIUM BROMIDE; RAPLON	5569772	JUN 07, 2015			
020903 001	RIBAVIRIN; REBETOL	5629425	SEP 19, 2014			
020272 007	RISPERIDONE; RISPERDAL	5659087	JUN 07, 2015			
		5710285	MAR 31, 2015			
		5731327	MAR 24, 2015			
		5731342	JAN 27, 2017			
		5747510	MAR 02, 2014			
		5808061	MAR 31, 2014			
		5811120	MAR 02, 2014			
		5843984	APR 14, 2017			
		5972383	MAR 02, 2014	U-287		
020272 008	RISPERIDONE; RISPERDAL				PC	JAN 14, 2000
020680 001	RITONAVIR; NORVIR	5418226	APR 14, 2013		NCE	AUG 18, 2004
020865 001	RIZATRIPTAN BENZOATE; MAXALT-MLT	5418226	APR 14, 2013		NCE	AUG 18, 2004
020865 002	RIZATRIPTAN BENZOATE; MAXALT-MLT	5914128	DEC 22, 2017			
		5158952	OCT 27, 2009	U-90	D-37	OCT 17, 2000
		4804663	DEC 29, 2007	U-90		
		4804663	DEC 29, 2007	U-90		
		5158952	DEC 29, 2009	U-90	D-37	OCT 17, 2000
		5948436	SEP 13, 2013			
		5457895	OCT 01, 2013			
		5457895	OCT 01, 2013			

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021042 001	ROFECOXIB; VIOXX	5474995	JUN 24, 2013	U-266	NCE	MAY 20, 2004
021042 002	ROFECOXIB; VIOXX	5691374	NOV 25, 2017			
021052 001	ROFECOXIB; VIOXX	5474995	JUN 24, 2013	U-266	NCE	MAY 20, 2004
021052 002	ROFECOXIB; VIOXX	5691374	NOV 25, 2017			
021071 002	ROSIGLITAZONE MALEATE; AVANDIA	5474995	JUN 24, 2013	U-266	NCE	MAY 20, 2004
021071 003	ROSIGLITAZONE MALEATE; AVANDIA	5691374	NOV 25, 2017			
021071 004	ROSIGLITAZONE MALEATE; AVANDIA	5474995	JUN 24, 2013	U-266	NCE	MAY 20, 2004
019839 001	SERTRALINE HYDROCHLORIDE; ZOLOFT	5474995	JUN 24, 2013	U-266	NCE	MAY 20, 2004
019839 002	SERTRALINE HYDROCHLORIDE; ZOLOFT	5691374	NOV 25, 2017			
019839 003	SERTRALINE HYDROCHLORIDE; ZOLOFT	5474995	JUN 24, 2013	U-266	NCE	MAY 20, 2004
019839 004	SERTRALINE HYDROCHLORIDE; ZOLOFT	5691374	NOV 25, 2017			
019839 005	SERTRALINE HYDROCHLORIDE; ZOLOFT	5474995	JUN 24, 2013	U-266	NCE	MAY 20, 2004
020478 001	SEVOFLURANE; ULTANE	5474995	JUN 24, 2013	U-266	NCE	MAY 20, 2004
019766 001	SIMVASTATIN; ZOCOR	5691374	NOV 25, 2017			
		4536518	DEC 30, 2005	U-152	NCE	MAY 25, 2004
		4536518	DEC 30, 2005	U-152	NCE	MAY 25, 2004
		4536518	DEC 30, 2005	U-152	NCE	MAY 25, 2004
		4536518	DEC 30, 2005	U-152	NCE	MAY 25, 2004
		5990176	JAN 27, 2017			
019766 002	SIMVASTATIN; ZOCOR					
		5100899	JUN 06, 2009	U-290	I-277	NOV 22, 2002
		5212155	MAY 18, 2010	U-291	I-278	NOV 22, 2002
		5308847	MAY 03, 2011	U-292	I-273	AUG 05, 2002
		5403833	APR 04, 2012	U-293	I-277	NOV 22, 2002
		5536729	SEP 30, 2013		I-278	NOV 22, 2002
019766 003	SIMVASTATIN; ZOCOR				I-273	AUG 05, 2002
		5100899	JUN 06, 2009	U-290	I-278	NOV 22, 2002
		5212155	MAY 18, 2010	U-291	I-273	AUG 05, 2002
		5308847	MAY 03, 2011	U-292	I-277	NOV 22, 2002
		5403833	APR 04, 2012	U-293	I-278	NOV 22, 2002
		5536729	SEP 30, 2013		I-273	AUG 05, 2002
019766 004	SIMVASTATIN; ZOCOR				I-278	NOV 22, 2002
		5100899	JUN 06, 2009	U-290	I-273	AUG 05, 2002
		5212155	MAY 18, 2010	U-291	I-277	NOV 22, 2002
		5308847	MAY 03, 2011	U-292	I-278	NOV 22, 2002
		5403833	APR 04, 2012	U-293	I-273	AUG 05, 2002
		5536729	SEP 30, 2013		I-277	NOV 22, 2002
019766 005	SIMVASTATIN; ZOCOR				I-278	NOV 22, 2002
		5100899	JUN 06, 2009	U-290	I-273	AUG 05, 2002
		5212155	MAY 18, 2010	U-291	I-278	NOV 22, 2002
		5308847	MAY 03, 2011	U-292	I-273	AUG 05, 2002
		5403833	APR 04, 2012	U-293	I-277	NOV 22, 2002
		5536729	SEP 30, 2013		I-278	NOV 22, 2002
021083 001	SIROLIMUS; RAPAMUNE				I-273	AUG 05, 2002
		5100899	JUN 06, 2009	U-290	NCE	SEP 15, 2004
		5212155	MAY 18, 2010	U-291		
		5308847	MAY 03, 2011	U-292		
		5403833	APR 04, 2012	U-293		
		5536729	SEP 30, 2013			
019640 005	SOMATROPIN, BIOSYNTHETIC; HUMATROPE					
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
019640 006	SOMATROPIN, BIOSYNTHETIC; HUMATROPE					
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
019640 007	SOMATROPIN, BIOSYNTHETIC; HUMATROPE					
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
		4978655	JUN 24, 2008	U-94	I-182	DEC 30, 1999
		4978655	JUN 24, 2008	U-94	ODE	DEC 30, 2003
020412 001	STAVUDINE; ZERIT					
020412 002	STAVUDINE; ZERIT					
020412 003	STAVUDINE; ZERIT					
020412 004	STAVUDINE; ZERIT					
020412 005	STAVUDINE; ZERIT					
020413 001	STAVUDINE; ZERIT					

PATENT AND EXCLUSIVITY DATA

*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
>ADD>						
020579 001	TAMSULOSIN HYDROCHLORIDE; FLOMAX	4703063	OCT 27, 2009		NCE	AUG 03, 2004
021012 001	TECHNETIUM TC-99M DEPREOTIDE; NEO TECT KIT	5260291	NOV 09, 2010		NCE	AUG 11, 2004
021029 001	TEMOZOLOMIDE; TEMODAR	5260291	NOV 09, 2010		NCE	AUG 11, 2004
021029 002	TEMOZOLOMIDE; TEMODAR	5260291	NOV 09, 2010		NCE	AUG 11, 2004
021029 003	TEMOZOLOMIDE; TEMODAR	5260291	NOV 09, 2010		NCE	AUG 11, 2004
021029 004	TEMOZOLOMIDE; TEMODAR	5260291	NOV 09, 2010		NCE	AUG 11, 2004
074823 001	TERAZOSIN HYDROCHLORIDE; TERAZOSIN HCL	4680291	JUL 14, 2004	U-281	PC	FEB 09, 2000
074823 002	TERAZOSIN HYDROCHLORIDE; TERAZOSIN HCL	4680291	JUL 14, 2004	U-73	NCE	DEC 30, 1999
074823 003	TERAZOSIN HYDROCHLORIDE; TERAZOSIN HCL	4755534	DEC 30, 2006	U-73	NCE	DEC 30, 1999
074823 004	TERAZOSIN HYDROCHLORIDE; TERAZOSIN HCL				NP	APR 29, 2001
020539 001	TERBINAFINE HYDROCHLORIDE; LAMISIL				NCE	SEP 30, 2002
020749 001	TERBINAFINE HYDROCHLORIDE; LAMISIL					
020980 001	TERBINAFINE HYDROCHLORIDE; LAMISIL					
020846 001	TERBINAFINE; LAMISIL					
019762 001	TESTOSTERONE; TESTODERM	5840327	AUG 15, 2016			
019762 002	TESTOSTERONE; TESTODERM	5840327	AUG 15, 2016			
020646 005	TIAGABINE HYDROCHLORIDE; GABITRIL	5010090	OCT 07, 2008			
		5354760	MAR 24, 2012			
		4876248	JAN 30, 2010			
020707 001	TILUDRONATE DISODIUM; SKELID	5292756	MAY 14, 2012	U-230		
020912 001	TIROFIBAN HYDROCHLORIDE; AGGRASTAT	5880136	SEP 27, 2010	U-254		
		5965581	OCT 23, 2016			
		5972967	OCT 23, 2016			
020913 001	TIROFIBAN HYDROCHLORIDE; AGGRASTAT	5972967	OCT 23, 2016			
020505 001	TOPIRAMATE; TOPAMAX				I-266	JUL 23, 2002
020505 002	TOPIRAMATE; TOPAMAX				I-274	OCT 01, 2002
020505 003	TOPIRAMATE; TOPAMAX				I-274	OCT 01, 2002
020505 004	TOPIRAMATE; TOPAMAX				I-266	JUL 23, 2002
020505 005	TOPIRAMATE; TOPAMAX				I-274	OCT 01, 2002
020505 006	TOPIRAMATE; TOPAMAX				I-266	JUL 23, 2002
020844 001	TOPIRAMATE; TOPAMAX SPRINKLE				I-274	OCT 01, 2002
020844 002	TOPIRAMATE; TOPAMAX SPRINKLE				I-266	JUL 23, 2002

PRESCRIPTION AND OTC DRUG PRODUCT
PATENT AND EXCLUSIVITY DATA
*PED and PED represent Pediatric Exclusivity

APPL/PROD NUMBER	INGREDIENT NAME; TRADE NAME	PATENT NUMBER	PATENT/PED EXCL EXPIRES	USE CODE	EXCLUS CODE	EXCLUS EXPIRES
020844 003	TOPIRAMATE; TOPAMAX SPRINKLE					
020671 001	TOPOTECAN HYDROCHLORIDE; HYCAMTIN					
020137 002	TORSEMIDE; DEMADEX	RE34672	AUG 11, 2006		I-274	OCT 01, 2002
020468 001	TRIAMCINOLONE ACETONIDE; NASACORT AQ	5976573	JUL 03, 2016	U-295	I-266	JUL 23, 2002
020719 001	TROGLITAZONE; PRELAY				I-256	NOV 30, 2001
020719 002	TROGLITAZONE; PRELAY					
020719 003	TROGLITAZONE; PRELAY				I-275	JUN 16, 2002
020720 001	TROGLITAZONE; REZULIN				I-275	JUN 16, 2002
020720 002	TROGLITAZONE; REZULIN				I-276	JUN 16, 2002
020720 003	TROGLITAZONE; REZULIN				I-276	JUN 16, 2002
020759 001	TROVAFLOXACIN MESYLATE; TROVAN	5164402	NOV 17, 2009	U-282	I-276	JUN 16, 2002
		5763454	JUN 15, 2015	U-282		
		5164402	NOV 17, 2009	U-282		
020759 002	TROVAFLOXACIN MESYLATE; TROVAN	5763454	JUN 15, 2015	U-282		
020900 001	UREA, C-13; PYLORI-CHEK BREATH TEST				NCE	SEP 17, 2001
020699 001	VENLAFAXINE HYDROCHLORIDE; EFFEXOR XR				I-251	MAR 11, 2002
020699 002	VENLAFAXINE HYDROCHLORIDE; EFFEXOR XR				I-251	MAR 11, 2002
020699 003	VENLAFAXINE HYDROCHLORIDE; EFFEXOR XR				I-251	MAR 11, 2002
020699 004	VENLAFAXINE HYDROCHLORIDE; EFFEXOR XR				I-251	MAR 11, 2002
019614 004	VERAPAMIL HYDROCHLORIDE; VERELAN	4863742	JUN 19, 2007			
020943 001	VERAPAMIL HYDROCHLORIDE; VERELAN PM	4863742	JUN 19, 2007			
020943 002	VERAPAMIL HYDROCHLORIDE; VERELAN PM	4863742	JUN 19, 2007			
020943 003	VERAPAMIL HYDROCHLORIDE; VERELAN PM	4859692	SEP 26, 2010		I-268	SEP 17, 2002
020547 001	ZAFIRLUKAST; ACCOLATE	4626538	JUN 23, 2003		NCE	AUG 13, 2004
020859 001	ZALEPLON; SONATA	4626538	JUN 23, 2003		NCE	AUG 13, 2004
020859 002	ZALEPLON; SONATA	D379506	MAY 27, 2011		NCE	JUL 26, 2004
021036 001	ZANAMIVIR; RELENZA	4778054	OCT 18, 2005			
		4811731	JUL 29, 2006			
		5035237	JUL 30, 2008			
		5360817	NOV 01, 2011			
		5648379	JUL 15, 2014	U-274		
		4627432	DEC 09, 2003			

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