

APPROVED DRUG PRODUCTS

With Therapeutic Equivalence Evaluations



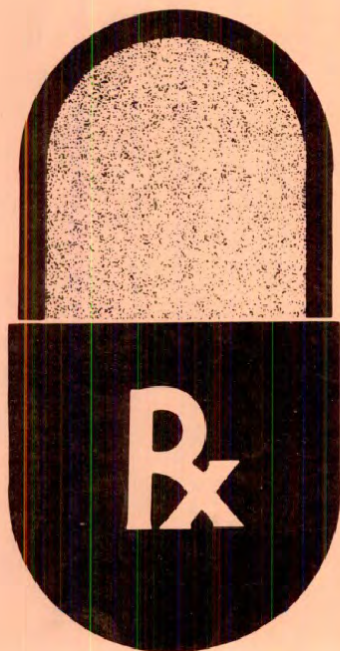
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**CUMULATIVE
SUPPLEMENT 5**
AUG'83 - JAN'84

SERIAL



APPROVED PRESCRIPTION DRUG PRODUCTS

WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS

4TH EDITION

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FOOD AND DRUG ADMINISTRATION
APPROVED PRESCRIPTION DRUG PRODUCTS
WITH
THERAPEUTIC EQUIVALENCE EVALUATIONS
CUMULATIVE SUPPLEMENT

I. PREFACE

This cumulative supplement is one of a series of monthly updates to the Approved Prescription Drug Products with Therapeutic Equivalence Evaluations, 4th Edition, (the List) to cover interim revisions to the annual publications of the List in its entirety. The List is comprised of several parts and some by their nature are identified by the term "List". The cumulative supplements routinely provide updates to two of these lists: The Drug Product List and the DESI Addendum.

The List cannot be used effectively without the current cumulative supplement. Users may wish to place an asterisk (*) in the List to the left of the ingredient(s) in the Drug Product List and the product name in the Addendum to indicate that changes to that entry appear in the cumulative supplement. It is also suggested that earlier cumulative supplements be discarded to avoid possible confusion. In this way, only the List and current cumulative supplement need be referenced.

A. DRUG PRODUCT LIST

The Drug Product List cumulative supplements include the changes made since August 1, 1983. Each subsequent cumulative supplement replaces the previous month's cumulative supplement.

Information in this cumulative supplement follows the format of the Drug Product List. 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.)

Context information on drug products is provided in each cumulative supplement for completeness to assist in locating the proper place in the Drug Product List for the revision. (Strength(s) which already exist in the publication will not be repeated for context.) A page number in parentheses referring to the Drug Product List is located to the right of the ingredient(s).

Additions to the Drug Product List are indicated by new information in the cumulative supplement. Additions new to the current cumulative supplement are indicated by the symbol >ADD> to the left of the line on which new information exists. The >ADD> symbol is dropped in subsequent cumulative supplements for that item.

Deletions from the Drug Product List are indicated by overstruck print in the cumulative supplement. Deletions new to the current cumulative supplement are indicated by the symbol >DELETED (DELETE) to the left of the line containing the overstruck print. The >DELETED symbol is dropped in subsequent cumulative supplements for that item.

A newly approved product is identified by the lozenge (◊) to the right of its strength. This identifier remains throughout all cumulative supplements for this edition.

B. ADDENDUM: DESI Pending List

Information in this cumulative supplement follows the format of the Addendum. Additions and deletions are indicated in the same manner as in the cumulative supplement to the Drug Product List. A change in Current Status of a DESI product is also indicated by an addition and a deletion.

II. SPECIAL NOTES

A. REPORT OF COUNTS FOR THE DRUG PRODUCT LIST

Categories of counts derived from product information in the Drug Product List and from this cumulative supplement are presented. The report includes counts of new molecular entities approved by the agency during the current month.

B. PRODUCTS CONTAINING PHENACETIN

The October 5, 1983, Federal Register (48FR45466) provides the following Summary: "The Food and Drug Administration (FDA) is withdrawing approval of new drug applications or parts of new drug applications that provide for drug products containing phenacetin, except for those drug products that are the subject of a hearing request. The basis of the withdrawal is phenacetin's high potential for misuse and its unfavorable benefit-to-risk ratio when incorporated in analgesic combinations which are then subject to excessive chronic use." The effective date of this withdrawal order is November 4, 1983.

Because the subject products are no longer approved, the cumulative supplement has identified them by deleting the applicable active ingredient headers followed by a reference to this Special Note.

C. THEOPHYLLINE CONTROLLED RELEASE

Two controlled released theophylline tablets are listed as therapeutically equivalent (AB). Because one of these products was recently approved for once-daily dosing, it is important to be aware that the therapeutic equivalence rating was made on the basis of 12-hour dosing intervals. The rating does not apply to once-daily dosing. To date, no data have been submitted upon which the Agency can base a therapeutic equivalence determination among any of the approved theophylline controlled-release products when dosed once-daily.

D. APPLICANT (NAME) CHANGES

Because 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, the cumulation of these transfers and name changes will be identified in this Special Notes section only. Where only partial approved product lines are transferred between applicants, each approved product involved will appear as an applicant name change in the cumulative supplement. The current list of applicant holder changes follows.

APPLICANT (NAME) CHANGES

<u>Former Applicant (Name)</u>	<u>New Applicant (Name)</u>	<u>New Abbreviated Name</u>
STERI-MED INC SUB KETCHUM LABORATORIES INC	QUANTUM PHARMICS LTD	QUANTUM PHARMICS
KETCHUM LABORATORIES INC	QUANTUM PHARMICS LTD	QUANTUM PHARMICS
PROFESSIONAL PHARMACAL	OPTOPICS LABORATORIES	OPTOPICS LABS
DIV STERI-MED INC	CORP	

The reader should consult the above cumulative list each month to become aware of such transfers and changes. By referring to the Applicant Index in the 4th Edition of the APDP, the transferred products can be identified. The reader might wish to flag these products in the list as a reminder that the applicant has been changed.

DESCRIPTION OF REPORT

The following report provides summary counts derived from product information in the Drug Product List and the current cumulative supplement. The counts appear in two sections. Section A. refers to the products in the List and Section B. to products in the current cumulative supplement. A new column of data will appear in Section A. each three-month period following July '83. Section A. therefore will provide baseline and quarterly data while Section B. provides monthly activity.

USE OF REPORT

From the data presented under Section B., users should be able to observe such things as (1) newly approved, DCSI effective and remarketed drug products which are added to the List; (2) products that are being removed from the List as the result of withdrawal of approval, changes from prescription to over-the-counter status and discontinued marketing of products; and, (3) trends in approval of products as either multi-source or single source during each month within the quarter. The report does not reflect category changes from multi-source to single source and vice versa. However, the net gain that results from all additions, deletions and category changes is reflected in the quarterly counts for multi-source and single source products.

Drug Product Definition

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

New Molecular Entity

The active moiety 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 part of a combination.

Drug Product Count

This report provides counts in several categories from the List composed of domestically marketed drug products approved for both safety and effectiveness under sections 505 and 507 of the Federal Food, Drug, and Cosmetic Act. Counts of products still pending in the DCSI review are not provided. Excluded also are those approved drug products marketed by distributors; those marketed solely abroad; and products now regarded as medical devices, biologics or foods. Also not included in the counts are those duplicate products of a given applicant whose only distinguishing characteristics are items such as package size, inactive ingredient(s), color and alternate manufacturing sites. These various counts are excluded because the Drug Product List itself excludes products from these categories.

REPORT OF COUNTS FOR THE DRUG PRODUCT LIST

A. COUNTS CUMULATIVE BY QUARTERS

<u>CATEGORIES COUNTED</u>	<u>JULY '83 (BASELINE)</u>	<u>OCT '83</u>
DRUG PRODUCTS LISTED	6679	6783
SINGLE SOURCE	1908 (28.6%)	1915 (28.2%)
MULTISOURCE ⁽¹⁾	4771 (71.4%)	4868 (71.8%)
THERAPEUTICALLY EQUIVALENT	3804 (57.0%)	3891 (57.4%)
NOT THERAPEUTICALLY EQUIVALENT	957 (14.3%)	967 (14.3%)
EXCEPTIONS ⁽²⁾	10 (0.1%)	10 (0.1%)
NEW MOLECULAR ENTITIES APPROVED	-	2
NUMBER OF APPLICANTS	304	310

B. ACTIVITY FOR SUPPLEMENT NUMBER 5

	<u>NOV '83</u>	<u>DEC '83</u>	<u>JAN '84</u>	<u>CUMULATIVE</u>
DRUG PRODUCTS ADDED:	43	39	32	114
NEWLY APPROVED	43	39	32	114
DESI EFFECTIVE	0	0	0	0
REMARKETED	0	0	0	0
DRUG PRODUCTS REMOVED:	36	6	2	44
WITHDRAWN APPROVAL	27	0	0	27
RX TO OTC SWITCH	0	0	0	0
DISCONTINUED MARKETING	9	6	2	17
NET GAIN IN DRUG PRODUCTS	7	33	30	70
SINGLE SOURCE PRODUCTS APPROVED	10	9	18	37
MULTISOURCE DRUG PRODUCTS APPROVED	33	30	14	77
NEW MOLECULAR ENTITIES APPROVED:	3	2	1	6
AS THE ENTITY	3	1	1	5
AS A SALT, ESTER OR DERIVATIVE				
OF THE ENTITY	0	1	0	1

(1) THERAPEUTIC EQUIVALENCE EVALUATIONS PROVIDED ONLY FOR MULTISOURCE PRODUCTS (I.E., AVAILABLE FROM MORE THAN ONE APPLICANT)

(2) AMINO ACID-CONTAINING PRODUCTS OF VARYING COMPOSITION (SEE PAGE 1-5 OF THE LIST)

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ACETAMINOPHEN; CODEINE PHOSPHATE (PAGE 3-1)

capsule; oral
ACETAMINOPHEN W/ CODEINE #3
 AA LEMMON 300MG;30MG*
 AA TYLENOL W/ CODEINE NO. 3
 AA MCNEIL PHARM 300MG;30MG

ACETAMINOPHEN; OXYCODONE HYDROCHLORIDE (PAGE 3-2)

TABLET; ORAL
CODACET
 AA HALSEY DRUG 325MG;5MG*
 AA PERCOCET-5
 AA PERCOCET

/ACETAMINOPHEN; PHENACETIN; PHENYLPROPANOLAMINE HYDROCHLORIDE;/
/PHENYLTOLOXAMINE CITRATE/ (PAGE 3-2)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

ALBUMIN, IODINATED, I-131, SERUM (PAGE 3-4)

INJECTABLE; INJECTION
RADIOIODINATED SERUM ALBUMIN (HUMAN) I-131
/MALLINCKRODT/ 16.7-250 UCL/ML

AMINO ACIDS (PAGE 3-6)

INJECTABLE; INJECTION
 > ADD > NEOPHAM 6.5%
 > ADD > CUTTER-VITRUM 6.5%*

AMINOPHYLLINE (PAGE 3-8)

/Liquibid; Oral/
 SOLUTION; ORAL
AMINOPHYLLINE
 AA BAY LABORATORIES 105MG/5ML*
 AA ROXANE LABORATORIES 105MG/5ML*

TABLET; ORAL
AMINOPHYLLINE
 BD BARR LABORATORIES 100MG*
 BD 200MG*
 AB VANGARD LABS/MHM 100MG*
 AB 200MG*

/AMPHETAMINE SULFATE/ (PAGE 3-13)

/CAPSULE; CONTROLLED RELEASE; ORAL
/BENZEDRINE/
/SALF LABORATORIES/ 15MG/
/TABLET; ORAL/
/BENZEDRINE/
/SALF LABORATORIES/ 15MG/
/15MG/

AMPICILLIN TRIHYDRATE; PROBENECID (PAGE 3-13)

POWDER FOR RECONSTITUTION; ORAL
POLYCELLIN-FRB
 AB BRISTOL LABS/B-M EQ 3.5GM BASE/BOT;1GM/BOT
PROBAMPACIN
 AB BIOCRRAFT LABS EQ 3.5GM BASE/BOT;1GM/BOT

ASPIRIN; BUTALBITAL (PAGE 3-15)

TABLET; ORAL
 AXOTAL
 ADRIA LABORATORIES 650MG;50MG*

ASPIRIN; BUTALBITAL; CAFFEINE (PAGE 3-15)

TABLET; ORAL
ASPIRIN AND CAFFEINE W/ BUTALBITAL
 > ADD > AB PUREPAC/KALIPHARMA 325MG;50MG;60MG*
 > ADD > AB BUTALBITAL W/ ASPIRIN & CAFFEINE
 > ADD > AB BOOTS LABORATORIES 325MG;50MG;60MG*

/ASPIRIN; BUTALBITAL; CAFFEINE; PHENACETIN/ (PAGE 3-15)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

ASPIRIN; CAFFEINE; DIHYDROCODEINE BITARTRATE (PAGE 3-15)

CAPSULE; ORAL
 SYNALGOS-DC
 IVES LABS/AMHO 356.4MG;30MG;16MG*

/ASPIRIN; CAFFEINE; PHENACETIN; PROPOXYPHENE HYDROCHLORIDE/
 (PAGE 3-15)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

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BROMPHENIRAMINE MALEATE (PAGE 3-22)

TABLET; ORAL

BROMPHENIRAMINE MALEATE
/AA/ /BOLAR PHARMACEUTICAL/ 4MG/

BUPIVACAINE HYDROCHLORIDE; EPINEPHRINE BITARTRATE (PAGE 3-23)

INJECTABLE; INJECTION

MARCAINE MQL W/ EPINEPHRINE
AP BREON LABS/STERLING 0.5%:0.0091MG/ML
AP 0.75%:0.0091MG/ML
SENSORCAINE
AP ASTRA HOSP PHARM 0.5%:0.0091MG/ML
AP 0.75%:0.0091MG/ML

BUTABARBITAL SODIUM (PAGE 3-24)

CAPSULE; ORAL

BUTICAPS
> DLT > /MCNEIL LABORATORIES/ 15MG/
> DLT > /30MG/
> DLT > /50MG/
> DLT > /100MG/
> ADD > WALLACE LABS/C-W 15MG
> ADD > 30MG
> ADD > 50MG
> ADD > 100MG

ELIXIR; ORAL

BUTABARBITAL SODIUM
/AA/ /REST-WARD/ 15MG/5ML/
/AA/ BUTALAN
/AA/ LANNETT 33.3MG/5ML

CAFFEINE; CARISOPRODOL; CODEINE PHOSPHATE; PHENACETIN (PAGE 3-24)

/TABLET; ORAL/
/SONA COMPOUND W/ CODEINE/
/WALLACE PHARM/C-W/ 32MG:100MG:16MG:160MG/

CAFFEINE; CARISOPRODOL; PHENACETIN (PAGE 3-25)

(ALL PRODUCTS - SEE SPECIAL NOTE B.)

CAFFEINE; ERGOTAMINE TARTRATE (PAGE 3-25)

SUPPOSITORY; RECTAL

CAFERGOT
BR SANDOZ PHARMS/SANDOZ 100MG;2MG
MIGRAINE
BR ORGANON/AKZONA 100MG;2MG

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE (PAGE 3-26)

SOLUTION; INTRAPERITONEAL

DIALYTE CONCENTRATE W/ DEXTROSE 30% IN PLASTIC CONTAINER
AM MCGAW/AM HOSP 510MG/100ML;30GM/100ML;
200MG/100ML;9.2GM/100ML;
9.6GM/100ML
510MG/100ML;30GM/100ML;
200MG/100ML;9.4GM/100ML;
11GM/100ML
DIALYTE CONCENTRATE W/ DEXTROSE 50% IN PLASTIC CONTAINER
AM MCGAW/AM HOSP 510MG/100ML;50GM/100ML
200MG/100ML;9.4GM/100ML;
9.6GM/100ML
510MG/100ML;50GM/100ML;
200MG/100ML;9.4GM/100ML;
11GM/100ML

CALCIUM CHLORIDE; DEXTROSE; MAGNESIUM CHLORIDE; SODIUM CHLORIDE; SODIUM LACTATE (PAGE 3-26)

SOLUTION; INTRAPERITONEAL

DIALYTE W/ DEXTROSE 2.5% IN PLASTIC CONTAINER
AM MCGAW/AM HOSP 26MG/100ML;2.5GM/100ML;
15MG/100ML;560MG/100ML;390MG/100ML

CALCIUM CHLORIDE; MAGNESIUM CHLORIDE; POTASSIUM CHLORIDE; SODIUM ACETATE; SODIUM CHLORIDE; SODIUM CITRATE (PAGE 3-27)

INJECTABLE; INJECTION

ISOLYTE E IN PLASTIC CONTAINER
AM MCGAW/AM HOSP 35MG/100ML;30MG/100ML;
74MG/100ML;640MG/100ML;
500MG/100ML;74MG/100ML

CEFTIZOXIME SODIUM (PAGE 3-30)

INJECTABLE; INJECTION

CEFIZOX
SK&F LABORATORIES EQ 1GM BASE/VIALM
EQ 2GM BASE/VIALM

CEFUROXIME SODIUM (PAGE 3-50)
INJECTABLE; INJECTION
EQ 750MG BASE/VIALM
GLAXO

CEPHALOTHIN SODIUM (PAGE 3-51)
INJECTION
KEPLIN
ELI LILLY
SEFTIN
AP GLAXO
EQ 1GM BASE/VIAL
EQ 2GM BASE/VIAL
EQ 1GM BASE/VIALM
EQ 2GM BASE/VIALM

CEPHALOTHIN SODIUM; DEXTROSE (PAGE 3-51)
INJECTABLE; INJECTION
CEPHALOTHIN SODIUM IN PLASTIC CONTAINER
TRAVENOL LABS
EQ 20MG BASE/NL;50MG/NLM
EQ 40MG BASE/NL;50MG/NLM

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

CEPHALOTHIN SODIUM; SODIUM CHLORIDE (PAGE 3-51)
INJECTABLE; INJECTION
CEPHALOTHIN SODIUM IN PLASTIC CONTAINER
TRAVENOL LABS
EQ 20MG BASE/NL;9MG/NLM
EQ 40MG BASE/NL;9MG/NLM

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

CHLORDIAZEPOXIDE (PAGE 3-53)
CAPSULE, CONTROLLED RELEASE; ORAL
LIBREASE
HOFFMAN-LA ROCHE
306MG

CHLORDIAZEPOXIDE HYDROCHLORIDE (PAGE 3-53)
CAPSULE; ORAL
CHLORDIAZEPOXIDE HCL
B&B PHARMACEUTICALS INC
18MG
10MG
25MG
40MG

CORTICOTROPIN (PAGE 3-43)

INJECTABLE; INJECTION
~~REPOSITORY CORTICOTROPIN~~
 /BC/ /MYETH. LABS/AMMO/ /40 UNITS/ML/
 /BC/ / / /80 UNITS/ML/

CYANOCOBALAMIN (PAGE 3-44)

INJECTABLE; INJECTION
DODECAMIN
 AP MAURRY BIOLOGICAL 1MG/MLM
 /AB/ REDISOL /0.1MG/ML/
 /AB/ MSD/MERCK /0.1MG/ML/
 /AB/ SYTOREX /0.1MG/ML/
 /AB/ PARKE DAVIS/M-L /0.1MG/ML/

CYCLOSPORINE (PAGE 3-46)

INJECTABLE; INJECTION
 SANDIMMUNE
 SANDOZ PHARMS/SANDOZ 50MG/MLM
 SOLUTION; ORAL
 SANDIMMUNE
 SANDOZ PHARMS/SANDOZ 100MG/MLM

CYPROHEPTADINE HYDROCHLORIDE (PAGE 3-46)

TABLET; ORAL
CYPROHEPTADINE HCL
 AA DURAMED PHARMS 4MGM

DAPSONE (PAGE 3-47)

TABLET; ORAL
~~ABOTT LABS/AMMO/~~
 /BP/ /MYERS. LABS/AMMO/ /25MG/
 /BP/ / / /100MG/
 DAPSONE
 /BP/ JACOBUS PHARM 25MG
 /BP/ / / 100MG

DESOXIMETASONE (PAGE 3-49)

ointment; TOPICAL
 TOPICORT
 HOECHST-ROUSSEL 0.25%M

DEXAMETHASONE (PAGE 3-49)

SOLUTION; ORAL
 DEXAMETHASONE
 ROXANE LABORATORIES 0.5MG/5MLM
 DEXAMETHASONE INTENSOL
 ROXANE LABORATORIES 0.5MG/0.5MLM

TABLET; ORAL
 DECADRON
 BP MSD/MERCK 6MG
 DEXAMETHASONE
 BP PAR PHARMACEUTICAL 6MGM
 BP ROXANE LABORATORIES 6MGM
 1MGH

DEXAMETHASONE SODIUM PHOSPHATE (PAGE 3-50)

INJECTABLE; INJECTION
DEXAMETHASONE
 > ADD > AP INVENEX LABS/DEXTER EQ 6MG PHOSPHATE/MLM
 > ADD > AP EQ 10MG PHOSPHATE/MLM
 /HEXADROL PHOSPHATE/
 AP HEXADROL EQ 6MG PHOSPHATE/ML
 EQ 20MG PHOSPHATE/ML

SOLUTION/DROPS; OPHTHALMIC

DEXAIR
 AT PHARMAFAIR EQ 0.1% PHOSPHATEM
DEXAMETHASONE SODIUM PHOSPHATE
 AT BARNES-HIND PHARMS EQ 0.1% PHOSPHATE

DEXTROSE; DOPAMINE HYDROCHLORIDE (PAGE 3-53)

INJECTABLE; INJECTION
~~DOPAMINE HCL IN DEXTROSE 5%~~
DOPAMINE HCL
 AP ABBOTT LABORATORIES 5GM/100ML;80MG/100ML
 AP 5GM/100ML;160MG/100ML
DOPAMINE HCL IN PLASTIC CONTAINER
 AP ABBOTT LABORATORIES 5GM/100ML;80MG/100MLM
 AP 5GM/100ML;160MG/100MLM
 5GM/100ML;320MG/100MLM

DEXTROSE; HEPARIN SODIUM (PAGE 3-53)

INJECTABLE; INJECTION
 HEPARIN SODIUM 20,000 UNITS AND DEXTROSE 5% IN PLASTIC
 CONTAINER
 TRAVENOL LABS 5GM/100ML;4,000 UNITS/100MLM

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ERYTHROMYCIN (PAGE 3-67)

OINTMENT; OPHTHALMIC
ERYTHROMYCIN
 AT E FOUGERA/BYK-GLDN 5MG/GM
 AT PHARMADERM/BYK-GLDN 5MG/GM
 AT ILOTYGIN
 AT DISTA PRODS/LILLY 5MG/GM

ETOPOSIDE (PAGE 3-74)

INJECTABLE; INJECTION
 VEPESIO
 BRISTOL LABS/B-M 20MG/MLM

FENTANYL CITRATE (PAGE 3-75)

INJECTABLE; INJECTION
 SUBLIMAZE
 JANSSEN PHARMA/ 15.0MG/ML/
 JANSSEN PHARMA EQ 0.05MG BASE/ML

FLUOCINOLONE ACETONIDE (PAGE 3-76)

CREAM; TOPICAL
 >_ADD_> FLUCCEY
 >_ADD_> AT NMC LABORATORIES 0.012M
 >_ADD_> AT 0.0252M

SOLUTION; TOPICAL
FLUCCINOLONE ACETONIDE
 >_ADD_> AT BAY LABORATORIES 0.012M

FLUOXHYMESTERONE (PAGE 3-77)

TABLET; ORAL
 FLUOXHYMESTERONE
 BP BOLAR PHARMACEUTICAL 2MG
 BP 5MG
 BP 10MG
 BP COLMED LABORATORIES 10MG

FUROSEMIDE (PAGE 3-79)

INJECTABLE; INJECTION
FUROSEMIDE
 AP NATCON CHEMICAL 10MG/MLM

FUROSEMIDE (PAGE 3-79)

TABLET; ORAL
FUROSEMIDE
 >_ADD_> AB BARR LABORATORIES 40MG
 AB ROXANE LABORATORIES 20MG
 AB 40MG
 AB ZENITH LABORATORIES 20MG
 AB 40MG

GENTAMICIN SULFATE (PAGE 3-79)

CREAM; TOPICAL
GENTAFAIR
 AT PHARMAFAIR EQ 1MG BASE/GM
 AT GENTAMICIN SULFATE
 AT NMC LABORATORIES EQ 1MG BASE/GM

INJECTABLE; INJECTION
GENTAMICIN SULFATE
 AP ABBOTT LABORATORIES EQ 60MG BASE/100MLM
 AP EQ 70MG BASE/100MLM
 AP EQ 80MG BASE/100MLM
 AP EQ 90MG BASE/100MLM
 AP EQ 100MG BASE/100MLM
 AP EQ 1.2MG BASE/MLM
 AP EQ 1.4MG BASE/MLM
 AP EQ 1.6MG BASE/MLM
 AP EQ 1.8MG BASE/MLM
 AP EQ 2MG BASE/MLM
 AP EQ 10MG BASE/MLM
 AP EQ 40MG BASE/MLM
 AP EQ 40MG BASE/MLM
 AP LYPHO-MED
GENTAMICIN SULFATE IN PLASTIC CONTAINER
 AP ABBOTT LABORATORIES EQ 60MG BASE/100MLM
 AP EQ 70MG BASE/100MLM
 AP EQ 80MG BASE/100MLM
 AP EQ 90MG BASE/100MLM
 AP EQ 100MG BASE/100MLM
 AP EQ 1.2MG BASE/MLM
 AP EQ 1.4MG BASE/MLM
 AP EQ 1.6MG BASE/MLM
 AP EQ 1.8MG BASE/MLM
 AP EQ 2MG BASE/MLM

OINTMENT; TOPICAL
GENTAMICIN SULFATE
 AT THAMES PHARMACAL EQ 1MG BASE/GM

OXTRIPHYLLINE (PAGE 3-131)

TABLET, ENTERIC COATED; ORAL

CHOLEDYL
 AB PARKE-DAVIS/W-L 100MG
 AB 200MG
OXTRIPHYLLINE
 AB BULAR PHARMACEUTICAL 100MG
 AB 200MG

OXYTETRACYCLINE HYDROCHLORIDE (PAGE 3-132)

CAPSULE; ORAL

OXLOPAR
 /AB/ /PARKE-DAVIS/W-L/ /EQ 250MG BASE/

PENICILLIN V POTASSIUM (PAGE 3-135)

POWDER FOR RECONSTITUTION; ORAL

Uticillin VK
 /AA/ /UP-PH/V /EQ 125MG BASE/ML/
 /AA/ /EQ 250MG BASE/5ML/

PHENDIMETRAZINE TARTRATE (PAGE 3-138)

TABLET; ORAL

PHENDIMETRAZINE TARTRATE
 AA FERDALE LABS 35MG

PHENTERMINE HYDROCHLORIDE (PAGE 3-139)

CAPSULE; ORAL

ADIPEX-P
 AA LEWTON 37.5MG
DAPEX-37.5
 AA FERDALE LABS 37.5MG
PHENTERMINE HCL
 AA CAMALL 37.5MG

POTASSIUM CHLORIDE (PAGE 3-143)

INJECTABLE; INJECTION

POTASSIUM CHLORIDE
 /AB/ TRAVENOL LABS /1MG/ML/
 /AB/ /5MG/ML/
 /AB/ /4MG/ML/

PREDNISOLONE (PAGE 3-145)

TABLET; ORAL

PREDNISOLONE
 /BP/ /FERDALE LABS/ /5MG/

PREDNISOLONE ACETATE (PAGE 3-146)

INJECTABLE; INJECTION

FERNISOLONE
 /BP/ /FERDALE LABS/ /25MG/ML/

PREDNISOLONE SODIUM PHOSPHATE (PAGE 3-147)

SOLUTION/DROPS; OPHTHALMIC

HYDELTRASOL
 /AT/ /NSD/HERCK/ /EQ 0.5% PHOSPHATE/
 > DLT > /AT/ /INFLAMASE/
 > DLT > /AT/ /STAP/PR/COOPERVISION/ /EQ 0.1% BASE/
 > ADD > /AT/ /INFLAMASE MILD/
 > ADD > /AT/ /COOPERVISION PHARMS EQ 0.1% PHOSPHATE
 > DLT > /AT/ /INFLAMASE FORTE/
 > ADD > /AT/ /STAP/PR/COOPERVISION/ /EQ 0.02% BASE/
 > ADD > /AT/ /COOPERVISION PHARMS EQ 0.02% PHOSPHATE
 > DLT > /AT/ /METRETON/
 > ADD > /AT/ /SCHERING EQ 0.5% PHOSPHATE
 > DLT > /AT/ /PREDAIR FORTE/
 > ADD > /AT/ /PHARMAFAIR EQ 0.9% PHOSPHATE
 > DLT > /AT/ /PREDNISOLONE SODIUM PHOSPHATE
 > DLT > /AT/ /BARNES-HIND PHARMS /EQ 0.1% PHOSPHATE/
 > ADD > /AT/ /BARNES-HIND PHARMS EQ 0.1% PHOSPHATE
 > DLT > /AT/ /BARNES-HIND PHARMS EQ 0.2% PHOSPHATE
 > DLT > /AT/ /BARNES-HIND PHARMS EQ 0.1% PHOSPHATE
 > ADD > /AT/ /MAURY BIOLOGICAL EQ 0.1% PHOSPHATE
 > ADD > /AT/ /EQ 0.9% PHOSPHATE

PREDNISONE (PAGE 3-147)

TABLET; ORAL

PREDNISONE
 BX DURAMED PHARMS 5MG
 BX 10MG
 BX 20MG

PROCAINAMIDE HYDROCHLORIDE (PAGE 3-149)

CAPSULE; ORAL

PROCAN
 /AB/ PARKE-DAVIS/W-L /250MG/
 /AB/ /500MG/

SOMATROPIN (PAGE 3-165)

INJECTABLE; INJECTION
ASELLACRIN 2
SERONO LABS 2 IU/VIALM
~~/ASELLACRIN/~~
ASELLACRIN 10

SPIRONOLACTONE (PAGE 3-166)

TABLET; ORAL
SPIRONOLACTONE
AB PUREPAC/KALIPHARMA 25MG

SULFACETAMIDE SODIUM (PAGE 3-167)

SOLUTION/DROPS; OPHTHALMIC
SODIUM SULFACETAMIDE
~~/AT/~~ /KETCHUM LABORATORIES/ 100/
~~/AT/~~ /KETCHUM LABORATORIES/ 300/
AT OPTOPICS LABS 100
AT 300
SULFACEL-15
~~/AT/~~ /STERIL-NEP/KETCHUM/ 150/
AT OPTOPICS LABS 150
SULFAIR FORTE
AT PHARMAFAIR 300

TESTOSTERONE (PAGE 3-173)

INJECTABLE; INJECTION
TESTOSTERONE
CARTER-GLOBAU LABS 50MG/MLM

TETRACYCLINE HYDROCHLORIDE (PAGE 3-174)

CAPSULE; ORAL
TETRACYCLINE HCL
~~/AB/~~ /LEITCH/ 150MG/
TABLET; ORAL
PANNYCIN
~~/AB/~~ UPJOHN 150MG/
SUNYCIN
~~/AB/~~ ER SQUIBB AND SONS 250MG

THALLOUS CHLORIDE, TL-201 (PAGE 3-175)

INJECTABLE; INJECTION
THALLOUS CHLORIDE TL 201
MEDI-PHYSICS 2MCI/ML

THEOPHYLLINE (PAGE 3-176)

(FOR CONTROLLED RELEASE PRODUCTS - SEE SPECIAL NOTE C.)

CAPSULE, CONTROLLED RELEASE; ORAL
SOMOPHYLLIN-CRT
BC FISOHS 100MG
THEO-24
BC SEARLE/SEARLE PHARMS 100MG
200MG
300MG

ELIXIR; ORAL
> ADD > ELIXONCH
> ADD > AA HR CENCI LABS 80MG/15MLM

SOLUTION; ORAL
THEOLATE
AA RIKER LABS/3M 80MG/15ML
THEOPHYLLINE
AA ROXANE LABORATORIES 80MG/15MLM

TABLET, CONTROLLED RELEASE; ORAL
THEOCONTIN
BC PURDUE FREDERICK 200MG
> DLT > /500MG/
THEO-DUR
BC KEY PHARMACEUTICALS 200MG
> ADD > UNIPHYL
> ADD > PURDUE FREDERICK 400MG

THIORIDAZINE HYDROCHLORIDE (PAGE 3-178)

CONCENTRATE; ORAL
THIORIDAZINE HCL
AA CORD LABORATORIES 30MG/MLM
AA 100MG/MLM
AA NATL PHARM MFG/BARRE 100MG/MLM

TABLET; ORAL
MELLARIL
AB SANDOZ PHARMS/SANDOZ 100MG
THIORIDAZINE HCL
AB BARR LABORATORIES 10MG
AB 15MG
AB 25MG
AB 50MG
AB 100MG
AB BOLLAR PHARMACEUTICAL 100MG
AB 100MG
AB CORD LABORATORIES 10MG
AB 15MG
AB 25MG
AB 50MG

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ASCORBIC ACID; BIOTIN; CYANOCOBALAMIN; DEXPANTHENOL;
ERGOCALCIFEROL; FOLIC ACID; NIACINAMIDE; PYRIDOXINE
HYDROCHLORIDE; RIBOFLAVIN PHOSPHATE SODIUM; THIAMINE
HYDROCHLORIDE; VITAMIN A; VITAMIN E (PAGE AD2)

/METHANODSTENOLONE/ (PAGE AD6)

/TABLET; ORAL/
/METHANODSTENOLONE/
/PAR. PHARMACEUTICAL/ 12.5mg/
5mg/

INJECTABLE; INJECTION

MVC PLUS

ASCOT HOSP PHARMS 10MG/ML; 0.006MG/ML; 0.5 UGM/ML;
 1.5MG/ML; 20 IU/ML; 0.04MG/ML;
 4MG/ML; 0.4MG/ML; 0.36MG/ML;
 0.3MG/ML; 330 IU/ML; 1 IU/ML

/BROMPHENIRAMINE MALEATE; PHENYLEPHRINE HYDROCHLORIDE;
/PHENYLEPROPANOLAMINE HYDROCHLORIDE/ (PAGE AD3)

/ELIXIR; ORAL/
/ELIXIR; DINEYAPP/
/AM. ROBIN/ 4mg/5ml; 5mg/5ml; 5mg/5ml/

/TABLET; CONTROLLED RELEASE; ORAL/
/DINEYAPP/
/AM. ROBIN/ 12mg; 15mg; 15mg/

/CARBAMPHEN EPHEDRINE MALEATE; CHLORPHENIRAMINE MALEATE; ISOPROPANOL/
/XOLOL; PHENYLEPROPANOLAMINE HYDROCHLORIDE/ (PAGE AD3)

/CAPSULE; CONTROLLED RELEASE; ORAL/
/XOLOL; ORNAGE/
/SKF. LABORATORIES/ 12mg; 15mg; 15mg; 5mg/

/SOLUTION; ORAL/
/XOLOL; ORNAGE/
/SKF. LABORATORIES/ 5mg/5ml; 5mg/5ml; 0.75/5ml; 15mg/5ml/

DICYCLOMINE HYDROCHLORIDE (PAGE AD3)

SYRUP; ORAL

BAYCICLOMINE

BAY LABORATORIES 10MG/5ML

DIPYRIDAMOLE (PAGE AD4)

TABLET; ORAL

DIPYRIDAMOLE

ASCOT HOSP PHARMS 50MG

DURAMED PHARMS 25MG

50MG

75MG

HALSEY DRUG 50MG

SUPERPHARM 50MG

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