105 CMR 720.000: LIST OF INTERCHANGEABLE DRUG PRODUCTS

Section

720.001: Purpose

The purpose of 105 CMR 720.000 is to establish a drug formulary, or list of interchangeable drug products, for use by physicians, other practitioners, and pharmacists licensed to practice within the commonwealth, so that consumers of prescription drug products may realize cost savings by buying less expensive, safe drug products.

720.002: Citation

105 CMR 720.000 shall be known as the 105 CMR 720.000: *Massachusetts List of Interchangeable Drug Products.*

720.010: Scope and Application

105 CMR 720.000 establishes the list of interchangeable drug products from which a pharmacist must interchange a reasonably available less expensive drug product than that written, when a prescription written by a practitioner indicates "interchange". 105 CMR 720.000 also establishes criteria and procedures for inclusion of drug products on this list.

720.020: Definitions

The terms used herein shall have the meanings set forth below. Terms defined in M.G.L. c. 112, § 12D and c. 94C, § 1, and not defined herein shall have the meanings set forth therein when used in 105 CMR 720.000, unless the context clearly requires a different interpretation.

Bioequivalent Drug Products means drug products whose rate and extent of absorption do not show a significant difference when administered at the same molar dose of therapeutic moiety under similar conditions. Some drug products may be equivalent in the extent of their absorption but not in their rate of absorption and yet may be considered therapeutically equivalent because such differences in the rate of absorption are not essential to the attainment of effective body drug concentrations or are considered medically insignificant for the particular drug product studies.

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Drug products for which bioequivalence is considered essential are those whose bioinequivalence would have therapeutic significance, *i.e.* use of different brands of the same drug product or different batches of the same drug product would result in therapeutic failure or a hazard to the patient. This is most critical in a drug product that has a narrow therapeutic-toxicity range which requires careful patient titration and monitoring for safe and effective use.

Commissioner means the commissioner of public health appointed under M.G.L. c. 17, § 2.

Department means the Department of Public Health established under M.G.L. c. 17 as an agency within the Executive Department of the Commonwealth of Massachusetts.

Drug Product means a product which contains an active drug ingredient and is in a dosage form, *e.g.* tablet, capsule, or solution, generally, but not necessarily in combination with other substances included in the manufacturing process. An active drug ingredient is that portion of a drug product intended to produce a therapeutic effect.

FDA means the Food and Drug Administration of the United States Department of Health and Human Services.

Generic name means a non-proprietary (common) name used to identify a drug product as listed by the United States Adopted Names Council and the United States Pharmacopeia in the USAN/USP *Dictionary of Drug Names*.

Interchangeable Drug Product means a product containing a drug in the same amounts of the same active ingredients in the same dosage form as other drug products with the same generic or chemical name.

Pharmaceutically equivalent drug products means drug products which contain the same active ingredients, and are identical in strength or concentration, dosage form, and route of administration.

Public Health Council means the Department's governing body established under M.G.L. c. 17, § 3. *See* also M.G.L. c. 111, § 3.

Therapeutically equivalent drug products means drug products which are pharmaceutically equivalent; meet applicable standards for strength, quality, purity and identity; are bioequivalent in that:

(a) they do not present a known or potential bioequivalence problem, and they do meet an acceptable in vitro standard; or

(b) if they do present such a known or potential problem, they are shown to meet an appropriate bioequivalence standards matching both rate and extent of absorption; are adequately labeled; and are manufactured in compliance with current Good Manufacturing Practice regulations.

720.040: Commission Review of Relevant Drug Products

In preparing the List of Interchangeable Drug Products and amendments thereto, the Drug Formulary Commission shall determine whether drug products meet the standards set forth in 105 CMR 720.050. In making this determination, the Commission shall assess and evaluate pertinent data, including, but not limited to, the United States Pharmacopeia and its supplements, additional pertinent listings of the FDA, other state formularies, formularies of various hospitals of the commonwealth, and data submitted by manufacturers and other interested persons, including chemical and laboratory listing data and clinical evidence concerning bioequivalence and therapeutic equivalence where available. In reviewing this material, the Commission shall utilize the pharmaceutical and medical expertise of its members.

720.050: List of Interchangeable Drug Products

The Massachusetts List of Interchangeable Drug Products (MLID) shall consist of:

(1) drug products which are considered by FDA to be therapeutically equivalent to other pharmaceutically equivalent products listed with the same generic or chemical name according to the most recent edition of "Approved Drug Products with Therapeutic Equivalence Evaluations" and its supplements as published by the United States Department of Health and Human Services:

(2) drug products specified on a list established by the Department and set forth in 105 CMR 720.200, for which the Commission has determined that the bioequivalence is not essential, or if the Commission has determined that the bioequivalence may be essential, bioequivalence has been established. The list may include the following categories of drug products:

(a) drug products which hold New Drug Applications (NDAs) or Abbreviated New Drug Applications (ANDAs) approved by the FDA, which FDA does not consider to be therapeutically equivalent to other pharmaceutically equivalent products listed with the same generic or chemical name; and

(b) drug products exempt from the Food, Drug and Cosmetic Act of 1962, and included in the Drug Efficacy Study Implementation (DESI) done by the National Academy of Sciences/National Research Council; and

(c) frequently prescribed drug products which were manufactured prior to 1938 and meet the FDA Good Manufacturing Practices Requirements; and

(d) frequently prescribed over-the-counter drug products which contain the same amounts of active ingredients, in the same dosage forms, as other drug products with the same general or chemical name.

720.060: Drug Products Excluded

The following categories of drug products are excluded from the list of interchangeable drug products:

(a) drug products for which the Commonwealth has determined that bioequivalence may be essential, but for which bioequivalence has not been established; and

(b) drug products which are the subject matter of patent rights issued by the U.S. Patent Office, for which provision by other than the patent-holder would violate the patent; and

(c) drug products available from only one manufacturer at one price.

720.070: Amendments to the Massachusetts List of Interchangeable Drugs

(1) Drug products which meet the criteria specified in 105 CMR 720.050(1) shall be deemed interchangeable and added to the Massachusetts List of Drugs upon publication by the United States Department of Health and Human Services of the most recent edition of "Approved Drug Products with Therapeutic Equivalence Evaluations" and its supplements.

(2) Drug products which meet criteria specified in 105 CMR 720.050(2) shall be deemed interchangeable and added to the Massachusetts List of Interchangeable Drugs in accordance with procedures set forth in 105 CMR 720.080.

720.080: Procedures for Amending the Massachusetts List of Interchangeable Drugs

The Department, working with the Commission, shall review at least one a year and revise as necessary the list of interchangeable drug products adopted pursuant to 105 CMR 720.050(2), and shall have the authority to review and revise the list of interchangeable drug products adopted pursuant to 105 CMR 720.050(1) as necessary. The revisions to 105 CMR 720.050(1) shall be specified on an exception list established by the Department and set forth in 105 CMR 720.200. The revisions will add and delete drug products, based on current information concerning therapeutic efficacy and interchangeability of drug products.

720.081: Petition to Amend List of Interchangeable Drug Products

Any person who desires a drug product or products to be added to or deleted from the List of Interchangeable Drug Products, shall file a written petition with the Department to amend the List, pursuant to M.G.L. c. 30A, § 4. Each petition shall be in such form as the Department may require and shall be submitted to the Drug Formulary Commission.

720.082: Commission Review of Petition

Upon receipt of a petition, the Department shall submit the petition and the supporting information to the Commission for review. The Commission shall make a preliminary determination whether the List of Interchangeable Drug Products should be amended as proposed.

720.083: Notice of Public Comment Period

Upon completion of the review of all relevant information, including petitions, by the Commission, the Department shall propose amendments to the List of Interchangeable Drug Products by issuing a Notice of Public Comment Period pursuant to M.G.L. c. 30A, §§ 2 and 3. The Department shall mail a Notice of Public Comment Period to each person who filed a petition during the period ending 30 days before the Notice of Public Comment Period is issued. In addition, the Department shall mail a Notice of the Public Comment Period to each person who has filed a written request therefore with the Department during December of the previous year pursuant to M.G.L. c. 30A, § 2.

720.084: Commission Recommendation of Amendments to Department

Following the comment period Department staff shall review all evidence and commentary concerning the proposed amendments, and shall report its recommendation to the Commission. The Commission shall consider the staff recommendations, make such revisions as it deems appropriate, and shall recommend Amendments to the List of Interchangeable Drug Products for adoption by the Commissioner and the Public Health Council.

720.090: Department Adoption of Amendments

The Commissioner and the Public Health Council shall consider the recommendations of the Drug Formulary Commission, and shall adopt Amendments to the List of Interchangeable Drug Products.

720.100: Severability

The provisions of 105 CMR 720.000 are severable. If any provision shall be declared invalid by any court, such provision shall be null and void and such determination shall not affect or impair any of the remaining provisions.

REGULATORY AUTHORITY

105 CMR 720.000: M.G.L. c. 17, § 13; c. 112, § 12D.

720.200: Appendix A

MASSACHUSETTS
LIST OF
INTERCHANGEABLE DRUGS

Department of Public Health regulation 105 CMR 720.050 describes the *Massachusetts List of Interchangeable Drugs*.

105 CMR 720.050(a) calls for the automatic adoption of all "A" rated drug products listed in the "*Approved Drug Products with Therapeutic Equivalence Evaluations*" and its supplements as published by the U.S. Food and Drug Administration (FDA), Department of Health and Human Services. This publication is commonly referred to as the "*Orange Book*". It is reprinted by the U.S. Pharmacopeial Convention Inc. (USP) as *Volume III* of the *USP DI*.

105 CMR 720.050(b) allows for the establishment of the *Massachusetts Additional List of Interchangeable Drugs* (*Additional List*), and provides the criteria upon which these drug products are approved.

All prescriptions written by generic name can be interchanged if the drug is multi-source. To determine if a prescription written for a brand name drug product is interchangeable in Massachusetts:

1. Look up the drug product by the brand name in the index or by generic name in the "*Approved Drug Products with Therapeutic Equivalence Evaluations*" ("*Orange Book*"). The drug products are arranged alphabetically.

2. Compare the dosage form and strength of the drug product prescribed with the dosage form and strength of the same drug product in the "*Orange Book*".

3. If the same drug product, dosage form and strength has been assigned an "A" rating by FDA and is <u>not</u> listed on the *Exception List* contained within 105 CMR 720.050, the drug product is interchangeable.

4. If the drug product is not listed in the "*Orange Book*", refer to 105 CMR 720.050(b), the *Massachusetts Additional List of Interchangeable Drugs* (*Additional List*).

5. Look up the drug product by the generic name in the *Additional List*. The drug products are arranged alphabetically.

6. Compare the dosage form and strength of the drug product prescribed with the dosage form and strength of the same drug product listed on the *Additional List*.

7. If the same drug product, dosage form and strength are listed, the drug product is interchangeable.

Copies of the "*Approved Drug Products with Therapeutic Equivalence Evaluations*" and its supplements ("*Orange Book*") are available from the:

U.S. Food and Drug Administration
Department of Health and Human Services
Government Printing Office
Washington, D.C. 20402-9371
OPC 6768
(202) 783-3238
and www.fda.gov/cder/drug

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Copies of the *USP DI* (third volume of *USP DI* is the "*Orange Book*") are available from:

The United States Pharmacopeial Convention, Inc.,
12601 Twinbrook Parkway
Rockville, MD 20852
(301) 881-0666

Copies of the *Massachusetts Additional List of Interchangeable Drug Products* (document number 105 CMR 720.000) are available from:

The State House Bookstore
Room 116
Boston, MA 02133
(617) 727-2834
and www.magnet.state.ma.us/dph/dcp/Drug Formulary/Drug Interchange

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FOREWORD

The *Massachusetts List of Interchangeable Drugs*, is prepared by the Drug Formulary Commission (DFC) and the Department of Public Health. The DFC is comprised of nine men and women appointed by the Governor for the express purpose of developing a list of those drug products that are safely interchangeable -- that is, equivalent to each other in all significant respects. The DFC was established by M.G.L. c. 17, § 13. This law was enacted with the intent of saving money for consumers of prescription drugs, since drug products that are marketed under trademark or proprietary names are often available in the generic forms from competing manufacturers at substantially lower prices. M.G.L. c. 112, § 12D mandates prescription forms that allow practitioners to prescribe interchangeable drug products by simply signing the signature line. If a practitioner determines that a brand name drug product should be dispensed, he/she must sign the signature line and write the words "**no substitution**" in his/her own handwriting in the space provided below the signature line.

The regulations call for the automatic adoption of "A" rated drug products listed in the "*Approved Drug Products with Therapeutic Equivalence Evaluations*" and its supplements (commonly referred to as the "*Orange Book*") as published by the U.S. Food and Drug Administration, Department of Health and Human Services, plus a list of additional drug products, the *Massachusetts Additional List of Interchangeable Drugs* ("*Additional List*"), individually reviewed and approved by the DFC and the Department. The regulations provide the criteria upon which the drug products listed on the *Additional List* are approved for interchange. The regulations also provide the DFC and the Department with the authority to review any "A" rated drug product listed in the "*Orange Book*" or drug product approved for interchange on the *Additional List* and delete it from the list of interchangeable drug products if deemed appropriate. Drug products assigned an "A" rating by FDA which are deleted from the Massachusetts *List* are placed on the *Exception List*. Drug products listed on the *Additional List* which are subsequently deleted are removed from the *Additional List*.

Of the many factors considered by the Commission in determining which drugs to include on the *List*, equivalent safety and effectiveness are paramount. The Commission reviews evidence on bioequivalence and pharmaceutical equivalence and includes on the *List* only those drug products determined to be fully interchangeable and whose manufacturers are approved by the U.S. Food and Drug Administration. Practitioners may prescribe any drug that appears on the *List* with confidence that it is as safe and effective as its brand name counterpart.

The efforts of the Commission in the assessment and evaluation of data and the preparation of the *List* are to be commended. The Department presents the *Massachusetts List of Interchangeable Drugs* with pride and with confidence that the *List* will greatly benefit consumers throughout the Commonwealth.

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INTRODUCTION

INTERCHANGEABLE (GENERIC) DRUG LAW

In 1976 the Massachusetts Legislature passed an Act Further Regulating the Establishment of a Formulary of Interchangeable Drug Products (St. 1976, c. 470, § 13), commonly known as the Generic Drug Law. This law, enacted to promote and regulate the use of generic drugs, created the Drug Formulary Commission to develop a list of interchangeable drug products and also required the use of a standard prescription form to encourage practitioners to prescribe generic drugs.

PRESCRIPTION FORM

M.G.L. c. 112, § 12D mandates prescription forms with one signature line. If the prescriber signs the prescription form and writes the words "**no substitution**" in his/her own handwriting in the space provided below the signature line, the pharmacist must fill the prescription exactly as indicated, with no interchange permitted. However, if the prescriber signs the prescription and does not write "**no substitution**" under his/her signature, the pharmacist is legally required to dispense a less expensive, equivalent interchangeable drug product listed in the *Massachusetts List of Interchangeable Drugs* if one is reasonably available.

MASSACHUSETTS LIST OF INTERCHANGEABLE DRUGS

The *Massachusetts List of Interchangeable Drugs* (MLID) consists of the "A" rated drug products listed in the "*Approved Drug Products with Therapeutic Equivalence Evaluations*" and its supplements as published by the U.S. Food and Drug Administration, Department of Health and Human Services ("*Orange Book*") and the *Massachusetts Additional List of Interchangeable Drugs* (*Additional List*). The *Additional List* is developed by the Drug Formulary Commission. The Commission determines drug products to be interchangeable only when they meet certain criteria:

(a) the drug product is available from more than one source, with the same active ingredient in the same dosage form and strength;

(b) its manufacturer is approved by the U.S. Food and Drug Administration (FDA); and

(c) when essential to therapeutic outcome, the manufacturer of the drug has documented clinical evidence of bioequivalence.

The Commission judges that all the drugs included on the MLID meet these standards and are bioequivalent, if essential, based on assessment and evaluation of the U.S. Pharmacopeia and its supplements, other state and hospital formularies, listings of the U.S. Department of Health and Human Services of the FDA, and on the expertise of its members.

The *List* does not include:

(a) drugs that are protected by patent rights or available from only one source;

(b) many controlled-release and enteric coated drug products since they may not consistently deliver the same quantities of their active ingredients;

(c) those drugs for which the Commission had any significant doubt about safe interchange between manufacturers; and

(d) any drug for which bioequivalence is considered essential but for which bioequivalence has not been demonstrated or an appropriate standard for bioequivalence has not been established.

Bioequivalence is determined to be necessary for a particular drug when bioinequivalence might result in therapeutic failure or hazard to the patient. Bioequivalent drug products do not show a significant difference in the rate and extent of absorption when administered at the same dosage under similar conditions. Drugs that are equivalent in the extent to which they are absorbed into a patient's body that differ in the rate of absorption may be therapeutically equivalent -- having the same medical effect -- either because the rate of absorption is not essential to the attainment of effective body concentrations of the drug, or because the difference in the rate is otherwise considered medically insignificant. Bioequivalence is a primary

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consideration for those drug products with a narrow therapeutic/toxic dosage range (when variation in the rate or extent of absorption could have a critical effect) where careful determination of the correct dosage and monitoring of the patient is essential to safe and effective use. To determine for which drugs bioequivalence is essential, the Commission relies on expert medical testimony, studies done by the pharmaceutical industry, the knowledge and expertise of the individual members of the Commission, and advice from the FDA.

All drug products manufactured by FDA approved firms are considered safe and effective for their intended use, even if the product has not been included in the MLID. A practitioner may begin a patient's therapy with a drug product from any manufacturer who has been approved by the FDA, even though interchange of the drug once the dosage has been calculated for the individual is not advised.

Several generic drug products are manufactured under the same new drug application (NDA) as the brand name drug products. According to section 1.6 of the *Orange Book*, drug products with the same NDA are therapeutically equivalent. Massachusetts regulations allow the interchange of these products. Distributors or repackagers of drug products manufactured under the same NDA as the brand name product are not identified in the *Orange Book*. Pharmacists who may not be able to determine if drug products are interchangeable should contact the manufacturers, distributors or repackagers. In addition the Department maintains an unofficial list of these products.

Information relative to the Interchangeable (Generic) Drug Law may be obtained from the Department of Public Health, Division of Food and Drugs, 305 South Street, Jamaica Plain, MA 02130, telephone number (617) 727-2670, and from the Boards of Registration in Medicine, Dentistry and Pharmacy.

DRUG PRODUCT PROBLEM REPORTING INSTRUCTIONS

Since 1971 the United States Pharmacopeia (USP), in cooperation with various professional associations and the Food and Drug Administration (FDA), has operated the *Drug Product Problem Reporting Program*. This program can be utilized by pharmacists, physicians, or consumers to report any product problems encountered when using drugs interchanged under the Massachusetts generic drug law. The program is product oriented, and patient identification not requested. Should you prefer to remain anonymous, so indicate to the USP and your name will be withheld from the manufacturer and the FDA. Your participation in reporting problems will help to ensure that the drug products prescribed and dispensed in Massachusetts are of continued high quality.

Reports should be sent to The United States Pharmacopeial Convention, Inc., 12601 Twinbrook Parkway, Rockville, MD 20852, (301) 881-0666. The USP is an impartial, non-governmental organization concerned with drug standards and quality control. After USP receives a report, copies are forwarded to the FDA and to the manufacturer of the product involved. Either the FDA or the manufacturer may act to investigate or correct problems.

EXCEPTION LIST

Orange Book "A" rated drug products not approved for interchange.

There are currently no products designated to be listed on the Exception List.

ADDITIONAL LIST

The *Massachusetts Additional List of Interchangeable Drugs* (*Additional List*) has been printed in a format designed to be concise and understandable. Interchangeable drugs are listed alphabetically according to their official (chemical or generic) names, and separate sections in each listing show dosage forms, strengths, FDA approved manufacturers, and categories.

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DRUG

Drugs are listed in alphabetical order by their generic names and are printed in capital letters. Drug products containing more than one active ingredient (for example, CODEINE PHOSPHATE, GUAIFENESIN) are listed in the conventional order of ingredients.

Only drug products grouped under single headings are to be interchanged.

DOSAGE FORM

Under the generic names are listed the various multisource dosage forms in which a drug product is available. Abbreviations used for dosage forms and approved manufacturers are found in the front of the *Additional List*.

Only identical dosage forms and strengths of identical drugs are to be interchanged.

STRENGTH

The approved strengths of the drug products are listed under the heading "Strength(s)." The "strengths" must be read along with the "dosage forms" since any strength shown is available only for the dosage form directly to its left. Dosage strength is in metric units that are sometimes rounded off from apothecary measures, which may introduce slight variations in the strength of certain products. Single ingredient drug product strengths are separated by commas. Combination drug products have a slash separating the strengths of the individual ingredients. If more than one strength of a single component of a combination drug product is approved, they will be separated by commas. For example, the strength of a tablet of aspirin with codeine phosphate is "325mg / 15mg, 30mg, 60mg" which means that the combination is available with 325 milligrams of aspirin and 15, 30, or 60 milligrams of codeine phosphate. Drug products with three or more components have their active ingredients listed individually in parentheses and have slashes separating the strengths of the individual ingredients.

MANUFACTURERS

Next to the heading "Manufacturers" are all approved manufacturers for the drug product in that group, listed by three letter abbreviations in capital letters. (See list of manufacturer abbreviations in front of the *Additional List*.) Listed manufacturers have met all legal requirements, including compliance with the FDA Good Manufacturing Practices for the production of the drug product indicated. Approved manufacturers hold current new drug applications (NDAs) or abbreviated new drug applications (ANDAs) when required by law.

NDA, ANDA APPLICANT (NAME) CHANGES

Because it is not practical to identify in the *Massachusetts Additional List of Interchangeable Drugs* (*Additional List*) 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, these transfers and name changes are identified in this section. In addition, the new manufacturers are listed in parenthesis beside the original manufacturer under the *Manufacturers' Abbreviations* section of the *Additional List*. Where only partial approved product lines are transferred between applicants, each approved product involved will appear with the manufacturer name change in the *Additional List* amendment.

Previously listed name changes have been incorporated into the revised Manufacturers' Abbreviations section.

ABBREVIATIONS

The following abbreviations are used in the *Massachusetts List of Interchangeable Drugs*.

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DOSAGE FORMS

aero	aerosol	ml	milliliter
amp	ampule	oint	ointment
cap	capsule	ophth	ophthalmic
conc	concentrate	oral gran	oral granules for reconstitution
e.c.	enteric-coated		
elix	elixir	oral powder	oral powder for reconstitution
g	gram		
Hbr	hydrobromide	oral sol	oral solution
HCl	hydrochloride	pow	powder
HC	hydrocortisone	sol	solution
inhl	inhalation	SR	sustained release
inhl liquid	inhalation liquid	subl tab	sublingual tablet
inhl sol	inhalation solution	supp	suppository
inj	injection	susp	suspension
irr sol	irrigating solution	syr	syrup
I.U.	international units	tab	tablet
liq	liquid	top aero	topical aerosol
lot	lotion	top swab	topical swab
mcg	microgram	U	units
mEq	milliequivalents	vag	vaginal
mg	milligram		

MANUFACTURERS' ABBREVIATIONS

3MP	3M Pharmaceutical		Inc
AAA	Alpha Therapeutic	ARC	Arcola Labs.
ABB	Abbott	APC	Arcum Pharmaceuticlal Corp.
ABI	Abic	ARP	Armenpharm
ABL	Able Laboratories	ARM	Armour Pharmaceuticals
ACI	ACIC Limited	ASC	Ascot Hospital Products
ACP	Advanced Care Prod.	ASA	Asta
ADV	Advanced Remedies	ASP	Astra Pharmaceuticals LP
AGV	Agvar Chemicals	ATH	Athena Neurosciences
AKO	Akorn	BAK	Baker Norton
AKZ	Akzona Inc.	BAN	Banner Pharmacaps
ALC	Alcon Labs	BAP	Barlan Pharmacal
ALL	Allergan Pharmaceuticals	B/I	Boehringer Ingelheim
ALI	Alliance Pharmaceutical	B/M	Boehringer Mannheim. Ther. Div
ALP	Alpharma	BAR	Barr Labs
APP	Alphapharm Party	BAS	Basel Pharmaceuticals
ALT	Altana	BAT	Bartor
ALZ	Alza Corp.	B&L	Bausch & Lomb
ALR	Alra Laboratories	BAY	Bayer Corp
AMA	Amaric	BEA	Beach Prod.
AMB	Ambix Labs	B-D	Becton,Dickinson & Co.
ACC	American Cyanamid Co.	BED	Bedford Laboratories
AHP	American Home Products	BEL	Bell
ARL	American Regent Labs.	BDP	Beta Derm Pharmaceuticals
AME	Amersham	BER	Berlex
AMG	Amgen	BFA	B.F. Asher
AMI	Amide Pharmaceuticals	BHC	B.H. Chemicals
AMT	American Therapeutics	BID	Biodevelopment
ANA	Anabolic	BIO	Bio Technology General
ANB	Anbex	BIV	Biovail
ANC	Angus Chemical	BLA	Blairex Laboratories
ANE	Anesta	BLO	Block Drug Co.
ANG	Angelini		
APO	Apothecon		
APK	Apothekernes		
APP	American Pharmaceutical Partners		

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BLU	Bluline
BRL	Blue Ridge Laboratories
BOC	Bock Pharmacal
BOW	Bowman Pharm.
BMS	Bristol Myers Squibb
BRC	Bracco Diagnostics
BRD	Bradley Pharm.
BRA	Braintree Laboratories
BRI	Bristol-Myers Prod.
BTG	BTG Pharmaceuticals
BVL	Ben-Venue Labs
BYR	Byron Chemical
C&M	C & M Pharmacal
CAD	Cadema Medical Products
CAG	Calgon Corp.
CDC	Carderm Capital
C-P	Chesebrough - Ponds
C-W	Cook - Waite
C/C	Chase Chemical
CSL	Chase Laboratories
C/P	Corvit Pharmaceuticals
CAL	Carlisle
CAM	Camall
CAR	Carnick Labs
CTW	CarterWallace
CEN	Century Pharmaceuticals
CBV	Cetus Ben Venue Therapeutics
CHA	Chamberlin Parenteral Corp.
CHE	Chelsea Laboratories
CHM	Chemed Corp.
CVO	Ciba Vision Ophthalmics
CIR	Circa Pharmaceuticals
CJD	Copanos, J.D.
CLA	Clay - Park
CLO	Clonmel Healthcare
CMB	C.M. Bundy
CMC	Consolidated Midland Corp.
C/T	Controlled Therapeutics
CMP	Carolina Medical Products
CNC	H.R. Cenci
COL	Colgate Palmolive
COM	Combe
CON	Connaught Laboratories
COO	Cooper Labs
COP	Copley Pharmaceutical
CPG	Consolidated Pharmacy Group
CRE	Creighton Products
CUM	Cumberland Swan
CUR	Curatex Pharmaceuticals
CTL	Central Pharmacal
D&G	Davis & Geck
D/L	DPT Laboratories
D-R	Del-Ray Laboratories Inc.
DAN	Danbury Pharmacal
DAR	Darby Group Companies
DEL	Dell
DEP	Deproco
DER	Dermik Laboratories
DES	Deseret Medical
DEY	Deylabs
DHL	DHL Laboratories
DIA	Dial Corp.
DIS	Dista
DMD	Duramed
DMG	D M Graham Laboratories
DOW	Dow Pharmaceutical
DPT	Dupont Pharmaceuticals
DPM	Dupont Merck
DUR	Dura Pharmaceuticals
DUN	Dunhall
DYN	Dynapharm
E/K	Eastman Kodak
EAT	Eaton Medical Corp.
ECR	ECR Pharmaceuticals
ELA	Elan Pharmaceuticals
ELL	Ellis Pharmaceuticals
EMP	EM Pharma
END	Endo
EON	Eon Laboratories
ENQ	Enquay Pharm
ENZ	Enzon
ESR	Elkins-Sinn/AH Robins
ERS	Ersana
ESI	ESI Lederle Generics
ESP	ESI Pharmacal
ETH	Ethicon Inc.
ETX	Ethex
ETK	Ethitek Pharmaceuticals
EVY	Everylife
EZC	E Z EM Co.
FAU	Faulding Pharm. Co.
FER	Ferndale
FRT	Ferrante
FRR	Ferring Labs
FIS	Fisons
FLE	Fleming & Co.
FOR	Forest
FOU	Fougera
FOY	Foy
FRE	Fresenius
G&W	G & W
GAL	Galderma
GEI	Geigy
GEC	Gencon
GED	Genderm
GET	Genentech
GEP	Genpharm
GEV	Geneva
GES	Gensia Sicor Pharmaceuticals Inc
GEZ	Genzyme
GIL	Gilbert Laboratories
GLD	Glades
GLW	GlaxoWellcome
GLE	Glenwood
GLO	Global Pharm.
G/P	Golden Pharms.
GOL	Goldine
GRE	Greenstone
GRI	Griffen,KW
G/L	Gruppo Lepetit
GUA	Guardian
GYM	GYMA Laboratories
GYN	Gynopharm
H-R	Holland-Rantos

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HAL	Halsey Labs
HAM	Hamilton Pharmaceuticals
HAN	Hnford GC
HEA	Heather
HEN	Heran Pharmaceutical
HEX	Hexcel Chemical Products
HER	Hermal Pharmaceutical
HIC	Hickam
H/D	Hill Dermaceuticals
HIR	Hirsch Industries
HIT	Hi Tech Pharma
HTP	High Technology Pharmacal
HLC	Halocarbon
HCC	Hoechst Celanese Corp.
HMR	Hoechst Marion Roussell
HOE	Hoechst-Rousel
HOR	Horus Therapeutics
HOY	Hoyt
HUD	Hudson Pharmaceuticals
HUN	Huntington
HYB	Hybritec Inc.
HYG	Hygenics
HYR	Hyrex
IMM	Immunex
IMP	IMP Inc.
ICN	ICN Pharmaceuticals
IMS	International Medication
INP	Interpharm
INV	Invamed, Inc.
INW	Inwood Labs
ICC	Interchem Corp.
ILC	International Latex Corp.
ING	Ingram Pharmaceutical
INH	Inhalon
IOL	Iolab
IOM	Iomed
IPR	IPR Pharm
IVA	IVAX
J&J	Johnson & Johnson
JAC	Jacobus
JAN	Janssen Pharmaceuticals
JER	Jerome Stevens Labs
JRW	Johnson RW
JON	Jones Pharma Inc
KAL	Kalapharm
KBP	Kabi Pharmacia
KEE	Keene
KED	Kendall
KEN	Kenwood
KIN	King Pharmaceuticals
KIR	Kirkman Sales
KNO	Knoll
KPI	Key Pharmaceuticals
KVP	KV Pharmaceutical Co.
L/F	Labs Fournier
L/A	Laboratories Atral
LAF	Lafayette Pharms
LAN	Lannett
LED	Lederle
LEI	Leiras
LEK	Lek Ljubliana
LEM	Lemmon
LEO	Leo Pharms
LIF	Life Labs
LIL	Lilly
LIP	Liposome
LIQ	Liquipharm
LNK	LNK International
LOC	Loch Pharmaceuticals
LOR	Lorex
LOT	Lotus Biochemical
LPI	LPI Holding
LUI	Luitpold
LUS	Lek USA Inc.
LYN	LYNE Laboratories
M/P	Mallinckrodt Pharmaceutical.
MAY	Mayrand
MAT	Matrix Labs
MCG	Mcgaw
MCN	McNeil Consumer Products
MDP	MD Pharmaceuticals
MEA	Mead Johnson
MJN	Mead Johnson Nutritionals
M/R	Medco Research
MVA	Medeva
MEP	Medics Pharmaceuticals
MPI	Medi Physics, Inc.
MAG	Mepha AG
MER	Mericon
MET	Metronic
MGI	MGI Pharma
MID	Midway Medical
MIK	Mikart Laboratories
MIS	Mission Pharmacol
MJP	MJ Pharmaceuticals
MKL	Moore Kirk Labs
MLI	Marchar Laboratories
MLP	Miller Pharmacal
MLX	Milex
MMD	Marion Merrell Dow
MOR	Morton Grove
MCK	Merck & Co.
MSM	Marsam
MSL	Marshall Pharmacal
MTC	Martec
MOV	Mova
MUR	Muro
MUT	Mutual Pharmaceuticals
MYL	Mylan Pharmaceuticals
NEP	Nephron Pharmaceuticals Inc
NEU	Neutrogena
NOR	Norbrook Laboratories
N/W	Norton Waterford
N/N	Novo Nordisk
NEW	Newtron Pharmaceuticals
NHN	Norton HN
NOV	Novocol
NVP	Novopharm Ltd.
NUM	Numark
NYC	Nycomed
NYL	Nylos Trading
ORI	Organon, Inc.
OCL	Oclassen
OHM	OHM Laboratories

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OMD	Ohmeda Pharmaceutical
OSA	On Site Azla
ODC	Ormont Drug & Chemical
OPC	Ortho Pharmaceuticals
OPT	Optopic Laboratories Corp.
ORG	Organics
OAP	Otsuka America Pharmaceutical
PAK	Pal-Pak
PAL	Palisades
P/D	Parke Davis
P/I	Plantex/Ikpharm
P/K	Purepac-Kalipharma
P/P	Parmed Pharmaceuticals
PAC	Paco Research
PAD	Paddock Labs
PHD	Pharmaderm
PAN	Panray
PAR	Par
PNL	Parnell
PER	Perrigo
PCE	Pharmachemie
PHC	Pharmics
PHK	Pharmakinetics Labs
PHM	Pharmeral
PHO	Phoenix Labs
PHS	Pharma Serve
PHT	Pharmaton
PFF	Pfeiffer
PFI	Pfizer
P/U	Pharmacia & Upjohn
P/A	Pharmaceutical Association
PIO	Pioneer Pharmaceutical Inc.
PPI	Physicians Products Inc.
PSA	Pharmaceutical Specialist Assoc
POH	Pohl Boskamp
POL	Polymedia
PGP	Prographarm
PRD	Professional Disposables
PRO	Proter Laboratory
PRI	Private Formulations
P&G	Proctor & Gamble
PRV	Pharmavite
PTK	Pharma-Tek
PUF	Purdue Frederick
PUR	Purepac
QUA	Quantum Pharmics Ltd.
QLT	QLT PhototherapeuticsInc.
RAN	Ranbaxy Pharmaceuticals
R/C	Reckitt & Colman
R&C	Reed & Carnrick
R/I	Research Industries
RXP	Rexar Pharmacal
RAC	Rachelle Labs
REN	Ren-Pharm Internatl. Ltd.
RHP	Rhone - Poulenc
RPR	Rhone-Poulenc Rorer
RAH	Robins, A.H.
REX	Rexall/Sundown
RIC	Richlyn Labs
ROC	Roche Labs.
RPF	Roerig/Pfizer
ROA	Roaco
RBP	Roberts Pharmaceutical
ROR	Rorer
ROS	Ross Labs
ROX	Roxane Labs
ROY	Royce Laboratories
RPC	Rosemont Pharmaceutical Corp.
RUG	Rugby Labs
S-M	Spencer-Mead
S/W	Sanofi Winthrop
S/L	Schmid Laboratories
SAK	Sankyo
SAV	Savage Labs/Altana
SAN	Sandoz
SCE	Scherer, R.P.
SPI	Schein Pharmaceutical, Inc.
SCH	Schering Corporation
S/P	Schering/Plough
SWZ	Schwarz Pharma
SZG	SchwarzGMBH
SCI	ScinoPharm International
SCS	SCS Pharmaceuticals
SEA	Searle
SER	Serono Laboratories
SEQ	Sequus Pharmaceuticals
SHM	Sherwood Medical
SHI	Shionogi USA
SID	Sidmak Laboratories
SIG	Sigma Tau
SIX	Silarx
SKB	Smith, Kline Beecham
SBH	Sola Barnes Hind
SOL	Solopak Laboratories
SLV	Solvay
SOM	Somerset
SBM	Sorin Biomedics
SDP	Sperti Drug Products
STI	Steifel
STL	Stanlabs Pharmaceutical Co.
STR	Star Pharmaceuticals
STS	Steris Laboratories
STZ	Storz Ophthalmics
SUP	Superpharm
SPP	Suppositoria
SUR	Survival Technology
SYN	Syntex
SYO	Syosset Labs
TAB	Tablicaps
TAG	Tag Pharmaceuticals
TAK	Takeda
TAP	Tap Holdings
TAR	Taro Pharmaceutical
TAY	Taylor Pharmaceuticals
TEC	Technilab
THE	Theratech
THK	Therakos
THA	Thames Phamacol Co. Inc.
TIC	Tican Pharmaceuticals
TOP	Topiderm
T/L	Torch Laboratories
TOR	Torigian Lab
UDL	UDL Laboratories
UMD	Unimed

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UPJ	Upjohn	WPP	West Point Pharma
USL	Upsher-Smith Labs	WES	Westwood Squibb Pharmaceuticals
VAL	Vale Chemical	W-W	West-Ward
VAN	Vangard	W/L	Wharton Labs
VES	Vestal	WBY	Whitby
VIC	Vicks Pharmacy Products	WWT	Whiteworth Towne
VIN	Vintage	WAL	Wallace Labs
VIR	Viratek	WAR	Warner-Lambert
VIS	Vistakon, Inc.	WAT	Watson Laboratories
VIV	Vivan Pharmacal	WOC	Wockhardt
W-A	Wyeth Ayerst	XTT	Xttrium Laboratories
W-C	Warner-Chilcott	YAM	Yamanouchi
WAW	Warner Wellcome	YOS	Yoshitomi Laboratories
WRR	Warrick Pharm	ZCA	Zeneca
WEP	WE Pharmaceuticals	ZGP	Zenith Goldline Pharmaceuticals
WEN	Wendt Laboratories		

MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

AMINOPHYLLINE, EPHEDRINE Hcl		
Dosage form(s):	TABLETS	Strength(s): 130mg/25mg
Manufacturers:	STL	
Category:	OTC	
AMYL NITRITE		
Dosage form(s):	INHALATION	Strength(s): 0.3ml
Manufacturers:	GLW, CMC	
Category:	PRE-38	
ASPIRIN W/ CODEINE PHOSPHATE		
Dosage form(s):	TABLET	Strength(s): 325mg/15mg, 30mg, 60mg
Manufacturers:	BAR, GLW, CHE, GEV, HAL, P/D, ZGP	
Category:	PRE-38	
ATROPINE SULFATE		
Dosage form(s):	OPHTHALMIC SOLUTION	Strength(s): 0.5%, 1%, 2%
	OPHTHALMIC OINTMENT	0.5%, 1%
Manufacturers:	ALC, ALL, ESR, FOU, INV, MUR, B&L, STS, SUR	
Category:	PRE-38	
ATROPINE SULFATE COMPOUND (ATROPINE SULFATE, SCOPOLAMINE HBr, HYOSCYAMINE SULFATE, PHENOBARBITAL)		
Dosage form(s):	TABLET	Strength(s): 0.0194mg/0.0065mg/
	0	1037mg/16.2mg
Manufacturers:	ALL, CHE, M/P, MAY, TAY, RAH, WES	
Category:	DESI	
BENZOCAINE, ANTIPYRINE		
Dosage form(s):	OTIC SOLUTION	Strength(s): 1.4%, 5.4%
Manufacturers:	AMB, W-A, CLA, RPC, S-M, THA	
Category:	PRE-38	
BENZOYL PEROXIDE		
Dosage form(s):	GEL Strength(s): 2.5%, 5%, 10%	
Manufacturers:	BMS, CLA, GAL, STI, SYO, VIC, WES	
Category:	OTC	
BENZTHIAZIDE		
Dosage form(s):	TABLET	Strength(s): 50mg
Manufacturers:	GEV, PFI, RAH,	
Category:	B	
BROMPHENIRAMINE MALEATE, DEXTROMETHORPHAN HBr, PSEUDOEPHEDRINE HCl		
Dosage form(s):	SYRUP	Strength(s): 2mg/10mg/30mg/5ml
Manufacturers:	RAH, RPC	
Category:	OTC	

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MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

BROMPHENIRAMINE MALEATE, PHENYLPROPANOLAMINE HCl, PHENYLEPHRINE, GUAIFENESIN COMBINATION		
Dosage form(s):	SYRUP	Strength(s): 4mg/5mg/5mg/100mg/5ml
Manufacturers:	HAL, LIF, RAH, RPC,	
Category:	OTC	
CAFFEINE AND SODIUM BENZOATE		
Dosage form(s):	INJECTION	Strength(s): 250mg/ml
Manufacturers:	TAY	
Category:	PRE-38	
CALCIUM GLUCONATE		
Dosage form(s):	INJECTION	Strength(s): 10%
Manufacturers:	ARL, APC, APP, CMC, CEN, BAY, ESR, GEN, HYR, KIR, LAN, LIL, M/P, P/D, RIC, STL, P/U ,	
Category:	PRE-38	
CARBINOXAMINE MALEATE, PSEUDOEPHEDRINE HCl, DEXTROMETHORPHAN HBr		
Dosage form(s):	DROPS	Strength(s): 2mg/25mg/4mg/ml
	SYRUP	4mg/60mg/15mg/5ml
Manufacturers:	ALP, ROS	
Category:	PRE-38	
CHLORAL HYDRATE		
Dosage form(s):	CAPSULE	Strength(s): 500mg
	SYRUP	250mg/5ml, 500mg/5ml
Manufacturers:	BMS, C/C, ALP, PHC, PUR, ROX, SCE, , ZEN	
Category:	PRE-38	
CHLORDIAZEPOXIDE W/ CLIDINIUM BROMIDE		
Dosage form(s):	CAPSULE	Strength(s): 5mg/2.5mg
Manufacturers:	BAR, CHE, EON, GEV, HAL, ROC, LEM, PAR, QUA, ZGP	
Category:	DESI	
CHLOROTHIAZIDE W/ RESERPINE		
Dosage form(s):	TABLET	Strength(s): 250mg/0.125mg, 500mg/0.125mg
Manufacturers:	MCK, MYL,	
Category:	B	
CHLORPHENIRAMINE MALEATE		
Dosage form(s):	TABLET	Strength(s): 4mg, 8mg, 12mg
	SYRUP	2mg/5ml
Manufacturers:	LAN, SCH	
Category:	OTC	
CHOLINE MAGNESIUM TRISALICYLATE (CHOLINE SALICYLATE, MAGNESIUM SALICYLATE)		
Dosage form(s):	TABLET	Strength(s): 500mg (=293mg/362mg), 750mg (=440mg/544mg), 1000mg (=587mg/725mg)
Manufacturers:	PUF, SID	
Category:	PRE-38	
CODEINE PHOSPHATE		
Dosage form(s):	INJECTION	Strength(s): 15mg, 30mg, 60mg/ml
Manufacturers:	ESR, KNO, STL, W-A	
Category:	PRE-38	
CODEINE PHOSPHATE /GUAIFENESIN LIQUID		
Dosage form(s):	LIQUID	Strength(s): 10mg/100mg/5ml
Manufacturers:	RAH, HAL	
Category:	PRE-38	
CYANOCOBALAMIN		
Dosage form(s):	TABLET	Strength(s): 10mcg, 25mcg,
	CAPSULE	50mcg, 100mcg, 250mcg
Manufacturers:	APP, BER, BMS, STS, DEL, ESR, INV, LEM, LIL, MMD, MCK, ALP, ORI, P/D, SAV, SOL, P/U, W-A	
Category:	OTC	

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MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

CYCLANDELATE			
	Dosage form(s):	CAPSULE	Strength(s): 200mg/400mg
	Manufacturers:	CHE, GEV, DAN, FOR, INW, W-A, LAN, LEM, MDP, PAR, PIO, ZGP	
	Category:	DESI	
DEXAMETHASONE			
	Dosage form(s):	TABLET	Strength(s):0.25mg, 0.5mg, 0.75mg, 1.5mg, 4mg
	Manufacturers:	GEV, DAN, MCK, MYL, ORI, PAR, PRI, RIC, SLV, ROX, , USL	
	Category:	B	
DIETHYLPROPION HCl			
	Dosage form(s):	TABLET	Strength(s): 25mg
		SUSTAINED RELEASE TABLET	75mg
	Manufacturers:	CAM, , LEM, MMD, MDP, 3MP	
	Category:	B	
DIETHYLSTILBESTROL			
	Dosage form(s):	TABLET	Strength(s): 0.1mg, 0.5mg, 1.5mg
		VAGINAL SUPPOSITORIES	
	Manufacturers:	BMS, LIL	
	Category:	B	
DIGOXIN			
	Dosage form(s):	TABLET	Strength(s): 0.125mg, 0.25mg, 0.5 mg
	Manufacturers:	GLW, AMI	
	Category:	PRE-38	
DIGOXIN			
	Dosage form(s):	INJECTION	Strength(s): 0.25mg/ml, 0.5mg/2ml
	Manufacturers:	GLW, ESR, EON, W-A	
	Category:	PRE-38	
DIMENHYDRINATE			
	Dosage form(s):	TABLET	Strength(s): 50mg
	Manufacturers:	ANA, STS, CHE, GEV, ESR, LEM, SEA, , W-A	
	Category:	OTC	
DIPHENHYDRAMINE HCl			
	Dosage form(s):	CAPSULE	Strength(s): 25mg, 50mg
		ELIXIR	12.5mg/5ml
	Manufacturers:	HAL, ICN, P/D	
	Category:	OTC	
DISULFIRAM			
	Dosage form(s):	TABLET	Strength(s): 250mg, 500mg
	Manufacturers:	W-A, DAN	
	Category:	B	
EPINEPHRINE HCl			
	Dosage form(s):	INJECTION	Strength(s): 0.01%, 0.1%
		OPHTHALMIC SOLUTION	0.1%, 0.25%, 0.5%, 1%, 2%
	Manufacturers:	ABB, ALC, ALL, ARL, ESR,IMS, INV, LAN, P/D, W-A	
	Category:	PRE-38	
ESTROGENS, ESTERIFIED			
	Dosage form(s):	TABLET	Strength(s): 0.3mg, 0.625mg, 1.25mg, 2.5mg
	Manufacturers:	BMS, PRI, SLV, SKB, SYN	
	Category:	B	
ETHAVERINE HCl			
	Dosage form(s):	CAPSULE	Strength(s): 100mg
		TABLET	
	Manufacturers:	BFA, KEN, LEM, MEP	
	Category:	PRE-38	
ETHINYL ESTRADIOL			
	Dosage form(s):	TABLET	Strength(s): 0.02mg, 0.05mg
	Manufacturers:	ORI, SCH, P/U	
	Category:	B	

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MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

FLUOXYMESTERONE		
Dosage form(s):	TABLET	Strength(s): 2mg, 5mg, 10mg
Manufacturers:	BMS, RPC, P/U	
Category:	B	
GLYBURIDE		
Dosage form(s):	TABLET	Strength(s): 1.25mg, 2.5mg, 5mg
Manufacturers:	HOE, P/U	
Category:	B	
HYDRALAZINE HCl, HYDROCHLOROTHIAZIDE, RESERPINE		
Dosage form(s):	TABLET	Strength(s): 25mg/15mg/0.1mg
Manufacturers:	DAN, GEI, LEM	
Category:	B	
HYDROCHLOROTHIAZIDE W/ RESERPINE		
Dosage form(s):	TABLET	Strength(s): 25mg/0.125mg, 50mg/0.125mg, 25mg/0.1mg, 50mg/0.1mg
Manufacturers:	KNO, CAM, GEI, GEV, DAN, LEM, MCK, PUR, ZGP	
Category:	B	
HYDROCODONE BITARTRATE W/ PHENYLPROPANOLAMINE		
Dosage form(s):	SYRUP	Strength(s): 5mg/25mg/5ml
Manufacturers:	DPT, ALP, RPC	
Category:	OTHER	
HYDROCORTISONE, IODOCHLORHYDROXYQUIN		
Dosage form(s):	CREAM OINTMENT	Strength(s): 0.5%, 1% / 3%
Manufacturers:	AMB, ALT, CLA, DER, BAY, DUR, GEI, LEM, ALP, SLV, THA,	
Category:	DESI	
HYDROFLUMETHIAZIDE, RESERPINE		
Dosage form(s):	TABLET	Strength(s): 25mg, 50mg/0.125mg
Manufacturers:	APO, RPC, ZGP	
Category:	B	
HYDROQUINONE 4% CREAM		
Dosage form(s):	TOPICAL CREAM	Strength(s): 4%
Manufacturers:	ICN, ETX	
Category:	PRE-62	
HYDROQUINONE CREAM 4%		
Dosage form(s):	CREAM	Strength(s): 4%
Manufacturers:	ICN; GLD	
Category:	PRE-1962	
HYDROQUINONE 4% CREAM with SUNCREENS		
Dosage form(s):	TOPICAL CREAM	Strength(s): 4%
Manufacturers:	ICN, ETX	
Category:	PRE-62	
HYDROQUINONE CREAM 4% with SUNCREEN		
Dosage form(s):	CREAM	Strength(s): 4%
Manufacturers:	ICN; GLD	
Category:	PRE-1962	
HYDROQUINONE TOPICAL SOLUTION		
Dosage form(s):	TOPICAL SOLUTION	Strength(s): 3%
Manufacturers:	NEU, GLD	
Category:	PRE-38	

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MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

ISOSORBIDE DINITRATE			
	Dosage form(s):	SUSTAINED RELEASE CAPSULE	Strength(s): 40mg
	Manufacturers:	ASC, GEV, FOR, W-A, SUP	
	Category:	B	
ISOXSUPRINE HCl			
	Dosage form(s):	TABLET	Strength(s): 10mg, 20mg
	Manufacturers:	GEV, MEA	
	Category:	PRE-38	
L-HYOSCYAMINE SULFATE			
	Dosage form(s):	TABLET	Strength(s): 0.125mg
	Manufacturers:	BFA, GLW, SWZ	
	Category:	PRE-38	
LEVODOPA			
	Dosage form(s):	TABLET	Strength(s): 250mg, 500mg
		CAPSULE	100, 250, 500mg
	Manufacturers:	ROC, P&G	
	Category:	B	
MAGNESIUM SALICYLATE			
	Dosage form(s):	TABLET	Strength(s): 600mg
	Manufacturers:	BFA, END, RBP, MLP, ,	
	Category:	OTHER	
MAGNESIUM SULFATE			
	Dosage form(s):	INJECTION	Strength(s): 10%, 12.5%, 50%
	Manufacturers:	ABB, APP, ARL, CMC, ESR, IMS, LIL	
	Category:	PRE-38	
MAZINDOL			
	Dosage form(s):	TABLET	Strength(s): 1mg
	Manufacturers:	SAN, W-A	
	Category:	B	
MEPHOBARBITAL			
	Dosage form(s):	TABLET	Strength(s): 32mg, 100mg, 200mg
	Manufacturers:	BOW, ICN, S/W	
	Category:	PRE-38	
METHENAMINE MANDELATE			
	Dosage form(s):	SUSPENSION	Strength(s): 0.25, 0.5g/5ml
		TABLET	0.25g, 0.5g, 1g
		ENTERIC COATED TABLET	0.25g, 0.5g, 1g
	Manufacturers:	GEV, HEA, ALP, P/D, RIC, SLV, TAB	
	Category:	PRE-38	
METHENAMINE COMBINATION (METHENAMINE, PHENYLSALICYLATE, ATROPINE SULFATE, HYOSCYAMINE, BENZOIC ACID, METHYLENE BLUE)			
	Dosage form(s):	TABLET	Strength(s): 40.8mg/18mg/0.03mg/ 0.03mg/4.5mg/5.4mg
	Manufacturers:	CHE, LEM, , S-M, STR	
	Category:	DESI	
METHYLENE BLUE			
	Dosage form(s):	INJECTION	Strength(s): 1%
		TABLET	
	Manufacturers:	ARL, CMC, ESR, TAY,	
	Category:	PRE-38	

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MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

METHYLTESTOSTERONE			
	Dosage form(s):	CAPSULE	Strength(s): 10mg
		SUBL. TABLET	10mg, 25mg
	Manufacturers:	DAN, INW, LAN, PUR, SCH	
	Category:	B	
MORPHINE SULFATE			
	Dosage form(s):	SUSTAINED RELEASE TABLET	Strength(s): 30mg, 60mg, 100mg
	Manufacturers:	PUF, ROX	
	Category:	B	
NEOSTIGMINE METHYLSULFATE			
	Dosage form(s):	INJECTION	Strength(s): 1-1000, 1-2000, 1-4000
	Manufacturers:	CMC, ESR, LAN, S-M,	
	Category:	PRE-38	
NITROGLYCERIN			
	Dosage form(s):	SUBLINGUAL TABLET	Strength(s): 0.3mg, 0.4mg, 0.6mg
	Manufacturers:	P/D ETX	
	Category:	PRE-38	
NORTRIPTYLINE HCl			
	Dosage form(s):	CAPSULE	Strength(s): 10mg, 25mg
	Manufacturers:	LIL, SAN	
	Category:	B	
NYLIDRIN			
	Dosage form(s):	TABLET	Strength(s): 6mg, 12mg
	Manufacturers:	GEV, C/P, DAN, ROR, ZGP	
	Category:	DESI	
OPIUM TINCTURE, DEODORIZED			
	Dosage form(s):	LIQUID	Strength(s): 10% OPIUM
	Manufacturers:	HAL, LIL	
	Category:	PRE-38	
PAPAVERINE HCl (NON-SUSTAINED RELEASE)			
	Dosage form(s):	INJECTION	Strength(s): 30mg/ml
		CAPSULE	75mg, 150mg, 300mg
		TABLET	30mg, 60mg, 100mg, 150mg, 200mg, 300mg
	Manufacturers:	CHE, CMC, GEV, DAN, HAL, HEA, LAN, LEM, MMD, MYL, PUR, REN, SLV, VAN, EON, ZGP	
	Category:	PRE-38	
PARALDEHYDE			
	Dosage form(s):	LIQUID	Strength(s): 100%
		INJECTION	
	Manufacturers:	CMC, ESR,	
	Category:	PRE-38	
PAREGORIC			
	Dosage form(s):	LIQUID	Strength(s): 2mg MORPHINE EQUIV./5ml
	Manufacturers:	APC, BOW, HAL, LAN, LIL, ALP, P/D, PUR, ROX, RPC, STL	
	Category:	OTC	
PENICILLIN G BENZATHINE			
	Dosage form(s):	INJECTION	Strength(s): 600, 000 UNITS/ml
	Manufacturers:	PFI, W-A	
	Category:	B	
PENTAERYTHRITOL TETRANITRATE			
	Dosage form(s):	TABLET	Strength(s): 10, 20
		SUSTAINED ACTION TABLET	80mg
	Manufacturers:	COO, GEV, DAN, INW, KIR, MER, P/D, PUR, STA, S-M, ZGP	
	Category:	DESI	

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MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

PHENAZOPYRIDINE HCl		
Dosage form(s):	TABLET	Strength(s): 100mg, 200mg
Manufacturers:	AMI, BAR, COP, C-P, LAN, P/D, QUA, RIC, S-M, TAB, VAN,	
Category:	PRE-38	
PHENOBARBITAL		
Dosage form(s):	ELIXIR	Strength(s): 20mg/5ml
	TABLET	15mg, 16mg, 30mg, 32mg, 60mg, 65mg, 100mg
Manufacturers:	APC, BAR, BOW, GEV, C/P, DAN, HAL, ICN, INW, LAN, LED, LEM, LIL, MMD, ALP, P/D, PUR, REX, ROX, RUG, STL, STA, TAB, EON, W-W, S/W, W-A, ZGP	
Category:	PRE-38	
PHENYLEPHRINE HCl		
Dosage form(s):	SOLUTION	Strength(s): 0.25%, 1%
	OPHTHALMIC SOLUTION	0.12, 2.5, 10%
Manufacturers:	AKO, ALC, ALL, ALP, MUR, B&L, PUF, RPC, STS, S/W	
Category:	PRE-38	
PHENYLPROPANOLAMINE HCl, PHENYLEPHRINE HCl, PHENYLTOLOXAMINE CITRATE, CHLORPHENIRAMINE MALEATE		
Dosage form(s):	PEDIATRIC DROPS	Strength(s): 5mg/1.25mg/2mg/0.5mg/ml
	PEDIATRIC SYRUP	5mg/1.25mg/2mg/0.5mg/5ml
Manufacturers:	ALP, APO	
Category:	PRE-38	
PHYTONADIONE		
Dosage form(s):	INJECTION	Strength(s): 2mg, 10mg/ml
Manufacturers:	ABB, ROC, IMS, MCK, SKB	
Category:	B	
PILOCARPINE HCl		
Dosage form(s):	OPHTHALMIC SOLUTION	Strength(s): 0.25%, 0.5%, 1%, 2%, 3%, 4%, 6%, 8%
Manufacturers:	ALC, CVO, W-A, OPT, B&L, PRO, STS	
Category:	PRE-38	
PIPERAZINE CITRATE		
Dosage form(s):	SYRUP	Strength(s): 500mg/5ml
	TABLET	250mg, 500mg
Manufacturers:	BLU,ALP, GLW, LAN, SLV, S/W	
Category:	PRE-38	
POTASSIUM GLUCONATE		
Dosage form(s):	LIQUID	Strength(s): 20mEq/15ml
Manufacturers:	ALP, CMC, GEV, LAN, LED, , RAH, ROX, RPC, S-M,	
Category:	B	
POTASSIUM IODIDE		
Dosage form(s):	LIQUID	Strength(s): SATURATED SOLUTION
Manufacturers:	ALP, CMC, GEN, GEV, P/R, ROX, RPC, STL, , USL	
Category:	PRE-38	
PREDNISOLONE ACETATE		
Dosage form(s):	INJECTION	Strength(s): 25, 50, 100mg/ml
Manufacturers:	ALC, ALL, CTL, STS, LEM, B&L, SCH	
Category:	B	
PREDNISOLONE TEBUTATE		
Dosage form(s):	INJECTION	Strength(s): 20mg/ml
Manufacturers:	ALP, FOY, RBP, MCK	
Category:	B	
PROBENECID W/ COLCHICINE		
Dosage form(s):	TABLET	Strength(s): 500mg/0.5mg
Manufacturers:	DAN, LEM, MCK, RIC, ZGP	
Category:	B	

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MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

PROMETHAZINE HCl			
	Dosage form(s):	TABLET	Strength(s): 12.5mg, 25mg, 50mg
	Manufacturers:	ALL, ALT, ALP, DAN, ESR, GEV, KNO, KVN, LEM, LIF, MSM, RPC, S/W, W-A	
	Category:	B	
PROPYLTHIOURACIL			
	Dosage form(s):	TABLET	Strength(s): 50mg
	Manufacturers:	ANA, KNO, CHE, DAN, LIL, HAL, LAN, LED, P/D, PUR, RIC, TAB, W-W, ZEN	
	Category:	B	
PSEUDOEPHEDRINE HCl			
	Dosage form(s):	LIQUID	Strength(s): 30mg/5ml
		TABLET	30mg, 60mg
		SUSTAINED RELEASE CAPSULE	120mg
	Manufacturers:	ALP, BOW, GLW, CEN, CHE, GEV, DAN, HAL, RBP, LEM, MMD, PAR, ROX, SLV, SUP	
	Category:	OTC	
PSEUDOEPHEDRINE HCl, CHLORPHENIRAMINE MALEATE			
	Dosage form(s):	TABLET	Strength(s): 60mg/4mg
	Manufacturers:	BER, GLW, ROR, SCH, SKB	
	Category:	OTC	
PSEUDOEPHEDRINE SULFATE, DEXBROMPHENIRAMINE MALEATE			
	Dosage form(s):	SUSTAINED RELEASE TABLET	Strength(s): 120mg/6mg
	Manufacturers:	COP, GEV, SCH	
	Category:	OTC	
QUININE SULFATE			
	Dosage form(s):	TABLET	Strength(s): 260mg
	Manufacturers:	CHE, GEV, MMD	
	Category:	PRE-38	
RAUWOLFIA SERPENTINA			
	Dosage form(s):	TABLET	Strength(s): 50mg, 100mg
	Manufacturers:	BOW, CMB, GEV, DAN, FER, HAL, ICN, KIR, PAN, PPI, PRI, BMS, SLV, RIC, TAB, VAL, ZEN	
	Category:	B	
RESERPINE			
	Dosage form(s):	TABLET	Strength(s): 0.1mg, 0.25mg, 0.5mg, 1.0mg
		INJECTION	
	Manufacturers:	BEL, BMS, BOW, CMB, GEV, DAN, FER, GEI, GEN, HAL, ICN, , KIR, LAN, LEM, LIL, MKL, MYL, PAN, PRV, PRI, PUR, SLV, REX, RIC, ROX, STL, TAB, P/U, VAL, ZEN	
	Category:	B	
SALSALATE			
	Dosage form(s):	TABLET	Strength(s): 500mg, 750mg
	Manufacturers:	GEV, 3MP, SID	
	Category:	Pre-38	
SODIUM FLUORIDE			
	Dosage form(s):	TABLET	Strength(s): 0.55mg, 1.1mg, 2.2mg (NaF)
		DROPS	0.125mg (F-)/DROP
		CHEWABLE TABLET	0.55mg, 1.1mg, 2.2mg (NaF)
	Manufacturers:	ABL, KNO, BOW, CHE, CMC, COP, C-P, GEN, HOY, KIR, RIC, RUG, STL,	
	Category:	PRE-38	
SULFAMETHOXAZOLE, PHENAZOPYRIDINE HCl			
	Dosage form(s):	TABLET	Strength(s): 500/100mg
	Manufacturers:	COP, RIC	
	Category:	DESI	
SULFASALAZINE			
	Dosage form(s):	ENTERIC COATED TABLETS	Strength(s): 500mg
	Manufacturers:	DAN, LED, LEM, MUT, KBP, ROW, SUP, VIP	
	Category:	B	

720.200: continued

MASSACHUSETTS ADDITIONAL LIST OF INTERCHANGEABLE DRUGS

SULFISOXAZOLE PHENAZOPYRIDINE HCl		
Dosage form(s):	TABLET	Strength(s): 500/50mg
Manufacturers:	COP, KAY, ROC, ROX, SLV, S-M, STN, VAN,	
Category:	DESI	
TERBUTALINE SULFATE		
Dosage form(s):	TABLET	Strength(s): 2.5mg, 5mg
Manufacturers:	GEI, MMD	
Category:	B	
TESTOSTERONE		
Dosage form(s):	INJECTION	Strength(s): 25mg/ml, 50mg/ml, 100mg/ml
Manufacturers:	BAT, STS,RBP, LIL, MAY	
Category:	PRE-38	
TETRACAINE HCl		
Dosage form(s):	OPHTHALMIC SOLUTION	Strength(s): 0.5%
Manufacturers:	ALC, GEN, B&L, S-M, S/W	
Category:	PRE-38	
THEOPHYLLINE (NON-SUSTAINED RELEASE)		
Dosage form(s):	CAPSULE	Strength(s): 100mg, 200mg
	TABLET	100, 125, 200, 225, 250mg
Manufacturers:	ALP, BEL, BER, CNC, CTL, FER, HAL, KNO, LAN, LIF, MMD, PAN, P/A, RIC, 3MP, ROR, ROX, RPC, SEA,	
Category:	B	
THEOPHYLLINE, GUAIFENESIN		
Dosage form(s):	ELIXIR	Strength(s): 150mg/90mg/15ml
	LIQUID	
Manufacturers:	MEA, RPC	
Category:	PRE-38	
THEOPHYLLINE, POTASSIUM IODIDE (THEOPHYLLINE, POTASSIUM IODIDE, ALCOHOL)		
Dosage form(s):	ELIXIR	Strength(s): 80mg/130mg/10%/15ml
Manufacturers:	FOR, ALP, RPC	
Category:	PRE-38	
THYROGLOBULIN		
Dosage form(s):	TABLET	Strength(s): 65mg
Manufacturers:	RIC, P/D, W-L	
Category:	B	
TRIAMCINOLONE DIACETATE		
Dosage form(s):	INJECTION	Strength(s): 25mg, 40mg/ml
Manufacturers:	BMS, LED, LEM, STS	
Category:	B	
TRICHLORMETHIAZIDE, RESERPINE		
Dosage form(s):	TABLET	Strength(s): 4mg/0.1mg
Manufacturers:	MMD, SCH	
Category:	B	
TRIPROLIDINE HCl, PSEUDOEPHEDRINE HCl		
Dosage form(s):	TABLET	Strength(s): 2.5mg/60mg
	SYRUP	1.25mg/30mg/5ml
Manufacturers:	GLW, ALP, CHE, CNC, GEV, HAL, ICN, LED, LEM, LIF, NEW, B&L, PRI, PUR, ROX, SLV, SUP, VAN, ZEN	
Category:	OTC	
TRYPSIN, BALSAM PERU, CASTOR OIL		
Dosage form(s):	TOPICAL AEROSOL	Strength(s): 0.1mg/72.5mg/650mg IN EACH 0.82CC SPRAY
Manufacturers:	COP, HIC	
Category:	PRE-38	

720.200: continued

PARTIAL PROPRIETARY BRAND CROSS-REFERENCE

Generically equivalent drug products in the same strength and dosage form listed in the *Additional List* are interchangeable if their respective manufacturers are listed for that product. This partial cross-reference section does not attempt to list all brand names which are approved for interchange. For most products only one, usually the innovator or most commonly prescribed brand, is listed below for quick reference purposes.

See page 4105 for precise instructions for determining the interchangeability of drug products.

BRAND	SEE
	(ATROPINE SULFATE, HYOSCINE HBr, HYOSCYAMINE HBr, PHENOBARBITAL)
	(THEOPHYLLINE, POTASSIUM IODIDE, ALCOHOL)
ACTIFED	TRIPROLIDINE HCl, PSEUDOEPHEDRINE HCl
ANDROID	METHYLTESTOSTERONE
	ANTABUSE DISULFIRAM
AQUAMEPHYTON	PHYTONADIONE
AURALGAN	BENZOCAINE, ANTIPYRINE
AZO-GANTANOL	SULFAMETHOXAZOLE, PHENAZOPYRIDINE
AZO-GANTRISIN	SULFISOXAZOLE PHENAZOPYRIDINE HCl
AZULFADINE	SULFASALAZINE
BENADRYL	DIPHENHYDRAMINE HCl
	CYANOCOBALAMIN
BICILLIN	PENICILLIN G BENZATHINE
BRETHINE	TERBUTALINE SULFATE
CHLOR-TRIMETON	CHLORPHENIRAMINE MALEATE
COL-BENEMID	PROBENECID W/ COLCHICINE
	CYCLANDELATE
DECADRON	DEXAMETHASONE
DESQUAM	BENZOYL PEROXIDE
DIMETANE DX	BROMPHENIRAMINE, DEXTROMETHORPHAN, PSEUDOEPHEDRINE
DIUPRESS	CHLOROTHIAZIDE W/ RESERPINE
DONNATAL	ATROPINE SULFATE COMPOUND
DRAMAMINE	DIMENHYDRINATE
DRIXORAL	(product reformulated)
ECONOPRED	PREDNISOLONE ACETATE
ELIXOPHYLLINE KI	THEOPHYLLINE, POTASSIUM IODIDE
EMPIRIN W/ CODEINE	ASPIRIN W/ CODEINE PHOSPHATE
ESTINYL	ETHINYL ESTRADIOL
EXNA	BENZTHIAZIDE
FEDAHIST	PSEUDOEPHEDRINE HCl, CHLORPHENIRAMINE MALEATE
GLAUCON	EPINEPHRINE HCl
GRANULEX	TRYPSIN, BALSAM PERU, CASTOR OIL
HALOTESTIN	FLUOX\YMESTERONE
HYCOMINE	HYDROCODONE BITARTRATE W/ PHENYLPROPANOLAMINE
HYDROPRES	HYDROCHLOROTHIAZIDE W/ RESERPINE
ISOPTO CARPINE	PILOCARPINE HCl
ISORDIL TEMBID	ISOSORBIDE DINITRATE
ISUPREL	ISOPROTERENOL HCl
KAON	POTASSIUM GLUCONATE
KENALOG	TRIAMCINOLONE DIACETATE
LARODOPA	LEVODOPA
LEVSIN	L-HYOSCYAMINE SULFATE
LIBRAX	CHLORDIAZEPOXIDE W/CLIDINIUM BROMIDE
MANDELAMINE	METHENAMINE MANDELATE
MEBARAL	MEPHOBARBITAL
MENEST	ESTROGENS, ESTERIFIED
METALONE T.B.A	PREDNISOLONE TEBUTATE
MICRONASE	GLYBURIDE
	MAGNESIUM SALICYLATE
NEO-SYNEPHRINE	PHENYLEPHRINE HCl

720.200: continued

BRAND	SEE
	CHLORAL HYDRATE
PAMELOR	NORTRIPTYLINE HCl
PHENERGAN PLAIN	PROMETHAZINE HCl
PONTOCAINE	TETRACAINE HCl
PROSED	METHENAMINE COMBINATION (METHENAMINE, PHENYLSALICYLATE, ATROPINE SULFATE, HYOSCYAMINE, BENZOIC ACID METHYLENE BLUE)
PROSTIGMINE	NEOSTIGMINE METHYLSULFATE
PYRIDIUM	PHENAZOPYRIDINE HCl
QUINAMM	QUININE SULFATE
SALUTENSIN	HYDROFLUMETHIAZIDE, RESERPINE
SANOREX	MAZINDOL
SERAPES	HYDRALAZINE HCl, HYDROCHLOROTHIAZIDE, RESERPINE
SLO-PHYLLIN GG	THEOPHYLLINE, GUAIFENESIN
SLO-PHYLLIN	THEOPHYLLINE (NON-SUSTAINED RELEASE)
SSKI	POTASSIUM IODIDE
SUDAFED	PSEUDOEPHEDRINE HCl
SYNTHROID	LEVOTHYROXINE SODIUM
TENUATE	DIETHYLPROPION HCl
	TESTOSTERONE
TUINAL	AMOBARBITAL SODIUM, SECOBARBITAL SODIUM
VASODILAN	ISOXSUPRINE

(PAGES 4127 THROUGH 4132 ARE RESERVED FOR FUTURE USE.)