



The Commonwealth of Massachusetts
Executive Office of Health and Human Services
Department of Public Health
Board of Registration in Pharmacy
Bureau of Health Professions Licensure
250 Washington Street, Boston, MA 02108-4619

Tel: 617-973-0800
TTY : 617-973-0988
www.mass.gov/dph/boards

TO: Pharmacist Licensure Applicants

FROM: Massachusetts Board of Registration in Pharmacy

RE: Multistate Pharmacy Jurisprudence Examination ("MPJE®")

The Board of Registration in Pharmacy ("Board") advises [applicants](#) for licensure as a pharmacist in the Commonwealth of Massachusetts to review the reference sources listed below to prepare for the [Multistate Pharmacy Jurisprudence Examination \("MPJE®"\)](#).

You may view websites as below or obtain copies of 247 CMR and 105 CMR 700.00 - 722.00 for a fee by requesting the "Pharmacy Package" at this link:

<https://www.sec.state.ma.us/spr/sprcat/catidx.htm>

Note: The Executive Office of Administration and Finance regulations 801 CMR Sections 1.00 – 3.00 that are included in this package are **not** included on the MPJE®.

Board Regulations - 247 CMR:

<https://www.mass.gov/law-library/247-cmr>

Board Policies:

Policies are adopted in order to provide clarification to statutes and/or regulations. Circular letters are similar to Board policies but are published by the Drug Control Program. Both policies and circular letters are enforceable and are available here:

<https://www.mass.gov/lists/policies-and-guidelines-of-the-board-of-registration-in-pharmacy>

Department of Public Health Regulations - 105 CMR §§ 700.00 - 722.00:

These regulations pertain to hospital pharmacies, mandated interchange, product labeling, prescription format and security standards:

<https://www.mass.gov/law-library/105-cmr>

Prescription Monitoring Program (“PMP”) Reporting Requirements:

<https://www.mass.gov/service-details/pharmacy-reporting-and-data-submission>

Massachusetts General Laws (M.G.L.) Chapters 13, 112 and 94C:

- M.G.L. c. 13 §§ 22 – 25A
<https://malegislature.gov/Laws/GeneralLaws/PartI/TitleII/Chapter13>
- M.G.L. c. 112 §§ 24 – 42D
<https://malegislature.gov/Laws/GeneralLaws/PartI/TitleXVI/Chapter112>
- M.G.L. c. 94C §§ 1 - 40 (Massachusetts Controlled Substances Act)
<https://malegislature.gov/Laws/GeneralLaws/PartI/TitleXV/Chapter94C>

Board of Registration in Medicine “Prescribing Practices: Policies & Guidelines”:

<http://www.mass.gov/eohhs/docs/borim/policies-guidelines/policy-15-05.pdf>

Federal Food, Drug, and Cosmetic Act (FDCA):

<https://www.ecfr.gov/current/title-21>

Federal Regulations (“CFR”) Part 1300 – 1321:

<https://www.ecfr.gov/current/title-21/chapter-II>

Drug Enforcement Administration (“DEA”) - Pharmacist’s Manual:

[https://www.deadiversion.usdoj.gov/GDP/\(DEA-DC-046\)\(EO-DEA154\)_Pharmacist_Manual.pdf](https://www.deadiversion.usdoj.gov/GDP/(DEA-DC-046)(EO-DEA154)_Pharmacist_Manual.pdf)

Poison Prevention Packaging Act:

<https://www.ecfr.gov/current/title-16/chapter-II/subchapter-E>

Prescription Drug Marketing Act:

<https://www.ecfr.gov/current/title-21/chapter-I/subchapter-C/part-203>

Medication Guides and Patient Package Inserts:

- <https://www.ecfr.gov/current/title-21/chapter-I/subchapter-C/part-208?toc=1>
- <https://www.ecfr.gov/current/title-21/chapter-I/subchapter-D/part-310/subpart-E/section-310.501>
- <https://www.ecfr.gov/current/title-21/chapter-I/subchapter-D/part-310/subpart-E/section-310.515>

Recalls:

<https://www.ecfr.gov/current/title-21/chapter-I/subchapter-A/part-7/subpart-C>

Misbranding:

<https://www.fda.gov/medical-devices/general-device-labeling-requirements/labeling-requirements-misbranding#misbrand>

Adulterated Drugs:

<https://uscode.house.gov/view.xhtml?path=/prelim@title21/chapter9/subchapter5&edition=prelim>

NABP Pharmacist Resources:

<https://nabp.pharmacy/resources/>

USP:

<https://www.usp.org/>