



MPJE Domain Crosswalk

Competency Statements to Revised Content Outline

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Introduction

Following the completion of the Multistate Pharmacy Jurisprudence Examination® (MPJE®) Practice Analysis, initiated in February 2026, the National Association of Boards of Pharmacy® (NABP®) undertook a comprehensive crosswalk activity to align the current MPJE competency statements with the updated (2027) content outline. This process involved leveraging a variety of expertise, both internally and from the pharmacy field, to systematically compare and map each competency area in the existing framework to one or more subdomains in the new content outline. By doing so, NABP's Exam Services team ensured a clear correlation between the established competency statements and the updated content outline.

The crosswalk illustrates how the current MPJE competency framework aligns with and supports the forthcoming updates to the exam. Every competency statement is associated with one or more subdomains in the upcoming 2027 content outline.

The updated MPJE Content Outline breaks down the overarching competency statements into specific content areas (domains, subdomains, and sub-subdomains). These subdomains ensure that candidates' competencies are assessed comprehensively by targeting knowledge, skills, and abilities. Domains, on the other hand, organize competency areas into specific content categories, grouping related tasks and responsibilities to systematically structure the exam framework.

Example:

1. **Pharmacy and Pharmacist Practice** (Domain)
 - B. Pharmacy Personnel (Subdomain)
 1. Licensures, registrations, and certifications (Sub-subdomain)

The following crosswalk table compares the current competency statements to the updated (2027) content outline.

Crosswalk Table

Original Competency & Sub-Competency	New Domain & Sub-Domain	Crosswalk Notes
1.1	1B2, 1B3	
Area 1: Licensure/Personnel 1.1: Pharmacist & non-pharmacist responsibilities (scope of duties, limitations, conditions to practice)	Domain 1: Pharmacy and Pharmacist Practice 1B: Pharmacy personnel – scope of practice & supervision (1.B.2-3)	Scope of duties maps to 1B2 (scope of practice) and 1B3 (supervision)
1.2	1B1, 1B4, 3C	
Area 1: Licensure/Personnel 1.2: Licensures, registrations, & certifications (qualifications, examinations, internships, renewals)	Domain 1: Pharmacy and Pharmacist Practice 1B1: Licensures, registrations, or certifications 1B4: CE	- Renewals & maintaining competency map to 1B1 - CE requirements map to 1B4
Area 1: Licensure/Personnel 1.2: Disciplinary actions & reporting impairment programs	Domain 3: Regulatory Authority and Legal Obligations 3C: Liability, malpractice, duty to report, & unprofessional conduct	Inability-to-practice reporting programs moved to Regulatory Authority domain
2.1	1A, 2A1, 2A2, 2A3	
Area 2: Pharmacist Practice 2.1: Requirements for issuing prescriptions/drug orders (non-controlled & controlled)	Domain 2: Medication Use Process 2A: Requirements for issuing prescriptions & medication orders (2A1-3)	Validity, changes, emergency prescriptions all captured under 2A sub-items
Area 2: Pharmacist Practice 2.1: Scope of authority & valid registration of prescribing practitioners	Domain 1: Pharmacy and Pharmacist Practice 1A: Practice of pharmacy – scope of authority, collaborative practice agreements, & delegative authorities	New outline explicitly names collaborative practice agreements; delegative authorities added

2.2	2C	
Area 2: Pharmacist Practice 2.2: Conditions for pharmacist participation in administration of drugs/patient therapy management	Domain 2: Medication Use Process 2C: Administration of medications, including training & adverse event reporting	- Adverse event reporting is newly explicit content - Training requirement now named
2.3	2B5	
Area 2: Pharmacist Practice 2.3: Counseling requirements (offering to counsel, documenting counseling)	Domain 2: Medication Use Process 2B5: Counseling	Documentation of counseling subsumed within 2B dispensing requirements
2.4	2B8	
Area 2: Pharmacist Practice 2.4: Returning or reusing drugs	Domain 2: Medication Use Process 2B8: Recalls, returns, & reuse (reverse distribution, charitable pharmacy, drug repository)	- Scope expanded - Specific channels (charitable pharmacy, drug repository) now named
2.5	1D, 1F, 3A1, 3A2	
Area 2: Pharmacist Practice 2.5: Regulations & agencies – promoting quality & safety of public health	Domain 1: Pharmacy and Pharmacist Practice 1D: Regulatory standards (NIOSH, OSHA, FDA, BOP, USP standards)	- Specific agencies now named - Agency authority/discipline roles also appear in 3A
Area 2: Pharmacist Practice 2.5: Protecting patient & health record confidentiality	Domain 1: Pharmacy and Pharmacist Practice 1F: Security of patient health information	Cybersecurity angle also covered in Domain 4 – 4B (pharmacy technology)
3.1	2A1	
Area 2: Pharmacist Practice 3.1: Determining legitimate medical purpose; corresponding responsibility	Domain 2: Medication Use Process 2A1: Validity of a prescription (authority, content, corresponding responsibility)	Corresponding responsibility now explicit sub-item; also supported by 3B record keeping

3.2	2B6	
Area 3: Dispensing Requirements 3.2: Transferring prescription/drug order information between pharmacies by authorized personnel	Domain 2: Medication Use Process 2B6: Transfers	Authorized personnel requirement still implied within 2B dispensing requirements
3.3	2B2, 2B7	
Area 3: Dispensing Requirements 3.3: Prospective drug utilization reviews & PMP reporting/access	Domain 2: Medication Use Process 2B2: Drug utilization review 2B7: PDMP requirements	- PMP terminology updated to PDMP - Reporting & access requirements retained
3.4	2A3, 2B3	
Area 3: Dispensing Requirements 3.4: Exceptions to dispensing or refilling prescriptions/drug orders	Domain 2: Medication Use Process 2A3: Emergency prescriptions & emergency refills 2B3: Fills, refills, & partial fills	- Partial fills now explicit sub-item - Emergency dispensing separated to 2A
3.5	2B1	
Area 3: Dispensing Requirements 3.5: Labeling of dispensed drugs	Domain 2: Medication Use Process 2B1: Pharmacy & manufacturer labeling, including OTC medications & patient education	OTC labeling & patient education material now explicitly named
3.6	2B4	
Area 3: Dispensing Requirements 3.6: Packaging of dispensed drugs	Domain 2: Medication Use Process 2B4: Packaging & repackaging	Repackaging added; also touches Domain 4 – 4C access & handling
3.7	1C, 1D, 1E, 2B8	
Area 3: Dispensing Requirements 3.7: Drug product conditions prohibiting dispensing	Domain 1: Pharmacy and Pharmacist Practice 1C: Adulterated or misbranded medications (recalls, expiration dates, counterfeit, EUA, importation, DSCSA)	- DSCSA & emergency use authorizations newly named - Importation added

3.8	1D, 1E, 2B1, 2B4, 2B9, 1E, 2B, 2C	
Area 3: Dispensing Requirements 3.8: Non-prescription products – dispensing, labeling, packaging; dispensing-restricted OTC drugs	Domain 2: Medication Use Process 2B1: Labeling (OTC) 2B4: Packaging 2B9: Drug product selection (substitutions, interchangeability)	- Hazardous drug handling moves to Domain 1 – 1E - Substitutions now explicit
Area 3: Dispensing Requirements 3.8: Hazardous drugs – dispensing, handling, labeling, packaging	Domain 1: Pharmacy and Pharmacist Practice 1D & 1E: State-specific standards on compounding & hazardous materials (BUD, competency training, environmental testing, record keeping)	- BUD, competency training, & environmental testing now explicit - NIOSH/OSHA in 1D
4.1	4C	
Area 4: Pharmacy Operations 4.1: Ordering, acquisition, & distribution of drugs – records maintenance & content	Domain 4: Pharmacy Operations 4C: Access, storage, handling, & security (facility, drugs, records)	- Distribution records retained - Acquisition & ordering records subsumed in 4C
4.2	3B, 3B1, 3B2, 3B3, 3B4, 3B5, 3B6, 4C	
Area 4: Pharmacy Operations 4.2: Record keeping – non-dispensing operations, storage, handling, hazardous drugs	Domain 3: Regulatory Authority and Legal Obligations 3B: CS – record keeping, inventory, order forms	- CS-specific records moved to Regulatory Domain 3B - General ops records remain in Domain 4
Area 4: Pharmacy Operations 4.2: CS inventory requirements	Domain 3: Regulatory Authority and Legal Obligations 3B1: Inventory 3B2: Record keeping 3B3: Order forms & ordering 3B4: DEA registration & power of attorney	DEA registration & power of attorney now explicit sub-items; not previously named
Area 4: Pharmacy Operations 4.2: Handling, storage, & disposal of hazardous drugs	Domain 3: Regulatory Authority and Legal Obligations 3B5: Theft & loss 3B6: Disposal	- Theft/loss reporting now a distinct sub-item - Disposal separated from general handling

4.3	4B, 4D	
Area 4: Pharmacy Operations	Domain 4: Pharmacy Operations	Devices explicitly added to scope of delivery requirements
4.3: Delivery of drugs	4D: Delivery of drugs & devices	
4.4	2B9	
Area 4: Pharmacy Operations	Domain 2: Medication Use Process	- Moved from Operations to Medication Use Process - Reflects clinical nature of selection decisions
4.4: Conditions for permitted or mandated product selection	2B9: Drug product selection (substitutions, interchangeability)	
4.5	1D, 1E	
Area 4: Pharmacy Operations	Domain 1: Pharmacy and Pharmacist Practice	- USP standards also in 1D - Compounding elevated from ops to practice standards
4.5: Compounding – sterile, nonsterile, hazardous, & non-hazardous preparations	1E: State-specific standards on compounding & hazardous materials	
4.6	4B	
Area 4: Pharmacy Operations	Domain 4: Pharmacy Operations	- Central-fill remapped to “shared services” - Technology & cyber security now explicit
4.6: Centralized prescription processing/central-fill pharmacy dispensing	4B: Pharmacy technology – record keeping, automation, e-prescribing, faxing, shared services	
4.7	4A, 3A1, 3A2	
Area 4: Pharmacy Operations	Domain 4: Pharmacy Operations	Disciplinary actions against practice settings moved to Domain 3 – 3A
4.7: Registration, licensure, certification, or permitting of practice setting (requirements, renewal, inspection)	4A: Requirements for registration, license, certification, or permitting of a practice setting or business entity	
Area 4: Pharmacy Operations	Domain 3: Regulatory Authority and Legal Obligations	Federal vs state authority split (FDA/DEA vs BOP) now explicitly distinguished
4.7: Disciplinary actions against a registered/licensed practice setting	3A: Role of regulatory authorities – authority, inspections, & discipline (federal: FDA, DEA; state: BOP, DOH)	