How to Study for the MPJE

Provided by former Commissioner Steven Anderson

1. Read and study the online (not PDF) version of the DEA Pharmacists Manual including the hyperlinks to Title 21. A full read through takes about 3 hours. If you know this well, you should be able to answer about half the MPJE exam questions. You may need to read this more than once. (See link in Federal Law Study Guide Links attachment)

2. Know federal regulations pertaining to the PPPA, general USP 795, 797 and 800 compounding requirements, FDA good manufacturing practices for 503A and 503B pharmacies as well as repackaging, FDA RX and OTC labeling requirements, misbranding vs. adulteration, and rules concerning pseudoephedrine sales. (See Federal Law Study Guide Links attachment)

3. Make an outline of Washington using the NABP Survey of Pharmacy Law, which you can purchase at https://nabp.pharmacy/publications-reports/publications/survey-of-pharmacy-law/ . Review applicable laws and rules on the Pharmacy law webpage highlighting details to fill in your outline (https://www.doh.wa.gov/LicensesPermitsandCertificates/ProfessionsNewReneworUpdate/PharmacyCommission/Laws).

This should take you about two days, and you can go back and review the highlighted information and your outline several times, including the day of the exam. The Survey is $195, but shows you the areas you need to focus on. It is definitely worth it if you plan to sit for MPJE's in multiple states. As Survey results are obtained from state boards of pharmacy, there are on rare occasion slight errors and omissions. You should be able to discern these while studying the law book.

4. The NAPLEX/MPJE Application Bulletin available on the NABP website contains an excellent list of competencies beginning on page 35. Answering these questions in relation to state law will definitely help you be prepared for the exam. The link is: https://nabp.pharmacy/programs/mpje/ . Also consider taking the pre/MPJE practice exam https://nabp.pharmacy/programs/mpje/pre-mpje/ .

5. Read the last two years' worth of the Washington PQAC Newsletters. Go to https://nabp.pharmacy/boards-of-pharmacy/washington/.

6. Go to https://quizlet.com/ and search Washington MPJE. Put in print format and use that as a pre-test. There is also a federal version that is very helpful. Please be aware that there may be errors or outdated answers to the questions. It is recommended that Quizlet be used after all other studying is complete.

Reminder:

- The MPJE is a computer-adaptive examination based in a nationally uniform content blueprint with questions that are tailored to assess the pharmacy jurisprudence requirements of individual states.
- The MPJE consists of 120 multiple-choice questions, 20 of which are designated as pre-test questions that do not affect the candidate's score. The examination content blueprint, which is the percentage of questions asked in each of the MPJE competency areas, is uniform for all candidates.
- The individual questions within each content area will differ from candidate to candidate, depending upon their ability level as estimated by the computer's technology. Therefore, candidate scores are not based solely on the number of correct answers, but on an estimate of the candidate's ability level based on the difficulty of the questions and the number of questions answered correctly. Taking a little extra time to answer the difficult questions will give a higher score than just guessing and moving on to the simpler questions.
- The questions may appear arbitrary and answers incomplete, requiring a judgment call. The idea is to mimic real practice settings. Acting in the patient's benefit is always a good choice if faced with a dilemma.
- Read the question and all the answers TWICE before selecting an answer. It is easy to misread a question and jump to an incorrect conclusion.

Washington Law Study Guide Links

Who Can Prescribe in Washington State?

https://www.doh.wa.gov/LicensesPermitsandCertificates/ProfessionsNewReneworUpdate/PharmacyCommission/WhoCanPrescribeandAdministerPrescriptions

Policies, Procedures, and Guidelines

https://www.doh.wa.gov/LicensesPermitsandCertificates/ProfessionsNewReneworUpdate/PharmacyCommission/PoliciesandProcedures

WA RCW's & WAC's

https://www.doh.wa.gov/LicensesPermitsandCertificates/ProfessionsNewReneworUpdate/PharmacyCommission/Laws

WA Pharmacy Commission Website

https://www.doh.wa.gov/LicensesPermitsandCertificates/ProfessionsNewReneworUpdate/PharmacyCommission/Laws

Bill Fassett's General Review Questions to Prepare for State Law Exams

https://pharmacy.wsu.edu/documents/2017/12/questions-state-pharmacy-law-exams.pdf/

Federal Law Study Guide References

Drug Enforcement Administration (DEA)

- Pharmacist's Manual: https://www.deadiversion.usdoj.gov/pubs/manuals/(DEA-DC-046)(EO-DEA154)_Pharmacist_Manual.pdf#search=pharmacist%20manual

Poison Prevent Packaging Act 16 CFR 1700 (refer to summary below)

- PPPA Guide for Healthcare Professionals: (https://www.cpsc.gov/PageFiles/113945/384.pdf)

Prescription Drug Marketing Act - may be accessed at:

- https://www.fda.gov/regulatoryinformation/lawsenforcedbyfda/significantamendmentstothefdcact/prescriptiondrugmarketingactof1987/default.htm

FDA statutes governing Recalls, Misbranding, Adulterated Drugs

- http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfCFR/CFRSearch.cfm?fr=7.3&SearchTerm=recalls
- https://www.gpo.gov/fdsys/pkg/USCODE-2010-title21/html/USCODE-2010-title21-chap9-subchapV-partA-sec352.htm
- https://www.gpo.gov/fdsys/pkg/USCODE-2010-title21/html/USCODE-2010-title21-chap9-subchapV-partA-sec351.htm

The Federal Food, Drug and Cosmetic Act (FDCA)

- FDA regulations governing labeling directions https://www.law.cornell.edu/uscode/text/21/chapter-9 http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfcfr/CFRSearch.cfm?fr=201.57

FDA regulations governing Patient Package Inserts

- http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfCFR/CFRSearch.cfm?fr=310.515
- http://www.accessdata.fda.gov/scripts/cdrh/cfdocs/cfCFR/CFRSearch.cfm?fr=310.501

FDA Compliance Policy Guides

- https://www.fda.gov/ICECI/ComplianceManuals/CompliancePolicyGuidanceManual/default.htm

Federal Limits on Pseudoephedrine Containing Products

- https://www.deadiversion.usdoj.gov/meth/cma2005.htm

Drug Supply Chain Security Act

- http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM453225.pdf
- http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM400470.pdf

Federal Law Study Guide References

FDA:

- Compounded Drug Products that are Essentially Copies of Approved Drug Products Under Section 503A: http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM510154.pdf

- Compounded Drug Products that are Essentially Copies of Approved Drug Products Under Section 503B: http://www.fda.gov/downloads/Drugs/GuidanceComplianceRegulatoryInformation/Guidances/UCM510153.pdf

Title	*Volume*	*Chapter*	*Browse Parts*	*Regulatory Entity*
Title 21 Food and Drugs	1	I	1-99	Food and Drug Administration, Department of Health and Human Services
	2		100-169	
	3		170-199	
	4		200-299	
	5		300-499	
	6		500-599	
	7		600-799	
	8		800-1299	
	9	II	1300-1399	Drug Enforcement Administration Department of Justice
		III	1400-1499	Office of National Drug Control Policy

DEA Pharmacist Manual Questions

https://www.deadiversion.usdoj.gov/pubs/manuals/pharm2/pharm_manual.pdf#search=pharmacist%20manuaL

Registration Requirements

1. What is required before any controlled substances can be dispensed, administered, possessed, or stored?
2. What does the DEA require of every potential registrant before issuing a DEA registration?
3. How often must a DEA registration be renewed?
4. If the pharmacy changes location, does the pharmacy need a new DEA Registration?
5. If the modification is approved, does the DEA issue a new certificate of registration and new DEA Form 222's?
6. What should the registrant do with the old active DEA Registration?
7. If a pharmacy closes or ceases dispensing controlled substances, what must be done with the DEA Registration and the unused 222 forms?
8. If a pharmacy transfers its business operations (sale) to another pharmacy, who does the pharmacy notify and when?
9. Does the transferring pharmacy need to take a controlled substance inventory and if so, when?
10. Does a copy of the inventory need to be sent to the DEA?

Transfer or Disposal of Controlled Substances

1. How does the transferring pharmacy transfer CII controlled substances to the receiving pharmacy?
2. What should the transferring pharmacy do with its controlled substance records?
3. To whom can a pharmacy transfer expired or unwanted controlled substances for destruction?
4. What is required to dispose of CII controlled substances?
5. How does a reverse distributor notify the DEA when controlled substances are destroyed?

Security Requirements

1. How and when should a pharmacy notify the DEA of controlled substance theft or significant loss?
2. Must a pharmacy also notify law enforcement or state regulatory agencies (i.e. Board of Pharmacy)?
3. Can the pharmacy have its corporate parent notify the DEA of the theft or loss?
4. What else must the pharmacy do with regard to report the theft or loss?
5. When must the pharmacy file the Form 106 with the DEA?
6. What if the pharmacy determines there is no theft or loss after notifying the DEA?

DEA Pharmacist Manual Questions

7. How does the pharmacy determine what is “Significant Loss”?
8. How is a loss reported if it occurs in transit, either to or from the pharmacy, and who reports it?
9. Who reports an in-transit loss from a Central Fill Pharmacy?
10. Does breakage and spillage of controlled substances constitute a loss?
11. What two ways may controlled substances be stored in the pharmacy?

Recordkeeping Requirements

1. How long must pharmacies maintain controlled substance records?
2. How must controlled substance records be maintained?
3. What controlled substance records are required to be maintained?
4. Can a pharmacy store records centrally? If so, how, and does that include all records?
5. How long does the pharmacy have to produce the records upon a request by the DEA?
6. How must pharmacies file controlled substance prescription hardcopies?
7. What if the controlled substance prescriptions are electronic?
8. Can the pharmacy keep both electronic and hardcopy prescriptions of controlled substance prescriptions transmitted electronically?

Inventory Requirements

1. What are the inventory requirements for CII, III, IV, and V controlled substances?
2. When must a pharmacy take a controlled substance inventory?
3. What must be included in the inventory?
4. Does the DEA require the pharmacy name, address, DEA number, and signature of the person responsible for taking the inventory?
5. On what date must the biennial inventory be taken?
6. When a drug not previously listed as a controlled substance is scheduled or a drug is rescheduled, when must it be inventoried by the pharmacy?

Ordering Controlled Substances

1. When is a DEA Form 222 required?
2. When ordering DEA 222 forms, how many forms come in a book and how many books may be ordered at one time by the pharmacy?
3. What are the colors of the three copies of DEA Form 222 and what happens to each copy in the order process?
4. Can a supplier accept a DEA Form 222 that contains minor misspellings of the drug name, or the size and strength have been entered incorrectly?
5. Can a supplier fill in sections on a DEA Form 222 that have been omitted by the purchaser?
6. Can the supplier substitute five packages of 100 if the purchaser requests one package of 500?
7. Can a purchaser void a line on a DEA Form 222?

DEA Pharmacist Manual Questions

8. Can a supplier void part or all of a purchaser's order on a DEA Form 222?
9. Who may sign a DEA Form 222?
10. To how many individuals can a registrant issue power of attorney?
11. When can a supplier not fill a DEA Form 222 submitted by a purchaser and what must the supplier do with the two copies of the form received?
12. What must the purchaser do if a DEA Form 222 is unacceptable to the supplier?
13. What must a pharmacy do if a completed DEA Form 222 that has been sent to the supplier is lost?
14. What must a pharmacy do if unused DEA Forms 222 are lost or stolen?
15. What records must a pharmacy keep after receiving controlled substances?

Valid Prescription Requirements

1. In addition to drug name, strength, dosage form, quantity, directions, and refills (if any), what four elements are required on a controlled substance prescription?
2. What are the requirements for a prescriber to issue controlled substance prescriptions?
3. What are the signature requirements for controlled substance prescriptions?
4. What is meant by the term "valid prescription"?
5. Can a practitioner write a prescription for controlled substances for office use?
6. What is meant by "corresponding responsibility"?
7. Can a practitioner **prescribe** or **dispense** a narcotic CIII - V controlled substance for maintenance of detoxification treatment? If so, under what conditions?
8. Can a practitioner prescribe a narcotic CII controlled substance for maintenance and detoxification treatment?
9. How can a practitioner write a controlled substance prescription without a personal DEA registration?

Dispensing Requirements

1. What warning is required by the FDA on prescription label for a controlled substance?
2. To whom can a pharmacist dispense a controlled substance?
3. Can a pharmacist dispense an emergency CII controlled substance without a written or electronic prescription?
4. Can a central fill pharmacy dispense an emergency CII prescription?
5. What are the special restrictions on the issuance of multiple CII controlled substance prescriptions for the same drug at the same time?
6. Explain the restrictions on partial filling of CII controlled substance prescriptions.
7. A CII controlled substance can be refilled under what specific circumstances?
8. Under what three circumstances does the DEA allow a facsimile of a non-emergency CII prescription to substitute for a written or electronic CII prescription?
9. What changes/additions may a pharmacist make on a CII controlled substance prescription?
10. What must be included on the prescription label for a controlled substance prescription filled at a central fill pharmacy?

DEA Pharmacist Manual Questions

11. What are the retail pharmacy record keeping requirements for controlled substance prescriptions transmitted by a retail pharmacy that are filled at a central fill pharmacy?
12. What are the central fill pharmacy record keeping requirements for controlled substance prescriptions transmitted by a retail pharmacy that are filled at a central fill pharmacy?
13. A CIII – V controlled substance prescription may be refilled a maximum of 5 times and expires in 6 months. How many times can the prescription be refilled if the quantity dispensed is less than the quantity written?
14. What information must be recorded on a prescription transferred verbally or by facsimile from another pharmacy?
15. What are the requirements for dispensing a CII – V controlled substance without a prescription?

Other Controlled Substance Regulations

1. What are the USPS packaging requirements for mailing controlled substances?
2. Can a controlled substance prescription be mailed to a person in a foreign country?
3. What must a pharmacy have to dispense controlled substances from an ADDD in a remote location?
4. What is the "Five Percent Rule"?

Combat Methamphetamine Epidemic Act of 2005

1. What are the requirements of retailers under the CMEA in regards to ephedrine and pseudoephedrine sales?
2. What are the recordkeeping requirements under the CMEA?
3. What are the purchasing requirements under the CMEA?
4. What are acceptable forms of purchaser ID under the CMEA?

Questions	Answer--WA	Reference
Pharmacy Quality Assurance Commission		
How many members serve on the PQAC? How many are public members?		
What are the qualifications to being a pharmacist on the PQAC?		
How long can you serve on the PQAC?		
Licensing		
What are the requirements to get licensing		
When do you renew your license		
What are the CE requirements?		
How do you inactivate a license?		
How do you reinstate an inactive license?		
Prescriptive Authority		
What is the prescribing authority of the following: 1. MD 2. DO 3. DVM 4. DPM (podiatry) 5. Naturopathy 6. Optometry 7. PA 8. ARNP 9.Chiropractor 10. Dentist		
Can a dentist prescribe fluconazole with an antibiotic for patients with frequent yeast infections?		
Can an Anesthesiologist prescribe atorvastatin?		
What prescriptions can you accept from out of state providers		
Who can prescribe a controlled substance		
Control Substances		
What are the requirements on a CII hard copy?		
How long are CII valid?		
How can a CII be sent to the pharmacy?		
Can you dispense a CII without a prescription in emergencies?		
How many times can CII be refilled?		
What is the CII day supply limitiation?		
What changes/additions can you make on a CII?		
Is the MD's name and DEA # on Rx written by midelvel practitioner required on the prescription?		
How many CS can be on one blank		
Is documentation of justification for early refills of CS required?		
Is ID required for pick up of CS?		
How long are CIII-IV valid?		
What is the days supply limitation on CIII-IV?		
How many times can CIII-IV be refilled?		
How many times can you transfer a CIII-IV		
How long are CV valid		
Can CV be OTC		

Questions	Answer--WA	Reference
Are there any scheduled drugs that differ from the federal schedule? (Ie: gabapentin)		
Can a doctor self prescribe?		
Pharmacy Requirements		
what are the requirements of the pharmacy practice site?		
Is a consultation area required?		
Who has access to the pharmacy?		
Who can purchase legend drugs for use in their practice?		
Who can possess CSA and keep them onsite?		
Who has dispensing authority?		
Which of the following can prescribe CSA drugs (and which class?): OD, NP, clinical nurse specialist, PA, midwife, nurse midwife, chiropractor, clinical psychologist, PT, RT, pharmacist		
What prescriptions can you accept from out of state providers?		
How long are prescriptions for legend drugs valid from the date issued		
Is there a maximum number of refills allowed for legend drugs? What is it?		
Is there a max # refills allowed for C5? What is it? Are the number of refills for C3 and C4 the same as federal or less		
Any special requirements for CII RX? If so, what?		
Which of the following must be on the label of an outpt rx: pt address, quantity, exp date, lot number, name of drug, RPH initials, technician initials, pharmacy phone number? other items?		
Can the prescribed brand name be placed on the label when a generic is dispensed? If so, what wording is required?		
How does a pharmacist determine whether a particular generic drug is suitable as a substitute for a brand name drug?		
Must generic savings be passed on to the patients? if so, what proportion and how determined?		
How can a prescriber indicate that substitution is or is not permitted?		
Do patients have the option of demanding the brand name when subsitution is permitted? how about pt whose rx are paid for by public funds?		
What documentation is required of the pharmacist when product interchange is performed? ▫ therapeutic drug interchange: a switch to a drug providing similar therapeutic response to the one prescribed (biosimilars)		
Under what conditions, if any, is therapeutic substitution allowed?		
What are the different classes of pharmacies?		
What minimum sets of references, products, or equipment are required for all community pharmacies?		
Do pharmacy hours have to be posted? where?		
Can the store be open but the pharmacy department be closed? under what requirements?		

Questions	Answer--WA	Reference
Do pharmacist licenses have to be on display? the original or official copy? is a photocopy ok? can you obscure your address on the copy displayed to the public?		
Are there any required notices that must be displayed to the public?		
How much counter space must a pharmacist/intern have for filling prescriptions or compounding?		
Is a separate pt counseling area required?		
Is OBRA counseling required for Medicaid only or for all pt?		
How must the offer to counsel be made?		
Does the offer to counsel need to be documented? Does the refusal of the offer need to be documented? how?		
Is counseling required on new rx, refills, or both?		
Can printed materials satisfy the counseling requirement? under what circumstances?		
Are there any special requirements for mail order pharmacies?		
Under what circumstances may a pharmacist fill prescriptions written in another state?		
Is there any provision for dispensing an emergency refill supply of legend drugs when there are no refills left and the prescriber cannot be contacted? what is allowed/required?		
Are pt profiles required for all pt?		
What information is required on the profile?		
What use of the profile is required? review prior to dispensing new rx? review prior to refills?		
Can a pt refuse to give certain info for the profile? if so, must the refusal be documented, and how?		
What are the requirements for use of computers to process rx?		
what must be done if the computer system goes down?		
Is there any requirement for use of computers to process prescriptions? Who must sign the printout?		
Can a pharmacist legally repackage another pharmacy's prescription?		
Can legend drugs prescriptions be faxed? Under what circumstances and requirements?		
Can legend drug prescriptions be transferred electronically (by computer)? Under what circumstances and requirements?		
Under what circumstances are faxed CSA prescriptions allowed?		
Can refills of legend drugs be transferred from one pharmacy to another? Can they be transferred back?		
Can an intern transfer refills?		
Must information regarding the transfer of legend drugs be recorded on the hard copy of the original rx, or can it merely be recorded on the computer?		
What information must be recorded by the transferring pharmacy?		
What information must be placed on the original copy of the transferred rx by the receiving pharmacy?		

Questions	Answer--WA	Reference
Is it possible for pharmacies under a common ownership to share a single pt/rx database? If so, what are the rules for rx transfer?		
How long must the pharmacy maintain the following records: original rx, refill records, drug purchase records, pt profiles? Are any of these different for CSAs? If any of these can be maintained on computer, how long do they need to be maintained 'online'?		
Must every pharmacy have a pharmacist in charge (PIC)?		
Are they any special requirements to be PIC?		
May 1 person be the PIC for more than one pharmacy?		
When must the BOP be notified of any of the following: change in PIC, change of pharmacy address, closing or sale of pharmacy, change of pharmacy phone number, change of an individual pharmacist's address, change of an intern's address, change of tech's address, change of person's name?		
What must be on the label of a unit-dose or single-dose package of a legend drug dispensed for a pt in a hospital or nursing home?		
What must be on the label of a multiple dose container dispensed for a pt in the hosp? In a nursing home?		
What is required on the label of a parenteral solution dispensed to a pt for home infusion therapy?		
What is required on the label of a radiopharmaceutical agent?		
What are special requirements, if any, to be a nuclear pharmacist?		
Can transfers of refill information be faxed between pharmacies?		
How rapidly must a pharmacy respond to a pt's request to amend the information in his or her medication record?		
May a pharmacy transmit a pt rx claim information electronically to a third party payor w/o the pt having provided written consent to the payor or the pharmacy?		
When must a pharmacy provide a copy of its Notice of Privacy Practices to a pt or other person requesting one?		
How can changes in the NOPP be communicated to pt?		
Can a non-custodial parent act on behalf of a minor to authorize disclosure of information in the pt medication record?		
What is the max amount of PSE that can be sold OTC in an individual in a given day?		
May pharmacists administer drugs? By which route(s)?		
What are the requirements for immunizations administered by pharmacists?		
To what extent are collaborative practice agreements allowed?		
What is the status of carisoprodol? Tramadol?		
Under what circumstances may a pharmacist refuse to fill a lawful rx?		
What is the max ratio of interns to pharmacists?		
What is the ratio of pharmacists to techs? Does it differ by setting?		
What are the requirements for storage and accountability for controlled substances in a hospital?		
What are the requirements for storage and accountability for CS in LTCF?		
Are tech-check-tech programs allowed? Under what circumstances?		

Questions	Answer--WA	Reference
Can techs fill IVs? Can techs fill or check drug dispensing machines?		
What are the requirements for remote processing of prescriptions?		
Is remote processing legal?		
What are the requirements of OTC labeling?		
When opening new pharmacy, does there have to be physical inspection before the pharmacy is licensed?		
If pharmacy uses central fill facility, is there any documentation required to be sent to patient?		
Is a patient profile needed for recipient of STD rx?		
Can you fill a rx from a doctor in canada? which province?		
Can a compounding pharmacy compound for vet office use?		
Going to central fill, any special requirments for rx?		
What is required for prescriber's agent to send a fax or call in an rx?		
Manual signature requirement for faxed rx?		
Can RPh administer drugs (b12, etc) w/o CDTA?		
What needs to be on OTC if put in rx vial?		
After hours access to pharmacy in hospital?		
When does RPh license expire?		
What are the requirements for a retired pharmacist license?		
OPTIONAL		
Continuing Pharmacy Education Requirements		
Discipline of Pharmacy License		
Licensure expiration		
Change on a C2?		
Pharmacist Self-Dispensing		
Valid DEA#		
DEA Letters		
Prescription records maintanied for 2 years		
Absence of pharmacist from hospital pharmacy		
Early Refill of opthalmic products		
Provision of drugs to ambulance and aid services		
Commission powers and Duties		
Schedule I		
Schedule II		
Schedule III		
Schedule IV		
Schedule V		
New Pharmacy registration		
Termination of Registration		
Transfer of Business		
Information included		
SLCP - Scheduled Listed Chemical Products		
Transfer of Controlled Substances		
Disposal of controlled substances		
Employment Waivers		
Theft or significant loss		

Questions	Answer--WA	Reference
Significant loss		
In-Transit Loss		
Breakage or Spillage		
Robberies and burglaries		
Maintaining Records		
Prescription records		
Inventory requirements		
CFR requires		
Biennial inventory		
Newly scheduled controlled substance		
Ordering controlled substances		
Completing order forms		
Refusing orders		
Cancelling and voiding of official order form		
Lost or stolen 222 forms		
Controlled substance Ordering System		
Ordering Schedule 3-5		
Valid Prescription requirements		
Paper or oral prescription		
Exemption from registration		
Fax C2 Prescriptions		
Exceptions for written C2 only		
recordkeeping for refill C3-C5		
Electronic recordkeeping of controlled substances		
Transfer of Schedule 3-5		
Dispensing Requirements		
Emergency dispensing		
Dispensing without prescription		
Ryan Haight Act		
Online Pharmacy		
Reporting		
Central fill pharmacy		
Long term Care		
5% rule		
USPS mailing requirements		
SLCP - Scheduled Listeed Chemical Products		
License Reinstatement requirements		
Approval of CE program		
CE Program Hours		
Presenting to health professionals		
Patient education training program		
Licensing Requirements		
Reciprocity applicants		
Temporary practice permit		
Foreign graduate		
Pharmacy manager		
Inactive Credential or Expired license		

Questions	Answer--WA	Reference
Retired Pharmacist license		
Professional responsibilities		
Unprofessional conduct		
Prescriptive Authority		
Monitoring of drug therapy by pharmacists		
AIDs prevention hours		
Emergency Drug Kit		
Drug facilities		
Pharmaceutical services		
Schedule 2 drugs nursing home		
Continuation of drug therapy		
Reporting dependencce		
Pharmacies' responsibilities		
Pharmacy license notification requirements		
New pharmacy registration		
Clinic dispensaries		
Emergency Action		
Refusal to permit inspection		
Returns or Exchange of drugs		
Exceptions to returns/exchange rule		
Prescription department		
Physical standards of pharmacies		
Inspections		
Poison control		
Prescription labeling		
Patient counseling		
Always child resistant except		
Closing a pharmacy		
Medication dose pack		
Electronic transfer of prescription information		
Schedule 2 electronic		
fax machines		
Electronic system board approval		
Electronic storage		
Class 100 environment		
Physical requirements of parenteral area		
Patient profile contains		
Antineoplastic medications		
Clinical Services		
Quality assurance		
Pharmacy responsibilities of drug distribution		
Facility responsibilities		
Absence of a pharmacist from hospital		
Emergency outpatient medications		
Schedule 2 in hospital		
Destruction of Schedule 2		
Administration of Meds		

Questions	Answer--WA	Reference
Medical system record		
Confidentiality and Security of data		
Drug Sample prohibitions		
Compounding Drugs		
Special preparation products		
Compounding controls		
Transferring drug supply		
Export Wholesaler		
Drug Price disclosure		
Drug samples		
Ephedrine exemption		
Theophylline restrictions		
OTC imprint regulation		
Euthanasia		
Pre-euthanasia sedation		
Record keeping and reports		
Chemical capture programs		
Utilization plans		
Non-Resident Pharmacies		
Extended Care (SNF)		
Destroying drugs		
OBRA Act		
Patient profile Home IV Therapy		
Nuclear Pharmacies		
Nuclear Labeling requirements		
Inner container		
Any Prescription in or out of state		
Mid-level practitioners, no out of state prescriptions		
Protocol		
Expired license or dead prescriber		
Adulterated drug		
Misbranded drug		
FDA Books		
Suboxone for addiction		