

Connecticut MPJE Practice Test (Sample)

Study Guide



Everything you need from our exam experts!

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Introduction

Preparing for a certification exam can feel overwhelming, but with the right tools, it becomes an opportunity to build confidence, sharpen your skills, and move one step closer to your goals. At Examzify, we believe that effective exam preparation isn't just about memorization, it's about understanding the material, identifying knowledge gaps, and building the test-taking strategies that lead to success.

This guide was designed to help you do exactly that.

Whether you're preparing for a licensing exam, professional certification, or entry-level qualification, this book offers structured practice to reinforce key concepts. You'll find a wide range of multiple-choice questions, each followed by clear explanations to help you understand not just the right answer, but why it's correct.

The content in this guide is based on real-world exam objectives and aligned with the types of questions and topics commonly found on official tests. It's ideal for learners who want to:

- Practice answering questions under realistic conditions,
- Improve accuracy and speed,
- Review explanations to strengthen weak areas, and
- Approach the exam with greater confidence.

We recommend using this book not as a stand-alone study tool, but alongside other resources like flashcards, textbooks, or hands-on training. For best results, we recommend working through each question, reflecting on the explanation provided, and revisiting the topics that challenge you most.

Remember: successful test preparation isn't about getting every question right the first time, it's about learning from your mistakes and improving over time. Stay focused, trust the process, and know that every page you turn brings you closer to success.

Let's begin.

How to Use This Guide

This guide is designed to help you study more effectively and approach your exam with confidence. Whether you're reviewing for the first time or doing a final refresh, here's how to get the most out of your Examzify study guide:

1. Start with a Diagnostic Review

Skim through the questions to get a sense of what you know and what you need to focus on. Your goal is to identify knowledge gaps early.

2. Study in Short, Focused Sessions

Break your study time into manageable blocks (e.g. 30 - 45 minutes). Review a handful of questions, reflect on the explanations.

3. Learn from the Explanations

After answering a question, always read the explanation, even if you got it right. It reinforces key points, corrects misunderstandings, and teaches subtle distinctions between similar answers.

4. Track Your Progress

Use bookmarks or notes (if reading digitally) to mark difficult questions. Revisit these regularly and track improvements over time.

5. Simulate the Real Exam

Once you're comfortable, try taking a full set of questions without pausing. Set a timer and simulate test-day conditions to build confidence and time management skills.

6. Repeat and Review

Don't just study once, repetition builds retention. Re-attempt questions after a few days and revisit explanations to reinforce learning. Pair this guide with other Examzify tools like flashcards, and digital practice tests to strengthen your preparation across formats.

There's no single right way to study, but consistent, thoughtful effort always wins. Use this guide flexibly, adapt the tips above to fit your pace and learning style. You've got this!

Questions

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- 1. What information must be included in an authorization form regarding HIPAA?**
 - A. Specific uses and disclosures of PHI allowed**
 - B. Patient's medication history**
 - C. Appointment scheduling preferences**
 - D. Insurance claim details**
- 2. Pharmacy technicians are required to be certified with DCP to practice. True or False?**
 - A. True**
 - B. False**
 - C. Depends on experience**
 - D. Not specified**
- 3. For patients to participate in investigational drug trials, they must be in what condition?**
 - A. Early stages of illness**
 - B. In an imminent life-threatening stage with no cure available**
 - C. Recovering from a recent treatment**
 - D. Stable condition with effective current treatments**
- 4. True or False: The total number of dosage units of all controlled substances distributed may not exceed 5% of all controlled substances dispensed in a calendar year.**
 - A. True**
 - B. False**
 - C. It depends on state law**
 - D. This rule only applies to Schedule II substances**
- 5. How long must a faxed prescription document be retained?**
 - A. 1 year**
 - B. 2 years**
 - C. 3 years**
 - D. 5 years**

6. Under what conditions can a faxed document for a C2 prescription be allowed?

- A. Only for patients with chronic conditions**
- B. If for hospice or LTC**
- C. In any situation with a prescriber's verbal approval**
- D. If it's a refill prescription**

7. What are the contents of a CDTM agreement?

- A. Only the type of prescriptive authorities**
- B. The types of patients they can see and required training**
- C. Type of prescriptive authorities, types of patients, procedures, and documentation requirements**
- D. No specific content is necessary**

8. What should a patient be informed about if a dose is missed?

- A. Nothing, as it will not affect therapy**
- B. Take two doses the following day**
- C. Action to take if a dose is missed**
- D. Contact the prescriber immediately**

9. What is required information for patient counseling?

- A. Only the name of the medication**
- B. Name, description, and dosage form**
- C. Common side effects only**
- D. Dosage and refill information only**

10. Must needles be stored in pharmacies so they are only available to authorized personnel?

- A. True**
- B. False**
- C. Only when shelves are open**
- D. Only during business hours**

Answers

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1. A
2. A
3. B
4. A
5. C
6. B
7. C
8. C
9. B
10. A

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Explanations

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1. What information must be included in an authorization form regarding HIPAA?

- A. Specific uses and disclosures of PHI allowed**
- B. Patient's medication history**
- C. Appointment scheduling preferences**
- D. Insurance claim details**

The correct answer is that an authorization form regarding HIPAA must include specific uses and disclosures of Protected Health Information (PHI) that are allowed. Under HIPAA regulations, an authorization form is a critical document that provides healthcare providers with the explicit permission to use or disclose a patient's PHI for particular purposes beyond treatment, payment, or healthcare operations. This requirement ensures transparency and allows patients to understand exactly how their information will be handled. When patients sign an authorization form, they should clearly see which types of disclosures are permitted and under what conditions. This could include, for instance, sharing information with insurance companies for billing purposes, or allowing information to be shared with third parties for research, among others. By including this information, the authorization form aligns with HIPAA's privacy rules, empowering patients to make informed choices about their personal health information. Other options, while related to healthcare, do not meet the criteria for what must be included in a HIPAA authorization form. For example, a patient's medication history, appointment scheduling preferences, and insurance claim details are not required to be outlined in an authorization form. Instead, they are typically managed and shared through other means within the healthcare system.

2. Pharmacy technicians are required to be certified with DCP to practice. True or False?

- A. True**
- B. False**
- C. Depends on experience**
- D. Not specified**

In Connecticut, pharmacy technicians are indeed required to be certified by the Department of Consumer Protection (DCP) to practice. This certification ensures that pharmacy technicians have met the necessary educational and competency standards to assist pharmacists effectively in their work. Certification typically involves passing an exam that assesses knowledge and skills related to pharmacy practice, which ultimately contributes to the safety and effectiveness of medication dispensing and patient care. Other options lack the clarity or accuracy regarding the certification requirement. For instance, saying it is false would imply that no certification is needed, which is incorrect as certification is a clear requirement. A statement about dependence on experience could mislead individuals into thinking that prior experience might exempt someone from certification, which is not the case in Connecticut. Similarly, stating that it is not specified does not align with the regulations currently set by the DCP for pharmacy technicians. Hence, the requirement for certification is a definitive aspect of pharmacy practice in Connecticut, reinforcing the importance of maintaining high professional standards.

3. For patients to participate in investigational drug trials, they must be in what condition?

- A. Early stages of illness**
- B. In an imminent life-threatening stage with no cure available**
- C. Recovering from a recent treatment**
- D. Stable condition with effective current treatments**

For patients to participate in investigational drug trials, being in an imminent life-threatening stage with no cure available is often a criterion. This condition allows researchers to assess the efficacy and safety of new treatments in populations that may not respond to existing therapies. In such cases, patients may be given the opportunity to try experimental drugs that could potentially save their lives, as the risk of participating in the trial may be outweighed by the potential for benefit when no established treatment options exist. This unique situation is particularly relevant in the context of clinical trials designed for conditions that are severely debilitating or life-threatening, where conventional treatments have failed, providing a critical avenue for exploration of new therapeutic options. Consequently, it is essential for participants to be in a challenging medical state to justify the participation in trials exploring their condition, thereby aligning with the regulatory goal of advancing medical knowledge while ensuring patient safety.

4. True or False: The total number of dosage units of all controlled substances distributed may not exceed 5% of all controlled substances dispensed in a calendar year.

- A. True**
- B. False**
- C. It depends on state law**
- D. This rule only applies to Schedule II substances**

The statement is true. According to federal regulations, specifically under the provisions governing the distribution of controlled substances, a registrant is allowed to distribute a limited quantity of controlled substances without registering as a distributor. This limit is set at 5% of the total number of dosage units of controlled substances dispensed in a calendar year. This rule helps ensure that those who are primarily practitioners and dispensers are not classified as distributors unless they exceed this threshold. While some might consider the variations in state laws, the federal limit is a general guideline that must be adhered to unless state laws are more stringent. The regulation ensures that distribution remains tightly controlled and monitored to prevent potential misuse or diversion of controlled substances. The option referring specifically to Schedule II substances is incorrect as the 5% rule applies to all controlled substances, not just those in a specific schedule.

5. How long must a faxed prescription document be retained?

- A. 1 year
- B. 2 years
- C. 3 years**
- D. 5 years

The correct answer is based on the requirement that a faxed prescription must be retained for a specific period, which in this case is three years. This retention period is consistent with various regulations governing the storage of prescription records to ensure proper oversight and accountability in pharmacies. By maintaining records for this duration, it allows for sufficient time to address any issues related to the prescriptions, such as audits, inquiries from regulatory authorities, or patient-related matters. The three-year requirement is significant as it aligns with best practices in pharmacy management and ensures compliance with both state and federal regulations regarding record-keeping. This duration helps facilitate proper monitoring of prescription dispensing and patient medication use, which is essential for patient safety and effective pharmacy practice. The other time frames provided do not match the established requirement for faxed prescriptions, highlighting the importance of adhering to the correct regulatory standards.

6. Under what conditions can a faxed document for a C2 prescription be allowed?

- A. Only for patients with chronic conditions
- B. If for hospice or LTC**
- C. In any situation with a prescriber's verbal approval
- D. If it's a refill prescription

The correct answer is that a faxed document for a Schedule II (C2) prescription is permitted if it is for hospice care or long-term care (LTC) facilities. This provision is based on federal and state regulations that allow certain exceptions for the dispensing of C2 medications in specific circumstances, such as for patients who are in hospice or residing in long-term care facilities. This helps to ensure that patients in these settings can receive necessary medications in a timely manner, especially in situations where immediate treatment is critical. In hospice settings, where patients may require rapid access to pain management and other controlled substances, the ability to fax prescriptions helps to expedite care. For long-term care facilities, the same principles apply, allowing for efficient medication management for residents who are often unable to visit a pharmacy in person. The other conditions presented do not meet the regulatory requirements set forth for C2 prescriptions. For example, faxing prescriptions for patients with chronic conditions or for refills does not generally align with the strict regulations surrounding C2 substances, which typically require written prescriptions. Similarly, verbal approval from a prescriber is insufficient for C2 prescriptions, which are governed by much stricter controls to prevent abuse and diversion.

7. What are the contents of a CDTM agreement?

- A. Only the type of prescriptive authorities
- B. The types of patients they can see and required training
- C. Type of prescriptive authorities, types of patients, procedures, and documentation requirements**
- D. No specific content is necessary

A Collaborative Drug Therapy Management (CDTM) agreement outlines the responsibilities and authorities shared between pharmacists and physicians to ensure comprehensive patient care. The correct answer includes a comprehensive list of elements that must be detailed in such agreements. The contents of a CDTM agreement typically cover several critical areas: the specific type of prescriptive authorities that the pharmacist will have, the types of patients they are permitted to manage, the procedures or protocols they are allowed to follow, and documentation requirements to ensure proper record-keeping and compliance. This structure ensures clarity in roles and responsibilities, safeguards patient safety, and aligns with regulatory standards. The other choices lack the full scope that's necessary for a comprehensive agreement. For instance, merely listing the type of prescriptive authorities does not encompass the operational framework and patient management aspects integral to a CDTM. Similarly, specifying only the types of patients and required training overlooks essential procedural details and the importance of documentation for accountability and quality of care. Lastly, stating that no specific content is necessary undermines the foundation of structured healthcare agreements which are designed to improve outcomes and clearly delineate responsibilities.

8. What should a patient be informed about if a dose is missed?

- A. Nothing, as it will not affect therapy
- B. Take two doses the following day
- C. Action to take if a dose is missed**
- D. Contact the prescriber immediately

When a patient has missed a dose of their medication, it is crucial that they are informed about the appropriate actions to take. This option emphasizes the need for clear guidance on how to manage a missed dose, which can vary depending on the medication and the timing of the next dose. Providing such information helps ensure that the patient understands whether they should take the missed dose as soon as they remember, skip it if it's almost time for the next dose, or take a specific action related to their therapy. Communicating action steps reinforces patient safety and encourages adherence to the medication regimen. It allows patients to make informed decisions and reduces the risk of potential complications from either taking the medication incorrectly or skipping doses without understanding the implications. In contrast to this, suggesting that nothing should be said (as in one of the other options) fails to address the importance of patient education regarding medication adherence and the implications of missed doses. Similarly, the options regarding taking two doses at once or contacting the prescriber immediately are not universally applicable or practical for all medications, which further underscores the need for tailored guidance specific to the situation.

9. What is required information for patient counseling?

- A. Only the name of the medication**
- B. Name, description, and dosage form**
- C. Common side effects only**
- D. Dosage and refill information only**

The requirement to provide comprehensive patient counseling focuses on ensuring that patients understand their medications for safe and effective use. The correct answer includes the name of the medication, a description of what the medication is intended for, and the dosage form, which provides critical information that aids in recognizing the medication and its appropriate use. Understanding the medication name helps prevent errors and ensures patients are taking the correct product. A description of the medication, including its intended purpose, allows patients to understand why they are taking it, which can improve adherence and therapeutic outcomes. Additionally, the dosage form (such as tablet, liquid, or injection) is essential information that guides the patient on how to administer the medication properly. Other options do not provide a comprehensive view that covers the essential aspects of patient counseling. Limiting information to just the name of the medication or focusing solely on common side effects or dosage and refill details does not equip the patient with the necessary knowledge for safe usage. Effective counseling must encompass a broader understanding to enhance patient engagement and awareness regarding their treatment.

10. Must needles be stored in pharmacies so they are only available to authorized personnel?

- A. True**
- B. False**
- C. Only when shelves are open**
- D. Only during business hours**

The correct answer is that needles must be stored in pharmacies so they are only available to authorized personnel. This requirement is in place to ensure safety and to prevent misuse or accidental injuries associated with needles, which are classified as biohazardous materials. By restricting access to authorized personnel, pharmacies adhere to regulations aimed at minimizing risk of injury, abuse, and contamination. This practice aligns with guidelines set by regulatory agencies, which mandate that potentially dangerous items, such as needles, be kept secure and accessible only to trained staff. This not only promotes safe handling of needles but also helps in maintaining proper inventory control and minimizing the potential for theft or abuse within the pharmacy environment. The other options do not represent best practices or legal requirements regarding the storage of needles. Storing them only during business hours would not provide constant security, while the idea of only allowing access when shelves are open does not adequately ensure that unauthorized individuals cannot access them at other times.

Next Steps

Congratulations on reaching the final section of this guide. You've taken a meaningful step toward passing your certification exam and advancing your career.

As you continue preparing, remember that consistent practice, review, and self-reflection are key to success. Make time to revisit difficult topics, simulate exam conditions, and track your progress along the way.

If you need help, have suggestions, or want to share feedback, we'd love to hear from you. Reach out to our team at hello@examzify.com.

Or visit your dedicated course page for more study tools and resources:

<https://connecticutmpje.examzify.com>

We wish you the very best on your exam journey. You've got this!

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