

Positron Emission Tomography-Computed Tomography (PET/CT) Fusion Practice Exam (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 kind of tumors generally show high FDG uptake?**
 - A. Benign tumors with low activity**
 - B. Slow-growing tumors**
 - C. Aggressive tumors with high metabolic rates**
 - D. Cysts with minimal metabolic activity**

- 2. Which range of Hounsfield units is associated with contrast material?**
 - A. 70**
 - B. 100 to 600**
 - C. 1**
 - D. -1000**

- 3. What should be done to correct for random events in PET imaging?**
 - A. Simultaneously acquire in 2 energy windows**
 - B. Create a map of density within the field of view**
 - C. Inject the proper dose**
 - D. Increase image time**

- 4. What can interfere with the results of a PET scan post-injection?**
 - A. Low blood pressure**
 - B. High blood glucose levels**
 - C. Insufficient hydration**
 - D. Increased exercise activity**

- 5. What is an important consideration regarding patient allergies before performing a PET scan?**
 - A. Assessment of other medical conditions**
 - B. Assessment of prior reactions to radiotracers**
 - C. Discussion on dietary restrictions**
 - D. Assessment of family medical history**

- 6. What is a common misconception about PET scans in relation to cancer detection?**
- A. They can identify early-stage diseases**
 - B. They are only for brain imaging**
 - C. They can reveal both structure and function**
 - D. They cannot differentiate between tumor types**
- 7. Which quality of crystals in PET systems impacts their sensitivity?**
- A. Higher density materials**
 - B. Faster decay times**
 - C. Stopping power**
 - D. Scintillation photon yield**
- 8. What is a primary limiting factor for PET imaging resolution?**
- A. Crystal construction**
 - B. Crystal stopping power**
 - C. Photo multiplier tubes**
 - D. Positron range**
- 9. How can obesity impact PET/CT imaging results?**
- A. It has no impact on imaging results.**
 - B. It may improve radiotracer uptake.**
 - C. Increased body fat may alter distribution and uptake of the radiotracer.**
 - D. It leads to better spatial resolution.**
- 10. What does the term 'system dead time' refer to in PET imaging?**
- A. The time taken for the system to reset after an event**
 - B. The period when the system cannot register incoming events**
 - C. The duration for signal processing**
 - D. The time taken for scintillation light to dissipate**

Answers

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

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Explanations

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1. What kind of tumors generally show high FDG uptake?

- A. Benign tumors with low activity
- B. Slow-growing tumors
- C. Aggressive tumors with high metabolic rates**
- D. Cysts with minimal metabolic activity

High FDG uptake in tumors is primarily associated with aggressive tumors that have high metabolic rates. This phenomenon is due to the enhanced glucose uptake and metabolism that characterizes rapidly dividing cancer cells. Positron Emission Tomography (PET) uses fluorodeoxyglucose (FDG), a radiolabeled glucose analog, to visualize areas of high metabolic activity, which often indicates tumor presence. Aggressive tumors, such as certain types of sarcomas, lymphomas, and carcinomas, typically exhibit this high glycolytic activity because they require more energy to support their rapid growth and proliferation. This increased metabolic rate leads to a greater uptake of FDG, making it an effective imaging tool for identifying and characterizing these malignancies. In contrast, benign tumors generally exhibit low metabolic activity, and their FDG uptake is usually minimal. Slow-growing tumors may also show reduced FDG uptake because their metabolic requirements are not as high as those of aggressive tumors. Cysts, which are fluid-filled spaces, typically have minimal metabolic activity, resulting in low FDG uptake as well. Therefore, the identification of aggressive tumors through high FDG uptake serves as an important clinical tool in diagnosis and treatment planning.

2. Which range of Hounsfield units is associated with contrast material?

- A. 70
- B. 100 to 600**
- C. 1
- D. -1000

The range of Hounsfield units associated with contrast material typically falls between 100 to 600. This range reflects the increased density that contrast agents introduce to tissues during imaging. By enhancing the contrast between surrounding structures, these agents provide better visibility of abnormalities or blood vessels on a CT scan. Hounsfield units (HU) are a measure of radiodensity used in computed tomography, where air is at -1000 HU and water is at 0 HU. Materials with HU values above that of water, such as contrast agents, indicate higher density and help delineate structures more clearly on imaging. The other options do not accurately represent the density levels of contrast materials. A singular value like 70 is too low to indicate any substantive contrast effect, while 1 is far less than water and would not be reflective of a contrast medium. The value of -1000 indicates air, which is not relevant to the use of contrast materials in imaging.

3. What should be done to correct for random events in PET imaging?

- A. Simultaneously acquire in 2 energy windows
- B. Create a map of density within the field of view
- C. Inject the proper dose**
- D. Increase image time

In the context of correcting for random events in PET imaging, the emphasis is primarily on ensuring optimal conditions for capturing accurate signals from tracer distribution. The correct choice involves injecting the proper dose of the radiotracer. Administering an appropriate amount of the radiotracer is crucial for distinguishing the signal from random coincidences, which are events that occur due to detector noise rather than actual positron emissions. When a sufficient dose is administered, it increases the likelihood of capturing genuine coincidences between detected photons from the annihilation events, thereby enhancing the signal-to-noise ratio. This helps in minimizing the impact of random events during the imaging process, leading to improved image quality and accuracy in quantifying radioactive uptake in tissues. Meanwhile, the other options, while they may seem applicable in different contexts, do not directly address the correction of random events related to tracer administration in the same effective manner as ensuring the correct dose does. Simultaneously acquiring in two energy windows is often used in advanced techniques but is not primarily aimed at correcting random events. Creating a density map is for different diagnostic purposes, and simply increasing the image time does not resolve the underlying issues related to random coincidences but rather may contribute to longer acquisition times without ensuring signal clarity.

4. What can interfere with the results of a PET scan post-injection?

- A. Low blood pressure
- B. High blood glucose levels**
- C. Insufficient hydration
- D. Increased exercise activity

High blood glucose levels can indeed interfere with the results of a PET scan post-injection. This is primarily due to the fact that PET scans often utilize a radiolabeled glucose analog, such as fluorodeoxyglucose (FDG), to assess metabolic activity in tissues. When blood glucose levels are elevated, normal cells may uptake the excess glucose instead of the radiolabeled tracer, which can lead to an underestimation of the abnormalities or lesions that are intended to be visualized. This results in reduced image specificity and sensitivity, potentially causing a misinterpretation of scans in conditions such as cancer, where increased metabolic activity is expected. For a successful PET scan, it's important to maintain appropriate blood glucose levels, typically fasting before the procedure, to ensure accurate imaging results. This understanding is crucial for clinical protocols to optimize PET scan effectiveness and diagnostic accuracy.

5. What is an important consideration regarding patient allergies before performing a PET scan?

- A. Assessment of other medical conditions**
- B. Assessment of prior reactions to radiotracers**
- C. Discussion on dietary restrictions**
- D. Assessment of family medical history**

Before performing a PET scan, it is crucial to assess a patient's prior reactions to radiotracers used in the procedure. This is because many PET scans involve the administration of a radiopharmaceutical that contains radioactively labeled substances, most commonly a glucose analog like fluorodeoxyglucose (FDG). Understanding a patient's allergy history, particularly in relation to these radiotracers, helps ensure patient safety and minimize the risk of adverse reactions during the scan. If a patient has experienced any allergic reactions to similar substances in the past, it may warrant further evaluation or even a decision to use a different radiotracer or to take a different approach altogether. This assessment is vital for the healthcare team's ability to prepare adequately for the scan and to manage any potential allergic responses should they occur. Thus, prior reactions to radiotracers play a significant role in the pre-scan evaluation process to ensure patient safety.

6. What is a common misconception about PET scans in relation to cancer detection?

- A. They can identify early-stage diseases**
- B. They are only for brain imaging**
- C. They can reveal both structure and function**
- D. They cannot differentiate between tumor types**

The notion that PET scans cannot differentiate between tumor types is a common misconception. In reality, PET scans do provide valuable insights into the metabolic activity of tumors, which can assist in differentiating between various types of cancer. Different types of tumors often have distinct metabolic profiles, which can be captured in the PET imaging process. For example, certain tumors may demonstrate higher glucose metabolism than others, allowing physicians to infer not only the presence of cancer but also details about tumor aggressiveness or type. In contrast to this misconception, options related to early-stage disease detection, brain imaging, and the ability to reveal both structure and function are more aligned with the capabilities of PET imaging. PET scans indeed have limitations in delineating between tumor types solely based on metabolism and may require correlation with other imaging modalities or histopathological evaluation for accurate diagnosis.

7. Which quality of crystals in PET systems impacts their sensitivity?

- A. Higher density materials**
- B. Faster decay times**
- C. Stopping power**
- D. Scintillation photon yield**

The attribute of crystals in PET systems that primarily influences their sensitivity is their ability to produce a significant amount of scintillation light in response to incident gamma photons. This characteristic is known as scintillation photon yield. When a gamma photon interacts with the crystal, it excites the molecules within the crystal, leading to the emission of light (scintillation). A high scintillation photon yield ensures that more photons are produced for each gamma interaction, effectively increasing the detection capability of the system. Higher sensitivity in PET systems allows for better image quality and improved detection of smaller lesions, as more emitted light can lead to increased signal and improved signal-to-noise ratios. Therefore, while other qualities such as higher density materials or faster decay times may contribute to the overall performance of the crystals, the scintillation photon yield directly correlates with the sensitivity of the PET system, making it a crucial factor in enhancing image acquisition and diagnostics.

8. What is a primary limiting factor for PET imaging resolution?

- A. Crystal construction**
- B. Crystal stopping power**
- C. Photo multiplier tubes**
- D. Positron range**

The primary limiting factor for PET imaging resolution is the positron range. In positron emission tomography, a positron is emitted during the decay of a radioactive isotope, and this positron travels a short distance in tissue before it annihilates with an electron, resulting in the emission of two gamma photons. The distance a positron can travel before annihilation is known as the positron range, and this range is influenced by the type of tissue and the energy of the emitted positron. When positrons travel varying distances before annihilation, they lead to uncertainty in the exact location of the source of the radiation, thereby degrading the spatial resolution of the resulting PET image. Shorter ranges typically correspond to higher resolution, as the location of the annihilation event can be more accurately determined. Other factors, while they play roles in the overall performance of the PET system, do not significantly influence resolution in the same way. Crystal construction and stopping power relate to how well the system can detect gamma photons, while photomultiplier tubes are responsible for the detection and amplification of these signals. However, the physical characteristics of the emitted positrons and their interactions with matter primarily dictate the spatial resolution constraints in PET imaging.

9. How can obesity impact PET/CT imaging results?

- A. It has no impact on imaging results.
- B. It may improve radiotracer uptake.
- C. Increased body fat may alter distribution and uptake of the radiotracer.**
- D. It leads to better spatial resolution.

Obesity can significantly impact PET/CT imaging results, particularly through the alteration of the distribution and uptake of the radiotracer. In obese individuals, the larger body mass can affect how the radiotracers are distributed within the body. Adipose tissue influences the metabolic pathways and may alter the biodistribution of certain tracers, leading to variations in uptake that could affect diagnostic accuracy. For instance, certain radiotracers might accumulate differently in adipose versus lean tissue, which could complicate the interpretation of scans. It may also influence how well a radiotracer is taken up by various tissues, which is vital in accurately characterizing and diagnosing tumors or other conditions. Understanding these factors is crucial for radiologists and nuclear medicine technicians when interpreting PET/CT scans in patients with obesity. Adjustments in the interpretation of results may be necessary to account for the potential changes in tracer behavior due to increased body fat.

10. What does the term 'system dead time' refer to in PET imaging?

- A. The time taken for the system to reset after an event
- B. The period when the system cannot register incoming events**
- C. The duration for signal processing
- D. The time taken for scintillation light to dissipate

In PET imaging, the term 'system dead time' specifically refers to the period when the detection system is unable to register incoming events following the detection of a previous event. During this interval, any new gamma photon events that occur are lost because the system is not capable of processing them. This loss can lead to inaccuracies in the measurement of radioactivity, affecting image quality and quantitation. Understanding dead time is crucial for interpreting PET data, as it impacts the overall performance of the imaging system. If the time taken to process each event is too long, or if the system finds itself in a situation where it is constantly detecting events, it can lead to a significant number of missed detections, ultimately compromising both temporal resolution and sensitivity. Other options may correlate with different aspects of the imaging process, but they do not address the concept of 'system dead time' accurately as it relates to event registration.

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://petctfusion.examzify.com>

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

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