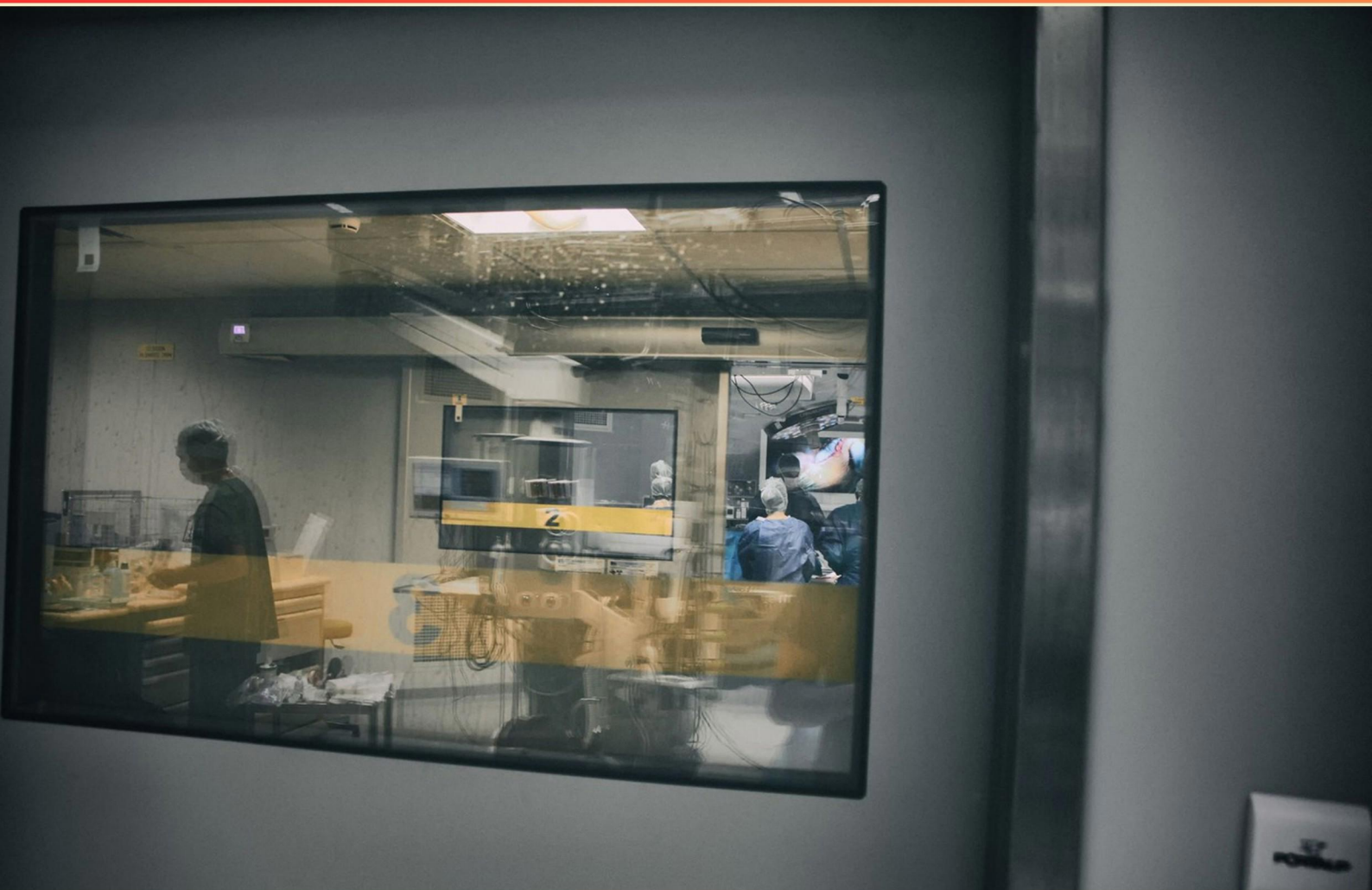


ACVIM SMALL ANIMAL INTERNAL MEDICINE (SAIM) SEMINAL PAPER PRACTICE EXAM (SAMPLE)

STUDY GUIDE



SAMPLE STUDY GUIDE WITH LIMITED QUESTIONS

VISIT US ONLINE FOR
THE FULL VERSION STUDY GUIDE

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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. Which of the following describes the fourth mucosal compartment?
 - A. Bacteria invasive within the mucosa
 - B. Bacteria circulating in the bloodstream
 - C. Bacteria within goblet cells
 - D. Bacteria in lymphoid tissue
2. After conjugated bile acids are secreted into the small intestine, what percentage is reabsorbed and where is the main site?
 - A. 70%, Jejunum
 - B. 90%, Ileum
 - C. 50%, Duodenum
 - D. 100%, Ileum
3. In Horn's Algorithm, the lower fence for outlier rejection is computed as:
 - A. $Q1 - 1.5 \times IQR$
 - B. $Q3 - 1.5 \times IQR$
 - C. $Q1 + 1.5 \times IQR$
 - D. $Q3 + 1.5 \times IQR$
4. Which description matches fast-growing endosteal bone formation?
 - A. Woven bone deposited perpendicular to trabecular surfaces, forming a web in intertrabecular space
 - B. Lamellar bone deposited on preexisting surfaces
 - C. Periosteal reaction along cortex
 - D. Cartilage formation within the marrow
5. What is the main location ASBT is found?
 - A. Stomach
 - B. Colon
 - C. Duodenum
 - D. Ileum

- 6. What is the recommended action if data are non-Gaussian and normality cannot be established after transformation?**
- A. Parametric methods cannot be used
 - B. Nonparametric methods cannot be used
 - C. Robust methods can be used only
 - D. Bootstrap methods are required
- 7. In ADDE, which description best reflects the effect on the tear film lipid layer?**
- A. Increased aqueous layer prevents adequate spreading of the lipid layer
 - B. Decreased aqueous layer prevents adequate spreading of the lipid layer
 - C. Decreased lipid layer thickness spreads
 - D. Tear film evaporation reduces
- 8. In the Garraway 2018 paper, which myeloid cell plays an important role in tumor progression and angiogenesis in colorectal cancer and was examined?**
- A. CD8+ T cells
 - B. Neutrophils
 - C. Monocyte-derived macrophages
 - D. Differentiated CD11b+
- 9. Why is radiographic surveillance valuable in cases where a bone biopsy returns a result of reactive bone?**
- A. It documents the biopsy technique used.
 - B. It indicates the biopsy was nondiagnostic.
 - C. It reconciles a potentially nondiagnostic biopsy with the lesion's ongoing imaging behavior over time.
 - D. It rules out malignancy.
- 10. Meibography is used to image which ocular structure?**
- A. Cornea
 - B. Lens
 - C. Meibomian glands
 - D. Retina

Answers

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

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Explanations

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1. Which of the following describes the fourth mucosal compartment?

- A. Bacteria invasive within the mucosa**
- B. Bacteria circulating in the bloodstream
- C. Bacteria within goblet cells
- D. Bacteria in lymphoid tissue

Think of bacteria as moving through a series of spaces from the lumen outward: free in the lumen, associated with the mucus on the surface, residing within epithelial cells, and finally taking up residence in the mucosal tissue itself. The fourth compartment refers to those organisms that have breached the epithelial barrier and are now inside the mucosa proper (the lamina propria/submucosa) where they can interact with immune cells. That's why invasive bacteria within the mucosa best fits as the fourth compartment. Bacteria circulating in the bloodstream are outside the mucosal layers entirely. Bacteria within goblet cells describe intracellular localization within a specific epithelial cell type but don't represent the broader mucosal-tissue compartment. Bacteria in lymphoid tissue correspond to immune tissue within the mucosa, not the mucosa itself as a barrier that has been breached.

2. After conjugated bile acids are secreted into the small intestine, what percentage is reabsorbed and where is the main site?

- A. 70%, Jejunum
- B. 90%, Ileum**
- C. 50%, Duodenum
- D. 100%, Ileum

Bile acids follow enterohepatic circulation, where after being secreted into the small intestine the vast majority are reabsorbed and returned to the liver. The terminal ileum is the primary site for this reabsorption of conjugated bile acids. About 90% are reabsorbed, helping to recycle bile acids efficiently and maintain the hepatic bile acid pool. In the ileal enterocytes, the apical transporter ASBT brings bile acids into the cell, and the basolateral OSTa/ β moves them into the portal circulation for return to the liver, where uptake occurs again by hepatic transporters. If ileal reabsorption is impaired, more bile acids reach the colon, causing diarrhea and a drop in bile acid pool, which the liver may compensate for by increasing synthesis. So, the best answer reflects that roughly 90% are reabsorbed, primarily in the ileum. The jejunum and duodenum reabsorb only a minority in comparison, and 100% reabsorption would be incorrect because a portion is normally lost in feces.

3. In Horn's Algorithm, the lower fence for outlier rejection is computed as:

- A. $Q1 - 1.5 \times IQR$**
- B. $Q3 - 1.5 \times IQR$
- C. $Q1 + 1.5 \times IQR$
- D. $Q3 + 1.5 \times IQR$

The idea is to define a range that separates the central 50% of the data from potential outliers using the interquartile range (IQR). Outliers are values that fall beyond the fences set 1.5 times the IQR away from the quartiles. For the lower side, you move 1.5 IQRs below the first quartile ($Q1$). That gives the lower fence as $Q1 - 1.5 \times IQR$. Any value smaller than that is considered a low outlier. The corresponding upper fence uses the third quartile plus $1.5 \times IQR$. So the correct expression for the lower fence is $Q1 - 1.5 \times IQR$. For example, if $Q1$ is 10 and IQR is 4, the lower fence is $10 - 1.5 \times 4 = 4$; values below 4 are flagged as outliers.

4. Which description matches fast-growing endosteal bone formation?

- A. Woven bone deposited perpendicular to trabecular surfaces, forming a web in intertrabecular space
- B. Lamellar bone deposited on preexisting surfaces
- C. Periosteal reaction along cortex
- D. Cartilage formation within the marrow

Rapid endosteal bone formation is typically composed of woven bone laid down quickly on inner bone surfaces. Woven bone has disorganized, randomly oriented collagen fibers and forms a dense, web-like network. When this occurs in trabecular bone, the new bone tends to be deposited perpendicular to existing surfaces, filling the intertrabecular spaces as a mesh. This rapid, scaffolding-like bone is then usually remodeled over time into the more organized lamellar bone. In contrast, lamellar bone is laid down in orderly, parallel layers on preexisting surfaces and represents slower, mature remodeling. A periosteal reaction occurs on the outer bone surface along the cortex, not endosteally. Cartilage formation within the marrow points to endochondral ossification, where a cartilage template is replaced by bone rather than direct endosteal deposition of bone tissue. So the description of woven bone forming a web in intertrabecular space on endosteal surfaces aligns with fast-growing endosteal bone formation.

5. What is the main location ASBT is found?

- A. Stomach
- B. Colon
- C. Duodenum
- D. Ileum

ASBT, the apical sodium-dependent bile acid transporter, is responsible for reclaiming bile acids from the gut lumen back into enterocytes. Its highest expression is in the ileum, especially the distal ileum, making that region the primary site of active bile acid reabsorption as part of the enterohepatic circulation. Most bile acids that reach the intestine are reabsorbed there and returned to the liver via the portal circulation. The stomach does not reabsorb bile acids, the duodenum reabsorbs only a small amount, and the colon mainly handles water absorption with minimal transporter-mediated bile acid uptake. So the main location is the ileum.

6. What is the recommended action if data are non-Gaussian and normality cannot be established after transformation?

- A. Parametric methods cannot be used
- B. Nonparametric methods cannot be used
- C. Robust methods can be used only
- D. Bootstrap methods are required

When data remain non-Gaussian and normality cannot be established after transformation, the assumptions that underlie parametric analyses are violated. Parametric methods rely on normality (or near-normality) of the data or residuals, so you cannot trust their results here. The appropriate path is to avoid parametric approaches and use alternatives that do not require strict normal distributions. Nonparametric tests are a natural choice because they do not assume a specific distribution. Robust methods can help with outliers or deviations but do not fix the fundamental normality requirement for all parametric tests, and bootstrap methods offer a resampling approach to inference without relying on normality, though they are not strictly required in every case.

7. In ADDE, which description best reflects the effect on the tear film lipid layer?

- A. Increased aqueous layer prevents adequate spreading of the lipid layer
- B. Decreased aqueous layer prevents adequate spreading of the lipid layer**
- C. Decreased lipid layer thickness spreads
- D. Tear film evaporation reduces

In aqueous-deficient dry eye the tear volume is reduced, so there is less aqueous surface for the lipid layer to spread over. The lipid layer sits on top of the aqueous layer and relies on that surface to spread evenly. When the aqueous layer is diminished, the lipid layer cannot distribute adequately, becoming uneven or patchy, which compromises the tear film's stability and promotes faster tear breakup and evaporation. This is why the statement describing decreased aqueous layer preventing adequate spreading of the lipid layer best fits the effect on the lipid layer in ADDE. Increasing the aqueous layer would improve spreading, decreasing lipid layer thickness isn't the primary description of this mechanism, and tear film evaporation is a consequence rather than a direct description of how the lipid layer spreads.

8. In the Garraway 2018 paper, which myeloid cell plays an important role in tumor progression and angiogenesis in colorectal cancer and was examined?

- A. CD8+ T cells
- B. Neutrophils
- C. Monocyte-derived macrophages
- D. Differentiated CD11b+**

Differentiated CD11b+ myeloid cells within the tumor microenvironment are key promoters of both tumor growth and new blood vessel formation in colorectal cancer. CD11b marks cells of the myeloid lineage (such as monocytes and macrophages that mature in the tumor context). When these cells differentiate in the tumor, they commonly adopt pro-angiogenic and pro-tumor functions, releasing VEGF, matrix metalloproteinases, and other factors that stimulate endothelial cells and remodeling of the surrounding tissue. The Garraway 2018 study specifically examined this differentiated CD11b+ population and found its involvement in driving angiogenesis and tumor progression in colorectal cancer, underscoring the significant role of this myeloid subset in shaping the tumor vasculature and growth.

9. Why is radiographic surveillance valuable in cases where a bone biopsy returns a result of reactive bone?

- A. It documents the biopsy technique used.
- B. It indicates the biopsy was nondiagnostic.
- C. It reconciles a potentially nondiagnostic biopsy with the lesion's ongoing imaging behavior over time.**
- D. It rules out malignancy.

When a bone biopsy shows reactive bone, the histology can be nondiagnostic or non-specific, making it hard to judge the true nature of the lesion from a single sample. Radiographic surveillance provides the longitudinal context by showing how the lesion behaves over time. If serial imaging remains stable or improves, that supports a benign, reactive process and helps avoid unnecessary further invasive testing. If the imaging changes—growth, new features, or progression—that raises concern for an underlying neoplasm or another pathology and prompts reevaluation, possible repeat biopsy, or additional workup. In short, imaging over time reconciles the uncertain biopsy result with what the lesion actually does on the body, guiding appropriate management.

10. Meibography is used to image which ocular structure?

- A. Cornea
- B. Lens
- C. Meibomian glands
- D. Retina

Meibography is imaging designed to visualize the Meibomian glands, which sit in the eyelid tarsal plates and produce the lipid layer of the tear film. Using infrared light, this method images the glands through the eyelids to assess their appearance, including dropout, atrophy, or distortion that can occur with meibomian gland dysfunction. This is particularly helpful for diagnosing evaporative dry eye related to MG dysfunction. The cornea, lens, and retina are examined with other modalities or structural assessments (such as slit-lamp examination with corneal assessment, lens imaging, or retinal fundus imaging), not with meibography. Therefore, the structure specifically imaged by meibography is the Meibomian glands.

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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:

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

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