

Infection and Response 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. Why should you finish a course of antibiotics?**
 - A. So that all bacteria are killed and no resistant strains remain**
 - B. So the medicine lasts longer**
 - C. To avoid taking other medicines**
 - D. Because it tastes better**

- 2. Which gene type commonly mediates resistance to beta-lactam antibiotics such as penicillin?**
 - A. Efflux pumps**
 - B. Carbohydrate modification**
 - C. Beta-lactamase genes**
 - D. Capsule formation genes**

- 3. What do white blood cells form after encountering a pathogen for the first time?**
 - A. Red blood cells.**
 - B. Platelets.**
 - C. Memory cells.**
 - D. Antibody-producing plasma cells.**

- 4. Why does an influenza vaccine require annual updates?**
 - A. Antigenic drift changes surface proteins targeted by vaccines.**
 - B. The virus mutates in predictable cycles unrelated to surface proteins.**
 - C. Annual updates are required to adjust for changes in manufacturing methods.**
 - D. Influenza vaccines aim to cover all subtypes in one shot.**

- 5. Whether or not a drug harms our body cells**
 - A. Toxicity**
 - B. Efficacy**
 - C. Dose**
 - D. Shelf life**

- 6. How is TMV spread?**
- A. Direct contact between plants**
 - B. Waterborne spread**
 - C. Airborne spores**
 - D. Soil contamination**
- 7. Which practice increases risk of HIV transmission?**
- A. Sharing needles during drug use**
 - B. Regular hand washing**
 - C. Vaccination**
 - D. Eating raw meat**
- 8. What is a non-communicable disease?**
- A. A disease that spreads rapidly.**
 - B. A disease that can be passed from person to person.**
 - C. A disease that cannot be passed from person to person.**
 - D. A disease that is caused by viruses only.**
- 9. Which of the following is a type of pathogen not bacteria, virus, or fungi?**
- A. Algae**
 - B. Virus**
 - C. Protist**
 - D. Bacteria**
- 10. How does vaccination provide memory without causing disease?**
- A. Vaccines cause disease to stimulate memory.**
 - B. Vaccines present antigens in a safe form to stimulate adaptive immunity, forming memory B and T cells without causing disease.**
 - C. Vaccines only produce temporary antibodies with no memory.**
 - D. Vaccines rely on innate immunity alone.**

Answers

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

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Explanations

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1. Why should you finish a course of antibiotics?

- A. So that all bacteria are killed and no resistant strains remain**
- B. So the medicine lasts longer**
- C. To avoid taking other medicines**
- D. Because it tastes better**

Finishing a course of antibiotics helps ensure the infection is truly cleared and reduces the chance of resistance developing. Antibiotics don't kill every bacterium instantly; some may survive as the drug levels wane. If you stop early, those survivors can regrow, causing the infection to come back and potentially spreading. Moreover, survival of bacteria that faced the antibiotic creates selective pressure for resistant strains, making future infections harder to treat. That's why completing the prescribed course is important, even if you start feeling better. It's not about making the medicine last longer, avoiding other medicines, or taste. If you feel unwell or have side effects, talk to a clinician rather than stopping early.

2. Which gene type commonly mediates resistance to beta-lactam antibiotics such as penicillin?

- A. Efflux pumps**
- B. Carbohydrate modification**
- C. Beta-lactamase genes**
- D. Capsule formation genes**

Bacteria commonly resist beta-lactam antibiotics through enzymatic inactivation. Beta-lactams, like penicillin, disrupt cell wall synthesis by blocking penicillin-binding proteins. Bacteria counter this by producing beta-lactamase enzymes that break open the beta-lactam ring, instantly neutralizing the drug. The genes encoding these enzymes—beta-lactamase genes—are often carried on transferable elements, which helps resistance spread among bacteria. Other mechanisms, such as efflux pumps, can contribute to resistance in some contexts by pumping drugs out of the cell, but they are not the primary way penicillins are inactivated. Modifications to surface carbohydrates or capsule formation relate more to virulence or immune evasion than to directly disabling beta-lactam antibiotics.

3. What do white blood cells form after encountering a pathogen for the first time?

- A. Red blood cells.**
- B. Platelets.**
- C. Memory cells.**
- D. Antibody-producing plasma cells.**

When the immune system first meets a pathogen, a branch of white blood cells called B cells is activated and can differentiate into two major outcomes. Some become plasma cells that rapidly produce antibodies to neutralize the pathogen right away. Others become memory cells that persist long term. These memory cells “remember” the specific pathogen so that if the same invader returns, the immune system can respond much faster and more effectively, often preventing illness. That’s why memory cells are the best choice here: they are the lasting product that provides future protection after the initial encounter. Red blood cells and platelets aren’t involved in forming immune memory, and while plasma cells do produce antibodies during the first response, memory cells are specifically the long-lasting form that remains for future encounters.

4. Why does an influenza vaccine require annual updates?

- A. Antigenic drift changes surface proteins targeted by vaccines.**
- B. The virus mutates in predictable cycles unrelated to surface proteins.**
- C. Annual updates are required to adjust for changes in manufacturing methods.**
- D. Influenza vaccines aim to cover all subtypes in one shot.**

Influenza vaccines are updated yearly because the virus changes its surface proteins through antigenic drift. The immune response from previous vaccines targets those surface proteins, mainly hemagglutinin, so small mutations can let the virus slip past antibodies. To keep protection as high as possible, scientists review global flu activity and reformulate the vaccine to match the strains predicted to be most common in the upcoming season. It’s not about changes in manufacturing methods, and vaccines don’t try to cover all subtypes in one shot—the goal is to align protection with the strains most likely to circulate.

5. Whether or not a drug harms our body cells

A. Toxicity

B. Efficacy

C. Dose

D. Shelf life

Toxicity describes the potential of a drug to harm body cells or tissues. It focuses on adverse effects and damage that can range from mild to severe, reflecting how a drug can be harmful even as it is used to achieve a benefit. This is different from efficacy, which is about whether the drug produces the desired therapeutic effect; dose, which is simply how much is given; and shelf life, which concerns how long the drug remains stable. So, when the question asks about whether a drug harms our body cells, toxicity is the term that directly captures that risk. Examples include liver toxicity from an overdose, kidney toxicity from certain medications, or bone marrow suppression from chemotherapy, illustrating how toxicity centers on harm rather than the intended benefit.

6. How is TMV spread?

A. Direct contact between plants

B. Waterborne spread

C. Airborne spores

D. Soil contamination

TMV spreads mainly through mechanical transmission, meaning virus particles move from an infected plant to a healthy one via contact with sap. This happens when plant tissue is damaged or handled with contaminated hands or tools—pruning, grafting, or even brushing against plants with dirty gloves can transfer the virus. The virus is very stable and can remain infective on surfaces for a long time, so cleaning and disinfecting tools and hands is essential to prevent spread. Waterborne spread isn't the primary route; while sap can move in water, irrigation or splash water is not the main way TMV travels between plants in typical settings. Airborne spread in the form of spores isn't applicable here since TMV is a virus without spores and isn't usually dispersed by wind. Soil contact alone doesn't directly infect plants unless the virus-bearing sap reaches a wound on the plant; soil mainly acts as a reservoir. So the most accurate description is that direct contact between plants, via contaminated hands or tools, best explains how TMV is spread.

7. Which practice increases risk of HIV transmission?

- A. Sharing needles during drug use**
- B. Regular hand washing**
- C. Vaccination**
- D. Eating raw meat**

HIV transmission occurs mainly through exposure to infected blood and certain body fluids, so activities that involve direct blood-to-blood contact or sharing infected fluids raise risk. Sharing needles during drug use creates a direct blood-to-blood pathway: if a needle or syringe carries HIV, using it transfers the virus from one person's blood to another, creating a high-risk route for infection with each use. Regular hand washing helps prevent many infections but doesn't establish a pathway for HIV spread, since casual contact and routine hygiene do not transmit the virus. Vaccination involves a clean, controlled exposure to a vaccine and is not a route for HIV transmission. Eating raw meat is not a transmission route for HIV either, as the virus does not survive in typical food handling once cooked and is not spread through eating. So, the practice that increases HIV transmission risk is sharing needles during drug use because it directly introduces blood-to-blood exposure.

8. What is a non-communicable disease?

- A. A disease that spreads rapidly.**
- B. A disease that can be passed from person to person.**
- C. A disease that cannot be passed from person to person.**
- D. A disease that is caused by viruses only.**

Non-communicable diseases are conditions that are not infectious and cannot be transmitted from one person to another. They are usually chronic, developing slowly and lasting a long time, and arise from a mix of factors such as lifestyle, genetics, and aging. Examples include heart disease, diabetes, cancer, and chronic lung conditions. Because these diseases aren't caused by an infectious agent that spreads between people, they don't pass from person to person. The other ideas describe infectious diseases or overly specific causes (like viruses only), which don't define non-communicable diseases.

9. Which of the following is a type of pathogen not bacteria, virus, or fungi?

- A. Algae**
- B. Virus**
- C. Protist**
- D. Bacteria**

Pathogens come from different groups, and among common disease-causing organisms, protists are a diverse set of single-celled eukaryotes that can cause illness in humans. This means protists are a type of pathogen even though they're not bacteria (prokaryotes), not viruses (acellular), and not fungi (a separate kingdom). For example, malaria is caused by Plasmodium, a protist, and amebic dysentery is caused by Entamoeba histolytica, illustrating that protists can be pathogenic. Algae are mostly photosynthetic organisms and aren't typical disease-causing pathogens in humans, while viruses are pathogens but belong to a different category altogether. So the best answer is protists.

10. How does vaccination provide memory without causing disease?

A. Vaccines cause disease to stimulate memory.

B. Vaccines present antigens in a safe form to stimulate adaptive immunity, forming memory B and T cells without causing disease.

C. Vaccines only produce temporary antibodies with no memory.

D. Vaccines rely on innate immunity alone.

Vaccination trains the adaptive immune system by presenting safe forms of pathogen antigens so the body can learn without getting sick. When these antigens appear, B cells respond by making antibodies and some become memory B cells, while T cells (helper and cytotoxic) form memory T cells. This creates a ready pool of antigen-specific cells that respond rapidly if the real pathogen is encountered later, producing antibodies and cellular responses quickly to prevent disease or lessen its severity. The safe forms used in vaccines—such as inactivated or attenuated organisms, subunit components, toxoids, or genetic vaccines that instruct cells to make the antigen—allow this memory to form without causing illness. Because vaccines are designed to avoid disease while still provoking a targeted, durable memory, they don't rely on causing infection, they don't depend on innate immunity alone, and they don't provide only temporary antibodies without lasting memory.

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://infectionandresponse.examzify.com>

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

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