

Medical Laboratory Professionals' Association of Ontario (MLPAO) Practice Exam (Sample)

Study Guide



Everything you need from our exam experts!

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SAMPLE

Questions

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- 1. What type of laboratory is equipped to handle pathogens that pose a moderate risk to personnel and the environment?**
 - A. Biosafety Level 1.**
 - B. Biosafety Level 2.**
 - C. Biosafety Level 3.**
 - D. Biosafety Level 4.**
- 2. What does the Coombs test screen for?**
 - A. Hemolytic anemia and blood type compatibility**
 - B. Diabetes and metabolic syndrome**
 - C. Coagulation disorders**
 - D. Immunodeficiency diseases**
- 3. Which is NOT a requirement of the Transportation of Dangerous Goods Regulation when shipping dangerous substances?**
 - A. A completed "shippers declaration" document**
 - B. Emergency contact information**
 - C. Specific package labeling and required hazard symbols**
 - D. Inclusion of "dry ice" in a sealed container**
- 4. What does a positive human chorionic gonadotropin (hCG) test indicate?**
 - A. Possible pregnancy or certain types of tumors**
 - B. Detection of hormonal imbalances**
 - C. Presence of genetic disorders**
 - D. Indicators of liver function**
- 5. In a blood transfusion context, which antigen typing is critical for preventing incompatibility reactions?**
 - A. Lewis**
 - B. Kell**
 - C. ABO**
 - D. Rh**

- 6. Which type of virus is detected using polymerase chain reaction (PCR) testing?**
- A. Only respiratory viruses**
 - B. Various viruses including HIV, influenza, and COVID-19**
 - C. Only bacterial infections**
 - D. Parasites and fungal infections**
- 7. Which risk is not associated with the WHMIS symbol described?**
- A. May cause environmental damage from fumes**
 - B. May react with water**
 - C. May be chemically unstable**
 - D. May burn unexpectedly**
- 8. At what temperature is sheep blood added to molten agar to produce chocolate agar?**
- A. 40-45°C**
 - B. 50-55°C**
 - C. 60-65°C**
 - D. 75-80°C**
- 9. Which statement about thermometers is incorrect?**
- A. The size is important**
 - B. Calibration must be verified**
 - C. Either alcohol or mercury is used as the indicator**
 - D. Records must be kept**
- 10. How many minutes should a urine specimen be centrifuged for microscopy analysis?**
- A. 3 minutes**
 - B. 5 minutes**
 - C. 7 minutes**
 - D. 10 minutes**

Answers

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

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Explanations

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1. What type of laboratory is equipped to handle pathogens that pose a moderate risk to personnel and the environment?

- A. Biosafety Level 1.**
- B. Biosafety Level 2.**
- C. Biosafety Level 3.**
- D. Biosafety Level 4.**

Biosafety Level 2 laboratories are specifically designed to handle pathogens that pose a moderate risk to personnel and the environment. At this level, standard laboratory practices and procedures are employed, along with specific safety equipment and protocols to mitigate the risk of exposure. This includes the use of personal protective equipment, such as gloves and lab coats, and implementing techniques to avoid generating aerosols that could spread infectious agents. Pathogens that may be handled in a Biosafety Level 2 laboratory include those that can cause disease in humans but are typically not transmitted by aerosols, making them less risky than those classified under higher biosafety levels. The nature of the containment equipment and safety measures in BSL-2 labs ensures that laboratory workers can safely conduct their work, while also protecting the surrounding environment from potential contamination. In contrast, the other biosafety levels correspond to varying degrees of risk associated with the pathogens they contain, each requiring more stringent safety measures and specialized equipment as the risk level increases. Thus, Biosafety Level 2 fulfills the requirement for handling moderate-risk pathogens effectively.

2. What does the Coombs test screen for?

- A. Hemolytic anemia and blood type compatibility**
- B. Diabetes and metabolic syndrome**
- C. Coagulation disorders**
- D. Immunodeficiency diseases**

The Coombs test is specifically designed to detect antibodies that may be present on the surface of red blood cells. This is particularly important in the context of hemolytic anemia, which can occur when the immune system mistakenly targets and destroys red blood cells. The test can either be direct or indirect. The direct Coombs test identifies antibodies that are bound to red blood cells in the patient's bloodstream, helping to diagnose conditions such as autoimmune hemolytic anemia. The indirect Coombs test, on the other hand, detects free antibodies in the serum and is often used to assess blood type compatibility prior to blood transfusions or organ transplants, making it crucial for ensuring patient safety during procedures that require blood products. Therefore, the Coombs test's primary applications in diagnosing hemolytic anemia and ensuring blood type compatibility make the first option correct. The other choices do not relate closely to the functions of the Coombs test and address different medical conditions or screening methods.

3. Which is NOT a requirement of the Transportation of Dangerous Goods Regulation when shipping dangerous substances?

- A. A completed "shippers declaration" document**
- B. Emergency contact information**
- C. Specific package labeling and required hazard symbols**
- D. Inclusion of "dry ice" in a sealed container**

The correct choice is based on the requirements set forth by the Transportation of Dangerous Goods Regulation, which outlines specific criteria for the safe shipment of dangerous substances. Inclusion of "dry ice" in a sealed container is indeed a necessary procedure for ensuring that dry ice, categorized as a dangerous good due to its sublimation into carbon dioxide gas, is transported safely. However, the regulation allows for certain exemptions or conditions under which dry ice can be shipped, including not needing to be placed in completely sealed containers as long as the packaging is designed to allow the gas to escape. On the other hand, the other requirements—having a completed "shipper's declaration" document, providing emergency contact information, and ensuring specific package labeling and hazard symbols—are critical and mandatory elements in the transportation of dangerous goods. The shipper's declaration formally indicates that the goods being shipped are classified as dangerous, emergency contact information ensures that immediate assistance can be obtained if needed, and proper labeling and hazard symbols visually communicate the risks associated with the material to handlers and emergency responders. Thus, while careful handling and transportation is critical, the requirement pertaining to dry ice does not align strictly with the overarching regulations as it maintains flexibility under certain stipulated conditions.

4. What does a positive human chorionic gonadotropin (hCG) test indicate?

- A. Possible pregnancy or certain types of tumors**
- B. Detection of hormonal imbalances**
- C. Presence of genetic disorders**
- D. Indicators of liver function**

A positive human chorionic gonadotropin (hCG) test primarily indicates the presence of hCG in the bloodstream, which is most commonly associated with pregnancy. After fertilization occurs, the developing placenta begins to produce hCG shortly after implantation, making it a reliable biomarker for confirming pregnancy. In addition to indicating pregnancy, elevated hCG levels can also suggest certain types of tumors, such as gestational trophoblastic disease or specific testicular cancers. This means that when evaluating a positive hCG result, a healthcare provider may consider both pregnancy and potential malignancies as possible causes. The other choices do not correctly align with what a positive hCG test signifies. While hormonal imbalances may be assessed through other tests, they are not related to hCG specifically. Additionally, genetic disorders and liver function indicators are outside the scope of what an hCG test measures; thus, they do not apply in this context.

5. In a blood transfusion context, which antigen typing is critical for preventing incompatibility reactions?

- A. Lewis
- B. Kell
- C. ABO**
- D. Rh

In the context of blood transfusions, the ABO antigen typing is critical for preventing incompatibility reactions. The ABO blood group system consists of antigens A and B, and individuals can have type A, type B, type AB, or type O blood based on the presence or absence of these antigens. When a person receives blood that contains antigens foreign to their own, the immune system may recognize these antigens as harmful, leading to a potentially life-threatening hemolytic transfusion reaction. Ensuring that the donor's and recipient's blood types are compatible is essential, which is why ABO typing precedes any transfusion. For example, a person with type A blood should not receive type B or AB blood, as their immune system would attack the foreign B antigens, resulting in serious complications. While other antigen typings, such as Rh factor, are also important in the context of blood transfusions (particularly in avoiding Rh incompatibility during pregnancy or subsequent transfusions), the initial compatibility primarily hinges on the ABO blood group system. Therefore, ABO typing directly addresses the most immediate compatibility issues encountered in blood transfusions, making it a critical focus for preventing adverse reactions.

6. Which type of virus is detected using polymerase chain reaction (PCR) testing?

- A. Only respiratory viruses
- B. Various viruses including HIV, influenza, and COVID-19**
- C. Only bacterial infections
- D. Parasites and fungal infections

Polymerase chain reaction (PCR) testing is a widely used molecular technique that amplifies specific segments of DNA or RNA, allowing for the detection of various pathogens. One of the significant advantages of PCR is its ability to identify viruses with high sensitivity and specificity, making it valuable in clinical diagnostics. When it comes to viruses, PCR can be used to detect a range of viral infections, including but not limited to HIV, influenza viruses, coronaviruses like SARS-CoV-2 (responsible for COVID-19), and many others. This versatility stems from PCR's capacity to target the genetic material of viruses directly, enabling quick and accurate diagnosis even in cases with low viral loads. In contrast, the other options presented do not encompass the range of pathogens that PCR can effectively detect. Limiting PCR testing solely to respiratory viruses excludes its application for numerous other viral infections. Suggesting that PCR is specific only to bacterial infections or to parasites and fungal infections is misleading, as traditional PCR is not typically the method of choice for these types of pathogens. Rather, PCR's broad applicability to various viruses is what makes it a critical tool in modern virology, underscoring the correctness of recognizing its use in detecting a wide range of viral pathogens.

7. Which risk is not associated with the WHMIS symbol described?

- A. May cause environmental damage from fumes**
- B. May react with water**
- C. May be chemically unstable**
- D. May burn unexpectedly**

The correct choice, which indicates that the risk is not associated with the WHMIS symbol, focuses on the symbol's specific implications. WHMIS, or the Workplace Hazardous Materials Information System, uses distinct symbols to quickly communicate the hazards of various substances. Each symbol corresponds to certain risks based on the properties of the materials they represent. The option stating "May cause environmental damage from fumes" does not align with the typical hazards indicated by most WHMIS symbols. While it's crucial to consider environmental impacts in a broader context, the primary symbols under WHMIS usually center on immediate health and safety risks in the workplace, such as flammability, toxicity, or reactivity, rather than environmental issues stemming from fumes. In contrast, the other options all reflect common risks associated with hazardous materials. The possibility of a substance reacting with water, being chemically unstable, or unexpectedly burning are all critical concerns directly related to the properties and behavior of specific substances as indicated by their corresponding WHMIS symbols. This highlights the importance of understanding not just the symbols, but also their specific indications regarding risks to ensure a safe working environment.

8. At what temperature is sheep blood added to molten agar to produce chocolate agar?

- A. 40-45°C**
- B. 50-55°C**
- C. 60-65°C**
- D. 75-80°C**

The appropriate temperature for adding sheep blood to molten agar for the production of chocolate agar is within the range of 40-45°C. At this temperature, the blood cells can lyse, which is crucial for forming chocolate agar. The heat from the agar is sufficient to promote the lysis of red blood cells, releasing hemoglobin and giving the medium its characteristic brown color. When blood is added at too high a temperature, such as in the range of 50-55°C or above, it risks denaturing the proteins and enzymes within the blood, which interferes with the intended use of the medium for growing specific bacteria that require the nutrients released from the lysed red cells. Thus, for producing chocolate agar effectively, it is essential to control the temperature to be between 40-45°C to ensure optimal growth conditions and nutrient availability for the microorganisms being cultured.

9. Which statement about thermometers is incorrect?

- A. The size is important
- B. Calibration must be verified
- C. Either alcohol or mercury is used as the indicator**
- D. Records must be kept

The statement regarding the use of either alcohol or mercury as an indicator in thermometers is incorrect. While both alcohol and mercury have historically been used in thermometers, the use of mercury has declined significantly due to safety and environmental concerns. Many modern thermometers utilize digital sensors, bimetallic strips, or other non-toxic substances instead of mercury or alcohol. When discussing thermometers, the importance of size pertains to their application; certain sizes are suited for specific tasks or environments. Calibration verification is critical to ensure accurate measurements, especially in laboratory settings where precision is vital. Keeping records of temperature readings is also essential for traceability and compliance with safety standards and regulations, particularly in medical and laboratory environments.

10. How many minutes should a urine specimen be centrifuged for microscopy analysis?

- A. 3 minutes
- B. 5 minutes**
- C. 7 minutes
- D. 10 minutes

For microscopy analysis of a urine specimen, centrifugation is typically performed for 5 minutes. This duration is considered optimal as it allows for sufficient sedimentation of the formed elements in the urine, such as red blood cells, white blood cells, epithelial cells, and crystals, without causing excessive disruption or lysing of these components. Centrifugation for this period strikes a balance, ensuring that all relevant elements are settled at the bottom of the tube, which can then be analyzed under the microscope. If centrifugation is shorter, the sediment may not adequately form, leading to potential loss of diagnostic information. On the other hand, longer centrifugation times beyond 5 minutes can lead to excessive pelleting and may cause the formation of artifacts, which could misinterpret the microscopic findings. Therefore, the 5-minute mark is a standard practice in laboratory settings to ensure accurate and reliable results during urine microscopy analysis.