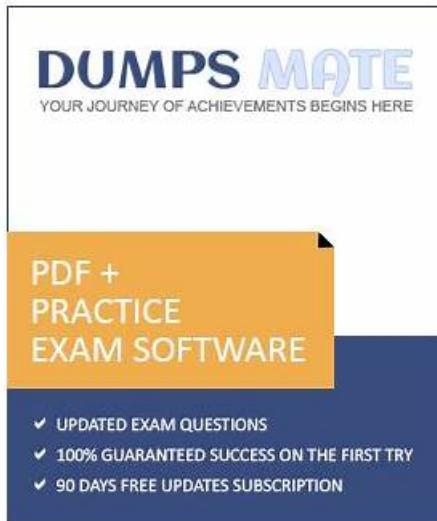


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American Society of Microbiology ABMM Sample Questions (Q202-Q207):

NEW QUESTION # 202

A traveler returning from sub-Saharan Africa presents with cyclical fevers, chills, and sweats. A Giemsa- stained peripheral blood smear reveals multiple ring forms within red blood cells, some applique forms, and occasional banana-shaped gametocytes. No Schüffner's dots or red blood cell enlargement is observed. Which *Plasmodium* species is the most likely cause?

- A. *Plasmodium malariae*
- B. *Plasmodium ovale*
- C. *Plasmodium falciparum*
- D. *Plasmodium vivax*

Answer: C

NEW QUESTION # 203

A patient develops a vesicular rash on their hands and feet, accompanied by fever and oral ulcers. The MOST likely causative agent is:

- A. Measles virus
- B. Varicella-zoster virus
- C. Herpes simplex virus
- D. Coxsackievirus A

Answer: D

NEW QUESTION # 204

Under the Clinical Laboratory Improvement Amendments (CLIA), participation in an approved external Proficiency Testing (PT) program serves primarily to:

- A. Provide an objective, ongoing assessment of the accuracy and reliability of the laboratory's testing compared to its peers.
- B. Fulfill requirements for laboratory accreditation by The Joint Commission only.
- C. Validate the performance characteristics (e.g., sensitivity, specificity) of newly implemented test methods.
- D. Determine the reference intervals (normal ranges) for quantitative assays performed by the laboratory.

Answer: A

NEW QUESTION # 205

A clinical microbiology laboratory is using flow cytometry to analyze lymphocyte subsets in a patient with suspected immunodeficiency. Which of the following cell surface markers is typically used to identify helper T cells?

- A. CD4
- B. CD19
- C. CD8
- D. CD56

Answer: A

NEW QUESTION # 206

A patient with a history of recurrent sinusitis and bronchitis is diagnosed with cystic fibrosis. Sputum cultures repeatedly grow a Gram-negative rod that is oxidase-positive and produces a blue-green pigment. This organism's chronic colonization of the respiratory tract in cystic fibrosis patients is primarily due to its ability to:

- A. Evade phagocytosis by alveolar macrophages through capsule production.
- B. Exhibit rapid intracellular replication within respiratory epithelial cells.
- C. Produce potent exotoxins that damage the ciliated epithelium.
- D. Form a biofilm that protects it from host defenses and antibiotics.

Answer: D

NEW QUESTION # 207

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