

ABMM Exam Lab Questions- Latest ABMM Guide Pass Success



American Board of Medical Microbiology (ABMM) Exam

Table of Contents

Eligibility Requirements	2
PLAN I:	2
PLAN II:	2
PLAN III:	2
EDUCATION	2
EXPERIENCE	2
NON-CPE RESIDENCIES	4
Exam Information	5
OBJECTIVE	5
EXAM FORMAT	5
ON EXAM DAY	5
RESPONSIBILITIES AND ROSTER OF SUBCOMMITTEES	5
ABOUT THE EXAM QUESTIONS	6
EXAM QUESTION (ITEM) DEVELOPMENT STAGES	8
SCORING	9
CRITERION-REFERENCING	9
DISCRIMINATION	10
CUT (PASSING) SCORE	10
RESULTS	10
PASS RATE (2016-2021)	11
FAQs	11
Preparing for the Exam	15
STUDY SUGGESTIONS	15
EXAM CONTENT: UPDATED JANUARY 2019	17
SAMPLE QUESTIONS	22
Contact Us	26

1

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American Society of Microbiology ABMM Sample Questions (Q10-Q15):

NEW QUESTION # 10

A clinical microbiology laboratory is preparing to validate a new molecular assay for the detection of methicillin-resistant *Staphylococcus aureus* (MRSA). To assess the positive predictive value (PPV) of the assay, they need to know:

- A. The number of true negative results divided by the total number of negative test results.
- **B. The number of true positive results divided by the total number of positive test results.**
- C. The number of true negative results divided by the total number of true negative and false positive results.
- D. The number of true positive results divided by the total number of true positive and false negative results.

Answer: B

NEW QUESTION # 11

A patient develops a localized skin infection after being bitten by a tick. The infection is characterized by a slowly expanding erythematous rash with central clearing (erythema migrans). The MOST likely causative agent is:

- A. *Anaplasma phagocytophilum*
- B. *Ehrlichia chaffeensis*
- **C. *Borrelia burgdorferi***
- D. *Rickettsia rickettsii*

Answer: C

NEW QUESTION # 12

A microbiology laboratory isolates a Gram-negative rod from a wound infection. The organism is oxidase- negative, ferments lactose on MacConkey agar, and produces indole. The MOST likely identification is:

- A. *Serratia marcescens*
- **B. *Escherichia coli***
- C. *Proteus mirabilis*
- D. *Klebsiella pneumoniae*

Answer: B

NEW QUESTION # 13

A traveler returning from Southeast Asia presents with fever, headache, and a maculopapular rash that started on the trunk and spread to the extremities, including the palms and soles. Serological testing is positive for *Orientia tsutsugamushi*. The vector responsible for transmitting this organism is a:

- A. Flea
- B. Mosquito
- **C. Mite (chigger)**
- D. Tick

Answer: C

NEW QUESTION # 14

A clinical microbiology laboratory is validating a new point-of-care test for the rapid detection of influenza A and B viruses. To accurately determine the clinical sensitivity of the test, it should be compared to a gold standard method using:

- A. A panel of known positive and negative samples stored at room temperature.
- B. Spiked samples containing known concentrations of influenza antigens.
- C. Samples from patients with other respiratory viral infections to assess cross-reactivity.
- **D. A large number of prospectively collected clinical samples from patients with suspected influenza, with results confirmed by RT-PCR.**

Answer: D

NEW QUESTION # 15

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