

Topic 'Transitioning DNA Analytic Devices and Technologies to the Field'

20 multiple-choice questions (MCQs) covering the key concepts. The questions range from basic recall to application and understanding.

Quiz: DNA Analytic Technologies for Field Applications

1. What is the main advantage of portable qPCR systems for field applications?

- A. They require a full laboratory setup.
- B. They eliminate the need for DNA extraction.
- C. They enable on-site DNA testing with rapid results.
- D. They can only detect bacterial DNA.

2. Why are portable glove boxes used during field DNA testing?

- A. To increase PCR speed.
- B. To prevent aerosol DNA contamination.
- C. To store DNA samples.
- D. To cool PCR reagents.

3. Which characteristic of field DNA extraction is considered most important?

- A. Highest DNA purity possible
- B. Long DNA fragments
- C. Speed, simplicity, and sufficient DNA quality
- D. Use of hazardous chemicals

4. Approximately how long can the complete field workflow (DNA extraction → PCR → result) be completed?

- A. 15 minutes
- B. 1 hour
- C. 3 hours
- D. Overnight

5. Which of the following is NOT a commonly used rapid DNA extraction method?

- A. Chelex® extraction
- B. Magnetic bead extraction
- C. Phenol-chloroform extraction
- D. Heat lysis

6. Why is high-purity DNA generally unnecessary for portable PCR systems?

- A. Portable PCR does not amplify DNA.
- B. Portable PCR amplifies short DNA fragments that tolerate partially degraded DNA.
- C. Portable PCR only detects RNA.
- D. DNA purity has no effect on PCR.

7. What is the primary purpose of fuzzy logic algorithms in field-deployed qPCR?

- A. Increase DNA extraction efficiency.
- B. Design PCR primers.
- C. Automatically interpret amplification results and reduce false positives.
- D. Increase PCR temperature.

8. Which technology amplifies DNA without repeated thermal cycling?

- A. Conventional PCR
- B. Sanger sequencing
- C. Isothermal amplification
- D. Gel electrophoresis

9. Recombinase Polymerase Amplification (RPA) typically operates at approximately:

- A. 25°C
- B. 39°C
- C. 63°C
- D. 95°C

10. Which statement best describes an advantage of RPA?

- A. Requires expensive laboratory equipment.
- B. Produces results in 10–20 minutes.
- C. Requires multiple thermal cycles.
- D. Requires ultra-pure DNA.

11. In the RPA-NALFIA system, which labels are attached to the primers?

- A. FITC and DAPI
- B. FAM and Biotin
- C. SYBR Green and Ethidium Bromide
- D. Gold nanoparticles and Cy5

12. A positive RPA lateral flow test is indicated by:

- A. A fluorescent glow only.
- B. Formation of bubbles.
- C. Appearance of a visible dark line on the test strip.
- D. Complete disappearance of the sample.

13. Loop-Mediated Isothermal Amplification (LAMP) generally operates at:

- A. 37°C
- B. 39°C
- C. 50°C
- D. 63°C

14. Why is LAMP considered highly specific?

- A. It amplifies RNA only.
- B. It uses six different polymerases.
- C. It employs 4–6 primers targeting multiple regions.
- D. It requires DNA sequencing.

15. Which enzyme is responsible for DNA synthesis in LAMP?

- A. Taq DNA polymerase
- B. Reverse transcriptase
- C. Bst DNA polymerase
- D. Cas12

16. How can LAMP results be detected visually?

- A. DNA bands on agarose gel only
- B. Blue-to-red fluorescence
- C. Color change of Hydroxynaphthol Blue (HNB)
- D. Silver staining

17. In CRISPR-based detection, what determines the specificity of target recognition?

- A. DNA polymerase
- B. Magnesium ions
- C. Guide RNA (gRNA)
- D. Ethanol

18. What happens after Cas12 recognizes the target porcine DNA?

- A. DNA replication stops.
- B. Cas12 performs collateral cleavage of nearby single-stranded DNA reporters.
- C. The DNA immediately degrades.
- D. PCR begins automatically.

19. Which combination is commonly used to improve CRISPR detection sensitivity?

- A. PCR followed by sequencing
- B. RPA or LAMP before CRISPR detection
- C. Gel electrophoresis before DNA extraction
- D. Western blot before PCR

20. According to the presentation, what is the recommended dual-tiered testing strategy for halal food authentication?

- A. Use PCR only for every sample.
- B. Use microscopy followed by culture.
- C. Use RPA/LAMP/CRISPR for rapid screening and qPCR for confirmation.
- D. Use DNA sequencing for every sample.

Bonus Higher-Order Thinking Question

21. A food inspector performing on-site halal verification requires a DNA detection method that is fast, battery-powered, requires minimal equipment, and provides results visible to the naked eye. Which technology would be the most suitable?

- A. Conventional laboratory qPCR
- B. RPA coupled with NALFIA lateral flow detection
- C. Sanger DNA sequencing
- D. Whole genome sequencing