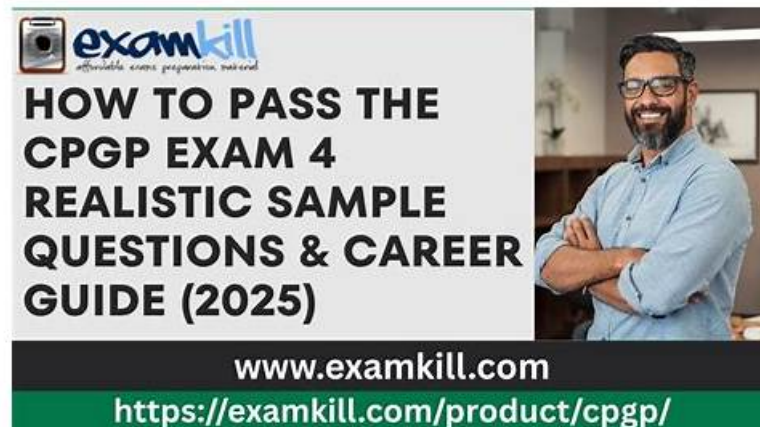


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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q324-Q329):

NEW QUESTION # 324

The purpose of retaining reserve samples is to:Response:

- A. Meet regulatory requirements and enable future testing if needed
- B. Use them for training new laboratory personnel
- C. Ensure that there is always an excess of product
- D. Have samples available for display purposes

Answer: A

NEW QUESTION # 325

What action should be taken for materials received with missing documentation?

Response:

- A. Proceed with using the materials, assuming documentation is not critical

- B. Discard the materials immediately without further verification
- C. Use the materials for non-critical applications only
- **D. Quarantine the materials until the documentation is reviewed and approved**

Answer: D

NEW QUESTION # 326

When conducting a needs analysis, it is crucial to identify:

Response:

- **A. Training requirements**
- B. Advertising channels
- C. Product prices
- D. Potential investors

Answer: A

NEW QUESTION # 327

The effectiveness of a QMS is assessed through QMM to ensure:

Response:

- **A. Legal compliance**
- B. Competitive pricing
- C. Effective advertising
- D. Process and product quality

Answer: A

NEW QUESTION # 328

What is the significance of utilities change control in pharmaceutical manufacturing?

Response:

- A. It allows for utilities to be frequently changed based on employee votes
- B. It ensures that utility systems are always the latest model
- **C. It maintains the integrity and qualified state of the utility systems**
- D. It focuses on making utility systems more aesthetically pleasing

Answer: C

NEW QUESTION # 329

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