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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q339-Q344):

NEW QUESTION # 339

The primary goal of applying phase-appropriate GMPs is to:

Response:

- A. Focus solely on the aesthetics of the final product
- B. Ensure the product meets predefined quality criteria throughout its lifecycle
- C. Increase the complexity of documentation

- D. Reduce the need for regulatory oversight

Answer: B

NEW QUESTION # 340

Selecting appropriate raw materials involves _____ studies to evaluate their quality.

Response:

- A. compatibility
- B. financial
- C. marketing
- D. development

Answer: A

NEW QUESTION # 341

Formal contracts or quality agreements with suppliers should include provisions for:

(Choose two)

Response:

- A. Continuous improvement and corrective action procedures
- B. Flexible payment terms based on the buyer's financial condition
- C. Audit and inspection rights
- D. Regular social events to maintain a good working relationship

Answer: A,C

NEW QUESTION # 342

The primary reason for the identification and labeling requirements for samples, reagents, and standards is to:

Response:

- A. Fulfill an arbitrary regulatory requirement
- B. Increase the workload of laboratory staff
- C. Enhance the laboratory's reputation
- D. Ensure the accuracy and reliability of test results

Answer: D

NEW QUESTION # 343

What does REMS stand for in pharmaceutical surveillance?

Response:

- A. Regulatory Evaluation and Mitigation Scheme
- B. Risk Evaluation and Management Strategy
- C. Risk Evaluation and Mitigation Strategy
- D. Regulatory Evaluation and Management Scheme

Answer: C

NEW QUESTION # 344

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