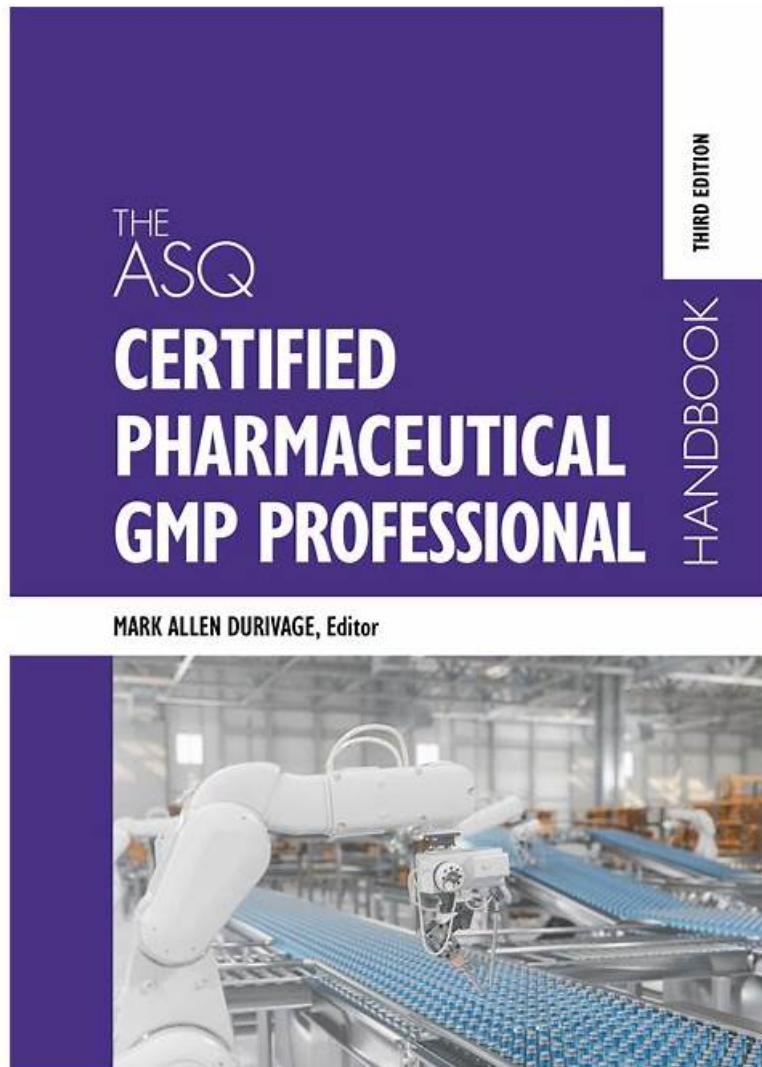


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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q280-

Q285):

NEW QUESTION # 280

Reviewing test methods against compendial requirements is important to ensure:

Response:

- A. Methods are chosen based on their complexity.
- B. Only non-compendial methods are used in the lab.
- C. The quickest method is chosen for all tests.
- D. **Compliance with both general and informational compendial chapters as needed.**

Answer: D

NEW QUESTION # 281

Configuration control in computerized systems ensures that:

Response:

- A. All users have equal administrative access
- B. **System settings and versions are properly documented and controlled**
- C. Configuration changes are decided by popular vote among users
- D. The system automatically updates without approval

Answer: B

NEW QUESTION # 282

In-process testing is important for:

Response:

- A. Marketing strategies
- B. Final product testing only
- C. Only at the beginning of the manufacturing process
- D. **Each step in the manufacturing process**

Answer: D

NEW QUESTION # 283

The layout of pharmaceutical manufacturing equipment must be designed to:

Response:

- A. **Minimize errors and facilitate cleaning and maintenance**
- B. Ensure that all equipment is visible from the facility's entrance
- C. Allow for easy photography and videography
- D. Reflect the hierarchical structure of the company

Answer: A

NEW QUESTION # 284

Validation studies for aseptic processing must include:

Response:

- A. **Temperature controls**
- B. **Process simulations (media fills)**
- C. Annual sales forecasts
- D. Taste testing

Answer: A,B

NEW QUESTION # 285

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