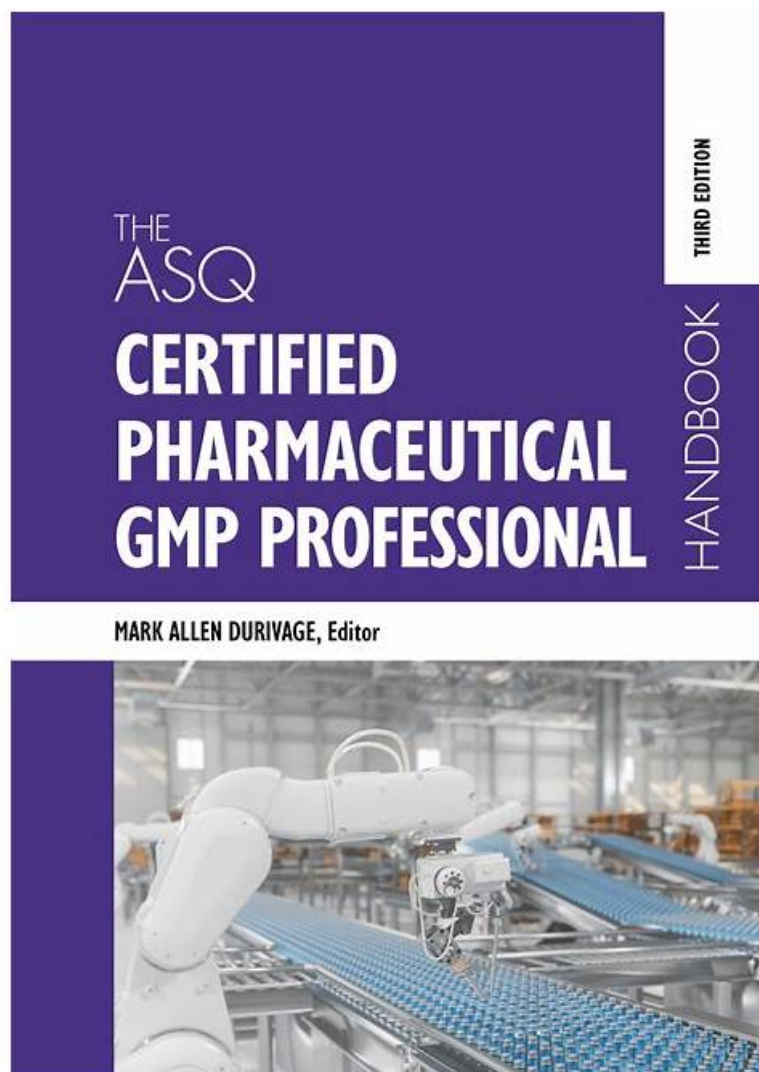


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## **ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q280-**

## 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

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Thanks for the helpful questions and answers, Each chapter discusses one particular CPGP element of the construction process, providing both conceptual information and useful tips on how to avoid common pitfalls in the construction process.

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