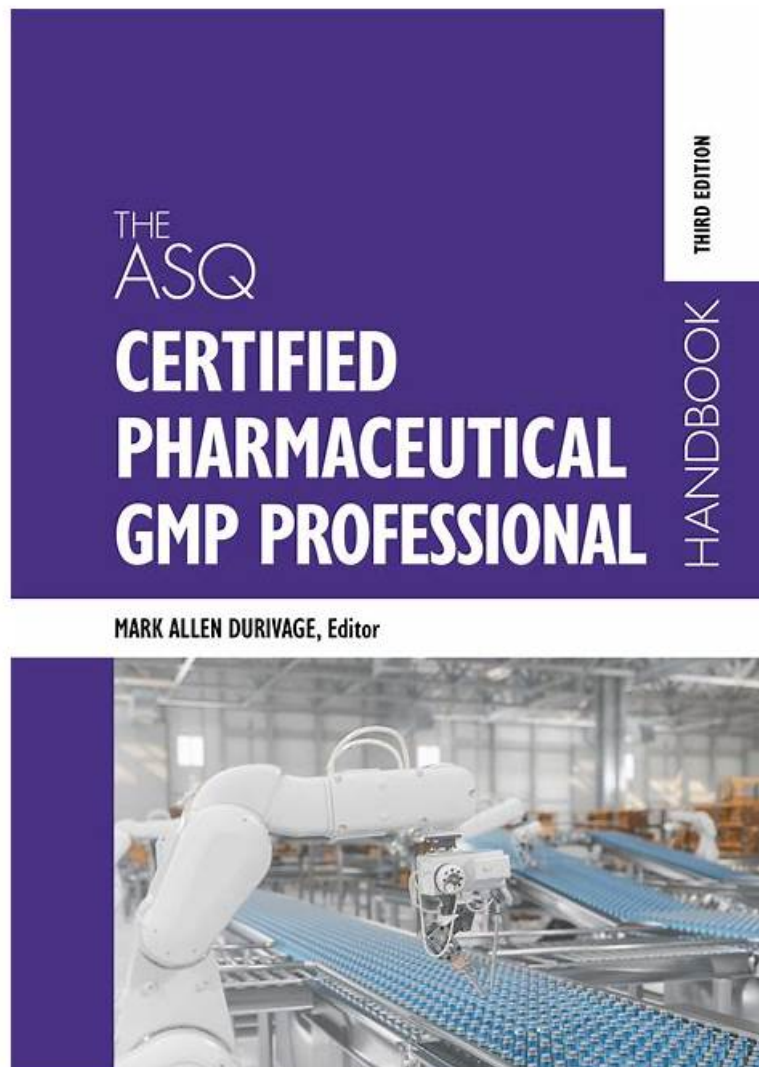


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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q33-Q38):

NEW QUESTION # 33

The process of _____ review involves evaluating test methods, qualifications, validation, and verification against required compendial chapters.

Response:

- A. administrative
- B. personnel performance
- C. financial
- **D. compendial or non-compendial requirements**

Answer: D

NEW QUESTION # 34

Biological agent traceability requirements are crucial to ensure:

Response:

- **A. Compliance with health and safety standards**
- B. The lowest production costs
- C. The fastest production time
- D. Easy advertisement

Answer: A

NEW QUESTION # 35

In the context of general cleaning and sanitization, what is the significance of using validated methods and materials? (Choose two)

Response:

- A. To align with the latest fashion trends in cleanliness
- **B. To guarantee the safety and compliance of the cleaning process**
- **C. To ensure the effectiveness and consistency of cleaning and sanitization**
- D. To increase the efficiency of social media marketing efforts

Answer: B,C

NEW QUESTION # 36

For how long should GMP-related records be retained, according to most regulatory guidelines?

Response:

- A. For the lifetime of the company for historical reference.
- B. Until the product expiration date plus one year.
- **C. For a minimum of 5 years after the product's commercial release.**
- D. Only during the period of product manufacturing.

Answer: C

NEW QUESTION # 37

What is a required element in both master and batch records?

Response:

- A. Future product development plans

- B. Personal employee information
- **C. Yield calculations**
- D. Marketing strategies

Answer: C

NEW QUESTION # 38

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