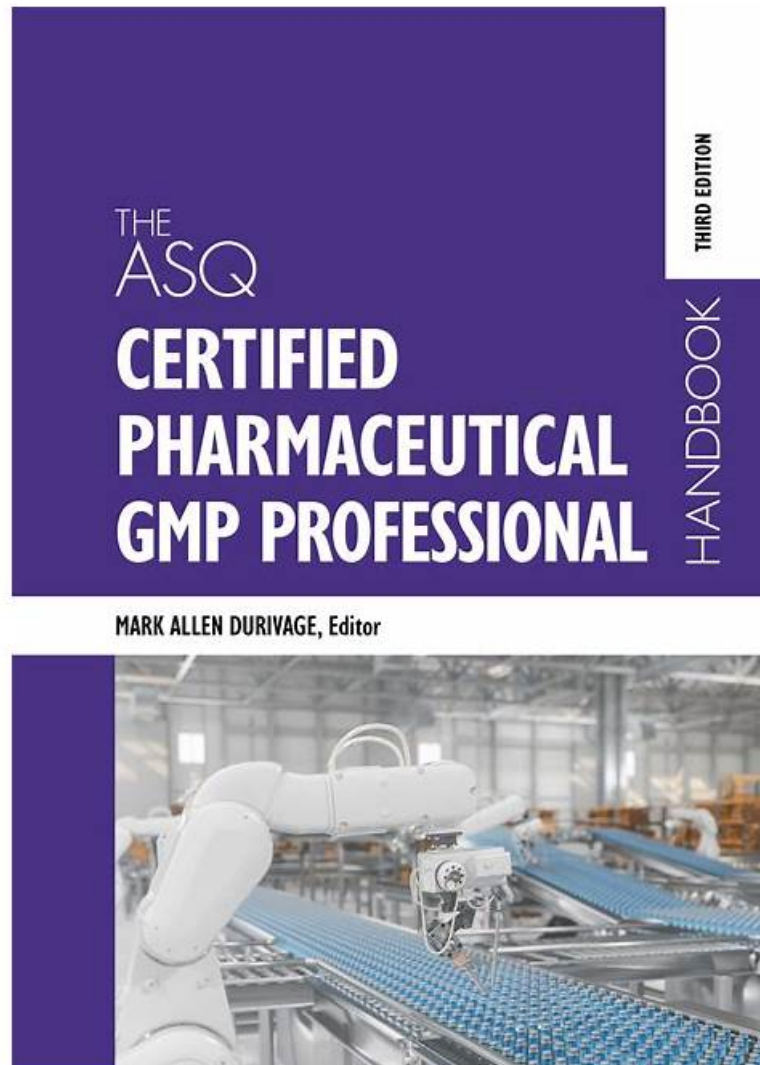


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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q297-

Q302):

NEW QUESTION # 297

Which document describes how a deviation was evaluated?

- A. Deviation investigation
- B. SOP
- C. CAPA report
- D. Change control form

Answer: A

NEW QUESTION # 298

When assessing the impact of proposed changes, what should be considered?

Response:

- A. Impact on all relevant aspects including products, processes, and systems
- B. The time taken for implementation
- C. Only direct costs
- D. The opinions of senior management only

Answer: A

NEW QUESTION # 299

Which of the following statements about sanitization and protection in manufacturing is true?

Response:

- A. Essential for maintaining product quality and safety
- B. Only required for liquid products
- C. Not necessary for solid dosage forms
- D. Only applies to nonsterile manufacturing environments

Answer: A

NEW QUESTION # 300

Which organization provides guidelines on veterinary pharmaceuticals and biologicals?

Response:

- A. USDA
- B. EMA
- C. FDA
- D. WHO

Answer: A

NEW QUESTION # 301

Master batch records are essential because they:

Response:

- A. Help in deciding the company's stock prices
- B. Provide a template for future marketing strategies
- C. Are used for employee training exclusively
- D. Serve as a standard for batch production to ensure consistency and compliance

Answer: D

NEW QUESTION # 302

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