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Certified Pharmaceutical GMP Professional

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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q502-Q507):

NEW QUESTION # 502

The implementation of corrective actions typically involves:

Response:

- A. Revising related procedures or processes
- B. Reducing the workforce
- C. Expanding office space
- D. Increasing product prices

Answer: A

NEW QUESTION # 503

Reserve samples must be retained to:

Response:

- A. Comply with regulatory requirements for post-market testing
- B. Enable future promotional activities
- C. Enhance the laboratory's collection
- D. Reduce manufacturing costs

Answer: A

NEW QUESTION # 504

Secure storage of reprocessed and reworked materials is important to prevent:

Response:

- A. Timely production schedules
- B. Employee satisfaction
- C. Cross-contamination
- D. Efficient use of space

Answer: C

NEW QUESTION # 505

When labeling reagents, solutions, and standards, which of the following information must be included?

Response:

- A. Opened-on date, expiry date, and quantity
- B. Color and viscosity
- C. Price and supplier name
- D. Physical appearance and smell

Answer: A

NEW QUESTION # 506

In inventory controls, what is the purpose of updating changes in material status?

Response:

- A. To reduce the space needed for storage
- B. To simplify the inventory process
- C. To maintain operational efficiency and traceability of materials
- D. To ensure accurate financial reporting

Answer: C

NEW QUESTION # 507

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