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>> CPGP Detail Explanation <<

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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q393-Q398):

NEW QUESTION # 393

The validation of automated or computerized systems should ensure that:

Response:

- A. Systems require minimal to no maintenance over their operational life
- B. Systems are capable of posting updates to social media automatically
- C. Systems are resistant to any form of cyber-attack, even hypothetical future threats
- D. Critical parameters are identified, controlled, and monitored

Answer: D

NEW QUESTION # 394

A pre-approval inspection in the pharmaceutical industry is conducted to:

Response:

- A. Assess the manufacturing site's compliance with regulatory standards before product approval.
- B. Evaluate the overall financial health of the manufacturing firm.
- C. Investigate a specific complaint or adverse event.
- D. Confirm the effectiveness of a new marketing strategy.

Answer: A

NEW QUESTION # 395

Change control for equipment in a pharmaceutical facility ensures that:

Response:

- A. Equipment can be replaced with newer models for aesthetic reasons
- B. All equipment is the latest technology, regardless of necessity
- C. Staff can request equipment based on personal preference
- D. The qualified state of equipment is maintained

Answer: D

NEW QUESTION # 396

The validation of sterilization processes is essential for:

(Choose two)

Response:

- A. Confirming the efficacy of sterilization methods
- B. Matching the sterilization process colors with the company logo
- C. Ensuring that product quality is not compromised
- D. Making the sterilization process more entertaining for employees

Answer: A,C

NEW QUESTION # 397

What are essential elements to review in Certificates of Analysis (COAs)?

(Choose two)

Response:

- A. Appropriate retention and accessibility
- B. Color of the product
- C. Completeness and internal review
- D. Marketing claims

Answer: A,C

NEW QUESTION # 398

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