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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q18-Q23):

NEW QUESTION # 18

The validation or verification of equipment cleaning processes is essential because:
Response:

- A. It determines which cleaning staff are most efficient
- **B. It ensures that cleaning procedures are effective and prevent cross-contamination**
- C. It provides an opportunity for staff to learn new cleaning techniques
- D. It is a requirement for receiving an interior design award

Answer: B

NEW QUESTION # 19

The FDA's regulations are crucial for pharmaceuticals in the USA. What does FDA stand for?

Response:

- A. Federal Drug Agency
- B. Food and Dietary Association
- C. Federal Department of Agriculture
- **D. Food and Drug Administration**

Answer: D

NEW QUESTION # 20

When assessing controls over supplier selection, what is considered a best practice?

Response:

- A. Selecting suppliers based solely on cost
- **B. Ensuring there are formal contracts or quality agreements in place**
- C. Choosing suppliers based on geographic proximity only
- D. Relying on supplier self-assessment reports without verification

Answer: B

NEW QUESTION # 21

ICH Q12's focus on product lifecycle management implies that risk management:

Response:

- A. Is a one-time activity
- B. Ends once the product is launched
- **C. Needs constant updating and application**
- D. Is only relevant for new products

Answer: C

NEW QUESTION # 22

What is the primary goal of conducting a pre-change analysis?

Response:

- A. To accelerate production
- B. To reduce costs
- **C. To minimize risk and ensure regulatory compliance**
- D. To simplify documentation

Answer: C

NEW QUESTION # 23

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