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**ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q183-Q188):**

**NEW QUESTION # 183**

Which of the following best describes the purpose of the GMP document system in a pharmaceutical manufacturing environment?

Response:

- A. To ensure efficient production schedules.
- **B. To document compliance with regulatory and corporate standards.**
- C. To monitor the stock levels of raw materials.
- D. To manage employee schedules and attendance.

**Answer: B**

**NEW QUESTION # 184**

Reviewing stability test data is crucial for:Response:

- A. Deciding the product's pricing strategy
- B. Determining the product's sales region
- **C. Establishing or challenging an expiry date based on trend analysis**
- D. Selecting the product's packaging color

**Answer: C**

**NEW QUESTION # 185**

The layout of pharmaceutical manufacturing equipment must be designed to:

Response:

- A. Reflect the hierarchical structure of the company
- B. Ensure that all equipment is visible from the facility's entrance
- **C. Minimize errors and facilitate cleaning and maintenance**
- D. Allow for easy photography and videography

**Answer: C**

**NEW QUESTION # 186**

Which document details post-approval changes in pharmaceutical manufacturing?

Response:

- **A. SUPAC guidelines**
- B. Marketing Authorization
- C. Clinical Trial Report
- D. Product Monograph

**Answer: A**

**NEW QUESTION # 187**

What are the goals of a metrology program in pharmaceutical manufacturing?

(Choose two)

Response:

- A. Showcasing the facility's technological advancements on social media
- **B. Ensuring measurement accuracy and reliability**
- **C. Maintaining compliance with regulatory requirements**
- D. Achieving celebrity endorsements for the manufacturing process

**Answer: B,C**

## NEW QUESTION # 188

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