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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q114-Q119):

NEW QUESTION # 114

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

Answer: C

NEW QUESTION # 115

When initiating an investigation for atypical results in a chemical test, which of the following is the first step?

Response:

- A. Reviewing the testing procedure and all associated records
- B. Adjusting the specification limits to accommodate the results
- C. Re-testing using the same method
- D. Discarding the atypical results as errors

Answer: A

NEW QUESTION # 116

Which of the following requirements is essential for the manufacture and storage environment in a pharmaceutical facility?

Response:

- A. Presence of a cafeteria within the production area
- B. Appropriate lighting, ventilation, and drainage
- C. Availability of a large parking lot
- D. High-tech security systems

Answer: B

NEW QUESTION # 117

A root cause analysis is essential for:

Response:

- A. Setting production targets
- B. Determining who is at fault
- C. Identifying the underlying cause of a detected problem
- D. Planning the company picnic

Answer: C

NEW QUESTION # 118

Different types of audits include:

(Choose two)

Response:

- A. Process audit
- B. Financial audit
- C. Product audit
- D. Systems audit

Answer: C,D

NEW QUESTION # 119

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