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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q271-Q276):

NEW QUESTION # 271

What is the primary goal of record retention policies in the pharmaceutical industry?

Response:

- A. To simplify employee training processes.
- **B. To ensure records are available for regulatory inspection and compliance verification.**
- C. To minimize the amount of paperwork.
- D. To reduce operational costs.

Answer: B

NEW QUESTION # 272

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

Response:

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

Answer: D

NEW QUESTION # 273

Which of the following is a potential source of contamination in a pharmaceutical manufacturing environment?

Response:

- A. Sealed product packaging
- B. Final product testing
- **C. Air handling systems**
- D. Properly cleaned equipment

Answer: C

NEW QUESTION # 274

During early development phases, GMP recommendations focus on:

Response:

- A. Final pricing strategies
- **B. Method qualification/validation**
- C. Final product packaging
- D. Mass marketing strategies

Answer: B

NEW QUESTION # 275

In laboratory investigations, the identification of unknowns typically involves:

Response:

- **A. Comparing test results to known standards or reference materials**
- B. Using a single test method for identification
- C. Consulting a psychic
- D. Guessing based on previous experience

Answer: A

NEW QUESTION # 276

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