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## ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q426-Q431):

### NEW QUESTION # 426

Quality unit (site) management responsibilities include:

Response:

- A. Ensuring proper batch release
- B. Setting sales targets
- C. Designing product packaging
- D. Organizing team-building events

**Answer: A**

**NEW QUESTION # 427**

The adoption of critical process parameters and specifications is crucial during:

Response:

- **A. All phases of development**
- B. Phase II clinical trials
- C. Marketing campaign development
- D. The conceptual design phase

**Answer: A**

**NEW QUESTION # 428**

What is a critical aspect of laboratory record-keeping requirements?

Response:

- A. Keeping records for no more than one year for space-saving
- B. Ensuring records are colorful and visually appealing
- C. Using only paper records to avoid electronic errors
- **D. Guaranteeing data integrity through proper data acquisition systems**

**Answer: D**

**NEW QUESTION # 429**

For combination products, GMP requirements include:

Response:

- **A. Studies specific to drug-device interactions**
- B. Excluding usability studies
- C. Ignoring device components
- D. Treating the product as a simple drug formulation

**Answer: A**

**NEW QUESTION # 430**

Release tests are primarily designed to:

Response:

- **A. Ensure a product meets the quality specifications at the time of release**
- B. Confirm the long-term stability of a product
- C. Assess the marketing potential of a product
- D. Determine the aesthetic appeal of a product

**Answer: A**

**NEW QUESTION # 431**

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