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## ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q286-Q291):

### NEW QUESTION # 286

Which of the following is NOT typically an in-process test?

Response:

- A. Sterility
- B. pH level
- C. Color of the product
- D. Weight uniformity

**Answer: C**

#### **NEW QUESTION # 287**

Record review processes are essential for ensuring:

Response:

- A. Quick disposal of outdated records
- B. Uniformity in record formatting
- C. Entertainment value of the data
- D. Data integrity and preventing data loss

**Answer: D**

#### **NEW QUESTION # 288**

A COA should be reviewed for completeness because:

Response:

- A. It ensures the material meets specified quality standards
- B. It influences the stock price of the company
- C. It provides historical data for marketing analysis
- D. It is a regulatory requirement

**Answer: A**

#### **NEW QUESTION # 289**

In assessing unit operations, deviations requiring investigation often relate to:

Response:

- A. Social media engagement rates
- B. Unexpected changes in process parameters
- C. Employee vacation schedules
- D. Branding decisions

**Answer: B**

#### **NEW QUESTION # 290**

The purpose of maintaining pedigree and sourcing details for APIs and other materials is to:

Response:

- A. Reduce the cost of goods sold
- B. Document the supply chain for quality and safety assurance
- C. Simplify the manufacturing process
- D. Enhance the appearance of the product

**Answer: B**

#### **NEW QUESTION # 291**

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