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## ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q315-Q320):

### NEW QUESTION # 315

The role of a Qualified Person includes ensuring:

Response:

- A. The effectiveness of marketing campaigns
- **B. Compliance with regulatory product quality standards**
- C. The efficiency of production processes
- D. The accuracy of financial reports

**Answer: B**

#### **NEW QUESTION # 316**

What is the significance of import alerts in the context of enforcement actions?

Response:

- A. They indicate exceptional product quality and compliance.
- B. They provide incentives for importing novel pharmaceuticals.
- **C. They notify about restrictions or bans on the importation of certain products.**
- D. They announce new partnership opportunities.

**Answer: C**

#### **NEW QUESTION # 317**

Which of the following statements about sanitization and protection in manufacturing is true?

Response:

- A. Only required for liquid products
- B. Not necessary for solid dosage forms
- C. Only applies to nonsterile manufacturing environments
- **D. Essential for maintaining product quality and safety**

**Answer: D**

#### **NEW QUESTION # 318**

GMP requirements for clinical trials material focus on:

(Select two)

Response:

- **A. Ensuring patient safety**
- **B. Guaranteeing the efficacy of the product**
- C. Enhancing the product's marketability
- D. Simplifying production processes

**Answer: A,B**

#### **NEW QUESTION # 319**

Which document describes how a deviation was evaluated?

- A. SOP
- **B. Deviation investigation**
- C. Change control form
- D. CAPA report

**Answer: B**

#### **NEW QUESTION # 320**

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