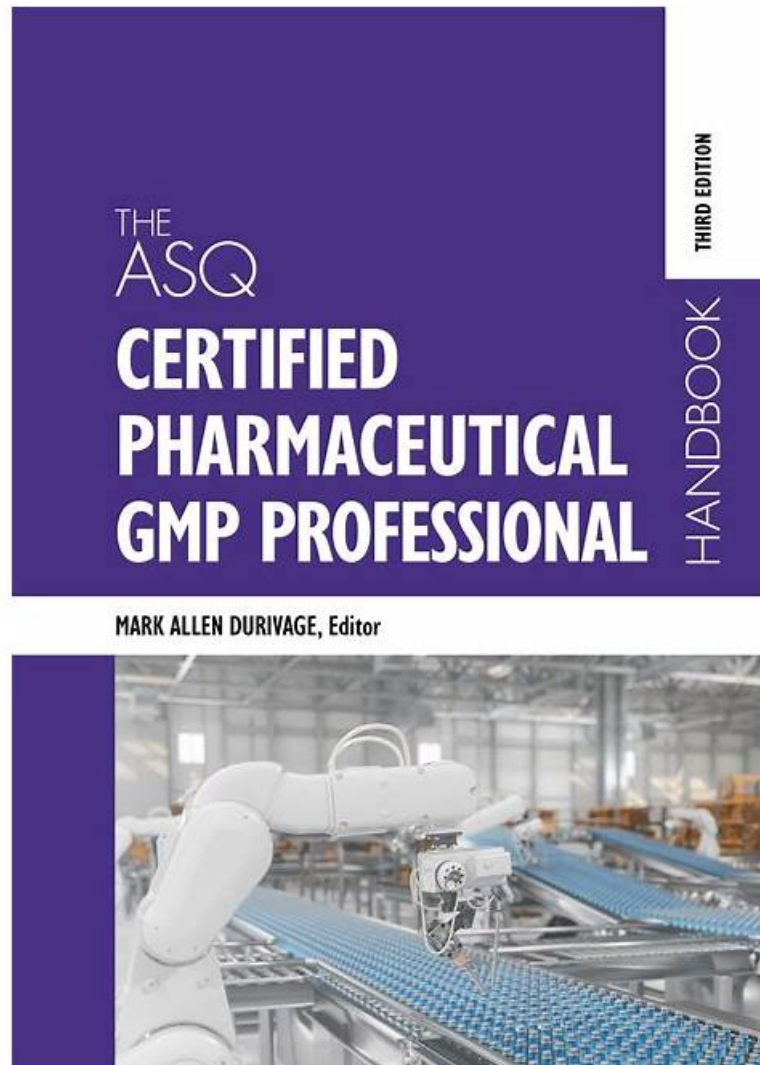


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## **ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q315-Q320):**

### **NEW QUESTION # 315**

What objectives are achieved through Performance Qualification (PQ) of equipment and utilities?

(Choose two)

Response:

- A. Proving that equipment can operate with zero noise levels
- **B. Demonstrating compliance with regulatory and user requirements over time**
- C. Ensuring equipment can be operated via remote control
- **D. Confirming consistent performance according to the intended use under real-world conditions**

**Answer: B,D**

### **NEW QUESTION # 316**

Formal contracts or quality agreements with suppliers should include provisions for:

(Choose two)

Response:

- **A. Audit and inspection rights**
- B. Regular social events to maintain a good working relationship
- **C. Continuous improvement and corrective action procedures**
- D. Flexible payment terms based on the buyer's financial condition

**Answer: A,C**

### **NEW QUESTION # 317**

CAPA effectiveness is confirmed by:

- A. QA approval alone
- B. End-of-shift reports
- **C. Trend analysis**
- D. Risk assessment only

**Answer: C**

### **NEW QUESTION # 318**

Why is it important to differentiate requirements for the sampling environment based on the type of material?

Response:

- A. To accommodate the personal preferences of the material suppliers
- B. It is solely for regulatory compliance, without practical benefits
- **C. Different materials require different ambient conditions to prevent alteration or contamination**
- D. It allows staff to choose their preferred working environment

**Answer: C**

### **NEW QUESTION # 319**

License withdrawal as an enforcement action results in:

Response:

- A. A temporary pause in production for reevaluation.
- **B. The loss of the right to produce or sell the product.**
- C. Increased scrutiny in unrelated business areas.
- D. The requirement to conduct more frequent advertising.

**Answer: B**

## NEW QUESTION # 320

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