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## ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q487-Q492):

### NEW QUESTION # 487

When handling atypical results, the investigation must include a review of \_\_\_\_\_ to identify possible deviations or errors.  
Response:

- A. the annual holiday schedule
- **B. test data and control samples**
- C. the cafeteria menu
- D. previous marketing campaigns

**Answer: B**

### NEW QUESTION # 488

ICH Q12 provides guidance on:  
Response:

- A. Validation of Analytical Procedures
- B. Quality Risk Management
- **C. Technical and Regulatory Considerations for Pharmaceutical Product Lifecycle Management**
- D. Pharmaceutical Quality System

**Answer: C**

**NEW QUESTION # 489**

In the process of equipment validation and qualification, which are key factors to assess for ensuring quality and compliance?  
(Choose two)

Response:

- A. The equipment's capacity for internet connectivity
- B. The color scheme of the equipment matching the company logo
- C. The robustness and reliability of the equipment under specified conditions
- D. The equipment's adherence to design and operational specifications

**Answer: C,D**

**NEW QUESTION # 490**

The use of cut-label procedures and splices is to:

Response:

- A. Reduce the effectiveness of inspections
- B. Ensure continuous operation and label integrity
- C. Increase label variety
- D. Decorate the packaging area

**Answer: B**

**NEW QUESTION # 491**

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 # 492**

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