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## ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q120-Q125):

### NEW QUESTION # 120

Successful technology transfer activities include all EXCEPT:

Response:

- A. Capability studies
- B. Ranging studies
- C. In-process control studies
- D. **Product renaming**

**Answer: D**

**NEW QUESTION # 121**

An adverse event is best described as:

Response:

- A. Any unexpected medical occurrence in a patient or clinical investigation subject administered a pharmaceutical product that does not necessarily have a causal relationship with this treatment.
- B. A minor complaint about product packaging.
- C. Any unsatisfactory customer service experience.
- D. A feedback suggesting improvements for product design.

**Answer: A**

**NEW QUESTION # 122**

Re-evaluation and revalidation frequency should be determined by:

Response:

- A. The phase of the moon
- B. The latest fashion trends
- C. Social media analytics
- D. Operational changes and process stability data

**Answer: D**

**NEW QUESTION # 123**

In business continuity planning, the supply chain impact analysis aims to:

Response:

- A. Determine the potential effects of supply chain disruptions on operations
- B. Ensure that the supply chain is environmentally friendly
- C. Prioritize suppliers based on their social media presence
- D. Focus on the most cost-effective suppliers without considering risk

**Answer: A**

**NEW QUESTION # 124**

Risk management principles in ICH Q9 should be applied:

Response:

- A. Only post-market
- B. Throughout the product lifecycle
- C. Exclusively during clinical trials
- D. Only at the beginning of product development

**Answer: B**

**NEW QUESTION # 125**

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