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ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q152-Q157):

NEW QUESTION # 152
Which of the following actions is typically NOT considered appropriate in response to an adverse event report?
Response:

- A. Reviewing and updating product information to include the adverse event.
- B. Reporting the event to regulatory authorities within the required timeframe.
- C. Immediate recall of all products potentially affected.
- D. Waiting for multiple reports of a similar event before taking any action.

Answer: D

NEW QUESTION # 153
Pedigree documentation for pharmaceutical products is used to:
Response:

- A. Enhance the brand image

- B. Provide a history of ownership and handling of a product
- C. Simplify the design process
- D. Promote sales

Answer: B

NEW QUESTION # 154

Regular _____ is crucial to ensuring the effectiveness of contamination control measures.
Response:

- A. Decrease in cleaning frequency
- B. Expansion of production lines
- C. Reduction of staff
- D. Monitoring

Answer: D

NEW QUESTION # 155

Critical Process Parameters (CPPs) are defined as:
Response:

- A. Parameters having an insignificant effect on CQAs
- B. Parameters that can be adjusted without impact on product quality
- C. Parameters only relevant during the initial stages of production
- D. Parameters that must be maintained to ensure CQAs are within specified limits

Answer: D

NEW QUESTION # 156

During product development, method qualification/validation is important because it:
Response:

- A. Is only a formal requirement without practical relevance
- B. Ensures the reliability and consistency of testing methods
- C. Determines the final product price
- D. Helps in designing the company logo

Answer: B

NEW QUESTION # 157

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