ASQ
CPGP
Certified Pharmaceutical GMP Professional

Latest Version: 6.3

Question: 1

The process of compendial methods review typically includes:
Response:

A. Ignoring the method's relevance to the product being tested.
B. Verifying the method's suitability and performance for the intended use.
C. Using the method directly from the compendium without any verification.
D. Choosing methods based on the shortest procedure.

Answer: B

Question: 2

The main reason for the traceability of intermediates in pharmaceutical manufacturing is to:
Response:

A. Allow for creative marketing
B. Ensure safety and efficacy through the supply chain
C. Make the packaging more attractive
D. Reduce the need for quality control

Answer: B

Question: 3

The design of water supply systems in pharmaceutical manufacturing should consider:
Response:

A. The color of the water
B. Taste preferences of the staff
C. Unit operations like dechlorination and reverse osmosis
D. The proximity to natural water bodies

Answer: C

Question: 4

Stability-indicating tests differ from release tests in that they:
Response:

A. Are used to determine the product's color and texture
B. Focus on the product's immediate quality upon release
C. Are designed to predict the product's stability over time
D. Assess the product's packaging integrity

Answer: C

Question: 5

The validation of sterilization processes is essential for:
(Choose two)
Response:

A. Confirming the efficacy of sterilization methods
B. Ensuring that product quality is not compromised
C. Matching the sterilization process colors with the company logo
D. Making the sterilization process more entertaining for employees

Answer: A,B

Question: 6

When should specifications be reviewed and updated?
Response:

A. Only when the product formula changes
B. Annually, as a standard procedure
C. When methods are revised or compendia are changed
D. When a new management team is appointed

Answer: C

Question: 7

The layout of pharmaceutical manufacturing equipment must be designed to:

Response:

A. Allow for easy photography and videography
B. Minimize errors and facilitate cleaning and maintenance
C. Reflect the hierarchical structure of the company
D. Ensure that all equipment is visible from the facility's entrance

Answer: B

Question: 8

The role of the supervisor in ensuring training effectiveness includes:
Response:

A. Enhancing their own skills
B. Ensuring staff understand and apply their training
C. Investing in new technologies
D. Networking with industry leaders

Answer: B

Question: 9

Remote audits are particularly useful when:
Response:

A. In-person travel is restricted
B. Auditors prefer not to travel
C. The organization is too large for in-person audits
D. The weather conditions are unfavorable

Answer: A

Question: 10

Calibration and engineering change control procedures are implemented to:
(Choose two)
Response:

A. Guarantee that modifications do not affect the precision and accuracy of measurements
B. Ensure equipment settings are optimized for the fastest production time
C. Maintain the integrity of manufacturing processes

D. Facilitate the live streaming of equipment operations

Answer: A,C

Question: 11

Audit follow-up actions are necessary to:
Response:

A. Check the effectiveness of implemented corrective actions
B. Reward the audited team
C. Finalize audit schedules for the next cycle
D. Conduct team-building activities

Answer: A

Question: 12

What is a critical consideration when designing a sampling plan for bulk chemicals?
Response:

A. The color of the chemicals
B. The representative nature of the sample and appropriate sample size
C. The preferences of the production department
D. The current stock price of the chemicals

Answer: B

Question: 13

Mutual recognition agreements impact the import and export of pharmaceuticals by:
Response:

A. Ensuring that product prices are consistent across borders.
B. Requiring additional testing for all imported or exported products.
C. Facilitating smoother regulatory approval processes between countries.
D. Increasing the number of required inspections for each product.

Answer: C

Question: 14

In the context of APR and PQR, what is the significance of reviewing product complaints?
Response:

A. To adjust the product price
B. To assess the product's packaging design
C. To evaluate and improve product quality and safety
D. To measure consumer brand loyalty

Answer: C

Question: 15

Quality agreements are essential because they:
Response:

A. Allow for the informal resolution of disputes
B. Clearly define product and service quality requirements and responsibilities
C. Serve as a formal record of the partnership for marketing purposes
D. Provide a platform for socializing with suppliers

Answer: B

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