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ASQ CPGP Exam Syllabus Topics:

Section	Weight	Objectives
Laboratory Systems	14%	- Testing methods, data integrity, and result reporting - Laboratory controls, calibration, and validation
Materials and Supply Chain Management	12%	- Inventory control, traceability, and distribution practices - Supplier qualification, material receipt, and storage
Infrastructure and Facilities	12%	- Equipment qualification, calibration, and cleaning validation - Facility design, maintenance, and utilities management
Regulatory Agency Governance	12%	- Inspection, enforcement, and compliance processes - Global regulatory framework and requirements
Quality Systems	18%	- Quality management system design and implementation - Documentation, change control, and records management - Audits, corrective and preventive actions
Filling, Packaging, and Labeling	13%	- Filling operations, controls, and verification - Packaging and labeling design and compliance
Product Development and Technology Transfer	9%	- Product lifecycle management and regulatory submissions - Technology transfer, scale-up, and validation
Manufacturing Systems	16%	- Contamination control, cleaning and hygiene - Process design, validation, and control - Sterile and non-sterile production requirements

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The CPGP certification verifies that you are a skilled professional. PrepAwayPDF product is designed by keeping all the rules and regulations in focus that ASQ publishes. Our main goal is that you can memorize the actual ASQ CPGP Exam Question to complete the Certified Pharmaceutical GMP Professional (CPGP) test in time with extraordinary grades.

ASQ Certified Pharmaceutical GMP Professional Sample Questions (Q95-Q100):

NEW QUESTION # 95

Pharmacovigilance primarily aims to:

Response:

- A. Monitor the long-term stability of pharmaceutical products
- B. Track the effectiveness of marketing strategies
- C. Oversee the distribution and retail of pharmaceutical products
- **D. Detect and assess adverse drug reactions**

Answer: D

NEW QUESTION # 96

The Site Master File (SMF) should contain information about:

Response:

- **A. The site's quality management system**
 - B. The company's annual financial reports
 - C. Upcoming product launches
 - D. The manufacturing site's personnel details
- The site's quality management system

Answer: A

NEW QUESTION # 97

The main elements of a quality agreement typically include:

Response:

- A. Guarantees of stock price performance
- B. Plans for joint marketing initiatives
- **C. Scope and purpose, roles and responsibilities, and dispute resolution procedures**
- D. Entertainment allowances for supplier visits

Answer: C

NEW QUESTION # 98

Technology transfer is crucial when there is a _____ change.

Response:

- A. marketing strategy
- B. sales team
- **C. manufacturing site**
- D. packaging color

Answer: C

NEW QUESTION # 99

The validation of automated or computerized systems should ensure that:

Response:

- **A. Critical parameters are identified, controlled, and monitored**
- B. Systems require minimal to no maintenance over their operational life
- C. Systems are resistant to any form of cyber-attack, even hypothetical future threats
- D. Systems are capable of posting updates to social media automatically

Answer: A

NEW QUESTION # 100

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