

# REGULATORY AFFAIRS

1. Which organization is responsible for setting food safety standards at the national level?
  - a) FDA (Food and Drug Administration)
  - b) USDA (United States Department of Agriculture)
  - c) FSSAI (Food safety standard authority of India)
  - d) FSIS (Food Safety and Inspection Service)
2. Which food safety regulation requires food manufacturers to provide accurate labeling information about allergens?
  - a) FDA Food Safety Modernization Act (FSMA)
  - b) USDA Organic Certification
  - c) European Union Food Information to Consumers Regulation
  - d) Canada's Food and Drug Regulations
3. What is the maximum allowable level of microbial contamination in fruit juices?
  - a) 10 colony-forming units (CFU) per milliliter
  - b) 100 CFU per milliliter
  - c) 1,000 CFU per milliliter
  - d) 10,000 CFU per milliliter
4. Which of the following is NOT a common allergen that must be declared on beverage labels?
  - a) Milk
  - b) Soy
  - c) Wheat
  - d) Caffeine
5. What is the minimum required pH level for acidified beverages to prevent microbial growth?
  - a) pH 2.5
  - b) pH 4.0
  - c) pH 5.5
  - d) pH 7.0
6. Which of the following statements about food allergen labeling is true?
  - a) All allergens must be listed in bold font on the label.
  - b) Allergen labeling is optional for small juice and b.
  - c) Only the top eight allergens need to be declared.
  - d) Allergen information must be listed in a separate document.
7. Which of these is a basic labeling requirement according to FOOD SAFETY AND STANDARDS (PACKAGING AND LABELLING) REGULATIONS, 2011
  - a) Date of manufacture
  - b) Veg/ Non- Vegetarian Food
  - c) Best before
  - d) All of these
8. ISO 9001: 2008 Quality management systems deals with
  - a) Fundamentals and vocabulary
  - b) Guidelines for performance improvements

- c) Customer Satisfaction
- d) Requirements for quality management
9. According to the FSS act, the full form of COA is:
  - a) Certificate of approval
  - b) Certificate of authority
  - c) Certificate of analysis
  - d) Certificate of Assessment
10. In food factory the ISO 22000 is for?
  - a) Increasing profits
  - b) Managing food safety
  - c) Employee welfare
  - d) None of the above
11. How can ISO 9001 assist food industry organizations in meeting regulatory requirements?
  - a) ISO 9001 provides a comprehensive list of all applicable regulations.
  - b) ISO 9001 offers a framework for organizations to establish and maintain regulatory compliance.
  - c) ISO 9001 exempts organizations from regulatory inspections.
  - d) ISO 9001 eliminates the need for organizations to comply with regulations.
12. Which regulatory aspect ensures that food packaging materials are safe for use?
  - a) Food Safety and Hygiene regulations
  - b) Quality Standards regulations
  - c) Import and Export regulations
  - d) Environmental and Sustainability regulations
13. Which regulatory category focuses on preventing deceptive or misleading advertising on food packaging?
  - a) Food Safety and Hygiene regulations
  - b) Quality Standards regulations
  - c) Labeling and Packaging regulations
  - d) Advertising and Marketing regulations
14. \_\_\_\_\_ of the Food Safety and Standard Regulations, 2011, specifies the standards for various infant products.
  - a) 2.1.9
  - b) 2.2.9
  - c) 2.4
  - d) 2.3
15. What does the term "net weight" refer to in food packaging regulations?
  - a) The total weight of the food product, including packaging
  - b) The weight of the food product without packaging
  - c) The weight of the food product after cooking or preparation
  - d) The weight of the packaging material used for the food product

16. The basic purpose of establishing food standards is \_\_\_\_\_.
  - a) To promote unfair trade in food.
  - b) To ensure food safety and quality.
  - c) Protection of public health and risk
  - d) To ensure fair trade in food.

# Regulatory Affairs Questions Answers

**United States. Congress. Senate.  
Committee on Homeland Security and  
Governmental Affairs. Permanent  
Subcommittee on Investigations**

## **Regulatory Affairs Questions Answers:**

*Certified In Healthcare Compliance Exam Success Guide 2025/2026* Lara Fitzroy, 2025-08-16 The Certified in Healthcare Compliance Exam Success Guide 2025 2026 is your trusted resource for mastering the CHC exam and advancing your career in healthcare compliance With 580 practice questions this guide delivers the tools and knowledge you need to confidently prepare for every exam domain Inside you ll find Comprehensive coverage of compliance standards regulations and best practices 580 exam style practice questions with detailed explanations to reinforce understanding Proven test taking strategies to reduce anxiety and maximize performance Up to date content aligned with the latest CHC exam format Whether you are new to compliance or a seasoned professional seeking certification this guide provides a structured path to success equipping you with the skills and confidence to excel on test day **FDA Regulatory Affairs** David Mantus, Douglas J.

Pisano, 2014-02-28 FDA Regulatory Affairs is a roadmap to prescription drug biologics and medical device development in the United States Written in plain English the concise and jargon free text demystifies the inner workings of the US Food and Drug Administration FDA and facilitates an understanding of how the agency operates with respect to compliance and product approval including clinical trial exemptions fast track status advisory committee procedures and more The Third Edition of this highly successful publication Examines the harmonization of the US Federal Food Drug and Cosmetic Act with international regulations on human drug biologics and device development research manufacturing and marketing Includes contributions from experts at organizations such as the FDA National Institutes of Health NIH and PAREXEL Focuses on the new drug application NDA process cGMPs GCPs quality system compliance and corresponding documentation requirements Provides updates to the FDA Safety and Innovation Act FDASIA incorporating pediatric guidelines and follow on biologics regulations from the 2012 Prescription Drug User Fee Act PDUFA V Explains current FDA inspection processes enforcement options and how to handle FDA meetings and required submissions Co edited by an industry leader Mantus and a respected academic Pisano FDA Regulatory Affairs Third Edition delivers a compilation of the selected US laws and regulations as well as a straightforward commentary on the FDA product approval process that s broadly useful to both business and academia

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Nomination of Richard G. Darman United States. Congress. Senate. Committee on Governmental Affairs, 1989

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*Friedman, Glauthier, and Groat Nominations* United States. Congress. Senate. Committee on Energy and Natural Resources, 1999

*Moodle 4 E-Learning Course Development* Susan Smith Nash, 2022-06-30

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