Foods Program Guidance Documents Under Development

(Expected to publish as drafts or finals by the end of December 2022.)

Introduction

The following list of guidance topics includes possible new topics for guidance documents or revisions to existing guidance documents that the FDA Foods Program is considering. We currently intend to develop guidance on each topic; however, the FDA Foods Program is neither bound by this list of topics, nor required to issue every guidance document on this list. Several factors may impact FDA's ability to issue the listed guidances, including, for example, new Administration priorities, emerging public health issues, or other extenuating circumstances. We are not precluded from issuing guidance documents on topics not on this list.

You may submit comments on the guidance topics at <u>www.regulations.gov</u> at Docket FDA-2021-N-0553.

Guidance for Industry

Title of Guidance	Category
Evaluating the Public Health Importance of Food Allergens Other Than the Major Food Allergens Defined in the Federal Food, Drug, and Cosmetic Act; Draft Guidance for FDA Staff and Stakeholders	Allergens
Questions and Answers Regarding Food Allergens, Including the Food Allergen Labeling and Consumer Protection Act of 2004 (Edition 5); Draft Guidance for Industry	Allergens
Testing Methods for Asbestos in Cosmetic Products Containing Talc; Draft Guidance for Industry	Cosmetics
Policy Regarding Certain New Dietary Ingredients and Dietary Supplements Subject to the Requirement for Pre-market Notifications; Draft Guidance for Industry	Dietary Supplements
New Dietary Ingredient (NDI) Notifications and Related Issues: NDI Notification Procedures and Timeframes; Guidance for Industry	Dietary Supplements
Best Practices for Convening a GRAS Panel; Guidance for Industry	Food Additives
Premarket Consultation on Cultured Animal Cell Foods: Draft Guidance for Industry	Food Additives
Foods Derived from Plants Produced Using Genome Editing; Draft Guidance for Industry	Food Safety

Food Safety
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Hazard Analysis and Risk-Based Preventive Controls for Human Food; Chapter 17: Classifying Food as Ready-To-Eat or Not Ready- to-Eat; Draft Guidance for Industry	FSMA
Hazard Analysis and Risk-Based Preventive Controls for Human Food; Chapter 18: Acidified Foods; Draft Guidance for Industry	FSMA
Questions and Answers Regarding the Accredited Third-Party Certification Program: Draft Guidance for Industry	FSMA
Refusal of Inspection by a Foreign Food Establishment or Foreign Government; Guidance for Industry	FSMA
Foreign Supplier Verification Programs for Importers of Food for Humans and Animals; Guidance for Industry	FSMA
Compliance with and Recommendations for Implementation of the Standards for the Growing, Harvesting, Packing, and Holding of Produce for Human Consumption for Sprout Operations: Guidance for Industry	FSMA
Labeling of Plant-based Milk Alternatives: Draft Guidance for Industry	Labeling
Labeling of Plant-Based Alternatives to Animal-Derived Foods; Draft Guidance for Industry	Labeling
Questions and Answers About Dietary Guidance Statements in Food Labeling; Draft Guidance for Industry	Labeling
Protein Efficiency Ratio (PER) Rat Bioassay Studies to Demonstrate that a New Infant Formula Supports the Quality Factor of Sufficient Biological Quality of Protein; Draft Guidance for Industry	Nutrition

ⁱFDA's Center for Veterinary Medicine also has published a list of <u>Guidances Under Development for 2022.</u>