

currently funded to carry out The Link Center Project for the period of September 1, 2022, through August 31, 2027. Much work has already been completed and further tasks are currently being accomplished. It would be unnecessarily time consuming and disruptive to the Link Center project and the beneficiaries being served for ACL to establish a new grantee at this time when critical services are presently being provided in an efficient manner. SAMHSA also has determined that the award of another contract or grant to provide these services would duplicate the activities carried out under this cooperative agreement. SAMHSA has further determined that a grant supplement to support the 988 State Policy Academy through this cooperative agreement is likely to be less expensive than a separate arrangement. This agreement promotes government efficiency and reduces the possibility of costly duplication of effort.

Dated: September 21, 2023.

Alison Barkoff,

Senior Official Performing the Duties of the Administrator and the Assistant Secretary for Aging.

[FR Doc. 2023-21046 Filed 9-26-23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2016-D-2343]

Hazard Analysis and Risk-Based Preventive Controls for Human Food; Draft Guidance for Industry; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of availability.

SUMMARY: The Food and Drug Administration (FDA, the Agency, or we) is announcing the availability of two additional draft chapters of a multichapter draft guidance for industry entitled “Hazard Analysis and Risk-Based Preventive Controls for Human Food.” This multichapter draft guidance, when finalized, will explain FDA’s current thinking on how to comply with the requirements for hazard analysis and risk-based preventive controls under FDA’s regulation entitled “Current Good Manufacturing Practice, Hazard Analysis, and Risk-Based Preventive Controls for Human Food.” The newly available draft chapters are entitled “Chapter 11—Food Allergen Program”

and “Chapter 16—Acidified Foods.” This draft guidance is not final nor is it in effect at this time.

DATES: Submit either electronic or written comments on the draft guidance by March 25, 2024 to ensure that the Agency considers your comment on this draft guidance before it begins work on the final version of the guidance.

ADDRESSES: You may submit comments on any guidance at any time as follows:

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else’s Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see “Written/Paper Submissions” and “Instructions”).

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in “Instructions.”

Instructions: All submissions received must include the Docket No. FDA-2016-D-2343 for “Hazard Analysis and Risk-Based Preventive Controls for Human Food.” Received comments will be placed in the docket and, except for those submitted as “Confidential Submissions,” publicly viewable at <https://www.regulations.gov> or at the

Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240-402-7500.

- *Confidential Submissions—*To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states “THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.” The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on <https://www.regulations.gov>. Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as “confidential.” Any information marked as “confidential” will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA’s posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: <https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf>.

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to <https://www.regulations.gov> and insert the docket number, found in brackets in the heading of this document, into the “Search” box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240-402-7500.

You may submit comments on any guidance at any time (see 21 CFR 10.115(g)(5)).

Submit written requests for single copies of the draft guidance to the Office of Food Safety (HFS-300), Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740. Send two self-addressed adhesive labels to assist that office in processing your requests. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance document.

FOR FURTHER INFORMATION CONTACT: Linda Kahl, Center for Food Safety and Applied Nutrition (HFS-300), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240-402-2784.

SUPPLEMENTARY INFORMATION:

I. Background

We are announcing the availability of two draft chapters entitled “Chapter

11—Food Allergen Program” and “Chapter 16—Acidified Foods” of a multichapter draft guidance for industry entitled “Hazard Analysis and Risk-

Based Preventive Controls for Human Food.” We previously announced the availability of several chapters of that draft guidance as shown in table 1.

TABLE 1—AVAILABLE DRAFT CHAPTERS IN HAZARD ANALYSIS AND RISK-BASED PREVENTIVE CONTROLS FOR HUMAN FOOD

Chapter No.	Chapter title	Publication
N/A	Introduction	81 FR 57816, August 24, 2016.
1	The Food Safety Plan	81 FR 57816, August 24, 2016.
2	Conducting a Hazard Analysis	81 FR 57816, August 24, 2016.
3	Potential Hazards Associated with the Manufacturing, Processing, Packing, and Holding of Human Food.	81 FR 57816, August 24, 2016.
4	Preventive Controls	81 FR 57816, August 24, 2016.
5	Application of Preventive Controls and Preventive Control Management Components	81 FR 57816, August 24, 2016.
6	Use of Heat Treatments as a Process Control	82 FR 41364, August 31, 2017.
14	Recall plan	84 FR 53347, October 7, 2019.
15	Supply-Chain Program for Human Food Products	83 FR 3449, January 25, 2018.
Appendix 1	Potential Hazards for Foods and Processes	81 FR 57816, August 24, 2016.
Appendix 2	Food Safety Plan Forms	81 FR 57816, August 24, 2016.
Appendix 3	Bacterial Pathogen Growth and Inactivation	81 FR 57816, August 24, 2016.

We also are announcing changes to the expected table of contents for the complete multichapter guidance.

We are issuing these chapters of the draft guidance consistent with our good guidance practices regulation (21 CFR 10.115). The draft guidance, when finalized, will represent the current thinking of FDA on how to comply with the requirements for hazard analysis and risk-based preventive controls under part 117 (21 CFR part 117), principally in subparts C and G. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternative approach if it satisfies the requirements of the applicable statutes and regulations.

The FDA Food Safety Modernization Act (FSMA) (Pub. L. 111–353) enables FDA to better protect public health by helping to ensure the safety and security of the food supply. FSMA enables FDA to focus more on preventing food safety problems rather than relying primarily on reacting to problems after they occur. FSMA recognizes the important role industry plays in ensuring the safety of the food supply, including the adoption of modern systems of preventive controls in food production.

Section 103 of FSMA amended the Federal Food, Drug, and Cosmetic Act (FD&C Act) by adding section 418 (21 U.S.C. 350g) with requirements for hazard analysis and risk-based preventive controls for establishments that are required to register as food facilities under our regulations in 21 CFR part 1, subpart H, in accordance with section 415 of the FD&C Act (21 U.S.C. 350d). We have established regulations to implement these requirements within part 117.

We intend to announce the availability for public comment of additional chapters of the draft guidance as we complete them. The titles of the additional chapters that we expect to make available for public comment are included in the table of contents for the complete multichapter guidance.

III. Paperwork Reduction Act of 1995

While this guidance contains no collection of information, it does refer to previously approved FDA collections of information. The previously approved collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (PRA) (44 U.S.C. 3501–3521). The collections of information in part 117 have been approved under OMB control number 0910–0751.

IV. Electronic Access

Persons with access to the internet may obtain the draft guidance at <https://www.fda.gov/FoodGuidances>, <https://www.fda.gov/regulatory-information/search-fda-guidance-documents>, or <https://www.regulations.gov>. Use the FDA website listed in the previous sentence to find the most current version of the guidance.

Dated: September 20, 2023.

Lauren K. Roth,

Associate Commissioner for Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2021–D–1158]

Cybersecurity in Medical Devices: Quality System Considerations and Content of Premarket Submissions; Guidance for Industry and Food and Drug Administration Staff; Availability

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of availability.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing the availability of a final guidance entitled “Cybersecurity in Medical Devices: Quality System Considerations and Content of Premarket Submissions.” As more medical devices are becoming interconnected, cybersecurity threats have become more numerous, more frequent, more severe, and more clinically impactful. As a result, ensuring medical device safety and effectiveness includes adequate medical device cybersecurity, as well as its security as part of the larger system. This final guidance supersedes the final guidance “Content of Premarket Submissions for Management of Cybersecurity in Medical Devices,” issued October 2, 2014.

DATES: The announcement of the guidance is published in the **Federal Register** on September 27, 2023.

ADDRESSES: You may submit either electronic or written comments on Agency guidances at any time as follows: