List of Subjects in 18 CFR Part 11

Dams, Electric power, Indians-lands, Public lands, Reporting and recordkeeping requirements.

By direction of the Commission.

Issued: August 17, 2017.

Nathaniel J. Davis, Sr.,

Deputy Secretary.

In consideration of the foregoing, the Federal Energy Regulatory Commission proposes to amend Part 11, Chapter I, Title 18, *Code of Federal Regulations,* as follows:

PART 11—ANNUAL CHARGES UNDER PART I OF THE FEDERAL POWER ACT

■ 1. The authority citation for part 11 continues to read as follows:

Authority: 16 U.S.C. 792–828c; 42 U.S.C. 7101–7352.

■ 2. In § 11.2, add paragraph (c)(1)(iv) to read as follows:

* * * * *

(c) * * *

(1) * * *

(iv) For all geographic areas in Alaska except for the Aleutian Islands Area, the Commission will calculate a statewide average per-acre land value based on the average per-acre land and building values published in the NASS Census for the Kenai Peninsula and the Fairbanks Areas. This statewide average per-acre value will be reduced by the sum of the state-specific modifier and seven percent. The resulting adjusted statewide average per-acre value will be applied to all projects located in Alaska, except for those projects located in the Aleutian Island Area.

* * * * *

[FR Doc. 2017–17846 Filed 8–30–17; 8:45 am]

BILLING CODE 6717–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 117

[Docket No. FDA–2016–D–2343]

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

AGENCY: Food and Drug Administration, HHS.

ACTION: Notification of availability.

SUMMARY: The Food and Drug Administration (FDA, we, or Agency) is announcing the availability of another draft chapter of a multichapter guidance for industry entitled ''Hazard Analysis and Risk-Based Preventive Controls for Human Food: Guidance for Industry.'' This multichapter draft guidance is intended to explain our current thinking on how to comply with the requirements for hazard analysis and risk-based preventive controls under our rule entitled ''Current Good Manufacturing Practice, Hazard Analysis, and Risk-Based Preventive Controls for Human Food.'' The newly available draft chapter is entitled ''Chapter Six—Use of Heat Treatments as a Process Control.''

DATES: Although you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that we consider your comment on this draft guidance before we issue the final version of the guidance, submit either electronic or written comments by February 27, 2018.

You may submit comments 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: Guidance for Industry.'' 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 office between 9 a.m. and 4 p.m., Monday through Friday.

- *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.gpo.gov/fdsys/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.

Submit written requests for single copies of the draft guidance to Office of Food Safety, Center for Food Safety and Applied Nutrition, Food and Drug Administration (HFS–300), 5001 Campus Dr., College Park, MD 20740. Send two self-addressed adhesive labels

to assist that office in processing your request. See the **SUPPLEMENTARY INFORMATION** section for electronic access to the draft guidance.

FOR FURTHER INFORMATION CONTACT: Jenny Scott, Center for Food Safety and Applied Nutrition (HFS–300), Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–2166.

SUPPLEMENTARY INFORMATION:

I. Background

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. It 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), in section 418 of the FD&C Act (21 U.S.C. 350g), by adding 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 (21 CFR part 117).

In the **Federal Register** of August 24, 2016 (81 FR 57816), we announced the availability of several chapters of a multichapter draft guidance for industry entitled ''Hazard Analysis and Risk-Based Preventive Controls for Human Food.'' We now are announcing the availability of an additional draft chapter of this multichapter guidance for industry. We are issuing 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 this topic. It does not establish any rights for any person and is not binding on FDA or the public. You can use an alternate approach if it satisfies the requirements of the applicable statutes and regulations. This guidance is not subject to Executive Order 12866.

The multichapter draft guidance for industry is intended to explain our current thinking on how to comply with the requirements for hazard analysis and risk-based preventive controls under part 117, principally in subparts C and G. The chapter that we are announcing in this document is entitled ''Chapter Six—Use of Heat Treatments as a Process Control.''

We intend to announce the availability for public comment of additional chapters of the draft guidance as we complete them.

II. Paperwork Reduction Act of 1995

This draft guidance refers to previously approved collections of information found in FDA regulations. These collections of information are subject to review by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in part 117 have been approved under OMB control number 0910–0751.

III. Electronic Access

Persons with access to the Internet may obtain the draft guidance at either *https://www.fda.gov/FoodGuidances* or *https://www.regulations.gov.* Use the FDA Web site listed in the previous sentence to find the most current version of the guidance.

Dated: August 22, 2017.

Anna K. Abram,

Deputy Commissioner for Policy, Planning, Legislation, and Analysis.

[FR Doc. 2017–18464 Filed 8–30–17; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF LABOR

Employee Benefits Security Administration

29 CFR Part 2550

[Application Number D–11712; D–11713; D–11850]

ZRIN 1210–ZA27

Extension of Transition Period and Delay of Applicability Dates; Best Interest Contract Exemption (PTE 2016–01); Class Exemption for Principal Transactions in Certain Assets Between Investment Advice Fiduciaries and Employee Benefit Plans and IRAs (PTE 2016–02); Prohibited Transaction Exemption 84–24 for Certain Transactions Involving Insurance Agents and Brokers, Pension Consultants, Insurance Companies, and Investment Company Principal Underwriters (PTE 84–24)

AGENCY: Employee Benefits Security Administration, Labor.

ACTION: Notice of proposed amendments to PTE 2016–01, PTE 2016–02, and PTE 84–24.

SUMMARY: This document proposes to extend the special transition period under sections II and IX of the Best Interest Contract Exemption and section VII of the Class Exemption for Principal Transactions in Certain Assets Between Investment Advice Fiduciaries and Employee Benefit Plans and IRAs. This document also proposes to delay the applicability of certain amendments to Prohibited Transaction Exemption 84–24 for the same period. The primary purpose of the proposed amendments is to give the Department of Labor the time necessary to consider possible changes and alternatives to these exemptions. The Department is particularly concerned that, without a delay in the applicability dates, regulated parties may incur undue expense to comply with conditions or requirements that it ultimately determines to revise or repeal. The present transition period is from June 9, 2017, to January 1, 2018. The new transition period would end on July 1, 2019. The proposed amendments to these exemptions would affect participants and beneficiaries of plans, IRA owners and fiduciaries with respect to such plans and IRAs.

DATES: Comments must be submitted on or before September 15, 2017.

ADDRESSES: All written comments should be sent to the Office of Exemption Determinations by any of the following methods, identified by RIN 1210–AB82:

Federal eRulemaking Portal: http://www.regulations.gov at Docket ID number: EBSA–2017–0004. Follow the instructions for submitting comments.

Email to: EBSA.FiduciaryRuleExamination@dol.gov.

Mail: Office of Exemption Determinations, EBSA, (Attention: D–11712, 11713, 11850), U.S. Department of Labor, 200 Constitution Avenue NW., Suite 400, Washington, DC 20210.

Hand Delivery/Courier: OED, EBSA (Attention: D–11712, 11713, 11850), U.S. Department of Labor, 122 C St. NW., Suite 400, Washington, DC 20001.

Comments will be available for public inspection in the Public Disclosure Room, EBSA, U.S. Department of Labor, Room N–1513, 200 Constitution Avenue NW., Washington, DC 20210. Comments will also be available online at *www.regulations.gov,* at Docket ID number: EBSA–2017–0004 and *www.dol.gov/ebsa,* at no charge. Do not include personally identifiable information or confidential business information that you do not want publicly disclosed. Comments online can be retrieved by most Internet search engines.