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:

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 This multichapter draft guidance is intended to explain our current thinking on how to comply with the requirements for hazard analysis and risk-based preventive controls under our rule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you can comment on any guidance at any time (see 21 CFR 10.115(g)(5)), to ensure that we consider your comment on this draft guidance before we issue the final version of the guidance, submit either electronic or written comments by February 27, 2018.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 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 described 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 laws. 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 instruct 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), HFA–305, 500 Campus Dr., College Park, MD 20740. Send two self-addressed adhesive labels.
announcing in this document is entitled “Chapter Six—Use of Heat Treatments as a Process Control.”

We intend to announce the availability of 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. You may obtain the draft guidance at either one of these Web sites (without charge) until November 21, 2017.

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.


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 proposed amendments 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–ABB2:


Email to: EBSA.FiduciaryRuleExamination@dol.gov.


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.