described in this section must be prepared and submitted in accordance with instructions (NUREG/BR–0007, NUREG/BR–0006 and NMMSS Report D–24 "Personal Computer Data Input for NRC Licensees"). Copies of these instructions may be obtained from the U.S. Nuclear Regulatory Commission, Division of Safeguards and Transportation, Washington, DC 20555. These prescribed computer readable forms replace the DOE/NRC Form 742, 742C, and 740M which have been submitted in paper form.

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PART 150—EXEMPTIONS AND CONTINUED REGULATORY AUTHORITY IN AGREEMENT STATES AND IN OFFSHORE WATERS UNDER SECTION 274

15. The authority citation for part 150 continues to read as follows:

Authority: Sec. 161, 68 Stat. 948, as amended, sec. 274, 73 Stat. 688 (42 U.S.C. 2201, 2021); sec. 201, 88 Stat. 1242, as amended (42 U.S.C. 5841).

Sections 150.3, 150.15, 150.15a, 150.31, 150.32 also issued under secs. 11e(2), 81, 68 Stat. 923, 935, as amended, secs. 83, 84, 92 Stat. 3033, 3039 (42 U.S.C. 2014e(2), 2111, 2113, 2114). Section 150.14 also issued under sec. 53, 68 Stat. 930, as amended (42 U.S.C. 2073). Section 150.15 also issued under secs. 135, 141, Pub. L. 97–425, 96 Stat. 2232, 2241 (42 U.S.C. 10155, 10161). Section 150.17a also issued under sec. 122, 68 Stat. 939 (42 U.S.C. 2152). Section 150.30 also issued under sec. 234, 83 Stat. 444 (42 U.S.C. 2282).

For the purposes of sec. 223, 68 Stat. 958, as amended (42 U.S.C. 2273); §§ 150.20(b)(2)–(5) and 150.21 are issued under sec.161b, 68 Stat. 948, as amended (42 U.S.C. 2201(b)); § 150.14 and 150.20(b)(5) are issued under sec. 161i, 68 Stat. 949, as amended (42 U.S.C. 2201(i)); and §§ 150.16–150.19 and 150.20(b)(1) are issued under sec. 161o, 68 Stat. 950, as amended (42 U.S.C. 2201(o)).

16. In § 150.16, paragraph (a) is revised to read as follows:

§ 150.16 Submission to Commission of nuclear material transfer reports.

(a) Each person who transfers and each person who receives special nuclear material pursuant to an Agreement State license shall complete and submit in computer readable form Nuclear Material Transaction Reports in accordance with instructions (NUREG/BR–0006 and NMMSS Report D–24 "Personal Computer Data Input for NRC Licensees") whenever he transfers or receives a quantity of special nuclear material of 1 gram or more of contained uranium-235, uranium-233, or plutonium. Each person who transfers this material shall submit in accordance with instructions the computer readable form promptly after the transfer takes place. Each person who receives special nuclear material shall submit in accordance with instructions the computer readable form within ten (10) days after the special nuclear material is received. Copies of the instructions may be obtained from the U.S. Nuclear Regulatory Commission, Division of Safeguards and Transportation, Washington, DC 20555. These prescribed computer readable forms replace the DOE/NRC Form 741 which have been submitted in paper form.

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17. In § 150.17, paragraph (a) is revised to read as follows:

§ 150.17 Submission to Commission of source material reports.

(a) Except as specified in paragraph (d) of this section and § 150.17a, each person who, pursuant to an Agreement State specific license, transfers or receives or adjusts the inventory in any manner by 1 kilogram or more of uranium or thorium source material of foreign origin or who imports 1 kilogram or more of uranium or thorium source material of any origin shall complete and submit in computer readable form Nuclear Material Transaction Reports in accordance with instructions (NUREG/BR–0006 and NMMSS Report D–24 "Personal Computer Data Input for NRC Licensees"). Copies of the instructions may be obtained from the U.S. Nuclear Regulatory Commission, Division of Safeguards and Transportation, Washington, DC 20555. Each person who receives the material shall submit in accordance with instructions the computer readable form within ten (10) days after the material is received.

* * * * *

Dated at Rockville, MD this 19th day of January 1993.

For the Nuclear Regulatory Commission.

Samuel J. Chilk,

Secretary of the Commission.

[FR Doc. 93–1809 Filed 1–25–93; 8:45 am]

BILLING CODE 7590–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 355

[Docket No. 80N–0042]

RIN 0905–AA06

Anticaries Drug Products for Over-the-Counter Human Use; Tentative Final Monograph; Reopening of Administrative Record; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of proposed rulemaking; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA) is extending to March 26, 1993, the comment period for the reopening of the administrative record for the proposed rulemaking for over-the-counter (OTC) anticaries drug products to obtain public comment on whether the labeling of OTC fluoride-containing drug products should include the quantity of fluoride, i.e., the specific amount of fluoride present in the product (57 FR 55199, November 24, 1992). This action is being taken because the agency recognizes the possible relevance to this issue of data and information presented at a National Institute of Dental Research (NIDR) workshop entitled "Methods for Assessing Fluoride Accumulation and Effects in the Body," held on January 13 to 15, 1993. This proposal is part of the ongoing review of OTC drug products conducted by FDA.

DATES: Written comments by March 26, 1993.

ADDRESSES: Written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, rm. 1–23, 12420 Parklawn Dr., Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: William E. Gilbertson, Center for Drug Evaluation and Research (HFD–810), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–295–8000.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of November 24, 1992 (57 FR 55199), FDA published a notice of proposed rulemaking reopening the administrative record for the rulemaking for OTC anticaries drug products. Interested persons were given until January 25, 1993 to respond.

In that document, the agency discussed recommendations made by the Public Health Service Ad Hoc Subcommittee on Fluoride of the

Committee to Coordinate Environmental Health and Related Programs (the Subcommittee) that the U.S. Public Health Service sponsor a scientific conference(s) to recommend both the optimal level of flouride exposure from all sources combined (including drinking water) and the appropriate usage of flouride-containing dental products. In considering the Subcommittee's recommendations, FDA requested input from three professional associations on the possibility of having OTC fluoride-containing drug products labeled to identify their fluoride levels. One association (dental group) submitted information in support of having fluoride levels listed in product labeling as percent weight/volume. The other two associations (trade groups) responded in opposition to changing the labeling of OTC fluoride-containing drug products to identify fluoride levels.

FDA has received a request from the latter two associations to extend the comment period for an additional 60 days to permit industry and other interested parties to prepare and submit additional information that may be developed for or in response to a NIDR workshop to be held on January 13 to 15, 1993. The request included a copy of the preliminary agenda for the workshop entitled "Methods for Fluoride Accumulation and Effects in the Body" (Ref. 1). The request stated its belief that important information may be forthcoming from the workshop.

FDA has carefully considered the request and believes that additional time for comment is in the public interest. The agency concurs with the request that important information may be forthcoming from the workshop. In fact, agency staff attended the workshop. Thus, the agency considers the limited extension of the comment period requested to be appropriate. Accordingly, the comment period is extended to March 26, 1993.

Interested persons may, on or before March 26, 1993, submit to the Dockets Management Branch (address above) written comments on whether the labeling of OTC fluoride-containing drug products should include the quantity of fluoride present in the product and how that information should be presented. Three copies of any comments are to be submitted, except that individuals may submit one copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday.

Reference

(1) Comment No. EXT 8, Docket No. 80N–0042, Docket Management Branch.

Dated: January 15, 1993.

Michael R. Taylor,

Deputy Commissioner for Policy.

[FR Doc. 93–1826 Filed 1–25–93; 8:45 am]

BILLING CODE 4160–01–M

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Part 1

[EE–62–92]

RIN 1545–AR09

Nondiscrimination Requirements for Qualified Plans; Hearing

AGENCY: Internal Revenue Service, Treasury.

ACTION: Notice of public hearing on proposed regulations.

SUMMARY: This document provides notice of a public hearing on proposed Income Tax Regulations requiring contributions or benefits provided under a tax-qualified retirement plan not discriminate in favor of highly compensated employees.

DATES: The public hearing will begin on Friday, April 23, 1993, and continue if necessary, on Monday, April 26, 1993, beginning each day at 10 a.m. Requests to speak and outlines of oral comments must be received by Friday, April 2, 1993.

ADDRESSES: The public hearing will be held in the IRS Auditorium, Seventh floor, 7400 Corridor, Internal Revenue Building, 1111 Constitution Avenue, NW., Washington, DC. Requests to speak and outlines of oral comments should be submitted to the Internal Revenue Service, P.O. Box 7604, Ben Franklin Station, Attn: CC:CORP:T:R [EE–62–92], room 5228, Washington, DC 20044.

FOR FURTHER INFORMATION CONTACT: Mike Slaughter of the Regulations Unit, Assistant Chief Counsel (Corporate), 202–622–7190 (not a toll-free number).

SUPPLEMENTARY INFORMATION: The subject of the public hearing is proposed amendments to the final regulations under section 401(a)(4) of the Internal Revenue Code of 1986. These proposed regulations were published in the **Federal Register** on Tuesday, January 12, 1993 (58 FR 3876).

The rules of § 601.601(a)(3) of the "Statement of Procedural Rules" (26 CFR part 601) shall apply with respect to the public hearing. Persons who have submitted written comments within the time prescribed in the notice of proposed rulemaking and who also desire to present oral comments at the hearing on the proposed regulations should submit not later than Friday, April 2, 1993, an outline of the oral comments/testimony to be presented at the hearing and the time they wish to devote to each subject.

Each speaker (or group of speakers representing a single entity) will be limited to 10 minutes for an oral presentation exclusive of the time consumed by the questions from the panel for the government and answers to these questions.

Because of controlled access restrictions, attendees cannot be admitted beyond the lobby of the Internal Revenue Building until 9:45 a.m.

An agenda showing the scheduling of the speakers will be made after outlines are received from the persons testifying. Copies of the agenda will be available free of charge at the hearing.

By direction of the Commissioner of Internal Revenue.

Dale D. Goode,

Federal Register Liaison Officer, Assistant Chief Counsel (Corporate).

[FR Doc. 93–1404 Filed 1–25–93; 8:45 am]

BILLING CODE 4830–01–M

OFFICE OF MANAGEMENT AND BUDGET

Office of Federal Procurement Policy

48 CFR Part 9904

Cost Accounting Standards Board; Cost Accounting Standards for Composition, Measurement, Adjustment, and Allocation of Pension Costs

AGENCY: Cost Accounting Standards Board, Office of Federal Procurement Policy, OMB.

ACTION: Advance notice of proposed rulemaking (ANPRM).

SUMMARY: The Office of Federal Procurement Policy, Cost Accounting Standards Board (CASB), proposes to revise the Cost Accounting Standards relating to accounting for pension costs under negotiated government contracts. Section 26(g)(1) of the Office of Federal Procurement Policy Act, 41 U.S.C. 422(g)(1), requires that the Board, prior to the promulgation of any new or revised Cost Accounting Standard, publish a report and an ANPRM. This ANPRM addresses certain problems that