

4.9(c).⁷ Your comment will be kept confidential only if the FTC General Counsel, in his or her sole discretion, grants your request in accordance with the law and the public interest.

Postal mail addressed to the Commission is subject to delay due to heightened security screening. As a result, we encourage you to submit your comments online. To make sure that the Commission considers your online comment, you must file it at <https://ftcpublic.commentworks.com/ftc/unavailabilityruleanpr>, by following the instructions on the Web-based form. If this Notice appears at <http://www.regulations.gov/#!home>, you also may file a comment through that Web site.

If you file your comment on paper, write "16 CFR Part 424—Retail Food Store Advertising Rule, Project No. P104203" on your comment and on the envelope, and mail or deliver it to the following address: Federal Trade Commission, Office of the Secretary, Room H-113 (Annex N), 600 Pennsylvania Avenue, NW., Washington, DC 20580. If possible, submit your paper comment to the Commission by courier or overnight service.

Visit the Commission Web site at <http://www.ftc.gov> to read this Notice and the news release describing it. The FTC Act and other laws that the Commission administers permit the collection of public comments to consider and use in this proceeding as appropriate. The Commission will consider all timely and responsive public comments that it receives on or before October 19, 2011. You can find more information, including routine uses permitted by the Privacy Act, in the Commission's privacy policy, at <http://www.ftc.gov/ftc/privacy.htm>.

List of Subjects in 16 CFR Part 424

Advertising, Foods, Trade practices.

Authority: 15 U.S.C. 41–58.

By direction of the Commission.

Donald S. Clark,
Secretary.

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⁷In particular, the written request for confidential treatment that accompanies the comment must include the factual and legal basis for the request, and must identify the specific portions of the comment to be withheld from the public record. See FTC Rule 4.9(c), 16 CFR 4.9(c).

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Part 51

[REG–112805–10]

RIN 1545–BJ39

Branded Prescription Drug Fee

AGENCY: Internal Revenue Service (IRS), Treasury.

ACTION: Notice of proposed rulemaking by cross-reference to temporary regulations.

SUMMARY: In the Rules and Regulations section of this issue of the **Federal Register**, the IRS is issuing temporary regulations relating to the branded prescription drug fee imposed by the Affordable Care Act (ACA). The regulations affect persons engaged in the business of manufacturing or importing certain branded prescription drugs. The text of the temporary regulations also serves as the text of the proposed regulations.

DATES: Written and electronic comments and requests for a public hearing must be received by November 16, 2011.

ADDRESSES: Send submissions to: CC:PA:LPD:PR (REG–112805–10), room 5205, Internal Revenue Service, P.O. Box 7604, Ben Franklin Station, Washington, DC 20044. Submissions may be hand-delivered Monday through Friday between the hours of 8 a.m. and 4 p.m. to: CC:PA:LPD:PR (REG–112805–10), Courier's Desk, Internal Revenue Service, 1111 Constitution Avenue, NW., Washington, DC, or sent electronically via the Federal eRulemaking Portal at <http://www.regulations.gov> (IRS REG–112805–10).

FOR FURTHER INFORMATION CONTACT: Concerning the proposed regulations, Celia Gabrysh at (202) 622–3130; concerning submissions of comments and requests for a hearing *Richard.A.Hurst@irs.counsel.treas.gov*, (202) 622–7180 (not toll free numbers).

SUPPLEMENTARY INFORMATION:

Paperwork Reduction Act

The collection of information contained in this notice of proposed rulemaking has been approved by the Office of Management and Budget in accordance with the Paperwork Reduction Act of 1995 (44 U.S.C. 3507(d)) and assigned control number 1545–2209.

Comments on the collection of information should be sent to the Office of Management and Budget, Attn: Desk

Officer for the Department of the Treasury, Office of Information and Regulatory Affairs, Washington, DC 20503, with copies to the Internal Revenue Service, *Attn:* IRS Reports Clearance Officer, SE:W:CAR:MP:T:T:SP, Washington, DC 20224. Comments on the collection of information should be received by October 17, 2011. Comments are specifically requested concerning:

Whether the proposed collection of information is necessary for the proper performance of the functions of the Internal Revenue Service, including whether the information will have practical utility;

The accuracy of the estimated burden associated with the proposed collection of information;

How the quality, utility, and clarity of the information to be collected may be enhanced;

How the burden of complying with the proposed collection of information may be minimized, including through the application of automated collection techniques or other forms of information technology; and

Estimates of capital or start-up costs of operation, maintenance, and purchase of service to provide information.

The collection of information in this proposed regulation is in § 51.7. This information is necessary to evaluate whether an error report regarding a preliminary fee calculation is valid and justifies an adjustment to the preliminary fee calculation. The likely respondents are manufacturers and importers of branded prescription drugs.

Estimated total annual reporting and/or recordkeeping burden: 1800 hours.

Estimated annual burden per respondent/recordkeeper: 40 hours.

Estimated number of respondents and/or recordkeepers: 45.

Estimated frequency of responses: Annually.

An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless the collection of information displays a valid control number.

Books or records relating to a collection of information must be retained as long as their contents may become material in the administration of any internal revenue law. Generally, tax returns and tax return information are confidential, as required by 26 U.S.C. 6103.

Background

Temporary regulations in the Rules and Regulations section of this issue of the **Federal Register** add a new part, part 51, to subchapter D, Miscellaneous

Excise Taxes. Part 51 provides guidance on the annual fee imposed on covered entities engaged in the business of manufacturing or importing branded prescription drugs by section 9008 of the ACA. The text of those regulations also serves as the text of these proposed regulations. The preamble to the temporary regulations explains the new part.

Special Analyses

It has been determined that this notice of proposed rulemaking is not a significant regulatory action as defined in Executive Order 12866, as supplemented by Executive Order 13563. Therefore, a regulatory flexibility assessment is not required. It also has been determined that section 553(b) of the Administrative Procedure Act (5 U.S.C. chapter 5) does not apply to these regulations. It is hereby certified that the collection of information in these regulations will not have a significant economic impact on a substantial number of small entities. This certification is based on the fact that these regulations primarily affect large corporations. Thus, Treasury Department and the IRS do not expect a substantial number of small entities to be effected. Therefore, a Regulatory Flexibility Analysis under the Regulatory Flexibility Act (5 U.S.C. chapter 6) is not required. Pursuant to section 7805(f) of the Internal Revenue Code, this notice of proposed rulemaking has been submitted to the Chief Counsel for Advocacy of the Small Business Administration for comment on its impact on small business.

Comments and Requests for a Public Hearing

Before these proposed regulations are adopted as final regulations, consideration will be given to any written comments (a signed original and eight (8) copies) or electronic comments that are submitted timely to the IRS. Comments are requested on all aspects of the proposed regulations. In addition, the IRS and the Treasury Department specifically request comments on the clarity of the proposed regulations and how they may be made easier to understand. All comments will be available for public inspection and copying. A public hearing may be scheduled if requested in writing by any person that timely submits written comments. If a public hearing is scheduled, notice of the date, time, and place for the hearing will be published in the **Federal Register**.

Drafting Information

The principal author of these regulations is Celia Gabrysh, Office of Associate Chief Counsel (Passthroughs and Special Industries). However, other personnel from the IRS and the Treasury Department participated in their development.

List of Subjects in 26 CFR Part 51

Drugs, Reporting and recordkeeping requirements.

Proposed Amendments to the Regulations

Accordingly, and under the authority of 26 U.S.C. 7805 (sec. 9008, Pub. L. 111-347 (124 Stat. 119)), 26 CFR part 51 is proposed to be added to read as follows:

PART 51—BRANDED PRESCRIPTION DRUGS

[The text of proposed §§ 51.1 through 51.11 is the same as the text of §§ 51.1T through 51.11T published elsewhere in this issue of the **Federal Register**.]

[The text of proposed § 51.6302-1 is the same as the text of paragraphs (a) and (b) of § 51.6302-1T published elsewhere in this issue of the **Federal Register**.]

Sarah Hall Ingram,

Deputy Commissioner for Services and Enforcement.

[FR Doc. 2011-21012 Filed 8-15-11; 11:15 am]

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POSTAL REGULATORY COMMISSION

39 CFR Part 3020

[Docket No. RM2011-8; Order No. 666]

Administrative Practice and Procedure, Postal Service

AGENCY: Postal Regulatory Commission.

ACTION: Notice of proposed rulemaking.

SUMMARY: A Federal statute directs the Commission to implement a modern classification system. This proposal responds to that directive by presenting a comprehensive Mail Classification Schedule. Issuance of this document will allow the Commission to consider comments and, if appropriate, to make revisions prior to adoption of a final schedule.

DATES: *Comments are due:* September 6, 2011.

ADDRESSES: Submit comments electronically by accessing the "Filing Online" link in the banner at the top of the Commission's Web site (<http://www.prc.gov>) or by directly accessing

the Commission's Filing Online system at <https://www.prc.gov/prc-pages/filing-online/login.aspx>. Commenters who cannot submit their views electronically should contact the person identified in the **FOR FURTHER INFORMATION CONTACT** section as the source for case-related information for advice on alternatives to electronic filing.

FOR FURTHER INFORMATION CONTACT:

Stephen L. Sharfman, General Counsel, at 202-789-6820 (case-related information) or DocketAdmins@prc.gov (electronic filing assistance).

SUPPLEMENTARY HISTORY: *Regulatory History:*

72 FR 29284, May 25, 2007;
72 FR 33261, June 15, 2007;
72 FR 50744, September 4, 2007.

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I. Introduction

The Postal Regulatory Commission (Commission) establishes a rulemaking docket pursuant to its responsibilities under the Postal Accountability and Enhancement Act (PAEA), Public Law 109-435, 120 Stat. 3198, December 20, 2006, to consider modifications to the Commission's rules governing the Mail Classification Schedule (MCS). Modifications are proposed to add material describing some Postal Service products and make conforming changes. The Commission provides this notice and opportunity for comment on whether the Commission should incorporate the proposed modifications by final rule into the Commission's rules at 39 CFR 3020, Subpart A—Mail Classification Schedule.

For products currently being offered by the Postal Service, this rulemaking does not add products to, remove products from, or transfer products between the existing market dominant or competitive product lists. However, this rulemaking does reorganize how products appear within each individual list. This reorganization is most apparent within the competitive product list where, at the suggestion of the Postal Service, the vestiges of "class" groupings have been replaced with functional product groupings.

Additionally, the currently published product lists require updating to remove