assumptions that support our conclusions.

- FAA experts, industry, and public participants are expected to hold a full discussion of all technical material presented at the meeting. If you present conclusions on this subject, you must submit data that supports your conclusions. All data will be part of the Rulemaking Dockets.
- We will try and accommodate all speakers. In order to do this, we may need to limit the time for presenters.
- We can make sign and oral interpretation available at the meeting, as well as an assistive listening device. If you need this assistance, make your request to FAA at least 10 days prior to the public meeting.
- A court reporter will record the discussions of the meeting. We will place the transcript of the meeting in the Rules Dockets. If you would like to purchase a copy of the transcript, you must contact the court reporter directly. We will provide further information at the meeting.
- We will review and consider all material presented. Position papers or materials that present views or information related to the proposed ADs may be accepted at the discretion of the presiding officer and placed in the Rules Dockets. The FAA requests that you provide 10 copies of all materials for distribution to the panel members. You have the choice on whether you want to present copies of the material to the audience.
- Panel member statements are intended to facilitate discussion of or to clarify issues. The FAA will consider comments made at this meeting before making a final decision on the issuance of any airworthiness directive.
- The meetings are designed to solicit public views and more complete information on the proposed ADs. Therefore, we will conduct the meeting in an informal and nonadversial manner.

Issued in Kansas City, Missouri, on January 15, 2004.

Dorenda D. Baker,

Manager, Small Airplane Directorate, Aircraft Certification Service.

[FR Doc. 04–1658 Filed 1–26–04; 8:45 am]

BILLING CODE 4910–13–P

FEDERAL TRADE COMMISSION

16 CFR Chapter I

Notice of Revised Regulatory Review Schedule

AGENCY: Federal Trade Commission.

ACTION: Notice of revised regulatory review schedule.

SUMMARY: The Federal Trade Commission (''Commission'') has a program of systematic review of all of its rules and guides. The Commission hereby gives notice that, based on its current ongoing review proceedings, as well as additional rulemaking proceedings required by new legislation, it does not intend to announce review of any additional rules or guides during 2004. The ten-year regulatory review schedule previously published by the Commission, 67 FR 9630 (Mar. 4, 2002), has been modified accordingly.

FOR FURTHER INFORMATION CONTACT: Neil Blickman, Federal Trade Commission, Bureau of Consumer Protection, Division of Enforcement, 600 Pennsylvania Ave., NW., Washington DC 20580, (202) 326–3038.

SUPPLEMENTARY INFORMATION: The Commission has decided not to initiate review of any additional rules or guides during 2004. Currently, the Commission has ongoing review or amendment proceedings that relate to a number of its rules and guides. In addition, during 2004, the Commission will be required to promulgate rules pursuant to the Fair and Accurate Credit Transactions Act of 2003, Pub. L. 108–159 (requiring at least 25 separate rules and 8 studies); the Fairness to Contact Lens Consumers Act of 2003, Pub. L. 108–164; and the Controlling the Assault of Non-Solicited Pornography and Marketing Act of 2003, Pub. L. 108–187. Accordingly, the Commission proposes a revised ten-year regulatory review schedule. A copy of this tentative schedule is appended. The Commission may, in its discretion, modify or reorder the schedule in the future to incorporate new legislative rules, or to respond to external factors (such as changes in the law) or other considerations.

Authority: 15 U.S.C. 41–58.

By direction of the Commission.

Donald S. Clark,

Secretary.

APPENDIX—REGULATORY REVIEW MODIFIED TEN-YEAR SCHEDULE

16 CFR part	Topic	Year to review
18	Guides for the Nursery Industry	2005
410	TV Picture Tube Size Rule	2005
424	Retail Food Store Advertising and Marketing Practices Rule	2005
14	Administrative Interpretations, General Policy Statements, and Enforcement Policy Statements	2006
311	Recycled Oil Rule	2006
312	Children's Online Privacy Protection Rule	2006
444	Credit Practices Rule	2006
455	Used Car Rule	2006
24	Guides for Select Leather and Imitation Leather Products	2007
435	Mail or Telephone Order Merchandise Rule	2007
500	Regulations Under Section 4 of the Fair Packaging and Labeling Act (''FPLA'')	2007
501	Exemptions from Part 500 of the FPLA	2007
502	Regulations Under Section 5(C) of the FPLA	2007
503	Statements of General Policy or Interpretations Under the FPLA	2007
305	Appliance Labeling Rule	2008
306	Automotive Fuel Ratings, Certification and Posting Rule	2008
429	Cooling Off Rule	2008
601	Summary of Consumer Rights, Notice of User Responsibilities, and Notice of Furnisher Responsibilities under the Fair Credit Reporting Act.	2008
254	Guides for Private Vocational and Distance Education Schools	2009
260	Guides for the use of Environmental Marketing Claims	2009
300	Rules and Regulations under the Wool Products Labeling Act	2009
301	Rules and Regulations under the Fur Products Labeling Act	2009
303	Rules and Regulations under the Textile Fiber Products Identification Act	2009
425	Rule Concerning the Use of Negative Option Plans	2009

APPENDIX—REGULATORY REVIEW MODIFIED TEN-YEAR SCHEDULE—Continued

16 CFR part	Topic	Year to review
239	Guides for the Advertising of Warranties and Guarantees	2010
433	Preservation of Consumers' Claims and Defenses Rule	2010
700	Interpretations of Magnuson-Moss Warranty Act	2010
701	Disclosure of Written Consumer Product Warranty Terms and Conditions	2010
702	Pre-sale Availability of Written Warranty Terms	2010
703	Informal Dispute Settlement Procedures	2010
23	Guides for the Jewelry, Precious Metals, and Pewter Industries	2011
423	Care Labeling Rule	2011
20	Guides for the Rebuilt, Reconditioned and Other Used Automobile Parts Industry	2012
233	Guides Against Deceptive Pricing	2012
238	Guides Against Bait Advertising	2012
240	Guides for Advertising Allowances and Other Merchandising Payments and Services	2012
251	Guide Concerning Use of the word ''Free'' and Similar Representations	2012
259	Guide Concerning Fuel Economy Advertising for New Automobiles	2012
310	Telemarketing Sales Rule	2013
801	Hart-Scott-Rodino Antitrust Improvements Act Coverage Rules	2013
802	Hart-Scott-Rodino Antitrust Improvements Act Exemption Rules	2013
803	Hart-Scott-Rodino Antitrust Improvements Act Transmittal Rules	2013

[FR Doc. 04–1690 Filed 1–26–04; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 101

[Docket No. 2003N–0496]

RIN 0910–AF09

Food Labeling: Health Claims; Dietary Guidance; Extension of Comment Period

AGENCY: Food and Drug Administration, HHS.

ACTION: Advance notice of proposed rulemaking; extension of comment period.

SUMMARY: The Food and Drug Administration (FDA) is extending to February 25, 2004, the comment period for the advance notice of proposed rulemaking (ANPRM) that appeared in the **Federal Register** of November 25, 2003 (68 FR 66040). In the ANPRM, FDA requested comments on alternatives for regulating qualified health claims in the labeling of conventional human foods and dietary supplements. FDA also solicited comments on various other issues related to health claims and on the appropriateness and nature of dietary guidance statements on conventional foods and dietary supplement labels. The agency is taking this action in response to requests for an extension to allow interested persons additional time to submit comments.

DATES: Submit written and electronic comments by February 25, 2004.

ADDRESSES: Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to *http://www.fda.gov/dockets/ecomments*.

FOR FURTHER INFORMATION CONTACT: Nancy T. Crane, Center for Food Safety and Applied Nutrition (HFS–830), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740–3835, 301–436–1456, or e-mail: *Nancy.Crane@cfsan.fda.gov*.

SUPPLEMENTARY INFORMATION:

I. Background

In the **Federal Register** of November 25, 2003 (68 FR 66040), FDA published an ANPRM with a 60-day comment period to request comments on alternatives for regulating qualified health claims in the labeling of conventional human foods and dietary supplements. FDA also solicited comments on various other issues related to health claims and on the appropriateness and nature of dietary guidance statements on conventional foods and dietary supplement labels. Comments on the regulatory alternatives and the additional topics will inform FDA's rulemaking to establish regulations for qualified health claims, as well as any policy initiative(s) that FDA may undertake to provide information to consumers to help them make wise food choices.

The agency has received multiple requests for either a 30-day or 60-day extension of the comment period for the ANPRM. Each request conveyed concern that the current 60-day comment period does not allow sufficient time to develop a meaningful or thoughtful response to the ANPRM. In addition, two requests noted that the current comment period occurred during a period of time that included the Thanksgiving and year-end holidays. All of the requests explained that an extension is necessary due to the complexity, implications, and/or importance of the rulemaking on health claims and dietary guidance in food and dietary supplement labeling.

FDA has considered the requests and is extending the comment period for the ANPRM for 30 days, until February 25, 2004. The agency believes that a 30-day extension allows adequate time for interested persons to submit comments without significantly delaying rulemaking on these important issues.

II. Request for Comments

Interested persons may, on or before February 25, 2004, submit to the Division of Dockets Management (see **ADDRESSES**) written or electronic comments on this ANPRM. Submit two copies of any comments, except that individuals may submit one copy. Identify comments with the docket number found in brackets in the heading of this document. Interested persons may review received comments in the office above between 9 a.m. and 4 p.m., Monday through Friday.

Dated: January 22, 2004.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 04–1772 Filed 1–23–04; 10:57 am]
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