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,

M , A i D t t , A t C t t .

[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.

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

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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
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