purposes (Chapter 8, Title VI of the Omnibus Budget Reconciliation Act of 1981, Pub. L. 97–35, 42 U.S.C. 9871 *et seq.*) and as amended now and hereafter.

3. Authority for the Child Care and Development Block Grants, under Section 5082 of OBRA 1990, (42 U.S.C. 9858 *et seq.*), and as amended now and hereafter.

4. Authority to administer the provisions of the Child Care and Development Block Grant Amendments of 1996, 42 U.S.C. 9801 note, under Sections 601–615 of the Personal Responsibility and Work Opportunity Reconciliation Act of 1996, 42 U.S.C. 1305 note, 42 U.S.C. 601 *et seq.*, and as amended now and hereafter.

(b) Limitations

1. This delegation shall be exercised under the Department's existing policies on delegations and regulations.

2. This delegation does not include the authority to submit reports to Congress and shall be exercised under financial and administrative requirements applicable to all Administration for Children and Families authorities.

3. The approval or disapproval of grant applications and the making of grant awards require concurrence of the appropriate Grants Officer. The approval or disapproval of contract proposals and awards are subject to the requirements of the Federal Acquisition Regulations and requires the concurrence of the Contracting Officer.

4. This delegation of authority does not include the authority to sign and issue notices of grant awards.

5. This delegation of authority does not include the authority to appoint Action Officials for Audit Resolution.

6. This delegation of authority does not include the authority to appoint Central Office or Regional Office Grant Officers for the administration of the child care related programs.

7. This delegation of authority does not include the authority to hold hearings.

8. This delegation of authority does not include the authority to approve or disapprove awards for grants or contracts for research, demonstration, or evaluations relating to child care.

9. Any redelegation shall be in writing and prompt notification must be provided to all affected managers, supervisors, and other personnel, and requires the concurrence of the Deputy Assistant Secretary for Administration.

(c) Effect on Existing Delegations

This delegation supersedes any previous delegation of authority pertaining to authorities delegated herein.

(d) Effective Date

This delegation was effective upon the date of signature.

I hereby affirm and ratify any actions taken by the Director, Office of Family Assistance, which involved the exercise of the authorities delegated herein prior to the effective date of this delegation.

Dated: February 16, 2007.

Wade F. Horn,

Assistant Secretary for Children and Families.

[FR Doc. E7–3306 Filed 2–26–07; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Statement of Organization, Functions and Delegation of Authority

Notice is hereby given that I have delegated to the Director, Office of Family Assistance, the following authorities vested in me by the Secretary of Health and Human Services in the memorandum dated August 20, 1991, pertaining to the Head Start Program and the Child Development Associate Scholarship Assistance Grants Program, in the memorandum dated August 20, 1991, pertaining to the Omnibus Budget Reconciliation Act of 1981, in the memorandum dated August 20, 1991, pertaining to the Omnibus Budget Reconciliation Act of 1990 (OBRA 1990, Pub. L. 101–508), and in the memorandum dated September 16, 1997, pertaining to the Personal Responsibility and Work Opportunity Reconciliation Act of 1996 (PRWORA, Pub. L. 104–193).

(a) Authorities Delegated

1. Authority to administer the provisions of the Child Development Associate Scholarship Assistance Act, 42 U.S.C. 10901–10905, and as amended now and hereafter.

2. Authority to administer the provisions of Subchapter D—Grants for Planning and Development of Dependent Care Programs and for other purposes (Chapter 8, Title VI of the Omnibus Budget Reconciliation Act of 1981, Pub. L. 97–35, 42 U.S.C. 9871 *et seq.*) and as amended now and hereafter.

3. Authority for the Child Care and Development Block Grants, under Section 5082 of OBRA 1990, (42 U.S.C. 9858 *et seq.*), and as amended now and hereafter.

4. Authority to administer the provisions of the Child Care and Development Block Grant Amendments of 1996, 42 U.S.C. 9801 note, under Sections 601–615 of the Personal Responsibility and Work Opportunity Reconciliation Act of 1996, 42 U.S.C. 1305 note, 42 U.S.C. 601 *et seq.*, and as amended now and hereafter.

(b) Limitations

1. This delegation shall be exercised under the Department's existing policies on delegations and regulations.

2. This delegation does not include the authority to submit reports to Congress and shall be exercised under financial and administrative requirements applicable to all Administration for Children and Families authorities.

3. The approval or disapproval of grant applications and the making of grant awards require concurrence of the appropriate Grants Officer. The approval or disapproval of contract proposals and awards are subject to the requirements of the Federal Acquisition Regulations and requires the concurrence of the Contracting Officer.

4. This delegation of authority does not include the authority to sign and issue notices of grant awards.

5. This delegation of authority does not include the authority to appoint Action Officials for Audit Resolution.

6. This delegation of authority does not include the authority to appoint Central Office or Regional Office Grant Officers for the administration of the child care related programs.

7. This delegation of authority does not include the authority to hold hearings.

8. This delegation of authority does not include the authority to approve or disapprove awards for grants or contracts for research, demonstration, or evaluations relating to child care.

9. Any redelegation shall be in writing and prompt notification must be provided to all affected managers, supervisors, and other personnel, and requires the concurrence of the Deputy Assistant Secretary for Administration.

(c) Effect on Existing Delegations

This delegation supersedes any previous delegation of authority pertaining to authorities delegated herein.

(d) Effective Date

This delegation was effective upon the date of signature.

I hereby affirm and ratify any actions taken by the Director, Office of Family Assistance, which involved the exercise of the authorities delegated herein prior to the effective date of this delegation.

Dated: February 16, 2007.

Wade F. Horn,

Assistant Secretary for Children and Families.

[FR Doc. E7–3325 Filed 2–26–07; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0053]

Agency Information Collection Activities; Proposed Collection; Comment Request; Food Labeling Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing an opportunity for public comment on the proposed collection of certain information by the agency. Under the Paperwork Reduction Act of 1995 (the PRA), Federal agencies are required to publish notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, and to allow 60 days for public comment in response to the notice. This notice solicits comments on the information collection provisions in FDA's food labeling regulations.

DATES: Submit written or electronic comments on the collection of information by April 30, 2007.

ADDRESSES: Submit electronic comments on the collection of information to: *http://www.fda.gov/dockets/ecomments*. Submit written comments on the collection of information to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. ''Collection of information'' is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal agencies to provide a 60-day notice in the **Federal Register** concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

With respect to the following collection of information, FDA invites comments on these topics: (1) Whether the proposed collection of information is necessary for the proper performance of FDA's functions, including whether the information will have practical utility; (2) the accuracy of FDA's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques, when appropriate, and other forms of information technology.

Food Labeling Regulations—21 CFR Parts 101, 102, 104, and 105 (OMB Control Number 0910–0381)—Extension

FDA regulations require food producers to disclose to consumers and others specific information about themselves or their products on the label or labeling of their products. Related regulations require that food producers retain records establishing the basis for the information contained in the label or labeling of their products and provide those records to regulatory officials. Finally, certain regulations provide for the submission of food labeling petitions to FDA. FDA's food labeling regulations under parts 101, 102, 104, and 105 (21 CFR parts 101, 102, 104, and 105) were issued under the authority of sections 4, 5, and 6 of the Fair Packaging and Labeling Act (the FPLA) (15 U.S.C. 1453, 1454, and 1455) and under sections 201, 301, 402, 403, 409, 411, 701, and 721 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 321, 331, 342, 343, 348, 350, 371, and 379e). Most of these regulations derive from section 403 of the act, which provides that a food product shall be deemed to be misbranded if, among other things, its label or labeling fails to bear certain required information concerning the food product, is false or misleading in any particular, or bears certain types of unauthorized claims. The disclosure requirements and other collections of information in the regulations in parts 101, 102, 104, and 105 are necessary to ensure that food products produced or sold in the United States are in compliance with the labeling provisions of the act and the FPLA.

Section 101.3 of FDA's food labeling regulations requires that the label of a food product in packaged form bear a statement of identity (i.e., the name of the product), including, as appropriate, the form of the food or the name of the food imitated. Section 101.4 prescribes requirements for the declaration of ingredients on the label or labeling of food products in packaged form. Section 101.5 requires that the label of a food product in packaged form specify the name and place of business of the manufacturer, packer, or distributor and, if the food producer is not the manufacturer of the food product, its connection with the food product. Section 101.9 requires that nutrition information be provided for all food products intended for human consumption and offered for sale, unless an exemption in § 101.9(j) applies to the product. Section 101.9(g)(9) also provides for the submission to FDA of requests for alternative approaches to nutrition labeling. Finally, § 101.9(j)(18) provides for the submission to FDA of notices from firms claiming the small business exemption from nutrition labeling.

Section 101.10 requires that restaurants provide nutrition information, upon request, for any food or meal for which a nutrient content claim or health claim is made. Section 101.12(b) provides the reference amount that is used for determining the serving sizes for specific products, including baking powder, baking soda, and pectin. Section 101.12(e) provides that a manufacturer that adjusts the reference amount customarily consumed (RACC) of an aerated food for the difference in density of the aerated food relative to the density of the appropriate nonaerated reference food must be prepared to show FDA detailed protocols and records of all data that were used to determine the density-adjusted RACC. Section 101.12(g) requires that the label or labeling of a food product disclose the serving size that is the basis for a claim made for the product if the serving size on which the claim is based differs from the RACC. Section 101.12(h) provides for the submission of petitions to FDA to request changes in the reference amounts defined by regulation.