

Dated: January 20, 2005.
Alvin Hall,
*Director, Management Analysis and Services
 Office, Centers for Disease Control and
 Prevention.*
 [FR Doc. 05-1389 Filed 1-25-05; 8:45 am]
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**DEPARTMENT OF HEALTH AND
 HUMAN SERVICES**

**Administration for Children and
 Families**

**Submission for OMB Review;
 Comment Request**

Title: Child Care and Development
 Fund Annual Financial Report for
 Tribes (ACF-696T).

OMB No.: 0970-0195.

Description: The Child Care and
 Development Fund (CCDF) annual
 financial reporting form (ACF-696T)
 provides a mechanism for Indian tribes
 to report expenditures under the CCDF
 program. The CCDF program provides

funds to tribes, as well as States and
 Territories, to assist low-income
 families in obtaining child care so that
 they can work or attend training/
 education, and to improve the quality of
 care. Information collected via the ACF-
 696T allows the Administration for
 Children and Families (ACF) to monitor
 expenditures and to estimate outlays,
 and may be used to prepare ACF budget
 submissions to Congress. The proposed
 information collection is identical to the
 currently used ACF-696T form for
 which the Office of Management and
 Budget (OMB) approval expires on
 March 31, 2005.

Respondents: Indian tribes and tribal
 organizations that are CCDF grantees.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
ACF-696T CCDF Financial Reporting Form for Tribes	232	1	7.5	1,740

*Estimated Total Annual Burden
 Hours:* 1,740.

Additional Information: Copies of the
 proposed collection may be obtained by
 writing to The Administration for
 Children and Families, Office of
 Information Services, 730 L'Enfant
 Promenade, SW., Washington, DC
 20447, Attn: ACF Reports Clearance
 Officer.

OMB Comment: OMB is required to
 make a decision concerning the
 collection of information between 30
 and 60 days after publication of this
 document in the **Federal Register**.
 Therefore, a comment is best assured of
 having its full effect if OMB receives it
 within 30 days of publication. Written
 comments and recommendations for the
 proposed information collection should
 be sent directly to the following: Office
 of Management and Budget, Paperwork
 Reduction Project, Attn: Desk Officer for

ACF, e-mail address:
Katherine_T.Astrich@omb.eop.

Dated: January 18, 2005.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05-1397 Filed 1-25-05; 8:45 am]

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**DEPARTMENT OF HEALTH AND
 HUMAN SERVICES**

**Administration for Children and
 Families**

**Proposed Information Collection
 Activity; Comment Request Proposed
 Projects**

Title: LIHEAP Quarterly Allocation
 Estimates.

OMB No.: 0970-0037.

Description: The Low Income Home
 Energy Assistance Program (LIHEAP)

Quarterly Allocation Estimates Form-
 535 is a one-page form that is sent to 50
 State grantees and the District of
 Columbia. It is also sent to tribal
 grantees that receive over \$1 million
 annually and that directly administer
 the LIHEAP Program. Grantees are asked
 to complete and submit the form in the
 4th quarter of every fiscal year. The data
 collected on the form are the grantee's
 estimates of obligations that they expect
 to make each quarter during the
 upcoming fiscal year. This is the only
 method used to request anticipate
 distribution of the grantee's LIHEAP
 funds for the program year. The
 information is used to disburse LIHEAP
 funds in accordance with grantee needs
 and to develop OMB apportionment
 requests.

Respondents: State, local or tribal
 Government.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Form ACF-535	55	1	.25	13.75

*Estimated Total Annual Burden
 Hours:* 13.75.

In compliance with the requirements
 of section 3506(c)(2)(A) of the
 Paperwork Reduction Act of 1995, the
 Administration for Children and
 Families is soliciting public comment
 on the specific aspects of the

information collection described above.
 Copies of the proposed collection of
 information can be obtained and
 comments may be forwarded by writing
 to the Administration for Children and
 Families, Office of Information Services,
 370 L'Enfant Promenade, SW.,
 Washington, DC 20447, Attn: ACF

Reports Clearance Officer. E-mail:
grjohnson@acf.hhs.gov. All requests
 should be identified by the title of the
 information collection.

The Department specifically requests
 comments on: (a) whether the proposed
 collection of information is necessary
 for the proper performance of the

functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: January 18, 2005.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 05-1398 Filed 1-25-05; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2005N-0012]

Agency Information Collection Activities; Proposed Collection; Comment Request; Allergen Labeling of Food Products Consumer Preference Survey and Experimental Study on Allergen Labeling of Food Products

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 a voluntary consumer survey entitled "Allergen Labeling of Food Products Consumer Preference Survey" and an

experimental study entitled "Experimental Study on Allergen Labeling of Food Products."

DATES: Submit written or electronic comments on the collection of information by March 28, 2005.

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>. All comments should be identified with the docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT:

Peggy Robbins, Office of Management Programs (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1223.

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.

Allergen Labeling of Food Products Consumer Preference Survey

Under section 903(b)(2) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 393(b)(2)), FDA is authorized to conduct research relating to foods and to conduct educational and public information programs relating to the safety of the Nation's food supply. FDA is planning to conduct a consumer survey about allergen labeling of food products under this authority. The Allergen Labeling of Food Products Consumer Preference Survey will collect information to gauge the impact of certain changes to the food label with respect to information about allergenic ingredients. This data collection is needed to satisfy some of the requirements of the Food Allergen Labeling and Consumer Protection Act (FALCPA) (Public Law 108-282, title II, section 204.4), including the requirement that FDA provide data on consumer preferences in a report to Congress. In particular, section 204.4 of the FALCPA asks FDA to describe in the report " * * * how consumers with food allergies or the caretakers of consumers would prefer that information about the risk of cross-contact be communicated on food labels as determined by using appropriate survey mechanisms." In addition, the survey will address other issues pertinent to allergen labeling changes mandated by the FALCPA. The data will be collected by means of a pool of people who will be screened (through self-report) for food allergy, and food allergy caregiver status. A balanced sample of 1,000 will be selected. Participation in the survey is voluntary.

FDA estimates the burden of the Allergen Labeling of Food Products Consumer Preference Survey collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN¹

Questionnaire	No. of Respondents	Annual Frequency per Response	Total Annual Responses	Hours per Response	Total Hours
Screener	35,000	1	35,000	.0055	193
Pre-test	30	1	30	.167	5