• Ask your doctor, nurse, or health department to file a Vaccine Adverse Event Reporting System (VAERS) form, or call VAERS yourself at 1–800–822–7967.

*6. The National Vaccine Injury Compensation Program*

In the rare event that you or your child has a serious reaction to a vaccine, a Federal program has been created to help you pay for the care of those who have been harmed.

For details about the National Vaccine Injury Compensation program, call 1–800–338–2382 or visit the program's website at http://www.hrsa.dhhs.gov/bhpr/vicp

*7. How Can I Learn More?*

• Ask your doctor or nurse. They can give you the vaccine package insert or suggest other sources of information.

• Call your local or state health department. They can give you the Parents Guide to Childhood Immunization, Immunization of Adults: A Call to Action, or other information.

• Contact the Centers for Disease Control and Prevention (CDC):

—Call 1–800–232–2522 (English)
—Call 1–800–232–0233 (Español)
—Visit the National Immunization Program's website at http://www.cdc.gov/nip

U.S. Department of Health & Human Services
Centers for Disease Control and Prevention
National Immunization Program
MMR (00/00/00) (Proposed)
Vaccine Information Statement
42 U.S.C. 300aa–26

Dated: August 28, 1998.

**Thena M. Durham,**

*Acting Associate Director for Management and Operations, Centers for Disease Control and Prevention (CDC).*

[FR Doc. 98–23736 Filed 9–2–98; 8:45 am]

**BILLING CODE 4163–18–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Children and Families

### Submission for OMB Review; Comment Request

*Title:* Child Care and Development Fund Annual Report on Services Provided (ACF–700).

*OMB No.:* 0980–0241.

*Description:* The Child Care and Development Fund (CCDF) Report is the required annual tribal aggregate information on services provided through the CCDF, which is required per Child Care and Development Block Grant (CCDBG) Final Rule 45 CFR Parts 98 and 99. Tribes are required to submit annual aggregate data appropriate to tribal programs on children and families receiving CCDF-funds or CCDBG funded Child care services. The CCDF regulations require Tribal Lead Agencies to report a supplemental narrative which describes general child care activities and actions in the Tribal Lead Agency's service area and is not limited to the CCDF-funded activities but addresses all child care in the Tribal Lead Agency's service area. This information will be included in the Secretary's report to Congress, as appropriate, and will be shared with all Tribal Lead Agencies to inform them of CCDF or CCDBG-funded activities in other tribal programs.

*Respondents:* Tribal Governments.

| Instrument | Number of respondents | Number of responses per respondent | Average burden hours per response | Total burden hours |
|---|---|---|---|---|
| CCDF Annual Report (ACF–700) | 224 | 1 | 40 | 9,760 |

Estimated Total Annual Burden Hours: 9,760.

*Additional Information:* Copies of the proposed collection may be obtained by writing to The Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, SW, Washington, DC 20447, Attn: ACF: ACF Reports Clearance Officer.

OMB Comment:

OMB is required to make a decision concerning the collection 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 followin Office of Management and Budget, Paperwork Reduction Project, 725 17th Street, NW, Washington, DC 20503, Attn: Ms. Laura Oliven.

Dated: August 28, 1998.

**Bob Sargis,**

*Acting Reports Clearance Officer.*

[FR Doc. 98–23745 Filed 9–2–98; 8:45 am]

**BILLING CODE 4184–01–M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. 98N–0698]**

### Agency Information Collection Activities: Proposed Collection; Comment Request; Survey of Consumer Attitudes Toward Potential Changes in Food Standards of Identity

**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, and to allow 60 days for public comment in response to the notice. This notice solicits comments on a voluntary survey of consumer attitudes toward potential changes in food standards of identity.

**DATES:** Submit written comments on the collection of information by November 2, 1998.

**ADDRESSES:** Submit written comments on the collection of information to the Dockets Management Branch (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:** Margaret R. Schlosburg, Office of Information Resources Management (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, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information listed below.

With respect to the following collection of information, FDA invites comments on: (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.

**Survey of Consumer Attitudes Toward Potential Changes in Food Standards of Identity**

Under section 903(b)(2) of the Federal Food, Drug, and Cosmetic 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 telephone-mail-telephone consumer survey about consumer attitudes towards potential changes in food standards of identity under this authority. A nationally representative sample of 600 adults, who regularly do the food shopping for their households, will be selected at random and asked if they would agree to complete a mail survey. Participation will be voluntary. Detailed information will be obtained about how consumers would be affected by changes to standards, and what their preferences are for retaining, revising, or eliminating standards. FDA is reviewing standard of identity regulations for foods in order to determine which elements of those regulations are most important to fulfilling the goals of those regulations. The information to be collected will address consumer attitudes toward potential changes in the standards of identity for particular products. The products will be chosen to represent general categories of products that share theoretically relevant characteristics. The changes will be chosen to represent general types of changes that might be made to standards of identity. Therefore, the information collected on particular changes in the standards of identity for particular products should provide information that can be generalized to other changes and other products. The information collected will be used to shape FDA's policy on revising standards of identity.

FDA estimates the burden of this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

| | No. of Respondents | Annual Frequency per Response | Total Annual Responses | Hours per Response | Total Hours |
|---|---|---|---|---|---|
| Receive initial recruiting telephone call | 600 | 1 | 600 | 0.8 | 48 |
| Read instructions and complete mail survey | 600 | 1 | 600 | 0.59 | 354 |
| Complete followup telephone interview | 600 | 1 | 600 | 0.08 | 48 |
| Total | | | | | 450 |

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.

The burden hours are based on two rounds of focus groups conducted to test the instrument. The length of the initial and followup interviews based on similar studies are based on similar studies that have been conducted.

Dated: August 26, 1998.

**William K. Hubbard,**

*Associate Commissioner for Policy Coordination.*

[FR Doc. 98–23752 Filed 9–2–98; 8:45 am]

**BILLING CODE 4160–01–F**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Health Resources and Services Administration

### Cooperative Agreement With the National Foundation for the Centers for Disease Control and Prevention

**AGENCY:** Health Resources and Services Administration (HRSA), DHHS.

**ACTION:** Notice of cooperative agreement award.

**SUMMARY:** The Health Resources and Services Administration (HRSA) announces its intent to award funds in fiscal year (FY) 1998 to provide support of a cooperative agreement with the National Foundation for the Centers for Disease Control and Prevention, Inc. (CDC Foundation), Atlanta, Georgia for the implementation of the Public Health Management Training Network.

Multiple reports cite a widening gap in the U.S. between the challenges to improve the health of Americans and the capacity of the public health workforce to meet these challenges. Given the increasing challenges facing our State and local agencies, this cooperative agreement is intended to examine curriculum development and strengthen the public health training and education infrastructure needed to maintain basic agency management competencies needed for the 21st century. Public health leaders and managers, their colleagues in health care organizations, community leaders and others critical to the public's health now work in an environment of change for which most have had little formal preparation. Most have strong technical training in public health, but few have the management skills and resources essential in the new era of Government downsizing, shrinking resources, managed care proliferation and emerging new health risks. This project will be coordinated with the ''Turning Point'' initiative, a collaboration of the Robert Wood Johnson Foundation, the W. K. Kellogg Foundation and the Centers for Disease Control and Prevention (CDC) to address management development. The cooperative agreement award will implement the Public Health Training Network through (1) the design and implementation of a model for improving the management capacity of