• 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