Dated: November 19, 1999.

J. Les Davison,

Acting Deputy Associate Administrator for Acquisition Policy.

[FR Doc. 99–30719 Filed 11–24–99; 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 Project

Title: Child Care and Development Fund Plan for States/Territories.

OMB No.: 0970–0114.

Description: The ACF–118, the Child Care and Development Fund (CCDF) Plan for States and Territories, is required from the child care lead agency by section 658E of the Child Care and Development Block Grant Act of 1990 (Pub. L. 101–508, 42 U.S.C. 9858). The implementing regulations for the statutorily required Plan is at 45 CFR 98.10 through 98.18. The Plan is required biennially and remains in effect for two years. States/Territories have completed the ACF–118 for the FFY 2000–2001 biennium. However, approval for the ACF–118 expires May 31, 2000. States and Territories may amend their Plans to reflect changes in their programs at any time during a biennium. Therefore, in order to provide continuity for the Plan process, ACF is requesting that the current approval of the ACF–118 be extended through the end of the biennium, i.e., September 30, 2001. The Tribal Plan (ACF–118A) is not affected by this notice.

Respondents: State, Local or Tribal Govt.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Child Care & Dev. Fund Plan for States/Terr.	56	.5	162.57	4,552

Estimated Total Annual Burden hours: 4,552.

In compliance with the requirements of section 3506(c)(2)(A) 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. 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 of 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: November 19, 1999.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 99–30765 Filed 11–24–99; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 99N–4933]

Agency Information Collection Activities: Proposed Collection; Comment Request; FDA Safety Alert/ Public Health Advisory Readership Survey

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 information collection, and to allow 60 days for public comment in response to the notice. This notice solicits comments on information collection requirements for FDA Safety Alert/Public Health Advisory Readership Survey.

DATES: Submit written comments on the collection of information by January 25, 2000.

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: Peggy 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, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this