

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request Proposed Projects

Title: Child Support Enforcement Program: State Plan Approval and Grant Procedures, State Plan Requirements, Standards for Program Operations, Federal Financial Participation, and Computerized Support Enforcement Systems.

OMB No.: 0970-0017.

Description: The State plan preprint and amendments serve as a contract with OCSE in outlining the activities the

States will perform as required by law in order for States to receive federal funds to meet the costs of these activities. The affected public is comprised of States receiving funds. Federal regulations require the States to amend their State plans only when necessary to reflect new or revised Federal statutes or regulations or material change in any State law, organization, policy or IV-D agency operations. OMB approved the Form OCSE-100, the IV-D State Plan. As a result of the Child Support Enforcement Amendments of 1988 (P.L. 98-378), the Omnibus Budget Reconciliation Act of 1987 (P.L. 100-203), the Family Support Act of 1988 (P.L. 100-485), the Omnibus Budget Reconciliation Act of 1993 (P.L. 103-66), the Social Security Act

Amendments of 1994 and related regulations, OCSE also received OMB approval for new and revised State plan pages. We are now requesting approval of 34 revised and new State plan preprint pages to reflect changes due to enactment of the Personal Responsibility and Work Opportunity Reconciliation Act of 1996 (PRWORA), P.L. 104-193, and the previous mentioned statutes. The information collected on the State plan pages is necessary to enable OCSE to monitor compliance with the requirements in Title IV-D of the Social Security Act and implementing regulations.

Respondents: States, Guam, Virgin Islands, Puerto Rico and District of Columbia.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
State Plan	54	1836	.717	1,316

Estimated Total Annual Burden Hours: 1,316.

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, Division of Information Resource Management Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 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: January 31, 1997.
Douglas J. Godesky,
Reports Clearance Officer.
[FR Doc. 97-2929 Filed 2-5-97; 8:45 am]
BILLING CODE 4184-01-M

Food and Drug Administration

Advisory Committee; Science Board to the Food and Drug Administration; Formation of a Subcommittee

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing the formation of a subcommittee of the Science Board to the Food and Drug Administration (Science Board). This subcommittee has been established to address issues related to toxicology testing methods. The subcommittee's recommendations will be presented to the Science Board for full public discussion at a future Science Board meeting.

FOR FURTHER INFORMATION CONTACT: Anita M. O'Connor, Office of Science (HF-33), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-3312.

SUPPLEMENTARY INFORMATION: FDA is announcing the formation of a

subcommittee to the Science Board. This subcommittee has been established to address issues related to toxicology testing methods. The subcommittee will meet several times over the next year to develop recommendations for the Science Board on the development and validation of new toxicology test methods. The subcommittee's recommendations will be presented to the Science Board for full public discussion at a future Science Board meeting. Opportunities for public comment will be announced in the Federal Register at least 15 days prior to the Science Board meeting. This notice is issued under the Federal Advisory Committee Act of October 6, 1972 (Pub. L. 92-463 (5 U.S.C. app. 2)).

Dated: January 30, 1997.
Michael A. Friedman,
Deputy Commissioner for Operations.
[FR Doc. 97-2924 Filed 2-5-97; 8:45 am]
BILLING CODE 4160-01-F

Health Care Financing Administration
[Doc. Identifier: HCFA-R-203]

Agency Information Collection Activities: Proposed Collection; Comment Request

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Health Care Financing Administration (HCFA), Department of Health and