

Annual burden estimates instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
OCSE-34 .....	54	4	8	1,728

*Estimated Total Annual Burden Hours: 1,728.*

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, 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: December 4, 1997.

**Bob Sargis,**

*Acting Reports Clearance Officer.*

[FR Doc. 97-32373 Filed 12-10-97; 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(s):*

*Title:* April 1998 Current Population Survey Supplement on Child Support.

*OMB No.:* New Collection.

*Description:* Collection of these data will assist legislators and policymakers in determining how effective their policymaking efforts have been over time in applying the various child support legislation to the overall child support enforcement picture. This information will help policymakers determine to what extent individuals on welfare would be removed from the welfare rolls as a result of more stringent child support enforcement efforts.

*Respondents:* Individuals or Households.

**ANNUAL BURDEN ESTIMATES**

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Supplement .....	47,000	1	.0241	1,136

*Estimated Total Annual Burden Hours: 1,136.*

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: Act Reports Clearance Officer. All requests should be identified by the title for 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: December 4, 1997.

**Bob Sargis,**

*Acting Reports Clearance Officer.*

[FR Doc. 97-32374 Filed 12-10-97; 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:* Revised Form OCSE-100, State Plan for Child Support Collection and Establishment of Paternity under Title IV-D of the Social Security Act.

*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. Due to enactment of HR2105, technical amendments for PRWORA, we are updating our State plan by revising 7 preprint pages. We are requesting

approval of the revised State plan preprint pages for Section 2.1, Establishing Paternity and Securing Support, Section 2.4, Collection and Distribution of Support Payments, Section 2.5, Services to Individuals Not Receiving Title IV-A and IV-E Foster

Care Assistance, Section 2.8, Medical Support Enforcement Activities, Section 2.12-16 State Law Authorizing Suspension of Licenses, Section 2.12-20, Adoption of Uniform State Laws, and Section 3.16, Cooperation by Applicants for and Recipients of Part A

Assistance. 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:* State governments.

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
State Plan .....	54	2.32	.717	90

*Estimated Total Annual Burden Hours:* 90.

*Additional Information:* Copies of the proposed collection may be obtained 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, Robert Driscoll.

*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, 725 17th Street, N.W., Washington, D.C. 20503, Attn: Ms. Wendy Taylor.

**Bob Sargis,**

*Acting Reports Clearance Officer.*

[FR Doc. 97-32372 Filed 12-10-97; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

[Docket No. 97N-0488]

#### Agency Information Collection Activities: Proposed Collection; Comment Request

**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 continuation of national surveys of prescription drug information provided to patients.

**DATES:** Submit written comments on the collection of information by February 9, 1998.

**ADDRESSES:** Submit written comments on the collection of information to the Dockets Management Branch (HFA-305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1-23, Rockville, MD 20857. All comments should be identified with the docket number found in brackets in the heading of this document.

**FOR FURTHER INFORMATION CONTACT:** Karen L. Nelson, Office of Information Resources Management (HFA-250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301-827-1482.

**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 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.

#### Years 1998 and 2000 Continuation of National Surveys of Prescription Drug Information Provided to Patients—(OMB Control Number 0910-0279—Extension)

FDA implements the provisions of the Federal Food, Drug, and Cosmetic Act (the act), designed to assure the adequate labeling of prescription (Rx) drugs. Under section 502(a) of the act (21 U.S.C. 352(a)), a drug product is misbranded if its labeling is false or misleading in any particular, and under section 201(n) of the act (21 U.S.C. 321(n)), a drug's labeling is misleading if its labeling or advertising fails to reveal material facts. FDA also has the authority to collect this information under Title VI of Pub. L. 104-180 (Related Agencies and Food and Drug Administration) section 601 (Effective Medication Guides), which directs the development of "a mechanism to assess periodically \* \* \* the frequency with which the [oral and written prescription] information is provided to consumers."

To assure that Rx drugs are not misbranded, FDA has historically asserted that adequate labeling requires certain information be provided to