the State operates on a July–June fiscal year, or September 1 if the State operates on a Federal fiscal year). No specific format is required for the intended use plan. The intended use of SSBG funds—including the types of activities to be supported and the categories and characteristics of individuals to be served—must be provided. States vary greatly in the information they provide and the structure of the report. States are required to submit a revised intended use plan if the planned use of SSBG funds changes during the year.

In order to provide a more accurate analysis of the extent to which funds are spent "in a manner consistent" with each of the States' plan for their use, as required by 42 USC 1397e(a), we are requesting that States voluntarily use the format of the post-expenditure reporting form to provide estimates of the amount of expenditures and the number of recipients, by service category, that the State plans to use SSBG funds to support as part of the intended use plan. Many States are already using the format of the post-expenditure reporting form as part of their intended use plan.

Respondents:

The post-expenditure reporting form and intended use plan are completed once annually by a representative of the agency that administers the Social Services Block Grant at the State level in each State.

Respondents:

State Governments

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Post-Expenditure Reporting Form	56	1	110	6,160
Use of Post-Expenditure Reporting Form as Part of the Intended Use Plan	56	1	2	112
Estimated Total Annual Burden Hours:				6,272

Additional Information:

Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Administration, 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. *E-mail address: infocollection@acf.hhs.gov*.

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, *Fax:* 202–395–7285, *E-mail: OIRA_SUBMISSION@OMB.EOP.GOV, Attn:* Desk Officer for the Administration for Children and Families.

Dated: January 31, 2011.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2011–2555 Filed 2–7–11; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007–D–0429; Formerly Docket No. 2007D–0496]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Labeling of Nonprescription Human Drug Products Marketed Without an Approved Application as Required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act: Questions and Answers

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled "Labeling of Nonprescription Human Drug Products Marketed Without an Approved Application as Required by the Dietary Supplement and Nonprescription Drug Consumer Protection Act: Questions and Answers" has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: Elizabeth Berbakos, Office of Information Management, Food and Drug Administration, 1350 Piccard Dr., PI50–400B, Rockville, MD 20850, 301–796–3792, *Elizabeth.Berbakos@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In the **Federal Register** of February 24, 2009 (74 FR 8264), the Agency announced that the proposed information collection had been submitted to OMB for review and clearance under 44 U.S.C. 3507. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0640. The approval expires on July 31, 2012. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: February 1, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–2662 Filed 2–7–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2011–N–0049]

Agency Information Collection Activities; Proposed Collection; Comment Request; Presubmission Conferences, New Animal Drug Applications and Supporting Regulations, and Food and Drug Administration Form 356V

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.