DATES: *Comments due within 30 days of publication.* OMB must make a decision about 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.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

SUPPLEMENTARY INFORMATION:

Description: The PI, prepared in response to the enactment of the CBCAP program, as set forth in Title II of the Child Abuse Prevention and Treatment Reauthorization Act of 2010 (Public Law 111–320) or CAPTA, provides direction to the states and territories to accomplish the purposes of (1) supporting community-based efforts to develop, operate, expand, and where appropriate to network, initiatives aimed at the prevention of child abuse and neglect, and to support networks of coordinated resources and activities to better strengthen and support families to reduce the likelihood of child abuse and neglect; and (2) fostering an understanding, appreciation, and knowledge of diverse populations in order to be effective in preventing and treating child abuse and neglect. This PI contains information collection requirements that are found in CAPTA and pursuant to receiving a grant award. The information submitted will be used by the agency to ensure compliance with the statute, complete the calculation of the grant award entitlement, and provide training and technical assistance to the grantee.

Respondents: State governments, quasi-public entities, and non-profit private agencies.

ANNUAL BURDEN ESTIMATES

Instrument	Annual number of respondents	Annual number of responses per respondent	Average burden hours per response	Total annual burden hours
Application	52	1	40	2,080
Annual Report	52	1	24	1,248

Estimated Total Annual Burden Hours: 3,328.

Authority: The CAPTA Reauthorization Act of 2010; Title II of the CAPTA, Pub. L. 115–271 (42 U.S.C. 5116 *et seq.*).

Mary B.Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–05411 Filed 3–15–21; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Submission for OMB Review; Social Services Block Grant (SSBG) Post-Expenditure Report, Pre-Expenditure Report, and Intended Use Plan (OMB #0970–0234)

AGENCY: Office of Community Services, Administration for Children and Families, HHS.

ACTION: Request for Public Comment.

SUMMARY: The Administration for Children and Families (ACF) is requesting a revision to the Social Services Block Grant (SSBG) Post-Expenditure Report, Pre-Expenditure Report, and Intended Use Plan (OMB #0970–0234, previously titled, ''Social Services Block Grant (SSBG) Post-Expenditure Report''). ACF is proposing to expand the information collection to include the collection of states' Intended Use Plans and retitle the information collection to clarify the role of the Pre-Expenditure Report.

DATES: *Comments due within 30 days of publication.* OMB must make a decision about 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.

ADDRESSES: Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain.* Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.

SUPPLEMENTARY INFORMATION:

Description: On an annual basis, states and territories are required to submit the following reports: (1) An Intended Use Plan that provides data and narrative descriptions related to the state's SSBG program. The Intended Use Plan includes details about the delivery of SSBG services, and the state agency administering the SBG Program. ACF is proposing to expand the currently approved information collection to include collection of states' Intended Use Plans. Grantees are required to submit their Pre-Expenditure Report no less than 30 days prior to the start of the period covered by the report. (2) A Pre-Expenditure Report that demonstrates the state's anticipated allocation of SSBG funding among the 29 pre-defined SSBG service categories. Historically, states have submitted this report using the Post-Expenditure Report Form, and the associated burden is included in the currently approved information collection. Grantees are required to submit their Intended Use Plan no less than 30 days prior to the start of the period covered by the report. (3) A Post-Expenditure Report that details the state's actual use of SSBG funding among each of the 29 service categories. Grantees are required to submit their Post-Expenditure Report within 6 months of the end of the period covered by the report.

Respondents: Agencies that administer the SSBG at the state or territory level, including the 50 states; District of Columbia; Puerto Rico; and the territories of American Samoa, Guam, the Virgin Islands, and the Commonwealth of Northern Mariana Islands.

Annual Burden Estimates: This request is specific to the Intended Use Plan. Currently approved materials and associated burden can be found at: *https://www.reginfo.gov/public/do/PRAViewICR?ref_nbr=202011-0970-006.*

Instrument	Annual number of respondents	Annual number of responses per respondent	Average burden hours per response	Total/annual burden hours
Intended Use Plan	56	1	40	2,240.

Estimated Total Annual Burden Hours: 2,240.

Authority: 42 U.S.C. 1397 through 1397e.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2021–05408 Filed 3–15–21; 8:45 am]

BILLING CODE 4184–24–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2012–N–0547]

Agency Information Collection Activities; Proposed Collection; Comment Request; Survey on the Occurrence of Foodborne Illness Risk Factors in Selected Retail and Foodservice Facility Types

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing an opportunity for public comment on the proposed collection of certain information by the Agency. Under the Paperwork Reduction Act of 1995 (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 ''Survey on the Occurrence of Foodborne Illness Risk Factors in Selected Retail and Foodservice Facility Types.''

DATES: Submit either electronic or written comments on the collection of information by May 17, 2021.

ADDRESSES: You may submit comments as follows. Please note that late, untimely filed comments will not be considered. Electronic comments must be submitted on or before May 17, 2021. The *https://www.regulations.gov* electronic filing system will accept comments until 11:59 p.m. Eastern Time at the end of May 17, 2021. Comments received by mail/hand delivery/courier (for written/paper submissions) will be considered timely if they are postmarked or the delivery service acceptance receipt is on or before that date.

Electronic Submissions

Submit electronic comments in the following way:

- *Federal eRulemaking Portal: https://www.regulations.gov.* Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to *https://www.regulations.gov* will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on *https://www.regulations.gov.*
- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see ''Written/Paper Submissions'' and ''Instructions'').

Written/Paper Submissions

Submit written/paper submissions as follows:

- *Mail/Hand Delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in ''Instructions.''

Instructions: All submissions received must include the Docket No. FDA–2012–N–0547 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Survey on the Occurrence of Foodborne Illness Risk Factors in Selected Retail and Foodservice Facility Types.'' Received comments, those filed in a timely manner (see **ADDRESSES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Dockets Management Staff between 9 a.m. and 4 p.m., Monday through Friday, 240–402–7500.

- Confidential Submissions—To submit a comment with confidential information that you do not wish to be made publicly available, submit your comments only as a written/paper submission. You should submit two copies total. One copy will include the information you claim to be confidential with a heading or cover note that states ''THIS DOCUMENT CONTAINS CONFIDENTIAL INFORMATION.'' The Agency will review this copy, including the claimed confidential information, in its consideration of comments. The second copy, which will have the claimed confidential information redacted/blacked out, will be available for public viewing and posted on *https://www.regulations.gov.* Submit both copies to the Dockets Management Staff. If you do not wish your name and contact information to be made publicly available, you can provide this information on the cover sheet and not in the body of your comments and you must identify this information as ''confidential.'' Any information marked as ''confidential'' will not be disclosed except in accordance with 21 CFR 10.20 and other applicable disclosure law. For more information about FDA's posting of comments to public dockets, see 80 FR 56469, September 18, 2015, or access the information at: *https://www.govinfo.gov/content/pkg/FR-2015-09-18/pdf/2015-23389.pdf.*

Docket: For access to the docket to read background documents or the electronic and written/paper comments received, go to *https://www.regulations.gov* and insert the docket number, found in brackets in the heading of this document, into the ''Search'' box and follow the prompts and/or go to the Dockets Management Staff, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240–402–7500.

FOR FURTHER INFORMATION CONTACT: Ila S. Mizrachi, Office of Operations, Food and Drug Administration, Three White Flint North, 10A–12M, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–7726, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Under the PRA (44 U.S.C. 3501–3521), Federal