Agency Contact: For further information or comments regarding this supplemental action, contact Cynthia LaCounte, U.S. Department of Health and Human Services, Administration for Community Living, Administration on Aging, 330 C Street SW., Washington DC 20201; telephone 202–795–7380; email *Cynthia.LaCounte@acl.hhs.gov*.

Dated: June 13, 2017.

Daniel P. Berger,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2017–12753 Filed 6–19–17; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Public Comment Request; Revision of a Currently Approved Information Collection (ICR-Rev); Title III Supplemental Form to the Financial Status Report (SF–425)

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing an opportunity for the public to 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 a 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 proposed action. This notice solicits comments on a proposed revision of an existing data collection regarding the information collection requirements relating to the Title III Supplemental Form to the Financial Status Report for all ACL/AoA Title III Grantees.

DATES: Submit written or electronic comments on the collection of information by August 21, 2017.

ADDRESSES: Submit electronic comments on the collection of information to: *jesse.more@acl.hhs.gov*. Submit written comments on the collection of information to the U.S. Department of Health and Human Services, Administration for Community Living, Washington, DC 20201, Attention: Jesse E. Moore, Jr.

FOR FURTHER INFORMATION CONTACT: Jesse E. Moore, Jr., Aging Services Program Specialist, Administration for Community Living, Washington, DC, 20201, 202–795–7578.

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 or update of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, ACL is publishing the notice of the proposed collection of information set forth in this document. With respect to the following collection of information, ACL invites comments on: (1) Whether the proposed collection of information is necessary for the proper performance of ACL's functions, including whether the information will have practical utility; (2) the accuracy of ACL'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.

Purpose

The *Title III Supplemental Form to the Financial Status Report* (SF–425) is used by ACL/AoA for all grantees to obtain a more detailed understanding of how projects funded under Title III of the Older Americans Act (OAA) of 1965, as amended, are being administered, and to ensure compliance with legislative requirements, pertinent Federal regulations and other applicable instructions and guidelines issued by the ACL. The level of data detail necessary is not available through the Federal Financial Status Report (SF–425) form. The Title III Supplemental Form provides necessary details on non-federal required match, administration expenditures, Older Relative Caregivers expenditures, and Long Term Care Ombudsman expenditures.

In addition to renewing OMB approval of this data collection, minor changes are being proposed to it to reflect changes in statutory language that occurred as a result of the 2016 reauthorization of the OAA. Specifically, the term ''Grandparents Only'' has been changed to ''Older Relative Caregivers,'' the new term in the OAA that describes this population of eligible service recipients. Similarly, the accompanying instructions for completing the Title III Supplemental Form to the Financial Status Report were also modified to include this same language. References in the Code of Federal Regulation (CFR) have been updated addressing financial reporting requirements and non-substantive technical edits have been made to the instructions.

Data Burden

ACL estimates the burden of this collection of information as follows: 56 State Units on Aging (SUA) respond semi-annually which should have an average estimated burden of 2 hours per grantee for a total of 112 hours per submission.

The proposed data collection tool may be found on the ACL Web site for review at: *https://www.acl.gov/sites/default/files/about-acl/2017-06/ACL%20Title%20III%20Supplemental%20Form%20and%20Instructions%202017.pdf*.

Respondent/data collection activity	Number of respondents	Responses per respondent	Hours per response	Total annual burden hours
Title III Supplemental Form to the Financial Status Report	56	2/yr	2	224
Total	56	2/yr	2	224

Dated: June 13, 2017.

Daniel P. Berger,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2017–12755 Filed 6–19–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2013–N–1119]

Agency Information Collection Activities; Proposed Collection; Comment Request; Food Canning Establishment Registration, Process Filing, and Recordkeeping for Acidified Foods and Thermally Processed Low-Acid Foods in Hermetically Sealed Containers

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 the information collection provisions of reporting and recordkeeping requirements for firms that process acidified foods and thermally processed low-acid foods in hermetically sealed containers.

DATES: Submit either electronic or written comments on the collection of information by August 21, 2017.

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 August 21, 2017. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of August 21, 2017. 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–2013–N–1119 for ''Agency Information Collection Activities; Proposed Collection; Comment Request; Food Canning Establishment Registration, Process Filing, and Recordkeeping for Acidified Foods and Thermally Processed Low-Acid Foods in Hermetically Sealed Containers.'' 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.

- 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.fda.gov/regulatoryinformation/dockets/default.htm.*

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.

FOR FURTHER INFORMATION CONTACT: JonnaLynn Capezzuto, Office of Operations, Food and Drug Administration, Three White Flint North, 10A63, 11601 Landsdown St., North Bethesda, MD 20852, *301–796–3794,* PRAStaff@*fda.hhs.gov.*

SUPPLEMENTARY INFORMATION:

I. Background

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,