ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
LIHEAP Leveraging Report	70	1	38	2,660

Estimated Total Annual Burden Hours: 2,660.

Additional Information: Copies of the proposed collection may be obtained by writing to the Administration for Children and Families, Office of Planning, Research and Evaluation, 330 C Street SW., Washington, DC 20201. Attention Reports Clearance Officer. All requests should be identified by the title of the information collection. Email 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, Email: *OIRA_SUBMISSION@OMB.EOP.GOV,* Attn: Desk Officer for the Administration for Children and Families.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2017–04882 Filed 3–10–17; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Public Comment Request; Proposed Extension With No Changes of a Currently Approved Collection; National Survey of Older Americans Act Participants

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 notice. This notice solicits comments on the information collection requirements relating to consumer assessment surveys that are used by ACL to measure program performance for programs funded under Title III of the Older Americans Act. This notice solicits comments on a proposed extension with no changes of a currently approved collection.

DATES: Submit written or electronic comments on the collection of information by May 12, 2017.

ADDRESSES: Submit electronic comments on the collection of information to: *Heather.Menne@acl.hhs.gov.* Submit written comments on the collection of information to: Heather Menne, U.S. Department of Health and Human Services: Administration for Community Living, Washington, DC 20201.

FOR FURTHER INFORMATION CONTACT: Heather Menne at 202–795–7733 or *Heather.Menne@acl.hhs.gov.*

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 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.

The National Survey of Older Americans Act (OAA) Participants information collection will include consumer assessment surveys for the Congregate and Home-delivered meal nutrition programs; Case Management, Homemaker, and Transportation Services; and the National Family Caregiver Support Program. This survey builds on earlier national pilot studies and surveys, as well as performance measurement tools developed by ACL grantees in the Performance Outcomes Measures Project (POMP). Changes identified as a result of these initiatives were incorporated into the last data collection package that was approved by OMB and are included in this proposed extension of a currently approved collection. This information will be used by ACL to track performance outcome measures; support budget requests; comply with the GPRA Modernization Act of 2010 (GPRAMA) reporting requirements; provide national benchmark information; and inform program development and management initiatives.

Descriptions of previous National Surveys of OAA Participants can be found under the section on OAA Performance Information on ACL's Web site at: *https://aoa.acl.gov/Program_Results/OAA_Performance.aspx.* Copies of the survey instruments and data from previous National Surveys of OAA Participants can be found and queried using the AGing Integrated Database (AGID) at: *http://www.agid.acl.gov/.* The proposed National Survey entitled National Survey of Older Americans Act Participants 2017 Draft may be found on the ACL Web site at: *https://aoa.acl.gov/Program_Results/OAA_Performance.aspx.*

ACL estimates the burden of this collection of information as follows:

Respondent/data collection activity	Number of respondents	Responses per respondent	Hours per response	Annual burden hours
Area Agency on Aging: Respondent selection process	250	1	4.0	1,000
Service Recipients (*i.e.,* Congregate and Home-delivered meal nutrition programs; Case Management, Homemaker, and Transportation Services).	4,000	1	.6667	2,666.80
National Family Caregiver Support Program Clients	2,000	1	.6667	1,333.40
Total	6,250	1	.80 (weighted mean)	5,000

Daniel Berger,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2017–04843 Filed 3–10–17; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2016–N–1112]

United States Food and Drug Administration and Health Canada Joint Regional Consultation on International Council for Harmonisation of Technical Requirements for Pharmaceuticals for Human Use; Public Meeting and Webcast; Request for Comments

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public meeting; request for comments.

SUMMARY: The Food and Drug Administration (FDA or the Agency) is announcing a regional public meeting entitled ''U.S. Food and Drug Administration and Health Canada Joint Regional Consultation on International Council for Harmonisation of Technical Requirements for Pharmaceuticals for Human Use (ICH).'' The goal of this meeting is to provide information and receive comments on the current activities of ICH, as well as the upcoming ICH meetings in Montreal. The topics to be covered in the public meeting are the topics for discussion at the forthcoming ICH Assembly Meeting in Montreal. The purpose of this public meeting is also to solicit public input prior to the ICH Assembly meeting and the Expert Working Group meetings in Montreal, Canada, scheduled for May 28 through June 1, 2017.

DATES: The public meeting will be held on April 24, 2017, from 11 a.m. to 2 p.m., Eastern Time. Submit either electronic or written comments on this public meeting by May 12, 2017. Late, untimely filed comments will not be considered. Electronic comments must be submitted on or before May 12, 2017. The *https://www.regulations.gov* electronic filing system will accept comments until midnight Eastern Time at the end of May 12, 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. See the **SUPPLEMENTARY INFORMATION** section for registration date and information. Registration to attend the meeting and requests for oral presentations must be received by April 19, 2017; see the **SUPPLEMENTARY INFORMATION** section for information on how to register for the meeting.

ADDRESSES: The public meeting will be held at 10903 New Hampshire Ave., Bldg. 31, Rm. 1503 Section A, Silver Spring, MD 20993. It will also be broadcast on the Web allowing participants to join in person OR via the Web. Entrance for the public meeting participants (non-FDA employees) is through Building 1 where routine security check procedures will be performed. For parking and security information, please refer to *http://www.fda.gov/AboutFDA/WorkingatFDA/BuildingsandFacilities/WhiteOakCampusInformation/ucm241740.htm.*

You may submit comments as follows:

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):* Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.
- For written/paper comments submitted to the Division of Dockets Management, 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–2016–N–1112 for the U.S. Food and Drug Administration and Health Canada Joint Public Consultation on International Council on Harmonisation of Technical Requirements for Pharmaceuticals for Human Use; Public Meeting. Received comments, those filed in a timely manner (see **DATES**), will be placed in the docket and, except for those submitted as ''Confidential Submissions,'' publicly viewable at *https://www.regulations.gov* or at the Division of Dockets Management 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