inclusion of refugee resettlement programs in pandemic influenza emergency plans and a basic description of providers conducting medical screening. ORR proposes to remove a requirement that all states describe a plan for the care, supervision of, and legal responsibility for, refugee children who become unaccompanied in the state. ORR also proposes to remove requirements specific to the Cuban/Haitian entrants and replace them with an assurance that states will provide all ORR-eligible populations with the benefits and services described in the State Plan.

ORR proposes adding language to clarify the following requirements related to the Unaccompanied Refugee Minors (URM) program: State policy on education and training vouchers, medical coverage, the location of URM providers, monitoring procedures, the process for establishing legal responsibility, and information about sub-contractors.

States must use a pre-print format for required components of State Plans for ORR- funded refugee resettlement services and benefits prepared by the Office of Refugee Resettlement (ORR) of the Administration for Children and Families (ACF).

States must submit by August 15 each year new or amended State Plan for the next Federal fiscal year. For previously approved plan, States must certify no later than October 31 each year that the approved State plan is current and continues in effect.

Respondents: State Agencies, Replacement Designees under 45 CFR 400.301(c), and Wilson-Fish Grantees (State 2 Agencies) administering or supervising the administration of programs under Title IV of the Act.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Title IV State Plan	55	1	15	750

Estimated Total Annual Burden Hours: 750.

In compliance with the requirements of the Paperwork Reduction Act of 1995 (Pub. L. 104–13, 44 U.S.C. Chap 35), 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 Planning, Research and Evaluation, 330 C Street SW., Washington DC 20201. Attn: ACF Reports Clearance Officer. Email address: *infocollection@acf.hhs.gov.* 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.

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2017–13033 Filed 6–21–17; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Comment Request; Revision of a Currently Approved Collection (ICR Rev); National Survey of Older Americans Act Participants (NSOAAP)

AGENCY: Administration for Community Living, HHS.

ACTION: Notice.

SUMMARY: The Administration for Community Living (ACL) is announcing that the proposed collection of information listed below has been submitted to the Office of Management and Budget (OMB) for review and clearance as required under the Paperwork Reduction Act of 1995 (the PRA). This 30-Day notice collects comments on the information collection requirements related to a Revision of a Currently Approved Collection (ICR Rev) (OMB approval number 0985–0023).

DATES: Submit written comments on the collection of information by July 24, 2017.

ADDRESSES: Submit written comments on the collection of information by fax 202–395–5806 or by email to *OIRA_submission@omb.eop.gov,* Attn: OMB Desk Officer for ACL.

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

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, ACL has submitted the following proposed collection of information to OMB for review and clearance. ACL is requesting approval for three years of an extension of the currently approved data collection with modifications.

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 with modifications 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.

Comments in Response to the 60 Day Federal Register Notice

A 60-Day notice was published in the **Federal Register** in Vol. 82, No. 13457 on March 13, 2017. A Notice of Correction was published in the **Federal Register** in Vol. 82, No. 15062 on March 24, 2017, announcing that ACL was requesting approval of a proposed extension with modifications of a currently approved data collection. A second Notice of Correction was published in the **Federal Register** in Vol. 82, No. 20896 on May 4, 2017, announcing that the web location of the proposed information collection would change due to an update of the *ACL.gov* Web site.

ACL received comments from eighty-nine (89) organizations and just over 13,900 individuals about the National Survey of Older Americans Act Participants (NSOAAP). ACL reviewed all of the comments. Eight (8) of the comments were deemed not relevant because they were: (a) Programmatic in nature and not survey-related; (b) referencing other data collections and not the NSOAAP (*e.g.,* Census); or (c) commentary without reference to the NSOAAP. The majority of the comments that ACL received expressed the need to retain demographic questions on sexual orientation/gender identity. In addition, comments addressed: (a) Methodological, survey design, and sampling considerations; (b) concern about the survey length; and (c) recommendations to modify and/or add clarifying questions throughout the survey. ACL has made minor changes to the survey based on some suggested changes, including retaining the primary question regarding sexual orientation. This survey has remained essentially the same since the last OMB approval on 7/17/2014 (OMB Control Number 0985–0023), and the sampling methodology and the data collection procedures are identical to the previous survey approved in 2014.

Burden Estimates

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://www.acl.gov/programs/oaa-performance-information.* 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 *https://agid.acl.gov/.* The proposed National Survey entitled National Survey of Older Americans Act Participants 2017 Revised may be found on the ACL Web site at: *https://www.acl.gov/about-acl/public-input.* The revisions, including the reinstatement of the primary question on sexual orientation, represent minor changes in terms of data collection burden that do not change the overall estimated burden on respondents.

ANNUAL BURDEN ESTIMATES

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

Estimated Total Annual Burden Hours: 5,000.

Dated: June 16, 2017.

Daniel P. Berger,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2017–13030 Filed 6–21–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0086]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Potential Tobacco Product Violations Reporting Form

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by July 24, 2017.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–7285, or emailed to *oira_submission@omb.eop.gov.* All comments should be identified with the OMB control number 0910–0716. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Amber Sanford, Office of Operations, Food and Drug Administration, Three White Flint North, 10A63, 11601 Landsdown St., North Bethesda, MD 20852, 301–796–8867, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Potential Tobacco Product Violations Reporting Form OMB Control Number 0910–0716—Extension

On June 22, 2009, the President signed the Family Smoking Prevention and Tobacco Control Act (the Tobacco Control Act) (Pub. L. 111–31) into law. The Tobacco Control Act amended section 201 *et seq.* of the Federal Food, Drug, and Cosmetic Act (the FD&C Act) (21 U.S.C. 321 *et seq.*) by adding a new chapter granting FDA important new authority to regulate the manufacture, marketing, and distribution of tobacco products to protect the public health generally and to reduce tobacco use by minors. FDA is requesting an extension of OMB approval for the collection of information to accept consumer and