under the Paperwork Reduction Act (PRA) would inhibit the ability to collect information to inform these activities. Therefore, an umbrella generic is requested to allow for quick turnaround requests for similar information collections related to these activities.

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. You can also obtain copies of the proposed collection of information by emailing *OPREinfocollection@acf.hhs.gov.* Identify all requests by the title of the information collection.

SUPPLEMENTARY INFORMATION:

Description: ACF hosts a variety of gatherings for many different purposes. This may include large scale conferences, meetings for grantees or contractors, workshops, trainings, poster sessions, and other in-person and virtual gatherings for individuals with interest in ACF programs (clients, researchers, policymakers, etc.), among others. To ensure ACF has adequate information to plan these activities, the Agency must often collect information from potential participants such as basic contact information, preferences for attendance (mode, special requests, etc.), organizational affiliation, feedback about meeting content, etc. Additionally, some activities require ACF to have additional information to have the means to select the most appropriate participants for attendance according to the type or purpose of a given activity, or to group participants into the most appropriate category or activity during an event. This may include information about poster presentations, speaking panels, training courses, professional perspectives, or experiences, etc. In addition, attendees may be asked to submit an application or abstract for prescreening to be selected for attendance.

The purposes of the collections under this umbrella generic information collection are to gather appropriate information to plan ACF gatherings. Example information collection activities could include:

- Registration forms
  - Information collected on these types of forms could include name, contact information, organization/affiliation, basic demographics, attendance needs, etc.
- Applications for panels, posters, or other presentation formats
  - Information collected on these types of applications could include title, author(s), institution/organization, abstract describing presentation or poster, instructions, etc.
- Pre-meeting surveys
  - Information collected on these types of surveys could include content preferences, scheduling needs and preferences, pre-meeting knowledge, etc.
- Post-Meeting/-Workshop/-Training Evaluation Surveys
  - Information collected on these types of surveys could include requests for feedback on the overall activity, feedback on content, post-meeting knowledge, post-meeting uses of content, preferences for future activities, etc.

As part of this generic, ACF requests OMB provide a response on individual generic information collections within 5 business days.

Note that this generic is primarily for information collected in connection with closed ACF meetings, as information collected in connection with public ACF meetings are not considered ''information'' under PRA per 44 U.S.C., 5 CFR ch. 11 (1–1–99 edition), 1320.3: Definitions.

Respondents: Potential respondents may include researchers, individuals with expertise in ACF program areas, individuals with interest in ACF program areas, those receiving ACF services, ACF grantees or contractors, among others with involvement or interest in ACF activities.

TOTAL BURDEN ESTIMATES

Example types of information collections	Total number of respondents	Total number of responses per respondent	Average burden hours per response	Total burden hours	Average hourly wage	Total annual cost
Registration Forms	30,000	1	.167	5,010	$64	$320,640
Applications	5,000	1	1.5	7,500	64	480,000
Pre- and Post-activity Surveys	20,000	1	.5	10,000	64	640,000
Other Activities	14,000	1	.5	7,000	64	448,000
Estimated Totals	69,000		*.428	29,510		1,888,640

* Average.

Mary B. Jones,

ACF/OPRE Certifying Officer.

[FR Doc. 2023–16436 Filed 8–1–23; 8:45 am]

BILLING CODE 4184–79–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Proposed Collection; Comment Request; National Survey of Older Americans Act Participants [OMB 0985–0023]

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 information listed above. 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 Proposed Extension without Change and the information collection requirements related to the National Survey of Older Americans Act Participants.

DATES: Comments on the collection of information must be submitted electronically by 11:59 p.m. (EST) or postmarked by October 2, 2023.

ADDRESSES: Submit electronic comments on the collection of information to *Kristen.Robinson@acl.hhs.gov.* Submit written comments on the collection of information to Administration for Community Living, Washington, DC 20201, Attention: Kristen Robinson.

FOR FURTHER INFORMATION CONTACT: Kristen Robinson, Administration for Community Living, Washington, DC 20201, by email at *Kristen.Robinson@acl.hhs.gov,* or by telephone at 202–795–7428.

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 of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, ACL is publishing a notice of the proposed collection of information set forth in this document. With respect to the following collection of information, ACL invites comments on our burden estimates or any other aspect of this collection of information, including:

(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 to determine burden estimates;

(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 using automated collection techniques when appropriate, and other forms of information technology. The National Survey of Older Americans Act (OAA) Participants information collection includes 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 (GPRMA) reporting requirements; provide national benchmark information; and inform program development and management initiatives.

Copies of the survey instruments and data from previous National Surveys of OAA Participants can be found and queried using the Aging, Independence, and Disability (AGID) Program Data Portal at *https://agid.acl.gov/.* The proposed data collection tools may be found on the ACL website at *https://www.acl.gov/about-acl/public-input.*

Estimated Program Burden: ACL estimates the annual burden associated with 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	300	1	4.0	1,200
Service recipients (i.e., Congregate and Home-Delivered Meal nutrition programs, Case Management, Homemaker, Transportation services) + Rotating Module	4,000	1	0.75	3,000
National Family Caregiver Support Program clients + Rotating Module	2,000	1	0.75	1,500
Total	6,300	1	*0.90	5,700

*(weighted mean).

Dated: July 28, 2023.

Alison Barkoff,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2023–16419 Filed 8–1–23; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2004–N–0451]

Food and Drug Administration Modernization Act of 1997: Modifications to the List of Recognized Standards, Recognition List Number: 059

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA or Agency) is announcing a publication containing modifications the Agency is making to the list of standards FDA recognizes for use in premarket reviews (FDA Recognized Consensus Standards). This publication, entitled ''Modifications to the List of Recognized Standards, Recognition List Number: 059'' (Recognition List Number: 059), will assist manufacturers who elect to declare conformity with consensus standards to meet certain requirements for medical devices.