

**ADDRESSES:** Submit written comments on the collection of information by:

(a) Email to: [OIRA\\_submission@omb.eop.gov](mailto:OIRA_submission@omb.eop.gov), Attn: OMB Desk Officer for ACL;

(b) fax to 202.395.5806, Attn: OMB Desk Officer for ACL; or

(c) by mail to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW, Rm. 10235, Washington, DC 20503, Attn: OMB Desk Officer for ACL.

**FOR FURTHER INFORMATION CONTACT:**

Leslie Green, Administration for Community Living, [leslie.green@acl.hhs.gov](mailto:leslie.green@acl.hhs.gov), 202-868-9384.

**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 responsible for administering the Title VI Program Performance Report. The purpose of this data collection is to fulfill the annual programmatic reporting required by the Title VI Part A/B and C grants to American Indians, Alaskan Native and Native Hawaiian Programs to provide nutrition, supportive services and caregiver services to elders and their caregivers.

**Comments in Response to the 60-Day Federal Register Notice**

ACL published a 60-day **Federal Register** Notice in the **Federal Register** soliciting public comments on this

reinstatement without change request. The 60-day FRN published on July 15, 2020 Volume 85, Number 136, pages 42857-42858; ACL did not receive any public comments during the 60-day FRN.

The proposed data collection tools may be found on the ACL website for review at <https://www.acl.gov/about-acl/public-input>.

**Estimated Program Burden**

There are 282 respondents taking 3.49 hours each to complete the response.

ACL estimates the 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
Title VI PPR .....	282	1	3.49	984
Total: .....	.....	.....	.....	984

Dated: October 15, 2020.

**Mary Lazare,**

*Principal Deputy Administrator.*

[FR Doc. 2020-23471 Filed 10-22-20; 8:45 am]

**BILLING CODE 4154-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Administration for Community Living**

**Agency Information Collection Activities; Proposed Collection; Comment Request; State Health Insurance Assistance Program Annual Sub-Recipients Report; OMB #0985-New**

**AGENCY:** Administration for Community Living, HHS.

**ACTION:** Notice.

**SUMMARY:** The Administration for Community Living is announcing that the proposed collection of information listed above has been submitted to the Office of Management and Budget (OMB) for review and clearance as required under section 506(c)(2)(A) of the Paperwork Reduction Act of 1995. This 30-Day notice collects comments on the information collection requirements related to the proposed information collection requirements related to the State Health Insurance Assistance Program Annual Sub-Recipients Report.

**DATES:** Submit written comments on the collection of information by November 23, 2020.

**ADDRESSES:** Submit written comments and recommendations for the proposed information collection within 30 days of publication of this notice to [www.reginfo.gov/public/do/PRAMain](http://www.reginfo.gov/public/do/PRAMain). Find the information collection by selecting “Currently under 30-day Review—Open for Public Comments” or by using the search function. By mail to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW, Rm. 10235, Washington, DC 20503, Attn: OMB Desk Officer for ACL.

**FOR FURTHER INFORMATION CONTACT:** Margaret Flowers, Administration for Community Living, Washington, DC 20201, 202-795-7315, [Margaret.Flowers@acl.hhs.gov](mailto:Margaret.Flowers@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. The Administration for Community Living (ACL) is requesting approval for a revised data collection associated with the State Health Insurance Assistance Program Annual Sub-Recipients Report. The purpose of this data collection is to collect sub-award data from grantees, including agency name, address, and annual federal funds received. Congress requires this data collection for program monitoring for the State Health Insurance Assistance Program (SHIP) under the Bipartisan Budget Act of 2018, SEC. 50207(b). This data collection allows the Administration for

Community Living (ACL) and the Center for Innovation and Partnership (CIP) to communicate with Congress and the public on the SHIP network of agencies. This is a new data collection requiring State SHIP grantees to provide the amount of federal funds provided annually to each sub-contractor and sub-grantee that are delivering SHIP services. The data collected will be will be electronically posted on the ACL website to educate the network on who the SHIP state sub-recipients are and how much money they are receiving. SHIP grantees are located in each of the 50 states, the District of Columbia, Puerto Rico, Guam and the U.S. Virgin Islands. The respondents for this data collection are grantees who meet with Medicare beneficiaries and older adults’ in-group settings and in one-on-one sessions to educate them on Medicare.

**Comments in Response to the 60-Day Federal Register Notice**

A notice published in the **Federal Register** on July 19, 2020 in FR 85 No. 148 pages 46123-46124. There were no public comments received during the 60-day FRN comment period.

To review the proposed data collection please visit the ACL website at <https://www.acl.gov/about-acl/public-input>.

**Estimated Program Burden**

ACL estimates the 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
	54	1	1	54
Total .....	54	1	1	54

Dated: October 15, 2020.

**Mary Lazare,**

*Principal Deputy Administrator.*

[FR Doc. 2020–23468 Filed 10–22–20; 8:45 am]

**BILLING CODE 4154–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Administration for Community Living

#### Agency Information Collection Activities; Submission for OMB Review; Alzheimer's and Dementia Program Data Reporting Tool (ADP–DRT); OMB #0985–0022

**AGENCY:** Administration for Community Living, HHS.

**ACTION:** Notice.

**SUMMARY:** The Administration for Community Living is announcing that the proposed collection of information listed above has been submitted to the Office of Management and Budget (OMB) for review and clearance as required under section 506(c)(2)(A) of the Paperwork Reduction Act of 1995. This 30-Day notice collects comments on the information collection requirements related to the Information Collection tools for information collection requirements related to Alzheimer's and Dementia Program Data Reporting Tool (ADP–DRT).

**DATES:** Submit written comments on the collection of information by 11:59 p.m. (EST) or postmarked by November 23, 2020.

**ADDRESSES:** Submit written comments on the collection of information by:

(a) Email to: *OIRA\_submission@omb.eop.gov*, Attn: OMB Desk Officer for ACL;

(b) fax to 202.395.5806, Attn: OMB Desk Officer for ACL; or

(c) by mail to the Office of Information and Regulatory Affairs, OMB, New Executive Office Bldg., 725 17th St. NW, Rm. 10235, Washington, DC 20503, Attn: OMB Desk Officer for ACL.

**FOR FURTHER INFORMATION CONTACT:** Erin Long. Submit written comments on the collection of information to Administration for Community Living, Washington, DC 20201 Attention: Erin

Long Phone: 202–795–7389 *Erin.Long@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. The Older American's Act requires ACL to evaluate “demonstration projects that support the objectives of this Act, including activities to bring effective demonstration projects to scale with a prioritization of projects that address the needs of underserved populations, and promote partnerships among aging services, community-based organizations, and Medicare and Medicaid providers, plans, and health (including public health) systems. (Section 201 (42 U.S.C. 3011) Sec. 127. Research and Evaluation). To fulfill the evaluation requirements and allow for optimal federal and state-level management of ACL's Alzheimer's Disease Program, specific information must be collected from grantees.

The current reporting tool is set to expire December 30, 2020. The Alzheimer's and Dementia Program (ADP) Project Officer has reviewed the current data collection procedures to ensure the acceptability of these items as appropriate and thorough evaluation of the program, while minimizing burden for grantees. The result of this process is the proposed modifications to the existing data collection tool. ACL is aware that different grantees have different data collection capabilities. Following the approval of the modified data collection tool, ACL will work with its grantees to offer regular training to ensure minimal burden.

#### Comments in Response to the 60-Day Federal Register Notice

ACL published both a 60-day and 30-day **Federal Register** Notice in the **Federal Register** soliciting public comments on this revision request. The 60-day FRN published on July 20, 2020 in volume 85 No. 137 pages 43241–43242. ACL received comments from one individual.

*Comments on Proposed Collection:* Alzheimer's and Dementia Program Data Reporting Tool (ADP–DRT) OMB #0985–0022.

#### General

It would be helpful if the explanation of categories and definitions for all data elements were part of this information collection (*i.e.*, PRA process). It is difficult to comment on estimated burden and utility of the information collection when the information being collected hasn't been fully explained. Also, definitions and data elements should be synchronized or crosswalked to those in the American Community Survey or another national collection to facilitate analyses across data collections.

#### PLWD & CG Served

**CG data points**—It is important to get a more fulsome profile of the caregivers to assess the impact caregiving has on their lives, their families, and those they care for. Understanding this data collection may not be for this purpose, a few extra data points could shed help expand the CG profile: employment status, # of chronic diseases, # of people cared for, # recent traumas experienced (*e.g.*, emotional, physical, etc.), etc.

There are sections on race and ethnicity. It's not clear what is meant by “Minority Status” or why it's needed. This section should be deleted to reduce burden.

**Living arrangement**—This section describes who the PLWD lives with but doesn't identify where the person is living. It would be helpful to know whether these individuals are living in a private home setting, an institutional setting such as a nursing home, supportive housing, or if they are experiencing homelessness. It would also be helpful to know where they are receiving most of their care—*i.e.*, in the home or outside of the home. Where people are receiving their care is relevant to the workforce and services needed to support them.

#### Professionals Trained

The note at the bottom states that “Persons trained should not include . . . Caregivers . . .” but there are caregivers who are trained and licensed and some family caregivers who receive stipends from Medicaid and other programs. It's not clear if they would be excluded. Also, in the middle of the sheet there's a section on “Total Units of Direct Service Delivered.” How does