projects funded through the CED program. The legislative requirement for this program is in Title IV of the Community Opportunities, Accountability and Training and Educational Services Act (COATS Human Services Reauthorization Act) of October 27, 1998, Public Law 105–285, section 680(b) as amended. The PPR collects information regarding the outcomes and management of CED projects. OCS will use the data to critically review the overall design and effectiveness of the program.

The PPR will continue to be administered to all active grant recipients of the CED program. Grant recipients will be required to use this reporting tool for their semi-annual reports to be submitted twice a year. Through a previous renewal, the current PPR replaced both the annual questionnaire and other semi-annual reporting formats, which resulted in an overall reduction in burden for the grant recipients, significantly improved the quality of the data collected by OCS, and allowed grant recipients to become accustomed to this format. OCS seeks to renew this PPR to continue to collect quality data from grant recipients. To ensure the burden on grant recipients is not increased, but that the information collected demonstrates the full impact of the program, OCS has conducted an in-depth review of the forms and requests minor changes to the PPR to provide clarity to users filling out the form.

Respondents: Active CED Grant Recipients.

ANNUAL BURDEN ESTIMATES

Instrument	Total number of respondents	Annual number of responses per respondent	Average burden hours per response	Annual burden hours
PPR for Current OCS–CED Grant Recipients	91	2	1.5	273

Estimated Total Annual Burden Hours: 273.

Authority: Section 680(a)(2) of the Community Services Block Grant (CSBG) Act, 42 U.S.C. 9921.

Mary B. Jones,

ACF/OPRE Certifying Officer.

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BILLING CODE 4184–24–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Community Living

Agency Information Collection Activities; Submission for OMB Review; Public Comment Request; Older Americans Act, Application for Title VI Parts A/B and C Grants OMB Control Number 0985–0064

AGENCY: Administration for Community Living, Department of Health and Human Services.

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 the Paperwork Reduction Act of 1995. This 30-day notice collects comments on the information collection requirements related to the information collection requirements for the Older Americans Act, Title VI Parts A/B and C Grants PPR OMB Control Number 0985–0064.

DATES: Submit written comments on the collection of information by October 31, 2022.

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.* 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: Jasmine Aplin, Administration for Community Living, *Jasmine.Aplin@acl.hhs.gov* (202) 795–7453.

SUPPLEMENTARY INFORMATION: This is an extension to a currently approved information collection (IC). ACL is responsible for administering the Title VI A/B (Nutrition and Supportive Service) and C (Caregiver) grants. The purpose of this data collection is to improve and standardize the format of the application. The instrument will collect data as prescribed by the Older Americans Act Section 612(a), 614(a) and 45 CFR 1326.19 related to the eligibility of Federally recognized Tribes and Native Hawaiian organizations for grant funds under this program and their capacity to deliver services to elders.

The Older Americans Act, Application for Title VI Parts A/B and C Grants collects information on the ability of federally recognized American Indian, Alaskan Native and Native Hawaiian organizations to provide nutrition, supportive, and caregiver services to elders within their service area. Applicants are required to provide a description of their organization's service area, the number of eligible elders in their service area, and their ability to deliver services and sign assurances that the organization will comply with all applicable laws and regulations.

This is an extension of a currently approved information collection. The proposed data collection materials have been updated to better align with the requirements of the Older Americans Act and Federal regulations, as well as to improve data quality and grantee accountability. Furthermore, this grantee application will better line up with the Title VI Program Performance Report under 0985–0007. This data collection will also support ACL in tracking performance outcomes and efficiency measures with respect to the annual and long-term performance targets established in compliance with the Government Performance Results Modernization Act (GPRMA).

Comments in Response to the 60-Day Federal Register Notice

A notice published in the **Federal Register** *87, No. 126 on Friday, July 1, 2022.* There were No public comments in response to the 60-day notice.

Estimated Program Burden: Title VI funding is broken into three categories. Parts A and B are for nutritional and supportive programming, with Part A being restricted to American Indian and Alaska Native grantees, and Part B restricted to Native Hawaiian grantees. Part C is for caregiver programming. All Part C grantees must have Part A/B funding, but not all Part A/B grantees will have Part C programs. Therefore, there are likely to be 295 unique respondents, but only 250 will have to complete all three portions of the application. This application covers all three parts of Title VI.

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 Application Part A/B	295	1	2.75	270.4
Title VI Application Part C	250	1	1.5	125
Total			4.25	395.4

The number of burden hours associated with the Title VI, Part C, data collection was calculated as 811.25.

However, since this instrument is used only once every three years results in an annualized number of 270.4 hours. Similarly, the total hours associated with the Title VI, Part C, application is 375.

Dated: September 23, 2022.

Alison Barkoff,

Acting Administrator and Assistant Secretary for Aging.

[FR Doc. 2022–21080 Filed 9–28–22; 8:45 am]

BILLING CODE 4154–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Proposed Collection: Public Comment Request; Information Collection Request Title: Data System for Organ Procurement and Transplantation Network, OMB No. 0915–0157—Revision

AGENCY: Health Resources and Services Administration (HRSA), Department of Health and Human Services (HHS).

ACTION: Notice.

SUMMARY: In compliance with the requirement for opportunity for public comment on proposed data collection projects of the Paperwork Reduction Act of 1995, HRSA announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the ICR to OMB, HRSA seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on this ICR should be received no later than November 28, 2022.

ADDRESSES: Submit your comments to *paperwork@hrsa.gov* or by mail to the HRSA Information Collection Clearance Officer, Room 14N136B, 5600 Fishers Lane, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of the data collection plans and draft instruments, email *paperwork@hrsa.gov* or call Samantha Miller, the HRSA Information Collection Clearance Officer at (301) 443–9094.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the information collection request title for reference.

Information Collection Request Title: Data System for Organ Procurement and Transplantation Network (OPTN), OMB No. 0915–0157—Revision.

Abstract: Section 372 of the Public Health Service Act requires that the Secretary of HHS, by contract, provide for the establishment and operation of a private, non-profit entity the OPTN, which on behalf of HRSA, oversees the U.S. donation and transplantation system. The OPTN Board of Directors (BOD) determines what data must be collected to appropriately fulfill their responsibilities pursuant to their regulatory authority in 42 CFR 121.11 of the OPTN Final Rule. HRSA, on behalf of the OPTN BOD and in alignment with the Paperwork Reduction Act of 1995, submits OPTN BOD-approved data elements for collection to OMB for official federal approval.

Need and Proposed Use of the Information: HRSA and the OPTN BOD use data to develop transplant, donation, and allocation policies; to determine whether institutional members are complying with policy; to determine member-specific performance; to ensure patient safety, and to fulfill the requirements of the OPTN Final Rule. In addition, the regulatory authority in 42 CFR 121.11 of the OPTN Final Rule requires the OPTN data to be made available, consistent with applicable laws, for use by OPTN members, the Scientific Registry of Transplant Recipients, HHS, and members of the public for evaluation, research, patient information, and other important purposes.

This is a request to revise the current OPTN data collection which includes time-sensitive, life-critical data on transplant candidates and donors, the organ matching process, histocompatibility results, organ labeling and packaging, and pre- and post-transplantation data on recipients and donors. This revision also includes OPTN BOD-approved changes to the existing OMB data collection forms. The OPTN collects these specific data elements from transplant hospitals, organ procurement organizations, and histocompatibility laboratories. The OPTN uses this information to (1) facilitate organ placement and match donor organs with recipients, (2) monitor compliance of member organizations with federal laws and regulations and with OPTN requirements, (3) review and report periodically to the public on the status of organ donation and transplantation in the United States, (4) provide data to researchers and government agencies to study the scientific and clinical status of organ transplantation, and (5) perform transplantation-related public health surveillance including the possible transmission of donor disease.

HRSA is requesting to make the following OPTN BOD-approved changes to improve the OPTN organ matching and allocation process and improve OPTN member compliance with OPTN requirements:

(1) Adding data collection forms from the OPTN donor management and organ matching system to the existing OMB-approved information collection. The system allows an organ procurement organization to add donors, run the donor/potential transplant recipients matches, and place a donated organ(s) with a computer-matched potential transplant recipient. Transplant centers will access the system to view posted donor information to assist them with accepting decisions, along with other donor/potential transplant recipient functions such as entering offer responses and verifying organ offer refusals. The OPTN donor management and organ matching system is comprised of eight data collection forms: