

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN <sup>1</sup>—Continued

| Participant subgroup  | Number of respondents | Number of responses per respondent | Total annual responses | Average burden per response | Total hours <sup>1</sup> |
|---|-----------------------|------------------------------------|------------------------|-----------------------------|--------------------------|
| <b>Number to complete consent (5 min) and main study (85 min)</b> |                       |                                    |                        |                             |                          |
| Youth (aged 13–17) .....  | 50                    | 1                                  | 50                     | 1.5 .....                   | 75                       |
| Young adults (aged 18–24) .....                                   | 50                    | 1                                  | 50                     | 1.5 .....                   | 75                       |
| Total .....   |                       |                                    |                        |                             | 150                      |
| Total .....   |                       |                                    |                        |                             | 193                      |

<sup>1</sup> There are no capital costs or operating and maintenance costs associated with this collection of information.

FDA’s burden estimate is based on prior experience with research that is similar to this proposed study. Applying assumptions from previous experience in conducting similar studies, approximately 150 youth and 150 young adults would take the eligibility screener, which is estimated to take 5 minutes to read and respond. An estimated 75 parents of youth participants will provide parental permission and schedule a site visit (10 minutes total); and an estimated 50 young adults will schedule a site visit (5 minutes). Finally, approximately 50 youth and 50 young adults will complete an in-person study visit that consists of the consent/assent (5 minutes) and complete the main study (85 minutes) to yield the desired sample size of 100 total. The total estimated burden for the data collection is 193 hours. Table 1 details these estimates.

Dated: November 16, 2022.

**Lauren K. Roth,**

*Associate Commissioner for Policy.*

[FR Doc. 2022–25406 Filed 11–21–22; 8:45 am]

**BILLING CODE 4164–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Health Resources and Services Administration**

**Solicitation of Nominations for Membership To Serve on the Advisory Committee on Infant and Maternal Mortality**

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

**ACTION:** Request for nominations.

**SUMMARY:** HRSA is seeking nominations of qualified candidates for consideration for appointment as members of the Advisory Committee on Infant and Maternal Mortality (ACIMM or Committee). ACIMM advises the Secretary of HHS (Secretary) on

department activities, partnerships, policies, and programs directed at reducing infant mortality, maternal mortality and severe maternal morbidity, and improving the health status of infants and women before, during, and after pregnancy. HRSA is seeking nominations of qualified candidates to fill open positions on the ACIMM.

**DATES:** Written nominations for membership on the ACIMM must be received on or before January 23, 2023.

**ADDRESSES:** Nomination packages must be submitted electronically as email attachments to Vanessa Lee, MPH, the ACIMM’s Designated Federal Official, at: [SACIM@hrsa.gov](mailto:SACIM@hrsa.gov).

**FOR FURTHER INFORMATION CONTACT:** Vanessa Lee, MPH, Designated Federal Official, Maternal and Child Health Bureau, HRSA, 5600 Fishers Lane, Room 18N84, Rockville, Maryland 20857; 301–443–0543; or [SACIM@hrsa.gov](mailto:SACIM@hrsa.gov). A copy of the ACIMM charter and list of the current membership may be obtained by accessing the ACIMM website at <https://www.hrsa.gov/advisory-committees/infant-mortality/index.html>.

**SUPPLEMENTARY INFORMATION:** The ACIMM was established in 1991 and advises the Secretary on department activities, partnerships, policies, and programs directed at reducing infant mortality, maternal mortality and severe maternal morbidity, and improving the health status of infants and women before, during, and after pregnancy. The Committee provides advice on how to coordinate federal, state, local, tribal, and territorial governmental efforts designed to improve infant mortality, related adverse birth outcomes, and maternal health, as well as influence similar efforts in the private and voluntary sectors. The Committee provides guidance and recommendations on the policies, programs, and resources required to address the disparities and inequities in infant mortality, related adverse birth

outcomes and maternal health outcomes, including maternal mortality and severe maternal morbidity. With its focus on underlying causes of the disparities and inequities seen in birth outcomes for women and infants, the Committee advises the Secretary on the health, social, economic, and environmental factors contributing to the inequities and proposes structural, policy, and/or systems level changes. The ACIMM shall meet approximately four times per year, or at the discretion of the Designated Federal Officer in consultation with the Chair.

**Nominations:** HRSA is requesting nominations for voting members to serve as Special Government Employees (SGEs) on the ACIMM to fill open positions. The Secretary appoints ACIMM members with the expertise needed to fulfill the duties of the Advisory Committee. Information about SGE membership on the ACIMM is set forth in the ACIMM charter. Nominees sought are medical, technical, or scientific professionals with special expertise in the field of maternal and child health, in particular infant and/or maternal mortality and related health disparities; members of the public having special expertise about or concern with infant and/or maternal mortality; and/or representatives from such public health constituencies, consumers, and medical professional societies. Interested applicants may self-nominate or be nominated by another individual or organization.

ACIMM consists of up to 21 members appointed by the Secretary for a term of up to 4 years. Individuals selected for appointment to the Committee will be invited to serve for up to 4 years. Members appointed as SGEs receive a stipend and reimbursement for per diem and travel expenses incurred for attending ACIMM meetings and/or conducting other business on behalf of the ACIMM, as authorized by 5 U.S.C. 5703 for persons employed intermittently in government service.

The following information must be included in the package of materials submitted for each individual nominated for consideration: (1) A statement that includes the name and affiliation of the nominee and a clear statement regarding the basis for the nomination, including the area(s) of expertise and/or experience that may qualify a nominee for service on the ACIMM, as described above; (2) confirmation the nominee is willing to serve as a member of the ACIMM; (3) the nominee's contact information (please include home address, work address, daytime telephone number, and an email address); and (4) a current copy of the nominee's curriculum vitae or resume. Nomination packages may be submitted directly by the individual being nominated or by the person/organization recommending the candidate.

HHS endeavors to ensure that the membership of the ACIMM is fairly balanced in terms of points of view represented and that individuals from a broad representation of geographic areas, gender, and ethnic and minority groups, as well as individuals with disabilities, are considered for membership. Appointments shall be made without discrimination on the basis of age, ethnicity, gender, sexual orientation, or cultural, religious, or socioeconomic status.

Individuals who are selected to be considered for appointment will be required to provide detailed information regarding their financial holdings, consultancies, and research grants or contracts. Disclosure of this information is required in order for HRSA ethics officials to determine whether there is a potential conflict of interest between the SGE's public duties as a member of the ACIMM and their private interests, including an appearance of a loss of impartiality as defined by federal laws and regulations, and to identify any required remedial action needed to address the potential conflict.

**Authority:** ACIMM is authorized by section 222 of the Public Health Service Act (42 U.S.C. 217a), as amended. The Committee is governed by provisions of Public Law 92-463, as amended, (5 U.S.C. app. 2).

**Maria G. Button,**

*Director, Executive Secretariat.*

[FR Doc. 2022-25435 Filed 11-21-22; 8:45 am]

**BILLING CODE 4165-15-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: Ryan White HIV/AIDS Program: Allocations Forms, OMB No. 0915-0318—Revision**

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

**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 January 23, 2023.

**ADDRESSES:** Submit your comments to [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or mail the HRSA Information Collection Clearance Officer, Room 14N39, 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, mail [paperwork@hrsa.gov](mailto:paperwork@hrsa.gov) or call Samantha Miller, the acting HRSA Information Collection Clearance Officer, at (301) 443-9094.

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

*Information Collection Request Title:* Ryan White HIV/AIDS Program: Allocations Forms, OMB No. 0915-0318—Revision.

*Abstract:* HRSA's HIV/AIDS Bureau administers the Ryan White HIV/AIDS Program (RWHAP) authorized under Title XXVI of the Public Health Service Act as amended by the Ryan White HIV/AIDS Treatment Extension Act of 2009. The RWHAP Allocations and Expenditures Reports (A&E Reports) allow HRSA to monitor and track the use of grant funds for compliance with program and grants policies, and requirements as outlined in the legislation. To avoid duplication and reduce recipient reporting burden, HRSA created an electronic grantee

contract management system (GCMS) that includes data required for various reports, including the Allocations Reports and other HRSA data reports, such as the RWHAP Services Report. Recipients can access GCMS year-round to upload or manually enter data on their service provider contractors or subrecipients, the RWHAP core medical and support services provided, and their funding amounts. Data required for Allocations Reports and other reports are automatically prepopulated from GCMS. Expenditures Report data are not auto-populated in the GCMS, and are thus still manually entered into the data reporting system.

**Allocations and Expenditures (A&E) Reports**

Recipients funded under RWHAP Parts A, B, C, and D are required to report financial data to HRSA at the beginning (Allocations Report) and at the end of their grant budget period (Expenditures Report). The A&E Reports request information recipients already collect, including the use of RWHAP grant funds for core medical and support services; and on various program components, such as administration, planning and evaluation, and clinical quality management. RWHAP Parts A and B recipients funded under the Ending the HIV Epidemic Initiative (EHE) are also required to report EHE services allocations and corresponding EHE award expenditures in the A&E Reports. This allows HRSA to track and report progress toward meeting the EHE goals. The reports are similar in content; however, in the first report, recipients document the allocation of their RWHAP grant award at the beginning of their grant budget period. In the second report, recipients document actual expenditures of their RWHAP grant award (including any carryover dollars) at the end of their grant budget period.

HRSA is proposing the following updates to the RWHAP Allocation Reports.

*RWHAP Part A Allocations Report*

- Revising row and column headers and other language for clarity and alignment with RWHAP requirements;
- Combining the columns for RWHAP Part A Formula and Supplemental Allocation amounts and updating the title;
- Moving the RWHAP Part A Minority AIDS Initiative (MAI) Award Amount row after the RWHAP Part A Supplemental Award Amount row;
- Changing the calculation for Service Allocation Subtotal percent in the Total