Dated: October 9, 2024.

Eric Flamm,

Acting Associate Commissioner for Policy.

[FR Doc. 2024–23967 Filed 10–16–24; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities; Proposed Collection; Public Comment Request; Generic Information Collection Request for Health Resources and Services Administration Stakeholder Gatherings

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

ACTION: Notice.

SUMMARY: In compliance with the Paperwork Reduction Act of 1995, HRSA submitted an Information Collection Request (ICR) to the Office of Management and Budget (OMB) for review and approval. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public during the review and approval period. OMB may act on HRSA's ICR only after the 30-day comment period for this notice has closed.

DATES: Comments on this ICR should be received no later than November 18, 2024.

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 Review—Open for Public Comments'' or by using the search function.

FOR FURTHER INFORMATION CONTACT: To request a copy of the clearance requests submitted to OMB for review, email Joella Roland, the HRSA Information Collection Clearance Officer, at *paperwork@hrsa.gov* or call (301) 443–3983.

SUPPLEMENTARY INFORMATION:

Information Collection Request Title: Umbrella Generic Information Collection Request for Information Collections Related to HRSA Gatherings, OMB No. 0906–xxxx—New.

Abstract: HRSA conducts gatherings for various purposes, including conferences, meetings, workshops, webinars, trainings, communities of practice, focus groups, and other in-person or virtual gatherings for individuals and organizations that are interested in HRSA programs. To ensure that HRSA has sufficient information to plan, convene, administer, and evaluate the effectiveness of these gatherings, HRSA must collect information from potential attendees, such as contact information, organizational information, logistical information (*e.g.,* preferred delivery methods), accommodation needs, and feedback about the gathering's content. Furthermore, HRSA may conduct a test of knowledge to see what attendees know about the subject matter before or during the meeting or focus group. After the gathering concludes, attendees may be asked to complete an evaluation form and/or a test of knowledge to measure the gathering's effectiveness. In some instances, attendees may also apply and/or submit an abstract for prescreening to be selected for attendance.

An illustrative, but not exhaustive, list of examples of standardized information collection activities related to HRSA gatherings include:

- *Registration Forms:* Information collected includes name, contact information, organization/affiliation, demographic information (age, race or ethnicity, occupation, and location), and attendee accommodation needs.
- *Application Forms for panels, posters, or other presentation formats:* For application forms, information collected also includes title, author(s), organization/affiliation, and presentation abstract, in addition to the information contained in the registration form.
- *Focus Groups:* Information collected includes attendee/presenter responses to standard questions regarding topics posed to smaller groups during HRSA gatherings.
- *Pre-/Post-gathering Forms:* Information collected includes attendee/presenter preferences, feedback, pre-/post-meeting questions, and tests of knowledge in response to standard questions.

A 60-day notice published in the **Federal Register** on June 21, 2024, 89 FR 52067–68. HRSA received one comment that was outside the scope of the proposed information collection.

Need and Proposed Use of the Information: The purpose of collections under this umbrella generic information collection is to gather appropriate information to plan, administer, and evaluate HRSA gatherings. While HRSA can evaluate the general need for and the overall practical utility of such information collection in advance, HRSA may not be able to determine the details of the specific individual collections until a later time. The planning for these gatherings is often on a quick timeline and the standard timeline to comply with a full request under the Paperwork Reduction Act could inhibit HRSA's ability to collect information to inform these activities. The information collected is expected to be voluntary, low-burden, and uncontroversial. Therefore, an umbrella generic is requested to allow for quick turnaround requests for similar information collections related to these activities.

As this Generic ICR for HRSA Stakeholder Gatherings will focus on the awareness, understanding, attitudes, preferences, or experiences of HRSA customers or other stakeholders (*e.g.,* funding recipients and their delivery partners, potential funding applicants) relating to existing or future services, products, or communication materials, the Fast Track Process should apply to this information collection. Therefore, HRSA requests OMB provide a response on individual generic information collections within 5 business days.

Likely Respondents: Attendees and presenters at HRSA conferences, meetings, workshops, webinars, trainings, communities of practice, and other in-person, virtual, or hybrid gatherings. Attendees and presenters may include HRSA funding recipients, individuals seeking to participate in a HRSA-funded program, members of the public who utilize HRSA-funded resources, contractors, researchers, and other members of the public. Responses to any information collections under this Generic ICR for HRSA Stakeholder Gatherings are not required to obtain or retain any benefit.

Burden Statement: Burden in this context means the time expended by persons to generate, maintain, retain, disclose, or provide the information requested. This includes the time needed to review instructions; to develop, acquire, install, and utilize technology and systems for the purpose of collecting, validating, and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information; to search data sources; to complete and review the collection of information; and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below. HRSA conducted this estimate based on reviewing burden estimates of forms

from previous HRSA gatherings, which were approved under other Umbrella or Regular packages.

TOTAL ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
Registration Forms	100,000	1	100,000	0.5	50,000
Applications	10,000	1	10,000	1.0	10,000
Pre- and Post-Gathering Forms	200,000	1	200,000	0.5	100,000
Focus Groups	100,000	3	300,000	3.0	900,000
Total	410,000	.....	610,000	.....	1,060,000

Maria G. Button,

Director, Executive Secretariat.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), Title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Topics in Viral Pathogenesis and Immunity.

Date: November 1, 2024.

Time: 9:30 a.m. to 11:00 a.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Neerja Kaushik-Basu, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3198, MSC 7808, Bethesda, MD 20892, (301) 435–1742, *kaushikbasun@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR Panel: Fogarty International Research Training.

Date: November 7, 2024.

Time: 2:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Aruna K. Behera, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4211, MSC 7814, Bethesda, MD 20892, (301) 435–6809, *beheraak@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Cell and Molecular Biology.

Date: November 12–13, 2024.

Time: 9:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Megan L. Goodall, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 594–8334, *megan.goodall@nih.gov*.

Name of Committee: Population Sciences and Epidemiology Integrated Review Group; Cardiovascular and Respiratory Diseases Study Section.

Date: November 13–14, 2024.

Time: 9:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: In Person and Virtual Meeting.

Contact Person: Raquel L. Velazquez-kronen, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Dr., Bethesda, MD 20892, (513) 301–9047, *velazquezrl@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Program Project—Biopsychosocial Mechanisms of Substance Use after Bariatric Surgery.

Date: November 13, 2024.

Time: 1:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Sara Louise Hargrave, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3170, Bethesda, MD 20892, (301) 443–7193, *hargravesl@mail.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Cell and Developmental Biology of Eye.

Date: November 13, 2024.

Time: 2:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Rass M. Shayiq, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2182, MSC 7818, Bethesda, MD 20892, (301) 435–2359, *shayiqr@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowships: Sensory and Motor Neurosciences, Cognition and Perception.

Date: November 14–15, 2024.

Time: 8:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Address: National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD 20892.

Meeting Format: Virtual Meeting.

Contact Person: Simon Peter Peron, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Dr., Room 1009K, Bethesda, MD 20892, (301) 594–6236, *peronsp@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Small Business: Cardiovascular Sciences Activities.

Date: November 14–15, 2024.

Time: 8:00 a.m. to 8:00 p.m.

Agenda: To review and evaluate grant applications.

Address: Hilton Garden Inn Bethesda Downtown, 7301 Waverly Street, Bethesda, MD 20814.

Meeting Format: In Person.

Contact Person: Dmitri V. Gnatenko, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of