

Transplant Recipients, and members of the public for evaluation, research, patient information, and other purposes.

This is a request to expand the current OPTN data collection, approved under OMB No. 0915–0157. HRSA is submitting this new data collection, separate from OMB No. 0915–0157, since it includes new forms developed in response to an HHS Secretarial Data Directive that are not in use by OPTN. HRSA believes that separating these data collections will minimize confusion, increase clarity among OPTN members and stakeholders, and enable more direct feedback on the new forms. Both data collections include time-sensitive, life-critical data on transplant candidates and potential organ donor patients, the organ matching process, histocompatibility results, organ labeling and packaging, as well as pre- and post-transplantation data on recipients and donors. The OPTN collects these specific data elements from transplant centers.

HRSA and the OPTN use 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, procurement, 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.

This new collection consists of three new data forms as directed by the Secretary of HHS, which were developed to improve the OPTN organ matching and allocation process and OPTN member compliance with OPTN requirements:

- One new form will collect data from the point of referral of a patient to an Organ Procurement Organization (OPO) for potential deceased organ donation. These data will provide a more objective source of information on procurement practices, the management of potential donor patients, and how these practices inform the supply of deceased donor organs available for transplant. These data may also help improve the monitoring of OPO performance, facilitating quality assurance and performance improvement efforts to reduce variation in the quality of care that OPOs provide to prospective donors and donor families.

- Two new forms will expand data collection from the point of patient registration, referral, and evaluation at transplant centers. These data will enable the collection of data from the point of referral. Pre-waitlisting data will provide insight into who is referred and by whom, who is evaluated, and who is placed on the organ transplantation waiting list. These data will also facilitate the OPTN's ability to address disparities in processes of care, improve access to organ transplantation, and assess overall system performance.

Once this collection is approved, HRSA will cease use of the Death Notification Registration and the Deceased Donor Death Referral forms that are included within the existing OMB-approved Data System for Organ Procurement and Transplantation Network OMB No. 0915–0157. HRSA decided to decommission these forms to avoid unnecessary burden and redundancy in the data collected by this package and the existing OMB data collection instrument.

Likely Respondents: Transplant Centers, OPOs, and Histocompatibility Laboratories.

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.

The total estimated burden hours for this Secretary of HHS directed data collection is 252,216.84 hours.

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Form No.	Form name	Number of respondents	Number of responses per respondent	Total responses	Average burden per response (in hours)	Total burden hours
1	Pre-Waitlist Transplant Referral Form	248	1,164.68	288,840.64	0.35	101,094.22
2	Pre-Waitlist Transplant Evaluation Form	248	594.22	147,366.56	0.40	58,946.62
3	Ventilated Patient Form	56	3,292.00	184,352.00	0.47	86,645.44.00
	Total	552	620,559.20	246,686.28

The average burden estimates for both new pre-waitlist forms are based on the 2023 burden estimates of existing OMB-approved Transplant Candidate Registration forms, which were approved under OMB control number 0915–0157. The average burden estimate of the Ventilated Patient Form is based on the average burden estimate of the 2024 burden estimates of the existing OMB-approved Death Notification Registration form, with an additional 0.08 hour per collected form burden to reflect an increase in total

data fields. The burden estimate for the Ventilated Patient Form has decreased since the publication of the 60-day **Federal Register** Notice due to the removal of measures from the form in response to public comment.

Maria G. Button,

Director, Executive Secretariat.

[FR Doc. 2025–12211 Filed 6–30–25; 8:45 am]

BILLING CODE 4165–15–P

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: Population Sciences and Epidemiology Integrated Review Group, Population based Research in Infectious Disease Study Section.

Date: July 30–31, 2025.

Time: 10:00 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: James T. Snyder, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities/ Room 3G31B, National Institutes of Health, NIAID, 5601 Fishers Lane MSC 9834, Bethesda, MD 20892–9834, (240) 669–5060, james.snyder@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Program Projects: Limited Competition: National Primate Research Centers (P51).

Date: July 30–August 1, 2025.

Time: 12:00 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: Elaine Sierra-Rivera, Ph.D., IRG Chief, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6182, Bethesda, MD 20892, (301) 435–2514, riverase@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Language, Cognition, Learning and Rehabilitation.

Date: July 31, 2025.

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: DeAnna L. Adkins, Ph.D., Scientific Review Officer, Scientific Review Branch, NSC Building, Bethesda, MD 20892, 301–496–9223, deanna.adkins@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Macromolecular Biophysics & Biological Chemistry SEP.

Date: July 31, 2025.

Time: 9:30 a.m. to 6:30 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: Shan Wang, Ph.D., Scientific Review Officer, Center for

Scientific Review, National Institute of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 496–4390, shan.wang@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Program Projects: Centers of Biomedical Research Excellence (COBRE) Phase 1.

Date: July 31–August 1, 2025.

Time: 10:00 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: Kaitlyn N. Hardell, MPH, Ph.D., Scientific Review Officer, SRB, NIA, Scientific Review Branch, 5601 Fishers Lane, Suite 8B, Rockville, MD 20892, (301) 594–7945, kaitlyn.hardell@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; HIV and AIDS Research.

Date: July 31–August 1, 2025

Time: 10:00 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: Louis A. Rosenthal, Ph.D., AB Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Rm 3G42B, National Institutes of Health/NIAID, 5601 Fishers Lane, MSC 9834, Bethesda, MD 20892–9834, (240) 669–5070 rosenthalla@niaid.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Topics in basic neurovascular biology, neurodegeneration, and neurological disorders.

Date: August 1, 2025.

Time: 10:00 a.m. to 5:30 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: Gagan Deep Bajaj, Scientific Review Officer, National Institute on Drug Abuse, NIH, Scientific Review Branch, 11601 Landsdown Street, 3WF Room 09A01, Bethesda, MD 20892, (301) 402–6965, gagan.bajaj@nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: June 26, 2025.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

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Name of Committee: Center for Scientific Review Special Emphasis Panel; Vascular and Circulatory Sciences.

Date: July 30, 2025.

Time: 9:00 a.m. to 6:30 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: Imoh Sunday Okon, Ph.D., Scientific Review Officer, Office of Scientific Review, National Center for Advancing Translational Sciences (NCATS), National Institutes of Health, 9609 Medical Center Drive, Suite 1E504, Bethesda, MD 20892, (301) 347–8881, imoh.okon@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; PAR–23–110: Biomedical Technology Optimization and Dissemination Center (BTOD) (RM1).

Date: July 30, 2025.

Time: 10:00 a.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: Editorial Board.

Contact Person: Zarana Patel, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, 301–496–9295, zarana.patel@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Career Transition and Development Awards.

Date: July 31–August 1, 2025.

Time: 10:00 a.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: Katherine M. Malinda, Ph.D., Scientific Review Officer, Center for