log-in/out processes for both computer and paper files.

Case managers, should they need to use paper records into the field, will take only those records needed to complete field activities, and all paper files will be kept in a locking file box while in transport and kept in a controlled facility when not being directly used for case management functions. Records in electronic format are accessible only to authorized users using two-factor authentication and password protection through a secured system protected by encryption, firewalls, and intrusion detection systems that require additional encryption for records stored on removable media. Records that become eligible for destruction are disposed of in alignment with the destruction methods prescribed by the NIST Special Publication (SP) 800–88. The associated information technology (IT) system(s) receive Authority to Operate (ATO) under the guidance of NIST SP 800–37.

RECORD ACCESS PROCEDURES:

Individuals seeking access to records about them in this system of records must submit a written access request to the System Manager identified in the ''System Manager(s)'' section of this SORN, in accordance with the Department's Privacy Act implementation regulations in 45 CFR. The request must contain the requester's full name, address, telephone number and/or email address, date and place of birth, and signature, and should identify the repatriation program or the applicable disaster, or otherwise provide enough information to enable OHSEPR to locate the requested records.

So that HHS may verify the requester's identity, the requester's signature must be notarized or the request must include the requester's written certification that the requester is the individual who the requester claims to be and that the requester understands that the knowing and willful request for or acquisition of a record pertaining to an individual under false pretenses is a criminal offense subject to a fine of up to $5,000.

You may request that a copy of the record be sent to you, or you may request an appointment to review the record in person (including with a person of your choosing, if you provide written authorization for agency personnel to discuss the record in that person's presence). You may also request an accounting of disclosures that have been made of the record, if any.

CONTESTING RECORDS PROCEDURES:

Individuals seeking to amend records about them in this system of records must submit a written amendment request to the System Manager identified in the ''System Manager(s)'' section of this SORN, in accordance with the Department's Privacy Act implementation regulations in 45 CFR. The request must contain the same information required for an access request. The request must include verification of the requester's identity in the same manner required for an access request; must reasonably identify the record and specify the information contested, the corrective action sought, and the reasons for requesting the correction; and should include supporting information to show how the record is factually inaccurate, incomplete, untimely, or irrelevant.

NOTIFICATION PROCEDURES:

Individuals who wish to know if this system contains records about them should submit a written notification request to the System Manager identified in the ''System Manager(s)'' section of this SORN, in accordance with the Department's Privacy Act implementation regulations in 45 CFR. The request must contain the same information required for an access request and must include verification of the requester's identity in the same manner required for an access request.

EXEMPTIONS PROMULGATED FOR THE SYSTEM:

None.

HISTORY:

81 FR 46687 (July 18, 2016), 83 FR 6591 (Feb. 14, 2018).

[FR Doc. 2023–19875 Filed 9–13–23; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0438]

Agency Information Collection Request; 30-Day Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with the requirement of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, is publishing the following summary of a proposed collection for public comment.

DATES: Comments on the ICR must be received on or before October 16, 2023.

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

FOR FURTHER INFORMATION CONTACT: When submitting comments or requesting information, please include the document identifier 0990–0438–30D and project title for reference, to Sherrette A. Funn, email: *Sherrette.Funn@hhs.gov, PRA@HHS.GOV* or call (202) 264–0041 the Reports Clearance Officer.

SUPPLEMENTARY INFORMATION: Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Title of the Collection:

Type of Collection: Extension.

OMB No.: 0990–0438.

Abstract: The Office of Population Affairs (OPA), in the Office of the Assistant Secretary for Health (OASH), U.S. Department of Health and Human Services (HHS), requests clearance for the collection of performance measures specifically for FY2020 Teen Pregnancy Prevention (TPP) Program grantees. Collection of performance measures is a requirement of all TPP awards and is included in the NOFOs. The data collection will allow OPA to comply with federal accountability and performance requirements, inform stakeholders of grantee progress in meeting TPP program goals, provide OPA with metrics for monitoring TPP grantees, and facilitate individual grantees' continuous quality improvement efforts within their projects. OPA requests clearance for one year to cover reporting during the no-cost extension period of the awards.

ESTIMATED ANNUALIZED BURDEN TABLE

Form	Type of respondent	Number of respondents	Number responses per respondent	Average burden per response (in hours)	Total burden hours
Partners and sustainability	All TPP grantees	90	2	15/60	45
Training	All TPP Grantees	90	2	15/60	45
Dissemination	All TPP Grantees	90	2	30/60	90
Stakeholder Engagement	All TPP Grantees	90	2	15/60	45
Reach and Demographics	Tier 1 and Tier 2 Phase II Grantees	64	2	3	384
Dosage	Tier 1 and Tier 2 Phase II Grantees	64	2	2	256
Fidelity and Quality	Tier 1 and Tier 2 Phase II Grantees	64	2	2	256
Tier 2 Innovation Network	Tier 2 Innovation Network Grantees	14	2	15/60	7
Supportive Services (Tier 1)	Tier 1 Grantees	54	2	15/60	27
Total			2		1,155

Sherrette A. Funn,

Paperwork Reduction Act Reports Clearance Officer, Office of the Secretary.

[FR Doc. 2023–19848 Filed 9–13–23; 8:45 am]

BILLING CODE 4168–11–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Notice of Closed Meeting

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

The meeting 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: Heart, Lung, and Blood Initial Review Group; NHLBI Mentored Patient-Oriented Research Study Section.

Date: October 26–27, 2023.

Time: 10:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Rockledge I, 6705 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Fungai Chanetsa, Ph.D., MPH, Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, National Institutes of Health, 6705 Rockledge Drive, Room 206–B, Bethesda, MD 20817, (301) 402–9394. *fungai.chanetsa@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–19907 Filed 9–13–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Notice of Closed Meeting

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

The meeting 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: Heart, Lung, and Blood Initial Review Group; Clinical Trials Review Study Section.

Date: October 23, 2023.

Time: 9:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: North Bethesda Marriott Hotel & Conference Center, Montgomery County Conference Center Facility, 5701 Marinelli Road, North Bethesda, MD 20852.

Contact Person: Keary A Cope, Ph.D., Scientific Review Officer, Office of Scientific Review/DERA, National Heart, Lung, and Blood Institute, National Institutes of Health, 6705 Rockledge Drive, Room 209–A, Bethesda, MD 20892–7924, (301) 827–7912, *copeka@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.233, National Center for Sleep Disorders Research; 93.837, Heart and Vascular Diseases Research; 93.838, Lung Diseases Research; 93.839, Blood Diseases and Resources Research, National Institutes of Health, HHS)

Dated: September 11, 2023.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–19905 Filed 9–13–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute; Notice of Closed Meeting

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

The meeting 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: Heart, Lung, and Blood Initial Review Group; NHLBI Single-Site and Pilot Clinical Trials Study Section.

Date: October 25–26, 2023.

Time: 8:30 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road NW, Washington, DC 20015.