Sherrette A. Funn,

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

[FR Doc. 2024–27723 Filed 11–26–24; 8:45 am]

BILLING CODE 4151–AE–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–4040–0013]

Agency Information Collection Request. 60-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 January 27, 2025.

ADDRESSES: Submit your comments to *sagal.musa@hhs.gov* or by calling (202) 205–2634.

FOR FURTHER INFORMATION CONTACT: When submitting comments or requesting information, please include the document identifier 4040–0013–60D and project title for reference, to Sagal Musa, email: *sagal.musa@hhs.gov,* or call (202) 205–2634 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: Disclosure of Lobbying Activities (SF–LLL) and Certification Regarding Lobbying.

Type of Collection: Extension.

OMB No. 4040–0013.

Abstract: Disclosure of Lobbying Activities (SF–LLL) and Certification Regarding Lobbying are OMB-approved collections (4040–0013). These information collections are used by grant applicants. This IC expires on February 28, 2025. We are requesting a three-year clearance of these collections.

Type of respondent: The Disclosure of Lobbying Activities (SF–LLL) and Certification Regarding Lobbying forms are used by organizations to apply for Federal financial assistance in the form of grants. These forms are submitted to the Federal grant-making agencies for evaluation and review.

ANNUALIZED BURDEN HOUR TABLE

Forms (if necessary)	Respondents (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
Disclosure of Lobbying Activities (SF–LLL).	Grant Applicants	12,675	1	1	12,675
Certification Regarding Lobbying	Grant Applicants	3,952	1	0.5	1,976
Total		12,675	1		14,651

Sherrette A. Funn,

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

[FR Doc. 2024–27714 Filed 11–26–24; 8:45 am]

BILLING CODE 4151–AE–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–0434]

Agency Information Collection Request. 60-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 January 27, 2025.

ADDRESSES: Submit your comments to *Sherrette.Funn@hhs.gov* or by calling (202) 264–0041 and *PRA@HHS.GOV.*

FOR FURTHER INFORMATION CONTACT: When submitting comments or requesting information, please include the document identifier 0990–0434–60D 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: Meetings, Conferences, and Seminars—Public Accommodations and Commercial Facilities—Funding and Sponsorship.

Type of Collection: Extension.

OMB No. 0990–0434—Office of the Assistant Secretary for Financial Resources, Office of Acquisitions.

Abstract: Meetings, Conferences, and Seminars—Public Accommodations and Commercial Facilities—Funding and Sponsorship: Performance of HHS mission requires the support of contractors. In some circumstances, depending on the requirements of the specific contract, the contractor is tasked to conduct meetings, conferences, and seminars. HHSAR 311.7101(a) (Responsibilities) and the clause at HHSAR 352.211–1 (Accessibility of meetings, conferences, and seminars to persons with disabilities) require contractors to provide a plan describing the contractor's ability to meet the accessibility standards in 28 CFR part 36. HHSAR 311.7202(b) (Responsibilities) and the clause at HHSAR 352.211–2 (Conference sponsorship request and conference materials disclaimer) require contractors to provide funding disclosure and a content disclaimer statement on conference materials. As a result of these clauses, HHS contractors

providing conference, meeting, or seminars services are required to provide specific information to HHS as stated in the HHS Acquisition Regulation.

The Agency is requesting a 3-year extension to collect this information from public or private businesses.

ANNUALIZED BURDEN HOUR TABLE

Forms (if necessary)	Respondents (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
	Business (Contractor)	1,067	1	1	1,067
Total		1,067	1	1	1,067

Sherrette A. Funn,

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

[FR Doc. 2024–27806 Filed 11–26–24; 8:45 am]

BILLING CODE 4150–24–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Heart, Lung, and Blood Institute, National Institutes of Health; Notice of Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the Sickle Cell Disease Advisory Committee.

The meeting will be held as a virtual meeting and open to the public, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting. The meeting can be accessed from the NIH Videocast at the following link: *https://videocast.nih.gov/watch=55419.*

Name of Committee: Sickle Cell Disease Advisory Committee.

Date: January 14, 2025.

Time: 10:00 a.m. to 3:00 p.m.

Agenda: NHLBI Sickle Cell Disease Program Updates and Long Term Follow-up of Participants undergoing gene therapy for SCD.

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

Contact Person: julie.panepinto@nih.gov.

Meeting Format: Virtual Meeting.

Any member of the public interested in presenting oral comments to the committee may notify the Contact Person listed on this notice at least 10 days in advance of the meeting. Interested individuals and representatives of organizations may submit a letter of intent, a brief description of the organization represented, and a short description of the oral presentation. Only one representative of an organization may be allowed to present oral comments and if accepted by the committee, presentations may be limited to five minutes. Both printed and electronic copies are requested for the record. In addition, any interested person may file written comments with the committee by forwarding their statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when applicable, the business or professional affiliation of the interested person.

Information is also available on the Institute's/Center's home page: *https://www.nhlbi.nih.gov/advisory-and-peer-review-committees/nhlbi-sickle-cell-disease-advisory-committee* where an agenda and any additional information for the meeting will be posted when available.

(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: November 22, 2024.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2024–27878 Filed 11–26–24; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Draft Revised Human Immunodeficiency Virus (HIV) Organ Policy Equity Act Safeguards and Research Criteria for Transplantation of Organs Infected With HIV

AGENCY: National Institutes of Health, Department of Health and Human Services.

ACTION: Request for comments.

SUMMARY: The HOPE Act requires the Secretary of Health and Human Services (the Secretary) to develop and publish criteria for research involving the transplantation of organs from donors with HIV to recipients with HIV. In 2015, the National Institutes of Health (NIH), and the U.S. Department of Health and Human Services (HHS) published research criteria applicable to such transplants, which have been in effect for all transplants involving organs from donors with HIV as authorized by the HOPE Act. As amended in an HHS final rule published elsewhere in this issue of the **Federal Register**, the Secretary determined that participation in clinical research should no longer be a requirement for the transplantation of kidneys and livers from donors with HIV to recipients with HIV and amended the HHS regulations governing the operation of the Organ Procurement and Transplantation Network (OPTN) to reflect this determination. As a result, HOPE Act transplants involving kidneys and livers from donors with HIV no longer need to comply with the research criteria. Given this regulatory change, NIH proposes to delete aspects of the research criteria that are specific to kidney and liver transplantation. NIH proposes additional changes to the research criteria based on its review of scientific evidence and in consideration of prior public feedback concerning the criteria, including comments provided in the recent rulemaking procedure that modified the OPTN regulations. NIH invites the public to submit comments regarding the proposed changes to the research criteria.

DATES: To ensure that comments will be considered, comments must be received no later than 5 p.m. on December 12, 2024.

ADDRESSES: Comments may be submitted by any of the following methods:

- *Email: HOPEAct@mail.nih.gov.*
- *Fax:* 301–451–5671.
- *Regular Mail:* Dr. Jonah Odim, 5601 Fishers Lane, Room 6B21, MSC 9827, Bethesda, MD 20892–9827.
- *Hand Delivery, Overnight Mail, FedEx, and UPS:* Dr. Jonah Odim, 5601 Fishers Lane, Room 6B21, MSC 9827, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Dr. Jonah Odim, Chief Clinical Transplantation Section, Transplantation Branch, 5601 Fishers