

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Institute on Aging; 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:* National Institute on Aging Special Emphasis Panel; Postbaccalaureate and Summer Research Education in AD/ADRD.

*Date:* October 29, 2024.

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

*Agenda:* To review and evaluate grant applications.

*Place:* National Institute on Aging, 5601 Fishers Ln., Rockville, MD 20852 (Virtual Meeting).

*Contact Person:* Lisa-Marie Tisdale Rowell, Ph.D., Scientific Research Officer, Scientific Review Branch, National Institute of Health, National Institute on Aging, 5601 Fishers Lane, Rockville, MD 20852, RM 1007G, (301) 594-5622, [wigfallt@mail.nih.gov](mailto:wigfallt@mail.nih.gov).

(Catalogue of Federal Domestic Assistance Program Nos. 93.866, Aging Research, National Institutes of Health, HHS)

Dated: July 26, 2024.

**Miguelina Perez,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2024-16947 Filed 7-31-24; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### Proposed Collection; 60-Day Comment Request; Electronic Individual Development Plan (eIDP) (National Eye Institute)

**AGENCY:** National Institutes of Health, HHS.

**ACTION:** Notice.

**SUMMARY:** In compliance with the requirement of the Paperwork

Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Eye Institute of the National Institutes of Health will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

**DATES:** Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

**FOR FURTHER INFORMATION CONTACT:** To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Dr. Cesar E. Perez-Gonzalez, Training Director, Office of the Scientific Director, National Eye Institute, NIH, Building 31, Room 6A22, MSC 0250, Bethesda, Maryland 20892 or call non-toll-free number (301) 451-6763 or Email your request, including your address to: [cesarp@nei.nih.gov](mailto:cesarp@nei.nih.gov). Formal requests for additional plans and instruments must be requested in writing.

**SUPPLEMENTARY INFORMATION:** Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 requires: written comments and/or suggestions from the public and affected agencies are invited to address one or more of the following points: (1) Whether the proposed collection of information is necessary for the proper performance of the function of the agency, including whether the information will have practical utility; (2) The accuracy of the agency's estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (3) Ways to enhance the quality, utility, and clarity of the information to be collected; and (4) Ways to minimize the burden of the collection of information on those who are to respond, including the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology.

*Proposed Collection Title:* Electronic Individual Development Plans, 0925-0772 extension, expiration date 10/31/2024, National Eye Institute (NEI), National Institutes of Health (NIH).

*Need and Use of Information Collection:* The National Eye Institute's (NEI) Office of the Scientific Director (OSD) goal is to train the next generation of vision researchers and

ophthalmologists. Trainees who participate in NEI research come with different levels of education (student, postbaccalaureate, predoctoral including graduate and medical students, postdoctoral fellows) and for different amounts of time (6 months to 5 years). Training at the NEI focuses on scientific and professional skill development. To enhance their chances of obtaining their ideal career, completing an annual Individual Development Plan (IDP) is an important step in helping a trainee's career and professional development and is standard practice in graduate and postdoctoral education. An IDP is an effective tool for trainees to think about their career goals and skills needed to achieve them during their time at the NEI. Trainees work together with their research mentor to organize and summarize their research projects, consider career goals, and set training goals and expectations, both for the mentee and mentor.

This information collection request is to implement an electronic Individual Development Plan (eIDP). The data collected comes from a detailed questionnaire focused on responses to professional goals and expectations while they are at the NEI. It is expected that the trainees will complete the eIDP annually and by doing so, it will help enhance the effectiveness of their training by setting clear goals that can be monitored not only by the trainee themselves but also by their mentor, the Training Director, and their Administrative Officer. In addition to this eIDP, the system will also implement an electronic exit survey. The data collected comes from a detailed questionnaire focused on responses to questions focused on trainee mentoring and professional experiences at the NEI as well as their plans after they depart. It is expected that the trainees will complete at the end of their tenure and that by doing so, the NEI Training Program can learn about ways to improve career development opportunities for future trainees as well as learn more about trainee job choices to better advise fellows. Additionally, we can use the survey to help determine mentor effectiveness and help identify problems in mentoring at the NEI.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 450.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
eIDP .....	150	1	2	300
Exit Survey Part 1 .....	150	1	30/60	75
Exit Survey Part 2 .....	150	1	30/60	75
Total .....	150	150	3	450

Dated: July 25, 2024.  
**Cesar E. Perez-Gonzalez,**  
*Training Director, National Eye Institute,  
National Institutes of Health.*  
[FR Doc. 2024–16917 Filed 7–31–24; 8:45 am]  
**BILLING CODE 4140–01–P**

DEPARTMENT OF HEALTH AND  
HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Notice of  
Meeting

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the President’s Cancer Panel. The meeting will be held as a virtual meeting and will be open to the public as indicated below. Individuals who plan to view the virtual meeting and need special assistance or other reasonable accommodations to view the meeting, should notify the Contact Person listed below in advance of the meeting. The meeting can be accessed by clicking on the links below.

*Name of Committee:* President’s Cancer Panel.  
*Date:* September 12, 2024.  
*Time:* 11:00 a.m. to 4:30 p.m.  
*Agenda:* Developing and Retaining a Robust and Diverse Cancer Workforce: Challenges and Opportunities Across the National Cancer Program.  
*Place:* National Institutes of Health, 31 Center Drive, Building 31, Room 11A48, Rockville, MD 20850 (Virtual Meeting), Access to Meeting: <https://nci.rev.vbrick.com/#/webcasts/presidentscancerpanel-meet1>.  
*Date:* September 13, 2024.  
*Time:* 11:00 a.m. to 5:30 p.m.  
*Agenda:* Developing and Retaining a Robust and Diverse Cancer Workforce: Challenges and Opportunities Across the National Cancer Program.  
*Place:* National Institutes of Health, 31 Center Drive, Building 31, Room 11A48, Rockville, MD 20850 (Virtual Meeting), Access to Meeting: <https://nci.rev.vbrick.com/#/webcasts/presidentscancerpanel-meet2>.

*Contact Person:* Samantha L. Finstad, Ph.D., Executive Secretary, President’s Cancer Panel, Office of the Director, National Cancer Institute, NIH, 31 Center Drive, Room

11A30B, MSC 2590, Bethesda, MD 20892, 240–276–6460, [samantha.finstad@nih.gov](mailto:samantha.finstad@nih.gov).  
Any interested person may file written comments with the committee by forwarding the 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: <http://deainfo.nci.nih.gov/advisory/pcp/index.htm>, where an agenda and any additional information for the meeting will be posted when available.  
(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)  
Dated: July 26, 2024.  
**Lauren A. Fleck,**  
*Program Analyst, Office of Federal Advisory Committee Policy.*  
[FR Doc. 2024–16949 Filed 7–31–24; 8:45 am]  
**BILLING CODE 4140–01–P**

DEPARTMENT OF HEALTH AND  
HUMAN SERVICES

Substance Abuse and Mental Health  
Services Administration

Current List of HHS-Certified  
Laboratories and Instrumented Initial  
Testing Facilities Which Meet Minimum  
Standards To Engage in Urine and Oral  
Fluid Drug Testing for Federal  
Agencies

**AGENCY:** Substance Abuse and Mental Health Services Administration, HHS.  
**ACTION:** Notice.

**SUMMARY:** The Department of Health and Human Services (HHS) notifies Federal agencies of the laboratories and Instrumented Initial Testing Facilities (IITFs) currently certified to meet the standards of the Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines) using Urine and the laboratories currently

certified to meet the standards of the Mandatory Guidelines using Oral Fluid.  
**FOR FURTHER INFORMATION CONTACT:** Anastasia Flanagan, Division of Workplace Programs, SAMHSA/CSAP, 5600 Fishers Lane, Room 16N06B, Rockville, Maryland 20857; 240–276–2600 (voice); [Anastasia.Flanagan@samhsa.hhs.gov](mailto:Anastasia.Flanagan@samhsa.hhs.gov) (email).  
**SUPPLEMENTARY INFORMATION:** The Department of Health and Human Services (HHS) publishes a notice listing all HHS-certified laboratories and Instrumented Initial Testing Facilities (IITFs) in the **Federal Register** during the first week of each month, in accordance with Section 9.19 of the Mandatory Guidelines for Federal Workplace Drug Testing Programs (Mandatory Guidelines) using Urine and Section 9.17 of the Mandatory Guidelines using Oral Fluid. If any laboratory or IITF certification is suspended or revoked, the laboratory or IITF will be omitted from subsequent lists until such time as it is restored to full certification under the Mandatory Guidelines.  
If any laboratory or IITF has withdrawn from the HHS National Laboratory Certification Program (NLCP) during the past month, it will be listed at the end and will be omitted from the monthly listing thereafter.  
This notice is also available on the internet at <https://www.samhsa.gov/workplace/drug-testing-resources/certified-lab-list>.  
HHS separately notifies Federal agencies of the laboratories and IITFs currently certified to meet the standards of the Mandatory Guidelines using Urine and of the laboratories currently certified to meet the standards of the Mandatory Guidelines using Oral Fluid.  
The Mandatory Guidelines using Urine were first published in the **Federal Register** on April 11, 1988 (53 FR 11970), and subsequently revised in the **Federal Register** on June 9, 1994 (59 FR 29908); September 30, 1997 (62 FR 51118); April 13, 2004 (69 FR 19644); November 25, 2008 (73 FR 71858); December 10, 2008 (73 FR 75122); April 30, 2010 (75 FR 22809); January 23,