

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
The National Survey of Organ Donation Attitudes and Practices—Telephone (English and Spanish versions) .....	2,000	1	2,000	0.37	740
The National Survey of Organ Donation Attitudes and Practices—Web (English and Spanish versions) .....	8,000	1	8,000	0.27	2,160
<b>Total</b> .....	<b>10,000</b>	.....	.....	.....	<b>2,900</b>

Amy P. McNulty,  
Deputy Director, Executive Secretariat.  
[FR Doc. 2024–11246 Filed 5–21–24; 8:45 am]  
BILLING CODE 4165–15–P

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Submission for OMB Review; 30-Day Comment Request; Application and Impact of Clinical Research Training on Healthcare Professionals in Academia and Clinical Research (Office of the Director)**

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

**ACTION:** Notice.

**SUMMARY:** In compliance with the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below.

**DATES:** Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

**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](http://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:** To request more information on the proposed project or to obtain a copy of the data collection plans and instruments, contact: Dr. Anne Zajicek, Program Director, Office of Clinical Research Education and Collaboration Outreach, OD, NIH, Building 1, Room 201, 1 Center Drive, Bethesda, MD 20892, or call non-toll-free number (301) 480–9913 or Email your request, including your address to: [zajiceka@mail.nih.gov](mailto:zajiceka@mail.nih.gov).

**SUPPLEMENTARY INFORMATION:** This proposed information collection was previously published in the **Federal Register** on Monday, July 31, 2023, Volume 88, pages 49472–49473 (64 FR 16184) and allowed 60 days for public comment. No public comments were received. The purpose of this notice is to allow an additional 30 days for public comment. The Office of Clinical Research Education and Collaboration Outreach, Office of the Director, National Institutes of Health, may not conduct or sponsor, and the respondent is not required to respond to, an information collection that has been extended, revised, or implemented on or after October 1, 1995, unless it displays a currently valid OMB control number.

In compliance with Section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the National Institutes of Health (NIH) has submitted to the Office of Management and Budget (OMB) a request for review and approval of the information collection listed below.

*Proposed Collection:* Application and Impact of Clinical Research Training on Healthcare Professionals in Academia and Clinical Research, 0925–0764-expiration date, 02/28/2026, Office of Clinical Research Education and Collaboration Outreach (OCRECO), National Institutes of Health (NIH), Office of the Director (OD).

*Need and Use of Information Collection:* The purpose of this survey is to assess the long-term impact and outcomes of clinical research training programs provided by the Office of Clinical Research Education and Collaboration Outreach located in the NIH Office of the Director (OD) over a ten-year follow-up period. The information received from respondents will provide insight on the following: impact of the courses on (a) promotion of professional competence, (b) research productivity and independence, and (c) future career development within clinical, translational and academic research settings. These surveys will provide preliminary data and guidance in (1) developing recommendations for collecting outcomes to assess the effectiveness of the training courses, and (2) tracking the impact of the curriculum on participants’ ability to perform successfully in academic, non-academic, research, and non-research settings.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 1,773.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondents	Estimated number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
OCRECO Learning Portal Registration (Attachment 1)	Healthcare Professionals .....	2,000	1	5/60	167
	Students .....	2,000	1	5/60	167
	General Public .....	1,000	1	5/60	83
IPPCR Lecture Evaluation (Attachment 2) .....	Healthcare Professionals .....	1,000	1	5/60	83
	Students .....	2,000	1	5/60	167
	General Public .....	1,000	1	5/60	83

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Form name	Type of respondents	Estimated number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
IPPCR Final Course Evaluation (Attachment 4) .....	Healthcare Professionals .....	1,000	1	5/60	83
	Students .....	2,000	1	5/60	167
	General Public .....	1,000	1	5/60	83
PCP Lecture Evaluation (Attachment 3) .....	Healthcare Professionals .....	1,000	1	5/60	83
	Students .....	2,000	1	5/60	167
	General Public .....	1,000	1	5/60	83
PCP Final Course Evaluation (Attachment 5) .....	Healthcare Professionals .....	1,000	1	5/60	83
	Students .....	2,000	1	5/60	167
	General Public .....	1,000	1	5/60	83
NIH Summer Course in Clinical and Translational Research Course Evaluation (Attachment 6). Sabbatical in Clinical Research Management Course Evaluation (Attachment 7).	Healthcare Professionals .....	20	1	5/60	2
Ethical and Regulatory Aspects of Clinical Research (Asynchronous/Online) Course (Attachment 8).	Healthcare Professionals .....	100	1	5/60	8
	Students .....	50	1	5/60	4
	General Public .....	100	1	5/60	8
<b>Total</b> .....	.....	.....	21,290	.....	1,773

Dated: May 16, 2024.  
**Lawrence A. Tabak,**  
*Principal Deputy Director, National Institutes of Health.*  
 [FR Doc. 2024-11257 Filed 5-21-24; 8:45 am]  
**BILLING CODE 4140-01-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Proposed Collection; 60-Day Comment Request; Post-Award Reporting Requirements Including Research Performance Progress Report Collection (OD)**

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

**SUMMARY:** In compliance with the requirement of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Institutes of Health (NIH) 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: Ms. Mikia P. Currie, Program

Analyst, Office of Policy for Extramural Research Administration, 6705 Rockledge Drive, Suite 350, Bethesda, Maryland 20892, or call a non-toll-free number 301-435-0941 or email your request, including your address to [ProjectClearanceBranch@mail.nih.gov](mailto:ProjectClearanceBranch@mail.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 on 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:* Public Health Service (PHS) Post-award Reporting Requirements Revision, OMB 0925-0002, Expiration Date 01/31/2026, Office of the Director (OD), National Institutes of Health (NIH).

*Need and Use of Information Collection:* Starting in 2025, NIH will

require applicable recipients to address progress in association with their approved Data Management and Sharing Plans within the Research Performance Progress Report (RPPR) in accordance with the final NIH Policy for Data Management and Sharing to promote the management and sharing of scientific data generated from NIH-funded or conducted research. The progress report forms will be updated to align with this requirement. The Training Data Tables will also be updated to reduce the burden and promote consistent information collection, including limiting the scope of information collection to data only relevant to the training stage(s) of the proposed program in Table 1 and removing instructions in Table 8 that are reported within the RPPR. Effective May 2025, NIH will be adopting the Common Forms for Biographical Sketch and Current and Pending (Other) Support as part of the directive from Guidance for Implementing National Security Presidential Memorandum (NSPM)-33. The Common Forms are part of a separate OMB collection, currently approved under 3145-0279. As such, elements collected within the Common Forms will be removed from NIH's current NIH Biosketch and Other Support formats. NIH will continue to collect additional information not captured on the Common Forms to adhere to the agency's implementation of the NIH Peer Review Regulations at 42 CFR part 52 as part of the NIH Biosketch form, which will be renamed the NIH Biosketch Supplement to reflect the supplemental information requested. The application and progress