ANNUALIZED BURDEN HOUR TABLE—Continued

Forms (if necessary)	Respondents (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response	Total burden hours
Research & Related Subaward Budget (Total Fed + Non-Fed) 5 YR 30 ATT.	Grant Applicants	303	1	1	303
Research & Related Senior/Key Person Profile	Grant Applicants	265	1	1	265
Research & Related Personal Data	Grant Applicants	24,012	1	1	24,012
Research & Related Multi-Project 10 Year Budget	Grant Applicants	4,606	1	1	4,606
Research & Related Budget 10YR	Grant Applicants	20,652	1	1	20,652
R&R Subaward Budget Attachment(s) Form 5 YR 30 ATT	Grant Applicants	37,048	1	1	37,048
R&R Subaward Budget Attachment(s) Form 10 YR 30 ATT	Grant Applicants	2,707	1	1	2,707
R&R Subaward Budget Attachment(s) Form	Grant Applicants	3,138	1	1	3,138
R&R R Multi-Project Subaward Budget Attachment(s) Form 10 YR 30 ATT.	Grant Applicants	4,606	1	1	4,606
Total		704,377	1	1	704,377

Catherine Howard,

Paperwork Reduction Act Reports Clearance Officer, Department of Health and Human Services, Office of the Secretary.

[FR Doc. 2025–12799 Filed 7–9–25; 8:45 am]

BILLING CODE 4151–AE–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Office of the Director, Notice of Charter Renewal

In accordance with Title 41 of the U.S. Code of Federal Regulations, Section 102–3.65(a), notice is hereby given that the charter for the Advisory Committee to the Director, National Institutes of Health, was renewed for an additional two-year period on May 31, 2027.

It is determined that the Advisory Committee to the Director, National Institutes of Health, is in the public interest in connection with the performance of duties imposed on the National Institutes of Health by law, and that these duties can best be performed through the advice and counsel of this group.

Inquiries may be directed to Patricia Brandt Hansberger, Acting Director, Office of Federal Advisory Committee Policy, Office of the Director, National Institutes of Health, 6701 Democracy Boulevard, Suite 1000, Bethesda, Maryland 20892 (Mail code 4875), Telephone (301) 594–2492, or *Patricia.Hansberger@nih.gov.*

Dated: July 8, 2025.

David W. Freeman,

Supervisory Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2025–12898 Filed 7–9–25; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Promoting Objectivity in Research 42 CFR Part 50 Subpart F and Responsible Prospective Contractors 45 CFR Part 94 (NIH/OD)

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 Institutes of Health, Office of Policy and Extramural Research Administration (OPERA), Office of Extramural Research (OER) 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: Brian E. Sass-Hurst, Acting Director, Division of Grants Compliance and Oversight, Office of Policy for Extramural Research Administration, National Institutes of Health, 6705 Rockledge Drive, Suite 800, or call non-toll-free number (301) 827–7581 or Email your request, including your address to *Brian.Sass-Hurst@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: Promoting Objectivity in Research 42 CFR part 50 Subpart F and Responsible Prospective Contractors 45 CFR part 94, 0925–0417, expiration date 06/30/2025, EXTENSION, Office of Policy and Extramural Research Administration (OPERA), Office of Extramural Research (OER), Office of the Director (OD), National Institutes of Health (NIH).

Need and Use of Information Collection: This request is for an Extension of a currently approved collection resulting from regulations regarding Promoting Objectivity in Research (42 CFR part 50, subpart F) and Responsible Prospective Contractors (45 CFR part 94). The purpose of these regulations is to promote objectivity in research by requiring institutions to establish standards to ensure that there is no

reasonable expectation that the design, conduct, or reporting of Public Health Service (PHS) funded research will be biased by any Investigator financial conflict of interest (FCOI).

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 680,473.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondents based on applicable section of regulation	Number of respondents	Number of responses per respondent	Average burden per response (in hrs.)	Total annual burden hours
Reporting				
Initial Reports under 42 CFR 50.605(b)(1) and (b)(3) or 45 CFR 94.5(b)(1) and (b)(3) from awardee Institutions.	1,128	1	2	2,256
Subsequent Reports under 42 CFR 50.605(a)(3)(iii) and (b)(2) or 45 CFR 94.5(a)(3)(iii) and (b)(2) from awardee Institutions.	50 FCOI reports as in 42 CFR 50.605(a)(3)(ii) and 45 CFR 94.5(a)(3)(ii). 5 mitigation reports	1 1	2 2	100 10
Annual Report under 42 CFR 50.605(b)(4) or 45 CFR 94.5(b)(4) from awardee Institutions.	2,712	1	1	2,712
Subsequent Reports under 42 CFR 50.606(a) or 45 CFR 94.6 from awardee Institutions.	20	1	10	200
Record Keeping				
Under 42 CFR 50.604(i) or 45 CFR 94.4(i) from awardee institutions.	2,000	1	4	8,000
Disclosure				
Under 42 CFR 50.604(a) or 45 CFR 94.4 for Investigators.	3,000	1	81	243,000
Under 42 CFR 50.604(b) or 45 CFR 94.4(e)(1) for Investigators.	38,000	1	30/60	19,000
Under 42 CFR 50.604(b) or 45 CFR 94.4(e)(1) for Institutions.	2,000	1	6	12,000
Under 42 CFR 50.604(c)(1) or 45 CFR 94.4(c)(1) from subrecipients.	500	1	1	500
Under 42 CFR 50.604(d) or 45 CFR 94.4 for Institutions.	[1] 3,000	1	1	3,000
Under 42 CFR 50.604(e)(1) or 45 CFR 94.4(e)(1) for Investigators.	38,000	1	4	152,000
Under 42 CFR 50.604(e)(2) or 45 CFR 94.4(e)(2) for Investigators.	38,000	1	1	38,000
Under 42 CFR 50.604(e)(3) or 45 CFR 94.4(e)(3) for Investigators.	1,128	1	30/60	564
Under 42 CFR 50.604(f) or 45 CFR 94.4(f) for institutions.	2,000	1	1	2,000
Under 42 CFR 50.605(a)(1) or 45 CFR 94.5(a)(1) for Institutions.	[2] 2,000	1	82	164,000
Under 42 CFR 50.605(a)(3) or 45 CFR 94.5(a)(3) for Institutions.	[3] 500	1	3	1,500
Under 42 CFR 50.605(a)(3)(i) or 45 CFR 94.5(a)(3)(i)	[4] 50	1	80	4,000
Under 42 CFR 50.605(a)(3)(ii) or 45 CFR 94.5(a)(3)(ii).	[5] 50	1	80	4,000
Under 42 CFR 50.605(a)(3)(iii) or 45 CFR 94.5(a)(3)(iii).	50	1	1	50
Under 42 CFR 50.605(a)(4) or 45 CFR 94.5(a)(4)	1,128	1	12	13,536
Public Website Posting under 42 CFR 50.605(a)(5) or 45 CFR 94.5(a)(5) from awardee Institutions.	2,000	1	5	10,000
Under 42 CFR 50.606(c) or 45 CFR 94.6(c)	[6] 50	[7 6] 3	18/60	45
Total	137,371	137,371		680,473

[1] Assuming that 3000 institutions solicit disclosures on an annual basis to all Investigators.

[2] Although an estimated 1,128 reports of Financial Conflict of Interest are expected annually, the 2,000 responding Institutions must review all financial disclosures associated with PHS-funded awards to determine whether any financial conflicts of interest exist. Thus, the review burden of 76,000 hours is based upon estimates that it will take on the average 2 hours for an institutional official(s) to review each of 38,000 financial disclosures associated with PHS funded awards. The burden for developing a management plan for identified FCOI is estimated at 80 hours × 1,128 cases = 90,240 hours.

[3] Assuming that this is a rare occurrence based on prior experience.

[4] Assuming only a fraction of the newly identified SFIs will constitute FCOI.

[5] Assuming only a fraction of the newly identified SFIs will constitute FCOI.

[6] Number based on 50.605/94.5(a)(3)(i)—of those only a fraction will relate to a project of clinical research whose purpose is to evaluate the safety or effectiveness of a drug, medical device, or treatment, but we are calculating the maximum estimated burden.

[7] Assuming an average of 3 publications annually.

Dated: July 3, 2025.

Jon Lorsch,

Acting NIH Deputy Director for Extramural Research, National Institutes of Health.

[FR Doc. 2025–12897 Filed 7–9–25; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-day Comment Request; Generic Clearance for Conferences, Meetings, Workshops, Poster Sessions and Registrations (Office of the Director)

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 Institutes of Health Office of the Director (OD) 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, Chief, Project Clearance Branch (PCB), Office of Policy for Extramural Research Administration, 6705 Rockledge Drive, Suite 803–B, Bethesda, Maryland 20892 or call non-toll-free number (301) 435–0941 or Email your request, including your address to: *curriem@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 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 minimizes 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: Generic Clearance for Conferences, Meetings, Workshops, Poster Sessions and Registrations, exp., date 09/30/2025 Extension, Office of the Director (OD), National Institutes of Health (NIH).

Need and Use of Information Collection: The information collection encompassed by this generic clearance continues to allow NIH to select the most appropriate participants for non-grantee activities sponsored, organized, and run by NIH staff, according to the type and purpose of the activity. For example, NIH may develop an application process or information collection to select a limited number of researchers to participate in a poster session, identify speakers and panelists with desired expertise on a specific topic to be covered at a meeting, or determine which researchers would mostly likely benefit from a training course or other opportunity. For NIH to plan and conduct activities that are timely for participants in their field of research, it is often necessary for such information to be collected within a relatively short turnaround time. In general, submitted abstracts or other application materials will be reviewed by an internal NIH committee responsible for planning the activities. This committee will be responsible for selecting and notifying participants. The information collected for these activities generally include title, author(s), and institution/organization, poster size and character limitations along with other requirements. This information is necessary to identify attendees eligible, present research, speak on panels, and discuss innovative approaches to science and technology for poster presentations among their peers. The registration form collects information from interested parties to register them and obtain the necessary qualifications for conferences, meetings, workshops, poster sessions, presentations and panels.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 8,075.

ESTIMATED ANNUALIZED BURDEN TABLE

Type of activity/form	Number of respondents	Number of responses per respondent	Average burden (in hours) per response	Total burden hours
Conferences	2,000	1	1	2,000
Meetings	2,000	1	45/60	1,500
Workshops	2,000	1	30/60	1,000
Poster Session	1,000	1	1	1,000
Panels	1,000	1	30/60	500
Presentations	1,000	1	1	1,000
Common Registration Form	1,500	1	3/60	75
Common Abstract Form	1,000	1	1	1,000
Total		11,500		8,075