would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Institute of Neurological Disorders and Stroke Special Emphasis Panel; Phase 2 Clinical Trials in Neurology.

Date: August 14, 2023.

Time: 3:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Neuroscience Center, 6001 Executive Boulevard, Rockville, MD 20852 (Virtual Meeting).

Contact Person: Iqbal Sayeed, Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH NSC, 6001 Executive Blvd., Suite 3208, MSC 9529, Rockville, MD 20852, 301–496–9223, *iqbal.sayeed@nih.gov*.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research Related to Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS.)

Dated: July 25, 2023.

Tyeshia M. Roberson-Curtis,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–16134 Filed 7–28–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-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 requirement of the Paperwork Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the Office of Clinical Research Education, Collaboration, and Outreach (OCRECO), Office of the Director (OD), 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. Anne Zajicek, M.D., Pharm.D., Program Director, Office of Clinical Research Education, Collaboration, and Outreach, NIH Office of the Director, Building 1, Room 201, MSC–0155, Bethesda, Maryland 20892 or call non-toll-free number (301) 480–9913 or email your request, including your address to: *zajiceka@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: Application and Impact of Clinical Research Training on Healthcare Professionals in Academia and Clinical Research, 0925–0764, exp., date 02/28/2026, Revision 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 revision is to: update the name of the office responsible for these on-line training programs (from Office of Clinical Research to Office of Clinical Research Education and Collaboration Outreach); revise the course evaluation survey questions; add an additional on-line course, ''Ethical and Regulatory Aspects of Clinical Research''; change the course opening and close dates from Oct–June to Sept–July. The survey will continue to assess the long-term impact and outcomes of clinical research training programs provided by the newly formed Office of Clinical Research Education, Collaboration, and Outreach (previously the Office of Clinical Research) 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 820.

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	2000	1	5/60	167
	Students	1000	1	5/60	83
	General Public	500	1	5/60	42
IPPCR Lecture Evaluation (Attachment 2)	Healthcare Professionals	750	1	5/60	63
	Students	500	1	5/60	42
	General Public	250	1	5/60	21
IPPCR Final Course Evaluation(Attachment 4)	Healthcare Professionals	750	1	5/60	63
	Students	500	1	5/60	42

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
	General Public	250	1	5/60	21
PCP Lecture Evaluation (Attachment 3)	Healthcare Professionals	750	1	5/60	63
	Students	500	1	5/60	42
	General Public	250	1	5/60	21
PCP Final Course Evaluation (Attachment 5)	Healthcare Professionals	750	1	5/60	63
	Students	500	1	5/60	42
	General Public	250	1	5/60	21
NIH Summer Course in Clinical and Translational Research Course Evaluation (Attachment 6).	Healthcare Professionals	20	1	5/60	2
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
Total			9,790		820

Dated: July 25, 2023.

Tara A. Schwetz,

Acting Principal Deputy Director, National Institutes of Health.

[FR Doc. 2023–16184 Filed 7–28–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Human Genome Research 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 National Advisory Council for Human Genome Research.

This is a hybrid meeting held in-person and virtually and is open to the public as indicated below. Individuals who plan to attend in-person or view the virtual meeting and need special assistance or other reasonable accommodations, should notify the Contact Person listed below in advance of the meeting. The meeting will be videocast and can be accessed from *https://www.genome.gov/about-nhgri/Institute-Advisors/National-Advisory-Council-for-Human-Genome-Research.*

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 Advisory Council for Human Genome Research.

Date: September 18–19, 2023.

Closed: September 18, 2023, 9:00 a.m. to 10:00 a.m.

Agenda: To review and evaluate grant applications.

Place: National Human Genome Research Institute, National Institutes of Health, 6700B Rockledge Drive, Suite 1100, Bethesda, MD 20892.

Open: September 18, 2023, 10:30 a.m. to 6:30 p.m.

Agenda: Report of Institute Director and Institute Staff.

Place: National Human Genome Research Institute, 6700B Rockledge Drive, Suite 1100, Bethesda, MD 20892.

Closed: September 19, 2023, 10:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Human Genome Research Institute, National Institutes of Health, 6700B Rockledge Drive, Suite 1100, Bethesda, MD 20892.

Contact Person: Rudy O. Pozzatti, Ph.D., Scientific Review Officer, Scientific Review Branch, National Human Genome Research Institute, 6700B Rockledge Drive, Suite 1100, Rockville, MD 20892, (301) 402–0838, *pozzattr@mail.nih.gov*.

Any interested person may file written comments with the committee within 15 days after the meeting 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://www.genome.gov/council,* where an agenda and any additional information for the meeting will be posted when available.

(Catalogue of Federal Domestic Assistance Program Nos. 93.172, Human Genome Research, National Institutes of Health, HHS)

Dated: July 26, 2023.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–16147 Filed 7–28–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Generic Clearance for NIH Citizen Science and Crowdsourcing Projects (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/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, Attention: Desk Officer for NIH.

FOR FURTHER INFORMATION CONTACT: To request more information on the proposed project or to obtain a copy of