

remove the IRB roster membership information from the IRB registration form. This change will align the IRB registration form with the 2018 Requirements at 45 CFR 46.103. The change, when implemented, is anticipated to result in a shorter, simplified IRB registration process for respondents. Updates to the software applications OHRP uses to manage IRB registration will be deployed to enable such changes.

The current form is approved through June 30, 2025. The purpose of the form is to provide a simplified procedure for: (1) IRBs to satisfy the requirements for IRB registration at 45 CFR part 46, subpart E; and (2) IRBs in the United

States (US) to satisfy the FDA requirements for IRB registration at 21 CFR 56.106.

Institutions engaged in nonexempt human subjects research conducted or supported by HHS, or another Common Rule department or agency, are required by the terms of the Federalwide Assurance (FWA) to rely upon only IRBs registered with OHRP for review of research to which the FWA applies, and must designate a registered IRB on the institution's FWA submission to OHRP. In this way, OHRP's FWA submission process, established pursuant to the requirements for assurances at 45 CFR 46.103, is linked to the regulatory requirements for IRB registration.

The respondents for this information collection are institutions or organizations operating IRBs that review human subjects research conducted or supported by HHS; or, in the case of FDA's requirements, each IRB in the United States that reviews clinical investigations regulated by FDA under sections 505(i) or 520(g) of the Federal Food, Drug and Cosmetic Act; and each IRB in the United States that reviews clinical investigations that are intended to support applications for research or marketing permits for FDA-regulated products. Many of the IRBs also review research conducted or supported by other Common Rule departments and agencies.

ANNUALIZED BURDEN HOUR TABLE

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
IRB Registration 0990–0279 Update and Renew	5,350	1	0.33	1,766
IRB-Registration 0990–0279 Initial and Update	350	2	0.5/0.33	291
Total				2,057

Susan R. Little,

Department Information Collection Clearance Officer, Paperwork Reduction Act Program, Department of Health and Human Services.

[FR Doc. 2025–09001 Filed 5–20–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

[Document Identifier: OS–0990–new]

Agency Information Collection Request; 60-Day Public Comment Request

AGENCY: Office of the Assistant Secretary for Technology Policy/Office of the National Coordinator for Health IT, 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 July 21, 2025.

ADDRESSES: Submit your comments to ASTP_Data@hhs.gov.

FOR FURTHER INFORMATION CONTACT: When submitting comments or

requesting information, please include the document identifier 0990–New–60D and project title for reference to ASTP_Data@hhs.gov and Meghan Gabriel at Meghan.Gabriel@hhs.gov, PRA@hhs.gov or call 202–465–0597.

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: National Survey of Digital Health Companies.

Type of Collection: New Data Collection.

OMB No.

Abstract:

The 21st Century Cures Act (Cures Act) aimed to advance the exchange of electronic health information by promoting patient access through standardized application programming interfaces (APIs). Digital health companies develop apps and health IT

tools that enable human interaction with APIs to exchange electronic health information. Prior studies indicate widespread adoption of standardized APIs for interoperability with electronic health records (EHRs). Ongoing assessment of these technologies is crucial to examining the impacts of the Cures Act's health IT provisions and is critical to informing the Assistant Secretary for Technology Policy/Office of the National Coordinator for Health IT's (ASTP/ONC's) policy efforts. With ASTP/ONC's support, the University of California, San Francisco (UCSF) conducted a 2022 survey of digital health companies assessing implementation of and experiences with healthcare APIs; findings from this survey work are published in the Journal of the American Medical Informatics Association. ASTP/ONC finds it essential to continue efforts to survey digital health companies to assess ASTP/ONC's implementation of statutorily mandated information blocking (42 U.S.C. 300jj–52) and APIs “without special effort” policies (42 U.S.C. 300jj–11) under the Cures Act. Information gathered from this effort will help inform ongoing ASTP/ONC efforts to help nurture an ecosystem of innovation and transparency in health care.

ANNUALIZED BURDEN HOUR TABLE

Forms (if necessary)	Respondents (if necessary)	Number of respondents	Number of responses per respondents	Average burden per response (minutes)	Total burden hours
Total	U.S. based Digital Health Companies	282	1	60	282
Total	282

Susan R. Little,

Department Information Collection Clearance
Officer, Paperwork Reduction Act Program,
Department of Health and Human Services.

[FR Doc. 2025–09004 Filed 5–20–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Notice of Closed Meetings

Pursuant to section 1009 of the Federal Advisory Committee Act, as amended, notice is hereby given of the following meetings.

The meetings 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: Center for Scientific Review Special Emphasis Panel; Imaging and Bioengineering Technology for Visual Systems (IBV).

Date: June 17–18, 2025.

Time: 8:30 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

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

Meeting Format: Virtual Meeting.

Contact Person: Susan Gillmor, Ph.D., Scientific Review Officer, National Institutes of Health, Center for Scientific Review, 6701 Rockledge Drive, Bethesda, MD 20892, 240–762–3076, susan.gillmor@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; RFA Panel: HIV and Substance Use.

Date: June 17, 2025.

Time: 9:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

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

Meeting Format: Virtual Meeting.

Contact Person: Kaitlyn N. Hardell, MPH, Ph.D., Scientific Review Officer, SRB, NIA, Scientific Review Branch, 5601 Fishers Lane, Suite 8B, Rockville, MD 20892, (301) 594–7945, kaitlyn.hardell@nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Program Projects: Cancer Biology.

Date: June 17–18, 2025.

Time: 9:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

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

Meeting Format: Virtual Meeting.

Contact Person: E. Tian, Ph.D., Scientific Review Officer, Research Program Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W618, ROCKVILLE, MD 20850, (240) 276–7246, tiane@mail.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; P50 SPORE's in Human Cancer.

Date: June 17–18, 2025.

Time: 9:30 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

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

Meeting Format: Virtual Meeting.

Contact Person: John Paul Cairns, Ph.D., Scientific Review Officer, Research Programs Review Branch, Division of Extramural Activities, 9609 Medical Center Drive, Room 7W244, National Cancer Institute, NIH, Bethesda, MD 20892, 240–276–5415, paul.cairns@nih.gov.

Name of Committee: Biological Chemistry and Macromolecular Biophysics Integrated Review Group, Macromolecular Structure and Function B Study Section.

Date: June 17–18, 2025.

Time: 9:30 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

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

Meeting Format: Virtual Meeting.

Contact Person: Alexei A Yeliseev, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 443–0552, yeliseeva@mail.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Special Topics: Institutional Training and Education Study Section.

Date: June 17–18, 2025.

Time: 10:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

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

Meeting Format: Virtual Meeting.

Contact Person: Adriana Stoica, Ph.D., Scientific Review Officer, Resources and Training Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, 7W234, Bethesda, MD 20892–9750, 240–276–6368, Stoica2@mail.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Bidirectional Influences Between Adolescent Social Media Use and Mental Health.

Date: June 18, 2025.

Time: 12:00 p.m. to 4:30 p.m.

Agenda: To review and evaluate grant applications.

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

Meeting Format: Virtual Meeting.

Contact Person: Regina T. Dolan-Sewell, Ph.D., Scientific Review Officer, Division of Extramural Activities, National Institute of Mental Health, NIH, Neuroscience Center, 6001 Executive Boulevard, Bethesda, MD 20852, (240) 796–6785, regina.dolan-sewell@nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: May 15, 2025.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2025–09074 Filed 5–20–25; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Center for Scientific Review Special Emphasis Panel Small Business Activities: Cardiovascular and Hematology, June 19, 2025, 09:00 a.m. to June 20, 2025, 06:00 p.m., National