TABLE 1—ESTIMATED ANNUAL RECORDKEEPING BURDEN [1]—Continued

21 CFR Section; [2] activity	Number of record-keepers	Number of records per recordkeeper [3]	Total annual records	Average burden per recordkeeping [4]	Total hours
123.12(a)(2); Prepare new written verification procedures to verify compliance of imports.	41	1	41	4	164
Total					1,930,264

[1] There are no capital costs or operating and maintenance costs associated with this collection of information.
[2] These estimates include the information collection requirements in the following sections:
§ 123.16—Smoked Fish—process controls (see § 123.6(b));
§ 123.28(a)—Source Controls—molluscan shellfish (see § 123.6(b));
§ 123.28(c) and (d)—Records—molluscan shellfish (see § 123.6(c)(7).
[3] Based on an estimated 280 working days per year.
[4] Estimated average time per 8-hour workday unless one-time response.

Based on a review of the information collection since its last OMB approval, we have made no adjustments to our burden estimate.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2026–03311 Filed 2–18–26; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Center for Scientific Review; 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: Center for Scientific Review Special Emphasis Panel; Endocrinology and Metabolism Topics.

Date: March 11, 2026.

Time: 1:00 p.m. to 3:00 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: Elaine Sierra-Rivera, Ph.D., IRG Chief, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6182, Bethesda, MD 20892, (301) 435–2514, *riverase@csr.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: February 13, 2026.

Bruce A. George,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2026–03247 Filed 2–18–26; 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; Information Program on Clinical Trials: Maintaining a Registry and Results Databank (National Library of Medicine)

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.* 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: Vivian Le, Office of Administration, National Library of Medicine, 8600 Rockville Pike, Building 38A, 4N401Q5, Bethesda, Maryland 20894 or call non-toll-free number 301–827–6328 or Email your request, including your address to: *vivian.le@nih.gov.*

SUPPLEMENTARY INFORMATION: This proposed information collection was previously published in the **Federal Register** on November 26, 2025, page 54340 (90 FR 54340) 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 National Library of Medicine (NLM), 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: Information Program on Clinical Trials: Maintaining a Registry and Results Databank, 0925–0586, Expiration Date 03/31/2026, Revision, National Library of Medicine (NLM), National Institutes of Health (NIH).

Need and Use of Information Collection: The National Institutes of Health operates *ClinicalTrials.gov*, which was established as a clinical trial registry under section 113 of the Food and Drug Administration Modernization

Act of 1997 (Pub. L. 105–115) and was expanded to include a results data bank by Title VIII of the Food and Drug Administration Amendments Act of 2007 (FDAAA) and by the Clinical Trials Registration and Results Information Submission regulations at 42 CFR part 11. *ClinicalTrials.gov* collects registration and results information for clinical trials and other types of clinical studies (*e.g.,* observational studies and patient registries) with the objectives of enhancing patient enrollment and providing a mechanism for tracking subsequent progress of clinical studies to the benefit of public health. It is widely used by patients, physicians, and medical researchers; in particular those involved in clinical research. While many clinical studies are registered and results information submitted voluntarily, 42 CFR part 11 requires the registration and submission of results information for certain applicable clinical trials of drug, biological, and device products whether or not they are approved, licensed, or cleared by the Food and Drug Administration.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 1,411,181.

ESTIMATED ANNUALIZED BURDEN HOURS

Submission type	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hour
Registration—Attachment 2				
Initial	7,400	1	8	59,200
Updates	7,400	8	2	118,400
Triggered, voluntary	182	1	8	1,456
Initial, non-regulated, NIH Policy	1,200	1	8	9,600
Updates, non-regulated, NIH Policy	1,200	8	2	19,200
Initial, voluntary and non-regulated	23,130	1	8	185,040
Updates, voluntary and non-regulated	23,130	8	2	370,080
Results Information Submission—Attachment 5				
Initial	7,400	1	40	296,000
Updates	7,400	2	10	148,000
Triggered, voluntary—also attachment 2	61	1	45	2,745
Initial, non-regulated, NIH Policy	1,200	1	40	48,000
Updates, non-regulated, NIH Policy	1,200	2	10	24,000
Initial, voluntary and non-regulated	2,100	1	40	84,000
Updates, voluntary and non-regulated	2,100	2	10	42,000
Other				
Certification to delay results—attachment 6	5,150	1	30/60	2,575
Extension requests and Appeals—attachment 7	175	1	2	350
Initial, expanded access—attachment 3	213	1	2	426
Updates, expanded access—attachment 3	213	2	15/60	107
Waiver requests and appeals—attachment 10	1	1	2	2
Total		323,878		1,411,181

Vivian K. Le,

Project Clearance Liaison, National Library of Medicine, National Institutes of Health.

[FR Doc. 2026–03222 Filed 2–18–26; 8:45 am]

BILLING CODE 4140–01–P

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: Integrative, Functional and Cognitive Neuroscience Integrated Review Group; Behavioral Neuroendocrinology, Neuroimmunology, Rhythms, and Sleep Study Section.

Date: March 23–24, 2026.

Time: 9:00 a.m. to 7:00 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: Simon Peter Peron, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 1009K, Bethesda, MD 20892, (301) 594–6236, *peronsp@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Fellowships: Brain Disorders and Related Neurosciences.

Date: March 23, 2026.

Time: 9:30 a.m. to 7:00 p.m.

Agenda: To review and evaluate grant applications.

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

Meeting Format: Virtual Meeting.

Contact Person: Vanessa S. Boyce, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4185, MSC 7850, Bethesda, MD 20892, (301) 402–3726, *boycevs@csr.nih.gov*.