

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 |
|--|-----------------------|------------------------------------|--|--------------------------|
| <b>Registration—Attachment 2</b>                   |                       |                                    |  |                          |
| 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                  |
| <b>Results Information Submission—Attachment 5</b> |                       |                                    |  |                          |
| 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                   |
| <b>Other</b>                                       |                       |                                    |  |                          |
| 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                        |
| <b>Total .....</b>                                 |                       | <b>323,878</b>                     |  | <b>1,411,181</b>         |

**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](mailto: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](mailto:boycevs@csr.nih.gov).

*Name of Committee:* Oncology 1—Basic Translational Integrated Review Group; Cancer Cell Biology Study Section.

*Date:* March 23–24, 2026.

*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:* Alyssa Diane Gregory, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 480–4906, [alyssa.gregory@nih.gov](mailto:alyssa.gregory@nih.gov).

*Name of Committee:* Cell Biology Integrated Review Group; Maximizing Investigators’ Research Award—D Study Section.

*Date:* March 23–24, 2026.

*Time:* 10:00 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:* Anne Marie Strohecker, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (202) 924–4186, [stroheckeram@csr.nih.gov](mailto:stroheckeram@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.

**Margaret N. Vardanian,**

*Program Analyst, Office of Federal Advisory Committee Policy.*

[FR Doc. 2026–03221 Filed 2–18–26; 8:45 am]

**BILLING CODE 4140–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**Proposed Collection; 60-Day Comment Request; Chimpanzee Research Use Form (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 Laboratory Animal Welfare (OLAW), Office of the Director (OD), 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 by April 20, 2026.

**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: Jane J. Na, Director, Division of Assurances, Office of Laboratory Animal Welfare, NIH, or call non-toll-free number (301) 496–7163 or email your request, including your address to: [olawdoa@mail.nih.gov](mailto:olawdoa@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 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:* Chimpanzee Research Use Form, 0925–0705, exp., date 7/31/2026, EXTENSION, Office of Laboratory Animal Welfare (OLAW), Office of the Director (OD), National Institutes of Health (NIH).

*Need and Use of Information Collection:* The purpose of this form is to continue to obtain information needed by the NIH to assess whether the proposed research satisfies the agency’s policy for permitting only noninvasive research involving chimpanzees. The NIH will consider the information submitted through this form prior to the agency making funding decisions or otherwise allowing the research to begin. Completion of this form is a mandatory step toward continuing to receive NIH support or approval for noninvasive research involving chimpanzees. The NIH does not fund any research involving chimpanzees proposed in new or other competing projects (renewals or revisions) unless the research is consistent with the definition of “noninvasive research,” as described in the “Standards of Care for Chimpanzees Held in the Federally Supported Chimpanzee Sanctuary System” (42 CFR part 9). Also see NOT–OD–16–095 at <https://grants.nih.gov/grants/guide/notice-files/NOT-OD-16-095.html> and 81 FR 6873. OMB approval is requested for three years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 15.

**ESTIMATED ANNUALIZED BURDEN HOURS**

| Form name                          | Type of respondent       | Number of respondents | Number of responses per respondent | Average burden per response (in hours) | Total annual burden hours |
|------------------------------------|--------------------------|-----------------------|------------------------------------|--|---------------------------|
| Chimpanzee Research Use Form ..... | Research Community ..... | 30                    | 1                                  | 30/60                                  | 15                        |
| Total .....                        | .....                    | .....                 | 30                                 | .....                                  | 15                        |