Neuropathophysiology of Decision Making and Chemobrain.

*Date:* October 26, 2022.

*Time:* 1:00 p.m. to 7:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

*Contact Person:* Aleksey Gregory Kazantsev, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5201, Bethesda, MD 20817, (301) 435–1042, *aleksey.kazantsev@nih.gov*.

*Name of Committee:* Endocrinology, Metabolism, Nutrition and Reproductive Sciences Integrated Review Group; Pathophysiology of Obesity and Metabolic Disease Study Section.

*Date:* October 27–28, 2022.

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

*Agenda:* To review and evaluate grant applications.

*Place:* Washington Marriott Georgetown, 1221 22nd Street NW, Washington, DC 20037.

*Contact Person:* Raul Rojas, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6185, Bethesda, MD 20892, (301)451–6319, *rojasr@mail.nih.gov*.

*Name of Committee:* Oncology 1—Basic Translational Integrated Review Group; Biochemical and Cellular Oncogenesis Study Section.

*Date:* October 27–28, 2022.

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

*Agenda:* To review and evaluate grant applications.

*Place:* Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road NW, Washington, DC 20015.

*Contact Person:* Jian Cao, MD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 827–5902, *caojn@csr.nih.gov*.

*Name of Committee:* Cardiovascular and Respiratory Sciences Integrated Review Group; Cardiovascular Differentiation and Development Study Section.

*Date:* October 27, 2022.

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

*Agenda:* To review and evaluate grant applications.

*Place:* Westin Grand, 2350 M Street NW, Washington, DC 20037.

*Contact Person:* Sara Ahlgren, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, RM 4136, Bethesda, MD 20892, (301) 435–0904, *sara.ahlgren@nih.gov*.

*Name of Committee:* Genes, Genomes, and Genetics Integrated Review Group; Prokaryotic Cell and Molecular Biology Study Section.

*Date:* October 27–28, 2022.

*Time:* 8:00 a.m. to 8:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Bethesdan Hotel, Tapestry Collection by Hilton, 8120 Wisconsin Ave., Bethesda, MD 20814.

*Contact Person:* Rebecca C Burgess, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 480–8034, *rebecca.burgess@nih.gov*.

*Name of Committee:* Biology of Development and Aging Integrated Review Group; Radiation Therapeutics and Biology Study Section.

*Date:* October 27–28, 2022.

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

*Agenda:* To review and evaluate grant applications.

*Place:* Hyatt Regency Bethesda, 1 Bethesda Metro Center, Bethesda, MD 20814.

*Contact Person:* Bo Hong, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6194, MSC 7804, Bethesda, MD 20892, 301–996–6208, *hongb@csr.nih.gov*.

*Name of Committee:* Infectious Diseases and Immunology A Integrated Review Group; Cellular and Molecular Immunology—B Study Section.

*Date:* October 27–28, 2022.

*Time:* 8:00 a.m. to 7:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* Residence Inn Bethesda, 7335 Wisconsin Avenue, Bethesda, MD 20814.

*Contact Person:* Liying Guo, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4198, MSC 7812, Bethesda, MD 20892, (301) 827–7728, *lguo@mail.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: September 28, 2022.

**Victoria E. Townsend,**

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

[FR Doc. 2022–21429 Filed 10–3–22; 8:45 am]

**BILLING CODE 4140–01–P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

### National Center for Complementary & Integrative Health; Notice of Closed Meeting

Pursuant to section 10(d) 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:* National Center for Complementary and Integrative Health Special Emphasis Panel; NCCIH Training and Education Review Panel (CT).

*Date:* November 9–10, 2022.

*Time:* 10:00 a.m. to 4:00 p.m.

*Agenda:* To review and evaluate grant applications.

*Place:* National Center for Complementary and Integrative, Democracy II, 6707 Democracy Blvd., Bethesda, MD 20892 (Virtual Meeting).

*Contact Person:* Jessica Marie McKlveen, Ph.D., Scientific Review Officer, Office of Scientific Review, Division of Extramural Activities, NCCIH, NIH, 6707 Democracy Boulevard, Suite 401, Bethesda, MD 20892–547, *jessica.mcklveen@nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.213, Research and Training in Complementary and Alternative Medicine, National Institutes of Health, HHS)

Dated: September 28, 2022.

**Victoria E. Townsend,**

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

[FR Doc. 2022–21426 Filed 10–3–22; 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; PHS Research Performance Progress Report and Other Post-Award Reporting (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 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: Ms. Mikia P. Currie, Program Analyst, 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: *ProjectClearanceBranch@mail.nih.gov*.

**SUPPLEMENTARY INFORMATION:** This proposed information collection was previously published in the **Federal Register** on June 8, 2022, pages 34888/34889 (87 FR 34888) 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 Office of the Director, Office of Extramural Research (OER), National Institutes of Health (NIH), 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:* Public Health Service (PHS) Research Performance Progress Report and Other Post-award Reporting –0925–0002 expiration date–09/30/2024, REVISION, Office of the Director, National Institutes of Health (NIH).

*Need and Use of Information Collection:* This collection under 0925–0002 expiration date 09/30/2024 is being revised to update the non-competing progress report collections for the implementation of the final National Institutes of Health (NIH) Policy for Data Management and Sharing (DMS Policy) to promote the management and sharing of scientific data generated from NIH-funded or conducted research. Starting in January 2023, NIH will require applicants and recipients to submit and address Data Management and Sharing (DMS) Plans within the SF424 Research and Related (R&R) application and the Research Performance Progress Report (RPPR) in accordance with the DMS Policy. The application and progress report forms will be updated to align with this requirement. NIH is also introducing a new, optional Data Management and Sharing (DMS) Plan format page that applicants may use to develop their DMS Plan. This collection will also update the PHS 2271 Statement of Appointment for trainees appointed institutional training awards to report on childcare cost support received. This collection is also updated to remove the iEdison instrument in accordance with the transfer of the iEdison system to the National Institute of Standards and Technology (NIST) under the Department of Commerce (DOC). NIST will maintain OMB clearance under 0693–0090.

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

ESTIMATED ANNUALIZED BURDEN HOURS

| Information collection forms | Number of respondents | Number of responses per respondent | Average burden per response (in hours) | Total annual burden hours |
|---|---|---|---|---|
| **REPORTING** | | | | |
| PHS 416–7 | 12,580 | 1 | 30/60 | 6,290 |
| PHS 6031–1 | 1,778 | 1 | 20/60 | 593 |
| PHS 568 | 11,180 | 1 | 5/60 | 932 |
| PHS 2271 | 22,035 | 1 | 15/60 | 5,509 |
| PHS 2590 | 243 | 1 | 18 | 4,374 |
| RPPR – | | | | |
| Core Data | 32,098 | 1 | 8 | 256,784 |
| Biosketch (Part of RPPR) | 2,544 | 1 | 2 | 5,088 |
| Data Tables (Part of RPPR) | 758 | 1 | 4 | 3,032 |
| Trainee Diversity Report (Part of RPPR) | 480 | 1 | 15/60 | 120 |
| PHS Human Subjects and Clinical Trial Information (Part of RPPR, includes inclusion enrollment report) | 6,420 | 1 | 4 | 25,680 |
| Publication Reporting | 97,023 | 3 | 5/60 | 24,256 |
| Final RPPR—Core Data | 18,000 | 1 | 10 | 180,000 |
| Data Tables (Part of Final RPPR) | 758 | 1 | 4 | 3,032 |
| Trainee Diversity Report (Part of Final RPPR) | 480 | 1 | 15/60 | 120 |
| PHS Human Subjects and Clinical Trial Information (Part of Final RPPR, includes inclusion/enrollment) | 3,600 | 1 | 4 | 14,400 |
| PHS 3734 | 479 | 1 | 30/60 | 240 |
| Data Management and Sharing Plan (Part of RPPR) | 15,649 | 1 | 2 | 31,298 |
| Data Management and Sharing Plan (Part of Final RPPR) | 8,621 | 1 | 2 | 17,242 |
| Reporting Burden Total | ..... | ..... | ..... | 578,990 |
| **RECORDKEEPING** | | | | |
| SBIR/STTR Life Cycle Certification | 1,500 | 1 | 15/60 | 375 |
| Total | 236,226 | 430,272 | ..... | 579,365 |

Dated: September 27, 2022.

**Tara A. Schwetz,**

*Acting Principal Deputy Director, National Institutes of Health.*

[FR Doc. 2022–21547 Filed 10–3–22; 8:45 am]

**BILLING CODE 4140–01–P**

## DEPARTMENT OF HOMELAND SECURITY

### U.S. Customs and Border Protection

**[1651–0077]**

### Customs-Trade Partnership Against Terrorism (CTPAT) and CTPAT Trade Compliance Program

**AGENCY:** U.S. Customs and Border Protection (CBP), Department of Homeland Security.

**ACTION:** 30-Day notice and request for comments; revision of an existing collection of information.

**SUMMARY:** The Department of Homeland Security, U.S. Customs and Border Protection will be submitting the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995 (PRA). The information collection is published in the **Federal Register** to obtain comments from the public and affected agencies.

**DATES:** Comments are encouraged and must be submitted no later than November 3, 2022 to be assured of consideration.

**ADDRESSES:** Written comments and/or suggestions regarding the item(s) contained in this notice 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:** Requests for additional PRA information should be directed to Seth Renkema, Chief, Economic Impact Analysis Branch, U.S. Customs and Border Protection, Office of Trade, Regulations and Rulings, 90 K Street NE, 10th Floor, Washington, DC 20229–1177, telephone number 202–325–0056 or via email *CBP_PRA@cbp.dhs.gov.* Please note that the contact information provided here is solely for questions regarding this notice. Individuals seeking information about other CBP programs should contact the CBP National Customer Service Center at 877–227–5511, (TTY) 1–800–877–8339, or CBP website at *https://www.cbp.gov/.*

**SUPPLEMENTARY INFORMATION:** CBP invites the general public and other Federal agencies to comment on the proposed and/or continuing information collections pursuant to the Paperwork Reduction Act of 1995 (44 U.S.C. 3501 *et seq.*). This proposed information collection was previously published in the **Federal Register** (87 FR 12473) on March 04, 2022, allowing for a 60-day comment period. This notice allows for an additional 30 days for public comments. This process is conducted in accordance with 5 CFR 1320.8. Written comments and suggestions from the public and affected agencies should address one or more of the following four points: (1) whether the proposed collection of information is necessary for the proper performance of the functions 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) suggestions to enhance the quality, utility, and clarity of the information to be collected; and (4) suggestions to minimize the burden of the collection of information on those who are to respond, including through the use of appropriate automated, electronic, mechanical, or other technological collection techniques or other forms of information technology, *e.g.,* permitting electronic submission of responses. The comments that are submitted will be summarized and included in the request for approval. All comments will become a matter of public record.

#### Overview of This Information Collection

*Title:* Customs-Trade Partnership Against Terrorism (CTPAT) and CTPAT Trade Compliance Program.

*OMB Number:* 1651–0077.

*Form Number:* N/A.

*Current Actions:* Revision of an existing information collection.

*Type of Review:* Revision.

*Affected Public:* Businesses.

*Abstract:* The CTPAT Program comprises of two different program divisions, CTPAT Security and CTPAT Trade Compliance. The CTPAT Security program is designed to safeguard the world's trade industry from terrorists and smugglers by prescreening its participants. The CTPAT Security program applies to United States and nonresident Canadian importers, United States exporters, customs brokers, consolidators, ports and terminal operators, carriers of cargo in air, sea and land, third party logistics providers, Mexican long haul highway carriers, and Canadian and Mexican manufacturers. The Trade Compliance program division is only available for U.S. and nonresident Canadian importers.

The CTPAT Program application requests an applicant's contact and business information, including the number of company employees, the number of years in business, and a list of company officers. CBP is adding the following data elements for all CTPAT partners to improve the screening of companies. This will ensure that CBP is confident that companies in the program are low risk:

- Date of Birth (DOB)
- Country of Birth
- Country of Citizenship
- Travel Document number (*e.g.,* visa or passport number)
- Immigration status information (*e.g.,* Alien Registration Number, Naturalization number)
- Driver's license information (*e.g.,* state and country of issuance, number, date of issuance/expiration)
- Social Security Number
- Trusted Traveler membership type and number (*e.g.,* FAST/NEXUS/SENTRI/Global Entry ID)
- Registro Federal de Contribuventes (RFC) Persona Fisica (needed for Mexican Foreign Manufacturers, Highway Carriers, and Long-Haul Carriers Only)

This collection of information is authorized by the SAFE Port Act (Pub. L. 109–347).

The CTPAT Trade Compliance program is an optional component of the CTPAT program and adds trade compliance aspects to the supply chain security aspects of the CTPAT Security program. The CTPAT Security program is a prerequisite to applying to the CTPAT Trade Compliance program. Current CTPAT importers are given the opportunity to receive additional benefits in exchange for a commitment to assume responsibility for monitoring their own compliance by applying to the CTPAT Trade Compliance program. After a company has completed the security aspects of the CTPAT Security program and is in good standing, it may opt to apply to the CTPAT Trade Compliance component. The CTPAT Trade Compliance program strengthens security by leveraging the CTPAT supply chain requirements, identifying low-risk trade entities for supply chain security, and increasing the overall efficiency of trade by segmenting risk and processing by account.

The CTPAT Trade Compliance program is open to U.S. and non-resident Canadian importers that have