

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:* NCI Genomic Data Commons (GDC) Data

Submission Request Form, 0925–0752, Expiration Date 03/31/2023, EXTENSION, National Cancer Institute (NCI), National Institutes of Health (NIH).

*Need and Use of Information Collection:* The purpose of the NCI Genomic Data Commons (GDC) Data Submission Request Form is to provide a vehicle for investigators to request the submission of their cancer genomic data into the GDC in support of data sharing. The purpose is also to provide a mechanism for the GDC Data Submission Review Committee to review and assess the data submission request for applicability to the GDC mission. The scope of the form involves

obtaining information from investigators that: (1) would like to submit data about their study into the GDC, (2) are affiliated with studies that adhere to GDC data submission conditions. The benefits of the collection are that it provides the needed information for investigators to understand the types of studies and data that the GDC supports and that it provides a standard mechanism for the GDC to assess incoming data submission requests.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 50 hours.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hour
Investigator .....	200	1	15/60	50
Total .....	.....	200	.....	50

Dated: November 2, 2022.

**Diane Kreinbrink,**

*Project Clearance Liaison, National Cancer Institute, National Institutes of Health.*

[FR Doc. 2022–24186 Filed 11–4–22; 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 10(d) 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; RFA: Countermeasures Against Chemical Threats Exploratory/Developmental Projects.

*Date:* December 5, 2022.

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

*Agenda:* To review and evaluate grant applications.

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

*Contact Person:* Jodie Michelle Fleming, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 812R, Bethesda, MD 20892, (301) 867–5309, [flemingjm@csr.nih.gov](mailto:flemingjm@csr.nih.gov).

*Name of Committee:* Center for Scientific Review Special Emphasis Panel; Small Business: Microbial Vaccine Development.

*Date:* December 5–6, 2022.

*Time:* 10:00 a.m. to 8:30 p.m.

*Agenda:* To review and evaluate grant applications.

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

*Contact Person:* Subhamoy Pal, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (301) 594–0926, [subhamoy.pal@nih.gov](mailto:subhamoy.pal@nih.gov).

*Name of Committee:* Center for Scientific Review Special Emphasis Panel; Fellowships: Oncology.

*Date:* December 6–7, 2022.

*Time:* 10:00 a.m. to 8:00 p.m.

*Agenda:* To review and evaluate grant applications.

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

*Contact Person:* Nywana Sizemore, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6189, MSC 7804, Bethesda, MD 20892, 301–408–9916, [sizemoren@csr.nih.gov](mailto:sizemoren@csr.nih.gov).

*Name of Committee:* Center for Scientific Review Special Emphasis Panel; Epigenomics of Neurodevelopment.

*Date:* December 6, 2022.

*Time:* 2:00 p.m. to 3:00 p.m.

*Agenda:* To review and evaluate grant applications.

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

*Contact Person:* Mary G Schueler, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5214, MSC 7846, Bethesda, MD 20892, 301–915–6301, [marygs@csr.nih.gov](mailto:marygs@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: November 2, 2022.

**David W. Freeman,**

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

[FR Doc. 2022–24204 Filed 11–4–22; 8:45 am]

**BILLING CODE 4140–01–P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**National Institutes of Health**

**National Heart, Lung, and Blood Institute; Amended Notice of Meeting**

Notice is hereby given of a change in the meeting of the Sleep Disorders Research Advisory Board, December 01, 2022, 12 p.m. to December 01, 2022, 4 p.m., National Institutes of Health, Rockledge II, 6701 Rockledge Drive, Bethesda, MD, 20892 which was

published in the **Federal Register** on November 01, 2022, 87 FR 65786.

Meeting is being amended to change the meeting time from 12 p.m. to 4 p.m. to 12 p.m. to 5 p.m. Also to change the agenda from, “The purpose of this meeting is to update the Advisory Board and public stakeholders on the research agenda across NIH for the upcoming fiscal year, and the activities of professional societies.” to “The purpose of this meeting is to discuss with the Advisory Board timely research opportunities in sleep and circadian biology. Updates on the research agenda across the NIH for the upcoming fiscal year and the activities of professional societies will be provided to all stakeholders.” The meeting is open to the public.

Dated: November 2, 2022.

**David W. Freeman,**

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

[FR Doc. 2022-24203 Filed 11-4-22; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

[Docket No. FR-6331-N-08]

### Public Interest De Minimis, Small Grants, and Minor Components Waiver of Build America, Buy America Provisions as Applied to Certain Recipients of HUD Federal Financial Assistance

**AGENCY:** Office of the Secretary, U.S. Department of Housing and Urban Development (HUD).

**ACTION:** Notice.

**SUMMARY:** In accordance with the Build America, Buy America Act (“BABA” or “the Act”) this notice advises that HUD is proposing a departmentwide public interest *de minimis*, Small Grants, and Minor Components waiver to the Buy America Domestic Content Procurement Preference (“Buy America Preference,” or “BAP”) as applied to the iron, steel, manufactured products, and construction materials requirement of the Act for recipients of Federal Financial Assistance. For the purposes of this proposed waiver, HUD is proposing to waive the application of the BAP for infrastructure projects whose total cost is an amount equal to or less than the Simplified acquisition threshold, which is currently \$250,000. HUD is also proposing to waive the application of the BAP for all Small Grants of Federal Financial Assistance that are equal to or below the Simplified acquisition threshold, which is

currently \$250,000. Additionally, HUD is proposing to waive the application of the BAP for Minor Components of an infrastructure project, such that a cumulative total of no more than a total of 5 percent of the total cost of the iron, steel, manufactured products, and construction materials used in and incorporated into the infrastructure project, up to a maximum of \$1 million. In accordance with the Act, HUD has found that such proposed De Minimis, Small Grants and Minor Components waivers are in the public interest. The waiver will assist HUD and its grantees and funding recipients in preventing immediate delays to critically important projects that serve to ensuring the safety and health of HUD constituents and continuing to provide economic opportunity through housing and community development projects. Moreover, this waiver will assist HUD in working to strengthen the housing market to bolster the economy and protect consumers, meet the need for quality affordable rental homes, utilize housing as a platform for improving quality of life, and build inclusive and sustainable communities free from discrimination.

**DATES:** HUD published this proposed waiver on its website on October 31, 2022. Comments on the proposed waiver set out in this document are due on or before November 15, 2022.

**ADDRESSES:** Interested persons are invited to submit comments on this proposed general applicability waiver. Copies of all comments submitted are available for inspection and downloading at [www.regulations.gov](http://www.regulations.gov).

To receive consideration as public comments, comments must be submitted through one of two methods, specified below. All submissions must refer to the above docket number and title.

1. *Electronic Submission of Comments.* Interested persons may submit comments electronically through the Federal eRulemaking Portal at [www.regulations.gov](http://www.regulations.gov).

HUD strongly encourages commenters to submit comments electronically. Electronic submission of comments allows the commenter maximum time to prepare and submit a comment, ensures timely receipt by HUD, and enables HUD to make them immediately available to the public. Comments submitted electronically through the [www.regulations.gov](http://www.regulations.gov) website can be viewed by other commenters and interested members of the public. Commenters should follow the instructions provided on that site to submit comments electronically.

2. *Submission of Comments by Mail.* Comments may be submitted by mail to the Regulations Division, Office of General Counsel, Department of Housing and Urban Development, 451 7th Street SW, Room 10276, Washington, DC 20410-0500.

No Facsimile Comments. Facsimile (FAX) comments will not be accepted.

3. *Public Inspection of Comments.* All properly submitted comments and communications submitted to HUD will be available for public inspection and copying between 8:00 a.m. and 5:00 p.m. weekdays at the above address. Due to security measures at the HUD Headquarters building, an advance appointment to review the submissions must be scheduled by calling the Regulations Division at (202) 708-3055 (this is not a toll-free number).

#### FOR FURTHER INFORMATION CONTACT:

Joseph Carlile, Department of Housing and Urban Development, 451 Seventh Street SW, Room 10226, Washington, DC 20410-5000, at (202) 402-7082 (this is not a toll-free number). HUD welcomes and is prepared to receive calls from individuals who are deaf or hard of hearing, as well as individuals with speech and communication disabilities. To learn more about how to make an accessible telephone call, please visit <https://www.fcc.gov/consumers/guides/telecommunications-relay-service-trs>. HUD encourages submission of questions about this document be sent to [BuildAmericaBuyAmerica@hud.gov](mailto:BuildAmericaBuyAmerica@hud.gov).

#### SUPPLEMENTARY INFORMATION:

##### I. Build America, Buy America

The Build America, Buy America Act (“BABA” or “the Act”) was enacted on November 15, 2021, as part of the Infrastructure Investment and Jobs Act (IIJA). Public Law 117-58. The Act establishes a domestic content procurement preference, the BAP, for Federal infrastructure programs. Section 70914(a) of the Act establishes that no later than 180 days after the date of enactment, HUD must ensure that none of the funds made available for infrastructure projects may be obligated by the Department unless it has taken steps to ensure that the iron, steel, manufactured products, and construction materials used in a project are produced in the United States. In section 70912, the Act further defines a project to include “the construction, alteration, maintenance, or repair of infrastructure in the United States” and includes within the definition of infrastructure those items traditionally included along with buildings and real property. Thus, starting May 14, 2022,