

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)

Dated: March 10, 2022.

**Melanie J. Pantoja,**

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

[FR Doc. 2022-05464 Filed 3-14-22; 8:45 am]

**BILLING CODE 4140-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### National Institutes of Health

#### National Cancer Institute; Notice of Meeting

Pursuant to section 10(a) of the Federal Advisory Committee Act, as amended, notice is hereby given of a meeting of the National Cancer Institute Board of Scientific Advisors.

The meeting will be held as a virtual meeting and is open to the public. Individuals who plan to view the virtual meeting and need special assistance or other reasonable accommodations to view the meeting, should notify the Contact Person listed below in advance of the meeting. The meeting will be videocast and can be accessed from the NIH Videocasting and Podcasting website (<http://videocast.nih.gov/>).

*Name of Committee:* National Cancer Institute Board of Scientific Advisors.

*Date:* March 28, 2022.

*Time:* 1:00 p.m. to 5:30 p.m.

*Agenda:* Director's Report; RFA, RFP, and PAR Concept Reviews; and Scientific Presentations.

*Place:* National Cancer Institute—Shady Grove, 9609 Medical Center Drive, Rockville, MD 20850 (Virtual Meeting).

*Date:* March 29, 2022.

*Time:* 1:00 p.m. to 5:30 p.m.

*Agenda:* RFA, RFP, and PAR Concept Reviews; and Scientific Presentations.

*Place:* National Cancer Institute—Shady Grove, 9609 Medical Center Drive, Rockville, MD 20852 (Virtual Meeting).

*Contact Person:* Paulette S. Gray, Ph.D., Director, Division of Extramural Activities, National Cancer Institute—Shady Grove, National Institutes of Health, 9609 Medical Center, Drive, 7th Floor, Room 7W444, Bethesda, MD 20892, 240-276-6340, [grayp@mail.nih.gov](mailto:grayp@mail.nih.gov).

Any interested person may file written comments with the committee by forwarding the statement to the Contact Person listed on this notice. The statement should include the name, address, telephone number and when

applicable, the business or professional affiliation of the interested person.

Information is also available on the Institute's/Center's home page: BSA: <http://deainfo.nci.nih.gov/advisory/bsa/bsa.htm>, where an agenda and any additional information for the meeting will be posted when available.

This notice is being published less than 15 days prior to the meeting due to scheduling difficulties.

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)

Dated: March 10, 2022.

**Melanie J. Pantoja,**

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

[FR Doc. 2022-05467 Filed 3-14-22; 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; NIH COVID-19 Vaccination Status Form Extension

**AGENCY:** National Institutes of Health, Department of Health and Human Services (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](http://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: Dr. Jessica McCormick-Ell, Ph.D., SM (NRCM), CBSP, RBP, NIH/ORS/SR/DOHS, Bldg. 13/3W80, Bethesda, MD, 20892-5760 or call non-toll-free number (301) 496-0590 or email your request, including your address to: [jessica.mccormick-ell@nih.gov](mailto:jessica.mccormick-ell@nih.gov).

**SUPPLEMENTARY INFORMATION:** This proposed information collection was previously published in the **Federal Register** on November 22, 2021 (86 FR 66319), 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 Research Services, 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:* NIH COVID-19 Vaccination Status Form, REVISION, 0925-0771, exp., 3/31/2022, Office of Research Services (ORS), National Institutes of Health (NIH).

*Need and Use of Information Collection:* This revision request includes two new forms as NIH is implementing a building entry protocol where all people, Federal Government employees, contractors, patients, and visitors, will answer questions about COVID exposure every day. The NIH COVID-19 Vaccination Status Form will continue to ensure the safety of the Federal workplace consistent with Executive Order 14042, Ensuring Adequate COVID Safety Protocols for Federal Contractors, Executive Order 14043, Requiring Coronavirus Disease 2019 Vaccination for Federal Employees, the COVID-19 Workplace Safety: Agency Model Safety Principles established by the Safer Federal Workforce Task Force, and guidance from the Centers for Disease Control and Prevention (CDC) and the Occupational Safety and Health Administration (OSHA). The proposed information collection will continue to be used to ensure compliance with vaccination requirements in the authorities above, generate the list of persons required to be tested on a routine basis, and will provide important information

regarding safety frameworks, guidance, and procedures.

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

estimated annualized burden hours are 8,499.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of collection	Number of respondents	Number responses per respondent	Average burden per response (in hours)	Total burden hours
NIH COVID-19 Vaccine Status Form .....	31,000	1	5/60	2,583
Screening Questionnaire .....	35,500	1	5/60	2,958
Building Access Form .....	35,500	1	5/60	2,958
Total .....	.....	102,000	.....	8,499

Dated: March 9, 2022.  
**Tara A. Schwetz,**  
*Acting Principal Deputy Director, National Institutes of Health.*  
 [FR Doc. 2022-05475 Filed 3-14-22; 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 NIH Electronic Application System for Certificates of Confidentiality**

**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](http://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 Dr. Pamela Reed Kearney, Division of Human Subjects Research, OER, NIH, 6705 Rockledge Dr., Building Rockledge 1, Room 812-C, Bethesda, MD 20817, or call non-toll-free number (301) 402-2512 or email your request, including your address to: [NIH-CoC-Coordinator@mail.nih.gov](mailto:NIH-CoC-Coordinator@mail.nih.gov).

**SUPPLEMENTARY INFORMATION:** This proposed information collection was previously published in the **Federal Register** on December 17, 2021, page 71650 (FR 86 pages 71650-71651) 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 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 February 28, 2023, 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:* Electronic Application for NIH Certificates of Confidentiality (CoC E-application System), 0925-0689, REVISION, exp., date 02/28/2023, Office of Extramural Research (OER), National Institutes of Health (NIH).

*Need and Use of Information Collection:* The current CoC system sends system communications and the approved Certificate to the Principal Investigator and the Institutional Official. The optional data fields will allow the requester to identify another person that receives CoC system

communications and the approved Certificate. This request system provides one electronic form to be used by all non-NIH funded research organizations that request a discretionary Certificates of Confidentiality (CoC) from NIH. As described in the authorizing legislation (Section 301(d) of the Public Health Service Act, 42 U.S.C. 241(d)), CoCs are issued by the agencies of the Department of Health and Human Services (HHS), including NIH, to authorize researchers to protect the privacy of human research subjects by prohibiting them from releasing names and identifying characteristics of research participants to anyone not connected with the research, except in limited circumstances specified in the statute. At NIH, the issuance of discretionary CoCs has been delegated to the OER in the NIH Office of the Director. NIH received 795 requests for CoCs from January 2020 through December 2020 and expects to receive approximately the same number of requests in subsequent years. The NIH has been using an online CoC system to review requests and issue CoCs since 2015. The current CoC request form includes six sections of information collected from research organizations. The information provided is used to determine eligibility for a CoC and to issue the CoC to the requesting organization. Eligible requesting organizations that provide legally binding affirmations that they will abide by the terms of the CoC are issued a Certificate of Confidentiality. This system has increased efficiency and reduced burden for both requesters and NIH staff who currently process these requests. OMB approval is requested for three years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 1193.