Reduction Act of 1995 to provide opportunity for public comment on proposed data collection projects, the National Institutes of Health (NIH), Office of Research Services (ORS), Division of Occupational Health and Safety (DOHS) 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 within 60 days of the date of this publication.

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: 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*. 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 minimizes 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: NIH COVID–19 Vaccination Status Form EXTENSION, 0925–0771, exp., 3/31/2022, Office of Research Services (ORS), National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of the NIH COVID–19 Vaccination Status Form is 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 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 2,583.

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
Certified nurse coaches	31,000	1	5/60	2,583
Total		31,000		2,583

Dated: November 16, 2021.

Lawrence A. Tabak,

Principal Deputy Director, National Institutes of Health.

[FR Doc. 2021–25412 Filed 11–19–21; 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 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: Center for Scientific Review Special Emphasis Panel; Small Business: Computational, Modeling, and Biodata Management.

Date: December 3, 2021.

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: Marie-Jose Belanger, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Rm. 6188, MSC 7804, Bethesda, MD 20892, (301) 435–1267, *belangerm@csr.nih.gov*.

This notice is being published less than 15 days prior to the meeting due to the timing limitations imposed by the review and funding cycle.

(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 17, 2021.

David W. Freeman,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021–25386 Filed 11–19–21; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Allergy and Infectious Diseases; 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: Microbiology, Infectious Diseases and AIDS Initial Review Group; Acquired Immunodeficiency Syndrome Research Study Section.

Date: December 13, 2021.

Time: 11:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3F40A, Rockville, MD 20892 (Virtual Meeting).

Contact Person: Dimitrios Nikolaos Vatakis, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, MSC–9823, Bethesda, MD 20892, (301) 761–7176, *dimitrios.vatakis@nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.855, Allergy, Immunology, and Transplantation Research; 93.856, Microbiology and Infectious Diseases Research, National Institutes of Health, HHS)

Dated: November 16, 2021.

Tyeshia M. Roberson-Curtis,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2021–25335 Filed 11–19–21; 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; Federal COVID Response—Audience Feedback To Inform Ongoing Messaging and Strategies for ''Combat COVID''

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, Office of Policy for Extramural Research Administration, 6705 Rockledge Drive, Suite 350, Bethesda, Maryland 20892, or call a 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 September 7, 2021, page 50143 (86 FR 50143) 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, any 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: Audience Feedback to Inform Ongoing Messaging and Strategies for ''Combat COVID,'' OMB #0925–0769, exp., date 12/31/2021, EXTENSION National Institutes of Health (NIH).

Need and Use of Information Collection: The purpose of the information collection is to collect routine feedback from the Combat COVID Initiative's two target audiences (the general public and healthcare providers) to identify evolving needs and better disseminate relevant information as it relates to COVID–19 treatment and Accelerating COVID–19 Therapeutic Interventions and Vaccines (ACTIV) clinical trial resources, specifically. Data collected will be used to inform the development and broad dissemination of Combat COVID resources, including new or enhanced message and material concepts (*e.g.,* social media ads, digital display ads, out-of-home ads), and/or web pages (*combatcovid.hhs.gov*). Because the COVID–19 treatment landscape continues to evolve, new evidence-based information continues to come to the forefront, and audience needs continue to change, it is critical for the Federal COVID Response (FCR) Team to collect quick audience feedback from the general public (especially from groups who have not historically been well-represented in clinical trials) and healthcare providers to identify these evolving needs. By understanding target audience needs, the FCR team will be able to properly develop and broadly disseminate relevant COVID–19 treatment and ACTIV clinical trial resources. A change request was recently submitted to update the recall stimuli (still images, audio, and video or animated images) for questions about exposure to the Combat COVID message and materials. It also removed questions that are no longer relevant and replaced them with more relevant questions about the latest COVID–19 treatment options and clinical trials.

OMB approval is requested for 1 year. There are no costs to respondents other than their time. The total estimated annualized burden hours are 3,528.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hours
Consumer Audience Feedback Team Screener	120	1	5/60	10
HCP Audience Feedback Team Screener	40	1	5/60	3
Consumer Audience Feedback Activity	60	8	1	480
HCP Audience Feedback Activity	20	8	1	160