

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 Institute of Allergy and Infectious Diseases Special Emphasis Panel; NIAID Clinical Trial Implementation Cooperative Agreement (U01 Clinical Trial Required).

Date: May 19, 2023.

Time: 11:00 a.m. to 1:00 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 3G58, Rockville, MD 20892 (Virtual Meeting).

Contact Person: Anuja Mathew, 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, Room 3G58, Rockville, MD 20852, 301-761-6911, anuja.mathew@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: April 20, 2023.

Tyeshia M. Roberson-Curtis,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023-08778 Filed 4-25-23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney 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: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel; Ancillary Studies to the IBD Genetics Consortium.

Date: June 20, 2023.

Time: 12:00 p.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, National Institute of Diabetes and Digestive and Kidney Diseases, Democracy II, 6707 Democracy Blvd., Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Maria E. Davila-Bloom, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 7017, 6707 Democracy Boulevard, Bethesda, MD 20892-5452, (301) 594-7637, davila-bloom@extra.nidDK.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.847, Diabetes, Endocrinology and Metabolic Research; 93.848, Digestive Diseases and Nutrition Research; 93.849, Kidney Diseases, Urology and Hematology Research, National Institutes of Health, HHS)

Dated: April 20, 2023.

Miguelina Perez,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023-08748 Filed 4-25-23; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute on Aging; 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 Institute on Aging Special Emphasis Panel; Clinical trials for age-related conditions affecting mobility.

Date: June 6, 2023.

Time: 11:00 a.m. to 1:30 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, National Institute on Aging, Gateway Building, 7201 Wisconsin Avenue, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Maurizio Grimaldi, M.D., Ph.D., Scientific Review Officer, National

Institute on Aging, National Institutes of Health, 7201 Wisconsin Avenue, Room 2C218, Bethesda, MD 20892, 301-496-9374, grimaldim2@mail.nih.gov.

(Catalogue of Federal Domestic Assistance Program Nos. 93.866, Aging Research, National Institutes of Health, HHS)

Dated: April 20, 2023.

Miguelina Perez,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023-08747 Filed 4-25-23; 8:45 am]

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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: National Institute of Allergy and Infectious Diseases Special Emphasis Panel; NIAID Investigator Initiated Program Project Applications (P01 Clinical Trial Not Allowed).

Date: May 11, 2023.

Time: 12:00 p.m. to 5:00 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 3G22, Rockville, MD 20892 (Virtual Meeting).

Contact Person: Richard G. Kostriken, 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, Room 3G22, Rockville, MD 20852, 240-669-2075, richard.kostriken@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: April 20, 2023.
Tyeshia M. Roberson-Curtis,
*Program Analyst, Office of Federal Advisory
 Committee Policy.*
 [FR Doc. 2023-08776 Filed 4-25-23; 8:45 am]
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**DEPARTMENT OF HEALTH AND
 HUMAN SERVICES**

National Institutes of Health

**Proposed Collection; 60-Day Comment
 Request; ABCD Study® Data Use
 Certification (National Institute on Drug
 Abuse)**

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
 National Institute on Drug Abuse
 (NIDA) 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. Elizabeth A. Hoffman,
 Senior Scientific Program Manager,
 Division of Extramural Research,
 National Institute on Drug Abuse, 3WFN
 Room 09C75 MSC 6021, Gaithersburg,
 MD 20877, or call non-toll-free number
 (301) 594-2265 or Email your request,
 including your address to:
elizabeth.hoffman@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: ABCD
 Study® Data Use Certification, 0925–
 NEW, exp., date XX/XX/XXXX,
 National Institute on Drug Abuse
 (NIDA), National Institutes of Health
 (NIH).

*Need and Use of Information
 Collection:* The purpose of this proposal
 is to inform data requestors about terms
 and conditions for using data generated
 by the Adolescent Brain Cognitive
 Development^(SM) (ABCD) Study and to
 obtain signed agreements from
 requestors and their institutional
 officials attesting to their commitment
 to abide by the data use terms and
 conditions. These include using data for
 research purposes; adhering to human
 subjects research requirements; not
 distributing the data to non-authorized
 users; minimizing risk of participant
 identifiability; using data ethically and
 responsibly; and keeping the data
 secure. Users must include a brief
 description of their research project and
 submit their signed data use agreements
 to the data repository to gain access to
 the ABCD Study® data. Users who plan
 to conduct research studies specifically
 on American Indian/Alaska Native (AI/
 AN) populations must submit an
 additional signed data use certification.

OMB approval is requested for 3
 years. There are no costs to respondents
 other than their time. The total
 estimated annualized burden hours are
 2,000.

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
Individuals (standard DUC form)	1800	1	1	1800
Individuals (additional AI/AN DUC form when needed)	200	1	1	200
Total		2000		2,000

Lanette A. Palmquist,
*Project Clearance Liaison, National Institute
 on Drug Abuse, National Institutes of Health.*
 [FR Doc. 2023-08743 Filed 4-25-23; 8:45 am]
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**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
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 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
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 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; Special
 Topic: Brain Imaging, Vision, Bioengineering
 and Low Vision Technology Development.

Date: May 25–26, 2023.

Time: 8:30 a.m. to 7:00 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: Susan Gillmor, Ph.D.,
 Scientific Review Officer, Center for
 Scientific Review, National Institutes of
 Health, 6701 Rockledge Drive, Bethesda, MD
 20892, (240) 762-3076, *susan.gillmor@
 nih.gov*.