Need and Use of Information
Collection: The NIMH Data Archive
(NDA) is an infrastructure that allows
for the submission and storage of human
subjects' data from researchers
conducting studies related to many
scientific domains, regardless of the
source of funding. The NIH and NIMH
developed this resource to allow for the
public collection of information from:
(1) Individuals who seek permission to
access data from the NDA for the
purpose of scientific investigation,
scholarship or teaching, or other forms
of research and research development,

via the Data Use Certification (DUC), and (2) individuals who request permission to submit data to the NDA for the purpose of scientific investigation, scholarship or teaching, or other forms of research and research development, via the Data Submission Agreement (DSA). The extensive information stored in the NDA continues to provide a rare and valuable scientific resource to the field and plays an integral part in fulfilling research objectives in multiple scientific domains. The NIH and the NIMH seek to encourage use of the NDA by

investigators in the field of multiple scientific research domains to achieve rapid scientific progress. In order to take full advantage of this resource and maximize its research value, it is important that data are made broadly available, on appropriate terms and conditions, to the largest possible number of investigators.

OMB approval is requested for three years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 1875.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
NDA Data Submission Agreement (DSA).	Researchers submitting data	300	1	90/60	450
NDA Data Use Certification (DUC)	Researchers requesting access to data.	950	1	90/60	1,425
Total			1,250		1,875

Dated: June 11, 2020.

Melba O. Rojas,

Project Clearance Liaison, National Institute of Mental Health, National Institutes of Health.

[FR Doc. 2020–13136 Filed 6–17–20; 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

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 contract proposals and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the contract proposals, 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; Development of Radiation/ Nuclear Medical Countermeasures (MCMs). Date: July 14, 2020. Time: 10:00 a.m. to 5:00 p.m. Agenda: To review and evaluate contract

Agenda: To review and evaluate contra proposals.

Place: National Institute of Allergy and Infectious Diseases, National Institutes of Health, 5601 Fishers Lane, Room 3G42, Rockville, MD 20892, (Telephone Conference Call).

Contact Person: Sandip Bhattacharyya, 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 3G42, Rockville, MD 20852, sandip.bhattacharyya@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: June 12, 2020.

Tyeshia M. Roberson,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2020–13098 Filed 6–17–20; 8:45 am]

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Nursing Research; 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 Nursing Research Initial Review Group. Date: June 25–26, 2020.

Time: 8:00 a.m. to 12:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institute of Nursing Research, 6701 Rockledge Drive, Room 6187, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Cheryl Nordstrom, Ph.D., Scientific Review Officer, National Institute of Nursing Research, National Institute of Health, 6701 Democracy Blvd., Suite 703H, Bethesda, MD 20892, (301) 827–1499, cheryl.nordstrom@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. 93.361, Nursing Research, National Institutes of Health, HHS) Dated: June 12, 2020.

Miguelina Perez,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2020-13095 Filed 6-17-20; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Neurological Disorders and Stroke; 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: National Institute of Neurological Disorders and Stroke Special Emphasis Panel; NSD–B Conflict SEP.

Date: July 1, 2020.

Time: 2:00 p.m. to 4:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Neuroscience Center, 6001 Executive Blvd., North Bethesda, MD 20852 (Video Assisted Meeting).

Contact Person: Joel A. Saydoff, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH NSC, 6001 Executive Blvd., Room 3205, MSC 9529, Bethesda, MD 20892, (301)–496–9223, joel.saydoff@nih.gov.

Name of Committee: National Institute of Neurological Disorders and Stroke Special Emphasis Panel; R13 Review.

Date: July 6, 2020.

Time: 10:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Neuroscience Center, 6001 Executive Blvd., North Bethesda, MD 20852 (Video Assisted Meeting).

Contact Person: Li Jia, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Research, NINDS/ NIH, 6001 Executive Boulevard, Room 3208D, Rockville, MD 20852, 301–451–2854, li.jia@nih.gov.

Name of Committee: National Institute of Neurological Disorders and Stroke Special Emphasis Panel; NSD–K: Clinical Trials in Neurological Disorders.

Date: July 7, 2020. Time: 9:00 a.m. to 3:00 p.m. Agenda: To review and evaluate grant applications.

Place: Neuroscience Center, 6001 Executive Blvd., North Bethesda, MD 20852 (Video Assisted Meeting).

Contact Person: Shanta Rajaram, Ph.D., Scientific Review Officer, Scientific Review Branch, Division of Extramural Activities, NINDS/NIH NSC, 6001 Executive Blvd., Suite 3208, MSC 9529, Bethesda, Md 20892, (301) 435–6033, rajarams@mail.nih.gov. (Catalogue of Federal Domestic Assistance Program Nos. 93.853, Clinical Research Related to Neurological Disorders; 93.854, Biological Basis Research in the Neurosciences, National Institutes of Health, HHS)

Dated: June 12, 2020.

Tyeshia M. Roberson,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2020-13097 Filed 6-17-20; 8:45 am]

BILLING CODE 4140-01-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Specimen Resource Locator (National Cancer Institute)

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 Institutes of Health, National Cancer Institute (NCI) will publish periodic summaries of propose 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: ${\rm To}$

obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: Joanne Demchok, Program Director, Cancer Diagnosis Program, Division of Cancer Treatment and Diagnosis, 9609 Medical Center Drive, Rockville, Md 20892 or call non-toll-free number 240–276–5959 or Email your request, including your address to: peterjo@mail.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: Specimen

Proposed Collection Title: Specimen Resource Locator (NCI), 0925–0703: Expiration Date 11/30/2020, REVISION, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information
Collection: The availability of
specimens and associated data is critical
to increase our knowledge of cancer
biology, and to translate important
research discoveries to clinical
application. The development of
molecular technologies in cancer
patients, with defined molecular
abnormalities, advances identification
and development of clinically useful
biomarkers and diagnostic assays that
guide treatment.

The discovery and validation of cancer prevention markers require access, by researchers, to quality clinical biospecimens. In response, to this need, NCI's Cancer Diagnosis Program developed, and is expanding, a searchable database: Specimen Resource Locator (SRL) https:// specimens.cancer.gov/tissue/ default.htm. The SRL allows scientist in the research community and the NCI to locate specimens needed for their research. The SRL lists all NCI supported and non-NCI supported biospecimens repositories and their links. It is not NCI's intent to collect the biospecimens; rather the collections are descriptions of the available data that can act as a resource and be shared with researchers and scientists who are interested. This submission does not involve any analysis.

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