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: The National Institute of Mental Health Data Archive (NDA), REVISION, OMB Control Number 0925–0667, National Institute of Mental Health (NIMH), National Institutes of Health (NIH).

Need and Use of Information Collection: This REVISION request seeks approval of updates to the previously approved National Database for Autism Research Data Access Request and Data Use Certification documents. The NIMH Data Archive (NDA), formerly known as the National Database for Autism Research (NDAR), 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 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 1000.

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	250	1	1 hour	250
NDA Data Use Certification (DUC)	Researchers requesting access to data.	750	1	1 hour	750
Total	.....	1000	1000	.....	1000

Melba Rojas,

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

[FR Doc. 2017–14451 Filed 7–10–17; 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 (5 U.S.C. App.), 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; T Cell Reagent Resource for the Study of Allergic Diseases (U19).

Date: August 1–2, 2017.

Time: 9:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 5601 Fishers Lane, Rockville, MD 20892 (Telephone Conference Call).

Contact Person: Thomas F. Conway, Ph.D., Scientific Review Officer, Scientific Review Program, Division of Extramural Activities, Room 3G51, National Institutes of Health, NIAID, 5601 Fishers Lane, MSC 9823, Bethesda, MD 20892–9823, 240–507–9685, *thomas.conway@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: July 5, 2017.

Natasha M. Copeland,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–14433 Filed 7–10–17; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of