knowledge of product analyses used by tobacco product manufacturers to ensure product consistency.

II. Description of Site Tours Program

In the Laboratory Site Tours Program, small groups of CTP staff plan to observe the operations of laboratories that perform testing and analyses of tobacco products and tobacco smoke relative to analytical chemistry, microbiology, toxicology, biomarkers of exposure or risk, and analytical method development. Please note that the Laboratory Site Tours Program is not intended to include official FDA inspections of facilities to determine compliance with the FD&C Act or for the purposes of developing Tobacco Product Manufacturing Practice regulations; rather, the program is meant to educate CTP staff and improve their understanding of laboratory testing and analyses used by the tobacco industry.

III. Site Selection

CTP plans to select a wide variety of laboratories that include academic, private, and those affiliated with tobacco manufacturers. All travel expenses associated with the site tours will be the responsibility of CTP. Final site selections will be based on the availability of CTP funds and resources for the relevant fiscal year, as well as the following factors, if applicable: (1) Compliance status of the requesting facility and affiliated firm, (2) whether the requesting facility is in arrears for user fees, and (3) whether the requesting facility or affiliated firm has a significant request or marketing application or submission pending with FDA.

IV. Requests for Participation

Identify requests for participation with the docket number found in brackets in the heading of this document. Received requests are available for public examination in the Division of Dockets Management (see **ADDRESSES**) between 9 a.m. and 4 p.m., Monday through Friday.

Dated: September 30, 2014.

Leslie Kux,

Assistant Commissioner for Policy.

[FR Doc. 2014–23844 Filed 10–6–14; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; NIMH Database of Cognitive Training and Remediation Studies (DCTRS) (NIMH)

SUMMARY: Under the provisions of 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. This proposed information collection was previously published in the **Federal Register** on April 15, 2014, pages 21250–21252 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 Institute of Mental Health (NIMH), 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.

Direct Comments to OMB: Written comments and/or suggestions regarding the item(s) contained in this notice, especially regarding the estimated public burden and associated response time, should be directed to the: Office of Management and Budget, Office of Regulatory Affairs, *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974, Attention: NIH Desk Officer.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 30-days of the date of this publication.

FOR FURTHER INFORMATION CONTACT: To obtain a copy of the data collection plans and instruments or request more information on the proposed project contact: Keisha Shropshire, NIMH Project Clearance Liaison, Science Policy and Evaluation Branch, OSPPC, NIMH, NIH, Neuroscience Center, 6001 Executive Boulevard, MSC 9667, Rockville Pike, Bethesda, MD 20892, or call 301–443–4335 or Email your request, including your address to: *nimhprapubliccomments@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection

NIMH Database of Cognitive Training and Remediation Studies (DCTRS)—New—National Institute of Mental Health (NIMH), National Institute of Health (NIH).

Need and Use of Information Collection: The NIMH Database of Cognitive Training and Remediation Studies (DCTRS) is an integrated database that includes study- and subject-level data from studies of cognitive remediation (CR) in schizophrenia. DCTRS will allow NIMH staff and interested investigators to examine the ways in which various patient characteristics, intervention approaches and features, and treatment combinations affect responses to remediation. The DCTRS Study Information Form and Data Submission Agreement are necessary for the ''Submitter'' to request permission to submit study data to the NIMH DCTRS for general research purposes. The primary use of this information is to collect submitter information and study information for inclusion in the NIMH DCTRS database. The DCTRS data submission agreement includes two forms: (1) The data submission form that includes the terms, agreement, submitter information and certifications, and (2) the study information form which collects de-identified data for each study.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 60.

ESTIMATED ANNUALIZED BURDEN HOURS

Form	Type of respondent	Number of respondents	Frequency of response	Average time per response (in hours)	Annual hour burden
Data Submission Agreement	Principal Investigators/Physicians	12	1	5	60

Dated: September 29, 2014.

Keisha Shropshire,

NIMH Project Clearance Officer, NIMH, NIH.

[FR Doc. 2014–23938 Filed 10–6–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; NIMH Data Repositories Data Submission Request; NIMH Data Repositories Data Access and Use Certification

SUMMARY: In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Institute of Mental Health (NIMH), National Institutes of Health (NIH), will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Written comments and/or suggestions from the public and affected agencies are invited on 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.

To Submit Comments and For Further Information: To obtain a copy of the data collection plans and instruments, submit comments in writing, or request more information on the proposed project, contact: NIMH Project Clearance Liaison, Science Policy and Evaluation Branch, OSPPC, NIMH, NIH, Neuroscience Center, 6001 Executive Boulevard, MSC 9667, Rockville Pike, Bethesda, MD 20892, or call 301–443–4335 or Email your request, including your address to: *nimhprapubliccomments@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Comment Due Date: Comments regarding this information collection are best assured of having their full effect if received within 60 days of the date of this publication.

Proposed Collection: NIMH Data Repositories (NDR) Data Submission Request, the NIMH Data Repositories Data Access and Use Certification, 0925–0667 Revision; National Institute of Mental Health (NIMH), National Institutes of Health (NIH).

Need and Use of Information Collection: The National Institutes of Mental Health (NIMH) Data Repositories are a group of Federal data repositories based on an informatics platform for human-subjects research domains related to mental health, initially established as the National Database for Autism Research (NDAR) to support autism-related research. In 2013, NIMH received approval from OMB for use of the NIMH Data Access Request and Use Certification (DUC) Form to meet the unique data access needs of all existing NIMH data repositories, which at the time consisted of NDAR, Pediatric MRI (PedsMRI), and the NIMH Clinical Research Datasets (NCRD)—OMB# 0925–0667 (Expiration: 09/30/2016). Now in 2014, two new databases have been added and integrated into the NDAR infrastructure, NDCT and RDoCdb. At this time, NIMH is seeking OMB approval to add an all-purpose NIMH Data Repositories Data Submission Request Form and to add a revised all-purpose NIMH Data Repositories Data Access and Use Certification Form. As the data repositories have matured, and with the introduction of the new databases—namely NDCT and RDoCdb—the information being collected for data submission has become more complex, rendering an OMB-approved submission form a new necessity.

OMB approval is requested for three years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 221.

ESTIMATED ANNUALIZED BURDEN HOURS

Form	A. Estimates annual burden hours			
	Number of respondents	Frequency of response	Average time per response (in hours)	Annual burden hour
NIMH Data Repositories Data Submission Request Form	40	1	95/60	63
NIMH Data Repositories Data Access and Use Certification Form	100	1	95/60	158

Dated: September 29, 2014.

Keisha L Shropshire,

Project Clearance Liaison, NIMH, NIH.

[FR Doc. 2014–23959 Filed 10–6–14; 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 Meetings

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 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: Center for Scientific Review Special Emphasis Panel Fellowships: Synthetic and Biological Chemistry.

Date: November 4–5, 2014.

Time: 8:00 a.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.