section 487F (42 U.S.C. 288–6); the Extramural Clinical Research LRP for Individuals from Disadvantaged Backgrounds (ECR–LRP) is authorized by an amendment to section 487E (42 U.S.C. 288–5); the Contraception and Infertility Research LRP (CIR–LRP) is authorized by section 487B (42 U.S.C. 288–2); and the Health Disparities Research Loan Repayment Program (HD–LRP) is authorized by section 485G (42 U.S.C. 287c–33).

The Loan Repayment Programs can repay up to $35,000 per year toward a participant's extant eligible educational loans, directly to financial institutions. The information proposed for collection will be used by the Division of Loan Repayment to determine an applicant's eligibility for participation in the program.

Frequency of Response: Initial application and one- or two-year renewal application.

Affected Public: Individuals or households; Nonprofits; and Businesses or other for-profit.

Type of Respondents: Physicians, other scientific or medical personnel, and institutional representatives.

Questions, required information, and requested documents remain largely unchanged. Improvements were made to the structure and appearance of online forms to provide applicants with a better user experience. Recommenders will no longer be asked to complete a recommendation form, but to write a reference letter that comments on the research skills and the abilities of the applicant. A general eligibility checklist (NIH 2674–20) was added at the start of the application to reduce the likelihood of ineligible individuals working through the application only to learn of their disqualification after submitting the application. Redundant questions or statements were eliminated. OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 33,242.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Estimated number of responses per respondent	Average burden hours per response	Annual burden hours requested
Intramural LRPs				
Initial Applicants	40	1	10	400
Advisors/Supervisors	40	1	1	40
Recommenders	120	1	30/60	60
Financial Institutions	8	1	15/60	2
Subtotal	208			502
Extramural LRPs				
Initial Applicants	1,650	1	11	18,150
Advisors/Supervisors	1,480	1	1	1,480
Recommenders	4,950	1	30/60	2,475
Financial Institutions	100	1	15/60	25
Subtotal	8,180			22,130
Intramural LRPs				
Renewal Applicants	40	1	7	280
Advisors/Supervisors	40	1	2	80
Subtotal	80			360
Extramural LRPS				
Renewal Applicants	1,000	1	8	8,000
Advisors/Supervisors	750	1	1	750
Recommenders	3,000	1	30/60	1,500
Subtotal	4,750			10,250
Total	13,218			33,242

Dated: July 1, 2016.

Lawrence A. Tabak,

Deputy Director, National Institutes of Health.

[FR Doc. 2016–16227 Filed 7–7–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Diabetes and Digestive and Kidney Diseases: 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: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel, NIDDK Training Grants Review.

Date: July 26, 2016.

Time: 4:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: CAROL J. GOTER-ROBINSON, PH.D., SCIENTIFIC REVIEW OFFICER, REVIEW BRANCH, DEA, NIDDK, NATIONAL INSTITUTES OF HEALTH, ROOM 7347, 6707 DEMOCRACY BOULEVARD, BETHESDA, MD 20892–5452, (301) 594–7791, *goterrobinsonc@extra.niddk.nih.gov.*

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel, Program Project on Gut Microbial Host Interactions.

Date: August 5, 2016.

Time: 1:00 p.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, 6707 Democracy Boulevard, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: MARIA E. DAVILA-BLOOM, PH.D., SCIENTIFIC REVIEW OFFICER, REVIEW BRANCH, DEA, NIDDK, ROOM 7017, 6707 DEMOCRACY BOULEVARD, BETHESDA, MD 20892, (301) 594–7637, *davila-bloomm@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: July 1, 2016.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2016–16142 Filed 7–7–16; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request: National Institute of Neurological Disorders and Stroke Federal Interagency Traumatic Brain Injury Research (FITBIR) Data Access Request

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 Neurological Disorders and Stroke (NINDS), 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: Dr. Sophia Jeon, Health Science Policy Analyst, Office of Science Policy and Planning (OSPP), NINDS, NIH, 31 Center Drive, Building 31, Room 8A03, Bethesda, MD 20892, or call non-toll-free number (301) 435–7571, or Email your request, including your address to: *sophia.jeon@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: National Institute of Neurological Disorders and Stroke Federal Interagency Traumatic Brain Injury Research (FITBIR) Data Access Request, 0925–0677, Expiration Date 08/31/2016—Reinstatement without change, National Institute of Neurological Disorders and Stroke (NINDS), National Institutes of Health (NIH).

Need and Use of Information Collection: The FITBIR Informatics System Data Access Request form is necessary for ''Recipient'' Principal Investigators and their organization or corporations with approved assurance from the DHHS Office of Human Research Protections to access data or images from the FITBIR Informatics System for research purposes. The primary use of this information is to document, track, monitor, and evaluate the use of the FITBIR datasets, as well as to notify interested recipients of updates, corrections or other changes to the database. Type of respondents affected by this information collection are researchers, such as Principal Investigators (PI), who are interested in obtaining access to study data and images from the FITBIR Informatics System for research purposes.

There are two scenarios for completing the form. The first is where the Principal Investigator (PI) completes the entire FITBIR Informatics System Data Access Request form, and the second where the PI has the Research Assistant begins filling out the form and PI provides the final reviews and signs it. The estimated annual burden hours to complete the data request form are listed below.

OMB approval reinstatement is requested for 3 years. The total estimated annualized burden hours are 63.

ESTIMATED ANNUALIZED BURDEN

Estimated annual burden hours for respondents

Form	Type of respondent	Number of respondents	Annual frequency per response	Hours per response	Total hours
FITBIR Informatics System Data Access Request.	Individuals (Principal Investigators)	40	1	95/60	63
Total		40	40		63

Dated: July 1, 2016.

Paul Scott,

Project Clearance Liaison Officer, National Institute of Neurological Disorders and Stroke, NIH.

[FR Doc. 2016–16256 Filed 7–7–16; 8:45 am]

BILLING CODE 4140–01–P