application, will not be treated confidentially, and may be made publicly available.

License applications submitted in response to this Notice will be presumed to contain business confidential information and any release of information in these license applications will be made only as required and upon a request under the Freedom of Information Act, 5 U.S.C. 552.

Dated: January 13, 2023.

Richard U. Rodriguez,

Associate Director, Technology Transfer Center, National Cancer Institute.

[FR Doc. 2023–01019 Filed 1–19–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Evaluation of the Enhancing Diversity of the NIH-Funded Workforce Program (National Institute of General Medical Sciences)

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 of General Medical Sciences (NIGMS) 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. Alison Gammie, Director, Division of Training, Workforce Development, and Diversity, NIGMS, 45 Center Drive, Room 2AS43J, Bethesda, MD 20892, or call non-toll-free number (301) 496–7301 or Email your request, including your address to: *alison.gammie@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: Evaluation of the Enhancing the Diversity of the NIH-funded Workforce Program Consortium (DPC), 0925–0747, 06/30/2024, EXTENSION, National Institute of General Medical Sciences (NIGMS), National Institutes of Health (NIH).

Need and Use of Information Collection: This request is for an Extension of a currently approved collection. The goal of the DPC is to address a unique and compelling need identified by NIH, namely to enhance the diversity of well-trained biomedical research scientists who can successfully compete for NIH research funding and/or otherwise contribute to the NIH-funded scientific workforce. The DPC is a national collaborative through which awardee institutions, in partnership with NIH, aim to enhance diversity in the biomedical research workforce through the development, implementation, assessment and dissemination of innovative and effective approaches to: (a) student outreach, engagement, training, and mentoring, (b) faculty development, and (c) institutional research training infrastructure. The Coordination and Evaluation Center (CEC) will evaluate the efficacy of the training and mentoring approaches implemented across a variety of contexts and populations and will disseminate information to the broader research community. The planned consortium-wide data collection and evaluation will provide comprehensive information about the multi-dimensional factors (individual, institutional, and faculty/mentor) that influence student and faculty success, professional development, and persistence within biomedical research career paths across a variety of contexts. The planned data collection, and the resulting findings, is projected to have a sustained, transformative effect on biomedical research training and mentoring nationwide.

OMB approval is requested for an extension of 13 months beyond the currently approved collection, until June 2024. There are no costs to respondents other than their time. The total estimated annualized burden hours are 11,730.

ESTIMATED ANNUALIZED BURDEN HOURS

Form name	Type of respondents	Number of respondents	Number of responses per respondent	Average time per response (in hours)	Total annual burden hours
Student Annual Follow-up survey (Attachment 13).	Non-BUILD Student and BUILD student.	15,000	1	45/60	11,250
BUILD Institutional Research & Program Data Requests (Attachment 19).	Personnel and Administrators at BUILD Institutions.	10	3	16	480
Total			15,030		11,730

Dated: January 13, 2023.

David N. Bochner,

Project Clearance Liaison, National Institute of General Medical Sciences, National Institutes of Health.

[FR Doc. 2023–00998 Filed 1–19–23; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Center for Advancing Translational Sciences; 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 cooperative agreement applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the cooperative agreement applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Center for Advancing Translational Sciences Special Emphasis Panel; NCATS CTSA UM1 Review Special Emphasis Panel.

Date: February 21, 2023.

Time: 11:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate cooperative agreement applications.

Place: National Center for Advancing Translational Sciences, National Institutes of Health, 6701 Democracy Boulevard, Room 1037, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Victor Henriquez, Ph.D., Scientific Review Officer, Office of Scientific Review, National Center for Advancing Translational Sciences, National Institutes of Health, 6701 Democracy Boulevard, Room 1037, Bethesda, MD 20892, (301) 435–0813, *henriquv@mail.nih.gov.*

(Catalogue of Federal Domestic Assistance Program Nos. 93.859, Pharmacology, Physiology, and Biological Chemistry Research; 93.350, B—Cooperative Agreements; 93.859, Biomedical Research and Research Training, National Institutes of Health, HHS)

Dated: January 13, 2023.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2023–01017 Filed 1–19–23; 8:45 am]

BILLING CODE 4140–01–P

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 information collection requests under OMB review, in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (240) 276–0361.

Project: SAMHSA's Publications and Digital Products Website Registration Surveys (OMB No. 0930–0313)—Revision

The Substance Abuse and Mental Health Services Administration (SAMHSA) is requesting OMB approval for a revision of SAMHSA's Publications and Digital Products website Registration Survey (OMB No. 0930–0313). SAMHSA is authorized under section 501(d)(16) of the Public Health Service Act (42 U.S.C. 290aa(d)(16)) to develop and distribute materials for the prevention, treatment, and recovery from mental and substance use disorders. To improve customer service and lessen the burden on the public to locate and obtain these materials, SAMHSA has developed a website that includes more than 500 free publications from SAMHSA and its component Agencies. These products are available to the public for ordering and download. When a member of the public chooses to order hard-copy publications, it is necessary for SAMHSA to collect certain customer information in order to fulfill the request. To further lessen the burden on the public and provide the level of customer service that the public has come to expect from product websites, SAMHSA has developed a voluntary registration process for its publication website that allows customers to create accounts. Through these accounts, SAMHSA customers are able to access their order histories and save their shipping addresses. During the website registration process, SAMHSA will also ask customers to provide optional demographic information that helps SAMHSA to evaluate the use and distribution of its publications and improve services to the public.

SAMHSA employs a web-based form for information collection to avoid duplication and unnecessary burden on customers who register for an account. Customer information is submitted electronically via web forms on the samhsa.gov domain. Customers can submit the web forms at their leisure or call SAMHSA's toll-free Call Center and an information specialist will submit the forms on their behalf. The electronic collection of information reduces the burden on the respondent and streamlines the data-capturing process. The following revisions were made to the SAMHSA Publications and Digital Products website Registration Survey:

- Revision of the SAMHSA Publications website Registration Survey Questions
- Addition of a SAMHSA Main Site Survey version
- Addition of a SAMHSA Store Survey version

SAMHSA estimates the burden of this information collection as follows:

TABLE 1—ESTIMATED ANNUAL REPORTING BURDEN

	Number of respondents	Annual frequency per response	Total annual responses	Hours per response	Total hours
Website Registration Survey	21,082	1	21,082	.033 (2 min.)	696

Written comments and recommendations for the proposed information collection should be sent within 30 days of publication of this notice to *www.reginfo.gov/public/do/PRAMain* . Find this particular information collection by selecting ''Currently under 30-day Review—Open for Public Comments'' or by using the search function.