long term sustainability of the Title X network.

Data collected from this effort will be used to inform the work of the training centers so they can better support the Title X grantees. This data will help OPA better understand challenges affecting Title X centers in order to better work with HHS entities and national stakeholders to provide resources to Title X centers. Data will be collected through an online data collection tool directly from grantees and from Title X centers.

Likely Respondents: This annual reporting requirement is service sites that receive funding (either directly from OPA or through a sub recipient or grantee agency) for family planning services authorized and funded by the Title X Family Planning Program [''Population Research and Voluntary Family Planning Programs'' (91)], which was enacted in 1970 as Title X of the Public Health Service Act (Section 1001 of Title X of the Public Health Service Act, 42 United States Code [U.S.C.] 300).

TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS

Type of respondent	Form name	Number of respondents	Number of responses per respondent	Average annualized burden per response (hours)	Annual total burden (hours)
Grantees	Sustainability Assessment—Grantees	92	1	0.66	60.72
Service Sites	Sustainability Assessment—Sites	4,168	1	0.66	2750.88
Total		4,260			2,811.60

Terry S. Clark,

Asst Information Collection Clearance Officer.

[FR Doc. 2015–14850 Filed 6–16–15; 8:45 am]

BILLING CODE 4150–48–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Submission for OMB Review; 30-Day Comment Request; Population Assessment of Tobacco and Health (PATH) Study—3rd Wave (NIDA)

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 February 11, 2015, pages 7619–7620 and allowed 60-days for public comment. One public comment was received. The purpose of this notice is to allow an additional 30 days for public comment. The National Institute on Drug Abuse (NIDA), 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.

DATES: *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: Dr. Kevin P. Conway, Deputy Director, Division of Epidemiology, Services, and Prevention Research, NIDA, NIH, 6001 Executive Boulevard, Room 5185, Rockville, MD 20852; or call non-toll-free number (301) 443–8755 or Email your request, including your address to: *PATHprojectofficer@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

Proposed Collection: Population Assessment of Tobacco and Health (PATH) Study—Third Wave of Data Collection—0925–0664—Revision, National Institute on Drug Abuse (NIDA), National Institutes of Health (NIH), in partnership with the Food and Drug Administration (FDA).

Need and Use of Information Collection: This is a revision request (OMB 0925–0664, expires 9/30/2016) for the Population Assessment of Tobacco and Health (PATH) Study to conduct the third wave of data collection. The PATH Study is a national longitudinal cohort study of tobacco use behavior and health among the U.S. household population of adults age 18 and older and youth ages 12 to 17. The Study conducts annual interviews and collects biospecimens from adults to assess within-person changes and between-person differences in tobacco-product use behaviors and related health conditions over time. Its longitudinal, population-based data will help to enhance the evidence base that informs FDA's regulatory actions under the Family Smoking Prevention and Control Act to protect the Nation's public health and reduce its burden of tobacco-related morbidity and mortality.

OMB approval is requested for 3 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 54,434.

ESTIMATED ANNUALIZED BURDEN HOURS

Form or activity name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Adult Extended Interview	Adults	25,444	1	1	25,444
Consent for Adult Extended Interview	Adults	2,046	1	4/60	136

ESTIMATED ANNUALIZED BURDEN HOURS—Continued

Form or activity name	Type of respondent	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total annual burden hours
Adult Extended Interview (Aged-up)	Adults	1,780	1	68/60	2,017
Consent for Biological Samples	Adults	1,780	1	5/60	148
Biospecimen Collection: Urine	Adults	13,805	1	10/60	2,301
Biospecimen Collection: Blood	Adults	765	1	18/60	230
Tobacco Use Form	Adults	14,570	1	5/60	1,214
Follow-up/Tracking Participant Information Form for Adults	Adults	27,224	2	8/60	7,260
Youth Extended Interview	Youth	9,625	1	35/60	5,615
Assent for Youth Extended Interview	Youth	1,923	1	3/60	96
Youth Extended Interview (Aged-up)	Youth	1,923	1	45/60	1,442
Parent Interview	Parents	9,818	1	16/60	2,618
Parent Permission and Consent for Parent Interview	Parents	2,161	1	5/60	180
Parent Interview (Aged-up)	Parents	1,961	1	19/60	621
Verification Interview	Adults	35,564	1	2/60	1,185
Validation Interview	Adults	3,579	1	4/60	239
Follow-up/Tracking Participant Information Form for Youth	Parents	11,548	2	8/60	3,079
Follow-up/Tracking Participant Information Form for Sample Shadow Youth.	Parents	2,282	2	8/60	609

Dated: June 12, 2015.

Genevieve deAlmeida-Morris,

Project Clearance Liaison, NIDA, NIH.

[FR Doc. 2015–14902 Filed 6–16–15; 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 Member Conflict: Neurobiology of Psychiatric Disorders and Addictions.

Date: June 24, 2015.

Time: 12:00 p.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Telephone Conference Call).

Contact Person: Boris P. Sokolov, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5217A, MSC 7846, Bethesda, MD 20892, 301–408–9115, *bsokolov@csr.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.

Name of Committee: Center for Scientific Review Special Emphasis Panel Small Business: Non HIV Microbial Vaccines and Countermeasures.

Date: July 13–14, 2015.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Wyndham Grand Chicago Riverfront, 71 E. Wacker Drive, Chicago, IL 60601375.

Contact Person: Andrea Keane-Myers, BS, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4218, Bethesda, MD 20892, 3014351221, *andrea.keane-myers@nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Fellowships: Oncological Sciences.

Date: July 13–14, 2015.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Ola Mae Zack Howard, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Dr., Room 4192, MSC 7806, Bethesda, MD 20892, 301–451–4467, *howardz@mail.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Member Conflict: Vascular Regulation and Diseases.

Date: July 16–17, 2015.

Time: 10:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892 (Virtual Meeting).

Contact Person: Ai-Ping Zou, MD, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4118, MSC 7814, Bethesda, MD 20892, 301–408–9497, *zouai@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Fellowships: Oncology.

Date: July 20–21, 2015.

Time: 8:00 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Embassy Suites at the Chevy Chase Pavilion, 4300 Military Road NW., Washington, DC 20015.

Contact Person: Juraj Bies, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Dr., Rm. 4158, MSC 7806, Bethesda, MD 20892, 301 435 1256, *biesj@mail.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Member Conflict: Auditory Mechanisms.

Date: July 20–21, 2015.

Time: 8:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD (Virtual Meeting).

Contact Person: John Bishop, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5182, MSC 7844, Bethesda, MD 20892, (301) 408–9664, *bishopj@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel Member Conflict: Topics in Virology.

Date: July 21, 2015

Time: 10:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.