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—Second Wave of Data Collection—0925–0664–Revision— National Institutes of Health (NIH), National Institute on Drug Abuse (NIDA), in partnership with the Food and Drug Administration (FDA).

Need and Use of Information Collection: This is a revision request (OMB 0925–0664, expires 11/30/2015) for the Population Assessment of Tobacco and Health (PATH) Study to conduct the second wave of data collection. The PATH Study is a large

national longitudinal cohort study on tobacco use behavior and health among the U.S. household population of adults age 18 and older and youth ages 12 to 17. The PATH Study conducts annual interviews and collects biospecimens from adults to help inform the development, implementation, and evaluation of tobacco-product regulations by FDA in meeting its mission under the Family Smoking Prevention and Tobacco Control Act (TCA) to regulate tobacco products, including tobacco-product advertising, labeling, marketing, constituents, ingredients, and additives. The longitudinal design of the PATH Study enables it to measure and report withinperson changes and between-person differences in tobacco product use

### ESTIMATED ANNUALIZED BURDEN HOURS

behaviors and health effects within the cohort over time. These data will help to inform regulatory decisions and actions by FDA.

OMB approval is requested for 3 years. There are no capital, operating, or maintenance costs to report. There are no costs to respondents other than their time. The total estimated annualized burden hours are 56,939. Factors accounting for the difference between the baseline and Wave 2 total hours include the following: (1) Wave 2 does not have a screening phase; (2) as indicated in Supporting Statement B, a 86 percent response rate for adult interviews and a 90 percent response rate for youth interviews are projected for Wave 2; and (3) fewer biological samples will be collected in Wave 2.

| Type of respondent and instrument                                       | Estimated<br>number of<br>respondents | Estimated<br>number of<br>responses per<br>respondent | Average<br>burden hours<br>per response | Estimated<br>total annual<br>burden hours<br>requested |
|---|---------------------------------------|---|---|--|
| Adults—Extended Interview   | *27,113                               | 1   | 1                                       | 27,113   |
| Adults-Baseline youth respondents who age into adult cohort-Consent     |                                       |   |   |  |
| for Extended Interview  | 2,295                                 | 1   | 2/60                                    | 77   |
| Adults-Baseline youth respondents who age into adult cohort-Extended    |                                       |   |   |  |
| Interview   | * 1,990                               | 1   | 68/60                                   | 2,255  |
| Adults-Baseline youth respondents who age into the adult cohort-Con-    | 1 000                                 |   | 4/00                                    | 100  |
| sent for Biological Samples   | 1,990<br>11.373                       |   | 4/60<br>10/60                           | 133<br>1.896   |
| Adults—Biospecimen Collection: Urine                                    |                                       |   | 18/60                                   | 269  |
| Adults—Biospecimen Collection: Blood<br>Adults—Tobacco Use Form         | 12,269                                |   | 4/60                                    | ≥69<br>818   |
| Adults—Follow-up/Tracking Participant Information Form                  | 33,615                                | 2   | 4/60<br>8/60                            | 8.964  |
| Youth—Extended Interview  | ** 10,537                             | <u></u>   | 32/60                                   | 5,620  |
| Youth-Shadow youth who age into youth cohort-Assent for Extended        | 10,337                                | I   | 52/00                                   | 5,020  |
| Interview   | 2,338                                 | 1   | 2/60                                    | 78   |
| Youth—Shadow youth who age into youth cohort—Extended Interview         | ** 2,105                              | 1   | 42/60                                   | 1,474  |
| Parent Interview  | 10,748                                | 1   | 14/60                                   | 2,508  |
| Parents of Shadow youth who age into youth cohort-Parent Permission     |                                       |   | ,                                       | 2,000  |
| and Consent for Parent Interview  | 2,338                                 | 1   | 2/60                                    | 78   |
| Parents of Shadow youth who age into youth cohort—Parent Interview      | 2,147                                 | 1   | 17/60                                   | 608  |
| Parents of youth—Follow-up/Tracking Participant Information Form for    | ,                                     |   |   |  |
| Youth   | 14,165                                | 2   | 8/60                                    | 3,777  |
| Adults-Follow-up/Tracking Participant Information Form for sample Shad- |                                       |   |   |  |
| ow youth (completed by parents)   | 4,772                                 | 2   | 8/60                                    | 1,273  |

\* Estimated total number of adult extended interview respondents is 27,113 adults from Wave 1 + 1,990 youth from Wave 1 who turn 18 by Wave 2 = 29,103.

\*\*Estimated total number of youth extended interview respondents is 10,537 youth from Wave 1+ 2,105 shadow youth who turn 12 by Wave 2 = 12,642.

#### Dated: June 25, 2014.

Genevieve R. deAlmeida,

Project Clearance Liaison, National Institute on Drug Abuse.

[FR Doc. 2014–15584 Filed 7–1–14; 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, Kidney Disease Ancillary Studies. Date: July 23, 2014. Time: 2:00 p.m. to 4:00 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: Michele L. Barnard, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 753, 6707 Democracy Boulevard, Bethesda, MD 20892–2542, (301) 594–8898, barnardm@extra.niddk.nih.gov.

Name of Committee: National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel, Time-Sensitive Obesity Prevention.

Date: July 30, 2014.

*Time:* 2:00 p.m. to 3:30 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: Michele L. Barnard, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 753, 6707 Democracy Boulevard, Bethesda, MD 20892–2542, (301) 594–8898, barnardm@extra.niddk.nih.gov.

*Name of Committee:* National Institute of Diabetes and Digestive and Kidney Diseases Special Emphasis Panel, RFA–DK13–017 Human Islet Research Network Consortium on Modeling Autoimmune Interactions (HIRN–CMAI).

Date: August 4, 2014.

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

*Agenda:* To review and evaluate cooperative agreement applications.

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

Contact Person: Ann A. Jerkins, Ph.D., Scientific Review Officer, Review Branch, DEA, NIDDK, National Institutes of Health, Room 759, 6707 Democracy Boulevard, Bethesda, MD 20892–5452, 301–594–2242, *jerkinsa@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: June 26, 2014.

## David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2014-15424 Filed 7-1-14; 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; RFA Panel: Ethical Issues Related to Central IRBs and Research Using Clinical Records.

Date: July 8, 2014.

*Time:* 12:00 p.m. to 6:00 p.m. *Agenda:* To review and evaluate grant applications.

<sup>^</sup>*Place:* National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Denise Wiesch, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3138, MSC 7770, Bethesda, MD 20892, (301) 437– 3478, wieschd@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; RFA Panel: Ethical Issues Related to Central IRBs and Research Using Clinical Records.

*Date:* July 8, 2014. *Time:* 12:00 p.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:* Karin F Helmers, Ph.D., Scientific Review Officer, Center for Scientific Review National Institutes of Health, 6701 Rockledge Drive, Room 3144, MSC 7770, Bethesda, MD 20892, (301) 254– 9975, helmersk@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.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research, 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS). Dated: June 26, 2014 Carolyn A. Baum, Program Analyst, Office of Federal Advisory Committee Policy. [FR Doc. 2014–15422 Filed 7–1–14; 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 Member

Conflict: Topics in Virology. Date: July 22–23, 2014.

*Time:* 9:00 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: Marci Scidmore, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3192, MSC 7808, Bethesda, MD 20892, 301–435– 1149, marci.scidmore@nih.gov.

*Name of Committee:* Center for Scientific Review Special Emphasis Panel Member Conflict: Neurological, Aging and Musculoskeletal Epidemiology.

*Date:* July 22, 2014.

*Time:* 2:00 p.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: George Vogler, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3140, MSC 7770, Bethesda, MD 20892, (301) 237– 2693, voglergp@csr.nih.gov.

*Name of Committee:* AIDS and Related Research Integrated Review Group AIDS Discovery and Development of Therapeutics Study Section.

Date: July 25, 2014.

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