agencies, and other entities participating in the voluntary paternity establishment program.

Respondents: State and Tribal IV–D agencies, hospitals, birth record agencies, and other entities participating in the voluntary paternity establishment program.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
None	1,113,719	1	0.17	189,332.23

Estimated Total Annual Burden Hours: 189,332.23.

OMB Comment: OMB is required to make a decision concerning the collection of information between 30 and 60 days after publication of this document in the **Federal Register**. Therefore, a comment is best assured of having its full effect if OMB receives it within 30 days of publication. Written comments and recommendations for the proposed information collection should be sent directly to the following:

Office of Management and Budget
Paperwork Reduction Project
Email: OIRA_SUBMISSION@OMB.EOP.GOV
Attn: Desk Officer for the Administration for Children and Families

Robert Sargis,

Reports Clearance Officer.

[FR Doc. 2014–01097 Filed 1–21–14; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Proposed Collection; 60-Day Comment Request; Next Series of Tobacco Use Supplements to the Current Population Survey

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 Cancer Institute (NCI), 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: Anne Hartman, Health Statistician, Risk Factor Monitoring and Methods Branch, National Cancer Institute, NIH, MSC 9762, 9609 Medical Center Drive, Bethesda, MD or call non-toll-free number 240–276–6704 or Email your request, including your address to: *hartmana@mail.nih.gov.* Formal requests for additional plans and instruments must be requested in writing.

DATES: *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: Next Series of Tobacco Use Supplements to the Current Population Survey (TUS–CPS), 0925–0368, Expiration Date 03/31/2013, Reinstatement with Change, National Cancer Institute (NCI), National Institutes of Health (NIH).

Need and Use of Information Collection: The 2014–15 Tobacco Use Supplement-Current Population Survey (TUS–CPS) will be conducted by the Census Bureau and is co-sponsored by the National Cancer Institute (NCI) and the Food and Drug Administration (FDA). Fielded since 1992, most recently in 2010–11, this survey is part of a continuing series of surveys (OMB No. 0925–0368) sponsored by NCI that has been administered triennially as part of the Census Bureau's and the Bureau of Labor Statistics' CPS. For the TUS–CPS, data will be collected from the U.S. civilian non-institutionalized population on smoking, other tobacco use, including switching, flavors, dependence, cessation attempts, and policy and social norms. The TUS–CPS has been a key source of national, state, some local-level, and health disparity data on these topics in U.S. households because it uses a large, nationally representative sample. The 2014–15 TUS–CPS is designed to meet both NCI's and FDA's goals. The NCI and FDA are co-sponsoring the 2014–15 TUS–CPS through parallel, but separate interagency agreements with the Census Bureau. The NCI is particularly focused on policy information such as home and workplace smoking policies, cigarette price, and impact of these on subsequent purchase and use behavior; and changes in smoking norms and attitudes. The FDA aims to support research to aid the development and evaluation of tobacco product regulations. The research findings generated from this program are expected to provide data to inform FDA regulation of the manufacture, distribution, and marketing of tobacco products to protect public health. A unique feature is the ability to link other social and economic Census Bureau and Bureau of Labor Statistics data, other sponsor-supported supplement data, and the National Longitudinal Mortality Study cancer incidence and cause-specific mortality data to the TUS–CPS data. Data will be collected in July 2014, January 2015, and May 2015 from about 255,000 respondents.

OMB approval is requested for 2 years. There are no costs to respondents other than their time. The total estimated annualized burden hours are 12,750.

ESTIMATED ANNUALIZED BURDEN HOURS

Type of respondent	Number of respondents	Responses per respondent	Average burden per response (in hour)	Annual burden hours
Individuals	127,500	1	6/60	12,750

Dated: January 15, 2014.

Vivian Horovitch-Kelley,

NCI Project Clearance Liaison, National Institutes of Health.

[FR Doc. 2014–01230 Filed 1–21–14; 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: Population Sciences and Epidemiology Integrated Review Group; Infectious Diseases, Reproductive Health, Asthma and Pulmonary Conditions Study Section.

Date: February 12–13, 2014.

Time: 8:30 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: The Dupont Circle Hotel, 1500 New Hampshire Ave. NW., Washington, DC 20036.

Contact Person: Lisa Steele, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3139, MSC 7770, Bethesda, MD 20892, (301) 257–2638, *steeleln@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Chronic Dysfunction and Integrative Neurodegeneration Study Section.

Date: February 18, 2014.

Time: 11:00 a.m. to 11:30 a.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: Samuel C Edwards, Ph.D., IRG CHIEF, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5210, MSC 7846, Bethesda, MD 20892, (301) 435–1246, *edwardss@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member conflict: Alcohol and Drugs.

Date: February 19–20, 2014.

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

Agenda: To review and evaluate grant applications.

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

Contact Person: Michael Selmanoff, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5164, MSC 7844, Bethesda, MD 20892, 301–435–1119, *selmanom@csr.nih.gov*.

Name of Committee: Genes, Genomes, and Genetics Integrated Review Group; Genomics, Computational Biology and Technology Study Section.

Date: February 19–20, 2014.

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

Agenda: To review and evaluate grant applications.

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

Contact Person: Barbara J Thomas, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2218, MSC 7890, Bethesda, MD 20892, 301–435–0603, *bthomas@csr.nih.gov*.

Name of Committee: Bioengineering Sciences & Technologies Integrated Review Group; Instrumentation and Systems Development Study Section.

Date: February 19–20, 2014.

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

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Kathryn Kalasinsky, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5158 MSC 7806, Bethesda, MD 20892, 301–402–1074, *kalasinskyks@mail.nih.gov*.

Name of Committee: Cell Biology Integrated Review Group; Intercellular Interactions Study Section.

Date: February 19, 2014.

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

Agenda: To review and evaluate grant applications.

Place: Hotel Palomar, 2121 P Street NW., Washington, DC 20037.

Contact Person: Wallace Ip, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5128, MSC 7840, Bethesda, MD 20892, 301–435–1191, *ipws@mail.nih.gov*.

Name of Committee: Biological Chemistry and Macromolecular Biophysics Integrated Review Group; Synthetic and Biological Chemistry B Study Section.

Date: February 19–20, 2014.

Time: 8:30 a.m. to 5:00 p.m.

Agenda: To review and evaluate grant applications.

Place: Serrano Hotel, 405 Taylor Street, San Francisco, CA 94102.

Contact Person: Kathryn M Koeller, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4166, MSC 7806, Bethesda, MD 20892, 301–435–2681, *koellerk@csr.nih.gov*.

Name of Committee: Surgical Sciences, Biomedical Imaging and Bioengineering Integrated Review Group; Surgery, Anesthesiology and Trauma Study Section.

Date: February 19–20, 2014.

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

Agenda: To review and evaluate grant applications.

Place: Hilton Hotel and Executive Meeting Center, 1750 Rockville Pike, Rockville, MD 20852.

Contact Person: Weihua Luo, MD, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5114, MSC 7854, Bethesda, MD 20892, (301) 435–1170, *luow@csr.nih.gov*.

(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: January 15, 2014.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2014–01119 Filed 1–21–14; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Mental Health; Notice of Closed Meeting

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 meeting.