6. Investigate associations between ZIKV infection in utero or in infancy and hearing loss and other physical, neurologic, and neurodevelopmental outcomes at 6 months of age.

7. Estimate survival of infants born to ZIKV infected mothers.

Secondary research questions we aim to address with the ZEN Colombia study are:

1. Identify risk factors for ZIKV infection in pregnant women, partners and infants. A spectrum of risk factors will be explored, including mosquito bites and mosquito bite preventive measures, sexual transmission, sociodemographic characteristics, and medical risk factors. The results of this analysis will provide information on the reduction in risk associated with adherence to recommended preventive measures and risk factors for infection in pregnant women.

2. Identify characteristics associated with taking preventive measures (mosquito bite prevention, sexual transmission) against contracting Zika virus among pregnant women and their partners. The results of this analysis will assist in targeting education or intervention to individuals at greatest risk for Zika infection.

3. Describe symptoms associated with ZIKV and estimate the positive predictive value of certain symptoms or constellations of symptoms in pregnant women, men, and infants to allow for refinement of clinical diagnosis of ZIKV infection in a setting in which testing and/or results might not be readily available.

4. Assess the duration of viremia following ZIKV infection and investigate risk factors (such as sociodemographics, comorbidities, and co-infections) associated with prolonged viremia among pregnant women, men, and infants with laboratory-confirmed ZIKV infection in blood.

The project aims to enroll approximately 5,000 women, 1,250 male partners, and 4,500 newborns. Pregnant women will be recruited in the first trimester of pregnancy for study enrollment, followed by assessments during pregnancy (every other week until 32 weeks gestation and monthly thereafter), and at or within 72 hours of delivery. At all visits, participants will complete visit-specific questionnaires. In addition to the questionnaires, at all pregnancy and delivery visits, participants will receive Colombian national recommended clinical care and provide samples for laboratory testing.

Male partners will be recruited around the time of the pregnant partners' study enrollment, followed by monthly visits until his pregnant partner reaches the third trimester (approximately 27 weeks gestation). If the male partner contracts ZIKV during this time, visits will occur every other week until the partner has two negative consecutive tests for ZIKV or the pregnancy ends. At all study visits, male partners will complete visit-specific questionnaires and provide samples for laboratory testing.

All newborns of mothers participating in the study will be followed every other week from birth to 6 months of age. At all visits, infants will receive national recommended clinical care (at birth and clinic visits at 1, 2, 3, and 6 months), provide samples for laboratory testing, and mothers will complete study-specific questionnaires about infant ZIKV symptoms. Infants will also have cranial ultrasounds at birth, their head circumference measured (birth, 72 hours, 1, 2, 3, and 6 months of age), and enhanced hearing/vision tests at 1 and 6 months old. For mothers and their infants, relevant information collected as part of clinical care will be abstracted from medical records. Study results will be used to guide recommendations made by both INS and CDC to prevent ZIKV infection; to improve counseling of patients about risks to themselves, their pregnancies, their partners, and their infants; and to help agencies prepare to provide services to affected children and families. Participation in this study is voluntary. The total estimated annualized burden hours are 19,415, and there are no costs to participants other than their time.

ESTIMATED ANNUALIZED BURDEN HOURS

Respondents	Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)
Pregnant women	Pregnant women eligibility questionnaire	3,125	1	5/60
	Pregnant women enrollment questionnaire	2,500	1	35/60
	Adult symptom questionnaire	2,500	15	10/60
	Pregnant women follow-up questionnaire	2,500	8	15/60
	Infant symptoms questionnaire	2,250	14	10/60
Male partners	Male partner eligibility questionnaire	2,500	1	5/60
	Male enrollment questionnaire	625	1	25/60
	Adult symptom questionnaire	625	7	10/60

Leroy A. Richardson,

Chief, Information Collection Review Office, Office of Scientific Integrity, Office of the Associate Director for Science, Office of the Director, Centers for Disease Control and Prevention.

[FR Doc. 2017–07879 Filed 4–18–17; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; 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/contract proposals 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/contract proposals, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Cancer Institute Special Emphasis Panel; HPV Review.

Date: May 10, 2017.

Time: 11:00 a.m. to 2:00 p.m.
Agenda: To review and evaluate contract proposals.
Place: National Cancer Institute Shady Grove, 9609 Medical Center Drive, Room 7W530, Rockville, MD 20850 (Telephone Conference Call).
Contact Person: Shamala K. Srinivas, Ph.D., Associate Director, Office of Referral, Review, and Program Coordination, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W530, Bethesda, MD 20892–9750, 240–276–6442, *ss537t@nih.gov*.

Name of Committee: National Cancer Institute Special Emphasis Panel; Cancer Immunotherapy Trials Network.
Date: May 18, 2017.
Time: 2:00 p.m. to 5:00 p.m.
Agenda: To review and evaluate grant applications.
Place: National Cancer Institute Shady Grove, 9609 Medical Center Drive, Room 7W102, Rockville, MD 20850 (Telephone Conference Call).
Contact Person: Shakeel Ahmad, Ph.D., Scientific Review Officer, Research Programs Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W102, Bethesda, MD 20892–9750, 240–276–6349, *ahmads@mail.nih.gov*.

Name of Committee: National Cancer Institute Special Emphasis Panel; Technical Evaluation Panel #1.
Date: May 23, 2017.
Time: 10:00 a.m. to 5:00 p.m.
Agenda: To review and evaluate contract proposals.
Place: National Cancer Institute Shady Grove, 9609 Medical Center Drive, Room 7W260, Rockville, MD 20850 (Telephone Conference Call).
Contact Person: Nadeem Khan, Ph.D., Scientific Review Officer, Research Technology and Contract Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W260, Bethesda, MD 20892–9750, 240–276–7684, *nadeem.khan@nih.gov*.

Name of Committee: National Cancer Institute Special Emphasis Panel; Technical Evaluation Panel #2.
Date: May 24, 2017.
Time: 10:00 a.m. to 5:00 p.m.
Agenda: To review and evaluate contract proposals.
Place: National Cancer Institute Shady Grove, 9609 Medical Center Drive, Room 7W260, Rockville, MD 20850 (Telephone Conference Call).
Contact Person: Nadeem Khan, Ph.D., Scientific Review Officer, Research Technology and Contract Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W260, Bethesda, MD 20892–9750, 240–276–7684, *nadeem.khan@nih.gov*.

Name of Committee: National Cancer Institute Special Emphasis Panel; Cancer Drug Resistance 1.
Date: June 27–28, 2017.
Time: 8:00 a.m. to 5:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Gaithersburg Marriott Washingtonian Center, 9751 Washingtonian Boulevard, Gaithersburg, MD 20878.
Contact Person: Michael B. Small, Ph.D., Chief, Program and Review Extramural Staff Training Office, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W412, Bethesda, MD 20892–9750, 240–276–6438, *smallm@mail.nih.gov*.

Name of Committee: National Cancer Institute Special Emphasis Panel; Cancer Drug Resistance 2.
Date: June 27–28, 2017.
Time: 8:00 a.m. to 3:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Gaithersburg Marriott Washingtonian Center, 9751 Washingtonian Boulevard, Gaithersburg, MD 20878.
Contact Person: Wlodek Lopaczynski, MD, Ph.D., Assistant Director, Office of the Director, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W514, Bethesda, MD 20892–9750, 240–276–6340, *lopacw@mail.nih.gov*.

Name of Committee: National Cancer Institute Initial Review Group; Subcommittee I—Transition to Independence.
Date: June 14–15, 2017.
Time: 8:00 a.m. to 1:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Crown Plaza National Airport, 1480 Crystal Drive, Arlington, VA 22202.
Contact Person: Delia Tang, MD, Scientific Review Officer, Research Technology and Contract Review Branch, Division of Extramural Activities, National Cancer Institute, NIH, 9609 Medical Center Drive, Room 7W602, Bethesda, MD 20892–9750, 240–276–6456, *tangd@mail.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.392, Cancer Construction; 93.393, Cancer Cause and Prevention Research; 93.394, Cancer Detection and Diagnosis Research; 93.395, Cancer Treatment Research; 93.396, Cancer Biology Research; 93.397, Cancer Centers Support; 93.398, Cancer Research Manpower; 93.399, Cancer Control, National Institutes of Health, HHS)

Dated: April 13, 2017.

Melanie J. Pantoja,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2017–07842 Filed 4–18–17; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

[Docket No. DHS–2017–0015]

Notice of Request for Revision to and Extension of a Currently Approved Information Collection for Chemical-Terrorism Vulnerability Information

AGENCY: National Protection and Programs Directorate, DHS.

ACTION: 60-Day Notice and request for comments; Revision of Information Collection Request: 1670–0015.

SUMMARY: The Department of Homeland Security (DHS or the Department), National Protection and Programs Directorate (NPPD), Office of Infrastructure Protection (IP), Infrastructure Security Compliance Division (ISCD), will submit the following Information Collection Request to the Office of Management and Budget (OMB) for review and clearance in accordance with the Paperwork Reduction Act of 1995. DHS proposes to remove five of the six instruments previously approved to support the Chemical-terrorism Vulnerability Information (CVI) program under the Chemical Facility Anti-terrorism Standards (CFATS) regulations, 6 CFR 27.400. DHS also proposes to extend this collection with revisions to reduce the estimated burden for the remaining instrument in this collection.

DATES: Comments are encouraged and will be accepted until June 19, 2017. This process is conducted in accordance with 5 CFR 1320.8.

ADDRESSES: Interested persons are invited to submit comments on the proposed revision to, and extension of, this approved information collection through the Federal eRulemaking Portal at *http://www.regulations.gov.* All submissions received must include the words ''Department of Homeland Security'' and the docket number DHS–2017–0015. Except as provided below, comments received will be posted without alteration at *http://www.regulations.gov,* including any personal information provided.

Comments that include trade secrets, confidential commercial or financial information, CVI,[1] Sensitive Security Information (SSI),[2] or Protected Critical Infrastructure Information (PCII) [3] should not be submitted to the public regulatory docket. Please submit such comments separately from other comments in response to this notice. Comments containing trade secrets, confidential commercial or financial information, CVI, SSI, or PCII should be appropriately marked and packaged in

[1] For more information about CVI see 6 CFR 27.400 and the CVI Procedural Manual at *http://www.dhs.gov/xlibrary/assets/chemsec_cvi_proceduresmanual.pdf.*

[2] For more information about SSI see 49 CFR part 1520 and the SSI Program Web page at *http://www.tsa.gov.*

[3] For more information about PCII see 6 CFR part 29 and the PCII Program Web page at *http://www.dhs.gov/protected-critical-infrastructure-information-pcii-program.*