information will be reported for all sites in their grantee network.

**Need and Proposed Use of the Information:**

The Title X Family Planning Program (“Title X program” or “program”) is the only Federal grant program dedicated solely to providing individuals with comprehensive family planning and related preventive health services (e.g., screening for breast and cervical cancer, sexually transmitted diseases (STDs), and human immunodeficiency virus [HIV]). By law, priority is given to persons from low-income families (Section 1006[c] of Title X of the Public Health Service Act, 42 USC 300). The Office of Population Affairs (OPA) within the Office of the Assistant Secretary for Health administers the Title X program.

In fiscal year 2013, Congress appropriated approximately $296.8 million for Title X family planning activities. In accordance with the statute and regulations (42 Code of Federal Regulations [CFR] part 59), at least 90% of the appropriation is used for clinical family planning services. In 2012, 98 Title X grantees provided family planning services to five million women and men through a network of 4,400 community-based clinics that include state and local health departments, tribal organizations, and other public and private nonprofit agencies. There is at least one clinic that receives Title X funds and provides services as required under the Title X statute in 73% of U.S. counties.

Sixty percent of the clients seen at Title X funded service sites self-identify as being uninsured. Seventy percent of the total clients are under the age 30. Thus Title X service sites see a large proportion of young and uninsured individuals. Over the past years, OPA has encouraged grantees to develop enrollment programs to ensure that clients who are currently uninsured understand new health insurance options that are available as a result of the ACA. Some sites already assist individuals with enrolling in Medicaid and other public insurance programs. With the availability of the health insurance marketplace, many more service delivery sites are assisting clients enroll in health insurance programs.

OPA does not have any data on how many sites are assisting and enrolling clients into health insurance programs. Thus we seek to collect this data in order to understand the impact of Title X funded service sites on assisting and enrolling clients into insurance programs. We will utilize this information to guide strategic planning around how Title X service sites and prepare for, and assist with, the full implementation of the ACA. Through a separate data collection process called the Family Planning Annual Report (FPAR) (OMB No. 0990–0221, expiration January 31, 2016), OPA collects information on the insurance status of the clients served. With the implementation of the ACA, many of the traditional clients served by Title X service sites will qualify for health insurance.

**Likely Respondents:**

This annual reporting requirement is for family planning services delivery projects authorized and funded by the Title X Family Planning Program.

**Burden Statement:**

Burden in this context means the time expended by persons to generate, maintain, retain, disclose or provide the information requested. This includes the time needed to review instructions, to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information, to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information, and to transmit or otherwise disclose the information. The total annual burden hours estimated for this ICR are summarized in the table below.

**TOTAL ESTIMATED ANNUALIZED BURDEN—HOURS**

<table>
<thead>
<tr>
<th>Form name</th>
<th>Number of respondents</th>
<th>Number of responses per respondent</th>
<th>Average burden per response (in hours)</th>
<th>Total burden hours</th>
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<td>4200 service sites</td>
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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Agency for Toxic Substances and Disease Registry**

[Docket No. ATSDR–2014–0002]

**Proposed Substances to be Evaluated for Set 28 Toxicological Profiles**

**AGENCY:**

Agency for Toxic Substances and Disease Registry (ATSDR), Department of Health and Human Services (HHS).

**ACTION:**

Request for comments on the proposed substances to be evaluated for Set 28 toxicological profiles.

**SUMMARY:**

ATSDR is initiating the development of its 28th set of toxicological profiles (CERCLA Set 28). This notice announces the list of proposed substances that will be evaluated for CERCLA Set 28 toxicological profile development. ATSDR’s Division of Toxicology and Human Health Sciences is soliciting public nominations from the list of proposed substances to be evaluated for toxicological profile development. ATSDR also will consider the nomination of any additional, non-CERCLA substances that may have public health implications, on the basis of ATSDR’s authority to prepare toxicological profiles for substances not found at sites on the National Priorities List. The agency will do so in order to “. . . establish and maintain inventory of literature, research, and studies on the health effects of toxic substances” under CERCLA Section 104(i)(1)(B), to respond to requests for consultation under section 104(i)(4), and to support the site-specific response actions conducted by ATSDR, as otherwise necessary.

**DATES:**

Nominations from the Substance Priority List and/or additional substances must be submitted no later than June 20, 2014.

**ADDRESSES:**

You may submit nominations, identified by Docket No. ATSDR–2014–0002, by any of the following methods:

Substances to be Evaluated for Set 28 Toxicological Profiles

Each year, ATSDR develops a list of substances to be considered for toxicological profile development. The Set 28 nomination process includes consideration of all substances on ATSDR’s Priority List of Hazardous Substances, also known as the Substance Priority List (SPL), as well as other substances nominated by the public. The 275 substances on the SPL will be considered for Set 28 Toxicological Profile development. This list may be found at the following Web site: http://www.atstdr.cdc.gov/SPL/resources.

www.atstdr.cdc.gov/toxprofiles/index.asp

Submission of Nominations for the Evaluation of Set 28 Proposed Substances: Today’s notice invites voluntary public nominations for substances included on the SPL and for substances not listed on the SPL. All nominations should include the full name of the nominator, affiliation, email address. When nominating a non-SPL substance, please include the rationale for the nomination. Please note that email addresses will not be posted on regulations.gov.

ATSDR will evaluate all data and information associated with nominated substances and will determine the final list of substances to be chosen for toxicological profile development. Substances will be chosen according to ATSDR’s specific guidelines for selection. These guidelines can be found in the Selection Criteria announced in the Federal Register on May 7, 1993 (58FR27286–27287). A hard copy of the selection criteria is available upon request or may be accessed at the following Web site: http://www.atstdr.cdc.gov/toxprofiles/guidance/criteria_for_selecting_tox_support.pdf

Please ensure that your comments are submitted within the specified nomination period. Nominations received after the closing date will be marked as late and may be considered only if time and resources permit.

Sascha Chaney,
Acting Director, Office of Policy, Planning and Evaluation, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry.

Date: June 19, 2014.
Time: 12:00 p.m. to 1:30 p.m.
Agenda: To review and evaluate grant applications.
Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892
Contact Person: Michelle Trout, Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2014–11807 Filed 5–21–14; 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: Infectious Diseases and Microbiology Integrated Review Group Clinical Research and Field Studies of Infectious Diseases Study Section.

Date: June 16, 2014.
Time: 8:00 a.m. to 6:00 p.m.
Agenda: To review and evaluate grant applications.
Place: Hotel Kabuki, 1625 Post Street, San Francisco, CA 94115.
Contact Person: Soheyla Saadi, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3211, MSC 7808, Bethesda, MD 20892, (301) 455–0903, sisadieh@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel Primate Skeletal Database.

Date: June 19, 2014.
Time: 12:00 p.m. to 1:30 p.m.
Agenda: To review and evaluate grant applications.
Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892
Contact Person: Daniel F. McDonald, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4110, MSC 7814, Bethesda, MD 20892, (301) 435–1215, mcdonald@csr.nih.gov.


Dated: May 16, 2014.

Michelle Trout,
Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2014–11848 Filed 5–21–14; 8:45 am]
BILLING CODE 4163–70–P