

Information.CollectionClearance@hhs.gov or (202) 690-6162.

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the OMB control number 0990-0313 and document identifier HHS-OS-0990-0313-30D for reference.

Information Collection Request Title: National Blood Collection and Utilization Survey OMB No.: 0990-0313.

Abstract: The National Blood Collection & Utilization Survey (NBCUS) is a biennial survey of the blood collection and utilization community (industry) to produce reliable and accurate estimates of national and regional collections, utilization, safety, and availability of all blood products, some cellular therapeutic products, as well as information on bacterial testing and human tissue transplantation that are of interest to the transfusion medicine community. The 2013 NBCUS shall be funded by the U.S. Department of Health and Human Services (DHHS) and performed by (contactor, to be determined). In previous years, the NBCUS program was performed under the auspices of the National Blood Data

Resource Center (NBDRC), a private subsidiary of AABB (formerly known as the American Association of Blood Banks), with private funding.

The survey includes a core of standard questions on blood collection, processing, and utilization practices to allow for comparison with data from previous surveys; additionally, questions to specifically address emerging and developing issues and technologies in blood collection and utilization are included. Biovigilance remains a key theme for the 2013 survey, as continued from the 2007, 2009, and 2011 iterations. To that end, questions on transfusion transmitted infections, transfusion associated circulatory overload, acute hemolysis, delayed hemolysis, and severe allergic reactions are included in the survey.

Need and Proposed Use of the Information: Under the authority of Section 301 of the Public Health Service Act (42 U.S.C. 241), as identified in the 1997 HHS Blood Action Plan, and twice in the Advisory Committee on Blood & Tissue Safety & Availability's (ACBTSA) recommendations to the Secretary, there is a need to provide national policy makers with current supply and demand data.

Likely Respondents: Respondents will include approximately 3,000 institutions that include U.S. blood collection and processing facilities, hospital-based transfusion blood banks, and cord blood banks. Participating institutions will be selected from the American Hospital Association (AHA) annual survey database and AABB member list of blood collection facilities.

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

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
National Blood Collection and Utilization Survey	3,000	1	1	3,000
Total	3,000	1	1	3,000

Darius Taylor,

Information Collection Clearance Officer.

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier HHS-OS-0990-New-30D]

Agency Information Collection Activities; Proposed Collection; Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3507(a)(1)(D) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, has submitted an

Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB) for review and approval. The ICR is for a new collection. Comments submitted during the first public review of this ICR will be provided to OMB. OMB will accept further comments from the public on this ICR during the review and approval period.

DATES: Comments on the ICR must be received on or before June 23, 2014.

ADDRESSES: Submit your comments to *OIRA_submission@omb.eop.gov* or via facsimile to (202) 395-5806.

FOR FURTHER INFORMATION CONTACT: Information Collection Clearance staff, *Information.CollectionClearance@hhs.gov* or (202) 690-6162.

Information Collection Request Title: Title X Family Planning Outreach and Enrollment Data Collection Form.

Abstract: The Office of Population Affairs within the Office of the Assistant

Secretary for Health seeks to collect data from the Title X service delivery grantees on efforts related to outreach and enrollment to assist individuals in obtaining health insurance available as a result of the Affordable Care Act (ACA). Grantees will be asked to collect and report information on the numbers of individuals who are; (1) assisted by a trained health center worker; (2) number of individuals who receive an eligibility determination for the marketplace, Medicaid or CHIP with the assistance of a trained worker; and (3) number of individuals who enroll in an insurance program with the assistance of a trained worker. For each of the data points above, respondents will have the option to break out the data for partial Medicaid (i.e. waiver programs), full Medicaid, and private marketplace plans. The detailed data will be optional for those who are able to provide it. The

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 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

Form name	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden hours
Outreach and Enrollment Activities	4200 service sites	1	0.20	840

Darius Taylor,
Information Collection Clearance Officer.
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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:

**Internet:* Access the Federal eRulemaking portal at <http://www.regulations.gov>. Follow the instructions for submitting comments.