submitted to the Commission pursuant to Subpart D of Part 14 of the Commission's rules or to any other request by the Commission may be submitted pursuant to a request for confidentiality in accordance with 47 CFR 0.459 of the Commission's rules.

Privacy Impact Assessment: The FCC completed a Privacy Impact Assessment (PIA) on June 28, 2007. The PIA may be reviewed at *http://www.fcc.gov/omd/privacyact/Privacy_Impact_Assessment.html.* The FCC is in the process of updating the PIA to incorporate various revisions made to the SORN.

Needs and Uses: On October 7, 2011, in document FCC 11–151, the FCC released a Report and Order adopting final rules to implement sections 716 and 717 of the Communications Act of 1934 (the Act), as amended, which were added to the Act by the Twenty-First Century Communications and Video Accessibility Act of 2010 (CVAA). *See* Public Law 111–260, 104. Section 716 of the Act requires providers of advanced communications services and manufacturers of equipment used for advanced communications services to make their services and equipment accessible to individuals with disabilities, unless doing so is not achievable. 47 U.S.C. 617. Section 717 of the Act establishes new recordkeeping requirements and enforcement procedures for service providers and equipment manufacturers that are subject to sections 255, 716, and 718 of the Act. 47 U.S.C. 618. Section 255 of the Act requires telecommunications and interconnected VoIP services and equipment to be accessible, if readily achievable. 47 U.S.C. 255. Section 718 of the Act requires web browsers included on mobile phones to be accessible to and usable by individuals who are blind or have a visual impairment, unless doing so is not achievable. 47 U.S.C. 619.

Among other things, the FCC established procedures in document FCC 11–151 to facilitate the filing of formal and informal complaints alleging violations of sections 255, 716, or 718 of the Act. Those procedures include a nondiscretionary pre-filing notice procedure to facilitate dispute resolution. As a prerequisite to filing an informal complaint, complainants must first request dispute assistance from the Consumer and Governmental Affairs Bureau's Disability Rights Office.

Pursuant to the new enforcement rules that will go into effect on October 8, 2013, these informal complaints will be filed on a new FCC Form 2000H. In addition, a new Request for Dispute Assistance form (FCC Form RDA) will be used to initiate the 30-day period which must precede the filing of an informal complaint. The burdens associated with filing the new 2000H and Request for Dispute Assistance forms are contained in the collection found in OMB control number 3060–0874. Therefore, the Commission extracted those burdens from the collection found in OMB control number 3060–1167. In addition, the Commission has revised its estimate of the number of requests for dispute assistance and the number of informal complaints that it expects to receive and the burdens associated with the processing and handling of those requests and complaints.

Federal Communications Commission.

Gloria J. Miles,

Federal Register Liaison, Office of the Secretary, Office of Managing Director.

[FR Doc. 2013–11879 Filed 5–17–13; 8:45 am]

BILLING CODE 6712–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS–OS–19606–60D]

Agency Information Collection Activities; Proposed Collection; Public Comment Request

AGENCY: Office of the Secretary, HHS.

ACTION: Notice.

SUMMARY: In compliance with section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Office of the Secretary (OS), Department of Health and Human Services, announces plans to submit an Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). The ICR is for extending the use of the approved information collection assigned OMB control number 0990–0221, which expires on January 31, 2014. Prior to submitting that ICR to OMB, OS seeks comments from the public regarding the burden estimate, below, or any other aspect of the ICR.

DATES: Comments on the ICR must be received on or before July 19, 2013.

ADDRESSES: Submit your comments to *Information.CollectionClearance@hhs.gov* or by calling (202) 690–6162.

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

SUPPLEMENTARY INFORMATION: When submitting comments or requesting information, please include the document identifier HHS–OS–19606–60D for reference.

Information Collection Request Title: Family Planning Annual Report: Forms and Instructions

OMB No.: 0990–0221

Abstract: The Office of Population Affairs (OPA), Office of the Assistant Secretary for Health (OASH), U.S. Department of Health and Human Services (HHS) administers and oversees the Title X Family Planning Program. The Family Planning Annual Report (FPAR) is an annual reporting requirement for family planning services delivery projects (''Title X service grantees'') authorized and funded by the Title X Family Planning Program [''Population Research and Voluntary Family Planning Programs'' (Pub. L. 91–572)], which was enacted in 1970 as Title X of the Public Health Service Act (Section 1001 of Title X of the Public Health Service Act, 42 United States Code 300). The Title X Family Planning Program is the only Federal grant program dedicated solely to providing individuals with comprehensive family planning and related preventive health services. The program's purpose is to assist individuals in determining the number and spacing of their children and is designed to provide access to contraceptive services, supplies, and information to all who want and need them. By law, priority is given to persons from low-income families (Section 1006[c] of Title X of the Public Health Service Act, 42 U.S.C. 300). The FPAR is the only source of annual, uniform reporting by all Title X service grantees. The FPAR provides consistent, national-, regional-, state-, and grantee-level data on the services provided and the characteristics of the individuals served. Note that there are no changes to the FPAR except minor corrections or clarifications to submission and reporting instructions or definitions. The estimated average hour burden has been reduced to 36 hours, which is 4 hours lower than the 40-hour estimate of the previous OMB submission.

Need and Proposed Use of the Information: OPA uses FPAR data to monitor compliance with statutory requirements and accountability and federal performance requirements for Title X family planning funds as required by the 1993 Government Performance and Results Act (GPRA) and HHS, to guide financial and program planning and evaluation, and to respond to inquiries about the program from policymakers and Congress.

Likely Respondents: Title X service grantees.

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
Family Planning Annual Report: Forms and Instructions	93	1	36	3,348
Total	93	1	36	3,348

OS specifically requests comments on (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions, (2) the accuracy of the estimated burden, (3) ways to enhance the quality, utility, and clarity of the information to be collected, and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Keith A. Tucker,

Information Collection Clearance Officer.

[FR Doc. 2013–11949 Filed 5–17–13; 8:45 am]

BILLING CODE 4150–28–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Informational Meeting Concerning Compliance With the Centers for Disease Control and Prevention's Import Permit Program; Public Webcast

AGENCY: Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS).

ACTION: Notice of public webcast.

SUMMARY: The Centers for Disease Control and Prevention (CDC), Department of Health and Human Services (HHS) announces a public webcast that will address the new import permit regulations for infectious biological agents, standards, and vectors; import permit inspections; and import permit exemptions. The purpose of this notice is to inform all interested parties, including those individuals and entities already possessing an import permit of the webcast.

DATES: The webcast will be held on Friday, July 12, 2013 from 1 p.m. to 5 p.m. EST. Those wishing to join the webcast are encouraged to register by July 5, 2013. Registration instructions are found on the HHS/CDC's Import Permit Program Web site, *http://www.cdc.gov/od/eaipp/index.htm.*

ADDRESSES: The webcast will be broadcast from the Centers for Disease Control and Prevention, 1600 Clifton Road NE., Atlanta, Georgia 30329.

FOR FURTHER INFORMATION CONTACT: Von McClee, Division of Select Agents and Toxins, Office of Public Health Preparedness and Response, Centers for Disease Control and Prevention, 1600 Clifton Road NE., MS A–46, Atlanta, GA 30333; phone: 404–718–2000; email: *lrsat@cdc.gov.*

SUPPLEMENTARY INFORMATION: On February 4, 2013, the Centers for Disease Control and Prevention (CDC) in the Department of Health and Human Services (HHS) published a final rule (78 FR 7674) amending etiological importation regulations to (1) clarify import permit regulatory definitions, (2) increase oversight by implementing inspections, (3) address exemptions and (4) describe the appeal process.

This webcast is an opportunity for the regulated community (i.e., academic institutions and biomedical centers, commercial manufacturing facilities, federal, state, and local laboratories, including clinical and diagnostic laboratories, research facilities, exhibition facilities, and educational facilities) and other interested individuals to obtain specific regulatory guidance and information regarding the newly amended regulations. The webcast will also provide assistance to those interested in applying for an etiological agent import permit. Representatives from HHS/CDC will be present during the webcast to address questions and concerns from the web participants.

Topics to be discussed during the webcast include: The new import permit regulations for infectious biological agents, standards, and vectors; import permit inspections; and import permit exemptions. A question and answer session will take place after each topic.

Individuals wishing to join the webcast are encouraged to register by July 5, 2013. Instructions for registration are found on the HHS/CDC's Import Permit Program Web site, *http://www.cdc.gov/od/eaipp/index.htm.* This is a webcast only event and there will be no on-site participation at the HHS/CDC broadcast facility. In-person participation cannot be accommodated. Closed-captioning video of the webcast will be available at *http://www.cdc.gov/od/eaipp/index.htm* after the webcast.

Dated: May 14, 2013.

Tanja Popovic,

Deputy Associate Director for Science, Centers for Disease Control and Prevention.

[FR Doc. 2013–11895 Filed 5–17–13; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

Eunice Kennedy Shriver National Institute of Child Health & Human Development; 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 meeting.

The meeting 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