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