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: Center for Scientific Review Special Emphasis Panel; Member Conflict: Neuroblastomas, Glioblastomas, and Multiple Sclerosis and Viruses.

Date: April 2, 2015.

Time: 10:00 a.m. to 11:30 a.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Telephone Conference Call).

Contact Person: Samuel C Edwards, Ph.D., IRG CHIEF, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5210, MSC 7846, Bethesda, MD 20892, (301) 435–1246, edwardss@csr.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Neuroblastomas, Glioblastomas, and Multiple Sclerosis and Viruses.

Date: April 9, 2015.

Time: 11:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Mark Caprara, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5156, MSC 7844, Bethesda, MD 20892, 301–435–1042, capramm@mail.nih.gov.

Name of Committee: Center for Scientific Review Special Emphasis Panel; Member Conflict: Reproductive Biology.

Date: April 10, 2015.

Time: 12:00 p.m. to 3:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, 6701 Rockledge Drive, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Michael Knecht, Ph.D., Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6176, MSC 7892, Bethesda, MD 20892, (301) 435–1046, knechtm@csr.nih.gov.


Dated: March 6, 2015.

Anna Snouffer,
Deputy Director, Office of Federal Advisory Committee Policy.

[SFR Doc. 2015–05566 Filed 3–11–15; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

[Document Identifier: HHS–OS–0990—New–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 a new Information Collection Request (ICR), described below, to the Office of Management and Budget (OMB). Prior to submitting the 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 May 11, 2015.

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–0990—New–60D for reference.

Information Collection Request Title: Privacy and Security Capacity Assessment of the Title X Network.

Abstract: The Office of the Assistant Secretary for Health Office of Population Affairs, (OPA) is requesting an approval by Office of Management and Budget (OMB) for a new information collection (Privacy and Security Capacity Assessment) which seeks to collect feedback from the Title X network regarding Title X grantees' and service sites' current privacy and security capabilities for health information exchange. This voluntary form will be administered at most annually and enable the Title X network to share important information to critically inform OPA's development of Family Planning Annual Report (FPAR 2.0), as well as identify any training assistance and inform guidance that OPA may offer in the future. OPA will solicit feedback from Title X agencies to advise our work on privacy and security, and proposes to make this data collection form available for up to 3 years so that OPA can accept feedback from the network regarding any changes or trends that might alter our approach to privacy and security as we proceed through the design and build process for the planned FPAR 2.0 data repository.

Likely Respondents: Title X Grantees, Sub recipients, and Service Sites.

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>273</td>
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<td>Total</td>
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<td>1</td>
<td>20/60</td>
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OPA 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.

Terry S. Clark,
Deputy Information Collection Clearance Officer.

[FR Doc. 2015–05577 Filed 3–11–15; 8:45 am] BILLING CODE 4150–28–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Disease, Disability, and Injury Prevention and Control Special Emphasis Panel (SEP): Initial Review

In accordance with Section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92–463), the Centers for Disease Control and Prevention (CDC) announces a meeting for the initial review of applications in response to [FOA] GH15–001, Conducting Public Health Research in Kenya, Funding Opportunity Announcement.

Time and Date: 12:00 p.m.—5:00 p.m., April 1, 2015 (Closed).

Place: Teleconference.

Status: The meeting will be closed to the public in accordance with provisions set forth in Section 552b(c)(4) and (6), Title 5 U.S.C., and the Determination of the Director, Management Analysis and Services Office, CDC, pursuant to Public Law 92–463.

Matters for Discussion: The meeting will include the initial review, discussion, and evaluation of applications received in response to “Conducting Public Health Research in Kenya, FOA GH15–001”.

Contact Person for More Information: Hylan Shoob, Scientific Review Officer, Center for Global Health (CGH) Science Office, CGH, CDC, 1600 Clifton Road, NE., Mailstop D–69, Atlanta, Georgia 30033, Telephone: (404) 639–4796.

The Director, Management Analysis and Services Office, has been delegated the authority to sign Federal Register notices pertaining to announcements of meetings and other committee management activities, for both the Centers for Disease Control and Prevention and the Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,
Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2015–05577 Filed 3–11–15; 8:45 am] BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Food and Drug Administration Joint Conference

[DOCKET NO. FDA–2015–N–0001]

2015 Parenteral Drug Association/Food and Drug Administration Joint Conference

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice of public conference.

The Food and Drug Administration (FDA), is announcing a public conference, to be held in co-sponsorship with the Parenteral Drug Association (PDA), entitled “Mission Possible: Patient-Focused Manufacturing, Quality, and Regulatory Solutions.” The conference will cover current issues affecting the industry as well as explore strategies to facilitate the development and continuous improvement of safe and effective medical products. The conference establishes a unique forum to discuss the foundations, emerging technologies, and innovations in regulatory science, as well as the current quality and compliance areas of concerns. Meeting participants will hear from FDA and industry speakers about the requirements and best practices to consider while implementing robust quality systems in order to deliver the best quality product.

Date and Time: The public conference will be held on September 28, 2015, from 7 a.m. to 7:30 p.m.; September 29, 2015, from 7 a.m. to 9:30 p.m.; and September 30, 2015, from 7 a.m. to 12:30 p.m.


Contact: Wanda Neal, Parenteral Drug Association, PDA Global Headquarters, Bethesda Towers, 4350 East West Hwy., Suite 150, Bethesda, MD 20814, 301–656–5900, ext. 111, FAX: 301–986–1093, email: info@pda.org; or Ken Nolan, Office of Communications, Food and Drug Administration, 10903 New Hampshire Ave., Silver Spring, MD 20993, 301–796–8629, email: kenneth.nolan@fda.hhs.gov.

Accommodations: Attendees are responsible for their own accommodations. To make reservations, contact the Renaissance Washington Hotel (see Location) and reference “the 2015 FDA/PDA Joint Regulatory Conference” to receive the PDA group rate. Room rates are: Single: $305 plus 14.5 percent State and local taxes. Requests will be processed on a first-come, first-served basis.

Registration: Attendees are encouraged to register at their earliest convenience. The PDA registration fees cover the cost of facilities, materials, and refreshments. Seats are limited; please submit your registration as soon as possible. Conference space will be filled in order of receipt of registration. Those accepted for the conference will receive confirmation. Registration will close after the conference is filled. Onsite registration will be available on a space available basis beginning at 1 p.m. on September 27, 2015, and at 7 a.m. from September 28 through 30, 2015. The cost of registration is as follows:

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Premier Package (Includes Conference and Workshop Registration)

Conference Only

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