the Paperwork Reduction Act of 1995 (44 U.S.C. 3501–3520). The collections of information in 21 CFR 601.12 have been approved under OMB control number 0910–0338; the collections of information in 21 CFR 606.171 have been approved under OMB control number 0910–0458; and the collections of information in 21 CFR 610.46, 630.6, 640.3 and 640.63 have been approved under OMB control number 0910–0116.

III. Comments

The draft guidance is being distributed for comment purposes only and is not intended for implementation at this time. Interested persons may submit either electronic comments regarding this document to *http://www.regulations.gov* or written comments to the Division of Dockets Management (see **ADDRESSES**). It is only necessary to send one set of comments. Identify comments with the docket number found in brackets in the heading of this document. Received comments may be seen in the Division of Dockets Management between 9 a.m. and 4 p.m., Monday through Friday, and will be posted to the docket at *http://www.regulations.gov.*

IV. Electronic Access

Persons with access to the Internet may obtain the draft guidance at either *http://www.fda.gov/BiologicsBloodVaccines/GuidanceComplianceRegulatoryInformation/Guidances/default.htm* or *http://www.regulations.gov.*

Dated: May 11, 2015.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2015–11690 Filed 5–14–15; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA–2014–N–0487]

Agency Information Collection Activities; Announcement of Office of Management and Budget Approval; Generic Clearance for the Collection of Qualitative Feedback on Food and Drug Administration Service Delivery

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a collection of information entitled, ''Generic Clearance for the Collection of Qualitative Feedback on Food and Drug Administration Service Delivery'' has been approved by the Office of Management and Budget (OMB) under the Paperwork Reduction Act of 1995.

FOR FURTHER INFORMATION CONTACT: FDA PRA Staff, Office of Operations, Food and Drug Administration, 8455 Colesville Rd., COLE–14526, Silver Spring, MD 20993–0002, *PRAStaff@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: On September 24, 2014, the Agency submitted a proposed collection of information entitled, ''Generic Clearance for the Collection of Qualitative Feedback on Food and Drug Administration Service Delivery'' to OMB for review and clearance under 44 U.S.C. 3507. An Agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number. OMB has now approved the information collection and has assigned OMB control number 0910–0697. The approval expires on November 30, 2017. A copy of the supporting statement for this information collection is available on the Internet at *http://www.reginfo.gov/public/do/PRAMain.*

Dated: May 11, 2015.

Leslie Kux,

Associate Commissioner for Policy.

[FR Doc. 2015–11689 Filed 5–14–15; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

U.S. National Authority for the WHO Global Code of Practice on the International Recruitment of Health Personnel; Notice of Public Meeting

Time and date: Meeting will be held on June 10th, 2015, 10:00 a.m. to 12:00 p.m. EDT.

Place: Room 405A, U.S. Department of Health & Human Services, 200 Independence Ave. SW., Washington, DC 20201.

Status: Open, but requiring RSVP to *us.who.irhp@hhs.gov* by June 3rd, 2015.

Purpose: The purpose of the World Health Organization (WHO) Global Code of Practice on International Recruitment of Health Personnel (Global Code) is ''to establish and promote voluntary principles and practices for the ethical international recruitment of health personnel and to facilitate the strengthening of health systems.'' The United States Government has designated the Office of Global Affairs (OGA) and the Health Resources and Services Administration (HRSA) as co-National Authority to be the point of contact for implementation activities. The Global Code encourages WHO member states to cooperate with all relevant stakeholders in their implementation efforts. This meeting is thus intended to provide an update to all interested stakeholders on U.S. Global Code implementation efforts to date and to provide a forum for questions on activities related to implementation of the Global Code.

The meeting will be open to the public as indicated above, with attendance limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify within their RSVP at least 10 business days prior to the meeting. Foreign nationals planning to attend the session in person will require additional paperwork for security clearance and that this clearance process requires a minimum of 10 business days.

RSVP: Due to security restrictions for entry into the HHS Humphrey Federal Building, we will need to receive RSVPs for this event. Please send your full name and organization to *us.who.irhp@hhs.gov.* If you are *not* a U.S. citizen, you must RSVP no later than May 26th, 2015. Please note this in the subject line of your RSVP, and our office will contact you to gain additional biographical information for your clearance. For U.S. citizens, please RSVP no later than Friday, June 3rd, 2015. Written comments are welcome and encouraged, even if you are planning to attend in person. Please send these to the email address: *us.who.irhp@hhs.gov.*

Dated: May 7, 2015.

Jimmy Kolker,

Assistant Secretary for Global Affairs, Department of Health and Human Services.

[FR Doc. 2015–11785 Filed 5–14–15; 8:45 am]

BILLING CODE 4150–38–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: Center for Scientific Review Special Emphasis Panel; Collaborative Applications: Behavioral Genetics and Epidemiology.

Date: June 2–3, 2015.

Time: 8:30 a.m. to 5:00 p.m.