provide an update on the current process and to gather additional input on such a program.

Date and Time: The public meeting will be held on May 10, 2011, from 2 p.m. to 3:30 p.m.

Location: The public meeting will be held at FDA's White Oak Campus, 10903 New Hampshire Ave., Bldg. 1, Conference Rooms 4101, 4103, and 4105, Silver Spring, MD 20993–0002.

Contact Person: Mari Long, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 4237, Silver Spring, MD 20993–0002, 301–796–7574, Fax 301–847–3541, *mari.long@fda.hhs.gov;* or

Peter C. Beckerman, Office of Policy, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 32, rm. 4238, Silver Spring, MD 20993–0002, 301–796–4830, Fax 301–847–3541, *peter.beckerman@fda.hhs.gov.*

Registration and Requests for Oral Presentations: If you wish to attend and/or present at the meeting, please e-mail your registration information to *GDUFA_Meeting2@fda.hhs.gov* by May 3, 2011. Your e-mail should contain complete contact information for each attendee, including name, title, affiliation, address, e-mail address, and telephone number. Registration is free and will be on a first-come, first-served basis. Early registration is recommended because seating is limited. FDA may limit the number of participants from each organization as well as the total number of participants, based on space limitations. Registrants will receive confirmation once they have been accepted. Onsite registration on the day of the meeting will be based on space availability. We will try to accommodate all persons who wish to make a presentation. The time allotted for presentations may depend on the number of persons who wish to speak, and if the entire meeting time is not needed for presentations, FDA reserves the right to terminate the meeting early.

If you need special accommodations because of disability, please contact Mari Long or Peter Beckerman (*see Contact Person*) at least 7 days before the meeting.

Comments: Regardless of attendance at the public meeting, interested persons may submit either electronic or written comments regarding this document by June 10, 2011. To ensure consideration, all comments must be received by June 10, 2011. Submission of comments prior to the meeting is strongly encouraged. Submit any comments that you plan to present at the public meeting to the docket by the date of the public meeting, but note that either electronic or written comments generally may be submitted until June 10, 2011.

Submit electronic comments to *http://www.regulations.gov.* Submit written comments to the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. It is only necessary to send one set of comments. It is no longer necessary to send two copies of mailed 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.

SUPPLEMENTARY INFORMATION:

I. Background

FDA is announcing its intention to hold a public meeting related to generic drug user fees. The Agency continues to solicit comment on whether to seek a user fee program that would provide additional resources for the review of human generic drug applications, as well as what such a program should look like. New legislation would be required for FDA to establish and collect user fees for generic drugs, and FDA is currently engaged in negotiations with industry over aspects of a joint proposal for a generic drug user fee program, including fees and performance goals. Because FDA can only negotiate with trade organizations, not individual companies, but remains interested in hearing from non-affiliated companies in addition to patient and consumer stakeholders, the Agency will hold a public meeting. The public meeting will provide a status update and seek input from stakeholders on generic drug user fees. In addition, FDA continues to encourage all interested stakeholders to submit either electronic or written comments to the docket (*see Comments*).

II. What information should you know about the public meeting, when and where will the public meeting occur, and what format will FDA use?

Through this notice, we are announcing a public meeting to update stakeholders and hear stakeholder views on what features FDA should propose for a generic drug user fee program. We will conduct the meeting on May 10, 2011, from 2 p.m. to 3:30 p.m. at FDA's White Oak Campus, 10903 New Hampshire Ave., Bldg. 1, Conference Rooms 4101, 4103, and 4105, Silver Spring, MD 20993–0002. In general, the meeting format will include a presentation by FDA and presentations by stakeholders and members of the public who have registered in advance to present at the meeting. The amount of time available for presentations will be determined by the number of people who register to make a presentation. We will also provide an opportunity for organizations and individuals to submit either electronic or written comments to the docket after the meeting (*see Comments*). FDA policy issues are beyond the scope of this initiative. Accordingly, the presentations should focus on process and funding issues, and not focus on policy.

Dated: April 26, 2011.

Leslie Kux,

Acting Assistant Commissioner for Policy.

[FR Doc. 2011–10382 Filed 4–28–11; 8:45 am]

BILLING CODE 4160–01–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, Brain Disorders Pathology and Treatment.

Date: May 26, 2011.

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

Agenda: To review and evaluate grant applications.

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

Contact Person: Dan D. Gerendasy, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3218, MSC 7843, Bethesda, MD 20892. 301–408–9164. *gerendad@csr.nih.gov.*

Name of Committee: Center for Scientific Review Special Emphasis Panel, Member Conflict: Cancer Biology and Therapeutics.

Date: June 1–2, 2011.

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

Agenda: To review and evaluate grant applications.

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

Contact Person: Malaya Chatterjee, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 6192, MSC 7804, Bethesda, MD 20892. (301) 806–2515. *chatterm@csr.nih.gov*.

Name of Committee: Surgical Sciences, Biomedical Imaging and Bioengineering Integrated Review Group, Clinical Molecular Imaging and Probe Development.

Date: June 1–2, 2011.

Time: 7 p.m. to 1 p.m.

Agenda: To review and evaluate grant applications.

Place: Hotel Valencia Riverwalk, 150 East Houston Street, San Antonio, TX 78205.

Eileen W. Bradley, DSC, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5100, MSC 7854, Bethesda, MD 20892. (301) 435–1179. *bradleye@csr.nih.gov*.

Name of Committee: Musculoskeletal, Oral and Skin Sciences Integrated Review Group, Musculoskeletal Tissue Engineering Study Section.

Date: June 6–7, 2011.

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

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Jean D. Sipe, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 4106, MSC 7814, Bethesda, MD 20892. 301/435–1743. *smithbf@auburn.edu*.

Name of Committee: Integrative, Functional and Cognitive Neuroscience Integrated Review Group, Somatosensory and Chemosensory Systems Study Section.

Datea: June 7–8, 2011.

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

Agenda: To review and evaluate grant applications.

Place: Latham Hotel, 3000 M Street, NW., Washington, DC 20007.

Contact Person: M. Catherine Bennett, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5182, MSC 7846, Bethesda, MD 20892. 301–435–1766. *bennettc3@csr.nih.gov*.

Name of Committee: Healthcare Delivery and Methodologies Integrated Review Group, Biomedical Computing and Health Informatics Study Section.

Date: June 7–8, 2011.

Time: 8 a.m. to 4 p.m.

Agenda: To review and evaluate grant applications.

Place: Sheraton Delfina Santa Monica Hotel, 530 West Pico Boulevard, Santa Monica, CA 90405.

Contact Person: Melinda Jenkins, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3156, MSC 7770, Bethesda, MD 20892. 301–437–7872. *jenkinsml2@mail.nih.gov*.

Name of Committee: Digestive, Kidney and Urological Systems Integrated Review Group, Pathobiology of Kidney Disease Study Section.

Date: June 7–8, 2011.

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

Agenda: To review and evaluate grant applications.

Place: Fairmont San Francisco Hotel, 950 Mason Street, San Francisco, CA 94108.

Contact Person: Atul Sahai, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2188, MSC 7818, Bethesda, MD 20892. 301–435–1198. *sahaia@csr.nih.gov*.

Name of Committee: Integrative, Functional and Cognitive Neuroscience Integrated Review Group, Sensorimotor Integration Study Section.

Date: June 7, 2011.

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

Agenda: To review and evaluate grant applications.

Place: One Washington Circle Hotel, One Washington Circle, NW., Washington, DC 20037.

John Bishop, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 5182, MSC 7844, Bethesda, MD 20892. (301) 408–9664. *bishopj@csr.nih.gov*.

Name of Committee: Endocrinology, Metabolism, Nutrition and Reproductive Sciences Integrated Review Group, Pregnancy and Neonatology Study Section.

Date: June 7, 2011.

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

Agenda: To review and evaluate grant applications.

Place: Hyatt Regency Bethesda, One Bethesda Metro Center, 7400 Wisconsin Avenue, Bethesda, MD 20814.

Contact Person: Michael Knecht, PhD, 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*.

Name of Committee: Genes, Genomes, and Genetics Integrated Review Group, Genetics of Health and Disease Study Section.

Date: June 7–8, 2011.

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

Agenda: To review and evaluate grant applications.

Place: The Fairmont Washington, DC, 2401 M Street, NW., Washington, DC 20037.

Contact Person: Richard Panniers, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2212, MSC 7890, Bethesda, MD 20892. (301) 435–1741. *pannierr@csr.nih.gov*.

Name of Committee: Healthcare Delivery and Methodologies Integrated Review Group, Nursing and Related Clinical Sciences Study Section.

Date: June 7–9, 2011.

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

Agenda: To review and evaluate grant applications.

Place: Sheraton Delfina Santa Monica Hotel, 530 West Pico Boulevard, Santa Monica, CA 90405.

Contact Person: Priscah Mujuru, RN, DRPH, COHNS, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3139, MSC 7770, Bethesda, MD 20892. 301–594–6594. *mujurup@mail.nih.gov*.

Name of Committee: Healthcare Delivery and Methodologies Integrated Review Group, Dissemination and Implementation Research in Health Study Section.

Date: June 7, 2011.

Time: 8:30 a.m. to 6 p.m.

Agenda: To review and evaluate grant applications.

Place: Sheraton Delfina Santa Monica Hotel, 530 West Pico Boulevard, Santa Monica, CA 90405.

Contact Person: Jacinta Bronte-Tinkew, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3164, MSC 7770, Bethesda, MD 20892. (301) 806–0009. *brontetinkewjm@csr.nih.gov*.

Name of Committee: Center for Scientific Review Special Emphasis Panel, Collaborative Applications in Child Psychopathology.

Date: June 7, 2011.

Time: 11 a.m. to 5 p.m.

Agenda: To review and evaluate grant applications.

Place: Doubletree Hotel Washington, 1515 Rhode Island Avenue, NW., Washington, DC 20005.

Contact Person: Jane A. Doussard-Roosevelt, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3184, MSC 7848, Bethesda, MD 20892. (301) 435–4445. *doussarj@csr.nih.gov*.

Name of Committee: Digestive, Kidney and Urological Systems Integrated Review Group, Xenobiotic and Nutrient Disposition and Action Study Section.

Date: June 8, 2011.

Time: 8 a.m. to 6 p.m.

Agenda: To review and evaluate grant applications.

Place: Sir Francis Drake Hotel, 450 Powell Street, San Francisco, CA 94102.

Contact Person: Patricia Greenwel, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 2172, MSC 7818, Bethesda, MD 20892. 301–435–1169. *greenwep@csr.nih.gov*.

Name of Committee: Healthcare Delivery and Methodologies Integrated Review Group, Community Influences on Health Behavior.

Date: June 8–9, 2011.

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

Agenda: To review and evaluate grant applications.

Place: Sheraton Delfina Santa Monica Hotel, 530 West Pico Boulevard, Santa Monica, CA 90405.

Contact Peson: Wenchi Liang, PhD, Scientific Review Officer, Center for Scientific Review, National Institutes of Health, 6701 Rockledge Drive, Room 3150, MSC 7770, Bethesda, MD 20892. 301–435–0681. *liangw3@csr.nih.gov*.

(Catalogue of Federal Domestic Assistance Program Nos. 93.306, Comparative Medicine; 93.333, Clinical Research; 93.306, 93.333, 93.337, 93.393–93.396, 93.837–93.844, 93.846–93.878, 93.892, 93.893, National Institutes of Health, HHS)

Dated: April 22, 2011.

Anna P. Snouffer,

Deputy Director, Office of Federal Advisory Committee Policy.

[FR Doc. 2011–10297 Filed 4–28–11; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

[Docket ID FEMA–2011–0012]

Agency Information Collection Activities: Proposed Collection; Comment Request, 1660–NEW; Level 1 Assessment and Level III Evaluations for the Center for Domestic Preparedness (CDP)

AGENCY: Federal Emergency Management Agency, DHS.

ACTION: Notice; 60-day notice and request for comments; new information collection; OMB No. 1660–NEW; FEMA Form 092–0–2, Level 1 Assessment Form; FEMA Form 092–0–2A, Level 3 Evaluation Form for Students; FEMA Form 092–0–2B, Level 3 Evaluation Form for Supervisors.

SUMMARY: The Federal Emergency Management Agency, as part of its continuing effort to reduce paperwork and respondent burden, invites the general public and other Federal agencies to take this opportunity to comment on a proposed new information collection. In accordance with the Paperwork Reduction Act of 1995, this notice seeks comments concerning course evaluations for the Center for Domestic Preparedness.

DATES: Comments must be submitted on or before June 28, 2011.

ADDRESSES: To avoid duplicate submissions to the docket, please use only one of the following means to submit comments:

(1) *Online.* Submit comments at *http://www.regulations.gov* under Docket ID FEMA–2011–0012. Follow the instructions for submitting comments.

(2) *Mail.* Submit written comments to Docket Manager, Office of Chief Counsel, DHS/FEMA, 500 C Street, SW., Room 835, Washington, DC 20472–3100.

(3) *Facsimile.* Submit comments to (703) 483–2999.

(4) *E-mail.* Submit comments to *FEMA-POLICY@dhs.gov.* Include Docket ID FEMA–2011–0012 in the subject line.

All submissions received must include the agency name and Docket ID. Regardless of the method used for submitting comments or material, all submissions will be posted, without change, to the Federal eRulemaking Portal at *http://www.regulations.gov,* and will include any personal information you provide. Therefore, submitting this information makes it public. You may wish to read the Privacy Act notice that is available via the link in the footer of *http://www.regulations.gov.*

FOR FURTHER INFORMATION CONTACT: Linda S. Pressley, Assistant Director of Analysis and Evaluation, Center for Domestic Preparedness, 256–847–2685 for additional information. You may contact the Records Management Division for copies of the proposed collection of information at facsimile number (202) 646–3347 or e-mail address: *FEMA-Information-Collections-Management@dhs.gov.*

SUPPLEMENTARY INFORMATION: The Center for Domestic Preparedness (CDP) is required by Congress to identify, develop, test, and deliver training to State, local, and Tribal emergency response providers, provide on-site and mobile training at the performance and management and planning levels, and facilitate the delivery of training by the training partners of the Department of Homeland Security pursuant to Section 1204 of the Implementing Recommendations of the 9/11 Commission Act of 2007, Public Law 110–53, 121 Stat. 266, August 3, 2007 (codified at 6 U.S.C. 1102). The collection of this data will help facilitate that Congressional mandate.

Collection of Information

Title: Level 1 Assessment and Level III Evaluations for the Center for Domestic Preparedness (CDP).

Type of Information Collection: New information collection.

OMB Number: OMB No. 1660–NEW.

Form Titles and Numbers: FEMA Form 092–0–2, Level 1 Assessment Form; FEMA Form 092–0–2A, Level 3 Evaluation Form for Students; FEMA Form 092–0–2B, Level 3 Evaluation Form for Supervisors.

Abstract: The forms in this collection of information will be used to survey the Center for Domestic Preparedness (CDP) students (and their supervisors) enrolled in CDP courses. The survey will collect information regarding quality of instruction, course material, and impact of training on their professional employment.

Affected Public: State, local or Tribal government.

Estimated Total Annual Burden Hours: 18,000 hours.

ANNUAL HOUR BURDEN

Data collection activity/instrument	Number of respondents	Frequency of responses	Hour burden per response	Annual responses	Total annual burden hours
Level 1 Assessment Form, FEMA Form 092–0–2	52,000	1	15 minutes (.25 hours)	52,000	13,000
Level 1 Assessment Form, FEMA Form 092–0–2	13,000	1	15 minutes (.25 hours)	13,000	3,250
Level 3 Evaluation Form for Students, FEMA Form 092–0–2A.	1,500	1	15 minutes (.25 hours)	1,500	375
Level 3 Evaluation Form for Students, FEMA Form 092–0–2A.	2,000	1	15 minutes (.25 hours)	2,000	500
Level 3 Evaluation Form for Supervisors, FEMA Form 092–0–2B.	1,500	1	15 minutes (.25 hours)	1,500	375
Level 3 Evaluation Form for Supervisors, FEMA Form 092–0–2B.	2,000	1	15 minutes (.25 hours)	2,000	500
Level 1 Assessment Form, FEMA Form 092–0–2	52,000	1	15 minutes (.25 hours)	52,000	13,000
Total	72,000				18,000