DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Clinical Laboratory Improvement Advisory Committee (CLIAC)

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 the following meeting of the aforementioned committee:

Times and Dates: 8:30 a.m.–5 p.m., August 31, 2011.
8:30 a.m.–12 p.m., September 1, 2011.

Place: CDC, 1600 Clifton Road, NE., Tom Harkin Global Communications Center, Building 19, Room 232, Auditorium B, Atlanta, Georgia 30333.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 100 people.

Purpose: This Committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services, the Assistant Secretary for Health, and the Director, CDC, regarding the need for, and the nature of, revisions to the standards under which clinical laboratories are regulated; the impact on medical and laboratory practice of proposed revisions to the standards; and the modification of the standards to accommodate technological advances.

Matters to be Discussed: The agenda will include agency updates from the CDC, the Centers for Medicare & Medicaid Services (CMS), and the Food and Drug Administration (FDA); presentations and discussions on the laboratory’s role in the development and use of electronic health records, electronic laboratory reporting for notifiable diseases, and meaningful use; and presentations and discussion on current practices in gynecologic cytology testing. Agenda items are subject to change as priorities dictate.

Online Registration Required: In order to expedite the security clearance processing at the CDC Royal Chemical Laboratory in Clifton, all CLIAC attendees are required to register for the meeting online at least 14 days in advance at http://www.cdc.gov/cliac/default.aspx by clicking the “Register for a Meeting” link and completing all forms according to the instructions given. Please complete all the required fields before submitting your registration and submit no later than August 16, 2011.

Providing Oral or Written Comments: It is the policy of CLIAC to accept written public comments and provide a brief period for oral public comments whenever possible. Oral Comments: In general, each individual or group requesting to make an oral presentation will be limited to a total time of five minutes (unless otherwise indicated). Speakers must also submit their comments in writing for inclusion in the meeting’s Summary Report. To assure adequate time is scheduled for public comments, individuals or groups planning to make an oral presentation should, when possible, notify the contact person below at least one week prior to the meeting date. Written Comments: For individuals or groups unable to attend the meeting, CLIAC accepts written comments until the date of the meeting (unless otherwise stated); however, it is requested that comments be submitted at least one week prior to the meeting date so that the comments may be made available to the Committee for their consideration and public distribution. Written comments, one hard copy with original signature, should be provided to the contact person below. Written comments will be included in the meeting’s Summary Report.

Contact Person for Additional Information: Nancy Anderson, Chief, Laboratory Practice Standards Branch, Division of Laboratory Science and Standards, Laboratory Science, Policy and Practice Program Office (LSPPPO), Office of Surveillance, Epidemiology and Laboratory Services, CDC, 1600 Clifton Road, NE., Mailstop F–11, Atlanta, Georgia 30333; telephone (404) 498–2741; fax (404) 498–2219; or via e-mail at Nancy.Anderson@cdc.gov.

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 the Centers for Disease Control and Prevention, and the Agency for Toxic Substances and Disease Registry.

Dated: June 30, 2011.

Elizabeth Millington,
Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2011–17009 Filed 7–6–11; 8:45 am]
BILLING CODE 4163–18–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

The meeting announced below concerns Human Immunodeficiency Virus (HIV) Prevention Projects for Young Men of Color Who Have Sex with Men and Young Transgender Persons of Color, Funding Opportunity Announcement (FOA) PS11–1113, 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 the aforementioned meeting:

Times and Dates: 8 a.m.–7 p.m., July 22, 2011 (Closed).
Place: Corporate Square, Building 8, Atlanta, Georgia 30333.

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 to be Discussed: The meeting will include the initial review, discussion, and evaluation of applications received in response to “HIV Prevention Projects for Young Men of Color Who Have Sex with Men and Young Transgender Persons of Color, FOA PS11–1113.” This subsequent meeting to the July 10–13, 2011 meeting published in the Federal Register on February 22, 2011, Volume 76, Number 35, Pages 9763–9786 has been scheduled due to the high volume of applications received and unanticipated scheduling conflicts for a significant number of the appointed reviewers.

Contact Person for More Information: Harriette Lynch, Public Health Analyst, Extramural Programs, National Center for HIV, Hepatitis and Sexually Transmitted Diseases Prevention, CDC, 1600 Clifton Road, NE., Mailstop E–60, Atlanta, Georgia 30333, Telephone (404) 498–2726, E-mail HILynch@cdc.gov.

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.

Dated: June 30, 2011.

Elizabeth Millington,
Acting Director, Management Analysis and Services Office, Centers for Disease Control and Prevention.

[FR Doc. 2011–17008 Filed 7–6–11; 8:45 am]
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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention (CDC)

Request for Nominations of Candidates To Serve on the Board of Scientific Counselors, National Center for Environmental Health/Agency for Toxic Substances and Disease Registry (BSC, NCEH/ATSDR), Centers for Disease Control and Prevention, Department of Health and Human Services (HHS)

The NCEH/ATSDR is soliciting nominations for consideration of membership on the BSC. The BSC, NCEH/ATSDR provides advice and guidance to the Secretary, HHS; the Director, CDC; and the Director, NCEH/
impartiality, as defined by Federal activities, or the appearance of a lack of Employee and private interests and responsibilities as a Special Government Officers Serving on Financial Disclosure Form for Special.

Centers for Disease Control and Administration asked to submit the “Confidential Government Employees Serving on financial information has been submitted to the Paperwork Reduction Act of 1995 (PRA).

The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995 (PRA).