

Dated: February 25, 2016.

Lorin S. Curit,

Director, Federal Acquisition Policy Division,
Office of Governmentwide Acquisition Policy,
Office of Acquisition Policy, Office of
Governmentwide Policy.

[FR Doc. 2016-04485 Filed 3-1-16; 8:45 am]

BILLING CODE 6820-EP-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 Childhood Obesity Research Demonstration 2.0, FOA DP 16-004, initial review.

SUMMARY: This document corrects a notice that was published in the **Federal Register** on February 10, 2016, Volume 81, Number 27, pages 7123-7124. The meeting time and date should read as follows:

Time and Date: 10:00 a.m.–6:00 p.m., EDT, March 15, 2016 (Closed).

FOR FURTHER INFORMATION CONTACT: Jaya Raman, Ph.D., Scientific Review Officer, CDC, 4770 Buford Highway NE., Mailstop F80, Atlanta, Georgia 30341, Telephone: (770) 488-6511, KVA5@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.

Elaine L. Baker,

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

[FR Doc. 2016-04591 Filed 3-1-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Clinical Laboratory Improvement Advisory Committee

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 committee meeting.

Times And Dates: 8:30 a.m.–5:00 p.m., April 13, 2016; 8:30 a.m.–12:00 p.m., April 14, 2016.

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

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 100 people. This meeting will also be webcast, please see information below.

Purpose: This Committee is charged with providing scientific and technical advice and guidance to the Secretary of Health and Human Services (HHS); the Assistant Secretary for Health; the Director, Centers for Disease Control and Prevention; the Commissioner, Food and Drug Administration (FDA); and the Administrator, Centers for Medicare and Medicaid Services (CMS). The advice and guidance pertain to general issues related to improvement in clinical laboratory quality and laboratory medicine practice and specific questions related to possible revision of the Clinical Laboratory Improvement Amendment (CLIA) standards. Examples include providing guidance on studies designed to improve safety, effectiveness, efficiency, timeliness, equity, and patient-centeredness of laboratory services; revisions to the standards under which clinical laboratories are regulated; the impact of proposed revisions to the standards on medical and laboratory practice; and the modification of the standards and provision of non-regulatory guidelines to accommodate technological advances, such as new test methods, the electronic transmission of laboratory information, and mechanisms to improve the integration of public health and clinical laboratory practices.

Matters For Discussion: The agenda will include agency updates from CDC, CMS, and FDA. Presentations and discussions will include methods for improving the effectiveness/efficiency of CLIAC meetings; an overview of the CMS Advisory Panel on Clinical Diagnostic Laboratory Tests; laboratory interoperability including the Office of the National Coordinator for Health Information Technology (ONC) policies and engagement with clinical laboratories; update on the cytology workload project; update on laboratory biosafety in clinical laboratories; and future CLIAC topics.

Agenda items are subject to change as priorities dictate.

Webcast: The meeting will also be webcast. Persons interested in viewing

the webcast can access information at: <http://cdclabtraining.adobeconnect.com/aprilcliac/>.

In-Person Attendance Online Registration Required: All people attending the CLIAC meeting in-person are required to register for the meeting online at least 5 business days in advance for U.S. citizens and at least 10 business days in advance for international registrants. Register at: <http://www.cdc.gov/cliac/Meetings/MeetingDetails.aspx#>.

Register by scrolling down and clicking the "Register for this Meeting" button and completing all forms according to the instructions given. Please complete all the required fields before submitting your registration and submit no later than April 7, 2016 for U.S. registrants and March 31, 2016 for international registrants.

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 on agenda items whenever possible.

Oral Comments: In general, each individual or group requesting to make oral comments 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, speakers should 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 for public distribution. Written comments, one hard copy with original signature, should be provided to the contact person listed below, and will be included in the meeting's Summary Report.

Availability of Meeting Materials: To support the green initiatives of the federal government, the CLIAC meeting materials will be made available to the Committee and the public in electronic format (PDF) on the internet instead of by printed copy. Check the CLIAC Web site on the day of the meeting for materials: http://wwwn.cdc.gov/cliac/cliac_meeting_all_documents.aspx.

Note: If using a mobile device to access the materials, please verify that the device's browser is able to download the files from the CDC's Web site before the meeting. Alternatively, the files can be downloaded to

a computer and then emailed to the portable device. An internet connection, power source, and limited hard copies may be available at the meeting location, but cannot be guaranteed.

Contact Person for Additional Information: Nancy Anderson, Chief, Laboratory Practice Standards Branch, Division of Laboratory Systems, Center for Surveillance, Epidemiology and Laboratory Services, Office of Public Health Scientific Services, Centers for Disease Control and Prevention, 1600 Clifton Road NE., Mailstop F-11, Atlanta, Georgia 30329-4018; telephone (404) 498-2741; or via email at NAnderson@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 CDC 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. 2016-04590 Filed 3-1-16; 8:45 am]

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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 Funding Opportunity Announcement (FOA) GH16-003, Technical collaboration with the Ministry of Public Health in the Kingdom of Thailand (MOPH)-Research in the conduct of research to assess, prevent, and mitigate public health threats of national and global importance.

Time and Date: 9:00 a.m.–1:00 p.m., EDT, March 23, 2016 (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 “Technical collaboration with the Ministry of Public Health in the Kingdom of Thailand (MOPH)-Research in the conduct of research to assess, prevent, and mitigate public health threats of national and global importance, GH16-003, initial review.”

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.

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Elaine L. Baker,

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

[FR Doc. 2016-04592 Filed 3-1-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Subcommittee on Procedures Review (SPR), Advisory Board on Radiation and Worker Health (ABRWH or the Advisory Board), National Institute for Occupational Safety and Health (NIOSH)

Notice of Cancellation: A notice was published in the **Federal Register** on February 4, 2016 Volume 81, Number 23, Page 6007, announcing an Audio Conference Call of the ABRWH-SPR on February 24, 2016. This meeting was canceled due to a lack of quorum for the meeting. Notice will be provided when the meeting is rescheduled in accordance with section 10(a)(2) of the Federal Advisory Committee Act (Pub. L. 92-463).

Contact Person For More Information: Theodore Katz, Designated Federal Officer, NIOSH, CDC, 1600 Clifton Road, Mailstop E-20, Atlanta, Georgia 30333, Telephone (513)533-6800, Toll Free 1(800)CDC-INFO, Email ocas@cdc.gov.

This notice is published less than the required 15 days prior to the start of the announced meeting, in accordance with

Section 102-3.150(b) of the GSA Final Rule (2001) that allows for exceptions to the meeting notification time requirement. Section 102-3.150(b) states the following: “In exceptional circumstances, the agency or an independent Presidential advisory committee may give less than 15 calendar days notice, provided that the reasons for doing so are included in the advisory committee meeting notice published in the **Federal Register**.”

In this case, the agency is giving less than 15 days’ notice due to the inability to have quorum for the meeting.

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Elaine L. Baker,

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

[FR Doc. 2016-04587 Filed 3-1-16; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30-Day-16-0841]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) has submitted the following information collection request to the Office of Management and Budget (OMB) for review and approval in accordance with the Paperwork Reduction Act of 1995. The notice for the proposed information collection is published to obtain comments from the public and affected agencies.

Written comments and suggestions from the public and affected agencies concerning the proposed collection of information are encouraged. Your comments should address any of the following: (a) Evaluate whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information will have practical utility; (b) Evaluate the accuracy of the agencies estimate of the burden of the proposed collection of information, including the validity of the methodology and assumptions used; (c) Enhance the quality, utility, and