

DEPARTMENT OF HEALTH AND HUMAN SERVICES**National Institutes of Health****National Institute of Biomedical Imaging and Bioengineering; 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: National Institute of Biomedical Imaging and Bioengineering Special Emphasis Panel; P41 BTRC review (2015/05).

Date: May 27, 2015.

Time: 9:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, Suite 920, 6707 Democracy Boulevard, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Manana Sukhareva, Ph.D., Scientific Review Officer, National Institute of Biomedical Imaging and Bioengineering, 6707 Democracy Boulevard, Suite 959, Bethesda, MD 20892, (301) 451-3397, sukharem@mail.nih.gov.

Name of Committee: National Institute of Biomedical Imaging and Bioengineering Special Emphasis Panel; 2015-10 SBIR HD Review.

Date: June 24, 2015.

Time: 10:00 a.m. to 6:00 p.m.

Agenda: To review and evaluate grant applications.

Place: National Institutes of Health, Two Democracy Plaza, Suite 920, 6707 Democracy Boulevard, Bethesda, MD 20892, (Virtual Meeting).

Contact Person: Ruth Grossman, DDS, Scientific Review Officer, National Institute of Biomedical Imaging and Bioengineering, National Institute of Health, 6707 Democracy Boulevard, Rm. 960, Bethesda, MD 20892, (301) 496-8775, grossmanrs@mail.nih.gov.

Dated: March 19, 2015.

David Clary,

Program Analyst, Office of Federal Advisory Committee Policy.

[FR Doc. 2015-06698 Filed 3-23-15; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES**Centers for Disease Control and Prevention****Advisory Committee on Breast Cancer in Young Women (ACBCYW)**

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:

Time and Date: 1:00 p.m.–4:00 p.m. EST, April 17, 2015.

Place: This meeting is accessible by teleconference and web access.

Teleconference and web access login information is as follows:

Toll-Free Telephone: 1-800-369-1856, Participant passcode: 1927394.

Net Conference and Web URL:

For Participants:

URL: <https://www.mymeetings.com/nc/join/>.

Conference number: PW1978421,

Audience passcode: 1927394 and:

<https://www.mymeetings.com/nc/join.php?i=PW1978421&p=1927394&t=c>.

Status: Open to the public, limited only by the net conference and audio phone lines available.

Purpose: The committee provides advice and guidance to the Secretary, HHS; the Assistant Secretary for Health; and the Director, CDC, regarding the formative research, development, implementation and evaluation of evidence-based activities designed to prevent breast cancer (particularly among those at heightened risk) and promote the early detection and support of young women who develop the disease. The advice provided by the Committee will assist in ensuring scientific quality, timeliness, utility, and dissemination of credible appropriate messages and resource materials.

Matters For Discussion: The agenda will include discussions on the current and emerging topics related to breast cancer in young women. These may include risk communication and health education, as well as approaches to increase awareness of clinicians/practitioners regarding topics such as breast cancer risk, breast health, symptoms, diagnosis, and treatment of breast cancer in young women. Discussions for formal recommendations will take place.

Agenda items are subject to change as priorities dictate.

Online Registration Required: All ACBCYW Meeting participants must register for the meeting online at least 3

business days in advance at http://www.cdc.gov/cancer/breast/what_cdc_is_doing/meetings.htm. Please complete all the required fields before submitting your registration and submit no later than April 14, 2015.

Contact Person For More Information:

Temeika L. Fairley, Ph.D., Designated Federal Officer, National Center for Chronic Disease Prevention and Health Promotion, CDC, 4770 Buford Highway NE., Mailstop K52, Atlanta, Georgia, 30341, Telephone (770) 488-4518, Fax (770) 488-4760 Email: acbcyw@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 Agency for Toxic Substances and Disease Registry.

Elaine L. Baker,

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

[FR Doc. 2015-06644 Filed 3-23-15; 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:

12:30 p.m.–5:00 p.m., April 15, 2014
8:30 a.m.–12:00 p.m., April 16, 2014

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 and the electronic transmission of laboratory information.

Matters For Discussion: The agenda will include agency updates from CDC, CMS, and FDA. Presentations and discussions will focus on laboratory information exchange in health information technology; and laboratory safety and quality: lessons learned through the Ebola response.

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://wwwn.cdc.gov/cliac/default.aspx>

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://wwwn.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 8, 2015 for U.S. registrants and April 1, 2015 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 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 public distribution. Written comments, one hard copy with original signature, should be provided to the contact person 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. 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. http://wwwn.cdc.gov/cliac/cliac_meeting_all_documents.aspx 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 Programs, Standards, and Services, Center for Surveillance, Epidemiology and Laboratory Services, Office of Public Health Scientific Services, CDC, 1600 Clifton Road NE., Mailstop F-11, Atlanta, Georgia 30329-4018; telephone (404) 498-2741; or via email at NAAnderson@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. 2015-06648 Filed 3-23-15; 8:45 am]

BILLING CODE 4163-18-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Substance Abuse and Mental Health Services Administration (SAMHSA) will publish a summary of information collection requests under OMB review, in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these documents, call the SAMHSA Reports Clearance Officer on (240) 276-1243.

Project: National Evaluation of the Comprehensive Community Mental Health Services for Children and Their Families Program: Phase VI (OMB No. 0930-0307)—REVISION

The Substance Abuse and Mental Health Services Administration (SAMHSA), Center of Mental Health Services is responsible for the national evaluation of the Comprehensive Community Mental Health Services for Children and Their Families Program (Children's Mental Health Initiative—CMHI) that will collect data on child mental health outcomes, family life, and service system development. Data will be collected on nine (9) service systems, and approximately 2,106 children and families and providers/administrators, using 26 instruments. Data collection will be decreased by 26,960 hours due to program changes resulting from the closing of 19 communities funded in FY 2009 that no longer require data collection and data collection for the Sector and Comparison Study.

Data collection for this evaluation will be conducted over the next 3-year period. Child and family outcomes of interest will be collected at intake and at 6-month follow-up. The individual families will participate in the study for the remaining 12 months. The outcome measures include the following: Child symptomatology and functioning, family functioning, satisfaction, and caregiver strain. The service system data will be collected every 6 months during the remaining 3 years of the evaluation. Service utilization and cost data will be tracked and submitted to the national evaluation every 6 months using two tools—the Flex Fund Tool and the Services and Costs Data Tool—to estimate average cost of treatment per child, distribution of costs, and allocation of costs across service categories. Service delivery and system variables of interest include the