

- September 22, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- October 28, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- November 24, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- December 16, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time

For meeting locations, web conference information, and the most up-to-date information, please visit the calendar on the ONC Web site, <http://www.healthit.gov/FACAS/calendar>.

Contact Person: Michelle Consolazio, Office of the National Coordinator, HHS, 355 E Street SW., Washington, DC 20024, 781-710-0786, Fax: 202-690-6079, email: michelle.consolazio@hhs.gov. Please email Michelle Consolazio for up-to-date information on these meetings. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice.

Agenda: The committee will hear reports from its workgroups and updates from ONC and other Federal agencies. ONC intends to make background material available to the public no later than 24 hours prior to the meeting start time. If ONC is unable to post the background material on its Web site prior to the meeting, it will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on ONC's Web site after the meeting, at <http://www.healthit.gov/FACAS/health-it-standards-committee>

Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the Committee. Written submissions may be made to the contact person prior to the meeting date. Oral comments from the public will be scheduled prior to the lunch break and at the conclusion of each meeting. Time allotted for each presentation will be limited to three minutes. If the number of speakers requesting to comment is greater than can be reasonably accommodated during the scheduled open public session, ONC will take written comments after the meeting.

Persons attending ONC's advisory committee meetings are advised that the agency is not responsible for providing wireless access or access to electrical outlets.

ONC welcomes the attendance of the public at its advisory committee meetings. Seating is limited at the location, and ONC will make every effort to accommodate persons with physical disabilities or special needs. If

special accommodations are required, please contact Michelle Consolazio at least seven (7) days in advance of the meeting.

Notice of this meeting is given under the Federal Advisory Committee Act (Pub. L. 92-463, 5 U.S.C., App. 2).

Dated: October 22, 2014.

Michelle Consolazio,

FACA Lead, Office of Policy, Office of the National Coordinator for Health Information Technology.

[FR Doc. 2014-26059 Filed 10-31-14; 8:45 am]

BILLING CODE 4150-45-P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Information Technology Policy Committee Advisory Meeting; Notice of Meeting

AGENCY: Office of the National Coordinator for Health Information Technology, HHS.

ACTION: Notice of meeting.

This notice announces forthcoming meetings of a public advisory committee of the Office of the National Coordinator for Health Information Technology (ONC). These meetings will be open to the public.

Name of Committee: Health IT Policy Committee.

General Function of the Committee: To provide recommendations to the National Coordinator on a policy framework for the development and adoption of a nationwide health information technology infrastructure that permits the electronic exchange and use of health information as is consistent with the Federal Health IT Strategic Plan and that includes recommendations on the areas in which standards, implementation specifications, and certification criteria are needed.

2015 Meeting Dates and Times:

- January 13, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- February 10, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- March 10, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- April 7, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- May 12, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- June 9, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- July 14, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- August 11, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- September 9, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time

- October 14, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- November 17, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time
- December 8, 2015, from 9:30 a.m. to 3:00 p.m./Eastern Time

For meeting locations, web conference information, and the most up-to-date information, please visit the calendar on the ONC Web site, <http://www.healthit.gov/FACAS/calendar>.

Contact Person: Michelle Consolazio, Office of the National Coordinator, HHS, 355 E Street SW., Washington, DC 20024, 781-710-0786, Fax: 202-690-6079, email: michelle.consolazio@hhs.gov. Please email Michelle Consolazio for up-to-date information on these meetings. A notice in the **Federal Register** about last minute modifications that impact a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice.

Agenda: The committee will hear reports from its workgroups and updates from ONC and other Federal agencies.

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Procedure: Interested persons may present data, information, or views, orally or in writing, on issues pending before the Committee. Written submissions may be made to the contact person prior to the meeting date. Oral comments from the public will be scheduled prior to the lunch break and at the conclusion of each meeting. Time allotted for each presentation will be limited to three minutes. If the number of speakers requesting to comment is greater than can be reasonably accommodated during the scheduled open public session, ONC will take written comments after the meeting.

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Michelle Consolazio at least seven (7) days in advance of the meeting.

ONC is committed to the orderly conduct of its advisory committee meetings. Please visit our Web site at <http://healthit.hhs.gov> for procedures on public conduct during advisory committee meetings.

Notice of this meeting is given under the Federal Advisory Committee Act (Pub. L. No. 92-463, 5 U.S.C., App. 2).

Dated: October 22, 2014.

Michelle Consolazio,

FACA Lead, Office of Policy, Office of the National Coordinator for Health Information Technology.

[FR Doc. 2014-26058 Filed 10-31-14; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[60Day-15-15CT]

Proposed Data Collections Submitted for Public Comment and Recommendations

The Centers for Disease Control and Prevention (CDC), as part of its continuing effort to reduce public burden, invites the general public and other Federal agencies to take this opportunity to comment on proposed and/or continuing information collections, as required by the Paperwork Reduction Act of 1995. To request more information on the below proposed project or to obtain a copy of the information collection plan and instruments, call 404-639-7570 or send comments to Leroy A. Richardson, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an email to omb@cdc.gov.

Comments submitted in response to this notice will be summarized and/or included in the request for Office of Management and Budget (OMB) approval. Comments are invited on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) ways to enhance the quality, utility, and clarity of the information to be collected; (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology; and (e) estimates of capital or start-up costs and costs of operation, maintenance, and purchase of services

to provide information. Burden means the total time, effort, or financial resources expended by persons to generate, maintain, retain, disclose or provide information to or for a Federal agency. This includes the time needed to review instructions; to develop, acquire, install and utilize technology and systems for the purpose of collecting, validating and verifying information, processing and maintaining information, and disclosing and providing information; to train personnel and to be able to respond to a collection of information, to search data sources, to complete and review the collection of information; and to transmit or otherwise disclose the information. Written comments should be received within 60 days of this notice.

Proposed Project

Sudden Death in the Young Registry—New—National Center for Chronic Disease Prevention and Health Promotion (NCCDPHP), Centers for Disease Control and Prevention (CDC).

Background and Brief Description

Every year, infants, children and youth die suddenly and unexpectedly from previously undiagnosed conditions. Little is known about the risk factors leading to SDY, although underlying genetic conditions contribute in some cases. For example, the risk of SDY is known to be higher for infants and children with heart-related conditions or neurological conditions such as epilepsy. However, estimates of the annual incidence of Sudden Death in the Young (SDY) vary broadly due to differences in definitions, inconsistencies in classifying cause of death on death certificates, differing case ascertainment methodologies, and other factors.

To address this gap in knowledge, the Centers for Disease Control and Prevention (CDC), the National Heart, Lung and Blood Institute (NHLBI), and the National Institute for Neurological Disorders and Stroke (NINDS) are collaborating with selected states and partner organizations to create the Sudden Death in the Young Registry (SDYr). The SDYr will establish the first uniform reporting system for SDY based on shared standards for reviewing and classifying deaths involving infants and children <19 years of age, and the systematic participation of individuals with expertise in key medical specialties. Up to 15 state health departments will receive funding from CDC to report standardized information to a central database for compilation and analysis. In addition, each awardee

will establish an advanced clinical review team consisting of a forensic pathologist, neurologist, epileptologist, and pediatric cardiologist to supplement its existing Child Death Review (CDR) process. Specialized medical expertise is essential to the accurate determination of cause of death.

The SDYr will build on ongoing collaborations involving state-based child death review teams, CDC, the National Institutes of Health (NIH), the Health Resources and Services Administration (HRSA), and the National Center for the Review and Prevention of Child Deaths (NCRPCD) at the Michigan Public Health Institute (MPHI). The MPHI-NCRPCD is a non-profit organization dedicated to preventing child deaths and serious injuries. The MPHI-NCRPCD provides technical assistance and training on death scene investigation, and prevention strategies, to partner organizations including state and local governments. The MPHI-NCRPCD also supports a Web-based National Child Death Review Case Reporting System that states can use on a voluntary basis to manage their state-specific CDR data. The system is currently used by 43 states free of charge. Due to variability in case definitions and reporting procedures, the system does not produce national estimates, but serves as a vital repository of information to facilitate state-based surveillance and public health activities.

Utilizing MPHI-NCRPCD Case Reporting System (CRS) infrastructure, MPHI and CDC have previously collaborated with nine states to strengthen case definition and surveillance for sudden unexpected infant death (SUID) involving infants < one year of age. The SDYr will build on this collaboration by increasing the number of participating states, systematically engaging key medical experts in the review of deaths of children <19 years of age, and requiring the collection and reporting of standardized data elements. States participating in the SDYr will report information through a special module called the NCRPCD-CRS-SDY Module. This information collection procedure was selected due to its familiarity to the majority of states, MPHI's successful collaborations with multiple partners, and shared interest in developing and promulgating best practices for the identification, classification, and prevention of SDY.

De-identified information collected through the SDYr will be used to generate estimates of the prevalence of SDY due to cardiac and neurologic conditions; to elucidate risk factors; and