

please contact Judy Sparrow 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. 92-463, 5 U.S.C., App. 2).

Dated: April 12, 2011.

**Judith Sparrow,**

*Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.*

[FR Doc. 2011-9691 Filed 4-20-11; 8:45 am]

**BILLING CODE 4150-45-P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**HIT Policy Committee's Workgroup Meetings; Notice of Meetings**

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

**ACTION:** Notice of meetings.

This notice announces forthcoming subcommittee meetings of a Federal advisory committee of the Office of the National Coordinator for Health Information Technology (ONC). The meetings will be open to the public via dial-in access only.

*Name of Committees:* HIT Policy Committee's Workgroups: Meaningful Use, Privacy & Security Tiger Team, Quality Measures, Governance, Adoption/Certification, and Information Exchange workgroups.

*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.

*Date and Time:* The HIT Policy Committee Workgroups will hold the following public meetings during May 2011: May 2nd Meaningful Use Workgroup, 9 a.m. to 11 a.m./ET; May 3rd Meaningful Use Workgroup in-person meeting, location—TBD, 9 a.m. to 3 p.m./ET; May 4th Privacy & Security Tiger Team,

2 p.m. to 4 p.m./ET; May 5th Quality Measures Workgroup, 10 a.m. to 11:30 a.m./ET; May 10th Meaningful Use Workgroup, 9 a.m. to 12 p.m./ET; May 13th Meaningful Use Workgroup hearing, location—TBD, 9 a.m. to 3 p.m./ET; May 16th Privacy & Security Tiger Team, 2 p.m. to 4 p.m./ET; May 19th Quality Measures Workgroup hearing, location—TBD, 9 a.m. to 3 p.m./ET; and May 20th Meaningful Use Workgroup, 10 a.m. to 1 p.m./ET.

*Location:* All workgroup meetings will be available via webcast; for instructions on how to listen via telephone or Web visit <http://healthit.hhs.gov>. Please check the ONC Web site for additional information or revised schedules as it becomes available.

*Contact Person:* Judy Sparrow, Office of the National Coordinator, HHS, 330 C Street, SW., Washington, DC 20201, 202-205-4528, Fax: 202-690-6079, e-mail: [judy.sparrow@hhs.gov](mailto:judy.sparrow@hhs.gov). Please call the contact person for up-to-date information on these meetings. A notice in the **Federal Register** about last minute modifications that affect a previously announced advisory committee meeting cannot always be published quickly enough to provide timely notice.

*Agenda:* The workgroups will be discussing issues related to their specific subject matter, e.g., meaningful use, information exchange, privacy and security, quality measures, governance, or adoption/certification. If background materials are associated with the workgroup meetings, they will be posted on ONC's Web site prior to the meeting at <http://healthit.hhs.gov>.

*Procedure:* Interested persons may present data, information, or views, orally or in writing, on issues pending before the workgroups. Written submissions may be made to the contact person on or before two days prior to the workgroup's meeting date. Oral comments from the public will be scheduled at the conclusion of each workgroup 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 until close of business on that day.

If you require special accommodations due to a disability, please contact Judy Sparrow at least seven (7) days in advance of the meeting.

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Dated: April 12, 2011.

**Judith Sparrow,**

*Office of Programs and Coordination, Office of the National Coordinator for Health Information Technology.*

[FR Doc. 2011-9694 Filed 4-20-11; 8:45 am]

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

**[60Day-11-0773]**

**Proposed Data Collections Submitted for Public Comment and Recommendations**

In compliance with the requirement of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404-639-5960 and send comments to Daniel Holcomb, CDC Reports Clearance Officer, 1600 Clifton Road, MS-D74, Atlanta, GA 30333 or send an e-mail to [omb@cdc.gov](mailto:omb@cdc.gov).

*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; and (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. Written comments should be received within 60 days of this notice.

**Proposed Project**

National Surveillance for Severe Adverse Events Associated with Treatment of Latent Tuberculosis Infection—(0920-0773 exp. 04/31/2011)—Reinstatement with change—Division of Tuberculosis Elimination (DTBE), National Center for HIV, Viral

Hepatitis, STD, and TB Prevention (NCHHSTP), Centers for Disease Control and Prevention (CDC).

**Background and Brief Description**

As part of the national tuberculosis (TB) elimination strategy, the American Thoracic Society and CDC have published recommendations for targeted testing for TB and treatment for latent TB infection (LTBI)(Morbidity and Mortality Weekly Report 2000;49[RR06];1–54). However, between October 2000 and September 2004, the CDC received reports of 50 patients with severe adverse events (SAEs) associated with the use of the two or three-month regimen of rifampin and pyrazinamide (RZ) for the treatment of LTBI; 12 (24%) patients died (Morbidity and Mortality Weekly Report 2003;52[31]:735–9). In 2004, CDC began collecting reports of SAEs associated with any treatment regimen for LTBI. For surveillance purposes, an SAE was defined as any drug-associated reaction resulting in a patient’s hospitalization or death after at least one treatment dose for LTBI. During 2004 – 2008, CDC received 17 reports of SAEs in 15 adults and two children; all patients had received isoniazid (INH) and had experienced severe liver injury (Morbidity and Mortality Weekly Report 2010; 59:224–9).

Reports of SAEs related to RZ and INH have prompted a need for this project—a national surveillance system of such events. The objective of the project is to determine the annual number and temporal trends of SAEs associated with any treatment for LTBI in the United States. Surveillance of such events will provide data to support periodic evaluation of guidelines for treatment of persons with LTBI and revision.

The Centers for Disease Control and Prevention request approval for a 3-year reinstatement with change of the previously approved National Surveillance for Severe Adverse Events Associated with Treatment of Latent Tuberculosis Infection—(OMB No. 0920–0773, expires April 31, 2011). The changes include a shortened data collection form and an increase in the number of respondents. This project will continue the passive reporting system for SAEs associated with therapy for LTBI. The system will rely on medical chart review and/or onsite investigations by TB control staff.

Potential respondents are any of the 60 reporting areas for the national TB surveillance system (the 50 states, the District of Columbia, New York City, Puerto Rico, and 7 jurisdictions in the Pacific and Caribbean). Data will be collected using the data collection form

for SAEs associated with LTBI treatment. Based on previous reporting, CDC anticipates receiving an average of 10 responses per year from the 60 reporting areas. The data collection form is completed by healthcare providers and health departments for each reported hospitalization or death related to treatment of LTBI and contains demographic, clinical, and laboratory information. CDC will analyze and periodically publish reports summarizing national LTBI treatment adverse events statistics and also will conduct special analyses for publication in peer-reviewed scientific journals to further describe and interpret these data.

The Food and Drug Administration (FDA) collects data on adverse events related to drugs through the FDA MedWatch Program. CDC is collaborating with FDA in the reporting of SAEs. Reporting will be conducted through telephone, e-mail, or during CDC site visits. In this request, CDC is requesting approval for approximately 60 burden hours annually, an estimated increase of 36 hours. This is due to an estimated increase of reports of SAEs after the publication of the MMWR report on SAEs in 2010. There are no costs to respondents other than their time.

ESTIMATE OF ANNUALIZED BURDEN TABLE

Type of respondents	Number of respondents	Number of responses per respondent	Average burden per response (in hours)	Total burden (in hours)
Physicians .....	10	1	1	10
Nurses .....	10	1	4	40
Medical Clerk .....	10	1	1	10
Total .....	.....	.....	.....	60

**Daniel Holcomb,**

Reports Clearance Officer, Centers for Disease Control and Prevention.

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**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Centers for Disease Control and Prevention**

[60 Day-11–0792]

**Proposed Data Collections Submitted for Public Comment and Recommendations**

In compliance with the requirement of Section 3506(c)(2)(A) of the

Paperwork Reduction Act of 1995 for opportunity for public comment on proposed data collection projects, the Centers for Disease Control and Prevention (CDC) will publish periodic summaries of proposed projects. To request more information on the proposed projects or to obtain a copy of the data collection plans and instruments, call 404–639–5960 and send comments to Daniel Holcomb, CDC Reports Clearance Officer, 1600 Clifton Road, MS–D74, Atlanta, GA 30333 or send an e-mail to [omb@cdc.gov](mailto:omb@cdc.gov).

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; and (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. Written comments should be received within 60 days of this notice.

**Proposed Project**

Environmental Health Specialists Network (EHS–Net) National Voluntary Environmental Assessment Information System (NVEAIS)—New—National Center for Environmental Health