

5. Following a structured process (e.g., modified Delphi), provide specific input to support or modify the proposed mappings (about 2.5 hours).

6. Participate in teleconference to discuss areas of disagreement among workgroup members, and to achieve consensus when possible (1.5 hours).

7. Following a structured process (e.g., modified Delphi), provide specific input to support or modify the proposed mappings, incorporating changes accepted in previous steps (about 1.0 hour).

8. Participate in final (optional) teleconference to review final recommendations and discuss contextual issues (1.0 hour).

Please note that should additional conference calls be necessary, workgroup members are expected to make every effort to participate. The workgroups will conduct business by telephone, email, or other electronic means as needed.

Background

The AHRQ Quality Indicators (AHRQ QIs) are a unique set of measures of health care quality that make use of readily available hospital inpatient administrative data. The QIs have been used for various purposes. Some of these include tracking, hospital self-assessment, reporting of hospital-specific quality or pay for performance. The AHRQ QIs are provider- and area-level quality indicators and currently consist of four modules: the Prevention Quality Indicators (PQI), the Inpatient Quality Indicators, the Patient Safety Indicators (PSI), and the Pediatric Quality Indicators (PedQIs). AHRQ is committed to converting the QIs from ICD-9-CM to ICD-10-CM/PCS in an accurate and transparent manner, taking advantage of the additional specificity of ICD-10-CM/PCS to improve the validity and usefulness of the QIs, from October 2014 onward.

Dated: July, 2, 2012.

Carolyn M. Clancy,
AHRQ Director.

[FR Doc. 2012-16734 Filed 7-9-12; 8:45 am]

BILLING CODE 4160-90-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

Mine Safety and Health Research Advisory Committee, National Institute for Occupational Safety and Health (MSHRAC, NIOSH)

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 for the aforementioned committee:

Times and Dates:

8:30 a.m.–5:15 p.m., August 20, 2012

8:30 a.m.–4:30 p.m., August 21, 2012

Place: Hilton Garden Inn Pittsburgh/Southpointe, 1000 Corporate Drive, Canonsburg, Pennsylvania 15317. Telephone: (724) 743-5000, Fax: (724) 743-5010.

Status: Open to the public, limited only by the space available. The meeting room accommodates approximately 50 people.

Purpose: This committee is charged with providing advice to the Secretary, Department of Health and Human Services; the Director, CDC; and the Director, NIOSH, on priorities in mine safety and health research, including grants and contracts for such research, 30 U.S.C. 812(b)(2), Section 102(b)(2).

Matters To Be Discussed: The meeting will focus on engineering noise controls, reducing coal dust exposures, reducing injuries through improved illumination, demographics survey of the mining industry, implementation of the National Academy of Science's recommendations, oxygen supply partnership, safety culture, occupational health and safety management systems, preventing coal dust explosions, and reducing silica exposures.

Agenda items are subject to change as priorities dictate.

Contact Person for More Information: Jeffery L. Kohler, Ph.D., Designated Federal Officer, MSHRAC, NIOSH, CDC, 626 Cochrans Mill Road, Mailstop P05, Pittsburgh, Pennsylvania 15236, telephone (412) 386-5301, fax (412) 386-5300.

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: July 2, 2012.

Elaine L. Baker,

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

[FR Doc. 2012-16776 Filed 7-9-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. FDA-2012-N-0001]

Anti-Infective Drugs Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). The meeting will be open to the public.

Name of Committee: Anti-Infective Drugs Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the Agency on FDA's regulatory issues.

Date and Time: The meeting will be held on September 5, 2012, from 8 a.m. to 5 p.m.

Location: FDA White Oak Campus, 10903 New Hampshire Ave., Building 31 Conference Center, the Great Room (rm. 1503), Silver Spring, MD 20993-0002. Information regarding special accommodations due to a disability, visitor parking, and transportation may be accessed at: <http://www.fda.gov/AdvisoryCommittees/default.htm>; under the heading "Resources for You," click on "Public Meetings at the FDA White Oak Campus." Please note that visitors to the White Oak Campus must enter through Building 1.

Contact Person: Diane Goyette, Center for Drug Evaluation and Research, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 31, Rm. 2417, Silver Spring, MD 20993-0002, 301-796-9001, FAX: 301-847-8533, email: AIDAC@fda.hhs.gov, or FDA Advisory Committee Information Line, 1-800-741-8138 (301-443-0572 in the Washington, DC area) to find out further information regarding FDA advisory committee information. 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. Therefore, you should always check the Agency's Web site at <http://www.fda.gov/AdvisoryCommittees/default.htm> and scroll down to the appropriate advisory committee meeting link, or call the appropriate advisory committee information line to learn about possible modifications before coming to the meeting.

Agenda: The committee will discuss new drug application (NDA) 201688,