

respondent from making such representations unless the representation is non-misleading, and, at the time of making such representation, respondent possesses and relies upon competent and reliable scientific evidence that substantiates that the representation is true. For purposes of Part II, the proposed order defines "competent and reliable scientific evidence" as at least two randomized, double-blind, placebo-controlled human clinical studies that are conducted by independent, qualified researchers and that conform to acceptable designs and protocols, and whose results, when considered in light of the entire body of relevant and reliable scientific evidence, are sufficient to substantiate that the representation is true.

Part III of the proposed order prohibits respondent from making representations, other than representations covered under Parts I or II, about the health benefits of any drug, dietary supplement, or cosmetic, unless the representation is non-misleading, and, at the time of making such representation, respondent possesses and relies upon competent and reliable scientific evidence that is sufficient in quality and quantity based on standards generally accepted in the relevant scientific fields, when considered in light of the entire body of relevant and reliable scientific evidence, to substantiate that the representation is true. For purposes of Part III, the proposed order defines "competent and reliable scientific evidence" as "tests, analyses, research, or studies that have been conducted and evaluated in an objective manner by qualified persons, and that are generally accepted in the profession to yield accurate and reliable results."

Part IV of the proposed order states that the order does not prohibit respondent from making representations for any drug that are permitted in labeling for that drug under any tentative or final standard promulgated by the Food and Drug Administration ("FDA"), or under any new drug application approved by the FDA. This part of the proposed order also states that the order does not prohibit respondent from making representations for any product that are specifically permitted in labeling for that product by regulations issued by the FDA under the Nutrition Labeling and Education Act of 1990.

Part V of the proposed order requires respondent to pay nine hundred thousand dollars (\$900,000) to the Commission to be used for equitable relief, including restitution, and any

attendant expenses for the administration of such equitable relief.

Parts VI, VII, VIII, and IX of the proposed order require respondent to keep copies of relevant advertisements and materials substantiating claims made in the advertisements; to provide copies of the order to its personnel; to notify the Commission of changes in corporate structure that might affect compliance obligations under the order; and to file compliance reports with the Commission. Part X provides that the order will terminate after twenty (20) years, with certain exceptions.

The purpose of this analysis is to facilitate public comment on the proposed order, and it is not intended to constitute an official interpretation of the agreement and proposed order or to modify their terms in any way.

By direction of the Commission.

**Donald S. Clark,**

*Secretary.*

[FR Doc. 2011-16739 Filed 7-1-11; 8:45 am]

**BILLING CODE 6750-01-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Meeting of the Advisory Committee on Minority Health

**AGENCY:** Office of Minority Health, Office of the Assistant Secretary for Health, Office of the Secretary, Department of Health and Human Services.

**ACTION:** Notice of meeting.

**SUMMARY:** As stipulated by the Federal Advisory Committee Act, the Department of Health and Human Services (DHHS) is hereby giving notice that the Advisory Committee on Minority Health (ACMH) will hold a meeting. This meeting is open to the public. Preregistration is required for both public attendance and comment. Any individual who wishes to attend the meeting and/or participate in the public comment session should e-mail [acmh@osophs.dhhs.gov](mailto:acmh@osophs.dhhs.gov).

**DATES:** The meeting will be held on Monday, August 29, 2011 from 9 a.m. to 5 p.m. and Tuesday, August 30, 2011 from 9 a.m. to 1 p.m.

**ADDRESSES:** The meeting will be held at the Doubletree Hotel, 1515 Rhode Island Ave., NW., Washington, DC 20005.

**FOR FURTHER INFORMATION CONTACT:** Ms. Monica A. Baltimore, Tower Building, 1101 Wootton Parkway, Suite 600, Rockville, Maryland 20852. Phone: 240-453-2882 Fax: 240-453-2883.

**SUPPLEMENTARY INFORMATION:** In accordance with Public Law 105-392,

the ACMH was established to provide advice to the Deputy Assistant Secretary for Minority Health in improving the health of each racial and ethnic minority group and on the development of goals and specific program activities of the Office of Minority Health.

Topics to be discussed during this meeting will include increasing the health care workforce and strategies to improve the health of racial and ethnic minority populations through the development of health policies and programs that will help eliminate health disparities, as well as other related issues.

Public attendance at the meeting is limited to space available. Individuals who plan to attend and need special assistance, such as sign language interpretation or other reasonable accommodations, should notify the designated contact person at least fourteen (14) business days prior to the meeting. Members of the public will have an opportunity to provide comments at the meeting. Public comments will be limited to three minutes per speaker. Individuals who would like to submit written statements should mail or fax their comments to the Office of Minority Health at least seven (7) business days prior to the meeting. Any members of the public who wish to have printed material distributed to ACMH committee members should submit their materials to the Executive Secretary, ACMH, Tower Building, 1101 Wootton Parkway, Suite 600, Rockville, Maryland 20852, prior to close of business August 24, 2011.

Dated: June 20, 2011.

**Garth N. Graham,**

*Deputy Assistant Secretary for Minority Health, Office of Minority Health, Office of the Assistant Secretary for Health, Office of the Secretary, U.S. Department of Health and Human Services.*

[FR Doc. 2011-16744 Filed 7-1-11; 8:45 am]

**BILLING CODE 4150-29-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### HIT Standards 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 Standards Committee's Workgroups: Clinical Operations, Vocabulary Task Force, Clinical Quality, Implementation, and Privacy & Security Standards workgroups.

*General Function of the Committee:* To provide recommendations to the National Coordinator on standards, implementation specifications, and certification criteria for the electronic exchange and use of health information for purposes of adoption, consistent with the implementation of the Federal Health IT Strategic Plan, and in accordance with policies developed by the HIT Policy Committee.

*Date and Time:* The HIT Standards Committee Workgroups will hold the following public meetings during July 2011: July 8th Vocabulary Task Force, 11 a.m. to 12 p.m./ET; and July 12th Implementation Workgroup, 12 to 1:30 p.m./ET.

*Location:* All workgroup meetings will be available via webcast; visit <http://healthit.hhs.gov> for instructions on how to listen via telephone or Web. Please check the ONC Web site for additional information 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., clinical operations vocabulary standards, clinical quality, implementation opportunities and challenges, and privacy and security standards activities. 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 workgroups' meeting dates. 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.

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: June 24, 2011.

**Judith Sparrow,**

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

[FR Doc. 2011-16755 Filed 7-1-11; 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 July 2011: July 8th Privacy & Security Tiger Team, 2 to 4 p.m./ET; and July 22nd Privacy & Security Tiger Team, 2 to 4 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.

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: June 24, 2011.

**Judith Sparrow,**

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

[FR Doc. 2011-16749 Filed 7-1-11; 8:45 am]

**BILLING CODE 4150-45-P**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### HIT Policy Committee Advisory Meeting; Notice of Meeting

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

**ACTION:** Notice of meeting.