A 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.

**Date and Time:** The HIT Standards Committee Workgroups will hold the following public meetings during September 2011: September 22nd Implementation Workgroup, 3 to 5 p.m./ET; TBD other Workgroups’ calls.

**Location:** All workshop meetings will be available via Webcast; visit [http://healthit.hhs.gov](http://healthit.hhs.gov) for instructions on how to listen via telephone or Web. Please check the ONC Web site for additional information and schedules as they become 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 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 workshop meetings, they will be posted on ONC’s Web site prior to the meeting at [http://healthit.hhs.gov](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 workshop 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](http://healthit.hhs.gov) for procedures on public conduct during advisory committee meetings.

**Notice:** This notice is given under the Federal Advisory Committee Act (Pub. L. 92–463, 5 U.S.C., App. 2).

**Dates:** Dated: August 8, 2011.

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

**SUPPLEMENTARY INFORMATION:** Under the PRA (44 U.S.C. 3501–3520), Federal agencies must obtain approval from the Office of Management and Budget (OMB) for each collection of information they conduct or sponsor. “Collection of information” is defined in 44 U.S.C. 3502(3) and 5 CFR 1320.3(c) and includes Agency requests or requirements that members of the public submit reports, keep records, or provide information to a third party. Section 3506(c)(2)(A) of the PRA (44 U.S.C. 3506(c)(2)(A)) requires Federal Agencies to provide a 60-day notice in the Federal Register concerning each proposed collection of information, including each proposed extension of an existing collection of information, before submitting the collection to OMB for approval. To comply with this requirement, FDA is publishing notice of the proposed collection of information set forth in this document.

**Agency Information Collection Activities; Proposed Collection; Comment Request; Extra Label Drug Use in Animals**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notice.