

considered a high priority for interlaboratory studies based upon the lack of adequately validated test methods and the regulatory and public health need for such test methods. Based on this evaluation, ICCVAM recommended that:

- The BG1Luc ER TA test method should be considered a high priority for interlaboratory validation studies as an *in vitro* test method for the detection of test substances with ER agonist and antagonist activity.
- Validation studies should include coordination and collaboration with the European Centre for the Validation of Alternative Methods (ECVAM) and the Japanese Center for the Validation of Alternative Methods (JaCVAM) and include one laboratory in each of the three respective geographic regions (United States, Europe, and Japan).
- In preparation for the interlaboratory validation study, XDS should conduct protocol standardization studies with an emphasis on filling data gaps in the antagonist protocol for the BG1Luc ER TA.

The NIEHS subsequently agreed to support the validation study in light of its role as one of the three NTP agencies, whose mission includes the development and validation of improved testing methods. Based on the results of this study, ICCVAM is now reviewing the validation status of this test method for identification of substances with *in vitro* ER agonist or antagonist activity. NICEATM and the ICCVAM Interagency Endocrine Disruptors Working Group prepared a draft BRD that provides a comprehensive description and the data from the validation study used to assess the accuracy and reliability of the BG1Luc ER TA test method. ICCVAM also developed draft recommendations for its use.

#### Availability of the Peer Panel Report

The Panel's conclusions and recommendations are detailed in the *Independent Scientific Peer Review Panel Report: Evaluation of the Validation Status of the BG1Luc4E2 ER TA (LUMICELL), an In Vitro Transcriptional Activation Assay Used to Identify Chemicals That Can Interact with Human Estrogen Receptors* which is available along with the draft documents reviewed by the Panel and the draft ICCVAM test method recommendations at <http://iccvam.niehs.nih.gov/methods/endocrine/PeerPanel11.htm>.

#### Request for Public Comments

NICEATM invites the submission of written comments on the Panel report. When submitting written comments, please refer to this **Federal Register** notice and include appropriate contact information (name, affiliation, mailing address, phone, fax, e-mail, and sponsoring organization, if applicable). All comments received will be made publicly available via the NICEATM–ICCVAM Web site at <http://iccvam.niehs.nih.gov/methods/endocrine/PeerPanel11.htm>. ICCVAM will consider the Panel report along with public comments and comments made by SACATM at their June 16–17, 2011 meeting (67 FR 23323) when finalizing test method recommendations. Final ICCVAM recommendations will be published in an ICCVAM test method evaluation report, which will be forwarded to relevant Federal agencies for their consideration. The evaluation report will also be available to the public on the NICEATM–ICCVAM Web site at <http://iccvam.niehs.nih.gov/methods/endocrine/ERTA-TMER.htm> and by request from NICEATM (see **ADDRESSES** above).

#### Background Information on ICCVAM, NICEATM, and SACATM

ICCVAM is an interagency committee composed of representatives from 15 Federal regulatory and research agencies that require, use, generate, or disseminate toxicological and safety testing information. ICCVAM conducts technical evaluations of new, revised, and alternative safety testing methods with regulatory applicability and promotes the scientific validation and regulatory acceptance of toxicological and safety testing methods that more accurately assess the safety and hazards of chemicals and products and that refine (decrease or eliminate pain and distress), reduce, and replace animal use. The ICCVAM Authorization Act of 2000 (42 U.S.C. 285l–3) established ICCVAM as a permanent interagency committee of the NIEHS under NICEATM. NICEATM administers ICCVAM and provides scientific and operational support for ICCVAM-related activities and conducts independent validation studies to assess the usefulness and limitations of new, revised, and alternative test methods and strategies. NICEATM and ICCVAM welcome the public nomination of new, revised, and alternative test methods and strategies applicable to the needs of U.S. Federal agencies. Additional information about ICCVAM and NICEATM can be found on the

NICEATM–ICCVAM Web site (<http://iccvam.niehs.nih.gov>).

SACATM was established in response to the ICCVAM Authorization Act [Section 285l-3(d)] and is composed of scientists from the public and private sectors. SACATM advises ICCVAM, NICEATM, and the Director of the NIEHS and NTP regarding statutorily mandated duties of ICCVAM and activities of NICEATM. SACATM provides advice on priorities and activities related to the development, validation, scientific review, regulatory acceptance, implementation, and national and international harmonization of new, revised, and alternative toxicological test methods. Additional information about SACATM, including the charter, roster, and records of past meetings, can be found at <http://ntp.niehs.nih.gov/go/167>.

#### References

Denison MS, Heath-Pagliuso S. 1998. The Ah receptor: A regulator of the biochemical and toxicological actions of structurally diverse chemicals. *Bull Environ Contam Toxicol* 61(5): 557–568.

ICCVAM. 2003a. ICCVAM Guidelines for the Nomination and Submission of New, Revised, and Alternative Test Methods. NIH Publication No. 03–4508. Research Triangle Park, NC.

ICCVAM. 2003b. ICCVAM Evaluation of In Vitro Test Methods For Detecting Potential Endocrine Disruptors: Estrogen Receptor and Androgen Receptor Binding and Transcriptional Activation Assays.

Dated: May 11, 2011.

**John R. Bucher,**

*Associate Director, National Toxicology Program.*

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**BILLING CODE 4140–01–P**

#### DEPARTMENT OF HEALTH AND HUMAN SERVICES

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

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

**ACTION:** Notice of meeting.

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

**Name of Committee:** HIT Standards Committee.

**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 meeting will be held on June 22, 2011, from 9 a.m. to 3 p.m./Eastern Time.

**Location:** To Be Determined. For up-to-date information, go to the ONC Web site, <http://healthit.hhs.gov>.

**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 this meeting. 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, including the Clinical Operations, Vocabulary Task Force, Clinical Quality, Implementation, and Enrollment Workgroups. ONC intends to make background material available to the public no later than two (2) business days prior to the meeting. 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://healthit.hhs.gov>.

**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 on or before June 25, 2011. Oral comments from the public will be scheduled between approximately 2 and 3 p.m./Eastern Time. Time allotted for each presentation will be limited to three minutes each. If the number of speakers requesting to comment is greater than can be reasonably accommodated during the scheduled open public hearing session, ONC will take written comments after the meeting until close of business.

Persons attending ONC's advisory committee meetings are advised that the agency is not responsible for providing 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 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: May 9, 2011.

**Judith Sparrow,**

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

[FR Doc. 2011-12107 Filed 5-17-11; 8:45 am]

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## 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.

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

**Name of Committee:** HIT 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.

**Date and Time:** The meeting will be held on June 8, 2011, from 10 a.m. to 4 p.m./Eastern Time.

**Location:** Renaissance Dupont Circle Hotel, 1143 New Hampshire Ave., NW., Washington, DC. For up-to-date information, go to the ONC Web site, <http://healthit.hhs.gov>.

**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 this meeting. 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, including the Meaningful Use Workgroup, the Privacy & Security Tiger Team, the Information Exchange Workgroup, and the Quality Measures Workgroup. ONC intends to make background material available to the public no later than two (2) business days prior to the meeting. 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://healthit.hhs.gov>.

**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 on or before June 2, 2011. Oral comments from the public will be scheduled between approximately 3 and 4 p.m. Time allotted for each presentation is limited to three minutes. If the number of speakers requesting to comment is greater than can be reasonably accommodated during the scheduled open public hearing session, ONC will take written comments after the meeting until close of business.

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

Dated: May 9, 2011.

**Judith Sparrow,**

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

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