private citizen burden hours (20 minutes/communication): 8.

B. Model Qualified Blind Trust: total users (executive branch): 2; private citizen users (100%): 2; private citizen burden hours (100 hours/model): 200.

C. Model Qualified Diversified Trust: total users (executive branch): 1; private citizen users (100%): 1; private citizen burden hours (100 hours/model): 100.

D.–H. Of the five remaining model qualified trust documents: total users (executive branch): 2; private citizen users (100%): 2; private citizen burden hours (100 hours/model): 200.

I.–J. Of the two model confidentiality agreements: total users (executive branch): 1; private citizen users (100%): 1; private citizen burden hours (50 hours/agreement): 50.

However, the total annual reporting hour burden on filers themselves is zero and not the 563 hours estimated above because OGE's estimating methodology reflects the fact that all respondents hire private trust administrators or other private representatives to set up and maintain the qualified blind and diversified trusts. Respondents themselves, typically incoming private citizen Presidential nominees, therefore incur no hour burden. The estimated total annual cost burden to respondents resulting from the collection of information is $1,000,000. Those who use the model documents for guidance are private trust administrators or other private representatives hired to set up and maintain the qualified blind and diversified trusts of executive branch officials who seek to establish qualified trusts. The cost burden figure is based primarily on OGE's knowledge of the typical trust administrator fee structure (an average of 1 percent of total assets) and OGE's experience with administration of the qualified trust program. The $1,000,000 annual cost figure is based on OGE's estimate of an average of five active trusts anticipated to be under administration for each of the next two years with combined total assets of $100,000,000. However, OGE notes that the $1,000,000 figure is a cost estimate for the overall administration of the trusts, only a portion of which relates to information collection and reporting. For want of a precise way to break out the costs directly associated with information collection, OGE is continuing to report to OMB the full $1,000,000 estimate for paperwork clearance purposes.

Public comment is invited on each aspect of the model qualified trust certificates and model trust documents, and underlying regulatory provisions, as set forth in this notice, including specific views on the need for and practical utility of this set of collections of information, the accuracy of OGE's burden estimate, the potential for enhancement of quality, utility and clarity of the information collected, and the minimization of burden (including the use of information technology).

Comments received in response to this notice will be summarized for, and may be included with, the OGE request for extension of the OMB paperwork approval for the set of the various existing qualified trust model certificates, the model communications package, and the model trust documents. The comments will also become a matter of public record.

Approved: May 3, 2007.

Robert I. Cusick,

Director, Office of Government Ethics.

[FR Doc. E7–9162 Filed 5–11–07; 8:45 am]

BILLING CODE 6345–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Toxicology Program (NTP); Liaison and Scientific Review Office; Meeting of the NTP Board of Scientific Counselors

AGENCY: National Institute of Environmental Health Sciences (NIEHS), National Institutes of Health.

ACTION: Meeting announcement and request for comments.

SUMMARY: Pursuant to Public Law 92–463, notice is hereby given of a meeting of the NTP Board of Scientific Counselors (NTP BSC). The NTP BSC is composed of scientists from the public and private sectors and provides primary scientific oversight to the NTP Director and evaluates the scientific merit of the NTP's intramural and collaborative programs.

DATES: The NTP BSC meeting will be held on June 22, 2007. The deadlines for submission of written comments and for pre-registration for the meeting are June 8 and June 15, 2007, respectively. Persons needing special assistance, such as sign language interpretation or other reasonable accommodation in order to attend, should contact 919–541–2475 (voice), 919–541–4644 TTY (text telephone), through the Federal TTY Relay System at 800–877–8339, or by e-mail to *niehsoeeo@niehs.nih.gov.* Requests should be made at least 7 days in advance of the event.

ADDRESSES: The NTP BSC meeting will be held in the Rodbell Auditorium, Rall Building at the NIEHS, 111 T.W. Alexander Drive, Research Triangle Park, NC 27709. Public comments and any other correspondence should be submitted to Dr. Barbara Shane, Executive Secretary for the NTP BSC (NTP Liaison and Scientific Review Office, NIEHS, P.O. Box 12233, MD A3–01, Research Triangle Park, NC 27709; fax: 919–541–0295; or e-mail: *shane@niehs.nih.gov*).

FOR FURTHER INFORMATION CONTACT: Dr. Barbara Shane (telephone: 919–541–4253 or e-mail: *shane@niehs.nih.gov*).

SUPPLEMENTARY INFORMATION:

Preliminary Agenda Topics and Availability of Meeting Materials

Preliminary agenda topics include:

- Update of NTP activities.
- Implementation of workshop and NTP retreat recommendations.
- Review of NTP contracts.
- NTP testing nominations.
- Five-year plan for the NTP Interagency Center for the Evaluation of Alternative Toxicological Methods (NICEATM)—Interagency Coordinating Committee on the Validation of Alternative Methods (ICCVAM).

A copy of the preliminary agenda, committee roster, and any additional information, when available, will be posted on the NTP Web site (*http://ntp.niehs.nih.gov/go/165*) or may be requested in hardcopy from the Executive Secretary for the NTP BSC (see **ADDRESSES** above). Following the meeting, summary minutes will be prepared and made available on the NTP Web site.

Attendance and Registration

The meeting is scheduled for June 22, 2007, from 8:30 a.m. to adjournment and is open to the public with attendance limited only by the space available. Individuals who plan to attend are encouraged to register online at the NTP Web site by June 15, 2007 to facilitate planning for the meeting. Please note that a photo ID is required to access the NIEHS campus. The NTP is making plans to videocast the meeting through the Internet at *http://www.niehs.nih.gov/external/video.htm.*

Request for Comments

Time is allotted during the meeting for the public to present comments to the NTP BSC on the agenda topics. Each organization is allowed one time slot per agenda topic. At least 7 minutes will be allotted to each speaker, and if time permits, may be extended to 10 minutes at the discretion of the NTP BSC chair. Registration for oral comments will also be available on-site, although time allowed for presentation by on-site registrants may be less than that for pre-registered speakers and will be

determined by the number of persons who register at the meeting.

Persons registering to make oral comments are asked, if possible, to send a copy of their statement to the Executive Secretary for the NTP BSC (see **ADDRESSES** above) by June 8, 2007, to enable review by the NTP BSC prior to the meeting. Written statements can supplement and may expand the oral presentation. If registering on-site and reading from written text, please bring 40 copies of the statement for distribution to the NTP BSC and NIEHS/NTP staff and to supplement the record. Written comments received in response to this notice will be posted on the NTP Web site and persons identified by their name and affiliation and/or sponsoring organization, if applicable. Persons submitting written comments should include their name, affiliation (if applicable), phone, e-mail, and sponsoring organization (if any) with the document.

Background Information on the NTP Board of Scientific Counselors

The NTP BSC is a technical advisory body comprised of scientists from the public and private sectors who provide primary scientific oversight to the overall program and its centers. Specifically, the NTP BSC advises the NTP on matters of scientific program content, both present and future, and conducts periodic review of the program for the purposes of determining and advising on the scientific merit of its activities and their overall scientific quality. Its members are selected from recognized authorities knowledgeable in fields such as toxicology, pharmacology, pathology, biochemistry, epidemiology, risk assessment, carcinogenesis, mutagenesis, molecular biology, behavioral toxicology and neurotoxicology, immunotoxicology, reproductive toxicology or teratology, and biostatistics. Members serve overlapping terms of up to four years. NTP BSC meetings are held annually or biannually.

Dated: May 3, 2007.

Samuel H. Wilson,

Deputy Director, National Institute of Environmental Health Sciences and National Toxicology Program.

[FR Doc. E7–9174 Filed 5–11–07; 8:45 am]

BILLING CODE 4140–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Agency for Healthcare Research and Quality

Notice of Meeting

In accordance with section 10(d) of the Federal Advisory Committee Act (5 U.S.C., Appendix 2), announcement is made of a Health Care Policy and Research Special Emphasis Panel (SEP) meeting.

A Special Emphasis Panel is a group of experts in fields related to health care research who are invited by the Agency for Healthcare Research and Quality (AHRQ), and agree to be available, to conduct on an a needed basis, scientific reviews of applications of AHRQ support. Individual members of the Panel do not attend regularly-scheduled meetings and do not serve for fixed terms or a long period of time. Rather, they are asked to participate in particular review meetings which require their type of expertise.

Substantial segments of the upcoming SEP meeting listed below will be closed to the public in accordance with the Federal Advisory Committee Act, section 10(d) of 5 U.S.C., Appendix 2 and 5 U.S.C. 552b(c)(6). Grant applications for Ambulatory Safety and Quality: ''Enabling Patient-Centered Care through Health IT (R18),'' are to be reviewed and discussed at this meeting. These discussions are likely to reveal personal information concerning individuals associated with the applications. This information is exempt from mandatory disclosure under the above-cited statutes.

SEP Meeting on: Ambulatory Safety and Quality: Enabling Patient-Centered Care through Health IT (R18).

Date: June 14–15, 2007 (Open on July 14 from 8 a.m. to 8:15 a.m. and closed for the remainder of the meeting).

Place: Hilton Washington DC/Rockville Executive Meeting (Formerly the Doubletree Hotel), 1750 Rockville Pike, Rockville, Maryland 20852.

Contact Person: Anyone wishing to obtain a roster of members, agenda or minutes of the non-confidential portions of this meeting should contact Mrs. Bonnie Campbell, Committee Management Officer, Office of Extramural Research, Education and Priority Populations, AHRQ, 540 Gaither Road, Room 2038, Rockville, Maryland 20850, Telephone (301) 427–1554.

Agenda items for this meeting are subject to change as priorities dictate.

May 7, 2007.

Carolyn M. Clancy,

Director.

[FR Doc. 07–2365 Filed 5–11–07; 8:45am]

BILLING CODE 4160–90–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2007N–0053]

Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Food Labeling Regulations

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a proposed collection of information has been submitted to the Office of Management and Budget (OMB) for review and clearance under the Paperwork Reduction Act of 1995.

DATES: Fax written comments on the collection of information by June 13, 2007.

ADDRESSES: To ensure that comments on the information collection are received, OMB recommends that written comments be faxed to the Office of Information and Regulatory Affairs, OMB, Attn: FDA Desk Officer, FAX: 202–395–6974. All comments should be identified with the OMB control number 0910–0381. Also include the FDA docket number found in brackets in the heading of this document.

FOR FURTHER INFORMATION CONTACT: Jonna Capezzuto, Office of the Chief Information Officer (HFA–250), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4659.

SUPPLEMENTARY INFORMATION: In compliance with 44 U.S.C. 3507, FDA has submitted the following proposed collection of information to OMB for review and clearance.

Food Labeling Regulations (OMB Control Number 0910–0381)—Extension

FDA regulations require food producers to disclose to consumers and others specific information about themselves or their products on the label or labeling of their products. Related regulations require that food producers retain records establishing the basis for the information contained in the label or labeling of their products and provide those records to regulatory