GSA has published rules in the **Federal Register** that fall under information collection 3090–0250. The rule that prescribed clause 552.238–70 ''Identification of Electronic Office Equipment Providing Accessibility for the Handicapped'' was published at 56 FR 29442, June 27, 1991, titled ''Implementation of Public Law 99–506'', with an effective date of July 8, 1991; and Clause 552.238–74 ''Industrial Funding Fee and Sales Reporting'' published at 68 FR 41286, July 11, 2003.

**B. Annual Reporting Burden**

None.

*OBTAINING COPIES OF PROPOSALS:* Requesters may obtain a copy of the information collection documents from the General Services Administration, Regulatory Secretariat (VIR), 1800 F Street, NW., Room 4035, Washington, DC 20405, telephone (202) 501–4755. Please cite OMB Control No. 3090–0250, Zero Burden Information Collection Reports, in all correspondence.

Dated: October 5, 2007.

**Al Matera,**

*Director, Office of Acquisition Policy.*

[FR Doc. E7–20255 Filed 10–12–07; 8:45 am]

**BILLING CODE 6820–61–S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Agency for Healthcare Research and Quality

### Meeting of the National Advisory Council for Healthcare Research and Quality

**AGENCY:** Agency for Healthcare Research and Quality (AHRQ), HHS.

**ACTION:** Notice of public meeting.

**SUMMARY:** In accordance with section 10(a) of the Federal Advisory Committee Act, this notice announces a meeting of the National Advisory Council for Healthcare Research and Quality.

**DATES:** The meeting will be held on Friday, November 9, 2007, from 9 a.m. to 3 p.m.

**ADDRESSES:** The meeting will be held at the Eisenberg Conference Center, Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland 20850.

**FOR FURTHER INFORMATION CONTACT:** Deborah Queenan, Coordinator of the Advisory Council, at the Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland 20850, (301) 427–1330. For press-related information, please contact Karen Migdail at (301) 427–1855.

If sign language interpretation or other reasonable accommodation for a disability is needed, please contact Mr. Donald L. Inniss, Director, Office of Equal Employment Opportunity Program, Program Support Center, on (301) 443–1144, no later than November 2, 2007. The agenda, roster, and minutes are available from Ms. Bonnie Campbell, Committee Management Officer, Agency for Healthcare Research and Quality, 540 Gaither Road, Rockville, Maryland 20850. Her phone number is (301) 427–1554.

**SUPPLEMENTARY INFORMATION:**

**I. Purpose**

The National Advisory Council for Healthcare Research and Quality was established in accordance with Section 921 (now Section 931) of the Public Health Service Act (42 U.S.C. 299c). In accordance with its statutory mandate, the Council is to advise the Secretary of the Department of Health and Human Services and the Director, Agency for Healthcare Research and Quality (AHRQ), on matters related to actions of the Agency to enhance the quality, improve the outcomes, reduce the costs of health care services, improve access to such services through scientific research, and to promote improvements in clinical practice and in the organization, financing, and delivery of health care services. The Council is composed of members of the public, appointed by the Secretary, and Federal ex-officio members.

**II. Agenda**

On Friday, November 9, the Council meeting will convene at 9 a.m., with the call to order by the Council Chair and approval of previous Council minutes. The Director, AHRQ, will present her update on AHRQ's current research, programs, and initiatives. The agenda will include a discussion of the National Healthcare Quality and Disparities Reports, needed research on Health Care Value and Capacity Building. The official agenda will be available on AHRQ's Web site at *http://www.ahrq.gov* no later than November 2, 2007.

Dated: October 5, 2007.

**Carolyn M. Clancy,**

*Director.*

[FR Doc. 07–5057 Filed 10–12–07; 8:45 am]

**BILLING CODE 4160–90–M**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

**[Docket No. 2007N–0098]**

### Agency Information Collection Activities; Submission for Office of Management and Budget Review; Comment Request; Focus Groups as Used by the Food and Drug Administration

**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 November 14, 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, or e-mailed to *baguilar@omb.eop.gov*. All comments should be identified with the OMB control number 0910–0497. 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.

**Focus Groups as Used by the Food and Drug Administration—(OMB Control Number 0910–0497)—Extension**

Focus groups provide an important role in gathering information because they allow for a more in-depth understanding of consumers' attitudes, beliefs, motivations, and feelings than do quantitative studies. Focus groups serve the narrowly defined need for direct and informal opinion on a specific topic and as a qualitative research toolhave three major purposes:

- To obtain consumer information that is useful for developing variables and measures for quantitative studies,

• To better understand consumers' attitudes and emotions in response to topics and concepts, and

• To further explore findings obtained from quantitative studies.

FDA will use focus group findings to test and refine their ideas, but will generally conduct further research before making important decisions such as adopting new policies and allocating or redirecting significant resources to support these policies.

In the **Federal Register** of March 27, 2007 (72 FR 14279), FDA published a 60-day notice requesting public comment on the information collection provisions. No comments were received.

FDA estimates the burden for completing the forms for this collection of information as follows:

TABLE 1.—ESTIMATED ANNUAL REPORTING BURDEN[1]

| FDA Center | Subject | No. of Focus Groups per Study | No. of Focus Groups Sessions Conducted Annually | No. of Participants per Group | Hours of Duration for Each Group (Includes Screening) | Total Hours |
|---|---|---|---|---|---|---|
| Center for Biologics Evaluation and Research | May use focus groups when appropriate | 1 | 5 | 9 | 1.58 | 71 |
| Center for Drug Evaluation and Research | Varies (e.g., direct-to-consumer Rx drug promotion, physician labeling of Rx drugs, medication guides, over-the-counter drug labeling, risk communication) | 10 | 200 | 9 | 1.58 | 2,844 |
| Center for Devices and Radiological Health | Varies (e.g., FDA Seal of Approval, patient labeling, tampons, online sales of medical products, latex gloves) | 4 | 16 | 9 | 2.08 | 300 |
| Center for Food Safety and Applied Nutrition | Varies (e.g., food safety, nutrition, dietary supplements, and consumer education) | 8 | 40 | 9 | 1.58 | 569 |
| Center for Veterinary Medicine | Varies (e.g., animal nutrition, supplements, labeling of animal Rx) | 5 | 25 | 9 | 2.08 | 468 |
| Total | | 28 | 286 | 9 | 1.78 | 4,252 |

[1]There are no capital costs or operating and maintenance costs associated with this collection of information.

Annually, FDA projects about 28 focus group studies using 186 focus groups lasting an average of 1.78 hours each. FDA has allowed burden for unplanned focus groups to be completed so as not to restrict the agency's ability to gather information on public sentiment for its proposals in its regulatory as well as other programs.

Dated: October 9, 2007.

**Jeffrey Shuren,**

*Assistant Commissioner for Policy.*

[FR Doc. E7–20291 Filed 10–12–07; 8:45 am]

**BILLING CODE 4160–01–S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

### Pediatric 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*: Pediatric Advisory Committee.

*General Function of the Committee*: To provide advice and recommendations to the agency on FDA's regulatory issues. The committee also advises and makes recommendations to the Secretary of Health and Human Services under 45 CFR 46.407 on research involving children as subjects that is conducted or supported by the Department of Health and Human Services, when that research is also regulated by FDA.

*Date and Time*: The meeting will be held on Tuesday, November 27, 2007, from 8 a.m. to 5:30 p.m. and Wednesday, November 28, 2007, from 8 a.m. to 6 p.m.

*Location*: Hilton, Washington DC North/Gaithersburg, Grand Ballroom, 620 Perry Pkwy., Gaithersburg, MD.

*Contact Person*: Carlos Peña, Office of Science and Health Coordination, Office of the Commissioner (HF–33), Food and Drug Administration, 5600 Fishers Lane (for express delivery, rm. 14B–08), Rockville, MD 20857, 301–827–3340, e-mail:*Carlos.Peña@fda.hhs.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 8732310001. Please call the Information Line 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. Therefore, you should always check the agency's Web site and call the appropriate advisory committee hot line/phone line to learn about possible modifications before coming to the meeting.

*Agenda*: On November 27, 2007, in response to the Pediatric Advisory Committee's 2005 request for specific updates after 2 additional years of influenza seasons, the committee will receive information on adverse event reports, focusing on neuropsychiatric and behavioral events, for Tamiflu (OSELTAMIVIR). On November 28, 2007, the Pediatric Advisory Committee will hear and discuss reports by the agency, as mandated in Section 17 of