DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Pediatric Advisory Committee; Amendment of Notice

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

The Food and Drug Administration (FDA) is announcing an amendment to the notice of meeting of the Pediatric Advisory Committee. This meeting was originally announced in the **Federal Register** of January 25, 2008 (73 FR 4581). The amendment is being made to reflect a change in the *Agenda* portion of the document. There are no other changes.

FOR FURTHER INFORMATION CONTACT: Carlos Peña, Office of the Commissioner (HF–33), Food and Drug Administration, 5600 Fishers Lane, 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.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of January 25, 2008, FDA announced that a meeting of the Pediatric Advisory Committee would be held on March 25, 2008. On page 4581, in the third column, the *Agenda* portion of document is changed to read as follows:

Agenda: On March 25, 2008, the Pediatric Advisory Committee will hear and discuss reports by the agency, as mandated in section 17 of the Best Pharmaceuticals for Children Act, on adverse event reports for TOPROL XL (metoprolol), BREVIBLOC (esmolol HCl), LOTENSIN (benazepril), COREG (carvedilol), COLAZAL (balsalazide), ELOXATIN (oxaliplatin), CELEBREX (celecoxib), and SUPRANE (desflurane). The Pediatric Advisory Committee will also hear an update on TRILEPTAL (oxcarbazepine) and the FDA Amendments Act of 2007.

FDA intends to make background material available to the public no later than 2 business days before the meeting. If FDA is unable to post the background material on its Web site prior to the meeting, the background material will be made publicly available at the location of the advisory committee meeting, and the background material will be posted on FDA's Web site after the meeting. Background material is available at *http://www.fda.gov/ohrms/dockets/ac/acmenu.htm*, click on the year 2008 and scroll down to the appropriate advisory committee link.

This notice is issued under the Federal Advisory Committee Act (5 U.S.C. app. 2) and 21 CFR part 14, relating to the advisory committees.

Dated: February 26, 2008.

Randall W. Lutter,

Deputy Commissioner for Policy.

[FR Doc. E8–4156 Filed 3–4–08; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

Agency Information Collection Activities: Submission for OMB Review; Comment Request

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301) 443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: The National Sample Survey of Registered Nurses 2008 (OMB No. 0915–0276)—Reinstatement With Change

The National Sample Survey of Registered Nurses (NSSRN) is carried out to assist in fulfilling the Congressional mandate of Section 806(f) of the Public Health Service Act (42 U.S.C. 296e) which requires that discipline-specific workforce information and analytical activities are carried out as part of the advanced nursing education, workforce diversity, and basic nursing education and practice programs.

Government agencies, legislative bodies and health professionals used data from previous national sample surveys of registered nurses to inform workforce policies. The information from this survey will continue to serve policy makers, and other consumers. The data collected in this survey will provide information on employment status of registered nurses (RNs), the setting in which they are employed and the proportion of RNs who are employed either full-time and part-time in nursing. The data will also indicate the number of RNs who are employed in jobs unrelated to nursing.

The proposed survey design for the 2008 NSSRN updates that of the previous eight surveys. A probability sample is selected from a sampling frame compiled from files provided by the State Boards of Nursing in the 50 States and the District of Columbia. These files constitute a multiple sampling frame of all RNs licensed in the 50 States and the District of Columbia. Sampling rates are set for each State based on considerations of statistical precision of the estimates and the costs involved in obtaining reliable national and State-level estimates.

Each sampled nurse will be asked to complete a self-administered questionnaire, which includes items on educational background, duties, employment status and setting, geographic mobility, and income. An electronic version was offered in the 2004 survey and will be again considered as a mode for response.

Estimated burden is as follows:

Form	Number of respondents	Responses per respondent	Total responses	Hours per response	Total burden hours
Nursing Survey	39,984	1	39,984	.33	13,195

Written comments and recommendations concerning the proposed information collection should be sent within 30 days of this notice to the desk officer for HRSA, either by e-mail to *OIRA_submission@omb.eop.gov* or by fax to 202–395–6974. Please direct all correspondence to the ''attention of the desk officer for HRSA.''