21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093) Rockville, MD 20857, 301–827–7001, e-mail: *Cathy.Groupe@fda.hhs.gov*, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 3014512533. Please call the Information Line for up-to-date information on this meeting.

Agenda: On April 25, 2006, the committee will meet between 8 a.m. to 5 p.m., to discuss new drug application (NDA) 21–359 CELLEGESIC (nitroglycerin [NTG] ointment), 0.4% intra-anal, Cellegy Pharmaceuticals, Inc., for the proposed indication of relief of pain associated with anal fissures. On April 26, 2006, the committee will meet between 8 a.m. to 12 noon, to discuss the agency's draft recommendations for relabeling of antihypertensive drugs for outcome claims, as a followup to the committee's meeting on June 15, 2005, where the committee discussed class labeling of antihypertensive drugs based on the proximity of their data to outcome trials. Following this, from approximately 1 p.m. to 5 p.m., the committee will discuss the ''Placebo in Hypertension Adverse Reaction Meta-Analysis'' Study, a meta-analysis of more than 80,000 patients in placebo-controlled trials of antihypertensive medications, which evaluated the risk of irreversible harm in conducting placebo-controlled trials in patients with hypertension. The background material will become available no later than the day before the meeting and will be posted on FDA's Web site at *http://www.fda.gov/orhms/dockets/ac/acmenu.htm* under the heading ''Cardiovascular and Renal Drugs Advisory Committee.'' (Click on the year 2006 and scroll down to the above named committee).

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 by April 14, 2006. On April 25, 2006, oral presentations from the public will be scheduled between approximately 8:15 a.m. to 8:45 a.m. On April 26, 2006, oral presentations from the public will be scheduled between approximately 8:15 a.m. to 8:45 a.m. and 1 p.m. to 1:30 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before April 14, 2006, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants and an indication of the approximate time requested to make their presentation.

Persons attending FDA's advisory committee meetings are advised that the agency is not responsible for providing access to electrical outlets.

FDA welcomes the attendance of the public at its advisory committee meetings and will make every effort to accommodate persons with physical disabilities or special needs. If you require special accommodations due to a disability, please contact John Lauttman at least 7 days in advance of the meeting at 301–827–7001.

Notice of this meeting is given under the Federal Advisory Committee Act (5 U.S.C. app. 2).

Dated: February 15, 2006.

Jason Brodsky,

Acting Associate Commissioner for External Relations.

[FR Doc. E6–2542 Filed 2–22–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Peripheral and Central Nervous System Drugs 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 Peripheral and Central Nervous System Drugs Advisory Committee. This meeting was announced in the **Federal Register** of January 27, 2006 (71 FR 4593). The amendment is being made to reflect a change in *Date and Time* and *Procedure* portions of the document. An additional day is being added to this meeting and the length of time allotted for the open public hearing portion is being extended. There are no other changes.

FOR FURTHER INFORMATION CONTACT: Sohail Mosaddegh, Center for Drug Evaluation and Research (HFD–21), Food and Drug Administration, 5600 Fishers Lane (for express delivery, 5630 Fishers Lane, rm. 1093), Rockville, MD 20857, 301–827–7001, FAX: 301–827–6776, e-mail: *sohail.mosaddegh@fda.hhs.gov*, or the FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington DC area), code 3014512543. Please call the Information Line for up-to-date information on this meeting.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of January 27, 2006, FDA announced that a meeting of the Peripheral and Central Nervous System Drugs Advisory Committee would be held on March 7, 2006, from 8 a.m. to 5 p.m., and the open public hearing portion scheduled between approximately 1 p.m. and 2 p.m. On page 4593, in the third column, the *Date and Time* portion of the document is amended to read as follows:

Date and Time: The meeting will be held on March 7 and 8, 2006, from 8 a.m. to 5 p.m.

On page 4594, in the first column, in the *Procedure* portion of the document, the third sentence is amended to read as follows:

Procedure: Oral presentations from the public will be scheduled between approximately 1 p.m. and 5 p.m. on March 7, 2006.

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 15, 2006.

Jason Brodsky,

Acting Associate Commissioner for External Relations.

[FR Doc. E6–2541 Filed 2–22–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Institute of Environmental Health Sciences; Proposed Collection; Comment Request; The Sister Study: A Prospective Study of the Genetic and Environmental Risk Factors for Breast Cancer

SUMMARY: In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, for opportunity for public comment on proposed data collection projects, the National Institute of Environmental Health Sciences (NIEHS), the National Institutes of Health (NIH) will publish periodic summaries of proposed projects to be submitted to the Office of Management and Budget (OMB) for review and approval.

Proposed Collection: Title: The Sister Study: A Prospective Study of the Genetic and Environmental Risk Factors for Breast Cancer. *Type of Information Collection Request:* Revision of OMB No. 0925–0522 and expiration date 31 July 2006. *Need and Use of Information Collection:* The purpose of the Sister Study is to study genetic and environmental risk factors for the