in care and custody decisions. It implements placement decisions, develops facilities for care through grants and contracts, and utilizes the foster care system in place for unaccompanied refugee children. The Division, working with other Federal agencies, reunites children with a parent abroad in appropriate cases. The Division conducts investigations and inspections of facilities and placement locations in which unaccompanied children reside. The Division compiles, and updates at least annually, a state-by-state list of professionals or entities qualified to provide the children guardian and attorney representation services. The Division prepares a plan to be submitted to Congress on how to ensure timely appointment of such representation. The Division also maintains statistical information and data on each child, and any actions concerning the child taken by relevant Federal entities while the child is under the Director's care.

Dated: February 28, 2003.

Tommy G. Thompson,

Secretary.

[FR Doc. 03–5720 Filed 3–10–03; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 03F–0048]

BASF Corp.; Filing of Food Additive Petition (Animal Use)—Conjugated Linoleic Acid

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that BASF Corp. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of conjugated linoleic acid in animal feed.

DATES: Submit written or electronic comments by May 26, 2003.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852. Submit electronic comments to http://www.fda.gov/dockets/ecomments.

FOR FURTHER INFORMATION CONTACT: Sharon A. Benz, Center for Veterinary Medicine (HFV–228), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6656, e-mail: sbenz@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (sec. 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 2250) has been filed by BASF Corp., 3000 Continental Dr.-North, Mount Olive, NJ 07828–1234. The petition proposes to amend the food additive regulations in part 573 *Food Additives Permitted in Feed and Drinking Water of Animals* (21 CFR part 573) to provide for the safe use of conjugated linoleic acid (CLA) as a source of fatty acids in swine diets at levels not to exceed 1 percent in complete feed.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations issued under the National Environmental Policy Act (40 CFR 1501.4(b)), the agency is placing the environmental assessment submitted with the petition that is the subject of this notice on public display at the Dockets Management Branch (see **ADDRESSES**) for public review and comment.

Interested persons may submit to the Dockets Management Branch (see **ADDRESSES**) written or electronic comments regarding this document. Submit a single copy of electronic comments to http://www.fda.gov/dockets/ecomments or two hard copies of any written comments, except that individuals may submit one hard copy. Comments are to be identified with the docket number found in brackets in the heading of this document. Received comments may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday. FDA will also place on public display any amendments to, or comments on, the petitioner's environmental information without further announcement in the **Federal Register**. If, based on its review, FDA finds that an environmental impact statement is not required and this petition results in a regulation, the notice of availability of the agency's finding and the evidence supporting that finding will be published with the regulation in the **Federal Register** in accordance with 21 CFR 25.40(c).

Dated: February 27, 2003.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 03–5641 Filed 3–10–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Cancer Institute; Amended Notice of Meeting

Notice is hereby given of a change in the meeting of the Board of Scientific Advisors, March 3, 2003, 8 a.m. to March 4, 2003, 1 p.m. Building 31, C Wing, 6th Floor, Conference Room 10, 9000 Rockville Pike, Bethesda MD 20892 which was published in the **Federal Register** on February 5, 2003, 68 FR 5901.

This meeting is amended to change the time of the open session of the Joint Meeting of the NCI, Board of Scientific Advisors and NCI Board of Scientific Counselors on March 3, 2003 from 8 a.m. to 10:45 a.m. The meeting was originally scheduled to be held from 8 a.m. to 10:15 a.m.

Dated: March 3, 2003.

LaVerne Y. Stringfield,

Director, Office of Federal Advisory Committee Policy.

[FR Doc. 03–5686 Filed 3–10–03; 8:45 am]

BILLING CODE 4140–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

National Institutes of Health

National Human Genome Research Institute; Notice of Closed Meeting

Pursuant to section 10(d) of the Federal Advisory Committee Act, as amended (5 U.S.C. Appendix 2), notice is hereby given of the following meeting.

The meeting will be closed to the public in accordance with the provisions set forth in sections 552b(c)(4) and 552b(c)(6), title 5 U.S.C., as amended. The grant applications and the discussions could disclose confidential trade secrets or commercial property such as patentable material, and personal information concerning individuals associated with the grant applications, the disclosure of which would constitute a clearly unwarranted invasion of personal privacy.

Name of Committee: National Human Genome Research Institute Special Emphasis Panel, Model Organism Database Review.

Date: March 25, 2003.

Time: 11:30 a.m. to 2:30 p.m.

Agenda: To review and evaluate grant applications.

Place: NIH, Bldg 31, Bethesda, MD 20892. (Telephone conference call.)

Contact Person: Ken D. Nakamura, PhD, Scientific Review Administrator, Office of