following the deadline for submission of abstracts. We expect that full planning proposals will be required to be submitted within 45 days of the date of the notification letter informing the applicant that their abstract has been accepted.

Paperwork Reduction Act of 1995 (Pub. L. 104–13)

Public reporting burden for this collection of information is estimated to average 10 hours per response, including the time for reviewing instructions, gathering and maintaining the data needed and reviewing the collection information.

The project description is approved under OMB control number 0970–0139 which expires 12/31/2003. An agency may not conduct or sponsor, and a person is not required to respond to, a collection of information unless it displays a currently valid OMB control number.

Required Notification of the State Single Point of Contact

This program is covered under Executive Order 12372, Intergovernmental Review of Federal Programs, and 45 CFR part 100, Intergovernmental Review of Department of Health and Human Services Program and Activities. Under the Order, States may design their own processes for reviewing and commenting on proposed Federal assistance under covered programs.

All States and Territories except Alabama, Alaska, Colorado, Connecticut, Hawaii, Idaho, Kansas, Louisiana, Massachusetts, Minnesota, Montana, Nebraska, New Jersey, New York, Ohio, Oklahoma, Oregon, Palau, Pennsylvania, South Dakota, Tennessee, Vermont, Virginia, and Washington have elected to participate in the Executive Order process and have established Single Points of Contact (SPOCs). Applicants from these 24 jurisdictions need take no action regarding E.O. 12372. Applicants for projects to be administered by Federally recognized Indian tribes are also exempt from the requirements of E.O. 12372. Otherwise, applicants should contact their SPOCs as soon as possible to alert them of the prospective applications and receive any necessary instructions. Applicants must submit any required material to the SPOCs as soon as possible so that the program office can obtain and review SPOC comments as part of the award process. It is imperative that the applicant submit all required materials, if any, to the SPOC and indicate the date of this submittal (or the date of contact if no submittal is required) on the Standard Form 424, item 16a.

Under 45 CFR 100.8(a)(2), a SPOC has 60 days from the application deadline to comment on proposed new or competing continuation awards.

SPOCs are encouraged to eliminate the submission of routine endorsements as official recommendations.

Additionally, SPOCs are requested to clearly differentiate between mere advisory comments and those official State process recommendations which may trigger ''the accommodate or explain'' rule. When SPOC comments are submitted directly to ACYF, they should be addressed to: William Wilson, ACYF's Office of Grants Management, Room 2220, Switzer Building, 330 C Street, SW., Washington, DC 20447. A list of the Single Points of Contact for each State and Territory can be found on the Web site *http://www.whitehouse.gov/omb/grants/spoc.html.*

Dated: March 5, 2003.

Joan E. Ohl,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 03–5721 Filed 3–10–03; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Statement of Organization, Functions, and Delegations of Authority

This notice amends Part K of the Statement of Organization, Functions, and Delegations of Authority of the Department of Health and Human Services (DHHS), Administration for Children and Families (ACF) as follows: Chapter KR, the Office of Refugee Resettlement (ORR), as last amended, July 15, 2002. This notice reflects the establishment of a new Division to incorporate the functions under the immigration laws of the United States, with respect to the care of unaccompanied alien children that were vested by statute in, or performed by, the Commissioner of the Immigration and Naturalization Service which were transferred to the Office of Refugee Resettlement by the Homeland Security Act of 2002, Public Law 107–296.

1. This Chapter is amended as follows:

A. *KR.10 Organization.* Delete in its entirety and replace with the following:

KR.10 Organization. The Office of Refugee Resettlement is headed by a Director who reports directly to the Assistant Secretary for Children and Families and consists of: Office of the Director (KRA), Division of Refugee Assistance (KRE), Division of Community Resettlement (KRF), Division of Budget, Policy and Data Analysis (KRG), Division of Unaccompanied Children's Services (KRH).

B. Delete KR.20 Functions, Paragraph A, Office of the Director in its entirety and replace with the following:

KR.20 Functions. A. The Office of the Director is directly responsible to the Assistant Secretary for Children and Families for carrying out ORR's mission and providing guidance and general supervision to the components of ORR. The Office provides direction in the development of program policy and budget and in the formulation of salaries and expense budgets. Staff also provide administrative and personnel support services. The Office is responsible for implementing certain provisions of the Trafficking Victims Protection Act.

The Office coordinates with the lead refugee and entrant program offices of other federal departments; provides leadership in representing refugee and entrant programs, policies and administration to a variety of governmental entities and other public and private interests; and acts as the coordinator of the total refugee and entrant resettlement effort for ACF and the Department. The Office coordinates the certification of, and services to, victims of severe forms of trafficking. It also coordinates with other Federal government agencies on certification activities and policy issues related to the trafficking law. In consultation with appropriate juvenile justice professionals and Federal immigration services and border security agencies, the Director makes placement determinations and coordinates care and placement services for unaccompanied alien children who are in Federal custody by reason of their immigration status.

C. Establish KR.20 Functions, Paragraph E, the Division of Unaccompanied Children's Services.

E. The Division of Unaccompanied Children's Services develops programs and guidance for the coordination and implementation of care and placement services for unaccompanied alien children who are in Federal custody by reason of their immigration status. The Division recommends placement determinations to the Director in consultation with appropriate juvenile justice professionals and Federal immigration services and border security agencies. The Division ensures consideration of the child's best interest

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