care field, and is lead by a very experienced team of researchers in child care policy research. The data collected through this study will provide information urgently needed by policymakers in the current environment of the next phase of welfare reform.

The Urban Institute is in a unique position to conduct this much-needed research because:

- They have developed a network of State and local connections and knowledge base while conducting their work on the Assessing the New Federalism Project, as well as a previous project on the experiences of families with the subsidy system, funded by ACF; and
- They have started the planning phase and ground work for the proposed project with funding secured through a foundation.

The Agency is providing members of the public, including qualified organizations which would be interested in competing for the funding if a competition were held, an opportunity to comment on the planned action.

Statutory Authority: This award will be made pursuant to the Child Care and Development Block Grant Act of 1990 as amended (CCDBG Act); section 418 of the Social Security Act; Consolidated Appropriations Act, 2001 (Pub. L. 106–554). The Catalog of Federal Domestic Assistance is 93.647.

DATES: In order to be considered, comments on this planned action must be received on or before September 9, 2002.

ADDRESSES: Interested parties, including qualified organizations which would be interested in competing for the funding if a competition were held, should write to: Karen Tvedt, Child Care Bureau, Administration on Children, Youth and Families (ACYF), Administration for Children and Families (ACF), Department of Health and Human Services, 330 C Street SW., Room 2046, Washington, DC 20447.

FOR FURTHER INFORMATION CONTACT: Karen Tvedt, Child Care Bureau, at (202) 401–5130.

Catalog of Federal Domestic Assistance Program Number 93.647, Child Care Research Discretionary Grants

Dated: July 29, 2002.

Joan E. Ohl,

Commissioner, Administration on Children, Youth and Families.

[FR Doc. 02–21980 Filed 8–27–02; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02F–0327]

ADM Alliance Nutrition, Inc.; Filing of Food Additive Petition (Animal Use)-Feed-Grade Biuret

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that ADM Alliance Nutrition, Inc. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of feed-grade biuret in lactating dairy cattle feed.

DATES: Submit written or electronic comments on the petitioner's environmental assessment by November 11, 2002.

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 Benz, Center for Veterinary Medicine (HFV–228), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6656.

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 2248) has been filed by ADM Alliance Nutrition, Inc., 1000 North 30th St., P.O. Box C1., Quincy, IL 62305–7100. 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 feed-grade biuret in lactating dairy cattle 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 information 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. Two copies of any comments are to be submitted, except individuals may submit one 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 (see **ADDRESSES**) 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 assessment without further announcement in the **Federal Register**. If, based on its review, the agency 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 of no significant impact 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: August 5, 2002.

Linda Tollefson,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. 02–21698 Filed 8–27–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00D–0053]

Determining Hospital Procedures for Opened-But-Unused, Single-Use Medical Devices; Request for Comments and Information

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is providing an opportunity for interested persons to submit comments about current practices with respect to opened-but-unused, single-use medical devices. FDA is publishing this notice in order to gather informed comment from individuals, professional organizations, original equipment manufacturers, reprocessors, and hospitals as it examines its policy with respect to opened-but-unused, single-use medical devices.

DATES: Submit written comments by November 26, 2002.

ADDRESSES: Submit written comments and information 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: Larry Spears, Center for Devices and Radiological Health (HFZ–300), Food and Drug Administration, 2094 Gaither Rd., Rockville, MD 20850, 301–594–4692.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of August 14, 2000 (65 FR 49583), FDA published a guidance entitled ''Enforcement Priorities for Single-Use Devices Reprocessed by Third Parties and Hospitals.'' The guidance defined ''opened-but-unused'' devices as:

> Single-use, disposable devices whose sterility has been breached or compromised, or whose sterile package was opened but not been used on a patient, that is, they have not been in contact with blood or bodily fluids.

The guidance did not apply the agency's enforcement priorities for reprocessed devices to opened-but-unused, single-use medical devices reprocessed in hospitals. The guidance did state, however, that the agency would examine its policy with respect to opened-but-unused, single-use medical devices. In doing so, FDA is soliciting information about current practices regarding this issue. A copy of the guidance is available on FDA's Web site at http://www.fda.gov/cdrh/reuse/1168.html.

FDA is interested in comments related to: (1) Whether or not hospitals have a written policy or procedure for handling sterile, single-use medical devices that are opened, for whatever reason, but are unused; (2) how hospitals determine if a single-use medical device that has been opened but unused is contaminated; and (3) what types of single-use medical devices are resterilized because they are opened but unused.

Interested persons may submit to the Dockets Management Branch (see **ADDRESSES**), written or electronic comments or information regarding this issue by [*insert date 90 days after date of publication in the* **Federal Register**]. Two copies of any comments are to be submitted, except that individuals may submit one 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.

Dated: August 19, 2002.

Linda S. Kahan,

Deputy Director, Center for Devices and Radiological Health.

[FR Doc. 02–21891 Filed 8–27–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Substance Abuse and Mental Health Services Administration

Center for Mental Health Services; Notice of Meeting

Pursuant to Pub. L. 92–463, notice is hereby given of the meeting of the Center for Mental Health Services (CMHS) National Advisory Council in September 2002.

A portion of the meeting will be open and will include a roll call, general announcements, and discussion about the Substance Abuse and Mental Health Service's matrix program areas, the New Freedom Commission on Mental Health, the Center for Mental Health Service's disparities grant program, and consumer affairs.

Public comments are welcome. Please communicate with the individual listed as contact below for guidance. If anyone needs special accommodations for persons with disabilities please notify the contact listed below.

The meeting will also include the review, discussion, and evaluation of grant applications. Therefore a portion of the meeting will be closed to the public as determined by the SAMHSA Administrator, in accordance with Title 5 U.S.C. 552b(c)(6) and 5 U.S.C. App. 2. & 10 (d).

A summary of the meeting and a roster of Council members may be obtained from Ms. Eileen Pensinger, Executive Secretary, CMHS, Room 15–99, Parklawn Building, Rockville, Maryland 20857, telephone (301) 443–4823.

Committee Name: CMHS National Advisory Council.

Meeting Date: September 5–6, 2002.

Place: The Double Tree Hotel, 1750 Rockville Pike, Rockville, Maryland.

Type: Closed: September 5, 2002—9:15 a.m.—11 a.m.

Open: September 5, 2002—11:15 a.m.—5:30 p.m.

Open: September 6, 2002—9 a.m.—12:15 p.m.

Contact: Eileen S. Pensinger, M.Ed., Executive Secretary, 5600 Fishers Lane, Parklawn Building, Room 15–99, Rockville, Maryland 20857, Telephone: (301) 443–4823 and FAX (301) 443–5163.

Dated: August 12, 2002.

Toian Vaughn,

Committee Management Officer, Substance Abuse and Mental Health Services Administration.

[FR Doc. 02–21935 Filed 8–27–02; 8:45 am]

BILLING CODE 4162–20–P

DEPARTMENT OF THE INTERIOR

Fish and Wildlife Service

Notice of Availability of Recovery Goals for Four Endangered Fishes of the Colorado River Basin

AGENCY: Fish and Wildlife Service, Interior.

ACTION: Notice of document availability.

SUMMARY: To further the recovery of humpback chub (*Gila cypha*), bonytail (*Gila elegans*), Colorado pikeminnow (*Ptychocheilus lucius*), and razorback sucker (*Xyrauchen texanus*), we, the Fish and Wildlife Service announce the availability of recovery goals for these endangered fishes of the Colorado River Basin. This information will serve as an amendment and supplement to the respective existing recovery plans for each species. The recovery goals for each species provide objective, measurable recovery criteria for downlisting and delisting that identify levels of demographic and genetic viability needed for self-sustaining populations and site-specific management actions/tasks needed to minimize or remove threats.

SUPPLEMENTARY INFORMATION: The purpose of these recovery goals is to describe site-specific management actions/tasks; provide objective, measurable recovery criteria; and provide estimates of the time required to achieve recovery of each of the four endangered fish species. Recovery of the humpback chub, bonytail, and razorback sucker is considered in two recovery units, *i.e.*, upper basin (upstream of Glen Canyon Dam, Arizona) and lower basin. Recovery of the Colorado pikeminnow is considered only in the upper basin recovery unit. Downlisting and delisting criteria by listing factors and management actions, as well as demographic criteria, are presented for populations of each species within the recovery units. In addition, updated life-history information and estimated time to achieve the downlisting and delisting requirements are also presented. These recovery goals serve as amendments and supplements to the recovery plans by providing more specific objective and measurable criteria to recover each of the four fish species.

Draft recovery goals were made available for public comment on September 10, 2001, through a **Federal Register** Notice of Availability (66 FR 47033–47034) and on November 23, 2001, through a **Federal Register** Notice of Reopening (66 FR 58748). Comments were categorized by topic and