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.