call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 30 days of this notice directly to the OMB desk officer: OMB Human Resources and Housing Branch, Attention: Brenda Aguilar, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: August 13, 2002.

John P. Burke, III,

Paperwork Reduction Act Team Leader, CMS Reports Clearance Officer, CMS Office of Information Services, Security and Standards Group, Division of CMS Enterprise Standards.

[FR Doc. 02–21825 Filed 8–27–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Medicare and Medicaid Services

[Document Identifier: CMS–1856/1893]

Agency Information Collection Activities: Submission for OMB Review; Comment Request

AGENCY: Centers for Medicare and Medicaid Services.

In compliance with the requirement of section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Centers for Medicare and Medicaid Services (CMS) (formerly known as the Health Care Financing Administration (HCFA), Department of Health and Human Services, is publishing the following summary of proposed collections for public comment. Interested persons are invited to send comments regarding this burden estimate or any other aspect of this collection of information, including any of the following subjects: (1) The necessity and utility of the proposed information collection for the proper performance of the agency's functions; (2) the accuracy of the estimated burden; (3) ways to enhance the quality, utility, and clarity of the information to be collected; and (4) the use of automated collection techniques or other forms of information technology to minimize the information collection burden.

Type of Information Collection Request: Extension of a currently approved collection; *Title of Information Collection:* Request for Certification in the Medicare/Medicaid Program to Provide Outpatient Physical Therapy and/or Speech-Language Pathology and the Outpatient Physical Therapy and/or Speech-Language Pathology Survey Report Form and Supporting Regulations in 42 CFR 485.701–485.729; *Form No.:* CMS–1856/1893 (OMB# 0938–0065); *Use:* The form CMS–1856 is utilized as an application to be completed by suppliers of OPT/SP services requesting participation in the Medicare/Medicaid programs. This form initiates the process of obtaining a decision as to whether the conditions of coverage are met as an OPT/SP supplier. It is used by the CMS Regional Offices (ROs) to enter the new supplier into the Online Survey, Certification and Reporting System (OSCAR). The survey report form CMS–1893 is an instrument used by the State survey agency to record data collected during an on-site survey of a supplier of OPT/SP services to determine compliance with the applicable conditions of participation and to report this information to the Federal Government. The form is primarily a coding worksheet designed to facilitate data reduction and retrieval into the OSCAR system at the CMS ROs. The form includes basic information on compliance (*i.e.,* met, not met, explanatory statements) and does not require any descriptive information regarding the survey activity itself.; *Frequency:* On occasion; *Affected Public:* Business or other for profit; *Number of Respondents:* 1,700; *Total Annual Responses:* 255; *Total Annual Hours:* 446.

To obtain copies of the supporting statement and any related forms for the proposed paperwork collections referenced above, access CMS Web site address at *http://www.hcfa.gov/regs/prdact95.htm,* or E-mail your request, including your address, phone number, OMB number, and CMS document identifier, to *Paperwork@hcfa.gov,* or call the Reports Clearance Office on (410) 786–1326. Written comments and recommendations for the proposed information collections must be mailed within 30 days of this notice directly to the OMB desk officer: OMB Human Resources and Housing Branch, Attention: Brenda Aguilar, New Executive Office Building, Room 10235, Washington, DC 20503.

Dated: August 20, 2002.

John P. Burke, III,

Paperwork Reduction Act Team Leader, CMS Reports Clearance Officer, Office of Strategic Operations and Regulatory Affairs, Division of Regulations Development and Issuances.

[FR Doc. 02–21826 Filed 8–27–02; 8:45 am]

BILLING CODE 4120–03–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Notice of Award of Non-Competitive Grant

AGENCY: Administration on Children, Youth and Families (ACYF), ACF, DHHS.

ACTION: Notice; Opportunity to Comment.

SUMMARY: Notice is hereby given that ACYF is considering awarding discretionary research grant funds without competition to The Urban Institute at 2100 M St. NW., Washington, DC, for up to $375,000 of Child Care and Development Block Grant funds in FY 2002. And, pending the availability of Federal funds, and the continuing non-Federal support of the project from other sources, ACYF will award up to $375,000 of Child Care and Development Block Grant funds for one additional fiscal year. The two-year project period would begin on September 30, 2002 and end on September 29, 2004. This award will be made to The Urban Institute to provide Federal support for a research project that will examine the interaction of child care providers and child care subsidy policies and practice.

The Urban Institute's research project addresses many questions of relevance to the child care field, to ACF, and to the Child Care Bureau in particular. It will fill a gap in the information currently available about the characteristics of subsidized and unsubsidized providers, and how implementation of subsidy policies affects the experiences of those providers. In addition, the study will explore the rate of participation of faith-based organizations in the child care subsidy system, addressing one of the Administration's priorities. It will also explore the occurrence of activities supporting children's early learning and literacy in diverse child care settings, as well as providers' characteristics that may be related to the likelihood of those activities being present in child care settings. The study answers a call for needed research on providers as expressed by researchers and policymakers in the most recent meeting of the Child Care Research Consortium held in Washington, DC on April 17–19, 2002.

The proposed project has a strong research design and methodology, builds on a solid understanding of the current state of research in the child

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.