Correction

In the **Federal Register** issue of May 21, 1998, in FR Doc. 98–13473, on page 27975, in the first column, correct the first sentence of the ''Eligible Applicants'' section to read:

Given the nature of the research involved, competition is open only to State agencies that administer TANF programs and to counties with populations greater than 500,000 that administer TANF programs.

Dated: May 28, 1998.

Margaret A. Hamburg,

Assistant Secretary for Planning and Evaluation.

[FR Doc. 98–14838 Filed 6–3–98; 8:45 am]

BILLING CODE 4151–04–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Proposed Information Collection Activity; Comment Request; Proposed Projects

Title: Appeal Procedures for Head Start Grantees and Current or Former Delegate Agencies.

OMB No.: 0980–0242.

Description: Section 646 of the Head Start Act requires the Secretary to prescribe procedures insuring that an agency or organization which desires to serve as a delegate agency under the Head Start Act will receive special notice and an opportunity for a timely appeal when an application has been wholly or substantially rejected or when such application has not been acted upon within a period of time deemed reasonable by the Secretary. The rule also describes the actions available prior to the suspension, termination, or reduction of financial assistance or when an application for refunding is denied.

Respondents: State, Local or Tribal Government.

ANNUAL BURDEN ESTIMATES

Instrument	Number of respondents	Number of responses per respondent	Average burden hours per response	Total burden hours
Appeal	10	1	16	160

Estimated Total Annual Burden Hours: 160.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and Families, Office of Information Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) Whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: June 1, 1998.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 98–14836 Filed 6–3–98; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Administration for Children and Families

Agency Information Collection Under OMB Review

Title: Head Start Grants Administration.

OMB No.: 0980–0243.

Description: This part establishes regulations applicable to program administration and grant management for grants under the Head Start Act. The regulations clarify definitions of terms applicable to the administration of the Head Start program. In addition the regulations establish a requirement for grantees to have student accidents insurance and bonding for certain officials. The regulations also require funding recipients to establish written personnel policies, and clarify the limitations on costs of development and administration of Head Start programs.

Respondents: State, Local or Tribal Governments.

Title	Number of respondents	Number of responses per respondent	Average burden per response	Burden
45 CFR 1301	2186	1	40	87,440

Estimated Total Annual Burden: 87,440.

In compliance with the requirements of Section 3506(c)(2)(A) of the Paperwork Reduction Act of 1995, the Administration for Children and Families is soliciting public comment on the specific aspects of the information collection described above. Copies of the proposed collection of information can be obtained and comments may be forwarded by writing to the Administration for Children and

Families, Office of Information Services, 370 L'Enfant Promenade, S.W., Washington, D.C. 20447, Attn: ACF Reports Clearance Officer. All requests should be identified by the title of the information collection.

The Department specifically requests comments on: (a) whether the proposed collection of information is necessary for the proper performance of the functions of the agency, including whether the information shall have practical utility; (b) the accuracy of the agency's estimate of the burden of the proposed collection of information; (c) the quality, utility, and clarity of the information to be collected; and (d) ways to minimize the burden of the collection of information on respondents, including through the use of automated collection techniques or other forms of information technology. Consideration will be given to comments and suggestions submitted within 60 days of this publication.

Dated: June 1, 1998.

Bob Sargis,

Acting Reports Clearance Officer.

[FR Doc. 98–14837 Filed 6–3–98; 8:45 am]

BILLING CODE 4184–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 98F–0342]

Alcide Corp.; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Alcide Corp. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of acidified sodium chlorite solutions in poultry processing.

DATES: Written comments on the petitioner's environmental assessment by July 6, 1998.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Robert L. Martin, Center for Food Safety and Applied Nutrition (HFS–217), Food and Drug Administration, 200 C St. SW., Washington, DC 20204–0001, 202–418–3074.

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 petition (FAP 8A4591) has been filed by Alcide Corp., 8561 154th Ave., NE., Redmond, WA 98052. The petition proposes to amend the food additive regulations in 21 CFR 173.325 to provide for a lower pH in the use of acidified sodium chlorite solutions in poultry processing.

The potential environmental impact of this action is being reviewed. To encourage public participation consistent with regulations promulgated 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 (address above) for public review and comment. Interested persons may, on or before July 6, 1998, submit to the Dockets Management Branch (address above) written comments. 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 office above 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: May 14, 1998.

Laura M. Tarantino,

Acting Director, Office of Premarket Approval, Center for Food Safety and Applied Nutrition.

[FR Doc. 98–14761 Filed 6–3–98; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

Neurological Devices Panel of the Medical Devices Advisory Committee; Notice of Meeting

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

This notice announces a forthcoming meeting of a public advisory committee of the Food and Drug Administration (FDA). At least one portion of the meeting will be closed to the public.

Name of Committee: Neurological Devices Panel of the Medical Devices Advisory Committee.

General Function of the Committee: To provide advice and recommendations to the agency on FDA regulatory issues.

Date and Time: The meeting will be held on June 12, 1998, 10:30 a.m. to 5 p.m.

Location: Corporate Bldg., conference room 020B, 9200 Corporate Blvd., Rockville, MD.

Contact Person: Janet L. Scudiero, Center for Devices and Radiological Health (HFZ–410), Food and Drug Administration, 9200 Corporate Blvd., Rockville, MD 20850, 301–594–1184, or FDA Advisory Committee Information Line, 1–800–741–8138 (301–443–0572 in the Washington, DC area), code 12513. Please call the Information Line for up-to-date information on this meeting.

Agenda: On June 12, 1998, the committee will discuss and make recommendations on reclassification of preamendment class III artificial embolism devices for neurological use based on information received from a call for safety and effectiveness information, under section 515(i) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360e), published in the **Federal Register** of August 14, 1995, and June 13, 1997 (60 FR 41984 and 62 FR 32352, respectively).

Procedure: On June 12, 1998, from 11 a.m. to 5 p.m., the meeting is open to the public. Interested persons may present data, information, or views, orally or in writing, on issues pending before the committee. Written submissions may be made to the contact person by June 9, 1998. Oral presentations from the public will be scheduled between approximately 11 a.m. and 12 m. and between approximately 3:45 p.m. and 4:15 p.m. Time allotted for each presentation may be limited. Those desiring to make formal oral presentations should notify the contact person before June 9, 1998, and submit a brief statement of the general nature of the evidence or arguments they wish to present, the names and addresses of proposed participants, and an indication of the approximate time requested to make their presentation.

Closed Committee Deliberations: On June 12, 1998, from 10:30 a.m. to 11 a.m., the meeting will be closed to permit FDA to present to the committee