morbidity, and social problems among both youth and adults in the U.S. OMB clearance for the 1999 survey expired January 2000 (OMB No. 0920–0258, expiration 01/00). Data on the health risk behaviors of adolescents is the focus of approximately 40 national health objectives in Healthy People 2010. The Youth Risk Behavior Survey provides data to measure at least 10 of these health objectives and 3 of the 10 Leading Health Indicators. In addition, the Youth Risk Behavior Survey can identify racial and ethnic disparities in health risk behaviors. No other national source of data measures as many of the 2010 objectives that address behaviors of adolescents. The data also will have significant implications for policy and program development for school health programs nationwide. The annualized burden is 9,173 hours.

Respondents	Number of respondents	Number of responses per respondent	Burden per response (hours)
High school students	12,000	1	0.75
School administrators	345	1	0.50

Dated: September 5, 2000.

Nancy Cheal,

Acting Associate Director for Policy, Planning and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 00–23203 Filed 9–8–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Centers for Disease Control and Prevention

[30DAY–67–00]

Agency Forms Undergoing Paperwork Reduction Act Review

The Centers for Disease Control and Prevention (CDC) publishes a list of information collection requests under review by the Office of Management and Budget (OMB) in compliance with the Paperwork Reduction Act (44 U.S.C. Chapter 35). To request a copy of these requests, call the CDC Reports Clearance Officer at (404) 639–7090. Send written comments to CDC, Desk Officer, Human Resources and Housing Branch, New Executive Office Building, Room 10235, Washington, DC 20503. Written comments should be received within 30 days of this notice.

Proposed Project

2001 National Health Interview Survey, Basic Module (0920–0214)—Revision—The National Center for Health Statistics (NCHS)—The annual National Health Interview Survey (NHIS) is a basic source of general statistics on the health of the U.S. population. Due to the integration of health surveys in the Department of Health and Human Services, the NHIS also has become the sampling frame and first stage of data collection for other major surveys, including the Medical Expenditure Panel Survey, the National Survey of Family Growth, and the National Health and Nutrition Examination Survey. By linking to the NHIS, the analysis potential of these surveys increases. The NHIS has long been used by government, university, and private researchers to evaluate both general health and specific issues, such as cancer, AIDS, and childhood immunizations. Journalists use its data to inform the general public. It will continue to be a leading source of data for the Congressionally-mandated ''Health US'' and related publications, as well as the single most important source of statistics to track progress toward the National Health Promotion and Disease Prevention Objectives, ''Healthy People 2000.''

Because of survey integration and changes in the health and health care of the U.S. population, demands on the NHIS have changed and increased, leading to a major redesign of the annual core questionnaire, or Basic Module, and a redesign of the data collection system from paper questionnaires to computer assisted personal interviews (CAPI). Those redesigned elements were partially implemented in 1996 and fully implemented in 1997 and are expected to be in the field until 2006. This clearance is for the fifth full year of data collection using the Basic Module on CAPI, and for implementation of the second ''Periodic Module'', which include additional detail questions on conditions, access to care, disabilities, and health care utilization. The ''Periodic Module'' will repeat a similar survey conducted in 1992, and will help track many of the Health People 2010 objectives. This data collection, planned for January–December 2001, will result in publication of new national estimates of health statistics, release of public use micro data files, and a sampling frame for other integrated surveys. The annualized burden is 48,600 hours.

Questionnaire (respondent)	Number of respondents	Number of responses per respondent	Average burden per respondent (in hours)
Family Core (adult family member)	42,000	1	21/60
Adult Core (sample adult)	42,000	1	21/60
Child Core (adult family member)	18,000	1	15/60
Periodic Module (sample adult)	42,000	1	21/60
All households	42,000	1	1 10/60

Dated: September 5, 2000.

Nancy Cheal,

Acting Associate Director for Policy, Planning, and Evaluation, Centers for Disease Control and Prevention (CDC).

[FR Doc. 00–23204 Filed 9–8–00; 8:45 am]

BILLING CODE 4163–18–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00F–1487]

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 as a component of a post-chill carcass spray or dip when applied to poultry meat, organs, or related parts or trim.

DATES: Submit written comments on the petitioner's environmental assessment by October 11, 2000.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Robert L. Martin, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 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 food additive petition (FAP 0A4–722) has been filed by Alcide Corp., 8561 154th Ave. NE., Redmond, WA 98052. The petition proposes to amend the food additive regulations in § 173.325 *Acidified sodium chlorite solutions* (21 CFR 173.325) to provide for the safe use of acidified sodium chlorite solutions as a component of a post-chill carcass spray or dip when applied to poultry meat, organs, or related parts or trim.

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 (address above) for public review and comment. Interested persons may submit to the Dockets Management Branch written comments by October 11, 2000. 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. 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 23, 2000.

Alan M. Rulis,

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

[FR Doc. 00–23161 Filed 9–8–00; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 00F–1488]

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 as an antimicrobial agent on processed, comminuted, or formed meat products prior to packaging.

DATES: Submit written comments on the petitioner's environmental assessment by October 11, 2000.

ADDRESSES: Submit written comments to the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

FOR FURTHER INFORMATION CONTACT: Robert L. Martin, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 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 food additive petition (FAP 0A4724) has been filed by Alcide Corp., 8561 154th Ave. NE., Redmond, WA 98052. The petition proposes to amend the food additive regulations in § 173.325 *Acidified sodium chlorite solutions* (21 CFR 173.325) to provide for the safe use of acidified sodium chlorite solutions as an antimicrobial agent on processed, comminuted, or formed meat products prior to packaging.

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 (address above) for public review and comment. Interested persons may submit to the Dockets Management Branch written comments by October 11, 2000. 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. 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 24, 2000.

Alan M. Rulis,

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

[FR Doc. 00–23162 Filed 9–8–00; 8:45 am]

BILLING CODE 4160–01–F