epilepsy; (3) directs and administers the development of a national, state, and local surveillance system of tracking health-related quality of life (HRQOL) among U.S. residents; (4) administers grants, cooperative agreements, contracts, and other procurement requests to implement evidence-based health promotion interventions and disseminate arthritis prevention and epilepsy education messages; (5) develops, validates, and refines HRQOL measure for use in tracking and prevention research at each life stage; (6) directs and coordinates the evaluation of community and state-based intervention programs for arthritis and epilepsy; (7) developes arthritis epidemiology capacity and other arthritis programmatic capabilities in state health department settings; and (8) disseminates health promotion and disease prevention information through national advocacy partners for arthritis and epilepsy.

Dated: October 12, 2006.

William H. Gimson,

Chief Operating Officer, Centers for Disease Control and Prevention (CDC).

[FR Doc. 06–8869 Filed 10–24–06; 8:45 am]

BILLING CODE 4160–18–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 2006F–0409]

Safe Foods Corporation; Filing of Food Additive Petition

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Safe Foods Corporation has filed a petition proposing that the food additive regulations be amended to expand the conditions for the safe use of cetylpyridinium chloride as an antimicrobial agent in a pre-chiller or post-chiller solution for application to raw poultry carcasses.

DATES: Submit written or electronic comments on the petitioner's environmental assessment by November 24, 2006.

ADDRESSES: Submit written comments to the Division of Dockets Management (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: Raphael Davy, Center for Food Safety and Applied Nutrition (HFS–265), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 301–436–1272.

SUPPLEMENTARY INFORMATION: Under the Federal Food, Drug, and Cosmetic Act (section 409(b)(5) (21 U.S.C. 348(b)(5))), notice is given that a food additive petition (FAP 6A4767) has been filed by Safe Foods Corporation, c/o Keller and Heckman LLP, 1001 G St. NW., suite 500 West, Washington, D.C. 20001. The petition proposes to amend the food additive regulations in § 173.375 *Cetylpyridinium chloride* (21 CFR 173.375) to expand the conditions for the safe use of cetylpyridinium chloride as an antimicrobial agent in a pre-chiller or post-chiller solution for application to raw poultry carcasses.

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 Division of Dockets Management (see **ADDRESSES**) for public review and comment. Interested persons may submit to the Division of Dockets Management written or electronic comments by (see **DATES**). Two copies of any written 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 Division of Dockets Management 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 25.51(b).

Dated: October 17, 2006.

Laura M. Tarantino,

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

[FR Doc. E6–17834 Filed 10–24–06; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Health Resources and Services Administration

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

Periodically, the Health Resources and Services Administration (HRSA) publishes abstracts of information collection requests under review by the Office of Management and Budget (OMB), in compliance with the Paperwork Reduction Act of 1995 (44 U.S.C. Chapter 35). To request a copy of the clearance requests submitted to OMB for review, call the HRSA Reports Clearance Office on (301)–443–1129.

The following request has been submitted to the Office of Management and Budget for review under the Paperwork Reduction Act of 1995:

Proposed Project: Grants for Hospital Construction and Modernization—Federal Right of Recovery and Waiver of Recovery (42 CFR Part 124, Subpart H) (OMB No. 0915–0099Extension)

The regulation known as ''Federal Right of Recovery and Waiver of Recovery,'' provides a means for the Federal Government to recover grant funds and a method of calculating interest when a grant-assisted facility under Titles VI and XVI of the Public Health Service Act is sold or leased, or there is a change in use of the facility. It also allows for a waiver of the right of recovery under certain circumstances. Facilities are required to provide written notice to the Federal Government when such a change occurs and to provide copies of sales contracts, lease agreements, estimates of current assets and liabilities, value of equipment, expected value of land on the new owner's books and remaining depreciation for all fixed assets involved in the transactions, and other information and documents pertinent to the change of status.

Estimates of annualized burden are as follows: