Attachment J.—Certification Regarding Environmental Tobacco Smoke

Public Law 103227, Part C Environmental Tobacco Smoke, also known as the Pro Children Act of 1994, requires that smoking not be permitted in any portion of any indoor routinely owned or leased or contracted for by an entity and used routinely or regularly for provision of health, day care, education, or library services to children under the age of 18, if the services are funded by Federal programs either directly or through State or local governments, by Federal grant, contract, loan, or loan guarantee. The law does not apply to children's services provided in private residences, facilities funded solely by Medicare or Medicaid funds, and portions of facilities used for inpatient drug or alcohol treatment. Failure to comply with the provisions of the law may result in the imposition of a civil monetary penalty of up to $1000 per day and/or the imposition of an administrative compliance order on the responsible entity. By signing and submitting this application the applicant/grantee certifies that it will comply with the requirements of the Act.

The applicant/grantee further agrees that it will require the language of this certification be included in any subawards which contain provisions for the children's services and that all subgrantees shall certify accordingly.

Attachment K.—

Title 45 of the Code of Federal Regulations

Part 16—Department of Grant Appeals Process
Part 74—Administration of Grants (grants and subgrants to entities)
Part 75—Informal Grant Appeal Procedures
Part 76—Debarment and Suspension from Eligibility for Financial Assistance

Subpart F—Drug Free Workplace Requirements

Part 80—Non-Discrimination—Under Programs Receiving Federal Assistance through the Department of Health and Human Services Effectuation of Title VI of the Civil Rights Act Of 1964
Part 81—Practice and Procedures for Hearings Under Part 80 of this Title
Part 83—Regulation for the Administration and Enforcement of Section 799A and 845 of the Public Health Service Act
Part 84—Non-discrimination on the Basis of Handicap in Programs and Activities Receiving Federal Financial Assistance
Part 85—Enforcement of Non-Discrimination on the Basis of Handicap in Programs or Activities Conducted by the Department of Health and Human Services
Part 86—Nondiscrimination on the Basis of Sex in Education Programs and Activities Receiving or Benefiting from Federal Financial Assistance
Part 91—Non-discrimination on the Basis of Age in Health and Human Services Programs or Activities Receiving Federal Financial Assistance
Part 92—Uniform Administrative Requirements for Grants and Cooperative Agreements to States and Local Governments (**Federal Register**, March 11, 1988)
Part 93—New Restrictions on Lobbying
Part 100—Intergovernmental Review of Department of Health and Human Services Programs and Activities

[FR Doc. 02–11217 Filed 5–6–02; 8:45 am]

BILLING CODE 4184–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 02F–0181]

Safe Foods Corp.; 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 Corp. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of cetylpyridinium chloride as an antimicrobial agent in poultry processing.

DATES: DATES: Submit written comments on the petitioner's environmental assessment by June 6, 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: Andrew D. Laumbach, Center for Food Safety and Applied Nutrition (HFS–265), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 202–418–3071.

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 2A4736) has been filed by Safe Foods Corp., c/o Keller and Heckman LLP, 1001 G St. NW., Washington, DC 20001. The petition proposes to amend the food additive regulations in part 173 Secondary Direct Food Additives Permitted in Food for Human Consumption (21 CFR part 173) to provide for the safe use of cetylpyridinium chloride as an antimicrobial agent in poultry processing.

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 (see **ADDRESSES**) for public review and comment. Interested persons may submit to the Dockets Management Branch written comments by June 6, 2002. 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 25.51(b).

Dated: April 11, 2002.

Laura M. Tarantino,

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

[FR Doc. 02–11255 Filed 5–6–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

[Docket No. 01P–0315]

Determination That Acetaminophen and Codeine Phosphate Tablets, 500 Milligrams (mg)/15 mg, 500 mg/30 mg, and 500 mg/60 mg, Were Not Withdrawn From Sale for Reasons of Safety or Effectiveness

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing its determination that acetaminophen and codeine phosphate tablets, 500 mg/15 mg, 500 mg/30 mg, and 500 mg/60 mg, were not withdrawn from sale for reasons of safety or effectiveness. This determination will allow FDA to approve abbreviated new drug applications (ANDAs) for acetaminophen and codeine phosphate