

practices in violation of Section 5(a) of the FTC Act. However, in *United States v. Hopkins Dodge, Inc.*, 849 F.2d 311 (8th Cir. 1988), the United States Court of Appeals for the Eighth Circuit held that the Commission's failure to explicitly state in these two cases that credit advertising violations of the TILA and Regulation Z are unfair or deceptive acts or practices precluded use of the two cases as predicates to civil penalty enforcement actions pursuant to section 5(m)(1)(B) of the FTC Act.

On January 31, 1989, the Commission issued an order against Reliable Mortgage Corp. *et al.* to show cause why the proceedings against them should not be reopened to consider modification of the decision therein to clarify that respondents' credit advertising violations of the TILA were also unfair and deceptive acts or practices in violation of the FTC Act. On January 31, 1989, the Commission also issued an order against Seekonk Freezer Meats *et al.* to show cause why the proceedings against them should not be reopened to consider modification of the decision therein to clarify that respondents' credit advertising violations of the TILA were either unfair or unfair and deceptive acts or practices in violation of the FTC Act. The Commission also issued a press release and published a notice in the *Federal Register* on February 17, 1989, announcing a 30-day period for public comment on the proposed reopenings.

Respondents failed to answer the show cause orders. (Respondent Reliable did, however, informally reply by letter to the FTC staff that it had no objections to the proposed reopening). No comments were received in response to the *Federal Register* notice.

As a result, on September 25, 1989, the Commission issued orders reopening the proceedings in *Reliable* and *Seekonk*, and directing briefs to consider modification of the decisions contained therein. The Commission now intends to decide whether to modify the decisions in *Reliable* and *Seekonk* to state expressly that the credit advertising practices addressed in *Reliable* constitute unfair and deceptive acts or practices in violation of section 5(a) of the FTC Act and that the credit advertising practices addressed in *Seekonk* constitute either unfair or unfair and deceptive acts or practices in violation of section 5(a) of the FTC Act. Interested parties may submit amicus curiae briefs as noted above.

By the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 89-27089 Filed 11-16-89; 8:45 am]

BILLING CODE 6750-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Office of the Secretary

Agency Forms Submitted to the Office of Management and Budget for Clearance

On Fridays, the Department of Health and Human Services, Office of the Secretary publishes a list of information collections it has submitted to the Office of Management and Budget (OMB) for clearance in compliance with the Paperwork Reduction Act (44 U.S.C. chapter 35). The following are those information collections recently submitted to OMB.

1. Teenage Parent Demonstration: 24-Month Follow-up—New—This survey is part of the impact evaluation of the Teenage Parent Demonstration Projects being conducted in Illinois and New Jersey. The survey will gather information on outcomes from demonstration participants which will contribute to the evaluation of the demonstrations. The purpose of the demonstration projects is to identify interventions which promote self-sufficiency among teenage parents. *Respondents:* individuals; *Number of Respondents:* 4,774; *Frequency of Response:* One time; *Average Burden per Response:* 1.31 Hours; *Total Burden:* 6,244 Hours.

OMB Desk Officer: Shannah Koss-McCallum.

Copies of the information collection packages listed above can be obtained by calling the OS Reports Clearance Officer on (202) 245-6511. Written comments and recommendations for the proposed information collection should be sent directly to the OMB desk officer designated above at the following address: OMB Reports Management Branch, New Executive Office Building, Room 3208, Washington, DC 20503.

Dated: November 8, 1989.

James F. Trickett,

Deputy Assistant Secretary for Management and Acquisition.

[FR Doc. 89-26945 Filed 11-16-89; 8:45 pm]

BILLING CODE 4150-04-M

Alcohol, Drug Abuse, and Mental Health Administration

Advisory Committee Meeting in December

AGENCY: Alcohol, Drug Abuse, and Mental Health Administration, HHS.

ACTION: Notice of meeting.

SUMMARY: This notice sets forth the schedule and proposed agenda of the forthcoming meeting of one of the agency's advisory committees in the month of December 1989.

The initial review group will be performing review of applications for Federal assistance. Therefore, portions of the meetings will be closed to the public as determined by the Administrator, ADAMHA, in accordance with 5 U.S.C. 552(b) (6) and 5 U.S.C. app. 2 10(d).

Notice of this meeting is required under the Federal Advisory Committee Act, Public Law 92-463.

Committee Name: Mental Health Acquired Immunodeficiency Syndrome Research Review Committee, NIMH.

Date and Time: December 5-6: 8:00 a.m.

Place: The Canterbury Hotel, 1733 N Street NW., Washington, DC 20036.

Status of Meeting: Open—December 5: 8:30-9:15 a.m. Closed—Otherwise

Contact: Regina Thomas, Room 9C-15, Parklawn Building, 5600 Fishers Lane, Rockville, MD 20857, (301) 443-6470.

Purpose: The Committee is charged with the initial review of applications for assistance from the National Institute of Mental Health for support of activities in the fields of research and psychoneuroimmunological, psychosocial, behavioral, and psychological aspects of AIDS as they relate to mental health, with recommendations to the National Advisory Mental Health Council for final review.

Substantive information, a summary of the meeting, and a roster of committee members may be obtained from Ms. Joanna Kieffer, NIMH Committee Management Officer, Room 9-105, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857, (301) 443-4333.

Dated: November 13, 1989.

Peggy W. Cockrill,

Committee Management Officer, Alcohol, Drug Abuse, and Mental Health Administration.

[FR Doc. 89-26992 Filed 11-16-89; 8:45 am]

BILLING CODE 4160-20-M

Food and Drug Administration

[Docket No. 89F-0462]

The Dow Chemical Co., E.I. Du Pont de Nemours and Co., B.F. Goodrich; Filing of Food Additive Petition**AGENCY:** Food and Drug Administration.**ACTION:** Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that the Dow Chemical Co., E.I. du Pont de Nemours and Co., and B.F. Goodrich have filed a petition proposing that the food additive regulations be amended to provide for the safe use of polyurethane resins, derived from reactions of diphenylmethane diisocyanate with 1,4-butanediol and polytetramethylene ether glycol, as rubber articles intended for repeated use in contact with food.

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C Street, SW., Washington, DC 20204, 202-472-5690.

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 OB4182) has been filed by the Dow Chemical Co., 1803 Bldg., Door 7, Midland, MI 48674, E.I. Du Pont de Nemours and Co., 1007 Market St., Wilmington, DE 19898, and B.F. Goodrich, 3925 Embassy Pkwy., Akron, OH 44313, proposed that § 177.2600 Rubber articles intended for repeated use (21 CFR 177.2600) of the food additive regulations be amended to provide for the safe use of polyurethane resins, derived from the reaction of diphenylmethane diisocyanate with 1,4-butanediol and polytetramethylene ether glycol, as rubber articles intended for repeated use in contact with food.

The potential environmental impact of this action is being reviewed. If 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: November 8, 1989.

Richard J. Ronk,

Deputy Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89-26990 Filed 11-16-89; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 89F-0452]

Enzyme Bio-Systems Ltd.; Filing of Food Additive Petition**AGENCY:** Food and Drug Administration.**ACTION:** Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that Enzyme Bio-Systems Ltd. has filed a petition proposing that the food additive regulations be amended to provide for the safe use of dimethylamine-epichlorohydrin and acrylamide acrylic acid resins as fixing agents for immobilizing glucose isomerase enzyme.

FOR FURTHER INFORMATION CONTACT: Geraldine E. Harris, Center for Food Safety and Applied Nutrition (HFF-334), Food and Drug Administration, 200 C Street SW., Washington, DC 20204, 202-426-9463.

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 Enzyme Bio-Systems Ltd., International Plaza, Route 9W, Englewood Cliffs, NJ 07632, has filed a petition (FAP 9A4175), proposing that the food additive regulations be amended to provide for the safe use of dimethylamine-epichlorohydrin and acrylamide acrylic acid resins as fixing agents for immobilizing glucose isomerase enzyme.

The potential environmental impact of this action is being reviewed. If 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: November 8, 1989.

Richard J. Ronk,

Deputy Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89-26988 Filed 11-16-89; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 89F-0451]

ICI Americas, Inc.; Filing of Food Additive Petition**AGENCY:** Food and Drug Administration.**ACTION:** Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that ICI Americas, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of polyethylene glycol (MW 1500-4000)/poly(12-hydroxystearic acid)

copolymers as a stabilizer in the preparation of polyacrylamide retention and drainage aids used in the manufacture of paper and paperboard intended to contact aqueous and fatty food.

FOR FURTHER INFORMATION CONTACT: Andrew D. Laumbach, Center for Food Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C Street, SW., Washington, DC 20204, 202-472-5690.

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 OB4179) has been filed by ICI Americas, Inc., Concord Pike & Murphy Rd., Wilmington, DE 19877, proposing that § 176.170 Components of paper and paperboard in contact with aqueous and fatty foods (21 CFR 176.170) be amended to provide for the safe use of polyethylene glycol (MW 1500-4000)/poly(12-hydroxystearic acid) copolymers as a stabilizer in the preparation of polyacrylamide retention and drainage aids used in the manufacture of paper and paperboard intended to contact aqueous and fatty foods.

The potential environmental impact of this action is being reviewed. If 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: November 8, 1989.

Richard J. Ronk,

Deputy Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89-26989 Filed 11-16-89; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 89F-0450]

ICI Americas, Inc.; Filing of Food Additive Petition**AGENCY:** Food and Drug Administration.**ACTION:** Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that ICI Americas, Inc., has filed a petition proposing that the food additive regulations be amended to provide for the safe use of poly(isobutene)/maleic anhydride, diethanolamine reaction product as a surfactant in the preparation of polyacrylamide retention and drainage aids used in the

manufacture of paper and paperboard intention to contact aqueous and fatty foods.

FOR FURTHER INFORMATION CONTACT: Andrew D. Laumbach, Center for Food Safety and Applied Nutrition (HFF-335), Food and Drug Administration, 200 C Street SW., Washington, DC 20204, 202-472-5690.

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 OB4178) has been filed by ICI Americas, Inc., Concord Pike and Murphy Rd., Wilmington, DE 19897, proposing that § 176.170 Components of paper and paperboard in contact with aqueous and fatty foods (21 CFR 176.170) be amended to provide for the safe use of poly(isobutene)/maleic anhydrides, diethanolamine reaction product as a surfactant in the preparation of polyacrylamide retention and drainage aids used in the manufacture of paper and paperboard intended to contact aqueous and fatty foods.

The potential environmental impact of this action is being reviewed. If 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: November 8, 1989.

Richard J. Ronk,

Deputy Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89-26991 Filed 11-16-89; 8:45 am]

BILLING CODE 4160-01-M

[Docket No. 89P-0225]

Ice Cream Deviating From the Standard of Identity; Temporary Permit for Market Testing

AGENCY: Food and Drug Administration, HHS.

ACTION: Notice.

SUMMARY: The Food and Drug Administration (FDA) is announcing that a temporary permit has been issued to the Hershey Creamery Co. to market test a product designated as "light ice cream" that deviates from the U.S. standard of identity for ice cream (21 CFR 135.110). The purpose of the temporary permit is to allow the applicant to measure consumer acceptance of the product, identify mass production problems, and assess commercial feasibility.

DATES: This permit is effective for 15 months, beginning on the date the food is introduced or caused to be introduced into interstate commerce, but no later than February 15, 1990.

FOR FURTHER INFORMATION CONTACT: Joanne Travers, Center for Food Safety and Applied Nutrition (HFF-414), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-485-0324.

SUPPLEMENTARY INFORMATION: In accordance with 21 CFR 130.17 concerning temporary permits to facilitate market testing of foods deviating from the requirements of standards of identity promulgated under section 401 of the Federal Food, Drug and Cosmetic Act (21 U.S.C. 341), FDA is giving notice that a temporary permit has been issued to the Hershey Creamery Co., 301 South Cameron St., P.O. Box 1821, Harrisburg, PA 17105.

The permit covers limited interstate marketing tests of a product that deviates from the U.S. standard of identity for ice cream in 21 CFR 135.110 in that: (1) The milkfat content of the product is reduced by at least 50 percent; and (2) sufficient vitamin A palmitate is added in a suitable carrier to ensure that a ½-cup serving of the product contains 8 percent of the U.S. Recommended Daily Allowance for vitamin A. The product meets all requirements of the standard with the exception of these deviations. The purpose of this variation is to offer the consumer a product that is nutritionally equivalent to ice cream but contains fewer calories and less fat.

For the purpose of this permit, the name of the product is "light ice cream." The principal display panel of the label must include the statements "reduced calorie" and "reduced fat" following the name. In addition, the label must bear the comparative statements "⅓ fewer calories than our regular ice cream" and "50 percent less fat than our regular ice cream."

The product complies with reduced calories labeling requirements in 21 CFR 105.66(d). In accordance with FDA's current views, reduced fat food labeling is acceptable because there is a 50 percent reduction in the fat content of the product. The information panel of the label must bear nutrition labeling in accordance with 21 CFR 101.9.

The permit provides for the temporary marketing of a total of 500,000 cases of two half-gallon containers. The test product will be produced and packaged at the Hershey Creamery Co., Harrisburg, PA 17105, and will be distributed in Connecticut, Delaware, Maine, Maryland, Massachusetts, New Hampshire, New Jersey, New York,

North Carolina, Ohio, Pennsylvania, Rhode Island, Vermont, Virginia, and West Virginia.

Each of the ingredients used in the food must be stated on the label as required by the applicable sections of 21 CFR part 101. This permit is effective for 15 months, beginning on the date the food is introduced into interstate commerce, but no later than _____.

Dated: November 8, 1989.

Richard J. Ronk,

Deputy Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 89-27056 Filed 11-16-89; 8:45 am]

BILLING CODE 4160-01-M

Health Resources and Services Administration

Filing of Annual Report of Federal Advisory Committee; Advisory Council on Nurses Education

Notice is hereby given that pursuant to section 13 of Public Law 92-463, the Annual Report for the following Health Resources and Services Administration's Federal Advisory Committee has been filed with the Library of Congress:

Advisory Council on Nurses Education

Copies are available to the public for inspection at the Library of Congress Newspaper and Current Periodical Reading Room, Room 1026, Thomas Jefferson Building, Second Street and Independence Avenue SE., Washington, DC, or weekdays between 9:00 a.m. and 4:30 p.m. at the Department of Health and Human Services, Department Library, HHS North Building, Room G-400, 330 Independence Avenue SW., Washington, DC, telephone (202) 245-6791. Copies may be obtained from: Dr. Mary S. Hill, Executive Secretary, Advisory Council on Nurses Education, Room 5C-14, Parklawn Building, 5600 Fishers Lane, Rockville, Maryland 20857, Telephone (301) 443-6193.

Dated: November 13, 1989.

Jackie E. Baum,

Advisory Committee Management Officer, HRSA.

[FR Doc. 89-26986 Filed 11-16-89; 8:45 am]

BILLING CODE 4160-15-M

National Institutes of Health

National Biotechnology Policy Board; Establishment

Pursuant to the Federal Advisory Committee Act of October 6, 1972 (Pub.