119°13′00″ W; to lat. 38°10′00″ N, long. 118°34′00″ W, thence to the point of beginning.

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Issued in Los Angeles, California, on November 21, 1995.

James H. Snow,

Acting Manager, Air Traffic Division, Western-Pacific Region.

[FR Doc. 95–29348 Filed 11–30–95; 8:45 am]

BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 177

[Docket No. 93F–0166]

Indirect Food Additives: Polymers

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of pyromellitic dianhydride as a modifier in the manufacture of polyethylene terephthalate copolymers intended for food-contact applications. This action is in response to a petition filed by M. & G. Ricerche S.p.A.

DATES: Effective December 1, 1995; written objections and requests for a hearing by January 2, 1996.

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

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3081.

SUPPLEMENTARY INFORMATION: In a notice published in the Federal Register of August 12, 1993 (58 FR 42975), FDA announced that a food additive petition (FAP 3B4375) had been filed by M. & G. Ricerche, S.p.A., c/o Keller and Heckman, 1001 G St. NW., suite 500 West, Washington, DC 20001 (formerly c/o Delta Analytical Corp., 7910 Woodmont Ave., suite 1000, Bethesda, MD 20814). The petition proposed to amend the food additive regulations in § 177.1630 *Polyethylene phthalate polymers* (21 CFR 177.1630) to provide for the safe use of pyromellitic dianhydride as a modifier in the manufacture of polyethylene terephthalate copolymers intended for food-contact applications.

FDA has evaluated data in the petition and other relevant material. The agency concludes that the proposed food additive use in polyethylene terephthalate food-contact articles is safe, and the regulation in § 177.1630 should be amended as set forth below.

In accordance with § 171.1(h) (21 CFR 171.1(h)), the petition and the documents that FDA considered and relied upon in reaching its decision to approve the petition are available for inspection at the Center for Food Safety and Applied Nutrition by appointment with the information contact person listed above. As provided in 21 CFR 171.1(h), the agency will delete from the documents any materials that are not available for public disclosure before making the documents available for inspection.

The agency has carefully considered the potential environmental effects of this action. FDA has concluded that the action will not have a significant impact on the human environment, and that an environmental impact statement is not required. The agency's finding of no significant impact and the evidence supporting that finding, contained in an environmental assessment, may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

Any person who will be adversely affected by this regulation may at any time on or before January 2, 1996, file with the Dockets Management Branch (address above) written objections thereto. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provisions of the regulation to which objection is made and the grounds for the objection. Each numbered objection on which a hearing is requested shall specifically so state. Failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is requested shall include a detailed description and analysis of the specific factual information intended to be presented in support of the objection in the event that a hearing is held. Failure to include such a description and analysis for any particular objection shall constitute a waiver of the right to a hearing on the objection. Three copies of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this document. Any objections received in response to the regulation may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 177

Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Director, Center for Food Safety and Applied Nutrition, 21 CFR part 177 is amended as follows:

PART 177—INDIRECT FOOD ADDITIVES: POLYMERS

1. The authority citation for 21 CFR part 177 continues to read as follows:

Authority: Secs. 201, 402, 409, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 342, 348, 379e).

2. Section 177.1630 is amended in paragraph (e)(4) by alphabetically adding a new substance to paragraph (i) in the ''List of Substances and Limitations'' to read as follows:

§ 177.1630 Polyethylene phthalate polymers.

* * * * *

(e) * * *

(4) * * *

List of Substances and Limitations

(i) * * *

Ethylene terephthalate copolymers: Prepared by the condensation of dimethyl terephthalate or terephthalic acid with ethylene glycol, modified with one or more of the following: Azelaic acid, dimethyl azelate, dimethyl sebacate, sebacic acid, pyromellitic dianhydride. The level of pyromellitic dianhydride shall not exceed 0.5 percent by weight of the finished copolymer which may be used under conditions of use E through H as described in Table 2 of § 176.170(c) of this chapter.

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Dated: November 21, 1995.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 95–29219 Filed 11–30–95; 8:45 am]

BILLING CODE 4160–01–F

LIBRARY OF CONGRESS

Copyright Office

37 CFR Part 253

[Docket No. 95–3 CARP]

Cost of Living Adjustment for Performance of Musical Compositions by Colleges and Universities

AGENCY: Copyright Office, Library of Congress.

ACTION: Final rule.

SUMMARY: The Copyright Office of the Library of Congress announces a cost of