The Commission orders

The October 31, 1985 order is amended to extend the term of MichCon's transportation service for ANR through October 9, 1987. In all other respects said order shall remain in full force and effect.

By the Commission.

Lois D. Cashell,

Acting Secretary.

[FR Doc. 85-30133 Filed 12-19-85; 8:45 am]

BILLING CODE 6717-01-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 175

[Docket No. 85F-0478]

Indirect Food Additives; Adhesives and Components of Coatings

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of polyamides consisting of the homopolymer of Nylon 12 derived from *omega*-aminododecanoic acid as side seam cements that may contact food. This action responds to a petition filed by Springborn Regulatory Services, Inc., on behalf of UBE Industries, Ltd.

DATES: Effective December 20, 1985; objections by January 21, 1986.

ADDRESS: Written objections to the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857.

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

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of March 8, 1985 (50 FR 9521), FDA announced that a petition (FAP 4B3796) had been filed by Springborn Regulatory Services, Inc., Enfield, CT 06082, on behalf of UBE Industries, Ltd., proposing that the food additive regulations be amended to provide for the safe use of polyamides consisting of the homopolymer of Nylon 12 derived from *omega*-aminododecanoic acid as side seam cements that may contact food.

FDA has evaluated the data in the petition and other relevant material. The agency concludes that the proposed food additive use is safe, and that the regulations 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 (address above) 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 and 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 may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday. FDA's regulations implementing the National Environmental Policy Act (21 CFR Part 25) have been replaced by a rule published in the **Federal Register** of April 26, 1985 (50 FR 16636, effective July 25, 1985). Under the new rule, an action of this type would require an abbreviated environmental assessment under 21 CFR 25.31a(b)(1).

Any person who will be adversely affected by this regulation may at any time on or before January 21, 1986, submit to the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857, written objections thereto and may make a written request for a public hearing on the stated objections. Each objection shall be separately numbered and each numbered objection shall specify with particularity the provision of the regulation to which objection is made. 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 regulation. Received objections may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 175

Adhesives, 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, Part 175 is amended as follows:

PART 175—INDIRECT FOOD ADDITIVES: ADHESIVES AND COMPONENTS OF COATINGS

1. The authority citation for 21 CFR Part 175 continues to read as follows:

Authority: Secs. 201(s), 409, 72 Stat. 1784-1788 as amended (21 U.S.C. 321(s), 348); 21 CFR 5.10 and 5.61.

2. In § 175.300(b)(3) (xxxii) by alphabetically inserting a new item under "Polyamides consisting of the following" to read as follows:

§ 175.300 Resinous and polymeric coatings.

* * * * *

(b) * * *

(3) * * *

(xxxii) * * *

Polyamides consisting of the following:

Homopolymer of *omega*-aminododecanoic acid, CAS Reg. No. 24937-16-4.

* * * * *

Dated: December 10, 1985.

Sanford A. Miller,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 85-30101 Filed 12-19-85; 8:45 am]

BILLING CODE 4160-01-M

21 CFR Part 177

[Docket No. 85F-0374]

Indirect Food Additives; Polymers

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of trimethyleneglycol di(*p*-aminobenzoate) as a curing agent in the manufacture of polyurethane resins intended for use in contact with dry food. This action responds to a petition filed by Polaroid Corp.

DATES: Effective December 20, 1985; objections by January 21, 1986.

ADDRESS: Written objections to the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857.

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

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of September 19, 1985 (50 FR 38036), FDA announced that a petition (FAP 4B3765) had been filed by Polaroid Corp., 238 South Main St., Assonet, MA 02702, proposing that § 177.1680 (21 CFR 177.1680) be amended to provide for the safe use of trimethyleneglycol di(*p*-aminobenzoate) in the manufacture of polyurethane resins intended to contact dry food.

FDA has evaluated data in the petition and other relevant material. The agency concludes that the proposed food additive use is safe, and that the regulations 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 (address above) by appointment with the information contact person listed above. As provided in 21 CFR 171.1(h)(2), 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 previously considered the environmental effects of this rule as announced in the notice of filing for FAP 4B3765 (September 19, 1985; 50 FR 38036). No new information or comments have been received that would affect the agency's previous determination that there is no sigificant impact on the human environment and that an environmental impact statement is not required.

Any person who will be adversely affected by this regulation may at any time on or before January 21, 1986, submit to the Dockets Management Branch (address above) written objections thereto and may make a written request for a public hearing on the stated objections. Each objection shall be separately numbered and each numbered objection shall specify with particularity the provision of the regulation to which ojection is made. 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 regulation. Received objections may be seen in the office above between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects in 21 CFR Part 177

Food additives, Polymeric 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 Center for Food Safety and Applied Nutrition, 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(s), 409, 72 Stat. 1784-1788 as amended (21 U.S.C. 321(s), 348); 21 CFR 5.10 and 5.61.

2. Part 177 is amended in § 177.1680(b) by alphabetically inserting a new item in the list of substances to read as follows:

§ 177.1680 Polyurethane resins.

* * * * *

(b) * * *

List of substances	Limitations
* * * * *	*
Trimethyleneglycol di (*p*-aminobenzoate) (CAS Reg. No. 57609-64-0)	As a curing agent
* * * * *	*

* * * * *

Dated: December 10, 1985.

Sanford A. Miller,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 85-30100 Filed 12-19-85; 8:45 am]

BILLING CODE 4160-01-M

21 CFR Part 178

[Docket No. 84F-0427]

Indirect Food Additives: Adjuvants, Production Aids, and Sanitizers

AGENCY: Food and Drug Administration.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of hydrogen peroxide for sterilizing food-contact surfaces prepared from polystyrene and rubber-modified polystyrene resins. This action responds to a petition filed by The Dow Chemical Co.

DATES: Effective December 20, 1985, objections by January 21, 1986.

ADDRESS: Written objections may be sent to the Dockets Management Branch (HFA-305), Food and Drug Administration, Rm. 4-62, 5600 Fishers Lane, Rockville, MD 20857.

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

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of January 9, 1985 (50 FR 1132), FDA announced that a food additive petition (FAP 5B3841) had been filed by The Dow Chemical Co., 1803 Building, Door 7, Midland, MI 48640, proposing that the food additive regulations be amended in 21 CFR 178.1005 to provide for the safe use of hydrogen peroxide for sterilizing food-contact surfaces prepared from polystyrene and rubber-modified polystyrene resins complying with 21 CFR 177.1640.

FDA has evaluated the data in the petition and other relevant material. The agency concludes that the proposed use is safe, and that the regulations 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 (address above) 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 and 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 may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday. FDA's regulations implementing the National Environmental Policy Act (21 CFR Part 25) have been replaced by a rule published in the Federal Register of April 26, 1985 (50 FR 16636, effective July 25, 1985). Under the new rule, an action of this type would require an abbreviated environmental assessment under 21 CFR 25.31a(b)(5).

Any person who will be adversely affected by this regulation may at any time on or before January 21, 1986, 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 178

Food Additives, Food packaging, Sanitizing solutions.

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, Part 178 is amended as follows:

PART 178—INDIRECT FOOD ADDITIVES: ADJUVANTS, PRODUCTION AIDS, AND SANITIZERS

1. The authority citation for 21 CFR Part 178 continues to read as follows:

Authority: Secs. 201(s), 409, 72 Stat. 1784–1788 as amended (21 U.S.C. 321(s), 348); 21 CFR 5.10 and 5.61.

2. In § 178.1005 by revising paragraph (e)(1) to read as follows:

§ 178.1005 Hydrogen peroxide solution.

* * * * *

(e) *Conditions of use.* (1) Hydrogen peroxide solution identified in and complying with the specifications in this section may be used by itself or in combination with other processes to treat food-contact surfaces prepared from ionomeric resins complying with § 177.1330 of this chapter, ethylene-methyl acrylate copolymer resins complying with § 177.1340 of this chapter, ethylene-vinyl acetate copolymers complying with § 177.1350 of this chapter, olefin polymers complying with § 177.1520 of this chapter, polyethylene terephthalate polymers complying with § 177.1630 of this chapter (excluding polymers described in § 177.1630(c)), and polystyrene and rubbers-modified polystyrene polymers complying with § 177.1640 to attain commercial sterility at least equivalent to that attainable by thermal processing for metal containers as provided for in Part 113 of this chapter.

* * * * *

Dated: December 10, 1985.

Sanford A. Miller,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 85-30102 Filed 12-19-85; 8:45 am]

BILLING CODE 4160-01-M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Office of the Assistant Secretary for Housing—Federal Housing Commissioner

24 CFR Parts 232 and 235

[Docket No. R-85-1269; FR-2195]

Mortgage Insurance—Changes in Interest Rates

AGENCY: Office of the Assistant Secretary for Housing—Federal Housing Commissioner, HUD.

ACTION: Final rule.

SUMMARY: This change in the regulations decreases the maximum allowable interest rate on section 232 (Mortgage Insurance for Nursing Homes) and on section 235 (Homeownership for Lower Income Familes) insured loans. This final rule is intended to bring the maximum permissible financing charges for these programs into line with competitive market rates.

EFFECTIVE DATE: December 13, 1985.

FOR FURTHER INFORMATION CONTACT: John N. Dickie, Chief Mortgage and Capital Market Analysis Branch, Office of Financial Management, Department of Housing and Urban Development, 451 Seventh Street, SW., Washington, DC 20410. Telephone (202) 755-7270. (This is not a toll-free number.)

SUPPLEMENTARY INFORMATION: The following amendments to 24 CFR Chapter II have been made to decrease the maximum interest rate which may be charged on loans insured by this Department under section 232 (fire safety equipment) and section 235 of the National Housing Act. The maximum interest rate on the HUD/FHA section 232 (fire safety equipment) and section 235 insurance programs has been lowered from 11.00 percent to 10.50 percent.

The Secretary has determined that this change is immediately necessary to meet the needs of the market and to prevent speculation in anticipation of a change.

As a matter of policy, the Department submits most of its rulemaking to public comment, either before or after effectiveness of the action. In this instance, however, the Secretary has determined that advance notice and public comment procedures are unnecessary and that good cause exists for making this final rule effective immediately.

HUD regulations published at 47 FR 56266 (1982), amending 24 CFR Part 50, which implement section 102(2)(C) of the National Environmental Policy Act of 1969, contain categorical exclusions from their requirements for the actions, activities and programs specified in § 50.20. Since the amendments made by this rule fall within the categorical exclusions set forth in paragraph (*l*) of § 50.20, the preparation of an Environmental Impact Statement or Finding of No Significant Impact is not required for this rule.

This rule does not constitute a "major