Issued in Seattle, Washington, on May 14, 1986.

Temple H. Johnson, Jr.,

Acting Manager, Air Traffic Division, Northwest Mountain Region.

[FR Doc. 86-11598 Filed 5-23-86; 8:45 am]

BILLING CODE 4910-13-M

DEPARTMENT OF COMMERCE

International Trade Administration

15 CFR Part 385

[Docket No. 60581-6081]

Export Licensing; Controls on Exports to South Africa; Information Collection Control Numbers

AGENCY: Export Administration, Commerce.

ACTION: Final rule; notice of OMB approval.

SUMMARY: The Export Administration Regulations (EAR) contain provisions relating to the licensing of all exports to the South African military and police and prohibits the exports of all computers, computer software, or goods or technology to service computers to all apartheid enforcing entities of the South African Government. The final rule containing these provisions was published in the **Federal Register** on November 18, 1985 (50 FR 47363) with an effective date of October 11, 1985. This document adds the Office of Management and Budget (OMB) control number 0625-0156 to the information collection requirements of the rule.

DATE: The regulation became effective on October 11, 1985.

FOR FURTHER INFORMATION CONTACT: Betty Ferrell, Regulations Branch, Export Administration, Telephone (202)388-3856.

SUPPLEMENTARY INFORMATION: OMB control number 0625-0156.

As required by the Paperwork Reduction Act of 1980, 44 U.S.C. 3501 *et seq.*, Export Administration submitted this regulation to OMB for approval of the reporting requirement on prohibiting the export of computers, computer software, or goods or technology to service computers to all apartheid enforcing entities of the South African Government, even though the export could be made under general license to other parties in South Africa. A license authorizing resale within the Republic of South Africa or Namibia, may also require identification of customers to whom the ultimate consignees have sold the computers, with further identification of customers at six month intervals until all computers exported under the license have been sold. As stated in the final rule, the approval of this reporting requirement was pending with OMB. On December 30, 1985, OMB approved this collection of information assigned control number 0625-0156.

List of Subjects in 15 CFR Part 385

Communist countries, Exports, Reporting and recordkeeping requirements.

Accordingly, Part 385 of the Export Administration Regulations (15 CFR Parts 368-399) is amended as follows:

PART 385—(AMENDED)

1. The authority citation for 15 CFR Part 385 continues to read as follows:

Authority: Pub. L. 96-72, 93 Stat. 503, 50 U.S.C. app 2401 et. seq., as amended by Pub. L. 97-145 of December 29, 1981 and by Pub. L. 99-64 of July 12, 1985; E.O. 12525 of July 12, 1985 (50 FR 28757, July 16, 1985); Pub. L. 95-223, 50 U.S.C. 1701 *et seq.*; E.O. 12532 of September 9, 1985 (50 FR 36861, September 10, 1985).

2. OMB control number 0625-0156 is added to the end of § 385.4 to read as follows:

§ 385.4 Country Groups T & V.

* * * * *

(Information collection requirements in paragraph (a) approved by the Office of Management and Budget under control number 0625-0156)

Dated: May 20, 1986.

Walter J. Olson,

Deputy Assistant Secretary for Export Administration.

[FR Doc. 86-11741 Filed 5-23-86; 8:45 am]

BILLING CODE 3510-DT-M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 176

[Docket No. 85F-0288]

Indirect Food Additives; Paper and Paperboard Components; Slimicide

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 *n*-dodecylguanidine hydrochloride as a slimicide in paper and paperboard intended for use in contact with food. This action responds to a petition filed by Betz Laboratories, Inc.

DATES: Effective May 27, 1986; objections by June 26, 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: Hortense S. Macon, 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 13, 1985 (50 FR 37437), FDA announced that a petition (FAP 5B3847) had been filed by Betz Laboratories, Inc., Trevose, PA 19047, proposing that § 176.300 *Slimicides* (21 CFR 176.300) be amended to provide for the safe use of *n*-dodecylguanidine hydrochloride as a slimicide in paper and paperboard intended for use in contact with food.

FDA has evaluated data in the petition and other relevant material. The agency concludes that the proposed use of the food additive is safe, and that the regulation 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 June 26, 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 176

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 of the Center for Food Safety and Applied Nutrition, Part 176 is amended as follows:

PART 176—INDIRECT FOOD ADDITIVES: PAPER AND PAPERBOARD COMPONENTS

1. The authority citation for 21 CFR Part 176 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 § 176.300(c) by alphabetically inserting a new item in the list of substances to read as follows:

§ 176.300 Slimicides.

* * * * *

(c) * * *

List of substances	Limitations
. . . . .	
n-Dodecylguanidine hydrochloride.	At a maximum level of 0.20 pound per ton of dry weight fiber.
. . . . .	

Dated: May 15, 1986.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 86–11765 Filed 5–23–86; 8:45 am]

BILLING CODE 4160-01-M

21 CFR PART 177

[Docket No. 85F-0518]

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 ethyleneacrylic acid copolymers containing up to 25 weight-percent of polymer units derived from acrylic acid in contact with food. This action responds to a petition filed by the The Dow Chemical Co.

DATES: Effective May 27, 1986; objections by June 26, 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: Hortense S. Macon, 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 December 9, 1985 (50 FR 50233), FDA announced that a petition (FAP 5B3877) had been filed by The Dow Chemical Co., Midland, MI 48674, proposing that § 177.1310 *Ethylene acrylic acid copolymers* (21 CFR 177.1310) be amended to provide for the safe use of ethylene-acrylic copolymers containing up to 25 weight-percent of polymer units derived from acrylic acid in contact with food.

FDA has evaluated data in the petition and other relevant material. The agency concludes that the proposed use of the food additive is safe, and that the regulation 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 June 26, 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 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, 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. In § 177.1310 by revising paragraphs (a) and (b), by redesignating existing paragraph (c) as paragraph (d), and by adding new paragraph (c) to read as follows:

§ 177.1310 Ethylene-acrylic acid copolymers.

* * * * *

(a) The ethylene-acrylic acid copolymers consist of basic copolymers produced by the copolymerization of ethylene and acrylic acid such that the finished basic copolymers contain no more than:

(1) 10 weight-percent of total polymer units derived from acrylic acid when used in accordance with paragraph (b) of this section; and

(2) 25 weight-percent of total polymer units derived from acrylic acid when used in accordance with paragraph (c) of this section.

(b) The finished food-contact articles made with no more than 10 percent total polymer units derived from acrylic acid, when extracted with the solvent or solvents characterizing the type of food and under the conditions of its intended use as determined from tables 1 and 2 of § 176.170(c) of this chapter, yield net acidified chloroform-soluble extractives not to exceed 0.5 milligram per square inch of food-contact surface when tested by the methods prescribed in § 177.1330(c), except that net acidified chloroform-soluble extractives from paper and paperboard complying with § 176.170 of this chapter may be corrected for wax, petrolatum, and mineral oil as provided in § 176.170(d)(5)(iii)(*b*) of this chapter. If the finished food-contact article is itself the subject of a regulation in Parts 174, 175, 176, 177, 178 and § 179.45 of this chapter, it shall also comply with any specifications and limitations prescribed for it by that regulation.

Note.—In testing the finished food-contact article, use a separate test sample for each extracting solvent.

(c) The finished food-contact layer made with basic copolymers containing more than 10 weight-percent but no more than 25 weight-percent of total polymer units derived from acrylic acid and with a maximum thickness of 0.0025 inch (2.5 mils) may be used in contact with food types I, II, IVB, VIA, VIB, VIIB, and VIII identified in table 1 of § 176.170(c) of the chapter under conditions of use B through H as described in table 2 of § 176.170(c) of this chapter, and in contact with food types III, IVA, V, VIIA, and IX identified in table 1 of § 176.170(c) of this chapter under conditions of use E through G as described in table 2 of § 176.170(c) of this chapter.

* * * * *

Dated: May 15, 1986.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 86–11763 Filed 5–23–86; 8:45 am]

BILLING CODE 4160–01–M

21 CFR Part 178

[Docket No. 81F–0350]

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 butyric acid, 3,3-bis(3-*tert*-butyl-4-hydroxyphenyl)ethylene ester as an antioxidant for olefin polymers. This action responds to a petition filed by American Hoechst Corp.

DATES: Effective May 27, 1986. Objections by June 26, 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: 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 November 24, 1981 (46 FR 57645), FDA announced that a petition (FAP 1B3584) had been filed by American Hoechst Corp., Somerville, NJ 08876, proposing that 21 CFR 178.2010 of the food additive regulations be amended to provide for the safe use of butyric acid, 3,3-bis(3-*tert*-butyl-4-hydroxyphenyl)ethylene ester as an antioxidant and/or stabilizer for olefin polymers by removing current restrictions on food type and temperature.

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, 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. This action was considered under FDA's final rule implementing the National Environmental Policy Act (21 CFR Part 25) that was 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 environmental assessment under 21 CFR 25.31a(a).

Any person who will be adversely affected by this regulation may at any time on or before June 26, 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 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 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 of the 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 § 178.2010(b) by revising the entry for "Butyric acid, 3,3-bis(3-*tert*-butyl-4-hydroxyphenyl)ethylene ester" to read as follows:

§ 178.2010 Antioxidants and/or stabilizers for polymers.

* * * * *

(b) * * *

Substances	Limitations
Butyric acid, 3,3-bis(3-*tert*-butyl-4-hydroxyphenyl)ethylene ester (CAS Reg. No. 32509-66-3).	For use only: 1. At levels not to exceed 0.5 percent by weight of olefin copolymers complying with § 177.1520(c) of this chapter, items 3.1 and 3.2 except that when used in contact with foods described as types III, IV-A, V, VII-A, and IX in Table 1 of § 176.170(c) of this chapter, the olefin copolymers may only be used under conditions of use E, F, and G set forth in Table 2 of § 176.170(c) of this chapter. 2. At levels not to exceed 0.5 percent by weight of olefin polymers complying with § 177.1520(c) of this chapter, item 1.1, 3.1, or 3.2 (where the copolymers complying with items 3.1 and 3.2 contain not less than 85 weight-percent of polymer units derived from propylene). 3. At levels not to exceed 0.2 percent by weight of olefin polymers complying with § 177.1520(c) of this chapter, items 2.1, 2.2, 3.1, and 3.2.

* * * * *

Dated: May 15, 1986.

Richard J. Ronk,

Acting Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 86-11764 Filed 5-23-86; 8:45 am]

BILLING CODE 4160-01-M

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Office of Assistant Secretary for Housing—Federal Housing Commissioner

24 CFR Part 883

[Docket No. R-86-1292; FR-1675]

Section 8 Housing Assistance Payments Program—State Housing Agencies

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

ACTION: Final rule; technical amendment.

SUMMARY: This rule corrects an erroneous cross reference in 24 CFR Part 883—Section 8 Housing Assistance Program—State Housing Agencies.

DATE: June 27, 1986.

FOR FURTHER INFORMATION CONTACT: Grady Norris, Assistant General Counsel for Regulations, Office of General Counsel, Room 10276, Department of Housing and Urban Development, 451 Seventh Street SW., Washington, DC 20410, telephone (202) 755-7055. (This is not a toll-free number).

SUPPLEMENTARY INFORMATION: 24 CFR 883.408(b) deals with the possible effect that delays during the period an assisted project is being constructed or rehabilitated may have on section 8 contract rents. Through an error, reference is made in the paragraph to "paragraph (d) of this section". The reference should be to "§ 883.305(b)(2)".

This rule makes the necessary correction, which is technical only and is not intended to have any substantive effect, since the rule as amended on March 28, 1983 (48 FR 12709) was originally intended to make the same cross reference as this document supplies. (This conclusion is supported by the parallel provisions found in Part 880. In § 880.401(b), a cross-reference correctly identifies § 880.204(b)(2) as the provision stating the grounds for allowing an increase in contract rents where there are construction delays. Section 880.401(b) is parallel to § 883.408(b), the rule corrected here, and § 880.204(b)(2) is parallel to the corrected cross-reference provision—§ 883.305(b)(2).)

Procedural Requirements

This rule does not constitute a "major rule" as that term is defined in section 1(b) of Executive Order 12291 on Federal Regulation issued by the President on February 17, 1981. Analysis of the proposed rule indicates that it does not: (1) Have an annual effect on the economy of $100 million or more; (2) cause a major increase in costs or prices for consumers, individual industries, Federal, State or local government agencies, or geographic regions; or (3) have a significant adverse effect on competition, employment, investment, productivity, innovation, or the ability of United States-based enterprises to compete with foreign-based enterprises in domestic or export markets.

This rule was not listed in the Department's Semiannual Agenda of Regulations published on April 21, 1986 (51 FR 12036); under Executive Order 12291 and the Regulatory Flexibility Act.

The catalog of Federal Domestic Assistance program number is 14.156.

Under 5 U.S.C. 605(b) (The Regulatory Flexibility Act), the Undersigned hereby certifies that this rule would not have a significant economic impact on a substantial number of small entities.

List of Subjects in 24 CFR Part 883

Grant programs—Housing and community development, Rent subsidies, New construction and substantial rehabilitation.

Accordingly, 24 CFR Part 883 is amended to read as follows:

PART 883—SECTION 8 HOUSING ASSISTANCE PAYMENTS PROGRAM—STATE HOUSING AGENCIES

1. Paragraph (b) of § 883.408 is revised to read as follows:

§ 883.408 Construction or rehabilitation period.

* * * * *

(b) Delays. Extensions of time may be granted for the reasons specified in the Agreement. However, contract rents will be increased only for the reasons stated in § 883.305(b)(2) of this part.

* * * * *