19 CFR Section	Description		OMB con- trol No.	
*	*	*	*	*
§123.73	ticipa Land	ition to par ite in the Border er Initiative ram.		1515–0217
*	*	*	*	*

#### Raymond W. Kelly,

Commissioner of Customs.

Approved: November 4, 1998.

#### Dennis M. O'Connell,

Acting Deputy Assistant Secretary of the Treasury.

[FR Doc. 98-34675 Filed 12-31-98; 8:45 am]

BILLING CODE 4820-02-P

# **DEPARTMENT OF HEALTH AND HUMAN SERVICES**

# Food and Drug Administration

21 CFR Part 178

[Docket No. 97F-0504]

# Indirect Food Additives: Adjuvants, **Production Aids, and Sanitizers**

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 expanded safe use of the butylated reaction product of p-cresol and dicyclopentadiene for use as an antioxidant in acrylonitrile/butadiene/ styrene copolymers in contact with food. This action is in response to a petition filed by The Goodyear Tire and Rubber Co.

**DATES:** The regulation is effective January 4, 1999; submit written objections and request for a hearing by February 3, 1999.

**ADDRESSES:** Submit written objections to the Dockets Management Branch (HFA-305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852.

#### FOR FURTHER INFORMATION CONTACT:

Hortense S. Macon, Center for Food Safety and Applied Nutrition (HFS-206), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202-418-3086.

**SUPPLEMENTARY INFORMATION:** In a notice published in the Federal Register of December 10, 1997 (62 FR 65084), FDA announced that a food additive petition

(FAP 8B4561) had been filed by The Goodyear Tire and Rubber Co., c/o Keller and Heckman LLP, 1001 G St. NW., suite 500 West, Washington, DC 20001. The petition proposed to amend the food additive regulations in § 178.2010 Antioxidants and/or stabilizers (21 CFR 178.2010) to provide for the expanded safe use of butylated reaction product of p-cresol and dicyclopentadiene for use as an antioxidant in acrylonitrile/butadiene/ styrene copolymers in contact with food.

In the notice of filing for this additive, FDA announced that it had determined under § 25.32(i) (21 CFR 25.32(i)) that this action was of a type that did not individually or cumulatively have a significant effect on the human environment. Subsequently, during FDA's indepth review of the petition, the agency determined that the proposed use of the subject additive was for both single service food-packaging materials and repeat use articles. Therefore, at the agency's request, the petitioner provided an amended claim of categorical exclusion from the requirement to prepare an environmental assessment under both § 25.32(i) (single service food packaging) and (j) (repeated use articles).

FDA has evaluated data in the petition and other relevant material. Based on this information, the agency concludes that: (1) The proposed use of the additive is safe, (2) the additive will achieve its intended technical effect, and (3) the regulations in § 178.2010 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 § 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 determined under § 25.32(i) and (j) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This final rule contains no collections of information. Therefore, clearance by the Office of Management and Budget under the Paperwork Reduction Act of 1995 is not required.

Any person who will be adversely affected by this regulation may at any time on or before February 3, 1999, 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.

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 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: 21 U.S.C. 321, 342, 348, 379e.

2. Section 178.2010 is amended in the table in paragraph (b) in the entry for "Butylated reaction product of *p*-cresol and dicyclopentadiene \* \* \*" by revising the entry under the heading "Limitations" to read as follows:

# § 178.2010 Antioxidants and/or stabilizers for polymers.

(b) \* \* \*

Substances			Limitations			
by reacting ratio of 1.5 t	<ul><li>p-cresol and dicyclopenta o 1, respectively, followed</li></ul>	* nd dicylopentadiene produced diene in an approximate mole d by alkylation with isobutylene product is not less than 18 per-	177.260 2. At level diene/st contact under c of § 176 VI-C, V	aponents of nonfood art 10(c)(4)(iii) of this chapt Is not to exceed 1.0 pertyrene copolymers. The with food of Types I, II, 10 conditions of use B through 11 conditions of use B through 12 conditions of use B through 13 conditions of use B through 14 conditions of use B through 15 conditions of use B through 16 conditions of use B through 17 conditions of use B through 18 conditions of use B through 18 conditions of use B through 19 conditions of use B through 10 conditions of use B through 11 conditions of use B through 12 conditions of use B through 13 conditions of use B through 14 conditions of use B through 15 conditions of use B through 16 conditions of use B through 17 conditions of use B through 18 conditions of use B throug	* icles complying with §§ 1 er. reent by weight of acrylo finished copolymers ma IV-B, VI-A, VI-B, VII-E ugh H, as described in ta and with food of Types ditions of use C through 76. 170(c) of this chapte	nitrile/buta- by be used in 3, and VIII ables 1 and 2 III, IV-A, V, G as de-

Dated: December 21, 1998.

#### L. Robert Lake,

Director, Office of Policy, Planning and Strategic Initiatives, Center for Food Safety and Applied Nutrition.

[FR Doc. 98-34734 Filed 12-31-98; 8:45 am]

BILLING CODE 4160-01-F

#### **DEPARTMENT OF STATE**

#### 22 CFR Part 41

[Public Notice 2939]

Bureau of Consular Affairs; VISAS: Passports and Visas Not Required for Certain Nonimmigrants—VWPP

**AGENCY:** Bureau of Consular Affairs, DOS.

**ACTION:** Final rule.

SUMMARY: This final rule adopts the interim rules which added Andorra, Argentina, Australia, Belgium, Brunei, Denmark, Finland, Iceland, Ireland, Liechtenstein, Luxembourg, Monaco, New Zealand, Norway, San Marino, Slovenia and Spain as participating countries in the Visa Waiver Pilot Program (VWPP), eliminated probationary entry status.

**DATES:** The rule takes effect January 4, 1999

FOR FURTHER INFORMATION CONTACT: H. Edward Odom, Chief, Legislation and Regulations Division, Visa Office, Room L603–C, SA–1, Department of State, Washington, D.C. 20520–0106, (202) 663–1203.

SUPPLEMENTARY INFORMATION: This final rule amends Part 41, Title 22 of the Code of Federal Regulations concerning visas for nonimmigrants pursuant to section 217 of the Immigration and Nationality Act. Over the past several years the Department published the following interim rules amending 22 CFR 42.1(1):

(1) 56 FR 46716, September 13, 1991, which removed the eight-country cap and added Andorra, Austria, Belgium, Denmark, Finland, Iceland, Liechtenstein, Luxembourg, Monaco, New Zealand, Norway, San Marino and Spain to the list of participating countries and extended the VWPP Program through September 30, 1994;

(2) 58 FR 40585, July 29, 1993, which added Brunei as a participating country;

- (3) 60 FR 15872, March 28, 1995, which extended the program through September 30, 1996, created a probationary status for the VWPP and added Ireland as a country with probationary status;
- (4) 61 FR 35628, July 8, 1996, which added Argentina;
- (5) 61 FR 39318, July 29, 1996, which added Australia; and
- (6) 62 FR 51030, September 30, 1997, which eliminated probationary entry status for countries, designated Ireland as a permanent participating country and added Slovenia to the VWPP.

Pub. L. 105–173, enacted on April 27, 1998, extends the Program through April 30, 2000.

# **Comments**

Each of the six interim rules invited interested persons to submit written comments concerning these amendments. No comments were received. This rule makes final the above-listed interim rules.

#### **Final Rule**

This final rule implements the regulation as published on September 30, 1997 [62 FR 51030]. This regulation is being promulgated in conjunction with the Immigration and Naturalization Service (INS) because action by the Attorney General in consultation with the Secretary of State is required under section 217 of the INA, as amended. (See INS Rule also published in the **Federal Register** on December 30, 1998.)

As no comments were received, the interim rule published on September 30, 1997 [62 FR 51030] is incorporated herein as a final rule.

# PART 41—VISAS: DOCUMENTATION OF NONIMMIGRANTS UNDER THE IMMIGRATION AND NATIONALITY ACT, AS AMENDED

Accordingly, the interim rule amending 22 CFR part 41 which was published at 62 FR 51030 is adopted as a final rule without change.

Dated: November 16, 1998.

# Nancy H. Sambaiew,

Acting Assistant Secretary for Consular Affairs.

[FR Doc. 98-34783 Filed 12-31-98; 8:45 am] BILLING CODE 4710-06-P

#### **DEPARTMENT OF TRANSPORTATION**

### **Coast Guard**

33 CFR Part 117 [CGD 08-98-078]

RIN 2115-AE47

# Drawbridge Operating Regulation; Bayou Lacombe. LA

**AGENCY:** Coast Guard, DOT.

**ACTION:** Notice of temporary deviation

from regulations.

SUMMARY: The Commander, Eighth Coast Guard District has issued a temporary deviation from the regulation governing the operation of the Tammany Trace swing span drawbridge across Bayou Lacombe, mile 5.2, at Lacombe, St. Tammany Parish, Louisiana. This deviation allows the St. Tammany Parish Police Jury to close the bridge to navigation continuously from 6 a.m. on January 4, 1999 until 6 a.m. on January 16, 1999 and from 6 a.m. on January 18 until 12 p.m. on January 22,