revoked, in whole or in part, by the port director, the procedures described in paragraph (b)(5) of this section shall apply.

(7) Application for and approval of a reporting program shall not limit or restrict the use of other alternative 16 means for obtaining the release of holders, containers and shipping devices.

* * * * *

PART 113—CUSTOMS BONDS

1. The general authority citation for part 113 continues to read as follows:

Authority: 19 U.S.C. 66, 1623, 1624.

* * * * *

2. Section 113.66 is amended by redesignating paragraph (c) as (d) and by adding a new paragraph (c) to read as follows:

§ 113.66 Control of containers and instruments of international traffic bond conditions.

* * * * *

(c) *Agreement to comply with application approved under 19 CFR 10.41b(b).* If the principal establishes a program for the cross-border movements of shipping devices based upon an application approved as provided in § 10.41b(b) of this chapter (19 CFR 10.41b(b)), the principal agrees:

(1) To timely file complete and accurate reports on the shipping devices, and to pay any applicable duty due on the devices and repairs made to such devices, as provided in the approved application;

(2) To retain complete and accurate records regarding the shipping devices, and to make such records available to Customs for inspection and audit upon reasonable notice, as also required in the approved application; and

(3) To otherwise comply with every other condition of the approved application.

Approved: January 31, 1996.

George J. Weise,

Commissioner of Customs.

Dennis M. O'Connell,

Acting Deputy Assistant Secretary of the Treasury.

[FR Doc. 96–4797 Filed 2–29–96; 8:45 am]

BILLING CODE 4820–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 73

[Docket No. 95C–0091]

Listing of Color Additives Exempt From Certification; Fruit Juice Color Additive and Vegetable Juice Color Additive; Confirmation of Effective Date

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; confirmation of effective date.

SUMMARY: The Food and Drug Administration (FDA) is confirming the effective date of November 13, 1995, of the final rule published in the Federal Register of October 10, 1995 (60 FR 52628), that amended the color additive regulations to provide for the safe use in food of dried fruit juice color additive, dried vegetable juice color additive, and vegetable juice color additive prepared by water infusion of the dried vegetable.

DATES: Effective date confirmed: November 13, 1995.

FOR FURTHER INFORMATION CONTACT: Aydin Örstan, Center for Food Safety and Applied Nutrition (HFS–217), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3076.

SUPPLEMENTARY INFORMATION: In the Federal Register of October 10, 1995 (60 FR 52628), FDA amended the color additive regulations in § 73.250 *Fruit juice* (21 CFR 73.250) to provide for the safe use of dried fruit juice color additive and in § 73.260 *Vegetable juice* (21 CFR 73.260) to provide for the safe use of dried vegetable juice color additive and vegetable juice color additive prepared by water infusion of the dried vegetable.

FDA gave interested persons until November 9, 1995, to file objections or requests for a hearing. The agency received no objections or requests for a hearing on the final rule. Therefore, FDA finds that the final rule published in the Federal Register of October 10, 1995, should be confirmed.

List of Subjects in 21 CFR Part 73

Color additives, Cosmetics, Drugs, Medical devices.

Therefore, under the Federal Food, Drug, and Cosmetic Act (secs. 201, 401, 402, 403, 409, 501, 502, 505, 601, 602, 701, 721 (21 U.S.C. 321, 341, 342, 343, 348, 351, 352, 355, 361, 362, 371, 379e)) and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Director, Center for Food Safety and Applied Nutrition, notice is given that no objections or requests for a hearing were filed in response to the October 10, 1995, final rule. Accordingly, the amendments promulgated thereby became effective November 13, 1995.

Dated: February 12, 1996.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 96–4717 Filed 2–29–96; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 180

[Docket No. 94F–0152]

Food Additives Permitted in Food on an Interim Basis or in Contact With Food Pending Additional Study; Mannitol

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to permit the manufacture of mannitol by fermentation of sugars or sugar alcohols such as glucose, sucrose, fructose, or sorbitol by the action of the yeast *Zygosaccharomyces rouxii.* This action is in response to a petition filed by Roquette America, Inc.

DATES: Effective March 1, 1996; written objections and requests for a hearing by April 1, 1996.

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

FOR FURTHER INFORMATION CONTACT: Rosalie M. Angeles, Center for Food Safety and Applied Nutrition (HFS–207), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3107.

SUPPLEMENTARY INFORMATION: In a notice published in the Federal Register of December 13, 1994 (59 FR 64207), FDA announced that a food additive petition (FAP 4A4412) had been filed by Roquette America, Inc., c/o Keller and Heckman, 1001 G St. NW., Washington, DC 20001. The petition proposed to amend the food additive regulations in § 180.25 *Mannitol* (21 CFR 180.25) to permit the manufacture of mannitol by fermentation of sugars or sugar alcohols such as glucose, sucrose, fructose, or sorbitol by the action of the yeast *Z. rouxii.*