Dated: June 9, 2000. **Kenneth S. Apfel**,

Commissioner of Social Security.

PART 404-FEDERAL OLD-AGE, SURVIVORS AND DISABILITY INSURANCE (1950—)

Accordingly, the interim final rules amending 20 CFR Part 404 published at 64 FR 57774 on October 27, 1999, are adopted as final without change.

[FR Doc. 00–15644 Filed 6–20–00; 8:45 am] BILLING CODE 4191–02–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 178

[Docket No. 99F-1421]

Indirect Food Additives: Adjuvants, Production Aids, and Sanitizers; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical

amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending its food additive regulations that provide for the safe use of tetradecanoic acid, lithium salt as a stabilizer for polypropylene and certain polypropylene copolymers intended for use in contact with food. When the regulation was last amended, the regulation published with some errors. This document corrects those errors.

DATES: This rule is effective June 21, 2000.

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–215), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3081.

SUPPLEMENTARY INFORMATION: FDA has discovered that two errors have become incorporated into the agency's current food additive regulations. In an amendment to 21 CFR 178.2010, published in the Federal Register of December 27, 1999 (64 FR 72273), there were errors regarding the food type VI–B. This document corrects those errors. Publication of this document constitutes final action under the Administrative Procedure Act (5 U.S.C. 553). FDA has determined that notice and public

comment are unnecessary because this amendment is nonsubstantive.

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) under the heading "Limitations" by revising the entry for "Tetradecanoic acid, lithium salt" to read as follows:

§ 178.2010 Antioxidants and/or stabilizers for polymers.

* * * * * (b) * * *

Substances Limitations

Tetradecanoic acid, lithium salt (CAS Reg. No. 20336-96-3)

For use only at levels not to exceed 0.15 percent by weight of polypropylene and polypropylene copolymers complying with § 177.1520(c) of this chapter, items 1.1a, 1.1b, 3.1a, 3.1b, 3.1c, 3.2a, and 3.2b. The finished polymers may only be used in contact with food of Types I, II, IV–B, VI–B, VII–B, and VIII as described in table 1 of § 176.170(c) of this chapter under conditions of use B through H as described in table 2 of § 176.170(c) of this chapter, and with food of Types III, IV–A, V, VI–A, VI–C, VII–A, and IX described in table 1 of § 176.170(c) of this chapter under conditions of use C through G as described in table 2 of § 176.170(c) of this chapter.

Dated: June 7, 2000.

L. Robert Lake,

Director of Regulations and Policy, Center for Food Safety and Applied Nutrition. [FR Doc. 00–15561 Filed 6–20–00; 8:45 am]

BILLING CODE 4160-01-F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 349

[Docket No. 98N-0002]

RIN 0910-AA01

Ophthalmic Drug Products for Overthe-Counter Human Use; Amendment of Final Monograph

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the monograph for over-the-counter (OTC) ophthalmic drug products (the regulation that establishes conditions under which these drug products are generally recognized as safe and effective and not misbranded). The amendment adds a new warning and revises an existing warning for ophthalmic vasoconstrictor drug products. These products contain the ingredients ephedrine hydrochloride, naphazoline hydrochloride, or