June 9, 1999, at 64 FR 30898, the Commission rescinded the Guides for the Watch Industry, 16 CFR part 245. The Commission's Guides for the Jewelry, Precious Metals, and Pewter Industries (''Jewelry Guides''), 16 CFR part 23, refer to the Watch Guides in footnote 1 in § 23.0. Because the Watch Guides have been rescinded, the Commission is amending the Jewelry Guides to remove the reference to the Watch Guides in footnote 1 in § 23.0.

List of Subjects in 16 CFR Part 23

Advertising, Jewelry, Labeling, Trade practices, Watch bands.

The Commission, under the authority of section 18 of the Federal Trade Commission Act, 15 U.S.C. 57a, amends 16 CFR part 23 as follows:

1. The authority citation for part 23 continues to read as follows:

Authority: Sec. 6, 5, 38 Stat. 721, 719; 15 U.S.C. 46, 45.

§ 23.0 [Amended]

2. Section 23.0 is amended by removing and reserving footnote 1.

By direction of the Commission.

Donald S. Clark,

Secretary.

[FR Doc. 99–15840 Filed 6–21–99; 8:45 am]

BILLING CODE 6750–01–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 5

Delegation of Authority and Organization; Center for Food Safety and Applied Nutrition

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the general redelegation of authority from the Commissioner of Food and Drugs to other officers of FDA. The amendment delegates to the Director and Deputy Director, Center for Food Safety and Applied Nutrition (CFSAN); the Director, Office of Regulations and Policy, CFSAN; and the Director, Office of Premarket Approval, CFSAN authority to implement the Federal Food, Drug, and Cosmetic Act (the act), as amended hereafter. This redelegation is necessary to improve the efficiency of program operations.

EFFECTIVE DATE: June 22, 1999.

FOR FURTHER INFORMATION CONTACT: Louis B. Brock, Regulation Coordination Staff (HFS–24), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–205–4273, or Loretta W. Davis, Division of Management Systems and Policy (HFA–340), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–827–4809.

SUPPLEMENTARY INFORMATION: Section 309 of the Food and Drug Administration Modernization Act of 1997 (Pub. L. 105–115) amended section 409 of the act (21 U.S.C. 348). New section 409(h) of the act requires manufacturers or suppliers of food-contact substances to notify the Secretary of Health and Human Services (and by delegation, the Commissioner of Food and Drugs), at least 120 days prior to the introduction or delivery for introduction into interstate commerce, of the identification and use of food-contact substances, and to provide information showing that the substance is safe according to the standards of section 409(c)(3)(A) of the act.

FDA is amending the general redelegation of authority from the Commissioner of Food and Drugs to the Director and Deputy Director, Center for Food Safety and Applied Nutrition (CFSAN); the Director, Office of Regulations and Policy, CFSAN; and the Director, Office of Premarket Approval, CFSAN authority to implement the act, as amended hereafter. This redelegation is necessary to improve the efficiency of program operations. Further redelegation of the authorities is not authorized at this time. Authority delegated to a position may be exercised by a person officially designated to serve in such position in an acting capacity or on a temporary basis.

List of Subjects in 21 CFR Part 5

Authority delegations (Government agencies), Imports, Organization and functions (Government agencies).

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR part 5 is amended as follows:

PART 5—DELEGATIONS OF AUTHORITY AND ORGANIZATION

1. The authority citation for 21 CFR part 5 continues to read as follows:

Authority: 5 U.S.C. 504, 552, App. 2; 7 U.S.C. 138a, 2271; 15 U.S.C. 638, 1261–1282, 3701–3711a; 15 U.S.C. 1451–1461; 21 U.S.C. 41–50, 61–63, 141–149, 321–394, 467f, 679(b), 801–886, 1031–1309; 35 U.S.C. 156; 42 U.S.C. 241, 242, 242a, 2421, 242n, 243, 262, 263, 264, 265, 300u–300u–5, 300aa–1; 1395y, 3246b, 4332, 4831(a), 10007–10008; E.O. 11921, 41 FR 24294, 3 CFR, 1977 Comp., p. 124–131; E.O. 12591, 52 FR 13414, 3 CFR, 1988 Comp., p. 220–223.

2. Section 5.61 is amended by adding paragraph (i) to read as follows:

§ 5.61 Food standards, food additives, generally recognized as safe (GRAS) substances, color additives, nutrient content claims, and health claims.

* * * * *

(i) The following officials are authorized to perform all the functions of the Commissioner of Food and Drugs under section 409(h) of the act, excluding the duties set out in section 409(h)(5) of the act, regarding premarket notification of food-contact substances:

(1) The Director and Deputy Director, Center for Food Safety and Applied Nutrition (CFSAN).

(2) The Director, Office of Regulations and Policy, CFSAN.

(3) The Director, Office of Premarket Approval, CFSAN.

Dated: June 11, 1999.

Margaret M. Dotzel,

Acting Associate Commissioner for Policy Coordination.

[FR Doc. 99–15753 Filed 6–21–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF THE TREASURY

Internal Revenue Service

26 CFR Parts 1, 20, and 25

[TD 8819]

RIN 1545–AX14

Use of Actuarial Tables in Valuing Annuities, Interests for Life or Terms of Years, and Remainder or Reversionary Interests; Correction

AGENCY: Internal Revenue Service (IRS), Treasury.

ACTION: Correction to final regulations.

SUMMARY: This document contains corrections to final regulations that were published in the **Federal Register** on Friday, April 30, 1999 (64 FR 23187) relating to the use of actuarial tables in valuing annuities, interests for life or terms of years, and remainder or reversionary interests.

DATES: This correction is effective May 1, 1999.

FOR FURTHER INFORMATION CONTACT: William L. Blodgett (202) 622–3090 (not a toll-free number).

SUPPLEMENTARY INFORMATION:

Background

The final regulations that are the subject of these corrections are under