

**PART 680—ADDITIONAL STANDARDS FOR MISCELLANEOUS PRODUCTS**

87. The authority citation for 21 CFR part 680 continues to read as follows:

Authority: Secs. 201, 501, 502, 503, 505, 510, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 351, 352, 353, 355, 360, 371); secs. 215, 351, 352, 353, 361 of the Public Health Service Act (42 U.S.C. 216, 262, 263, 263a, 264).

88. The heading for Subpart A—Allergenic Products is removed.

**Subpart B [Removed]**

89. Subpart B, consisting of §§ 680.10 through 680.16, is removed.

**Subpart C [Removed]**

90. Subpart C, consisting of §§ 680.20 through 680.26, is removed.

**PART 700—GENERAL**

91. The authority citation for 21 CFR part 700 continues to read as follows:

Authority: Secs. 201, 301, 502, 505, 601, 602, 701, 704 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 331, 352, 355, 361, 362, 371, 374).

**§ 700.10 [Removed]**

92. Section 700.10 *Shampoo preparations containing eggs as one of the ingredients* is removed.

**PART 801—LABELING**

93. The authority citation for 21 CFR part 801 continues to read as follows:

Authority: Secs. 201, 301, 501, 502, 507, 519, 520, 701, 704 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 331, 351, 352, 357, 360i, 360j, 371, 374).

**§ 801.403 [Removed]**

94. Section 801.403 *Specific medical devices; recommended warning and caution statements* is removed from subpart H.

**§ 801.408 [Removed]**

95. Section 801.408 *Pessaries for intracervical and intrauterine use* is removed from subpart H.

**§ 801.427 [Removed]**

96. Section 801.427 *Professional and patient labeling for intrauterine contraceptive devices* is removed from subpart H.

Dated: October 6, 1995.

William B. Schultz,

*Deputy Commissioner for Policy.*

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