

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**21 CFR Parts 211 and 720**

[Docket No. 00N-1217]

**Code of Federal Regulations; Technical Amendments**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Final rule; technical amendments.

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**SUMMARY:** The Food and Drug Administration (FDA) is amending its regulations to reflect a correct footnote and a part heading. This action is being taken to improve the accuracy of the regulations.

**EFFECTIVE DATE:** *[Insert date of publication in the Federal Register.]*

**FOR FURTHER INFORMATION CONTACT:** LaJuana D. Caldwell, Office of Policy, Planning, and Legislation (HF-27), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857.

**SUPPLEMENTARY INFORMATION:** FDA has discovered that errors have been incorporated into the agency's codified regulations for 21 CFR parts 211 and 720. 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**

*21 CFR Part 211*

Drugs, Labeling, Laboratories, Packaging and containers, Prescription drugs, Reporting and recordkeeping requirements, Warehouses.

DMB

Display Date	4-7-00
Publication Date	4-10-00
Certifier	SNK/PSB

*21 CFR Part 720*

Confidential business information, Cosmetics.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR parts 211 and 720 are amended as follows:

**PART 211—CURRENT GOOD MANUFACTURING PRACTICE FOR FINISHED PHARMACEUTICALS**

1. The authority citation for 21 CFR part 211 continues to read as follows:

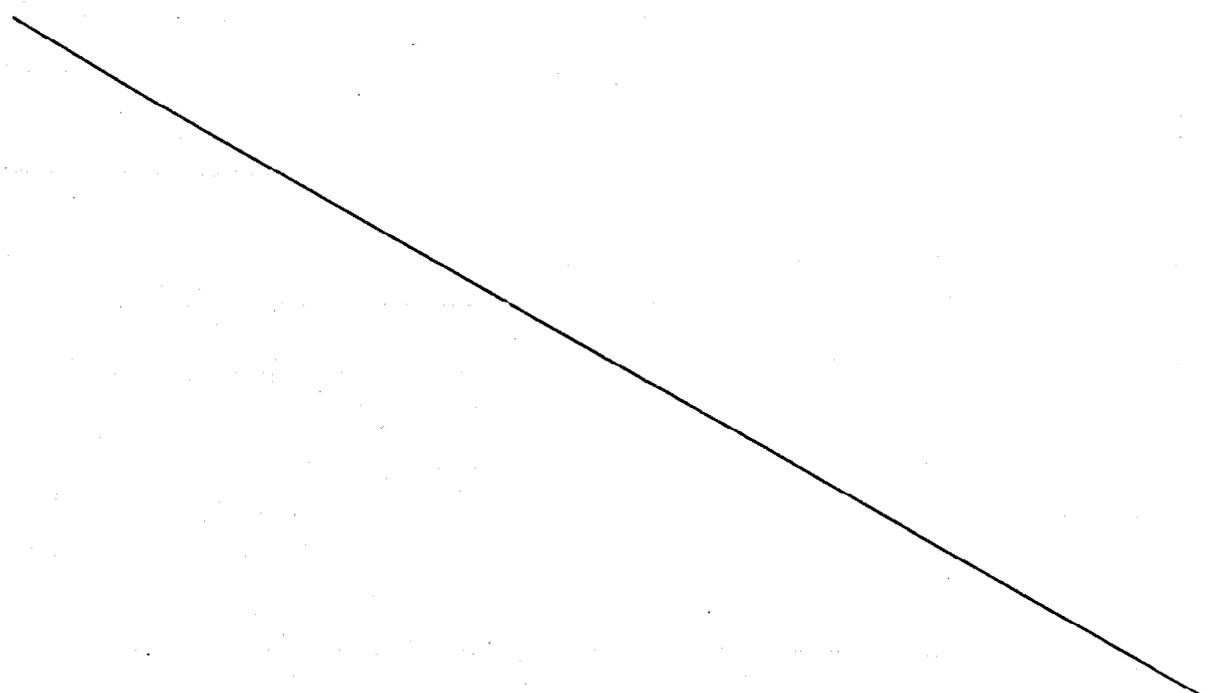
**Authority:** 21 U.S.C. 321, 351, 352, 355, 360b, 371, 374.

**§ 211.194 [Amended]**

2. Section 211.194 *Laboratory records* is amended by removing in paragraph (a)(2) and its footnote the number “2” and by adding in their place the number “1”.

**PART 720—VOLUNTARY FILING OF COSMETIC PRODUCT INGREDIENT COMPOSITION STATEMENTS**

3. The authority citation for 21 CFR part 720 continues to read as follows:



**Authority:** 21 U.S.C. 321, 331, 361, 362, 371, 374.

4. The heading for part 720 is revised to read as set forth above.

Dated: 3/31/00  
March 31, 2000



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Margaret M. Dotzel  
Acting Associate Commissioner for Policy

[FR Doc. 00-???? Filed ??-??-00; 8:45 am]

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