an insured depository institution) that is a savings and loan holding company. A company has control over a saving association if it: directly or indirectly, or acting through one or more other persons owns, controls, or has the power to vote 25 percent or more of any class of voting securities; or would be deemed to control the company under § 574.4(a) of this chapter or presumed to control the company under § 574.4(b) of this chapter, and in the latter case, control has not been rebutted. Notwithstanding any other provision of this section, no company shall be deemed to own or control another by virtue of its ownership or control of shares in a fiduciary capacity. When used to refer to a subsidiary of a savings association, the term *subsidiary* means a ''subsidiary'' that is controlled by the savings association within the meaning of 12 CFR part 574 of this chapter.

(e) References to the Reserve Bank or the Comptroller shall be deemed to include the Director of OTS; and

* * * * *

Dated: September 29, 2003.

By the Office of Thrift Supervision.

James E. Gilleran,

Director.

[FR Doc. 03–25217 Filed 10–6–03; 8:45 am]

BILLING CODE 6720–01–P

CONSUMER PRODUCT SAFETY COMMISSION

16 CFR Part 1000

Statement of Organization and Functions

AGENCY: Consumer Product Safety Commission.

ACTION: Final rule.

SUMMARY: The Consumer Product Safety Commission is amending its statement of organization and functions to reflect the transfer of the National Injury Information Clearinghouse from the Directorate for Epidemiology to the Office of the Secretary.

EFFECTIVE DATE: October 7, 2003.

FOR FURTHER INFORMATION CONTACT: Stephen Lemberg, Office of the General Counsel, Consumer Product Safety Commission, Washington, DC 20207, telephone (301) 504–7630, email *slemberg@cpsc.gov.*

SUPPLEMENTARY INFORMATION: The reference to the Clearinghouse in section 1000.27, Directorate for Epidemiology, is being moved to section 1000.16, Office of the Secretary.

Since this rule relates solely to internal agency management, pursuant to 5 U.S.C. 553(b) notice and other public procedures are not required and it is effective immediately upon publication in the **Federal Register**. Further, this action is not a rule as defined in the Regulatory Flexibility Act, 5 U.S.C. 601–612, and, thus, is exempt from the provisions of the Act.

List of Subjects in 16 CFR Part 1000

Organization and functions (government agencies).

■ Accordingly, part 1000 is amended as follows:

PART 1000—[AMENDED]

■ 1. The authority citation for part 1000 continues to read as follows:

Authority: 5 U.S.C. 552(a).

§ 1000.27 [Amended]

■ 2. In § 1000.27, remove the last sentence.

§ 1000.16 [Amended]

■ 3. In § 1000.16, add at the end the sentence ''It administers the National Injury Information Clearinghouse.''

Dated: September 30, 2003.

Todd A. Stevenson,

Secretary, Consumer Product Safety Commission.

[FR Doc. 03–25297 Filed 10–6–03; 8:45 am]

BILLING CODE 6335–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 172

[Docket No. 1987F–0179]

Food Additives Permitted for Direct Addition to Food for Human Consumption; Olestra; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA) is correcting a final rule that appeared in the **Federal Register** of August 5, 2003 (68 FR 46403). The document denied the requests for a hearing and response to objections it has received on the final rule that amended the food additive regulations to provide for the safe use of sucrose esterified with medium and long chain fatty acids (olestra) as a replacement for fats and oils in savory snacks. The document was published with inadvertent errors. This document corrects those errors.

FOR FURTHER INFORMATION CONTACT: Mary Ditto, Center for Food Safety and Applied Nutrition (HFS–255), Food and Drug Administration, 5100 Paint Branch Pkwy., College Park, MD 20740, 202–418–3102.

SUPPLEMENTARY INFORMATION: In FR Doc. 03–19509, appearing on page 46403 in the **Federal Register** of Tuesday, August 5, 2003, the following corrections are made:

1. On page 46408, in the second column, under the heading ''*D. Adequacy of Olestra's Label Statement* [33]'' the first sentence is corrected to read ''In its fifth objection and request for a hearing, CSPI challenges the label statement required by the 1996 final rule, claiming that it is not sufficient to protect the public from adverse effects associated with consumption of olestra.''

2. On page 46408, in the third column, under the heading ''*E. Alleged Procedural Problems in the Olestra Proceeding*'' the first sentence is corrected to read ''In its sixth objection and hearing request, CSPI claims that there were a number of problems with the procedures utilized by FDA to reach a decision about the safety of olestra.''

3. On page 46408, in the third column, under heading ''*E. Alleged Procedural Problems in the Olestra Proceeding*'' the second to the last sentence on that page is corrected to read ''As in the case with its fifth objection and hearing request, CSPI specifically identifies no factual issue underlying any of its six procedural complaints.''

Dated: September 30, 2003.

Jeffrey Shuren,

Assistant Commissioner for Policy.

[FR Doc. 03–25198 Filed 10–6–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF JUSTICE

Drug Enforcement Administration

21 CFR Parts 1300, 1309, and 1310

[Docket No. DEA–210F]

RIN 1117–AA69

Implementation of the Methamphetamine Anti-Proliferation Act; Thresholds for Retailers and for Distributors Required To Submit Mail Order Reports; Changes to Mail Order Reporting Requirements

AGENCY: Drug Enforcement Administration (DEA), Justice.

ACTION: Final rule.