§ 368.1 Introduction.

This part implements Public Law 104–52, the “Prohibition of Cigarette Sales to Minors in Federal Buildings and Lands Act,” which prohibits the sale of tobacco products through vending machines and the distribution of free samples of tobacco products on Federal property.

§ 368.2 Definitions.

As used in this part—

Building and real property occupied and maintained by the Board.

Minor means an individual under the age of 18 years.

Tobacco product means cigarettes, cigars, little cigars, pipe tobacco, smokeless tobacco, snuff, and chewing tobacco.

§ 368.3 Vending machines.

The sale of tobacco products in vending machines is prohibited in or around Federal property occupied and maintained by the Railroad Retirement Board.

§ 368.4 Concession stands.

Tobacco products may be sold on property occupied and maintained by the Railroad Retirement Board only as authorized by the Railroad Retirement Board or the General Services Administration or other Federal agency. Concession stands may not sell tobacco products to minors.

§ 368.5 Free tobacco samples.

The distribution of free samples of tobacco products is prohibited in or around Federal property occupied and maintained by the Railroad Retirement Board.

Dated: February 21, 1996.

By Authority of the Board.

Beatrice Ezerski,
Secretary to the Board.

[FR Doc. 96–4676 Filed 3–1–96; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 5

Delegations of Authority and Organization; Issuance of Notices Relating to Debarment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the regulations for delegations of authority to FDA officials in the Center for Drug Evaluation and Research (CDER), the Center for Veterinary Medicine (CVM), and the Center for Biologics Evaluation and Research (CBER) by adding a new delegations section concerning the issuance of notices relating to proposals and orders for debarment and denial of an application to terminate debarment. Additionally, FDA is amending the regulations regarding petitions so that certain officials of CDER, CVM, and CBER are authorized to respond to petitions concerning debarment and refusal to terminate debarment. This action will make the process of issuing such notices and responses to petitions more efficient.

EFFECTIVE DATE: March 4, 1996.

FOR FURTHER INFORMATION CONTACT: Ellen Rawlings, Division of Management Systems and Policy (HFA–340), Food and Drug Administration, 5600 Fishers Lane, Rockville, MD 20857, 301–443–4976.

SUPPLEMENTARY INFORMATION: New section 306 of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 335a), created by the Generic Drug Enforcement Act of 1992, authorizes the Secretary of Health and Human Services and, by previous delegation, the Commissioner of Food and Drugs (the Commissioner) to take actions relating to debarment proposals and orders as well as proposals and orders to deny an application to terminate a debarment order. Certain aspects of this authority are being redelegated in new § 5.98 from the Commissioner to the Directors of CDER, CVM, and CBER, to the Deputy Directors of CDER and CVM, and the Associate Director for Policy Coordination and Public Relations, CBER, as appropriate. In addition, FDA is amending § 5.31 (21 CFR 5.31) by delegating authority to the Directors of CDER, CVM, and CBER, to the Deputy Directors of CDER and CVM, and the Associate Director for Policy Coordination and Public Relations of CBER to respond to petitions concerning actions they are authorized to take under new § 5.98. The redelegations will make the process of issuing such notices and responses to petitions more efficient.

Further redelegation of the authority delegated is not authorized. Authority delegated to a position by title 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:

Section 5.31 is amended by adding new paragraphs (f)(1)(vi), (f)(2)(x), and (f)(8) to read as follows:

§ 5.31 Petitions under part 10.

(f) * * * *  * * * *  * * * *  * * * *  * * * *  * * * *  * * * *

(1) * * * *  * * * *  * * * *  * * * *  * * * *  * * * *  * * * *

(vi) Section 5.98 Issuance of notices relating to proposals and orders for debarment and denial of an application to terminate debarment.

(2) * * * *  * * * *  * * * *  * * * *  * * * *  * * * *  * * * *

(x) Section 5.98 Issuance of notices relating to proposals and orders for debarment and denial of an application to terminate debarment.

§ 5.98 Issuance of notices relating to proposals and orders for debarment and denial of an application to terminate debarment.

The Director and Deputy Director, Center for Drug Evaluation and Research (CDER), the Director and Deputy Director, Center for Veterinary Medicine (CVM), and the Director and Associate Director for Policy Coordination and Public Relations, Center for Biologics Evaluation and Research (CBER) are authorized to issue the following notices under section 306 of the Federal Food, Drug, and Cosmetic Act (the act) which relate to the assigned functions of their organizations:

(a) Notices of opportunity for hearing on proposals for mandatory or permissive debarment.

(b) Notices ordering debarment when opportunity for a hearing has been waived.

(c) Notices ordering debarment where the person notifies the agency that the person acquires debarment under section 306(c)(2)(B) of the act.

(d) Notices of opportunity for hearing on proposals denying an application to terminate debarment under section 306(d)(3) of the act.

(e) Orders denying an application to terminate debarment under section 306(d)(3) of the act when opportunity for a hearing has been waived.

Dated: February 26, 1996.

William K. Hubbard, Associate Commissioner for Policy Coordination.

EFFECTIVE DATE: This policy statement is effective March 4, 1996.

FOR FURTHER INFORMATION CONTACT: Stanley S. Colvin, Assistant General Counsel, United States Information Agency.

The Agency specifically reserves the right to limit the number of au pair participants to not more than 22,720. The Agency does not believe that currently designated sponsors, and those organizations receiving new designations, will be affected by this numerical limitation. This belief is based upon the past history of au pair activities and the Agency's knowledge of the growth rates of similar programs overseen by the Agency. The Agency specifically reserves the right to limit the number of participants sponsored by an individual organization. Participant levels for newly designated au pair sponsors will be determined by the Agency in consultation with the sponsor. The organization's prior experience, organizational capacity, and resources will be specifically considered in determining participant levels.

List of Subjects in 22 CFR Part 514

Cultural exchange programs.