[FR Doc. 03–3268 Filed 2–10–03; 8:45 am]
BILLING CODE 4910–13–M

## RAILROAD RETIREMENT BOARD

### 20 CFR Parts 260 and 320

**RIN 3220–AB03**

### Requests for Reconsideration and Appeals Within the Board; Correction

**AGENCY:** Railroad Retirement Board.

**ACTION:** Final rule; correction.

**SUMMARY:** The Railroad Retirement Board (Board) published in the **Federal Register** of December 17, 2002, a document that simplified the procedures that govern requests for reconsideration and appeals within the Board. Sections 260.9(b) and 320.39(a) inadvertently contained inaccurate terminology. This document corrects that terminology.

**DATES:** Effective on February 11, 2003.

**FOR FURTHER INFORMATION CONTACT:** Marguerite P. Dadabo, Assistant General Counsel, Railroad Retirement Board, (312) 751–4945, TDD (312) 751–4701.

**SUPPLEMENTARY INFORMATION:** The Railroad Retirement Board published a document in the **Federal Register** of December 17, 2002 (67 FR 77152). That document simplified the procedures that govern requests for reconsideration and appeals within the Board. We discovered that inaccurate terminology was contained in § 260.9(b) and § 320.39(a). This document corrects that terminology.

In rule FR Doc. 02–31640 published on December 17, 2002 (67 FR 77152), make the following corrections. On page 77156 in § 260.9(b), in the first column (line 4 thereof), and on page 77157 in § 320.39(a), in the third column (line 21 thereof), remove the word ''reconsideration'' and insert in their place the words ''hearings officer's''.

By Authority of the Board.

Dated: February 4, 2003.

For the Board.

**Beatrice Ezerski,**

*Secretary to the Board.*

[FR Doc. 03–3308 Filed 2–10–03; 8:45 am]
BILLING CODE 7905–01–P

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

### 21 CFR Part 558

### New Animal Drugs for Use in Animal Feeds; Salinomycin

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

**ACTION:** Final rule.

**SUMMARY:** The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental abbreviated new animal drug application (ANADA) filed by Intervet, Inc. The supplemental ANADA provides for use of a salinomycin Type A medicated article to make Type C medicated feeds used for the prevention of coccidiosis in roaster and replacement (breeder and layer) chickens and for the prevention of coccidiosis in quail.

**DATES:** This rule is effective February 11, 2003.

**FOR FURTHER INFORMATION CONTACT:** Lonnie W. Luther, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–8549, e-mail: lluther@cvm.fda.gov.

**SUPPLEMENTARY INFORMATION:** Intervet, Inc., P.O. Box 318, 405 State St., Millsboro, DE 19966, filed a supplement to ANADA 200–075 that provides for use of SACOX (salinomycin) Type A medicated article to make Type C medicated feeds used for the prevention of coccidiosis in roaster and replacement (breeder and layer) chickens and for the prevention of coccidiosis in quail. The supplemental ANADA is approved as of November 8, 2002, and the regulations are amended in 21 CFR 558.550 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

FDA has determined under 21 CFR 25.33(a)(1) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

**List of Subjects in 21 CFR Part 558**

Animal drugs, Animal feeds.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 558 is amended as follows:

## PART 558--NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

1. The authority citation for 21 CFR part 558 continues to read as follows:

**Authority:** 21 U.S.C. 360b, 371.

**§ 558.550 [Amended]**

2. Section 558.550 *Salinomycin* is amended in paragraph (a)(2) by adding ''(d)(2)(i),'' numerically.

Dated: January 21, 2003.

**Steven D. Vaughn,**

*Director, Office of New Animal Drug Evaluation, Center of Veterinary Medicine.*

[FR Doc. 03–3351 Filed 2–10–03; 8:45 am]
BILLING CODE 4160–01–S

## DEPARTMENT OF THE TREASURY

### Office of Foreign Assets Control

### 31 CFR Part 501

### Rules Governing Availability of Information

**AGENCY:** Office of Foreign Assets Control, Treasury.

**ACTION:** Final rule.

**SUMMARY:** The Office of Foreign Assets Control (''OFAC'') of the U.S. Department of the Treasury is issuing a final rule concerning the disclosure of certain civil penalties information. OFAC intends to publish information about civil penalties imposed and informal settlements on a weekly basis. If the publication falls on a holiday, or if required by an emergency, publication may be postponed to the following week.

**DATES:** This rule is effective February 11, 2003.

**FOR FURTHER INFORMATION CONTACT:** Chief of Civil Penalties, tel.: 202/622–6140, or Chief Counsel, tel. 202/622–2410.