

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Lasalocid; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correcting amendments.

SUMMARY: The Food and Drug Administration (FDA) is correcting a document amending the animal drug regulations to reflect approval of an original new animal drug application (NADA) that appeared in the **Federal Register** of April 27, 2006 (71 FR 24814). FDA is correcting a paragraph designation in the table for lasalocid cattle feeds which was drafted in error. This correction is being made to improve the accuracy of the animal drug regulations.

DATES: This rule is effective [*insert date of publication in the Federal Register*].

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV-6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240-276-9019, e-mail: george.haibel@fda.hhs.gov.

SUPPLEMENTARY INFORMATION: For the reasons set forth in the preamble, FDA is correcting 21 CFR part 558 to read as follows:

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.

§ 558.311 [Amended]

- 2. Section 558.311 is corrected in the table in the “Lasalocid sodium in grams per ton” column, in the entry for use of lasalocid at 30 to 600 grams per ton in combination with chlortetracycline at 500 to 4000 grams per ton, by removing the second paragraph designation “(xxiii)” and by adding in its place the paragraph designation “(xxviii)”.

Dated: June 1, 2006.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

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