

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Food and Drug Administration

## 21 CFR Part 522

## Animal Drugs, Feeds, and Related Products; Change of Sponsor

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

**SUMMARY:** The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor for a new animal drug application (NADA) from Schering-Plough Animal Health Corp. to Walco International, Inc.

**EFFECTIVE DATE:** May 27, 1997.

**FOR FURTHER INFORMATION CONTACT:** Thomas J. McKay, Center for Veterinary Medicine (HFV-102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301-827-0213.

**SUPPLEMENTARY INFORMATION:** Schering-Plough Animal Health Corp., P.O. Box 529, Kenilworth, NJ 07033, has informed FDA that it has transferred ownership of, and all rights and interests in, approved NADA 031-971 (cupric glycinate injection) to Walco International, Inc., 15 West Putnam, Porterville, CA 93257. Accordingly, the agency is amending the regulations in 21 CFR 522.518 to reflect the change of sponsor.

**List of Subjects in 21 CFR 522**

Animal drugs.

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 522 is amended as follows:

**PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS**

1. The authority citation for 21 CFR part 522 continues to read as follows:

**Authority:** Sec. 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b).

**§ 522.518 [Amended]**

2. Section 522.518 *Cupric glycinate injection* is amended in paragraph (b) by removing "000061" and adding in its place "No. 049185".

Dated: May 6, 1997.

**Robert C. Livingston,**

*Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.* [FR Doc. 97-13822 Filed 5-23-97; 8:45 am]

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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

## Food and Drug Administration

## 21 CFR Part 558

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

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 new animal drug application (NADA) filed by Hoffmann-La Roche, Inc. The supplemental NADA provides for removal of the international feed number (IFN) for an ingredient in free-choice, lasalocid, liquid Type C feed.

**EFFECTIVE DATE:** May 27, 1997.

**FOR FURTHER INFORMATION CONTACT:** Jack Caldwell, Center for Veterinary Medicine (HFV-126), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301-594-1638.

**SUPPLEMENTARY INFORMATION:** Hoffmann-La Roche, Inc., 340 Kingsland St., Nutley, NJ 07110-1199, filed supplemental NADA 96-298, which provides for removing the IFN for the condensed molasses fermentation solubles ingredient of the free-choice, lasalocid, liquid Type C feed. The molasses solubles described by the IFN refer to those solubles from sugar cane molasses. The liquid Type C feed contains beet molasses solubles that do not have an IFN.

The supplemental NADA is approved as of May 27, 1997, and the regulations are amended in 21 CFR 558.311(e)(3)(i) to reflect the approval.

This action does not affect the safety and effectiveness upon which the application was approved. Therefore, a freedom of information summary is not required.

The agency has determined under 21 CFR 25.24(a)(9) 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.

**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:** Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

**§ 558.311 [Amended]**

2. Section 558.311 *Lasalocid* is amended in the table in paragraph (e)(3)(i), in the entry for "Condensed Molasses Fermentation Solubles", in the third column by removing "5-25-399" and adding in its place "N/A".

Dated: May 7, 1997.

**Robert C. Livingston,**

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

[FR Doc. 97-13825 Filed 5-23-97; 8:45 am]

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## DEPARTMENT OF THE TREASURY

## Internal Revenue Service

## 26 CFR Parts 1 and 601

[TD 8719]

RIN 1545-AU41 and 1545-AV19

**Requirements Respecting the Adoption or Change of Accounting Method; Extensions of Time To Make Elections; Correction**

AGENCY: Internal Revenue Service (IRS), Treasury.

ACTION: Correction to temporary regulations.

**SUMMARY:** This document contains corrections to the temporary regulations (TD 8719) which were published in the **Federal Register** for Thursday, May 15, 1997 (62 FR 26740). The regulations relate to the procedure for requesting a change in accounting method and the standards for granting an extension of time to request a change in accounting method. The regulations provide for a longer period of time for filing an application for change in accounting method with the Commissioner.

**EFFECTIVE DATE:** May 15, 1997.

**FOR FURTHER INFORMATION CONTACT:** Cheryl L. Oseekey at (202) 622-4970 (not a toll-free number).

**SUPPLEMENTARY INFORMATION:****Background**

The temporary regulations that are the subject of this correction are under section 446 of the Internal Revenue Code.