

Ractopamine in grams/ton	Combination in grams/ton	Indications for use	Limitations	Sponsor
(ii) 4.5	Tylosin 40	Finishing swine: As in paragraph (e)(1)(i) of this section; and for prevention of swine dysentery (vibriotic).	Feed continuously as sole ration until market weight following the use of tylosin at 100 grams per ton (g/t) for at least 3 weeks.	000986
(iii) 4.5	Tylosin 100	1. Finishing swine: As in paragraph (e)(1)(i) of this section; and for prevention and/or control of porcine proliferative enteropathies (ileitis) associated with <i>Lawsonia intracellularis</i> . 2. Finishing swine: As in paragraph (e)(1)(i) of this section; and for prevention of swine dysentery (vibriotic).	Feed continuously as sole ration for 21 days. Feed continuously as sole ration for at least 3 weeks followed by tylosin at 40 g/t until market weight.	000986 000986
(iv) 4.5 to 18		For improved feed efficiency and increased carcass leanness in finishing swine fed a complete ration containing at least 16 percent crude protein from 150 lb (68 kg) to 240 lb (109 kg) body weight.	Feed continuously as sole ration.	000986
(v) 4.5 to 18	Tylosin 40	Finishing swine: As in paragraph (e)(1)(iv) of this section; and for prevention of swine dysentery (vibriotic).	Feed continuously as sole ration until market weight following the use of tylosin at 100 g/t for at least 3 weeks.	000986
(vi) 4.5 to 18	Tylosin 100	1. Finishing swine: As in paragraph (e)(1)(iv) of this section; and for prevention and/or control of porcine proliferative enteropathies (ileitis) associated with <i>Lawsonia intracellularis</i> . 2. Finishing swine: As in paragraph (e)(1)(iv) of this section; and for prevention of swine dysentery (vibriotic).	Feed continuously as sole ration for 21 days. Feed continuously as sole ration for at least 3 weeks followed by tylosin at 40 g/t until market weight.	000986 000986

(2) [Reserved]

Dated: November 8, 2002.

**Steven D. Vaughn,**

*Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.*  
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## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

#### 21 CFR Part 558

#### New Animal Drugs; 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 an approved new animal drug application (NADA) from Boehringer Ingelheim Vetmedica, Inc., to Pennfield Oil Co.

**DATES:** This rule is effective December 3, 2002.

**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:

Boehringer Ingelheim Vetmedica, Inc., 2621 North Belt Hwy., St. Joseph, MO 64506-2002, has informed FDA that it has transferred ownership of, and all rights and interest in, NADA 128-550 for ANCHOR Zinc Bacitracin Type A medicated article to Pennfield Oil Co., 14040 Industrial Rd., Omaha, NE 68137. Accordingly, the agency is amending the regulations in 21 CFR 558.78 to reflect the transfer of ownership.

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.

#### § 558.78 [Amended]

2. Section 558.78 *Bacitracin zinc* is amended in paragraph (a)(2) by removing "To 000010" and by adding in its place "No. 053389".

Dated: November 8, 2002.

**Steven D. Vaughn,**

*Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine.*  
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## DEPARTMENT OF THE TREASURY

### Internal Revenue Service

#### 26 CFR Part 1

[TD 9021]

RIN 1545-AX68

#### Loans From a Qualified Employer Plan to Plan Participants or Beneficiaries

**AGENCY:** Internal Revenue Service (IRS), Treasury.

**ACTION:** Final regulations.

**SUMMARY:** This document contains final regulations relating to loans made from a qualified employer plan to plan participants or beneficiaries. These final regulations affect administrators of, participants in, and beneficiaries of qualified employer plans that permit