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 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:** 21 U.S.C. 360b.

■ 2. In § 522.2471, revise paragraphs (e)(1)(i), (e)(1)(ii), and (e)(1)(iii) to read as follows:

**§ 522.2471 Tilmicosin.**

* * * * *

(e) * * *

(1) * * *

(i) *Amount.* 10 to 20 milligrams per kilograms (mg/kg) of body weight as a single subcutaneous injection.

(ii) *Indications for use.* For the treatment of bovine respiratory disease (BRD) associated with *Mannheimia haemolytica*, *Pasteurella multocida*, and *Histophilus somni*. For the control of respiratory disease in cattle at high risk of developing BRD associated with *M. haemolytica*.

(iii) *Limitations.* Do not use in female dairy cattle 20 months of age or older. Use of this antibiotic in this class of cattle may cause milk residues. Do not slaughter within 42 days of last treatment.

* * * * *

Dated: February 16, 2010.

**Steven D. Vaughn,**

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

[FR Doc. 2010–4206 Filed 3–1–10; 8:45 am]

**BILLING CODE 4160–01–S**

## DEPARTMENT OF HEALTH AND HUMAN SERVICES

### Food and Drug Administration

### 21 CFR Part 558

**[Docket No. FDA–2010–N–0002]**

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

**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 ADM Alliance Nutrition, Inc. The supplemental NADA provides for use of a higher concentration chlortetracycline Type A medicated article for the manufacture of medicated feeds for livestock and poultry.

**DATES:** This rule is effective March 2, 2010.

**FOR FURTHER INFORMATION CONTACT:** Cindy L. Burnsteel, Center for Veterinary Medicine (HFV–130), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 240–276–8341, e-mail: *cindy.burnsteel@fda.hhs.gov*.

**SUPPLEMENTARY INFORMATION:** ADM Alliance Nutrition, Inc., 1000 North 30th St., Quincy, IL 62305–3115, filed a supplement to NADA 48–480 that provides for the use of CHLORATET 50 (chlortetracycline), a Type A medicated article containing 50 grams of chlortetracycline per pound, for the manufacture of medicated feeds for livestock and poultry. The supplement provides for use of Type A medicated articles containing 90 or 100 grams of chlortetracycline per pound. The supplemental NADA is approved as of January 7, 2010, and the regulations are amended in 21 CFR 558.128 to reflect the approval.

Approval of this supplemental NADA did not require review of additional safety or effectiveness data or information. Therefore, a freedom of information summary is not required.

FDA has determined under 21 CFR 25.33 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 Subject 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.

■ 2. In § 558.128, revise paragraph (b)(2) to read as follows:

**§ 558.128 Chlortetracycline.**

* * * * *

(b) * * *

(2) No. 012286: 50, 90, or 100 grams per pound of Type A medicated article.

* * * * *

Dated: February 16, 2010.

**Steven D. Vaughn,**

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

[FR Doc. 2010–4205 Filed 3–1–10; 8:45 am]

**BILLING CODE 4160–01–S**

## DEPARTMENT OF LABOR

### Employee Benefits Security Administration

### 29 CFR Part 2520

**RIN 1210–AB21**

### Multiemployer Pension Plan Information Made Available on Request

**AGENCY:** Employee Benefits Security Administration, Labor.

**ACTION:** Final rule.

**SUMMARY:** This document contains a final rule implementing section 101(k) of the Employee Retirement Income Security Act of 1974, as amended by the Pension Protection Act of 2006. Section 101(k) requires the administrator of a multiemployer plan to provide copies of certain actuarial and financial documents about the plan to participants, beneficiaries, employee representatives and contributing employers upon request. The final rule affects plan administrators, participants and beneficiaries and contributing employers of multiemployer plans.

**DATES:** This final rule is effective on April 1, 2010.

**FOR FURTHER INFORMATION CONTACT:** June Solonsky or Stephanie L. Ward, Office of Regulations and Interpretations, Employee Benefits Security Administration, (202) 693–8500. This is not a toll-free number.

**SUPPLEMENTARY INFORMATION:**

### A. Background

Section 101(k) of the Employee Retirement Income Security Act