From	To	MEA
§ 95.6001 VICTOR ROUTES—U.S. § 95.6437 VOR FEDERAL AIRWAY V437 Is Amended To Read in Part		
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[FR Doc. 04–23375 Filed 10–21–04; 8:45 am]
BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 529

Certain Other Dosage Form New Animal Drugs; Oxytetracycline

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 Phoenix Scientific, Inc. The supplemental ANADA provides for use of oxytetracycline hydrochloride soluble powder for skeletal marking of finfish fry and fingerlings by immersion.

DATES: This rule is effective October 22, 2004.

FOR FURTHER INFORMATION CONTACT: Joan C. Gotthardt, Center for Veterinary Medicine (HFV–130), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–7571, e-mail: *joan.gotthardt@fda.gov*.

SUPPLEMENTARY INFORMATION: Phoenix Scientific, Inc., 3915 South 48th Street Ter., St. Joseph, MO 64503, filed a supplement to ANADA 200–247 that provides for use of Oxytetracycline HCl Soluble Powder–343 for skeletal marking of finfish fry and fingerlings by immersion. The approval of this supplemental ANADA relied on publicly available safety and effectiveness data contained in Public Master File (PMF) 5667 which were compiled under National Research Support Project 7 (NRSP–7), a national agricultural research program for obtaining clearances for use of new drugs in minor animal species and for special uses. The supplemental ANADA is approved as of September 15, 2004, and the regulations are amended in 21 CFR 529.1660 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 Division of Dockets Management (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(d)(4) 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 529

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 529 is amended as follows:

PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 529 continues to read as follows:

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

■ 2. Section 529.1660 is amended by revising paragraphs (a) and (b) to read as follows:

§ 529.1660 Oxytetracycline.

(a) *Specifications*—(1) Each gram of powder contains 366 milligrams (mg) oxytetracycline hydrochloride.

(2) Each gram of powder contains 753 mg oxytetracycline hydrochloride.

(b) *Sponsors.* See sponsors in § 510.600(c) of this chapter for use of products described in paragraph (a) of this section as in paragraph (d) of this section.

(1) No. 046573 for use of product described in paragraph (a)(1) of this section.

(2) No. 059130 for use of product described in paragraph (a)(2) of this section.

* * * * *

Dated: October 14, 2004.

Steven D. Vaughn,

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

[FR Doc. 04–23686 Filed 10–21–04; 8:45 am]
BILLING CODE 4160–01–S

FEDERAL COMMUNICATIONS COMMISSION

47 CFR Part 54

[CC Docket Nos. 96–45, 97–21, and 02–6; FCC 04–181]

Federal-State Joint Board on Universal Service; Changes to the Board of Directors for the National Exchange Carrier Association, Inc.; and Schools and Libraries Universal Service Support Mechanism

AGENCY: Federal Communications Commission.

ACTION: Final rule, correction.

SUMMARY: This document corrects an error in the dates and supplementary sections of a **Federal Register** document regarding the Commission addressing pending petitions for reconsideration filed by Sprint Corporation, United States Telecom Association, Inc., and MCI Worldcom, Inc. The Commission agreed with petitioners that the Commission should seek recovery from schools and libraries in certain instances, and therefore grants their petitions in part. The Commission resolved the limited question raised in the *Second Further Notice of Proposed Rulemaking (Second FNPRM)* in CC Docket No. 02–06 of from whom the Commission will seek recovery of schools and libraries funds disbursed in violation of the statute or a rule. The Commission modified its requirements in this area so that recovery will be sought from whichever party or parties has committed the statutory or rule violation. The summary was published in the **Federal Register** on September 17, 2004.

DATES: Effective October 17, 2004.