Directorate of Defense Trade Controls (DDTC). The office at the Department of State, formerly known as the Office of Defense Trade Controls and before that as the Office of Munitions Control, responsible for reviewing applications to export and reexport items on the U.S. Munitions List. (See 22 CFR parts 120 through 130.)

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PART 774—[AMENDED]

■ 11. The authority citation for part 774 continues to read as follows:

Authority: 50 U.S.C. app. 2401 *et seq.*; 50 U.S.C. 1701 *et seq.*; 10 U.S.C. 7420; 10 U.S.C. 7430(e); 18 U.S.C. 2510 *et seq.*; 22 U.S.C. 287c, 22 U.S.C. 3201 *et seq.*, 22 U.S.C. 6004; 30 U.S.C. 185(s), 185(u); 42 U.S.C. 2139a; 42 U.S.C. 6212; 43 U.S.C. 1354; 46 U.S.C. app. 466c; 50 U.S.C. app. 5; Sec. 901–911, Pub. L. 106–387; Sec. 221, Pub. L. 107–56; E.O. 13026, 61 FR 58767, 3 CFR, 1996 Comp., p. 228; E.O. 13222, 66 FR 44025, 3 CFR, 2001 Comp., p. 783; Notice of August 6, 2004, 69 FR 48763 (August 10, 2004).

■ 12. In Supplement No. 1 to Part 774, revise all references to the ''Office of Defense Trade Controls'' to read ''Directorate of Defense Trade Controls''; revise all references to ''Directorate of Defense Trade Control'' to read ''Directorate of Defense Trade Controls''; and revise all references to ''DTC'' to read ''DDTC''.

Dated: October 4, 2004.

Peter Lichtenbaum,

Assistant Secretary for Export Administration.

[FR Doc. 04–22861 Filed 10–8–04; 8:45 am]

BILLING CODE 3510–33–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 520 and 558

New Animal Drugs; Change of Sponsor; Sulfaquinoxaline

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 Hess & Clark, Inc., to Phoenix Scientific, Inc.

DATES: This rule is effective October 12, 2004.

FOR FURTHER INFORMATION CONTACT: David R. Newkirk, Center for Veterinary Medicine (HFV–100), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6967, e-mail: *david.newkirk@fda.gov*.

SUPPLEMENTARY INFORMATION: Hess & Clark, Inc., 944 Nandino Blvd., Lexington, KY 40511, has informed FDA that it has transferred ownership of , and all rights and interest in, the following three approved NADAs, to Phoenix Scientific, Inc., 3915 South 48th Street Ter., St. Joseph, MO 64503:

NADA Number	Trade Name
6–391	S.Q. (sulfaquinoxaline) 40% Medicated Feed
6–677	S.Q. (sulfaquinoxaline) 20% Solution
7–087	Sulfaquinoxaline Solubilized

Accordingly, the agency is amending the regulations in 21 CFR 520.2325a and 558.586 to reflect the transfer of ownership and a current format.

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

21 CFR Part 520

Animal drugs.

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 parts 520 and 558 are amended as follows:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 520 continues to read as follows:

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

§ 520.2325a [Amended]

■ 2. Section 520.2325a is amended in paragraph (a)(1) by removing ''050749'' and by adding in its place ''059130''.

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

■ 3. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

■ 4. Section 558.586 is amended by revising the section heading; by removing paragraphs (c) and (d); by redesignating paragraphs (e) and (f) as paragraphs (c) and (d); and by revising paragraph (a) and adding paragraph (b) to read as follows:

§ 558.586 Sulfaquinoxaline.

(a) *Specifications.* Type A medicated articles containing 40 percent sulfaquinoxaline.

(b) *Approvals.* See No. 059130 in § 510.600(c) of this chapter.

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Dated: September 27, 2004.

Steven D. Vaughn,

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

[FR Doc. 04–22760 Filed 10–8–04; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF DEFENSE

Office of the Secretary

32 CFR Part 199

RIN 0720–AA89

TRICARE; Changes Included in the National Defense Authorization Act for Fiscal Year 2002, (NDAA–02), and a Technical Correction Included in the NDAA–03

AGENCY: Office of the Secretary, DoD.

ACTION: Final rule.

SUMMARY: This rule makes several changes to the TRICARE program authorized by Congress in the NDAA–02. Specifically, revisions to the definition of durable medical equipment (DME); adoption of the same pricing methods for durable medical equipment, prosthetics, orthotics and supplies (DMEPOS) as are in effect for the Centers for Medicare & Medicaid Services (CMS); clarification that rehabilitative therapy is a TRICARE benefit; addition of augmentative communication devices (ACD)/speech generating devices (SGDs) as a TRICARE benefit; addition of hearing aids for family members of active duty members as a TRICARE Basic Program benefit; revisions to the definition of prosthetics; permanent authority for transitional health care for certain members separated from active duty; and revisions to the time period of eligibility for transitional health care.

This final rule also addresses a technical correction found in section 706 of the Bob Stump NDAA–03, relating to transitional health care for dependents of certain members separated from active duty.

DATES: This rule is effective December 13, 2004. Actual implementation will coincide with the transition in each TRICARE Region to the next generation