introductory text, and the first sentence of paragraph (e)(i), remove ''duplicate or corrected'' and add in its place ''replacement.''

■ b. In the third and fourth sentences of paragraph (a), the first sentence of paragraph (d) introductory text, and the first sentence of paragraph (g), remove '',duplicate, or corrected'' and add in its place ''or replacement.''

■ 4. Section 422.110 is revised to read as follows:

§ 422.110 Individual's request for change in record.

(a) *Form SS–5.* If you wish to change the name or other personal identifying information you previously submitted in connection with an application for a social security number card, you must complete and sign a Form SS–5 except as provided in paragraph (b) of this section. You must prove your identity, and you may be required to provide other evidence. (See § 422.107 for evidence requirements.) You may obtain a Form SS–5 from any local Social Security office or from one of the sources noted in § 422.103(b). You may submit a completed request for change in records to any Social Security office, or, if you are outside the U.S., to the Department of Veterans Affairs Regional Office, Manila, Philippines, or to any U.S. Foreign Service post or U.S. military post. If your request is for a change of name on the card, we may issue you a replacement card bearing the same number and the new name. We will grant an exception from the limitations specified in § 422.103(e)(2) for replacement social security number cards representing a change in name or, if you are an alien, a change to a restrictive legend shown on the card. (See § 422.103(e)(3) for the definition of a change to a restrictive legend.)

(b) *Assisting in enumeration.* We may enter into an agreement with officials of the Department of State and the Department of Homeland Security to assist us by collecting, as part of the immigration process, information to change the name or other personal identifying information you previously submitted in connection with an application or request for a social security number card. If your request is to change a name on the card or to correct the restrictive legend on the card to reflect a change in alien status, we may issue you a replacement card bearing the same number and the new name or legend. We will grant an exception from the limitations specified in § 422.103(e)(2) for replacement social security number cards representing a change of name or, if you are an alien, a change to a restrictive legend shown on the card. (See § 422.103(e)(3) for the definition of a change to a restrictive legend.)

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 558

Animal Drugs, Feeds, and Related Products; Withdrawal of Approval of New Animal Drug Applications

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations by removing those portions that reflect approval of 15 new animal drug applications (NADAs) because the products are no longer manufactured or marketed. In a notice published elsewhere in this issue of the **Federal Register**, FDA is withdrawing approval of the NADAs.

DATES: This rule is effective December 27, 2005.

FOR FURTHER INFORMATION CONTACT: Pamela K. Esposito, Center for Veterinary Medicine (HFV–212), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9067, e-mail: *pesposit@cvm.fda.gov*.

SUPPLEMENTARY INFORMATION: The following sponsors have requested that FDA withdraw approval of the 15 NADAs listed in table 1 of this document because the products are no longer manufactured or marketed:

TABLE 1.

Sponsor	NADA Number, Product (Drug)	21 CFR Section Affected (Sponsor Drug Labeler Code)
Bioproducts, Inc., 320 Springside Dr., Suite 300, Fairlawn, OH 44333–2435	NADA 119–063, Pyrantel Tartrate Ton Pack (pyrantel tartrate)	558.485 (051359)
Farmland Industries, Inc., Kansas City, MO 64116	NADA 138–656, BN Wormer—19.2 BANMINTH Premix (pyrantel tartrate)	558.485 (021676)
I.M.S. Inc., 13619 Industrial Rd., Omaha, NE 68137	NADA 129–395, HYGROMIX 0.6 Premix (hygromycin B)	558.274 (050639)
	NADA 129–646, TYLAN 10 Sulfa-G (tylosin, sulfamethazine)	558.630 (050639)
	NADA 136–601, Swine Guard BN (pyrantel tartrate)	558.485 (050639)
J. & R. Specialty Supply Co., 310 Second Ave., SW., P.O. Box 506, Waseca, MN 56093	NADA 96–780, TYLAN 10; TYLAN 40 (tylosin)	n/a (049768)
Kerber Milling Co., Box 152, 1817 E. Main St., Emmetsburg, IA 50536	NADA 98–687, Hy-Test Hy-Boost TY 5 Medicated (tylosin)	558.625 (029341)
M & M Livestock Products Co., Eagle Grove, IA 50533	NADA 96–837, M & M Tylosin Premix (tylosin)	558.625 (026282)
Nutra-Blend Corp., P.O. Box 485, Neosho, MO 64850	NADA 129–161, Nutra-Blend TYLAN 10 Sulfa Premix (tylosin, sulfamethazine)	558.630 (050568)
	NADA 136–384, Swine Wormer-BN BANMINTH (pyrantel tartrate)	558.485 (050568)

TABLE 1.—Continued

Sponsor	NADA Number, Product (Drug)	21 CFR Section Affected (Sponsor Drug Labeler Code)
South St. Paul Feeds, Inc., 500 Farwell Ave., South St. Paul, MN 55075	NADA 136–369, Custom Ban Wormer 9.6 (pyrantel tartrate)	558.485 (001800)
Stockton Hay & Grain Co.	NADA 49–462, Rainbrook Broiler Premix No. 1 (ampolium, arsanilic acid, ethopabate, penicillin G procaine, streptomycin) NADA 91–646, Rainbow Broiler Base Concentrate (ampolium, bacitracin zinc, ethopabate) NADA 91–647, Rainbow Broiler Base Concentrate (ampolium, chlortetracycline, ethopabate)	n/a (036541) n/a (036541) n/a (036541)
Triple ''F'', Inc., 10104 Douglas Ave., Des Moines, IA 50322	NADA 131–146, FLAVOMYCIN 0.4 (bambermycins)	558.95 (011490)

Following the withdrawal of approval of these NADAs, Kerber Milling Co., M & M Livestock Products Co., Nutra-Blend Corp., and South St. Paul Feeds, Inc., are no longer sponsors of an approved application. Therefore, we are removing entries for these four sponsors from 21 CFR 510.600(c).

As provided below, the animal drug regulations are amended to reflect the withdrawal of approvals.

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 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

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 510 and 558 are amended as follows:

PART 510—NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

§ 510.600 [Amended]

■ 2. Section 510.600 is amended in the table in paragraph (c)(1) by removing the entries for ''Kerber Milling Co.'', ''M & M Livestock Products Co.'', ''Nutra-Blend Corp.'', and ''South St. Paul Feeds, Inc.''; and in the table in paragraph (c)(2) by removing the entries for ''001800'', ''026282'', ''029341'', and ''050568''.

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.

§ 558.95 [Amended]

■ 4. Section 558.95 is amended by removing and reserving paragraph (a)(3).

§ 558.274 [Amended]

■ 5. Section 558.274 is amended in paragraph (a)(4) by removing '', 043733, and 050639'' and by adding in its place ''and 043733''; and in the table in paragraphs (c)(1)(i) and (c)(1)(ii) in the ''Sponsor'' column by removing '', 050639''.

§ 558.485 [Amended]

■ 6. Section 558.485 is amended by removing and reserving paragraphs (b)(2) and (b)(4); and in paragraph (b)(3) by removing '', 049685, 050568, 050639, and 051359'' and by adding in its place ''and 049685''.

§ 558.625 [Amended]

■ 7. Section 558.625 is amended by removing and reserving paragraphs (b)(22), (b)(31), (b)(52), and (b)(79).

§ 558.630 [Amended]

■ 8. Section 558.630 is amended in paragraph (b)(10) by removing '', 050568, 050639''.

Dated: December 7, 2005.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

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DEPARTMENT OF JUSTICE

Drug Enforcement Administration

21 CFR Parts 1300 and 1308

[Docket No. DEA–264]

RIN 1117–AA95

Implementation of the Anabolic Steroid Control Act of 2004

AGENCY: Drug Enforcement Administration (DEA), Department of Justice.

ACTION: Final rule.

SUMMARY: The purpose of this rulemaking is to conform the Drug Enforcement Administration's (DEA) regulations to the provisions of the Anabolic Steroid Control Act of 2004. Effective January 20, 2005, the Act amended the Controlled Substances Act (CSA) and replaced the existing definition of ''anabolic steroid'' with a new definition. This new definition altered the basis for all future administrative scheduling actions relating to the control of anabolic steroids as Schedule III controlled substances by eliminating the requirement to prove muscle growth. Additionally, the Act lists 59 specific substances as being anabolic steroids. As such, these substances and their salts, esters and ethers are Schedule III controlled substances. This rulemaking amends 21 CFR Parts 1300 and 1308 to reflect these changes.

The Act also amends the CSA by revising the language requiring exclusion of certain over the counter products from regulation as controlled substances. The Act clarifies that the exclusionary language in 21 U.S.C. 811(g)(1) pertains only to non-narcotic ''drugs'' that may, under the Federal Food, Drug, and Cosmetic Act (FDCA),