80a–2(a)(3)(C)), of the issuer of, or any insurer or provider of credit support for, the security.

(6) *Requisite NRSROs* means:

(i) Any two NRSROs that have issued a rating with respect to a security or class of debt obligations of an issuer; or

(ii) If only one NRSRO has issued a rating with respect to such security or class of debt obligations of an issuer at the time the investment company acquires the security, that NRSRO.

(7) *Resale Price* means the acquisition price paid to the seller of the securities plus the accrued resale premium on such acquisition price. The accrued resale premium is the amount specified in the repurchase agreement or the daily amortization of the difference between the acquisition price and the resale price specified in the repurchase agreement.

(8) *Unrated Securities* means securities that have not received a rating from the Requisite NRSROs.

4. Section 270.12d3–1 is amended by removing the note following paragraph (d)(8).

PART 274—FORMS PRESCRIBED UNDER THE INVESTMENT COMPANY ACT OF 1940

5. The authority citation for Part 274 continues to read as follows:

Authority: 15 U.S.C. 77f, 77g, 77h, 77j, 77s, 78c(b), 781, 78m, 78n, 78o(d), 80a–8, 80a–24, and 80a–29, unless otherwise noted.

Note: The text of Form N–SAR does not, and this amendment will not, appear in the *Code of Federal Regulations*.

6. Form N–SAR (referenced in 17 CFR 274.101) is amended by revising the second sentence in the first paragraph of the Instructions to Specific Items 24 and 25 to read as follows:

FORM N–SAR

* * * * *

Instructions to Specific Items

* * * * *

ITEMS 24 and 25: Acquisition of securities of registrant's regular brokers or dealers

* * * *See* Rule 12d3–1, Investment Company Act Release No. 14036, dated July 13, 1984, adopting Rule 12d3–1, and Investment Company Act Release No. 25058, dated July 5, 2001, amending Rule 12d3–1. * * *

* * * * *

Dated: July 5, 2001.

By the Commission.

Margaret H. McFarland,

Deputy Secretary.

[FR Doc. 01–17302 Filed 7–10–01; 8:45 am]

BILLING CODE 8010–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 558

Animal Drugs, Feeds, and Related Products; Tylosin; Withdrawal of Approval of NADAs

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to remove those portions that reflect approval of two new animal drug applications (NADAs) listed below. In a notice published elsewhere in this issue of the **Federal Register**, FDA is withdrawing approval of the NADAs.

DATES: This rule is effective July 23, 2001.

FOR FURTHER INFORMATION CONTACT: Pamela K. Esposito, Center for Veterinary Medicine (HFV–210), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–5593.

SUPPLEMENTARY INFORMATION: Heinold Feeds, Inc., P.O. Box 377, Kouts, IN 46347, has requested that FDA withdraw approval of NADA 95–628 for Tylosin® Antibiotic Premix and NADA 127–506 for Tylan® Sulfa-G Premixes because the products are no longer manufactured or marketed.

Following the withdrawal of approval of these NADAs, Heinold Feeds, Inc., is no longer the sponsor of any approved applications. Therefore, 21 CFR 510.600(c) is amended to remove entries for this sponsor.

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 *Names, addresses, and drug labeler codes of sponsors of approved applications* is amended in the table in paragraph (c)(1) by removing the entry for ''Heinold Feeds, Inc.,'' and in the table in paragraph (c)(2) by removing the entry for ''043727''.

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.

§ 558.625 [Amended]

4. Section 558.625 *Tylosin* is amended by removing and reserving paragraph (b)(9).

§ 558.630 [Amended]

5. Section 558.630 *Tylosin and sulfamethazine* is amended in paragraph (b)(10) by removing ''043727,''; and by removing ''and 051359, 053389'' and by adding in its place ''051359, and 053389''.

Dated: July 2, 2001.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 01–17407 Filed 7–10–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF TRANSPORTATION

Coast Guard

33 CFR Part 117

[CGD08–01–014]

Drawbridge Operating Regulation; Green River, Spottsville, Kentucky

AGENCY: Coast Guard, DOT.

ACTION: Notice of temporary deviation from drawbridge regulations.