Dated: February 3, 1995.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 95–3804 Filed 2–14–95; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Part 558

New Animal Drugs for Use in Animal Feeds; Tylosin and Virginiamycin

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 reflecting approval of four new animal drug applications (NADA's) held by Premiere Agri Technologies, Inc. The NADA's provide for use of Type A medicated articles and Type B medicated feeds containing tylosin and Type B medicated feeds containing virginiamycin. In a notice published elsewhere in this issue of the **Federal Register**, FDA is withdrawing approval of the NADA's.

EFFECTIVE DATE: February 27, 1995.

FOR FURTHER INFORMATION CONTACT: Mohammad I. Sharar, Center for Veterinary Medicine (HFV–216), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1722.

SUPPLEMENTARY INFORMATION: In a notice published elsewhere in this issue of the **Federal Register**, FDA is withdrawing approval of the following NADA's:

NADA No.	Drug name	Sponsor name and address
45–690	Tylosin Type B medicated feeds and Type A medicated article.	Premiere Agri Technologies, Inc., P.O. Box 2508, Fort Wayne, IN 46801–2508 (former sponsor Henwood Feed Additives)
97–289	Tylosin Type B medicated feeds and Type A medicated article.	Do. (Former sponsor Feed Specialties Co., Inc.)
133–361	Virginiamycin Type B medicated feed.	Do. (Former sponsor Feed Specialties Co., Inc.)
133–839	Virginiamycin Type B medicated feed.	Do. (Former sponsor MacPage, Inc.)

The sponsor requested withdrawal of approval of the NADA's. This final rule removes 21 CFR 558.625(b)(11) and (b)(15) and amends 21 CFR 558.635(b)(2) to reflect the withdrawal of approval of these NADA's.

List of Subjects 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: Secs. 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b, 371).

§ 558.625 [Amended]

2. Section 558.625 *Tylosin* is amended by removing and reserving paragraphs (b)(11) and (b)(15).

3. Section 558.635 *Virginiamycin* is amended by revising paragraph (b)(2) to read as follows:

§ 558.635 Virginiamycin

* * * * *

(b) * * *

(2) 2.2 percent activity (10 grams per pound) to 011490, 016968, and 017790 in § 510.600(c) of this chapter for use as in paragraphs (f)(1)(iv) and (f)(1)(v) of this section.

* * * * *

Dated: January 6, 1995.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 95–3802 Filed 2–14–95; 8:45 am]

BILLING CODE 4160–01–F

UNITED STATES INFORMATION AGENCY

22 CFR Part 514

[Rulemaking No. 110]

Exchange Visitor Program

AGENCY: United States Information Agency.

ACTION: Final rule.

SUMMARY: The Agency hereby adopts as final with modifications the interim rule governing its oversight and administration of au pair programs. Au pair programs permit foreign nationals to enter the United States for a period of one year for the purpose of residing with an American host family while participating directly in the home life of the family and providing limited child care services. The foreign national also attends a United States accredited post-secondary educational institution. These rules are promulgated pursuant to Public Law 103–415 which authorizes the continued operation, until September 30, 1995, of au pair programs currently designated by the Agency.

DATES: Effective date: These rules are effective February 15, 1995.

Applicability dates: With the exceptions of § 514.31(j) (1) and (4), and § 514.31(k), these rules apply to all au pair placements and operations as of February 15, 1995. The provisions set forth at § 514.31(j) (1) and (4) and § 514.31(k) shall apply only to au pair participants placed after date of publication.

Compliance date: Sponsor implementation of the provisions set forth at § 514.31(g) (1) and (2) will not be expected before March 31, 1995.

FOR FURTHER INFORMATION CONTACT: Stanley S. Colvin, Assistant General Counsel, United States Information Agency, 301 4th Street, SW., Washington, DC 20547; Telephone, (202) 619–6829.

SUPPLEMENTARY INFORMATION: First begun pursuant to the provisions of the United States Information and Educational Exchange Act of 1948 (''Smith-Mundt''), and subsequently incorporated into and broadened under the Fulbright-Hays Act, educational and cultural exchange activities have, over the past forty years, exposed millions of foreign nationals to the United States, its peoples, cultures, skills, business techniques, educational institutions, and way of life. The Fulbright-Hays Act mandates reciprocal exchange and Americans traveling abroad have, in similar fashion, developed an enhanced awareness of foreign people, their cultures and societies. Thus, Fulbright-Hays programs further one of the Agency's primary missions: increasing mutual understanding between Americans and others through people-to-people contact. Originally conducted by the Department of State, oversight of exchange activities, occurring under the umbrella of the Exchange Visitor Program, has been the responsibility of the Agency since 1978.

The Fulbright-Hays Act sets forth certain parameters which all exchange activities must meet. With an eye towards ensuring that these parameters were being met and acting in response to a Congressional request, the General Accounting Office (''GAO'') investigated Agency oversight and administration of the Exchange Visitor Program and its attendant utilization of the J visa. In its report to Congress, dated February 5, 1990 and entitled ''Inappropriate Uses