Substances	Limitations
* * * * * * *	
Di-*tert*-butyl-*m*-cresyl phosphonite condensation product with biphenyl (CAS Reg. No. 178358–58–2) produced by the condensation of 2,4-di-*tert*-butyl-*m*-cresol with the Friedel-Crafts addition product (phosphorus trichloride and biphenyl) so that the food additive has a minimum phosphorus content of 5.0 percent.	For use only: 1. At levels not to exceed 0.1 percent by weight of olefin polymers complying with § 177.1520(c) of this chapter, items 1.1, 2.1, 2.2, 3.1(a), 3.1(b), 3.2(a), or 3.2(b).
* * * * * * *	

Dated: January 6, 1999.

L. Robert Lake,

Director, Office of Policy, Planning and Strategic Initiatives, Center for Food Safety and Applied Nutrition.

[FR Doc. 99–1032 Filed 1–15–99; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

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

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 new animal drug application (NADA) filed by Elanco Animal Health, A Division of Eli Lilly and Co. The supplemental NADA provides for use of monensin and tylosin Type A medicated articles for making Type B and C cattle feeds, the Type C cattle feed to be fed at a range of 60 to 90 milligrams of tylosin per head per day (mg/hd/day) rather than the currently approved 90 mg/hd/day.

EFFECTIVE DATE: January 19, 1999.

FOR FURTHER INFORMATION CONTACT: Jack Caldwell, Center for Veterinary Medicine (HFV–126), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0217.

SUPPLEMENTARY INFORMATION: Elanco Animal Health, A Division of Eli Lilly and Co., Lilly Corporate Center, Indianapolis, IN 46285, filed supplemental NADA 104–646 that provides for combining Rumensin® (80 grams per pound (g/lb) monensin sodium) and Tylan® (40 or 100 g/lb tylosin phosphate) Type A medicated articles to make Type B and C medicated cattle feeds. The Type C medicated cattle feeds are to be fed to cattle fed in confinement for slaughter at 50 to 360 mg/hd/day monensin and 60 to 90 mg/hd/day tylosin for improved feed efficiency and reduction of incidence of liver abscesses caused by *Fusobacterium necrophorum* and *Actinomyces pyogenes*. The tylosin feeding level is the same as currently approved under 21 CFR 558.625(f)(1)(i)(*c*) for use of tylosin Type C cattle feeds. The supplemental NADA is approved as of November 19, 1998, and the regulations are amended in 21 CFR 558.355(f)(3)(ii)(*b*) to reflect the approval.

A summary of data and information submitted to support approval of this supplemental application may be seen in the Dockets Management Branch (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.

The agency has determined under 21 CFR 25.33(a)(2) 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.

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

§ 558.355 [Amended]

2. Section 558.355 *Monensin* is amended in paragraph (f)(3)(ii)(*b*) by removing ''90'' and adding in its place ''60 to 90.''

Dated: December 17, 1998.

Andrew J. Beaulieu,

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

[FR Doc. 99–1037 Filed 1–15–99; 8:45 am]

BILLING CODE 4160–01–F

POSTAL SERVICE

39 CFR Part 20

Global Direct—Canada Admail Service

AGENCY: Postal Service.

ACTION: Final rule.

SUMMARY: The Postal Service published an interim rule and request for comment on a new service, Global Direct—Canada Admail, in the **Federal Register** on August 21, 1998, (63 FR 44789). The Postal Service hereby gives notice that it is adopting the interim rule.

EFFECTIVE DATE: January 19, 1999.

FOR FURTHER INFORMATION CONTACT: Walter J. Grandjean, (202) 314–7256.

SUPPLEMENTARY INFORMATION: In cooperation with Canada Post Corporation (CPC), the Postal Service introduced, on an interim basis, Global Direct—Canada Admail. This international mail service is primarily intended for major printing firms, direct marketers, mail order companies, and other high-volume mailers seeking easier access to the Canadian domestic postal system. It is intended to provide mail delivery in an average of 5 to 10 business days in major urban areas throughout Canada. Ancillary services for local business reply and the return of undeliverable mail are also introduced for use with Global Direct—Canada Admail.

On August 21, 1998, the Postal Service published in the **Federal Register** (63 FR 44789) an interim rule and request for comment on this new service, Global Direct—Canada Admail. Comments were requested on or before September 21, 1998.

The Postal Service did not receive any written comments on the interim rule