PART 178—INDIRECT FOOD ADDITIVES: ADJUVANTS, PRODUCTION AID, AND SANITIZERS

1. The authority citation for 21 CFR part 178 continues to read as follows:

Authority: Secs. 201, 402, 409, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 342, 348, 379e).

2. Section 178.3400 is amended in the table in paragraph (c) by alphabetically adding a new entry under the headings ''List of Substances'' and ''Limitations'' to read as follows:

§ 178.3400 Emulsifiers and/or surface-active agents.

* * * * *

(c) * * *

List of Substances	Limitations
* * *	* * * *
Polyethyleneglycol alkyl(C_{10}–C_{12}) ether sulfosuccinate, disodium salt (CAS Reg. No. 68954–91–6).	For use only at levels not to exceed 5 percent by weight of total monomers used in the emulsion polymerization of polyvinyl acetate, acrylic, and vinyl/acrylic polymers intended for use as coatings for paper and paperboard.
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Dated: June 5, 1997.

Fred R. Shank,

Director, Center for Food Safety and Applied Nutrition.

[FR Doc. 97–16399 Filed 6–23–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 524 and 556

Animal Drugs, Feeds, and Related Products; Eprinomectin

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 new animal drug application (NADA) filed by Merck Research Laboratories, Division of Merck & Co., Inc. The NADA provides for use of eprinomectin on cattle for treatment and control of certain gastrointestinal roundworms, lungworms, cattle grubs, lice, mange mites, and flies. The regulations are also amended to provide for a tolerance for residues of the drug in milk and in edible tissues.

EFFECTIVE DATE: June 24, 1997.

FOR FURTHER INFORMATION CONTACT: Melanie R. Berson, Center for Veterinary Medicine (HFV–135), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1643.

SUPPLEMENTARY INFORMATION: Merck Research Laboratories, Division of Merck & Co., Inc., P.O. Box 2000, Rahway, NJ 07065–0914, filed NADA 141–079, which provides for the use of Ivomec® Eprinex™ Pour-On (5 milligrams per milliliter eprinomectin) on cattle for the treatment and control of gastrointestinal roundworm, lungworm, cattle grub, lice, mange mite, and fly infections. The NADA is approved as of April 16, 1997, and the regulations are amended by adding new § 524.814 to reflect the approval. The regulations are also amended to provide for a tolerance for eprinomectin residues in milk and edible cattle tissues in new § 556.227. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part 20 and 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

Under section 512(c)(2)(F)(i) of the Federal Food, Drug, and Cosmetic Act, this approval qualifies for a 5-year period of marketing exclusivity beginning April 16, 1997, because no active ingredient (including any ester or salt of the active ingredient) of the drug has been approved in any other application filed under section 512(b)(1) of the act.

FDA has carefully considered the potential environmental effects of this action. FDA has concluded that the action will not have a significant impact on the human environment, and that an environmental impact statement is not required. FDA's finding of no significant impact and the evidence supporting that finding, contained in an environmental assessment, may be seen in the Dockets Management Branch (address above) between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects

21 CFR Part 524

Animal drugs.

21 CFR Part 556

Animal drugs, Foods.

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 524 and 556 are amended as follows:

PART 524—OPHTHALMIC AND TOPICAL DOSAGE FORM NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 524 continues to read as follows:

Authority: Sec. 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b).

2. New § 524.814 is added to read as follows:

§ 524.814 Eprinomectin.

(a) *Specifications.* Each milliliter contains 5 milligrams of eprinomectin.

(b) *Sponsor.* See No. 000006 in § 510.600(c) of this chapter.

(c) *Related tolerances.* See § 556.227 of this chapter.

(d) *Conditions of use*—(1) *Amount.* One milliliter (5 milligrams) per 10 kilograms of body weight (500 micrograms per kilogram).

(2) *Indications for use.* The drug used in beef and dairy cattle for the treatment and control of adult and fourth stage larvae (L4) gastrointestinal nematodes (*Haemonchus placei, Ostertagia*

ostertagi (including inhibited L4)), *Trichostrongylus axei, T. colubriformis, Cooperia oncophora, C. punctata, C. surnabada, Nematodirus helvetianus, Bunostomum phlebotomum, Oesophagostomum radiatum, Trichuris* spp. (adults); lungworms (adult and L4) (*Dictyocaulus viviparus*); cattle grubs (all parasitic stages) (*Hypoderma lineatum, H. bovis*); lice (*Damalinia bovis, Linognathus vituli, Haematopinus eurysternus, Solenopotes capillatus*); mange mites (*Chorioptes bovis, Sarcoptes scabiei*), and flies (*Haematobia irritans*). Controls *H. irritans* for 7 days and *D. vivaparus* for 21 days after treatment.

(3) *Limitations.* Apply topically along backbone from withers to tailhead. Consult your veterinarian for assistance in the diagnosis, treatment, and control of parasitism.

PART 556—TOLERANCES FOR RESIDUES OF NEW ANIMAL DRUGS IN FOOD

3. The authority citation for 21 CFR part 556 continues to read as follows:

Authority: Secs. 402, 512, 701 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 342, 360b, 371).

4. New § 556.227 is added to subpart B to read as follows:

§ 556.227 Eprinomectin.

Tolerances are established for residues of eprinomectin B1a (marker residue) in milk of 12 parts per billion and in liver (target tissue) of 4.8 parts per million.

Dated: June 5, 1997.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 97–16398 Filed 6–23–97; 8:45 am]

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DEPARTMENT OF DEFENSE

Department of the Army

32 CFR Part 552

[APG Reg 1–1]

Protests, Picketing, and Other Similar Demonstrations on the Installation of Aberdeen Proving Ground, MD

AGENCY: Department of the Army, DoD.

ACTION: Final Rule.

SUMMARY: This final rule establishes 32 CFR Part 552, Subpart P, Protests, Picketing, and Other similar Demonstrations, and authenticates Aberdeen Proving Ground Regulation, APG Reg. 1–1. This subpart implements policies, responsibilities, and procedures for protests, picketing, and other similar demonstration on the Aberdeen Proving Ground military reservation. This regulation is applicable to all personnel assigned, residing, working, or visiting on the Aberdeen Proving Ground reservation.

EFFECTIVE DATE: June 24, 1997.

ADDRESSES: Commander, U.S. Army Test and Evaluation Command, Office of the Chief Counsel and Staff Judge Advocate, Aberdeen Proving Ground, Maryland 21005.

FOR FURTHER INFORMATION CONTACT: Laura R. Haug, Deputy Chief Counsel, telephone (410) 278–1105 or 1107.

SUPPLEMENTARY INFORMATION: Supplementation of this subpart by subordinate units is prohibited.

On April 2, 1997, we published the proposed rule in the Notice of Proposed Rulemaking section of the **Federal Register** (Vol. 62, No. 63, pages 15639–15640) with the comment period ending on May 2, 1997.

We did not receive any objections to the proposed rule. We did, however, receive a comment from a citizen who indicated that the wording in § 552.213(a) that Aberdeen Proving Ground ''is NOT open for expressive activity'' is inconsistent with regulation. We agree with this comment since expressive activity may be permitted in certain circumstances with the Commander's approval based on the Commander's concerns for discipline, mission accomplishment, protection of property, and the health, morale, and welfare of the Aberdeen Proving Ground community. Therefore, § 552.213 is amended to indicate that Aberdeen Proving Ground ''is a non-public forum and is open for expressive activity only under certain circumstances.''

Executive Order 12291

This rule is not a major rule as defined by Executive Order 12291.

Regulatory Flexibility Act

The Regulatory Flexibility Act has no bearing on this rule.

Paperwork Reduction Act

This rule does not contain reporting or recordkeeping requirements subject to the Paperwork Reduction Act.

List of Subjects in 31 CFR Part 552

Federal buildings and facilities.

32 CFR part 552 is amended by adding and reserving subpart O and adding a new subpart P as follows:

Subpart P—Protests, Picketing, and Other Similar Demonstrations on the Installation of Aberdeen Proving Ground, Maryland

Authority: 18 U.S.C. Sec. 1382.

Subpart P—Protests. Picketing, and Other Similar Demonstrations on the Installation of Aberdeen Proving Ground, Maryland

§ 552.211 Purpose.

This subpart establishes policies, responsibilities, and procedures for protests, pickeing, and other similar demonstrations on the Aberdeen Proving Ground installation.

§ 552.212 Scope.

(a) The provisions of this subpart apply to all elements of U.S. Army Garrison, Aberdeen Proving Ground (USAGAPG), and the supported organizations and activities on the Aberdeen and Edgewood Areas of Aberdeen Proving Ground.

(b) The provisions of this subpart cover all public displays of opinions made by protesting, picketing, or any other similar demonstration.

(c) The provisions of this subpart are applicable to all people, military and civilian employees, and all visitors, family members, or others, entering, upon or present at Aberdeen Proving Ground.

§ 552.213 Policy.

(a) Aberdeen Proving Ground is a non-public forum and is open for expensive activity only under certain circumstances. Aberdeen Proving Ground is a military installation under the exclusive federal jurisdiction at which official business of the federal government is conducted, including military training, testing of weapon systems and other military equipment, and other official business.

(b) On Aberdeen Proving Ground, except for activities authorized under 5 United States Code Chapter 71, Labor Management Relations, it is unlawful for any person to engage in any public displays of opinions made by protesting, picketing or any other similar demonstration without the approval of the Commander, U.S. Army Garrison, Aberdeen Proving Ground. Therefore, unless prior approval has been obtained as outlined below in 32 CFR 552.214, it will be unlawful for any person on Aberdeen Proving Ground to: