* * * *Effective July 17, 1997*

Kake, AK, Kake, NDB or GPS RWY 10, Orig Cancelled
Kake, AK, Kake, NDB RWY 10, Orig
Arkadelphia, AR, Arkadelphia Muni, NDB or GPS RWY 4, Amdt 6 Cancelled
Arkadelphia, AR, Arkadelphia Muni, NDB RWY 4, Amdt 6
Burlington, CO, Kit Carson County, NDB or GPS RWY 15, Orig Cancelled
Burlington, CO, Kit Carson County, NDB RWY 15, Orig
Hayden, CO, Yampa Valley, VOR/DME or GPS–B, Orig Cancelled
Hayden, CO, Yampa Valley, VOR/DME–B, Orig
Hayden, CO, Yampa Valley, VOR or GPS–A, Amdt 3 Cancelled
Hayden, CO, Yampa Valley, VOR–A, Amdt 3
Lake City, FL, Lake City Muni, NDB or GPS RWY 28, Amdt 1 Cancelled
Lake City, FL, Lake City Muni, NDB RWY 28, Amdt 1
Marco Island, FL, Marco Island, VOR/DME or GPS RWY 17, Amdt 6 Cancelled
Marco Island, FL, Marco Island, VOR/DME RWY 17, Amdt 6
Tampa, FL, Tampa Intl, VOR or GPS RWY 9, Amdt. 7B Cancelled
Tampa, FL, Tampa Intl, VOR RWY 9, Amdt. 7B
Brunswick, GA, Malcolm McKinnon, VOR or GPS RWY 4, Amdt 14B Cancelled
Brunswick, GA, Malcolm McKinnon, VOR RWY 4, Amdt 14B
Macon, GA, Middle Georgia Regional, VOR or GPS RWY 13, Amdt 7B Cancelled
Macon, GA, Middle Georgia Regional, VOR RWY 13, Amdt 7B
Macon, GA, Middle Georgia Regional, VOR or GPS RWY 23, Amdt 1A Cancelled
Macon, GA, Middle Georgia Regional, VOR RWY 23, Amdt 1A
Macon, GA, Middle Georgia Regional, NDB or GPS RWY 5, Amdt 20A Cancelled
Macon, GA, Middle Georgia Regional, NDB RWY 5, Amdt 20A
Newnan, GA, Newnan-Coweta County, NDB or GPS RWY 32, Amdt 3 Cancelled
Newnan, GA, Newnan-Coweta County, NDB RWY 32, Amdt 3
Crawfordsville, IN, Crawfordsville Muni, NDB or GPS RWY 4, Amdt 4 Cancelled
Crawfordsville, IN, Crawfordsville Muni, NDB RWY 4, Amdt 5
Huntington, IN, Huntington Muni, NDB or GPS RWY 9, Orig Cancelled
Huntington, IN, Huntington Muni, NDB RWY 9, Orig
Sullivan, IN, Sullivan County, NDB or GPS RWY 36, Amdt 6 Cancelled
Sullivan, IN, Sullivan County NDB RWY 36, Amdt 6
Washington, IN, Daviess County, NDB or GPS RWY 18, Amdt 5 Cancelled
Washington, IN, Daviess County, NDB RWY 18, Amdt 5
Mount Sterling, KY, Mount Sterling-Montgomery County, NDB or GPS RWY 21, Amdt 1 Cancelled
Mount Sterling, KY, Mount Sterling-Montgomery County, NDB RWY 21, Amdt 1
Monticello, KY, Wayne County, NDB or GPS RWY 21, Amdt 1 Cancelled
Monticello, KY, Wayne County, NDB RWY 21, Amdt 1
Vandalia, IL, Vandalia Muni, VOR or GPS RWY 18, Amdt 11 Cancelled
Vandalia, IL, Vandalia Muni, VOR RWY 18, Amdt 11
Houlton, ME, Houlton Intl, VOR or GPS RWY 5, Amdt 9A Cancelled
Houlton, ME, Houlton Intl, VOR RWY 5, Amdt 9A
Drummond Island, MI, Drummond Island, NDB or GPS RWY 26, Amdt 1 Cancelled
Drummond Island, MI, Drummond Island, NDB RWY 26, Amdt 1
Chadron, NE, Chadron Muni, VOR/DME or GPS RWY 2, Amdt 1A Cancelled
Chadron, NE, Chadron Muni, VOR/DME RWY 2, Amdt 1A
Fremont, NE, Fremont Muni, NDB or GPS RWY 13, Amdt 2A Cancelled
Fremont, NE, Fremont Muni, NDB RWY 13, Amdt 2A
Valentine, NE, Miller Field, NDB or GPS RWY 31, Amdt 6B Cancelled
Valentine, NE, Miller Field, NDB RWY 32, Amdt 6B
Wahoo, NE, Wahoo Muni, NDB or GPS RWY 20, Amdt 2 Cancelled
Wahoo, NE, Wahoo Muni, NDB RWY 20, Amdt 2
Raton, NM, Raton Municipal/Crews Field, NDB or GPS RWY 2, Amdt 3A Cancelled
Raton, NM, Raton Municipal/Crews Field, NDB RWY 2, Amdt 4
Battle Mountain, NV, Battle Mountain, VOR/DME or GPS RWY 3, Amdt 4 Cancelled
Battle Mountain, NV, Battle Mountain, VOR/DME RWY 3, Amdt 4
Johnstown, NY, Fulton County, NDB or GPS RWY 10, Amdt 1 Cancelled
Johnstown, NY, Fulton County, NDB RWY 10, Amdt 1
Olean, NY, Cattaraugus County-Olean, RNAV or GPS RWY 22, Amdt 4A Cancelled
Olean, NY, Cattaraugus County-Olean, RNAV RWY 22, Amdt 4A
Schenectady, NY, Schenectady County, NDB or GPS RWY 22, Amdt 14 Cancelled
Schenectady, NY, Schenectady County, NDB RWY 22, Amdt 14
Schenectady, NY, Schenectady County, NDB or GPS RWY 28, Amdt 9 Cancelled
Schenectady, NY, Schenectady County, NDB RWY 28, Amdt 9
Delaware, OH, Delaware Muni, VOR or GPS RWY 28, Amdt 5 Cancelled
Delaware, OH, Delaware Muni, VOR RWY 28, Amdt 5
Delaware, OH, Delaware Muni, NDB or GPS RWY 10, Amdt 4 Cancelled
Delaware, OH, Delaware Muni, NDB RWY 10, Amdt 4
Marysville, OH, Union County, NDB or GPS RWY 27, Amdt 5 Cancelled
Marysville, OH, Union County, NDB RWY 27, Amdt 5
Duncan, OK, Halliburton Field, VOR or GPS RWY 35, Amdt 10 Cancelled
Duncan, OK, Halliburton Field, VOR RWY 35, Amdt 10
Idabel, OK, Idabel, NDB or GPS RWY 17, Amdt 3 Cancelled
Idabel, OK, Idabel, NDB RWY 17, Amdt 3
Clearfield, PA, Clearfield-Lawrence, VOR or GPS RWY 30, Amdt 4 Cancelled
Clearfield, PA, Clearfield-Lawrence, VOR RWY 30, Amdt 4
Lafayette, TN, Lafayette Muni, NDB or GPS RWY 19, Amdt 2B Cancelled
Lafayette, TN, Lafayette Muni, NDB RWY 19, Amdt 2B
Richmond/Ashland, VA, Hanover County Muni, NDB or GPS RWY 16, Orig Cancelled
Richmond/Ashland, VA, Hanover County Muni, NDB RWY 16, Orig
Suffolk, VA, Suffolk Muni, NDB or GPS RWY 7, Amdt. 1B Cancelled
Suffolk, VA, Suffolk Muni, NDB RWY 7, Amdt. 1B
Everett, WA, Snohomish County (Paine Fld), NDB or GPS RWY 16R, Amdt 12A Cancelled
Everett, WA, Snohomish County (Paine Fld), NDB RWY 16R, Amdt 12A
Manitowish Waters, WI, Manitowish Waters, NDB or GPS RWY 32, Orig Cancelled
Manitowish Waters, WI, Manitowish Waters, NDB RWY 32, Orig

[FR Doc. 97–16528 Filed 6–23–97; 8:45 am]
BILLING CODE 4910–13–M

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 175 and 178

[Docket No. 96F–0292]

Indirect Food Additives: Adhesives and Components of Coatings; and Adjuvants, Production Aids, and Sanitizers

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the food additive regulations to provide for the safe use of polyethyleneglycol alkyl(C_{10}–C_{12}) ether sulfosuccinate, disodium salt as a component of adhesives and as an emulsifier and/or surface-active agent in the manufacture of articles or components of articles intended for use in contact with food. This action is in response to a petition filed by Cytec Industries, Inc.

DATES: Effective June 24, 1997; written objections and requests for a hearing by July 24, 1997.

ADDRESSES: Submit written objections to the Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857.

FOR FURTHER INFORMATION CONTACT: Vir D. Anand, Center for Food Safety and Applied Nutrition (HFS–216), Food and Drug Administration, 200 C St. SW., Washington, DC 20204, 202–418–3081.

SUPPLEMENTARY INFORMATION: In a notice published in the **Federal Register** of August 26, 1996 (61 FR 43771), FDA

announced that a food additive petition (FAP 6B4518) had been filed by Cytec, Industries Inc., c/o Keller and Heckman, 1001 G St. NW., suite 500 West, Washington, DC 20001. The petition proposed to amend the food additive regulations in § 175.105 *Adhesives* (21 CFR 175.105) and § 178.3400 *Emulsifiers and/or surface-active agents* (21 CFR 178.3400) to provide for the safe use of polyethyleneglycol alkyl(C_{10}–C_{12}) ether sulfosuccinate, disodium salt as a component of adhesives and as an emulsifier and/or surface-active agent in the manufacture of articles or components of articles intended for use in contact with food.

Previously, the subject additive was listed in §§ 175.105 (43 FR 16311, April 18, 1978) and 178.3400 (58 FR 26684, May 5, 1993) with an alternative name (sulfosuccinic acid 4-ester with polyethylene glycol dodecyl ether, disodium salt with a corresponding CAS Reg. No. of 39354–45–5). Subsequently, the petitioner found that the ethoxylated alcohol used to synthesize the polyethylene glycol dodecyl ether portion of the additive contained not a single C_{12} species, but was a mixture consisting predominantly of C_{10}–C_{12} alcohols. This final rule, in effect, lists the correct chemical description and the corresponding CAS Reg. No. of the additive.

All of the chemistry and safety data presented in the earlier petitions (FAP's 8B3350 and 9B4120) resulting in the regulations cited above are incorporated in the present petition and remain unchanged. There are no compositional changes to the additive, its method of manufacture, use level, or technical effect. Further, there is no change in the safety evaluation. (For a full discussion of the safety evaluation, see 43 FR 16311 and 58 FR 26684.)

FDA has evaluated data in the petition and other relevant material. The agency concludes that the proposed use of the additive in adhesives and as an emulsifier and/or surface-active agent is safe, that the additive will have the intended technical effect, and, therefore, that §§ 175.105 and 178.3400 should be amended as set forth below.

In accordance with § 171.1(h) (21 CFR 171.1(h)), the petition and the documents that FDA considered and relied upon in reaching its decision to approve the petition are available for inspection at the Center for Food Safety and Applied Nutrition by appointment with the information contact person listed above. As provided in § 171.1(h), the agency will delete from the documents any materials that are not available for public disclosure before making the documents available for inspection.

The agency 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. The agency'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. No comments were received during the 30-day comment period specified in the filing notice for comments on the environment assessment submitted with the petition.

Any person who will be adversely affected by this regulation may at any time on or before July 24, 1997, file with the Dockets Management Branch (address above) written objections thereto. Each objection shall be separately numbered, and each numbered objection shall specify with particularity the provisions of the regulation to which objection is made and the grounds for the objection. Each numbered objection on which a hearing is requested shall specifically so state. Failure to request a hearing for any particular objection shall constitute a waiver of the right to a hearing on that objection. Each numbered objection for which a hearing is requested shall include a detailed description and analysis of the specific factual information intended to be presented in support of the objection in the event that a hearing is held. Failure to include such a description and analysis for any particular objection shall constitute a waiver of the right to a hearing on the objection. Three copies of all documents shall be submitted and shall be identified with the docket number found in brackets in the heading of this document. Any objections received in response to the regulation may be seen in the Dockets Management Branch between 9 a.m. and 4 p.m., Monday through Friday.

List of Subjects

21 CFR Part 175

Adhesives, Food additives, Food packaging.

21 CFR Part 178

Food additives, Food packaging.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs and redelegated to the Director, Center for Food Safety and Applied Nutrition, 21 CFR parts 175 and 178 are amended to read as follows:

PART 175—INDIRECT FOOD ADDITIVES: ADHESIVES AND COMPONENTS OF COATINGS

1. The authority citation for 21 CFR part 175 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 175.105 is amended in the table in paragraph (c)(5) by alphabetically adding a new entry under the heading ''Substances'' to read as follows:

§ 175.105 Adhesives.

* * * * *

(c) * * *

(5) * * *

Substances	Limitations
* * * * * * *	
Polyethyleneglycol alkyl(C_{10}–C_{12}) ether sulfosuccinate, disodium salt (CAS Reg. No. 68954–91–6).	
* * * * * * *	

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.
* * *	* * * *

* * * * *

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*