Substances	Limitations
* * * *	* * *
Periodic acid (CAS Reg. No. 10450–60–9).	
Polyethylenimine reaction product with 1,2-dichloroethane (CAS Reg.No. 68130–97–2) is the reaction product of homopolymerization of ethylenimine in aqueous hydrochloric acid at 100 °C and of cross-linking with 1,2-dichloroethane. The finished polymer has an average molecular weight of 50,000 to 70,000 as determined by gel permeation chromatography. The analytical method is entitled ''Methodology for Molecular Weight Detection of Polyethylenimine,'' which is incorporated by reference in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Copies may be obtained from the Division of Petition Control, Center for Food Safety and Applied Nutrition (HFS–215), 200 C St. SW., Washington, DC 20204, and may be examined at the Center for Food Safety and Applied Nutrition's Library, 200 C St. SW., rm. 3321, Washington, DC, or at the Office of the Federal Register, 800 North Capitol St. NW., suite 700, Washington, DC.	May be used as a fixing material in the immobilization of glucoamylase enzyme preparations from *Aspergillus niger* for use in the manufacture of beer. May be used as a fixing material in the immobilization of: 1. Glucose isomerase enzyme preparations for use in the manufacture of high fructose corn syrup, in accordance with § 184.1372 of this chapter. 2. Glucoamylase enzyme preparations from *Aspergillus niger* for use in the manufacture of beer. Residual ethylenimine in the finished polyethylenimine polymer will be less than 1 part per million as determined by gas chromatography-mass spectrometry. The residual ethylenimine is determined by an analytical method entitled ''Methodology for Ethylenimine Detection in Polyethylenimine,'' which is incorporated by reference in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Residual 1,2-dichloroethane in the finished polyethylenimine polymer will be less than 1 part per million as determined by gas chromatography. The residual 1,2-dichloroethane is determined by an analytical method entitled, ''Methodology for Ethylenedichloride Detection in Polyethylenimine,'' which is incorporated by reference in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Copies may be obtained from the Division of Petition Control, Center for Food Safety and Applied Nutrition (HFS–215), 200 C St. SW., Washington, DC 20204, or may be examined at the Center for Food Safety and Applied Nutrition's Library, 200 C St. SW., rm. 3321, Washington, DC, or the Office of the Federal Register, 800 North Capitol St. NW., suite 700, Washington, DC.

* * * * *

Dated: January 17, 1996.

William K. Hubbard,

Associate Commissioner for Policy Coordination.

[FR Doc. 96–2747 Filed 2–8–96; 8:45 am]

BILLING CODE 4160–01–F

21 CFR Parts 520, 522, and 558

New Animal Drugs; Change of Sponsor

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor name for SmithKline Animal Health Products, Division of SmithKline Beckman Corp. to SmithKline Beecham Animal Health due to a merger with Beecham Laboratories, Division of Beecham, Inc., and to reflect a change of sponsor for approved new drug applications (NADA's) previously held by SmithKline Beecham Animal Health to Pfizer, Inc.

EFFECTIVE DATE: February 9, 1996.

FOR FURTHER INFORMATION CONTACT: Judith M. O'Haro, Center for Veterinary Medicine (HFV–238), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1737.

SUPPLEMENTARY INFORMATION: In the Federal Register of October 8, 1991 (56 FR 50652), the animal drug regulations were amended to reflect the change of sponsor name for SmithKline Animal Health Products, Division of SmithKline Beckman Corp. to SmithKline Beecham Animal Health due to a merger with Beecham Laboratories, Division of Beecham, Inc. The regulations were amended to reflect the change of sponsor for 28 new animal drug applications (NADA's) from Beecham Laboratories, Division of Beecham Inc., to SmithKline Beecham Animal Health, and the change of sponsor for 22 NADA's from Norden Laboratories, Inc., to SmithKline Beecham Animal Health also due to the merger. The new company was assigned a new sponsor labeler code. The amended regulations did not reflect SmithKline Beecham Animal Health as the new sponsor in §§ 558.58, 558.311, 558.355, and 558.625. The sponsor currently listed for these products is Pfizer, Inc. Accordingly, the agency is amending these sections to reflect the change of sponsor.

In the Federal Register of November 2, 1995 (60 FR 55657), FDA published a document that amended the animal drug regulations to reflect a change of sponsor for 62 NADA's from SmithKline Beecham Animal Health to Pfizer, Inc. FDA inadvertently amended the regulations in 21 CFR 520.2260a, 520.2260b, and 520.2260c to reflect Pfizer, Inc. as the sponsor. However, Solvay Animal Health remains the sponsor of these sulfamethazine containing applications. The codified sections that should have been amended are 520.2220a, 520.2220b, 520.2220c, 520.2220d, and 522.2220. In addition, the agency omitted an amendment to 21 CFR 520.45a(a)(2). Accordingly, the agency is amending these sections to reflect this change of sponsor.

List of Subjects

21 CFR Parts 520 and 522

Animal drugs.

21 CFR Part 558

Animal drugs, Animal feeds.

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

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 520, 522, and 558 are amended as follows:

1. The authority citation for 21 CFR part 520 continues to read as follows:

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

§ 520.45a [Amended]

2. Section 520.45a *Albendazole suspension* is amended in paragraph (a)(2) by removing ''053571'' and adding in its place ''000069''.

§ 520.2220a [Amended]

3. Section 520.2220a *Sulfadimethoxine oral solution and soluble powder* is amended in paragraph (b) by removing ''053571'' and adding in its place ''000069''.

§ 520.2220b [Amended]

4. Section 520.2220b *Sulfadimethoxine tablets and boluses* is amended in paragraph (b)(1) by removing ''053571'' and adding in its place ''000069''.

§ 520.2220c [Amended]

5. Section 520.2220c *Sulfadimethoxine oral suspension* is amended in paragraph (c) by removing ''053571'' and adding in its place ''000069''.

§ 520.2220d [Amended]

6. Section 520.2220d *Sulfadimethoxine-ormetoprim tablets* is amended in paragraph (b) by removing ''053571'' and adding in its place ''000069''.

§ 520.2260a [Amended]

7. Section 520.2260a *Sulfamethazine oblets and boluses* is amended in paragraph (b)(1) by removing ''000069'' and adding in its place ''053501''.

§ 520.2260b [Amended]

8. Section 520.2260b *Sulfamethazine sustained-release boluses* is amended in paragraph (b)(1) by removing ''000069'' and adding in its place ''053501''.

§ 520.2260c [Amended]

9. Section 520.2260c *Sulfamethazine sustained-release tablets* is amended in paragraph (a) by removing ''000069'' and adding in its place ''053501''.

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

10. The authority citation for 21 CFR part 522 continues to read as follows:

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

§ 522.2220 [Amended]

11. Section 522.2220 *Sulfadimethoxine injection* is amended in paragraph (a)(2)(i) by removing ''053571'' and adding in its place ''000069''.

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

12. 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).

§ 558.58 [Amended]

13. Section 558.58 *Amprolium and ethopabate* is amended in the table in paragraph (d)(1), in item (iii), in the entry for virginamycin,15, under the ''Limitations'' and the ''Sponsor'' columns by removing ''000007'' and adding in its place ''000069''.

§ 558.311 [Amended]

14. Section 558.311 *Lasalocid* is amended in paragraph (b)(2) by removing ''000007'' and adding in its place ''000069''.

§ 558.355 [Amended]

15. Section 558.355 *Monensin* is amended in paragraph (b)(5) by removing ''000007'' and adding in its place ''000069''.

§ 558.625 [Amended]

16. Section 558.625 *Tylosin* is amended in paragraph (b)(25) by removing ''000007'' and adding in its place ''000069''.

Dated: February 1, 1996.

Robert C. Livingston

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

[FR Doc. 96–2846 Filed 2–8–96; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

Office of the Secretary

24 CFR Part 86

[Docket No. FR–4009–F–01]

Elimination of Requirements Governing the Lobbying of HUD Personnel; Removal of 24 CFR Part 86

AGENCY: Office of the Secretary, HUD.

ACTION: Final rule.

SUMMARY: This final rule removes part 86 from title 24 of the Code of Federal Regulations. Part 86, which was promulgated to comply with section 13 of the Department of Housing and Urban Development Act (42 U.S.C. 3537b), established recordkeeping, reporting, and registration requirements governing attempts to influence HUD programs. It also placed limitations on the fees paid to consultants who are engaged to influence the award or allocation of the Department's financial assistance. Effective January 1, 1996, the Lobbying Disclosure Act of 1995 (Pub. L. 104–65, approved December 19, 1995) repealed the authority for part 86—section 13 of the Department of Housing and Urban Development Act. This rule conforms the Code of Federal Regulations to this repeal.

EFFECTIVE DATE: March 11, 1996.

FOR FURTHER INFORMATION CONTACT: Aaron Santa Anna, Assistant General Counsel, Ethics Law Division; Office of General Counsel; Room 2158; U.S. Department of Housing and Urban Development; 451 Seventh Street, SW., Washington, DC 20410–0500; telephone (202) 708–0836. Hearing or speech-impaired individuals may call HUD's TDD number (202) 708–0113, or 1–800–877–8399 (Federal Information Relay Service TDD). (Other than the ''800'' number, these are not toll-free numbers.)

SUPPLEMENTARY INFORMATION: Section 112 of the Department of Housing and Urban Development Reform Act of 1989 (Pub. L. 101–235, approved December 15, 1989) added a new section 13 to the Department of Housing and Urban Development Act, 42 U.S.C. 3531, *et seq*. Section 13 contained two principal features. The first established the standards under which:

—Persons that make expenditures to influence a HUD officer or employee in the award of financial assistance or the taking of a management action by the Department must keep records, and report to HUD, on the expenditures; and