rate reports, entitled ''Instruction Manual for Electronic Filing of the Index of Customers,'' and ''Instruction Manual for Electronic Filing of the Discount Transportation Rate Report,'' respectively, and attached to this notice as Appendices A and B, are hereby adopted. The first electronic filing of the Index of Customers under sections 284.106(c) and 284.223(b) will be April 1, 1996. April 1 is one of the four scheduled filing dates provided for in the referenced regulations. The first discount rate reports to be filed electronically will be the reports due for the month of March 1996. Those reports are due within 15 days of the close of the March billing period.

By direction of the Commission.

Lois D. Cashell,

Secretary.

[FR Doc. 96–5166 Filed 3–5–96; 8:45 am]

BILLING CODE 6717–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510, 520, 522, and 524

Animal Drugs, Feeds, and Related Products; 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 the change of sponsor for 28 approved new animal drug applications (NADA's) from Coopers Animal Health, Inc., to Mallinckrodt Veterinary, Inc.

EFFECTIVE DATE: March 6, 1996.

FOR FURTHER INFORMATION CONTACT: Thomas J. McKay, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0213.

SUPPLEMENTARY INFORMATION: Coopers Animal Health, Inc., 1201 Douglas Ave., Kansas City, KS 66103–1438, has informed FDA that it has transferred the ownership of, and all rights and interests in, the following approved NADA's to Mallinckrodt Veterinary, Inc., Mundelein, IL 60060.

NADA No.	Trade name	Active ingredient
6–602	A–H Tablets 25 milligrams (mg)/100 mg.	Doxylamine Succinate
6–983	A–H Injection	Doxylamine Succinate, Chlorobutanol
10–987	Butazolidin Tablets/Bolus.	Phenylbutazone
11–222	Diquel Tablets	Ethylisobutrazine Hydrochloride
11–575	Butazolidin Injection 20%.	Phenylbutazone
11–877	Jenotone Tablets.	Aminopromazine Fumarate
11–893	Dermathycin Injection.	Thyroid Stimulating Hormone
15–182	Canopar Tablets.	Thenium Closylate
13–181	Jenomycin Tablets.	Aminopromazine Fumarate, Neomycin Sulfate
34–477	Jenotone Solution.	Aminopromazine Fumarate
35–016	Scolaban 400	Bunamidine Hydrochloride
35–265	Diquel Solution.	Ethylisobutrazine Hydrochloride
38–800	Butazolidin Granules.	Phenylbutazone
44–757	Prolate I–E	Phosmet
48–913	Halox Wormer Drench.	Haloxon
65–476	Cortisporin Veterinary Ophthalmic Ointment.	Bactricin ZN, Neomycin Sulfate, Polymyxin B Sulfate, Hydrocortisone Acetate
65–485	Neosporin Ophthalmic Ointment.	Bactricin ZN, Neomycin Sulfate, Polymyxin B Sulfate
92–483	Halox Bolus ..	Haloxon
95–614	Tribrissen 30/120/480/960 Tablets.	Sulfadiazine, Trimethoprim
97–288	Imizol Equine Injection.	Imidocarb Dipropionate
101–161	Thenatol PW Tablets.	Thenium Closylate, Piperazine Phosphate
105–093	Tribrissen 24% Injection.	Trimethoprim, Sulfadiazine Sodium
106–965	Tribrissen 48% Injection.	Trimethoprim, Sulfadiazine
116–087	Burazolidin Paste/Butazolidin/Phenylzone/Bute.	Phenylbutazone
120–326	Filban Chewable Wafers.	Diethylcarbamazine Citrate
124–842	Filban Tablets	Diethylcarbamazine Citrate
131–918	Tribrissen 400 Oral Paste.	Trimethoprim, Sulfadiazine
136–741	Tribrissen 60 Oral Suspension.	Trimethoprim, Sulfadiazine

The agency is amending 21 CFR 510.600(c)(1) and (c)(2), and parts 520, 522, and 524 to reflect the change of sponsor.

List of Subjects

21 CFR Part 510

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

21 CFR Parts 520, 522, and 524

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 510, 520, 522, and 524 are amended as follows:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: Secs. 201, 301, 501, 502, 503, 512, 701, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e).

§ 510.600 [Amended]

2. Section 510.600 is amended in the table in paragraph (c)(1) by removing the entry for ''Coopers Animal Health, Inc.''; and in the table in paragraph (c)(2) by removing the entry for ''017220''.

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

3. 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) .

§ 520.82a [Amended]

4. Section 520.82a *Aminopropazine fumarate tablets* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.82b [Amended]

5. Section 520.82b *Aminopropazine fumarate, neomycin sulfate tablets* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.222 [Amended]

6. Section 520.222 *Bunamidine hydrochloride* is amended in paragraph (c) by removing ''017220'' and adding in its place ''011716''.

§ 520.622c [Amended]

7. Section 520.622c *Diethylcarbamazine citrate chewable tablets* is amended in paragraph (b)(5) by removing ''017220'' and adding in its place ''011716''.

§ 520.784 [Amended]

8. Section 520.784 *Doxylamine succinate tablets* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.863 [Amended]

9. Section 520.863 *Ethylisobutrazine hydrochloride tablets* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.1120a [Amended]

10. Section 520.1120a *Haloxon drench* is amended in paragraph (c) by removing ''017220'' and adding in its place ''011716''.

§ 520.1120b [Amended]

11. Section 520.1120b *Haloxon boluses* is amended in paragraph (c) by removing ''017220'' and adding in its place ''011716''.

§ 520.1720a [Amended]

12. Section 520.1720a *Phenylbutazone tablets and boluses* is amended in paragraph (b)(1) by removing ''017220'' and adding in its place ''011716''.

§ 520.1720b [Amended]

13. Section 520.1720b *Phenylbutazone granules* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.1720c [Amended]

14. Section 520.1720c *Phenylbutazone paste* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.1805 [Amended]

15. Section 520.1805 *Piperazine phosphate with thenium closylate tablets* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.2362 [Amended]

16. Section 520.2362 *Thenium closylate tablets* is amended in paragraph (c) by removing ''017220'' and adding in its place ''011716''.

§ 520.2610 [Amended]

17. Section 520.2610 *Trimethoprim and sulfadiazine tablets* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.2611 [Amended]

18. Section 520.2611 *Trimethoprim and sulfadiazine oral paste* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 520.2612 [Amended]

19. Section 520.2612 *Trimethoprim and sulfadiazine oral suspension* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

20. The authority citation for 21 CFR Part 522 continues to read as follows:

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

§ 522.82 [Amended]

21. Section 522.82 *Aminopropazine fumarate sterile solution injection* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 522.784 [Amended]

22. Section 522.784 *Doxylamine succinate injection* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 522.863 [Amended]

23. Section 522.863 *Ethylisobutrazine hydrochloride injection* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 522.1155 [Amended]

24. Section 522.1155 *Imidocarb dipropionate sterile powder* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

§ 522.1720 [Amended]

25. Section 522.1720 *Phenylbutazone injection* is amended in paragraph (b)(1) by removing ''017220'' and adding in its place ''011716''.

§ 522.2610 [Amended]

26. Section 522.2610 *Trimethoprim and sulfadiazine sterile suspension* is amended in paragraphs (a)(2) and (b)(2) by removing ''017220'' and adding in its place ''011716''.

PART 524—OPHTHALMIC AND TOPICAL DOSAGE FORM NEW ANIMAL DRUGS

27. 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).

§ 524.154 [Amended]

28. Section 524.154 *Bacitracin or bacitracin zinc-neomycin sulfate-polymyxin B sulfate ophthalmic ointment* is amended in paragraph (a)(2) by removing ''017220'' and adding in its place ''011716''.

§ 524.155 [Amended]

29. Section 524.155 *Bacitracin zinc-polymyxin B sulfate neomycin sulfate-hydrocortisone or hydrocortisone acetate ophthalmic ointment* is amended in paragraph (a)(1) by removing ''017220'' and adding in its place ''011716''.

§ 524.1742 [Amended]

30. Section 524.1742 *N-(Mercaptomethyl) phthalimide S-(O,O-dimethyl phosphorodithioate) emulsifiable liquid* is amended in paragraph (b) by removing ''017220'' and adding in its place ''011716''.

Dated: February 28, 1996.

Robert C. Livingston,

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

[FR Doc. 96–5213 Filed 3–5–96; 8:45 am]

BILLING CODE 4160–01–F

ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[KY–71–2–6062a; FRL–5427–4]

Approval and Promulgation of Implementation Plans—Kentucky: Approval of Revision To The State Implementation Plan

AGENCY: Environmental Protection Agency (EPA).

ACTION: Direct final rule.

SUMMARY: This action approves a revision to the Kentucky State Implementation Plan (SIP) adopted by the Kentucky Natural Resources and Environmental Protection Cabinet (KNREP) on March 4, 1993, for the