adequate controlled airspace to accommodate a GPS SIAP to RWY 20 at Nut Tree Airport, Vacaville, CA.

Interested parties were invited to participate in this rulemaking proceeding by submitting written comments on the proposal to the FAA. No comments to the proposal were received. Class E airspace designations are published in paragraph 6005 of FAA Order 7400.9C dated August 17, 1995, and effective September 16, 1995, which is incorporated by reference in 14 CFR 71.1. The E airspace designation listed in this document will be published subsequently in this Order.

The Rule

This amendment to part 71 of the Federal Aviation Regulations (14 CFR part 71) amends the Class E airspace area at Vacaville, CA. The development of a GPS SIAP to RWY 20 has made this action necessary. The intended effect of this action is to provide adequate controlled airspace for aircraft executing the GPS RWY 20 SIAP at Nut Tree Airport, Vacaville, CA.

The FAA has determined that this regulation only involves an established body of technical regulations for which frequent and routine amendments are necessary to keep them operationally current. Therefore, this regulation—(1) is not a ''significant regulatory action'' under Executive Order 12866; (2) is not a ''significant rule'' under DOT Regulatory Policies and Procedures (44 FR 10034; February 26, 1979); and (3) does not warrant preparation of a Regulatory Evaluation as the anticipated impact is so minimal. Since this is a routine matter that will only affect air traffic procedures and air navigation, it is certified that this rule will not have a significant economic impact on a substantial number of small entities under the criteria of the Regulatory Flexibility Act.

List of Subjects in 14 CFR Part 71

Airspace, Incorporation by reference, Navigation (air).

Adoption of the Amendment

In consideration of the foregoing, the Federal Aviation Administration amends 14 CFR part 71 as follows:

PART 71—[AMENDED]

1. The authority citation for 14 CFR part 71 continues to read as follows:

Authority: 49 U.S.C. 106(g), 40103, 40113, 40120; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389; 14 CFR 11.69.

§ 71.1 [Amended]

2. The incorporation by reference in 14 CFR 71.1 of the Federal Aviation Administration Order 7400.9C, Airspace Designations and Reporting Points, dated August 17, 1995, and effective September 16, 1995, is amended as follows:

Paragraph 6005 Class E airspace areas extending upward from 700 feet or more above the surface of the earth.

* * * * *

AWP CA E5 Vacaville, CA [Revised]

Nut Tree Airport, CA
(Lat. 38°22′37″ N, long. 121°57′45″ W)
Sacramento VORTAC
(Lat. 38°38′26″ N, long. 121°33′06″ W)

That airspace extending upward from 700 feet above the surface within a 5-mile radius of Nut Tree Airport and within 2.2 miles each side of the Sacramento VORTAC 259° radial, extending from the 5-mile radius to 11.3 miles west of the VORTAC and within 2.6 miles each side of the 034° bearing from the Nut Tree Airport, extending from the 5-mile radius to 10.5 miles northeast of the airport.

* * * * *

Issued in Los Angeles, California, on January 30, 1996.

James H. Snow,

Acting Management, Air Traffic Division, Western-Pacific Region.

[FR Doc. 96–3175 Filed 2–12–96; 8:45 am]

BILLING CODE 4910–13–M

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 a change of sponsor for 35 approved new animal drug applications (NADA's) from Syntex Animal Health, Division of Syntex Agri-business, Inc., to Fort Dodge Laboratories, Division of American Home Products.

EFFECTIVE DATE: February 13, 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: Syntex Animal Health, Division of Syntex Agri-business, Inc., 3401 Hillview Ave., Palo Alto, CA 94303, has informed FDA that it has transferred the ownership of, and all rights and interests in, the following approved NADA's to Fort Dodge Laboratories, Division of American Home Products Corp., 800 Fifth St. NW., Fort Dodge, IA 50501:

NADA No.	Drug Name
9–576	Synovex® S and Synovex® C Implants (progesterone and estradiol benzoate)
11–427	Synovex® H Implants (testosterone propionate and estradiol benzoate)
15–126	Spectinomycin Injectable & Tablets (spectinomycin)
30–414	Flucort® (V Solution (flumethasone)
30–415	Flucort® Tablets (flumethasone)
32–168	DOMOSO® Solution (dimethyl sulfoxide)
36–211	Anaprime® Suspension (flumethasone)
36–212	Fluosmin® Suspension (flumethasone acetate)
37–586	Erythromast® '36' Solution (erythromycin)
38–801	Anaprime® Ophthalmic Solution (flumethasone)
41–629	Spectinomycin® Oral Solution and Spectogard® Oral Solution (spectinomycin)
41–665	Tranvet® Chewable Tablets (propiopromazine hydrochloride)
45–512	Synotic® Otic Solution (fluocinolone acetonide and dimethyl sulfoxide)
45–716	Tranvet® Injectable Solution (propiopromazine hydrochloride)
47–334	Synsac® Solution (flucinolone acetonide and dimethyl sulfoxide)
47–925	DOMOSO® Gel (dimethyl sulfoxide)
49–725	Anaprime® Opthakote® Ophthalmic Solution (flumethasone with neomycin sulfate and polymyxin B sulfate)

NADA No.	Drug Name
49–726	Optiprime® Opthakote® Ophthalmic Solution (neomycin sulfate and polymyxin B sulfate)
96–674	Equiproxen® Granules (naproxen)
96–675	Equiproxen® Injectable (naproxen)
100–254	Synchrocept® (prostalene)
110–776	Benzelmin® Powder for Suspension (oxfendazole)
110–777	Benzelmin® Top Dress Pellets (oxfendazole)
115–578	Di-Trim® Tablets (trimethoprim and sulfadiazine)
128–549	Bovilene® Sterile Solution (fenprostalene)
128–967	Repose® (sodium secobarbital and dibucaine hydrochloride)
132–105	Benzelmin® 37.5% Paste (oxfendazole)
132–486	Di-Trim® 24% Injection (trimethoprim and sulfadiazine)
133–841	Benzelmin® 9.06% Suspension (oxfendazole)
134–778	Di-Trim® 48% Injection (trimethoprim and sulfadiazine)
136–342	Di-Trim® 400 Oral Paste (trimethoprim and sulfadiazine)
136–740	Benzelmin® Plus Paste (oxfendazole plus trichlorfon)
138–903	Porcilene® Sterile Solution (fenprostalene)
140–854	Synanthic® 9.06% and 22.5% Suspension(oxfendazole)
140–892	Synanthic® 18.5% Paste (oxfendazole)

Accordingly, the agency is amending 21 CFR parts 510, 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 *Names, addresses, and drug labeler codes of sponsors of approved applications* is amended in the table in paragraph (c)(1) by revising the entry for ''Fort Dodge Laboratories'' to read ''Fort Dodge Laboratories, Division of American Home Products Corp., 800 Fifth St. NW., Fort Dodge, IA 50501''; and in the table in paragraph (c)(2) in the entry for ''000856'' by revising the sponsor name and address to read, ''Fort Dodge Laboratories, Division of American Home Products, 800 Fifth St. NW., Fort Dodge, IA 50501.''

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) .

§ 520.960 [Amended]

4. Section 520.960 *Flumethasone tablets* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 520.1468 [Amended]

5. Section 520.1468 *Naproxen granules* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 520.1628 [Amended]

6. Section 520.1628 *Oxfendazole powder and pellets* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 520.1629 [Amended]

7. Section 520.1629 *Oxfendazole paste* is amended in paragraph (a)(2) and (b)(2) by removing ''000033'' and adding in its place ''000856''.

§ 520.1630 [Amended]

8. Section 520.1630 *Oxfendazole suspension* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 520.1631 [Amended]

9. Section 520.1631 *Oxfendazole and trichlorfon paste* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 520.2002 [Amended]

10. Section 520.2002 *Propiopromazine hydrochloride* is amended in paragraph (c) by removing ''000033'' and adding in its place ''000856''.

§ 520.2122 [Amended]

11. Section 520.2122 *Spectinomycin dihydrochloride oral solution* is amended in paragraph (b)(2) by removing ''000033'' and adding in its place ''000856''.

§ 520.2610 [Amended]

12. Section 520.2610 *Trimethoprim and sulfadiazine tablets* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 520.2611 [Amended]

13. Section 520.2611 *Trimethoprim and sulfadiazine oral paste* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

14. 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).

§ 522.842 [Amended]

15. Section 522.842 *Estradiol benzoate and testosterone propionate in combination* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 522.914 [Amended]

16. Section 522.914 *Fenprostalene solution* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 522.960a [Amended]

17. Section 522.960a *Flumethasone suspension* is amended in paragraph (c)

by removing ''000033'' and adding in its place ''000856''.

§ 522.960b [Amended]

18. Section 522.960b *Flumethasone acetate injection* is amended in paragraph (c) by removing ''000033'' and adding in its place ''000856''.

§ 522.960c [Amended]

19. Section 522.960c *Flumethasone solution* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 522.1468 [Amended]

20. Section 522.1468 *Naproxen for injection* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 522.1940 [Amended]

21. Section 522.1940 *Progesterone and estradiol benzoate in combination* is amended in paragraph (b) and (d)(1)(iii) by removing ''000033'' and adding in its place ''000856''.

§ 522.2002 [Amended]

22. Section 522.2002 *Propiopromazine hydrochloride injection* is amended in paragraph (c) by removing ''000033'' and adding in its place ''000856''.

§ 522.2012 [Amended]

23. Section 522.2012 *Prostalene solution* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 522.2120 [Amended]

24. Section 522.2120 *Spectinomycin injection* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 522.2610 [Amended]

25. Section 522.2610 *Trimethoprim and sulfadiazine sterile suspension* is amended in paragraph (a)(2) and (b)(2) by removing ''000033'' and adding in its place ''000856''.

PART 524—OPHTHALMIC AND TOPICAL DOSAGE FORM NEW ANIMAL DRUGS

26. 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).

§ 524.660a [Amended]

27. Section 524.660a *Dimethyl sulfoxide solution* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 524.660b [Amended]

28. Section 524.660b *Dimethyl sulfoxide gel* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 524.960 [Amended]

29. Section 524.960 *Flumethasone, neomycin sulfate, and polymyxin B sulfate ophthalmic solutions* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 524.981d [Amended]

30. Section 524.981d *Fluocinolone acetonide, dimethyl sulfoxide solution* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

§ 524.1484e [Amended]

31. Section 524.1484e *Neomycin sulfate and polymyxin B sulfate ophthalmic solution* is amended in paragraph (b) by removing ''000033'' and adding in its place ''000856''.

Dated: February 1, 1996.

Robert C. Livingston,

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

[FR Doc. 96–3078 Filed 2–12–96; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF LABOR

Occupational Safety and Health Administration

29 CFR Parts 1910, 1915, 1917, 1918, 1919, 1926, and 1928

Office of Management and Budget Control Numbers Under the Paperwork Reduction Act

AGENCY: Occupational Safety and Health Administration, Labor.

ACTION: Final rule.

SUMMARY: The Occupational Safety and Health Administration (OSHA) is adding new sections to its Safety and Health Regulations for General Industry, Construction and Shipyard Employment. These new sections will be used to consolidate and display all of the control numbers assigned by the Office of Management and Budget (OMB) for ''approved'' information collection requirements. OSHA is also identifying information collection requirements found in certain of its other regulations and displaying the OMB control number at the end of each section containing a collection of information. None of the requirements are new; they have been promulgated by OSHA at various times over the past 25 years. The display of OMB control numbers is required under the implementing rules and regulations of OMB and under the Paperwork Reduction Act of 1995.

EFFECTIVE DATE: February 13, 1996.

FOR FURTHER INFORMATION CONTACT: Ms. Anne Cyr, Office of Information and Consumer Affairs, U.S. Department of Labor, Occupational Safety and Health Administration, Room N3647; 200 Constitution Avenue, NW, Washington, DC 20210 (202–219–8148, FAX 202–219–5986).

SUPPLEMENTARY INFORMATION:

I. Background

OSHA has a number of provisions within its occupational safety and health standards that require employers to collect or prepare information. These types of provisions are broadly classified as ''information collection requirements.'' All information collection requirements are subject to review and approval by OMB on not more than a three-year cycle. It should be noted that OSHA cannot impose a penalty on employers for violating collection of information (recordkeeping, reporting, etc.) requirements if the agency has failed to obtain OMB approval of the requirement. When OMB approves collection of information requirements, it issues a ''control number'' for the collection of information provision. All agencies are required to display [show to the public] the OMB control numbers so the public will know that OMB has given the agency approval to require the information [report, record, documentation, form, etc.] to be collected. In the past, OSHA has displayed the OMB control number by printing it at the end of each section or subpart in which the requirement to collect information appeared. However, to enable the public easily and readily to identify all of the collection of information requirements, OSHA is dedicating one section in part 1910 (Safety and Health Standards for General Industry), one section in part 1915 (Shipyard Employment Standards), and one section in part 1926 (Construction Safety and Health Standards) to list those requirements and show the OMB control number. As a result of this new format, the parenthetical notes and approval/control numbers now printed at the end of the individual sections or subparts can be removed. This procedure is not being used for collections of information in 29 CFR Parts 1917, 1918, 1919, and 1928 because there are only one, two, or three collection of information