new address is 5100 Paint Branch Pkwy., College Park, MD 20740.

Publication of this document constitutes final action on these changes under the Administrative Procedure Act (5 U.S.C. 553). Notice and public procedure are unnecessary because FDA is merely correcting nonsubstantive errors.

Therefore, under the Federal Food, Drug, and Cosmetic Act and under authority delegated to the Commissioner of Food and Drugs, 21 CFR chapter I is amended as follows:

1. Parts 1, 5, 10, 70, 71, 73, 80, 100, 101, 102, 106, 107, 108, 109, 110, 130, 161, 165, 170, 172, 173, 175, 176, 177, 178, 180, 181, 184, 189, 190, 211, 701, 1240, and 1250 are amended by removing ''200 C Street, SW., Washington, DC 20204'' or ''200 C St. SW., Washington, DC 20204'' wherever they appear and by adding in their place ''5100 Paint Branch Pkwy., College Park, MD 20740.''

Dated: October 31, 2001.

Margaret M. Dotzel,

Associate Commissioner for Policy.

[FR Doc. 01–27811 Filed 11–5–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 510

New Animal Drugs; Change of Sponsor's Name

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's name from Marsam Pharmaceuticals, Inc., to Marsam Pharmaceuticals, LLC.

DATES: This rule is effective November 6, 2001.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209, e-mail: lluther@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: Marsam Pharmaceuticals, Inc., Bldg. 31, 24 Olney Ave., Cherry Hill, NJ 08034, has informed FDA of a change of sponsor's name to Marsam Pharmaceuticals, LLC. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) to reflect the change of sponsor's name.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801-808.

List of Subjects in 21 CFR Part 510

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

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 part 510 is amended as follows:

PART 510—NEW ANIMAL DRUGS

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

Authority: 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) in the entry for ''Marsam Pharmaceuticals, Inc.'' and in the table in paragraph (c)(2) in the entry for ''000209'' by removing ''Inc.'' and by adding in its place ''LLC''.

Dated: October 26, 2001.

Claire M. Lathers,

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

[FR Doc. 01–27813 Filed 11–5–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 522

Implantation or Injectable Dosage Form 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 for an approved new animal drug application (NADA) from Elanco Animal Health, A Division of Eli Lilly & Co., to Ivy Laboratories, Div. of Ivy Animal Health, Inc.

DATES: This rule is effective November 6, 2001.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209.

SUPPLEMENTARY INFORMATION: Elanco Animal Health, A Division of Eli Lilly & Co., Lilly Corporate Center, Indianapolis, IN 46285, has informed FDA that it has transferred to Ivy Laboratories, Div. of Ivy Animal Health, Inc., 8857 Bond St., Overland Park, KS 66214, ownership of, and all rights and interests in NADA 118–123 for COMPUDOSE 200 (estradiol) and COMPUDOSE 400 implants for cattle. Accordingly, the agency is amending the regulations in 21 CFR 522.840 to reflect the transfer of ownership.

This rule does not meet the definition of ''rule'' in 5 U.S.C. 804(3)(A) because it is a rule of ''particular applicability.'' Therefore, it is not subject to the congressional review requirements in 5 U.S.C. 801–808.

List of Subjects in 21 CFR Part 522

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 part 522 is amended as follows:

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

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

Authority: 21 U.S.C. 360b.

§ 522.840 [Amended]

2. Section 522.840 *Estradiol* is amended in paragraph (b) by removing ''000986'' and by adding in its place ''No. 021641''.

Dated: October 26, 2001.

Claire M. Lathers,

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

[FR Doc. 01–27812 Filed 11–5–01; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF TRANSPORTATION

Coast Guard

33 CFR Parts 100 and 165

[USCG–2001–10936]

Safety Zones, Security Zones, and Special Local Regulations

AGENCY: Coast Guard, DOT.

ACTION: Notice of temporary rules issued.