(i) *Official number.* A vessel for which a permit has been issued under § 622.4 must display its official number--

* * * * *

7. In § 622.31, paragraph (k) is added to read as follows:

§ 622.31 Prohibited gear and methods.

* * * * *

(k) *Traps for royal red shrimp in the Gulf EEZ and transfer at sea.* A trap may not be used to fish for royal red shrimp in the Gulf EEZ. Possession of a trap and royal red shrimp on board a vessel is prohibited. A trap used to fish for royal red shrimp in the Gulf EEZ may be disposed of in any appropriate manner by the Assistant Administrator or an authorized officer. In addition, royal red shrimp cannot be transferred in the Gulf EEZ, and royal red shrimp taken in the Gulf EEZ cannot be transferred at sea regardless of where the transfer takes place.

[FR Doc. 02–19977 Filed 8–6–02; 8:45 am]

BILLING CODE 3510–22–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 Blue Ridge Pharmaceuticals, Inc., to IDEXX Pharmaceuticals, Inc.

DATES: This rule is effective August 7, 2002.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–101), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209, e-mail: *lluther@cvm.fda.gov*.

SUPPLEMENTARY INFORMATION: Blue Ridge Pharmaceuticals, Inc., 4249–105 Piedmont Pkwy., Greensboro, NC 27410, has informed FDA of a change of name to IDEXX Pharmaceuticals, Inc. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) to reflect the change.

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.

2. Section 510.600 is amended in the table in paragraph (c)(1) by removing the entry for ''Blue Ridge Pharmaceuticals, Inc.'' and by alphabetically adding an entry for ''IDEXX Pharmaceuticals, Inc.''; and in the table in paragraph (c)(2) by revising the entry for ''065274'' to read as follows.

§ 510.600 Names, addresses, and drug labeler codes of sponsors of approved applications.

* * * * *

(c) * * *

(1) * * *

Firm name and address	Drug labeler code
* * * * *	* *
IDEXX Pharmaceuticals, Inc., 4249–105 Piedmont Pkwy., Greensboro, NC 27410	065274
* * * * *	* *

(2) * * *

Drug labeler code	Firm name and address
* * * * *	* *
065274	IDEXX Pharmaceuticals, Inc., 4249–105 Piedmont Pkwy., Greensboro, NC 27410
* * * * *	* *

Dated: July 19, 2002.

Andrew J. Beaulieu,

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

[FR Doc. 02–19906 Filed 8–6–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 and 529

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 DEC International, Inc., to Pharmacia & Upjohn Co.

DATES: This rule is effective August 7, 2002.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–101), Food and Drug Administration, 7500 Standish Pl.,

Rockville, MD 20855, 301–827–0209, e-mail: lluther@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: DEC International, Inc., 1919 South Stoughton Rd., P.O. Box 8050, Madison WI 53708–8050, has informed FDA that it has transferred ownership of, and all rights and interests in, NADA 141–200 for EAZI–BREED CIDR Progesterone Intravaginal Inserts to Pharmacia & Upjohn Co., 7000 Portage Rd., Kalamazoo, MI 49001–0199. Accordingly, the agency is amending the regulations in 21 CFR 529.1940 to reflect the transfer of ownership.

Following this change of sponsorship, DEC International, Inc., is no longer the sponsor of any approved application. Accordingly, 21 CFR 510.600(c) is being amended to remove the entries for DEC International, Inc.

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

21 CFR Part 510

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

21 CFR Part 529

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 and 529 are 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) by removing the entry for ''DEC International, Inc.'' and in the table in paragraph (c)(2) by removing the entry for ''067080''.

PART 529—CERTAIN OTHER DOSAGE FORM NEW ANIMAL DRUGS

3. The authority citation for 21 CFR part 529 continues to read as follows:

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

§ 529.1940 [Amended]

4. Section 529.1940 *Progesterone intravaginal inserts* is amended in paragraph (b) by removing ''067080'' and by adding in its place ''000009''.

Dated: July 17, 2002.

Alan Rudman,

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

[FR Doc. 02–19862 Filed 8–6–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 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 for an approved abbreviated new animal drug application (ANADA) from Equi Aid Products, Inc., to Farnam Companies, Inc.

DATES: This rule is effective August 7, 2002.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–101), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209, e-mail: lluther@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: Equi Aid Products, Inc., 1517 West Knudsen Dr., Phoenix, AZ 85027, has informed FDA that it has transferred ownership of, and all rights and interest in, ANADA 200–168 for CW 48 (pyrantel tartrate) Type A medicated article to Farnam Companies, Inc., 301 West Osborn, Phoenix, AZ 85013–3928. Accordingly, the agency is amending the regulations in § 558.485 (21 CFR 558.485) to reflect the transfer of ownership.

Following this change of sponsorship, Equi Aid Products, Inc., is no longer the sponsor of any approved application. Accordingly, § 510.600(c) is amended to remove the entries for Equi Aid Products, Inc.

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

21 CFR Part 510

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

21 CFR Part 558

Animal drugs, Animal feeds.

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 and 558 are 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) by removing the entry for ''Equi Aid Products, Inc.'' and in the table in paragraph (c)(2) by removing the entry for ''062240''.

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

3. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

§ 558.485 [Amended]

4. Section 558.485 *Pyrantel tartrate* is amended in paragraph (b)(7) by removing '', 017135, and 062240'' and by adding in its place ''and 017135''.

Dated: July 17, 2002.

Alan Rudman,

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

[FR Doc. 02–19861 Filed 8–6–02; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Oxytetracycline Hydrochloride Soluble Powder

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.