PART 246—SPECIAL SUPPLEMENTAL NUTRITION PROGRAM FOR WOMEN, INFANTS AND CHILDREN

■ 1. The authority citation for part 246 continues to read as follows:

Authority: 42 U.S.C. 1786.

■ 2. In § 246.26, a new paragraph (j) is added to read as follows:

§ 246.26 Other provisions.

* * * * *

(j) *Data collection related to local agencies.* (1) Each State agency must collect data related to local agencies that have an agreement with the State agency to participate in the program for each of Federal fiscal years 2006 through 2009, including those local agencies that participated only for part of the fiscal year. Such data shall include:

(i) The name of each local agency;

(ii) The city in which each local agency was headquartered and the name of the state;

(iii) The amount of funds provided to the participating organization, i.e., the amount of federal funds provided for nutrition services and administration to each participating local agency; and

(iv) The type of participating organization, e.g., government agency, educational institution, non-profit organization/secular, non-profit organization/faith-based, and ''other.''

(2) On or before August 31, 2007, and each subsequent year through 2010, State agencies must report to FNS data as specified in paragraph (j)(1) of this section for the prior Federal fiscal year. State agencies must submit this data in a format designated by FNS.

PART 247—COMMODITY SUPPLEMENTAL FOOD PROGRAM

■ 1. The authority citation for part 247 continues to read as follows:

Authority: Sec. 5, Pub. L. 93–86, 87 Stat. 249, as added by Sec. 1304(b)(2), Pub. L. 95–113, 91 Stat. 980 (7 U.S.C. 612c note); sec. 1335, Pub. L. 97–98, 95 Stat. 1293 (7 U.S.C. 612c note); sec. 209, Pub. L. 98–8, 97 Stat. 35 (7 U.S.C. 612c note); sec. 2(8), Pub. L. 98–92, 97 Stat. 611 (7 U.S.C. 612c note); sec. 1562, Pub. L. 99–198, 99 Stat. 1590 (7 U.S.C. 612c note); sec. 101(k), Pub. L. 100–202; sec. 1771(a), Pub. L. 101–624, 101 Stat. 3806 (7 U.S.C. 612c note); sec. 402(a), Pub. L. 104–127, 110 Stat. 1028 (7 U.S.C. 612c note); Pub. L. 107–171.

■ 2. In § 247.29, a new paragraph (d) is added to read as follows:

§ 247.29 Reports and recordkeeping.

* * * * *

(d) *What data must the State agency collect related to local agencies?* (1) Each State agency must collect data related to local agencies that have an agreement with the State agency to participate in the program for each of Federal fiscal years 2006 through 2009; including those local agencies that participated only for part of the fiscal year. Such data shall include:

(i) The name of each local agency;

(ii) The city in which each participating local agency was headquartered and the name of the state;

(iii) The amount of funds provided to the participating organization, i.e., the amount of federal administrative funds provided to each participating local agency; and

(iv) The type of participating organization, e.g., government agency, educational institution, non-profit organization/secular, non-profit organization/faith-based, and ''other.''

(2) On or before August 31, 2007, and each subsequent year through 2010, State agencies must report to FNS data as specified in paragraph (d)(1) of this section for the prior Federal fiscal year. State agencies must submit this data in a format designated by FNS.

PART 251—THE EMERGENCY FOOD ASSISTANCE PROGRAM

■ 1. The authority citation for part 251 continues to read as follows:

Authority: 7 U.S.C. 7501–7516.

■ 2. In § 251.10, a new paragraph (i) is added to read as follows:

§ 251.10 Miscellaneous provisions.

* * * * *

(i) *Data collection related to eligible recipient agencies.* (1) Each State agency must collect data related to eligible recipient agencies that have an agreement with the State agency to participate in the program for each of Federal fiscal years 2006 through 2009, including those eligible recipient agencies that participated only for part of the fiscal year. Such data shall include:

(i) The name of each eligible recipient agency;

(ii) The city in which each participating eligible recipient agency was headquartered and the name of the state;

(iii) The amount of funds provided to the participating organization, i.e., the sum of the amount of federal administrative funds plus the value of the commodities purchased under Section 214 of the Emergency Food Assistance Act of 1983 provided to each participating eligible recipient agency; and

(iv) The type of participating organization, e.g., government agency, educational institution, non-profit organization/secular, non-profit organization/faith-based, and ''other.''

(2) On or before August 31, 2007, and each subsequent year through 2010, State agencies must report to FNS data as specified in paragraph (i)(1) of this section for the prior Federal fiscal year. State agencies must submit this data in a format designated by FNS.

Dated: April 23, 2007.

Nancy Montanez Johner,

Under Secretary, Food, Nutrition, and Consumer Services.

[FR Doc. 07–2173 Filed 5–1–07; 8:45 am]

BILLING CODE 3410–30–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 510

New Animal Drugs; Change of Sponsor's Address

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 address for Alpharma, Inc.

DATES: This rule is effective May 2, 2007.

FOR FURTHER INFORMATION CONTACT: David R. Newkirk, Center for Veterinary Medicine (HFV–100), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–6967, e-mail: *david.newkirk@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Alpharma, Inc., One Executive Dr., Ft. Lee, NJ 07024, has informed FDA of a change of address to 440 Rte. 22, Bridgewater, NJ 08807. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c) 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. In § 510.600, in the table in paragraph (c)(1), revise the entry for ''Alpharma, Inc.''; and in the table in paragraph (c)(2), revise the entry for ''046573'' 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
* * * * *	
Alpharma, Inc., 440 Rte. 22, Bridgewater, NJ 08807.	046573
* * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * *	
046573	Alpharma, Inc., 440 Rte. 22, Bridgewater, NJ 08807
* * * * *	

Dated: April 24, 2007.

Bernadette Dunham,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. E7–8322 Filed 5–1–07; 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; Fenbendazole Paste

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to correct an inadvertent error in the conditions of use of fenbendazole paste in horses and cattle. This action is being taken to improve the accuracy of the animal drug regulations.

DATES: This rule is effective May 2, 2007.

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9019, e-mail: *george.haibel@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: FDA is amending the animal drug regulations in 21 CFR 558.95 to correct an inadvertent error in the conditions of use of fenbendazole paste in horses and cattle. The error in the agency's regulations was introduced in a final rule reflecting the approval of a supplemental new animal drug application that published in the **Federal Register** on March 9, 2007 (72 FR 10595). This action is being taken to improve the accuracy of the animal drug regulations.

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 520

Animal drugs.

■ Therefore, under the Federal Food, Drug, and Cosmetic Act and under the authority delegated to the Commissioner of Food and Drugs and redelegated to the Center for Veterinary Medicine, 21 CFR part 520 is amended as follows:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

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

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

■ 2. Section 520.905c is revised to read as follows:

§ 520.905c Fenbendazole paste.

(a) *Specifications.* Each gram of paste contains 100 milligrams (mg) fenbendazole (10 percent).

(b) *Sponsor.* See No. 057926 in § 510.600(c) of this chapter.

(c) *Related tolerances.* See § 556.275 of this chapter.

(d) *Special considerations.* See § 500.25 of this chapter.

(e) *Conditions of use*—(1) *Horses*—(i) *Indications for use and amounts*—(A) For control of large strongyles (*Strongylus edentatus*, *S. equinus*, *S. vulgaris*), small strongyles, pinworms (*Oxyuris equi*), and ascarids (*Parascaris equorum*): 2.3 mg per pound (/lb) of body weight, or for foals and weanlings (less than 18 months of age), 4.6 mg/lb of body weight. Retreatment at intervals of 6 to 8 weeks may be required.

(B) For control of arteritis caused by the fourth-stage larvae of S. vulgaris: 4.6 mg/lb of body weight daily for 5 days. Treatment should be initiated in the spring and repeated in 6 months.

(C) For treatment of encysted mucosal cyathostome (small strongyle) larvae including early third-stage (hypobiotic), late third-stage, and fourth-stage larvae: 4.6 mg/lb of body weight daily for 5 consecutive days.

(D) Fenbendazole paste 10 percent may be used concomitantly with approved forms of trichlorfon for the indications provided in paragraph (e)(1)(i)(A) of this section and for treating infections of stomach bots as provided in § 520.2520.

(ii) *Limitations.* Do not use in horses intended for human consumption.

(2) *Cattle*—(i) *Amount.* 2.3 mg/lb of body weight. Retreatment may be needed after 4 to 6 weeks.

(ii) *Indications for use.* For the removal and control of lungworms (*Dictyocaulus viviparus*), stomach worms (*Haemonchus contortus*, *Ostertagia ostertagi*, *Trichostrongylus axei*), and intestinal worms (*Bunostomum phlebotomum*, *Nematodirus helvetianus*, *Cooperia punctata*, *C. oncophora*, *Trichostrongylus colubriformis*, and *Oesophagostomum radiatum*).

(iii) *Limitations.* Cattle must not be slaughtered within 8 days following last treatment.

Dated: April 24, 2007.

Bernadette Dunham,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. E7–8391 Filed 5–1–07; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

33 CFR Part 165

[CGD05–06–112]

RIN 1625–AA87

Security Zone; Severn River and College Creek, Annapolis, MD

AGENCY: Coast Guard, DHS.

ACTION: Final rule.

SUMMARY: The Coast Guard is establishing a permanent security zone on certain waters of the Severn River and College Creek. This action is necessary to ensure the security of high-ranking public officials and safeguard