LIST OF COMMENTERS

Abbreviation	Name
Allegheny	Allegheny Power and Allegheny Energy Supply Company, L.L.C.
AGA	American Gas Association.
APPA and TAPS	American Public Power Association and Transmission Access Policy Study Group.
Butte County	Butte County, California.
California Resources	California Resources Agency.
California State Agencies	California Coastal Commission, California Energy Commission, California Electricity Oversight Board, and California State Lands Commission.
Dominion	Dominion Transmission Inc., Dominion Cove Point, LNG, LP, and Dominion South Pipeline Company, LP.
EEI	Edison Electric Institute.
INGAA	Interstate Natural Gas Association of America.
MidAmerican	MidAmerican Energy Company.
NARUC	National Association of Regulatory Utility Commissioners.
NHA	National Hydropower Association.
SCE	Southern California Edison Company.
Western Energy Board	Western Interstate Energy Board and Committee on Regional Electric Power Cooperation.
Williston Basin	Williston Basin Interstate Pipeline Company.
Department of the Interior	United States Department of the Interior.

[FR Doc. E7–22141 Filed 11–13–07; 8:45 am]
BILLING CODE 6717–01–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 address for IDEXX Pharmaceuticals, Inc.

DATES: This rule is effective November 14, 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: IDEXX Pharmaceuticals, Inc., 4249–105 Piedmont Pkwy., Greensboro, NC 27410, has informed FDA of a change of address to 7009 Albert Pick Rd., Greensboro, NC 27409. 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 ''IDEXX Pharmaceuticals, Inc.''; and in the table in paragraph (c)(2) revise 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., 7009 Albert Pick Rd., Greensboro, NC 27409.	065274
* * *	* *

(2) * * *

Drug labeler code	Firm name and address
* * *	* *
065274	IDEXX Pharmaceuticals, Inc., 7009 Albert Pick Rd., Greensboro, NC 27409
* * *	* *

Dated: November 6, 2007.

Bernadette Dunham,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. E7–22210 Filed 11–13–07; 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; Chlortetracycline Powder

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect approval of a supplemental new animal drug application (NADA) filed by Fort Dodge Animal Health, Division of Wyeth Holdings Corp. The supplemental NADA provides for label revisions for chlortetracycline soluble powder.

DATES: This rule is effective November 14, 2007.

FOR FURTHER INFORMATION CONTACT: Joan C. Gotthardt, Center for Veterinary Medicine (HFV–130), Food and Drug Administration, 7500 Standish Pl.,