regulated organizations will not be compromised.

After considering these factors, the Commission has determined to amend Part 171, as set forth below.

List of Subjects in 17 CFR Part 171

Administrative practice and procedure, Commodity exchanges, Commodity futures.

■ In consideration of the following, the Commission hereby amends chapter I of title 17 of the Code of Federal Regulations as follows:

PART 171—RULES RELATING TO REVIEW OF NATIONAL FUTURES ASSOCIATION DECISIONS IN DISCIPLINARY, MEMBERSHIP DENIAL, REGISTRATION AND MEMBER RESPONSIBILITY ACTIONS

■ 1. The authority citation for part 171 continues to read as follows:

Authority: 7 U.S.C. 4a, 12a, and 21.

■ 2. Section 171.1(b) is amended in paragraph (b)(4) by adding ‘‘, Hearing Committee’’ between ‘‘Business Conduct Committees’’ and ‘‘or arbitration panels’’; and replacing ‘‘.’’ with ‘‘;’’ at the end of (b)(4); and by adding new paragraph (b)(5):

§ 171.1 Scope of rules.

* * * * *

(b) * * *

(5) Suspension of a member or a person associated with a member based solely on that person’s failure to pay an arbitration award or a settlement agreement resulting from an arbitration action brought pursuant to section 17(b)(10) of the Act or rules and regulations of the National Futures Association, or a settlement agreement resulting from a mediation proceeding sponsored by the National Futures Association, unless there are extraordinary circumstances that involve something more than the ministerial application of a predetermined sanction, or raise a colorable claim that the National Futures Assocaition has acted arbitrarily.

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Issued in Washington, DC on the 10th day of January 2005, by the Commission.

Jean A. Webb,

Secretary of the Commission.

[FR Doc. 05–709 Filed 1–12–05; 8:45 am]

BILLING CODE 6351–01–M

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 Alstoe, Ltd.

DATES: This rule is effective January 13, 2005.

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.gov*.

SUPPLEMENTARY INFORMATION: Alstoe, Ltd., Animal Health, Granary Chambers, 37–39 Burton St., Melton Mowbray, Leicestershire LE13 1AF, England has informed FDA of a change of address to Pera Innovation Park, Nottingham Rd., Melton Mowbray, Leicestershire, England LE13 0PB. 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. Section 510.600 is amended in the table in paragraph (c)(1) by revising the entry for ‘‘Alstoe, Ltd.’’; and in the table in paragraph (c)(2) by revising the entry for ‘‘062408’’ 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
* * * * *	
Alstoe, Ltd., Animal Health, Pera Innovation Park, Nottingham Rd., Melton Mowbray, Leicestershire, England LE13 0PB	062408
* * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * *	
062408	Alstoe, Ltd., Animal Health, Pera Innovation Park, Nottingham Rd., Melton Mowbray, Leicestershire, England LE13 0PB
* * * * *	

Dated: January 3, 2005.

Steven D. Vaughn,

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

[FR Doc. 05–697 Filed 1–12–05; 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; Levamisole Powder for Oral Solution

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 an abbreviated new animal drug application (ANADA) filed by Phoenix Scientific, Inc. The ANADA provides for use of levamisole hydrochloride soluble powder to make a drench solution for oral administration to cattle and sheep which is effective against various internal parasites.

DATES: This rule is effective January 13, 2005.