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.

* * * * *

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.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV 104), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 301–827–8549, e-mail: *lonnie.luther@fda.gov*.

SUPPLEMENTARY INFORMATION: Phoenix Scientific, Inc., 3915 South 48th St. Terrace, St. Joseph, MO 64503, filed ANADA 200–386 for Levamisole Hydrochloride Soluble Drench Powder used to make a drench solution for oral administration to cattle and sheep which is effective against various internal parasites. Phoenix Scientific's Levamisole Hydrochloride Soluble Drench Powder is approved as a generic copy of Schering-Plough Animal Health Corp.'s, LEVASOL (levamisole hydrochloride) Soluble Drench Powder, approved under NADA 112–051. The ANADA is approved as of December 17, 2004, and the regulations are amended in 21 CFR 520.1242a to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In accordance with the freedom of information provisions of 21 CFR part 20 and 21 CFR 514.11(e)(2)(ii), a summary of safety and effectiveness data and information submitted to support approval of this application may be seen in the Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

FDA has determined under 21 CFR 25.33(a)(1) that this action is of a type that does not individually or cumulatively have a significant effect on the human environment. Therefore, neither an environmental assessment nor an environmental impact statement is required.

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 Subject in 21 CFR Part 520

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 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 520.1242a is amended by revising paragraphs (a) and (b)(4) to read as follows:

§ 520.1242a Levamisole powder for oral solution.

(a) *Specifications*. Each package of powder contains 9.075, 11.7, 18.15, 46.8, 362.7, or 544.5 grams (g) levamisole hydrochloride.

(b) * * *

(4) No. 059130 for use of 46.8-, 362.7-, and 544.5-g packages as in paragraphs (e)(1)(i), (e)(1)(ii)(B), (e)(1)(iii), (e)(2)(i), (e)(2)(ii)(B), and (e)(2)(iii) of this section; and for use of an 18.15-g package as in paragraph (e)(3) of this section.

* * * * *

Dated: January 6, 2005.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 05–675 Filed 1–12–05; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

33 CFR Part 110

[CGD01–04–004]

1625–AA01

Anchorage Grounds; Buzzards Bay, MA

AGENCY: Coast Guard, DHS.

ACTION: Final rule.

SUMMARY: The Coast Guard hereby establishes anchorage regulations for Buzzards Bay, Nantucket Sound, and adjacent waters of Massachusetts by relocating anchorage ground ''L'' in Buzzards Bay to an area near Naushon Island, MA. This action is intended to increase the safety of life and property on Buzzards Bay, improve the safety of anchored vessels in anchorage ''L'', and provide for the overall safe and efficient flow of vessel traffic and commerce along the newly established Recommended Traffic Route for Deep Draft Vessels. This regulation will maintain the original shape and dimension of anchorage ''L'' but move the anchorage to a new location within Buzzards Bay.

DATES: This rule is effective February 14, 2005.

ADDRESSES: Comments and material received from the public, as well as documents indicated in this preamble as being available in the docket, are part of docket [CGD01–02–027] and are available for inspection or copying at First Coast Guard District, 408 Atlantic Ave., Boston, Massachusetts 02110 between 8 a.m. and 3 p.m., Monday through Friday, except Federal holidays.

FOR FURTHER INFORMATION CONTACT: Mr. John J. Mauro, Commander (oan), First Coast Guard District, 408 Atlantic Ave., Boston, MA 02110, Telephone (617) 223–8355, e-mail: *jmauro@d1.uscg.mil.*

SUPPLEMENTARY INFORMATION:

Regulatory Information

On April 16, 2004, we published a notice of proposed rulemaking (NPRM) entitled Anchorage Grounds; Buzzards Bay, MA in the **Federal Register** (69 FR 20568). We received one comment on the proposed rule. No public hearing was requested and none was held.

Background and Purpose

In light of significant oil spills in Rhode Island Sound in 1996 and Buzzards Bay in 2003, the Coast Guard investigated methods of improving navigational safety in Buzzards Bay. The Coast Guard conducted a Port and Waterways Safety Assessment (PAWSA) to collect input on potential navigational safety improvements in Buzzards Bay from the local maritime community. After studying the issue and collecting mariner input, the Coast Guard concluded that a Recommended Traffic Route for Deep Draft vessels in Buzzards Bay should be implemented to improve navigation safety in this area.

Presently, there are two designated anchorage grounds in Buzzards Bay; anchorage ''L'' and anchorage ''M'', whose locations are described in 33 CFR 110.140(b)(3) and 33 CFR 110.140(b)(4), respectively. The present location of anchorage ''L'' puts it directly in the path of the Recommended Route for Deep Draft vessels entering or leaving the Cape Cod Canal via Cleveland Ledge Channel depicted on current versions of NOAA nautical charts in the area. Thus, this rule is needed to move anchorage ''L'' to a new and safer location. Although the location of anchorage ''L'' will change, its size and shape will remain the same.

The Coast Guard has defined the anchorage areas contained herein with the advice and consent of the Army Corps of Engineers, New England District, located at 696 Virginia Rd., Concord, MA 01742.

This regulation will not exclude fishing activity or the transit of vessels in the anchorage grounds. The Coast Guard expects no increase in the amount of vessels utilizing anchorage ''L'' as a result of this change in its location.