

(g) *Can I comply with this AD in any other way?* You may use an alternative method of compliance or adjust the compliance time if:

(1) Your alternative method of compliance provides an equivalent level of safety; and
(2) The Manager, Fort Worth Airplane Certification Office (ACO), approves your alternative. Submit your request through an FAA Principal Maintenance Inspector. The inspector may add comments before sending it to the Manager, Fort Worth ACO.

(3) Alternative methods of compliance approved in accordance with AD 2002-13-02, which is superseded by this AD, are not approved as alternative methods of compliance with this AD.

Note: This AD applies to each airplane identified in paragraphs (a)(1), (a)(2), and (a)(3) of this AD, regardless of whether it has been modified, altered, or repaired in the area subject to the requirements of this AD. For airplanes that have been modified, altered, or repaired so that the performance of the requirements of this AD is affected, the owner/operator must request approval for an alternative method of compliance in accordance with paragraph (g) of this AD. The request should include an assessment of the effect of the modification, alteration, or repair on the unsafe condition addressed by this AD; and, if you have not eliminated the unsafe condition, specific actions you propose to address it.

(h) *What if I need to fly the airplane to another location to comply with this AD?* The FAA can issue a special flight permit under §§ 21.197 and 21.199 of the Federal Aviation Regulations (14 CFR 21.197 and 21.199) to operate your airplane to a location where you can accomplish the requirements of this AD provided that the following is adhered to:

(1) Operate in day visual flight rules (VFR) only.
(2) Ensure that the hopper is empty.
(3) Limit airspeed to 135 miles per hour (mph) indicated airspeed (IAS).
(4) Avoid any unnecessary g-forces.
(5) Avoid areas of turbulence.
(6) Plan the flight to follow the most direct route.

(i) *Are any service bulletins incorporated into this AD by reference?* Actions required by this AD must be done in accordance with Snow Engineering Co. Service Letter #55, Revised October 23, 2002; Snow Engineering Co. Service Letter #70, Revised October 23, 2002; Snow Engineering Co. Service Letter #226, dated December 17, 2002; Snow Engineering Process Specification Number 197, Revised June 4, 2002; and Snow Engineering Co. Service Letter #220, dated December 17, 2002. The Director of the Federal Register approved this incorporation by reference under 5 U.S.C. 552(a) and 1 CFR part 51. You may get copies from Air Tractor, Inc., P.O. Box 485, Olney, Texas 76374. You may view copies at FAA, Central Region, Office of the Regional Counsel, 901 Locust, Room 506, Kansas City, Missouri, or at the Office of the Federal Register, 800 North Capitol Street, NW., suite 700, Washington, DC.

(j) *Does this AD action affect any existing AD actions?* This amendment supersedes AD 2002-13-02, Amendment 39-12789.

(k) *When does this amendment become effective?* This amendment becomes effective on April 4, 2003.

Issued in Kansas City, Missouri, on March 11, 2003.

David R. Showers,

Acting Manager, Small Airplane Directorate, Aircraft Certification Service.

[FR Doc. 03-6262 Filed 3-18-03; 8:45 am]

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DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 71

[Docket No. FAA-2003-14457; Airspace Docket No. 03-ACE-10]

Modification of Class E Airspace; Herington, KS

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Direct final rule; request for comments; correction.

SUMMARY: This action corrects a direct final rule; request for comments that was published in the **Federal Register** on Tuesday, February 25, 2003, (68 FR 8704). It corrects an error in the location of the Herington Regional Airport, KS in the legal description of the Herington, KS Class E airspace.

DATES: This direct final rule is effective on 0901 UTC, May 15, 2003.

Comments for inclusion in the Rules Docket must be received on or before March 25, 2003.

FOR FURTHER INFORMATION CONTACT: Kathy Randolph, Air Traffic Division, Airspace Branch, ACE-520C, DOT Regional Headquarters Building, Federal Aviation Administration, 901 Locust, Kansas City, MO 64106; telephone: (816) 329-2525.

SUPPLEMENTARY INFORMATION:

History

Federal Register document 03-4322 published on Tuesday, February 25, 2003, (68 FR 8704) modified Class E airspace at Herington, KS. The modification was to correct the Herington Regional Airport, KS airport reference point used in the legal description of the Herington, KS Class E airspace area. The latitude of the Herington Regional Airport, KS airport reference point was published incorrectly.

Accordingly, pursuant to the authority delegated to me, the Herington, KS Class E airspace, as published in the **Federal Register** on Tuesday, February 25, 2003, (68 FR

8704), (FR Doc. 03-4322), is corrected as follows:

§ 71.1 [Corrected]

On page 8705, Column 1, second paragraph from the bottom, change “Herington Regional Airport, KS (lat. 39°41’41”N., long. 96°48’29”W.)” to read “Herington Regional Airport, KS (lat. 38°41’41”N., long. 96°48’29”W.)”

Issued in Kansas City, MO, on March 7, 2003.

Herman J. Lyons, Jr.,

Manager, Air Traffic Division, Central Region.

[FR Doc. 03-6623 Filed 3-18-03; 8:45 am]

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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 Vetrepharm Research, Inc., to Bioniche Animal Health USA, Inc.

DATES: This rule is effective March 19, 2003.

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: dnewkirk@cvm.fda.gov.

SUPPLEMENTARY INFORMATION:

Vetrepharm Research, Inc., 119 Rowe Rd., Athens, GA 30601, has informed FDA of a change of name to Bioniche Animal Health USA, Inc. Accordingly, the agency is amending the regulations in 21 CFR 510.600(c) to reflect this 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