

**Adoption of the Amendment**

■ Accordingly, the Federal Aviation Administration amends 14 CFR part 71 as follows:

**PART 71—DESIGNATION OF CLASS A, CLASS B, CLASS C, CLASS D, AND CLASS E AIRSPACE AREAS; AIRWAYS; ROUTES; AND REPORTING POINTS**

■ 1. The authority citation for part 71 continues to read as follows:

**Authority:** 49 U.S.C. 106(g), 40103, 40113, 40120; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., p. 389.

**§ 71.1 [Amended]**

■ 2. The incorporation by reference in 14 CFR 71.1 of Federal Aviation Administration Order 7400.9K, dated August 30, 2002, and effective September 16, 2002, is amended as follows:

*Paragraph 6005 Class E airspace areas extending upward from 700 feet or more above the surface of the earth.*

\* \* \* \* \*

**ACE NE E5 Valentine, NE**

Valentine, Miller Field, NE

(Lat. 42°51'28" N., long. 100°32'51" W.)

Valentine NDB

(Lat. 42°51'42" N., long. 100°32'59" W.)

That airspace extending upward from 700 feet above the surface within a 6.5-mile radius of Miller Field; and within 2.6 miles each side of the 149° bearing from the Valentine NDB extending from the 6.5-mile radius to 7.9 miles southeast of the airport; and within 1.8 miles each side of the 325° bearing from the airport extending from the 6.5-mile radius to 9.2 miles northwest of the airport.

\* \* \* \* \*

Issued in Kansas City, MO, on May 8, 2003.

**David W. Hope,**

*Acting Manager, Air Traffic Division, Central Region.*

[FR Doc. 03–12379 Filed 5–16–03; 8:45 am]

**BILLING CODE 4910–13–M**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**21 CFR Parts 510 and 522**

**Injectable or Implantable Dosage Form New Animal Drugs; Zinc Gluconate**

**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 new animal drug application (NADA) filed by Technology Transfer, Inc. The NADA provides for use of zinc gluconate solution for chemical sterilization of dogs by intratesticular injection.

**DATES:** This rule is effective May 19, 2003.

**FOR FURTHER INFORMATION CONTACT:**

Melanie R. Berson, Center for Veterinary Medicine (HFV–110), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–7543, e-mail: [mberson@cvm.fda.gov](mailto:mberson@cvm.fda.gov).

**SUPPLEMENTARY INFORMATION:**

Technology Transfer, Inc., 33 East Broadway, suite 190, Columbia, MO 65203, filed NADA 141–217 that provides for use of NEUTERSOL (zinc gluconate neutralized by arginine) Injectable Solution for chemical sterilization of 3- to 10-month-old male dogs by intratesticular injection. The NADA is approved as of March 17, 2003, and the regulations are amended in 21 CFR part 522 by adding new § 522.2690 to reflect the approval. The basis of approval is discussed in the freedom of information summary.

In addition, Technology Transfer, Inc., has not been previously listed in the animal drug regulations as a sponsor of an approved application. At this time, 21 CFR 510.600(c) is being amended to add entries for the firm.

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 Dockets Management Branch (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.

Under section 512(c)(2)(F)(i) of the Federal Food, Drug, and Cosmetic Act (the act) (21 U.S.C. 360b(c)(2)(F)(i)), this approval qualifies for 5 years of marketing exclusivity beginning March 17, 2003.

The agency has determined under 21 CFR 25.33(d)(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 Subjects**

**21 CFR Part 510**

Administrative practice and procedure, Animal drugs, Labeling, Reporting and recordkeeping requirements.

**21 CFR Part 522**

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 parts 510 and 522 are 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 alphabetically adding an entry for “Technology Transfer, Inc.” and in the table in paragraph (c)(2) by numerically adding an entry for “067647” 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
* * *	* *
Technology Transfer, Inc., 33 East Broadway, suite 190, Columbia, MO 65203.	067647
* * *	* *

(2) \* \* \*

Drug labeler code	Firm name and address
* *	* *
067647	Technology Transfer, Inc., 33 East Broadway, suite 190, Columbia, MO 65203.
* *	* *

**PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS**

■ 3. The authority citation for 21 CFR part 522 continues to read as follows:

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

■ 4. Section 522.2690 is added to read as follows: