| From                                    | To                    | Changeover points |         |  |  |
|---|-----------------------|-------------------|---------|--|--|
| FIOITI                                  | То                    | Distance          | From    |  |  |
| § 95.8005 Jet Routes Changeover Points  |                       |                   |         |  |  |
| J–58 Is Amended To Add Changeover Point |                       |                   |         |  |  |
| Milford, UT VORTAC                      | Farmington, NM VORTAC | 92                | Milford |  |  |

[FR Doc. 02–32412 Filed 12–23–02; 8:45 am] BILLING CODE 4910–13–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,

ппъ.

ACTION: Final rule.

**SUMMARY:** The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of address for Phoenix Scientific, Inc.

**DATES:** This rule is effective December 24, 2002.

#### 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; email: dnewkirk@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: Phoenix Scientific, Inc., 3915 South 48th St. Terrace, P.O. Box 6457, St. Joseph, MO 64506–0457, has informed FDA of a change of address to 3915 South 48th St. Terrace, St. Joseph, MO 64503. Accordingly, the agency is amending the regulations in 21 CFR 510.600 to reflect the change of sponsor's address.

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 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 "Phoenix Scientific, Inc." and in the table in paragraph (c)(2) by revising the entry for "059130" 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 |     |   |
|---------------------------|---------|------|-------------------|-----|---|
|                           | *       | *    | *                 | *   | * |
| Phoenix Scientific, Inc., |         |      | 059130            |     |   |
| 3915                      | South   | 48th | St. T             | er- |   |
| race,                     | St. Jo: | seph | , MO              |     |   |
| 64503                     | 3.      |      |                   |     |   |
|                           | *       | *    | *                 | *   | * |

| (2) * * *         |   |                       |      |                 |                             |          |
|-------------------|---|-----------------------|------|-----------------|-----------------------------|----------|
| Drug labeler code |   | Firm name and address |      |                 |                             |          |
|                   | * | *                     | *    | *               | *                           |          |
| 059130            |   |                       | 3915 | South<br>St. Jo | ntific,<br>n 48th<br>oseph, | St. Ter- |
|                   | * | *                     | *    | *               | *                           |          |

Dated: December 4, 2002.

#### Steven D. Vaughn,

Director, Office of New Animal Drug Evaluation, Center for Veterinary Medicine. [FR Doc. 02–32346 Filed 12–23–02; 8:45 am] BILLING CODE 4160–01–8

### DEPARTMENT OF HEALTH AND HUMAN SERVICES

**Food and Drug Administration** 

21 CFR Parts 510, 520, 522, and 524

**New Animal Drugs; Change of Sponsor** 

**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 for 25 approved new animal drug applications (NADAs) from American Cyanamid to Fort Dodge Animal Health.

**DATES:** This rule is effective December 24, 2002.

#### 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, email: lluther@cvm.fda.gov.

SUPPLEMENTARY INFORMATION: American Cyanamid, Division of American Home Products, P.O. Box 1339, Fort Dodge, IA 50501, has informed FDA that it has transferred ownership of, and all rights and interest in, the following 25 approved NADAs to Fort Dodge Animal Health, Division of American Cyanamid Co., P.O. Box 1339, Fort Dodge, IA 50501:

| 30301.         |  |
|----------------|--|
| NADA<br>Number | Trade Name                               |
| 006–084        | SULMET Drinking Water Solution           |
| 008-774        | SULMET Solution Injectable               |
| 011-582        | VETAMOX Soluble Powder                   |
| 011-644        | FELAC                                    |
| 013-957        | S.E.Z. Drinking Water 6.25%              |
| 015-160        | Sodium Sulfachloropyrazine               |
|                | Solution                                 |
| 033-342        | PROBAN Cythioate Tablets 30              |
|                | mg                                       |
| 033–606        | PROBAN Oral Liquid                       |
| 033–653        | S.E.Z. Drinking Water Solution           |
| 033–654        | S.E.Z. Oblets 15 g                       |
| 033–655        | S.E.Z. Intravenous Solution              |
| 047–033        | S.E.Z. C–R Oblets 15 g                   |
| 055–012        | AUREOMYCIN Sulmet Soluble                |
|                | Powder                                   |
| 055–018        | AUREOMYCIN Tablets 25 mg                 |
| 055–020        | AUREOMYCIN Soluble Powder                |
| 055–039        | AUREOMYCIN Soluble Oblets                |
| 065–071        | AUREOMYCIN Soluble Powder                |
| 065–269        | POLYOTIC Soluble Powder                  |
| 065–270        | POLYOTIC Oblets                          |
| 065–313        | BACIFERM Soluble 50                      |
| 065–440        | AUREOMYCIN Soluble Powder                |
| OCE 444        | Concentrate POLYOTIC Soluble Powder Con- |
| 065–441        | centrate                                 |
| 122–271        | SULMET Oblets                            |
| 122-271        | SULMET Oblets SULMET Soluble Powder      |
| 122-212        | SULMET SUMBLE FOWARI                     |