By the National Credit Union Administration Board on July 23, 1997.

Becky Baker,

Secretary of the Board.

Accordingly, NCUA amends 12 CFR chapter VII as follows:

PART 701—[AMENDED]

1. The authority citation for Part 701 is revised to read as follows:

Authority: 12 U.S.C. 1752(5), 1755, 1756, 1757, 1759, 1761a, 1761b, 1766, 1767, 1782, 1784, 1787, 1789. Section 701.6 is also authorized by 15 U.S.C. 3717. Section 701.31 is also authorized by 15 U.S.C. 1601 *et seq.*; 42 U.S.C. 1981 and 3601–3610. Section 701.35 is also authorized by 42 U.S.C. 4311–4312.

2. Section 701.21(c)(7)(ii)(C) is revised to read as follows:

§ 701.21 Loans to members and lines of credit to members.

* * * * *

(c) * * *

(7) * * *

(ii) * * *

(C) *Expiration.* After March 8, 1999, or as otherwise ordered by the NCUA Board, the maximum rate on federal credit union extensions of credit to members shall revert to 15 percent per year. Higher rates may; however, be charged, in accordance with paragraphs (c)(7)(ii) (A) and (B) of this section, on loans and line of credit balances existing on or before March 8, 1999.

* * * * *

[FR Doc. 97–19935 Filed 7–30–97; 8:45 am]

BILLING CODE 7535–01–U

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 510

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 46 new animal drug applications (NADA's) from Solvay Animal Health, Inc., to Fort Dodge Animal Health, A Division of American Cyanamid Co.

EFFECTIVE DATE: July 31, 1997.

FOR FURTHER INFORMATION CONTACT: Thomas J. McKay, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0213.

SUPPLEMENTARY INFORMATION: Solvay Animal Health, Inc., 1201 Northland Dr., Mendota Heights, MN 55120, has informed FDA that it has transferred the ownership of, and all rights and interests in, the following approved NADA's to Fort Dodge Animal Health, A Division of American Cyanamid Co., P.O. Box 1339, Fort Dodge, IA 50501:

NADA No.	Drug Name
006–417	Recovr
006–103	Follutein
006–707	Sulquin 6–50
008–274	Pig Scour Tablets
009–035	Ophtaine
011–141	Unistat-2
011–482	Vetame Tabs and Injection
011–879	Rubfrafer Injection
012–198	Vetalog Parenteral
012–258	Panalog Ointment (Solvaderm)
013–624	Vetalog Tabs
014–250	Novastat
031–448	Rheaform Bolus
031–553	Esb3 Powder & Solution
032–319	Furox Aerosol Spray
032–738	Pacitran Soluble
033–127	Vetisulid Bolus
033–318	Vetisulid Injectable
033–319	Vetisulid Tabs
033–373	Vetisulid Powder
034–536	Aklomix
034–537	Novastat-3
034–705	Equipoise
035–388	Novastat-W
039–666	Unistat-3
040–181	Vetisulid Oral Suspension
046–146	Vetalog Cream
046–147	Dirocide Syrup
049–892	Spanbolet II
055–060	Potassium G penicillin
055–064	Redicillin (Princillin)
055–066	Redicillin (Princillin)
055–071	Redicillin (Princillin)
065–130	Crystaline Pro Penicillin
065–174	Crysticillin 300 A.S. Vet
065–410	Tetra–Sal Soluble
091–192	Renografin 76
091–240	Renovist
091–327	Gastrogratin
093–512	Dirocide Tabs
096–676	Panalog Cream
099–388	Vetalog Oral Powder
126–232	Calfspan
131–808	Dirocide Syrup

NADA No.	Drug Name
139–913	Equron
140–909	Sulka-S-Bolus

Accordingly, the agency is amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) to reflect the change of sponsor. The drug labeler code assigned to Solvay Animal Health is being retained as the drug labeler code for the new sponsor.

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: Secs. 201, 301, 501, 502, 503, 512, 701, 721 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e).

§ 510.600 [Amended]

2. Section 510.600 *Names, addresses, and drug labeler codes of sponsors of approved applications* is amended in the table in paragraph (c)(1) by removing the entry for ''Solvay Animal Health, Inc.'' and by alphabetically adding a new entry for ''Fort Dodge Animal Health, A Division of American Cyanamid Co., P.O. Box 1339, Fort Dodge, IA 50501'' and in the table in paragraph (c)(2) in the entry for ''053501'' by removing the sponsor name and address ''Solvay Animal Health, Inc., 1201 Northland Dr., Mendota Heights, MN 55120'' and adding in its place ''Fort Dodge Animal Health, A Division of American Cyanamid Co., P.O. Box 1339, Fort Dodge, IA 50501''.

Dated: July 22, 1997.

Robert C. Livingston,

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

[FR Doc. 97–20249 Filed 7-30-97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510 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 three approved new animal drug applications (NADA's) from Syntex Animal Health, Inc., Division of Syntex Agribusiness, Inc., to Medicis Dermatologics, Inc.

EFFECTIVE DATE: July 31, 1997.

FOR FURTHER INFORMATION CONTACT: Thomas J. McKay, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0213.

SUPPLEMENTARY INFORMATION: Syntex Animal Health, Inc., Division of Syntex Agribusiness, Inc., 3401 Hillview Ave., P.O. Box 10850, Palo Alto, CA 94303, has informed FDA that it has transferred ownership of, and all rights and interests in NADA's 15–151 (*fluocinolone acetonide, neomycin sulfate cream*), 15–152 (*fluocinolone acetonide cream*), and 15–298 (*fluocinolone acetonide solution*) to Medicis Dermatologics, Inc., 4343 East Camelback Rd., suite 250, Phoenix, AZ 85018–2700. Accordingly, the agency is amending the regulations in 21 CFR 524.981a, 524.981b, and 524.981c to reflect the transfer of ownership. The agency is also amending the regulations in 21 CFR 510.600(c)(1) and (c)(2) by alphabetically adding a new listing for Medicis Dermatologics, Inc.

List of Subjects

21 CFR Part 510

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

21 CFR Part 524

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 parts 510 and 524 are amended as follows:

PART 510—NEW ANIMAL DRUGS

1. The authority citation for 21 CFR part 510 continues to read as follows:

Authority: Secs. 201, 301, 501, 502, 503, 512, 701, 721 of the Federal Food, Drug, and Cosmetic Act (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 a new entry for ''Medicis Dermatologics, Inc.'' and in the table in paragraph (c)(2) by numerically adding a new entry for ''099207'' 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
* * *	* * * *
Medicis Dermatologics, Inc., 4343 East Camelback Rd., suite 250, Phoenix, AZ 85018–2700.	099207
* * *	* * * *

(2) * * *