administrators of the 4 orders to ascertain whether at least two-thirds of the producers marketing their milk under each of the orders approved the issuance of the amended orders. The final decision concluded that amended orders were needed to effectuate the declared policy of the Agricultural Marketing Agreement Act. That Act requires that at least two-thirds of the producers voting in a referendum must vote affirmatively before an order can be issued.

Less than two-thirds of the producers whose milk is pooled under the Tennessee Valley order approved the issuance of the proposed amended order. Consequently, on July 3 the Department issued a notice of proposed termination of the Tennessee Valley order. It is now evaluating comments received in response to that notice.

At the present time, the Tennessee Valley milk order is being administered under the interim provisions adopted in August 1996, whereas the surrounding orders with transportation credit provisions are being administered with revised provisions that became effective on August 1, 1997.

In July 1997, an extraordinary volume of supplemental milk was received in the neighboring Southeast order. As a result of these receipts, the transportation credit balancing fund for that order was virtually depleted in July. There is now good reason to believe that shipments of supplemental milk may be rerouted to handlers under the Tennessee Valley order in September since that order still has the interim provision allowing unlimited payments for transportation credits even if the money to pay for the credits must come from the producer-settlement fund. Although the Tennessee Valley order has a viable balance in the TCBF at the present time, it is likely that funds from the producer-settlement fund will be necessary for transportation credit payments for September's milk. Were this to happen, it would reduce blend prices to producers in the Tennessee Valley order while their counterparts in the surrounding markets with transportation credit provisions would suffer no such reduction under the revised August 1997 amendments. This situation would be inconsistent with the premises upon which the psf transfer provision was included in the Tennessee Valley order.

This suspension is necessary to ensure that producers' milk will not be moved in an uneconomic and inefficient manner simply to obtain unlimited transportation credits under the Tennessee Valley order and to ensure that producers in the Tennessee Valley order will be treated in an equitable manner in relation to producers supplying the adjacent Southeast, Carolina, and Louisville-Lexington-Evansville orders.

Accordingly, it is appropriate to suspend the aforesaid provisions during the period of consideration of terminating the Tennessee Valley milk order.

It is hereby found and determined that thirty days' notice of the effective date hereof is impractical, unnecessary and contrary to the public interest in that:

(a) The suspension is necessary to reflect current marketing conditions and to assure orderly marketing conditions in the marketing area, in that such rule is necessary to permit the continued pooling of the milk of dairy farmers who have historically supplied the market without the need for making costly and inefficient movements of milk; and

(b) This suspension does not require of persons affected substantial or extensive preparation prior to the effective date.

Therefore, good cause exists for making this order effective less than 30 days from the date of publication in the **Federal Register**.

List of Subjects in 7 CFR Part 1011

Milk marketing orders.

For the reasons set forth in the preamble, 7 CFR Part 1011 is amended as follows:

PART 1011—MILK IN THE TENNESSEE VALLEY MARKETING AREA

1. The authority citation for 7 CFR Part 1011 continues to read as follows:

Authority: 7 U.S.C. 601–674.

§ 1011.61 [Suspended in part]

2. In § 1011.61, paragraph (a)(4) is suspended.

§ 1011.81 [Suspended in part]

3. In § 1011.81, paragraph (b) is suspended.

Dated: August 29, 1997.

Lon Hatamiya,

Administrator.

[FR Doc. 97–23568 Filed 9–3–97; 8:45 am]

BILLING CODE 3410–02–P

CONSUMER PRODUCT SAFETY COMMISION

16 CFR Parts 1000, 1014, 1021, 1051, 1115, 1211, 1402, 1406, 1500, 1502, 1700, and 1702

Address and Telephone Number Corrections

AGENCY: Consumer Product Safety Commission.

ACTION: Final rule.

SUMMARY: The Commission is amending 16 CFR chapter II to correct errors in addresses and telephone numbers.

EFFECTIVE DATE: September 4, 1997.

FOR FURTHER INFORMATION CONTACT: Joseph F. Rosenthal, Office of the General Counsel, Consumer Product Safety Commission, Washington, D.C. 20207, telephone 301-504-0980.

SUPPLEMENTARY INFORMATION: Some addresses, office designations, and telephone numbers in various parts of 16 CFR chapter II are obsolete as a result of the Consumer Product Safety Commission's relocation to new headquarters in 1994. This rule makes the necessary corrections. It also revises some authority citations to conform to **Federal Register** recommendations.

Since this rule relates solely to internal agency management, pursuant to 5 U.S.C. 553(b), notice and other public procedures are not required and it is effective immediately on the specified effective date. Further, this action is not a rule as defined in the Regulatory Flexibility Act, 5 U.S.C. 601-612 and, thus, is exempt from the provisions of the Act. This action will have no effect on the environment.

List of Subjects

16 CFR Part 1000

Organization and functions (Government agencies).

16 CFR Part 1014

Privacy

16 CFR Part 1021

Environmental impact statements.

16 CFR Part 1051

Administrative practice and procedure, consumer protection.

16 CFR Part 1115

Administrative practice and procedure, business and industry, consumer protection, reporting and recordkeeping requirements.

16 CFR Part 1211

Consumer protection, imports, labeling, reporting and recordkeeping requirements.

16 CFR Part 1402

Consumer protection, labeling, radio, television.

16 CFR Part 1406

Consumer protection, fire prevention, flammable materials, heaters, household appliances, labeling, reporting and recordkeeping requirements.

16 CFR Part 1500

Consumer protection, hazardous materials, imports, infants and children, labeling, law enforcement, reporting and recordkeeping requirements, toys.

16 CFR Part 1502

Administrative practice and procedure, consumer protection, hazardous substances, poison prevention.

16 CFR Part 1700

Consumer protection, drugs, infants and children, packaging and containers, poison prevention, reporting and recordkeeping requrements.

16 CFR Part 1702

Administrative practice and procedure, consumer protection, drugs, infants and children, packaging and containers, poison prevention.

Accordingly, 16 CFR chapter II is amended as follows:

PART 1000—[AMENDED]

1. The authority citation for part 1000 continues to read as follows:

Authority: 5 U.S.C. 552(a).

2. In section 1000.8 remove the words ''5401 Westbard Avenue'' and add, in their place, ''4330 East West Highway''.

PART 1014—[AMENDED]

1. The authority citation for part 1014 continues to read as follows:

Authority: 5 U.S.C. 552a.

2. In section 1014.3(a) remove the words ''5401 Westbard Avenue'' and add, in their place ''4330 East West Highway''.

3. In section 1014.3(c) remove the words ''Division of Personnel's Processing Unit in Room 337, 5401 Westbard Avenue'' and add, in their place, ''Office of Human Resources Management, Room 523, 4330 East West Highway''.

PART 1021—[AMENDED]

1. The authority citation for part 1021 is revised to read as follows:

Authority: 42 U.S.C. 4321-4347; 40 CFR part 1500 et seq.

2. In section 1021.11 remove the words ''(301-492-6550)'' and add, in their place, ''(301-504-0550)''.

PART 1051—[AMENDED]

1. The authority citation for part 1051 continues to read as follows:

Authority: 5 U.S.C. 553(e), 5 U.S.C. 555(e).

2. In section 1051.3 remove the words ''either, 5401 Westbard Avene (third floor) Bethesda, Maryland or 1111 18th Street, NW, (eighth floor), Washington, DC'' and add, in their place ''4330 East West Highway, Bethesda, Maryland''.

PART 1115—[AMENDED]

1. The authority citation for part 1115 continues to read as follows:

Authority: 15 U.S.C. 2061, 2064, 2065, 2066(a), 2068, 2069, 2070, 2071, 2073, 2076, 2079, and 2084.

2. In section 1115.10(a) remove the words ''and Enforcement''.

3. In section 1115.10(a) remove the words ''301-492-6608''and add, in their place, ''301-504-0608''.

PART 1211—[AMENDED]

1. The authority citation for part 1211 is revised to read as follows:

Authority: 15 U.S.C. 2063 and 2065.

2. In section 1211.2(c) remove the words ''5401 Westbard Avenue'' and add, in their place ''4330 East West Highway''.

3. In section 1211.4(c) remove the words ''5401 Westbard Avenue'' and add, in their place, ''4330 East West Highway''.

4. In section 1211.5(b)(3) remove the words ''5402 Westbard Avenue'' and add, in their place, ''4330 East West Highway''.

5. In section 1211.10(d) remove the words ''5401 Westbard Avenue'' and add, in their place, ''4330 East West Highway''.

6. In section 1211.12(c)(2) remove the words ''5401 Westbard Avenue'' and add, in their place, ''4330 East West Highway''.

PART 1402—[AMENDED]

1. The authority citation for part 1402 is revised to read as follows:

Authority: 15 U.S.C. 2051, 2076.

2. In section 1402.4(b)(1) remove the words ''Associate Executive Director for Compliance and Enforcement, Consumer Product Safety Commission, 5401 Westbard Avenue'' and add, in their place, ''Assistant Executive Director for Compliance, Consumer Product Safety Commission, 4330 East West Highway''.

PART 1406—[AMENDED]

1. The authority citation for part 1406 is revised to read as follows:

Authority: 15 U.S.C. 2051, 2076.

2. In section 1406.5(d)(2) remove the words ''Associate Executive Director for Compliance and Administrative Litigation, Consumer Product Safety Commission, 5401 Westbard Avenue, Bethesda, Maryland 20207'' and add, in their place, ''Assistant Executive Director for Compliance, Consumer Product Safety Commission, Washington, DC 20207''.

PART 1500—[AMENDED]

1. The authority citation for part 1500 continues to read as follows:

Authority: 15 U.S.C. 1261-1278, 2079.

2. In section 1500.42, footnote 1, remove the words ''Directorate for Health Sciences, CPSC, Washington, D.C. 20207, (301) 492-6957''and add, in their place, '' Directorate for Epidemiology and Health Sciences, CPSC, Washington, DC 20207, (301) 504-0957''.

PART 1502—[AMENDED]

1. The authority citation for part 1502 continues to read as follows:

Authority: 15 U.S.C. 1261(q)(1)(B), 1262(a), 1262(e), 1269(a); 15 U.S.C. 1474(a); 21 U.S.C. 371(e)-(g).''

2. In section 1502.4(b) remove the words ''Room 420, 5401 Westbard Avenue, Bethesda, Maryland 20816'' and add, in their place ''Room 502, 4330 East West Highway, Bethesda, Maryland 20814''.

3. In section 1502.4(c) remove the words ''(301) 492-6800'' and add, in their place, ''(301) 504-0800''.

4. In section 1502.16(a) remove the words ''Room 420, 5401 Westbard Ave.'' and add, in their place ''Room 502, 4330 East West Highway''.

5. In section 1502.17(a) remove the words ''(301) 492-6800'' and add, in their place ''(301) 504-0800''.

PART 1700—[AMENDED]

1. The authority citation for part 1700 continues to read as follows:

Authority: 15 U.S.C. 1471-1476. Secs. 1700.1 and 1700.14 also issued under 15 U.S.C. 2079(a).

2. In section 1700.14(b) remove the words ''Attention: Bureau of Compliance, 5401 Westbard Avenue'' and add, in their place, ''Office of Compliance, 4330 East West Highway''.

PART 1702—[AMENDED]

1. The authority citation for part 1702 is revised to read as follows:

Authority: 15 U.S.C. 1471(4), 1472, 1474, 1269(a), 2079(a); 21 U.S.C. 371(a).

2. Section 1702.2(a)(1) is revised to read as follows:

§ 1702.2 Procedural requirements and recommendations.

(a) * * *

(1) Be mailed to the Office of the Secretary, Consumer Product Safety Commission, Washington, DC 20207, or delivered to the Office of the Secretary, Consumer Product Safety Commission, 4330 East West Highway, Bethesda, MD 20814.

* * * * *

Sadye E. Dunn,

Secretary, Consumer Product Safety Commission.

[FR Doc. 97–23498 Filed 9–3–97; 8:45 am]

BILLING CODE 6355–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Tetracycline Hydrochloride Soluble Powder

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 Med-Pharmex, Inc. The ANADA provides for oral use of tetracycline hydrochloride soluble powder in the drinking water of swine and calves for control and treatment of certain diseases caused by pathogens susceptible to tetracycline, and of chickens and turkeys for control of certain diseases caused by pathogens susceptible to tetracycline.

EFFECTIVE DATE: September 4, 1997.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center For Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209.

SUPPLEMENTARY INFORMATION: Med-Pharmex, Inc., 2727 Thompson Creek Rd., Pomona, CA 91767–1861, filed ANADA 200–234, which provides for oral use of tetracycline hydrochloride soluble powder in the drinking water of calves, swine, chickens, and turkeys, as follows: (1) For calves for control and treatment of bacterial enteritis (scours) caused by *Escherichia coli*, and bacterial pneumonia (shipping fever complex) associated with *Pasteurella* spp., *Actinobacillus pleuropneumoniae* (*Hemophilus* spp.), and *Klebsiella* spp. susceptible to tetracycline; (2) for swine for control and treatment of bacterial enteritis (scours) caused by *E. coli*, and bacterial pneumonia associated with *Pasteurella* spp., *Actinobacillus pleuropneumoniae* (*Hemophilus* spp.), and *Klebsiella* spp. susceptible to tetracycline; (3) for chickens for control of chronic respiratory disease (CRD or air-sac disease) caused by *Mycoplasma gallisepticum* and *E. coli*; infectious synovitis caused by *M. synoviae* susceptible to tetracycline; and (4) for turkeys for control of infectious synovitis caused by *M. synoviae* and bluecomb (transmissible enteritis or coronaviral enteritis) complicated by bacterial organisms susceptible to tetracycline.

Approval of Med-Pharmex's ANADA 200–234 tetracycline hydrochloride soluble powder is as a generic copy of Fermenta's NADA 65–496 tetracycline hydrochloride soluble powder. ANADA 200–234 is approved as of July 22, 1997, and the regulations are amended in 21 CFR 520.2345d(a)(1) 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 Dockets Management Branch (HFA–305), Food and Drug Administration, 12420 Parklawn Dr., rm. 1–23, Rockville, MD 20857, between 9 a.m. and 4 p.m., Monday through Friday.

The agency has determined under 21 CFR 25.24(d)(1)(i) 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.

List of Subjects 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: Sec. 512 of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b).

§ 520.2345d [Amended]

2. Section 520.2345d *Tetracycline hydrochloride soluble powder* is amended in paragraph (a)(1) by removing ''047864, 000010, 057561, and 059130'' and adding in its place ''047864, 051259, 054273, 057561, and 059130''.

Dated: August 22, 1997.

Michael J. Blackwell,

Deputy Director, Center for Veterinary Medicine.

[FR Doc. 97–23372 Filed 9–3–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Clindamycin Hydrochloride Liquid

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 veterinary prescription use in dogs of clindamycin hydrochloride liquid for therapy of wounds, abscesses, and dental infections, and therapy of osteomyelitis.

EFFECTIVE DATE: September 4, 1997.

FOR FURTHER INFORMATION CONTACT: Lonnie W. Luther, Center for Veterinary Medicine (HFV–102), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0209.

SUPPLEMENTARY INFORMATION: Phoenix Scientific, Inc., 3915 South 48th St. Terrace, P.O. Box 6457, St. Joseph, MO 64506–0457, filed ANADA 200–193 that provides for veterinary prescription use in dogs of clindamycin hydrochloride liquid for therapy of wounds, abscesses, and dental infections when administered orally at 2.5 milligrams per pound (mg/lb) every 12 hours, and for therapy of osteomyelitis when administered orally at 5.0 mg/lb every 12 hours.

Phoenix Scientific, Inc.'s, ANADA 200–193 clindamycin hydrochloride liquid is approved as a generic copy of Pharmacia & Upjohn's NADA 135–940 Antirobe Aquadrops®. The ANADA is approved as of August 1, 1997, and the regulations are amended in 21 CFR 520.447(b) to reflect the approval. The