SUPPLEMENTARY INFORMATION:

Background

On June 4, 1997, Customs published in the **Federal Register** (62 FR 30448) interim regulations (T.D. 97–45) which amended § 24.24 of the Customs Regulations (19 CFR 24.24) to update the list of ports that process commercial vessels that transport cargo that are subject to the Water Resources Development Act of 1986. That document contained several typographical errors in the columns headed ''Port code, port name and state'' and ''Port descriptions and notations'', both of which may be relied on by importers in the preparation of necessary entry documentation. The errors identified are under the headings for Delaware, the District of Columbia, Illinois, Massachusetts, and Michigan, and consist of incomplete State abbreviations (for Maryland and Illinois), incorrect port codes (for East Chicago and Escanaba), and incomplete port descriptions (for Delaware and Massachusetts). Accordingly, this document corrects those typographical errors.

Corrections to Publication

The document (FR Doc. 97–14409) published in the **Federal Register** (62 FR 30448) on June 4, 1997, is corrected as follows:

1. On page 30450, under the heading for ''Delaware'', in the column headed ''Port descriptions and notations'', in the second line the word ''lower'' is added after the words ''all points on the'';

2. Also on page 30450, under the heading for ''District of Columbia'', in the column headed ''Port code, port name and state'', in the first line the capital letter ''D'' is removed and the designation ''MD'' is added in its place;

3. On page 30451, under the heading for ''Illinois'', in the column headed ''Port code, port name and state'', in the third line the numbers ''3902'' are removed and the numbers ''3904'' are added in their place; and in the column headed ''Port descriptions and notations'', in the first line the designation ''II.'' is removed and the designation ''IL'' is added in its place;

4. Also on page 30451, under the heading for ''Massachusetts'', in the column headed ''Port descriptions and notations'', in the second line the word ''River'' is removed and the word ''Rivers'' is added in its place; and

5. On page 30452, under the heading for ''Michigan'', in the column headed ''Port code, port name and state'', in the fifth line the number ''3803'' is removed and the number ''3808'' is added in their place.

Dated: August 21, 1997.

Harold M. Singer,

Chief, Regulations Branch.

[FR Doc. 97–22639 Filed 8–25–97; 8:45 am]

BILLING CODE 4820–02–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 522

Implantation or Injectable Dosage Form New Animal Drugs; Gentamicin Injection; Technical Amendment

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations for gentamicin injection. A document which published in the **Federal Register** of May 15, 1996 (61 FR 24440), inadvertently resulted in the 1997 edition of the Code of Federal Regulations not containing reference to two gentamicin injection approvals. This document amends the gentamicin injection regulations to reflect the approvals.

EFFECTIVE DATE: August 26, 1997.

FOR FURTHER INFORMATION CONTACT: David L. Gordon, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1739.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of May 15, 1996 (61 FR 24440), FDA published a document to reflect approval of Schering-Plough's supplemental NADA 101–862. In amending the regulations to reflect the supplemental approval, FDA provided amendatory instructions which resulted in two paragraphs inadvertently being removed. This document reestablishes those paragraphs in 21 CFR 522.1044(b)(3) and (b)(4).

List of Subjects in 21 CFR Part 522

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 522 is amended as follows:

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

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

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

2. Section 522.1044 is amended by adding paragraphs (b)(3) and (b)(4) to read as follows:

§ 522.1044 Gentamicin sulfate injection.

* * * * *

(b) * * *

(3) See No. 054273 for use of 50 milligrams-per-milliliter solution in dogs as in paragraph (d)(5) of this section.

(4) See No. 050604 for use of 100 milligrams-per-milliliter solution in chickens as in paragraph (d)(3) of this section.

* * * * *

Dated: August 18, 1997.

Stephen F. Sundlof,

Director, Center for Veterinary Medicine.

[FR Doc. 97–22622 Filed 8–25–97; 8:45 am]

BILLING CODE 4160–01–F

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 522

Implantation or Injectable Dosage Form New Animal Drugs; Polysulfated Glycosaminoglycan

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 Luitpold Pharmaceuticals, Inc. The NADA provides for intramuscular injection of polysulfated glycosaminoglycan for dogs for control of signs associated with noninfectious degenerative and/or traumatic arthritis of canine synovial joints.

EFFECTIVE DATE: August 26, 1997.

FOR FURTHER INFORMATION CONTACT: Ellen M. Buck, Center For Veterinary Medicine (HFV–114), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–594–1617.

SUPPLEMENTARY INFORMATION: Luitpold Pharmaceuticals, Inc., Animal Health Division, 1 Luitpold Dr., Shirley, NY 11967, filed NADA 141–038 that