of 1939,[12] and Sections 8, 30, 31, and 38 of the Investment Company Act of 1940.[13]

List of Subjects in 17 CFR Part 232

Incorporation by reference, Reporting and recordkeeping requirements, Securities.

Text of the Amendment

■ In accordance with the foregoing, Title 17, Chapter II of the Code of Federal Regulations is amended as follows:

PART 232—REGULATION S–T—GENERAL RULES AND REGULATIONS FOR ELECTRONIC FILINGS

■ 1. The authority citation for Part 232 continues to read as follows:

Authority: 15 U.S.C. 77f, 77g, 77h, 77j, 77s(a), 77sss(a), 78c(b), 78l, 78m, 78n, 78o(d), 78w(a), 78ll(d), 79t(a), 80a–8, 80a–29, 80a–30 and 80a–37.

■ 2. Section 232.301 is revised to read as follows:

§ 232.301 EDGAR Filer Manual.

Filers must prepare electronic filings in the manner prescribed by the EDGAR Filer Manual, promulgated by the Commission, which sets out the technical formatting requirements for electronic submissions. The requirements for filers using modernized EDGARLink are set forth in the EDGAR Release 8.6 EDGARLink Filer Manual Volume I, dated July 2003. Additional provisions applicable to Form N–SAR filers and Online Forms filers are set forth in the EDGAR Release 8.6 Filer Manual Volume II N–SAR Supplement, dated July 2003, and the EDGAR Release 8.6 OnlineForms Filer Manual Volume III, dated July 2003. All of these provisions have been incorporated by reference into the Code of Federal Regulations, which action was approved by the Director of the Federal Register in accordance with 5 U.S.C. 552(a) and 1 CFR Part 51. You must comply with these requirements in order for documents to be timely received and accepted. You can obtain paper copies of the EDGAR Filer Manual from the following address: Public Reference Room, U.S. Securities and Exchange Commission, 450 Fifth Street, NW., Washington, DC 20549–0102 or by calling Thomson Financial Inc at (800) 638–8241. Electronic format copies are available on the Commission's Web site. The address for the Filer Manual is *http://www.sec.gov/info/edgar.shtml.* You can also photocopy the document at the Office of the Federal Register, 800 North Capitol Street, NW., Suite 700, Washington, DC.

By the Commission.

Dated: July 22, 2003.

Jill M. Peterson,

Assistant Secretary.

[FR Doc. 03–19087 Filed 7–30–03; 8:45 am]

BILLING CODE 8010–01–P

[12] 15 U.S.C. 77sss.

[13] 15 U.S.C. 80a–8, 80a–29, 80a–30, and 80a–37.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 526

Intramammary Dosage Forms; 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 (NADAs) from Pfizer, Inc., to Schering-Plough Animal Health Corp.

DATES: This rule is effective July 31, 2003.

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, e-mail: *dnewkirk@cvm.fda.gov.*

SUPPLEMENTARY INFORMATION: Pfizer, Inc., 235 East 42d St., New York, NY 10017, has informed FDA that it has transferred ownership of, and all rights and interest in, the following three approved NADAs to Schering-Plough Animal Health Corp., 1095 Morris Ave., Union, NJ 07083:

NADA No.	Trade Name
55–069	ORBENIN DC (cloxacillin benzathine)
55–070	DARICLOX (cloxacillin sodium)
55–100	AMOXI–MAST (amoxicillin trihydrate)

Accordingly, the agency is amending the regulations in 21 CFR 526.88, 526.464b, and 526.464c to reflect the transfer of ownership.

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 in 21 CFR Part 526

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 526 is amended as follows:

PART 526—INTRAMAMMARY DOSAGE FORMS

■ 1. The authority citation for 21 CFR part 526 continues to read as follows:

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

§ 526.88 [Amended]

■ 2. Section 526.88 *Amoxicillin trihydrate for intramammary infusion* is amended in paragraph (b) by removing ''000069'' and by adding in its place ''000061''.

§ 526.464b [Amended]

■ 3. Section 526.464b *Cloxacillin benzathine for intramammary infusion, sterile* is amended in paragraph (d) by removing ''000069'' and by adding in its place ''000061''.

§ 526.464c [Amended]

■ 4. Section 526.464c *Cloxacillin sodium for intramammary infusion, sterile* is amended in paragraph (b) by removing ''000069'' and by adding in its place ''000061''.

Dated: July 18, 2003.

Steven D. Vaughn,

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

[FR Doc. 03–19445 Filed 7–30–03; 8:45 am]

BILLING CODE 4160–01–S

DEPARTMENT OF DEFENSE

Office of the Secretary

32 CFR Part 199

RIN 0720–AA79

TRICARE; Elimination of Nonavailability Statement and Referral Authorization Requirements and Elimination of Specialized Treatment Services Program

AGENCY: Office of the Secretary, DoD

ACTION: Interim final rule.

SUMMARY: This rule implements Section 735 of the National Defense Authorization Act for Fiscal Year 2002 (NDAA–02) (Public Law 107–107). It also implements Section 728 of the National Defense Authorization Act for Fiscal Year 2001 (NDAA–01) (Public Law 106–398). Section 735 of NDAA–02