(d) This section does not require the Bureau to take any action that would result in a fundamental alteration in the nature of a program or activity or in undue financial and administrative burdens.

§ 1072.112. Compliance procedures.

(a) Except as provided in paragraph (b) of this section, this section applies to all allegations of discrimination on the basis of disability in programs and activities conducted by the Bureau and denial of access to electronic and information technology.

(b) The Bureau shall process complaints alleging violations of section 504 with respect to employment according to the procedures established by the Equal Employment Opportunity Commission in 29 CFR part 1614 pursuant to section 501 of the Rehabilitation Act of 1973 (29 U.S.C. 791).

(c) All other complaints alleging violations of section 504 or section 508 may be sent to Labor and Employee Relations, Office of the Chief Human Capital Officer Consumer Financial Protection Bureau, 1700 G Street NW., Washington, DC 20052. The Office of the Chief Human Capital Officer shall be responsible for coordinating implementation of this section.

(d) *Complaint-filing procedures.* (1) Any person who believes that he or she has been subjected to discrimination prohibited by this part may by himself or herself or by his or her authorized representative file a complaint. Any person who believes that any specific class of persons has been subjected to discrimination prohibited by this part and who is a member of that class or the authorized representative of a member of that class may file a class complaint.

(2) The Bureau shall accept and investigate each timely filed, complete complaint over which it has jurisdiction.

(3) A complete complaint must be filed within 180 days of the alleged act of discrimination. A complaint submitted to the Bureau via first-class mail will be deemed to have been filed when postmarked. A complaint submitted to the Bureau via any other means of delivery will be deemed to have been filed when received by the Bureau. The Bureau may extend this time period for good cause.

(e) If the Bureau receives a complaint over which it does not have jurisdiction, it shall promptly notify the complainant and shall make reasonable efforts to refer the complaint to the appropriate government entity.

(f) The Bureau shall notify the Architectural and Transportation Barriers Compliance Board upon receipt of any complaint alleging that a building or facility that is subject to the Architectural Barriers Act of 1968, as amended (42 U.S.C. 4151–4157), is not readily accessible to and usable by individuals with disabilities.

(g)(1) Within 180 days of the receipt of a timely filed, complete complaint over which it has jurisdiction, the Bureau shall notify the complainant of the results of the investigation in a letter containing:

(i) Findings of fact and conclusions of law;

(ii) A description of a remedy for each violation found; and

(iii) A notice of the right to appeal.

(2) Bureau employees are required to cooperate in the investigation and attempted resolution of complaints. Employees who are required to participate in any investigation under this section shall do so as part of their official duties and during the course of regular duty hours.

(3) If a complaint is resolved informally, the terms of the agreement shall be reduced to writing and made part of the complaint file, with a copy of the agreement provided to the complainant. The written agreement shall describe the subject matter of the complaint and any corrective action to which the parties have agreed.

(h) Appeals of the findings of fact and conclusions of law or remedies must be filed by the complainant within 30 days of receipt from the Bureau of the letter required by § 1072.112(g). The Bureau may extend this time for good cause.

(i) Timely appeals shall be accepted and processed by the Chief Human Capital Officer, who will issue the final agency decision which may include appropriate corrective action to be taken by the Bureau.

(j) The Bureau shall notify the complainant of the results of the appeal within 60 days of the receipt of the timely appeal. If the Bureau determines that it needs additional information from the complainant, it shall have 60 days from the date it received the additional information to make its determination on the appeal.

(k) The time limits cited in paragraphs (g) and (j) of this section may be extended for an individual case when the Chief Human Capital Officer determines there is good cause, based on the particular circumstances of that case, for the extension.

(l) The Bureau may delegate its authority for conducting complaint investigations to other federal agencies or may contract with a nongovernment investigator to perform the investigation, but the authority for making the final determination may not be delegated to another entity.

Dated: June 18, 2012.

Richard Cordray,

Director, Bureau of Consumer Financial Protection.

[FR Doc. 2012–18827 Filed 8–3–12; 8:45 am]

BILLING CODE 4810–AM–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Parts 510, 522, and 524

[Docket No. FDA–2012–N–0002]

New Animal Drugs; Change of Sponsor; Change of Sponsor Address; Azaperone; Miconazole, Polymyxin B, and Prednisolone Suspension

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 two new animal drug applications (NADAs) from Janssen Pharmaceutica NV, to Elanco Animal Health, a Division of Eli Lilly & Co. FDA is also amending the animal drug regulations to reflect a change of sponsor's address for Veterinary Service, Inc.

DATES: This rule is effective August 6, 2012.

FOR FURTHER INFORMATION CONTACT: Steven D. Vaughn, Center for Veterinary Medicine (HFV–100), Food and Drug Administration, 7520 Standish Pl., Rockville, MD 20855, 240–276–8300, email: *steven.vaughn@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Janssen Pharmaceutica NV, Turnhoutseweg 30, B–2340 Beerse, Belgium, has informed FDA that it has transferred ownership of, and all rights and interest in, NADA 115–732 for STRESNIL (azaperone) Injection and NADA 141–298 for SUROLAN (miconazole nitrate, polymyxin B sulfate, prednisolone acetate) Otic Suspension to Elanco Animal Health, a Division of Eli Lilly & Co., Lilly Corporate Center, Indianapolis, IN 46285. Following these changes of sponsorship, Janssen Pharmaceutica NV will no longer be the sponsor of an approved application. Accordingly, the Agency is amending the regulations in 21 CFR 510.600, 522.150, and 524.1445 to reflect the transfer of ownership.

In addition, Veterinary Service, Inc., 416 North Jefferson St., P.O. Box 2467,

Modesto, CA 95354 has informed FDA of a change of address to 4100 Bangs Ave., Modesto, CA 95356. Accordingly, the Agency is amending the regulations in 21 CFR 510.600 to reflect these changes.

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

21 CFR Part 510

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

21 CFR Parts 520 and 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, 522, 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: 21 U.S.C. 321, 331, 351, 352, 353, 360b, 371, 379e.

■ 2. In § 510.600, in the table in paragraph (c)(1), remove the entry for ''Janssen Pharmaceutica NV'' and revise the entry for ''Veterinary Service, Inc.''; and in the table in paragraph (c)(2), remove the entry for ''012758'' and revise the entry for ''033008'' 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
* * * * *	
Veterinary Service, Inc., 4100 Bangs Ave., Modesto, CA 95356	033008
* * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * *	
033008	Veterinary Service, Inc., 4100 Bangs Ave., Modesto, CA 95356.
* * * * *	

PART 522—IMPLANTATION OR INJECTABLE DOSAGE FORM NEW ANIMAL DRUGS

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

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

§ 522.150 [Amended]

■ 4. In paragraph (b) of § 522.150, remove ''012578'' and in its place add ''000986''.

PART 524—OPHTHALMIC AND TOPICAL DOSAGE FORM NEW ANIMAL DRUGS

■ 5. The authority citation for 21 CFR part 524 continues to read as follows:

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

■ 6. In § 524.1445, revise paragraph (b) to read as follows:

§ 524.1445 Miconazole, polymixin B, and prednisolone suspension.

* * * * *

(b) *Sponsor.* See No. 000986 in § 510.600(c) of this chapter.

* * * * *

Dated: August 1, 2012.

Elizabeth Rettie,

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

[FR Doc. 2012–19147 Filed 8–3–12; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

33 CFR Part 165

[Docket Number USCG–2012–0223]

RIN 1625–AA00

Safety Zone; 2012 Ironman US Championship Swim, Hudson River, Fort Lee, NJ

AGENCY: Coast Guard, DHS.

ACTION: Final rule.

SUMMARY: The Coast Guard is establishing a temporary safety zone on the navigable waters of the Hudson River in the vicinity of Englewood Cliffs and Fort Lee, NJ for the 2012 Ironman US Championship swim event. This temporary safety zone is necessary to protect the maritime public and event participants from the hazards associated with swim events. This rule is intended to restrict all vessels and persons from entering into, transiting through, mooring, or anchoring within the safety zone unless authorized by the Captain of the Port (COTP) New York or a designated representative.

DATES: This rule is effective from 6 a.m. until 10 a.m. on August 11, 2012.

ADDRESSES: Documents mentioned in this preamble are part of docket [USCG–2012–0223]. To view documents mentioned in this preamble as being available in the docket, go to *http://www.regulations.gov,* type the docket number in the ''SEARCH'' box and click ''SEARCH.'' Click on Open Docket Folder on the line associated with this rulemaking. You may also visit the Docket Management Facility in Room W12–140 on the ground floor of the Department of Transportation West Building, 1200 New Jersey Avenue SE., Washington, DC 20590, between 9 a.m. and 5 p.m., Monday through Friday, except Federal holidays.

FOR FURTHER INFORMATION CONTACT: If you have questions on this rule, call or email Ensign Kimberly Farnsworth, Coast Guard; Telephone (718) 354–4163, email *Kimberly.A.Farnsworth@uscg.mil.* If you have questions on viewing or submitting material to the docket, call Renee V. Wright, Program Manager, Docket Operations, telephone (202) 366–9826.

SUPPLEMENTARY INFORMATION:

Table of Acronyms

DHS Department of Homeland Security
FR **Federal Register**
CFR Code of Federal Regulations
NPRM Notice of Proposed Rulemaking
COTP Captain of the Port

A. Regulatory History and Information

On June 8, 2012, we published a notice of proposed rulemaking (NPRM) entitled 2012 Ironman US Championship Swim, Hudson River, Fort Lee, NJ in the **Federal Register** (77 FR 34285). We received no comments on the proposed rule. No public meeting was requested and none was held.

Under 5 U.S.C. 553(d)(3), the Coast Guard finds that good cause exists for making this rule effective less than 30 days after publication in the **Federal Register**. This event will occur before 30 days has elapsed after the publication of the rule. The event sponsor is unable and unwilling to postpone this event because the date of this event was