(d) Inspect and modify the routing of the electrical wiring and replace any electrical parts in accordance with the specified portions of Eurocopter Alert Service Bulletin EC155 No. 24A011, Revision 1, dated May 14, 2004. The Director of the Federal Register approved this incorporation by reference in accordance with 5 U.S.C. 552(a) and 1 CFR part 51. Copies may be obtained from American Eurocopter Corporation, 2701 Forum Drive, Grand Prairie, Texas 75053–4005, telephone (972) 641–3460, fax (972) 641–3527. Copies may be inspected at the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, call 202–741–6030, or go to: *http://www.archives.gov/federal_register/code_of_federal_regulations/ibr_locations.html.*

(e) This amendment becomes effective on April 18, 2006.

Note 2: The subject of this AD is addressed in Direction Generale de l'Aviation Civile (France) AD F–2004–057 R1, dated July 21, 2004.

Issued in Fort Worth, Texas, on February 23, 2006.

David A. Downey,

Manager, Rotorcraft Directorate, Aircraft Certification Service.

[FR Doc. 06–2357 Filed 3–13–06; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF ENERGY

Federal Energy Regulatory Commission

18 CFR Part 35

[Docket No. RM06–13–000; Order No. 674]

Conditions for Public Utility Market-Based Rate Authorization Holders

AGENCY: Federal Energy Regulatory Commission, DOE.

ACTION: Final rule: clarification.

SUMMARY: This document clarifies a correction that was published in the **Federal Register** on March 7, 2006. That action amended an effective date for a Final Rule that published in the **Federal Register** on February 27, 2006. The correction document referenced the wrong **Federal Register** page number.

DATES: *Effective Date:* February 27, 2006.

FOR FURTHER INFORMATION CONTACT: Frank Karabetsos, Office of General Counsel, Federal Energy Regulatory Commission, 888 First Street, NE., Washington, DC 20426, (202) 502–8133, *Frank.Karabetsos@ferc.gov.*

SUPPLEMENTARY INFORMATION: In FR Doc. 06–2155, published in the **Federal Register** on March 7, 2006 (71 FR 11304), the correction language cited the wrong page number for the original **Federal Register** document. FR Doc. 06–2155 is clarified and corrected as follows:

On page 11304, column 1, under **SUPPLEMENTARY INFORMATION**, change ''(71 FR 9698)'' to ''(71 FR 9695)'' and ''On page 9698 * * *'' to ''On page 9695''.

Magalie R. Salas,

Secretary.

[FR Doc. 06–2404 Filed 3–13–06; 8:45 am]

BILLING CODE 6717–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

Oral Dosage Form New Animal Drugs; Sulfamerazine, Sulfamethazine, and Sulfaquinoxaline 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 a new animal drug application (NADA) filed by Alpharma Inc. The NADA provides revised labeling for a soluble powder containing sulfamerazine, sulfamethazine, and sulfaquinoxaline used in drinking water of chickens and turkeys as an aid in the control of coccidiosis and acute fowl cholera.

DATES: The rule is effective March 14, 2006.

FOR FURTHER INFORMATION CONTACT: Dianne T. McRae, Center for Veterinary Medicine (HFV–104), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–0161, e-mail: *dianne.mcrae@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Alpharma Inc., One Executive Drive, Fort Lee, NJ 07024, filed NADA 100–094 for POULTRYSULFA (sulfamerazine, sulfamethazine, and sulfaquinoxaline) Antimicrobial Soluble Powder, an over-the-counter product used in the drinking water of chickens and turkeys as an aid in the control of coccidiosis and acute fowl cholera. The NADA relies on the National Academy of Sciences/National Research Council (NAS/NRC), Drug Efficacy Study Group's (DESI) effectiveness evaluation and subsequent FDA conclusions. The findings were published in the **Federal Register** of July 5, 1984 (49 FR 27543).

Using the official analytical method of detection, residues of sulfamethazine and sulfamerazine in edible tissues co-elute and cannot be quantified individually. There are no products containing only sulfamerazine approved for use in chickens or turkeys. Therefore, a tolerance for sulfamerazine residues in edible tissues of chickens or turkeys is not established at this time.

Products that comply with the NAS/NRC findings and FDA's conclusions regarding those findings are eligible for immediate copying under the Generic Animal Drug and Patent Term Restoration Act (GADPTRA) (see the eighth in a series of policy letters issued to facilitate implementation of GADPTRA that published in the **Federal Register** of August 21, 1991 (56 FR 41561), available online at *http://www.fda.gov/cvm/Documents/8thltr.doc*).

The NADA is approved as of February 2, 2006, and part 520 (21 CFR part 520) is amended by adding new § 520.2218 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 Division of Dockets Management (HFA–305), Food and Drug Administration, 5630 Fishers Lane, rm. 1061, Rockville, MD 20852, between 9 a.m. and 4 p.m., Monday through Friday.

FDA has determined under 21 CFR 25.33(a)(1) 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.

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 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: