§ 240.12a–11 Exemption of security-based swaps sold in reliance on Securities Act of 1933 Rule 240 (§ 230.240) from section 12(a) of the Act.

(a) The provisions of Section 12(a) of the Act (15 U.S.C. 78l(a)) do not apply to any security-based swap offered and sold in reliance on Rule 240 under the Securities Act of 1933.

(b) This rule will expire on the compliance date for final rules that the Commission may adopt further defining both the terms security-based swap and eligible contract participant. In such event, the Commission will publish a rule removing this section from 17 CFR part 240 or modifying it as appropriate.

■ 5. Section 240.12h–1 is amended by adding paragraph (i) to read as follows:

§ 240.12h–1 Exemptions from registration under section 12(g) of the Act.

* * * * *

(i) Any security-based swap offered and sold in reliance on Rule 240 under the Securities Act of 1933. This rule will expire on the compliance date for final rules that the Commission may adopt further defining both the terms *security-based swap* and *eligible contract participant.* In such event, the Commission will publish a rule removing this paragraph (i) from 17 CFR part 240 or modifying it as appropriate.

PART 260—GENERAL RULES AND REGULATIONS, TRUST INDENTURE ACT OF 1939

■ 6. The authority citation for Part 260 continues to read as follows:

Authority: 15 U.S.C. 77eee, 77ggg, 77nnn, 77sss, 78ll(d), 80b–3, 80b–4, and 80b–11.

■ 7. Section 260.4d–12 is added to read as follows:

§ 260.4d–12 Exemption for security-based swaps offered and sold in reliance on Securities Act of 1933 Rule 240 (§ 230.240).

Any security-based swap offered and sold in reliance on Rule 240 of this chapter (17 CFR 230.240), whether or not issued under an indenture, is exempt from the Act. This rule will expire on the compliance date for final rules that the Commission may adopt further defining both the terms *security-based swap* and *eligible contract participant.* In such event, the Commission will publish a rule removing this section from 17 CFR part 260 or modifying it as appropriate.

By the Commission.

Dated: July 1, 2011.

Elizabeth M. Murphy,

Secretary.

[FR Doc. 2011–17039 Filed 7–8–11; 8:45 am]

BILLING CODE 8011–01–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 510

[Docket No. FDA–2011–N–0003]

New Animal Drugs; Change of Sponsor's Name and Address

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's name from Alpharma, LLC, to Alpharma, LLC, a wholly owned subsidiary of Pfizer, Inc. The sponsor's mailing address will also be changed.

DATES: This rule is effective July 11, 2011.

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, *e-mail: steven.vaughn@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: Alpharma, LLC, 400 Crossing Blvd., Bridgewater, NJ 08807 has informed FDA of a change of name and mailing address to Alpharma, LLC, a wholly owned subsidiary of Pfizer, Inc., 235 East 42d St., New York, NY 10017. Accordingly, the Agency is amending the regulations in 21 CFR 510.600(c) 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 in 21 CFR Part 510

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

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 510 is 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), revise the entry for ''Alpharma LLC''; and in the table in paragraph (c)(2), revise the entry for ''046573'' 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
* * * * *	
Alpharma, LLC, a wholly owned subsidiary of Pfizer, Inc., 235 East 42d St., New York, NY 10017	046573
* * * * *	

(2) * * *

Drug labeler code	Firm name and address
* * * * *	
046573	Alpharma, LLC, a wholly owned subsidiary of Pfizer, Inc., 235 East 42d St., New York, NY 10017
* * * * *	

Dated: July 1, 2011.

Elizabeth Rettie,

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

[FR Doc. 2011–17292 Filed 7–8–11; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

24 CFR Part 3500

[Docket No. FR–5180–F–07]

RIN 2502–AH85

Real Estate Settlement Procedures Act (RESPA): Technical Corrections and Clarifying Amendments

AGENCY: Office of the Assistant Secretary for Housing—Federal Housing Commissioner, HUD.

ACTION: Final rule.

SUMMARY: This final rule makes technical corrections and certain clarifying amendments to HUD's RESPA regulations promulgated by a final rule published on November 17, 2008. The majority of the regulations promulgated by the November 17, 2008, final rule became applicable on January 1, 2010. Now that the regulations have been in