method to be approved, the repair must meet the certification basis of the airplane, and the approval must specifically refer to this AD.

(4) AMOCs approved previously in accordance with AD 2010–20–08, Amendment 39–16442 (75 FR 61337, October 5, 2010), are approved as AMOCs for the corresponding provisions of paragraphs (g) through (m) of this AD.

(5) AMOCs approved previously in accordance with AD 2010–20–08, Amendment 39–16442 (75 FR 61337, October 5, 2010), that have post-repair inspections are approved as AMOCs for the corresponding provisions of paragraph (o) of this AD for the repaired area only.

(t) Related Information

For more information about this AD, contact Nathan Weigand, Aerospace Engineer, Airframe Branch, ANM–120S, FAA, Seattle Aircraft Certification Office (ACO), 1601 Lind Avenue SW., Renton, WA 98057–3356; phone: 425–917–6428; fax: 425–917–6590; email: *Nathan.P.Weigand@faa.gov.*

(u) Material Incorporated by Reference

(1) The Director of the Federal Register approved the incorporation by reference (IBR) of the service information listed in this paragraph under 5 U.S.C. 552(a) and 1 CFR part 51.

(2) You must use this service information as applicable to do the actions required by this AD, unless the AD specifies otherwise.

(3) The following service information was approved for IBR on October 22, 2013.

(i) Boeing Alert Service Bulletin 747–53A2450, Revision 7, dated November 2, 2011.

(ii) Reserved.

(4) The following service information was approved for IBR on November 9, 2010 (75 FR 61337, October 5, 2010).

(i) Boeing Alert Service Bulletin 747–53A2450, Revision 5, dated January 29, 2009.

(ii) Reserved.

(5) The following service information was approved for IBR on September 12, 2001 (66 FR 441440, August 8, 2001).

(i) Boeing Alert Service Bulletin 747–53A2450, Revision 2, including Appendix A, dated January 4, 2001.

(ii) Reserved.

(6) For Boeing service information identified in this AD, contact Boeing Commercial Airplanes, Attention: Data & Services Management, P.O. Box 3707, MC 2H–65, Seattle, WA 98124–2207; telephone 206–544–5000, extension 1; fax 206–766–5680; Internet *https://www.myboeingfleet.com.*

(7) You may view this service information at FAA, Transport Airplane Directorate, 1601 Lind Avenue SW., Renton, WA. For information on the availability of this material at the FAA, call 425–227–1221.

(8) You may view this service information that is incorporated by reference 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/cfr/ibr-locations.html.*

Issued in Renton, Washington, on August 16, 2013.

Jeffrey E. Duven,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. 2013–22408 Filed 9–16–13; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

[Docket No. FDA–2013–N–0002]

Oral Dosage Form New Animal Drugs; Amprolium; Meloxicam

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 actions for new animal drug applications (NADAs) and abbreviated new animal drug applications (ANADAs) during August 2013. FDA is also informing the public of the availability of summaries of the basis of approval and of environmental review documents, where applicable.

DATES: This rule is effective September 17, 2013.

FOR FURTHER INFORMATION CONTACT: George K. Haibel, Center for Veterinary Medicine (HFV–6), Food and Drug Administration, 7519 Standish Pl., Rockville, MD 20855, 240–276–9019, *george.haibel@fda.hhs.gov.*

SUPPLEMENTARY INFORMATION: FDA is amending the animal drug regulations to reflect approval actions for NADAs and ANADAs during August 2013, as listed in table 1. In addition, FDA is informing the public of the availability, where applicable, of documentation of environmental review required under the National Environmental Policy Act (NEPA) and, for actions requiring review of safety or effectiveness data, summaries of the basis of approval (FOI Summaries) under the Freedom of Information Act (FOIA). These public documents 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. Persons with access to the Internet may obtain these documents at the CVM FOIA Electronic Reading Room: *http://www.fda.gov/AboutFDA/CentersOffices/OfficeofFoods/CVM/CVMFOIAElectronicReadingRoom/default.htm.*

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.

TABLE 1—ORIGINAL AND SUPPLEMENTAL NADAS AND ANADAS APPROVED DURING AUGUST 2013

NADA/ ANADA	Sponsor	New Animal Drug Product Name	Action	21 CFR Section	FOIA Summary	NEPA Review
200–514 ...	Phibro Animal Health Corp., GlenPointe Centre East, 3d floor, 300 Frank W. Burr Blvd., Suite 21, Teaneck, NJ 07666.	BOVIPROL (amprolium) 9.6% Oral Solution.	Original approval as a generic copy of NADA 13–149.	520.100	Yes	CE [1].
200–550 ...	Ceva Sante Animale, 10 Avenue de la Ballastiére 33500 Libourne, France.	MELOXIDYL (meloxicam) Oral Suspension.	Original approval as a generic copy of NADA 141–213.	520.1350	Yes	CE [1].

[1] The Agency has determined under 21 CFR 25.33(a)(1) that this action is categorically excluded (CE) from the requirement to submit an environmental assessment or an environmental impact statement because it is of a type that does not individually or cumulatively have a significant effect on the human environment.

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:

PART 520—ORAL DOSAGE FORM NEW ANIMAL DRUGS

■ 1. The authority citation for 21 CFR part 520 continues to read as follows:

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

■ 2. In § 520.100, revise paragraph (b)(2) to read as follows:

§ 520.100 Amprolium.

* * * * *

(b) * * *

(2) No. 066104 for use of product described in paragraph (a)(1) of this section as in paragraph (d) of this section.

* * * * *

§ 520.1350 [Redesignated as § 520.1367]

■ 3. Redesignate § 520.1350 as § 520.1367.

■ 4. Amend newly redesignated § 520.1367 by revising paragraphs (a) and (b) to read as follows:

§ 520.1367 Meloxicam.

(a) *Specifications*—(1) Each milliliter of suspension contains 0.5 milligrams (mg) meloxicam.

(2) Each milliliter of suspension contains 1.5 mg meloxicam.

(b) *Sponsors.* See sponsors in § 510.600(c) of this chapter for uses as in paragraph (c) of this section:

(1) No. 000010 for use of the products described in paragraph (a) of this section; and

(2) No. 013744 for use of the product described in paragraph (a)(2) of this section.

* * * * *

Dated: September 11, 2013.

Bernadette Dunham,

Director, Center for Veterinary Medicine.

[FR Doc. 2013–22523 Filed 9–16–13; 8:45 am]

BILLING CODE 4160–01–P

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

24 CFR Parts 5 and 202

[Docket No. FR–5536–F–02]

RIN 2502–AJ00

Federal Housing Administration (FHA) Approval of Lending Institutions and Mortgagees: Streamlined Reporting Requirements for Small Supervised Lenders and Mortgagees

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

ACTION: Final rule.

SUMMARY: This rule streamlines the FHA financial statement reporting requirements for lenders and mortgagees who are supervised by federal banking agencies and whose consolidated assets do not meet the thresholds set by their supervising federal banking agencies for submission of audited financial statements (currently set at $500 million in consolidated assets). HUD's regulations currently require all supervised lenders and mortgagees to submit annual audited financial statements as a condition of FHA lender approval and recertification. Through this rule, in lieu of the annual audited financial statements, small supervised lenders and mortgagees would be required to submit their unaudited financial regulatory reports that align with their fiscal year ends and are required to be submitted to their supervising federal banking agencies. Small supervised lenders and mortgagees would only be required to submit audited financial statements if HUD determines that the supervised lenders or mortgagees pose heightened risk to the FHA insurance fund.

This rule does not impact FHA's annual audited financial statements submission requirement for nonsupervised and large supervised lenders and mortgagees. The rule also does not impact those supervised lenders and mortgagees with consolidated assets in an amount that requires that lenders or mortgagees submit audited financial statements to their respective supervising federal banking agencies. Additionally, this final rule, consistent with the proposed rule, makes three technical changes to current regulations regarding reporting requirements for FHA-approved supervised lenders and mortgagees.

DATES: *Effective Date:* October 17, 2013.

FOR FURTHER INFORMATION CONTACT: Richard Toma, Deputy Director, Office of Lender Activities and Program Compliance, Office of Housing, Department of Housing and Urban Development, 490 L'Enfant Plaza East SW., Room P3214, Washington, DC 20024–8000; telephone number 202–708–1515 (this is not a toll-free number). Persons with hearing or speech impairments may access this number through TTY by calling the toll-free Federal Relay Service at 800–877–8339.

SUPPLEMENTARY INFORMATION:

I. Background

On April 18, 2013 (78 FR 23178), HUD published for public comment a proposed rule that would streamline reporting requirements and relieve burden on small supervised lenders and mortgagees.[1] HUD's regulations, at 24 CFR 202.5(g), require that all lenders and mortgagees provide annual audited financial statements within 90 days of their fiscal year ends. Small supervised lenders and mortgagees, however, are not required by their supervising federal banking agencies to submit audited financial statements, but are permitted to submit unaudited financial regulatory reports. These unaudited financial regulatory reports currently include a consolidated or fourth quarter Report of Condition and Income (Federal Financial Institutions Examination Council forms 031 and 041, also known as the ''Call Report''), a consolidated or fourth quarter Thrift Financial Report, and a consolidated or fourth quarter NCUA Call Report (NCUA Form 5300 or 5310). The HUD requirement is therefore inconsistent with that of the federal banking agencies, and has the potential to impose a potentially financially prohibitive requirement on small supervised lenders and mortgagees who wish to participate in FHA programs. While HUD takes its counterparty risk management responsibilities seriously, HUD also seeks to balance its management of risk with the execution of its mission.

Upon reconsideration, HUD has determined that the financial regulatory reports required by the federal banking agencies contain sufficient information for HUD to ensure that small supervised lenders and mortgagees are suitably capitalized to meet potential needs associated with their participation in

[1] The term ''small supervised lenders and mortgagees'' refers to those lenders and mortgagees supervised by the Board of Governors of the Federal Reserve System; the Federal Deposit Insurance Corporation (FDIC); and the National Credit Union Administration (NCUA) (collectively, the ''federal banking agencies'') whose consolidated assets do not meet the thresholds set by their supervising federal banking agencies for submission of audited financial statements (currently set at $500 million in consolidated assets).