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DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 39

[Docket No. FAA-2011-0959; Directorate Identifier 2011-NE-25-AD; Amendment 39-16970; AD 2012-04-14]

RIN 2120-AA64

Airworthiness Directives; Rolls-Royce plc Turbofan Engines

AGENCY: Federal Aviation Administration (FAA), DOT.

ACTION: Final rule; correction.

SUMMARY: The FAA is correcting an airworthiness directive (AD) that published in the **Federal Register**. That AD applies to RB211-Trent 800 series turbofan engines. The last comment response in the preamble and the first sentence of regulatory text paragraph (g)(1) are incorrect. The repetitive inspection interval should be 2,000 flight cycles, not 1,000 flight cycles. This document corrects those errors. In all other respects, the original document remains the same.

DATES: This final rule is effective April 11, 2012.

ADDRESSES: You may examine the AD docket on the Internet at <http://www.regulations.gov>; or in person at the Docket Management Facility between 9 a.m. and 5 p.m., Monday through Friday, except Federal holidays. The AD docket contains this AD, the regulatory evaluation, any comments received, and other information. The address for the Docket Office (phone: 800-647-5527) is Document Management Facility, U.S. Department of Transportation, Docket Operations, M-30, West Building Ground Floor, Room W12-140, 1200 New Jersey Avenue SE., Washington, DC 20590.

FOR FURTHER INFORMATION CONTACT:

Alan Strom, Aerospace Engineer, Engine

Certification Office, FAA, Engine & Propeller Directorate, 12 New England Executive Park, Burlington, MA 01803; email: alan.strom@faa.gov; phone: 781-238-7143; fax: 781-238-7199.

SUPPLEMENTARY INFORMATION: AD 2012-04-14, Amendment 39-16970 (77 FR 13485, March 7, 2012), currently requires inspecting the front combustion liner head section for cracking, and if found cracked, removing the front combustion liner head section from service at the next shop visit.

As published, the last comment response in the preamble, and the first sentence of regulatory text paragraph (g)(1), are incorrect. No other part of the preamble or regulatory text has been changed; therefore, only the changed portions of the final rule is being published in the **Federal Register**.

The effective date of this AD remains April 11, 2012.

Correction of Non-Regulatory Text

In the **Federal Register** of March 7, 2012, AD 2012-04-14; Amendment 39-16970, is corrected to read as follows:

On page 13486, in the 3rd column, under the heading Need to Show All Acceptable Means of Completing the On-Wing Inspection, the 2nd sentence in the 1st paragraph is corrected to read "We changed the 2nd sentence of paragraphs (f)(1) and (g)(1) of the proposed AD from:"

On page 13486, in the 3rd column, under the heading Need to Show All Acceptable Means of Completing the On-Wing Inspection, the 1st sentence in the 2nd and 3rd paragraphs, is deleted.

Correction of Regulatory Text

§ 39.13 [Corrected]

■ In the **Federal Register** of March 7, 2012, AD 2012-04-14; Amendment 39-16970, on page 13487, in the first column, in paragraph (g)(1), the first sentence is corrected to read as follows:

* * * * *

(g)(1) At intervals not to exceed 2,000 FCs, inspect the front combustion liner head section for cracking.

* * * * *

Issued in Burlington, Massachusetts, on March 30, 2012.

Colleen D'Alessandro,

Assistant Manager, Engine & Propeller Directorate, Aircraft Certification Service.

[FR Doc. 2012-8289 Filed 4-6-12; 8:45 am]

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DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 520

[Docket No. FDA-2012-N-0002]

Oral Dosage Form New Animal Drugs; Change of Sponsor; Lincomycin Hydrochloride Soluble Powder; Penicillin G Potassium in Drinking Water; Tetracycline Powder

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; technical amendment.

SUMMARY: The Food and Drug Administration (FDA) is amending the animal drug regulations to reflect a change of sponsor for three abbreviated new animal drug applications (ANADAs) for lincomycin hydrochloride; penicillin G potassium, USP; and tetracycline hydrochloride soluble powders administered in drinking water from Teva Animal Health, Inc., to Quo Vademus, LLC.

DATES: This rule is effective April 9, 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: Teva Animal Health, Inc., 3915 South 48th Street Ter., St. Joseph, MO 64503, has informed FDA that it has transferred ownership of, and all rights and interest in, ANADA 200-136 for Tetracycline Hydrochloride Soluble Powder 324; ANADA 200-303 for Lincomycin Hydrochloride Soluble Powder; and ANADA 200-347 for Penicillin G Potassium, USP, all soluble powders administered in drinking water to Quo Vademus, LLC, 277 Faison West McGowan Rd., Kenansville, NC 28349. Accordingly, the Agency is amending the regulations in part 520 (21 CFR part 520) to reflect the transfer of ownership and a current format.

In addition FDA has noticed two errors in § 520.1696 *Penicillin oral dosage forms*. At this time, § 520.1696a is being removed because no sponsor is listed, and an obsolete drug labeler code is being removed from § 520.1696d.

These actions are being taken to improve the accuracy of the regulations.

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:

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 § 520.1263, revise the section heading to read as follows:

§ 520.1263 Lincomycin.

* * * * *

■ 3. In § 520.1263c, revise the section heading and paragraph (b) to read as follows:

§ 520.1263c Lincomycin powder.

* * * * *

(b) *Sponsors.* See Nos. 000009, 046573, 054925, 061623, and 076475 in § 510.600(c) of this chapter for use as in paragraph (d) of this section.

* * * * *

■ 4. In § 520.1696, revise the section heading to read as follows:

§ 520.1696 Penicillin.

* * * * *

§ 520.1696a [Removed and Reserved]

■ 5. Remove and reserve § 520.1696a.

■ 6. In § 520.1696b, revise the section heading, paragraphs (a) and (b), and the heading for paragraph (c) to read as follows:

§ 520.1696b Penicillin G powder.

(a) *Specifications.* Each gram of powder contains penicillin G potassium equivalent to 1.54 million units of penicillin G.

(b) *Sponsors.* See Nos. 010515, 046573, 053501, 059320, 061623 and 076475 in § 510.600(c) of this chapter.

(c) *Conditions of use in turkeys—*

* * * * *

■ 7. In § 520.1696c, revise the section heading and remove and reserve paragraph (c) to read as follows:

§ 520.1696c Penicillin V powder.

* * * * *

■ 8. In § 520.1696d, revise the section heading and paragraph (b) and remove and reserve paragraph (c) to read as follows:

§ 520.1696d Penicillin V tablets.

* * * * *

(b) *Sponsors.* See Nos. 050604 and 053501 in § 510.600(c) of this chapter.

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■ 9. In § 520.2345, revise the section heading to read as follows:

§ 520.2345 Tetracycline.

* * * * *

■ 10. In § 520.2345d, revise paragraph (b)(4) to read as follows:

§ 520.2345d Tetracycline powder.

* * * * *

(b) * * *

(4) Nos. 054925, 057561, 061623, and 076475: 324 grams per pound as in paragraph (d) of this section.

* * * * *

Dated: March 27, 2012.

William T. Flynn,

Acting Director, Center for Veterinary Medicine.

[FR Doc. 2012–8322 Filed 4–6–12; 8:45 am]

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37 CFR Parts 201 and 202

[Docket No. 2011–8]

Discontinuance of Form CO in Registration Practices

Correction

In rule document 2012–7429 appearing on pages 18705–18707 in the issue of March 28, 2012, make the following corrections:

1. On page 18706, in the third column, in the 17th line from the bottom, “□.” should read “Ⓢ.”.

§ 202.2 [Corrected]

■ 2. On page 18707, in § 202.2, in the first column, in amendatory instruction 4, in the third line, “□” should read “Ⓢ”.

[FR Doc. C1–2012–7429 Filed 4–6–12; 8:45 am]

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DEPARTMENT OF HOMELAND SECURITY

Federal Emergency Management Agency

44 CFR Part 64

[Docket ID FEMA–2012–0003; Internal
Agency Docket No. FEMA–8225]

Suspension of Community Eligibility

AGENCY: Federal Emergency
Management Agency, DHS.

ACTION: Final rule.

SUMMARY: This rule identifies communities where the sale of flood insurance has been authorized under the National Flood Insurance Program (NFIP) that are scheduled for suspension on the effective dates listed within this rule because of noncompliance with the floodplain management requirements of the program. If the Federal Emergency Management Agency (FEMA) receives documentation that the community has adopted the required floodplain management measures prior to the effective suspension date given in this rule, the suspension will not occur and a notice of this will be provided by publication in the **Federal Register** on a subsequent date.

DATES: The effective date of each community’s scheduled suspension is the third date (“Susp.”) listed in the third column of the following tables.

FOR FURTHER INFORMATION CONTACT: If you want to determine whether a particular community was suspended on the suspension date or for further information, contact David Stearrett, Federal Insurance and Mitigation Administration, Federal Emergency Management Agency, 500 C Street SW., Washington, DC 20472, (202) 646–2953.

SUPPLEMENTARY INFORMATION: The NFIP enables property owners to purchase Federal flood insurance that is not otherwise generally available from private insurers. In return, communities agree to adopt and administer local floodplain management aimed at protecting lives and new construction from future flooding. Section 1315 of the National Flood Insurance Act of 1968, as amended, 42 U.S.C. 4022, prohibits the sale of NFIP flood insurance unless an appropriate public body adopts adequate floodplain management measures with effective enforcement measures. The communities listed in this document no longer meet that statutory requirement for compliance with program regulations, 44 CFR part 59.