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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.

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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.