part 51. Contact British Aerospace Regional Aircraft American Support, 13850 Mclearen Road, Herndon, Virginia 20171, for a copy of this service information. You may review copies at the Docket Management Facility, U.S. Department of Transportation, 400 Seventh Street, SW., Room PL–401, Nassif Building, Washington, DC; on the Internet at *http://dms.dot.gov*; or at the National Archives and Records Administration (NARA). For information on the availability of this material at the NARA, call (202) 741–6030, or go to *http://www.archives.gov/federal_register/code_of_federal_regulations/ibr_locations.html.*

Issued in Renton, Washington, on December 1, 2006.

Kevin M. Mullin,

Acting Manager, Transport Airplane Directorate, Aircraft Certification Service.

[FR Doc. E6–20952 Filed 12–11–06; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF TRANSPORTATION

Federal Aviation Administration

14 CFR Part 39

[Docket No. 2004–NE–19–AD; Amendment 39–13197; AD 2004–26–05]

RIN 2120–AA64

Airworthiness Directives; Rolls-Royce plc RB211–524 Series Turbofan Engines; Correction

AGENCY: Federal Aviation Administration, DOT.

ACTION: Final rule; correction.

SUMMARY: This document makes a correction to airworthiness directive (AD) 2004–26–05 applicable to certain Rolls-Royce plc (RR) RB211–524 series turbofan engines that was published in the **Federal Register** on January 5, 2005. The part number UL29916 in the Applicability section is incorrect. This document corrects that part number. In all other respects, the original document remains the same.

DATES: *Effective Date:* December 12, 2006.

FOR FURTHER INFORMATION CONTACT: Ian Dargin, Aerospace Engineer, Engine Certification Office, FAA, Engine and Propeller Directorate, 12 New England Executive Park, Burlington, MA 01803–5299; telephone (781) 238–7178; fax (781) 238–7199.

SUPPLEMENTARY INFORMATION: A final rule airworthiness directive FR Doc, 05–85 applicable to RR RB211–524 series turbofan engines, was published in the **Federal Register** on January 5, 2005 (70 FR 681). The following correction is needed:

§ 39.13 [Corrected]

■ On page 682, in the first column, in the PART 39—AIRWORTHINESS DIRECTIVES Section, in the Applicability paragraph, in the second line, ''UL29916'' is corrected to read ''UL26916''.

Issued in Burlington, Massachusetts, on December 5, 2006.

Diane M. Cook,

Acting Manager, Engine and Propeller Directorate, Aircraft Certification Service.

[FR Doc. E6–21122 Filed 12–11–06; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 558

New Animal Drugs For Use in Animal Feeds; Tylosin

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 supplemental new animal drug application (NADA) filed by Elanco Animal Health, A Division of Eli Lilly & Co. The supplemental NADA provides for an alternate feeding regimen for tylosin phosphate in Type C medicated swine feeds used for the control of swine proliferative enteropathies.

DATES: This rule is effective December 12, 2006.

FOR FURTHER INFORMATION CONTACT: Joan C. Gotthardt, Center for Veterinary Medicine (HFV–130), Food and Drug Administration, 7500 Standish Pl., Rockville, MD 20855, 301–827–7571, e-mail: *joan.gotthardt@fda.hhs.gov*.

SUPPLEMENTARY INFORMATION: Elanco Animal Health, A Division of Eli Lilly & Co., Lilly Corporate Center, Indianapolis, IN 46285, filed a supplement to NADA 12–491 that provides for use of TYLAN (tylosin phosphate) Type A medicated articles. The supplement provides for an alternate feeding regimen for the control of swine proliferative enteropathies (ileitis) associated with *Lawsonia intracellularis*. In addition, Elanco Animal Health revised the names of other enteric pathogens of swine to reflect changes in the scientific nomenclature for these bacteria. The supplemental NADA is approved as of November 7, 2006, and the regulations in 21 CFR 558.625 are amended 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.

Under section 512(c)(2)(F)(iii) of the Federal Food, Drug, and Cosmetic Act (21 U.S.C. 360b(c)(2)(F)(iii)), this approval qualifies for 3 years of marketing exclusivity beginning November 7, 2006.

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 558

Animal drugs, Animal feeds.

■ 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 558 is amended as follows:

PART 558—NEW ANIMAL DRUGS FOR USE IN ANIMAL FEEDS

■ 1. The authority citation for 21 CFR part 558 continues to read as follows:

Authority: 21 U.S.C. 360b, 371.

■ 2. In § 558.625, revise paragraphs (f)(1)(i)(*b*), (f)(1)(vi)(*b*)(*1*), (f)(1)(vi)(*c*)(*1*), and (f)(1)(vi)(*e*)(*1*) to read as follows:

§ 558.625 Tylosin.

* * * * *

(f) * * *

(1) * * *

(i) * * *

(*b*) *Indications for use.* For reduction of incidence of liver abscesses caused by *Fusobacterium necrophorum* and *Arcanobacterium (Actinomyces) pyogenes*.

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(vi) * * *