

Listed diseases, infections, and infestations.

(3) Changes are made in Veterinary Services (VS) regulations.

(4) Changes are made on the National Veterinary Stockpile (NVS) list, USDA Select Agents and Toxins List, or Centers for Disease Control and Prevention (CDC) Category A, B, or C Bioterrorism Agents/Diseases list.

(5) Changes or additions are requested by stakeholders. Stakeholders must submit change requests in writing via postal mail or email using the contact information provided in paragraph (a) of this section. Written requests must include a justification for the proposed change. Examples of justifications can be found in the NLRAD System Standards Document, available on the APHIS website at [https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/nlrاد/ct\\_national\\_list\\_reportable\\_animal\\_diseases](https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/nlrاد/ct_national_list_reportable_animal_diseases).

(d) *Reporting.* The following reporting procedures will be required:

(1) *Notifiable diseases.* Any animal health professional with knowledge of occurrence or suspected occurrence of an animal disease, disease agent, or condition listed as notifiable in the NLRAD must immediately report such identification or suspicion to both APHIS and the State where the livestock is located. Reporting to APHIS may be accomplished as described on the NLRAD website available at: [https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/nlrاد/ct\\_national\\_list\\_reportable\\_animal\\_diseases](https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/nlrاد/ct_national_list_reportable_animal_diseases), or by contacting a local APHIS office.<sup>1</sup> Reporting to the State should be to the State animal health official listed at [https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/sa\\_disease\\_reporting/ct\\_usda\\_aphis\\_animal\\_health](https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/sa_disease_reporting/ct_usda_aphis_animal_health) for the State in question.

(2) *Monitored diseases.* (i) Laboratories must report occurrence information of confirmed cases of an animal disease or condition listed as monitored in the NLRAD on a monthly basis to the State where the animal is located by contacting the State animal health official listed at [https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/sa\\_disease\\_reporting/ct\\_usda\\_aphis\\_animal\\_health](https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/sa_disease_reporting/ct_usda_aphis_animal_health).

<sup>1</sup> Contact information for APHIS offices can be found on the APHIS website at <https://www.aphis.usda.gov/aphis/banner/contactus>, or in the local phone directory (listed under Animal and Plant Health Inspection Service (APHIS), Veterinary Services).

(ii) States must report information of confirmed cases of an animal disease or condition listed as monitored in the NLRAD on a monthly basis to APHIS through the Designated Information Technology System available on the APHIS website at: [https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/sa\\_disease\\_reporting/ct\\_usda\\_aphis\\_animal\\_health](https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/sa_disease_reporting/ct_usda_aphis_animal_health).

(3) *Additional guidance.* Additional reporting information, including background and resources to support reporting efforts, can be found in the NLRAD System Standards Document available on the APHIS website at: [https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/nlrاد/ct\\_national\\_list\\_reportable\\_animal\\_diseases](https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/nlrاد/ct_national_list_reportable_animal_diseases). Revisions to the NLRAD System Standards Document, other than updates to the NLRAD described in paragraph (c) of this section, will be announced to the public as needed through the publication of a notice in the **Federal Register**. The notice will also provide for a public comment period.

(4) *Alternative reporting structures.* The regulations in this paragraph (d) do not supersede any alternative reporting structure that APHIS may require through issuance of a general regulation or Federal Order.

#### **PART 161—REQUIREMENTS AND STANDARDS FOR ACCREDITED VETERINARIANS AND SUSPENSION OR REVOCATION OF SUCH ACCREDITATION**

■ 2. The authority for part 161 continues to read as follows:

**Authority:** 7 U.S.C. 8301–8317; 15 U.S.C. 1828; 7 CFR 2.22, 2.80, and 371.4.

■ 3. In § 161.4, paragraph (f) is revised to read as follows:

##### **§ 161.4 Standards for accredited veterinarian duties.**

\* \* \* \* \*

(f) An accredited veterinarian shall immediately report all diagnosed or suspected cases of any animal disease, disease agent, or condition classified as notifiable in the National List of Reportable Animal Diseases (NLRAD) in accordance with reporting provisions set forth in § 57.2 of this chapter. The NLRAD can be viewed on the APHIS website at: [https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/sa\\_disease\\_reporting/ct\\_usda\\_aphis\\_animal\\_health](https://www.aphis.usda.gov/aphis/ourfocus/animalhealth/monitoring-and-surveillance/sa_disease_reporting/ct_usda_aphis_animal_health).

\* \* \* \* \*

Done in Washington, DC, this 26th day of March 2020.

**Mark Davidson,**

*Acting Administrator, Animal and Plant Health Inspection Service.*

[FR Doc. 2020–06697 Filed 4–1–20; 8:45 am]

**BILLING CODE 3410–34–P**

## **NUCLEAR REGULATORY COMMISSION**

### **10 CFR Chapter I**

[NRC–2017–0214]

#### **Retrospective Review of Administrative Requirements**

**AGENCY:** Nuclear Regulatory Commission.

**ACTION:** Request for comment; extension of comment period.

**SUMMARY:** On February 4, 2020, the U.S. Nuclear Regulatory Commission (NRC) requested input from its licensees and members of the public to identify outdated or duplicative administrative requirements that may be eliminated without an adverse effect on public health or safety, common defense and security, protection of the environment, or regulatory efficiency and effectiveness. The public comment period originally was scheduled to close on April 6, 2020. The NRC has decided to extend the public comment period to allow more time for members of the public to develop and submit their comments.

**DATES:** The comment period for the document published on February 4, 2020 (85 FR 6103) is extended. Comments should be filed no later than May 6, 2020. Comments received after this date will be considered, if it is practical to do so, but the NRC is able to ensure consideration only for comments received on or before this date.

**ADDRESSES:** You may submit comments by any of the following methods:

- *Federal Rulemaking website:* Go to <https://www.regulations.gov> and search for Docket ID NRC–2017–0214. Address questions about NRC dockets to Carol Gallagher; telephone: 301–415–3463; email: [Carol.Gallagher@nrc.gov](mailto:Carol.Gallagher@nrc.gov). For technical questions contact the individuals listed in the **FOR FURTHER INFORMATION CONTACT** section of this document.

- *Email comments to:* [Rulemaking.Comments@nrc.gov](mailto:Rulemaking.Comments@nrc.gov). If you do not receive an automatic email reply confirming receipt, then contact us at 301–415–1677.

- *Fax comments to:* Secretary, U.S. Nuclear Regulatory Commission at 301–415–1101.

- *Mail comments to:* Secretary, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001, ATTN: Rulemakings and Adjudications Staff.

- *Hand deliver comments to:* 11555 Rockville Pike, Rockville, Maryland 20852, between 7:30 a.m. and 4:15 p.m. (Eastern Time) Federal workdays; telephone: 301–415–1677.

For additional direction on obtaining information and submitting comments, see “Obtaining Information and Submitting Comments” in the **SUPPLEMENTARY INFORMATION** section of this document.

**FOR FURTHER INFORMATION CONTACT:** Andrew G. Carrera, telephone: 301–415–1078; email: [Andrew.Carrera@nrc.gov](mailto:Andrew.Carrera@nrc.gov); or Pamela Noto, telephone: 301–415–6795; email: [Pamela.Noto@nrc.gov](mailto:Pamela.Noto@nrc.gov). Both are staff of the Office of Nuclear Material Safety and Safeguards, U.S. Nuclear Regulatory Commission, Washington, DC 20555–0001.

**SUPPLEMENTARY INFORMATION:**

**I. Obtaining Information and Submitting Comments**

*A. Obtaining Information*

Please refer to Docket ID NRC–2017–0214 when contacting the NRC about the availability of information for this action. You may obtain publicly available information related to this action by any of the following methods:

- Federal Rulemaking website: Go to <https://www.regulations.gov> and search for Docket ID NRC–2017–0214.

- NRC’s Agencywide Documents Access and Management System (ADAMS): You may obtain publicly available documents online in the ADAMS Public Documents collection at <https://www.nrc.gov/reading-rm/adams.html>. To begin the search, select “Begin Web-based ADAMS Search.” For problems with ADAMS, please contact the NRC’s Public Document Room (PDR) reference staff at 1–800–397–4209, 301–415–4737, or by email to [pdr.resource@nrc.gov](mailto:pdr.resource@nrc.gov).

- NRC’s PDR: You may examine and purchase copies of public documents at the NRC’s PDR, Room O1–F21, One White Flint North, 11555 Rockville Pike, Rockville, Maryland 20852.

*B. Submitting Comments*

Please include Docket ID NRC–2017–0214 in your comment submission. When preparing and submitting your comments, see “Tips for Submitting Effective Comments” (ADAMS Accession No. ML20014E720).

The NRC cautions you not to include identifying or contact information that you do not want to be publicly disclosed in your comment submission. The NRC will post all comment submissions at <https://www.regulations.gov> as well as enter the comment submissions into ADAMS. The NRC does not routinely edit comment submissions to remove identifying or contact information.

If you are requesting or aggregating comments from other persons for submission to the NRC, then you should inform those persons not to include identifying or contact information that they do not want to be publicly disclosed in their comment submission. Your request should state that the NRC does not routinely edit comment submissions to remove such information before making the comment submissions available to the public or entering the comment into ADAMS.

**II. Discussion**

On February 4, 2020, the NRC published a document in the **Federal Register** (85 FR 6103) requesting input from its licensees and members of the public to identify outdated or duplicative administrative requirements that may be eliminated without an adverse effect on public health or safety, common defense and security, protection of the environment, or regulatory efficiency and effectiveness. The public comment period was originally scheduled to close on April 6, 2020. By letter dated March 12, 2020 (ADAMS Accession No. ML20084Q158), the Nuclear Energy Institute requested that the NRC extend the comment period by 30 days. The NRC is granting this request and will extend the public comment period until May 6, 2020, to allow more time for members of the public to submit their comments.

Dated at Rockville, Maryland, this 26th day of March 2020.

For the Nuclear Regulatory Commission.

**John R. Tappert,**

*Director, Division of Rulemaking, Environmental, and Financial Support, Office of Nuclear Material Safety and Safeguards.*

[FR Doc. 2020–06682 Filed 4–1–20; 8:45 am]

**BILLING CODE 7590–01–P**

**DEPARTMENT OF TRANSPORTATION**

**Federal Aviation Administration**

**14 CFR Part 39**

[Docket No. FAA–2020–0293; Project Identifier MCAI–2019–00122–E]

**RIN 2120–AA64**

**Airworthiness Directives; Rolls-Royce Deutschland Ltd & Co KG (Type Certificate Previously Held by Rolls-Royce plc) Turbofan Engines**

**AGENCY:** Federal Aviation Administration (FAA), DOT.

**ACTION:** Notice of proposed rulemaking (NPRM).

**SUMMARY:** The FAA proposes to adopt a new airworthiness directive (AD) for certain Rolls-Royce Deutschland Ltd & Co KG Trent 1000–AE3, Trent 1000–CE3, Trent 1000–D3, Trent 1000–G3, Trent 1000–H3, Trent 1000–J3, Trent 1000–K3, Trent 1000–L3, Trent 1000–M3, Trent 1000–N3, Trent 1000–P3, Trent 1000–Q3, Trent 1000–R3, Trent 7000–72, and Trent 7000–72C model turbofan engines. This proposed AD was prompted by a report of a crack finding of the front air seal on the intermediate-pressure compressor (IPC) shaft assembly during the stripping of a flight test engine. This proposed AD would require initial and repetitive borescope inspections (BSIs) of the IPC shaft assembly and, depending on the results of the inspection, replacement of the IPC shaft assembly with a part eligible for installation. The FAA is proposing this AD to address the unsafe condition on these products.

**DATES:** The FAA must receive comments on this proposed AD by May 18, 2020.

**ADDRESSES:** You may send comments, using the procedures found in 14 CFR 11.43 and 11.45, by any of the following methods:

- *Federal eRulemaking Portal:* Go to <https://www.regulations.gov>. Follow the instructions for submitting comments.

- *Fax:* 202 493 2251.

- *Mail:* U.S. Department of Transportation, Docket Operations, M–30, West Building Ground Floor, Room W12 140, 1200 New Jersey Avenue SE, Washington, DC 20590.

- *Hand Delivery:* Deliver to Mail address above between 9 a.m. and 5 p.m., Monday through Friday, except Federal holidays.

For service information identified in this NPRM, contact Rolls-Royce Deutschland Ltd. & Co KG, Eschenweg 11, 15827 Blankenfelde-Mahlow, Germany; phone: +49 (0) 33 708 6 0; email: <https://www.rolls-royce.com/>