

Authority: 49 U.S.C. 106(f), 40113, 44701.

§ 39.13 [Amended]

■ 2. The FAA amends § 39.13 by adding the following new airworthiness directive:

2025–02–14 Rolls-Royce Deutschland Ltd & Co KG: Amendment 39–22947; Docket No. FAA–2024–2414; Project Identifier MCAI–2024–00530–E.

(a) Effective Date

This airworthiness directive (AD) is effective April 1, 2025.

(b) Affected ADs

None.

(c) Applicability

This AD applies to Rolls-Royce Deutschland Ltd & Co KG Model 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 engines.

(d) Subject

Joint Aircraft System Component (JASC) Code 7230, Turbine Engine Compressor Section.

(e) Unsafe Condition

This AD was prompted by reports of cracked intermediate pressure compressor (IPC) shaft assembly front air seals. The FAA is issuing this AD to prevent an IPC stage 1 disk burst or failure of the IPC front seal. The unsafe condition, if not addressed, could result in an IPC stage 1 disk burst with consequent release of high energy debris and damage to the airplane or failure of the IPC front seal and release of debris, which could lead to an engine in-flight shutdown (IFSD) and in the case of a dual IFSD could result in reduced control of the airplane.

(f) Compliance

Comply with this AD within the compliance times specified, unless already done.

(g) Required Actions

Except as specified in paragraphs (h), and (i) of this AD: Perform all required actions within the compliance times specified in, and in accordance with, European Union Aviation Safety Agency AD 2024–0178, dated September 12, 2024 (EASA AD 2024–0178).

(h) Exceptions to EASA AD 2024–0178

(1) Where EASA AD 2024–0178 refers to its effective date, this AD requires using the effective date of this AD.

(2) Where the service information referenced in EASA AD 2024–0178 specifies to reject the engine, this AD requires removing the affected part from service.

(3) This AD does not adopt the Remarks paragraph of EASA AD 2024–0178.

(i) No Reporting Requirement

Although the service information referenced in EASA AD 2024–0178 specifies to submit certain information to the

manufacturer, this AD does not include that requirement.

(j) Alternative Methods of Compliance (AMOCs)

(1) The Manager, AIR–520 Continued Operational Safety Branch, FAA, has the authority to approve AMOCs for this AD, if requested using the procedures found in 14 CFR 39.19. In accordance with 14 CFR 39.19, send your request to your principal inspector or local Flight Standards District Office, as appropriate. If sending information directly to the Manager, AIR–520 Continued Operational Safety Branch, send it to the attention of the person identified in paragraph (k) of this AD and email to: AMOC@faa.gov.

(2) Before using any approved AMOC, notify your appropriate principal inspector, or lacking a principal inspector, the manager of the local flight standards district office/certificate holding district office.

(k) Additional Information

For more information about this AD, contact Barbara Caufield, Aviation Safety Engineer, FAA, 2200 South 216th Street, Des Moines, WA 98198; phone: (781) 238–7146; email: barbara.caufield@faa.gov.

(l) Material Incorporated by Reference

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

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

(i) European Union Aviation Safety Agency (EASA) AD 2024–0178, dated September 12, 2024.

(ii) [Reserved]

(3) For EASA material identified in this AD, contact EASA, Konrad-Adenauer-Ufer 3, 50668 Cologne, Germany; phone: +49 221 8999 000; email: ADs@easa.europa.eu; website: easa.europa.eu. You may find this EASA AD on the EASA website at ad.easa.europa.eu.

(4) You may view this material at the FAA, Airworthiness Products Section, Operational Safety Branch, 1200 District Avenue, Burlington, MA 01803. For information on the availability of this material at the FAA, call (817) 222–5110.

(5) You may view this material at the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, visit www.archives.gov/federal-register/cfr/ibr-locations or email fr.inspection@nara.gov.

Issued on February 19, 2025.

Suzanne Masterson,

Deputy Director, Integrated Certificate Management Division, Aircraft Certification Service.

[FR Doc. 2025–03009 Filed 2–24–25; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 101

[Docket No. FDA–2016–D–2335]

RIN 0910–AI13

Food Labeling: Nutrient Content Claims; Definition of Term “Healthy”

AGENCY: Food and Drug Administration, Department of Health and Human Services (HHS).

ACTION: Final rule; delay of effective date.

SUMMARY: In accordance with the memorandum of January 20, 2025, from the President, entitled “Regulatory Freeze Pending Review,” the effective date of the final rule entitled “Food Labeling: Nutrient Content Claims; Definition of Term ‘Healthy,’” is delayed until April 28, 2025.

DATES: As of February 25, 2025, the effective date for the final rule published December 27, 2024 (89 FR 106064), is delayed until April 28, 2025.

FOR FURTHER INFORMATION CONTACT: Vincent de Jesus, Office of Nutrition and Food Labeling Human Foods Program, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–1450; Denise See or Barbara Little, Office of Policy, Regulations, and Information (HFS–024), Human Foods Program, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–2378.

SUPPLEMENTARY INFORMATION:

I. Electronic Access and Filing

A copy of the notice of proposed rulemaking (87 FR 59168, September 29, 2022), all comments received, the final rule (89 FR 106064, December 27, 2024), and all background material may be viewed online at <https://www.regulations.gov> using the docket number listed above. A copy of this document will be placed in the docket. Electronic retrieval help and guidelines are available on the website. It is available 24 hours each day, 365 days each year. An electronic copy of this document may also be downloaded from the Office of the Federal Register’s website at <https://www.federalregister.gov> and the Government Publishing Office’s website at <https://www.gpo.gov>.

II. Background

FDA published a final rule, entitled “Food Labeling: Nutrient Content Claims; Definition of Term ‘Healthy,’”

in the **Federal Register** on December 27, 2024 (89 FR 106064). The final rule was published with an effective date of February 25, 2025. On January 20, 2025, the President issued a memorandum entitled “Regulatory Freeze Pending Review” (90 FR 8249 (January 28, 2025)). With respect to rules that have been published in the **Federal Register**, but have not taken effect, the memorandum orders Agencies to consider postponing the rules’ effective dates for 60 days (*i.e.*, until April 28, 2025) for the purpose of reviewing any questions of fact, law, and policy the rules may raise.

In accordance with this direction, FDA is delaying the effective date of the final rule “Food Labeling: Nutrient Content Claims; Definition of Term ‘Healthy’” (89 FR 106064), until April 28, 2025. We note that the compliance date remains unchanged at this time. The final rule:

- updates the requirements for when the term “healthy” can be used as an implied nutrient content claim in the labeling of human food products to help consumers identify foods that can serve as the foundation of a nutritious diet that is consistent with current dietary recommendations;
- establishes parameters for use of the term “healthy” or derivative terms “health,” “healthful,” “healthfully,” “healthfulness,” “healthier,” “healthiest,” “healthily,” and “healthiness” as an implied nutrient content claim on the label or in labeling of a food that suggests that a food, because of its nutrient content, may help consumers maintain healthy dietary practices, where there is also implied or explicit information about the nutrition content of the food on the label or in the labeling;
- establishes a framework based on food groups and nutrients to limit (NTL) for the “healthy” claim;
- establishes that “food group,” for the purposes of the “healthy” claim, refers to the groups of foods recommended in the *Dietary Guidelines, 2020–2025* (for adults and children 2 years of age and older; available at <https://www.dietaryguidelines.gov>), which are vegetables, fruits, dairy, grains, protein foods, as well as oils;
- establishes food group equivalents (FGEs) that identify qualifying amounts of foods from each food group based on nutritional content;
- requires that, to bear a claim subject to the rule, individual food products, mixed products, main dishes, and meals must meet FGEs and specific limits for added sugars, saturated fat, and sodium based on a percentage of the Daily Value for these nutrients;

- provides that individual foods or mixed products that are comprised of one or more of the following foods encouraged by the *Dietary Guidelines, 2020–2025*, with no other added ingredients except for water: vegetable; fruit; whole grains; fat-free and low-fat dairy; lean meat, seafood, eggs, beans, peas, lentils, nuts and seeds, automatically qualify (*i.e.*, without having to meet the FGE and NTL requirements) for the “healthy” claim because of their nutrient profile and positive contribution to an overall healthy diet;

- provides that all water, tea, and coffee with less than 5 calories per reference amount customarily consumed and per labeled serving automatically qualify for the “healthy” claim; and

- requires the establishment and maintenance of certain records for foods bearing the “healthy” claim where the FGE contained in the product is not apparent from the label of the food. The records must be kept for a period of at least 2 years after introduction or delivery for introduction of the food into interstate commerce. During an inspection, such records must be provided to FDA upon request for official review and photocopying or other means of reproduction.

To the extent that 5 U.S.C. 553 applies to this action, it is exempt from notice and comment because it constitutes a rule of procedure under 5 U.S.C. 553(b)(A). Alternatively, FDA’s implementation of this action without opportunity for public comment, effective immediately, is based on the good cause exceptions in 5 U.S.C. 553(b)(B) and (d)(3). Seeking public comment is impracticable, unnecessary, and contrary to the public interest. The temporary delay in the effective date until April 28, 2025, is necessary to give Agency officials the opportunity for further review and consideration of the new regulation, consistent with the memorandum described previously. Seeking public comment is unnecessary because of the limited impact of the delayed effective date; the compliance date, and not the effective date, controls when parties must comply with this rule, and the compliance date in the final rule is not until 2028. FDA also stated in the final rule that parties are free to begin implementing the rule earlier than the compliance date. In addition, given the imminence of the effective date and the brief length of the extension of the effective date, seeking prior public comment on this temporary delay would have been impracticable, as well as contrary to the public interest in the orderly promulgation and

implementation of regulations. FDA also recognizes that certain affected entities would benefit from being informed as soon as possible of the extension and its length in order to plan and adjust their implementation process accordingly.

Robert F. Kennedy, Jr.,

Secretary, Department of Health and Human Services.

[FR Doc. 2025–03118 Filed 2–24–25; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HOUSING AND URBAN DEVELOPMENT

24 CFR Parts 3280, 3282, 3285, and 3286

[Docket No. FR–6233–F–03]

RIN 2502–AJ58

Manufactured Home Construction and Safety Standards; Postponing Effective Date

AGENCY: Office of the Assistant Secretary for Housing—Federal Housing Commissioner, Department of Housing and Urban Development (HUD).

ACTION: Final rule; delay of effective date.

SUMMARY: On September 16, 2024, HUD published the “Manufactured Home Construction and Safety Standards” final rule (MHCSS 4th and 5th Sets) in the **Federal Register**. The MHCSS 4th and 5th Sets final rule established a March 17, 2025, effective date. Consistent with the President’s January 20, 2025, memorandum titled “Regulatory Freeze Pending Review”, this document announces that HUD is delaying the effective date for the MHCSS 4th and 5th Sets final rule until September 15, 2025.

DATES: As of February 25, 2025, the effective date for the MHCSS 4th and 5th Sets, published at 89 FR 75704 (September 16, 2024), is delayed from March 17, 2025, until September 15, 2025.

FOR FURTHER INFORMATION CONTACT:

Teresa B. Payne, Deputy Assistant Secretary—Administrator, Office of Manufactured Housing Programs, Office of Housing, Department of Housing and Urban Development, 451 7th Street SW, Washington, DC 20410; telephone 202–402–5365 (this is not a toll-free number). HUD welcomes and is prepared to receive calls from individuals who are deaf or hard of hearing, as well as individuals with speech or communication disabilities. To learn more about how to make an accessible telephone call, please visit: