

**(i) Credit for Previous Actions**

This paragraph provides credit for the actions specified in paragraph (g) of this AD, if those actions were performed before the effective date of this AD using Boeing Alert Service Bulletin 777-53A0081, dated September 8, 2016, or Boeing Alert Service Bulletin 777-53A0081, Revision 1, dated May 1, 2017.

**(j) Alternative Methods of Compliance (AMOCs)**

(1) The Manager, Seattle ACO 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 of the certification office, send it to the attention of the person identified in paragraph (k)(1) of this AD. Information may be emailed to: [9-ANM-Seattle-ACO-AMOC-Requests@faa.gov](mailto:9-ANM-Seattle-ACO-AMOC-Requests@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.

(3) An AMOC that provides an acceptable level of safety may be used for any repair, modification, or alteration required by this AD if it is approved by The Boeing Company Organization Designation Authorization (ODA) that has been authorized by the Manager, Seattle ACO Branch, FAA, to make those findings. To be approved, the repair method, modification deviation, or alteration deviation must meet the certification basis of the airplane, and the approval must specifically refer to this AD.

(4) AMOCs approved previously for AD 2017-16-10 are approved as AMOCs for the corresponding provisions of Boeing Alert Service Bulletin 777-53A0081, Revision 2, dated March 29, 2019, that are required by paragraph (g) of this AD.

(5) Except as specified by paragraph (h)(2) of this AD: For service information that contains steps that are labeled as Required for Compliance (RC), the provisions of paragraphs (j)(5)(i) and (j)(5)(ii) of this AD apply.

(i) The steps labeled as RC, including substeps under an RC step and any figures identified in an RC step, must be done to comply with the AD. If a step or substep is labeled "RC Exempt," then the RC requirement is removed from that step or substep. An AMOC is required for any deviations to RC steps, including substeps and identified figures.

(ii) Steps not labeled as RC may be deviated from using accepted methods in accordance with the operator's maintenance or inspection program without obtaining approval of an AMOC, provided the RC steps, including substeps and identified figures, can still be done as specified, and the airplane can be put back in an airworthy condition.

**(k) Related Information**

(1) For more information about this AD, contact Eric Lin, Aerospace Engineer, Airframe Section, FAA, Seattle ACO Branch, 2200 South 216th St., Des Moines, WA

98198; phone and fax: 206-231-3523; email: [eric.lin@faa.gov](mailto:eric.lin@faa.gov).

(2) For service information identified in this AD, contact Boeing Commercial Airplanes, Attention: Contractual & Data Services (C&DS), 2600 Westminister Blvd., MC 110-SK57, Seal Beach, CA 90740-5600; phone: 562-797-1717; internet: <https://www.myboeingfleet.com>. You may view this referenced service information at the FAA, Transport Standards Branch, 2200 South 216th St., Des Moines, WA. For information on the availability of this material at the FAA, call 206-231-3195.

**(l) Material Incorporated by Reference**

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

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

(i) Boeing Alert Service Bulletin 777-53A0081, Revision 2, dated March 29, 2019.

(ii) [Reserved]

(3) For service information identified in this AD, contact Boeing Commercial Airplanes, Attention: Contractual & Data Services (C&DS), 2600 Westminister Blvd., MC 110-SK57, Seal Beach, CA 90740-5600; phone: 562-797-1717; internet: <https://www.myboeingfleet.com>.

(4) You may view this service information at the FAA, Transport Standards Branch, 2200 South 216th St., Des Moines, WA. For information on the availability of this material at the FAA, call 206-231-3195.

(5) You may view this service information that is incorporated by reference at the National Archives and Records Administration (NARA). For information on the availability of this material at NARA, call 202-741-6030, or go to: <http://www.archives.gov/federal-register/cfr/ibr-locations.html>.

Issued in Des Moines, Washington, on June 5, 2019.

**Michael Kaszycki,**

*Acting Director, System Oversight Division,  
Aircraft Certification Service.*

[FR Doc. 2019-13058 Filed 6-19-19; 8:45 am]

**BILLING CODE 4910-13-P**

**RAILROAD RETIREMENT BOARD****20 CFR Part 200****RIN 3220-AB67****General Administration: Designation of Central and Field Organization; Internal Organization**

**AGENCY:** Railroad Retirement Board.

**ACTION:** Final rule.

**SUMMARY:** The Railroad Retirement Board (Board) amends its regulations to update the members of the Executive Committee, update the responsibilities of the Executive Committee members, and update office titles.

**DATES:** This rule becomes effective June 20, 2019.

**ADDRESSES:** Stephanie Hillyard, Secretary to the Board, Railroad Retirement Board, 844 N Rush Street, Chicago, Illinois 60611-1275.

**FOR FURTHER INFORMATION CONTACT:** Marguerite P. Dadabo, Assistant General Counsel, (312) 751-4945, TTD (312) 751-4701.

**SUPPLEMENTARY INFORMATION:** The Railroad Retirement Board (Board) amends its regulations in regard to the Board's policy on internal organization. The regulations amended are all contained in § 200.1(b). In § 200.1(b)(1) of the Board's regulations, the Board removes the language that states "the General Counsel also serves as the Senior Executive Officer," and increases the number of members of the Executive Committee from six to seven members by adding as a member the Director of Field Service. A description of the Director of Field Service's responsibilities is added to § 200.1(b)(2). Finally, under § 200.1(b)(3), the office name of the Washington/Legislative Office is changed to the Office of Legislative Affairs. Section 200.1(b)(3) of the regulation also removes the Office of Planning, and renames the Bureau of Quality Assurance to the Program Evaluation and Management Services (PEMS).

This change was published as a proposed rule on April 27, 2017, and comments were invited to be submitted by June 26, 2017. See 82 FR 19330 (April 27, 2017). No comments were submitted, and the final rule is essentially the same as the proposed rule.

The Board, with the concurrence of the Office of Management and Budget, has determined that this is not a significant regulatory action under Executive Order 12866, as amended. Therefore, no regulatory impact analysis is required. There are no changes to the information collections associated with § 200.1(b).

**List of Subjects in 20 CFR Part 200**

Railroad employees, Railroad retirement, General administration.

For the reasons set out in the preamble, the Railroad Retirement Board amends title 20, chapter II, subchapter A, part 200 of the Code of Federal Regulations as follows:

**PART 200—GENERAL ADMINISTRATION**

■ 1. The authority citation for part 200 continues to read as follows:

**Authority:** 45 U.S.C. 231f (b)(5) and 45 U.S.C. 362; § 200.4 also issued under 5 U.S.C. 552; § 200.5 also issued under 5 U.S.C. 552a; § 200.6 also issued under 5 U.S.C. 552b; and § 200.7 also issued under 31 U.S.C. 3717.

■ 2. Section 200.1 is amended by revising paragraph (b) to read as follows:

**§ 200.1 Designation of central and field organization.**

\* \* \* \* \*

(b) *Internal organization.* (1) Reporting directly to the Board Members is the seven member Executive Committee. The Executive Committee is comprised of the General Counsel, the Director of Administration, the Director of Programs, the Chief Financial Officer, the Chief Information Officer, and the Director of Field Service. The Chief Actuary is a non-voting member. The Board members will designate a member of the Executive Committee as Senior Executive Officer.

(2) The Executive Committee is responsible for the day to day operations of the agency. The Senior Executive Officer is responsible for the direction and oversight of the Executive Committee. The General Counsel is responsible for advising the Board Members on major issues, interpreting the Acts and regulations administered by the Board, drafting and analyzing legislation, representing the Board in litigation and administrative forums and planning, directing, and coordinating the work of the Office of General Counsel, the Office of Secretary to the Board, the Bureau of Hearings and Appeals, and the Office of Legislative Affairs through their respective directors. The Director of Programs is responsible for managing, coordinating, and controlling the program operations of the agency which carry out provisions of the Railroad Retirement and Railroad Unemployment Insurance Acts. The Director of Administration is responsible for managing, coordinating and controlling certain administrative operations of the Board including the Division of Acquisition Management, the Bureau of Human Resources, the Office of Public Affairs, and the Division of Real Property Management. The Chief Financial Officer is responsible for the financial management of the agency, and the Chief Information Officer is responsible for coordinating the agency's information resources management program.

The Chief Actuary is responsible for the actuarial program of the Board, and for maintaining statistical and financial information. The Director of Field Services is responsible for the oversight of the agency's nationwide field offices.

(3) The Office of Equal Employment Opportunity is responsible for equal employment opportunity and affirmative employment programs.

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By Authority of the Board

**Stephanie Hillyard,**

*Secretary to the Board.*

[FR Doc. 2019-13050 Filed 6-19-19; 8:45 am]

**BILLING CODE P**

**DEPARTMENT OF HEALTH AND HUMAN SERVICES**

**Food and Drug Administration**

**21 CFR Part 101**

[Docket No. FDA-2018-D-0075]

**The Declaration of Added Sugars on Honey, Maple Syrup, Other Single-Ingredient Sugars and Syrups, and Certain Cranberry Products; Guidance for Industry; Availability**

**AGENCY:** Food and Drug Administration, HHS.

**ACTION:** Notification of availability.

**SUMMARY:** The Food and Drug Administration (FDA or we) is announcing the availability of a final guidance for industry entitled "The Declaration of Added Sugars on Honey, Maple Syrup, Other Single-Ingredient Sugars and Syrups, and Certain Cranberry Products." This guidance provides clarification on the labeling requirements for single-ingredient packages and/or containers of pure honey, pure maple syrup, and other pure sugars and syrups, which are not required to bear the words "Includes Xg Added Sugars," but must still include the percent Daily Value for added sugars on their labels. This guidance is also intended to advise food manufacturers of our intent to exercise enforcement discretion related to the use of a "+" symbol immediately following the percent Daily Value for added sugars on single-ingredient packages and/or containers of pure honey, pure maple syrup, and other pure sugars and syrups; the "+" symbol would lead the consumer to a statement that is truthful and not misleading in a footnote at the bottom of the Nutrition Facts label. The guidance also advises food manufacturers of our intent to exercise enforcement discretion with respect to the use of a "+" symbol immediately after the added sugars percent Daily Value information that leads the consumer to a statement that is truthful and not misleading outside of the Nutrition Facts label on certain dried

cranberry and cranberry beverage products that are made up of cranberry juice sweetened with added sugars and that contain total sugars at levels no greater than comparable products with endogenous (inherent) sugars, but no added sugars. Further, this guidance advises of our intent to exercise enforcement discretion regarding compliance with Nutrition Facts label final rule and Serving Size final rule requirements until July 1, 2021, for the single-ingredient sugars and syrups as well as the cranberry products discussed in the guidance document.

**DATES:** The announcement of the guidance is published in the **Federal Register** on June 20, 2019.

**ADDRESSES:** You may submit either electronic or written comments on FDA guidances at any time as follows:

*Electronic Submissions*

Submit electronic comments in the following way:

- *Federal eRulemaking Portal:* <https://www.regulations.gov>. Follow the instructions for submitting comments. Comments submitted electronically, including attachments, to <https://www.regulations.gov> will be posted to the docket unchanged. Because your comment will be made public, you are solely responsible for ensuring that your comment does not include any confidential information that you or a third party may not wish to be posted, such as medical information, your or anyone else's Social Security number, or confidential business information, such as a manufacturing process. Please note that if you include your name, contact information, or other information that identifies you in the body of your comments, that information will be posted on <https://www.regulations.gov>.

- If you want to submit a comment with confidential information that you do not wish to be made available to the public, submit the comment as a written/paper submission and in the manner detailed (see "Written/Paper Submissions" and "Instructions").

*Written/Paper Submissions*

Submit written/paper submissions as follows:

- *Mail/Hand delivery/Courier (for written/paper submissions):* Dockets Management Staff (HFA-305), Food and Drug Administration, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852.

- For written/paper comments submitted to the Dockets Management Staff, FDA will post your comment, as well as any attachments, except for information submitted, marked and identified, as confidential, if submitted as detailed in "Instructions."