7400.11J is publicly available online at *www.faa.gov/air_traffic/publications/.* You may also contact the Rules and Regulations Group, Office of Policy, Federal Aviation Administration, 600 Independence Avenue SW, Washington, DC 20597; telephone: (202) 267–8783.

FAA Order JO 7400.11J lists Class A, B, C, D, and E airspace areas, air traffic service routes, and reporting points.

Good Cause for No Notice and Comment

Section 553(b)(3)(B) of Title 5, United States Code, (the Administrative Procedure Act) authorizes agencies to dispense with notice and comment procedures for rules when the agency for ''good cause'' finds that those procedures are ''impracticable, unnecessary, or contrary to the public interest.'' Under this section, an agency, upon finding good cause, may issue a final rule without seeking comment prior to the rulemaking. The FAA finds that prior notice and public comment to this final rule is unnecessary due to the brief length of the extension of the effective date and the fact that there is no substantive change to the rule.

Delay of Effective Date

■ Accordingly, pursuant to the authority delegated to me, the effective date of the final rule for Airspace Docket 23–AGL–26, as published in the **Federal Register** on August 19, 2024 (89 FR 66987), FR Doc. 2024–18431, and corrected on September 30, 2024 (89 FR 79429), FR Doc. 2024–22253, is hereby delayed until December 26, 2024.

Authority: 49 U.S.C. 106(f), 106(g); 40103, 40113, 40120; E.O. 10854, 24 FR 9565, 3 CFR, 1959–1963 Comp., P. 389.

Issued in Washington, DC, on October 8, 2024.

Brian Eric Konie,

Acting Manager, Rules and Regulations Group.

[FR Doc. 2024–23612 Filed 10–11–24; 8:45 am]

BILLING CODE 4910–13–P

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 820

[Docket No. FDA–2021–N–0507]

RIN 0910–AH99

Medical Devices; Quality System Regulation Amendments; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Final rule; correction.

SUMMARY: The Food and Drug Administration (FDA or Agency) is correcting a final rule that appeared in the **Federal Register** on February 2, 2024. In that final rule, FDA amended the device current good manufacturing practice (CGMP) requirements of the Quality System (QS) regulation to harmonize and modernize the device CGMP. FDA is correcting an editorial error that inadvertently omitted a definition in the codified of the final rule. This action is editorial in nature and is intended to ensure the accuracy and clarity of the Agency's regulations.

DATES: Effective February 2, 2026.

FOR FURTHER INFORMATION CONTACT: Laurie Sternberg, Center for Devices and Radiological Health, Food and Drug Administration, 10903 New Hampshire Ave., Bldg. 66, Rm. 5517, Silver Spring, MD 20993–0002, 240–402–0425.

SUPPLEMENTARY INFORMATION: In the **Federal Register** of February 2, 2024 (89 FR 7496), FDA published a final rule that amended the device CGMPs requirements in 21 CFR part 820. The preamble indicated that the definition for ''batch'' or ''lot'' was set forth at § 820.3 (21 CFR 820.3) but the definition for ''batch'' or ''lot'' was inadvertently omitted from the codified portion of § 820.3 in the final rule. FDA is, therefore, correcting the codified for § 820.3 to include the definition of ''batch'' or ''lot'' as was intended and to be consistent with the preamble of the final rule.

In FR Doc. 2024–01709 appearing on page 7524 in the **Federal Register** of Friday, February 2, 2024 (89 FR 7496), the following correction is made:

§ 820.3 [Corrected]

■ 1. On page 7524, in amendment number 4, in the first column, in paragraph (a) of § 820.3, the definition for ''Batch or lot'' is added in alphabetical order to read as follows:

''*Batch* or *lot* means one or more components or finished devices that consist of a single type, model, class, size, composition, or software version that are manufactured under essentially the same conditions and that are intended to have uniform characteristics and quality within specified limits.''

Dated: October 7, 2024.

Kimberlee Trzeciak,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2024–23701 Filed 10–11–24; 8:45 am]

BILLING CODE 4164–01–P

DEPARTMENT OF HOMELAND SECURITY

Coast Guard

33 CFR Part 117

[Docket No. USCG–2024–0845]

RIN 1625–AA09

Drawbridge Operation Regulation; Wappinger Creek, New Hamburg, New York

AGENCY: Coast Guard, DHS.

ACTION: Final rule.

SUMMARY: The Coast Guard is removing the existing drawbridge operation regulation for the Metro-North Commuter Railroad Bridge, mile 0.0 across the Wappinger Creek at New Hamburg, New York. In 1991, the Metro-North Railroad Bridge was allowed to no longer be maintained as a movable structure and in 2004, the bridge was converted to a fixed bridge. The operating regulation is no longer applicable or necessary.

DATES: This rule is effective October 15, 2024.

ADDRESSES: To view documents mentioned in this preamble as being available in the docket, go to *https://www.regulations.gov.* Type the docket number (USCG–2024–0845) in the ''SEARCH'' box and click ''SEARCH''. In the Document Type column, select ''Supporting & Related Material.''

FOR FURTHER INFORMATION CONTACT: If you have questions on this rule, call or email Ms. Judy Leung-Yee, Project Officer, First Coast Guard District, telephone 212–514–4336, email *Judy.K.Leung-Yee@uscg.mil.*

SUPPLEMENTARY INFORMATION:

I. Table of Abbreviations [Delete/Add Any Abbreviations Not Used/Used in This Document]

CFR Code of Federal Regulations
DHS Department of Homeland Security
FR Federal Register
Pub. L. Public Law
§ Section
U.S.C. United States Code

II. Background Information and Regulatory History

The Coast Guard is issuing this final rule without prior notice and opportunity to comment pursuant to authority under section 4(a) of the Administrative Procedure Act (APA) (5 U.S.C. 553(b)). This provision authorizes an agency to issue a rule without prior notice and opportunity to comment when the agency for good cause finds that those procedures are