Proposed Rules

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This section of the FEDERAL REGISTER contains notices to the public of the proposed issuance of rules and regulations. The purpose of these notices is to give interested persons an opportunity to participate in the rule making prior to the adoption of the final rules.

DEPARTMENT OF HEALTH AND HUMAN SERVICES

Food and Drug Administration

21 CFR Part 146

[Docket No. FDA–2022–P–1668]

Food Standards of Identity Modernization; Pasteurized Orange Juice; Proposed Rule; Correction

AGENCY: Food and Drug Administration, HHS.

ACTION: Proposed rule.

SUMMARY: The Food and Drug Administration (FDA or we) is correcting the proposed rule entitled ''Food Standards of Identity Modernization; Pasteurized Orange Juice; Proposed Rule'' (90 FR 37817, August 6, 2025). In the proposed rule, FDA proposed to amend the standard of identity (SOI) for pasteurized orange juice (POJ) by lowering the minimum orange juice soluble solids content from 10.5° to 10° Brix. The proposed rule inadvertently included an additional summary of benefits table, an additional summary table and an extraneous paragraph and sentence. This document corrects those errors.

DATES: August 14, 2025.

FOR FURTHER INFORMATION CONTACT: Keronica Richardson, Office of Policy, Regulations, and Information, Human Foods Program, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, 240–402–2378.

SUPPLEMENTARY INFORMATION: In FR Doc. 2025–14949, appearing on pages 37822 to 37823 in the **Federal Register** of Wednesday, August 6, 2025, the following corrections are made:

1. In the **SUPPLEMENTARY INFORMATION** section, in subsection VII: Preliminary Economic Analysis of Impact, on page 37822, remove the extraneous paragraph starting with the sentence, ''The proposed rule, if finalized, would not require firms in the POJ industry to change their manufacturing practices or behavior in any way'' and ending with the sentence, ''We request comment on our described benefits and costs of the proposed rule.''

2. In the **SUPPLEMENTARY INFORMATION** section, in subsection VII: Preliminary Economic Analysis of Impact, on page 37823, remove ''Table 1—Summary of Benefits, Costs and Distributional Effects of the Proposed Rule''.

3. In the **SUPPLEMENTARY INFORMATION** section, in subsection VII: Preliminary Economic Analysis of Impact, on page 37823, remove the extraneous sentence ''In line with Executive Order 14192, in 2 we estimate present and annualized values of cost, cost savings, and net costs over an infinite time horizon.''

4. In the **SUPPLEMENTARY INFORMATION** section, in subsection VII: Preliminary Economic Analysis of Impact, on page 37823, remove ''Table 2—E.O. 14192 Summary Table''.

The full preliminary analysis of economic impacts remains available in the docket for this proposed rule (Ref. 2) and at *https://www.fda.gov/about-fda/economics-staff/regulatory-impact-analyses-ria.*

Dated: August 11, 2025.

Grace R. Graham,

Deputy Commissioner for Policy, Legislation, and International Affairs.

[FR Doc. 2025–15473 Filed 8–13–25; 8:45 am]

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ENVIRONMENTAL PROTECTION AGENCY

40 CFR Part 52

[EPA–R01–OAR–2024–0188; FRL–12928–01–R1]

Air Plan Approval; Rhode Island; Decommissioning of Stage II Vapor Recovery Systems

AGENCY: Environmental Protection Agency (EPA).

ACTION: Proposed rule.

SUMMARY: The Environmental Protection Agency (EPA) is proposing to approve a State Implementation Plan (SIP) revision submitted by the State of Rhode Island. This revision removes requirements for Stage II vapor recovery equipment at gasoline dispensing facilities (GDFs). This revision also includes minor updates to Stage I vapor recovery regulatory amendments. The intended effect of this action is to propose approval of Rhode Island's revised vapor recovery regulations. This action is being taken under the Clean Air Act.

DATES: Written comments must be received on or before September 15, 2025.

ADDRESSES: Submit your comments, identified by Docket ID No. EPA–R01–OAR–2024–0188 at *https://www.regulations.gov,* or via email to *martinelli.ayla@epa.gov.* For comments submitted at *Regulations.gov,* follow the online instructions for submitting comments. Once submitted, comments cannot be edited or removed from *Regulations.gov.* For either manner of submission, the EPA may publish any comment received to its public docket. Do not submit electronically any information you consider to be Confidential Business Information (CBI) or other information whose disclosure is restricted by statute. Multimedia submissions (audio, video, etc.) must be accompanied by a written comment. The written comment is considered the official comment and should include discussion of all points you wish to make. The EPA will generally not consider comments or comment contents located outside of the primary submission (*i.e.,* on the web, cloud, or other file sharing system). For additional submission methods, please contact the person identified in the **FOR FURTHER INFORMATION CONTACT** section. For the full EPA public comment policy, information about CBI or multimedia submissions, and general guidance on making effective comments, please visit *https://www.epa.gov/dockets/commenting-epa-dockets.* Publicly available docket materials are available at *https://www.regulations.gov* or at the U.S. Environmental Protection Agency, EPA Region 1 Regional Office, Air and Radiation Division, 5 Post Office Square—Suite 100, Boston, MA. EPA requests that, if at all possible, you contact the contact listed in the **FOR FURTHER INFORMATION CONTACT** section to schedule your inspection.

FOR FURTHER INFORMATION CONTACT: Ayla Martinelli, Air Quality Branch, U.S. Environmental Protection Agency, EPA Region 1, 5 Post Office Square—Suite 100, (Mail code 5–MI), Boston, MA 02109–3912, tel. (617) 918–1057, email: *martinelli.ayla@epa.gov.*