



19 August 2025

(25-5148)

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Committee on Technical Barriers to Trade

Original: English

## NOTIFICATION

### *Corrigendum*

The following communication, dated 18 August 2025, is being circulated at the request of the delegation of the United States of America.

#### Food Standards of Identity Modernization; Pasteurized Orange Juice

Reason for corrigendum:	
<input type="checkbox"/>	Notification circulated in error
<input type="checkbox"/>	Clerical error in notification
<input checked="" type="checkbox"/>	Clerical error in notified measure/reference document
<input type="checkbox"/>	Other

#### **Description:**

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](#), 6 August 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.

90 Federal Register (FR) 39139, 14 August 2025; [Title 21 Code of Federal Regulations \(CFR\) Part 146](#):

<https://www.govinfo.gov/content/pkg/FR-2025-08-14/html/2025-15473.htm>

<https://www.govinfo.gov/content/pkg/FR-2025-08-14/pdf/2025-15473.pdf>

This correction and the proposed rule are identified by Docket Number FDA-2022-P-1668. The Docket Folder is available on [Regulations.gov](#) at <https://www.regulations.gov/docket/FDA-2022-P-1668/document> and provides access to [primary](#) and [supporting](#) documents as well as comments received. Documents are also accessible from Regulations.gov by searching the Docket Number. WTO Members and their stakeholders are asked to submit comments to the [USA TBT Enquiry Point](#) by or before [4pm Eastern Time](#) on 4 November 2025. [Comments](#) received by the USA TBT Enquiry Point from WTO Members and their stakeholders will be shared with FDA and will also be submitted to the [Docket](#) on Regulations.gov if received within the comment period.

[https://members.wto.org/crnattachments/2025/TBT/USA/25\\_05326\\_00\\_e.pdf](https://members.wto.org/crnattachments/2025/TBT/USA/25_05326_00_e.pdf)