WORLD TRADE ORGANIZATION

G/TBT/N/USA/1889/Add.3

28 January 2025

(25-0670)

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Committee on Technical Barriers to Trade

Original: English

NOTIFICATION

Addendum

The following communication, dated 28 January 2025, is being circulated at the request of the delegation of the United States of America.

Title: Nonprescription Drug Product With an Additional Condition for Nonprescription Use

Reason for Addendum:	
[ ]	Comment period changed - date:
[ ]	Notified measure adopted - date:
[ ]	Notified measure published - date:
[X]	Notified measure enters into force - date: 21 March 2025; The effective date for the final rule published 26 December 2024, (89 FR 105288), is delayed until 21 March 2025.
[X]	Text of final measure available from[1]: https://www.govinfo.gov/content/pkg/FR-2024-12-26/html/2024-30261.htm https://www.govinfo.gov/content/pkg/FR-2024-12-26/pdf/2024-30261.pdf
[ ]	Notified measure withdrawn or revoked - date: Relevant symbol if measure re-notified:
[ ]	Content or scope of notified measure changed and text available from[1]: New deadline for comments (if applicable):
[ ]	Interpretive guidance issued and text available from[1]:
[X]	Other: The effective date for the final rule published 26 December 2024, (89 FR 105288), is delayed until 21 March 2025. https://members.wto.org/crnattachments/2025/TBT/USA/25_00971_00_e.pdf

Description: In accordance with the memorandum of 20 January 2025, from the President, entitled ''Regulatory Freeze Pending Review,'' the effective date of the final rule, (notified as G/TBT/N/USA/1889/Add.2) entitled Nonprescription Drug Product With an Additional Condition for Nonprescription Use,'' (ACNU) is delayed until 21 March 2025.

The effective date for the final rule published 26 December 2024, (89 FR 105288), is delayed until 21 March 2025.

[1] This information can be provided by including a website address, a pdf attachment, or other information on where the text of the final/modified measure and/or interpretive guidance can be obtained.

90 Federal Register (FR) 8173, 27 January 2025; Title 21 Code of Federal Regulations (CFR) Parts 201 and 314:

https://www.govinfo.gov/content/pkg/FR-2025-01-27/html/2025-01840.htm

https://www.govinfo.gov/content/pkg/FR-2025-01-27/pdf/2025-01840.pdf

This action and previous actions notified under the symbol G/TBT/N/USA/1889 are identified by Docket Number FDA-2021-N-0862. The Docket Folder is available from Regulations.gov at https://www.regulations.gov/docket/FDA-2021-N-0862/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.