NOTIFICATION

Addendum

The following communication, dated 27 March 2023, is being circulated at the request of the delegation of the United States of America.

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**Title:** Current Good Manufacturing Practices, Quality Control Procedures, Quality Factors, Notification Requirements, and Records and ReportMercis, for Infant Formula

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| **Reason for Addendum:** | |
| [ ] | Comment period changed - date: |
| [ ] | Notified measure adopted - date: |
| [X] | Notified measure published - date: 24 March 2023 |
| [X] | Notified measure enters into force - date: 24 March 2023 |
| [X] | Text of final measure available from[[1]](#footnote-1):  <https://www.govinfo.gov/content/pkg/FR-2023-03-24/html/2023-05418.htm>  <https://www.govinfo.gov/content/pkg/FR-2023-03-24/pdf/2023-05418.pdf>  <https://members.wto.org/crnattachments/2023/TBT/USA/final_measure/23_8450_00_e.pdf> |
| [ ] | Notified measure withdrawn or revoked - date:  Relevant symbol if measure re-notified: |
| [ ] | Content or scope of notified measure changed and text available from1:  New deadline for comments (if applicable): |
| [ ] | Interpretive guidance issued and text available from1: |
| [ ] | Other: |

**Description:** TITLE: Food Labeling, Infant Formula Requirements, Food Additives and Generally Recognized as Safe Substances, New Dietary Ingredient Notification; Technical Amendments

AGENCY: Food and Drug Administration, HHS

ACTION: Final rule; technical amendments

SUMMARY: The Food and Drug Administration (FDA or we) is amending its regulations that pertain to food labeling, infant formula requirements, food additives, direct food substances affirmed as generally recognized as safe (GRAS), and new dietary ingredient (NDI) notifications. These amendments correct typographical errors, correct errors in sample labels, restore inadvertent omissions, and update office and organization names, addresses, and other references. This action is ministerial or editorial in nature.

DATES: This rule is effective on 24 March 2023.

This final rule; technical amendments is identified by Docket Number FDA-2022-N-2898. The Docket Folder is available on Regulations.gov at <https://www.regulations.gov/docket/FDA-2022-N-2898/document> and provides access to primary documents as well as comments received. Documents are also accessible from Regulations.gov by searching the Docket Number.

Previous actions notified under the symbol [G/TBT/N/USA/885](https://eping.wto.org/en/Search?domainIds=1&documentSymbol=usa%2F885) are identified by Docket Number FDA-1995-N-0036. The Docket Folder is available on Regulations.gov at [https://www.regulations.gov/docket/FDA-1995-N-0036/document](https://www.regulations.gov/docket/FDA-1995-N-0063/document) and provides access to primary and supporting documents as well as comments received. Documents are also accessible from [Regulations.gov](http://www.regulations.gov/) by searching the Docket Number.

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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. [↑](#footnote-ref-1)