NOTIFICATION

Addendum

The following communication, dated 25 January 2023, is being circulated at the request of the delegation of the United States of America.

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**Title:** Color Additive Certification; Increase in Fees for Certification Services

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| **Reason for Addendum:** |
| [X] | Comment period changed - date: 10 March 2023; Proposed rule; reopening of the comment period until 10 March 2023<https://www.govinfo.gov/content/pkg/FR-2023-01-24/html/2023-01361.htm><https://www.govinfo.gov/content/pkg/FR-2023-01-24/pdf/2023-01361.pdf> |
| [ ] | Notified measure adopted - date:  |
| [ ] | Notified measure published - date:  |
| [ ] | Notified measure enters into force - date:  |
| [ ] | Text of final measure available from[[1]](#footnote-1):  |
| [ ] | Notified measure withdrawn or revoked - date: Relevant symbol if measure re-notified:  |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: Color Additive Certification; Increase in Fees for Certification Services; Reopening of the Comment Period

AGENCY: Food and Drug Administration, HHS

ACTION: Proposed rule; reopening of the comment period

SUMMARY: The Food and Drug Administration (FDA or we) is reopening the comment period for the proposed rule, "Color Additive Certification; [Increase in Fees for Certification Services](https://www.fda.gov/food/cfsan-constituent-updates/fda-proposes-increase-color-certification-fees)," which published in the Federal Register of 2 November 2022. We are taking this action in response to a request from stakeholders to extend the comment period to allow additional time for interested parties to collect, analyze, and incorporate data to develop comments for this proposed rule.

DATES: FDA is reopening the comment period on the proposed rule "Color Additive Certification; Increase in Fees for Certification Services," which published in the Federal Register on 2 November 2022 (87 FR 66116). Either electronic or written comments must be submitted by 10 March 2023.

This proposed rule; reopening of the comment period and the proposed rule notified as [G/TBT/N/USA/1937](https://eping.wto.org/en/Search?domainIds=1&documentSymbol=usa%2F1937) are identified by Docket Number FDA-2022-N-1635. The Docket Folder is available on Regulations.gov at <https://www.regulations.gov/docket/FDA-2022-N-1635/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. WTO Members and their stakeholders are asked to submit comments to the USA TBT Enquiry Point by or before [4pm](http://time-time.net/times/time-zones/usa-canada/current-eastern-time-est.php) [Eastern Time](https://24timezones.com/time-zone/et) on 10 March 2023. Comments received by the USA TBT Enquiry Point from WTO Members and their stakeholders will be shared with the regulator and will also be submitted to the [Docket](https://www.regulations.gov/docket/FDA-2022-N-1635/document) on Regulations.gov if received within the comment period.

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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)