



23 November 2022

(22-8700)

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Committee on Sanitary and Phytosanitary Measures

Original: English

## NOTIFICATION

### Addendum

The following communication, received on 22 November 2022, is being circulated at the request of the Delegation of the United States of America.

#### Requirements for Additional Traceability Records for Certain Foods; Final Rule

The Food and Drug Administration (FDA, the Agency, or we) is issuing a final rule establishing additional recordkeeping requirements for persons who manufacture, process, pack, or hold foods the Agency has designated for inclusion on the Food Traceability List (FTL). The final rule adopts provisions requiring these entities to maintain records containing information on critical tracking events in the supply chain for these designated foods, such as initially packing, shipping, receiving, and transforming these foods. The requirements established in the final rule will help the Agency rapidly and effectively identify recipients of foods to prevent or mitigate foodborne illness outbreaks and address credible threats of serious adverse health consequences or death resulting from foods being adulterated or misbranded. We are issuing this regulation in accordance with the FDA Food Safety Modernization Act (FSMA).

<https://www.govinfo.gov/content/pkg/FR-2022-11-21/pdf/2022-24417.pdf>  
[https://members.wto.org/crnattachments/2022/SPS/USA/22\\_7958\\_00\\_e.pdf](https://members.wto.org/crnattachments/2022/SPS/USA/22_7958_00_e.pdf)

#### **This addendum concerns a:**

- ☐ Modification of final date for comments
- ☒ Notification of adoption, publication or entry into force of regulation
- ☐ Modification of content and/or scope of previously notified draft regulation
- ☐ Withdrawal of proposed regulation
- ☐ Change in proposed date of adoption, publication or date of entry into force
- ☐ Other:

**Comment period: (If the addendum extends the scope of the previously notified measure in terms of products and/or potentially affected Members, a new deadline for receipt of comments should be provided, normally of at least 60 calendar days. Under other circumstances, such as extension of originally announced final date for comments, the comment period provided in the addendum may vary.)**

- ☐ Sixty days from the date of circulation of the addendum to the notification and/or (dd/mm/yy): Not applicable

**Agency or authority designated to handle comments:** ☐ National Notification Authority, ☐ National Enquiry Point. Address, fax number and e-mail address (if available) of other body:

For access to the docket to read background documents or comments received, go to <https://www.regulations.gov> and insert the docket number, found in brackets in the heading of this

final rule, into the "Search" box and follow the prompts, and/or go to the Dockets Management Staff (HFA-305), 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, +(240) 402 7500.

For further information contact: Katherine Vierk, Office of Analytics and Outreach, Center for Food Safety and Applied Nutrition, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, +(240) 402 2122, [Katherine.Vierk@fda.hhs.gov](mailto:Katherine.Vierk@fda.hhs.gov).

**Text(s) available from: ☐ National Notification Authority, ☐ National Enquiry Point. Address, fax number and e-mail address (if available) of other body:**

Text can be found in the Federal Register, Vol. 87, No. 223, page 70910 or on the internet at <https://www.govinfo.gov/content/pkg/FR-2022-11-21/pdf/2022-24417.pdf>.

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