NOTIFICATION

Addendum

The following communication, received on 7 December 2021, is being circulated at the request of the Delegation of the United States of America.

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| Laboratory Accreditation for Analyses of Foods; Final Rule |
| The Food and Drug Administration (FDA, the Agency, or we) is amending its regulations to establish a program for the testing of food in certain circumstances by accredited laboratories, as required under the Federal Food, Drug, and Cosmetic Act (FD&C Act). Establishing this program will help FDA improve the safety of the US food supply and protect US consumers by helping ensure that certain food testing of importance to public health is conducted subject to appropriate oversight and in accordance with appropriate model standards to produce reliable and valid test results.This rule is effective 1 February 2022. The incorporation by reference of certain publications listed in the rule is approved by the Director of the Federal Register as of 1 February 2022.<https://members.wto.org/crnattachments/2021/SPS/USA/21_7601_00_e.pdf> |
| **This addendum concerns a:** |
| [ ] Modification of final date for comments |
| [**X**] 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, 5630 Fishers Lane, Rm. 1061, Rockville, MD 20852, 240 402 7500.For further information contact: With regard to the final rule: Stacie Hammack, Chemist, Food and Feed Laboratory Operations, Office of Regulatory Affairs, Food and Drug Administration, 60 8th Street NE, Atlanta, GA 30309, 301 796 5817; Stacie.Hammack@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 US Federal Register, Vol. 86, No. 230, page 68728 or on the Internet at: <https://www.govinfo.gov/content/pkg/FR-2021-12-03/pdf/2021-25716.pdf>. |

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