WORLD TRADE ORGANIZATION

G/SPS/N/USA/3310/Add.1

20 January 2025

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Committee on Sanitary and Phytosanitary Measures Original: English

NOTIFICATION

Addendum

The following communication, received on 20 January 2025, is being circulated at the request of the Delegation of the United States of America.

Listing of Color Additives Exempt From Certification; Myoglobin; Final Amendment; Order

The Food and Drug Administration (FDA or we) is amending the color additive regulations to provide for the safe use of myoglobin as a color additive in ground meat and ground poultry analogue products. We are taking this action in response to a color additive petition (CAP) submitted by Motif FoodWorks, Inc. (Motif FoodWorks or petitioner). This order is effective 19 February 2025.

https://www.federalregister.gov/d/2025-01239
https://members.wto.org/crnattachments/2025/SPS/USA/25_00792_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 the electronic and written/paper comments received, go to https://www.regulations.gov and insert the docket number, found in brackets in the heading of this document, (i.e. FDA–2022–C–0098) 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: Ellen Anderson, Office of Pre-market Additive Safety, Human Foods Program, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740– 3835, +(240) 402 1309 or Keronica Richardson, Office of Policy, Regulations, and Information, Human Foods Program, Food and Drug Administration, 5001 Campus Dr., College Park, MD 20740, +(240) 402 1262.

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. 90, No. 11, Page 5590 or on the Internet at: https://www.federalregister.gov/d/2025-01239.