NOTIFICATION

The following notification is being circulated in accordance with Article 10.6

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| **1.** | **Notifying Member:** EUROPEAN UNION  **If applicable, name of local government involved (Article 3.2 and 7.2):** |
| **2.** | **Agency responsible:**  European Commission  **Name and address (including telephone and fax numbers, email and website addresses, if available) of agency or authority designated to handle comments regarding the notification shall be indicated if different from above:**  European Commission,  EU-TBT Enquiry Point,  Fax: +(32) 2 299 80 43,  E-mail: [grow-eu-tbt@ec.europa.eu](mailto:grow-eu-tbt@ec.europa.eu)  Website: <http://ec.europa.eu/growth/tools-databases/tbt/en/> |
| **3.** | **Notified under Article 2.9.2 [****X],** **2.10.1 [****],** **5.6.2 [****],** **5.7.1 [****], 3.2 [****], 7.2 [****],** **other****:** |
| **4.** | **Products covered (HS or CCCN where applicable, otherwise national tariff heading. ICS numbers may be provided in addition, where applicable):** Food |
| **5.** | **Title, number of pages and language(s) of the notified document:** Draft Commission Delegated Regulation amending the Annex to Regulation (EU) No 609/2013 of the European Parliament and of the Council to allow the use of nicotinamide riboside chloride as a source of niacin in food for special medical purposes and total diet replacement for weight control; (5 page(s), in English), (2 page(s), in English) |
| **6.** | **Description of content:** This draft Commission Delegated Regulation concerns the authorisation of the addition of nicotinamide riboside chloride, as a source of niacin to total diet replacement for weight control and food for special medical purposes in line with EFSA's relevant scientific opinion. |
| **7.** | **Objective and rationale, including the nature of urgent problems where applicable:** The proposed measure aims at allowing the addition of nicotinamide riboside chloride, as a source of niacin to total diet replacement for weight control and food for special medical purposes. The draft measure reflects the favourable outcome of the safety assessment by EFSA. In order to allow the addition of the substance to the mentioned categories of foods it is necessary to amend the Annex to Regulation (EU) No 609/2013 on food intended for infants and young children, food for special medical purposes and total diet replacement for weight control. |
| **8.** | **Relevant documents:**  Regulation (EU) No 609/2013 of the European Parliament and of the Council of 12 June 2013 on food intended for infants and young children, food for special medical purposes and total diet replacement for weight control  <http://eur-lex.europa.eu/LexUriServ/LexUriServ.do?uri=OJ:L:2013:181:0035:0056:en:PDF> |
| **9.** | **Proposed date of adoption:** 4nd quarter 2022  **Proposed date of entry into force:** The proposed measure shall enter into force on the 20th day following its publication in the Official Journal of the EU. |
| **10.** | **Final date for comments:** 60 days from notification |
| **11.** | **Texts available from: National enquiry point [****]** **or address, telephone and fax numbers and email and website addresses, if available, of other body:**  European Commission,  EU-TBT Enquiry Point,  Fax: + (32) 2 299 80 43,  E-mail: [grow-eu-tbt@ec.europa.eu](mailto:grow-eu-tbt@ec.europa.eu)  The text is available on the EU-TBT Website : <http://ec.europa.eu/growth/tools-databases/tbt/en/>  <https://members.wto.org/crnattachments/2022/TBT/EEC/22_6844_00_e.pdf>  <https://members.wto.org/crnattachments/2022/TBT/EEC/22_6844_01_e.pdf> |