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: Preventing International Trade Barriers | TBT - European Commission |
| **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):** Biocidal products |
| **5.** | **Title, number of pages and language(s) of the notified document:** Draft Commission Implementing Regulation renewing the approval of dazomet as an active substance for use in biocidal products of product-type 8 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council; (3 page(s), in English), (2 page(s), in English) |
| **6.** | **Description of content:** This draft Commission Implementing Regulation renews the approval of dazomet as an active substance for use in biocidal products of product-type 8, subject to compliance with certain conditions, including a condition for placing on the market of treated articles treated with or incorporating dazomet.  The opinion of the European Chemicals Agency can be found on its website ([Biocidal Products Committee opinions on active substance approval - ECHA (europa.eu)](https://echa.europa.eu/regulations/biocidal-products-regulation/approval-of-active-substances/bpc-opinions-on-active-substance-approval?p_p_id=viewsubstances_WAR_echarevsubstanceportlet&p_p_lifecycle=0&p_p_state=normal&p_p_mode=view&_viewsubstances_WAR_echarevsubstanceportlet_delta=50&_viewsubstances_WAR_echarevsubstanceportlet_orderByCol=staticField_-104&_viewsubstances_WAR_echarevsubstanceportlet_orderByType=asc&_viewsubstances_WAR_echarevsubstanceportlet_resetCur=false&_viewsubstances_WAR_echarevsubstanceportlet_cur=3)). |
| **7.** | **Objective and rationale, including the nature of urgent problems where applicable:** Harmonisation of the EU market on biocidal products.; Protection of human health or safety; Protection of the environment; Harmonization |
| **8.** | **Relevant documents:**  Regulation (EU) No 528/2012 of the European Parliament and of the Council of 22 May 2012 concerning the making available on the market and use of biocidal products (OJ L 167, 27.6.2012, p. 1.). Available in all EU languages.  [EUR-Lex - 32012R0528 - EN - EUR-Lex (europa.eu)](https://eur-lex.europa.eu/legal-content/EN/TXT/?uri=CELEX%3A32012R0528&qid=1653319893936) |
| **9.** | **Proposed date of adoption:** August 2025  **Proposed date of entry into force:** 20 days from 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 : <https://technical-barriers-trade.ec.europa.eu/en/home>  <https://members.wto.org/crnattachments/2025/TBT/EEC/25_03471_00_e.pdf>  <https://members.wto.org/crnattachments/2025/TBT/EEC/25_03471_01_e.pdf> |