EUROPEAN
COMMISSION

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COMMISSION IMPLEMENTING DECISION (EU) …/…

of XXX

repealing Implementing Decision (EU) 2024/2930 postponing the expiry date of the approval of dazomet 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

(Text with EEA relevance)

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(Text with EEA relevance)

THE EUROPEAN COMMISSION,

Having regard to the Treaty on the Functioning of the European Union,

Having regard to 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[1], and in particular Article 14(5) thereof,

After consulting the Standing Committee on Biocidal Products,

Whereas:

(1) Dazomet was included in Annex I to Directive 98/8/EC of the European Parliament and of the Council[2] as an active substance for use in biocidal products of product-type 8. Pursuant to Article 86 of Regulation (EU) No 528/2012, it was therefore considered approved until 31 July 2022 under that Regulation subject to the conditions set out in Annex I to Directive 98/8/EC.

(2) On 26 January 2021, an application was submitted in accordance with Article 13(1) of Regulation (EU) No 528/2012 for the renewal of the approval of dazomet for use in biocidal products of product-type 8 ('the application'). The application was evaluated by the competent authority of Belgium.

(3) Pursuant to Commission Implementing Decision (EU) 2021/1289[3], the expiry date of approval of dazomet for use in biocidal products of product-type 8 was postponed to 31 January 2025. That expiry date was further postponed to 31 July 2026 by Commission Implementing Decision (EU) 2024/2930[4] in order to allow sufficient time for the examination of the application.

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1 OJ L 167, 27.6.2012, p. 1, ELI: http://data.europa.eu/eli/reg/2012/528/oj.

2 Directive 98/8/EC of the European Parliament and of the Council of 16 February 1998 concerning the placing of biocidal products on the market (OJ L 123, 24.4.1998, p. 1, ELI: http://data.europa.eu/eli/dir/1998/8/oj).

3 Commission Implementing Decision (EU) 2021/1289 of 2 August 2021 postponing the expiry date of approval of dazomet for use in biocidal products of product-type 8 (OJ L 279, 3.8.2021, p. 45, ELI: http://data.europa.eu/eli/dec_impl/2021/1289/oj).

4 Commission Implementing Decision (EU) 2024/2930 of 28 November 2024 postponing the expiry date of the approval of dazomet 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 (OJ L, 2024/2930, 2.12.2024; ELI: http://data.europa.eu/eli/dec_impl/2024/2930/oj).

(4) Commission Implementing Regulation (EU)[5] [*Publication office please add here the number of this Regulation* C(2025) 2500 *once it is associated, along the whole Decision*] renewed the approval of dazomet for use in biocidal products of product-type 8, subject to the conditions in the Annex to that Regulation, including the expiry date of approval. Therefore, it is appropriate to repeal Implementing Decision (EU) 2024/2930 postponing the expiry date of the approval of dazomet,

HAS ADOPTED THIS DECISION:

Article 1

Implementing Decision (EU) 2024/2930 is repealed.

Article 2

This Decision shall enter into force on the twentieth day following that of its publication in the *Official Journal of the European Union*.

Done at Brussels,

For the Commission
The President
Ursula VON DER LEYEN

[5] Commission Implementing Regulation (EU) XXX of XXX 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 (OJ L, …, …, p. …, ELI).