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DECISION OF THE EEA JOINT COMMITTEE No 73/2026

of 20 March 2026

amending Annex II (Technical regulations, standards, testing and certification) to the EEA Agreement [2026/1321]

THE EEA JOINT COMMITTEE,

Having regard to the Agreement on the European Economic Area ("the EEA Agreement"), and in particular Article 98 thereof,

Whereas:

(1) Commission Implementing Regulation (EU) 2025/2345 of 19 November 2025 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 ([1]) is to be incorporated into the EEA Agreement.

(2) Commission Implementing Decision (EU) 2025/2279 of 13 November 2025 postponing the expiry date of the approval of aluminium phosphide releasing phosphine for use in biocidal products of product-types 14, 18 and 20 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([2]) is to be incorporated into the EEA Agreement.

(3) Commission Implementing Decision (EU) 2025/2280 of 13 November 2025 on the unresolved objections regarding the conditions for granting an authorisation for the biocidal product Speed Easy Clean in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([3]) is to be incorporated into the EEA Agreement.

(4) Commission Implementing Decision (EU) 2025/2282 of 13 November 2025 postponing the expiry date of the approval of cholecalciferol for use in biocidal products of product-type 14 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([4]) is to be incorporated into the EEA Agreement.

(5) Commission Implementing Decision (EU) 2025/2283 of 13 November 2025 postponing the expiry date of the approval of lambda-cyhalothrin for use in biocidal products of product-type 18 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([5]) is to be incorporated into the EEA Agreement.

(6) Commission Implementing Decision (EU) 2025/2284 of 13 November 2025 postponing the expiry date of the approval of magnesium phosphide releasing phosphine for use in biocidal products of product-type 18 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([6]) is to be incorporated into the EEA Agreement.

(7) Commission Implementing Decision (EU) 2025/2297 of 13 November 2025 on the unresolved objections regarding the conditions for granting an authorisation for the biocidal product Saltidin 20 % Outdoor in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([7]) is to be incorporated into the EEA Agreement.

(8) Commission Implementing Decision (EU) 2025/2309 of 13 November 2025 on the unresolved objections regarding the conditions for granting an authorisation for the biocidal product ERO MP in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([8]) is to be incorporated into the EEA Agreement.

([1]) OJ L, 2025/2345, 20.11.2025, ELI: http://data.europa.eu/eli/reg_impl/2025/2345/oj.
([2]) OJ L, 2025/2279, 14.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2279/oj.
([3]) OJ L, 2025/2280, 17.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2280/oj.
([4]) OJ L, 2025/2282, 17.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2282/oj.
([5]) OJ L, 2025/2283, 17.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2283/oj.
([6]) OJ L, 2025/2284, 17.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2284/oj.
([7]) OJ L, 2025/2297, 17.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2297/oj.
([8]) OJ L, 2025/2309, 17.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2309/oj.

(9) Commission Implementing Decision (EU) 2025/2299 of 14 November 2025 postponing the expiry date of the approval of *Bacillus thuringiensis* subsp. *israelensis* Serotype H14, Strain AM65-52 for use in biocidal products of product-type 18 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([9]) is to be incorporated into the EEA Agreement.

(10) Commission Implementing Decision (EU) 2025/2301 of 14 November 2025 postponing the expiry date of the approval of deltamethrin for use in biocidal products of product-type 18 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council ([10]) is to be incorporated into the EEA Agreement.

(11) Commission Implementing Decision (EU) 2025/2344 of 19 November 2025 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 ([11]) is to be incorporated into the EEA Agreement.

(12) Implementing Decision (EU) 2025/2344 repeals Commission Implementing Decision (EU) 2024/2930 ([12]), which is incorporated into the EEA Agreement and which is consequently to be repealed under the EEA Agreement.

(13) Annex II to the EEA Agreement should therefore be amended accordingly,

HAS ADOPTED THIS DECISION:

Article 1

Chapter XV of Annex II to the EEA Agreement is amended as follows:

1. The following points are inserted after point 12nnna (Commission Implementing Decision (EU) 2024/816):

‘12nnnb. **32025 R 2345**: Commission Implementing Regulation (EU) 2025/2345 of 19 November 2025 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, 2025/2345, 20.11.2025).

12nnnc. **32025 D 2279**: Commission Implementing Decision (EU) 2025/2279 of 13 November 2025 postponing the expiry date of the approval of aluminium phosphide releasing phosphine for use in biocidal products of product-types 14, 18 and 20 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2279, 14.11.2025).

12nnnd. **32025 D 2280**: Commission Implementing Decision (EU) 2025/2280 of 13 November 2025 on the unresolved objections regarding the conditions for granting an authorisation for the biocidal product Speed Easy Clean in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2280, 17.11.2025).

12nnne. **32025 D 2282**: Commission Implementing Decision (EU) 2025/2282 of 13 November 2025 postponing the expiry date of the approval of cholecalciferol for use in biocidal products of product-type 14 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2282, 17.11.2025).

([9]) OJ L, 2025/2299, 18.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2299/oj.
([10]) OJ L, 2025/2301, 17.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2301/oj.
([11]) OJ L, 2025/2344, 20.11.2025, ELI: http://data.europa.eu/eli/dec_impl/2025/2344/oj.
([12]) OJ L, 2024/2930, 2.12.2024, ELI: http://data.europa.eu/eli/dec_impl/2024/2930/oj.

12nnnf. **32025 D 2283**: Commission Implementing Decision (EU) 2025/2283 of 13 November 2025 postponing the expiry date of the approval of lambda-cyhalothrin for use in biocidal products of product-type 18 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2283, 17.11.2025).

12nnng. **32025 D 2284**: Commission Implementing Decision (EU) 2025/2284 of 13 November 2025 postponing the expiry date of the approval of magnesium phosphide releasing phosphine for use in biocidal products of product-type 18 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2284, 17.11.2025).

12nnnh. **32025 D 2297**: Commission Implementing Decision (EU) 2025/2297 of 13 November 2025 on the unresolved objections regarding the conditions for granting an authorisation for the biocidal product Saltidin 20 % Outdoor in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2297, 17.11.2025).

12nnni. **32025 D 2309**: Commission Implementing Decision (EU) 2025/2309 of 13 November 2025 on the unresolved objections regarding the conditions for granting an authorisation for the biocidal product ERO MP in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2309, 17.11.2025).

12nnnj. **32025 D 2299**: Commission Implementing Decision (EU) 2025/2299 of 14 November 2025 postponing the expiry date of the approval of *Bacillus thuringiensis* subsp. *israelensis* Serotype H14, Strain AM65-52 for use in biocidal products of product-type 18 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2299, 18.11.2025).

12nnnk. **32025 D 2301**: Commission Implementing Decision (EU) 2025/2301 of 14 November 2025 postponing the expiry date of the approval of deltamethrin for use in biocidal products of product-type 18 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (OJ L, 2025/2301, 17.11.2025).

12nnnl. **32025 D 2344**: Commission Implementing Decision (EU) 2025/2344 of 19 November 2025 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 (OJ L, 2025/2344, 20.11.2025).'

2. The text of point 12zzzzzzzzzzzn (Implementing Decision (EU) 2024/2930) is deleted.

Article 2

The texts of Implementing Regulation (EU) 2025/2345 and Implementing Decisions (EU) 2025/2279, (EU) 2025/2280, (EU) 2025/2282, (EU) 2025/2283, (EU) 2025/2284, (EU) 2025/2297, (EU) 2025/2309, (EU) 2025/2299, (EU) 2025/2301 and (EU) 2025/2344 in the Icelandic and Norwegian languages, to be published in the EEA Supplement to the *Official Journal of the European Union*, shall be authentic.

Article 3

This Decision shall enter into force on 21 March 2026, provided that all the notifications under Article 103(1) of the EEA Agreement have been made (*).

(*) No constitutional requirements indicated.

Article 4

This Decision shall be published in the EEA Section of, and in the EEA Supplement to, the *Official Journal of the European Union*.

Done at Brussels, 20 March 2026.

For the EEA Joint Committee
The President
Nicolas VON LINGEN