Miljø- og Ligestillingsministeriet
Departementet

Vand og pesticider
J.nr. 2025 - 687
Ref. PIJEN
Den 10. juni 2025

Høring om foreløbige forslag til Kommissionens gennemførelsesforordninger om fornyet godkendelse og administrativ forlængelse af godkendelser af biocidaktivstoffer, jf. Europa-Parlamentets og Rådets forordning (EU) nr. 528/2012 om tilgængeliggørelse på markedet og anvendelse af biocidholdige produkter

Kommissionen har nedenstående forslag på dagsordenen til drøftelse i den stående komité for biocidholdige produkter den 20. juni 2025. Det forventes, at Kommissionen vil sætte forslagene til skriftlig afstemning i 3. kvartal 2025. Nærmere dato for afstemningerne kendes ikke.

Kommissionens forslag ligger inden for standardmandatet på biocidområdet, og det følger deraf, at Danmark agter at støtte forslagene. Standardmandatet kan findes på Folketingets hjemmeside: (https://www.ft.dk/samling/20231/almdel/MOF/bilag/621/2899620.pdf).

Forslag

B.01 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of lauric acid for use in biocidal products of product-type 19 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.02 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of ethyl butylacetylaminopropionate for use in biocidal products of product-type 19 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.03 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of transfluthrin 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

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.04 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of azoxystrobin for use in biocidal products of product-types 7, 9 and 10 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.05 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of zineb for use in biocidal products of product-type 21 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.06 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of copper pyrithione for use in biocidal products of product-type 21 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.07 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of copper thiocyanate for use in biocidal products of product-type 21 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.08 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of dicopper oxide for use in biocidal products of **product-type 21** in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.09 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of *Pythium oligandrum* strain M1 for use in biocidal products of product-type 10 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

B.10 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of imidacloprid 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

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)

Procedure: Rådgivningsprocedure

B.11 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of 4,5-Dichloro-2-octyl-2H-isothiazol-3-one for use in biocidal products of product-types 8 and 21 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

C.01 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) 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

Legal Basis: Regulation (EU) No 528/2012 - Article 14(4)(a)
Procedure: Undersøgelsesprocedure

C.02 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) 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

Legal Basis: Regulation (EU) No 528/2012 - Article 14(5)
Procedure: Rådgivningsprocedure

For punktet C.01 forligger der udtalelse fra Biocidkomiteen. Denne udtalelse har dannet den faglige baggrund for det tilknyttede forslag fra Kommissionen. Udtalelsen kan findes på følgende link:

C.01 dazomet	https://echa.europa.eu/documents/10162/39f22a7f-3697-1b44-a4ec-c1fa57e79e21

Miljø- og Ligestillingsministeriet skal bede om eventuelle bemærkninger senest **mandag d. 16. juni 2025**. Bemærkninger kan sendes til Pia Haugaard Jensen (pijen@mim.dk) med kopi til mim@mim.dk. Spørgsmål kan ligeledes rettes til Pia Haugaard Jensen på pijen@mim.dk.