



Biocider
J.nr. 2026 - 129287
Ref. NIFAL
Den 7. oktober 2026

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

Kommissionen har nedenstående forslag til skriftlig afstemning, som følge af drøftelse i den stående komité for biocidholdige produkter den 1. oktober 2026. Punkter vedr. medetomidin blev senest drøftet i den stående komité for biocidholdige produkter den 25. juni 2026.

Kommissionens afstemning er igangsat med deadline 21. oktober 2026.

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 of the Committee on a draft Commission Implementing Decision (EU) not approving 1,3-Bis(hydroxymethyl)-5,5-dimethylimidazolidine-2,4-dione (DMDMH) as an existing active substance for use in biocidal products of product types 6 and 13 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal basis: Regulation (EU) No 528/2012 – Article 89(1)

Procedure: Undersøgelsesprocedure

- B.02** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of indoxacarb 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

Kommissionen foreslag vil forlænge aktivstoffets optag til 31. december 2028.

- B.03** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of N,N-diethyl-meta-toluamide 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

Kommissionen foreslag vil forlænge aktivstoffets optag til 31. januar 2029

- B.04** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of hydrogen cyanide for use in biocidal products of

product-types 8, 14 and 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

Kommissionen foreslår vil forlænge aktivstoffets optag til 30. september 2029

- B.05** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of *Bacillus thuringiensis* subsp. *kurstaki*, serotype 3a3b, strain ABTS-351 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

Kommissionen foreslår vil forlænge aktivstoffets optag til 31. august 2029

- B.06** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) 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

Legal basis: Regulation (EU) No 528/2012 – Article 14(5)

Procedure: Rådgivningsprocedure

Kommissionen foreslår vil forlænge aktivstoffets optag til 31. december 2027

- B.07** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of brodifacoum, bromadiolone, chlorophacinone, coumatetralyl, difenacoum, difethialone and flocoumafen 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

Legal basis: Regulation (EU) No 528/2012 – Article 14(5)

Procedure: Rådgivningsprocedure

Kommissionen foreslår vil forlænge aktivstoffets optag til 31. december 2027

- C.02** Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) approving peanut butter as an existing active substance 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 89(1)

Procedure: Undersøgelsesprocedure

- C.04** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) on the non-approval of active bromine generated from sodium bromide by electrolysis, alkyl (C12-C14) dimethylbenzylammonium chloride (ADBAC (C12-C14)) and hydrogen peroxide as active substances for use in biocidal products of the respective product-types 11, 12, 22 and 12 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

Legal basis: Regulation (EU) No 528/2012 – Article 89(1)

Procedure: Undersøgelsesprocedure

- C.05** Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) renewing 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

Legal basis: Regulation (EU) No 528/2012 – Article 14(4)(a)

Procedure: Undersøgelsesprocedure

- C.06** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) repealing Implementing Decision (EU) 2025/2279 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

Legal basis: Regulation (EU) No 528/2012 – Article 14(5)

Procedure: Undersøgelsesprocedure

- C.07** Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) renewing 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

Legal basis: Regulation (EU) No 528/2012 – Article 14(4)(a)

Procedure: Undersøgelsesprocedure

- C.08** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) repealing Implementing Decision (EU) 2025/2284 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

Legal basis: Regulation (EU) No 528/2012 – Article 14(5)

Procedure: Undersøgelsesprocedure

- C.09** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) not renewing the approval of bromoacetic acid for use in biocidal products of product-type 4 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)b

Procedure: Undersøgelsesprocedure

- C.10** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) repealing Implementing Decision (EU) 2025/360 postponing the expiry date of the approval of bromoacetic acid for use in biocidal products of product-type 4 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: Undersøgelsesprocedure

- C.01** Exchange of views of the Committee on a draft Commission Implementing Decision not renewing the approval of medetomidine as an active substance 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(4)b

Procedure: Undersøgelsesprocedure

- C.02** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) repealing Implementing Decision (EU) 2026/1089 postponing the expiry date of the approval of medetomidine 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: Legal Basis: Regulation (EU) No 528/2012 – Article 14(5)

Procedure: Undersøgelsesprocedure

For punkterne B.01 og C.01 foreligger der udtalelser fra Biocidkomiteen. Disse udtalelser har dannet den faglige baggrund for de tilknyttede forslag fra Kommissionen. Udtalelserne kan findes på følgende links:

B.01 1,3-Bis(hydroxymethyl)-5,5-dimethylimidazolidine-2,4-dione (DMDMH) i produkttype 6	https://echa.europa.eu/documents/10162/c7654e23-783e-585b-2c50-e55fdf1d3ec9
B.01 1,3-Bis(hydroxymethyl)-5,5-dimethylimidazolidine-2,4-dione (DMDMH) i produkttype 13	https://echa.europa.eu/documents/10162/c77045d9-b93c-5a75-ee22-bd67eeb36d49
C.01 Medetomidin i produkttype 21	https://echa.europa.eu/documents/10162/c414113a-719e-cd77-89eb-ce37315fa2a2

Punkterne C.04 og C.09 omfatter forslag om ikke-optag, da der ikke længere er konkrete ansøgere, som støtter en ansøgning om hhv. godkendelse og fornyet godkendelse af de pågældende kombinationer af aktivstoffer og produkttyper. Information kan tilgås på Kommissionens hjemmeside:

C.04 active bromine generated from sodium bromide by electrolysis, alkyl (C12-C14) dimethylbenzylammonium chloride (ADBAC (C12-C14)) and hydrogen peroxide	https://ec.europa.eu/transparency/comitology-register/screen/documents/118242/1/consult
C.09 Bromoacetic acid	https://ec.europa.eu/transparency/comitology-register/screen/documents/118247/1/consult

Miljøstyrelsen skal bede om eventuelle bemærkninger senest fredag d. 16 oktober 2026. Bemærkninger kan sendes til biocides@mst.dk. Spørgsmål kan ligeledes rettes til biocides@mst.dk