



Kemikalier og biocider
J.nr. 2026 - 34160
Ref. nifal
Den 14. april 2026

Høring om foreløbige forslag til Kommissionens gennemførelsesforordninger om fornyet godkendelse, ikke-godkendelse, ikke-fornyelse og administrative 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 fremsat forslag til punkterne C.03, C.04, C.07-C.10 til skriftlig afstemning, som følge af drøftelse i den stående komité for biocidholdige produkter den 18. marts 2026. Skriftlig afstemning skal ske senest 24. april 2026.

Det forventes, at Kommissionen vil fremsætte forslag til punkterne C.01, C.02, C.05 og C.06 til skriftlig afstemning i 2. kvartal 2026. 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/almindel/MOF/bilag/621/2899620.pdf>

Forslag

C.01 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) not approving terbutryn, 1,2-Benzisothiazol-3(2H)-one (BIT), and tetrahydro 1,3,4,6-tetrakis(hydroxymethyl)imidazo[4,5-d]imidazole-2,5(1H,3H)-dione (TMAD) for use in biocidal products of product-types 9, 9, and 12, respectively, 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.02 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) not approving poly(dimethyloctadecyl[3-(trihydroxysilyl)propyl]ammonium chloride) generated from dimethyloctadecyl[3-(trimethoxysilyl)propyl]ammonium chloride as an existing active substance for use in biocidal products of product types 2, 7 and 9

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

Procedure: Undersøgelsesprocedure

- C.03** Exchange of views of the Committee on a draft Commission Implementing Decision not renewing the approval of cypermethrin 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)(b)
Procedure: Undersøgelsesprocedure
- C.04** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) repealing Implementing Decision (EU) 2025/362 postponing the expiry date of the approval of cypermethrin 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
- C.05** Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) renewing the approval of cis-tricos-9-ene as an 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 14(4)(a)
Procedure: Undersøgelsesprocedure
- C.06** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) repealing Implementing Decision (EU) 2024/1283 postponing the expiry date of the approval of cis-tricos-9-ene 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
- C.07** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) 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:** Regulation (EU) No 528/2012 – Article 14(5)
Procedure: Rådgivningsprocedure
- C.08** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of IPBC 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
- C.09** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of boric acid 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

- C.10** Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of disodium tetraborate 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 punkterne C.02 og C.05 foreligger der udtalelser fra Biocidkomiteen. Disse udtalelser har dannet den faglige baggrund for de tilknyttede forslag fra Kommissionen. Udtalelserne kan findes på følgende links:

C.02 poly(dimethyloctadecyl[3-(trihydroxysilyl)propyl]ammonium chloride) generated from dimethyloctadecyl[3-(trimethoxysilyl)propyl]ammonium chloride i produkttype 2	https://echa.europa.eu/documents/10162/cb5d4c1d-0c40-9e2e-21a4-10f70bdb48dc
C.02 poly(dimethyloctadecyl[3-(trihydroxysilyl)propyl]ammonium chloride) generated from dimethyloctadecyl[3-(trimethoxysilyl)propyl]ammonium chloride i produkttype 7	https://echa.europa.eu/documents/10162/241760da-1a3b-6dc3-5838-be9c3ff54cd3
C.02 poly(dimethyloctadecyl[3-(trihydroxysilyl)propyl]ammonium chloride) generated from dimethyloctadecyl[3-(trimethoxysilyl)propyl]ammonium chloride i produkttype 9	https://echa.europa.eu/documents/10162/f710e1e4-1973-350e-3f49-097f1030853b
C.05 cis-tricos-9-ene	https://echa.europa.eu/documents/10162/039c1f27-5c51-092d-c27d-bf573de9d4d3

Punkterne C.01 og C.03 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.01 terbutryn, 1,2-Benzisothiazol-3(2H)-one (BIT), and tetrahydro 1,3,4,6-tetrakis(hydroxymethyl)imidazo[4,5-d]imidazole-2,5(1H,3H)-dione (TMAD)	https://technical-barriers-trade.ec.europa.eu/en/notification/38895
C.03 cypermethrin	https://technical-barriers-trade.ec.europa.eu/en/notification/38733

Miljøstyrelsen skal bede om eventuelle bemærkninger senest tirsdag d. 21. april 2026. Bemærkninger kan sendes til biocides@mst.dk.

Spørgsmål kan ligeledes rettes til biocides@mst.dk.