EUROPEAN COMMISSION

Health and Food Safety Directorate General

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87th Standing Committee on Biocidal Products

19 March 2025

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SUMMARY REPORT

A.01 Adoption of the Agenda

The agenda was adopted.

A.02 Adoption of the minutes of the 86th SCBP meeting

The minutes of the 86th SCBP meeting were adopted.

A.03 Discussion on the application of approval of formaldehyde released from the reaction products of paraformaldehyde and 2-hydroxypropylamine (ratio 1:1), as an existing active substance for use in biocidal products of product-types 2, 6, 11 and 13 , and of reaction products of paraformaldehyde and 2-hydroxypropylamine (ratio 3:2) as an existing active substance for use in biocidal products of product-types 2, 6, 11, 12 and 13

A discussion took place on the application of approval of formaldehyde released from the reaction products of paraformaldehyde and 2-hydroxypropylamine (ratio 1:1), as an existing active substance for use in biocidal products of product-types 2, 6, 11 and 13, and of reaction products of paraformaldehyde and 2-hydroxypropylamine (ratio 3:2) as an existing active substance for use in biocidal products of product-types 2, 6, 11, 12 and 13.

A.04 Discussion on the application of approval of DBNPA as an existing active substance for use in biocidal products of product-type 11

A discussion took place on the application of approval of DBNPA as an existing active substance for use in biocidal products of product-type 11.

A.05 Discussion on the application of renewal of approval of medetomidine as an existing active substance for use in biocidal products of product-type 21

A discussion took place on the application of renewal of approval of medetomidine as an existing active substance for use in biocidal products of product-type 21.

A.06 Information on a draft Commission Implementing Regulation specifying a procedure for the authorisation of same biocidal products in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (repealing Commission Implementing Regulation (EU) No 414/201)

The Commission provided information on a draft Implementing Regulation specifying a procedure for the authorisation of same biocidal products in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council (repealing Commission Implementing Regulation (EU) No 414/2013).

A.07 Information on the unresolved objections regarding the conditions for granting an authorisation for the biocidal product families Cypermethrin liquids and Cypermethrin solids, and on the biocidal product Icon 10 CS, in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission informed that internal discussions are still on-going within the Commission.

A.08 Information on decisions on amendments to Union authorisations under preparation or published

The Commission informed on amendments to Union authorisations under preparation or published. The Commission recalled the importance of the quality check of the translations of the Summary of Products Characteristics annexed to the authorisation.

B.01 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision on the non-approval of ethylene oxide for use in biocidal products of product type 2 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Decision. A discussion took place and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.02 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation approving 2,2-Dibromo-2-cyanoacetamide (DBNPA) as an existing active substance for use in biocidal products of product-type 6 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place, and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.03 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation approving formic acid as an existing active substance for use in biocidal products of product-type 6 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place, and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.04 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation approving 1,2-Benzisothiazol-3(2H)-one (BIT) 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

The Commission introduced the draft Regulation. A discussion took place, and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.05 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) granting a Union authorisation for the biocidal product family 'B.Braun Medical Propanol Family' in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place, and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.06 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) granting a Union authorisation for the biocidal product family 'BPF Propan-2-ol Dr Deppe' in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place, and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.07 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) granting a Union authorisation for the single biocidal product 'Fernox Biocide AF10', in accordance with Regulation (EU) No 582/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place, and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.08 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) granting a Union authorisation for the single biocidal product 'Hydrocid 306', in accordance with Regulation (EU) No 582/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place, and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.09 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Regulation (EU) granting a Union authorisation for the biocidal product family 'PAA family UCO Group', in accordance with Regulation (EU) No 582/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place, and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

B.10 Exchange of views and possible opinion of the Committee on a draft Commission Implementing Decision concerning the extension of the action taken by the Environment Agency of Luxembourg permitting the making available on the market and use of the biocidal product Biobor JF in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Decision. A discussion took place and the Commission informed that the draft will be included in the next written consultation of the Standing Committee.

Vote Postponed

C.01 Exchange of views of the Committee on a draft Commission Implementing Regulation amending Implementing Regulation (EU) No 564/2013 as regards the adaptation of fees to inflation

The Commission introduced the draft Regulation. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.02 Exchange of views of the Committee on a draft Commission Implementing Regulation approving 2-methyl-2H-isothiazol-3-one (MIT) as an existing active substance for use in biocidal products of product-type 6 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.03 Exchange of views of the Committee on a draft Commission Implementing Regulation approving 2-methyl-2,3-dihydro-1,2-thiazol-3-one hydrochloride for use in biocidal products of product type 6 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.04 Exchange of views of the Committee on a draft Commission Implementing Regulation approving peracetic acid generated from 1,3-diacetyloxypropan-2-yl acetate and hydrogen peroxide as an existing active substance for use in biocidal products of product-type 2 in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.05 Exchange of views of the Committee on a draft Commission Implementing Regulation renewing the approval of epsilon-metofluthrin as an active substance 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

The Commission introduced the draft Regulation. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.06 Exchange of views of the Committee on a draft Commission Implementing Decision repealing Implementing Decision (EU) 2024/2460 postponing the expiry date of the approval of metofluthrin 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

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.07 Exchange of views of the Committee on a draft Commission Implementing Decision on the non-approval of epsilon-metofluthrin 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

As the draft Decision was not yet available, no discussion took place on this agenda item.

C.08 Exchange of views of the Committee on a draft Commission Implementing Decision not approving certain active substances for use in biocidal products in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.09 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of nonanoic acid for use in biocidal products of product-type 2

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.10 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval of bis(N-cyclohexyl-diazenium-dioxy)-copper (Cu-HDO) for use in biocidal products of product-type 8

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.11 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval decanoic acid for use in biocidal products of product-type 4

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.12 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval octanoic acid for use in biocidal products of product-type 8

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.13 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval iodine for use in biocidal products of product-type 3

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.14 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval polyvinylpyrrolidone iodine for use in biocidal products of product-types 1 and 3

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.15 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval 1R-trans phenothrin for use in biocidal products of product-type 18

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.16 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval S-methoprene for use in biocidal products of product-type 18

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.17 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval medetomidine for use in biocidal products of product-type 21

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.18 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval IPBC for use in biocidal products of product-type 8

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.19 Exchange of views of the Committee on a draft Commission Implementing Decision (EU) postponing the expiry date of the approval DDACarbonate for use in biocidal products of product-type 8

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.20 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) granting a Union authorisation for the biocidal product family 'desmanol pure' in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.21 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) granting a Union authorisation for the single biocidal product 'exeol air cid 01', in accordance with Regulation (EU) No 582/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.22 Exchange of views of the Committee on a draft Commission Implementing Regulation (EU) granting a Union authorisation for the biocidal product family 'PRODHYNET's Lactic acid based products', in accordance with Regulation (EU) No 582/2012 of the European Parliament and of the Council

The Commission introduced the draft Regulation. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.23 Exchange of views of the Committee on a draft Commission Implementing Regulation granting a Union authorisation for the biocidal product family 'CHLOROCRESOL BASED PRODUCTS-CID Lines NV' in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission explained that no documents were provided but that following the receipt of the Agency's opinion on an Article 75(1)(g) request a revised version will be prepared for the next meeting.

C.24 Exchange of views of the Committee on a draft Commission Implementing Decision concerning the extension of the action taken by the Ministry of Health of the Czech Republic permitting the making available on the market and use of the biocidal product Tandem in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.

C.25 Exchange of views of the Committee on a draft Commission Implementing Decision concerning the extension of the action taken by the Dutch Ministry of Infrastructure and Water Management permitting the making available on the market and use of the biocidal product Biobor JF in accordance with Regulation (EU) No 528/2012 of the European Parliament and of the Council

The Commission introduced the draft Decision. A discussion took place and the Committee agreed that the draft will be included in a written consultation of the Standing Committee as soon as possible.