04 December 2019
BPC-A-33-2019_rev2

Draft agenda

33rd meeting of the Biocidal Products Committee (BPC)

10 - 11 December 2019

ECHA Conference Centre, Annankatu 18, Helsinki

Starts on 10 December at 09:30, ends on 11 December at 18:00

1. – Welcome and apologies

2. – Agreement of the agenda

BPC-A-33-2019_rev2

For agreement

3. – Declarations of potential conflicts of interest to agenda items

4. – Agreement of the minutes and review of actions from BPC-31

BPC-M-31-2019

For agreement

5. – Administrative issues

5.1. Housekeeping issues

For information

5.2. Other administrative issues and report from other Committees

For information

5.3. ECHA Activities Coordination Tool (ACT)

For information

6. – Work programme for BPC

6.1. BPC Work Programme for active substance approval

BPC-33-2019-01

For information

6.2. BPC Work Programme for Union authorisation

BPC-33-2019-02

For information

6.3. Outlook for BPC

BPC-33-2019-03

For information

6.4. Status ED assessment for active substances

BPC-33-2019-04

For information

7. – Applications for approval of active substances*

7.1. Procedural and administrative aspects:

7.1.1. Catalogue of specific conditions and elements to be taken into account at the product authorisation stage for active substance approval

BPC-33-2019-05

For information

7.1.2. Revised opinion template for active substance approval

BPC-33-2019-06

For agreement

7.1.3. Opinion request Commission pursuant to Article 75(1)(g) on sodium chloride specifications and water quality for the generation of active chlorine by electrolysis

BPC-33-2019-07

For information

7.2. Follow-up BPC opinion on DBNPA for PT 4 following BPC-31

BPC-33-2019-08A,B

For agreement

* For the discussions of the draft BPC opinions at least the following documents will be distributed: a draft BPC opinion (denoted by A), a draft assessment report (AR) which may cover more than one PT (denoted by B) and a document containing open issues covering all the PTs to be discussed for that substance (denoted by C).

7.3. Draft BPC opinion on icaridin for PT 19

Previous discussion: BPC-28

BPC-33-2019-09A, B, C

For adoption

7.4. Draft BPC opinion on cyanamide for PT 3 and 18

Previous discussion: BPC-16

PT 3: BPC-33-2019-10A, B, C

PT 18: BPC-33-2019-11A and BPC-33-2019-10B, C

For adoption

7.5. Draft BPC opinion on formaldehyde for PT 2 and 3

Previous discussion: BPC-13

PT 2: BPC-33-2019-12A, B, C

PT 3: BPC-33-2019-13A, B, C

For adoption

7.6. Draft BPC opinion on carbendazim for PT 7 and 10

Previous discussion: BPC-25

PT 7: BPC-33-2019-14A, B, C

PT 10: BPC-33-2019-15A and BPC-33-2019-14 B, C

For adoption

8. – Union authorisation**

8.1 Update on Union authorisation

For information

8.1.1 Revised opinion template for Union authorisation

BPC-33-2019-19

For agreement

8.1.2 Introducing new data during the peer review phase for applications for Union authorisation

BPC-33-2019-20

For agreement

8.1.3 Implementation of CA document on "Addressing concerns of co-formulants that contribute significantly to a product's efficacy" (CA-Jan18-Doc.4.2_final)

BPC-33-2019-21

For discussion

** For the discussions of the draft BPC opinions at least the following documents will be distributed: a draft BPC opinion (denoted by A), a draft Summary of Product Characteristics (SPC) (denoted by B), a draft product assessment report (PAR) (denoted by C) and a document containing open issues to be discussed for the biocidal product or biocidal product familiy (denoted by D).

8.1.4 Revised procedure "Linguistic review of the translations of the summary of product characteristics (SPC) for Union authorisation applications

BPC-33-2019-22

For agreement

8.2 Draft BPC opinions on Union authorisation applications for a product family containing propan-2-ol

Previous discussion: WG-IV-2019

BPC-33-2019-23A, B, C, C1, D, E

For adoption

8.3 Draft BPC opinions on Union authorisation applications for a product family containing iodine / PVP-iodine

Previous discussion: WG-IV-2019

BPC-33-2019-24A, B, C, D

For adoption

9. - Any other business

9.1 Applying the ECHA Read-Across Assessment Framework in biocides

BPC-33-2019-25A, B

For agreement

9.2 Request to ECHA on guidance for risk assessment for bees

BPC-33-2019-26A, B

For information

10. - Action points and conclusions

For agreement

Provisional time schedule for the

33rd meeting of the Biocidal Products Committee (BPC)

ECHA Conference Centre, Annankatu 18, Helsinki

10 December 2019: starts at 09:30; 11 December 2019 ends at 18:00

Please note that the time schedule indicated below is provisional and subject to possible change. The schedule is distributed to participants on a preliminary basis. If needed, follow-up discussions may take place on the following day for BPC opinions.

Tuesday 10 December: morning session

Items 1-5	Opening items and administrative issues	
Item 5.3	ECHA Activities Coordination Tool (ACT)	
Item 6	Work programme for BPC	
	6.1.	BPC Work Programme for active substance approval
	6.2.	BPC Work Programme for Union authorisationl
	6.3.	Outlook for BPC
	6.4.	Status ED assessment for active substances
Item 7.1	Procedural and administrative aspects:	
	7.1.1.	Catalogue of specific conditions and elements to be taken into account at the product authorisation stage for active substance approval
	7.1.2.	Revised opinion template for active substance approval
	7.1.3.	Opinion request Commission pursuant to Article 75(1)(g) on sodium chloride specifications and water quality for the generation of active chlorine by electrolysis
Item 7.2	Follow-up BPC opinion on DBNPA for PT 4 following BPC-31	
Item 7.3	Draft BPC opinion on icaridin for PT 19	

Tuesday 10 December: afternoon session

Item 7.4	Draft BPC opinion on cyanamid for PT 3 and 18	
Item 7.5	Draft BPC opinion on formaldehyde for PT 2 and 3	
Item 7.6	Draft BPC opinion on carbendazim for PT 7 and 10	
Item 8.1	Update on Union authorisation	
	8.1.1.	Revised opinion template for Union authorisation
	8.1.2.	Introducing new data during the peer review phase for applications for Union authorisation
	8.1.3.	Implementation of CA document on “Addressing concerns of co-formulants that contribute significantly to a product’s efficacy” (CA-Jan18-Doc.4.2_final)
	8.1.4.	Revised procedure “Linguistic review of the translations of the summary of product characteristics (SPC) for Union authorisation applications

Wednesday 11 December: morning session

Item 8.2	Draft BPC opinions on Union authorisation applications for a product family containing propan-2-ol
Item 8.3	Draft BPC opinions on Union authorisation applications for a product family containing iodine / PVP-iodine

Wednesday 11 December: afternoon session

Item 9.1	Introduction of ECHA Read-across assessment framework in biocides
Item 9.2	Request to ECHA on guidance for risk assessment for bees
Item 10	Action points and conclusions

End of meeting

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