

21 November 2017 BPC-A-23-2017_rev1

Draft agenda

23rd meeting of the Biocidal Products Committee (BPC)

11 – 14 December 2017
ECHA Conference Centre, Annankatu 18, Helsinki
Starts on 11 December at 13:30,
ends on 14 December at 16:00

 – Welcome and 	l apo	loaies
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2. - Agreement of the agenda

BPC-A-23-2017

For agreement

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

4. - Agreement of the minutes and review of actions from BPC-22

BPC-M-22-2017

For agreement

5. - Administrative issues

5.1. Housekeeping issues

For information

5.2. Other administrative issues and report from other Committees

BPC-23-2017-01

For information

6. - Work programme for BPC

6.1. Revised BPC Work Programme 2017-2018

BPC-23-2017-02

For information

6.2. Outlook for BPC

BPC-23-2017-03

For information

7. - Applications for approval of active substances*

7.1. Templates and formats for active substance approval: catalogue of specific conditions and elements to be taken into account at the product authorisation stage for active substance approval

BPC-23-2017-04

For information

7.2. Draft BPC opinion on cholecalciferol for PT 14

Previous discussion(s): WG-I-2017; BPC-21

BPC-23-2017-05A, B, C and D

For adoption

7.3. Draft BPC opinion on formaldehyde for PT 2

Previous discussion(s): TM-I-2012; (withdrawn from BPC-13)

BPC-23-2017-06A, B and C

For adoption

7.4. Draft BPC opinion on empenthrin for PT 18

Previous discussion(s): WG-IV-2017

BPC-23-2017-07A, B and C

For adoption

7.5. Draft BPC opinion on cyphenothrin for PT 18

Previous discussion(s): WG-II-2017; WG-III-2017

BPC-23-2017-08A, B and C

For adoption

7.6. Draft BPC opinion on penflufen for PT 8

Previous discussion(s): WG-IV-2017

BPC-23-2017-09A, B and C

For adoption

^{*} 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).

7.7. Outcome written procedure on the adoption of the BPC opinion on acetamiprid for PT 18

BPC-23-2017-10

For information

7.8. Revised Assessment Report following the submission of data after active substance approval:

7.8.1. MBIT for PT 6

BPC-23-2017-11

For agreement

7.8.2. DCPP for PT 1, 2 and 4

BPC-23-2017-12

For agreement

7.8.3. CMK for PT 1, 2, 3, 6, 9 and 13

BPC-23-2017-19

For agreement

7.8.4. Cyfluthrin for PT 18

BPC-23-2017-21

For agreement

7.9. Question eCA (France) on the evaluation of carbon dioxide generated in-situ for PT 19

BPC-23-2017-22

For information

Item 8 - Union authorisation†

- 8.1 Update on Union authorisation
- 8.2 Draft BPC opinions on Union authorisation applications for product families containing iodine / PVP-iodine

BPC-23-2017-13A, B, C and D

For adoption

BPC-23-2017-14A, B, C and D

For adoption

[†] 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).

Item 9 - Article 38 opinions

9.1 Draft BPC opinion on unresolved objections during the mutual recognition of two insect repellents

BPC-23-2017-15A and B

For adoption

Item 10 - Article 75(1)(g) opinions

10.1 Draft BPC opinion on inclusion into Annex I of corn cob

BPC-23-2017-16A and B

For adoption

10.2 Draft BPC opinion on copper sulphate pentahydrate - PT 3

BPC-23-2017-17

For adoption

10.3 Draft BPC opinion on eligibility of certain food and feed active substances for inclusion into Annex I

BPC-23-2017-18

For adoption

Item 11 - Any other business

Item 12 - Action points and conclusions

For agreement



Provisional time schedule for the 23rd meeting of the Biocidal Products Committee (BPC)

ECHA Conference Centre, Annankatu 18, Helsinki 11 December 2017: starts at 13:30; 14 December ends at 16:00

Please note that the time schedule indicated below are 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.

Monday 11 December: afternoon session

Items 1-5	Opening items and administrative issues
Item 6	Work programme of the BPC 2017-18
Item 10.1	Draft BPC opinion on inclusion into Annex I of corn cob
Item 10.2	Draft BPC opinion on copper sulphate pentahydrate – PT 3
Item 7.9	Question eCA (France) on the evaluation of carbon dioxide generated in-situ for PT 19

Tuesday 12 December: morning session

Item 8.1	Update on Union authorisation
Item 8.2	Draft BPC opinion on Union authorisation applications for product families containing iodine / PVP-iodine

Tuesday 12 December: afternoon session

Item 8.2		opinion ntaining i			applications ed)	for	product
Item 9.1		opinion of two in		objectio	ns during	the	mutual

Wednesday 13 December: morning session

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Item 7.1	Templates and formats for active substance approval: catalogue of specific conditions and elements to be taken into account at the product authorisation stage for active substance approval
Item 7.2	Draft BPC opinion on cholecalciferol for PT 14
Item 7.3	Draft BPC opinion on formaldehyde for PT 2

Wednesday 13 December: afternoon session

Item 7.4	Draft BPC opinion on empenthrin for PT 18
Item 7.5	Draft BPC opinion on cyphenothrin for PT 18

Thursday 14 December: morning session

Item 7.6	Draft BPC opinion on penflufen for PT 8
Item 7.7	Outcome written procedure on the adoption of the BPC opinion on acetamiprid for PT 18 $$
Item 7.8	Revised Assessment Report following the submission of data after active substance approval:
Item 7.8.1	MBIT for PT 6
Item 7.8.2	DCPP for PT 1, 2 and 4
Item 7.8.3	CMK for PT 1, 2, 3, 6, 9 and 13
Item 7.8.4	Cyfluthrin for PT 18

Thursday 14 December: afternoon session

Item 10.3	Draft BPC opinion on eligibility of certain food and feed active substances for inclusion into Annex I
Item 11	AOB
Item 12	Action points and conclusions

End of meeting

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