29 September 2016
BPC-A-17-2016_rev2

Draft agenda

17th meeting of the Biocidal Products Committee (BPC)

11-12 October 2016

ECHA Conference Centre, Annankatu 18, Helsinki

Starts on 11 October at 09:30, ends on 12 October at 13:00

1. – Welcome and apologies

2. – Agreement of the agenda

BPC-A-17-2016-rev2
For agreement

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

4. – Agreement of the minutes and review of actions from BPC-16

BPC-M-16-2015
For agreement

5. – Administrative issues

5.1. Housekeeping issues

For information

5.2. Other administrative issues and report from other Committees

BPC-17-2016-01
For information

6. – Work programme for BPC

6.1. BPC Work Programme

BPC-17-2016-02 and BPC-17-2016-03
For information

7. – Applications for approval of active substances*

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

For information

7.2. Procedure for renaming of an active substance according to Article 13 of Regulation (EU) No 1062/2014

BPC-17-2016-05a and 5b

For agreement

7.3. Outcome of the e-consultation Ad-hoc Working Group on Environmental Exposure

BPC-17-2016-06

For agreement

7.4. Draft BPC opinion on dichlofluanid for PT 21

Previous discussion(s): WG-III-2016

BPC-17-2016-11A, B and C

For adoption

7.5. Draft BPC opinion on silicium dioxide (Kieselguhr) for PT 18

Previous discussion(s): WG-III-2016

BPC-17-2016-08A, B and C

For adoption

7.6. Draft BPC opinion on silicon dioxide (as a nanomaterial formed by aggregates and agglomerates) (Degussa/Evonik) for PT 18

Previous discussion(s): WG-III-2016

BPC-17-2016-09A, 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 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. Draft BPC opinion on PHMB (1660; 1.8) for PT 5

Previous discussion(s): WG-III-2016

BPC-17-2016-10A, B and C

For adoption

7.8. Working procedure for Union Authorisation

BPC-17-2016-12a and 12b

For agreement

7.9. Additional data submitted after active substance approval:

a. 1R-trans-phenothrin

BPC-17-2016-13a, 13b, 13c and 13d

For information

b. copper, granulated

For information

Item 8 – Any other business

a. Consultation BPC on ECHA report on the regulatory applicability of alternative and non-animal approaches (3R's)

Item 9 – Agreement of the action points and conclusions

For agreement

Provisional timeline for the

17th meeting of the Biocidal Products Committee (BPC)

ECHA Conference Centre, Annankatu 18, Helsinki

11 October 2016: starts at 09:30; 12 October 2016: ends at 13:00

Please note that the timings indicated below are provisional and subject to possible change. They are distributed to participants on a preliminary basis.

Tuesday 11 October: morning session

Items 1-5	Opening items and administrative issues
Item 6	Work programme for BPC 2016-2017
Item 7.1	Catalogue of specific conditions and elements to be taken into account at the product authorisation stage
Item 7.2	Procedure for renaming of an active substance according to Article 13 of Regulation (EU) No 1062/2014
Item 7.3	Outcome of the e-consultation Ad-hoc Working Group on Environmental Exposure
Item 7.4	Draft BPC opinion on dichlofluanid for PT 21

Tuesday 11 October: afternoon session

Item 7.5	Draft BPC opinion on silicium dioxide for PT 18
Item 7.6	Draft BPC opinion on silicon dioxide (as a nanomaterial formed by aggregates and agglomerates) for PT 18

Wednesday 12 October: morning session

Item 7.7	Draft BPC opinion on PHMB for PT 5
Item 7.8	Working procedure for Union Authorisation
Item 7.9	Additional data submitted after active substance approval for:
	a) 1R-trans-phenothrin
	b) copper, granulated
Item 8	AOB
Item 9	Agreement of the action points and conclusions

End of meeting

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