

**Revised draft agenda**  
**11<sup>th</sup> meeting of the Biocidal Products Committee (BPC)**  
**15-18 June 2015**  
**ECHA Conference Centre, Annankatu 18, Helsinki**  
**15 June: starts at 13:30**  
**18 June: ends at 13:00**

**Item 1 – Welcome and apologies**

**Item 2 – Agreement of the agenda**

BPC-A-11-2015\_rev2  
***For agreement***

**Item 3 – Declarations of potential conflicts of interest to agenda items**

**Item 4 – Agreement of the minutes and review of actions from BPC-10**

BPC-M-10-2015\_rev1  
***For agreement***

**Item 5 – Administrative issues**

**5.1 Housekeeping issues**

***For information***

**5.2 Other administrative issues and report from other Committees**

BPC-11-2015-01  
***For information***

**Item 6 – Work programme for BPC**

**6.1 Revised BPC Work Programme 2015-2016**

BPC-11-2015-02  
***For information***

**6.2 Outlook**

BPC-11-2015-03  
***For discussion***

## Item 7 – Applications for approval of active substances\*

### 7.1 Working procedure and templates: update from SECR

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

BPC-11-2015-04

**For information**

### 7.2 Draft BPC opinion on biphenyl-2-ol for PT 3, 4 and 6

*Previous discussion(s): WG-V-2014 and BPC-9*

**PT 3:** BPC-11-2015-05A, B and C

**PT 4:** BPC-11-2015-06A and B; BPC-11-2015-05C

**PT 6:** BPC-11-2015-07A and B; BPC-11-2015-05C

**For adoption**

### 7.3 Confirmation of the conclusions of the combined CAR for DDAC for PT 8

*Previous discussion(s): TM-III-2008, TM-IV-2008, TM-I-2009, TM-II-2009, TM-II-2013 and WG-II-2015*

BPC-11-2015-08A and B

**For agreement**

### 7.4 Confirmation of the conclusions of the combined CAR for ADBAC/BKC for PT 8

*Previous discussion(s): TM-III-2008, TM-IV-2008, TM-I-2009, TM-II-2009 and TM-II-2013*

BPC-11-2015-08A and BPC-11-2015-09

**For agreement**

### 7.5 Draft BPC opinion on PHMB for PT 1, 2, 3, 4, 6, 9 and 11

*Previous discussion(s): WG-III-2014 and WG-I-2015*

**PT 1:** BPC-11-2015-10A, B and C

**PT 2:** BPC-11-2015-11A and B; BPC-11-2015-10C

**PT 3:** BPC-11-2015-12A and B; BPC-11-2015-10C

**PT 4:** BPC-11-2015-13A and B; BPC-11-2015-10C

**PT 6:** BPC-11-2015-14A and B; BPC-11-2015-10C

**PT 9:** BPC-11-2015-15A and B; BPC-11-2015-10C

**PT 11:** BPC-11-2015-16A and B; BPC-11-2015-10C

**For adoption**

### 7.6 Draft BPC opinion on cybutryne for PT 21

*Previous discussion(s): TM-III-2011, TM-I-2012, BPC-8 and WG-II-2015*

BPC-11-2015-17A, A(2), B, C and D

**For adoption**

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\* 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 (may cover more than one PT) and a document containing open issues (covering all the PTs to be discussed for that substance).

**7.7 Draft BPC opinion on triclosan for PT 1**

*Previous discussion(s): WG-V-2014*

BPC-11-2015-18A, B and C

***For adoption***

**7.8 Draft BPC opinion on cyromazine for PT 18**

*Previous discussion(s): TM-II-2012, WG-I-2015*

BPC-11-2015-22A, B and C

***For adoption***

**7.9 Outcome of the written procedure for C(M)IT/MIT for PT 13**

BPC-11-2105-25

***For information***

**Item 8 – Any other business**

**8.1 Article 75(1)(g) request on sulfuryl fluoride**

BPC-11-2015-19

***For agreement***

**8.2 ARTFood ad-hoc Working Group guidance on “Estimating dietary risk from transfer of biocidal active substances into foods – non-professional uses”**

***For information***

**8.3 List of endpoints for imidacloprid**

BPC-11-2105-20

***For discussion***

**8.4 APCP Working Group guidance document on “‘Specification’, ‘Reference specification’, ‘Source’ and ‘Reference source’ - terminology used for processes under the Biocidal Products Regulation (BPR) (EU) No 528/2012”**

BPC-11-2015-21

***For information***

**8.5 Follow-up on the Workshop “Reviewing the active substance assessment process”**

BPC-11-2015-23

***For discussion***

**8.6 PAR template for Union Authorisation**

BPC-11-2015-24A and B

***For information***

**Item 9 – Agreement of the action points and conclusions**

***For agreement***

**Provisional timeline for the  
11<sup>th</sup> meeting of the Biocidal Products Committee (BPC)**

**ECHA Conference Centre, Annankatu 18, Helsinki**

**15 June 2015: starts at 13:30**

**18 June 2015: 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. Morning sessions usually start at 09:00.

**Monday 15 June: afternoon session**

Items 1-5	Opening items and administrative issues
Item 6	Work programme of the BPC 2015-16
Item 7	Applications for approval of active substances
Item 7.1	Working procedures and templates
Item 7.2	Draft BPC opinion on biphenyl-2-ol for PT 3, 4, 6
Item 7.3	Confirmation of the conclusions of the combined CAR for DDAC PT 8
Item 7.4	Confirmation of the conclusions of the combined CAR for ADBAC/BKC for PT8

**Tuesday 16 June: morning session**

Item 7	Follow up to previous discussions on draft substance opinions
Item 7.5	Draft BPC opinion on PHMB for PT 1, 2, 3, 4, 6, 9, 11

**Tuesday 16 June: afternoon session**

Item 7.5(cont'd)	Draft BPC opinion on PHMB for PT 1, 2, 3, 4, 6, 9, 11
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**Wednesday 17 June: morning session**

Item 7.	Follow up to previous discussions on draft substance opinions
Item 7.6	Draft BPC opinion on cybutryne for PT 21

**Wednesday 17 June: afternoon session**

Item 7.7	Draft BPC opinion on triclosan for PT 1
Item 7.8	Draft BPC opinion on cyromazine for PT 18
Item 7.9	Outcome of the written procedure for C(M)IT/MIT for PT 13

**Thursday 18 June: morning session**

Item 7	Follow up to previous discussions on draft substance opinions
Item 8	Any other business
Item 8.1	Article 75(1)(g) request on sulfuryl fluoride
Item 8.2	ARTFood ad-hoc Working Group guidance on "Estimating dietary risk from transfer of biocidal active substances into foods – non-professional uses"
Item 8.3	List of endpoints for imidacloprid
Item 8.4	APCP Working Group guidance document on "Specification', 'Reference specification', 'Source' and 'Reference source' - terminology used for processes under the Biocidal Products Regulation (BPR) (EU) No 528/2012"
Item 8.5	Follow-up on the Workshop "Reviewing the active substance assessment process"
Item 8.6	PAR template for Union Authorisation
Item 9	Agreement of the action points and conclusions