10 September 2013
RAC/A/26/2013

Final Agenda

26th meeting of the Committee for Risk Assessment

10-13 September 2013
ECHA Conference Centre (Annankatu 18, Helsinki)
10 September: starts at 9:00
13 September: ends at 13:00

Item 1 – Welcome and Apologies

Item 2 – Adoption of the Agenda

RAC/A/26/2013
For adoption

Item 3 – Declarations of conflicts of interest to the Agenda

Item 4 – Report from other ECHA bodies and activities

a) Report on RAC 25 action points, written procedures and other ECHA bodies

RAC/26/2013/01
For information

b) RAC workplan for all processes

For information

Item 5 – Harmonised classification and labelling (CLH)

5.1 CLH dossiers

a) Lead
b) Dodemorph
c) Dodemorph acetate
d) Phenol, dodecyl-, branched (tetrapropenylphenol (TPP))
e) Imidazole
f) Spirotetramat

g) Sulfoxaflor

h) 1,2-Epoxybutane

For discussion/adoption

i) Anti-coagulant rodenticides – general discussion

RAC/26/2013/02

For discussion/agreement

1. Chlorophacinone
2. Bromadiolone ketone
3. Difenacoum
4. Difethialone
5. Flocoumafen
6. Warfarin
7. Brodifacoum
8. Coumatetralyl

5.2 Appointment of RAC (co-)rapporteurs for CLH dossiers

RAC/26/2013/03 (room document)

For agreement

5.3 General and procedural CLH issues

For information

Item 6 – Restrictions

6.1 General restriction issues

For information

6.2 Restriction Annex XV dossiers

a) Lead in consumer articles – 2nd version of the draft opinion

For discussion

b) 1-Methylpyrrolidin-2-one (NMP) – outcome of the conformity check

For agreement

c) Nonyl phenol – outcome of the conformity check

For agreement

6.3 Appointment of (co-)rapporteurs for restriction dossiers

RAC/26/2013/05 (confidential room document)

For information

Item 7 – Authorisation

7.1 Authorisation application on phthalates – outcome of the conformity check and introductory presentation on the application

For agreement

7.2 Recommendation of the review period in applications for authorisation

RAC/26/2013/06

For agreement

7.3 Capacity building

a) DNEL setting (BBP)

RAC/26/2013/07

For discussion/agreement

b) ECHA project on carcinogenicity dose-response analysis of Cr(VI)- and As-containing substances

For information

7.4 Appointment of (co-) rapporteurs for authorisation applications

(*Closed session*)

RAC/26/2013/08 (confidential document)

For agreement

Item 9 – AOB

a) Report from the project on economic evaluation of environmental impacts

For information

Item 10 – Action points and main conclusions of RAC-26

Table with Conclusions and Action points from RAC-26

For adoption

PROVISIONAL TIMELINE FOR THE DISCUSSIONS AT RAC-26

Please note that this timeline is provisional. Changes can be made before and during the meeting in order to accommodate the discussions.

Tuesday, 10 September: Morning session

Item 1 – Welcome and Apologies

Item 2 – Adoption of the Agenda

Item 3 – Declarations of conflicts of interest to the Agenda

Item 4a – Report from other ECHA bodies and activities

Item 5.1 CLH dossiers

Tuesday, 10 September: Afternoon session

Item 5.1 CLH dossiers

Item 6.2 a - Lead in consumer articles

Evening: possible ad hoc working groups

Wednesday, 11 September: Morning session

Item 6.2 a - Lead in consumer articles

Item 6.2 b - NMP - outcome of conformity check

Item 6.2 c - Nonyl phenol – outcome of the conformity

Item 6.3 Appointment of (co-)rapporteurs for restriction dossiers

Item 5.1 CLH dossiers

Wednesday, 11 September: Afternoon session

Item 5.1 CLH dossiers

Item 5.2 - Appointment of (co-)rapporteurs for CLH dossiers

Item 5.3 General and procedural CLH issues

Evening: Formal Dinner

Thursday, 12 September: Morning session

Joint RAC/SEAC session:

Item 6.1 - General restriction issues;

Item 7.2 Recommendation of the review period in applications for authorisation

Item 7.1 Authorisation application on phthalates (introductory presentation)

Item 9 Report from the project on economic evaluation of environmental impacts

RAC session

Item 7.1 Authorisation application on phthalates (conformity check)

Item 7.3 Capacity building – Application for Authorisation

Item 7.4 Appointment of (co-)rapporteurs for authorisation applications (closed session)

Thursday, 12 September: Afternoon session

Item 5.1 CLH dossiers

Evening session for RAC and SEAC (co-) rapporteurs for the authorisation applications

Friday, 13 September: Morning session

Item 5.1 CLH dossiers

Item 9 – AOB

Item 10 – Action points and main conclusions of RAC-26