11 September 2012
RAC/A/22/2012

Final Agenda

22nd meeting of the Committee for Risk Assessment

11-14 September 2012
ECHA Conference Centre (Annankatu 18, Helsinki)
11 September: starts at 9:00
14 September: ends at 13:00

Item 1 – Welcome and Apologies

Item 2 – Adoption of the Agenda

RAC/A/22/2012
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 21 action points, written procedures and other ECHA bodies

RAC/22/2012/01
For information

b) Implementation of Conflict of Interest Policy
- General principles and guidance for Committee members

RAC/22/2012/02
For discussion

- Eligibility criteria

For information

Item 5 – Harmonised classification and labelling (CLH)

5.1 CLH dossiers

a) Fenoxycarb
b) Tralkoxydim

c) 4-Vinylcyclohexene (VCH)
d) Cymoxanil
e) 3-Iodo-2-propynylbutylcarbamate (IPBC)
f) Formaldehyde
g) Methyl-2,5-dichlorobenzoate
h) Tetrahydrofurfuryl alcohol (THFA)
i) Cycloxydim

For discussion/adoption

5.2 Requests under Article 77(3) (c) - CLH dossiers

a) Gallium arsenide
b) Epoxiconazole

For discussion/adoption

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

RAC/22/2012/03
For agreement

5.4 General and procedural CLH issues

a) State of play of CLH dossiers

b) Opinion development process

For information

Item 6 – Restrictions

6.1 General restriction issues

a) Update on intended restriction dossiers

b) Update on the review of restriction process

RAC/22/2012/04 ***(room document)***
For information

6.2 Restriction Annex XV dossiers

a) Chromium VI – 2nd version of the draft opinion

For discussion

b) Dichlorobenzene – 1st version of the draft opinion

For discussion

c) Nonylphenol – outcome of the conformity check

For agreement

6.3 Requests under Article 77(3)(c) - restriction dossiers

a) Non-classified phthalates (DINP and DIDP)

For discussion/adoption

6.4 Appointment of (co-)rapporteurs for restriction dossiers

RAC/22/2012/05 *(room document)*

For information

Item 7 – Authorisation

a) Capacity building
   - Establishing DNELs and dose-response functions

     RAC/22/2012/06 (*room document*)

     For discussion
   - Valuation of environmental impacts of PBTs

     For discussion
   - Proportionality in evaluating Applications for Authorisation (AfAs)

     For discussion
   - AfAs with 'multiple dimensions'

     RAC/22/2012/07 *(room document)*

     For discussion

b) Participation of case-owners and stakeholder observers in opinion development process

For discussion

Item 8 – AOB

a) C&L Inventory
b) Feedback on the first four restrictions from the Commission´s Impact Assessment point of view

For information

Item 9 – Action points and main conclusions of RAC-22

Table with Conclusions and Action points from RAC-22

For adoption