2 December 2013
RAC/A/27/2013

Final Agenda

27th meeting of the Committee for Risk Assessment

2-5 December 2013
ECHA Conference Centre (Annankatu 18, Helsinki)
2 December: starts at 9:00
5 December: ends at 16:00

Item 1 – Welcome and Apologies

Item 2 – Adoption of the Agenda

RAC/A/27/2013
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 26 action points, written procedures and other ECHA bodies

RAC/27/2013/01
For information

b) RAC workplan for all processes

For information

Item 5 – Harmonised classification and labelling (CLH)

5.1 CLH dossiers

a) Sulfoxaflor
b) Phenol, dodecyl-, branched (TPP)
c) Lead
d) Anti-coagulant rodenticides
    1. Flocoumafen
    2. Warfarin
    3. Brodifacoum

e) Triflusulfuron methyl

f) Bifenazate

g) Fenpyroximate

h) Lenacil

i) Tributyltin compounds

For discussion/adoption

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

RAC/27/2013/02 (confidential room document)

For agreement

5.3 General and procedural CLH issues

a) State of play of CLH dossiers

RAC/27/2013/03

For information

Item 6 – Restrictions

6.1 General restriction issues

a) Update on intended restriction dossiers

For information

b) Revision of the restriction process

RAC/27/2013/04

For discussion/agreement

RAC/27/2013/09

For information

6.2 Restriction Annex XV dossiers

a) Lead in consumer articles – 4th version of the draft opinion

For adoption

b) Nonyl phenol – 1st version of the draft opinion

For discussion

c) 1-Methyl-2-pyrrolidone (NMP) – 1st version of the draft opinion

For discussion

d) Cadmium in paints- outcome of conformity check

For agreement

6.3 Appointment of (co-)rapporteurs for restriction dossiers

RAC/27/2013/05 (confidential room document)

For information/agreement

Item 7 – Authorisation

7.1 Authorisation applications

a) Authorisation application on the use of DEHP in a stop-off formulation in manufacturing of aero engines – first version of the draft opinion

For discussion/agreement

b) Authorisation applications on Phthalates (submitted within the August submission window) - outcome of the conformity check

For agreement

7.2 Capacity building

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

RAC/27/2013/06

RAC/27/2013/07

For discussion/agreement

b) ECHA project on carcinogenicity dose response analysis of Trichloroethylene

For information

7.3 Appointment of (co-) rapporteurs for authorisation applications

RAC/27/2013/08 (confidential room document)

For agreement

Item 8 – AOB

- Update on Guidance activities

Item 9 – Action points and main conclusions of RAC-27

- Table with Conclusions and Action points from RAC-27

For adoption