04/06/2013
RAC/A/25/2013

Final Agenda

25th meeting of the Committee for Risk Assessment

4-7 June 2013
ECHA Conference Centre (Annankatu 18, Helsinki)
4 June: starts at 9:00
7 June: ends at 13:00

Item 1 – Welcome and Apologies

Item 2 – Adoption of the Agenda

RAC/A/25/2013
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 24 action points, written procedures and other ECHA bodies

RAC/25/2013/01
For information

b) RAC work plan for all processes

For information

Item 5 – Harmonised classification and labelling (CLH)

5.1 Sensitisation criteria following the 2nd ATP to the CLP Regulation

For discussion and agreement

5.2 CLH dossiers

a) Etridiazole
b) Metosulam
c) Organic acids
    a. Octanoid acid,

b. Nonanoic acid,
c. Decanoic acid

d) Diisohexylphthalate (DIHP)
e) Imazalil
f) Tebuconazole
g) Spirotetramat
h) Dimethenamid-P
i) Carvone
j) Tembotrione
k) Flonicamid

For discussion/adoption

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

RAC/25/2013/02 (confidential room document)
For agreement

5.4 General and procedural CLH issues

RAC/25/2013/05
For information/discussion

Item 6 – Restrictions

6.1 General restriction issues

For information

6.2 Restriction Annex XV dossiers

a) Lead in consumer articles – 1st version of RAC draft opinion
For discussion

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

6.3 Appointment of (co-)rapporteurs for restriction dossiers

RAC/25/2013/03(confidential)
For information/agreement

Item 7 – Authorisation

7.1 Capacity building

For information

7.2 Recommendation of the review period in applications for authorisation

RAC/25/2013/08
For discussion/agreement

7.3 Revised working procedure for appointment of (co-)rapporteurs for authorisation applications

RAC/25/2013/04
For discussion/agreement

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

RAC/25/2013/06 (confidential room document)
For agreement

Item 8 – RAC Manual of Conclusions and recommendations

RAC/25/2013/07
For information/agreement

Item 9 – AOB

a) Stockholm Convention decision on HBCDD - relevance for Annex XIV

Item 10 – Action points and main conclusions of RAC-24

Table with Conclusions and Action points from RAC-24

For adoption