25 May 2010
RAC/A/11/2010

Final Agenda

Eleventh meeting of the Committee for Risk Assessment

25 – 27 May 2010

Helsinki, Finland

25 May: starts at 9:00
27 May: ends at 12.30

Item 1 – Welcome & Apologies

Item 2 – Adoption of the Agenda

RAC/A/11/2010
For adoption

Item 3 – Declarations of conflicts of interest to the Agenda

Item 4 – Adoption of the draft minutes of RAC-10

- Adoption of the draft minutes

RAC/M/10/2010 draft final
For adoption

Item 5 – Administrative issues and information items

a. Status report on the RAC - 10 action points
b. Outcome of written procedures
c. Report from other ECHA bodies and activities

RAC/11/2010/32
ROOM DOCUMENT
For information

d. Follow-up of the 1st written procedure for adoption of a RAC opinion

RAC/11/2010/33
ROOM DOCUMENT

For agreement

Item 6 – CLH

6.1 CLH Dossiers

a. Gallium arsenide

For discussion and possible adoption

b. Tetrahydrofuran

For discussion and possible adoption

c. TDCP

For discussion and possible adoption

d. Cryolites (CAS13775-53-6 and CAS15096-52-3)

For discussion and possible adoption

e. HBCDD

For discussion and possible adoption

f. Fuberidazole

For initial discussion

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

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

RAC/11/2010/34
ROOM DOCUMENT
For agreement

6.3 General CLH issues

a. State of play of the submitted CLH dossiers

RAC/11/2010/35
ROOM DOCUMENT
For information

b. Revision of the working procedures on accordance check and on processing of dossiers for harmonised classification and labeling

RAC/11/2010/24
RAC/11/2010/25
For agreement

c. Feedback from the Commission on adopted CLH opinions and follow-up actions

RAC/11/2010/36
ROOM DOCUMENT
For agreement

Item 7 – Restrictions

7.1 Restriction Annex XV dossiers

a. DMF – conformity check

For decision

b. Lead and its compounds in jewellery – conformity check

For decision

7.2 General restriction issues

- Update on intended restriction dossiers

For information

Item 8 – Authorisation

a. Discussion note on the content of final Commission decisions and their effect on the format of the RAC and SEAC opinions

RAC/11/2010/26
For discussion

b. Elements of RAC and SEAC working procedure for developing opinions on the applications for authorisation

RAC/11/2010/27
For discussion

c. Discussion paper on the scope and content of conformity check of authorisation applications

RAC/11/2010/28
For discussion

d. WP on conformity check of authorisation applications

RAC/11/2010/29
For discussion and possible agreement

e. Terms of reference for authorisation (co-) rapporteurs

RAC/11/2010/30
For discussion

Item 9 – Guidance issues

a. Feedback from guidance consultations
b. Report on other guidance activities

For information

Item 10 – Any other business

a. RAC meeting calendar for 2011

RAC/11/2010/31
For information

b. Renewal of RAC membership

For information

c. Workshop on evaluation of two organic siloxane compounds

For information

d. Presentation on Extended One Generation Reproductive Toxicity Studies (EOGRTS)

For information

Item 11 – Main conclusions and Action Points of RAC-11

- Table with main conclusions and action points from RAC- 11

For adoption

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