6 September 2010
RAC/A/12/2010

Final Agenda

Twelfth meeting of the Committee for Risk Assessment

7 – 9 September 2010

Helsinki, Finland

7 September: starts at 9:00
9 September: ends at 16:00

Preceded by a Presentation to RAC of the results of the EU Project PHIME to be held on 6 September from 15:30 to 18:30

Item 1 – Welcome & Apologies

Item 2 – Adoption of the Agenda

RAC/A/12/2010
For adoption

Item 3 – Declarations of conflicts of interest to the Agenda

Item 4 – Adoption of the draft minutes of RAC-11

- Adoption of the draft minutes

RAC/M/11/2010 draft final
For adoption

Item 5 – Administrative issues and information items

a. Status report on the RAC - 11 action points
b. Outcome of written procedures
c. Report from other ECHA bodies and activities

RAC/12/2010/45
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For information

Item 6 – MSCA support to RAC and Renewal of RAC Membership

a. Update on the letters sent to MSCA and on the preparations for renewal of RAC Membership

For information

b. Role of (co-)rapporteurs if their RAC Membership is not renewed

RAC/12/2010/37

For discussion and possible agreement

Item 7 – CLH

7.1 CLH Dossiers

a. TDCP (adopted by written procedure before RAC-12)

For information

b. HBCDD

For adoption

c. Fuberidazole

For adoption

d. White spirit dossiers

For first discussion and possible adoption

e. Acequinocyl

For first discussion and possible adoption

f. TNPP

For first discussion and possible adoption

g. Bifenthrin

For first discussion

7.2 Appointment of RAC (co-) rapporteurs for CLH dossiers (if relevant)

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

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For decision

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For information

7.3 General CLH issues

a. State of play of the submitted CLH dossiers

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For information

b. Report from the discussions at the ad hoc meeting held after RAC-11 on criteria for assessing the reliability and relevance of the studies which support the RAC opinions

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For information

c. ECHA-EFSA cooperation on the classification and labelling of active substances in Plant Protection Products.

For information

Item 8 – Restrictions

8.1 Restriction Annex XV dossiers

a. DMFu – first draft opinion

For first discussion

b. Lead and its compounds in jewellery – first draft opinion

For first discussion

c. Phenylmercury compounds – conformity check

For decision

d. Mercury in measuring devices – conformity check

For decision

8.2 Appointment of RAC (co-) rapporteurs for restriction dossiers (if relevant)

For agreement

8.3 General restriction issues

- Update on intended restriction dossiers

For information

Item 9 – Authorisation

9.1 Content of an authorisation application

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For information

9.2 Conformity check

a. Scope and content of conformity check

RAC/12/2010/38
(Response to comments table) **RAC/12/2010/39**
For discussion

b. Second discussion on the working procedure for conformity check of authorisation applications

RAC/12/2010/40
(Response to comments table) **RAC/12/2010/41**
For discussion

9.3 Working procedure for developing opinions for authorisation applications

RAC/12/2010/42
(Response to comments table) **RAC/12/2010/43**
(Response to comments table) **RAC/12/2010/44**
For first discussion

9.4 Questions on alternatives

For discussion

Item 10 – Guidance issues

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

For information

Item 11 – Any other business

a. Workshop on non-testing methods

For information

b. Revision of the RAC Meeting calendar for 2011

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For information

c. 2nd International Conference on Risk Assessment

For information

d. Initial considerations on the use of the results of the Extended One Generation Reproductive Toxicity Studies (EOGRTS) in C&L and risk assessment processes

For information

Item 12 – Main conclusions and Action Points of RAC-12

- Table with main conclusions and action points from RAC- 12

For adoption

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