

Final Agenda
29th meeting of the Committee for Risk Assessment

2-6 June 2014

ECHA Conference Centre (Annankatu 18, Helsinki)

2 June: starts at 9:00
6 June: ends at 13:00

Item 1 – Welcome and Apologies

Item 2 – Adoption of the Agenda

RAC/A/29/2014
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 28 action points, written procedures and other ECHA bodies

RAC/29/2014/01
RAC/29/2014/02 (room document)
For information

- b) RAC workplan for all processes

For information

Item 5 – Harmonised classification and labelling (CLH)

5.1 CLH dossiers

- a) Bupirimate (ISO) (remaining health hazards)

- b) 1-methyl-2-pyrrolidone (NMP)
- c) Propylene oxide
- d) Glutaraldehyde
- e) Tinuvin 123
- f) Flumioxazin (ISO)
- g) 1,2-dichloropropane (PDC)

For discussion/adoption

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

RAC/29/2014/03 (restricted room document)

For agreement

5.3 General and procedural CLH issues

- a) New procedures for agreement seeking

RAC/29/2014/04

For information/agreement

Item 6 – Restrictions

6.1 General restriction issues

- a) Review of the restriction process:
Update from Task Force

For information

6.2 Restriction Annex XV dossiers

- a) Opinion development

- 1) Nonylphenol – 4th version of the draft opinion

For adoption

- 2) 1-Methyl-2-pyrrolidone (NMP) – 4th version of the draft opinion

For adoption

- 3) Cadmium and its compounds in paints – 2nd version of the draft opinion

For discussion

- 4) Cadmium and its compounds in artist paints – first plenary discussions on the key issues document

For discussion

- 5) Chrysotile - first plenary discussions on the key issues document

For discussion

b) Conformity check

- 1) Isopropylidenediphenol (Bisphenol A) – outcome of conformity check

For agreement

- 2) Ammonium salts - outcome of the conformity check

For agreement

6.3 Appointment of (co-)rapporteurs for restriction dossiers

RAC/29/2014/05 (restricted document)

For agreement

Item 7 – Authorisation

7.1 Authorisation applications

- a) Authorisation application on phthalates – 2nd version of the RAC draft opinions (applications submitted within the August 2013 submission window)

1. Two uses of DEHP submitted by ARKEMA FRANCE (DEHP 2a):

- i. Formulation of DEHP in compounds, dry-blends and Plastisol formulations
- ii. Industrial use in polymer processing by calendering, spread coating, extrusion, injection moulding to produce PVC articles

For discussion/agreement

2. Two uses of DEHP submitted by Grupa Azoty Zakłady Azotowe Kędzierzyn Spółka Akcyjna (DEHP 2b):

- i. Formulation of DEHP in compounds, dry-blends and Plastisol formulations
- ii. Industrial use in polymer processing by calendering, spread coating, extrusion, injection moulding to produce PVC articles

For discussion/agreement

3. Three uses of DEHP submitted by DEZA a.s. (DEHP 2c):

- i. Formulation of DEHP in compounds, dry-blends and Plastisol formulations
- ii. Industrial use in polymer processing by calendering, spread coating, extrusion, injection moulding to produce PVC articles

- iii. Use in ceramic sheets and printing pastes for production of capacitors and lambda sensor elements

For discussion/agreement

4. The third use of DBP and DEHP submitted by Roxel (UK Rocket Motors) Ltd (DEHP 3):

- i. Industrial use of DBP within a specialty paint in manufacture of motors for rockets and tactical missiles

For discussion/agreement

5. The second and the third uses of DBP submitted by DEZA a.s. (DBP 2):

- i. Use in propellants
- ii. Use in ceramic sheets and printing pastes for production of capacitors and lambda sensor elements

For discussion/agreement

6. Two uses of DEHP submitted by VINYLOOP FERRARA S.p.A., Stena Recycling AB and Plastic Planet srl (DEHP 4):

- i. Formulation of recycled soft PVC containing DEHP in compounds and dryblends
- ii. Industrial use of recycled soft PVC containing DEHP in polymer processing by calendaring, extrusion, compression and injection moulding to produce PVC articles

For discussion/agreement

- b) Authorisation application – 1st outline RAC draft opinions (applications submitted within the November 2013 submission window)

1. The use of diarsenic trioxide submitted by Boliden Kokkola Oy (Diarsenic trioxide 1):

- i. Use of diarsenic trioxide in the purification of metal impurities from the leaching solution in the zinc electrowinning process

For discussion

2. The use of diarsenic trioxide submitted by Nordenhamer Zinkhütte GmbH (Diarsenic trioxide 2):

- i. Industrial use of diarsenic trioxide to produce a copper concentrate in the purification of the leaching solution in a zinc electrowinning process

For discussion

3. Two uses of diarsenic trioxide submitted by Linxens France (Diarsenic trioxide 3):

- i. Formulation of diarsenic trioxide into a mixture
- ii. Industrial use of diarsenic trioxide as processing aid in gold electroplating

For discussion/agreement

4. Six uses of lead sulfochromate yellow (C.I. pigment yellow 34) and lead chromate molybdate sulphate red (C.I. pigment red 104) submitted by DCC Maastricht B. V. OR (Lead chromate pigments 2):
 - i. Distribution and mixing pigment powder in an industrial environment into solvent-based paints for non-consumer use
 - ii. Industrial application of paints on metal surfaces (such as machines vehicles, structures, signs, road furniture, coil coating etc.)
 - iii. Professional, non-consumer application of paints on metal surfaces (such as machines, vehicles, structures, signs, road furniture etc.) or as road marking
 - iv. Distribution and mixing pigment powder in an industrial environment into liquid or solid premix to colour plastic/plasticised articles for non-consumer use
 - v. Industrial use of solid or liquid colour premixes and pre-compounds containing pigment to colour plastic or plasticised articles for non-consumer use
 - vi. Professional use of solid or liquid colour premixes and pre-compounds containing pigment in the application of hotmelt road marking

For discussion

- c) Authorisation application - outcome of the conformity check
 1. HBCDD 1

For agreement

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

RAC/29/2014/06 (restricted room document)

For agreement

Item 8 – AOB

Item 9 – Action points and main conclusions of RAC-29

Table with Conclusions and Action points from RAC-29

For adoption

PROVISIONAL TIMELINE FOR THE DISCUSSIONS AT RAC-29

Please note that this timeline is provisional. Changes can be made before and during the meeting in order to accommodate the discussions.

Monday 2 June: Morning session

- Item 1 - Welcome and Apologies
- Item 2 - Adoption of the Agenda
- Item 3 - Declarations of conflicts of interest to the Agenda
- Item 4 - Report from other ECHA bodies and activities
- Item 7 - Session on common approach to the applications for authorisation
- Item 7.1 - Authorisation applications

Monday 2 June: Afternoon session

- Item 7.1 - Authorisation applications

Evening: possible ad hoc working groups

Tuesday 3 June: Morning session

- Item 7.1 - Authorisation applications

Tuesday 3 June: Afternoon session

- Item 7.1 - Authorisation applications
- Item 6.2.a - Restriction - opinion development

Evening: possible ad hoc working groups

Wednesday 4 June: Morning session

- Item 3 - Declarations of conflicts of interest to the Agenda (closed session)
- Item 6.2.b - Restriction - conformity check
- Item 5.1 - CLH dossiers
- Item 6.2.a - Restriction - opinion development

Wednesday 4 June: Afternoon session

- Item 6.2.a - Restriction - opinion development
- Item 6.2.b - Restriction - conformity check
- Item 6.1. - Review of the restriction process: update from Task Force

Evening: Informal Dinner

Thursday 5 June: Morning session

- Item 5.3 - General and procedural CLH issues
- Item 5.1 - CLH dossiers
- Item 6.2.a - Restriction - opinion development

Thursday 5 June: Afternoon session

- Item 5.1 - CLH dossiers

Evening: Session for (co-) rapporteurs AfA

Friday 6 June: Morning session

- Item 5.1 - CLH dossiers
- Item 7.2/5.2/6.3 Appointment of (co-) rapporteurs: Authorisation (closed session), CLH and Restriction
- Item 8 - AOB

END OF THE MEETING