Provisional Draft Agenda
Member State Committee & Committee for Risk Assessment
Joint Workshop

Fine tuning the testing requirements and evaluation of selected human health endpoints under REACH and CLP

11-12 October 2018
ECHA Conference Centre
Annankatu 18, in Helsinki, Finland

11 October: starts at 1 pm
12 October: ends at 1 pm

Item 1 – Welcome

Item 2 – Practicalities

Item 3 – Introduction of the Member State Committee and the Risk Assessment Committee

Presentation of the two committees' functions, regulatory processes and main areas of work, with special focus on the possible circumstances of further interaction.

Item 4 – Examples of interaction

Short introductory presentations will be given on each of the topics below by members of MSC and RAC:

- **Mutagenicity**
  Discussion on the ability of newer test methods (e.g. comet assay; genotoxicity test *in vivo*) to produce results that are suitable for classification, labelling and risk assessment.

- **Dose selection in systemic toxicology tests**
  Discussion on possible reasons and suitability of the results of low dose toxicity testing for classification, labelling and risk assessment.

- ***In vitro* testing for skin sensitisation**
  Discussion on the future role and interpretation of *in vitro* skin sensitization testing.

These will be followed by discussion of key questions in break-out groups.

Item 5 – Conclusions

- Conclusions with a short discussion on the developments in OECD and on their impact on the work of the two committees.
- Proposals for future cooperation between the two committees.