

Dear all:

Please find attached a summary of the pre-conference virtual meetings of Working Group 2 (AOPs) and 3 (standards). If you have addition points for the Dublin meeting, we are happy to include them.

Best.

Thomas

Short presentations:

- Overview of AOP-KB and its modules (Clemens)
- Overview of OpenTox standards and deliverables (Thomas?)
- Data format of AOP-KB Hub (i.e. the upcoming data interchange interface with AOP-KB – to be handled by Hristo Aladjov, who will be present in the session)
- Intermediate effects DB (IEDB) module, data format "OECD Harmonised Templates": Making science findings acceptable in the regulatory domain (Clemens)

Target outcome of discussion (in parentheses the WG(s) that contribute mostly or profit mostly from the outcome)

- Agreement on possible collaboration/synergy between Opentox and AOP-(KB) community, esp. consolidation of available formats and definition of recommendations (both WGs)
- Agreement on joint effort to create a closed vocabulary/ontology in the AOP domain, with a pilot exercise for the "Key Event" naming (mostly WG2)
- Regulatory acceptance of AOPs: How can our collaboration help? (mostly WG2)
- AOP -> consists of several Key Events -> each of them detectable/measurable with one or more test methods: How can our collaboration lead to a homogenous description of methods? (WG2 + WG3)
- More general: How can we define standards for data and metadata exchange (WG2 and WG3)
- Communication strategy with experts to understand the specific requirements of each experimental technique (WG3)
- Possible ways of harmonization with vendors of experimental equipment (WG3)
- Agreement on how to harmonize with other biological and biomedical domains for standards for data and metadata (WG3)
- Linking data and metadata to AOPs produced from them (standard format) (WG2 and WG3)
- Guidance on how to access AOP-KB data for further analysis and predictions in third party systems (WG2 and WG3)