How to bring your registration dossier in compliance with REACH – Tips and Hints (Part 3)

Conclusion

14th May 2013

Rupert Simon
European Chemicals Agency

Questions during the webinar

- **Submit your question**
  - via the Q&A panel (until 13:00h Helsinki time, UTC +2)
- **Then**
  - monitor the Q&A panel for our response
  - remain logged-in to the Webinar
  - we answer until 14:00h (one hour after the last presentation)

Questions after the webinar

- (Re)submit your question via the Helpdesk form at

  http://echa.europa.eu/en/web/guest/contact

  - if by 14:00h no answer is provided to your question;
  - if you have a question that requires a complex answer involving several experts

- When you use the ECHA contact form,
  - you will receive an acknowledgement of receipt and
  - we will answer your question as soon as possible

Use the correct format in IUCLID

- Report any information as **e**ndpoint **s**tudy **r**ecord (ESR)
- Report adaptation arguments to the testing regime separate from experimental or *in silico* results (i.e. different ESR)
- Report underpinning data in a proper ESR (e.g. hydrolytical instability under hydrolysis)
- Each argument for adaptation to the standard testing regime needs to be reported in an ESR separately

Improve your dossier quality now!

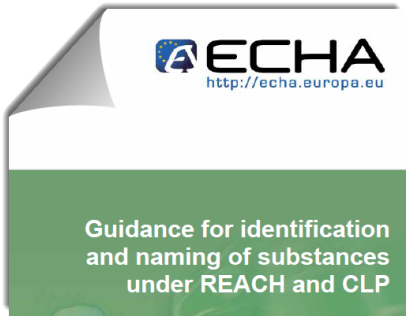
- Avoid common pitfalls
- Keep yourself up-to-date:
  - Evaluation Progress Reports
  - Follow our events
- Keep your dossiers up-to-date
  - Do not wait for an ECHA decision

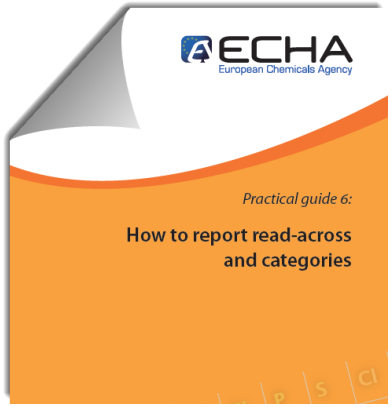
Tool to support improving dossier quality:

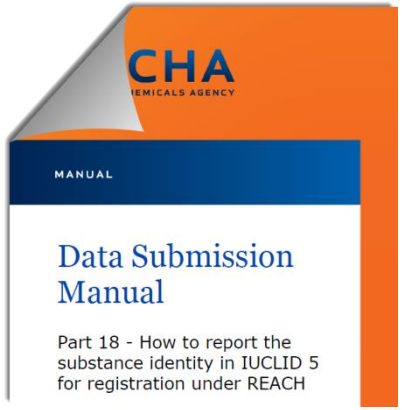
Plugin:

http://echa.europa.eu/view-article/-/journal_content/title/new-tool-to-support-registrants-in-improving-dossier-quality-now-available

Information on the ECHA website







Information material

- Evaluation progress reports

http://echa.europa.eu/regulations/reach/evaluation

- Targeted Compliance check

http://www.echa.europa.eu/view-article/-/journal_content/1a87ce8e-6286-4d1b-9dc2-b2d10d6f1d79

- Guidance document on Endpoint specific guidance (R.7)

http://echa.europa.eu/guidance-documents/guidance-on-information-requirements-and-chemical-safety-assessment

- Guidance document on QSARs and grouping (R.6)

http://echa.europa.eu/documents/10162/13632/information_requirements_r6_en.pdf

- Practical guides 1, 2, 3, 4, 5 and 6

http://echa.europa.eu/practical-guides

Upcoming events: Webinars

For all registrants

- **Part 4 – How to bring your registration dossier in compliance with REACH – Tips and Hints**
  11 September 2013

- **Part 5 – How to bring your registration dossier in compliance with REACH – Tips and Hints**
  27 November 2013

http://echa.europa.eu/support/training-material/webinars

Post event survey

- Once the event has ended, you will be directed to a post-event survey page.
- Your feedback is important to us.
- Your feedback helps us make the content of future webinars more relevant for your individual needs.
- Please take the time to fill out the survey.

Thank you