

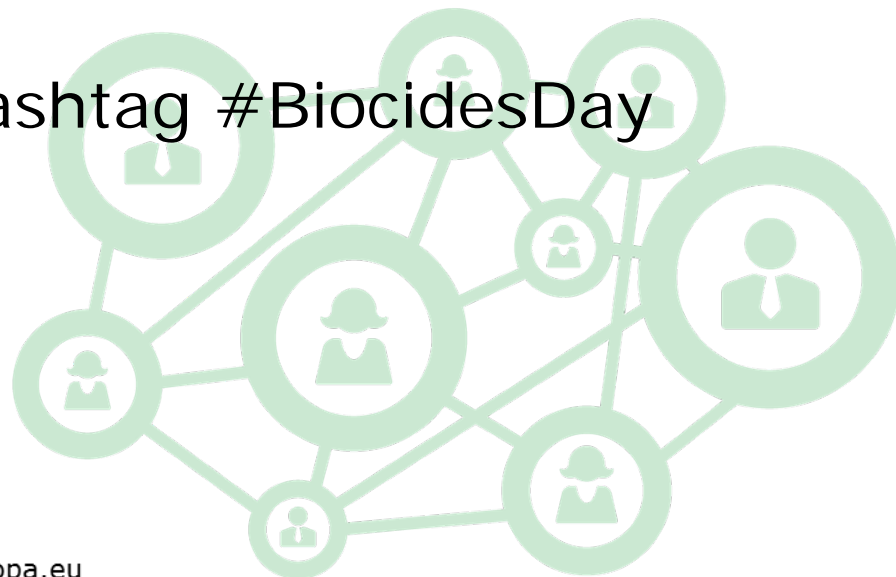
Closing remarks

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ECHA



Thank you for sharing with us

- **140** here in person from **30** countries
- **260** online
- **35** one-to-one sessions – use the networking session to ask more
- **15 000** reach for our hashtag #BiocidesDay
- **70** questions online



Today's material online

- Presentations there - video recording tomorrow
- Give us feedback today via Slido (BiocidesDay)
- For unanswered questions, contact our helpdesk

Tomorrow

- IT tool training – welcome back
- IUCLID, R4BP 3 and SPC Editor
- No live stream but main parts will be recorded and made available after the event

Complexities

- Legislation and implementing procedures
- IT tools and their further development
- Union authorisation – biocidal product family
- Working in consortia
- *In situ* generated substances
- Treated articles vs biocidal products



Challenges you raised

- Timelines for evaluations – how to speed up
- Endocrine disruptor criteria – how to adapt
- Risk assessment under biocides and REACH – how to avoid unnecessary overlap
- Who advises on what – Member State vs ECHA
 - Plea for open and clear communication between applicant and evaluating authority
- How to find consortia and deal with letters of access
- Level of the fees

Keep an eye out for these

- Public consultation for endocrine disruptor guidance
- Deadlines to apply for Union authorisation
- Brexit – check our website for Q&As



Note down in your calendar

**REACH 2018
conference**

30-31 January 2018

**Biocides 5 years
conference**

October 2018



Safe journey home

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