

Biocides Stakeholders' Day

26-27 SEPTEMBER 2017

HELSINKI, FINLAND



Tuesday 26 September

Plenary session 1

Biocides regulatory developments

Chair: Jack de Bruijn, Director of Risk Management, ECHA

08.00 REGISTRATION

09.00 OPENING
GEERT DANCET
Executive Director
ECHA

09.05 FUTURE OUTLOOK ON BIOCIDES
MARTINUS NAGTZAAM
European Commission

09.25 UPCOMING DEVELOPMENTS
HUGUES KENIGSWALD
ECHA

09.45 ENDOCRINE DISRUPTORS – WHERE ARE WE
SIMON GUTIERREZ ALONSO
ECHA

10.00 QUESTIONS AND ANSWERS

10.45 BREAK



Tuesday 26 September

Plenary session 2

Building a biocides application

Chair: Christel MUSSET, Director of Registration, ECHA


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- 11.15 IT TOOLS AND SUPPORT
VALERIO SPINOSI
ECHA
- 11.30 WORKING IN A CONSORTIUM
AN GHEKIERE
ARCHE Consulting
- 11.50 UNION AUTHORISATION FOR A PRODUCT FAMILY
CAROLINE HALL
Evonik Nutrition and Care
- 12.10 FREE RADICALS AND OTHER *IN SITU* GENERATED
SUBSTANCES: EXPERIENCES & CHALLENGES
BRIGITTE VAN NOORLOOS
Board for the Authorisation of Plant Protection Products and Biocides
the Netherlands
- 12.35 QUESTIONS & ANSWERS
- 13.10 BREAK

Tuesday 26 September

Plenary session 3

Tips from authorities

Chair: Jukka MALM, Deputy Executive Director, ECHA

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- 14.30 HOW TO DO A SUCCESSFUL UNION AUTHORISATION APPLICATION
CHIARA PECORINI
ECHA
- 14.50 TREATED ARTICLES – MEMBER STATE ADVICE
ULRIKE FRANK
Swedish Chemicals Agency, KEMI
- 15.10 WHAT TO EXPECT FROM ENFORCEMENT
FRANCESCA RAVAIOLI
Ministry of Health
General Directorate on medical devices and pharmaceutical service
Italy
- 15.30 QUESTIONS & ANSWERS
- 16.15 CLOSING REMARKS
JACK DE BRUIJN
Director of Risk Management
ECHA

14.30 - 18.30

ONE-TO-ONE SESSIONS WITH ECHA EXPERTS

16.30 - 18.30

NETWORKING

Wednesday 27 September

Training

Training on IT tools for biocides applications

9.00-10.30

IUCLID

11.00-14.00

R4BP 3

Summary of Product Characteristics Editor



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