



Part 1: BPR overviews

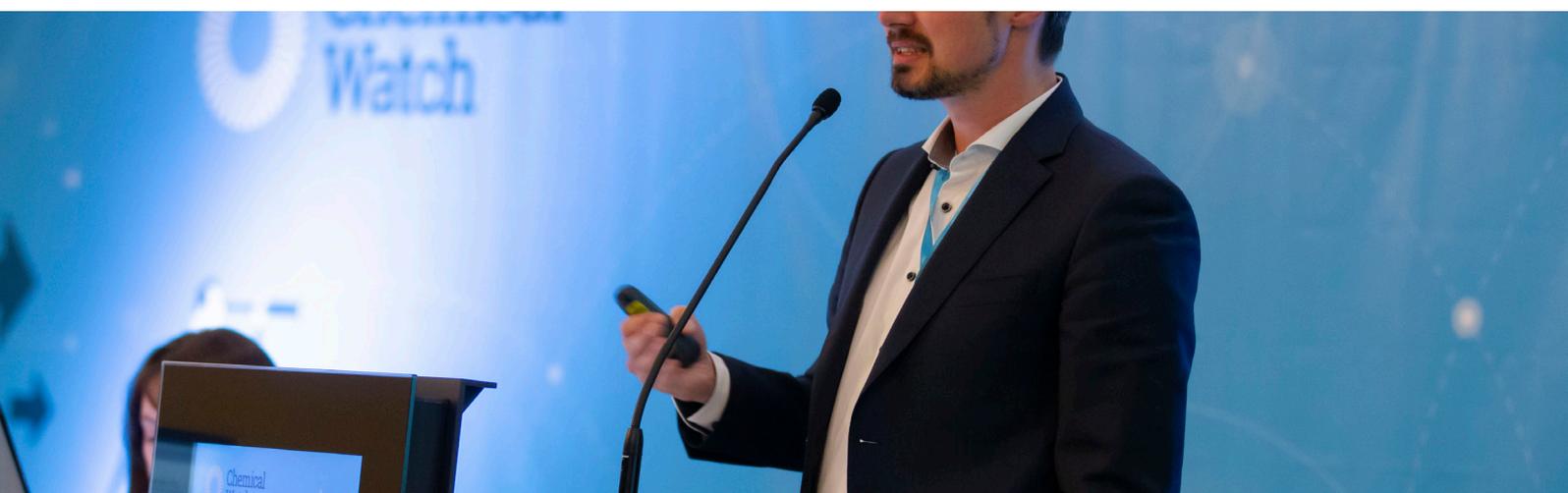
- 11.00-11.15 **Welcome and introduction**
David Dillon, Senior Regulatory and Compliance Analyst, Chemical Watch
- 11.15 -11.35 **EU Commission overview**
Marta Cainzos Garcia, DG Sante, EU Commission, Belgium
- 11.40 -12.00 **ECHA: Current priorities and work of the Environment Working Group**
- ECHAs involvement in COVID-19 actions
 - Progress of the review program
 - The Environment Working Group – introduction and current guidance and IT related developments
- Heike Schimmelpfennig, Chair BPC Environment Working Group, ECHA, Finland
- 12.05 - 12.25 **Member State overview**
Danish EPA response to COVID-19 with focus on:
- National legislation
 - Enforcement
 - Efficacy issues
- Henrik Svenstrup, Deputy Head of Division, Pesticides & Biocides, Environmental Protection Agency, Denmark
- 12.30 -12.50 **Industry overview**
- Green steps implemented
 - Why does paint need preservatives?
 - Future availability of biocides for PT6 and PT7 are at risk
 - What can be done?
- Hanne Jensen, R&D Manager Biology, Decorative Paints, Jotun A/S, Norway

12.50-13.15 **Q&A and panel session**

Part 2: The assessment of endocrine disrupting chemicals under the BPR

- 14.15 - 14.25 **Introduction**
David Dillon, Senior Regulatory and Compliance Analyst, Chemical Watch
- 14.25 - 14.45 **Experience & tips from an Evaluating Competent Authority**
- Active Substances, already approved and currently under assessment
 - Biocidal products
- Charline Azzopardi, Human Health Expert, Biocides Unit, Ministry of Health, Belgium
- 14.50 - 15.10 **Overview of EFSA endocrine disruptors' assessment of pesticides' active substances**
Alfonso Lostia, EFSA, Italy
- 15.15 - 15.35 **ED assessment experiences for active substances and biocidal products**
Martijn van Velthoven, Director Regulatory Affairs Europe, Troy, Netherlands
- 15.40 - 16.00 **What level of protection from EDCs do we need as a society?**
- ED identification across different EU regulations
 - Improvement of ED standard information requirements and regulatory measures
 - Increasing the level of protection in the short term
- Pia Juul Nielssen, Senior Advisor/Consultant, CHEM Trust

16.00 - 16.25 **Q&A and panel session**





Part 3: The product authorisation and active substance renewal processes

11.00 - 11.15

Introduction

David Dillon, Senior Regulatory and Compliance Analyst, Chemical Watch

11.15 - 11.35

Product authorisation processes: what is and is not possible

- Union Authorisation: The ironman amongst product authorisations?
- National Authorisation: One member state = fast lane?
- Biocidal Product Families: The option of having blue, green and yellow products?
- Single Product Authorisation: The hidden champion?
- COVID-19 impacts: Disinfectants, disinfectants, disinfectants!

Silvia Wagner, Managing Director, spectra Consult GmbH, Germany

11.40 - 12.00

The renewal process including finalised criteria for full/limited renewals

- Explain Ctgb's interpretation of principles of renewal of active substance approvals
- By guiding you through the general framework, including the latest criteria for limited/full renewals
- In order to collectively develop 'fit for purpose' renewals

Lucas Kalkers, Policy Officer, Ctgb, the Netherlands

12.05 - 12.25

Exclusion and substitution in the context of active substances renewals

- BPR Article 5 exclusion criteria
- BPR Article 10 substitution criteria
- Active substances meeting exclusion/substitution criteria - public consultation
- Approaches to Art 5(2) derogation
- Consequences of meeting substitution criteria

Karen Howard, Head of Biocides (Europe), Exponent International Ltd, UK

12.25 - 12.50

Q&A and panel session

Part 4: Sustainable use, innovation and benefits of biocides under the BPR and biocides topics overlapping with other legislation

14.00 - 14.10

Introduction

David Dillon, Senior Regulatory and Compliance Analyst, Chemical Watch

14.10 - 14.30

Sustainable use from Member State perspective

- User categories
- Best practices and Integrated Pest Management
- Control of the use of biocides
- Compliance monitoring
- Additionally: Covid-19 and biocide registration

María Luisa González Márquez, Ministerio de Sanidad, Spain

14.35 - 14.55

Microorganisms under the BPR

Adi Cornelese, Senior Regulatory Expert, APIS Applied Insect Science GmbH, Germany

15.00 - 15.20

REACH, MDR and RoHS overlaps with the BPR

Cristina Garcia, Senior Regulatory and Compliance Analyst, Chemical Watch, Portugal

15.20 - 16.00

Q&A and panel session





Part 5: Testing and modelling

11.00 - 11.15

Introduction

David Dillon, Senior Regulatory and Compliance Analyst, Chemical Watch

11.15 - 11.35

Disinfectant efficacy: experience of preparing efficacy for a product family and how the guidance works in practice

- eCA and peer-review role with efficacy;
- difficulties in matching generic guidance with a non-standard usage area i.e. cleanrooms; and
- keeping up with ongoing changes to the guidance

Siobhan Murphy, Regulatory Specialist, Contec Cleanroom Ltd, UK

11.40 - 12.00

Status and approaches in dietary risk and animal safety assessments

- Available and developing guidance in dietary risk and animals safety assessments
- Approaches for dietary risk assessment in professional vs. non-professional uses
- Currents issues in animal safety: Requirements for exposure and risk assessments

Michael Werner, Senior Regulatory Affairs Manager, Prosacon GmbH, Germany

12.00 - 12.15

Q&A and panel session

Part 6: COVID-19 related updates for disinfectants

13.15-13.25

Introduction

David Dillon, Senior Regulatory and Compliance Analyst, Chemical Watch

13.25-13.40

Covid 19 and the challenges it poses for the disinfection industry

David Ashworth, Managing Director, Klarus Consulting Ltd., UK

13.45-14.00

Industry Covid-19 mobilisation: Disinfectant Practical Guide

- Cross-Industry Alliance on Covid-19
- The practical guide on fast-tracking supply of disinfectants during Covid-19 and under the EU biocides rules
- Overview of derogations in member states and concrete examples
- Learnings and outlook

Nathalie Hanon, Manager CEHTRA SL - EVP, CEHTRA, Spain &

Gerard McElwee, Counsel, Member of the Brussels; Bar (E list), Solicitor, Law Society of Ireland, FieldFisher LLP, Belgium &

Eléonore Mullier, Senior Associate, Steptoe LLP, Belgium

14.00-14.30

Q&A and panel session

Close of conference



Prices

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Three ways to register

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Event timings

26 May 2020 11.00-16.25

27 May 2020 11.00-16.00

28 May 2020 11.00-14.30