

16th International Akademie Fresenius Conference

# **Endocrine Disruptors**

+++ HYBRID EDITION +++

# 25 and 26 November 2025 in Dusseldorf/Germany and via Live Stream

### Highlights

### **Regulatory Developments**

- ECHA on the implementation of the new CLP hazard classes for endocrine disruptors
- EDs under REACH and CLP: Experience from German UBA and BASF
- Outlook on REACH revision and GHS-related activities
- Pesticides: Updates and experience reports from the EU Commission, ANSES, and industry

#### **Next Generation Testing and Innovative Methods**

- · BfR on current regulatory acceptance of NAMs
- Readiness of new ED testing methods
- Thyroid: Increasing confidence in the human thyroid microtissue assay
- Using Al: From QSAR predictions to LLM evidence summarisation

#### **Developments in Ecotoxicology**

- The SETAC MAPPED initiative
- Tiered assessment schemes linking NAMs to adverse outcomes in aquatic vertebrates

### Beyond EATS: New Pathways and European Initiatives

- Next generation risk assessment of obesogenic chemicals: GOLIATH project findings and updates on PARC and AFARA initiatives
- Metabolism-disrupting chemical testing: Advances in methodology and future direction



### The Experts

Jürgen Arning German Environment Agency (UBA) | Arianna Bassan Innovatune | Denise Bloch German Federal Institute for Risk Assessment (BfR) | Pierre-François Chaton French Agency for Food, Environmental and Occupational Health & Safety (ANSES) | ZhiChao Dang Dutch National Institute for Public Health and the Environment (RIVM) | Chad Deisenroth formerly U.S. Environmental Protection Agency (EPA) | Valery E. Forbes Florida Atlantic University | Karma Fussell Wella | Jorke Kamstra Utrecht University | Laurent Lagadic Bayer | Emily McVey Dutch National Institute for Public Health and the Environment (RIVM) | Kirsi Myöhänen European Chemicals Agency (ECHA) | Torben Österlund Public-private platform for the validation of endocrine disruptors characterization methods (PEPPER) | Anette Thiel Ramboll | Helen Tinwell Bayer CropScience | Nikolay Tzvetkov European Commission | Markus Wahl BASF | Lennart Weltje BASF

INSTITUT FRESENIUS



### Tuesday, 25 November 2025

### Morning Session | 09:00 – 12:35 CET

08:30 On-site registration and opening of the virtual meeting room

09:00 Welcome address by the organisers and the Chairs

Lennart Weltje, BASF, Germany

Emily McVey, National Institute for Public Health and the Environment (RIVM), The Netherlands

### Regulatory Developments and Future Perspectives

### 09:10 Implementation of the new CLP hazard classes for endocrine disruptors

- Classification of endocrine disruptors under the CLP regulation
- · ECHA CLP guidance on endocrine disruptors
- Future work by the ECHA ED Expert Group and RAC on endocrine disruptors

Kirsi Myöhänen, European Chemicals Agency (ECHA), Finland (virtually)

### 09:45 Environmental EDs under REACH - Processes, integration of CLP and perspectives

- · Environmental ED assessment
- CLP implications
- Outlook: REACH revision and GHS-related activities

Jürgen Arning, German Environment Agency (UBA), Germany

#### 10:20 Case study on Geraniol: Critical assessment of potential endocrine activity on the thyroid modality and downstream adversities

- Human relevance of potential direct and indirect thyroid effects in vivo, with regulatory implications for REACH, CLP, Biocides, and Pesticides
- Species differences in liver enzyme induction and the question of human relevancy
- Guidance on assessing human relevance and species differences with NAMs and challenges in classification under the CLP ED criteria

Markus Wahl, BASF, Germany

#### 10:55 Coffee break

### 11:25 Accepting the challenge: Quantitative risk assessment for ED substances with a focus on human health - Initial results and potential way(s) forward

- Development of a guidance for quantitative human health risk assessment for ED substances by the CEFIC Sector **Group Biocides for Europe**
- · Presentation of initial results

Anette Thiel, Ramboll, Germany

#### Timings are in Central European Time CET.

### 12:00 Update on the regulatory processes under Regulation (EC) No 1107/2009

- Status of the active substances under the PPP Regulation, including the specific conditions of approvals
- Recent developments in the field, including developments related to ED

Nikolay Tzvetkov, European Commission, Belgium (virtually)

12:35 Lunch break

### Afternoon Session | 13:45 – 18:00 CET

#### 13:45 Member State experience with ecotoxicological ED assessments for pesticides

Pierre-François Chaton, French Agency for Food, Environmental and Occupational Health & Safety (ANSES), France

#### 14:20 Experience and perspectives from the pesticide industry

Helen Tinwell, Bayer CropScience, France

### 14:55 Environmental risk assessment of endocrine-active human pharmaceuticals

ZhiChao Dang, National Institute for Public Health and the Environment (RIVM), The Netherlands

15:30 Coffee break

### Developments in Ecotoxicology

### 16:00 Assessing population relevance of endocrine disrupting effects in fish an amphibians: The SETAC MAPPED initiative

- Population relevance under the ECHA/EFSA ED criteria
- · Applying population models to explore the relevance of individual-level responses in standard ecotoxicological tests from endocrine modalities in fish and amphibians
- Identification of focal species and practical applications of population modelling

Valery E. Forbes, Florida Atlantic University, United States of America

#### 16:35 Tiered assessment schemes linking NAMs to adverse outcomes to identify endocrine disruptors in aquatic vertebrates

- Linking in vitro responses to individual-level adverse outcomes, and further to population-level responses, using coherent, evidence-based tiered assessment schemes
- Using the assessment of the thyroid modality in amphibians as an example of decision logic to navigate through tier testing
- Forward-looking thinking towards a fish-based tiered assessment scheme for EAS (Estrogen/Androgen/ Steroidogenesis) modality

Laurent Lagadic, Bayer, Germany



## Next Generation Testing and Innovative Methods

### 17:10 Advancing the acceptance and use of the human thyroid microtissue assay

- Filling technology gaps for in vitro thyroid testing
- Increasing confidence in the human thyroid microtissue assay as a new approach method
- Integration of thyroid microtissues into an AOP-based tiered screening paradigm to support the context of use

**Chad Deisenroth,** formerly U.S. Environmental Protection Agency (EPA), United States of America (virtually)

17:45 Final discussion18:00 End of the first day18:45 Evening event



After the first conference day, you are welcome to attend our evening event for an unhurried evening of good food and leisure time. We invite you to join us for a little stroll through the famous Düsseldorf Christmas market. Come along to continue the day's interesting discussions in a relaxed and comfortable atmosphere.

### Wednesday, 26 November 2025

## Morning Session | 09:00 – 13:10 CET

09:00 Brief address by the Chairs

**Lennart Weltje,** BASF, Germany **Emily McVey,** National Institute for Public Health and the Environment (RIVM), The Netherlands

## Next Generation Testing and Innovative Methods

### **09:10** The PEPPER experiences with ring-trial validation of methods in the ED space

- Requirements and readiness of ED test methods
- Organising validation projects
- Submission and review under the OECD TG system

**Torben Österlund,** Public-private platform for the validation of endocrine disruptors characterization methods (PEPPER), France

### Timings are in Central European Time CET.

### **09:45** Current status of the regulatory acceptance of NAMs for ED assessment

- Applicability of read-across to evaluate adversity under plant protection and biocidal products regulation
- Further efforts needed for a harmonised and stand-alone NAM-based approach to predict adverse endocrine effects
- More research to improve the in vitro detection of non-EATS mechanisms

**Denise Bloch,** German Federal Institute for Risk Assessment (BfR), Germany

10:05 Case studies in assessing the endocrine disruption capabilities of cosmetics ingredients without new animal testing serving REACH and cosmetic product regulation requirements

Karma Fussell, Wella, Germany

10:40 Coffee break

### 11:10 From QSAR predictions to LLM evidence summarisation: Using AI for endocrine disruptor assessment

- Predictive AI: Use of QSARs and machine learning for predicting endocrine activity to safety assessment
- Large Language Models (LLMs): Role of LLMs in summarising scientific evidence, literature, and mechanistic data to support decision-making
- Addressing explainability, accountability, and transparency as critical elements for applying Al in toxicology

Arianna Bassan, Innovatune, Italy

# Beyond EATS: New Pathways and European Initiatives

### 11:45 From data gaps to detection: PARC's contributions to metabolism-disrupting chemical testing

- Critical data gaps related to key substance groups, including PFAS and bisphenol analogues
- Methodological advancements for the detection of metabolism-disrupting chemicals
- · Findings and way forward

**Denise Bloch,** German Federal Institute for Risk Assessment (BfR), Germany

### 12:20 Next generation risk assessment of obesogenic chemicals in chemical regulation

- Findings from the GOLIATH project (IATA and prevalidation work, chemical selection)
- Ongoing work in PARC and the AFARA project
- Moving further towards Defined Approaches or IATAs

Jorke Kamstra, Utrecht University, The Netherlands

12:55 Summary and final discussion
13:10 Lunch and end of the conference

## Registration

By web www.akademie-fresenius.com/endocrine By email registration@akademie-fresenius.com

Hotline +49 231 75896-50 Die Akademie Fresenius GmbH Alter Hellweg 46, 44379 Dortmund



### **Participation Fee:**

### € 1,995.00 plus VAT on-site in Dusseldorf

Representatives of an authority or a public university are eligible for a reduced fee of € 895.00 plus VAT per person (please provide evidence). The reduced fee cannot be combined with other rebates.

### € 1,195.00 plus VAT Live Stream

Representatives of an authority or a public university are eligible for a reduced fee of € 695.00 plus VAT per person (please provide evidence). The reduced fee cannot be combined with other rebates.

If you are unable to attend, you can order the event documentation for € 295.00 plus VAT.

### www.akademie-fresenius.com/endocrine



Book now

### Do you have any questions?



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### The Organisers

For 30 years, Akademie Fresenius has been your partner for practiceorientated training on all the latest topics surrounding the safety and quality of food, consumer goods and chemical products along the whole production chain. Our portfolio not only includes international conferences but also offers national trade meetings, intensive practical seminars and training in small work groups.

Our events are designed to promote an active exchange amongst our participants and offer the perfect platform for bringing the industry, the scientific sector, the authorities and the consulting field together. Excellent service, all-inclusive. Our wide-ranging advanced training opportunities contribute to giving our customers the competitive edge in all quality assurance, risk assessment, legal, production and

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#### Terms of Participation and Purchase: The registration fee includes the participation in the event, event documentation, and, in the case of on-site events, lunch, coffee breaks, beverages as well as the evening event. You will receive written confirmation of your registration. Upon receiving our invoice, please transfer the amount due without further deductions before the event begins. The price of the event documentation includes the login details for the secure download section of our webpage where you will find the presentations in a pdf format. The login details will be sent to you via email after the event.

Group Reductions: For joint bookings received from one company we grant a 15% discount from the third participant onwards

Terms of Cancellation: For our on-site events, written cancellations or transfers will be accepted free of charge up to four weeks prior to the start of the event. After this date and up to a week prior to the start of the event we will reimburse 50% of the registration fee. We cannot, unfortunately, provide refunds for later cancellations. For our online events, cancellations or rebookings to another event will be accepted free of charge up to one week before the start of the event. In the event of later cancellations or non-login to the online event, no participation fees can be refunded. In this case, however, you will receive access to the documentation download after the event.

### Who should attend this conference?

#### Professionals working in the fields of:

- Toxicology and ecotoxicology
- Research and development
- Registration and regulatory affairs
- Chemical risk assessment
- Legal and general counselling

#### Sectors that should take part:

- Chemical/biocide/agrochemical/pharmaceutical/ cosmetic industries
- Research institutes, Regulatory authorities
- Environmental and health risk consultants
- Testing laboratories and contract research organisations (CROs)

### **Conference Venue**

https://all.accor.com

Novotel Düsseldorf City West Niederkasseler Lohweg 179, 40547 Düsseldorf Phone: +49 211 52060-0 Email: h3279@accor.com

We have reserved a limited number of rooms for our participants at the hotel. These rooms can be booked up to 4 weeks prior to the start of the event. Please book early and directly through the hotel quoting "Akademie Fresenius" as reference.

Please note that you can name a substitute participant free of charge at any time.

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