



Online Training

A Practical Guide to the ECHA/EFSA Endocrine Disruptors Guidance: Biocides, REACH and PPPs

15 and 16 December 2026, Online 

Navigate the ECHA/EFSA guidance documents on endocrine disruptors with confidence. This Training will provide you with practical knowledge on how to implement the requirements for the different relevant regulations: REACH, BPR and PPPs

Learning Objectives:

- The most important aspects of EDs in general and the ECHA/EFSA guideline
- How the guideline impacts the BPR regulation
- How the guideline impacts REACH
- How the guideline impacts the PPP Regulation
- how the guideline impacts the PPP Regulation

Tuesday, 15 December 2026

09:00 – 15:00 CET

Introduction

- General overview ED (definition, biological/scientific background)
- General overview ED (short history)

General overview OECD 150

- General overview of the guideline
- Principles of OECD 150
- Methods for the evaluation of EDs properties

ECHA/EFSA Guideline and ECETOC implementation (the 7 steps)

- General overview of the two guidelines
- Methods for the evaluation of EDs properties
- Practical implementation of the guidelines for regulatory purposes
- General overview of the two guidelines (Human Health and Environmental)
- Case studies (Human Health and Environmental)

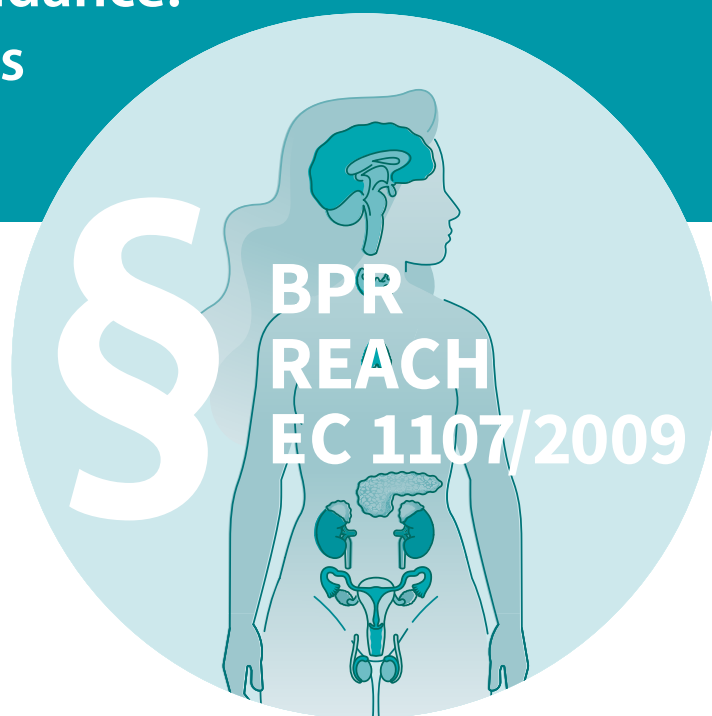
Timings are in **Central European Time CET**.

The Experts

Costanza Rovida, TEAM mastery

Libero Andrea Mazzucchelli, TEAM mastery

Chiara Marelli, TEAM mastery



Wednesday, 16 December 2026

09:00 – 15:00 CET

Impact on BPR

- Regulatory aspects of the ED properties
- Regulatory requirements for active substances and biocidal products
- Regulatory consequences for active substances and products
- Identification of ED in Biocides
- EDs assessment of biocidal products
- Case study

Impact on PPP Regulation

- Regulatory aspects of the ED properties
- Identification of EDs in PPP
- Insights and practical steps for active substances approval
- Case studies & Insights for co-formulants

Impact on REACH Regulation

- Regulatory aspects of the ED properties
- Overview of endocrine disruptor SVHC and listed in Annex XIV (REACH)
- Assessment of the toxicological endpoint triggering ED properties
- Risk assessment for the Application for Authorisation

Status of ED in other Regulations

- CLP: new risk classes

Participation Fee

€ 845.00 plus VAT

Registration

By web www.akademie-fresenius.com/4083
By email registration@akademie-fresenius.com

Participation Fee: € 845.00 plus VAT

Representatives of an authority or a public university are eligible for a reduced fee of € 595.00 plus VAT per person (please provide evidence). The reduced fee cannot be combined with other rebates.

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



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Do you have any questions?



Anne Möller
Phone: +49 231 75896-84
anne.moeller@akademie-fresenius.de

The Organisers

For more than 30 years, Akademie Fresenius has been your partner for practice-orientated 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 technical questions.

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You can find details on upcoming and new events at www.akademie-fresenius.com

Who should attend this training?

Professionals working in the fields of:

- ✓ Regulatory affairs
- ✓ Registration
- ✓ Risk assessment
- ✓ Toxicology, Ecotoxicology

Sectors that should take part:

- ✓ Plant Protection industry
- ✓ Biocide Producers
- ✓ Chemical Industry (REACH)
- ✓ Scientific and regulatory consultancies

How will this online training work?

Our online training will be live – with interactive participation – and will be held in the English language. The training will be conducted using the meeting tool Zoom. Prior to the training, we will provide you with your login details, which will allow you to participate and ask questions from your preferred location. Simply log in on the day of the training – and away you go!

Terms of Participation and Purchase: The registration fee includes the participation in the online event and the event documentation for download. You will receive written confirmation of your registration. Upon receiving our invoice, please transfer the amount due without further deductions before the event begins.

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

Terms of Cancellation: Cancellations or rebookings to another event are possible in writing without giving reasons and free of charge up to one week before the start of the online 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. Please note that you can name a substitute participant free of charge at any time.

General Terms and Conditions: By registering, you agree to our General Terms and

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