TRAINING

« COSMETICS REGULATION IN EUROPE AND THE LATEST NEWS »

3 Afternoons : 24/11, 25/11 et 27/11

This training allows you to develop or validate your knowledge in the field of cosmetics. You will find all the necessary information to better understand the constraints as well as the opportunities to bring a product to market that complies with regulations.

24/11, 25/11 et 27/11

MONDAY 24 & TUESDAY 25 NOVEMBER 2025

- Validate your regulatory knowledge concerning cosmetics.
- Understand the constraints.
- Study and experiment on one or more concrete cases.
- Q&A – Exchange and debate on the subject.

THURSDAY 27 NOVEMBER 2025

- Validate your regulatory knowledge concerning cosmetics.
- Understand the constraints.
- Study and experiment on one or more concrete cases.
- Q&A – Exchange and debate on the subject.

Public

- Anyone involved in the cosmetics development and/or marketing process (CEO, project manager, R&D, regulatory or marketing departments).

Prerequisites

- No special prerequisites.

Level

No level required

Terms

Remote Training (Webinar)

Accessibility

If a specific adaptation is needed, particularly in relation to a disability, please do not hesitate to let us know.

REGISTRATION FORM TO BE RETURNED TO
CONTACT@PHARMANAGER-DEVELOPMENT.COM

PHARMANAGER development

TRAINING

« COSMETICS REGULATION IN EUROPE AND THE LATEST NEWS »

2 AFTERNOONS : 24 & 25 NOVEMBER 2025

- Attendees' connection
- Presentations / Round table of everyone's expections and objectives

Cosmetic Product Regulation in Europe: Focus on the Product Information File

– The regulation of cosmetics in Europe
– Regulation 1223/2009 EC on cosmetic products:
– Objectives
– Definitions
– How to compile my product information file?
– File structure:
– Sections other than the safety report
– The safety report
– The 10 parts that make up section A of the report

Cosmetic Safety Assessment

Post-Marketing Surveillance: Cosmetovigilance, How to Implement Your Procedure?

- Regulatory obligations:
  – Safety concept: central aspect of the regulation
  – Regulatory obligations
  – Regulatory definitions, severity criteria, reporting to authorities
- Cosmetovigilance cases:
  -Definition of a case
  – Causality assessment methods
  – Responsibilities
  – Steps in case management
  – Different types of reactions to cosmetic products
- Case study:
  – Use of collected data / evaluation / decision-making
  – Integration of data into the Product Information File
  – Evaluation, reporting
- Market surveillance of cosmetic products:
  – General overview
  – Focus on France (ANSES and DGCCRF, inspections, and penalties...)

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1 Afternoon: 27 November (1.30pm- 5.00pm)

- Attendees' connection
- Presentations / Round table of everyone's expections and objectives

Cosmetic Claims, How to Communicate Safely?

- Context and regulatory texts:
  – Self-regulation regime
  – The six common criteria of the European Regulation: definition and examples
- Case study:
  – "Free-from" claims: the "spirit" of the text, what is allowed, what is prohibited
  – "Hypoallergenic" claim: the "spirit" of the text, how to justify this type of claim?
- How to justify a claim?
  – Regulatory context of clinical testing for cosmetic products
  – Types of clinical tests: safety, use, effectiveness
  – Methodologies of safety and efficacy tests, proof of claimed effects
- Website / Advertising / Influencers

=> Quiz, exercises: Validate your knowledge

TRAINING

« COSMETICS REGULATION IN EUROPE AND THE LATEST NEWS »

TRAINERS

Céline Pozza

Scientific and Regulatory Affairs Officer – Toxicology, Medicinal products, Biocides, Annex III Plants

11 years of experience

Geoffrey Tessier-Houeix

Scientific and Regulatory Affairs Officer – Cosmetics

7 years of experience

MEANS AND MODALITIES

The training will be conducted via the Teams video conferencing tool (a link will be sent to participants before the session for access). The training material (PowerPoint) will be presented through a combination of lectures and discussions with the trainers. Educational documentation will be provided in electronic format. A knowledge assessment must be completed at the end of the training. A certificate of attendance will be issued.

REGISTRATION FORM

« COSMETICS REGULATION IN EUROPE AND THE LATEST NEWS »

Company name:

Name of signatory:

Intracommunity VAT number:

Company Training Manager:

(email required - Otherwise: Person signing the training agreement)

PARTICIPANTS AND CHOICE OF SESSIONS

Name : .....
First name : .....
Function : .....
E-mail : .....

☐ 2 Afternoons: 24 & 25 November 2025 (1.30pm- 5.00pm) - 750€HT soit 900€TTC

Name : .....
First name : .....
Function : .....
E-mail : .....

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If a particular adaptation of the training is necessary, linked in particular to a particular disability situation, do not hesitate to let us know.

Before registering, make sure you have read the training prerequisites and validate that the objectives are in line with expectations. Do not hesitate to let us know your expectations before the training if necessary.

Individual expectations and objectives:

Total : **€ tax ex.**

Name and address of the company to be invoiced:

Date :

Signature :

Pharmanager Development reserves the right to cancel the training in the event of insufficient registrations. In the event of cancellation, Pharmanager Development will notify participants at least 5 working days before the scheduled training date.

IMPORTANT: Our training courses are eligible for support by an OPCO. However, the invoice will be established and due by the participating company, in the amount of the training.

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