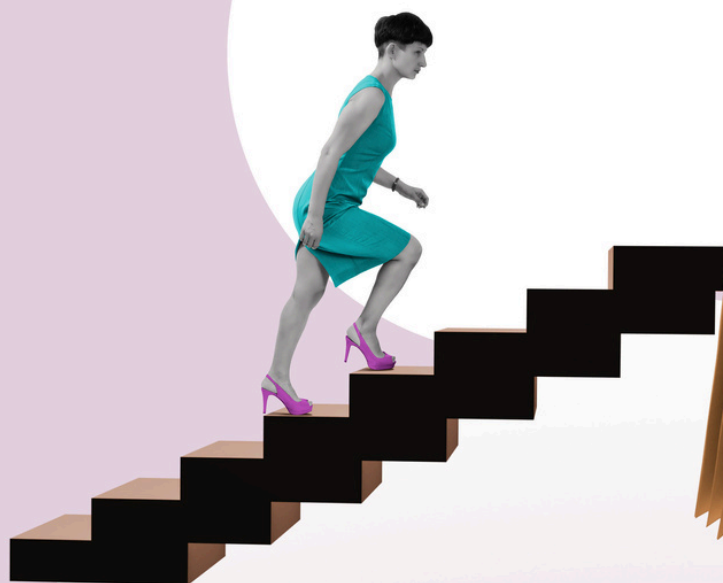


BUILD SKILLS WITH  
**COSMED ACADEMY**

INTER-COMPANY  
INTRA-COMPANY  
E-LEARNING



REGULATIONS  
EXPORT  
MICROBIOLOGY  
ENVIRONMENT

# Train *with Cosmed*

Cosmed Academy offers you a wide range of training in European and International cosmetic regulations.

Our training courses are designed to help you build your skills and support you in achieving regulatory compliance for your cosmetic products.

They are developed by experts in the cosmetics industry and address the latest regulatory developments.

Our organisation is registered under activity declaration number **93 13 216 36 13**.

## Level of satisfaction

*2023 total (January to December)*



## Rates

*Inter-company, remote,  
virtual classroom training*

	7H (1D)	14H (2D)	2H-3H30	4H-4H30
• Member	€420	€840	€200	€250
• Member (addit.pers)	€350	€700	-	-
• Non-member	€520	€1040	€300	€350



## Location & times *of training*

**All our trainings are available on-line**

**For on site trainings**, we welcome you to our premises located in Aix en Provence, 15 minutes by taxi from the Aix Tgv Train Station.

The 7-hour training sessions take place from 9 am to 5 pm. The training sessions lasting between 2 to 4 hours take place between 9 am and 2 pm.

## Catering *On-site*

Cosmed covers the costs of catering during training days. You can eat at our partner La Famille located near our premises.

## Accommodations

*Nearby*

**The Camp** - + 33 4 13 91 20 30

550 rue Denis Papin, 13100 Aix-en-Provence

**Hôtel de l'Arbois** - + 33 4 42 58 59 60

97 Rue du Dr Albert Aynaud, 13100 Aix-en-Provence



***Your contact person : Saloua Salhi***

Disability Expert : Saloua SALHI

+33 4 42 16 63 82 [formation@cosmed.fr](mailto:formation@cosmed.fr)

# REGULATIONS FINISHED PRODUCTS

## BASICS OF EU COSMETIC REGULATIONS AND GOOD MANUFACTURING PRATICES (2 DAYS)

### OBJECTIVES

- Understanding the basics of EU cosmetic regulations

### DAY 1 PROGRAM

#### 1-European regulations

- Definitions : cosmetic product, responsible person, manufacturer, distributor, making available on the market
- Safety evaluation assessor qualification, content of the Safety report
- Cosmetovigilance : what obligations ?
- Product Information Dossier
- Product notification : Simulating a notification
- Substance restrictions : Annexes II to VI
- Animal Experimentation
- labelling : Mandatory information
- Claims : Common criteria and guidelines

#### 2-Competent authorities in France : ANSM, DGCCRF

- Market Surveillance and control in France

### EDUCATIONAL METHODS

Presentation media provided prior to training  
 Alternation of theoretical, practical and concrete contributions  
 Technical means: video projector, whiteboards, LMS Software

### MONITORING/EVALUATION

Questionnaire expectations and level evaluation  
 Results : questionnaire at the beginning and end of the training  
 Satisfaction at the end of the training and 1 month after  
 Implementation : attendance sheet per half-day and delivery of a training certificate of completion.

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COSMED Academy



In person or remotely



7h (1 day)  
9am -5pm



R&D, marketing, Quality Departments

Participants : 3 to 8 max



No prerequisites



Cosmed  
495 rue René Descartes  
Les Ocres de l'Arbois, Bat B  
13100 Aix en Provence

# REGULATIONS FINISHED PRODUCTS

## BASICS OF EU COSMETIC REGULATIONS AND GOOD MANUFACTURING PRACTICES (GMP)

### OBJECTIVES

- Understanding the basics of GMP

### DAY 2 PROGRAM

- 1- ISO 22716 Scope
  - Terms of definition
  - Staff, Premises, Equipment
  - Raw Materials and packaging materials Production
  - Finished products
  - Quality Control Laboratory
  - Handling of out of specification products
  - Waste, Deviations
  - Complaints and Recalls
  - Change management
  - Internal audit, documentation
- 2 - The non conformities most encountered during an audit
  - Surveillance and control of the market in France
  - Update on inspections carried out by the authorities

### EDUCATIONAL METHODS

Presentation media provided prior to training  
 Alternation of theoretical, practical and concrete contributions  
 Technical means: video projector, whiteboards, LMS Software

### MONITORING/EVALUATION

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COSMED Academy



In person or remotely



7h (1day)  
9am - 5pm



R&D, marketing, Quality Departments

Participants : 3 to 8 max



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No prerequisites

## HOW TO GATHER THE PRODUCT INFORMATION DOSSIER?

### OBJECTIVES

- Knowing the building blocks of the product information dossier
- Understanding the purpose of each section of the safety report
- Knowing the data to be provided for each section according to product type
- Understanding how to integrate the safety and toxicology report into the product information dossier

### PROGRAM

- 1- Reminder of the regulatory context
  - 2- Dossier Structure
  - 3- Items other than the Safety report
  - 4- Safety report
- The 10 parts making up Part A of the report
- Objectives of each section
  - Required data
  - What to ask to suppliers
  - Explanation of key calculations

- 5- Part B of the report

The Training will be illustrated by an example from the product information dossier

### EDUCATIONAL METHODS

Presentation media provided prior to training  
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In person or remotely



7h (1day)  
9am - 5pm



R&D, Marketing, Regulatory  
Departments

Participants : 3 to 10 max



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Knowing the cosmetic products

## SAFETY EVALUATION IN COSMETICS

### OBJECTIVES

- Understanding the principles and concepts of the safety assessment
- knowing how to calculate the safety margin of a cosmetic ingredient and formula

### PROGRAM

1. Risk Management (hazard and exposure concepts)
2. Safety assesment and the European cosmetic regulations (dedicated articles and Annex I)
3. Guidelines and guiding values
4. Toxicological indexes and values (NOAEL, LOAEL, PoD)
5. Regulatory tools (data access)
6. Risk assessment of cosmetic ingredients
7. Technical and regulatory documents required
8. Perspectives on the European Green Deal
9. Examples and practical cases (MoS calculation)

### EDUCATIONAL METHODS

Presentation media provided prior to training  
 Alternation of theoretical, practical and concrete contributions  
 Technical means: video projector, whiteboards, LMS Software

### MONITORING/EVALUATION

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# REGULATIONS FINISHED PRODUCTS

## CLAIMS, ADVERTISING : REGULATORY FRAMEWORK

### OBJECTIVES

- Mastering the justification of claims
- Choosing your claims, their communication
- Preparing for future regulatory developments

### PROGRAM

#### 1-Regulatory context

- Regulatory texts
- Self regulation regime

#### 2-Claim definitions what -what the texts cover

#### 3- The six common criteria of European regulation

- Definition
- Examples

#### 4-"Without " claims

- the Spirit of the text
- What is permitted
- What is prohibited

#### 5-"Hypoallergenic" claims

- The spirit of the text
- How to justify this type of claim

#### 6- Environmental claims

- Organic, natural, recyclable... impact of the AGECE law

#### 7-How to justify a claim

#### 8-Workshop around labelling, participant communication Media

### EDUCATIONAL METHODS

Presentation media provided prior to training  
 Alternation of theoretical, practical and concrete contributions  
 Technical means: video projector, whiteboards, LMS Software

### MONITORING/EVALUATION

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COSMED **Academy**



in person or remotely



7h (1 day)  
9am - 5pm



Regulatory, Manufacturing, R&D

Participants 3 to 10 max



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No prerequisites



# REGULATIONS FINISHED PRODUCTS

## LABELLING YOUR COSMETIC PRODUCTS SAFELY

### OBJECTIVES

- Understanding legal requirements
- Knowing how to adapt your labelling according to the products

### PROGRAM

#### 1-General labelling rules

- Regulatory requirements in cosmetics
- Rules for ingredient lists (order, content, allergens, +/-...)
- Small products
- Unpackaged products (packaged at point of sale, bulk)
- Fragrance and flammable products- Aerosols and CLP
- Professional products
- Environmental labelling (TRIMAN, INFO TRI, AGECE) wipes

#### 2-Labelling workshop

- Workshop around labelling with participants

### EDUCATIONAL METHODS

Presentation media provided prior to training  
 Alternation of theoretical, practical and concrete contributions  
 Technical means: video projector, whiteboards, LMS Software

### MONITORING/EVALUATION

Questionnaire expectations and level evaluation  
 Results : questionnaire at the beginning and end of the training  
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# REGULATIONS FOR INGREDIENTS

## I HAVE A NEW COSMETIC INGREDIENT WHAT ARE MY OBLIGATIONS ?

### OBJECTIVES

- Understanding the definitions and regulations involved
- Mastering French specificities
- Knowing the obligations in Europe and for Export

### PROGRAM

1. **Définitions, relevant regulations, practical advice on implementation**
2. **Obligations in Europe :**
  - INCI repository
  - Summary of REACH AND CLP obligations : chemical and toxicological characterisations - SDS
  - Regulation 1223/2009 : Evidence of safety
  - Professional recommendations
3. **French specificity : nano declaration**

### EDUCATIONAL METHODS

Presentation media provided prior to training  
 Alternation of theoretical, practical and concrete contributions  
 Technical means: video projector, whiteboards, LMS Software

### MONITORING/EVALUATION

Questionnaire expectations and level evaluation  
 Results : questionnaire at the beginning and end of the training  
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 Implementation : attendance sheet per half-day and delivery of a training certificate of completion.

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COSMED Academy



In person or remotely



7h (1 day)  
9am - 5pm



regulatory, R&D Department

Participants : 3 to 10 max



Understanding basics of regulations

## INNOVATING WITH THE NAGOYA PROTOCOL

### OBJECTIVES

Knowing the definitions and regulations involved

- Understanding the obligations of a cosmetic company
- Practical advice on the implementation of Nagoya adapted to the cosmetic sector issues

### PROGRAM

1. Nagoya Protocol : an overview of the objectives and scope
2. Definitions and the international, European and French regulatory framework
3. ABS regime : Access to genetic resources and sharing of benefits from their use
4. Supplier/user relationship : Prior Consent and the terms and conditions of an agreement.
5. Implementation for a cosmetic company : key elements and practical advice

### EUDCATIONAL METHODS

Presentation media provided prior to training  
 Alternation of theoretical, practical and concrete contributions  
 Technical means: video projector, whiteboards, LMS Software

### MONITORING/EVALUATION

Questionnaire expectations and level evaluation  
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In person or remotely



7h (1 day)  
9am - 5pm



R&D, Marketing, Quality Departments

Participants : 3 to 10 max



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No prerequisites

# CONDITIONS

## General

### REGISTRATION PROCEDURES :

Any participant must be confirmed by sending the registration form and the payment (unless covered by a funding body). The payment will only be cashed after completion of the training. Upon receipt of the registration form, Cosmed will send you a training agreement. A notification will be sent you no later than 10 days before the training. A certificate of completion will be sent to you after the training, accompanied by a paid invoice.

### SUPPORT SPECIFIC TO FRANCE :

All our training can be covered as part of the continuing education plan for employees or company manager.

For trainees wishing to be supported by their OPCO, registration must be completed 3 weeks before the start of the training.

A bilateral training agreement will be sent to you, together with documents intended for the funding body, Cosmed is Qualiopi Certified.

### CANCELLATION CONDITIONS :

In the event of a withdrawal of the trainee notified in writing 15 calendar days before the date of the training, a fee amounting to 50 % of the total amount of the registration fee will be charged. After this deadline, the total amount of the registration fee will be due. Cosmed reserves the right to cancel a training session at least 8 working days before the start of training. In this case, the registration will be postponed to another session or the registration fee will be returned.

