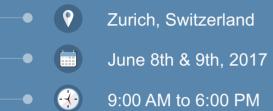


2-day In-person Seminar:

# Cosmetics Product Regulation – Four Years after its implementation





# **Annelie Struessmann**

Technical Director, CONUSBAT

Dr. Annelie Struessmann is the Technical & Regulatory Director with CONUSBAT Regulatory Services, a provider of internationalization compliance services for Cosmetics, Personal Care, Fine Chemicals and Borderline Industries: www.conusbat.com. In 2003, she joined the firm and, based on many years of experience gained while working with the international industry, she established the service area for regulatory affairs.

# **Overview** :

Since July 2013, Regulation (EC) No 1223/2009 on Cosmetic Products of the EU has been fully implemented and represents a modern regulatory framework grounded on state of the art of cosmetic science and product technology. It is structured in regulatory modules, which include the safety assessment and the Cosmetic Product Safety Report (CPSR), Product Information File (PIF), Responsible Person (RP), label information, cosmetovigilance, substance regulations, claims, etc. The legislator's existing goal is assurance of the safety for the ingredients and for the cosmetics products in use of consumers. Price

Price: **\$1,695.00** (Seminar for One Delegate)

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\*\*Please note the registration will be closed 2 days (48 Hours) prior to the date of the seminar.







# Agenda:

#### **Day One**

#### Day Two

Lecture 1: Welcome of Participants & Presenters -Introduction into the Topic

Lecture 2: The legislative Environment and Authority of the EU, Stakeholder Participation Processes

Lecture 3: Regulation (EC) 1223/2009 - The Cosmetics Product Regulation (CPR) - Framework Structure, Key Provisions, Regulatory Modules

Lecture 4: Roles and Responsibilities, the Responsible Person

#### Why should you attend:

The European Union's (EU) Cosmetics Products Regulation (CPR) 1223/2009 represents a newly re-casted piece of legislation, fully implemented since July 2013, and putting in place demanding provisions for those seeking compliance. Affected by the challenges of meeting compliance are the players in the world's biggest cosmetic market, the EU's internal market, as such similarly EU and non-EU manufactures of cosmetics as well as the suppliers of cosmetic ingredients requested to provide data on their chemicals.

Provisions of the CPR are formulated as modules and these include substance regulations, product information file, labeling, cosmetovigilance, claims, notification, responsible person, etc. Compliance with these modules requires know-how, diligence and ongoing adjustment to state of the art of knowledge and documentation. In this format, the EU Regulation represents not only the entry requirements for marketing of cosmetic product in the European Union;

- Lecture 1a: A look at the Cosmetic Sector, History, Compliance and other Issues
- Lecture 1b: The Animal Testing Ban
- Lecture 2: Outstanding Chapters of the CPR PIF and Cosmetic Product Safety Report, Substance Regulations
- Lecture 3: Outstanding Chapters of the CPR Criteria for Claims, Notification, Market Surveillance, Consumer Information
- Lecture 4: Regulatory compliance as a tool to improved performance

#### Areas Covered in the Session:

- Animal Testing Ban
- Roles & Responsibilities in the Supply Chain
- Product Information File (PIF)
- Safety Assessment
- Criteria for Claims
- Cosmetovigilence
- Substance Regulations
- Product Labeling
- Borderline Industry Legislation

#### Who Will Benefit:

- Managers in Cosmetics, Personal care and Consumer Health Product Industries
- Managers in Ingredient Suppliers, fine Chemicals
  Companies
- Managers in Regulatory Affairs, R&D, Formulators
- Government officials, KOLs
- International Sales & Marketing Representatives



	Group Participation
10%	2 Attendees to get offer
20%	3 to 6 Attendees to get offer
25%	7 to 10 Attendees to get offer
30%	10+ Attendees to get offer

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Look forward to meeting you at the seminar

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