

Workshop: The New European Regulation on Cosmetic Products

Part I: Overview of the legal situation

Welcome & Introduction

How the EU regulatory frameworks for cosmetics evolved in Europe

- The European Union
- The internal market
- EU bodies
- EU regulatory instruments
- Scientific committees
- Stakeholder impact & stakeholder organizations

The new EU regulatory approach for cosmetics

- Why a new EU Regulatory Framework
- Actual status of the implementation
- Transitional periods
- A synoptic view of the outline of the new regulation
- How and where to look for new contents & provisions
 - New definitions
 - New phrasings
 - New standards
 - New terms
 - New references

Q&A / Break

Part II: Safety assessment and product information files

Safety Requirements from the Cosmetics Regulation

- Contents of the Product Information File
- The Cosmetic Product Safety Report as part of the PIF
- EU Commission Guidance
- A Practical Approach towards the Safety Assessment

Replacement of animal tests & non-testing strategies

- Facts and Issues from the Animal Testing Bans
- The EFfCI Approach and Conclusions
- Status on Availability of Alternative Methods
- EFfCI Practical Advice For Industry

Q&A / Break

Part III: Selected chapters with major changes

Responsibilities for actors in the supply chain

- The position 'Responsible Person'
- Manufacturers, importers, distributors and their obligations

Changes to the notification procedure

- The central electronic procedure
- New information requirements
- Specific distributor responsibilities

Additional labelling provisions

Changes towards provisions for substances

- Nanomaterials
- CMR substances in cosmetics

Access to information for the public & market surveillance

- In-market control
- Communication of serious undesirable effects

Q&A / Break

Concluding summary