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REACH

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The essential update on REACH – analyse the legal implications of recent developments and what practical strategies to adopt throughout the supply chain

**SRA and BSB CPD:
12.5 hours**

DAY 1 Tuesday 23rd June 2015

08.30 Registration and Coffee

09.00 Chair's Welcome & Introduction



Gauthier van Thuyne

*Partner, Head of the Belgian Environmental Law Department,
Co-Chair of the Global REACH team*

ALLEN & OVERY LLP

09.10 Articles under REACH and the Impact of Latest Developments in the Guidance

- Update on the requirements
- What is the likely practical impact on supply chain management?
- How to calculate amount of SVHCs and articles?
- Ensuring compliance when the goalposts are moving



Dr. Barry Podd
CONSULTANT

10.05 Substance and Dossier Evaluations



- CoRAP listing and substance process
- Dossier evaluation processes
- Lessons learned including read-across, grouping, animal testing and intermediates
- Challenges for the industry including the new ECHA policies on updates and EOGRTs
- Legal remedies



Darren Abrahams
Partner

STEPTOE & JOHNSON



Jean-Philippe Montfort
Partner

MAYER BROWN

10.50 Morning coffee

11.10 Lessons Learnt from 2010 and 2013 Registrations and Practical Guidance to Prepare for 2018 Registrations



- REACH 2018 roadmap
- How to prepare for it now?
- What lessons can be learnt from 2010 and 2013 registrations?
- Illegal dossiers
- Letter of access
- Lead times in testing houses
- Will it lead to withdrawal of products?
- Data requirements
- How will authorities approach enforcement?
- Learning from dossier and substance evaluation
- Discussions on dossier quality
- What triggers a CSR?



Dr. Bernd Friede
*REACH Compliance
Manager*

**ELKEM AS, SILICON
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Matteo Dalla Valle
*European Regulatory
Specialist*

**CHEVRON ORONITE
S.A.S.**



Endwin Hensema
*Product Regulatory
Manager EU*

ARIZONA CHEMICAL

Linda-Jean Cockcroft
Associate Partner
EPPA



Sylvia Jacobi
*Corporate Toxicology
Director*

ALBEMARLE EUROPE

12.00 REACH for SMEs

- Specific problems with registrations 2018
- Data sharing
- Authorisation
- Recent case law on administrative fees
- IT tools/guidance



Douglas Leech
Technical Director

CHEMICAL BUSINESS ASSOCIATION

12.40 Lunch

13.40 Authorisation Experience: an Assessment of More than 20 Successful Authorisation Applications (AfAs) so Far



- What were the most convincing arguments to receive a positive response to an application?
- What drove ECHA committees to define review periods and authorisation conditions?
- Different challenges for manufacturers, formulators and downstream users when preparing for and submitting authorisation applications



Hugo Waeterschoot
REACH advisor
EUROMATEUX

14.25 How to Deal with the Threat of Authorisation: Practical Strategies to Assess Human Exposure Preceding Authorisation

- How to be prepared for authorisation: relevance of exposure information
- Human exposure: workers and consumers
- Use of (historical) exposure data, industry practice
- How can industry influence the choice for a risk management option?



Doeke van der Schaaf
Industrial Chemicals Program Manager
TNO TRISKELION

15.10 Afternoon tea

15.30 Update on Enforcement

- National interpretation of REACH
- How will authorities react to registrations 2018
- Enforcement of substances with authorisation requirements

Senior Representative
EUROPEAN CHEMICAL AGENCY

16.15 Review of Recent Case Law at the Board of Appeals and CJEU

- Disclosure of information cases
- Intermediate status of process agents
- Dossier evaluations
- Board of Appeal procedure and remedies



Gauthier van Thuyne
*Partner, Head of the Belgian Environmental Law Department,
Co-Chair of the Global REACH team*
ALLEN & OVERY LLP

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Julie Cheuy-Ruckect, Nintendo

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17.00 REACH and Biocides

- Interlink between REACH and biocides
 - Commodity chemicals
 - Speciality chemicals
 - In-situ generated chemicals
- REACH Registration versus Article 95 BPR Listing
 - Relevance for marketing / authorising chemicals
 - Data access and "cascade rights"
 - Mandatory data sharing
 - Scope
 - Principles
- Remaining issues with data sharing
- Conclusions



Koen Van Maldegem
Partner, Competition and
EU Regulatory
FIELD FISHER



Dr. Jeannette Paulussen
Head of Regulatory
Affairs
WIL RESEARCH

17.40 Chair's Closing Remarks

17.45 End of Day One

DAY 2 Wednesday 24th June 2015

08.30 Registration and Coffee

09.00 Chair's Welcome & Introduction



Cándido García Molyneux
Of Counsel
COVINGTON & BURLING

09.15 New Requirements under Classification and Labelling Regime (CLP)



- Challenges for companies, in particular SMEs
- New regulation and requirements, including ATP
- Trigger for regulation
- Rules for mixtures
- ECHA practices
- How CLP interacts with other regulations, including REACH



Dr. Roger Van der Linden
CONSULTANT
REPRESENTING
BOREALIS



Prof. Lucas Bergkamp
Partner
HUNTON & WILLIAMS

10.05 Data Sharing

- ECHA procedure and recent Board of Appeal case law
- Implementing the new European Commission regulation on transparency and cost sharing in SIEFs
- Pricing principles and structures
- Arbitration
- Data sharing and confidentiality considerations
- Global data sharing



Ursula Schliessner
Partner
JONES DAY

10.50 Morning coffee

11.10 Communication of Information in the Supply Chain: Safety Data Sheets



- Transitional rules
- Implications for downstream users
- ECHA's update on data requirements and assessment
- Data fraud
- Illegal dossiers



Dr. Bernd Friede
REACH Compliance
Manager
ELKEM AS, SILICON
MATERIALS



Cándido García Molyneux
Of Counsel
COVINGTON & BURLING

12.00 Regulating Nanomaterials

- Recent working group decision regarding the definition
- JRC report
- Current position on the mandatory reporting scheme
- Review of different approaches by different countries
- Revision of the REACH Annexes to address nanomaterials
- Nanomaterials under vertical legislation (biocides cosmetics, medical devices, food, food contact etc)



Steffi Friedrichs
Director General
NANOTECHNOLOGY
INDUSTRIES ASSOCIATION



Dr. Anna Gergely
Director, EHS Regulatory
STEPTOE & JOHNSON

12.50 Lunch

13.50 Obligations and Responsibilities throughout the Supply Chain

- Supply chain evaluation: know your supply chain for your different products - they're not all the same
- Transport regulation: which key aspects are important
- Customs: why this matters, its interface with your different types of customers and consequences for your REACH obligations
- Case study on monomers in imported polymers: what is needed; issues to obtain this
- Supply chain communications and dossiers: know what your customers do with your product
- Strategies: is it better to use products from outside the EU? Comparison of the amount of work vs freedom and being in control

Dr. Peter Douben
Founder & Director
REACHWISE

14.35 Global Compliance



- Review of relevant regulations in China, Korea and Turkey
- Legal registers in different countries
- What are the best strategies to adopt?
- Prospects for transatlantic regulation

Moderator:



Cándido García Molyneux
Of Counsel
COVINGTON & BURLING



Nick Choi
Junior Consultant
CHEMTOPIA CO. LTD.



Jean-Philippe Montfort
Partner
MAYER BROWN

Louise Halpin
Managing Director
CHEMICAL INSPECTION
AND REGULATION
SERVICES LTD.



Melih Babayigit
Owner
CRAD CEVRE RISK ANALIZ

15.40 Afternoon tea

16.00 Fair and Transparent Cost Sharing

- Distributors' view on fair and transparent cost sharing
- Challenges faced during the previous registration deadlines
- The ECHA Board of Appeal decision in *Vanadium*
- Reconciling treatment of consortium members and LoA recipients



Dr. Uta Jensen-Korte
Director General
EUROPEAN ASSOCIATION
OF CHEMICAL
DISTRIBUTORS (FECC)



Scott S. Megregian
Partner
K & L Gates

16.50 Chair's Closing Remarks

17.00 End of Conference

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