

Event programme

Updating your REACH registration dossier

29 February 2012 – CIA Offices, London, SW1P 3JJ



Who should attend?

Manufacturers, Importers and Only Representatives who have found errors in their registration dossier, who need to re-evaluate the classification, or have new customer uses to include need to update their registration. This workshop will look at why updates are needed, when and by whom – whether by the Lead Registrant only or also by the co-registrants.

You'll get the most from the day if you have a working knowledge of IUCLID and previously prepared registration dossiers. However, if you're not fully up to speed don't worry, you'll still really benefit from coming along to help determine if you need to take action or not.

Book before 29 January for an "early bird" discount.

Why attend?

Achieving your first registration was a significant step towards REACH compliance – but the duties do not stop there. More than a year has passed since the first phase-in deadline and you may need to update your registration to ensure it remains valid. Perhaps you have been asked by ECHA to provide more information, or you know you need to improve the quality of the dossier you submitted. Maybe you have new uses to be included in your Chemical Safety Report. Or perhaps you registered your substance as an intermediate under strictly controlled conditions but need to upgrade your dossier to that of a standard registration.

This one-day workshop will explore the requirements for dossier upgrades and updates and give you a practical insight into meeting your obligations including:

- ✓ What is needed to update a dossier, when and why
- ✓ Appreciating the practicalities for dossier updates
- ✓ Data-filling options when upgrading a registration of an intermediate to a full dossier
- ✓ Associated fees and costs

To do this we will use a combination of teaching and practical demonstrations. We have an informal style so there will be plenty of opportunity to quiz our experts throughout the day.

Programme

9:30	Coffee & Registration
	Welcome and Introduction
	Reasons for updating your dossier Change in legal status, tonnage band, new or improved information on risks, classification and labelling changes; further information at ECHA's request or spontaneous updates
	Information quality and consistency Substance identification, composition and impurities, quality of data set in terms of hazard and risk conclusions
	Reporting new uses and updating the CSR Developing new exposure scenarios, generic vs. specific, consistency with other use patterns, helping customers use your product
12:30	Lunch
	Upgrading an intermediate dossier to a full registration Data requirements, options for data-gap filling, exposure-based waiving, Risk Management Measures (RMM)
	Difficult DNELs and PNECs? What the values in the lead dossier mean for your registration and need for RMM
	Who needs to update the registration Only the Lead Registrant, only individual registrants, or all registrants?
	Practical issues and tips to help you IUCLID, REACH-IT, ECHA fees, data sharing, Letters of Access, timing, business rules, completeness checks
	Q & A
16:30	Close

