



A Short Course on Regulatory Affairs with emphasis on the
Understanding 'Small Molecule' Generic Medicines

21 - 22 July 2011

Legacy Falcon Hotel, Stratford-upon-Avon, United Kingdom

Our Experts will provide comprehensive and up-to-date training on EU pharmaceutical legislation, legal base and other official requirements and guidelines concerning appropriately-focused elements of pharmaceutical science and clinical investigation that are important to the development and licensing of generic medicines in the Europe.

This course is suitable for people (likely to be from analytical, formulation, pharmaceutical development, manufacturing and regulatory/registration departments) needing to understand how to prepare Abridged Marketing Authorisation Applications for generic medicines. It will also be useful to people wanting better to understand regulatory science and scientific writing and other skills needed for the creation of successful licensing dossiers.

EXCEPTIONAL VALUE: £995.00 + VAT High-quality, carefully-structured training, accommodation for one night (21 July)*, one breakfast, two lunches and a conference dinner (21 July), daytime in-course refreshments, a substantial delegates manual to reinforce the learning experience and a certificate of attendance

[14h Continuing Professional Development (CPD)]

* extra night(s) bookable if required, subject to availability