



# MANAGING A CTIMP

The aim of this workshop is to equip clinical research staff with an understanding of the key elements involved in conducting a clinical trial of an investigational medicinal product (CTIMPs). The workshop is relevant to those who are new to research and for those who would like to improve their understanding of clinical trials involving an investigational medicinal product.

- Define a clinical trial of an investigational medicinal product.
- Understand the role of the sponsor in commercial and non-commercial CTIMPs.
- Understand the essential documents included in a Study Site File.
- Plan a study visit in accordance with the protocol and study schedule.

**9<sup>th</sup> February 2011, 09.30 – 16.00**

For more information or to book a place contact Sandra Kane:

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