DIA EUROPE CONFERENCES AND WORKSHOPS 2013

For more information and a complete listing of all DIA events, please visit: www.diahome.org > click on Meetings & Training
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- **25th Annual EuroMeeting and Exhibition**
  4-6 March 2013 | Amsterdam, Netherlands | ID 13101
- **7th European Forum for Qualified Person for Pharmacovigilance (QPPV)**
  16-18 April 2013 I London, United Kingdom | ID 13104
- **Impact of the New Pharmacovigilance Legislation on Regulatory Affairs**
  4-5 June 2013 I London, United Kingdom | ID 13117
- **7th Annual Clinical Forum and Exhibition**
  8-9 October 2013 I Dublin, Ireland | ID 13103

In development:

- **10th Middle East Regulatory Conference - MERC 2013**
  September/October 2013 I Muscat, Oman | ID 13102
- **7th Joint DIA/EFGCP/Paediatrics Conference**
  September/October 2013 I London, United Kingdom | ID 13115
- **Statistics Workshop**
  October 2013 I Location to be confirmed | ID 13107
- **14th Conference on European Electronic Document Management (eDM) and Exhibition**
  November 2013 I Dublin, Ireland | ID 13110
- **HTA Forum**
  Autumn 2013 I Location to be confirmed | ID 13108
- **Combination Products Workshop**
  Autumn 2013 I Location to be confirmed | ID 13106

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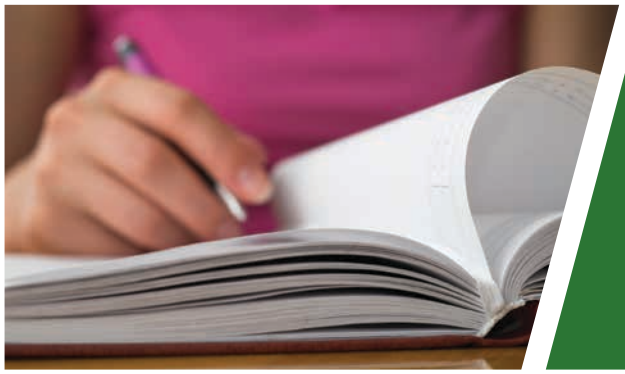

2013 TRAINING PROGRAMME DIA EUROPE

Chemistry, Manufacturing and Controls (CMC) / Quality

- **CTD Dossier Requirements: Focus on EU Module 1 and Quality Module 3**
  2013 dates and location to be confirmed
- **Quality by Design for Chemical and Biotech Products – A hands-on course for the pharmaceutical industry and regulators**
  2013 dates and location to be confirmed

Clinical Research

- **Advanced GCP Study Monitoring**
  5-6 June 2013 | Basel, Switzerland | ID 13549
- **Clinical Project Management – Part I**
  2013 dates and location to be confirmed
- **Clinical Project Management – Part II**
  20-22 February 2013 | Berlin, Germany | ID 13501
- **Clinical Statistics for Non-Statisticians**
  24-25 October 2013 | London, United Kingdom | ID 13551
- **Essentials of Clinical Study Management**
  17-19 April 2013 | Vienna, Austria | ID 13527
  20-22 November 2013 | Paris, France | ID 13554
- **Practical GCP Compliance Auditing of Trials and Systems**
  23-25 October 2013 | London, United Kingdom | ID 13548
- **Quality Risk Management (QRM)**
  Dates to be confirmed | Basel, Switzerland

Non-Clinical Safety Sciences

- **Non-Clinical Safety Sciences and Their Regulatory Aspects**
  November 2013 | Lisbon, Portugal

Regulatory Affairs

- **Authorisation of Biopharmaceuticals, Biosimilars and Advanced Therapies in Europe**
  18-20 September 2013 | Basel, Switzerland | ID 13546
- **European Regulatory Affairs: In-depth review of current registration procedures in the European Union**
  21-22 February 2013 | Berlin, Germany | ID 13525
  6-7 June 2013 | Basel, Switzerland | ID 13550
  21-22 November 2013 | Paris, France | ID 13553
- **Good Management of Medical Devices including In Vitro Diagnostics and Companion Diagnostics: Legal and practical aspects as used in personalised medicine**
  10-12 June 2013 | Amsterdam, Netherlands | ID 13547
- **Health Authority Interactions – Preparation, consultation and implementation**
  2013 dates and location to be confirmed
- **Health Technology Assessment (HTA)**
  2013 dates and location to be confirmed
- **Paediatric Investigation Plans (PIP)**
  15-16 April 2013 | Amsterdam, Netherlands | ID 13503
- **The Impact of Regulatory Affairs on Chemistry, Manufacturing & Controls (CMC)**
  5-7 June 2013 | Basel, Switzerland | ID 13532
- **US Regulatory Affairs: A comprehensive review of regulatory procedures for INDs and NDAs in the US**
  6-8 November 2013 | Paris, France | ID 13552

Safety and Pharmacovigilance

- **Benefit/Risk Management**
  13-14 May 2013 | Germany | ID 13523
  26-27 September 2013 | Prague, Czech Republic | ID 13524
- **Diagnosis and Management of Drug-Induced Liver Injury (DILI)**
  September 2013 | Paris, France
- **How to Prepare for Pharmacovigilance Audits and Inspections**
  11-12 June 2013 | Amsterdam, Netherlands
  7–8 November 2013 | Paris, France
- **Introduction to Signal Detection and Data Mining in Pharmacovigilance**
  10-11 June 2013 | Amsterdam, Netherlands
  6–7 November 2013 | Paris, France
- **Pre-Marketing Clinical Safety**
  18-19 April 2013 | Vienna, Austria | ID 13526

European Medicines Agency Information Days and Courses

- **EudraVigilance Information Day**
  17 May 2013 | London, United Kingdom | ID 13529
  22 October 2013 | London, United Kingdom | ID 13530
- **Excellence in Pharmacovigilance: Clinical trials and post-marketing**
  18-22 February 2013 | London, United Kingdom | ID 13502
  7-11 October 2013 | London, United Kingdom | ID 13522
- **IDMP International Standards ICH M5/M2 and the Implementation of eSubmission of MPIs in the EU, Article 57(2) Information Day**
  20 November 2013 | London, United Kingdom | ID 13531
- **The New Individual Case Safety Report (ICSR) International Standard and ICHE2B/M2 Information Day**
  5 February 2013 | London, United Kingdom | ID 13528
- **EudraVigilance courses:**
  EudraVigilance – Electronic reporting of ICSR
  eXtended EudraVigilance Medicinal Product Dictionary
  Introduction to Pharmacovigilance and Electronic Transmission of Individual Case Safety Reports (ICSR) for the Use of Eudravigilance at the European Medicines Agency

Courses throughout the year | European Medicines Agency, London, United Kingdom and selected European cities.

For course details on EV, please click here.

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