

Introduction to Signal Detection and Data Mining in Pharmacovigilance

Course #10558

7 October 2010

Park Inn Russell Square, London, UK



Course Faculty

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Former Qualified Person for Pharmacovigilance,
Pharmacovigilance Expert, London, United Kingdom

About the Drug Information Association

The DIA is a professional association of approximately 18,000 members worldwide who are involved in the discovery, development, regulation, surveillance or marketing of pharmaceuticals or related products. We are committed to the broad dissemination of information on the development of new medicines or generics and biosimilars, with continuously improved professional practice as the goal. The DIA is a financially independent non-profit organisation that funds itself from meeting and membership fees. The voluntary efforts of DIA members and speakers allow the DIA to organise conferences, workshops and training courses and provide publications at a reasonable, competitive cost.

**This course has limited capacity.
Register early.**

Course Overview

The World Health Organisation (WHO) defines the term *Signal* as "reported information on a possible causal relationship between an adverse event and drug, the relationship being unknown or incompletely documented previously". Adverse Drug Reactions (ADRs) may be identified as *Signals* for clinical and / or quantitative reasons. This course will cover the fundamentals of classical and statistical signal detection and data mining in Pharmacovigilance.

Who Should Attend

Professionals who work in:

- Pharmacovigilance (including QPPV)
- Clinical Development
- Risk Management
- Pharmacoepidemiology
- Information Technology
- Regulatory Affairs
- Quality and Compliance
- Legal

Learning Objectives

At the conclusion of this course, participants should be able to:

- Explain and apply the basic concepts and principles of signal detection in Pharmacovigilance
- Explain the role and differences of classical and statistical signal detection in the ongoing safety surveillance of medicinal products
- Outline how to apply signal detection within their function based on the possibilities and limitations of methodology and data
- Employ data mining techniques to analyse large volumes of adverse event report data
- Discuss key messages from the *EMA Guideline on the uses of statistical signal detection methods in the Eudravigilance Data Analysis System*

AGENDA

08:30	Registration
09:00	Start of Training Course
	Coffee Break
12:00 – 13:00	Lunch
13:00	Continuation of Training Course
	Coffee Break
17:30	End of Training Course

Session 1

**Signal Detection –
Theory, Methods and Regulatory Basis**

Session 2

Signal Detection – Application and Workshop

Session 3

Data Mining – Theory, Methods

Session 4

Data Mining – Application

Unless otherwise disclosed, DIA acknowledges that the statements made by speakers are their own opinion and not necessarily that of the organisation they represent, or that of the Drug Information Association.

Speakers and agenda are subject to change without notice. Recording of any DIA tutorial/workshop information in any type of media, is prohibited without prior written consent from DIA.

Hotel Information

The DIA has blocked a limited number of rooms at the:

Park Inn London Russell Square
92 Southampton Row
WC1B 4BH London, United Kingdom
Ph +44 207 400 3808 - Fax +44 207 400 3817
<http://www.london.russell-square.parkinn.co.uk/>

at the special rate of:
Room in single use GBP 140.00 - Room in double use GBP 150.00

These rates are per room and night and include service, taxes, VAT and buffet Breakfast.

To reserve a room, please use the hotel booking form on the DIA website or call the hotel.

IMPORTANT: To be assured of accommodation at the Park Inn Russell Square, registrants are recommended to complete their reservation by 7 September 2010 at the latest. Reservations received after that date are subject to availability.

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How you, your colleagues and company can benefit:

- Up-to-the-minute knowledge of important industry hot topics and latest developments
- Practical industry and agency case studies
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- Networking opportunities
- Limited attendance allows for a more personal quality learning experience
- Attend accredited training courses and earn continuing education credits
- Train your whole team: group discounts available

DIA UPCOMING TRAINING COURSES IN 2010

Clinical Research



Advanced GCP Study Monitoring

4 June 2010 | Prague, Czech Republic | ID 10560
19 November 2010 | Paris, France | ID 10561

Clinical Project Management in Europe – Part I

22-24 September 2010 | Basel, Switzerland | ID 10544

Clinical Statistics for Non-Statisticians

13-14 September 2010 | Paris, France | ID 10542

Essentials of Clinical Study Management

5-7 May 2010 | Vienna, Austria | ID 10527
10-12 November 2010 | Lisbon, Portugal | ID 10528

Practical GCP Compliance Auditing of Trials & Systems

6-8 October 2010 | London, United Kingdom | ID 10546

Regulatory Affairs



An Introduction to Product Information Management (PIM)

26-27 April 2010 | Vienna, Austria | ID 10541
28-29 October 2010 | Geneva, Switzerland | ID 10539

Building the eCTD

23-24 September 2010 | Basel, Switzerland | ID 10545

Comprehensive Training on European Regulatory Affairs including Different Registration Procedures and Variations: Expert Overview

4-6 October 2010 | Location to be confirmed

CTD Dossier Requirements: Focus on EU Module 1 and Quality Module 3

26-28 April 2010 | Vienna, Austria | ID 10529
5-7 December 2010 | United Arab Emirates | ID 10530

European Regulatory Affairs: Review of Current Registration Procedures in the EU

3-4 June 2010 | Prague, Czech Republic | ID 10538
18-19 November 2010 | Paris, France | ID 10540

Good Management of Medical Devices

26-28 April 2010 | Paris, France | ID 10543
27-29 October 2010 | Geneva, Switzerland | ID 10547

US Regulatory Affairs

18-21 October 2010 | Prague, Czech Republic | ID 10552

Quality by Design

Training Course is currently under development by the expert faculty: Dr. Fritz Erni and Professor Johannes Khinast

Safety and Pharmacovigilance



Excellence in Pharmacovigilance: Clinical Trials and Post Marketing

25-29 October 2010 | Vienna, Austria | ID 10533

Introduction to Signal Detection and Data Mining in Pharmacovigilance

26 April 2010 | Paris, France | ID 10550
7 October 2010 | London, United Kingdom | ID 10558

How to Prepare for Pharmacovigilance Audits and Inspections

27 April 2010 | Paris, France | ID 10551
8 October 2010 | London, United Kingdom | ID 10559

Medical Approach in Diagnosis and Management of ADRs

13-14 September 2010 | Paris, France | ID 10531

Practical Guide for Pharmacovigilance: Clinical Trials and Post Marketing

2-4 June 2010 | Prague, Czech Republic | ID 10525
1-3 December 2010 | Paris, France | ID 10526

The New Individual Case Safety Report (ICSR) International Standard and ICH E2B/M2 Information Day at the European Medicines Agency

25 June 2010 | London, United Kingdom | ID 10568

EudraVigilance Information Day at the European Medicines Agency

22 June 2010 | London, United Kingdom | ID 10534
19 October 2010 | London, United Kingdom | ID 10535

EudraVigilance (EV) and EudraVigilance Medicinal Product Dictionary (EVMPD)

Courses throughout the year | European Medicines Agency, London, UK and selected European cities
For course details on EV, please visit www.diahome.org > Training > EudraVigilance > Click on Related Courses

Non-Clinical Sciences



Non-Clinical Safety Sciences and Their Regulatory Aspects

22-26 November 2010 | Lisbon, Portugal | ID 10562

All Curricular Areas



Crisis Management

3-4 June 2010 | Basel, Switzerland | ID 10563
14-15 October 2010 | Paris, France | ID 10564

For more information and a complete listing of all training courses, please visit www.diahome.org and click on Training.

REGISTRATION FORM

Introduction to Signal Detection and Data Mining in Pharmacovigilance
7 October 2010 - Park Inn Russell Square, London, UK

ID# 10558



If DIA cannot verify your membership upon receipt of registration form, you will be charged the non-member fee. Registration fee includes course material. The fee is inclusive of lunch and coffee breaks of EUR 125.00 per day.

CATEGORY	MEMBER		NON-MEMBER (with optional membership)			NON-MEMBER (without optional membership)	
	FEE	TOTAL	FEE	Membership	TOTAL	FEE	TOTAL
Industry	€ 770.00	€ 920.92 <input type="checkbox"/>	€ 770.00	€ 115.00	€ 885.00 <input type="checkbox"/>	€ 885.00	€ 885.00 <input type="checkbox"/>
Government/Academia (Full-Time)	€ 385.00	€ 460.46 <input type="checkbox"/>	€ 385.00	€ 115.00	€ 500.00 <input type="checkbox"/>	€ 500.00	€ 500.00 <input type="checkbox"/>
TOTAL AMOUNT DUE:		€ _____	NOTE: Payment due 30 days after registration and must be paid in full by commencement of the course				

STUDENT RATES AND GROUP DISCOUNTS ARE AVAILABLE! PLEASE CONTACT DIA FOR MORE INFORMATION.

DISCOUNTS AVAILABLE IF BOTH PHARMACOVIGILANCE COURSES ARE ATTENDED ON THE 7TH AND 8TH OF OCTOBER 2010

10558DIA

RESPONSIBILITY/INTEREST AREA | Please select one Primary Interest Area (P) and one Secondary Interest Area (S) by placing a P or S on the appropriate line.

- | | | | |
|--|---|---|---|
| <input type="checkbox"/> Advertising & Promotion | <input type="checkbox"/> Medical Communications | <input type="checkbox"/> Pharmacology | <input type="checkbox"/> Regulatory Aff airs |
| <input type="checkbox"/> CMC | <input type="checkbox"/> Medical Writing | <input type="checkbox"/> Pricing/Reimbursement | <input type="checkbox"/> Research & Development |
| <input type="checkbox"/> Clinical Data Management/ eClinical | <input type="checkbox"/> Nonclinical | <input type="checkbox"/> Project Management | <input type="checkbox"/> Statistics |
| <input type="checkbox"/> Clinical Research | <input type="checkbox"/> Outsourcing | <input type="checkbox"/> Professional Education, Training & Development | <input type="checkbox"/> Strategic Planning |
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| <input type="checkbox"/> Document Management/ eSubmissions | | <input type="checkbox"/> Quality Assurance/Quality Control | |
| <input type="checkbox"/> Manufacturing | | | |

REGISTRANT

PLEASE COMPLETE IN BLOCK CAPITAL LETTERS OR MAKE REGISTRATION EVEN SIMPLER BY ATTACHING THE REGISTRANT'S BUSINESS CARD HERE

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Last Name

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Company

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Street Address / P.O. Box

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Telephone

Fax (Required for confirmation)

Email (Required to receive presentation download instructions)

Please indicate your professional category: ☐ Academia ☐ Government

☐ Industry ☐ Contract Service Organisation

PAYMENT METHODS - CREDIT CARD PAYMENT IS PREFERRED

☐ Please charge my credit card - credit card payments by VISA, Mastercard or AMEX can be made by completing the relevant details below. Please note that other types of credit card cannot be accepted.

☐ VISA ☐ MC ☐ AMEX

Card Number

Exp. Date

Cardholder's Name

Date

Cardholder's Signature

☐ Cheques should be made payable to: D.I.A. and mailed together with a copy of the registration form to facilitate identification to:

D.I.A., Elisabethenanlage 25, Postfach, 4002 Basel, Switzerland

☐ Bank transfers: When DIA completes your registration, an email will be sent to the address on the registration form with instructions on how to complete the bank transfer. Payments in EURO should be addressed to "Account Holder: DIA." including your name, company, Meeting ID# 10558 as well as the invoice number to ensure correct allocation of your payment.

Payments must be net of all charges and bank charges must be borne by the payer.

Persons under 18 are not allowed to attend DIA meetings.

CANCELLATION POLICY

Cancellations must be made in writing and be received at the DIA Europe office five working days prior to the course start

Cancellations are subject to an administrative fee: Full Meeting Cancellation: Industry (Member/Non-member) = € 200.00 - Government/Academia/Non-profit (Member/Non-member) = € 100.00

Registrants who do not cancel five working days prior to the course start date and do not attend, will be responsible for the full registration fee. DIA Europe reserves the right to alter the venue and dates if necessary. If an event is cancelled DIA Europe is not responsible for airfare, hotel or other costs incurred by registrants. Registrants are responsible for cancelling their own hotel and travel reservations.

Transfer Policy

You may transfer your registration to a colleague prior to the start of the event but membership is not transferable. Substitute registrants will be responsible for the non-member fee, if applicable. Please notify the DIA Europe office of any such substitutions as soon as possible.

IMPORTANT:

Hotel and travel reservations should be made ONLY after receipt of written registration confirmation from DIA.
If you have not received your confirmation within five working days, please contact DIA.

HOW TO REGISTER

The DIA Customer Services Team will be pleased to assist you with your registration.
Please call us on +41 61 225 51 51 from Monday to Friday between 08:00 and 17:00 CET.

Online www.diahome.org

Fax +41 61 225 51 52

Email diaeurope@diaeurope.org

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