Postmarketing Drug Safety and Pharmacovigilance

November 17-18, 2010, Horsham, PA

An In-depth Look at Practical Solutions to Postmarketing Safety and Pharmacovigilance

Learn the key tools available for pharmacovigilance, discuss their uses, and consider the future directions of the field.

WHAT YOU WILL LEARN

- How to make your pharmacovigilance program more efficient and compliant
- US, European, and other global requirements and standards in drug safety and pharmacovigilance
- Privacy, information technology, and data protection issues involved in drug safety
- How to speak the language of drug safety, signaling, risk management, and pharmacovigilance
- The future of drug safety and pharmacovigilance
- The basics of drug safety inspection

CONTINUING EDUCATION

The Drug Information Association is accredited by the Accreditation Council for Pharmacy Education as a provider of continuing pharmacy education. This program is designed for 14.5 contact hours or 1.45 continuing education units (CEUs). 286-000-10-034-L04-P

Type of Activity: Knowledge

If you would like to receive a statement of credit, you must attend the course, sign-in at the DIA registration desk each day of the course, and complete the on-line credit request process through My Transcript at www.diahome.org. Participants will be able to download a statement of credit upon successful submission of the credit request. My Transcript will be available for credit requests on November 25, 2010.

Disclosure Policy: It is Drug Information Association policy that anyone in a position to control the content of a continuing education activity must disclose to the program audience (1) any real or apparent conflict(s) of interest related to the content of their presentation and/or the educational activity, and (2) discussions of unlabeled or unapproved uses of drugs or medical devices. Faculty disclosures will be included in the course materials.

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

- Identify the legal and regulatory basis of postmarketing drug safety and pharmacovigilance
- Evaluate how your drug safety system will do on an audit or governmental inspection
- Explain the requirements, changes, pitfalls, and risks of drug safety and pharmacovigilance, applying the knowledge gained in your daily pharmacovigilance functions
- Appraise US, EU, and other countries handling of expedited reports, ICSRs, aggregate reports, safety labeling, risk management, causality determination, signaling, audits and inspections, quality, compliance, IT, basic epidemiology, and more
- Analyze your company’s drug safety systems and determine whether they are meeting the needs and requirements

WHO SHOULD ATTEND

Professionals involved in:

- Global drug safety, pharmacovigilance, and risk management
- Regulatory and legal affairs
- CROs, start-ups, small companies, generic drug companies, and anyone needing to get up to speed rapidly on the basics of drug safety and pharmacovigilance
- Training and teaching of drug safety and pharmacovigilance
- Clinical health care
- New drug development
- Outsourcing and offshoring of drug safety
- Supervising and dealing with drug safety

Faculty comprises professionals in the pharmaceutical and related industries who are experts actively practicing in their particular disciplines.

For detailed program information including faculty and topics, please contact Laura Parker at +1.215.442.6101 or Laura.Parker@diahome.org

Unless otherwise disclosed, DIA acknowledges that the statements made by speakers are their own opinion and not necessarily that of the organization they represent, or that of the Drug Information Association. Speakers and agenda are subject to change without notice. Recording of any DIA training material in any type of media is prohibited without prior written consent from DIA.
REGISTRATION FORM • Meeting ID# 10440 • Registration, limited to 50, is reserved for the first 50 registrants.

You may register online at www.diahomes.org or you may return this completed form by mail to DIA at the address below under check payment method, or by fax to +1.215.442.6199. Walk-in registration will NOT be accepted. Registration must be confirmed in writing by the DIA office. If you have not received confirmation within five business days, please telephone DIA at +1.215.442.6100 or email confirmationservices@diahomes.org.

![Image](image_url)

**POSTMARKETING DRUG SAFETY AND PHARMACOVIGILANCE**

November 17-18, 2010 • Meeting ID# 10440 • DIA Worldwide Headquarters, Horsham, PA, USA

To see if this course will be offered at other times, go to www.diahomes.org.

**TUITION/REGISTRATION FEES:** Registration fee includes continental breakfast(s), luncheon(s), reception (if applicable), and all course materials. If DIA cannot verify your membership upon receipt of this registration form, you will be charged the nonmember fee.

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* A one-year membership to DIA is available to those paying a NONMEMBER meeting registration fee. If paying a nonmember fee, please indicate if you do, or do not, want membership.

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Please check the applicable category below.

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- Industry
- CSO
- Student (Full-time, verification required)

**PAYMENT METHODS:** Register online at www.diahomes.org or check the appropriate payment method.

- CHECK drawn on a US bank payable to: Drug Information Association, mailed along with this form to: DIA, 800 Enterprise Road, Suite 200, Horsham, PA 19044-3595. Please include a copy of this registration form to facilitate identification of attendee.

- BANK TRANSFER 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. Payment should be made in US dollars. Your name and company, as well as the Meeting I.D. # must be included on the transfer document to ensure payment to your account.

- CREDIT CARD number may be faxed to: +1.215.442.6199. You may prefer to pay by check or bank transfer since non-US credit card payment will be subject to the currency conversion rate at the time of the charge.

**TRAINING COURSE LOCATION**

DIA Worldwide Headquarters
800 Enterprise Road, Suite 200
Horsham, PA 19044, USA

**HOTEL INFORMATION**

Hotel reservations should be made ONLY after receipt of written registration confirmation from DIA.

Joseph Ambler Inn
1005 Horsham Road
North Wales, PA 19454-1413, USA

Tucked away on 12 acres of rolling countryside, the Joseph Ambler Inn offers the exceptional features expected of a great country inn including an award-winning Pennsylvania restaurant and 52 pampering B&B guestrooms. Surrounded by peaceful meadows and comprising 5 historic buildings, the inn’s facilities and grounds are unmatched in all of Bucks and Montgomery counties.

Just 45 minutes from Philadelphia, the Joseph Ambler Inn is centrally located to the attractions and activities of the area. Valley Forge National Park, New Hope, and Bucks County are just a one-hour drive. Day trips can be made to Atlantic City casinos, Lancaster County Amish Country, and Reading, the shopping outlet headquarters.

A limited block of rooms has been reserved at the Joseph Ambler Inn at a low rate per night until the release date of November 1, 2010. If you are planning to stay at the hotel in order to facilitate interactive discussion with faculty and fellow participants.

Single $127 / Double $127

Attendees must make their own hotel reservations. To reserve your room, contact the Joseph Ambler Inn by telephone at +1.215.362.7500 and mention the DIA Training Course.