IPI Academy

MANAGEMENT forum



Presented by
Management Forum

Demystifying European Post-Marketing Pharmacovigilance

This course provides an overview for senior managers of the key post-market pharmacovigilance activities required to further understand the 2012 EU legislation that has undergone over 20 updates in the last 10 years.

Format:
Bespoke training

CPD:
18 hours for your records (depending on your requirements)

Certificate of completion

Course overview

This course provides a comprehensive overview for senior managers of the key post-market pharmacovigilance activities required to further understand the 2012 EU legislation that has undergone over 20 updates in the last 10 years.

The course intends to show how the safety of products is managed at the regulatory authority level, the interactions with regulatory bodies and pharma, and the internal pharma processes in managing the safety of the company products. From receiving safety cases and information to signal analysis and safety communication, all of this is viewed from a practical point of view for compliance and Key Performance Indicators (KPIs).

The European Medicines Agency (EMA) and the Pharmacovigilance Risk Assessment Committee (PRAC) deliberations and the UK Brexit implications will be discussed.

Benefits of attending

- **Explore** how the safety of products is managed at the regulatory authority level, the interactions with regulatory bodies and pharma
- **Learn** the internal pharma processes in managing the safety of the company products from receiving safety cases/information, all the way through to signal analysis and safety communication
- **Discuss** The European Medicines Agency (EMA) and the Pharmacovigilance Risk Assessment Committee (PRAC) deliberations and Brexit implications

Who should attend?

This three-day course is intended for senior management professionals from European and US pharmaceutical companies, who need to understand EU Pharmacovigilance. It will be particularly beneficial for those working in allied technical areas (regulatory, clinical, QA, and auditing) who require an overview of key pharmacovigilance activities and also need to understand the many complexities of EU pharmacovigilance, including the UK Brexit implications.

All of these activities are discussed in relation to Quality Management Systems (QMS), audits and inspections.

Reviews

★★★★☆

Graeme was a comfortable speaker that was open to questions throughout his presentations which was very positive.

Robin van Osch
Junior pharmacovigilance officer
Interdos
Jul 15 2024

Run this programme in-house for your whole team

Coming to IPI Academy for your in-house training provides an all-inclusive service which gives you access to a wide variety of content, learning platforms and delivery mechanisms as well as your own personal training adviser who will work with you from the initial enquiry through to feedback and follow-up after the programme.

With over 600 trainers, all practitioners and experts across a huge range of fields, we can provide the training you need, where you need it, when you need it, and at a price which suits your budget. Our approach to tailored learning and development consists of designing and delivering the appropriate solution for each client.

For your FREE consultation and to find out more about how we can work with you to solve your training needs, please contact our training advisers:



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IPI Academy

IPI Academy is a training initiative of Falconbury and Management Forum; leading providers of industry training for over 30 years, based in the UK.

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