



Presented by
Management Forum

Introduction to Veterinary Pharmacovigilance

3-4 July 2025
+ 5-6 November 2025

A basic training course for those working on drug safety in the EU. This course has been designed to provide basic training and a good introduction to those concerned with veterinary pharmacovigilance.



Format:
Live online



CPD:
12 hours for your records



Certificate of completion

Course overview

This two-day event has been designed to provide an essential overview of veterinary pharmacovigilance and will offer practical guidance and basic training for all those working in drug safety in the EU.

Our experienced trainer will clarify roles and responsibilities, explain commonly used terminology and take participants through all the key aspects of this complex subject. There will be plenty of time for interaction and questions and answers to enable participants to get a good understanding of the issues involved. The use of industry case studies will demonstrate real-life scenarios to help embed learning.

Benefits of attending

- **Gain** an overview of the European regulatory framework
- **Understand** the requirements of the new EU pharmacovigilance legislation
- **Learn** about VICH
- **Understand** adverse event reporting
- **Hear about** causality assessment
- **Minimise** the impact of data with errors
- **Get to grips** with literature searches
- **Understand** UK PV requirements post Brexit

Who should attend?

The course will be beneficial to those new to veterinary pharmacovigilance, support staff and experienced personnel who require a better understanding of drug safety in their current role. Adverse event monitoring and drug safety officers, together with regulatory affairs and personnel from registration departments, will also find this seminar useful.

Programme

Day 1

What is pharmacovigilance?

- Beneficial and harmful effects of veterinary medicinal products
- Key definitions

The current regulatory framework and its global impact

- Overview of European regulatory framework, the EU pharmacovigilance legislation
- Implications for global environment – link to VICH
- Practical applications of definitions
- Role of the MAH
- Role of the NCA/EMA
- Role of the QPPV

Adverse event reporting

- Definitions
- Impact of VICH guidelines
- Expedited vs periodic
- How to handle animal SARs
- Handling human SARs
- Understanding the wider scope of pharmacovigilance

Causality assessment

- The principles of causality assessment with practical examples
- Medical evaluation of individual reports of adverse events
- Strategies for follow-up

Identifiable Reporter

Day 2

Do vets report AEs and why?

Electronic communication in pharmacovigilance

- Reporting in EV Vet
- VEDDRA

EV Vet demo

Minimising the impact of data with errors

- Consistent assessment and coding

Clinical trial AE reporting requirements

- Post-authorisation safety studies
- Phase IV studies

Literature searches

- Peer-reviewed worldwide literature
- Local journals and magazines

PSURs

- VMD (UK), VDD (Canada) and APVMA (Australia) requirements
- Format and content of the PSUR
- Analysis of data
- Incidence calculation
- Compliance and the PSUR

Presenter



Declan O'Rourke

Declan O'Rourke has over 20 years' experience in industry where he has held technical, marketing, product development, clinical development, production and pharmacovigilance roles.

He is a veterinary surgeon, holds a Diploma in Marketing, a Master of Business Administration and a Fellowship of the Royal College of Veterinary Surgeons. He now directs Ortec PV Consultancy Ireland specialising in pharmacovigilance and represented IFAH-EU in the VICH Working Group on pharmacovigilance.

He is Honorary Associate Professor in Veterinary Pharmaceutical Development at Nottingham Veterinary School and Past President of British Cattle Veterinary Association.

Course dates

3-4 July 2025

Live online

09:00-16:30 **UK (London)** (UTC+01)

Course code 14777

GBP **1,299** ~~1,499~~

EUR **1,819** ~~2,099~~

USD **2,087** ~~2,399~~

Until 29 May

5-6 November 2025

Live online

09:00-16:30 **UK (London)** (UTC+00)

Course code 14960

GBP **1,299** ~~1,499~~

EUR **1,819** ~~2,099~~

USD **2,087** ~~2,399~~

Until 01 Oct

How to book



Online:

ipi.academy/2211

Alternatively contact us to book, or if you have any queries:



Email:

info@ipiacademy.com



Phone:

[+44 \(0\)20 7749 4749](tel:+442077494749)

Discounts

- Booking more than one delegate on any one date qualifies for a **15% discount** on the second and subsequent places.
- Most events qualify for an **early booking discount** prior to 6 weeks before the course date. Be sure to check on our website, where the latest discounts will be shown.

Further information

Fee

The fee includes all meals and refreshments for the duration of the course (for venue-based courses) and a complete set of course materials (provided electronically). If you have any particular requirements, please advise customer services when booking.

Please note

IPI Academy (and our training partners) reserve the right to change the content and timing of the programme, the speakers, the date and venue due to reasons beyond their control. In the unlikely event that the course is cancelled, we will refund the registration fee and disclaim any further liability.

Terms and conditions

The rest of our terms, the event cancellation policy and the terms and conditions are on our website, please visit ipi.academy/content/terms-and-conditions

Reviews



Excellent! [Speakers were] very knowledgeable with clear experience in a broad range of PV roles.



Fiona MacGillivray
Director and Owner
MacVet Limited
Oct 2 2024



The webinar was great!!!



Miguel Angel Ramirez Arredondo
Regulatory Affairs Manager
Sanfer Salud Animal
Jun 26 2024



The speaker was very knowledgeable, and his course was quite diverse, handling EU and UK topics, but also zoomed in a bit on USA market. I feel more confident handling PV.



Jeroen Lievens
Quality & Regulatory Manager
Vetoquinol
Oct 2 2024



I enjoyed all parts of the webinar. Sharing of information throughout made the webinar engaging.



Alesia Mitchell
Associate Director, Global AH PV and RA Quality Standard
Boehringer Ingelheim Animal Health
Jun 26 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:



ALEKSANDRA BEER
Tel: +44 (0)20 7749 4749
Email: inhouse@ipiacademy.com



YESIM NURKO
Tel: +44 (0)20 7749 4749
Email: inhouse@ipiacademy.com



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.

10-12 Rivington Street
London EC2A 3DU

ipi.academy
Tel: +44 (0)20 7749 4749
Email: info@ipiacademy.com