IPI Academy

MANAGEMENT forum

Presented by
Management Forum

Advanced Veterinary Pharmacovigilance

15-16 October 2025
+ 21-22 May 2026, 4-5 November 2026

Including the main requirements of Volume IXB, an update on the proposed EU Regulation (2019/6) on Veterinary Medicinal Products and the impact of Brexit.

Format:
Live online

CPD:
12 hours for your records

Certificate of completion

Course overview

The purpose of this seminar is to provide a comprehensive, yet practical, assessment of the requirements of EU Regulation (2019/6) on Veterinary Medicinal Products and Veterinary good pharmacovigilance practices (VGVP). The programme will also consider The Veterinary Medicines (Amendment etc.) Regulations 2024.

Interactive sessions throughout the two days will give delegates the opportunity to discuss key issues of current concern with our experienced trainer.

Benefits of attending

- **Explore** the European regulatory framework and the implications of the new EU pharmacovigilance legislation
- **Learn** the key requirements of VGVP and the pharmacovigilance system master file (PSMF)
- **Gain** PV training and PV reporting in licensing/distribution agreements
- **Understand** product safety reviews, signal detection and benefit-risk assessments
- **Look** at risk management and crisis management

Who should attend?

This seminar will be beneficial to those who have some experience of veterinary pharmacovigilance and is a good follow-on course from the Introduction to Veterinary Pharmacovigilance, which is also run by Management Forum. Adverse event monitoring and drug safety officers, including QPPVs and deputy QPPVs, together with personnel from regulatory affairs and registration departments, will find this event useful.

The course can be used as part of the training to become a QPPV or deputy QPPV.

Programme

Day 1

The DDPS and the PSMF

- The DDPS
- The DDPS – what happens now?
- The PSMF – purpose and maintenance
- Transition from DDPS to PSMF

PV training

- Requirements for PV training
- Who trains whom and when?
- Training records, maintenance and updates
- Role of QA and HR in training

Product safety reviews

- The safety review committee
- Timings for safety review
- Record keeping for safety review meetings

Company core safety information

- Core safety information
- How to determine what to include and what to exclude in a company core data sheet (CCDS)
- Maintenance and development of a CCDS

Compliance and PV

- Basic principles
- Measuring compliance
- Quality versus quantity in safety reports

Day 2

PV reporting in licensing/distribution agreements

- What types of agreements exist?
- Audits of pharmacovigilance capabilities in partners
- What agreements need to be in place for PV reporting?
- PV reporting agreements – what needs to be covered?
- Monitoring PV agreements – what happens if it goes wrong?

Signal detection/benefit-risk and risk management

- The pharmacovigilance system
- Signal detection and analysis: what is required?
- Benefit-risk assessments
- Risk management/minimisation
- Crisis management
- Communication with the public

Audits and inspections: are you ready?

- What are inspectors looking for?
- Inspection findings and outcomes
- Implications of the proposed EU pharmacovigilance legislation

Risk management dealing with an alert

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

15-16 October 2025	**Live online** 09:30-17:00 **UK (London)** (UTC+01) *Course code 14991*	GBP **1,299** ~~1,499~~ EUR **1,819** ~~2,099~~ USD **2,087** ~~2,399~~ **Until 10 Sep**
21-22 May 2026	**Live online** 09:30-17:00 **UK (London)** (UTC+01) *Course code 16134*	GBP **1,299** ~~1,499~~ EUR **1,819** ~~2,099~~ USD **2,087** ~~2,399~~ **Until 16 Apr**
4-5 November 2026	**Live online** 09:30-17:00 **UK (London)** (UTC+00) *Course code 16425*	GBP **1,299** ~~1,499~~ EUR **1,819** ~~2,099~~ USD **2,087** ~~2,399~~ **Until 30 Sep**

How to book

Online:
ipi.academy/2235

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

Email:
info@ipiacademy.com

Phone:
+44 (0)20 7749 4749

Discounts

- Booking more than one delegate on any one date qualifies for a **30% 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 the our terms, the event cancellation policy and the terms and conditions are on our website, please visit ipi.academy/content/terms-and-conditions

Reviews

★★★★★

Overall really enjoyed the webinar, found it very informative and useful. Would recommend.

Nikkita Evans
Senior Pharmacovigilance Safety Scientist
Propharma Group
Oct 9 2024

★★★★★

I participated to the course in order to verify if our way of working is aligned with the expert one. I also get some intersting different approach that for sure I'll apply in my system.

Leonardo Giraudo
QPPV, PV compliance expert
Pharma Quality Europe Srl
Oct 9 2024

★★★★★

Excellent webinar, very informative! This seminar exceeded my expectations!

Stella Giorgoudelli
Regulatory Manager
Hellafarm SA
Oct 9 2024

★★★★★

Thank you for the clear overview of how to maintain or set up a good PV system. Very good webinar, enough breaks in between presentations. I really thought the webinar was very accessible and [I] was able to ask all of my questions. Very complete programme. Useful webinar!

Sanne de Jong
deputy QPPV
Dopharma
Oct 9 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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