



Presented by
Management Forum

A Practical Guide to Producing and Maintaining the PSMF

3 October 2025

This course provides a practical guide to both producing and maintaining the PSMF to ensure compliance and has been fully revised to include the latest updates.



Format:
Live online



CPD:
6 hours for your records



Certificate of
completion

Course overview

The pharmacovigilance system master file (PSMF) is a legal requirement for any medicinal product authorised in the European Union. The PSMF provides the regulators with a detailed description and assessment of the entire pharmacovigilance system and the outputs contained in the annexes provide an understanding of a company's compliance. This course will provide a practical guide to planning, writing, maintaining and updating the PSMF to ensure compliance. The programme will cover the importance of the PSMF in regulatory inspections, including common PSMF inspection findings. You will discuss the processes and systems required to manage the PSMF as well as the latest advice on the impact of Brexit.

Benefits of attending

- **Understand** the regulatory requirements for the PSMF
- **Gain** an overview of the key issues in producing, maintaining and updating the PSMF
- **Discuss** the PSMF as a QMS document
- **Learn** about the roles of the QPPV and the PSMF
- **Review** common inspection findings and deficiencies related to the PSMF

Who should attend?

This course will be relevant for anyone working in pharmacovigilance who requires a comprehensive overview of the PSMF, including QPPVs and those responsible for safety assessments. It will also be of interest to those who work with pharmacovigilance, eg in regulatory affairs, clinical, sales and marketing, legal, commercial and quality, as well as the audit group.

Programme

Introduction and background to the PSMF

- The DDPS and the PSMF
- Objectives of the PSMF
- Registration of the PSMF
- Regulatory requirements and accessibility of the PSMF
- Responsibilities of the marketing authorisation holder, updates and the EU QPPV

The content of the PSMF

- The PSMF template
- The level of detail required by the PSMF
- Preparation of the annexes
- The PSMF log book

The sections of the PSMF

- The EU QPPV
- Sources of safety data
- IT and databases
- Regulatory timeline compliance
- The PSMF processes
- The PSMF and audits
- The company quality system and the PSMF

The annex requirements for the PSMF

- The company product list
- The EU QPPV list of delegated tasks
- The list of SOPs and procedures
- List of delegated activities to third-party partners
- A list of completed audits and schedules
- A list of performance indicators for the PSMF section
- The roles and responsibilities of the EU QPPV
- Master file number and version changes (audit trail)

The PSMF and inspections

- The PSMF and inspection findings
- Regulatory authority requests to view the PSMF
- Transfer of responsibility for a pharmacovigilance system to the QPPV
- Notifying the QPPV of changes to the PSMF
- PSMF responsibilities with shared marketing authorisation holders
- Change control, logbook, versions and archiving
- Audit trails and the PSMF
- The PSMF post-inspection

Final discussion session

Presenter



Graeme Ladds

Graeme Ladds, Director of PharSafer, has over 30 years' experience working in the pharmaceutical industry. Having started his career at Ashbourne Pharmaceuticals in 1989 as Head of Drug Safety & Medical Information, he went on to become Head of Global Pharmacovigilance at Shire Pharmaceuticals. He then set up his consultancy and specialist CRO company, PharSafer Associates Ltd, where he has been involved in establishing pharmacovigilance in companies, performing audits across Europe and the USA, SOP writing, acting as QP for companies, and helping with regulatory inspections.

Course date

3 October 2025

Live online

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

Course code 14994

GBP ~~649 749~~

EUR ~~909 1,049~~

USD ~~1,043 1,199~~

Until 29 Aug

How to book



Online:

ipi.academy/2194

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



Email:

info@ipiacademy.com



Phone:

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

Reviews



Very good seminar which covered all important points around the production and maintenance of the PSMF. The speaker was excellent with a profound knowledge of the topic.



Marie-Christine Klös
Global PV Specialist
Boehringer Ingelheim Vetmedica GmbH
Feb 14 2025



Informative speaker, really good [content] and thoroughly enjoyed the course.



Dominic O'Hare
Associate Director
Norgine
Aug 5 2024



In general, I was very satisfied with this webinar, I learned a lot.



Jasmina Roca
QPPV deputy
Isdin
Apr 24 2024



Excellent presenters and excellent presentations



Irina Dukovska Teova
Pharmacovigilance specialist
Alkaloid AD Skopje
May 17 2023

Run this programme in-house for your whole team

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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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