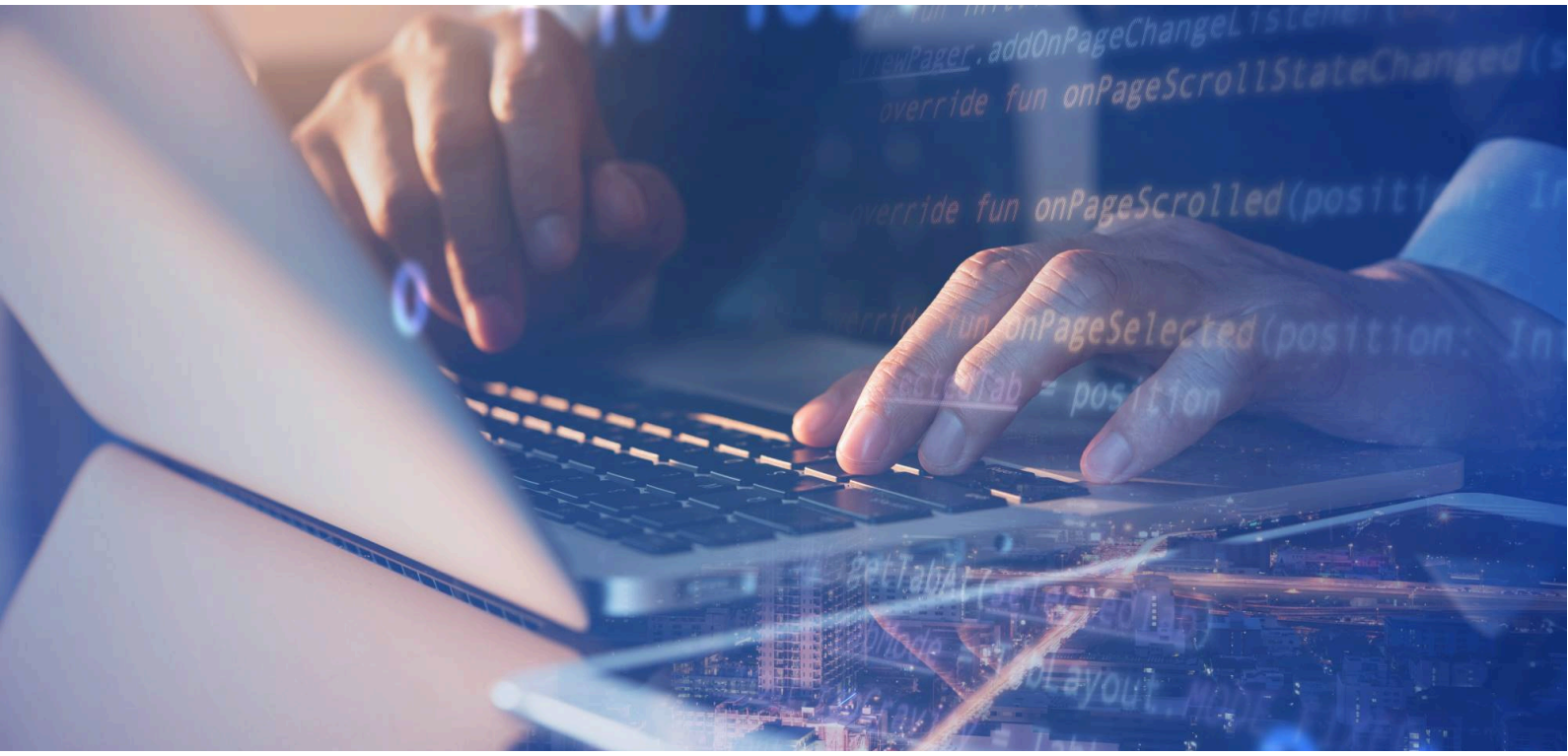
IPI Academy

MANAGEMENT forum



Presented by
Management Forum

Mastering Computer System Validation

16-17 June 27
+ 9-10 December 27

This course covers the key concepts of computer systems validation and provides practical guidance on how to comply with the regulations applicable to the pharmaceutical industry.

Format:
Live online

CPD:
12 hours for your records

Certificate of completion

Course overview

Ensure your computerised systems are fully compliant, reliable, and designed for long-term success. Mastering Computer System Validation provides a deep dive into the critical discipline that underpins trust in digital pharmaceutical operations. This course moves beyond simple testing to show how a well-structured validation lifecycle proves that your systems perform as intended, every time. You will learn how to establish and maintain systems that generate accurate data, maintain robust audit trails, and secure electronic records for the lifetime of your products.

Regulatory expectations have evolved, and so must your approach. In an era of increasing scrutiny, inadequate validation can lead not only to data integrity breaches but also to costly compliance failures and regulatory findings. Through this course, you will understand how to interpret and apply the principles of EU Annex 11, FDA 21 CFR Part 11, and GAMP®5 with confidence and clarity.

This training transforms CSV from a checkbox exercise into a strategic enabler. You'll explore how validation supports operational excellence, business continuity, and digital transformation initiatives from traditional on-premise systems to modern AI driven and cloud-based platforms. Real world case studies and interactive exercises ensure the lessons translate directly to your workplace.

By the end of the course, you will have mastered a practical, risk-based framework for computerised system validation that aligns with both regulatory requirements and business objectives. Most importantly, you'll gain the assurance and authority to lead validation activities that withstand inspection, safeguard data integrity, and inspire confidence across your organisation.

Benefits of attending

- **Gain** a comprehensive overview of computer system validation (CSV)
- **Enhance** your understanding of the key terms used in CSV
- **Recognise** how Good Practices apply to computerised systems
- **Understand** the different approaches which might be employed
- **Learn** how to manage the risk associated with data integrity
- **Master Regulatory Frameworks** - Gain expert-level understanding of EU Annex 11, FDA 21 CFR Part 11, and GAMP 5 principles
- **Practical Risk-Based Approach** - Learn to streamline validation without compromising compliance
- **Inspection Readiness** - Develop documentation and strategies that satisfy inspectors globally
- **Future-Proof Skills** - Understand CSV in the context of AI, SaaS, and data integrity challenges
- **Real-World Application** - Apply your knowledge to authentic case studies from pharmaceutical operations

Who should attend

- Quality management specialists
- Pharmaceutical validation professionals
- Pharmaceutical project managers
- Computer system engineering specialists

Presenters



Aneta Jell

Aneta Jell is a seasoned Quality Assurance professional with over 20 years of experience across manufacturing, contract operations, and quality leadership in the pharmaceutical industry. Currently serving as Quality and training manager at PaulPalmer limited she oversees global contract manufacturing quality, ensuring GxP compliance, regulatory alignment, and supply chain integrity. Her expertise spans GMP, ISO 9001, ISO 17025, and QMS administration, with proven strength in audits, CAPA, deviations, and technical agreements. A qualified QMS lead auditor with bilingual proficiency in English and Slovak, Aneta is recognised for driving continuous improvement, fostering compliance culture, and leading high-performing quality and HSE teams.



Farah Nadeem

Farah Nadeem is an Operations Manager at Paul R. Palmer Limited/ Pharmaceutical Consultant and a Trainee Qualified Person. Farah is a Pharmacist and recognised QA expert and has over 20 years of experience in the Pharmaceutical industry in the development, manufacture and supply of medicinal products.

Throughout her career, she has held various quality assurance roles and actively pursued new opportunities to expand her knowledge and expertise. As a result, she has gained a comprehensive understanding of manufacturing, storage, and distribution processes, as well as the facilities and services that support them. Involved in documentation management: writes, reviews, and audits SOPs and QMS implementations. Assists companies in meeting regulatory requirements.

In her current role as a QA Operations Manager for a Pharmaceutical Consultant Company, Farah Nadeem provides various solutions, including assistance in obtaining the first MIA(IMP) and MIA licenses, as well as preparation and hosting for regulatory inspections. Her work involves collaborating with Paul Palmer to offer services such as QP declarations for API manufacturers, regulatory inspection hosting, commercial and clinical batch release, compliance audits, and coaching and training in GxP.

Course dates

16-17 June 27	**Live online** 09:30-17:30 **UK (London)** (UTC+01) *Course code 17241*	GBP **1,299** ~~1,499~~ EUR **1,819** ~~2,099~~ USD **2,087** ~~2,399~~ **Until 21 Apr**
9-10 December 27	**Live online** 09:30-17:30 **UK (London)** (UTC+00) *Course code 17579*	GBP **1,299** ~~1,499~~ EUR **1,819** ~~2,099~~ USD **2,087** ~~2,399~~ **Until 14 Oct**

How to book

Online:
ipi.academy/2628

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

★★★★★

All good. Very experienced and knowledgeable speaker, explained every aspect in detail and made simpler.

Pramod Patil
Validation Specialist
MeiraGTx
Jul 9 2025

★★★★☆

Overall, the webinar was good. The content was informative, the presentation was clear.

Pavan Shetty
Senior Validation Specialist
Symbiosis
Nov 24 2025

★★★★☆

The session was highly interactive, and the speaker was very open to questions, providing detailed explanations and insights. Their willingness to dive deep into the topics made the session both engaging and informative.

Jayamol Thomas
Quality Manager
Moonlake
Jul 9 2025

★★★★★

Overall, the course was really good, the speakers were of a very high standard and knew their stuff super well. This really shone through their examples and diverse experience in pharmaceuticals.

Yasmeen Scotto
QA Validation Associate
Catalent Micron Technologies Limited
Mar 19 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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