





Presented by Management Forum

Effective Deviation Management Systems in Pharma

26 September 2025 + 20 March 2026

Master pharmaceutical deviation management in just 1 hour. Learn root cause analysis, CAPA processes, and compliance requirements to ensure product quality and safety.



Format:

Live online

(1)

CPD:

1 hours for your records

(j)

Certificate of completion

Course overview

Managing deviations effectively is a key component of a robust quality management system and supports the continuous improvement of pharmaceutical processes and product performance.

Occurring at any stage of a pharmaceutical product's lifecycle - development, manufacturing, or supply operations - deviations must be properly documented, investigated, and analysed to identify root causes and prevent recurrence. Failure to manage deviations adequately can pose risks to quality, compliance, safety, and the efficacy of pharmaceutical products intended to save lives.

This course will explore the crucial steps involved in the overall management of deviations in the pharmaceutical industry.

Benefits of attending

- Know the fundamentals of deviations management
- Understand the steps and tools involved in root cause analysis and Corrective and Preventive Action (CAPA)
- Learn about of the different components of a good deviation report
- Be aware of the risks involved in the inadequate deviation management

Who should attend

This course is ideal for professionals in the following industries:

- Pharma
- Biotech
- Medical Devices
- Life Sciences (particularly those involved in handling deviations)

Additionally, this course is well-suited for professionals in pharma and related industries who want to understand the fundamentals of deviation management and its impact on both products and business. It is also beneficial for anyone interested in staying updated on the latest regulations and industry trends affecting pharmaceutical deviation management.

Programme

Introduction to deviations

- Basics and examples
- Industry best practices

Deviation management system

- Steps and tools in the deviation investigation
- Corrective and Preventive Action (CAPA) process

Industry regulations and trends

- Health authority requirements
- Compliance risks and challenges



Presenter



Gurpal Singh

Dr. Gurpal Singh is an experienced Pharmaceutical Quality Management professional with more than 25 years of experience and a history of success in QA operations management, Quality systems implementation and Continuous improvement. He previously worked in global quality assurance roles within large pharmaceutical companies like MSD and Novartis where he led various quality projects and QA initiatives ensuring compliance with international GxP and regulatory requirements. Gurpal holds a doctorate in Chemistry from Magadh University of India and holds professional certificates in Six Sigma, Quality Auditing and Business Management. He is also a member of International Society of Pharmaceutical Engineering.

Course dates

26 September 2025

Live online

12:00-13:00 **UK (London)** (UTC+01)

Course code 16643

GBP 99 124

EUR **139** 174

USD 160 199

Until 22 Aug

20 March 2026

Live online

12:00-13:00 **UK (London)** (UTC+00)

Course code 16644

GBP 99 124

EUR **139** 174

USD 160 199

Until 13 Feb

How to book



Online:

ipi.academy/3246

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

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

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