IPI Academy

MANAGEMENT forum



Presented by
Management Forum

The Latest Updates in Global GMP

In this 90-minute webinar, you will gain an insight into the latest updates in GMP, as well as consolidating your existing knowledge on the subject.

Format:
Bespoke training

CPD:
1.5 hours for your records (depending on your requirements)

Certificate of completion

Course overview

Many pharmaceutical manufacturers tend to describe GMP practices as the rules personnel must follow during production and the recording of production steps. However, GMP is a comprehensive set of standards in which every step of all processes, including pharmaceutical production and R&D activities throughout the product life cycle, is validated and documented.

Historical GMP failures have caused significant tragedies for many companies. Today, any country that prioritises patient health and safety permits the production, import, and sale of pharmaceutical products only in compliance with internationally recognised current GMP standards. GMP (Good Manufacturing Practices) is often referenced as cGMP (current Good Manufacturing Practices) in alignment with FDA terminology. The 'c' in cGMP stands for 'current,' emphasising that these rules are continuously updated and dynamic, rather than following a 'produce what is available' approach. Health authorities remind manufacturers that staying up-to-date is a regulatory requirement.

This distinguished GMP Update training covers updates across all sections of the GMP Guide, incorporating innovations from EMA, PIC/S, FDA, WHO, ANVISA, MHRA, and other regulatory bodies. Participants will also explore the latest developments in concept papers, draft articles, and international GMP guidance, ensuring a complete and current understanding of global GMP expectations.

Benefits of attending

- **Gain** the advantage of refreshing your existing GMP knowledge and reinforcing key concepts
- **Learn** about new developments and topics often missed in more superficial GMP trainings
- **Discover** insights relevant not only to pharmaceutical and biotechnology companies but also to veterinary medical product manufacturers, suppliers, and medical device manufacturers
- **Apply** updated guidance and best practices directly to your work, ensuring compliance with the latest international standards
- **Understand** how evolving GMP regulations impact different sectors across the product life cycle, from R&D to production

Who should attend

- Quality Assurance
- Quality Control
- GMP Compliance
- Manufacturing
- Supply Chain/ Logistics
- Regulatory Affairs
- Engineering
- IT

This course will also be of interest to:

- Regulatory Authority Inspectors / Auditors (Human and Veterinary Medicinal Products)
- Pharmacy, chemistry, biopharmaceutical sciences, veterinary science, engineering students and new graduates

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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