IPI Academy

MANAGEMENT forum

Presented by
Management Forum

The Latest Updates in Global GMP

9 July 2024

In this free 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:
Live online

CPD:
1.5 hours for your records

Certificate of completion

Course overview

This webinar will provide the latest GMP updates which are supported by the following health regulatory authorities: EMA, PIC/S, FDA, WHO, ANVISA, MHRA, etc. and will address all sections that will be revised in the GMP Guide. Concept papers, draft articles, and the latest developments in the International GMP Guidelines will also be discussed.

Many pharmaceutical manufacturers tend to describe GMP practices as the rules that personnel must comply with throughout production and the recording of production steps. However, GMP is a comprehensive set of rules in which the steps to be followed in all processes, including the production of pharmaceutical products and R&D activities throughout the product life cycle, are validated and recorded.

GMP disasters throughout history have caused indelible tragedies in many companies. Every country that considers patient health and safety as the priority allows the production, import, and sale of pharmaceutical products per the internationally recognised current GMP rules. GMP, regularly appears as cGMP (current Good Manufacturing Practices) and is encouraged and supported by the regulatory authorities who often remind the manufacturer that they must constantly stay up-to-date.

This webinar will provide an essential update on the latest changes and ensure you are cGMP ready.

Benefits of attending

- **Refresh** your existing knowledge surrounding GMP practices
- **Learn** the latest updates in cGMP, including guidance from the EMA, PIC/S, FDA, WHO, ANVISA and MHRA
- **Explore** the latest innovations in the concept papers, draft articles and the latest developments in the International GMP Guides

Who should attend

- Quality Assurance Personnel
- Quality Control Personnel
- GMP Compliance Personnel
- Manufacturing Personnel
- Supply Chain/ Logistics Personnel
- Regulatory Affairs Personnel
- Engineering Personnel
- IT Personnel
- Regulatory Authority Inspectors/Auditors (Human and Veterinary Medicinal Products)

Programme

Brief History of GMP

APIC, USP, ICH Updates

PIC/S, FDA, EMA, MHRA, WHO and Other Health Authority Updates

Q&A Session

Presenter

Mustafa Edik

Mustafa Edik is an Independent GMP Consultant and Auditor.

After graduating as a Chemist from university, Mustafa began his 25 year plus career as a Laboratory Supervisor at Bayer, a German Pharmaceutical Company. After 15 years of working as a Quality Assurance Assistant Manager, Laboratory Supervisor, Pharmaceutical Quality Management Systems, and GMP Lead Auditor, he decided to continue his career as a Consultant. He has served the Turkish Atomic Energy Authority (TAEA) as Principal GMP Auditor and Consultant for 6 years. TAEA was audited by the Republic of Turkey Ministry of Health and granted GMP Certificate for 5 Radiopharmaceuticals. This success has won great acclaim from all health authorities and industry.

He has prepared and presented various training courses and workshops to more than 8000 individuals from 150 International and local Pharmaceutical, Medical Device, and Cosmetics companies on GMP, GDP and Pharmaceutical Quality Management Systems. He has taken part in several International Pharmaceutical Facility Establishment projects as GMP Consultant and has also set up various Quality Management Systems for Local Pharmaceutical and Medical Device Companies.

While he was the Vice President of Quality and Technical Operations at a Quality Academia Training and Consultancy firm, he acquired and converted it into a 100 % Turkish Company. As the only IRCA Certificated Pharmaceutical Quality Management Systems and GMP Lead Auditor in Turkey, he currently conducts API, Excipient, Packaging Materials Suppliers and Manufacturers, Third Party Logistics Service Providers, Sterile and Non-Sterile Manufacturing Facilities Audits according to FDA, EMA, PIC /S, TMMDA, MHRA, TGA Health Canada, and WHO regulations and guidelines.

He finished his second university degree in Biopharmaceutical Sciences BSc (Hons) at Atlantic Technological University - Ireland. He is the author of chapter 6 of the book published by PDA named “Good Distribution Practices” and his new book on ‘GMP Audits in Pharmaceutical and Biotechnology Industires’ will be published by Taylor & Francis in June 28, 2024.

Course date

9 July 2024

Live online
14:00-15:30 **UK (London)** (UTC+01)
Course code 15192

How to book

Online:
ipi.academy/2864

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

Email:
info@ipi.academy

Phone:
+44 (0)20 7749 4749

Discounts

- Booking more than one delegate on any one date qualifies for a **15% 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

Run this programme in-house for your whole team

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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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