



Speakers



Dr Heinrich Prinz
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GMP for Beginners

Understanding the importance of GMP



Live Online Training on 19/20 October 2021



Highlights

- GMP History & Trends
- Basic Principles of GMP
 - Personnel
 - Hygiene
 - Premises / Production
 - Documentation
 - Risk management
 - Qualification / Validation
 - Communication with clients/authorities
- Elements of a QA System
 - Change Control
 - Deviations
 - CAPA (Corrective Actions – Preventive Actions)
 - Failure Investigations
 - OOS (Out of Specification)
 - Audits – Inspections
 - Falsified Products


Objective


The Online Training is designed for people who have no or little knowledge of GMP.

- You get to know the most important pharmaceutical regulations and their importance,
- you get a basic overview of GMP requirements in pharmaceutical production, and
- you become familiar with technical terms from the field of GMP and their meaning

Background

In the manufacture and quality control of medicinal products, compliance with the GMP rules is the decisive aspect for manufacturing high-quality products. For this reason, every staff member in the pharmaceutical industry has to be familiar with the basic GMP requirements. The relevant European GMP regulations define the following prerequisites:

 **Commission directive 2003/94/EC**
The personnel shall receive initial and ongoing training, the effectiveness of which shall be verified, covering in particular the theory and application of the concept of quality assurance and good manufacturing practice

 **EudraLex Vol. 4 Good manufacturing practice (GMP) guidelines**
2.9 Besides the basic training on the theory and practice of Good Manufacturing Practice, newly recruited personnel should receive training appropriate to the duties assigned to them.....

In practice, many members of staff are often unaware of the contents and meaning of the different GMP requirements from Europe and US and their consequences for product quality. During this Online Training, speakers with long-standing experience in the training of employees introduce and explain the most important elements of a pharmaceutical GMP system in an easy-to-understand way.

Target Audience

The Online Training is directed to staff from the pharmaceutical industry having no or little experience with the current GMP requirements. This includes both employees who, after their vocational training, enter the pharmaceutical industry and experienced colleagues who work for the first time in a GMP-regulated environment. Participation is also recommended for personnel from suppliers who have to understand the quality requirements of their customers.



To ensure a high quality transmission the presentations of this online training were recorded in advance.
Two discussion panels: The online training includes live Q&A sessions with the speakers (each about ½ hour in the morning and at the end of the event days). This provides the opportunity to ask questions, which the speakers will then answer.

Programme 19 October 2021

Welcome & Introduction 09.00 – 09.15 h

GMP History & Trends 09.15 – 10.15 h

- Development of GMPs
- GMP: Goal and general ideas
- Types of regulatory documents and their meaning
- The Dossier
- GMP for APIs, medicinal products and excipients - a comparison
- Authorities and industry groups: ICH, PIC/S, EMA, FDA, WHO, APIC, ISPE, IPEC
- Ph. Eur. & USP
- GMP in the US: Comparison of US and EU regulations
- Differences between European and FDA view on GMP / GMP vs cGMP
- Typical expectations of FDA and European inspectors

Break 10.15 – 10.30 h

Hygiene / Personnel Hygiene 10.30 – 11.15 h

- General aspects and rules
- Hygiene program
- Personnel flow
- Medical examination
- Contamination
- Monitoring

Personnel and Training 11.15 – 12.00 h

- General aspects
- Qualification
- Key personnel
- Job descriptions
- Training (purpose, goals, contents, target groups)
- Planning and documentation of training



Q&A Session 1 12.00 – 12.30 h

Break 12.30 – 13.30 h

Documentation 13.30 – 15.00 h

- Structure of documentation
- Responsibilities for the documentation
- SOP
- Documentation in the manufacturing process
- Documentation in the quality control
- Batch record review
- Annual report / Product quality report
- Specifications

Break 15.00 – 15.15 h

Premises / Production 15.15 – 16.00 h

- Requirements for room and equipment
- Classification of rooms
- Sterile production/isolator
- Maintenance of hygiene
- How to behave during production

General & Specific Aspects of a QA System 16.00 – 17.00 h

- Quality Management System (QMS) cited in the regulations
- General aspects of Quality Assurance
- How to implement and structure a system
- Responsibilities
- Deviations
- CAPA
- Change Control
- OOS
- Failure Investigations
- Self-inspections



Q&A Session 2 17.00 – 17.30 h

Programme 20 October 2021

Risk Management 09.00 – 09.45 h

- Main topics of ICH Q 9 / Part 3 EU GMP Guideline
- Risk management tools
- Use of risk analysis during qualification
- How to handle FMEA?

Qualification/Calibration/Maintenance 09.45 – 10.45 h

- Definitions: Qualification, calibration, maintenance
- Organizing qualification
- Steps in Qualification studies: DQ, IQ, OQ, PQ
- Qualification parameters of typical types of equipment: Clean rooms, water systems, production equipment, analytical equipment
- Performing risk analysis: tools and practical tips
- Calibration: critical types of equipment
- How to build up a calibration system
- Maintenance: Requirements and system

Break 10.45 – 11.00 h

Validation 11.00 – 12.15 h

- Definitions
- Process Validation
- PAT
- Validation Master Plan (VMP)
- Cleaning Validation
- Computer Validation
- Validation of Analytical Methods



Q&A Session 3 12.15 – 12.45 h

Break 12.45 – 13.45 h

Audits and Self-Inspections 13.45 – 14.45 h

- Types of audits
- Requirements
- Dos and don'ts for the auditee - How to survive audits?
- Performing audits and self-inspections
- Good audit practices

Break 14.45 – 15.00 h

Packaging/Storage/Transportation 15.00 – 16.00 h

- Packaging/Storage/Transportation in the regulations
- Managing of packaging process
- What is necessary to regulate in a pharmaceutical company
- WHO good storage practice – elements and requirements
- Transportation as part of storage
- How to maintain the quality during transportation

Falsified Products 16.00 – 16.30 h

- Regulatory requirements
- Responsibilities of QP and the pharmaceutical industry
- What measures can be taken
- Strategies against falsified products



Q&A Session 4 16.30 – 17.00 h

Speakers



Dr Heinrich Prinz
PDM Consulting, Germany

Dr Prinz worked with Boehringer Mannheim before he joined Biotest where he was Head of Quality Assurance, responsible for both the pharmaceutical and the medical device division. Since 2003 he works as a freelance consultant.



Dr Gabriele Schönberger,
ZS.CTIS Consulting,
formerly Boehringer Ingelheim, Germany

Dr. Schönberger is a pharmacist. From 1989 to June 2001 she was employed at Asta Medica AG, among other things as plant manager for parenteralia, head of validation within pharmaceutical production, IPC, regulatory affairs and packaging development. From July 2001 until end of 2018 she was in the QA department of Boehringer Ingelheim GmbH. Currently she works as a consultant.



Dr Wolfgang Schumacher, formerly
F. Hoffmann-La Roche Ltd., Switzerland

Dr Schumacher studied chemistry and pharmacy. After entering Asta Medica, he headed different positions. From 2001 to 2016 he was Head of the department of Quality Computer Systems at F. Hoffmann-La Roche, Basle. He is a member of the ECA Advisory Board.



Participants' comments:

„From now on I will always remember the difference between qualification and validation! :) Excellent examples and lots of useful hands-on information.“

Anniina Pertovaara, Hankintatukku Oy, Finland

„To all presentations: Very good, clear, informative presentations, with the right amount of information and presented at the right place. Excellent presentations, with lots of information, transmitted in a clear way.“ Dr Fátima Sobral, Direção-Geral de Alimentação e Veterinária, Portugal

„Very informative.“

Jennifer Martin, Cultivation Sector Consulting, USA

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Reservation Form (Please complete in full)



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Title, first name, surname

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Company

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- Cancellation until 2 weeks prior to the conference 10%.

- Cancellation within 1 week prior to the conference 50%.

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Date of the Live Online Training

Tuesday, 19 October 2021, 09.00 h – 17.30 h CEST

Wednesday, 20 October 2021, 09.00 h – 17.00 h CEST

Technical Requirements

For our Live Online Training Courses and Webinars, we use Cisco WebEx, one of the leading suppliers of online meetings. At <http://www.webex.com/test-meeting.html> you can check if your system meets the necessary requirements for the participation at a WebEx meeting and at the same time install the necessary plug-in. Please just enter your name and e-mail address for the test. If the installation is not possible because of your rights for the computer system, please contact your IT department. WebEx is a standard nowadays and the necessary installation is fast and easy.

Fees (per delegate, plus VAT)

ECA Members € 1,490

APIC Members € 1,590

Non-ECA Members € 1,690

EU GMP Inspectorates € 845

The conference fee is payable in advance after receipt of invoice.

Registration

Via the attached reservation form, by e-mail or by fax message.

Or you register online at www.gmp-compliance.org.

Presentations/Certificate

The presentations will be made available to you prior to the Live Online Training as PDF files. After the event, you will automatically receive your certificate of participation.

Conference language

The official conference language will be English.

Ordering Recordings

Independent from the Live Online Training Courses, you can also order recordings of selected Live Online Training Courses at the same conditions – at www.gmp-compliance.org/recordings. These recordings will be provided on our media server. All you need to watch it is an Internet browser – no additional software.

Organisation and Contact

ECA has entrusted Concept Heidelberg with the organisation of this event.

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