

# GMP for Beginners

Understanding the importance of GMP

## SPEAKERS:



**Dr Bettina Pahlen**  
*Quality x Pharma Consulting*



**Dr Heinrich Prinz**  
*Apceth*



**Dr Wolfgang Schumacher**  
*formerly F. Hoffmann-  
La Roche*



Good  
Manufacturing  
Practices

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19-20 March 2019, Hamburg, Germany  
29-30 October 2019, Berlin, Germany

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## LEARNING OBJECTIVES:

- GMP: Where do we come from – where do we go?
- 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



# GMP for Beginners

19-20 March 2019, Hamburg, Germany | 29-30 October 2019, Berlin, Germany

## Objectives

The course 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:

### *Commisson 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 the US and their consequences for product quality. During this course, speakers with long-standing experience in the training of employees will introduce and explain the most important elements of a pharmaceutical GMP system in an easy-to-understand way.

## Target Group

The course 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.

## Programme

### GMP: where do we come from - where do we go to?

- Development of GMPs
- GMP: Goal and general ideas
- Types of regulatory documents and their meaning
- GMP regulation for APIs, medicinal products and excipients - a comparison
- Authorities and industry groups: ICH, PIC/S, EMA, FDA, WHO, APIC, ISPE, IPEC

### GMP in the US

- Comparison of US and EU regulations
- Differences between the European and the FDA view on GMP / GMP vs cGMP
- Typical expectations of FDA and European inspectors

### Quality Management System

- Quality Management System cited in the regulations
- General aspects of Quality Assurance
- How to implement and structure a system
- Responsibilities

### Personnel and Training

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

### Hygiene / Personal Hygiene

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

### Documentation

- 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

### Specific Aspects of a QA System

- Deviations
- CAPA
- Change Control
- OOS
- Failure Investigations
- Self-inspections

Participants' comments of March 2018 course:

*„In total very helpful and well presented/explained lectures. Thank you.“*

Katharina Bubb , Janssen Vaccines AG, Switzerland

*„Highly informative. Worthy & completed answers received for questions –*

*highly open discussion atmosphere. Highly informative for getting the overview of GMP Basics.“*

Laia Camprubi Gallet , Bayer Bitterfeld GmbH



## Risk Management

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

## Premises / Production

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

## Qualification/Calibration/Maintenance

- Definitions: Qualification, validation, calibration, maintenance, risk analysis
- Organizing qualification and validation: the validation master plan (VMP)
- 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
- Validation of computerised systems

## Process Validation, Computer Validation and Validation of Analytical Methods

- General aspects and requirements
- Process validation
- Documentation of process validation
- Validation of analytical methods
- Documentation of analytical methods validation

## Cleaning Validation

- Regulators requirements
- The cleaning procedure
- Building up a cleaning validation
- Sampling
- Analytical tests

## Audits and Inspections

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

## Packaging/Storage/Transportation

- 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

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

## Speakers



**Dr Bettina Pahlen**, *Quality x Pharma Consulting GmbH, Alling, Germany*

Bettina Pahlen, PhD, studied pharmacy at the University of Muenster, Germany, graduated in pharmaceutical chemistry and performed post-docs in US and Germany. During the last

15 years she worked at university, authority and in different areas of the pharmaceutical industry (R&D, manufacturing, quality control, quality assurance). Since July 2007, she has been working as a consultant in the pharmaceutical industry focussing on GxP Quality Assurance aspects.



**Dr Heinrich Prinz**, *Apceth, 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 has been working as a freelance consultant and part of his time he is the Senior Supervisor 'Production and Quality Assurance' at Apceth, a biotech company.



**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.

## Social Event

In the evening of the first course day, you are cordially invited to a social event. This is an excellent opportunity to share your experiences with colleagues from other companies in a relaxed atmosphere.





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**GMP for Beginners**

- 19-20 March 2019, Hamburg, Germany
- 29-30 October 2019, Berlin, Germany

- Mr  Ms

Title, first name, surname

Company

Department

**Important: Please indicate your company's VAT ID Number**

**P.O. Number (if applicable)**

Street/P.O. Box

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Zip Code

Country

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E-Mail (please fill in)

**Easy Registration**



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Germany



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+ 49 6221 84 44 34



e-mail:  
info@concept-heidelberg.de



Internet:  
www.gmp-compliance.org



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**General terms and conditions**  
If you cannot attend the conference you have two options:  
1. We are happy to welcome a substitute colleague at any time.  
2. If you have to cancel entirely we must charge the following processing fees. Cancellation:  
- until 1 week prior to the conference 10 %  
- until 1 week prior to the conference 50 %  
- within 1 week prior to the conference 100 %.

CONCEPT HEIDELBERG reserves the right to change the materials, instructors, or speakers without notice to cancel an event. If the event that you cancelled is cancelled, registrants will be notified as soon as possible and will receive a full refund of fees paid. CONCEPT HEIDELBERG will not be responsible for discount airfare penalties or other costs incurred due to a cancellation.  
**Terms of payment:** Payable without deductions within 10 days after receipt of invoice.  
German law shall apply. Court of jurisdiction is Heidelberg.

**Important:** This is a binding registration and above fees are due in case of cancellation or non-attendance. If you cannot take part, you have to inform us in writing. The cancellation fee will then be calculated according to the point of time at which we receive your message. In case you do not appear at the event without having informed us, you will have to pay the full registration fee, even if you have not made the payment yet. Only after we have received your payment, you are entitled to participate in the conference (receipt of payment will not be confirmed!) (As of January 2012)

**Privacy Policy:** By registering for this event, I accept the processing of my Personal Data. Concept Heidelberg will use my data for the processing of this order, for which I hereby declare to agree that my personal data is stored and processed. Concept Heidelberg will only send me information in relation with this order or similar ones. My personal data will not be disclosed to third parties (see also the privacy policy at [http://www.gmp-compliance.org/eca\\_privacy.html](http://www.gmp-compliance.org/eca_privacy.html)). I note that I can ask for the modification, correction or deletion of my data at any time via the contact form on this website.

**Date and Venue March 2019**

Tuesday, 19 March 2019, 09.00 h - 18.00 h  
(Registration and coffee 08.30 h - 09.00 h)  
Wednesday, 20 March 2019, 08.30 h - 17.00 h

Barcelo Hamburg  
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20095 Hamburg  
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**Date and Venue October 2019**

Tuesday, 29 October 2019, 09.00 h - 18.00 h  
(Registration and coffee 08.30 h - 09.00 h)  
Wednesday, 30 October 2019, 08.30 h - 17.00 h

Steigenberger Hotel Berlin  
Los-Angeles-Platz 1  
10789 Berlin  
Phone +49 (0)30 212 7 - 0  
Email berlin@steigenberger.de

**Fees (per delegate plus VAT)**

ECA Members € 1,390  
APIC Members € 1,490  
Non-ECA Members € 1,590  
EU GMP Inspectorates € 795  
The conference fee is payable in advance after receipt of invoice and includes conference documentation, dinner on the first day, lunch on both days and all refreshments. VAT is reclaimable.

**Accommodation**

CONCEPT HEIDELBERG has reserved a limited number of rooms in the conference hotel. You will receive a room reservation form/POG when you have registered for the course. Reservation should be made directly with the hotel. Early reservation is recommended.

**Registration**

Via the attached reservation form, by e-mail or by fax message. Or you register online at [www.gmp-compliance.org](http://www.gmp-compliance.org).

**Conference Language**

The official conference language will be English.

**Organisation and Contact**

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