



Speakers



David Abraham
Quality Resource
Solutions Associates, UK



Heike Gottschalg
Boehringer Ingelheim,
Germany



Alfred Hunt
Hunt Pharma Solutions,
Ireland



Savvas Koulouridas
Fagron BV, Netherlands



Dr Daniel Müller
GMP/GDP Inspector,
Germany



Robert Müller
Boehringer Ingelheim,
Germany

Supported by the
European GDP Association



An ECA Foundation Interest Group

The GDP-Compliance Manager



Live Online Training on 19/20 November 2024



Highlights

- Expectations of the Inspectorates
- Gap Analysis and Risk Assessment
- Implementation Planning
- Key Areas of the EU-GDP Guidelines:
 - Quality Management and Organisation
 - Deviations and Complaints
 - Premises and Equipment
 - Personnel
 - Supplier Selection and Qualification
 - Transport
 - Contracting
- 4 Q&A sessions

All participants will receive a Roadmap to Good
Distribution Practice:
- Overview of the designated responsibilities
- Checklist for the implementation of GDP principles

Objectives

This Live Online Training provides practical guidance to bring and keep your organisation in compliance with the GDP regulations.

Background

The globalisation of the pharmaceutical supply chain has created new challenges for the manufacture and supply of medicinal products in various markets, resulting in reduced control and increased security risk to the products.

The **EU-GDP Guidelines** have been extensively revised to take into account the changing nature of the globalised supply chain. The requirements have been effective since 2013. These requirements highlight the need for an effective quality management system supported by risk assessment and appropriate controls.

This two-day Live Online Training has been designed to bring you up-to-date with the current regulatory expectations and standards for Good Distribution Practice (GDP) and to provide you with **tools and guidance** to help you with **identifying the gaps in your quality systems and planning and implementing the actions required**.

Target Audience

GDP Compliance Managers and Responsible Persons from companies involved in the distribution and supply of medicinal products.

Moderator

Dr Markus Funk
CONCEPT HEIDELBERG (on behalf of ECA)



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Programme

Welcome and Introduction

The (new) GDP Guideline: What was it all about?

- Background to development and revision of the EU GDP Guidelines
- Overview of structure and content
- What is the impact on industry and other stakeholders?

Quality Management System (QMS)

- What is a QMS and why do we need it?
- What does an effective QMS look like?
- How to develop and implement an effective QMS

Operations

- Qualification of suppliers and customers
- Receipt, storage and return of medicinal products
- Deviation and Complaint Management in a wholesaler facility
- How to conduct a gap analysis, develop plans and implement the new requirements



Questions & Answers Session I

GDP Inspection Findings and what to learn from them

- Findings and their ratings
- Examples from manufacturers, wholesalers, storage facilities and transport deviations

Personnel

- Competency requirements for GDP personnel
- Overview of the role and responsibilities of the Responsible Person
- Necessary documentation
- Training matrix and managing continuous training



Questions & Answers Session II

Premises & Equipment

- What is a must for medicinal products
- How to plan and implement facility improvement ensuring compliance with the current requirements

Transportation

- Key requirements for transportation of medicines
- How to develop and implement a GDP-compliant and cost effective transportation network.

Contracts in the Global Supply Chain

- International laws and systems – how they work and fit together
- Jurisdictions and conflict of law provisions
- Contract law, Technical/ Quality Agreement, Supply Agreement
- 3PL Providers: two bilateral agreements or one tripartite agreement?
- When things go wrong



Questions & Answers Session III

Outsourced Activities

- What is an outsourced activity?
- How to set priorities to audit, approve and manage service providers
- How to develop and manage contracts and agreements

Case Study for a successful Implementation Approach

- How we approached the new requirements
- Challenges and best practice



Questions & Answers Session IV

Short Summary and Take Away Message

- Developing a take home action plan for the delegates

David Abraham, Quality Resource Solutions Associates, UK

David has extensive experience in both business and Quality Management. David's background has seen him working within Pharmaceutical and Healthcare arena from print and packaging, distribution as well as pharmaceutical manufacturing organizations. His work continues to see his engagement across the industry, supply chain and training organisations providing resource, awareness, training and consulting in quality management and the application of GXP as well as continue to provide input on a number of technical committees at a national, European and International level.

Heike Gottschalg, Boehringer Ingelheim Pharma GmbH & Co. KG, Germany

Heike Gottschalg is responsible for the Quality Oversight and Compliance for logistics from a global perspective in the Corporate Division Quality within Boehringer Ingelheim. Before that she has been responsible for the global implementation of the GDP requirements within Boehringer Ingelheim.

Alfred Hunt, Hunt Pharma Solutions, Ireland

Alfred Hunt is a consultant. From 2008 until 2015 he was an Inspector with the Health Products Regulatory Authority (HPRA), formerly the Irish Medicines Board (IMB). He was also key member of the European Medicines Agency (EMA) drafting group which developed the revised EU GDP Guidelines (2013/C 343/01).

Savvas Koulouridas, Fagron BV, Netherlands

Savvas Koulouridas is Global Innovation Director of Fagron. He is leading the innovation and global marketing department of the company. He is a lawyer in profession and has also worked as a consultant on pharmaceutical law (GMP regulations and Pharmaceutical Contracts).

Dr Daniel Müller

GMP/GDP Inspectorate, Local Government, Germany
Dr Daniel Müller is head of the GMP Inspectorate at the local competent authority in Tübingen, Germany. Since 2001 he has been working as GMGP inspector, conducting national inspections as well as EMA- and overseas inspections.

Robert Müller, Boehringer Ingelheim Pharma GmbH & Co. KG, Germany

Robert Müller is responsible for maintaining global standards for shipping and temperature monitoring in the Global Logistics group (Corporate Division Supply Network & Lifecycle Management). In collaboration with the colleagues of the Global Quality group he has been responsible for the global implementation of the GDP requirements within Boehringer Ingelheim.



4 Q&A sessions ensure interaction and that your questions are answered.

About the European GDP Association:

The European GDP Association aims to support Pharmaceutical Industry, Authorities and Logistic Providers with regard to the implementation of Good Distribution Practice.

More information at:

www.good-distribution-practice-group.org

If the bill-to-address deviates from the specifications on the right, please fill out here:

Reservation Form (Please complete in full)



The GDP-Compliance Manager, Live Online Training on 19/20 November 2024

Title, first name, surname

Department

Company

Important: Please indicate your company's VAT ID Number

Purchase Order Number, if applicable

CONCEPT HEIDELBERG
P.O. Box 101764
Fax +49 (0) 62 21/84 44 34

D-69007 Heidelberg
GERMANY

City

ZIP Code

Country

Phone / Fax

E-Mail (Please fill in)

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- If you cannot attend the conference you have two options:
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Date of the Live Online Training

Tuesday, 19 November 2024, 09.00 – 17.00 h
 Wednesday, 20 November 2024, 09.00 – 17.00 h
 All times mentioned are CET.

Technical Requirements

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Fees (per delegate, plus VAT)

ECA Members € 1,690
 European GDP Association Members € 1,690
 APIC Members € 1,790
 Non-ECA Members € 1,890
 EU GMP Inspectorates € 945
 The 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.

You cannot attend the Live Event?

We also offer many of the training courses and conferences as recordings. This means that you can watch the videos of the event „on demand“ – whenever it suits you – on our web server. It is quite uncomplicated and doesn't require any software – you simply watch the video on your browser. You can find all recorded events at www.gmp-compliance.org/recordings.

Organisation and Contact

ECA has entrusted Concept Heidelberg with the organisation of this event.
 CONCEPT HEIDELBERG
 P.O. Box 10 17 64
 D-69007 Heidelberg
 Phone: +49(0) 62 21/84 44-0
 Fax: 49(0) 62 21/84 44 34
info@concept-heidelberg.de
www.concept-heidelberg.com

For questions regarding content please contact:

Dr Markus Funk (Director Operations) at
 +49(0) 62 21/84 44 40, or per e-mail at
funk@concept-heidelberg.de

For questions regarding organisation please contact:

Ms Nicole Bach (Organisation Manager) at
 +49(0) 62 21/84 44 22, or per e-mail at
bach@concept-heidelberg.de

