

Speakers



Ciara Clarke Sumac Works, Ireland



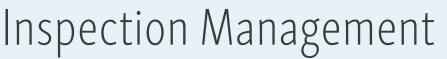
Alexander Kammerlocher GMP/GDP Inspectorate, Local Government, Germany



Katja Kotter Vetter Pharma-Fertigung, Germany



Dr Ralf Schreiner QProgress, Germany

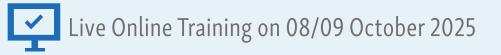


How to pass global GMP Inspections

GMP Certification Programme

Certified Quality Assurance Manager

GMP





Highlights

- Inspection Management
 - How Inspectors are trained
 - Adequate Preparation
 - Mock Inspection
 - Successful Inspection Management
 - Efficient Follow-up
- Experience from global Inspections
 - FDA
 - Brazil (ANVISA)
 - Mexico (COFEPRIS)
 - Turkey (MOH)
 - Russia (FSI SID&GP)
 - Eurasian Economic Union (EAEU)
 - China (NMPA)
 - South Korea (MFDS)
 - Taiwan (TFDA)

All participants receive a Checklist for FDA Inspection Preparation

Objectives

You will understand the purpose and organisation of regulatory inspections and you will learn how to prepare your company to pass an inspection or customer audit and how to assure the most positive outcome.

Get practical knowledge of:

- What inspectors are looking for
- Successful preparation and management of inspections
- Performing a MOCK-Inspection
- Latest trends (with a view on virtual/remote inspections)

In addition, you will hear examples from global inspections to gain a **better understanding of what is expected**.

Background

GMP audits and inspections are **fundamental elements of managing quality** in the pharmaceutical industry. On the one hand, pharmaceutical companies have to perform supplier audits. And on the other hand, the pharmaceutical companies as well as the suppliers are frequently inspected by the authorities (both national and international inspectorates like the FDA) as a central element of supervision.

For the company, an inspection can have a decisive influence on the daily work and its economic future. A sound and thorough preparation is an essential key to successfully pass an inspection.

Target Audience

This GMP Education Course is designed for all persons involved in preparing, managing and escorting audits and inspections.

Moderator

Dr Gerhard Becker CONCEPT Heidelberg (on behalf of ECA)



Stay informed with the GMP Newsletters from ECA

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To subscribe, simply scan the QR code on the right or visit www.gmp-compliance.org/ gmp-newsletter



Programme

- Inspection preparation, strategy and tactics
- Information transfer between inspectorates

Approach and Expectations of the Agencies

- What to expect, when being inspected in the near future
- Observations some practical examples

Preparing for a Regulatory Inspection

- Team building
- Gap analysis and action plan
- Roles and responsibilities
- Training of the staff
- Function of moderator, escorts and experts

Case Study:

Proactive Compliance and Inspection Management – it's more than Self Inspection

- How to increase inspection risk-awareness
- Risk categorisation and ranking
- Risk reduction prioritization
- Reporting of the results to senior management

The MOCK-Inspection: Auditing your Company to prepare for international Inspections

- Internal Audit and Mock-Inspection
- Audit strategy
- Roles and Responsibilities
- Communication and co-operation
- Sequence of preparation steps
- Co-operation with customers and external auditors

Expectations from Inspectorates worldwide

- Brazil (ANVISA)
- Mexico (COFEPRIS)
- Turkey (MOH)
- Russia (FSI SID&GP)
- Eurasian Economic Union (EAEU)
- China (NMPA)
- South Korea (MFDS)
- Taiwan (TFDA)

The FDA Approach

- The MRA between the U.S. and the EU and its consequences
- The FDA Inspection System
- What does FDA expect?

Responding to Audit and Inspection Findings

- How to reply to report and observations
- Dissent and dispute
- Proof of CAPA effectiveness
- Ensuring that measures are implemented company-wide
- What to do if a target date can not be achieved?

GMP/GDP Certification Scheme



Building on your education the ECA GMP/GDP certification programmes provide you with the appropriate supplement to acquire this

qualification. Simply choose any three courses within the programme according to your professional interest. Your certificate is then valid for two years. To renew it, you can pick any training from the ECA courses and conferences list within that two-years period – allowing you to broaden your knowledge in GMP and GDP compliance.

This could be of interest for you as well

Would you like to train a larger group of participants in your company?

We offer practice-oriented GMP/GDP training courses on:

- Basic GMP
 - APIs (ICH Q7)
 - Medicinal Products
 - Biopharmaceuticals
- Quality Assurance
- Quality Control
- Validation/Qualification
- Regulatory Affairs
- Sterile Manufacturing
- IT / Computer Validation
- Good Distribution Practice (GDP)
- Data Integrity

You will find a time schedule for each training course at www.gmp-compliance.org/training/gmp-gdp-in-house-trainings



Ciara Clarke Sumac Works, Ireland

Ciara Clarke started her consultancy business 2021. In her last role she was Senior QA Executive, QP and Deputy RP at Viatris (formerly Mylan). She was also Assistant Lecturer in Science at the Technological University Dublin.



Alexander Kammerlocher Regional Council Office, Germany

Alexander Kammerlocher is inspector at the Regional Council Office (Regierungspräsidium) in the

federal state of Baden-Württemberg.



Katja Kotter Vetter Pharma-Fertigung, Germany

Katja Kotter is Vice President Regulatory Affairs and Quality Compliance. She has broad experience in managing authority inspections and customer audits.



Dr Ralf Schreiner QProgress, Germany

Dr Ralf Schreiner started his consultancy business in 2018. Prior to that, he spent 20 years in various management positions in the pharmaceutical industry, most recently as Executive Director Quality Systems at Actavis/Allergan.

Your Benefit:

Internationally Acknowledged Certificate from ECA Academy

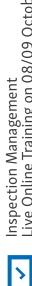
The EU GMP Guide requires:

"... All personnel should be aware of the principles of Good Manufacturing Practice that affect them and receive initial and continuing training,...".



This is why you receive an acknowledged participant certificate, which lists the contents of the seminar in detail and with which you document your training.

'ation Form (Please complete in full)



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			CONCEPT HEIDELBERG P.O. Box 101764
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Date of the Live Online Training

Wednesday, 08 October 2025, 09.00h - 15.30h Thursday, 09 October 2025, 09.00h - 15.30h All times mentioned are CEST.

Technical Requirements

We use Webex for our live online training courses and webinars. At www.gmp-compliance.org/training/online-training-technical-information you will find all the information you need to participate in our events and you can check if your system meets the necessary requirements to participate. If the installation of browser extensions is not possible due to your rights in the IT 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 € 1,890 APIC Members € 1,990 Non-ECA Members € 2.090 EU GMP Inspectorates € 1,045 The conference fee is payable in advance after receipt of invoice.

Registration

Via the attached reservation form, by e-mail or by fax – or search and register directly at www.gmp-compliance.org under the number 21973.

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 E-Mail: info@concept-heidelberg.de www.concept-heidelberg.com

For questions regarding content:

Mr Wolfgang Schmitt (Operations Director) at +49(0) 62 21/84 44 39, or per e-mail at w.schmitt@concept-heidelberg.de

For questions regarding organisation please contact:

Mr Niklaus Thiel (Organisation Manager) at +49(0) 62 21/84 44 43, or per e-mail at thiel@concept-heidelberg.de

writing. The cancellation fee will then be calculated accoraing to une point or 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 paymentyst. Only after we have received your payment, you are entitled to participate in the con-ference (receipt of payment will not be confirmed)! (As of July 2022). German law shall apply. Court of jurisdiction is Heidelberg.

Important: This is a binding registration and above fees are due in case of can-

invoice.