ECA Academy
Your GMP/GDP Information Source

ICH Q9 / ICH Q10 Training Courses

Speakers:

Richard M. Bonner
Formerly with Eli Lilly, UK

Dr Heinrich Prinz
Apceth GmbH, Germany

Dr Bernd Renger
Immediate Past Chair of the European QP Association; Renger Consulting, Germany

Dr Thomas Schneppe
Bayer Pharma AG, Germany

Dr Helene Zuurmond
Pfizer, Belgium

ICH Q 9 Training Course
19-20 October 2016, Berlin, Germany

ICH Q 10 Training Course
20-21 October 2016, Berlin, Germany

Free of charge Quality Manual on CD-ROM and ICH Q10/ ISO 9001 Matrix

Save money and book both courses for € 990,- each!

ICH Q9 Training Course

19-20 October 2016, Berlin, Germany

Objectives

The Guideline ICH Q9 „Quality Risk Management" was finalised in November 2005 (Step 5). Consequently, this guideline has been implemented in the EU, the US (FDA) and in Japan.

The ICH Q9 training course in hand deals with the **practical implementation of the requirements**. Individual examples help to show the application in the following GMP areas:

- Validation
- Change Control/Change Management
- Auditing/Inspections
- Quality Systems

As a complement to the lectures, the closing **workshop** offers the opportunity to practise Quality Risk Management techniques **with a case study**.

Target Audience

This event has been designed for quality assurance personnel. It also addresses staff from other departments, like e.g. production, engineering, quality control, and regulatory affairs.

Programme

ICH Q 9 Quality Risk Management

- Basic requirements
- Comparison to ISO 14971
- Practical Examples

How to Realise Quality Risk Management in a GMP Environment

- An overview of the diverse techniques (HACCP, FTA etc.)
- Access benefit of some tools
- Examples from the production of drug (medicinal) products and APIs

How to Apply Quality Risk Management in Validation

- Understand why the use of risk management in process validation is now expected
- What does the FDA expect for batch conformance prior to, and post, product approval
- What is the benefit of using the risk management approach versus the traditional 3 batch validation approach?
- What does ICH Q9 mean with respect to quality risk management in validation
- Learn why the quality risk management approach to validation will result in less ongoing process support during production

Design of an Event Handling System based on a Quality System and Quality Risk Management Approach

- The quality system
- Risk management principles
- Events (e. g. deviations, complaints etc.)
- Risk management application
- Outputs

How to implement Quality Risk Management in a Pharmaceutical Company

- The risk-based approach and its impact on key GMP processes (GAMP®5, FDA Guidance on Process Validation, etc)
- ICH Q9 in the context of ICH Q8 and ICH Q10
- The implementation of „quality risk management thinking and doing" on management and shop floor level (tools and experiences)

How to Make a Risk-based Audit Schedule

- Understand how to assess risk between different operations
- Identify priorities for the audit
- Learn how to use a point system to assign audit priorities based on risk
- How to use a template to make a risk-based audit schedule

Workshop Quality Risk Management in Practice

Learn how to **create an audit schedule by looking at the various risk categories** associated with the various operational activities within the differing units. This workshop will help you look across the different units from production operations, vendors, third-parties and laboratories, rank them by risk and then apply this to a template to create an audit schedule covering the next 3 years.

After the workshop you will be able to use or adapt the template to conduct a similar risk profile for your own facilities and third-party operations

Speakers

Richard M. Bonner, formerly with Eli Lilly, United Kingdom
Mr Bonner is currently located in the UK and works as a consultant to the Pharmaceutical Industry. Previous to his current role he was a Senior Quality Adviser for Eli Lilly and Company. He had 31 years experience within the pharmaceutical industry working in production, technical services and both Quality Control and Quality Assurance functions. He has been involved in multiple inspections from the MHRA, FDA and other authorities. Mr Bonner is a Qualified Person in Europe and Chairman of the European Compliance Academy.

Dr Helene Zuurmond, Pfizer, Belgium
Dr Helene Zuurmond studied Chemistry at Leiden University in the Netherlands. After working at a Pfizer site in Italy in the registration compliance and quality systems area, she is now working in the Global Quality Organisation within the same company, where she is responsible for design and implementation of compliant and efficient quality systems at the Pfizer manufacturing sites

Dr Heinrich Prinz, Apceth GmbH, 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 and part of his time he is the Senior Supervisor 'Production and Quality Assurance' at Apceth, a biotech company.

Dr Thomas Schneppe, Bayer Pharma AG, Berlin, Germany
Dr Thomas Schneppe worked for Klöckner Pentapack Contract Manufacturing, Asche AG and Schering AG. Since 2006 he is Head of the Department Integrated Quality Management Training at Bayer Pharma AG.

ICH Q 10 Training Course

20-21 October 2016, Berlin, Germany

Objectives

The International Conference on Harmonisation has published the Guidance ICH Q10 Pharmaceutical Quality Systems in June 2008. This Guideline has been transferred to European, US and Japanese regulation. Thus, all companies in these regions have to implement the key requirements of ICH Q10. However, the US FDA is still using their own Guidance for Industry (Quality System Approach to Pharmaceutical cGMP).

The implementation of these requirements have caused a number of questions. Among others ISO elements like continual improvement are new in the pharmaceutical industry. This training course has been developed to discuss the requirements and how they can be implemented in pharmaceutical industry.

Target Group

This event has been designed for quality assurance personnel. It also addresses staff from other departments, like e.g. production, engineering, quality control, and regulatory affairs.

In addition the topics will be further discussed in **interactive Workshops.**

Programme

Introduction to ICH Q10

- What are the key elements of ICH Q10?
- Comparison with the FDA Guidance for Industry
- How to comply with the Guidelines

Continual Improvement (Part 1)

- Process Monitoring
- Key Performance Indicators (KPIs)
- Process Performance and Capability (link to Q8, Q9 and Process Validation)

Continual Improvement (Part 2)

- CAPA Management as a tool for Continual Improvement
- Change Management
- Trending
- Annual Reviews

Monitoring Quality Process Performance and Quality System

- The Senior Management's responsibility
- How to perform Quality Management Reviews
- Key Elements of a Review System

Responsibility of Senior Management

- How to involve the management
- Management Review
- Practical Examples

Management of Outsourced Activities and Purchased Materials

- How to choose, qualify and monitor Suppliers and Providers
- Supplier Qualification as Part of the Quality System
- Change of Ownership
- Monitoring of the performance
- Implementation of QMS - some milestones

All presentations will also include interactive Workshops!

Speakers

Dr Heinrich Prinz,
Apceth GmbH , 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. In addition, he is the Senior Supervisor 'Production and Quality Assurance' at Apceth, a biotech company.

Dr Bernd Renger,
Immediate Past Chairman of the European QP Association;
Renger Consulting, Germany
Dr Bernd Renger is a member of the European Compliance Academy (ECA) Advisory Board and Immediate Past Chairman of the European QP Association. Since 2011, he is running his own consultancy business. Before that he was VP of Quality Control at Vetter Pharma-Fertigung. He started his career 1977 at Hoechst AG as a research and development chemist. Since then, he has held several positions at Mundipharma, Byk Gulden (now Takeda) and Baxter BioScience in Vienna.

Dr Thomas Schneppe,
Bayer Pharma AG, Berlin, Germany
Dr Thomas Schneppe worked for Klöckner Pentapack Contract Manufacturing, Asche AG and Schering AG. Since 2006 he is Head of the Department Integrated Quality Management Training at Bayer Pharma AG.

Quality Management Manual of a fictive Company on CD-ROM – free of charge for all participants of the ICH Q10 course

The Quality Management Manual of the fictive company "Example" does not only take into consideration the quality assurance system (QA System) as required by the GMP regulation but also the requirements of the international standards EN ISO 9001: 2000 on Quality Management Systems and EN ISO 13485: 2003 "Quality Management Systems – Medical Devices – Requirements for Regulatory Purposes". The Quality Manual was developed by a task force of the German Medicines Manufacturers Association. The content is structured according to ISO 9001. In the appendix of the publication you will find exemplary job descriptions, e.g. for the Qualified Person, Head of Production and Head of Quality Control. Further examples include forms for the review by the management and a process flow chart.

ICH Q10 versus ISO 9001-2015 Matrix
As part of the conference binder the participants will also receive a matrix which compares the ICH Q10 Guideline and the international standard on quality management ISO 9001. This matrix is helpful to identify areas that are not covered in one of the two documents.

Social Event

On **Wednesday evening you are cordially invited to a social event**. This is an excellent opportunity to share your own views and experiences with colleagues from other companies in a relaxed and casual atmosphere.
Join us for a guided city tour and a dinner in the heart of Berlin

Easy Registration

Reservation Form:
CONCEPT HEIDELBERG
P.O. Box 10 17 64
69007 Heidelberg, Germany

Reservation Form:
+ 49 6221 84 44 34

e-mail:
info@concept-heidelberg.de

Internet:
www.gmp-compliance.org

ICH Q9 Training Course
Wednesday, 19 October 2016, 10.00 - 17:45 h
(Registration and coffee 09.30 - 10.00 h)
Thursday, 20 October 2016, 09.00-12.15 h

Conference fees (per delegate plus VAT)
ECA Members € 1,090
APIC Members € 1,190
Non-ECA Members € 1,290
EU GMP Inspectorates € 645
The conference fee is payable in advance after receipt of invoice and includes conference documentation, lunch and dinner (Social Event) on the first day, and all refreshments. VAT is reclaimable.

ICH Q10 Training Course
Thursday, 20 October 2016, 13.30 - 17.45 h
(Registration and coffee 13.00 - 13.30 h)
Friday, 21 October 2016, 09.00 - 16.00 h

Venue of both courses
Steigenberger Hotel Berlin
Los-Angeles-Platz 1
10789 Berlin
Phone +49 (0)30 212 7 - 0
Fax +49 (0)30 212 7-799

Conference fees (per delegate plus VAT)
ECA Members € 1,090
APIC Members € 1,190
Non-ECA Members € 1,290
EU GMP Inspectorates € 645
The conference fee is payable in advance after receipt of invoice and includes conference documentation, lunch on the second day and all refreshments. VAT is reclaimable.

Save money and book both courses for € 990,- EACH!
If you book the "ICH Q9 Training Course" AND the „ICH Q10 Training Course" simultaneously, the fee for **EACH** conference reduces as follows:
ECA Members € 891
APIC Members € 940
Non-ECA Members € 990
EU GMP Inspectorates € 495

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

Registration
Via attached reservation form, by e-mail or by fax message. Or you register online at 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.

CONCEPT HEIDELBERG
P.O. Box 10 17 64
69007 Heidelberg, Germany
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.de

For questions regarding content:
Mr Oliver Schmidt, Operations Director, at
+49 (0) 62 21 / 84 44 23,
e-mail: schmidt@concept-heidelberg.de

For questions regarding reservation, hotel, organisation etc.:
Ms Susanne Ludwig, Organisation Manager, at
+49 (0) 62 21 / 84 44 44
e-mail: ludwig@concept-heidelberg.de

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

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CONCEPT HEIDELBERG
P.O. Box 10 17 64
Fax +49 (0) 6221/84 44 34

69007 Heidelberg
Germany

Registration form (please complete in full)

☐ **ICH Q 9 Training Course**
19-20 October 2016, Berlin, Germany

☐ **ICH Q 10 Training Course**
20-21 October 2016, Berlin, Germany

☐ Mr ☐ Ms Title _____

First name, surname

Company

Department

Important: Please indicate your company's VAT ID Number **Purchase Order No. (if applicable)**

Street / P.O. Box

City Zip Code Country

Phone / Fax

E-mail (please fill in)

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 2 weeks prior to the conference 10 %,
- until 1 weeks 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 or to cancel an event. If the event must be cancelled, registrants will be notified as soon as possible and will receive a full refund of fees paid. CONCEPT HEIDELBERGwill 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. **Important:** This is a binding registration and above fees are due in case of cancellation or non-appearance.

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). 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.