

Speakers



Dr Steffen Groß
PEI, Germany



Dr Line Lundsberg-Nielsen
Lundsberg Consulting, UK



Dr Martin Maus
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Dr Christian Palmes
Bayer, Germany



Dr Jörg Schiewe
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ICH Q8 Training Course

From QbD to Process Validation

Live Online Training on 9/10 May 2023



Development, Process Validation, Lifecycle Approach,
Control Strategy / PAT / RTRT

Highlights

- Quality by Design (QbD)
- Quality Target Product Profile (QTPP)
- Critical Quality Attributes (CQA)
- Critical Process Parameters (CPP)
- DoE Examples
- Design Space (DS)
- Control Strategy
- Process Analytical Technology (PAT)/RTRT
- ICH Q12: A Lifecycle Approach to Process Validation

Objectives

You will be updated on the latest regulatory developments and learn how to apply the respective paradigms in Pharmaceutical Development to be better able to design strategies for the implementation of Quality by Design (QbD) according to ICH Q8.

During this Live Online Training elements and methodologies associated with ICH Q8 will be discussed. All this will be illustrated with examples and case studies.

Background

The impact of ICH Q8, Q9, Q10, and Q11 is changing both the regulatory expectations and the strategies of Pharmaceutical Development, and this impact will continue to grow, especially in view of the new ICH Q12 Guideline.

The QbD concept described in ICH Q8 and ICH Q11 have to be seen as an overarching paradigm and an interdisciplinary approach across the product lifecycle. It also systematically emphasises enhanced product and process understanding throughout the product lifecycle.

Ideally, application of ICH Q8 and ICH Q11 elements already starts in the early design phase of a drug product where both patients needs and process design are considered. The QbD concept requires a comprehensive understanding of the chemical and physical nature of the individual active substance(s) and excipients, and of the way their attributes interact in the formulation and how they are impacted by the manufacturing process. During the design phase, it is important to establish the Quality Target Product Profile (QTPP), determine the Critical Quality Attributes (CQAs), identify Critical Process Parameters (CPPs) and Material Attributes (material CQAs) and to understand how the process parameters and material attributes affect the CQAs. The relationship between process inputs (material attributes and process parameters) and the CQAs is described in the Design Space and ensured during manufacturing with an enhanced control strategy, leading to improved process understanding, greater operational flexibility and opportunities for more efficient life cycle management activities.

ICH Q8 combined with the new Q12 will open the door to a powerful era of refined, modern and efficient pharmaceutical development and optimization for those companies who are ready to invest in this new paradigm.

Target Audience

This Live Online Training is designed for all scientists, engineers, managers and executives from Pharmaceutical and Biotech Development units and support functions to Manufacturing, including Quality Assurance and Technical/CMC Regulatory Affairs, who are involved in the implementation of ICH Q8 elements.

Programme Day 1

Introduction to QbD – setting the Scene

- Drug product development at a glance – from first in man to marketing authorization
- Pharmaceutical QbD: Quo vadis?
- Application of QbD principles to drug product development

QbD - Regulatory Perspective

- Current state of PAT & QbD implementation and regulatory challenges
- Quality by on-line (PAT) measurements
- Real time release testing: general considerations
- Going forward: ICH Q12 / Q13 / Q14

QbD Toolbox: DoE, PAT, Basic Statistics

- Value-added use of QbD tools – generic approaches and tailored solutions
- Examples for different unit operations



Q&A Session 1



Case Study DoE

- Why we use DoE in the pharmaceutical development
- Example: DoE for formulation selection / optimization
- Example: DoE for manufacturing process optimization
- DoE vs “traditional” approach – when to use which

Quality Risk Management and Control Strategy

- Objectives of Quality Risk Assessment (QRA) as part of development
- Overview on risk assessment tools
- Introduction to Process Risk Map
- Introduction of risk-based control strategy development



Q&A Session 2

GMP/GDP Certification Scheme

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Programme Day 2

Reports and Documentation

- Development Reports
- Transfer protocols and reports
- Control Strategy and link to the submission dossier

How the QbD derived Control Strategy defines Process Validation

- QbD and PAT as an enabler for gaining Process Understanding and designing the process and the control strategy, PV stage 1 (establishing the control strategy)
- Different approaches to PV/PPQ depending on the type of control strategy, PV stage 2 (traditional, continuous process verification or hybrid approach used to verify the control strategy)
- Ongoing/Continued Process verification, PV stage 3 (verifying the validity and robustness of the control strategy)



Process Validation - Case Study (Small Molecule Drug Product)

Case study related to the previous presentation (PV stage 1 and 2)

- Case example of a solid dosage form process enabled by a QbD approach
- Establishing the control strategy: Examples of the application of PAT/RTRT
- Validation of the process – verification of the control strategy

Ongoing Process Verification and Lifecycle Approach of a Process established from QbD Principles

- Continuous process verification versus continued/ongoing process verification (PV stage 3)
- Case study – the continuation of the example from above (PV stage 3)
- ICH Q12, performance-based control and the link to PAT
- Life cycle management of the product, process and control strategy opportunities for a product developed using of a QbD principles



Q&A Session 3

Participant's comment from the May 2018 course:

"Excellent speakers!"

Gordana Savi, Croatian Agency for Medicinal Products and Medical Devices

Speakers



Dr Steffen Groß, Paul-Ehrlich-Institut (PEI) Federal Institute for Vaccines and Biomedicines, Germany

Steffen Groß received his PhD degree from the Max Planck Research Unit Molecular Cell Biology Jena, Germany. Today, he is Head Section Monoclonal and Polyclonal Antibodies and Scientific Assessor (Quality, Pre-clinic) at the Paul-Ehrlich Institute (PEI).



Dr Line Lundsberg-Nielsen, Lundsberg Consulting Ltd., UK

Line is a scientist and runs her own consultancy business focusing on applying a science and risk based approach for pharmaceutical development, process design, technology transfer, qualification and process validation. She has many years of experience within the pharmaceutical industry and has a theoretical as well as practical approach to QbD, PAT and RTRT from working at Novo Nordisk and Lundbeck before being a consultant. Line is an active ISPE member and has had different chairing roles and is a well-recognized international speaker and instructor



Dr Martin Maus, Boehringer Ingelheim Pharma GmbH, Germany

Martin is a Principal Scientist at Boehringer Ingelheim in Biberach where he is working in late stage product development of solid oral dosage forms.



Dr Christian Palmes, Bayer AG, Germany

Christian is currently CMC Design Space Expert (Statistical Expert) in the Department of Digital Transformation in CPD, Pharmaceuticals, at Bayer. Previously he was Head of Statistics & Data Sciences (leadership and management of the department of statistics for laboratory diagnostics) at Siemens Healthineers and CMC Senior Statistician (statistical inference for large molecule biopharmaceuticals) at Boehringer Ingelheim.



Dr Jörg Schiewe, Boehringer Ingelheim Pharma GmbH, Germany

Jörg is currently Head of Global CMC Experts at Boehringer Ingelheim. In the past 20 years he was involved in the development of innovative inhalation products. In his current position Jörg provides scientific expertise and guidance for chemical, analytical and pharmaceutical development projects and is involved in international regulatory submissions.



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Date of the Live Online Training

Tuesday, 9 May 2023, 9.00 to 17.00 h CEST
 Wednesday, 10 May 2023, 9.00 to 13.30 h CEST

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Fees (per delegate, plus VAT)

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

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Organisation and Contact

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

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