

Speakers



Dr Ragna Hoffmann
Boehringer Ingelheim



Dr Gerd Michael Maier
Boehringer Ingelheim



Dr Uwe Muenster
Bayer



Dr Alexander Pontius
Bayer

Dissolution Testing

Development / Quality Control and *in vivo* Relevance



Live Online Training on 29/30 June 2022



Image: ERWEKA GmbH, Germany

Highlights

- Importance of Dissolution Testing in Drug Development and for a Commercial Product
- In vivo Relevant Dissolution Testing
- Discriminatory Power of a Dissolution Method
- Regulatory Requirements (Pharmacopoeias, Required Data for Application of Marketing Authorisation)
- Country-specific Challenges: Japan, Korea, Taiwan, China etc.
- Automation of Dissolution Methods
- Mechanical Qualification and Performance Verification Testing (PVT)
- Development of Dissolution Methods
 - How to Set Specifications?
 - Analytical Validation
 - Practical Recommendations
- OOS Results in Dissolution Testing
- Dissolution Profile Comparison



Q&A sessions and exercises ensure interaction

Objective

This Live Online Training on Dissolution Testing aims at providing delegates with a sound understanding of the principles and best practices in dissolution testing.

As Dissolution represents a very interdisciplinary topic, a broad variety of areas within the development and commercial phase will be discussed. You will get to know

- how to characterize formulations
- how to support formulation and process development:
- how to evaluate the impact of formulation and process parameters changes
- how to control the quality (QC tool) of clinical trial supplies and the commercial product
- how to support drug product stability testing
- how to justify formulation/production changes (e.g., according to SUPAC, Biowaivers)
- how to predict in vivo performance

Due to the wide range of applications and the sensitivity of dissolution testing, sound method development and validation is of essential importance. Furthermore also knowledge on dissolution apparatus qualification, dissolution specification setting, dissolution profile comparison and handling of OOS/OOE results will be trained and discussed.

Background

The dissolution test is a key test parameter for assessing the performance of solid and semi-solid dosage forms in both drug development and quality control. In these fields it is used to assure batch-to-batch quality as well as providing process control information as part of the approach to Process Validation.

Dissolution testing is usually connected to in vivo performance because the API must be released from the formulation in the gastro intestinal tract (GIT) before in vivo absorption can occur. Therefore dissolution testing is generally employed during Drug Product development and optimization. A dissolution test should therefore have adequate discriminatory power to detect relevant Drug Product changes.

Where dissolution testing data can be shown to be correlated to in vivo performance, clinical trials may be avoided by *in vitro* dissolution studies under certain circumstances, thereby reducing development time and costs.

There are many dissolution guidances and associated guidelines (e.g. FDA, EMA and the Pharmacopoeias) dealing with Scale-up and Post-Approval Changes, Bioequivalence studies, Waiver of in vivo Bioavailability and Bioequivalence Studies. Additionally, there are some country-specific dissolution requirements which are very challenging for global pharmaceutical companies.

This Live Online Training will, therefore, cover the following topics:

- physicochemical and biopharmaceutical foundations
- dissolution method development,
- validation of the dissolution methodology

- approaches for setting specifications
- OOS and OOE Results in dissolution testing
- statistical methods for comparing dissolution profiles
- approaches for substitution of BE-studies (biowaiver) and
- approaches to establish in vitro in vivo correlations (IVIVC)
- country-specific dissolution requirements and challenges

In addition, the expectations of the European Medicines Agency (EMA) and of the pharmacopoeias (Ph.Eur. 2.9.3 and USP Chapters <711> and <1092>) including USP Reference Standard Tablets and Mechanical calibration for the dissolution apparatus qualification will be discussed.

The objective of this course is to cover all aspects of dissolution testing with a focus on practical examples.

Programme

Fundamentals of Dissolution Testing: From Physicochemistry to Bioavailability

- Mechanism and theories of solid dissolution (e.g. diffusion layer model)
- Intrinsic dissolution rate
- Sink conditions
- Kinetics of drug release
- Relationship between dissolution and bioavailability
- Quality control dissolution testing and in vivo predictive dissolution testing
- Biopharmaceutics Classification System
- Fraction of a dose absorbed classification system
- Hurdles and limitations of dissolution testing

Dissolution Testing throughout the Drug Product Development Lifecycle

- Use of dissolution testing during Drug Product development
- What is Biorelevance? Meaning and Misconceptions
- How to establish a link between dissolution and bioavailability
- The role of IVIVC
- Setting biorelevant dissolution specifications
- BCS based biowaivers
- Waivers based on proportional similarity
- Country specific regulatory differences
- Case studies

Dissolution Testing – Regulatory Guidelines

- Prerequisites of international and mostly harmonized pharmacopoeias (USP, EP, Pharm Jap)
- Miniaturization of dissolution tests
- General guidelines for dissolution testing
- Contents and differences in Chinese pharmacopoeia
- Validation of dissolution test methods
- Bioequivalence considerations
- Special in vitro bioequivalence applications in Japan
- Waiving dissolution tests by disintegration tests

Development of Dissolution Methods - The balancing Act between Quality Control and Clinically Relevance

- Method development for Immediate Release, Extended Release and Delayed Release Formulations
- Regulatory recommendations
- Dissolution apparatus and medium selection
- Use of surfactants
- Adequate discriminatory capability
- Standard Dissolution Test Conditions
- Evaluation of bio-relevance
- Dissolution methods for developing an IVIVE/C to gain regulatory flexibility
- Case studies

Mechanical Calibration & Performance Verification Test (PVT)

- Regulatory basis
- Fundamentals of instrument qualification
- Qualification and calibration of dissolution apparatuses
- Mechanical calibration
- USP Performance Verification Test (PVT)
- Deviations and OOC

Setting Specifications for Dissolution Methods

- How to set adequate dissolution specifications for various types of formulations:
- Requirements of different Pharmacopoeias and Guidelines
- Specifics and exceptions

PRACTICAL EXAMPLES / EXERCISES I

How to Set Specifications: Sharing Information of the Learned Theories

- Presentation of Case Studies and discussion of potential results

OOS Results in Dissolution Testing

- Regulatory aspects
- Dissolution methods having appropriate discriminatory power
- General OOS procedure for dissolution testing
- Defining and handling of OOS results including CAPA
- OOS evaluation for immediate release products
- OOS evaluation for capsules
- OOS evaluation for modified-release products
- OOT/OOE results: Evaluating stability effects by applying dissolution testing

Automation in Dissolution Testing

- Why and when is automation valuable?
- Various types of dissolution systems
- New products on the market

Analytical Validation of Dissolution Testing Methods

- Pharmacopoeial and Regulatory Recommendations (e.g., ICH Q2 (R1), USP <1092>, RDC No. 166/2017)
- Validation characteristics:
 - Specificity, Linearity, Precision, Accuracy and Robustness
 - Validation of automated procedures
- Some practical recommendations for performing the validation and recommended acceptance criteria
- Dissolution method transfer

PRACTICAL EXAMPLES / EXERCISES II

Analytical Validation of Dissolution Methods

Putting theory to work (case studies):

- Develop validation protocol for validation of dissolution methods for solid oral dosage forms
- Pitfalls in performing the experiments

Dissolution Profile Comparison; Approaches and Issues

- Importance of dissolution profile comparisons during drug product development and for a commercial product
- Regulatory requirements concerning dissolution profile comparison
- Different approaches to compare dissolution profiles: Model dependent and independent approaches
- Examples

Case Study: Application of Dissolution Testing in Industrial Drug Product Development

Discussion of various case studies occurring during product development



Q&A sessions

Q&A sessions on both days ensure interaction and that your questions are answered.

Speakers



Dr Ragna Hoffmann
Boehringer Ingelheim Pharma GmbH & Co. KG,
Biberach, Germany

Dr Ragna Hoffmann is a pharmacist by training and received her PhD in pharmaceutical technology from the Eberhard-Karls University of Tuebingen. She joined Boehringer Ingelheim in 2007 and has since worked in different areas of drug product development (preformulation, formulation development, and analytical development). Dr. Hoffmann is currently heading a dissolution lab within the Analytical Development Department at Boehringer Ingelheim in Biberach, Germany.



Dr Gerd Michael Maier
Boehringer Ingelheim Pharma GmbH & Co. KG,
Biberach, Germany

Dr. Gerd-Michael Maier is a chemist by training and did his PhD graduation in bioinorganic chemistry at the University of Konstanz. He joined Boehringer Ingelheim in 1997 where he started in the Drug Regulatory Affairs department and worked as a regulatory affairs manager. After three years as team leader Regulations & Training in the Quality Systems group Gerd-Michael joined the department of Drug Discovery Support in research leading a CMC lab for almost 10 years. Since 2014 he holds the position of a dissolution lab head within the Analytical Development Department at Boehringer Ingelheim, Biberach, Germany.



Dr Uwe Muenster
Bayer AG, Wuppertal, Germany

Uwe has been interested in biopharmaceutical and dissolution topics for 15 years. He is heading the dissolution department within Chemical and Pharmaceutical Development at Bayer. Uwe started his career with Bayer in 2006 as Postdoc in research pharmacokinetics, dealing with absorption prediction and drug transport across barriers. Two years later he joined Bayer's Chemical and Pharmaceutical Development. As lab head his work was focused on generation, optimization and implementation into the R&D screening cascade of in vitro, in vivo and ex vivo biopharmaceutical prediction models. Uwe was involved in multiple research collaborations on biopharmaceutical and dissolution topics with Academia and Industry partners. Furthermore, he was running a dissolution lab within the department he is now heading. Uwe is a pharmacist by training. He obtained his PhD at the Free University of Berlin.



Dr Alexander Pontius
Bayer AG, Leverkusen, Germany

Alexander is a pharmacist by training and did his PhD graduation in biopharmaceutical analytics. He was heading a QC group in the global pharmaceutical development division and was responsible for all aspects of the dissolution methodology. This work covered the development and validation of dissolution methods, bio-pharmaceutical evaluations, dossier submission of innovative drug products, handling of post approval changes as well as support of life cycle management and patent protection of market products. At present, Alexander is working as Quality System Manager within the enterprise-wide Corporate Quality function. He is responsible for the regulation management within the overarching Quality Management System at Bayer.



Date of the Live Online Training

Wednesday, 29 June 2022, 9.00 h – 18.00 h

Thursday, 30 June 2022, 9.00 h – 16.30 h

All times mentioned are CEST.

Technical Requirements

We use Webex Events 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 € 1,590

APIC Members € 1,690

Non-ECA Members € 1,790

EU GMP Inspectorates € 895

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.

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

info@concept-heidelberg.de

www.concept-heidelberg.de

For questions regarding content please contact:

Dr Markus Funk (Operations Director) at

+49(0)62 21/84 44 40, or at

funk@concept-heidelberg.de

For questions regarding organisation etc. please contact:

Mr Niklaus Thiel (Organisation Manager) at

+49(0)62 21/84 44 43, or at

thiel@concept-heidelberg.de

GMP/GDP Certification Programme

This seminar is recognised within the GMP Certification Programme. By attending selected seminars, the participant can acquire an additional certificate. We offer the following modules:

ECA Certified Validation Manager

ECA Certified QA Manager

ECA Certified API Production Manager

ECA Certified Quality Control Manager

ECA Certified Technical Operations Manager

ECA Certified Computer Validation Manager

ECA Certified Regulatory Affairs Manager

ECA Certified Microbiological Laboratory Manager

ECA Certified Sterile Production Manager

ECA Certified Biotech Manager

ECA Certified Pharmaceutical Development Manager

ECA Certified GMP Auditor

ECA Certified GDP Compliance Manager

ECA Certified Packaging Manager

ECA Certified Data Integrity Manager



At www.gmp-compliance.org you will find a text explaining which seminars are recognised for which certificates. Or you send an e-mail to info@gmp-compliance.org or a fax to +49-(0)6221- 84 44 64 with the request for information about the GMP Certification Programme. We will then send you our brochure on the topic.

Your Benefits: 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.

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

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

D-69007 Heidelberg
GERMANY

Reservation Form (Please complete in full)



Dissolution Testing, Live Online Training on 29/30 June 2022

Title, first name, surname

Department

Company

Important: Please indicate your company's VAT ID Number

Purchase Order Number, if applicable

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 %
 - Cancellation until 1 week prior to the conference 50 %
 - Cancellation 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 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.
Important: This is a binding registration and above fees are due in case of can-

cellation 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). German law shall apply. Court of jurisdiction is Heidelberg.

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.