

Speakers:

Susanne Ding

Boehringer Ingelheim, Germany

Alessandro Giuseppe Giunta

AbbVie, Germany

Rebecca Haywood

Pfizer, United Kingdom

Katrien Himpens

I&J Innovative Medicine, Belgium

Patryk Jegorow

Takeda, Ireland

Shirley Murphy

Takeda, Ireland

Claudius Pop

Novartis, German

Andreas Schwinn

Roche Pharma AG, Germany

Niina Taylor

Brenda Van Assche

1& I Innovative Medicine, Belaium

Markus Zeitz

Takeda, Switzerland

IMP Working Group

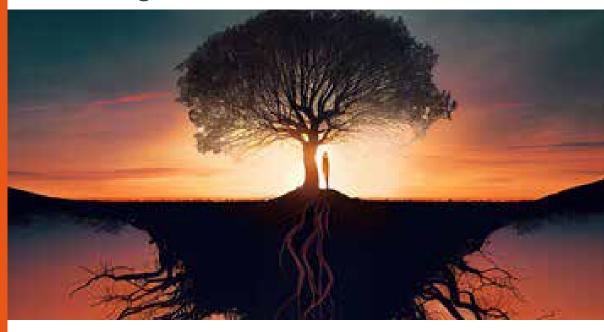
The Qualified Person Forum Pre-Conference Session

Specific Requirements for IMPs

Amsterdam

27 November 2024

Strengthen the IMP QP's Foundation



Legislation updates impacting IMP QPs,
Clinical Trial Regulation survey,
third country IMP GMP inspections first practical experiences,
regulatory view,
critical thinking as an IMP QP,
Quality culture in the IMP context,
Use of GenAl in GxP regulated environment: application & considerations,
Practical examples,
Q&A sessions

Background IMP Working Group

The IMP Working Group is a well-established part of the EQPA and its Pre-Conference Workshop an established piece of the annual QP Association event. The workshop is also facilitated as an annual meeting of the IMP Working Group and provides an excellent opportunity for networking and cultivating personal contacts.

Objectives

- Represent QPs within the European Qualified Person Association
- Maintain and grow a strong IMP network
- Share and enhance knowledge of IMP QP members by exchanging experiences, challenges, ideas and insights
- Discuss and share interpretation of regulatory guidance
- Identify and promote best practices by benchmarking across industry
- Influence new legislation by providing expert review and feedback to Regulatory Authorities
- Represent the IMP QP role to other stakeholders, e.g., industry colleagues and non-EU regulatory bodies

Management and Administration

The IMP Working Group is led by:

Susanne Ding, Boehringer Ingelheim, Germany **Katrien Himpens**, J&J Innovative Medicine, Belgium

Patryk Jegorow, Takeda, Ireland

Niina Taylor, EQPA

Administration work is provided by the EQPA Secretary. To become member of the IMP Working Group please contact us via eMail: info@ qp-association.eu. Please note that membership in this group will be granted to Members of the QP Association only.

Target Audience

- Qualified Persons and aspiring QPs for IMPs,
- Authority representatives involved with IMPs,
- · Regulatory Affairs,
- Senior Management & other IMP Industry stakeholders

Agenda IMP Pre-Conference

Welcome and Introduction

Lecture & discussion

Legislation updates impacting IMP QPs

Taskforce presentation survey

Clinical Trial Regulation – experience so far

Lecture & discussion

Third country IMP EU GMP inspections – first practical experiences

Lecture & discussion

Regulatory view

Interactive case studies - practical examples

Presentation & interactive discussion

Critical thinking as an IMP QP

Presentation & discussion

Quality culture in the IMP context

Presentation & discussion

Use of GenAl in GxP regulated environment - application and considerations

Q&A sessions (submit your questions in advance to impqp@qp-association.eu before 04 Nov 2024)

Welcome Reception QP Forum

Moderator IMP Pre-Conference

Patryk Jegorow, Head of Quality Compliance and Systems Biologics Operating Unit, Qualified Person, Takeda, Ireland

Agenda Parallel Session "Challenges for IMP QPs"

- ATIMP Release
- Phase appropriate Quality Management Systems (QMS)
- CTA / IMPD compliance check

Speakers Susanne Ding Qualified Person IMP, Boehringer Ingelheim, Germany Alessandro Giuseppe Giunta Qualified Person, Associate Director IMP Quality Management, AbbVie, Germany ☐ **QP Forum** 28-29 November 2024 Rebecca Haywood Qualified Person, Pfizer, United Kingdom Department \Box **Katrien Himpens** Qualified Person / Site Quality head, J&J Innovative Important: Please indicate your company's VAT ID Number Medicine, Belgium Reservation Form (Please complete in full) Patryk Jegorow Head of Quality Compliance and Systems $\stackrel{\times}{\leq}$ Biologics Operating Unit, Qualified Person, Takeda, ☐ Specific Requirements for IMPs 27 November 2024 Ireland **Shirley Murphy** Head of Knowledge Management, Global Quality Ms Compliance & Systems, Takeda, Ireland Title, first name, surname E-mail (Please fill in) **Claudius Pop** Head Clinical Supplies QA / Qualified Person, Street / P.O. Box Novartis, Germany Phone / Fax Company Ĭ **Andreas Schwinn** City П Senior Qualified Person, Roche Pharma AG, Germany Niina Taylor If the bill-to-address deviates from the specification to the **EOPA Brenda Van Assche** Site Quality Head / Qualified Person, J&J Innovative Medicine, Belgium Markus Zeitz Head of Digital Quality, Takeda, Switzerland CONCEPT HEIDELBERG D-69007 Heidelberg ⁻ax 06221/84 44 34 Postfach 10 17 64 ight, please fill out here:

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(As of July 2022),

CONCEPT HEDELBERG

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Date Pre-Conference

Wednesday, 27 November 2024, 09.00 – 18.00 h (Registration: 08.30 – 9.00)

Date QP Forum

Thursday, 28 November 2024, 9.00 – 18.00 (Registration: Wednesday, 27 November, 18.00 – 19.00 and Thursday 28 November, 08.30 – 9.00) Friday, 29 November 2024, 8.30 – 14.45

Fee Pre-Conference

€ 1.190,- per delegate plus VAT. The fee is payable in advance after receipt of invoice.

Fees OP Forum

EU GMP Inspectorates: € 995,- plus VAT QP Association Member: € 1.790,- plus VAT Non-QP Association Member: € 1.990,- plus VAT

Venue

Leonardo Royal Hotel Paul van Vlissingenstraat 24 1096 BK, Amsterdam The Netherlands

Phone: (+31) 20 250 0000

E-Mail: info.royalamsterdam@leonardo-hotels.nl

Accommodation

You will receive a room reservation link when you have registered for the conference.

Reservation should be made directly with the hotel. Early reservation is recommended

Registration

Via the attached reservation form, by e-mail to info@qp-association.eu or by fax to +49 6221 / 84 44 34 . Or you register online at www.qp-forum.org.

Conference language

The official conference language will be English.

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.

Organisation / Contact

CONCEPT HEIDELBERG PO Box 10 17 64

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

For questions regarding reservation, hotel, organisation etc:

Ms Marion Grimm (Organisation Manager) at +49 (0) 62 21 / 84 44 18, or per e-mail at marion.grimm@concept-heidelberg.de.

Saving Opportunities

Book both the QP Forum and a Pre-Conference Session: Delegates who attend the QP Forum and a Pre-Conference Session will get a **discount of 200** € on the QP Forum.

Important Information!

Download: The presentations of the QP Forum and the Pre-Conference Sessions will be available for download and your print-out before and after the conference.

Note: there will be no print-outs available during the conference.