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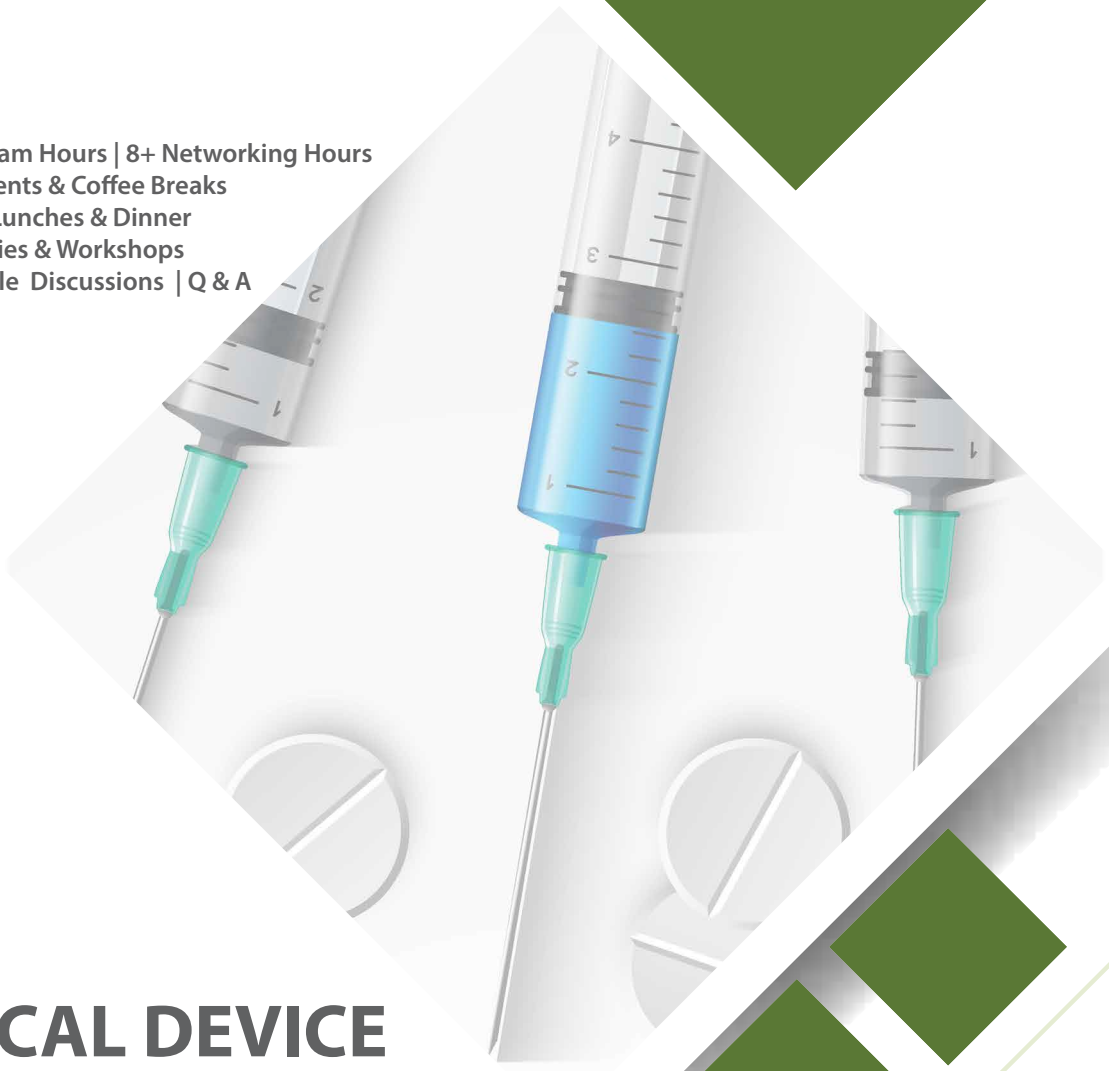
- 10+ Program Hours | 8+ Networking Hours
- Refreshments & Coffee Breaks
- Business Lunches & Dinner
- Case Studies & Workshops
- Roundtable Discussions | Q & A



MEDICAL DEVICE REGULATIONS SUMMIT 2020

September 22-23, 2020 | Prague, Czech Republic

SPEAKERS BOARD

Bijan Elahi
Award Winning International Medical Device Risk-Mgmt Educator, Consultant and Author
Medtronic, NL

Fayez Abou Hamad
MD Vigilance Expert – Pharmacist, Complaint & Vigilance Manager
Terumo Europe, BE

Dr. Daniel Latham
Head – Device Development & LCM
Novartis, CH

Dr. Clemens Günther
Director Nonclinical Safety Consumer Care
Bayer AG, DE

Dr. Silke Conrad
Quality Expert Medical Devices
Novartis, CH

Philippe Auclair
Senior Director. Regulatory Strategy and Advocacy
Abbott, BE

Dr. Max D. Singh
Global Director – Orthopedic Focus Team
TÜV SÜD Product Service GmbH, DE

Mark A. Chipperfield, M.Sc., B.Eng.(Hons), AMIMechE, MTOPRA
Company Director and Principal Consultant
Corvus Device , UK

Mariano Chiusano
EMEA Director of Regulatory Affairs
Johnson & Johnson, CH

Torsten Kneuss
Quality Assurance Manager Combination Products
Bayer Pharma, DE

Serge Mathonet
Global Regulatory Affairs CMC Biologics - Site Leader
Sanofi R&D, FR

WHO YOU WILL MEET

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SNAPSHOT OF ATTENDEES - Pre-Filled Syringes Summit 2018:

- AbbVie - Accord Healthcare - Aptar Pharma - Aristo Pharma - Bayer - Becton Dickinson - Bespak Europe - Bioton - Bristol-Myers Squibb - Cambridge Consultants - Celanese - Corvus Device Datwyler - F. Hoffmann-La Roche - Flex - GlaxoSmithKline - Hekuma - HTL-STREFA - Intertek - Janssen - Laboratoire Aguettant - Medac - Merck Group - Novartis - Novo Nordisk - Pall Life Sciences - RAUMEDIC - Sanofi - SCHOTT Pharmaceutical Packaging - SHL Group - Solvias - Sonceboz - STADA Arzneimittel - Stevanato Group - Terumo Europe - TOPAS Advanced Polymers - West Pharmaceutical Services - Worrell - Others

Agenda: https://qepler.com/pdf/pfs.pdf

SNAPSHOT OF ATTENDEES - Drug/Device Combination Products Summit 2018:

AbbVie - Ablynx - anteris medical - AOP Orphan Pharmaceuticals - Astellas Pharma - AstraZeneca - BIOCORP - Biogen - Boehringer Ingelheim microParts - Bristol-Myers Squibb - BSI Group - Corvus Device - Cytel - Design Science - Eli Lilly - Freelancer - GSK - H&B Electronic - H&T Presspart - Hanway Associates - Janssen - LEO Pharma - Maetrics - Medac - Medtronic - Nelson Labs - Nelson Labs Europe - Novartis - Orion Corporation - Orion Pharma - Pall Life Sciences - Pharmathen - Progress - PME - Regeneron Pharmaceuticals - Sanofi R&D - Sanofi-Aventis - sfm medical devices - Sharp Clinical Services - Spiegelberg - TERUMO EUROPE - tesa Labtec - Teva - UPM Raflatac - Others

Agenda: https://qepler.com/pdf/ddc.pdf

SNAPSHOT OF ATTENDEES - Pharmaceutical Lyophilization Summit 2019:

- Bachem AG - Bayer - Bayer Pharmaceuticals - BIOCAD - BioTestLab, Ltd - Boehringer Ingelheim - Boehringer Ingelheim Animal Health - CONTIPRO a.s. - CSL Behring AG - CSLBehring GmbH - Datwyler Pharma Packaging International NV - De Montfort University - Ghent University - GOETHE Biotechnology GmbH - iQ-mobil solutions GmbH/Tempris - Janssen Pharmaceutica NV - Kingston University London - Laboratoire Aguettant - Lonza AG - Lyofal - Martin Christ Gefriertrocknungsanlagen GmbH - MediWound - MSD International - Novartis - oncomed manufacturing a.s. - Patheon - PIGO srl - Sanofi Pasteur - sfm medical devices GmbH - Shire - Skan AG - SP Scientific - Takeda - Takeda GmbH - Takeda Vaccines - UCL School of Pharmacy - Vaxxinova Int. - Weibo Hi-tech Group - Others

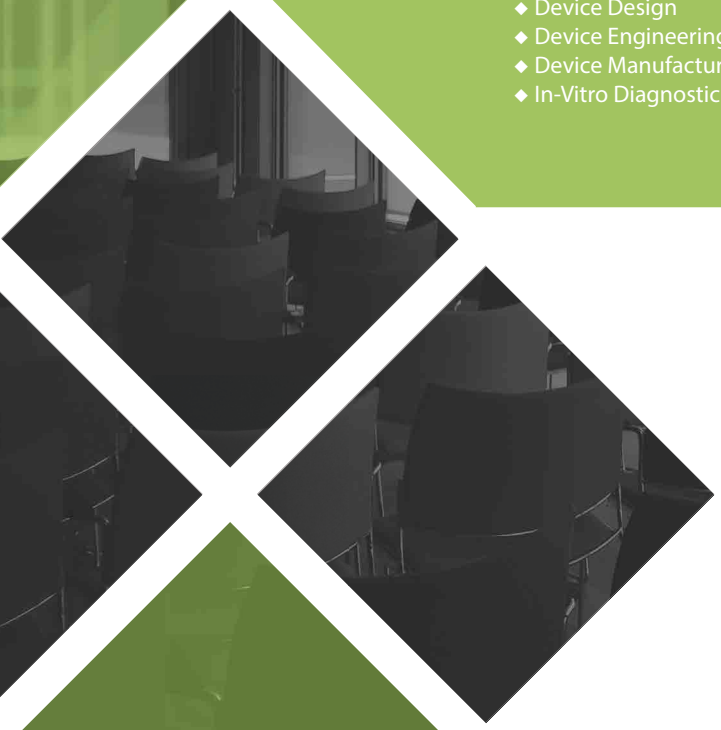
Agenda: http://qepler.com/pdf/lyo.pdf

POSITIONS

- C-Level, Presidents, Chairs, Members of the Board & VPs
- Vice presidents, Directors, & Heads
- Leaders & Managers
- Principals, Engineers, Analysts & Scientists
- Instructors & Trainers & Teachers
- Advisors, Coordinators, Auditors & Consultants
- Other Professionals, Experts & Specialists

DIVISIONS

- CMC
- Combination Products
- Compliance
- Design Controls
- Device Design
- Device Engineering
- Device Manufacturing
- In-Vitro Diagnostic
- IVDR
- MDR
- Medical Devices
- Packaging
- PFS
- Product Development
- QA
- QC
- R&D
- Regulatory Affairs
- Risk Management
- Safety Management
- Sterilization
- Traceability
- Validation
- Vigilance
- Other



INDUSTRIES

- Pharmaceutical
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- Regulatory Agencies
- Training providers
- Other

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MEDICAL DEVICE REGULATIONS SUMMIT 2020

September 22 | Prague, Czech Republic

07:30 - 08:00 — Registration and Welcome Coffee

08:00 - 08:10 — Opening Address from the Chairman

08:10 - 08:50 — Implementation of the MDR and effect on Combination Products

Mark A. Chipperfield, M.Sc., B.Eng.(Hons), AMIMechE, MTOPRA |
Company Director and Principal Consultant | Corvus Device , UK

08:50 - 09:30 — Speed Networking

09:30 - 10:10 — TBA

Serge Mathonet | Global Regulatory Affairs CMC Biologics - Site Leader | Sanofi R&D, FR

10:10 - 10:40 — Morning coffee and networking break

10:40 - 11:20 — Case Study 3 - RESERVED FOR

GlaxoSmithKline

11:20 - 12:00 — TBA

Dr. Daniel Latham | Head – Device Development & LCM | Novartis, CH

12:00 - 13:00 — Business lunch

13:00 - 13:30 — Slot Reserved for a Gold or Silver Sponsor

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MEDICAL DEVICE REGULATIONS SUMMIT 2020

September 22 | Prague, Czech Republic

13:30 - 14:10 — Updating the QMS to MDR

- MDR – Main changes
- QMS gap analysis on existing combination product/medical device processes and action plan
- Step wise implementation

Dr. Silke Conrad | Quality Expert Medical Devices | Novartis, CH

14:10 - 14:30 — Slot Reserved for a Booth or Bronze Sponsor

14:30 - 15:00 — Afternoon coffee and networking break

15:00 - 15:40 — TBA

Torsten Kneuss | Quality Assurance Manager Combination Products | Bayer Pharma, DE

15:40 - 16:40 — Workshop - Risk Management

Bijan Elahi | Award Winning International Medical Device Risk-Mgmt Educator, Consultant and author | Medtronic, NL

16:40 -17:20 — EU MDR Vigilance System – EU Vigilance Decision Tree and Optimal Set Up

Fayez Abou Hamad | MD Vigilance Expert – Pharmacist, Complaint & Vigilance Manager | Terumo Europe, BE

17:20 -18:10 — Roundtable Discussion

18:10 - 18:20 — Chairman's closing remarks and end of day one

19:00 - 21:00 — Business dinner

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MEDICAL DEVICE REGULATIONS SUMMIT 2020

September 23 | Prague, Czech Republic

08:00 - 08:30 — Registration and Welcome Coffee

08:30 - 08:40 — Opening Address from the Chairman

08:40 - 09:20

EUDAMED, The European Medical Device Database. Current Status and Future Developments

EUDAMED, the European Medical Device Database of Medical Devices is one of the most significant changes introduced by the new European Medical Device Regulations on Medical Devices (MDR) and In Vitro Diagnostics Medical Devices (IVDR). As the core of the new regulatory system of MDR and IVDR, a lot of resources and efforts have already been spent by the EU Commission and different stakeholders to design, build, validate and further develop the database. This presentation will analyze the current status of EUDAMED and future improvements of one of the most challenging European Initiatives ever introduced in the medical device sector.

Mariano Chiusano | EMEA Director of Regulatory Affairs | Johnson & Johnson, CH

09:20 - 10:00

Relation between manufacturers and Notified Bodies

- Implementation of the regulation is just starting
- The current environment is fluid
- Both manufacturers and Notified bodies are learning
- Review of the challenges seen
- How to overcome them

Philippe Auclair | Senior Director. Regulatory Strategy and Advocacy | Abbott, BE

10:00 - 10:30 — Morning coffee and networking break

10:30 - 10:50

Slot Reserved for a Speaking Sponsor

10:50 - 11:50

Workshop - EU MDR Post Market Surveillance – Requirements and Lessons Learned

Fayez Abou Hamad | MD Vigilance Expert – Pharmacist, Complaint & Vigilance Manager | Terumo Europe, BE

11:50 - 12:50 — Business lunch

12:50 - 13:30

Case Study - RESERVED FOR

Zimmer Biomet

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MEDICAL DEVICE REGULATIONS SUMMIT 2020

September 23 | Prague, Czech Republic

13:30 - 14:30

Workshop: General Safety and Performance Requirements (GSPR) according to the EU MDR

Dr. Max D. Singh | Global Director – Orthopedic Focus Team | TÜV SÜD Product Service GmbH, DE

14:30 - 15:00

Afternoon coffee and networking break

15:00 - 15:40

Biocompatibility testing to meet the requirements of the EU Medical Device Regulation

- Understanding the principles of nonclinical safety evaluation of medical devices
- Clarifying the requirements for biocompatibility testing according to ISO-10993
- Selecting an appropriate test strategy
- How to practically apply biocompatibility testing
- Examples and potential pitfalls

Dr. Clemens Günther | Director Nonclinical Safety Consumer Care | Bayer AG, DE

15:40 - 16:20

Case Study

16:20 - 16:40

Q & A

16:40 -16:50

Chairman's closing remarks and end of day two

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MEDICAL DEVICE
REGULATIONS
SUMMIT
2020

September 22-23 | Prague, Czech Republic

BIOGRAPHIES

Bijan Elahi
Award Winning International Medical Device Risk-Mgmt Educator, Consultant and Author
Medtronic, NL

Bijan Elahi has worked in risk management for medical devices for over 25 years at the largest medical device companies in the world, as well as small startups. He is currently employed at Medtronic as a Technical Fellow where he serves as the corporate expert on safety risk management of medical devices. In this capacity, he offers education and consulting on risk management to all Medtronic business units, worldwide. Bijan is also a lecturer at Delft University of Technology, and Eindhoven University of Technology in the Netherlands, where he teaches risk management to doctoral students in engineering. Bijan is a frequently invited speaker at professional conferences, and is also a contributor to ISO 14971, the international standard on the application risk management to medical devices. He is the author of the book Safety Risk Management for Medical Devices.

Fayez Abou Hamad
MD Vigilance Expert – Pharmacist, Complaint & Vigilance Manager
Terumo Europe, BE

Fayez is a pharmacist with more than 15 years experience in the quality assurance within the pharmaceutical and medical device industry. Since 2008, Fayez joined Terumo Europe where he held roles of increasing responsibility in maintaining quality, risk management and clinical evidence systems. In his current position, he is responsible for global complaint handling and vigilance systems. Fayez is also a certified lead auditor for medical device quality system regulations.

Dr. Daniel Latham
Head – Device Development & LCM
Novartis, CH

Daniel Latham is Head of Device Development Operations in Global Drug Development, Novartis, where he leads an organization responsible for developing delivery systems for combination products for new biologic entities, biosimilars and small molecules.
During the past 10 years at Novartis he has overseen significant device and primary packaging developments and launches and has significantly supported the growth of device development and combination products within the organization.
Prior to Novartis he working in a variety of roles in consumer healthcare focusing on the development of OTC medicines, transdermal patches and medical devices.
He has a PhD in controlled drug delivery from Queen Mary's, University of London and Bachelor and Master's degrees in Engineering from the University of Sheffield.

Dr. Clemens Günther
Director Nonclinical Safety Consumer Care
Bayer AG, DE

Dr. Clemens Günther received his diploma in biology and doctorate for natural sciences from the Free University, Berlin-Germany.
From 1990 to 2006 he started his professional career at Schering AG, Berlin-Germany.
From 2007 to 2013, Dr. Clemens Günther was Director and Head of Global Preclinical Development at Intendis GmbH, branded later-on as Bayer Dermatology. In this position, he was responsible for Nonclinical Safety for the marketed product portfolio of Bayer Dermatology as well as the global preclinical development strategy including human DMPK for development and life cycle management projects.
Since integration of Intendis into Bayer in 2013, he became Director Nonclinical Safety Consumer Care within the Division of Bayer Pharmaceuticals.
Meanwhile Dr. Clemens Günther has gained 29 years experience in nonclinical safety. He has been involved in nonclinical development and regulatory toxicology of small molecules, biologics, medical devices and drug device combination products.

Dr. Silke Conrad
Quality Expert Medical Devices
Novartis, CH

Silke Conrad is currently Quality Expert for Medical Device Development at Novartis Pharma AG. In her current role, she mainly supports medical devices and combination products from a QA perspective, negotiates and approves QA agreements and establishes and maintains QA processes for medical device development. Silke joined Novartis in 2011 as Regulatory CMC Associate Director for Biologics from the medical devices company Ypsomed AG/Switzerland, where she was leading the regulatory affairs department.
Prior to Ypsomed AG, Silke worked for Sanofi-Aventis in Frankfurt/Germany and Holmes Chapel/UK holding different positions in Regulatory Affairs and Quality Assurance with main focus on biotech/biological products and combination products.
Silke is a food chemist/environmental toxicologist from education and holds a doctorate degree in molecular biology/cancer research from the Technical University of Kaiserslautern/Germany.

Torsten Kneuss
Quality Assurance Manager Combination Products
Bayer Pharma, DE

Torsten Kneuss studied Business Administration and Engineering. Since 1999 he has been working with pharmaceutical packaging materials, medical devices and combination products, including several years within the field of quality control, development, operations, and pharmacovigilance. Since November 2017 he is, as a Quality Manager Combination Products, responsible for devices and combination products within Bayer AG.

Mariano Chiusano
EMEA Director of Regulatory Affairs
Johnson & Johnson, CH

Mariano Chiusano is a Chemical Engineer with more than 20 years of experience in the medical device sector covering different roles in Quality and Regulatory Affairs within Johnson & Johnson for different medical devices classes in Endo-surgery, Cardiology, Orthopedics. He has been spending the last 7 years in Diabetology expanding his experience to In vitro Medical Devices and Digital products covering the role of Legal Manufacturer for the systems for the measurement of blood glucose and insulin delivery.

BIOGRAPHIES

Mark A. Chipperfield, M.Sc., B.Eng.(Hons), AMIMechE, MTOPRA
Company Director and Principal Consultant
Corvus Device , UK

Mark serves as an independent consultant to the Pharma and Medical Device industries via his company Corvus Device Ltd.
He has over twenty years of experience in Medical Device, Drug Delivery Device and Combination Products across Development, Operations, Regulatory/Quality Compliance and product maintenance – from a range of roles with GSK, sanofi-aventis, Novartis and F. Hoffmann-La Roche.
Through his career to date he has been heavily involved in development of medical devices for combination products in several forms: syringes, pen injectors, auto-injectors, patch injectors, solution/suspension inhalers, multi-dose disposable and reusable dry powder inhalers, convenience kits, dispensers and special purpose applicators.
He has performed numerous due diligence and technical evaluations of novel delivery technologies; developed products through the full design control phases to market; and maintained marketed products.
Mark is a veteran of several successful IND/CTA/NDA/BLA/MAA submissions and approvals that have included drug delivery devices.
He has experienced many of the challenges associated with delivery device development and device product maintenance within large pharmaceutical companies and implemented Medical Device development guidance, quality systems and business processes.
Qualified with a Master's Degree in Engineering Management from Loughborough University and a Bachelor's Degree in Mechanical Engineering from London South Bank, he has maintained Continuous Professional Development with supplemental and progressive training in areas such as Technical, Manufacturing, Risk Management, Quality & Compliance, Technical Authorship, Project Management and Leadership.
He is an active presenter in the field and has co-authored a case study chapter for PDA's 2013 publication 'Combination Products: Implementation of cGMP requirements', and worked with RAPS to co-author the introduction for their 2016 publication; 'Global Medical Device Strategy'.

Serge Mathonet
Global Regulatory Affairs CMC Biologics - Site Leader
Sanofi R&D, FR

Serge is a global regulatory affairs group leader with more than 14 years of regulatory affairs experience in Chemistry, Manufacturing and Controls (CMC) focusing on biologics and biologics/device combination product development, licensing and launch/life cycle management activities. Prior to joining Global Regulatory Affairs, Serge has held various positions in Sanofi Industrial Quality Operations in API Regulatory Compliance activities either at corporate or industrial site level.
Serge is a core member of EBE Biomanufacturing group leading the combination product topic groups advocating in coordination with 7 Industry groups - Medtech and Pharma.

Philippe Auclair
Senior Director. Regulatory Strategy and Advocacy
Abbott, BE

Philippe represents European Industry in various European Commission expert groups and is the proud recipient of a US FDA CDRH Director's Special Citation for his engagement with GHTF. Since 2015, he serves as an advisor to the Asian Harmonization Working Party Technical Committee.
He chairs the Post Market Surveillance and the Notified Body Working Groups of association Medtech Europe.
Philippe received a "Global Leadership Award" from the Regulatory Affairs Professional Society (RAPS) in 2010, has been elected "RAPS Fellow" in 2012 and is co -chair of the RAPS European Advisory Committee.

REGISTRATION FORM

This registration form is editable.
When you have completed the form - please save and email it to register@qepler.com

SUMMIT NAME: Medical Device Regulations Summit 2020

REGISTRATION DATE:

TOTAL PRICE:

PROMOCODE:

PACKAGE NAME:

PACKAGE NAME	Register by 17.04..2020	Register by 19.06.2020	Register by 14.08.2020	Standard price
Individual ticket (2 Days)	€1395 (save €500) ☐	€1595 (save €300) ☐	€1695 (save €200) ☐	€1895 ☐
Individual ticket (1 Day)	€795 (save €200) ☐	€845 (save €150) ☐	€895 (save €100) ☐	€995 ☐
Group ticket (2-3 delegates)	€1095 (save €600) ☐	€1295 (save €400) ☐	€1395 (save €300) ☐	€1695 ☐
Group ticket (4+ delegates)	€895 (save €700) ☐	€995 (save €600) ☐	€1095 (save €500) ☐	€1595 ☐
Non-profit organizations	€695 (save €300) ☐	€795 (save €200) ☐	€895 (save €100) ☐	€995 ☐
Documentation €499 ☐	Promotional materials distribution €699 ☐	Speaker €2495 ☐	Pop up stand €3495 ☐	Bronze €4095 ☐
Booth €5495 ☐	Silver €6995 ☐	Gold €7995 ☐		

ATTENDEE DETAILS	1ST ATTENDEE	2ND ATTENDEE	3RD ATTENDEE	4TH ATTENDEE	5TH ATTENDEE	6TH ATTENDEE
Title:						
Name:						
Surname:						
Company:						
Country:						
Job Title:						
Direct phone:						
Email:						
Special Requirenments: (If you have any special dietary requirements or other needs that would enhance your enjoyment of this summit, please specify)						

INVOICE DETAILS

Title: Name: Surname:

Job Title:

Direct Phone: Mobile: Email:

Company: Country: City: EU VAT #:

Address: Postcode:

Payment Method:

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Signature:
«I agree to be bound by Terms and Conditions of registratin»

TERMS & CONDITIONS

REGISTRATION & PAYMENT

Upon receiving the signed registration form, we will process your application. The registration confirmation and the invoice will be sent to you within one (1) working day with the relevant payment instructions and terms. The registration fee includes access to all sessions, coffee breaks, lunches, dinner and conference materials. Payment is due 10 working days from the invoice date. Payment should be made by Credit Card, Pay Pall or by Bank Transfer. The delegate is responsible for any bank charges/fees associated with the payment.

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You may substitute a delegate at any time and at no extra cost. Cancellations must be in a written notice. Cancellations made 30 days or more before the event start date will be refunded less than 50% of the registration fee. Cancellations made less than 30 days before the event start date will receive no refund. If you cannot attend an event due to illness or other unforeseen circumstances, you may transfer your delegate pass to another upcoming event within one year from original event start date.

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PACKAGES

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PACKAGE NAME	PRICE
DOCUMENTATION Post-event presentations and other materials. Presentation content is subject to speaker's approval for distribution.	€499
PROMOTIONAL MATERIALS DISTRIBUTION (Distribution of your company's promotional materials to all attendees)	€699

SPONSORSHIP PACKAGES

BENEFITS	SPEAKER €2495	POP UP STAND €3495	BRONZE €4095	BOOTH €5495	SILVER €6995	GOLD €7995
Number of passes included	1	1	2	2	3	4
Registration fee for additional company representatives	€1295	€1295	€1195	€1195	€1095	€1095
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Pull-up banner at the entrance to the auditorium (to be provided by sponsor)					•	•
Speaking slot	20 min		20 min	20 min	30 min	30 min
Opening keynote presentation						15 min
Recognition in chairman's opening address	•	•	•	•	•	•
Seat on a panel discussion			•	•	•	•
Opening & closing speech						•
Chairman of Day 1						•
Chairman of Day 2					•	
Logo and URL on summit website, agenda and pre/post-summit communication activities	•	•	•	•	•	•
Recognition on Qepler social media channels (LinkedIn/Facebook/Twitter/Instagram)	•	•	•	•	•	•
Colour advert in placed in agenda			1/4 Page	1/4 Page	1/2 Page	1 Page
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Online distribution of your company's promotional materials to all attendees			•	•	•	•
Lanyards for summit badges, notepads, pens and other promotional materials (max. 5) given to all participants and speakers (to be provided by sponsor)						•

MARKETING CAMPAIGN

✓Website ✓Email Marketing ✓Digital Advertising ✓Social Marketing ✓Press ✓Direct Sales

PARTICIPATION FEE

Fees are inclusive of the 2-day summit, materials, online post-event documentation/presentation package, lunches, snacks, refreshments and business dinner.

TRAVEL AND ACCOMMODATION

Hotel accommodation and travel expenses are not included in the fee. Special rates for the event venue will be sent upon availability.

VENUE

Event venue will be announced online and sent to the delegates within a reasonable period before the summit start date.

POST-EVENT DOCUMENTATION

Presentations and other materials will be sent to the attendees within 72 hours after the event. Presentation content is subject to speaker's approval for distribution.

DISCOUNTS

Early booking discounts are not valid in conjunction with any other offer.

EVENTS CALENDAR 2019 - 2021

Please **choose** the **Summit / Summits** and **send** us suggested **Session Title** via email or webpage.
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DRUG DELIVERY - FORMULATION - R&D - DEVICES - COMBINATION PRODUCTS - PFS - LYOPHILIZATION - E & L - IMPURITIES - HPAPI - INHALATION - ASEPTIC PROCESSING - MDR - CMC - QRM - PFS - DDC - EM

PHARMACEUTICAL LYOPHILIZATION SUMMIT.....February 13-14, 2019
Prague, Czech Republic
Discuss best practices in tech & regulatory updates, process, formulation, testing, monitoring and new products development.
http://qepler.com/pdf/lyo.pdf

HIGHLY POTENT APIs SUMMIT.....February 20-21, 2019
Berlin, Germany
Assess and reduce manufacturing and handling challenges for highly potent active pharmaceutical ingredients.
https://qepler.com/pdf/hpapi.pdf

GENOTOXIC IMPURITIES IN PHARMACEUTICALS SUMMIT.....April 11-12, 2019
Berlin, Germany
GTI strategies & new methodologies: analysis, in silico & regulations. Challenges & opportunities.
https://qepler.com/pdf/genetoxic.pdf

INHALED DRUG DELIVERY SUMMIT.....May 23-24, 2019
Berlin, Germany
Assess and harness novel approaches to the development of inhaled drug products for enhanced patient care.
https://qepler.com/pdf/inhalation.pdf

2ND ANNUAL PRE-FILLED SYRINGES SUMMIT.....June 4-5, 2019
Barcelona, Spain
Enhance expertise sharing in developing pre-filled syringes and provide attendees with ample networking opportunities.
https://qepler.com/pdf/pfs.pdf

EXTRACTABLES & LEACHABLES SUMMIT.....October 15-16, 2019
Berlin, Germany
Get the latest updates in regulation, analytical testing, risk & safety assessment, biocompatibility.
https://qepler.com/pdf/enl.pdf
REGISTRATION IS OPEN NOW!

2ND ANNUAL DRUG/DEVICE COMBINATION PRODUCTS SUMMIT.....December 4-5, 2019
Prague, Czech Republic
Get up to date with the regulatory and quality compliance strategies for combination product development.
https://qepler.com/pdf/2ddc.pdf
REGISTRATION IS OPEN NOW!

CONTACTS

Please send your session title and summit name to:

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LinkedIn: https://www.linkedin.com/in/evgenia-makar/
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EVENTS CALENDAR 2020

Please **choose** the **Summit / Summits** and **send** us suggested **Session Title** via email or webpage.
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2ND ANNUAL PHARMACEUTICAL LYOPHILIZATION SUMMIT.....February 12-13, 2020
Berlin, Germany
Overviewing regulatory updates and advanced technologies, developing lyophilization process that ensure quality at all stages.
http://qepler.com/pdf/2lyo.pdf
THE CALL FOR SPEAKERS IS OPEN NOW!

2ND ANNUAL HIGHLY POTENT APIs SUMMIT.....February 19-21, 2020
Prague, Czech Republic
Enhancing HPAPIs manufacturing and handling via new technologies, engineering improvement and successful contamination control strategies.
https://qepler.com/pdf/2hpapi.pdf
THE CALL FOR SPEAKERS IS OPEN NOW!

PHARMACEUTICAL ASEPTIC PROCESSING SUMMIT.....May 19-20, 2020
Prague, Czech Republic
Ensuring safe highly potent aseptic production based on new regulations and technologies implementation.
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