



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



# PHARMACEUTICAL ASEPTIC PROCESSING SUMMIT 2020

**May 19-20, 2020** | Prague, Czech Republic

**Venue:**

**Novotel Praha Wenceslas Square Hotel**

Katerinska 38, 120 00 Prague





Klaus Ullherr  
Senior Product Manager  
Robert Bosch Packaging Technology, DE



**BOSCH**  
Invented for life



Mostafa Nakach  
Head of Formulation & Process  
Development / Biologics Drug Product  
Development  
Sanofi, FR



Dr. Thomas Schwarz  
Head Strategic Planning & PMO  
Novartis, CH



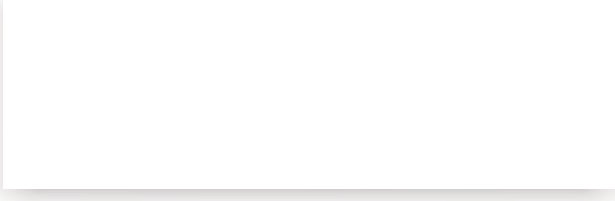
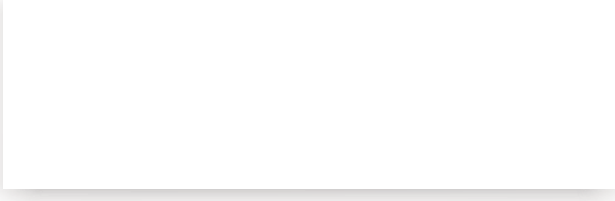
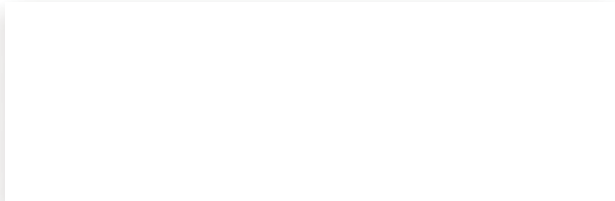
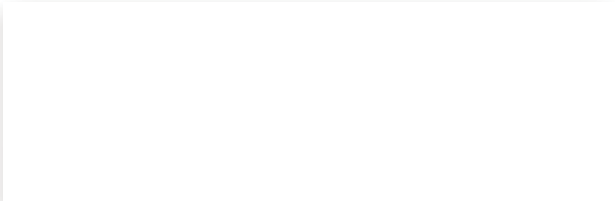
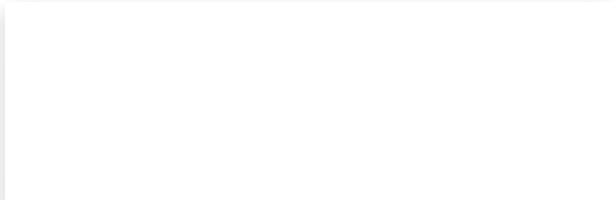
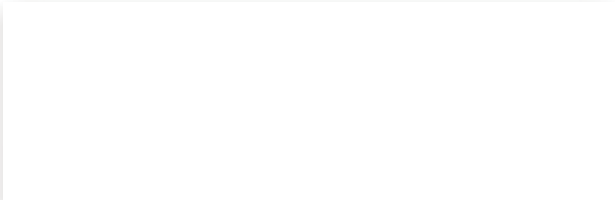
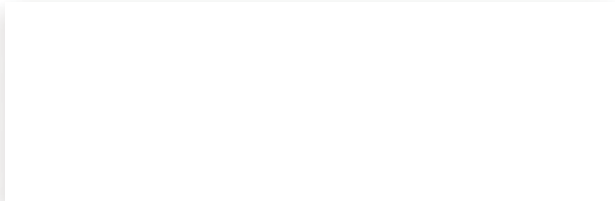
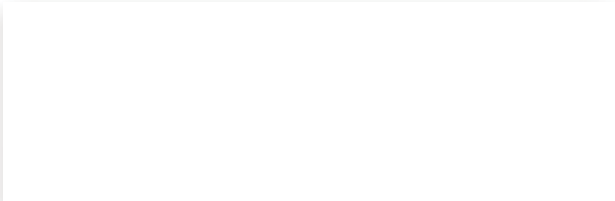
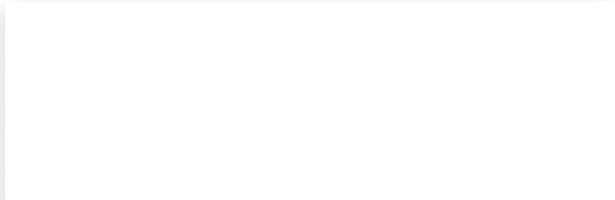
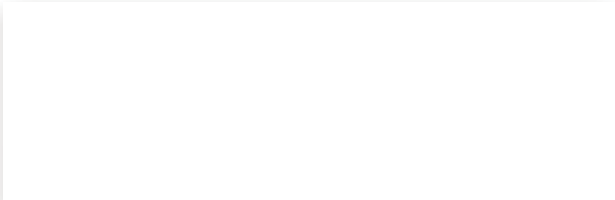
Dr. Benedikt Reichart  
Global Business Development Manager,  
Contract Manufacturing  
Fresenius Kabi, AT



**FRESENIUS  
KABI**  
caring for life



Dr. Olivier Chancel  
Sterility & Aseptic Practices Expert  
Boehringer Ingelheim, FR



## 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 - Vaxxinoa Int. - Weibo Hi-tech Group - Others

## SNAPSHOT OF ATTENDEES - Highly Potent APIs Summit 2019:

- Affygility Solutions - Ardena - ARTeSYN Biosolutions - Bayer AG - Bayer Consumer Care AG - Biocon - Bristol Myers Squibb - Celgene - ChargePoint Technology Ltd - EVER Pharma Jena GmbH - Gedeon Richter - GlaxoSmithKline - Guido Maik Consulting - Hebel Process Solution LLC - HPAPI Project Services Limited - ILC Dover - Ipsen - JLS International Germany - Krka, d.d., Novo mesto - Leeds Beckett University - Lonza - Lonza Pharma & Biotech - Lusochimica SpA - Merck Group - Minaken - Mitsui & Co. Italia S.P.A. - MSD - Novartis - Novasep - Oriento SA - Patheon, part of Thermo Fisher Scientific - Piramal Pharma Solutions - Pliva Croatia Ltd, TAPI Croatia - Praevena AG - Regeneron Pharmaceuticals, Inc - Skan AG - Takeda Ireland Limited - Umicore AG & Co KG - Others

## SNAPSHOT OF ATTENDEES - Pre-Filled Syringes Summit 2018:

- AbbVie - Accord Healthcare - Aptar Pharma - Aristo Pharma - Bayer - Becton Dickinson - Bepak 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

## 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

- ◆ ADC
- ◆ API/HPAPI
- ◆ Aseptic processing
- ◆ Cell and Gene Therapy
- ◆ CMC
- ◆ Combination Products
- ◆ Compliance
- ◆ Container/Closure Testing
- ◆ Containment
- ◆ Contamination Control
- ◆ Design Control
- ◆ Environmental Monitoring and Control
- ◆ Equipment Qualification
- ◆ Facility
- ◆ Fill Finish
- ◆ Filter integrity testing
- ◆ Formulation
- ◆ GMP
- ◆ HSE
- ◆ Industrial Hygiene
- ◆ Infrastructure management
- ◆ Isolators
- ◆ Lyophilization
- ◆ Maintenance
- ◆ Manufacturing
- ◆ Media Fills
- ◆ Medical Devices
- ◆ Microbiology
- ◆ Occupational Safety
- ◆ Packaging and Labeling
- ◆ PFS
- ◆ Plant
- ◆ Process Engineering
- ◆ Process Simulation
- ◆ Process Validation
- ◆ QA
- ◆ QbD
- ◆ QC
- ◆ Quality management
- ◆ Regulatory Affairs
- ◆ Restricted Access Barrier Systems (RABS)
- ◆ Risk Management
- ◆ Sanitization
- ◆ Sterile Products
- ◆ Sterilization
- ◆ Validation
- ◆ Velocity Testing
- ◆ Visual Inspection
- ◆ Other

## INDUSTRIES

- ◆ Pharmaceutical
- ◆ Biotechnology
- ◆ Chemical
- ◆ Medical Devices
- ◆ CDMOs
- ◆ CMOs
- ◆ CROs
- ◆ NOPS
- ◆ Regulatory Agencies
- ◆ Bioprocessing services and equipment
- ◆ Equipment suppliers
- ◆ Training providers
- ◆ Other

May 19-20 | Prague, Czech Republic

# PROPOSED TOPICS

Annex 1 and aseptic processing - compliance with the guide. Considerations for implementation. Regulatory framework. Aseptic Processing principles. Preparing for regulatory inspections.

Antibody Drug Conjugates / HPAPIs production: aseptic processing of ADCs / HPAPIs - complying with regulatory requirements, ensuring aseptic fill-finish, empowering cleaning, preventing cross contamination, providing occupational safety.

Lyophilization of highly potent products and aseptic processing.

Aseptic process development, validation and evaluation. QRM for aseptic processing. Reviewing main results, challenges and considerations achieved in aseptic manufacturing.

Aseptic processing simulation: concept creation based on current regulations, process simulation, media fills.

Pharma 4.0: Innovations and advanced aseptic manufacturing technologies. New ways for manufacturing excellence. Automated Solutions.

Highly potent facility design and engineering considerations: building safe aseptic production at each stage. Conceptual design, basic engineering, detail engineering, implementation, installation. Equipment and utilities overview, room installation, cleaning and waste management. Creating a new aseptic filling line.

Isolator Technology use in aseptic processing for loss minimization/elimination. Cycle time reduction.

Ensuring safe production for both sides (manufacturer and patient) through innovations and data sciences.

Eliminating operator impact on production via barrier systems. Restricted Access Barrier Systems.

Robotics use in aseptic processing. Modernisation of the traditional lines. New facility installation. Future, advantages and disadvantages.

Overcoming technical challenges in decontamination processes and material transfers. H2O2 decontamination – ensuring production robustness.

Ageing facilities: choosing between upgrade and renovation. Overviewing 2 cases and ways to ensure required aseptic manufacturing.

Contamination control program- cleaning and disinfection programs for aseptic facilities. Current regulations and guidance. Contamination control principles. Sterility and cleanliness of product/non-product contact surfaces. Ensuring controlled environments through disinfection and cleaning procedures. Validation protocols creation for cleaning systems implementation. Antimicrobial testing.

Airflow visualization for contamination risks assessment.

Implementing containment technologies in aseptic processing – step by step strategy.

Cross contamination prevention and control for highly potent products. VR use for cross contamination prevention.

Boosting sterilization: successful strategies, advanced technologies implementation. Sterility testing.

Sterility assurance in aseptic processing using microbial monitoring and advanced techs implementation. Regulations. Environmental monitoring design based on process analysis, risk assessment, new technologies and instruments. Components determination.

Applying QbD, ICH Q9, risk management to improve aseptic processing and sterility assurance. Aseptic process analysis and QbD based design.

Cleanroom examination, cleaning and disinfection procedures, mistakes identification. Cleaning automation and technologies.

Gowning: procedures, training, personnel qualification.

Microbiology techniques use in aseptic processing

Integrating sanitization techniques to empower contamination control.

Filtration: filter integrity testing in aseptic processing.

Single-use systems implementation.

May 19 | Prague, Czech Republic

08:00 - 08:30

 Registration and Welcome Coffee

08:30 - 08:40

 Opening Address from the Chairman

08:40 - 09:20

Case Study 1

09:20 - 10:00

 Speed Networking

10:00 - 10:40

Case Study 2

10:40 - 11:10

 Morning coffee and networking break

11:10 - 12:10

Workshop 1

12:10 - 12:50

Case Study 3

12:50 - 13:50

 Business lunch

13:50 - 14:20

Slot Reserved for a Gold or Silver Sponsor

May 19 | Prague, Czech Republic

14:20 - 15:00

Case Study 4 - Reserved



Mostafa Nakach | Head of Formulation & Process Development / Biologics Drug Product Development | **Sanofi, FR**

15:00 - 15:30

 Afternoon coffee and networking break

15:30 - 15:50

Slot Reserved for a Booth or Bronze Sponsor

15:50 - 16:30

Site Transformation Stein Steriles: From Small Molecule to Combination Product Processing

- ◆ 20 years of sterile manufacturing
- ◆ Evolution of filling technologies (ampoules, vials, prefilled syringes)
- ◆ Launch Excellence
- ◆ Development of technologies
- ◆ Change of portfolio and
- ◆ Combination products



Dr. Thomas Schwarz | Head Strategic Planning & PMO | **Novartis, CH**



16:30 - 17:00

Case Study 6 - Reserved



Dr. Benedikt Reichart | Global Business Development Manager, Contract Manufacturing | **Fresenius Kabi, AT**



17:00 - 18:00

 Roundtable Discussion

Topics

18:00 - 18:10

 Chairman's closing remarks and end of day one

19:00 - 21:00

 Business dinner

May 20 | Prague, Czech Republic

08:00 - 08:30

 Registration and Welcome Coffee

08:30 - 08:40

 Opening Address from the Chairman

08:40 - 09:20

Case Study 1 - Reserved



Klaus Ullherr | Senior Product Manager | [Robert Bosch Packaging Technology, DE](#)



**BOSCH**

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09:20 - 10:20

Workshop 1

10:20 - 10:50

 Morning coffee and networking break

10:50 - 11:10

Slot Reserved for a Speaking Sponsor

11:10 - 12:10

Workshop 2

12:10 - 13:10

 Business lunch

13:10 - 13:30

Slot Reserved for a Speaking Sponsor

May 20 | Prague, Czech Republic

13:30 - 14:10

Case Study 2 - Reserved



Dr. Olivier Chancel | Sterility & Aseptic Practices Expert | Boehringer Ingelheim, FR



14:10 - 14:40

 Afternoon coffee and networking break

14:40 - 15:20

Case Study 3

15:20 - 16:00

Case Study 4

16:00 - 16:20

 Q & A

16:20 - 16:30

 Chairman's closing remarks and end of day two

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May 19-20 | Prague, Czech Republic

## BIOGRAPHIES



**Dr. Thomas Schwarz**  
Head Strategic Planning &  
PMO  
Novartis, CH

Thomas Schwarz started his career in the pharmaceutical industry 20 years ago as a production manager at Sanofi in biotech production. He joined Novartis in 2006 and has held several positions with increasing responsibility (e.g. Process Manager, QC Team Head, Project Manager) in different technology areas like sterile manufacturing, QC operations and clinical supply management. In his current role, Thomas oversees the long-term planning and strategic projects of the Aseptics platform (6 sites worldwide). Thomas holds a PhD in Chemistry and has completed an executive MBA program.



**Klaus Ullherr**  
Senior Product Manager  
Robert Bosch Packaging  
Technology, DE

Klaus Ullherr (Senior Product Manager) has a degree in electrical engineering. After university he worked for several years as a project manager in the electrical industry. In March 2000 he joined Robert Bosch GmbH, Packaging Technology, Product Division Pharma Liquid. During the first two years he was project manager responsible for handling complex customer orders. Since 2002 he is product manager for the business fields syringes and filling systems with global product responsibility. His main functions are

- \* Market analysis
- \* Initiating new product developments
- \* Business development
- \* Expert for syringe processing

He is member of the «PDA Interest Group pre-filled syringes» and works as an expert in the DIN/ISO group for primary packaging. Klaus is also member of ISPE and co-track-leader of the yearly ISPE Aseptic Conference. He is a well-known speaker on conferences covering trends and solutions for fill/finish equipment especially for pre-filled syringes, other pre-sterilized containers and single-use-filling-systems.



**Dr. Benedikt Reichart**  
Global Business Development  
Manager,  
Contract Manufacturing  
Fresenius Kabi, AT

Benedikt acts as the first point of contact for pharmaceutical companies planning the outsourcing of manufacturing of sterile pharmaceuticals at Fresenius Kabi's manufacturing sites in Europe and Asia. Benedikt joined Fresenius Kabi as a Management Trainee after completing his PhD in Organic Chemistry from the Karl-Franzens University of Graz, Austria in 2014. He afterwards covered the role of executive assistant at Fresenius Kabi focusing on pharmaceutical manufacturing, global investments, strategy roll-out, capacity and supply chain management.



**Dr. Olivier Chancel**  
Sterility & Aseptic Practices  
Expert  
Boehringer Ingelheim, FR

Dr. Olivier Chancel is a doctor pharmacist, graduating in technological pharmacy, quality control and management. Currently, he is the sterility and aseptic practice assurance expert and formerly head of performance and pharmaceuticals in Merial, Toulouse, France. He provides corporate support for sterility and aseptic processes to Merial sites. He used to work in the pharmaceutical industry for 15 years in various positions including pharmaceutical research and development, quality assurance, quality control, project management and production for both solid and injectable dosage forms. He has published a number of papers in peerreviewed journals relating to cleanroom activities or aseptic practices and also presented/moderated at various industry forums in Europe and the US (PDA, A3P, SFSTP, SMI, ASPEC, ECA, PHARMIG...)

# REGISTRATION FORM

This registration form is editable.  
When you have completed the form - please save and email it to [register@qepler.com](mailto:register@qepler.com)

**SUMMIT NAME:** PHARMACEUTICAL ASEPTIC PROCESSING SUMMIT 2020

**REGISTRATION DATE:**

**TOTAL PRICE:**

**PROMOCODE:**

**PACKAGE NAME:**

PACKAGE NAME	Register by 08.11..2019	Register by 10.01.2020	Register by 03.04.2020	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)	€1195 (save €600)	€1395 (save €400)	€1495 (save €300)	€1795
Group ticket ( 4+ delegates )	€995 (save €700)	€1095 (save €600)	€1195 (save €500)	€1695
Non-profit organizations	€695 (save €300)	€795 (save €200)	€895 (save €100)	€995

**DELEGATE PACKAGES**

**SPONSORSHIP PACKAGES**


Documentation €499	Promotional materials distribution €699	Speaker €2495	Pop up stand €3495	Bronze €4095
Booth €5495	Silver €6995	Gold €7995		

**PLEASE MARK if you will join the Business Dinner on February 20th**  Business Dinner (Feb 20) €45

**ATTENDEES DETAILS:**

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 Requirements: (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:  
Bank Transfer  Credit Card  Pay Pall    
Signature: \_\_\_\_\_  
«I agree to be bound by Terms and Conditions of registration»

**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.

**CANCELLATION & SUBSTITUTION POLICY**

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.

**EVENT CHANGES & CANCELLATIONS**

While all effort will be made to adhere to the advertised package, Qepler s.r.o. reserves the right to change event dates, sites or location, omit event features, or merge the event with another event as it deems necessary without penalty. In such situations no refunds, part refunds or alternative offers will be made. In the event that Qepler s.r.o. permanently cancels the event for any reason, and provided that the event is not postponed to a later date nor is merged with another event, you will receive a credit note or refund for 100% of the conference fee paid. Please note, Qepler s.r.o. will not be held liable for any accommodation or associated travel costs should the event be canceled or rescheduled.

**DATA PROTECTION**

The personal information provided by you will be held in the Qepler database. For data update please write to [databasemanager@qepler.com](mailto:databasemanager@qepler.com).

# PACKAGES

## ONLINE PACKAGES

If you are unable to attend, you may purchase these packages:

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
Coupon (1 free pass for the other Qepler events)					•	•
Pop up stand in the break area (3m wide x 3m height; includes 1 table, chairs, 1 electrical socket)		•	•			
Exhibition booth with LCD monitor for video presentations in the break area (3m wide x 3m deep; includes 1 table, chairs, 1 electrical socket)				•	•	•
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
Company flyer/brochure included in conference folder (to be provided by sponsor)			•	•	•	•
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.



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**PHARMA - MEDICAL DEVICES - CHEMISTRY**



**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/genotoxic.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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 Executive Director  
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 Email: [denis.polikarpov@qepler.com](mailto:denis.polikarpov@qepler.com)  
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 Tel: +420 608 030 490  
 Email: [evgenia.makar@qepler.com](mailto:evgenia.makar@qepler.com)  
 LinkedIn: <https://www.linkedin.com/in/evgenia-makar/>  
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## PHARMA - MEDICAL DEVICES - CHEMISTRY



DRUG DELIVERY - FORMULATION - R&D - DEVICES - COMBINATION PRODUCTS - PFS - LYOPHILIZATION - E & L - IMPURITIES - HPAPI -  
 INHALATION - ASEPTIC PROCESSING - MDR - CMC - QRM - PFS - DDC - EM

- 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.  
<https://qepler.com/pdf/ap.pdf>  
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