GMP–PHARMA CONGRESS

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#sharing challenges and solutions in practice

8/9 April 2025, WIESBADEN
RheinMain CongressCenter

8 Conferences | **100** Speakers | **120** Exhibitors

- Aseptic Technologies/Annex 1 Conference
- Digitalisation & Artificial Intelligence
- Trends in Barrier Systems & Robotics
- Cleanroom Challenges
- Sustainability/Green GMP
- Single-Use Systems
- ATMPs
- Medical Cannabis

Please scan the code to read the **full agenda and details** of the GMP PharmaCongress or to **register directly online** – or visit www.pharma-congress.com

CONCEPT HEIDELBERG

Supported by ECA Academy *Your GMP/GDP Information Source*

OVERVIEW

The guiding theme of the GMP PharmaCongress 2025 on 8/9 April will once again be „users sharing challenges and solutions in practice“. Therefore, benefit from your colleagues‘ experience and from the direct information exchange.

The Conference Tracks

As a participant you can switch between any of the **8 conference tracks** any time and also visit the PharmaTechnica Expo with close to 120 international exhibitors.

The GMP PharmaCongress Conference Tracks	8 April 2025	9 April 2025
European Aseptic Technologies – Annex 1 Conference	✓	✓
Digitalisation & Artificial Intelligence	✓	✓
Trends in Barrier Systems & Robotics	✓	✓
Cleanroom Challenges in Ongoing Operations	✓	✓
Sustainability/Green GMP	✓	✓
Single-Use Systems in Sterile & Biomanufacturing	✓	✓
ATMPs – Hurdles & Achievements in Quality and Safety	✓	✓
Medical Cannabis – Cultivation, Processing, Systems & Technology	✓	✓
GMP PharmaTechnica Expo	✓	✓

KEYNOTE

Keynote 8 April 2025

Artificial Intelligence (AI) in Manufacturing and Quality at Sanofi

Dr Maite Durrenbach, Chief Quality Officer, SANOFI

Dr Maite Durrenbach is Chief Quality Officer at Sanofi. In her function she leads the Quality Function of the group for R&D, Industrial Affairs and Commercial Activities. She joined Sanofi in November 2001 and has worked in many management roles since then.

- Launch of the Integrated Quality System Management Platform with 80,000 users
- AI Applications in use and potential areas for AI
- Digitalisation of Quality Documentation and gradually replacing it with video SOPs, e-Forms, etc
- Key Performance Indicators: what has been achieved so far – what are the goals

European Aseptic Technologies – Annex 1 Conference

GMP-PHARMA CONGRESS

Fill & Finish of various Ready-to-Use Containers from clinical late Stage until commercial Launch of Biologicals through to high-potent Products

Stylianos Sampanis, *Sanofi-Aventis Deutschland GmbH*
Ralf Wagner, *Optima pharma*

Accelerating Pharmaceutical Manufacturing: A Case Study of entering Syringe and Cartridge Fill-Finish Production

Henning Austermann, *Siegfried Hameln*
Klaus Ullherr, *Syntegon Technology*

H2O2 Ingress Study Approach for Isolator Decontamination of the AT-Vials

Dr Maria Loos, *Johnson and Johnson*
Adrian Keller, *SKAN*

Container Closure Integrity Test

Luigi Scaffidi, *Boehringer Ingelheim Pharma*

Pain Points with Annex 1

Quality Risk Management in Aseptic Manufacturing: Reasonable Use

Dr Ingrid Walther, *ECA Working Group on Annex 1*

Sterile Filtration – PUPSIT and Requirements beyond

Dr Frank Sielaff, *Regional GMP Authority Darmstadt, Germany*

Practical Application of setting up an annual Contamination Control Strategy (CCS) Assessment

Ruben van der Galiën, *GE HealthCare*
Dr Prachi Sawant Raschdorf, *GE HealthCare*

RTU

Horst Koller, *HK Packaging Consulting*
Katharina Golly, *Novartis Pharma*

"RTU + RTS Materials – GMP Requirements for the Pharmaceutical Manufacturer and Supplier Qualification"

Dr Rainer Kahlich, *Local GMP Authority of Baden-Württemberg, Germany*

Digitalisation & Artificial Intelligence

Digitalisation and AI from the Inspector's Point of View
Dr Arno Terhechte, *Local GMP Authority Münster, Germany*

Accelerate Automation Project Implementation and Reduce Risk with a Digital Twin
Rasmus Wendelboe Jørgensen, *Novo Nordisk*
Vicky Athanasiou, *Emerson*

Case Study: Design & Implementation of a new highly-automated modular OSD Production Facility at Bayer Leverkusen
Andreas Bail, *Bayer*
Anton Kopitzsch, *Glatt Ingenieurtechnik*

Integrating Digitalisation & Robotics in Pharmaceutical Manufacturing: Strategies, Challenges, and Compliance in the Digital Era
Maja Karovic, *F. Hoffmann-La Roche*
Yvonne Duckworth, *CRB*

Smart Panel – The Future of the pharmaceutical Production Process
Christoph Dechow, *Boehringer Ingelheim Pharma*
Dr Sebastian Wibbeling, *Fraunhofer-Institut for Material Flow and Logistics IML*

AI in Medical Image Processing
Daniel Wolf, *Ulm University Medical Center*

Continued Process Verification Using Automated Data Assessment
Dr Philip Hörsch, *Vetter Pharma-Fertigung*

Application of dynamic Learning Systems to increase Efficiency in Pharma Production Lines
Felix Georg Müller, *plus10 GmbH*
Martin Heitmann, *d-fine GmbH*

Live Demos
- Virtual Pharma Campus
  Pharmaplan
- Digital Assistant for Maintenance at the Shopfloor
  ZETA
- Going to Market Faster in Life Sciences, Leveraging the Emerson Digital Twin
  Emerson Automation Solutions
- Kneat Gx: Live Digital Validation Software Demonstration
  Kneat Solutions

Trends in Barrier Systems & Robotics

GMP Risk Assessment for performing the Test for Sterility in an Isolator
Dr Bettina Rietz-Wolf, *Local GMP Authority Baden-Württemberg, Germany*
Dr Timo Krebsbach, *SKAN*

Aseptic Process Simulation with a gloveless robotic Filling Line
Thorsten Häfner, *PSM*
Sebastian Hillbrand, *SKAN*

Air Velocity at Working Position and other Cleanroom (Airflow) Challenges in new Annex 1
Jörg Zimmermann, *Vetter Pharma-Fertigung*
Dr Johannes Rauschnabel, *Syntegon Technology*

Case Studies and Future Trends for Zero Human Interactions in Aseptic Filling
Dr Arne Schröder, *Vetter Pharma-Fertigung*
Tobias Resch, *Stäubli Tec-Systems*

Upgrade of integrated H2O2 Bio-Decontamination System for Production of Vial Filling Line with oRABS – Part II
Pasquale Cataldo, *Roche Diagnostics*
Kenan Kanmaz, *Optima pharma containment*

Compliance of Annex 1 Requirements for Glove Integrity Testing
Jason Creek, *Roche Diagnostics*
Kenan Kanmaz, *Optima pharma containment*

A novel Way for Measuring H2O2 in an Isolator
Paul Devuyst, *GlaxoSmithKline*
Theresa Ladwig, *SKAN*

Implementation of RABS Systems in Small Volume Manufacturing
Marta Rodríguez Vélez, *Letipharma*

A Case Study highlighting the Validation of a closed gloveless Aseptic Filling Workcell
Joachim Vereecke, *White Raven*
Brent Lieffers, *Cytiva*

Cleanroom Challenges in Ongoing Operations

Cleanroom Garments – Clothing Basics, Barrier Functions and more
Jörg Mesenich, *Mesenich Consulting*
Gabi Schmeer-Lioe, *German Institutes for Textile and Fibre Research Denkendorf (DITF)*

Qualification Study for Cleanroom Garments
Carsten Moschner, *CMC3*

Consumables and Cleanroom – Expectations and Experiences of an Inspector
Dr Daniel Müller, *Local GMP Authority Baden-Württemberg, Germany*

Disposables – Face Masks and the Like – what Protection do they really provide?
Monika Lamprecht, *Lamprecht C&C*

Cleanroom Wipes – What is Cleanroom-Compatible really?
Carsten Moschner, *CMC3*

Mopping Systems and the associated Systems
Margarete Witt-Mäckel, *Witt Hygienemanagement*

Cleanroom Gloves – the Balancing Act between Cleanroom Suitability and Personnel Protection
Monika Lamprecht, *Lamprecht C&C*
Robert Sevdas, *Dastex*

Facility Monitoring with Single-Use active viable Sampling in the daily Practice of a Contract Developer and Manufacturer – Complete and simple Solution
Dr Thomas Müller, *Recipharm*
Ivan Spiro, *Particle Measuring Systems*

Pitfalls in microbiological Cleanroom Monitoring: Common Sources of Error in Qualification and Ongoing Monitoring
Melanie Braun, *Labor LS*

Live Demos

- Facility Monitoring Systems
  Particle Measuring Systems
- Continuous and rapid Monitoring of Bacteria in pharmaceutical Grade Water
  BWT Pharma & Biotech
- Surface Properities of Stainless Steel – how the Adhesion Behavior of Products can be influenced by modifying Stainless Steel Surfaces
  Bolz Intec

Case Study: Disinfectants and their Effectiveness on various Surfaces
Dr Hans-Joachim Anders, *Novartis Pharma*

Navigating the Challenges of implementing New Annex 1 in Non-Sterile Manufacturing
Martina Gjorgjevska, *The ForceCT*
Apostol Todorovski, *Sinceritas*

Sustainability/Green GMP

How can Sustainability be integrated into a pharmaceutical Quality Management System? ... or is it already included?
Dr Andrea Bauer, *ABC&Q*

Circular Economy Opportunities for the pharmaceutical Industry
Susana Santos, *BIAL Portela & C^a*

The new F-Gas Regulation and its Impact on pharmaceutical Freeze-Drying
Thomas Beutler, *GEA*

European Union – Supply Chain Directive, Green Deal and National Regulations like the German Supply Chain Due Diligence Act
Leonie Evans, *Meisterernst Attorneys*

From Compliance to Sustainability: The Green GMP Journey
Ana Cláudia Pinho, *BIAL Portela & C^a*

Greenfield Project – A sustainable Manufacturing Building
io consulting/Erbe Elektromedizin

Process Load Profiles as a Basis for a Cost-Efficient and sustainable Design of Utility Supply Systems
Bianca Bohrer, *PSM Saar*
Peter Gross, *PGC*

Sustainability Strategies at Pekana
Dr Marius Beyersdorf, *Pekana*

Recycling and Cleanroom Garments
Carsten Moschner, *CMC3*

Sustainable Refrigeration Technologies: Overview and Implementation of innovative Air-Cooling Technology for Freeze-Drying Processes
Fabian Plaum, *Hof*
Christian Sonntag, *Roche*

Sustainable Heat and Cooling Systems – the Luna Project
Thomas Frank, *Refolution*
N.N., *Abbvie*

Understanding the "Hidden Factory" in pharmaceutical manufacturing
Dr Frank Thielmann, *Takeda*

Single-Use Systems in Sterile & Biomanufacturing

Single-Use Systems – GMP Inspector's View
Dr Daniel Müller, *Local GMP Authority Baden-Württemberg, Germany*

Single-Use Technology in biopharmaceutical Production: An Overview from USP to Fill&Finish Technologies
Prof Dr Regine Eibl, *Zurich University of Applied Sciences*

Quality Approach in Manufacturing of Single-Use Systems: How to assure Performance, Robustness, and Sterility of Single-Use System
Dr Simone Biel, *Merck*

Pharma QA/QC when using Single-Use Equipment
Dr Alicja Sobantka, *Octapharma*

Case Study Merck: Single-Use Technology in Aseptic Drug Product Manufacturing
Nicola Rutigliani, *Merck*

Case Study BioNTech: CCS for Processing Frozen Sterile Drug Products in a Single-Use Assembly
Angus Liu, *BioNTech*

Case Study Sanofi: Optimization of Single-Use Systems for Fill-Finish Manufacturing Operations to the new Requirements
Dr Rebecca Geyer, *Sanofi*

E&L Testing of Process Materials used in Bioproduction – Case Studies on Study Design and showing the practical Hurdles when performing E&L Studies
Dr Koen Smets, *Nelson Labs*

Particle Cleanliness Assessment of SUS
Gerald Dallmann, *SGS INSTITUT FRESENIUS GmbH*

Case Study: Manufacturing of a Monoclonal Antibody with SUT
Jyotsna Agnihotry, *Flavine Europe*

Live Demos

- NEW Shadow board for sterile filtration of drug product (optimized for PUPSIT)
  Cytiva
- TBA
  Merck

ATMPs – Hurdles & Achievements in Quality and Safety

GMP-PHARMA CONGRESS

Lessons learned from EMA Filling for TEMP
Dr Katja Aschermann, *Astator*

Quality Assurance of mRNA Vaccines for Human Use: the Role of the European Pharmacopoeia
Prof Dr Gerit Borchard, *University in Geneva & member of the EDQM expert group*

Industrial Scale in Vitro Expression of Bacteriophages and other Proteins
Patrick Grossmann, *Invitris*

Viral Clearance
Sandra Zucchet, *CRL*

Visible and Subvisible Particle Control for Cell Therapy From Development to Commercialisation
Dr Roman Mathaes, *Clear Solutions Laboratories*

Flexibility and Redundancy in Visual Inspection Production Planning with Small-Scale Systems
Alexander Schäfer, *Wilco*

Gloveless aseptic Fillers for small Batches and Cell & Gene Therapy Sectors: a novel Approach utilizing modular Design and magnetic Levitation Conveyors
Giacomo Guidi, *IMA Life*

Process Validation Sterile Drug Products: Strategy, Execution and maintaining the validated State
Dr Anne Orillo, *Novartis Pharma*

Challenges on the Way to becoming a Contract Manufacturer
Dr Carolin Klemm, *DKMS Stem Cells*
Simone Sonnenberg, *DKMS Stem Cells*

Vector Safety Assessment in Cell and Gene Therapy by NGS/TGS Sequencing
Dr Richard Gabriel, *ProtaGene*

Challenges and Special Requirements for GMP Inspections of ATMPs
Alexander Kammerlocher, *Local GMP Authority Baden-Württemberg, Germany*

Medical Cannabis – Cultivation, Processing, Systems & Technology

Challenges and Experiences from current GMP Inspections
Dr Rainer Gnibl, *District Government of Upper Bavaria, Germany*

The Intersection between GMP and GACP
Luis Meirinhos Soares, *CANNAVIGIA*

Cannabis Cultivation under GACP
N.N.

Drying of Medical Cannabis – Challenges for Process Validation
Tina Cacanoska, *PharmaRolly*

Contamination Control Strategy in Cannabis Manufacturing
Martina Gjorgjevska, *The Force CT*
Apostol Todorovski, *Sinceritas*

Validation /Qualification – Experiences & Lessons learned
Dr Ingrid Walther, *ECA Cannabis Working Group*

Regulatory Status and Quality Standards of Cannabinoids Manufacture
Dr Giorgia Tossi, *Linnea*

Current Challenges in Microbiological Decontamination of Medicinal Cannabis
Dr David Surjo, *GO NEXUS*

Case Study – Startup Medical Cannabis
N.N.

Case Study – Dr Reckeweg – Project
N.N.

Digitalisation in Cannabis Production
N.N.

Israel Medical Cannabis Regulation
Dr Viviana Braude, *Cronos*

Medical Cannabis Manufacturing & Compounding: Regulatory Issues
Dr Hanneke Later-Nijland, *Genome Lawyers*
Dr Monika Hupfauf, *KOCH / HUPFAUF Attorneys-at-Law*

The following close to 100 speakers from industry and authorities already have confirmed their participation (constantly updated):

Jyotsna Agnihotry
Flavine Europe, Germany, *Head of QA and Regulatory Affairs*

Dr Hans-Joachim Anders
Novartis Pharma, *Teamlead Analytical Science and Technology*

Dr Katja Aschermann
Astator, Germany, *Consultanrt*

Vicky Athanasiou
Emerson, Netherlands, *Director Process Simulation Europe*

Henning Austermann
Siegfried Hameln, Germany, *Head of Engineering*

Andreas Bail
Bayer, Germany, *PCT Lead Engineer*

Dr Andrea Bauer
ABC&Q, *Owner*

Thomas Beutler
GEA, *Senior Director Lyophilization Technology Management*

Dr Markus Beyersdorf
Pekana, *General Manager*

Dr Simone Biel
Merck, *European Field Marketing Specialist*

Bianca Bohrer
PSM Saar, *CEO*

Prof Dr Gerit Borchard
University in Geneva, *Chair of the mRNA Vaccines for Human Use Working Party at the European Pharmacopoeia*

Dr Viviana Braude
Cronos, Israel, *VP Quality and Regulations & Member of ECA's Cannabis Working Group*

Melanie Braun
Labor LS, *Lead Microbiological Services*

Tina Cacanoska
PharmaRolly, North Macedonia, *Chief Quality Officer and QP & Member of ECA's Cannabis Working Group*

Pasquale Cataldo
Roche Diagnostics, Germany, *Innovation Project & Lab Lead*

Jason Creek
Roche Diagnostics, Germany, *Expert for sterile manufacturing and lab lead*

Gerald Dallmann
SGS INSTITUT FRESENIUS GmbH, *Division Manager*

Christoph Dechow
Boehringer Ingelheim Pharma, Germany, *Head of Digital Transformation Management*

Paul Devuyst
GlaxoSmithKline, Belgium, *Senior Manager Aseptic Technologies GSK Global MSAT*

Yvonne Duckworth
CRB, USA, *Fellow of Digital Technology*

Prof Dr Regine Eibl
Zurich University of Applied Science, *Professor at the Zürich University and the platform leader for "Single-use technology" of the Swiss Biotechnet*

Leonie Evans
Meisterernst Attorneys, *Attorney and Partner*

Thomas Frank
Refolution, *CEO*

Dr Richard Gabriel
ProtaGene, *Vice President R&D*

Ruben van der Galiën
GE HealthCare, Netherlands, *Qualified Person / Pharmacist*

Dr Rebecca Geyer
Sanofi, *Head of Implementation Management, Drug Product*

Martina Gjorgjevska
The Force CT, *Quality Manager*

Dr Rainer Gnibl
District Government of Upper Bavaria, Germany, *GMP Inspector*

Katharina Golly
Novartis Pharma, Switzerland, *Senior Expert Engineering*

Peter Gross
PGC, *CEO*

Patrick Grossmann
Invitris, *CEO & Co-Founder*

Giacomo Guidi
IMA Life, *R&D Isolation Technologies*

Thorsten Häfner
PSM, Germany, *VP Business Development*

Martin Heitmann
d-fine GmbH, Germany, *Senior Manager*

Sebastian Hillbrand
SKAN, Switzerland, *Strategic Product Manager*

Dr Philip Hörsch
Vetter Pharma-Fertigung, Germany, *Director QA - Validation/ Risk Management/Trending*

Dr Monika Hupfauf
KOCH / HUPFAUF Attorneys-at-Law, *Owner*

Dr Rainer Kahlich
Local GMP Authority Baden-Württemberg, Tübingen, Germany, *GMP Inspector*

Alexander Kammerlocher
Local GMP Authority Baden-Württemberg, Tübingen, Germany, *Surveillance of medical devices*

Kenan Kanmaz
Optima pharma containment, Germany, *Technical Sales Manager*

Maja Karovic
F. Hoffmann-La Roche, Switzerland, *Network Technology Lead Robotics*

SPEAKERS

Adrian Keller
SKAN, Switzerland, *Strategic Product Manager*

Dr Carolin Klemm
DKMS Stem Cells, *Head of Production*

Horst Koller
HK Packaging Consulting, Switzerland, *CEO*

Anton Kopitzsch
Glatt Ingenieurtechnik, Germany, *Team Lead Automation Engineering*

Dr Timo Krebsbach
SKAN, Switzerland, *Strategic Product Manager*

Theresa Ladwig
SKAN, Switzerland, *Sales Engineering*

Monika Lamprecht
Lamprecht C&C

Dr Hanneke Later-Nijland
Genome Lawyers, *Partner*

Brent Lieffers
Cytiva, Canada, Senior Director, *Innovation Advocacy*

Angus Liu
BioNTech, *Associate Director CMC*

Dr Maria Loos
Johnson and Johnson, *MSAT Senior Associate Scientist*

Dr Roman Mathaes
Clear Solutions Laboratories, *CEO*

Jörg Mesenich
Mesenich Consulting, *Owner*

Carsten Moschner
CMC3, *Founder*

Dr Daniel Müller
Local GMP Authority Baden-Württemberg, Tübingen, Germany, *Head of GMP Inspectorate*

Felix Georg Müller
plus10 GmbH, Germany, *CEO and Co-founder*

Dr Thomas Müller
Recipharm, *Sterility Assurance Manager*

Dr Anne Orillo
Novartis Pharma, *Senior Validation Lead*

Ana Cláudia Pinho
BIAL Portela &C^a, S.A, *Senior Manager, Sustainability*

Fabian Plaum
Hof, *Freeze-Drying and Freeze-Thaw applications*

Dr Johannes Rauschnabel
Syntegon Technology, Germany, *Head of the process development department*

Tobias Resch
Stäubli Tec-Systems, *Key Account Manager*

Dr Bettina Rietz-Wolf
Local GMP Authority Baden-Württemberg, Tübingen, Germany, *Inspector*

Marta Rodríguez Vélez
Letipharma, Spain, *Quality Assurance*

Nicola Rutigliani
Merck, *Aseptic Production Associate Director*

Stylianos Sampanis
Sanofi-Aventis Deutschland GmbH, Germany

Yves Samson
Kereon AG, Switzerland, *Founder and Chairman*

Susana Santos
BIAL Portela &C^a, S.A, *Environment Specialist*

Dr Prachi Sawant Raschdorf
GE HealthCare, Netherlands, *Qualified Person / Pharmacist*

Luigi Scaffidi
Boehringer Ingelheim Pharma, Germany

Alexander Schäfer
Wilco, *Technical Product Manager / Project Engineer AVI*

Gabriele Schmeer-Lioe
DITF - German Institutes for Textile and Fiber Research Denkendorf, *Scientist; Head of the Cleanroom Textiles and Electrostatics Laboratory*

Dr Koen Smets
Nelson Labs, *Scientific Expert E&L*

Dr Arne Schröder
Vetter Pharma-Fertigung, Germany, *Head of Production in the area of manufacturing and filling of sterile drug products*

Robert Sevdas
Dastex, *Key Account Manager*

Dr Frank Sielaff
Regional GMP Authority, Darmstadt, Germany, *GMP Inspector*

Dr Alicja Sobantka
Octapharma, *Head Corporate Material Qualification Group*

Luis Meirinhos Soares
CANNAVIGIA, former GMP Inspector at INFARMED, Portugal, *Head of Compliance and Regulatory Affairs & Member of ECA's Cannabis Working Group*

Simone Sonnenberg
DKMS Stem Cells, *Project Manager*

Christian Sonntag
Roche, *Senior Project Manager*

Ivan Spiro
Particle Measuring Systems, *Sales Manager*

Dr David Surjo
GOC NEXUS GMBH, Germany, *CEO*

Dr Arno Terhechte
Local GMP Authority Münster, Germany, *Medicines Inspector*

Dr Frank Thielmann
Takeda

Apostol Todorovski
Sinceritas, *Head of Quality Control*

Dr Giorgia Tossi
Linnea, Switzerland, *Chief Quality Officer & Member of ECA's Cannabis Working Group*

Klaus Ullherr
Syntegon Technology, Germany, *Senior Product Manager*

Joachim Vereecke
White Raven, Belgium, *COO and Co-Founder*

Ralf Wagner
Optima pharma, Germany, *Director Sales D/A/CH, Spain, Portugal*

Dr Ingrid Walther
Chairman of the ECA Working Group on Annex 1,
Leader of the ECA Cannabis Working Group, Consultant

Rasmus Wendelboe Jørgensen
Novo Nordisk, Denmark, *Automation Lead*

Dr Sebastian Wibbeling
Fraunhofer-Institut for Material Flow and Logistics IML, Germany, *Head of Health Care Logistics*

Margarete Witt-Mäckel
Witt Hygienemanagement, *Founder*

Daniel Wolf
Ulm University Medical Center

Jörg Zimmermann
Vetter Pharma-Fertigung, *Vice President Vetter Development Service, External Affairs*

Sandra Zucchet
CRL, *R&D Scientist for Viral Clearance*

ORGANISATIONAL ISSUES

Fees
€ 790,- for the one day ticket plus VAT. These one day tickets allow you to follow any track offered that day (you can also switch between the tracks any time). They include a lunch and beverages during the tracks and in breaks as well as the free visit of the PharmaTechnica Expo and the social event on the evening of the first congress day. Charges are payable after receipt of invoice. Please note that due to the special fees for the congress, ECA membership discounts are not applicable.

PharmaTechnica Expo
Parallel to the tracks there will be the PharmaTechnica Expo. Take advantage of this opportunity to get to know new technologies, products and services at the stands of the close to 120 international exhibitors.

Location
RheinMain CongressCenter (rmcc)
Friedrich-Ebert-Allee 1 | 65189 Wiesbaden
Phone: +49 (0) 611 1729-444
E-Mail: veranstaltungsservice-rmcc@wicm.de

Contacts – Conference Tracks
For questions regarding the content of the tracks:
Cleanroom Challenges in Ongoing Operations | Sustainability/Green GMP
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European Aseptic Technologies – Annex 1 Conference | Trends in Barrier Systems & Robotics | Digitalisation & Artificial Intelligence
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Single-Use Systems in Sterile & Biomanufacturing
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Medical Cannabis – Cultivation, Processing, Systems and Technology
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ATMPs – Hurdles & Achievements in Quality and Safety
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Please note
Exhibition Visit: The PharmaTechnica Expo will also be open to visitors on both days who are not attending the Congress. Please be aware, though, that you will need to register in advance of the visit. The visit of the exhibition does not entitle you to also attend any of the tracks.

Congress Materials: Please note that there will not be any print-outs at the Congress. Instead you will receive all available presentations prior to the Congress as Downloads.

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Easy Registration

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Internet:
www.pharma-congress.com

Congress Dates

Tuesday, 8 April 2025, 09.00 - 18.00 h
Wednesday 9 April 2025, 09.00 - 17.00 h
Registration
Tuesday & Wednesday, 8/9 April 2025, 08.00 - 09.00 h

Fees

The one day ticket is available for € 790,- plus VAT. It includes participation in any track on that day and the visit of the PharmaTechnica Expo. In addition, lunch and beverages during the tracks and in breaks as well as the social event on the evening of the first congress day, 8 April is included; please mark if you would like to attend the Social Event.

Only until 31 December 2024: Save up to € 200 Early Bird Discount

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Reservation Form (Please complete in full)
Please also mark the day you plan on attending the Congress. To be able to prepare the conference rooms, we would appreciate if you also marked the track you are interested in (Please only mark one track per day).

The GMP PharmaCongress Tracks	Participation on 8 April 2025 ☐	Participation on 9 April 2025 ☐
European Aseptic Technologies – Annex 1 Conference	☐	☐
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Sustainability/Green GMP	☐	☐
Single-Use Systems in Sterile & Biomanufacturing	☐	☐
ATMPs	☐	☐
Medical Cannabis	☐	☐
Participation in Social Event	☐	n.a.
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WH/30102024