## GMP PharmaCongress 2025 – Agenda 8 April 2025

8/9 April 2025, RheinMain CongressCenter, Wiesbaden

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Time	Aseptic Technologies & Annex 1 Conference	Digitalisation & Artificial Intelligence	Single-Use Systems in Ste- rile & Biomanufacturing	Cleanroom Challenges in Ongoing Operations	Trends in Barrier Systems & Robotics	Sustainability/Green GMP	Achievements	Medical Cannabis	Time
Room	Forum 1.1	Studio 1.3	Studio 1.5	Studio 1.2	Forum 1.2	Studio 1.1	Forum 1.3	Studio 1.4	Room 9:00 h
9:00 h 9:15 h			Artific		nufacturing and Quality at Sanofi Chief Quality Officer, Sanofi				
9:30 h				Wallhäußer Innovat	tion Award Ceremony				9:30 h
9:45 h					and Award Ceremony				9:45 h
10:00 h					Coffee Break				
10:15 h		Coffee	e Break						
									10:15 h
	Quality RISK Management in Aseptic Manufacturing: Reasonable Use Dr Ingrid Walther, Pharma Consulting Walther	Accelerate Automation Project Implementation and Reduce Risk with a Digital Twin Rasmus Wendelboe Jørgensen, Novo Nordisk Vicky Athanasiou, Emerson	Single-Use Systems - GMP	Cleanroom Garments - Clothing Basics, Barrier Functions and More Jörg Mesenich, Consultant Gabriele Schmeer-Lioe, Deutsche Institute für Textil- und Faserforschung (DITF)	Air Velocity at Working Position and other Cleanroom (Airflow) Challen- ges in new Annex 1 Jörg Zimmermann, Vetter Pharma-Fertigung Dr Johannes Rauschnabel, Syntegon Technology	How can Sustainability be integrated into a Pharmaceutical Quality Management System? or is it already included? Dr Andrea Bauer, ABC&Q	Navigating the EMA Process: Key Insights from Filing for a TEMP Dr Katja Aschermann, Astator	Challenges and Experiences from current GMP Inspections	10:30 h 10:45 h 11:00 h
11:30 h	Sterile Filtration - PUPSIT and Requirements beyond Dr Frank Sielaff, Regional Authority Darmstadt, Germany	Live Demos PHARMAPLAN AG ZETA GmbH Emerson Automation Solutions	Inspector's View Dr Daniel Müller, Local GMP Authority of Baden-Württemberg Germany	Qualification Study for Cleanroom Garments Carsten Moschner, CMC3	GMP Risk Assessment for Perfor- ming the Test for Sterility in an Isolator Dr Bettina Rietz-Wolf, Local GMP Authority of Baden-Württemberg, Germany Dr Timo Krebsbach, SKAN	Circular Economy Opportunities for the Pharmaceutical Industry Susana Lima Santos, Bial	Challenges and Special Require- ments for GMP Inspections of ATMPs Alexander Kammerlocher, Local GMP Authority of Baden-Württemberg, Germany	Dr Rainer Gnibl, District Government of Upper Bavaria, Germany	11:15 h 11:30 h 11:45 h
12:00 h 12:15 h 12:30 h		Kneat Solutions			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	The new F-Gas Regulation and its Impact on pharmaceutical Freeze- Drying Thomas Beutler, GEA	Quality Assurance of mRNA Vaccines for human use: the Role of the European Pharmacopoeia Prof Dr Gerrit Borchard, University Geneva & Member of the EDQM expert group	The Intersection between GMP and GACP Luis Meirinhos Soares, formerly INFARMED & Member of ECA's Cannabis Working Group	12:30 h
12:45 h 13:00 h 13:15 h			ı Break		Lunch Break				12:45 h 13:00 h 13:15 h
14.00 h	annual Contamination Control	Case Study: Design & Implementa- tion of a new highly-automated modular OSD Production Facility at Bayer Leverkusen Andreas Bail, Bayer Anton Kopitzsch, Glatt	Single-Use Technology: An overview from USP to Fill&Finish technologies Prof Dr Regine Eibl, Zurich University of Applied Sciences	Consumables and Cleanroom - Ex- pectations and Experiences of an Inspector Dr Daniel Müller, Local GMP Authority of Baden-Württemberg Germany					13:30 h 13:45 h 14:00 h
14:30 h 14:45 h	Requirements for the Pharmaceuti- cal Manufacturer and Supplier Qualification Dr Rainer Kahlich, Authority of Baden-	Integrating Digitalization & Robotics in Pharmaceutical Manufacturing: Strategies, Challenges, and Compli- ance in the Digital Era Maja Karovic, F. Hoffmann-La Roche Yvonne Duckworth, CRB	Live Demos	Disposables - Face Masks and the Like - what Protection do they Really Provide? Monika Lamprecht, Lamprecht Consulting and Coaching	Case Studies and future Trends for Zero Human Interactions in Aseptic Filling Dr Arne Schröder, Vetter Pharma-Fertigung Tobias Resch, Stäubli Tec-Systems	The EU Green Deal – Supply Chain Due Diligence Directive (CS3D) and the German Supply Chain Act (LkSG) Leonie Evans, Meisterernst Attorneys	Challenges on the Way to becoming a Contract Manufacturer Dr Carolin Klemm, DKMS Stem Cell	Case Study 1 Cannabis Cultivation under GACP Natalie Thurner, Chemgineering Dr Michal Wojcicki, Cannerald	14:15 h 14:30 h 14:45 h
15:00 h 15:15 h 15:30 h		Coffee	Innerspace Break		Case study: E-Beam used as transfer technology for RTU Pre-filled Syringes at Pfizer Puurs on multiple filling lines Marcus Hoppe, Pfizer Manfred Holzer, SKAN	From Compliance to Sustainability: The Green GMP Journey Ana Cláudia Pinho, Bial	Industrial Scale in Vitro Expression of Bacteriophages and other Proteins Matthias Steiger, Invitris Dr Frenk Smrekar, Jafral	Drying of Medical Cannabis – Challenges for Process Validation Tina Cacanoska, PharmaRolly	15:00 h 15:15 h 15:30 h
16:00 h 16:15 h	RTU-Packaging Systems in relation to Annex I Horst Koller, HK Packaging Consulting	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	sterility	Cleanroom Wipes - What is Cleanroom-Compatible really? Carsten Moschner, CMC3	Coffee Break				15:45 h 16:00 h 16:15 h
16:45 h 17:00 h	indirect Product Contact Surface Dilemma (bowls/lanes)	Continued Process Verification Using Automated Data Assessment Dr Philip Hörsch, Vetter Pharma-Fertigung Bettina Schroeder, Vetter Pharma-Fertigung	Case Study Roche: Exploring Alternative Media for Filter Flushing: Implications for Protein Concentra- tion and Product Quality Julia Mathy, Roche Diagnostics	Mopping Systems and the Associ- ated Systems Margarete Witt-Mäckel, Witt Hygienemanagement	Implementation of RABS systems in Small Volume Manufacturing Marta Rodríguez-Vélez, Letipharma	3R Initiative Within Roche's Global QC Network Dr Sven M. Deutschmann, Roche	The Challenge of GMP Manufactu- ring of innovative Exosome-based Therapies Sandrine Mores, ExoBiologics	Update from the German Cannabis Agency Dr Anne Wolf, German Cannabis Agency (BfArM)	16:30 h 16:45 h 17:00 h
17:50 11	Discussion	Discussion	Discussion	Discussion	Discussion	Discussion	Discussion	Validation /Qualification - Experiences & Lessons learned Dr Ingrid Walther, Walther Pharma Consulting	17:15 h 17:30 h
18:00 h		Socia	l Event			Socia	al Event		



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Time	Aseptic Technologies & Annex 1 Conference	Digitalisation & Artificial Intelligence	Single-Use Systems in Ste- rile & Biomanufacturing	Cleanroom Challenges in Ongoing Operations	Trends in Barrier Systems & Robotics	Sustainability/Green GMP	ATMPs – Hurdles & Achievements	Medical Cannabis	Time
Room	Forum 1.1	Studio 1.3	Studio 1.5	Studio 1.2	Forum 1.2	Studio 1.1	Forum 1.3	Studio 1.4	Room 9:00 h
9:00 h 9:15 h			Pharmaceutical Manuf		3D Printing Process and other New Technologies				
9:30 h				Di Kalijita Shegokal Salioo, Chief Fila	rma Innovation Officer (CPO), DiHeSys				9:30 h
9:45 h									
10:00 h	Coffee Break				Coffee Break				
10:15 h	Collee Break				Collee Break				10:15 h
	Containers in an aseptic Ísolator	Digitalisation and AI from the Inspector's Point of View Dr Arno Terhechte, GMP Inspectorate Münster, Germany	Case Study Merck: Single-Use Technology in Aseptic Drug Product Manufacturing Nicola Rutigliani, Merck Healthcare	Cleanroom Gloves - the Balancing Act Between Cleanroom Suitability and Personnel Protection Monika Lamprecht, Lamprecht Consulting and Coaching	Aseptic Transfers in Small Batch Filling - Industry standards and new approaches to meet Annex 1 Thorsten Haefner, PSM Sebastian Hillbrand, SKAN	Process Load Profiles as a Basis for a Cost efficient and sustainable Design of Utility Supply Systems Bianca Bohrer, PSM Saar Peter Gross, Consulting	From personalized medicine to flexible machine solutions Vilma Methner, Optima	Regulatory Status and Quality Standards of Cannabinoids Manu- facture Dr Giorgia Tossi, Linnea	10:30 h 10:45 h 11:00 h
11:30 h	Accelerating Pharmaceutical Manufacturing: A Case Study of entering Syringe and Cartridge Fill-Finish Production Henning Austermann, Siegfried Hameln Klaus Ullherr, Syntegon Technology	AI (Artificial Intelligence) in Manufacturing Dr Monika Hupfauf, KOCH/HUPFAUF Attorneys Amir Abou Elmagd, Genome Lawyers	Case Study BioNTech: CCS for Processing Frozen Sterile Drug Products in a Single-Use Assembly Dr Yuan-An (Angus) Liu, BioNTech	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 System	A Case Study highlighting the Validation of a closed gloveless Aseptic Filling Workcell Joachim Vereecke, White Raven Brent Lieffers, Cytiva	Sustainability Strategies at Pekana Dr Marius Beyersdorff, Pekana Gabriele Brutscher, Pekana	Visible and Subvisible Particle Control for Cell Therapy From Development to Commercialization Dr Roman Mathaes, Clear Solutions Laboratories	Challenges in Microbiological Decontamination of Medicinal Cannabis Dr David Surjo, GOC NEXUS	11:15 h 11:30 h 11:45 h
12:00 h 12:15 h 12:30 h		Tempri Bausch REA El	Demos s GmbH +Ströbel ektonik		Challenges and benefits for modern and state-of-the art fill & finish equipment to reduce glove interven- tions; Dr Christian Matz, Formerly F. Hoffmann-La Roche Patrick Wieland, Bausch + Ströbel	Recycling and Cleanroom Garments Carsten Moschner, CMC3	Process Validation Sterile Drug Products: Strategy, execution and maintaining the validated state Dr Anne Orillo, Novartis	Contamination Control Strategy in Cannabis Manufacturing Martina Gjorgjevska, The Force CT Apostol Todorovski, Sinceritas	12:00 h 12:15 h 12:30 h
12:45 h 13:00 h	Yokogawa Deutschland GmbH								12:45 h 13:00 h
13:15 h	Lunch Break								
13:30 h 13:45 h 14:00 h	<ul> <li>H2O2 Ingress Study Approach for Isolator Decontamination of the AT-Vials</li> <li>Dr Maria Loos, Johnson and Johnson Adrian Keller, SKAN</li> <li>Benefits and Challenges in Develop- ing a GenAI Solution for a GxP-rele- vant Process</li> <li>Case Study: Manufacturing of a Monicoring Europe</li> <li>Dr Maria Loos, Johnson and Johnson Adrian Keller, SKAN</li> </ul>							13:30 h 13:45 h 14:00 h	
14:15 h 14:30 h 14:45 h	Container Closure Integrity Test Luigi Scaffidi, Boehringer Ingelheim Pharma	Artificial Intelligence (AI) for Discrepancy Management Dr Philipp Fey, Boehringer Ingelheim Pharma Jorge Gil-Hernandez, Boehringer Ingelheim Pharma	Case Study Sanofi: Optimization of Single-Use Systems for Fill-Finish Manufacturing Operations to the new Requirements Dr Rebecca Geyer, Sanofi	Live Demos Particle Measuring Systems BWT Pharma & Biotech GmbH Bolz Intec GmbH	Benefits of Digitalization in Sterile Testing Katharina Schlereth, Labor L+S Harald Kiesel, SKAN	Quantifying the present and future environmental sustainability of cleanrooms Justin Z. Lian, University of Leiden	Vector Safety Assessment in Cell and Gene Therapy by NGS/TGS sequencing Dr Richard Gabriel, ProtaGene	Case Study 2 – Pharmaceutical Cannabinoid Extractions: Balancing Efficiency and Quality Dr Nikos Xynos, Nomad Lab Scientific	14:15 h 14:30 h 14:45 h
15:00 h 15:15 h 15:30 h		Coffee	e Break	AVEVA	Compliance of Annex 1 Require- ments for Glove Integrity Testing Jason Creek, Roche Diagnostics Kenan Kanmaz, Optima Pharma contain- ment	Sustainable Heat and Cooling Systems - the LUNA Project Michael Eberhard, Abbvie Thomas Frank, Refolution	Viral Clearance ATMPs – What if the Product is a Virus? Sandra Zucchet, CRL	Digitalization in Cannabis Production Hannes Schubert, NESS	15:00 h 15:15 h 15:30 h
15:45 h 16:00 h 16:15 h	Highest Flexibility meets latest Regulations of Annex 1 Sébastien Trichot, Sanofi	Application of dynamic learning systems to increase efficiency in pharma production lines Felix Georg Müller, plus10 Martin Heitmann, d-fine	Particle Cleanliness Assessment of SUS Gerald Dallmann, SGS Institut Fresenius	Case Study: Disinfectants and Their Effectiveness on Various Surfaces Dr Hans-Joachim Anders, Novartis	Their aces Coffee Break				15:45 h 16:00 h 16:15 h
		AI in Medical Image Processing Daniel Wolf, Ulm University Medical Center	E&L testing of process materials used in bioproduction Dr Koen Smets, Nelson Lab	Navigating the Challenges of Implementing New Annex 1 in Non-Sterile Manufacturing Martina Gjorgjevska, The ForceCT Apostol Todorovski, Sinceritas	Barrier Systems – Current GMP Requirements Dr Daniel Müller, Local GMP Authority of Baden-Württemberg, Germany	Sustainable Refrigeration Technolo- gies: Overview and Implementation of innovative Air-Cooling Technology for Freeze-Drying Processes Christian Sonntag, Roche Fabian Plaum, Hof	Gloveless aseptic Fillers for small Batches and Cell & Gene Therapy Sectors: a novel Approach utilizing Modular Design and magnetic Levitation Conveyors <i>Giacomo Guidi, IMA Life</i>	Israel Medical Cannabis Regulation Dr Viviana Braude, Cronos Israel	16:30 h 16:45 h 17:00 h
17:15 h 17:30 h	Discussion	Discussion	Discussion	Discussion	Discussion	Discussion	Discussion	Medical Cannabis Manufacturing & Compounding: Regulatory Issues Dr Hanneke Later-Nijland, Genome Lawyers Dr Monika Hupfauf, KOCH/HUPFAUF Attorney.	17:30 h
17:45 h		End of (	Congress		End of Congress				
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