Connecting People, Science and Regulation®

### PDA Good Aseptic Manufacturing Conference 2025

Excellence in Aseptic Manufacturing: Bridging Compliance, Innovation, and Sustainability

Basel, Switzerland 22-23 May 2025 Agenda Preview

Thurs	day, 22 May 2025	
9:00	Welcome and Introduction	Falk Klar, PDA Europe
	Welcome from the Co-Chairs	Simone Biel, Merck KGaA  David Keen, Ecolab
Opening P	lenary	
	Title to be announced	GSK
	Title to be announced	Invited Regulatory Speaker
	Advancing Aseptic Fill & Finish: A Collaborative Breakthrough in Next-Generation Production	Pfizer & Syntegon
10:35	Networking Coffee Break, Exhibition	
	How Can We Address Value Chain Emissions?	ten23 health
	Interactive Questionnaire Session	
	Plenary Discussion	



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12:20	Networking Lunch Break, Poster Session, Exhibition			
13:20	LIVE Guided Poster Walk Engage with our Poster Presenters in our Exhibition Hall			
Session 1	Track A Closed Systems		Track B Enablers for Aseptic Manufacturing	
	A Risk-Based Approach for Assuring the Integrity of Closed, Single-Use Systems in Aseptic Processing	MSD	Title to be announced	Invited Speaker
	Integrity Assurance of Final Sterile Filtration Process Step With SUS	Merck KGaA	Title to be announced	Invited Speaker
	Case Study of Aseptic Processing in a Closed System in Clean Room Class C	BioNTech	The Digital Transformation of Sterility Testing: The Case for Alternative Microbiological Methods With Complex Product Formulations	Charles River
	Q&A, Discussion		Q&A, Discussion	
14:55	Networking Coffee Break & Exhibition			
Session 2	Track A First Air		Track B Decontamination	
	Title to be announced	Invited Speaker	Application and Qualification of an Automated UV Disinfection	AstraZeneca



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			Robot for Aseptic Cleanrooms	
	Title to be announced	Invited Speaker	Replace Settle Plates via Continuous Active Air Monitoring	Takeda
	Q&A, Discussion		Q&A, Discussion	
17:45	End of Conference Day 1 & Networking Event			

## Friday, 23 May 2025

Session 3	Track A  Aseptic Process Simulation		Track B	
			Particle Risk Mitigation	
09:00	EU GMP Annex 1, QRM & APS: Practical Approach & Points to Consider	Minipharm	Total Particle Count: Two Options to Be Compliant to EU GMP Annex 1—3- Way Valve Redesign, Counter Resistant to VHP	GSK
	Aseptic Process Simulation: Pragmatic and Innovative Approach	A3P	Achieving Particle Reduction via Supplier Annex 1 Strategies and Innovation in RTU/RTF Packaging	Gerresheimer
	Q&A, Discussion		Q&A, Discussion	
10:10	Networking Coffee Break, Poster Session & Exhibition			
Session 4	Track B Application for Quality Risk Management			



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	Excellence in Manufacturing Execution				
	Case Study: Syringe and Cartridge Fill- Finish Production	Siegfried Hameln	$\mathcal{E}$		Human Error Solutions
	Case Study: E-Beam Used More Than 15 Years as Transfer Technology for RTU Pre-Filled Syringes at Pfizer Grange Castle Ireland on Multiple Filling Lines	Pfizer & SKAN	Title to be annou	unced	Invited Speaker
	Q&A, Discussion		Q&A, Discussion		
11:50	Networking Lunch Break, Poster Session & Exhibition				
Closing Ple	Closing Plenary				
	Interactive Questionnaire Session				
	Title to be announced			Invited Speaker	
	Title to be announced			Invited Regulatory Speaker	



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13:45	Plenary Discussion		
14:30	Networking Coffee Break, Poster Session & Exhibition		
15:00	Passport Raffle		
	Best Poster Presentation		
	Interactive Session		
	Co-Chairs Conference Summary	Simone Biel, Merck KGaA  David Keen, Ecolab	
	Closing Remarks & Farewell	Falk Klar, PDA Europe	
16:30	End of Conference		

The agenda is subject to change without notice. Speaker invited, pending confirmation.

