26 November 2025

GMP Compliance Trends in Analytical Laboratories

Time		Time
09:00 h	Welcome and Introduction (at the Plenum)	09:00 h
09:15 h	**KEYNOTE: Phage Therapy: History, Current Challenges and Perspectives** Dr Frédérique Vieville, CEO, 5QBD-Biotech	09:15 h
09:30 h		09:30 h
09:45 h		09:45 h
10:00 h	Coffee break / Conference Exhibition	10:00 h
10:15 h		10:15 h
10:30 h		10:30 h
10:45 h	**Set-up of a QC Laboratory as Part of Greenfield Facility** Ivana Heckel, ten23 health	10:45 h
11:00 h		11:00 h
11:15 h	**Why every Factor matters in a Contamination Investigation** Jeanne Moldenhauer, Excellent Pharma Consulting	11:15 h
11:30 h		11:30 h
11:45 h	**Small CROs and Validation & Conduction of (Bio)analytical Methods** Dr Timo G. Kretzschmar, TiKrESolution	11:45 h
12:00 h		12:00 h
12:15 h	Lunch Break / Conference Exhibition	12:15 h
12:30 h		12:30 h
12:45 h		12:45 h
13:00 h		13:00 h
13:15 h		13:15 h
13:30 h		13:30 h
13:45 h	**Universal Study Design for Instrument Changes in Pharmaceutical Release Analytics** Dr Anne Ries, BI Pharma	13:45 h
14:00 h		14:00 h
14:15 h	**Auditing Audit Trails - QA vs QC Perspective** Apostol Todorovski, GMP Insiders	14:15 h
14:30 h		14:30 h
14:45 h	**Audit Trail Requirements for Digitalised GMP Laboratories** Dr Bob McDowall & Mahboubeh Lotfinia	14:45 h
15:00 h		15:00 h
15:15 h	Coffee Break / Conference Exhibition	15:15 h
15:30 h		15:30 h
15:45 h		15:45 h
16:00 h	**Evaluation of Stability Data: Extrapolation of Shelf-Life by Statistical Analysis** Dr Joachim Ermer, Ermer Quality Consulting	16:00 h
16:15 h		16:15 h
16:30 h	**Improving Analytical Procedure Transfers in the Pharmaceutical Industry** Ulla Bondegaard, Novo Nordisk	16:30 h
16:45 h		16:45 h
17:00 h	Final Discussion & Q&A Session	17:00 h
17:15 h		17:15 h
17:30 h		17:30 h
17:45 h		17:45 h
18:00 h		18:00 h