

# Table Tutorials, 18 September 2019

Tutorial descriptions can be found in the userBridge event app

Room	Session 1 8:30 - 9:10	Session 2 9:25 - 10:05	Session 3 10:40 - 11:20	Session 4 11:35 - 12:15
1 <b>Aristotelis LEFT</b>	<input type="radio"/> What's New in LORENZ docuBridge? Manuel Stein & Christian Hapke, LORENZ	<input type="radio"/> What's New in LORENZ docuBridge? Manuel Stein & Christian Hapke, LORENZ	<input type="radio"/> Hyperlinking in LORENZ docuBridge - All You Should Know Christian Hapke, LORENZ	<input type="radio"/> Hyperlinking in LORENZ docuBridge - All You Should Know Christian Hapke, LORENZ
2 <b>Aristotelis RIGHT</b>	<input type="radio"/> Current Submission Challenges in Europe Karl-Heinz Loebel, PharmaLex	<input type="radio"/> Current Submission Challenges in Europe Karl-Heinz Loebel, PharmaLex	<input type="radio"/> LORENZ Validation Services: How They Can Add Value for Our Customers Massud Fazelly & Leonhard Keller, LORENZ	<input type="radio"/> Successful Management of Large Submissions Dr. Margaret Schubert, Cato Research
3 <b>Aphrodite A</b>	<input type="radio"/> Compliance and Pragmatism: How to Create eCTD DMF Dossiers for Suppliers in USA Dr. Miriam Essid, Schott	<input type="radio"/> LORENZ Automator - Towards a Standardized Submission Dispatch Solution Hanna Gnevko & Katharina Schmitz, LORENZ	<input type="radio"/> Types of Submissions in the EU and Their Lifecycles Dr. Michael Jandke, Miltenyi Biotec	<input type="radio"/> Registration of Medical Devices in Russia and EAEU Member States Edelgard Rehak PhD, Edelgard Rehak Consulting
4 <b>Aphrodite B</b>	<input type="radio"/> LORENZ docuBridge as Global Submission Management System Anne Nørskov-Nielsen & Sara Braunstein, Ferring	<input type="radio"/> Regulatory Process Optimization with LORENZ Solutions Stefan Peev, Acino	<input type="radio"/> LORENZ Automator at Bayer: Process Optimization & Complexity Reduction Dagmar Goehmann & Kevin Gerchufsky, Bayer	<input type="radio"/> LORENZ docuBridge Features Useful in Major Submissions Michael Josten & Birgit Sorgenfrei, Bayer
5 <b>Aphrodite C</b>	<input type="radio"/> The Global Dossier Concept & Strategies for International Same Day Filing Yongyan Zhang & Kim Hyung-Jin, Bayer	<input type="radio"/> What's New in LORENZ drugTrack? Michael Eggers & Markus Pfahlert, LORENZ	<input type="radio"/> What's New in LORENZ drugTrack? Michael Eggers & Markus Pfahlert, LORENZ	<input type="radio"/> Health Authority Commitment Tracking for Marketing Authorizations Dr. Grace Ng-Krülle & Markus Pfahlert, LORENZ
6 <b>Kleoniki A Basement</b>	<input type="radio"/> Pre-publishing Submission QC Using LORENZ docuBridge FIVE Almut Jenk & Jared Albright, LORENZ	<input type="radio"/> Best Practices for Resolving Errors and Warnings in Validation Reports Freddie Day-Robinson & Parul Patel, LORENZ	<input type="radio"/> Evaluating Your Applications - The Authority's View Parul Patel & Jared Lantzy, LORENZ	<input type="radio"/> Global Regulatory Update Jared Lantzy, LORENZ
7 <b>Kleoniki B Basement</b>	<input type="radio"/> Introducing LORENZ Cloud: Why it's Desirable Wladimir Getze, LORENZ	<input type="radio"/> Regulatory Intelligence Tool for Submission Validation Dr. Margaret Hurley, Hurley Consulting	<input type="radio"/> Making Sure Your Documents are eCTD Ready Marta Estruch, InvoFarma	<input type="radio"/> Making Sure Your Documents are eCTD Ready Marta Estruch, InvoFarma
8 <b>Conference Room</b>	<input type="radio"/> Agency Round Table Dr. med Klaus Menges, BfArM & Kent Briggs, VECTOR	<input type="radio"/> Agency Round Table Dr. med Klaus Menges, BfArM & Kent Briggs, VECTOR	<input type="radio"/> Agency Round Table Dr. med Klaus Menges, BfArM & Kent Briggs, VECTOR	<input type="radio"/> Managing License Combinations due to the EU eCTD Mandate Emma Gent & Yongyan Zhang, Bayer