

# Table Tutorials 2017

Tutorial descriptions can be found in the userBridge event app

		Session 1 8:30 - 9:10	Session 2 9:20 - 10:00	Session 3 10:45 - 11:25	Session 4 11:35 - 12:15
1	Baie de Agnes Room, Ground Floor	<input type="radio"/> LORENZ docuBridge: What's new? Elsmari Eggers & Fabian Witzel, LORENZ	<input type="radio"/> LORENZ docuBridge: What's new? Elsmari Eggers & Fabian Witzel, LORENZ	<input type="radio"/> LORENZ docuBridge: What's new? Elsmari Eggers & Fabian Witzel, LORENZ	<input type="radio"/> LORENZ docuBridge: What's new? Elsmari Eggers & Fabian Witzel, LORENZ
2	Galerie, Ground Floor	<input type="radio"/> Agency Round Table Kent Briggs, Vector & Matthias Schremser, LORENZ	<input type="radio"/> Agency Round Table Kent Briggs, Vector & Matthias Schremser, LORENZ	<input type="radio"/> LORENZ docuBridge Configuration Frank Schroer, LORENZ	<input type="radio"/> LORENZ docuBridge Configuration Frank Schroer, LORENZ
3	Matisse A, First Floor	<input type="radio"/> Update eCTD submissions in Europe: current EU Module 1 specifications, the HMA's eSubmission Roadmap 2.0 and Best Practice in creating baselines Karl-Heinz Loebel, PharmaLex	<input type="radio"/> EU IDMP SPOR around the corner – have you done your homework? Dr. Dieter Schlaps, IT-Consulting Life Science	<input type="radio"/> Near Future Changes: From CESP to CESSP including eAF Beate Kienzler, Dr. Regenold	<input type="radio"/> PSUR Repository: Used cases Anna Rubik, ARAC AT Regulatory Affairs Consulting
4	Matisse B, First Floor	<input type="radio"/> LORENZ authorBridge: Your key to regulatory submission templates Antoinette Azevedo, eSubmissionSolutions	<input type="radio"/> LORENZ docuBridge ONE: Empowering Submission of INDs and DMFs as eCTD Antoinette Azevedo, eSubmissionSolutions	<input type="radio"/> LORENZ docuBridge: Replicate sequence Michael Josten, Bayer	<input type="radio"/> Case Study - Regulatory Information Management Tool Implementation Emma Gent, Bayer
5	Mont Baron, First Floor	<input type="radio"/> Getting Documents Submission Ready Marianne Mowrer & Nora Keeling, Mentara	<input type="radio"/> Getting Documents Submission Ready Marianne Mowrer & Nora Keeling, Mentara	<input type="radio"/> IND Safety Reporting to FDA – Procedures and Timelines for eCTD Compilation Dr. Veronika Alt & Dr. Sandra Oetjen, ERA Consulting	<input type="radio"/> Biotech Companies and Their Expectations for Use of eSolutions Hillary Hafeken & Virginia Smith, United Therapeutics
6	Segurane, First Floor	<input type="radio"/> LORENZ Automator: The One-Click submission Hanna Gnevko, LORENZ	<input type="radio"/> LORENZ docuBridge: Feedback session Jason Berning & Manuel Stein, LORENZ	<input type="radio"/> Global e-Submission Regulatory Updates Jared Lantzy, LORENZ	<input type="radio"/> US Module 1 Version 3.3 Jared Lantzy, LORENZ
7	Mont Alban, First Floor	<input type="radio"/> LORENZ drugTrack: Where we are today Nicole Sagner, LORENZ	<input type="radio"/> LORENZ drugTrack: Where We will be tomorrow Dr. Sebastian Knieps, Lionpharm & Markus Pfahlert, LORENZ	<input type="radio"/> LORENZ drugTrack: Where we are today Nicole Sagner, LORENZ	<input type="radio"/> LORENZ drugTrack: Where We will be tomorrow Dr. Sebastian Knieps, Lionpharm & Markus Pfahlert, LORENZ
8	Cafe / Bar, Ground Floor	<input type="radio"/> LORENZ drugTrack: Feedback session Markus Pfahlert, LORENZ	<input type="radio"/> Validation Criteria unleashed Jan Wermusch, LORENZ	<input type="radio"/> eLearning for LORENZ products Tim Ivanić, LORENZ	<input type="radio"/> eLearning for LORENZ products Tim Ivanić, LORENZ