

## Pre-Conference Short Courses (1 June 2026)

Course 1	Course 2
<p><b>Practitioner's View of Adaptive Designs</b></p> <p>Tutors: Prof. Tim Friede (University of Göttingen) and Dr. Hiya Banerjee (Eli Lilly)</p> <p><i>Covers adaptive design methodology, ICH E20, Monte Carlo simulation, Clinical Scenario Evaluation framework, DMC roles.</i></p>	<p><b>Designing Better Trials: Advanced Randomization Methods for Modern Clinical Research</b></p> <p>Tutors: Dr. Alex Sverdlov (Novartis), Dr. Yevgen Ryznik (Uppsala University), Dr. Diane Uschner (Roche) — Randomisation Working Group</p> <p><i>Covers restricted randomization, multi-arm and platform trial randomization, Response-Adaptive Randomization (RAR), regulatory perspectives.</i></p>

## Scientific Programme (2–3 June 2026)

Time	Title	Presenter
<b>DAY 1 — Tuesday 2 June 2026</b>		
09:00–10:30	<b>KEYNOTE 1: Optimality, Loss and Stepped-wedge Designs</b>	Prof. Anthony Atkinson
<b>Session 1 — Platform Trials &amp; Operational/Regulatory Adaptations (11:00–12:30)</b>		
11:00	<b>Adaptive Designs in Fast-Track Registration Processes for Digital Health Applications</b>	Liane Kluge
	<b>Estimands and Estimation Strategies for Platform Trials with Time Trends</b>	Marta Bofill Roig
	<b>Adaptive Endpoint Change in Platform Trials: How to Change If You Must</b>	Tobias Mielke
	<b>Managing Imbalance in Randomisation with Delayed Eligibility Reveal</b>	Rajenki Das

**Session 2 — Borrowing Strength for Efficient Trials: Anytime-Valid Inference, External Controls and Covariate-Informed Design (13:30–14:45)**

13:30	<b>Dynamic Borrowing and Prognostic Covariate Adjustment for RCTs With External Controls: A Neutral Comparative Simulation Study</b>	Luzia Berchtold
	<b>Group Sequential Design with Prognostic Score Matching as an Adaptive Information Use</b>	Ajsi
	<b>Anytime-Valid Inference with E-values and Confirmatory Adaptive Designs</b>	Werner Brannath

**Session 3 — Interim Monitoring, Futility & Confirmatory Programme Design (15:15–17:00)**

15:15	<b>Optimising Log-Rank Test Power in Group Sequential Trials with Non-Proportional Hazards via Event-Balanced Randomisation</b>	Atanu Bhattacharjee
	<b>Group Sequential Designs and the Two Trials Rule — Can the Approaches Be Combined?</b>	Fredrik Öhrn
	<b>Interim Decision-Making in Two Parallel Pivotal Studies</b>	Saswati Saha
	<b>Futility Bounds for Drop-The-Losers Clinical Trial Designs</b>	Dominique-Laurent Couturier
	<b>Designing Event Re-estimation: Objectives and Flexibility</b>	Yevgen Tymofyeyev

**Workshop dinner**

**DAY 2 — Wednesday 3 June 2026**

09:00– 10:00	<b>KEYNOTE 2: Multiple Questions = Multiple Testing?</b>	Dr. Cornelia Kunz
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**Session 4 — Multiplicity, Multiple Endpoints & Reproducible Testing (10:15–11:45)**

10:15	<b>Assessing Multiple Endpoints Using a Novel Software Solution in a Late-Stage Oncology Study</b>	Valeria Mazzanti
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	<b>A Non-parametric Approach for Estimating Correlation Between Log-rank Test Statistics with Applications to Conjunctive Power Calculation</b>	Anne Lyngholm Sørensen
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	<b>Optimal Weighted Tests for Replication Studies and the Two-Trials Rule with Multiple Hypotheses</b>	David Robertson
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	<b>Automated Generation of SAP Multiplicity Appendices via the appendMCP R Package</b>	Michael Grayling
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**ICH E20 Panel Discussion (12:45–13:45)**

12:45	<b>Panel Discussion: ICH E20 on Adaptive Clinical Trials — Implications and Implementation</b>	Panellists: Alex Sverdlov, Ayon Mukherjee, Sofia Villar, Andrew Thomson
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**Session 5 — Dose Optimization & Precision Oncology Trial Designs (14:00–15:15)**

14:00	<b>PREDOSE: Pharmacometrically-Refined Early-phase Dose Optimization Design for Oncology Study Enhancement</b>	Damitri Kundu
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	<b>Bayesian Non-Inferiority Trials for Dose Finding</b>	Connor Fitchett
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	<b>Incorporating Clinical Belief and Restrictions in Umbrella Trial Designs</b>	Brian Terry
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**Session 6 — Algorithmic Adaptive Design (15:30–17:00)**

15:30	<b>Adaptive Clinical Trials Based on Design-Optimal E-values: A Novel Approach for Efficient and <math>n_{\max}</math> anytime-valid inference in single-arm trials with binary data</b>	Stef Baas
	<b>Tuning Bayesian Response-Adaptive Randomization Hyperparameters: Monte Carlo Evaluation and Optimization of Power and Patient Benefit</b>	Philipp Besendorfer
	<b>Adaptive Design Strategies for Ethical and Efficient Clinical Trials Using Simulated Annealing</b>	Rosamarie Frieri
	<b>Confidence Sets for a Level Set in Linear Regression</b>	Fang Wan