

Advances in Time-to-Event Analyses in Clinical Trials

8:15am-8:45am	Check-in and breakfast for in-person attendees
8:45am-9:00am	Opening/Logistics – Mary Putt, PhD, ScD, University of Pennsylvania Welcome – Susan Ellenberg, PhD, Professor Emerita, University of Pennsylvania
Morning Session: M	oderator Wei-Ting Hwang, PhD, <i>University of Pennsylvania</i>
9:00am-9:30am	Overview: Non-Proportional Hazards and Composite Endpoints Devan Mehrotra, PhD, <i>Merck and Co., Inc.</i>
9:30am	Lu Tian, ScD, Stanford University On estimating the survival distribution of the duration of response
9:55am	Zhenzhen Xu, PhD, Food and Drug Administration Design of immuno-oncology studies involving biomarker-defined subgroups
10:20am	Fan Li, PhD, <i>Yale University</i> Multiply robust estimation of causal effects with noncompliance and time-to-event outcomes
10:45am-11:15am	Break
11:15am-12:00pm	Panel Discussion: (Pamela Shaw, PhD, Kaiser Permanente WHRI – AM Online Facilitator) Pralay Mukhopadhyay, PhD, Otsuka Pharmaceutical Douglas Schaubel, PhD, University of Pennsylvania Mei-Cheng Wang, PhD, Johns Hopkins
12:00pm-12:30pm	Q&A Session
12:30pm-1:45pm	Lunch
Afternoon Session:	Moderator Yimei Li, PhD, <i>University of Pennsylvania</i>
1:45pm-1:50pm	Welcome back for PM session
1:50pm	Terry Therneau, PhD, <i>Mayo Clinic</i> Multi-state models for trial data
2:15pm	Lu Mao, PhD, <i>University of Wisconsin</i> Non- and semi-parametric analysis of composite time-to-event endpoints
2:40pm	Anne Eaton, PhD, <i>University of Minnesota</i> Statistical approaches for component-wise censored composite endpoints
3:05pm	Richard Cook, PhD, University of Waterloo
	Estimands in clinical trials with complex life history processes
3:30pm-3:45pm	Estimands in clinical trials with complex life history processes Break
3:30pm-3:45pm	Break Panel Discussion: (Bryan Blette, PhD, University of Pennsylvania – PM Online Facilitator) Ionut Bebu, PhD, George Washington University Rebecca Betensky, PhD, New York University