



# PSI One Day Meeting: Time-to-Event and Recurrent Event Endpoints

Novartis, Basel, Switzerland

Wednesday 14th September 2016, 08:30 - 16:00

This exciting one-day workshop will cover a wide range of statistical aspects relating to event-driven trials. We have assembled a group of very knowledgeable speakers, who will share their thoughts, ideas and experiences, including case studies, on a range of particular issues relating to planning, conduct and analysis of time-to-event and recurrent event trials.

The first half of the day will be dedicated to time-to-event endpoints, with the afternoon focusing on recurrent event endpoints. Each session will be concluded with a discussion by Prof. Dr. Armin Koch (Hannover Medical School, Germany).

**Informative censoring in a rare disease:  
a regulatory experience in PAH**

Lilla Di Scala, Actelion

**Unblinded sample-size reassessment in  
time-to-event clinical trials**

Dominic Magirr, Astra Zeneca

**Analyzing non-monotonous time-to-  
event outcome probabilities in  
randomized clinical trials**

Tobias Bluhmki, Universität Ulm

*Q&A with Armin Koch,  
Hannover Medical School*

**The Analysis of Recurrent Events:  
A Summary of Methodology**

Jennifer Rogers, University of Oxford

**Recurrent Event Data Endpoints in  
Chronic Heart Failure Studies:  
What is the Estimand of Interest?**

Mouna Akacha, Novartis

**Sample size & interim analysis  
considerations for recurrent event data  
analyses**

Ekkehard Glimm, Novartis

*Q&A with Armin Koch,  
Hannover Medical School*