Date	Topic
2 October	Quantitative Decision Making (QDM) in the Clinical Drug Development
9 October	SAS, R, Python – Differences, Similarities and Potentialities of Widely Used Stats Software
16 October	Clinical Trial Management during the COVID-19 Pandemic – Guidelines and Practical Impacts
23 October	Taxonomy of Subgroup-related Analyses and Issues in Clinical Trials
30 October	Data Visualization for a Better Communication with Non-Stats People

IBIG Forum

October 2020   GoToMeeting

The Friday Webinar h. 10,00 – 12,30

SIMeF
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IBIG FORUM AGENDA

2 October	Quantitative Decision Making (QDM) in the Clinical Drug Development Chair: Marco Costantini, GSK Vaccines	
10.00 – 11.00	Quantitative decision-making: rationale, frameworks and probabilities of success	**Gaëlle Saint Hilary, Servier & Politecnico Turin**
11.00 – 11.30	A GSK-developed R-Shiny app as an interactive tool for QDM	**Luca Grassano, GSK Vaccines**
11.30 – 12.00	A novel measure of drug benefit–risk assessment based on Scale Loss Score (SLoS)	**Pavel Mozgunov, Lancaster University**
12.00 – 12.30	Assurance in vaccine efficacy clinical trial design based on immunological responses	**Andrea Callegaro, GSK Vaccines**

9 October	SAS, R, Python – Differences, Similarities and Potentialities of Widely Used Stats Software Chairs: Angela Gambioli, GSK Vaccines – Veronica Sciannameo, University of Padua	
10.00 – 10.40	Using R for Reporting and Analyzing Clinical Trial Data	**Daniele Bottigliengo, University of Padua**
10.40 – 11.20	Learn, Prototype and Deploy with Ease: A Case for Python in Research	**Arjun Magge, University of Pennsylvania**
11.20 – 12.00	A population-based statistical analysis using SAS-EG: the case of COVID-19 in Friuli Venezia Giulia	**Michele Gobbato, ARCS Friuli Venezia Giulia**
12.00 – 12.30	Round table and discussion	**Angela Gambioli & Veronica Sciannameo**

16 October	Clinical Trial Management during the COVID-19 Pandemic – Guidelines & Practical Impacts Chair: Giulia Zigon, GSK Vaccines	
10.00 – 10.35	Clinical Monitoring during Covid Time: What we learnt to anticipate the Future	**Roberto Vallalta, GSK**
10.35 – 11.10	The Impact of COVID-19 on Estimands and Estimation	**Thomas Drury, GSK**
11.10 – 11.20	Break	
11.20 – 11.55	Impact of Covid-19 on Assessment of Safety in Clinical Trials	**Jennifer Shannon, GSK**
11.55 – 12.30	Mapping COVID-19 related data into SDTM	**Stefania Mirandola, Chiesi Farmaceutici**

23 October	**Taxonomy of Subgroup-related Analyses and Issues in Clinical Trials** Chair: Fabio Montanaro, Latis	
10.00 – 10.35	Taxonomy of exploratory subgroup-related analyses in clinical trials Part 1: Pre-specified analysis	**David Svensson, Astrazeneca**
10.35 – 11.10	Taxonomy of exploratory subgroup-related analyses in clinical trials Part 2: Data-driven ML analysis	**David Svensson, Astrazeneca**
11.10 – 11.20	Break	
11.20 – 11.55	All the things I wish I had known before I tried to detect a subgroup	**Nicole Kraemer, MorphoSys**
11.55 – 12.30	An introduction to Optimizing Treatment for a given patient: ITR analysis	**Paolo Eusebi, UCB**

30 October	**Data Visualization for a Better Communication with Non-Stats People** Chairs: Luca Grassano, GSK Vaccines – Andrea Nizzardo, Menarini Ricerche	
10.00 – 10.35	Clinical Data Reporting: from standard Tables, Listings and Figures to advanced and interactive Data Visualization	**Nicola Celli, Blue BI Anna Proserpi, Blue BI Alberto Romanelli, SAS**
10.35 – 11.10	A SAS graphical macro interface for descriptive tables in clinical trials	**Davide Soranna, Istituto Auxologico Italiano**
11.10 – 11.20	Break	
11.20 – 11.55	Innovative graphical solutions to explore and visualize clinical trial data with R Shiny	**Valentino Conti, GSK Vaccines Institute for Global Health**
11.55 – 12.30	An interactive platform to aggregate and display data from multiple sources	**Grace Hsu, Cytel**