

ConSPIC 2016 Agenda

September 22, 2016 - Day 1

Time	Track 1 Hall 1	Track 2 Hall 2	Track 3 MR-05	Track 4 MR-06
0900 - 1030	Registration			
1030 - 1100	Inaugural talk			
1100 - 1230	Plenary session- Jim Baker			
1230 - 1330	Lunch			
1330 - 1355	P001: Multivariate Analysis - A Statistical approach from a Non-Statistician <i>-Tamilselvi Senthilkumar</i>	P005: Associated Persons (AP) Domains: I collect what you have not noticed <i>-Padmasree Sirigireddy</i>	S001: Systematic Reviews: Has it been a game changer in clinical research? <i>-Dr.Sreekumar Nair (Invited talk)</i>	P009: Biomarkers as Clinical Endpoints <i>-Vamsidhar Terala</i>
1355 - 1420	P002: Incidence Rate in Patient Years <i>-Mohan K K</i>	P006: Multiple Baseline Calculation in Programmers Perspective: Traditional vs ADAM approach <i>-Sunil Ganeshna</i>	S002: LATENT CLASS ANALYSIS USING PROC LCA <i>-Anitha Cecelia</i>	P010: Adverse Event: Intensity behind the scenes <i>-Neha Srivastava</i>
1420 - 1445	P003: Database Go Live and SDTM Test Dataset ready (Study Startup) - Challenges and Options: <i>-Bhaskar Debnath</i>	P007: Vertical ADaM, a long look <i>-V Vishnu BabuNaidu</i>	S003: Forecasting the efficacy in Longitudinal studies using change point mixed models <i>-Priya Diana Dsilva</i>	P011: Implementation and Applicability of PROC FCMP in Creating Dynamic Custom SAS® Functions <i>-Vikramaditya Yandapalli, Ajay Yalwar, Prathamesh Athavale</i>
1445 - 1510	P004: Playing with Patients Diary Data <i>-Srihari Vadlakonda</i>	P008: An Innovative approach to automate ADaM dataset generation using Pseudocode <i>-Yajurvanan Chinnasamy</i>	S004 Insights on QT/QTc studies in clinical trials: Evolution, Elaboration and Evaluation <i>-Ramiya Ravindranath, Bhargav Modi, Jairaj Ramaiah</i>	P012: Overview of Reporting Adverse Events of Special Interest <i>-Meghana Kulkarni, Shelendra Singh Pawar</i>
1510 - 1540	Tea Break			
1540 - 1605	P013: PK/PD - Foremost Milestone in Drug development <i>-Vidya.R</i>	P017: Delivering SDTM Datasets by FPI- Challenges & Possible Solutions <i>-Pallavi Uttarwar</i>	S005- Beta regression models for analyzing QOL data <i>-Dr. Pritam Gupta</i>	P020: Challenges In Assignment Of 'EPOCH' In Partial Blinded Crossover Trials <i>-Sagar Das</i>

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1605 - 1630	P014: Exposure Response Analysis using Proc Logistic <i>-Harish Kumar</i>	P018: The BRIDG MODEL: DAM for regulated clinical research <i>-Naveenpal Arumugam</i>	S006: Sample size determination in clinical trials with multiple correlated co-primary endpoints <i>-Suryakant Somvanshi</i>	P021: Can we automate human life value? <i>-Rohith B.C</i>
1630 - 1655	P015: Study Data Reviewer's Guide (SDRG) - What is it? Why we need them? <i>-Dr Shanmugavel S Sundaram</i>	P019: SDTM Trial Design Models- A Sneak Peek <i>-Rathan Kumar Rangu, Lalitha Peddiboyena</i>	S007: Do we really need high sample size and power for BE studies? <i>-Chandrasekhar Bhupathi</i>	P022: Pharmacokinetics: Compartmental Model and Data handling, Challenges in CDISC world <i>-Marshal Chettiar, Ajay Daparthi, Prabhatha Mateti</i>
1655 - 1720	P016: Program DATA Vector (PDV) Essentials: How to Think Through the SAS® DATA Step <i>-Chirayu Patel</i>		S008: Dose Escalation Designs <i>-Ramakrishna Battula, Paridhi Jain, Shital Pokharkar</i>	P023: From Basic to Advance, ODS is all you need to know <i>-Abdulkadir Lokhandvala, Shabbir Bookseller</i>
1800 - 1845			IASCT Annual General Meeting	

September 23, 2016 - Day 2

Time	Track 1 Hall 1	Track 2 Hall 2	Track 3 MR-05	Track 4 MR-06
0915 - 1030	Plenary Session- Anandaraj Thangappan			
1030 - 1100	Tea Break			
1100 - 1125	P024: Beyond programming in survival and difference in proportion analysis <i>-Abhramoy Mandal, Avinash Bandi</i>	P028: Traceability Checks For CDISC ADaM <i>-Obulpathi Naidu K</i>	S009: Modelling Efficacy And Toxicity In Oncology Dose-Finding Studies <i>-Yajnaseni Chakraborti</i>	A001: Quantitative Signal Detection with Post-Marketing Data using R and Tableau <i>-Sridhar Punnamurthi</i>
1125 - 1150	P025: e-QC of Tables & Listings made validator Job easy! <i>-Muthukumaran Kannan</i>	P029: Baseline implementation challenges in ADAM BDS Datasets <i>-Asha Antony</i>	S010: Bayesian predictive approach to interim monitoring in clinical trials <i>-Ashwini Shenoy</i>	A002: Safety Signal - How far are we from detecting it? <i>-Pratibha Jalui</i>
1150 - 1215	P026: Statistical programming challenges in analyzing data from adaptive clinical trials <i>-Preeti Malik</i>	P030: SDTM mapping challenges in Multiple Myeloma trials: A Case Study <i>-Smitha Joseph</i>	S011: A Look at the Two Stage Multi-Arm Design <i>-Soorma Das</i>	A003: Using Analytics in Monitoring Patient Safety <i>-Arnab Sengupta</i>
1215 - 1240	P027: Landscape of IO Therapies in Clinical Trials <i>-Karthik Chandrasekhar.P, Aditya Shah</i>	P031: Evolution of SDTM, General Pit falls and Best practices <i>-Ranjan Routray, Shrishaila Patil K, Lekshmanan S</i>	S012: Seamless two stage adaptive design in clinical trials <i>-Gordhan Bagri</i>	A004: Risk Based Monitoring in Clinical Trials with Tools – Be SMART <i>-Venkatesh Krishnamurthy, SatyaVyshnavi Thondapu, Sravan Kumar Nagi Reddy, Milan Bhagat</i>
1240 - 1330	Lunch			
1330 - 1355	P032: What's my name? My name is _____ (Fun with SAS functions) <i>-Soujanya Konda, Suresh Kumar K</i>	P036: SDTM IG 3.3 - New Domains and Variables <i>-Prashant Suthar</i>	S013- Phase II Clinical Trial Design Incorporating Toxicity Monitoring <i>-Dr.Shesh Rai (Invited talk)</i>	A005: Predicting Future Course of Clinical Trials <i>-Rohan Sathe</i>
1355 - 1420	P033: End-to-End Information Flow in Clinical Endpoint Adjudication <i>-Pravinkumar D</i>	P037: The Metadata Driven Approach - When It Fails <i>-Tushar Sakpal</i>	S014: Two sample Comparisons Involving Zero-Inflated Continuous Data in Clinical Trials <i>-Dr. H. V. Kulkarni</i>	A006: Big Data Analysis in Clinical Trials <i>-Rajan Josephraj Paul</i>

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1420 - 1445	P034 Systole and Diastole of a Rescue Study - Programmers Monitoring Report <i>-Hari Prasad</i>	P038: Hidden Forces that boost the power of Open-CDISC: PERL , SAS and VBA <i>-Megha Agarwal</i>	S015: Comorbidity as prognostic factor in clinical trials in patients with hematologic indication <i>-Tuli De</i>	A007: Forecasting Trial Costs at pre-Clinical Development Stage <i>-Manigandan Ramkumar</i>
1445 - 1510	P035: Creating and Customizing the Kaplan-Meier Survival Plot in PROC LIFETEST <i>-Neha Shah, Pavan Kumar KP</i>	P039: A Practical Approach to create Define.xml and Define.PDF <i>-Samundeeswari, Hussain Sabir, Soma Sekhar K</i>	S016: Importance of Randomly selected blocks in block randomization <i>-Meera Mohan</i>	A008: Data Driven Approach to identify scientific misconduct in clinical operation <i>-Vikas Sharma, Deepali Pilankar, Swati Rizhwani</i>
1510 - 1610	Tea Break and Student Poster Competition			
1610 - 1635	P040: How statistical programmers can be prepared for the FDA Advisory Committee Meeting? <i>-Anshul Sinha</i>	P044: Jumpstart - A Quick Data Review Tool <i>-Rashmi Seta</i>	S017- A Spline Enhanced Population Pharmacokinetic Ordinal longitudinal Model subject to Informative Dropout <i>-Dr.Arindam Chakraborty (Invited talk)</i>	S021: Securing Privacy by Minimizing Disclosure Risk in De-identified Data <i>-Debasis Dey</i>
1635 - 1700	P041: RELREC - SDTM Programmer's Bermuda Triangle <i>-Charumathy Sreeraman</i>	P045: Utility to automate the creation of datasets with metadata from Specification and simplify the process of updating datasets for data transfers. <i>-Kiruthiga Mohan</i>	S018: Covariate Selection in Model Building <i>-Aditi Marathe, Garima Joshi</i>	S022: Modeling Unstructured Covariance Matrix: When it works and when it doesn't <i>-Hitendra Nath Pandey</i>
1700 - 1725	P042: Learning best practices in developing project level macros the hard way <i>-Kartik Rajan</i>	P046: Robust macro for issue tracking and to check the Data and Metadata updates between data transfers <i>-Shivaram Sharma</i>	S019: Tipping point as a sensitivity analysis <i>-Neha Pandey</i>	S023: Exploring the underlying distribution of Pharmacokinetic parameters <i>-Abhinandan Chakraborty</i>
1725 - 1750	P043: Handling missing data by adding dummy records or Imputation <i>-Sridevi, Vidya Mandavkar</i>	P047: A Dynamic Tool To Effectively interpret clinical outcomes <i>-Shilpakala Vasudevan, Mangala Hiriyannaiah</i>	S020: Statistical Method for Identification of Biomarker Driven Subgroups <i>-Chaitali Pisal</i>	S024: Different imputation methods use in Clinical Trials <i>-Arijit Sarkar, Bristi Bose, Preeti Kumari, Pankaj Tiwari</i>
1900 onwards	DJ Party and Dinner			

September 24, 2016 - Day 3

Time	Track 1 Hall 1	Track 2 Hall 2	Track 3 MR-05	Track 4 MR-06
0800 – 1000	Poster session			
1000 – 1130	Plenary session - Prof. Ashis SenGupta			
1130 – 1200	Tea Break			
1200 – 1225	P048: A peek behind the curtain: SAS system options <i>-Ajeet Raut</i>	P051: Strategies to Overcome the future resourcing demands of SAS Programmers in Clinical Research Industry - An Overview <i>-Dinesh Kumar</i>	S025: Gatekeeping Strategies <i>-Suresh Kumar Kothakonda</i>	P054: Exploring Oncology Domains <i>-Senthil Yuvaraja</i>
1225 – 1250	P049: Think beyond SAS macros start programming with Proc Lua <i>-Aravindan Karunakaran</i>	P052: The seven year itch <i>-Korak Datta</i>	S026: Different approaches for handling treatment switching for reporting in clinical studies <i>-Ravinder Arakati</i>	P055: Clinical Trial Data Anonymization Guidelines and Challenges <i>-Farooq Ali, Kedar Maheshwari</i>
1250 – 1315	P050: OSI-BIMO for FDA submission- Programmers approach in preparing it <i>-Puli Raju Mandati</i>	P053: Mindfulness - The transition from Tibetan monasteries to the corporate boardrooms, a roadmap for a better well-being in the age of disruption <i>-Nishanth N</i>	S027: Use of Propensity Score Matching in non-interventional study <i>-Surbhi Vijay, Ashutosh Mishra, Prashant Kulkarni, Shyam B Tiwari</i>	P056: The Tangled Tale - Painful process in laboratory data integration, reconciliation and review <i>-Manohar Naidu, Praveenraj Mathivanan</i>
1315 – 1400	Lunch			
1400 - 1425	P057: CLOPPER-PEARSON CONFIDENCE INTERVAL USING PROC FREQ - LIMITATION AND WORKAROUND <i>-Anik Chatterjee</i>	P060: Managing Non-CRF Data Efficiently - SDTM Perspective <i>-Kirti Srivastava</i>	S028- A class of Covariate-Adjusted Response-Adaptive Allocation Designs for Multi-treatment Binary Response Trials <i>-Dr.Atanu Biswas (Invited talk)</i>	
1425 - 1450	P058: Tag your TAUGS <i>-Praveenrao Polsani;Kumaran Selvaraj</i>	P061 Working with Define.XML V2.0.0 in SDTM environment <i>-Vijay Reddy</i>	S029: Analysis of Competing Risk Data with Masked Cause-of-Event <i>-Ashok Kumar Singh</i>	

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1450 - 1515	P059: R we ready for 'R'? Validation and review made easy - <i>Satheesh Kantam, Prajakta Anil Chavan</i>	P062: OpenCDISC to Enhanced Pinnacle 21 Reports using SAS - <i>Amitkumar Kawle, Kishore kumar Paramkusham, Abhinav Kumar, Nitin Suryawanshi</i>	S030: Modelling and analysis of recurrent event data - <i>Rajan Sareen, Sonam Singh, Keerthana Palwai</i>	
1515 - 1545	Tea Break			
1545 - 1715	Valedictory - Dr. Ramesh Hariharan			
1715 - 1745	Awards and Closing			