Disso India Hyderabad 2018
PROGRAMME

DAY 1 – June 28, 2018		
SESSION	**TIME**	**TITLE & SPEAKER**
	8.00 am -9.00 am	Registration
	9.00 am -10.00 am	Inauguration
SESSION I	10.00 am -11.00am	**Recent Drug Regulation and the Pharmacop(o)eias** Dr.Roger L.Willams, Ex-USFDA, EX CEO,USP,NDA Partners LLC, USA.
	11.00 am – 11.30am	**Application of USP 4 in dissolution testing of complex parenterals and IVIVC** Dr. Namita Tipnis,University of Connecticut, USA
	11.30 am-12.00 pm	**TEA BREAK AND POSTER SESSION**
SESSION II SPONSORED BY NOVARTIS	12.00 pm-12.30 pm	**A science based approach to simplify regulatory pathway for a complex generics using IVRT** Dr.Vinod Shah Ex-US FDA, Maryland, USA
	12.30 pm-1.00 pm	**IVRT for topical dosage forms as BA/BE waiver** Kailas Thakker COO & Co-founder, Tergus Pharma, USA
	12.30 pm-1.00 pm	**Challenges and development of Trans Dermal Patches in perspective of Dissolution Studies.** Dr. Dange Veerpaneni President/CEO, Sparsha Pharma, Hyderabad, India
	1.00 – 1.30 pm	Panel Discussion
	1.30 pm- 2.30 pm	**LUNCH BREAK AND POSTER SESSION**
SESSION III	2.30 pm- 3.00 pm	**Excipients Quality & trouble Shooting Topic to be received** Seema Trivedi, GM Technical,Anshul Life Sciences
	3.00 pm-3.30 pm	**Life cycle management of analytical methods** Saji Thomas, Director, Par Pharmaceutical, USA
	3.30 pm- 4.00 pm	**TEA BREAK AND POSTER SESSION**
SESSION IV	4.00 pm-4.30 pm	**Opportunities & Practical Challenges in Dissolution testing of Generic Drugs** Dr. Kalyanaraman Senior Director – AR&D, Dr. Reddy's Laboratories Ltd, India
	4.30 pm- 5.00 pm	**Pro-drugs Challenges In Dissolution** Chuei Wuei Leong, Founder and Principal consultant, CEXA Consultancy Sdn Bhd, Malaysia
	5.00 pm- 5.30 pm	Panel Discussion
	5.30pm- 6.00 pm	**POSTER SESSION**

Disso India Hyderabad 2018
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DAY 2 – June 29, 2018		
SESSION	**TIME**	**TITLE & SPEAKER**
SESSION V	9.30 am -10.00 am	**Tracking the effect of drug interactions with in vitro dissolution studies** Prof. Imre Klebovich Profesor of Pharmaceutics, Semmelweis University Department of Pharmaceutics, Budapest,Hungary
	10.00 am -10.30 am	**The Future of Bioequivalence: changing paradigms at the FDA** Prof. Jennifer Dressman, Professor and Director of the Institute of Pharmaceutical Technology, Goethe University, Germany
	10.30 am – 11.00 am	**TEA BREAK AND POSTER SESSION**
SESSION VI	11.00 am -11.30 am	**PBPK Modelling and Simulation: An In silico - In Vivo Bridge for Efficient Formulation Development** Dr. Anant Ketkar, Sun Pharma Advanced Research Company, India
	11.30am -12.00 pm	**In vitro testing for non-conventional dosage forms** Samir Haddouchi Managing Director, SPS Pharma Services, Orleans, France
	12.00 pm- 12.30 pm	**Current Regulatory Observations & Trends related to Dissolution Studies** Vijay Kshirsagar, Director and CEO, TRAC Pharma Consulting, Mumbai, India
	12.30 pm- 1.00 pm	Panel Discussion
	1.00 pm -2.00 pm	**LUNCH BREAK AND POSTER SESSION**
	2.00 pm- 2.30 pm	**Predicting Bioavailability from Dissolution: Unlocking the mystery(ies) !!** Umesh Banakar, Professor and President, Banakar Consulting Services, USA
SESSION VII	2.30 pm -3.00pm	**New Developments in Fully Automated Dissolution Technology** Grove Geoffrey , Product Manager and Application Scientist, SOTAX Corporation, USA
	3.00pm -3.30 pm	**Chromatography automations in the QC lab** Kunihiko Koriyama, Dy GM, Shimadzu India

Disso India Hyderabad 2018
PROGRAMME

	3.30pm- 4.00pm	**TEA BREAK AND POSTER SESSION**
SESSION VIII	4.00 pm – 4.30 pm	**Dissolution Failures: Interesting Scenarios** Sandip Tiwari, Fellow, Manufacturing Science and Technology, Actavis Laboratories FL, Inc., Florida, USA
	4.30 pm – 5.00 pm	Panel discussion
	5.00 pm - 5.15pm	**POSTER AWARDS &CONCLUDING REMARKS**