



See differences between formulations that apparatus 1 and 2 won't show



SOTAX CE 7smart USP apparatus 4

The Flow Through dissolution method is approved for use and accepted by the FDA and USP for a wide range of formulations and applications including :

Small Volume Dissolution • IVIVC Studies • XR/MR Solid Dose • APIs, Powders, Granules
Injectable Suspensions • Poorly Soluble Compounds • Soft Gels • Suppositories • Implants
Stents • Drug Coated Medical Devices • Liposomes • Microspheres • Nanoparticles

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Date : 12th & 13th September, 2019

Venue : Radisson® Hotel, Chandigarh Zirakpur

Programme Schedule

Disso India - Chandigarh 2019 Programme

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TIME	TITLE & SPEAKER
DAY 1 : 12/09/2019	
09.00 am - 10.00 am	Inaugural Session of Disso India 2019
Module 1 : Importance of Dissolution during formulation development	
10.00 am - 10.30 am	BCS, Dissolution and Biowaiver Dr. Vinod P. Shah , Ex-USFDA, Pharmaceutical Consultant, USA
10.30 am - 11.00 am	Dissolution enhancement by modulating the physical form of the API Dr. Raj Suryanarayanan , PhD, Professor and William and Mildred Peters Endowed Chair, Department of Pharmaceutics, College of Pharmacy, University of Minnesota
11.00 am - 11.30 am	MORNING TEA
11.30 am - 12.00 pm	Effect of surface anisotropy of crystal habits on dissolution performance Dr. Arvind Bansal , Professor & Head (Dept. of Pharmaceutics) NIPER SAS Nagar, Panjab
12.00 pm - 12.30 pm	PANEL DISCUSSION 1
12.30 pm - 01.00 pm	Excipients' Role in Modifying Dissolution Ms. Seema Trivedi , GM Technical, Anshul Life Sciences, Mumbai, India
01.00 pm - 02.00 pm	LUNCH (PARALLEL POSTER SESSION)
02.00 pm - 02.30 pm	Critical process parameters affecting dissolution Dr. Deo Narain Dikshit , Director, Aqex Pharmsolutions
02.30 pm - 03.00 pm	Implementation of QbD for dissolution testing Dr. B S Bhoop , Prof. Emeritus, Panjab University
03.00 pm - 03.30 pm	Solubility to permeability to bioavailability : Connecting the dots Dr. Namita Tipnis Varde , Ph.D., Application Scientist, Electrolab India Pvt. Ltd.
03.30 pm - 04.00 pm	EVENING TEA
04.00 pm - 04.30 pm	Dissolution Qualification: general concerns and indirect importance of automation Michel Magnier , Product Manager and Application specialist, SOTAX AG, Switzerland
04.30 pm - 05.00 pm	PANEL DISCUSSION 2

TIME	TITLE & SPEAKER
DAY 2 : 13/09/2019	
Module 2 - Dissolution of Novel Drug Delivery Systems	
09.00 am - 09.30 am	Poster Awards and Felicitation
09.30 am - 10.00 am	Topical drug classification system (TCS) Dr. Vinod P. Shah , Ex-USFDA, Pharmaceutical Consultant, USA
10.00 am - 10.30 am	Importance of dissolution studies in evaluation of DPIs Dr. Paul W S Heng , GEA-NUS Pharm Processing Res Lab, Dept of Pharmacy, National University
10.30 am - 11.00 am	Dissolution of topical products Dr. Rajeev Raghuvanshi , Dr. Reddy's Lab
11.00 am - 11.30 am	MORNING TEA
11.30 am - 12.00 am	Practical approaches for dissolution testing of nano formulations Prof. Dr. Padma Devarajan , Institute of Chemical Technology, (ICT), Mumbai, India
12.00 pm - 12.30 pm	Other NDDS / differentiated products + Long acting parenterals Samir Haddouchi , Managing Director, SPS Pharma Services, Orleans, France
12.30 pm - 01.00 pm	PANEL DISCUSSION 3
01.00 pm - 02.00 pm	LUNCH
Module 3 - Regulatory and IP considerations	
02.00 pm - 02.30 pm	Regulatory aspects of dissolution Vijay Kshirsagar , Director and CEO, TRAC Pharma Consulting, Mumbai, India
02.30 pm - 03.00 pm	Patent opportunities with dissolution studies Dr. Umesh Banakar , Professor and President, Banakar Consulting Services, USA
03.00 pm - 03.20 pm	Efficiently Automated UV/VIS Spectroscopy Atul Yelpale , Product Specialist- Anachem Mettler-Toledo India Pvt. Ltd.
03.20 pm - 03.50 pm	PANEL DISCUSSION 4
03.50 pm - 04.00 pm	VOTE OF THANKS
04.00 pm - 04.30 pm	EVENING TEA

• CONFERENCE PARTNERS •

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