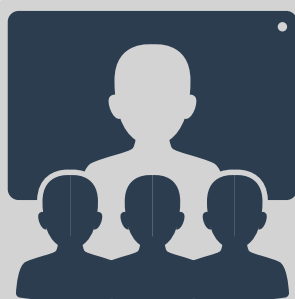


# Programme Schedule



**DISSO INDIA - 2020**  
**INTERNATIONAL**  
**ONLINE SYMPOSIUM**

**Date : 13th - 16th May, 2020**



# Disso India 2020

## International Online Conference on Dissolution Science and Applications

Theme : Dissolution as a Quality Assurance Tool for Product Development, Approval and Batch Releases

### PROGRAM SCHEDULE

Title and Topics	Speaker	Time	Partner
<b>May 13, 2020</b>			
<b>Inaugural Session of Disso India 2020 - Online International Conference</b> Address by Chief Guest, Dr. V. G. Somani, DCGI, New Delhi & by the Guest of Honour, Dr. B. Suresh, Pro Chancellor - JSS Academy of Higher Education and Research & President - Pharmacy Council of India.		2.30 - 3.30 PM	
<b>Module 1- Recent Advances in Dissolution Science and Testing</b>			
1. Dissolution and Bioclassification System (BCS)	Dr Vinod P Shah (Ex-USFDA) Consultant, USA <a href="mailto:dr.vpshah@comcast.net">dr.vpshah@comcast.net</a>	3.30- 4.05 PM (15 min Q&A)	
2. Automation in Dissolution	Kempf Juergen, Sotax AG, Switzerland <a href="mailto:Juergen.kempf@sotax.com">Juergen.kempf@sotax.com</a>	4.30- 5.05 PM (15 min Q&A)	
3. Role of Dissolution in Complex Generic Drug Products	Dr Vinod P Shah (Ex-USFDA) Consultant, USA <a href="mailto:dr.vpshah@comcast.net">dr.vpshah@comcast.net</a>	5.30-6.05 PM (15 min Q&A)	
<b>May 14, 2020</b>			
<b>Module 2 – Regulatory Aspects of Dissolution Testing and Method Development Strategy</b>			
1. Regulatory Compliance Challenges & Issues Related to Dissolution Testing & Studies	Vijay Kshirsagar, TRAC Consulting, India <a href="mailto:yukshirsagar@gmail.com">yukshirsagar@gmail.com</a>	2.30-3.05 PM (15 min Q&A)	
2. Dissolution Testing post COVID-19	Dr. Leong Chuei Wuei, CEXA Consultancy, Malaysia <a href="mailto:chueiwuei.leong@dreamcatcher.asia">chueiwuei.leong@dreamcatcher.asia</a>	3.30 - 4.05 PM (15 min Q&A)	
3. The Future of Clinically Relevant Dissolution Testing and Physiologically Based Biopharmaceutics Modeling (PBBM/PBPK) in Drug Product Development, Manufacturing Changes and Controls	Dr. Sandra Suarez-Sharp, Vice President - Regulatory Affairs, Simulations Plus, Inc., USA <a href="mailto:sandra.suarez@simulations-plus.com">sandra.suarez@simulations-plus.com</a>	4.30 – 5.05 PM (15 min Q&A)	
4. Harmonization of CMC area of dissolution, EU/US perspectives, including new FDA guidances & USP Chapters & Monographs	Vivian A. Gray V. A. Gray Consulting, Inc., USA. <a href="mailto:vagray@rcn.com">vagray@rcn.com</a>	5.30 – 6.05 PM (15 mts Q & A)	

• CONFERENCE PARTNERS •




# Disso India 2020

## International Online Conference on Dissolution Science and Applications

Theme : Dissolution as a Quality Assurance Tool for Product Development, Approval and Batch Releases

### PROGRAM SCHEDULE

Title and Topics		Speaker	Time	Partner
<b>May 15, 2020</b>				
<b>Module 3 – Dissolution for Novel Drug Delivery Systems</b>				
1.	Dissolution testing Methodologies for Nanomedicines	Prof. Padma Devarajan, ICT, Mumbai <a href="mailto:pv.devarajan@ictmumbai.edu.in">pv.devarajan@ictmumbai.edu.in</a>	2.30 - 3.05 PM (15 min Q&A)	
2.	In vitro release testing of conventional dosage forms	Samir Haddouchi, SPS Pharma Services, France <a href="mailto:samir.haddouchi@sps-pharma.com">samir.haddouchi@sps-pharma.com</a>	3.30 - 4.05 PM (15 min Q&A)	
3.	IVIVC for specific products Example of injectable SR formulations, Importance of setting the right dissolution method	Prof. Jean Michel Cardot, France <a href="mailto:j-michel.cardot@uca.fr">j-michel.cardot@uca.fr</a>	4.30 - 5.05 PM (15 min Q&A)	
4.	Development of IVIVCs for Complex parenteral products such as microspheres	Prof Diane Burgess, USA <a href="mailto:djbuconn@gmail.com">djbuconn@gmail.com</a>	5.30-6.05 PM (15 min Q&A)	
<b>May 16, 2020</b>				
<b>Module 4 – Material properties and formulation development</b>				
1.	Ternary Amorphous Solid Dispersions for Solubility Enhancement	Dr. Arvind K. Bansal, NIPER India <a href="mailto:akbansal@niper.ac.in">akbansal@niper.ac.in</a>	2.30-3.05 PM 15 min Q&A	<b>ACG</b>
2.	Influence of Particle Characteristics on Dissolution Testing	Sandeep Kulkarni . Director, Image provision Technology PVT Ltd, India <a href="mailto:sandeep@imageprovision.com">sandeep@imageprovision.com</a>	3.30 - 3.55 PM (10 min) Q&A	
3.	IDAS – In Vitro Dissolution and Absorption Systems to Evaluate Impact of Dissolution on Permeation/Absorption	Dr. Vatsala Nageshwaran, Absorption Systems, USA <a href="mailto:vnaageshwaran@absorption.com">vnaageshwaran@absorption.com</a>	4.15- 4.50 PM (15 min Q&A)	
4.	Dissolution Testing: Enhanced Learnings from DISSO2020 !!	Dr. Umesh Banakar, Prof.& President, Banakar Consulting, USA <a href="mailto:ubanakar@gmail.com">ubanakar@gmail.com</a>	5.10- 5.45 PM (15 min Q&A)	
5.	Conclude and Vote of Thanks	Dr. L. Ramaswamy, General Secretary, SPDS	6.00 - 6.10 pm	

• CONFERENCE PARTNERS •



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



SWISS  
QUALITY  
ISO 9001

### 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 • IVVC 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

#### SOTAX India Pvt. Ltd.

601, ECO House  
Vishweshwar Nagar  
Goregaon East  
Mumbai 400063

phone: +91-22-42950191/92  
fax: +91-22-40141332  
sotaxindia@sotax.com  
www.sotax.com