



SYMPOSIUM ON DISSOLUTION SCIENCE & DRUG DEVELOPMENT



Venue :

National Institute of Pharmaceutical Education and Research (NIPER), Auditorium Hall, Sector 67, Near PCA Stadium, S.A.S. Nagar 160 062, Punjab

-: Date :-
4th March 2025

-: Time :-
9.30 am to 5.30 pm

Industry Partners :-





Prof. Saranjit Singh
Ex. Professor & Head,
Pharmaceutical Analysis,
NIPER, SAS Nagar



Suhas Yewale
Associate Director,
Techno Commercial,
SOTAX India Pvt Ltd, Mumbai



Dr Ashutosh Sharma
AVP, Analytical Development
Sun Pharmaceuticals
India Ltd



Sudeep Ojha
General Manager,
Analytical Development,
Zydus Lifesciences Ltd



Dr Hargovind Seth
Head - QC,
Windlas Biotech
Limited



Dr Sanyog Jain
Professor,
Pharmaceutics,
NIPER, SAS Nagar



Raveendranath Govindaraj
Founder & Managing Partner,
Lab Iconics Technologies LLP

Agenda

09.30 - 10.30	Inauguration	State Drug Commissioners (Chief Guest), NIPER Director, Dr A K Bansal, Dr Ramaswamy, Dr Rajesh Gupta
10.30 - 11.00	Dissolution Chapter & Diff Types of Dissolution Apparatus as per IP including Type IV, an emerging tool	Suhas Yewale
11.00 - 11:20	Tea Break/Stall visits/Networking	
11.20 - 11.40	Session I: Chair Dr Rajiv Desai Importance of dissolution in defining BCS classification	Dr Sanyog Jain
11.40 - 12.10	Difference between QC and discriminatory dissolution (for in vitro bioequivalence) testing	Dr Ashutosh Sharma
12.10 - 12.40	'Effect and Causes of Coning' during dissolution of tablets	Sudeep Ojha
12.40 - 13.00	Question & Answers related to above Presentation	Chair & all above speakers
13.00 - 14.00	Lunch/Stall Visits/Networking	
14.00 - 14.30	Session II: Chair Dr A K Bansal Investigation of OOS results obtained during dissolution testing	Dr Hargovind Seth
14.30 - 15.00	Problems during dissolution of capsule formulations – cross linking of gelatin / interaction of drug with gelatin / interference with analytical method	Prof. Saranjit Singh
15.00 - 15.30	How digitalization and lab automation help in pharmaceutical and biotechnology industries to improve efficiency, productivity, and compliance	Raveendranath Govindaraj,
15.30 - 15.50	Question & Answers related to above Presentations of Session II	Chair & all Speakers
15.50 - 16.30	Panel Discussion / Q&A What care needs to be taken during dissolution testing? Common Flaws?	Moderator: Dr Rajeev Desai Panellists : • Harish Jain • Dr Rajesh Gupta • Dr Deo Narayan Dikshit
16.30 - 17.00	USP4 Live Demo	
16.30 - 17.00	Tea Break	

Fee payment details

Go To : <https://www.onlinesbi.sbi/sbicollect/icollecthome.htm>

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Registration fees Rs.1000 + gst (Rs.1180)

Register here : <https://forms.gle/PKT8BGUDFTnjghef6>

The Delegate registration shall be approved only against the receipt of the payment of the Delegate fee

Registration support

Dr U. R Lal, Asst Professor, NIPER, SAS Nagar, Mob : 8219487781 • Email : urlal@niper.ac.in



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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 :

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SOTAX India Pvt. Ltd.

Office No. 2-4, 3rd Floor,
Anupam Annapolis, Aarey Road,
Goregaon (East), Mumbai - 400063

Phone : 022-26851903
sotaxindia@sotax.com
www.sotax.com

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