



SOTAX

One Day Seminar on CGLP for the Pharma Industry

“Enhancing Quality Control in the Pharmaceutical Industry: Best Practices for Success”

PRIME SPONSOR : SOTAX INDIA LTD



Mr. Praveen Naik
AGM, QC, Unichem
Laboratories



Mr. Veerdhaval Kamble
GM Quality Assurance,
Glenmark
Pharmaceuticals.



Mr. Vinay Manerikar
Dy. General Manager QC
Zydus lifescience



Mr. Sandip Mahale
Manager Quality Assurance
Lupin Limited



Dr. Umesh Banakar
Pharmaceutical
Consultant

On Sunday, 19th January 2025
At Z Square, Cine Samrat, Panaji - Goa
Timing: 9.30 A.M. to 5 P.M.

About the Seminar

Quality Control (QC) is an essential part of the pharmaceutical industry, ensuring that products meet required quality standards. This seminar aims to provide QC personnel with the knowledge, skills, and best practices needed to maintain and improve quality control processes, ensuring the highest standards in pharmaceutical manufacturing.

This seminar is specially designed to provide key insights to QC professionals, ensuring that their activities meet the standards required by Indian regulatory authorities and align with global best practices. Whether you are a regulatory expert, quality control or quality assurance professional, this seminar is an invaluable opportunity to enhance your understanding and stay up-to-date with the latest requirements.

Join us for an in-depth seminar exploring the several updates and practical implications of current Good Laboratory Practices.

Why attend? - Key Highlights

Gain in-depth knowledge of QC best practices in the pharmaceutical sector

- Learn how to maintain compliance with global pharmaceutical regulations
- Understand new techniques and technologies transforming QC processes
- Network with industry professionals and regulatory experts
- Enhance your organisation's quality standards and minimise compliance risks

Who Should Attend?

- Quality Control Personnel
- Regulatory Affairs Professionals
- Pharmaceutical Quality Assurance Teams
- Compliance Managers, Production Managers, and Supervisors
- Anyone interested in understanding the key concepts of cGMP in the pharma industry.

Registration Details:

Category		Fees
1	IPA Member & Teaching faculty from pharmacy colleges	Rs. 1,000
2	Non-IPA Member	Rs. 1,500
3	M.Pharm Students from Goa College of Pharmacy & PES' Rajaram & Tarabai College of Pharmacy [limited seats available]	Rs. 700

Last date for Registration: 5th January 2025, by 5 p.m.

Only 100 seats on a first-come, first-serve basis.

Spot Registrations will not be entertained.

Registration process:

Please fill up the registration form using the link below and submit it.

<https://docs.google.com/forms/d/e/1FAIpQLSdhDBr-nnUA6-bQ7cMhSngAy6K3-43RDDvJlcbCRyECWyUNMw/viewform?usp=sharing>

1. You will receive a confirmation of your seat within 48 hours of submission.
2. Only after confirmation of your seat can you pay and send the proof of payment to ipagoa@gmail.com. Payment to be online on:

"The Indian Pharmaceutical Association",

SB Account No. 11031491092

State Bank of India, Panaji Branch

IFSC: SBIN0000509.

Sessions and Expert Speakers:

	Topic	Name of Speaker
1.	Lecture 1: Good QC Laboratory Practices	Mr. Praveen Naik
2.	Lecture 2: How to Reduce OOS / Incidents in Laboratory	Mr. Veerdhaval Kamble
3.	Lecture 3: Lean management and human errors in QC laboratory	Mr. Vinay Manerikar
4.	Lecture 4: Nitrosamine Impurities : Introduction, Generation & Control (Prevention).	Mr. Sandip Mahale
5.	Lecture 5: The In Vitro Dissolution Test: Overstretching the limits !!	Dr. Umesh Banakar
5.	Panel Discussion and Q & A	All the above speakers

Don't Miss Out on This Opportunity!

Stay ahead of the curve, ensure your facilities meet the highest standards, and build a network of peers equally dedicated to improving industry practices. Join Us and Take the First Step Toward Achieving Uncompromising Quality and Compliance in Pharmaceutical Manufacturing!

For any further details, please Email us at: ipagoa@gmail.com

Organised by:

The Indian Pharmaceutical Association, Goa State Branch

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