

REGISTRATION FORM

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REGISTRATION FEES

IPA Members : 2000/-

Academic / Research Student : 1000/-

Spot Registration / Non IPA Member : 2500/-

Note : No Cancellation. Please send your cheque / DD in favour of IPA – RAD payable at Mumbai to the following address

The Indian Pharmaceutical Association

Kalina, Santacruz (East), Mumbai – 400098.

Phone: +91-22-26671072 / 26670744

E-mail: ipacentre@ipapharma.org

Website : www.ipapharma.org



National Seminar on “Validation Requirements for Regulatory Compliance”

Venue

Sci Tech Centre
7, Prabhat Nagar, Jogeshwari (W), Mumbai, India

On 8th October 2011

Organised by

Indian Pharmaceutical Association

Regulatory Affairs Division

&

**Industrial Pharmacy
Division**

(The Regulatory Affairs Division and the Industrial Pharmacy Division of the Indian Pharmaceutical Association have been in the forefront conducting national & international Seminars, Workshops, Symposia for the professionals working in the pharma industry as part of continuing pharmacy educational program).

Co-ordinators

SP Manek

J Jayaseelan

K G Gadewar

Secretary IPA-RAD

Secretary IPA-IPD

Member IPA-RAD

PROGRAM:	Seminar Objectives
9.00 - 9.30 am : Registration	
9.30 - 9.45 am : Welcome address – Ram Banarase , Chairman, IPA-RAD	Validation is a regulatory requirement in Pharmaceutical industry for Processes, Analytical Methods, Cleaning Procedures and Computer systems to ensure consistent Quality, Safety and Efficacy for the drug products.
9.45 - 10.15 am : Overview of Validation requirements in Pharmaceutical Industry - Kaushik Desai , Chairman, IPA-IPD	The ONE DAY SEMINAR is designed to provide an unique platform to Pharmaceutical industry to review:
10.15 - 10.30am : Tea / Coffee Break	
10.30 - 11.15 am: Challenges of Validation of Stability Indicating Analytical Methods – R. Girijan , Toshvin	a) Importance and the requirements of the validation of critical manufacturing and the packaging processes to ensure consistent quality of the drug products.
11.15-12.00 noon : Validation of Pharmaceutical Packaging, Dr. C S Purushottam , Director, SIES School of Packaging	b) To explore and evaluate the challenges and various issues related to validation of analytical methods in general and stability indicating methods in particular, with focus on regulatory requirements.
12.00 - 12.45 pm: Validation of Computer Systems – Mr. Vasistha Mehta , Epitome Technologies, Gujarat	c) To understand needs of and how to validate computer systems increasingly being deployed in manufacturing, packaging and analytical areas.
12.45 - 1.45 pm : Lunch Break	
1.45 - 2.30 pm : Key Requirements for Qualification and on going Validation of Stability Chambers - Dr. Pramit Kumar , Technical Director, Thermolab Scientific Equipment Pvt. Ltd.	d) To learn frequently encountered observations related to validations during Regulatory audits of the facilities.
2.30 - 3.15 pm : Validation-common Audit observations - Dr. V. B. Malkar , Head Quality Management, Reliance Life Sciences.	e) The learned Speakers from the pharmaceutical industry will share their expertise and provide practical methodologies and options to meet the critical challenges of validation in the pharma industry.
3.15 - 3.30 pm : Tea / Coffee Break	
3.30- 4.30 pm : Panel discussion-Moderator: Dr. S.P. Manek & Panel of experts	<p>WHO SHALL ATTEND?</p> <p>Seminar will be very useful for senior and middle level executives working in Product Development, Analytical methods development, Quality Assurance, Regulatory affairs and those engaged in manufacture of various dosage forms as it will provide them with insights into practical methodologies and issues faced during execution of validation program.</p>