

Conference on		
Quality Management in Bulk Drug and Formulation Manufacturing		
Day 1 : 12th August, 2016		
10.00 A.M		<i>Welcoming the Guests, Invocation & Lighting of the Lamp</i>
10.05 A.M.		Welcome address & about conference Dr U.S.N.Murty , Project Director, NIPER Hyderabad
10.15 A.M		Address by Guest of Honour : Dr A.K.S.Bhujanga Rao , President, Natco Research Centre
10.25 A.M		Address by Guest of Honour : Dr P.G.Rao , Honorary Vice Chancellor Univ. of Science & Technology, Meghalaya
10.35 A.M		Address by Chief Guest : Dr K.V.Raghavan , INAE Distinguished Professor & Former Director, IICT
10.55 A.M.		Vote of Thanks
11.00 A.M - 11.45 A.M		Key Note Address : Dr K.V. Surendranath , United States Pharmacopeia, Hyderabad
		Regulatory Expectations on Pharmaceutical Quality System
Session I		
12.00 P.M - 12.45 P.M	IL1	Dr Premnath Shenoy , Former Director Regulatory Affairs, AstraZeneca
		QMS and draft revised Schedule M
12.45 P.M - 1.30 P.M	IL2	Sunil Singhai , President, Adept Pharma Consultants, Hyderabad
		Pharmaceutical Products - cGMP and Environment
Session II		
2.30 P.M - 3.15 P.M	IL3	Dr Vikaslata Jain , Head ESO India, Quality Assurance, Novartis
		Deviation Management and Root cause analysis.
3.15 P.M - 4.00 P.M	IL4	Madan Mohan Reddy , Director QA & regulatory Eisai, Visakha Patnam
		Key Quality and Regulatory considerations for supply of Pharmaceutical products to Japan
Session III		
4.30 P.M - 5.15 P.M	IL5	T. Lakshmana Murthy , Director, Quality Assurance United States Pharmacopeia (I) Pvt. Ltd, Hyderabad
		Documentation and Data Management

Day 2 : 13th August, 2016

Session IV		
9. 30 A.M - 10. 15 A.M	IL6	G. Sangeetha , General Manager Regulatory Affairs, Hetero Drugs (Pvt) Ltd., Hyderabad
		Update on today's regulatory environment - US Market
10.15 A.M - 11.00 A.M	IL7	Dr Vilas Dahunkar , Chief Scientist- Process R&D Dr. Reddy's Laboratories Ltd, Hyderabad
		QbD application in generic drug development
Session V		
11.30 A.M - 12. 15 P.M	IL8	Dr R. Srinivas , Dean, NIPER Hyderabad
		Characterization of Drug metabolites and degradation products by Liquid chromatography-electrospray ionization tandem mass spectrometry
12.15 P.M - 1.00 P.M	IL9	Dr B. Mahesh Kumar , Associate Director, Nektar Therapeutics (India) Private Limited, Hyderabad
		Associate Director- Quality Assurance Nektar Therapeutics (India) Private Limited Hyderabad
		Quality Metrics to monitor QMS
2.00 P.M - 2.30 P.M	IL10	Dr Nalini Shastri , NIPER Hyderabad
		Six Sigma for Pharmaceutical Industry – Importance and Implementation
2.30 P.M - 3.00 P.M	IL11	T.Lakshmana Murty , Director, Quality Assurance, USP India, Hyderabad
		Change Control Management