

ADVANCED GMP WORKSHOPS 2018

FOR MANUFACTURING, QUALITY & REGULATORY PERSONNEL

12-13 November: Goa

15-16 November: Hyderabad

19-20 November: Ahmedabad

HEAR FROM USFDA, MHRA, WHO, INDUSTRY AND OTHER EXPERTS ON

- ☐ Organisation Culture Expectations of a Regulator
- ☐ Good Data and Record Management Practices and Sharing of Experiences & Issues
- ☐ Key Trends in Quality and GMP Inspections
- ☐ FDA Inspection Process- What You Should Anticipate
- ☐ Next Gen Quality Impact of Digital & Advanced Analytics on Quality
- ☐ Quality Metrics
- Building Quality Culture & Capabilities



Dr S Eswara Reddy Drugs Controller General (India), CDSCO



Dr Letitia Robinson Director, USFDA India Office



Mr Ewan Norton Senior GMDP Inspector, MHRA



Mr Vimal Sachdeva Technical Officer (Senior Inspector), WHO



Mr Thomas Arista Deputy Director, USFDA India Office



Mr Robert Iser
Former Director, Office of Process
and Facilities CDER, USFDA
& Vice President, PAREXEL Consulting



Mr Daniele Iacovelli Expert Associate Partner, McKinsey & Company, Germany



Mrs Julie Rose Expert Associate Partner, McKinsey & Company, France



Dr Jaidev Rajpal Partner, McKinsey & Company, India



Mr Vivek Arora Associate Partner, McKinsey & Company, India

Industry Speakers

Goa Mr Indrajit Bose (Lupin) Dr Ranjana Pathak (Cipla)* Hyderabad Mr Sairam Philkana (Dr Reddy's Lab) Mr Ganadhish Kamat (Dr Reddy's Lab) Ahmedabad Mr Rakesh Sheth (Torrent)* Mr Vipul Doshi (Cadila Healthcare)

*Confirmation Awaited

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Registration Closes on 31 October 2018

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