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FOR MANUFACTURING, QUALITY & REGULATORY PERSONNEL

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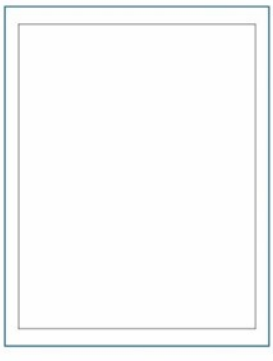
- ❑ *Key Trends in Quality and GMP Inspections*
- ❑ *FDA Inspection Process- What You Should Anticipate*



Mr Thomas Arista
Deputy Director, USFDA India Office

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

And Subject Matter Experts from Other Regulatory Agencies, Industry and Consultants on Topics of Relevance for Indian Pharmaceutical Industry

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Registration Closes on **12 October 2018**

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For registration write to archana.jatkar@ipa-india.org