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DAY 1 – 03.10.2016	DAY 2 – 04.10.2016
08.30 –09.00 Registration : Morning Coffee and Networking	***08.30 –09.00*** Registration : Morning Coffee and Networking
09.00 – 09.20 Welcome, Intro to Speakers	
09.20 – 10.30 Commissioning & Qualification of a pharmaceutical facility: A new approach as per latest EU GMP Annex 15-Part 1 By Mr. Paolo Curto' , DOC Managing Director	***09.00 – 10.30*** NEW EP Monograph for WFI Production : a New Challenge for Pharmaceutical Industry. By Mr. Paolo Curto' , DOC Managing Director
10.30 –10.45 Morning Break	***10.30 –10.45*** Morning Break
10.45 – 12.45 Commissioning & Qualification of a pharmaceutical facility: A new approach as per latest EU GMP Annex 15-Part 2 By Mr. Paolo Curto' , DOC Managing Director	***10.45 – 12.45*** : Only MASCO can: Turn-key approach in a pharmaceutical facility,
12.45 –14.00 Lunch Break	***12.45 –14.00*** Lunch Break
14.00 –15.30 Cleaning Validation of GMP Systems: A new approach as per latest EU GMP Annex 15 by Mr. Antonio Legnani-DOC Process Validation Director	***14.00 –16.30***: **Polysan**: VISIT OF THE FACTORY Case study in real time
15.30 -15.45 Afternoon Break and Networking	
15.45 –17.00 Process Validation and Material Qualification in light of New EU GMP Annex 15 and USP Chapters by Mr. Antonio Legnani-DOC Process Validation Director	***16.30 – 17.30*** Q & A and Networking
17.00 -17.30 Q & A, End of Session	END OF THE TRAINING PROGRAM