Quality Control for Pharmaceutical Manufacturers

⇒ Understand the essentials of Quality Control, Analytical Method, Method Validation, Verification

⇒ Get informed about Quality Control of Medicines in the life cycle of Medicines and the Supply Chain

⇒ Gain fundamentals on Analytical Validation and Method Verification

⇒ Learn how to compile QC data for regulatory Purposes (for marketing authorisation dossiers e.g. CTD)

⇒ Understand the Concept of Data Integrity

⇒Learn what international assessors want to know when doing audits in your premises

⇒ Understand the criticality of information given to the national regulatory authority

Join us for practical sessions, theoretical background and professional exchange!

Venue: Conference hall + on site visit in Pharmaceutical Quality Control Lab

Trainers: experienced trainers from international pharma industry

Duration: 2 days

Target group: professionals from pharmaceutical manufacturers (regulatory affairs managers, heads of Quality Assurance or heads of production), NMRA assessors

Can be combined with our training on CTD

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