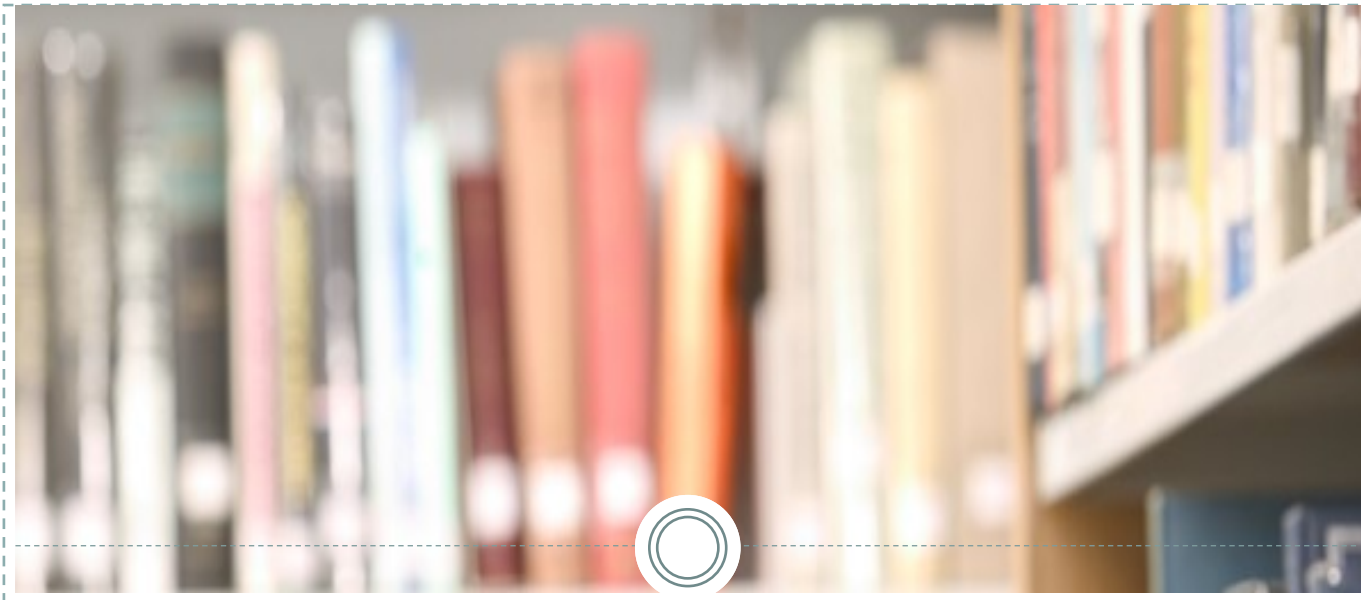

# Good Documentation Practices & Data integrity

⇒ Understand why good documentation is essential

⇒ Gain fundamentals on regulatory requirements

⇒ Learn how to prepare a good batch documentation and SOP

⇒ Get informed about Document Management Systems

⇒ Get acquainted with Data Integrity Standards

⇒ Learn how to design or improve data integrity process

⇒ Understand the criticality of data integrity from a regulatory point of view

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

| | |
|---|---|
| Venue: | Conference hall |
| Trainers: | experienced trainers from international pharma industry |
| Duration: | 3 days |
| Target group: | professionals from pharmaceutical manufacturers (Heads of Quality Control, Heads of production, Regulatory Affairs), NMRA inspectors |

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