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Webinar on

Proper Documentation and SOPs to Ensure Laboratory Compliance

Areas Covered

- *Laws, Regulations and guidance applicable to laboratory operations*
- *It's not just about meet the regulations, it's getting the processes to achieve business needs*
- *Process mapping approach to designing a sample executable laboratory operations SOP*
- *Steps necessary to ensure existing laboratory and new laboratories have processes implemented that ensure compliance*

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This presentation will review the regulations, citations, and typical procedures found in laboratory operations and then propose a way to build better documentation.

PRESENTED BY:

Over 30 years of diverse international industry experience in Quality Assurance, Quality Control, and Regulatory Affairs. Last 18 years as a consultant to biopharma, pharma and device industries.

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On-Demand Webinar

Duration : 60 Minutes

Price: $200

Webinar Description

This presentation will review the regulations, citations, and typical procedures found in laboratory operations and then propose a way to build better documentation.

- What laboratory documentation is required for compliance
- What are typical citations as it relates to laboratory documentation
- With forethought and design, good procedures are possible
- Good documentation and record integrity ensure good compliance

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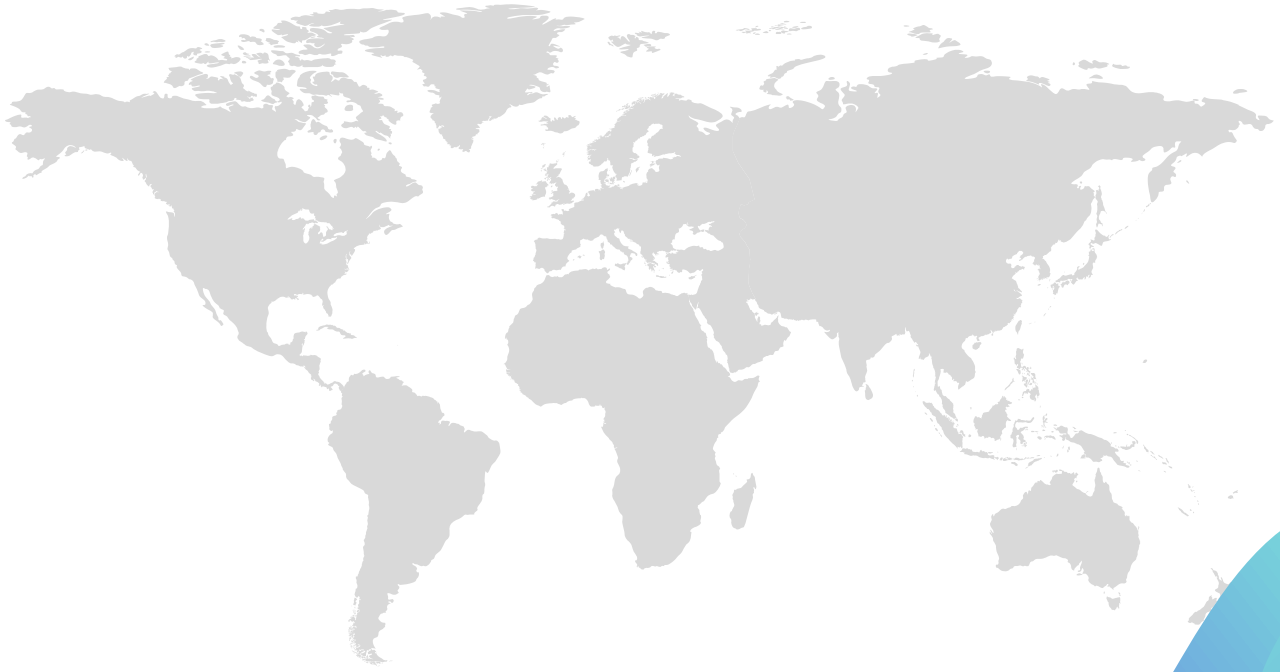
Who Should Attend ?

Laboratory operations management.
Quality Assurance Management.
Designers of laboratories



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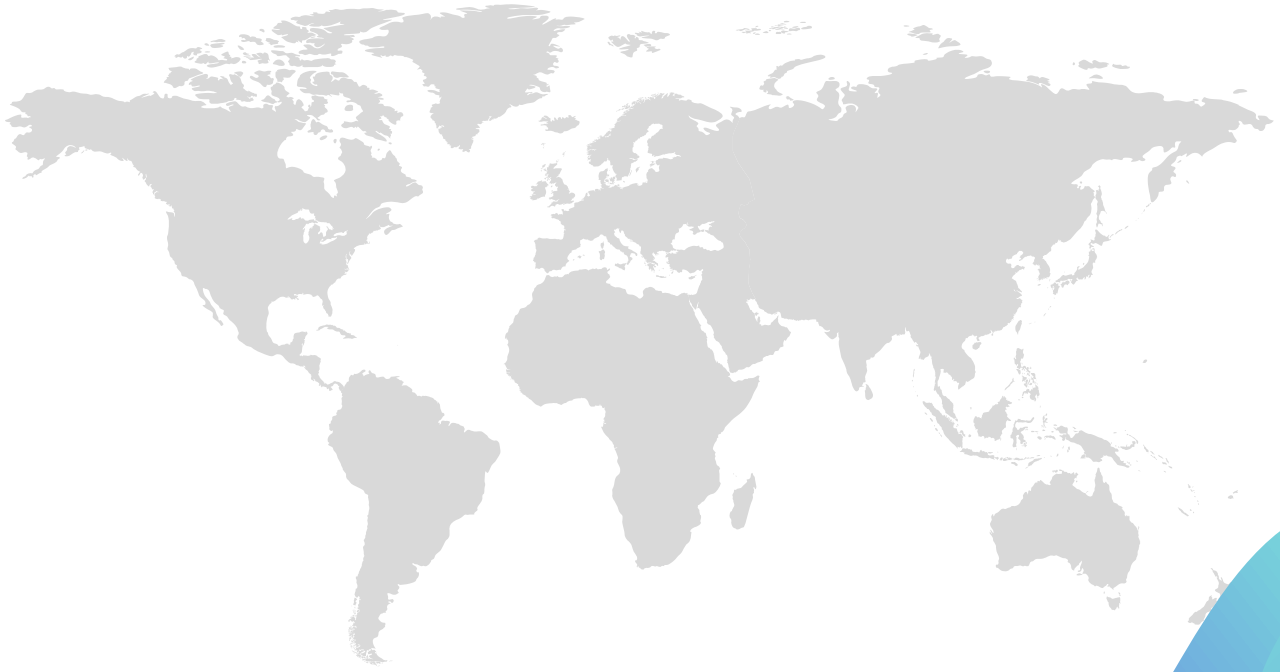
Why Should You Attend ?

The goal of laboratory documentation should be to ensure compliance. However, failures in documentation and procedures represent more than half of the top citations by FDA. FDA is telling us we just don't get it. As such we must look at what does proper documentation look like.



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