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Webinar on

Fundamentals of Good Manufacturing Practices (GMP) - Developing a Compliance Mindset

Areas Covered

- *Good Manufacturing Practice fundamentals*
- *GMP principals*
- *Data Integrity Principals found in GMPs*
- *Summary of specific areas covered throughout GMPs*
- *Case Study example scenarios with a discussion of best answers*

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Learning Objectives

- *Fundamental requirements of Good Manufacturing Practices (FDA based)*
- *Basic Principles the GMPs require*
- *Linkage to Data Integrity*
- *Learning how to approach operations from a compliance mindset to ensure success in any situation*

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Persons new to the industry or planning to enter the regulated pharmaceutical/biopharmaceutical industry.

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

A discussion of the basic principles and requirements for Good Manufacturing Practices. The presentation provides a brief history and basis for the existence of GMPs, a discussion of what they cover, and examples of real-life situations along with a discussion of possible responses and the reasons one or more responses may be appropriate or not.

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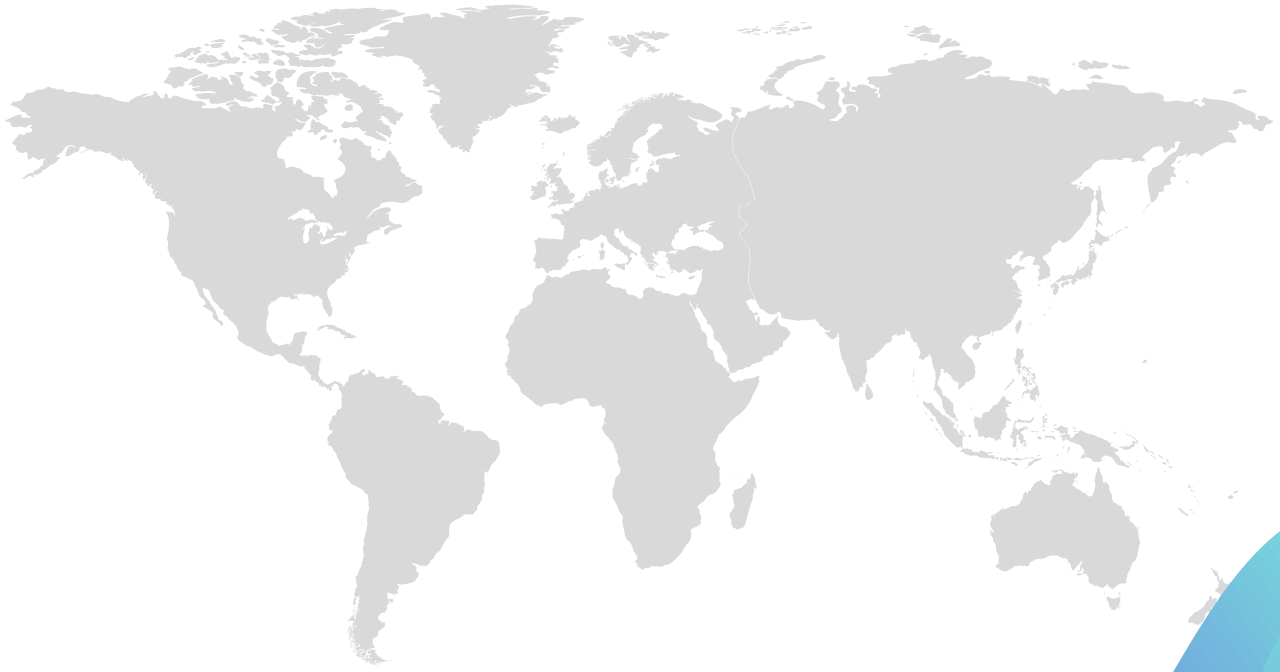
Who Should Attend ?

Students preparing to enter into regulated environments. New employees. Refresher for all levels.



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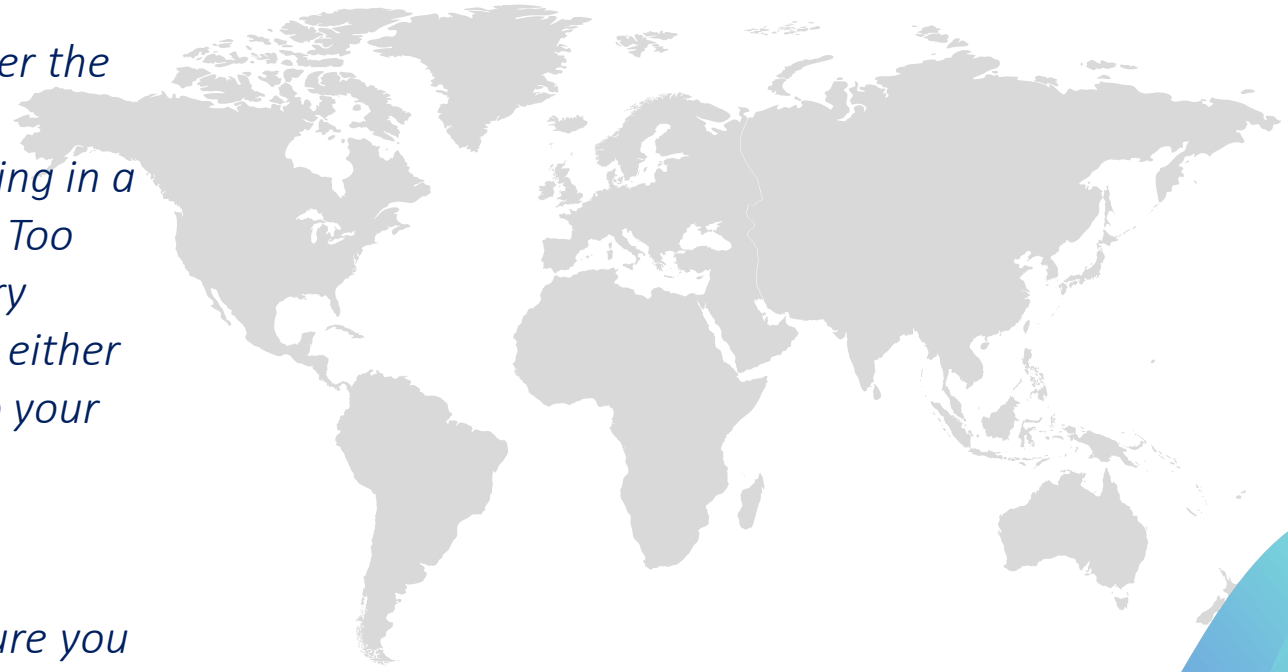
Why Should You Attend ?

Persons new to the industry or planning to enter the regulated pharmaceutical/biopharmaceutical industry. Equally valuable as a refresher. Working in a regulated industry requires a specific mindset. Too often personnel don't understand the necessary mindset or don't follow the necessary mindset either out of ignorance or other factors. Add value to your resume, ensure you have the right mindset.

In this webinar you will learn not just what is required, but how to think and operate to ensure you meet the intent of regulations applicable to the pharma/biopharma industry. We will translate the regulations into practical English – what do they mean? We will also explore different scenarios with possible actions and discuss the appropriateness or lack thereof for each action.



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