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Webinar on

Performing Effective and Compliant Sterility Failure Investigations

Date : April 02,2020

Learning Objectives

- *Regulations guiding manufactured product sterility testing and how to ensure adherence to program testing requirements*
- *Conducting sterility test failure investigations and what to look for during the investigation process*
- *Role of contamination such as microbial identification of contaminants and its importance*
- *Retest during a sterility failure investigation and how to apply corrective and preventative actions based on a sterility failure investigation*
- *Dispositioning impacted products based on investigational findings while avoiding common mistakes during product disposition*
- *Impact of sterility failure investigation for commercially distributed products under a stability testing program*

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This 90-minute webinar program will disclose how to prevent mistakes by implementing a robust CAPA program.

PRESENTED BY:

Upon earning a degree in Zoology at North Carolina State University, Joy began working in the pharmaceutical and biotech industries in 1992 at Pharmacia & UpJohn performing Environmental Monitoring and Sterility Testing.

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Date : April 02, 2020

Time : 01 : 00 PM EST

Duration : 90 Minutes

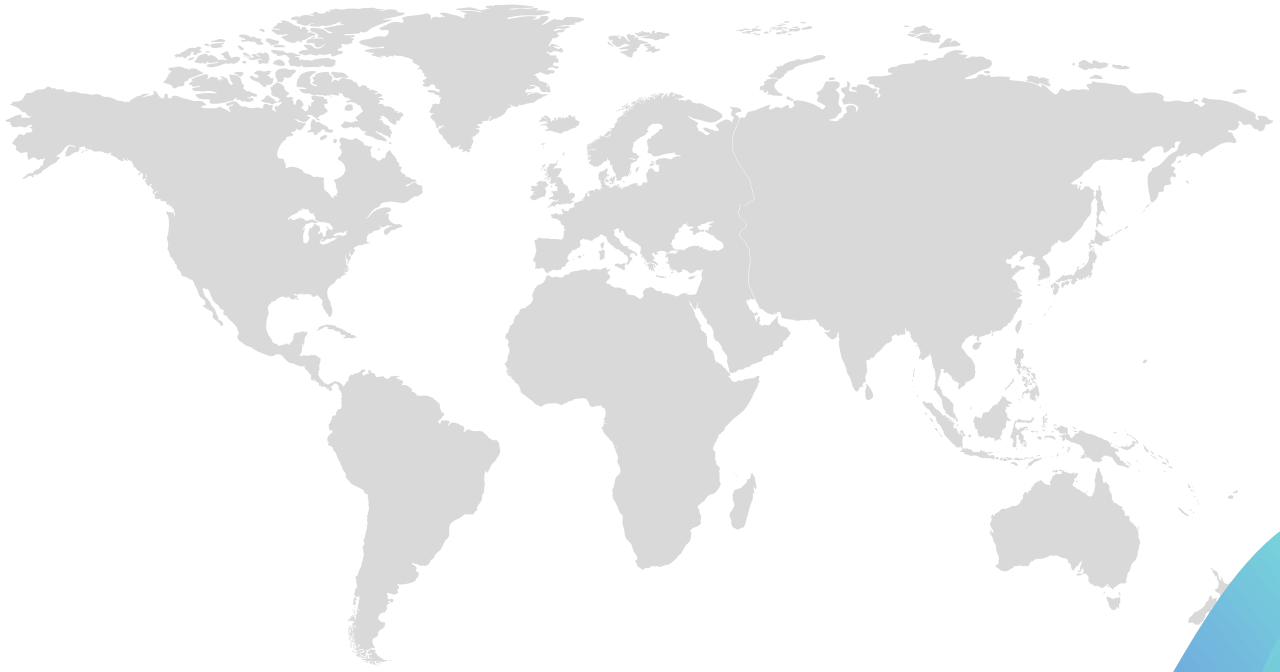
Price: $179

Webinar Description

This 90-minute webinar program will disclose how to prevent mistakes by implementing a robust CAPA program. Errors and inaccuracies made when corrective and preventative actions are not clearly recognized and applied during sterility test failure investigation will be discussed. Many times ineffective investigational procedures and tools are used to conduct a sterility test failure investigations. The webinar will also explain how avoiding common mistakes will ensure that manufacturers meet the sterility requirements USP <71> and other regulatory guidelines applicable to their finished products, bulk drug substance, raw materials or excipients

Who Should Attend ?

Quality Control and Quality Assurance Professionals
Production Personnel
Manufacturing Supervisors
Mangers
Microbiology Analysts



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Why Should Attend ?

The objective of this webinar is to provide an understanding of the regulations for sterility testing programs and the process of conducting an effective, robust and compliant sterility test investigations for various types of products.

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