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Webinar on

Cleaning Validation

Learning Objectives

- *Cleaning Validation Guidance*
- *Establishing Health-Based Limits*
- *Quality Risk Management*
- *Lifecycle Approach*
- *Cleaning Verification*
- *Continued Validation Monitoring*
- *Cleaning Analytical and Microbial Methods*

This webinar discusses global Cleaning Validation as it is implemented in the pharmaceutical and biopharmaceutical industries.

PRESENTED BY:

Igor Gorsky has been a pharmaceutical industry professional for over 30 years. He held multiple positions with increasing responsibility at Alpharma, Wyeth, and the Shire.

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On-Demand Webinar

Duration : 90 Minutes

Price: $200

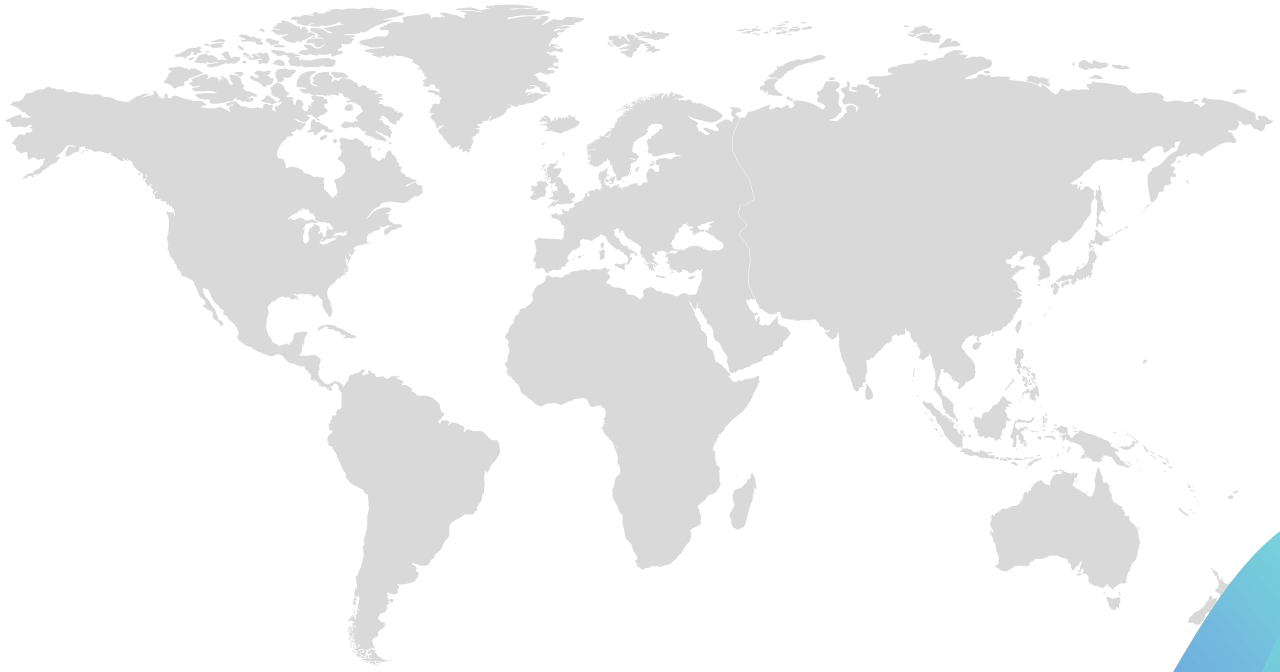
Webinar Description

This webinar discusses global Cleaning Validation as it is implemented in the pharmaceutical and biopharmaceutical industries. In addition, it describes the use of QRM (quality risk management) in Cleaning Validation and the growing trend of applying this approach to the lifecycle of the product manufacturing process. Establishing of health-based Cleaning Validation limits also discussed in this session. Additionally, recommendations on the compliant and sustainable programs will also be covered, as well as regulator's expectations for these programs.

Cleaning Validation is a regulatory requirement as well as expectations. In addition, a robust and compliant cleaning validation program makes perfect business sense for successful manufacturing facilities.

Who Should Attend ?

Pharmaceutical
Biopharmaceutical
Nutraceuticals
Medical Devices
APIC
Validation
Quality Assurance
R & D
Production
Quality Control
Microbiology
Engineering



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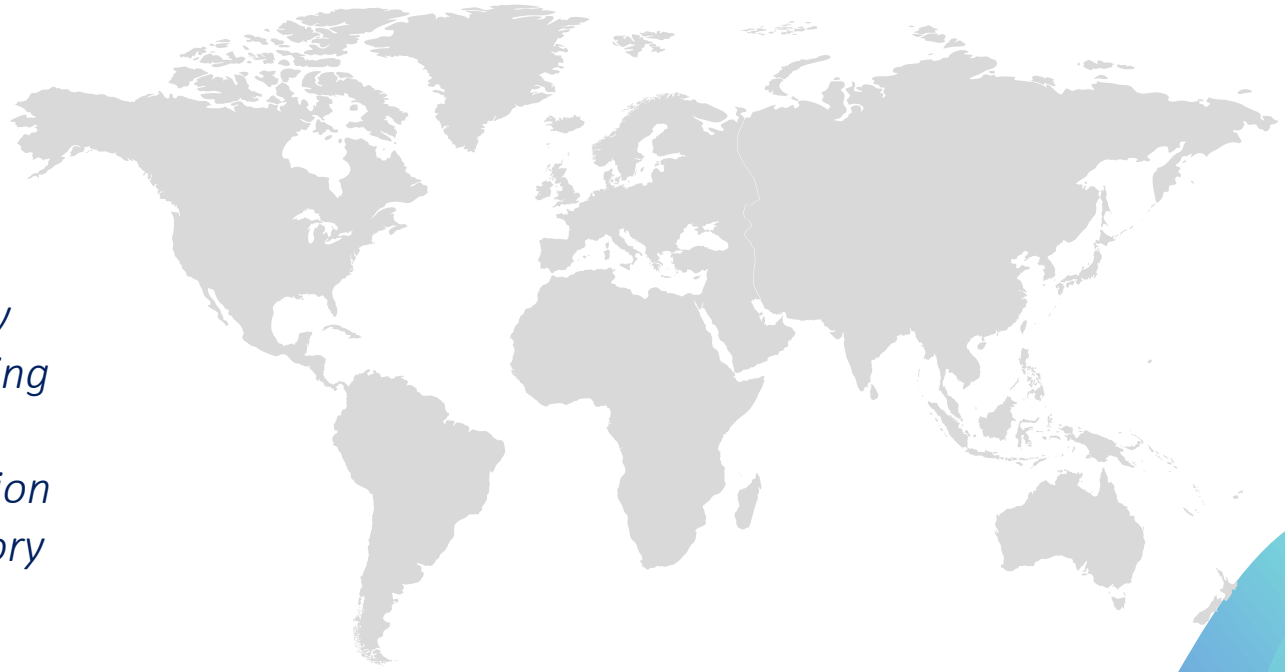
Why Should You Attend ?

If you are a professional who is involved in the development of pharmaceutical and biopharmaceutical products and processes' cleaning procedures throughout the continuous lifecycle of drug products this webinar is definitely for you. During this session, we will discuss Cleaning Validation that includes the development of cleaning process programs and cleaning verification and validation programs. We will discuss regulatory validation guidance and show how quality risk management should be utilized in the implementation of a risk-based lifecycle approach to cleaning validation. We will also touch upon the use of statistics in the cleaning validation process and talk about continued verification/monitoring. We will concentrate on a risk-based approach as it is a cornerstone of cleaning process validation principals and practice.



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