GRC Online Training Hub
EDUCATORS
Axona Technology and Solutions

Webinar on

Writing Validation Master Plans: Best Practices for Authoring a Compliant Document

Learning Objectives

- *What is a VMP and what is its intended use*
- *How is a VMP different that validation SOPs*
- *Components of a VMP*
- *Regulatory requirements for a VMP*
- *Team Writing a VMP*
- *Examples of VMPs*
- *Effective writing practices for an audit-ready document*

GRC Online Training Hub
EDUCATORS
Axons Technology and Solutions

Validation Master plans are written to assist an organization with validation strategies or to provide control over a specific process.

PRESENTED BY:

Upon earning a degree in Zoology at North Carolina State University, Joy made her debut in the pharmaceutical industry in 1992 at Pharmacia & Upjohn performing Environmental Monitoring and Sterility Testing. Her hard work allowed her to move into a supervisory role at Abbott Laboratories where she oversaw their Quality Control Lab. In 1998 Joy moved to Wyeth Lederle and worked in Quality Assurance, performing GMP Compliance audits, batch record reviews, and holding annual GMP training for new employees.

GRC Online Training Hub EDUCATORS Axens Technology and Solutions

On-Demand Webinar

Duration : 90 Minutes

Price: $200

Webinar Description

This webinar will discuss the major components of Validation Master Plans. It will discuss how the VMP is different from Validation SOPs. Various regulatory requirements for Validation Master Plans will be discussed as well as effective guidelines for authoring a VMP and a team writing approach to authoring a Validation Master Plans will be discussed. Various types and examples of VMPs will be given and discussed. Participants will learn how to effectively write a VMP most appropriate for their manufacturing organization.

Validation Master Plans discuss validation activities across an entire site or within an organization. The Validation Master Plan is a summary of validation strategies. The purpose of Validation Master Plans is to document the compliance requirements for the site and to ensure that sufficient resources are available for validation projects.

Validation Master Plans may be written to cover specific departmental validation activities or the validation process for a specific type of system. Validation Master plans are written to assist an organization with validation strategies or to provide control over a specific process.

Who Should Attend ?

Validation Engineers

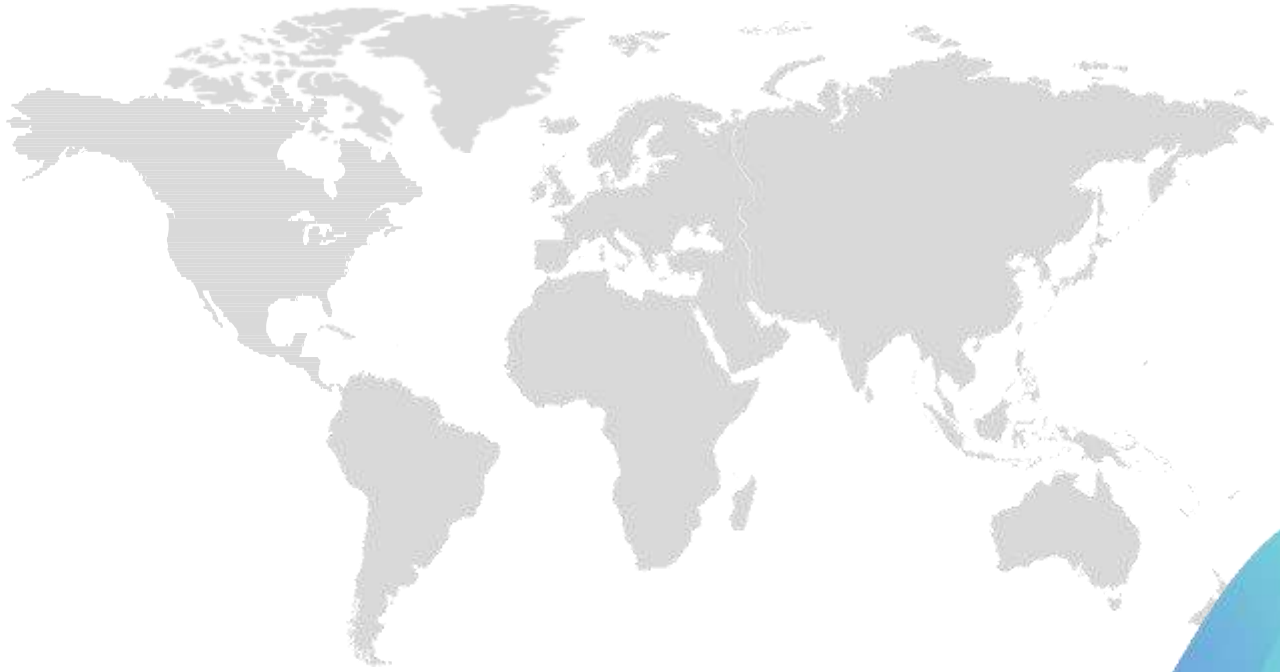
Supervisors and Managers

Quality Assurance personnel

Engineering Consultants

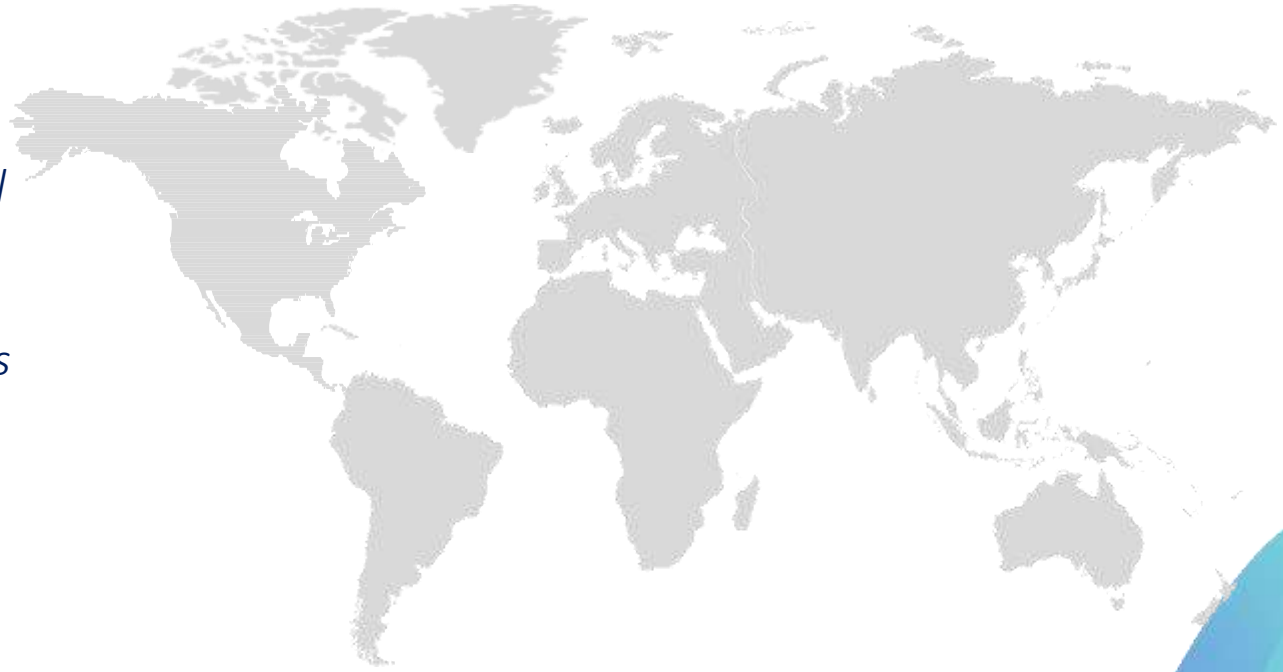
Regulatory Personnel

Auditors



Why Should Attend ?

Attendees should register to learn the major components of a Validation Master Plans. They will learn how the VMP is different from Validation SOPs. Various regulatory requirements for Validation Master Plans will be discussed as well as effective guidelines for authoring a VMP and a team writing approach to authoring a Validation Master Plans will be discussed.



GRC Online Training Hub
EDUCATORS
Access Technology and Solutions

To register please visit:

www.grceducators.com

support@grceducators.com

740 870 0321