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Webinar on

Regulatory Affairs Project Management

Learning Objectives

- *Setting tasks focused on project objectives, Re-evaluating resource needs and time Constraint*
- *Creating a project timeline and task tracker, Assigning managing resources*
- *Leading effective project team meetings , Tracking progress on tasks and overall project*
- *Using available tools to assist in project completion*
- *Ensuring effective team communication , Communicating progress to management*
- *Identifying and mitigating risks to project Completion,*

In this webinar will address approaches to regulatory affairs project management for clinical trial applications, marketing authorization applications, and ongoing management of regulatory obligations.

PRESENTED BY:

Peggy J. Berry, MBA, RAC, is the President & CEO at Synergy Consulting where she provides consulting services to companies in all aspects of drug development. She also provides group and one-on-one training in drug development, regulatory affairs, and project management topics. She is an editor-in-chief of Fundamentals of US Regulatory Affairs, 6th edition (RAPS, MD 2010).

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On-Demand Webinar

Duration : 90 Minutes

Price: $200

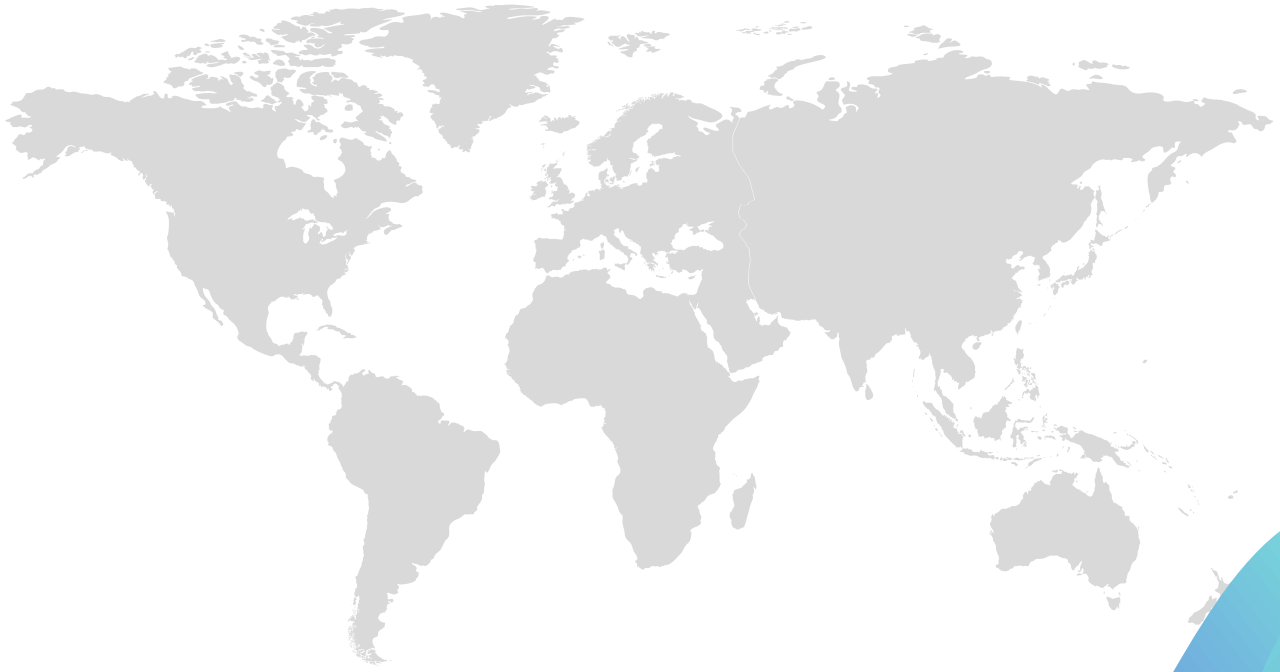
Webinar Description

This program will address approaches to regulatory affairs project management for clinical trial applications, marketing authorization applications, and ongoing management of regulatory obligations. The information obtained will enable effective management and tracking of time and resources to complete the project objectives and ensure regulatory compliance.

Who Should Attend ?

Regulatory Affairs, Regulatory Operations, Project Team Members across Disciplines.

This presentation is targeted toward the following organizational positions and discage understandable by all technically educated or trained individuals, regardless of specialty. This information would be useful at levels from senior management to operative and would be valuable for experience levels ranging from seasoned veterans to those newly assigned roles related to regulatory project management.



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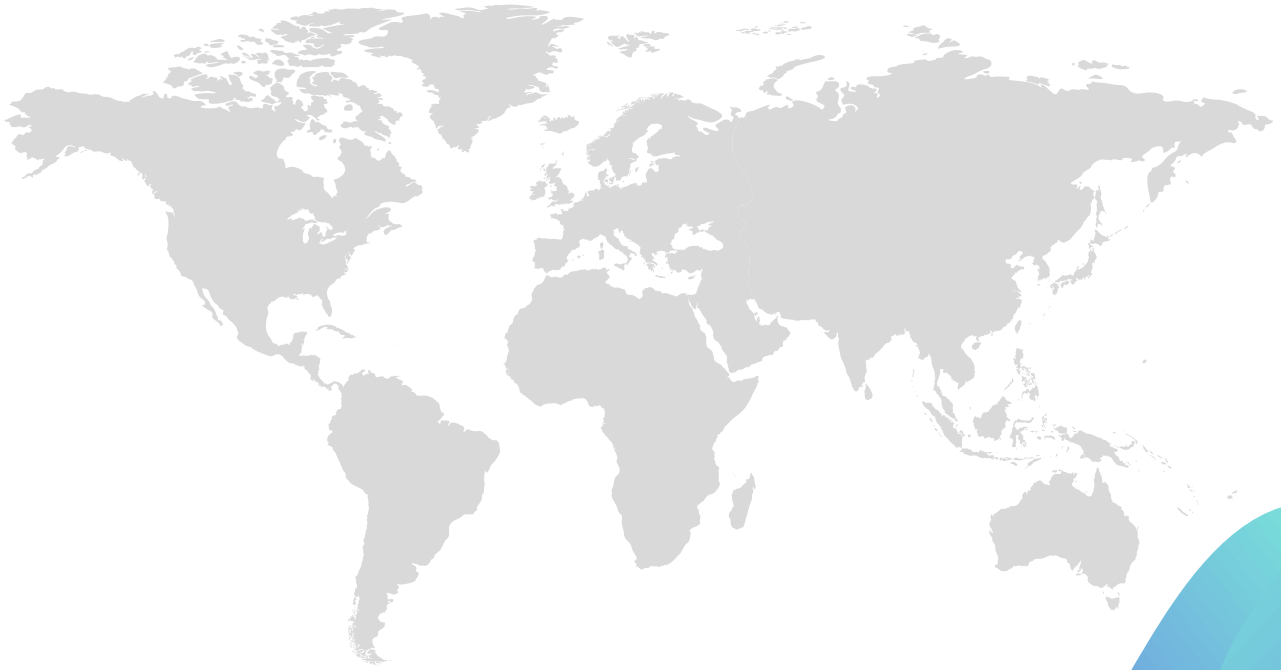
Why Should Attend ?

This program will address aspects of traditional project management with consideration of tailoring for regulatory affairs projects as well as tips for leading the team to completion and maintaining established timelines.



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