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Webinar on

Comparability Protocols For Approved Drugs

Learning Objectives

Attend this training to determine whether or not a comparability protocol will provide you with future advantage in product life-cycle management.

This webinar will address approaches to developing comparability protocols, including primary content considerations and timing of submission.

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 has also held Regulatory Affairs roles within two clinical contract research organizations and has worked in review divisions at the FDA.

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On-Demand Webinar

Duration : 90 Minutes

Price: $200

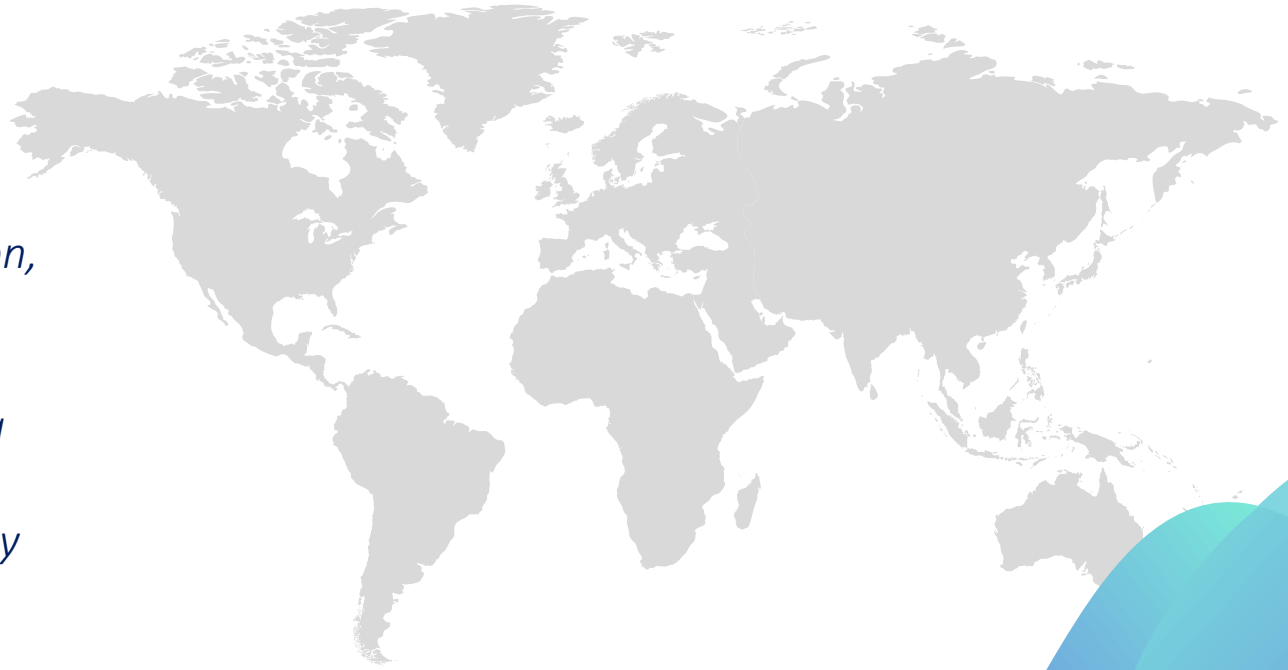
Webinar Description

This webinar will address approaches to developing comparability protocols, including primary content considerations and timing of submission. The information obtained will enable completion of appropriate testing to make changes to the drug substance or drug product formulation, manufacturing facility, and container closure which can be designed to save time and money in the future.

Who Should Attend ?

Quality Assurance, Quality Control (Chem and Micro), Process and Design Engineering, Process Automation, Manufacturing Operations, Validation, Regulatory Affairs

This presentation is targeted toward the following organizational positions and disciplines and is presented in practical language 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 to roles related to manufacturing.



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