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Webinar on

GMP's Applied to Medical Cannabis: All You Need to Know

Learning Objectives

- *Recent federal government position on ending Cannabis prohibition nationwide*
- *FDA position on CBD and other cannabis-derived products Background from a regulatory standpoint*
- *Approvals by states and what have we learned*
- *Challenges with the regulation of plant-based product*
- *Dosage challenges, HR policies and the new cannabis era (labor laws challenges)*
- *Common quality failures and threats*
- *Prepare for what's next in GMP compliance*

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Areas Covered

- *Current events associated with cannabis*
- *FDA, CANADA, and Europe regulations and standards*
- *What does it mean for drug manufacturing sites?*
- *Application from Pharma to Cannabis*
- *Our role from GMP standpoint*
- *Industry crossover*

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Attend the webinar to learn what the regulations say about medical cannabis, including GMP guidelines for pharmaceuticals, cannabis quality control regulations, and more.

PRESENTED BY:

Ginette Collazo, Ph. D. is an Industrial-Organizational Psychologist with 20 years of experience that specializes in Engineering Psychology and Human Reliability, disciplines that study the interaction between human behavior and productivity. She has held positions leading training and human reliability programs in the Pharmaceutical and Medical Device Manufacturing Industry.

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On-Demand Webinar

Duration : 90 Minutes

Price: $200

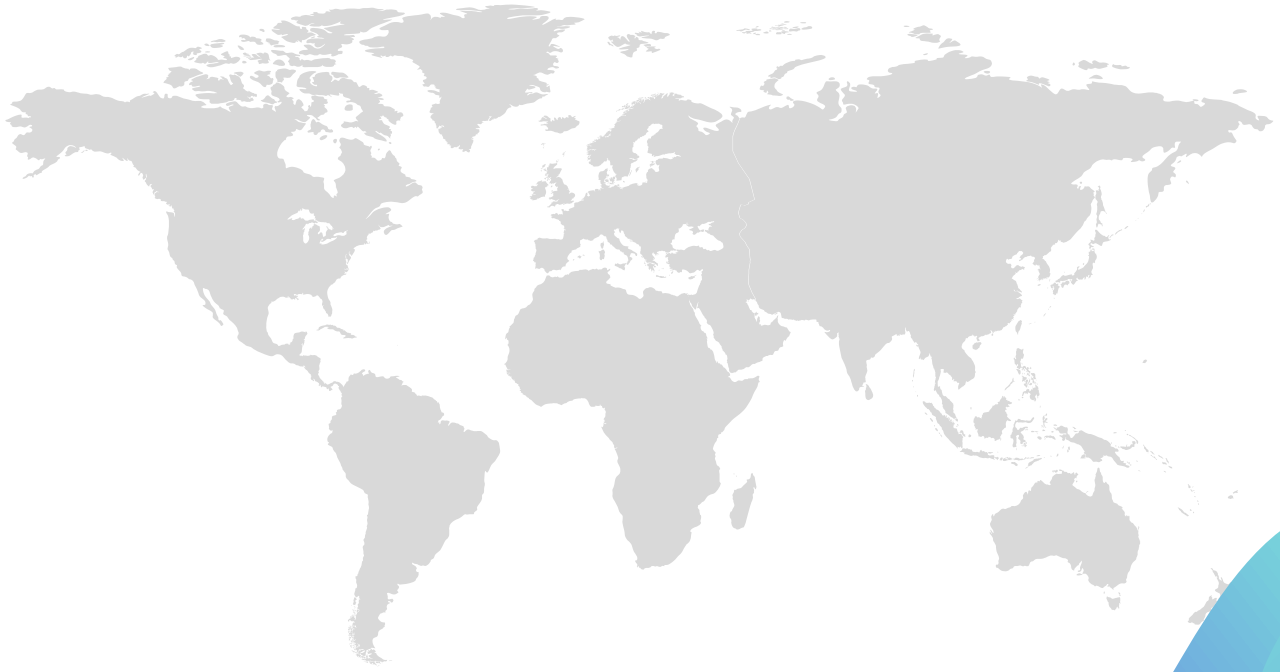
Webinar Description

When we discuss growing, producing, and manufacturing medical cannabis, we must think of it as a medicine. Medicine by definition is a substance intended to assist you with a medical condition, to help you feel better and not harm you. Drugs produced in the pharmaceutical industry go through extensive quality controls to ensure a level of safety for the consumer or patient. Yet when we talk process and quality controls in medical cannabis production, there is still a lot to learn.

With all-new events, the FDA has stated that it “recognizes the significant public interest in cannabis and cannabis-derived compounds, particularly CBD.” However, there are many unanswered questions about the science, safety, and quality of products containing CBD. The agency is working on answering these questions through ongoing efforts including feedback from a recent FDA hearing and information and data gathering through a public docket. In order to start understanding what it means for GMP regulated industries, we will discuss what we need to know, for now.

Who Should Attend ?

Training managers and coordinators
Operations
Manufacturing
Plant engineering
QA/QC staff
Process excellence/improvement professionals
Industrial/process engineers
Compliance officers
Regulatory/legislative affairs professionals
General/corporate counsel



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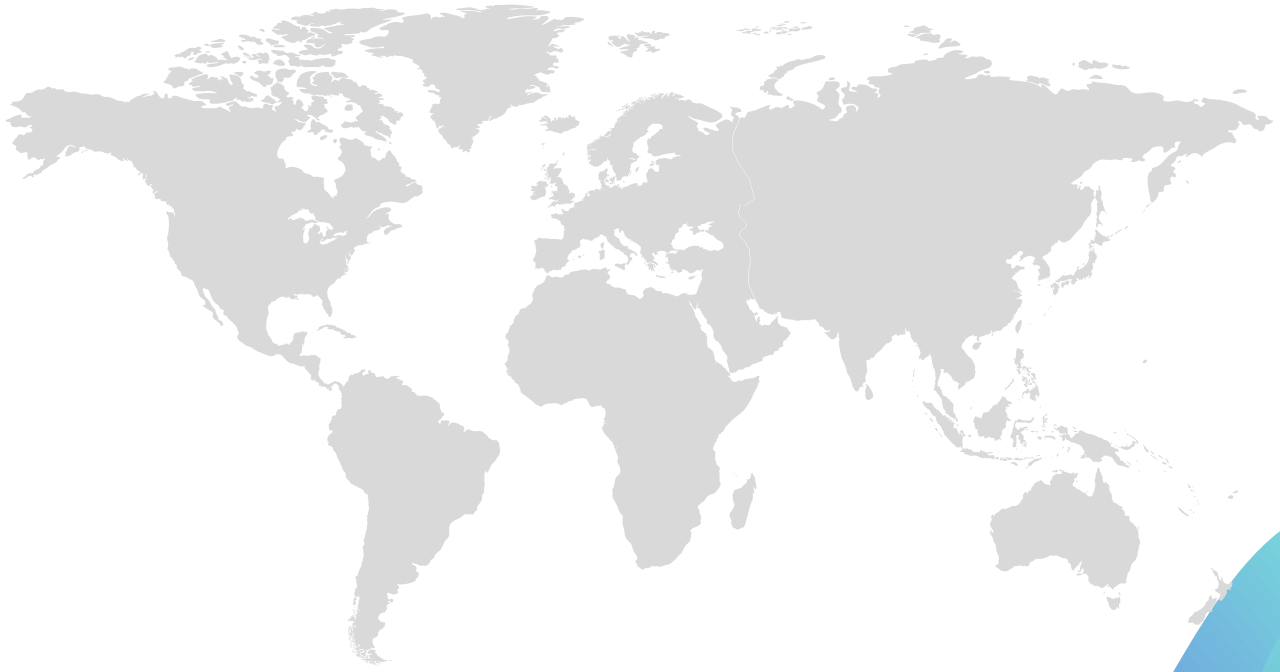

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www.grceducators.com
support@grceducators.com
740 870 0321