Basic GMP Training for the QC Laboratory

JAN 31- FEB 03 , 2022 AT 12:30 PM EDT-04:30 PM EDT 09:30AM- 01:30PM PDT

Learning Objectives

The Good Manufacturing Practice Certificate Program provides a comprehensive overview on the best practices regarding management of manufacturing and quality control testing for pharmaceutical products.
The Live Online Training is designed for people who have no or little knowledge of GMP.

- You get to know the most important pharmaceutical regulations and their importance,
- you get a basic overview of GMP requirements in pharmaceutical production, and
- you become familiar with technical terms from the field of GMP and their meaning
- Understand the regulatory requirements governing GMP compliant QC Labs.
- Understand how to apply those requirements
- Understand the regulatory expectations regarding laboratory design and utilities
- Understand the regulatory expectations regarding laboratory equipment / instrumentation

Who Will Benefit

- Research and Development
- Quality Control
- Quality Assurance
- Technical Operations
- Contract Laboratories
- Regulatory affairs personnel as well as individuals who are responsible for the review or audit of such laboratory data and reports should likewise consider the value of this comprehensive certification in their positions.

Training Overview

In the manufacture and quality control of medicinal products, compliance with the GMP rules is the decisive aspect for manufacturing high quality products. For this reason, every staff member in the pharmaceutical industry has to be familiar with the basic GMP requirements

Agenda

There are ten (10) areas of focus:

- An Introduction to Good Manufacturing Practice (GMP),
- CFR Title 21, Parts 58, 210, 211 Overview
  GMP in the Warehouse,
- Quality Control Laboratory Design
- Microbiology in the Workplace,
- Cleaning and Sanitation,
- Good Documentation Practices - GMP Laboratory
- Documentation and Record Keeping, Stability & Training
- Contamination Control
- Production Controls
- Packaging Controls
- Quality Assurance and Quality Control
- Pharmaceuticals Corrective Actions and Preventative Actions (CAPAs).
- The Regulatory Inspection
- Deviations / Non-conformances
- OOS / OOT
- How to write an effective investigation

REGISTRATION FORM

Name _____ **Email** _____

Organization _____ **Department** _____ **Position** _____

Phone _____ **Mobile** _____

One Dial-in One Attendee - Live	Live Group-Max. 05 Attendees
$1495	**$3995**

Register Online At www.worldcomplianceseminars.com

or by calling our offi ce : 844-267-7299

Terms And Condition

- Registration shall be reconfirmed only once payment has been made prior to the course.
- Cancellation of participant/s shall be submitted in writing to WCS ten days (10 days) before the course. There is a 50% liability on all bookings once made.
- WCS reserves the right to make alterations to the conference/executive briefing content, timing, speakers without notice. The event may be postponed or cancelled due to unforeseen events beyond the control of WCS. If such a situation arises, we will refund your registration fee or we will try to reschedule the event.