MAWA EVENTS

Address: No. 857, Block A2. Leisure Commerce Square - No 9., 46150 Petaling Jaya, Selangor, Malaysia

Phone: +601116373203 | **Email:** info@mawaevents.net

PHARMACEUTICAL GOOD MANUFACTURING PRACTICE

"Ensuring Quality, Safety, and Compliance in Pharmaceutical Production"

Schedule

Venue (InHouse)	Fees
At Your Organization Premises	Ask For The Quotation

► **Available delivery methods:** In-House Training

Introduction

This course provides an in-depth understanding of Pharmaceutical Good Manufacturing Practices (GMP), focusing on regulatory compliance, product quality, and safety. Participants will gain the knowledge and practical tools required to implement and maintain GMP standards throughout the manufacturing, packaging, and distribution processes to meet international quality benchmarks.

Objectives

- Understand the key principles and regulatory requirements of GMP.
- Learn how to design and maintain quality management systems in pharmaceutical production.
- Identify and control potential risks in manufacturing and supply chain processes.
- Ensure product consistency, safety, and compliance with regulatory authorities.
- Promote a culture of quality and continuous improvement in operations.

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Why Attend

This course is essential for professionals involved in pharmaceutical production and quality assurance, offering practical insights into maintaining GMP compliance and avoiding costly regulatory issues.

Target Audience

Pharmaceutical production managers, quality assurance and quality control personnel, regulatory affairs officers, laboratory staff, and anyone involved in manufacturing or auditing pharmaceutical operations.

Individual Benefits

- Gain practical knowledge of GMP principles and their application.
- Strengthen career prospects in pharmaceutical quality and compliance.
- Develop the ability to identify and resolve compliance issues effectively.

Organizational Benefits

- Improve product quality and ensure patient safety.
- Maintain compliance with international regulatory bodies such as WHO, FDA, and EMA.
- Minimize production errors, recalls, and compliance-related costs.

Instructional Methodology

Lectures, case studies, group discussions, practical workshops, and real-world examples from pharmaceutical manufacturing environments.

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Course Outline

Module 1: Introduction to GMP and Regulatory Framework

Module 2: Quality Management Systems in Pharmaceutical Manufacturing

Module 3: Documentation and Record-Keeping Requirements

Module 4: Personnel and Training in GMP Compliance

Module 5: Facility and Equipment Design for GMP Operations

Module 6: Production Controls and Validation Processes

Module 7: Handling of Deviations, Complaints, and Recalls

Module 8: Auditing and Continuous Improvement in GMP Environments

Certification

Participants who successfully complete the training will receive a Certificate of Completion in Pharmaceutical Good Manufacturing Practice (GMP), demonstrating their competence in maintaining compliance and quality within pharmaceutical operations.

Why Choose MAWA Events

- **Global Expertise:** More than 17 years of experience in professional training and consulting.
- **Industry-Leading Faculty:** Courses delivered by seasoned professionals with hands-on experience.
- **Practical Insights:** Learn to turn theory into actionable strategies for real-world business impact.
- **Client-Focused Solutions:** Customized programs designed to achieve your organisation’s unique goals.

In-House / Customized Training Interested in running this course for your team? Please contact us:	TEL: **+601116373203**	EMAIL: **info@mawaevents.net**