




Good Manufacturing Practice - The Essentials

 **Delivery:** Online, In-house, e-Learning

 **Duration:** 1 day

 **Cost:** £780 (+ VAT)

Course overview

This interactive course examines why we have Good Manufacturing Practice (GMP), its legal status, and the key GMP issues surrounding pharmaceutical manufacturing such as documentation, training, and facilities. This informative course introduces the EU Guide to GMP and considers its implementation in the modern pharmaceutical regulatory environment.

Who should attend?

This course is suitable for all pharmaceutical professionals, including newcomers to the industry, who wish to gain solid knowledge and understanding of the essentials of pharmaceutical GMP.

Course programme

This course covers the following topics:

- Expectations of medicines
- Why do we have GMP? What's involved?
- Pharmaceutical quality system
- The EU guide to GMP and the US CFRs
- GMP - who is involved? Roles of key personnel and their departments
- Documentation - what are the GMP requirements?
- GMP in the warehouse and dispensary
- GMP in production and packaging
- People and training
- Introduction to deviations

In addition, the course includes several interactive exercises

Learning outcomes

By the end of the course you will be able to:

- Find your way confidently around the EU Guide to EU GMP and how it is applied in the UK as the Orange Guide
- Understand the expectations of current Good Manufacturing Practices, and how these can be achieved at a practical level in the manufacture of pharmaceutical products

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