

pharmakon

Basic GMP

Understand GMP and learn how companies work with it in practice

Get a solid introduction to GMP and the EU's regulations for pharmaceutical manufacturing, and build a fundamental understanding of the requirements, responsibilities, and mindset needed to produce safe, high-quality medicines.

In this course, you will gain a basic knowledge of the rules and standards that form the foundation of Good Manufacturing Practice (GMP).

You will be introduced to the life sciences industry and the journey of a medicine to the market, so you understand the connection between legal requirements, quality, and patient safety. We will explore the requirements of the EU GMP guidelines (EudraLex Volume 4) in depth. You will work with key topics such as the pharmaceutical quality system, risk management, requirements for production, personnel, facilities and equipment, as well as documentation.

The training alternates between presentations, discussions, exercises, and GMP simulations. This approach provides both theoretical knowledge and practical understanding, as well as opportunities to network and exchange experiences with other participants. You will gain a broad understanding of why compliance with GMP is essential to ensure patients have access to safe, high-quality medicines.

This course can also be conducted on-site at the company - [see here](#)

You can find the course in Danish [here](#).

At the course you will meet:

Tine Gjerding Dahlberg, Educational Consultant, Pharmakon

Julie Dersch, Educational Consultant, Pharmakon

Sign-up dates

Hillerød

20. apr. — 21. apr.
2026

Course facts

FORMAT

2 days

RATING

4,77 out of 5

PRICE THIS YEAR

17.500 DKK excl. VAT,
overnight stay included

Do you have questions about this course?

Kontakt

Kursuskoordinator

Elizabeth Ruelykke

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Target group

The course is aimed at academics and technicians who are inexperienced with GMP, as well as new employees in the life sciences industry who need a basic understanding of GMP.

It is also suitable for those already working with GMP who want a deeper understanding of the rationale behind their work and the GMP regulations.

Your benefits

- you understand the fundamental principles of GMP and the structure of the EU GMP guidelines
- you can work more confidently with documentation, quality, and GMP requirements
- you can apply a basic GMP mindset and contribute to ensuring the quality of medicines

Your company's benefits

- your employee has an understanding of GMP, quality, GMP-mindset, and responsibility
- your employee works more consistently and correctly according to applicable regulations
- your employee contributes to improved quality and patient safety

Course agenda

The course is a two-day course. Accommodation is included.

Day 1: 09.00 – 21.00

- Introduction to the Life Sciences Industry
- The Path of a Medicinal Product to the Market
- Legal Framework and Structure of EudraLex Vol. 4
- Pharmaceutical Quality Systems and Risk Management
- Facilities and Equipment

Day 2: 09.00 – 16.00

- Production and Raw Materials
- Documentation and Good Documentation Practice
- Deviations and Changes
- Personnel, Behavior and Hygiene
- GMP Mindset in Practice
- Interactive GMP Game

The course consists of presentations, discussions and exercises.

Sign up online at

www.pharmakon.dk