pharmakon

Mastering GMP Inspections

Gain practical tools to prepare for, host and follow-up on regulatory GMP inspections

This course provides practical guidance on how to prepare your organisation, host the inspection process, and respond effectively after the inspection. A key strength of the course is the combination of practical industry experience and direct authority insight from an experienced GMP Inspector from the Danish Medicines Agency.

Regulatory GMP inspections place high demands on your organisation before, during, and after the inspection. This course gives you practical tools to prepare your organisation, host the inspection in a structured and professional way, and follow up effectively afterwards.

A key strength of the course is the combination of two perspectives. You will gain practical input from an experienced GMP and Quality consultant and direct authority insight from an experienced GMP Inspector from the Danish Medicines Agency. This gives you a stronger understanding of both inspection readiness and inspector expectations in practice.

You will learn how to prepare your organisation, documentation, systems, and facilities for inspection, how to communicate and act during the inspection, and how to respond clearly and effectively afterwards. Through expert input, real-life cases, discussions, and exercises, you will turn regulatory expectations into practical action you can use in your daily work.

The course language is English.

At the course you will meet:

Henrik Thrane, Principal Consultant, Compliance360 - Quality & GxP Consultancy

Thomas Vestergaard Pedersen, Team Leader & GMP Inspector, Danish Medicines Agency

Course facts

FORMAT
2 days

PRICE THIS YEAR
17.500 DKK excl. VAT,
Overnight stay included

Sign-up dates

Hillerød
30. sep. — 1. okt.
2026

Do you have questions about this course?

Kontakt
Kursuskoordinator
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Target group

This course is aimed at experienced professionals in GMP-regulated organisations who are involved in inspection readiness, inspection hosting, or follow-up. Relevant for QA, QC, RA, manufacturing, and site management. Not a basic GMP course.

Your benefits

- prepare your organisation for regulatory GMP inspections
- host inspections and communicate professionally with inspectors
- respond effectively to findings and follow-up requirements

Your company's benefits

- improve readiness for regulatory GMP inspections
- handle inspections in a more structured and professional way
- strengthen responses, follow-up, and organisational learning

Course agenda

The course duration is 2 days. Accommodation is included.

Day 1: 09.00 - 21.00
• The regulatory landscape of GMP inspections
• Preparation for the inspection
• Under inspection

Day 2: 09.00 - 15.45
• Time after the inspection
• DKMA inspections
• Follow-up to the inspection
• Time after inspection

The course combines expert presentations, authority and industry perspectives, practical cases, group exercises and Q&A.

Sign up online at
www.pharmakon.dk