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GMP Compliance Adviser

The most comprehensive GMP online knowledge portal worldwide.

What is the GMP Compliance Adviser?

The GMP Compliance Adviser is the world's largest online publication for quality management in the pharmaceutical industry, seamlessly combining theory and practice. It is trusted by more than 10,000 professionals in over 50 countries.

Content

GMP in Practice

Experts share their knowledge, empowering you to make confident decisions.

- Quality Management Systems
- Personnel
- Premises
- Facilities and Equipment
- Pharmaceutical Water
- Qualification
- Process Validation
- Cleaning Validation
- Computerised Systems
- Hygiene
- Production
- Sterile Production
- Packaging
- Laboratory Controls
- Documentation
- Storage and Transportation
- Outsourced Activities
- Audits and Supplier Qualification
- Quality Risk Management
- Active Pharmaceutical Ingredients
- Inspections and Drug Safety

GMP Regulations

Here you can find all relevant national and international GMP regulations.

- Addresses
- National Bodies and Pharmaceutical Associations
- EU Directives and Guidelines
- USA: CFR and FDA Guidelines
- ICH Guidelines
- PIC/S Guidelines
- GMP of other Regions
- WHO Guidelines

AI in the GMP Compliance Adviser

The AI assistant **GMP Chat** provides intelligent support. It is based exclusively on verified GMP Compliance Adviser knowledge and delivers precise, reliable, and well-founded answers – always transparent and fully referenced.

Your Benefits

- Stay up to date with all relevant GMP regulations at all times.
- Get specific instructions for action from our experts.
- Understand regulatory requirements from the authorities' perspective.
- Use ready-to-use resources such as templates, SOPs, checklists, flowcharts, and more.
- Face audits and inspections with confidence.
- Strengthen and continuously improve your GMP compliance.
- Receive regular updates and new content as regulations evolve.
- Get fast, reliable answers with the support of the GMP Chat AI assistant.

GOOD MANUFACTURING PRACTICE-KNOWLEDGE FOR YOUR SAFETY.

Contributors

Almost 100 experts from the pharmaceutical industry, authorities, suppliers and service providers share their expertise with users and help them to put GMP requirements into practice.

The contributors come from various disciplines such as pharmacy, chemistry and biology, engineering and technology.

Technical Requirements

All you need is a valid email address, an internet connection, and a modern web browser with the latest version installed.

Target Group

Managers in the pharmaceutical industry working in

- quality assurance,
- production,
- quality control,
- development.

Department managers from supplier industries also benefit, including those working in

- plant construction,
- consulting,
- design offices.

Licences and Prices

Named User Licence

Online access for one person
as a **3-month-subscription** 597,00 €*

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