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GMP:KnowHow Pharma Logistics (GDP)

The knowledge portal for the practical and up-to-date implementation of GDP-compliant storage and transport.

What is the GMP:KnowHow Pharma Logistics (GDP)?

The online knowledge portal GMP:KnowHow Pharma Logistics (GDP) offers comprehensive logistics knowledge for everyone involved in the supply chain of medicinal products and medical devices. It provides practical guidance on implementing the EU GDP guidelines and other relevant regulations. Editable sample documents such as SOPs and various quality documents are also available.

Background

The supply chain for medicinal products and medical devices is international and involves numerous actors. The understanding of GDP varies greatly depending on the country and industry. Different guidelines and laws apply and the interfaces to GMP must be considered.

To ensure patient safety, the supply chain of medicinal products and medical devices is subject to strict regulation and monitoring. In Europe, the EU GDP guidelines are the leading document.

Content

- Quality Management
- Personnel
- Facilities and Equipment
- Documentation
- Operation
- Complaints and Returns
- Outsourced Activities
- Self Inspection
- Transport

Tools

- Quality Agreement for GMP/GDP-compliant Logistics Services
- Quality Manual
- SOPs
- Job Descriptions
- GDP Questionnaire
- Site Master File with Organisational Chart
- Best Practice – the Peither Logistik GmbH

All documents can be edited individually.

Cited Regulations

- EU GDP guidelines (2013/C 343/01 and 2015/C 95/01)
- EU GMP Guide Parts I – III and Annexes
- Swiss TPA, MPLO
- PIC/S (PE 011-1, PI 047-1 Annex)
- WHO (GSDP and GTDP)
- DIN EN ISO 9001:2015, 13485:2016

Target Group

Employees in the supply chain of pharmaceuticals and medical devices, e.g.

- pharmaceutical companies
- medical device manufacturers
- wholesalers
- logistics service providers

Your Benefits

- Always up to date thanks to continuous updates
- Time savings through editable templates for SOPs, job descriptions, and other quality-relevant documents
- Direct comparison of topic-specific practical expertise and regulatory requirements
- Round-the-clock access (24/7) to all relevant content
- Clear and easy-to-understand presentation of complex topics using infographics

Author

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Simone Ferrante was previously Head of Quality Control and Responsible Person for the entire Grieshaber Group. Furthermore, she is a long-time author and GDP expert for the GMP-Verlag Peither AG.

Technical Requirements

You only need a valid email address, an internet connection, and a current version of a web browser.

Licences and Prices

Named User Licence

Online access for one person
as a **3-month-subscription** 460,00 €*

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