GxP Compliance Expert

m/f/d, Aargau

The GxP Compliance Expert is a member of the Network & Capabilities Chapter within the Clinical Distribution Team (Deliver), which is jointly accountable for all elements of clinical supply chain distribution for the global Roche/Genentech portfolio of >500 studies. This role drives GxP compliance and quality within Clinical Distribution (Deliver). Key duties include proactive GxP advice, managing deviations/CAPAs/improvements, acting as the audit/inspection SME, handling regulated documents, analysing quality trends, and developing the team's GxP training.

Start Date: Oct/Nov/Dec 2026
End Date: 12-month contract (with the option of extension)
Workload: 100%
Location: Kaiseraugst
Work Model: Hybrid, three days on-site and two days working from home possible

Tasks & Responsibilities:

- SME for all GxP and compliance-related topics (proactive advice on these matters) incl system administration (Veeva quality docs)
- Audit/Inspection SME for Deliver for non-study specific examinations
- Processing of deviations as well as CAPAs, Changes, Process Improvements
- Quality KPI Ownership for Deliver
- Responsibility for deviation management improvement processes
- Offer comprehensive compliance expertise to the business, conducting Root Cause Analysis (RCA) to identify and address quality, regulatory and compliance-related issues
- Managing the structure for the departments' SOPs and any other regulated documents (The business know-how is provided by the chapter's SMEs)
- Responsibility for trend reporting across Quality and Compliance requirements
- Implementation of GxP training and further development of employees' GxP knowledge

Job profile

Life Sciences - Supply Chain / Operations
Aargau
Contracting
Full Time
Oct/Nov/Dec 2026 - 12-month contract (with the option of extension)
Clinical Supply Chain, GxP

Sounds interesting?

Sophia Merkt
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Must Haves:

- You hold a degree (BA, MS, MBA) in a relevant field (e.g. Supply Chain Management & Logistics, Pharmacy, Biology, Chemistry, Industrial Engineering and Management), backed by significant expertise and extensive years of senior professional experience.
- At least 6 years of professional experience
- You possess extensive knowledge of cGMP standards alongside hands-on experience navigating internal (PTQ, Pharma-Technical Quality) and external (e.g., FDA, EMA) regulatory inspections.
- You bring a deep understanding of end-to-end supply chain management, specifically encompassing warehouse and transportation management, logistics, and clinical supply chain distribution processes.
- You demonstrate strong interpersonal, communication, organisational, and prioritisation skills with a steadfast focus on customer needs.
- You are fluent in both written and spoken German and English.
- You are proficient in using Google applications, MS Office (Word, Excel, PowerPoint), Visio, and key industry systems such as Veeva and Cornerstone.

Sounds like a great job?
Then we look forward to receiving your complete application documents through our online application form.

When applying by email, the sender agrees that his or her data will be used in accordance with our data privacy policy.

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