

QA Operations Expert (d/m/f)

Job ID
REQ-10068587

12月 15, 2025

Austria

摘要

负责管理责任范围内的质量方面, 并确保运营业务符合cGMP(现行良好生产规范) 质量保证协议, 监管要求和诺华质量手册, 并根据相关标准操作程序进行

About the Role

Major Accountabilities

- ~监督所有生产和测试活动, 确保符合cGxP, 包括数据完整性和电子合规性
- ~支持异常调查
- ~审查和批准生产、质量控制以及 AS 和 T 记录
- ~MBR审查
- ~支持运营支出改善项目

专业人员-根据注册执行批放行

收到后 24 小时内报告与诺华产品相关的技术投诉/不良事件/特殊情况
(营销样本的分发适用)

Key Performance Indicators

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Work Experience

职能广度
关键谈判
领导力
运营管理和执行
跨界协作
项目管理

Skills

员工绩效评估
人员管理
技术智能
QA(质量保证)
GMP程序
质量标准
质量控制 (QC)测试
处理歧义
自我意识
持续学习
技术专长

Language

英语

Why Novartis: Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients' lives. Ready to create a brighter future together? <https://www.novartis.com/about/strategy/people-and-culture>

Benefits and Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: <https://www.novartis.com/careers/benefits-rewards>

部门
Operations

Business Unit
Quality

地点
Austria

站点
Schaftenau

Company / Legal Entity
AT33 (FCRS = AT033) Novartis Pharmaceutical Manufacturing GmbH

Functional Area
Quality

Job Type
Full time

Employment Type
正式

Shift Work
No

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