

QC PLANNER - CDI (h/f)

Job ID
REQ-10023543

11月 07, 2024

France

摘要

管理责任范围内的质量方面和项目。
确保并支持整体GxP合规性和诺华质量管理体系的合规性。

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

管理责任范围内的质量方面和项目。
确保并支持整体GxP合规性和诺华质量管理体系的合规性。

Work Experience

在制药工业/生物技术领域从事环境监测和清洁区的QC/QA工作
职能广度
跨界协作

Skills

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>

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: <https://talentnetwork.novartis.com/network>

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
Innovative Medicines

地点
France

站点
Huningue

Company / Legal Entity
FR12 (FCRS = FR012) Novartis Pharma S.A.S.

Functional Area
Quality

Job Type
Full time

Employment Type
正式

Shift Work
No

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