

## 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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