

QC Analyst (H/F) - CDD

Job ID

REQ-10077358

May 14, 2026

LOC_FR

About the Role

Major Accountabilities

- ~ 为质量组织提供技术和/或行政支持。
- ~ 按照监管员的指示执行简单、常规、重复性活动。
- ~ 以有效、合规的方式遵循标准操作程序(SOP)和/或实验室方法。
- ~ 协助准备文件（如报告、记录、陈述等）。
- ~ 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

- ~ 客户满意度
- ~ 准点率
- ~ 按照指定的周期时间按时完成工作
- ~ 始终遵守GMP和健康、安全和环境指南以及标准操作程序
- ~ 监管检查零投诉

Work Experience

- ~ 技术知识
- ~ 运营管理和执行
- ~ 跨界协作

Skills

- ~ 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

Role Requirements

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: Learn about all the ways we'll help you thrive personally and professionally.

[Read our handbook \(PDF 30 MB\)](#)

Division

DIV_TO

Business Unit

Quality

Location

LOC_FR

Site

Huningue

Company / Legal Entity

FR12 (FCRS = FR012) Novartis Pharma S.A.S.

Functional Area

FCT_QA

Job Type

Full time

Employment Type

临时 (固定期限)

Shift Work

No

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