

QA Officer Operations

Job ID

REQ-10073737

May 15, 2026

LOC_ES

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

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

Production / Manufacturing

Location

LOC_ES

Site

Zaragoza

Company / Legal Entity

ES45 (FCRS = ES045) Advanced Accelerator Applications Iberica S.L.U.

Functional Area

FCT_QA

Job Type

Full time

Employment Type

正式

Shift Work

No

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