

## QC Analyst

Job ID  
REQ-10064103

10月 13, 2025

Singapore

### 摘要

此角色利用化学实验室技能测试和测量产品或材料，同时确保分析按照既定的标准操作程序 (SOP)、分析方法和当前的编年。

### About the Role

#### Major Accountabilities

- 样品存储和管理
- 药品/成品/投诉/稳定性/包装材料样品根据 GxP 标准进行分析测试/记录
- 测试/样品存储和管理。
- 稳定性样本与 GxP 标准的分析文档
- 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

此角色利用化学实验室技能测试和测量产品或材料,同时确保分析按照既定的标准操作程序SOP、分析方法和当前的编年。

## Work Experience

具备制药/化学分析/QC/同等的技术和科学知识  
在制药行业的实验室环境中的工作经验

## Skills

实验室设备  
质量控制 QC 测试  
质量控制抽样  
全面质量管理  
Gxp(吉特普)  
行业标准  
实验室卓越  
决策能力  
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  
Universal Hierarchy Node

地点  
Singapore

站点  
Tuas South Avenue

Company / Legal Entity  
SG12 (FCRS = SG012) Novartis Singapore Pharmaceutical Manufacturing Pte Ltd

Functional Area  
Quality

Job Type  
Full time

Employment Type  
正式

Shift Work  
No

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