# QC Specialist I - Analytical

Job ID
REQ-10042742

Apr 04, 2025

Singapore

## Summary

This role will be responsible to establish and ensure testing of drug substance release and stability testing including testing of intermediates in process control samples and lab operations are accordance with written testing SOP ' s and local/international regulations.

## About the Role

**Key Responsibilities:**

- Sample storage and management.
- Analytical testing and documentation of API / drug substance / drug product / finished product / Complaints / stability / packaging material samples
- Ensure all activities in compliance with cGxP, incl. data integrity
- Stability (when not centralized)
    - Testing/Sample storage and management

    - Analytical documentation of stability samples to cGxP standards
- Detect and report potential accident, risks and propose solutions

**Essential Requirements:**

- Preferred: Previous experience working in a laboratory environment in the pharmaceutical industry (quality assurance, production), aseptic technique.
- Administrative activities and GMP and HSE-compliant, efficient production and documentation of standardized tasks in the infrastructure
- Breakthrough Analysis; Being Resilient; Operational Excellence; Continuous Learning; Digital & Tech Savvy
- Laboratory equipment; Quality Control (QC) Testing; Quality Control Sampling; knowledge of TQM and related industry GxP standards and processes; Laboratory Excellence; Quality decision making
- Should be willing to work in shifts

**Desirable Requirements:**

- University degree or equivalent experience in Pharmacy or Chemistry or equivalent + 0-4 years working experience

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部門
Operations

部門
Innovative Medicines

国
Singapore

勤務地
Tuas South Avenue

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

Functional Area
Quality

雇用形態
Regular

Shift Work
No

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