QC Specialist I

Job ID
REQ-10004976

9月 03, 2024

Singapore

摘要

Skilled and experienced laboratory professional who contributes by performing analytical release testing, investigational support, research support, and stability testing.

About the Role

QC Specialist I

About the Role:

Skilled and experienced laboratory professional who contributes by performing analytical release testing, investigational support, research support, and stability testing.

Key Responsibilities:

- Maintain QC Raw Materials laboratory in full cGMP compliance.
- Lead raw material method validation/ verification and routine release testing
- Plan day to day operational activities in QCRM (i.e., housekeeping, release testing). Perform data entry, review and approval of RM packages for batch release.
- Lead improvement projects and perform technical reviews of procedures and testing monographs for raw materials.
- Lead laboratory investigations (e.g., OOS, deviation) and lead change controls for QC Raw Materials.
- Lead creation and revisions of RM testing monographs
- Prepare and participate in health authority inspections and internal audits
- Other duties or projects assigned by the QC Team Leader - Raw Materials

Essential Requirements:

- University degree in Pharmacy or Chemistry or equivalent
- 3-5 years relevant experience in Pharma/Manufacturing sector in analytical lab in a GMP environment
- Experience in Raw Material Lab and Method validation.
- Handling quality metrics & issues Knowledge of GMP Management of Quality Audit, Quality Change, Control Good Documentation

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部门
Operations

Business Unit
Innovative Medicines

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
Regular

Shift Work

No

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