QC Specialist I

Job ID
REQ-10004976
Sep 03, 2024
Singapore

Summary

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

About the Role

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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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Operations
Business Unit
Innovative Medicines
Location
Singapore
Site
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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