NOVARTIS

QC Specialist II

Job ID
REQ-10048888

4月 24, 2025

Singapore

摘要

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

About the Role

Key Responsibilities:

- OOx/Deviation handling .
- CAPA definition -KPI trending -Ensure all activities in compliance with cGxP, incl. data integrity review and approval of analytical data / tests (analytical release) Stability -Stability testing (Projects) - protocol preparation, evaluation, report preparation .
- Reporting (Stability plan preparation, trend analysis, evaluation) -Performance of Stability studies, protocols and comparative reports for supplier qualification -Review and approval of analytical tests (analytical release) -Microbiological QC -Perform Microbiological testing of

materials and utilities, environmental and personnel monitoring -Provide expert Support for site qualification and validation activities -Maintain and calibrate equipment incl. plan preparation -Support in supplier qualification -Trending and analysis of KPI/KQI -Support sample planning and sampling execution -Reporting of technical complaints / adverse events / special case scenarios related to Novartis products within 24 hours of receipt -Distribution of marketing samples (where applicable)

Essential Requirement:

- 3-5years experience in Pharma/Manufacturing sector in analytical lab in.
- Collaborating across boundaries.
- a GMP environment/equivalent.

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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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