QC Analyst IV - Analytical

Job ID REQ-10005670 Giu 04, 2024 Singapore

Sommario

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About the Role

QC Analyst IV

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Key Responsibilities:

- Ensure all activities in compliance with cGxP, incl. data integrity
- Review and approval of analytical data / tests (analytical release).
- Maintain and calibrate equipment incl. plan preparation
- Support in supplier qualification, trending and analysis of KPI/KQI.
- Support sample planning and sampling execution.
- 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.

Essential Requirements:

- Professional experience (3-5 years) in the pharmaceutical sector or in the manufacture of active substances in analytical laboratories in a GMP environment or equivalent; Collaborating across boundaries; Functional Breadth; efficient inter and intra-departmental communications.
- Breakthrough Analysis; Being Resilient; Operational Excellence; Continuous Learning; Digital & Tech Savvy.
- MS Office applications and other standard IT applications supporting Quality activities.
- Laboratory equipment; Quality Control (QC) Testing; Quality Control Sampling; Knowledge of TQM and related industry GxP standards and processes; Laboratory Excellence; Quality decision making

Desirable Requirements:

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

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Divisione

Operations

Business Unit

Innovative Medicines

Posizione

Singapore

Sito

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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