

Specialist - QA Ops - Manufacturing Management

Job ID REQ-10041626

2月 25, 2025

Singapore

摘要

This role support/provide quality oversight in ensuring a smooth manufacturing operation, new product launches/transfer in a compliant/timely manner, drug substance batch review/release are in full gmp compliance to regulatory standards and ensures quality strategy/continuous improvement are executed in alignment to site objective/s.

About the Role

Key Responsibilities:

- · Ensure all activities in compliance with cGxP, incl. data integrity
- Review and approval of analytical data / tests (analytical release)
- Oversight of all production and testing activities, ensures compliance with cGxP, incl. data integrity and eCompliance
- Support exception investigations

- · Review and approval of production, QC, and AS & T records
- MBR review. Support OpEx improvement projects. Executes batch release in compliance with registration (if Qualified Person)
- · Comply with all HSE guidelines. Detect and report potential accident, risks and propose solutions
- · Participate in HSE risk assessments. Preparation and participation to internal HSE audits

Role Requirements:

Essential Requirements:

- 5+ years of experience in pharmaceutical quality control, quality assurance or production
- Operations Management and Execution; Functional Breadth; Collaborating across boundaries;
 Applied Practice
- Collaboration; result-oriented. Good knowledge of GMP; Continuous Learning; Operational Excellence; Digital & Tech Savvy
- · MS Office applications and other standard IT applications supporting Quality activities
- Technological competence; Quality Assurance; Knowledge of GMP, Quality Standards; Quality Control (QC) Testing

Desirable Requirements:

• University degree with a scientific / technological background (e.g. Chemistry, Pharmacy, Biology, Biochemistry, or 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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