Specialist - QA Ops - Manufacturing Mgmt

Job ID
REQ-10007341
Sep 03, 2024
Singapour

Résumé

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 driven in alignment to site objective/s.

About the Role

Position Title : Specialist - QA Ops - Manufacturing Mgmt

Location – Singapore

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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)
• 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

Commitment to Diversity & Inclusion: :

We are committed to building an outstanding, inclusive work environment and diverse teams representative of the patients and communities we serve.

Role Requirements :

Essential Requirements:

• 3+ 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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Division
Operations
Business Unit
Innovative Medicines
Emplacement
Singapour
Site
Tuas South Avenue
Company / Legal Entity

SG12 (FCRS = SG012) Novartis Singapore Pharmaceutical Manufacturing Pte Ltd
Functional Area
Qualité
Job Type
Full time
Employment Type
Regular
Shift Work
No
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