U NOVARTIS

Specialist - QA Ops - Manufacturing Mgmt

Job ID REQ-10041624 Feb 24, 2025 Singapur

Summary

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

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:

- 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)

Why Novartis: Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients' lives. Ready to create a brighter future together? https://www.novartis.com/about/strategy/people-and-culture

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Benefits and Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: <u>https://www.novartis.com/careers/benefits-rewards</u>

Division Operations **Business Unit Innovative Medicines** Standort Singapur 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 Apply to Job

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

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List of links present in page

- 1. https://www.novartis.com/about/strategy/people-and-culture
- 2. https://talentnetwork.novartis.com/network
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- 4. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Tuas-South-Avenue/Specialist---QA-Ops---Manufacturing-Mgmt_REQ-10041624
- 5. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Tuas-South-Avenue/Specialist---QA-Ops---Manufacturing-Mgmt_REQ-10041624