U NOVARTIS

QC Analyst III

Job ID REQ-10048886 Apr 24, 2025 Singapore

Summary

-This role utilizes chemistry laboratory skills to test and measure product or materials while ensuring that analysis is performed according to established Standard Operating Procedures (SOPs), Analytical Methods & current Compendia.

About the Role

Key Responsibilities:

- Sample storage and management -Analytical testing/documentation of drug product / finished product / complaints / stability / packaging material samples to GxP standards Stability -Testing/Sample storage and management.
- Analytical documentation of stability samples to GxP standards -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:

- Sound technical & scientific knowledge of pharmaceutical/ chemical.
- Working experience in Laboratory environment in the Pharmaceutical.
- analytics/QC/ 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 Location Singapore 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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