

QC Analyst/ Specialist

Job ID
REQ-10022512
Sep 13, 2024
Singapur

Zusammenfassung

About the Role:

Execution of assigned tasks in the quality control laboratory in accordance with cGxP regulations.
Performance of laboratory specific activities such as analyses, maintenance, calibration and qualification of analytical equipment

About the Role

Position Title: QC Analyst / Specialist

Location – Singapore

Key Responsibilities:

- Sample storage and management
- Analytical testing and documentation of API / drug substance / drug product / finished product / Complaints / stability / packaging material samples
- Ensure all activities in compliance with cGxP, incl. data integrity
- Stability (when not centralized)
- o Testing/Sample storage and management
- o Analytical documentation of stability samples to cGxP standards
- Comply with all HSE guidelines
- Detect and report potential accident, risks and propose solutions
- Responsible for participating in initial training and retraining
- Able to support rotating shift hours (Day/night).

Role Requirements:

Essential Requirements:

- Preferred: Previous experience working in a laboratory environment in the pharmaceutical industry (quality assurance, production), aseptic technique.

- Administrative activities and GMP and HSE-compliant, efficient production and documentation of standardized tasks in the infrastructure
- Breakthrough Analysis; Being Resilient; Operational Excellence; Continuous Learning; Digital & Tech Savvy
- Laboratory equipment; Quality Control (QC) Testing; Quality Control Sampling; knowledge of TQM and related industry GxP standards and processes; Laboratory Excellence; Quality decision making
- Ensure proper maintenance of QC IPC/DS lab equipment and systems to ensure full cGMP-compliance as part of shift team.
- Perform product testing and analysis under cGMP to meet required timelines.
- Provide technical support to run and validate necessary test methods on lab equipment and in developing method transfer/validation protocols and reports.
- Perform routine testing for in process, release and stability test samples and validation samples.
- Support and validate necessary test methods on lab equipment under cGMP.
- Prioritizes workload to ensure documents are reviewed and testing is performed in a timely manner.
- Support and coordinate laboratory investigations and facilitates root cause finding.
- Prepare and check QC documents, including assays of least to average complexity, to ensure completeness, accuracy, consistency, and clarity and that materials or final products have been manufactured, tested, or inspected according to specification and cGMPs.
- Support the execution of improvements to optimize test procedures or efficiency whenever possible.
- Prepare and participate in health authorities inspections and internal audits in respective area.

Desirable Requirements:

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

Why Novartis: Our purpose is to reimagine medicine to improve and extend people's lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: <https://www.novartis.com/about/strategy/people-and-culture>

You'll receive: You can find everything you need to know about our benefits and rewards in the Novartis Life Handbook. <https://www.novartis.com/careers/benefits-rewards>

Join our Novartis Network: If this role is not suitable to your experience or career goals but you wish to stay connected to hear more about Novartis and our career opportunities, join the Novartis Network here: <https://talentnetwork.novartis.com/network>.

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>

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up:

<https://talentnetwork.novartis.com/network>

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

Abteilung

Operations

Business Unit

Innovative Medicines

Ort

Singapur

Website

Tuas South Avenue

Company / Legal Entity

SG12 (FCRS = SG012) Novartis Singapore Pharmaceutical Manufacturing Pte Ltd

Functional Area

Quality

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.

Job ID

REQ-10022512

QC Analyst/ Specialist

[Apply to Job](#)

Source URL: <https://www.adacap.com/careers/career-search/job/details/req-10022512-qc-analyst-specialist>

List of links present in page

1. <https://www.novartis.com/about/strategy/people-and-culture>
2. <https://talentnetwork.novartis.com/network>
3. <https://www.novartis.com/careers/benefits-rewards>
4. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Tuas-South-Avenue/QC-Analyst--

Specialist_REQ-10022512

5. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Tuas-South-Avenue/QC-Analyst--Specialist_REQ-10022512