

# Analyst – Quality Operations

Job ID  
REQ-10030263  
Nov 22, 2024  
India

## Summary

Provide quality services in compliance with cGMP requirements and Novartis Quality Management System as defined and agreed between QOP and business partners

## About the Role

### Major accountabilities:

- Perform and deliver Quality Operations services in support of product quality compliance and regulatory workflows
- Hold accounts in workflow applications (such as SAP, Dragon, SUBWAY, etc.) to ensure appropriate execution of service deliverables
- Support to Stability management eg: Draft reports/assessments of temperature excursion assessments (TEA), transport category assignment (risk assessment (TRA)).
- Support to QC release activities eg: Create, modify and review: Inspection Plans, Inspection Lot Numbers, Certificate Of Analysis, Certificate of Compliance, Specifications etc.
- Support to Testing Monograph management eg: Author testing monograph, Perform impact assessments etc.
- Authoring of risk evaluation reports for Nitrosamines both Step-1 & Step-2. Handling of risk evaluation reports with respect to country specific/local ones. Performing authoring activity in Subway software. Data collection and slides preparation which are required for weekly work stream leads call/Steerco meetings.
- Perform Statistical support, Performance trending and Business support.
- Ensure compliance to the Novartis internal quality standards, relevant regulatory requirements, filed product quality standards and service level agreements
- Comply with all internal functional operating procedures like time tracking, KPI reporting, ticket management tools and other internal systems and processes
- Learn & develop understanding to generate insights through data and digital
- Provide active support during internal and external audits.
- Adhere to the current GxP and compliance policies of Novartis

### Key performance indicators:

- On-time and GMP-compliant release of dosage forms -No complaints about inspections by authorities in your own area of responsibility without these being noticed and communicated beforehand -Successfully support continuous improvement projects

### Minimum Requirements:

- Pharmacy/ Science/ MBA / Engineering/ equivalent from a reputed institute
- Min 3 years of experience in Quality Assurance, Regulatory or in the manufacturing of pharmaceutical drug substances/ products/ medical devices
- GxP knowledge, Basic IT knowledge
- Good communication, presentation and interpersonal skills
- Experience of working closely with the global stakeholders

#### **Skills:**

- Continuous Learning.
- Dealing With Ambiguity.
- Gmp Procedures.
- QA (Quality Assurance).
- Quality Control (Qc) Testing.
- Self Awareness.
- Technological Expertise.
- Technological Intelligence.

#### **Languages :**

- Fluent in English (written and spoken)

**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>

Division

Operations

Business Unit

Innovative Medicines

Location

India

Site

Hyderabad (Office)

Company / Legal Entity

IN10 (FCRS = IN010) Novartis Healthcare Private Limited

Functional Area

Quality

Job Type

Full time

Employment Type

Regular

Shift Work

No

[Apply to Job](#)

## Accessibility and accommodation

Novartis is committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to perform the essential functions of a position, please send an e-mail to [diversityandincl.india@novartis.com](mailto:diversityandincl.india@novartis.com) and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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

Job ID

REQ-10030263

## Analyst – Quality Operations

[Apply to Job](#)

---

**Source URL:** <https://prod1.id.novartis.com/careers/career-search/job/details/req-10030263-analyst-quality-operations>

### 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/Hyderabad-Office/Analyst---Quality-Operations\\_REQ-10030263](https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Hyderabad-Office/Analyst---Quality-Operations_REQ-10030263)
5. <mailto:diversityandincl.india@novartis.com>
6. [https://novartis.wd3.myworkdayjobs.com/en-US/Novartis\\_Careers/job/Hyderabad-Office/Analyst---Quality-Operations\\_REQ-10030263](https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Hyderabad-Office/Analyst---Quality-Operations_REQ-10030263)