

Specialist-Quality Operations

Job ID
REQ-10024461
Oct 03, 2024
India

Summary

-Manages Quality aspects and projects within area of responsibility. -Ensures and supports overall GxP conformity and Compliance with the Novartis Quality Management Systems.

About the Role

Major accountabilities:

- Exposure in handling of customer/market complaints, field alerts, recalls and conducting investigations. Can independently handle meetings/ discussion with business partners/ suppliers.
- Experience in creating Quality assurance agreement and Quality risk assessment for Contract manufacturers / suppliers.
- Understanding on Audits & CAPA management for suppliers.
- Strong understanding on Quality Management System and GxP requirements.
- Provide support in preparation, review and approval of GMP documents including local and global SOPs, WP's, Investigation reports, etc., where required.
- Ensure implementation of applicable Novartis QMS requirements in the function.
- Provide support as key user/ Super user for IT tools used for Quality Management System.
- Provide timely and effective communication of any potential compliance gaps/ risks in respective function, the respective SPOCs or Team Lead and facilitate for resolution of identified gaps/ risks.
- Initiate, monitor and fulfill the timely review of APQR documents for all own site and Contract manufacturers.
- Provide quality support to Nitrosamine risk-based evaluation/ Changes, as required. Initiate and implement quality improvement/ simplification projects, wherever possible. Impart trainings on GMP/Data integrity and other relevant trainings, as required.
- Create various periodic performance metrics reports for respective GxP activities and share/ present to Business leaders/ partners.
- Author technical documents (like testing plans, testing monographs, Stability report etc.) and provide data driven technical and analytical insights to improve process understanding, quality and compliance of the product.
- Manage different type of change control like product stewardship/ Administration Stewardship /Asset Stewardship in electronic system like Agile PLM, from Change Initiation to closure as needed.
- Experience in Document Management tools and competency in MS office tools.
- Ensure all time readiness of the activities for internal/ Business partner audits (including data integrity audits), host audits, and manage audit action plans for timely closure of agreed CAPAs.

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>

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 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.

iframe{ width: 100%; margin-top: 3rem; } @media screen and (max-width: 767px){ iframe{ height: 30vh !important; } } @media screen and (min-width: 768px){ iframe{ height: 34vh !important; } }

Job ID

REQ-10024461

Specialist-Quality Operations

[Apply to Job](#)

Source URL: <https://prod1.id.novartis.com/id-en/careers/career-search/job/details/req-10024461-specialist-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/Specialist-Quality-Operations_REQ-10024461
5. <mailto:diversityandincl.india@novartis.com>
6. https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Hyderabad-Office/Specialist-Quality-Operations_REQ-10024461