U NOVARTIS

Specialist Quality Operations

Job ID REQ-10015901 Σεπ 03, 2024 India

Περίληψη

Provide quality support in compliance with cGMP requirements and Novartis Quality Management System. Manage Quality aspects & projects within area of responsibility.

About the Role

Major accountabilities:

Have expertise in Supplier Quality management and QMS activities. Drafting of QRA, QAA and AMR documents. Handling Supplier Qualifications and change notification.

Interpret and compile APQR and/ or extracted data from Internal Novartis systems into a pre-defined template and draft conclusion of product quality review.

Create and review GxP documents including SOPs, working procedures, trend reports, qualification

reports and technical investigations, as and when needed.

• Provide active support during internal and external audits by collecting and presenting the requested process/ data and reports

· Adherence to the current GxP and compliance policies of Novartis 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, TEDI etc.) to ensure

appropriate execution of service deliverables

• Generate and analyze predefined and ad-hoc reports in various applications (such as AGILE PLM, AQWA etc.) and perform follow-up actions if required

· Ensure compliance to the Novartis internal quality standards, relevant regulatory requirements, filed

product quality standards and service level agreements

· Support implementing service quality and process improvement projects, CAPA management within

Quality Service Centers

· Comply with all internal functional operating procedures like time tracking, KPI reporting, ticket

management tools and other internal systems and processes

Requirements for the role

- Minimum 6 years of experience in Quality assurance activities in pharmaceutical company.
- 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.
- Quality Standards.
- Self Awareness.
- Technological Expertise.
- Technological Intelligence.

Languages :

• English.

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: <u>https://www.novartis.com/careers/benefits-rewards</u>

Toμέας Operations Business Unit Innovative Medicines Τοποθεσία India Τοποθεσία 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 <u>Apply to Job</u>

Accessibility and accommodation

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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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