

Associate Manager - MS&T

Job ID
REQ-10015033
Jul 11, 2024
India

About the Role

Specialist / Associate Manager - MS&T

Location - Hyderabad

About the Role:

The purpose of the Investigation & Deviation Owner role is to work collaboratively with process experts and the multifunctional operations teams in Steriles/large molecules platform and take ownership of the deviation management for the site. The individual shall actively participate in investigations of Deviations/Complaints/OOXs by interacting with Cross Functional Teams (CFT) and implementation of Corrective and Preventive Actions (CAPA), Effectiveness Check (EC), Risk assessment and Quality management. The individual plays a key role in facilitating effective communication between teams and supporting problem-solving activities. The individual shall contribute to the enhancement of quality, productivity, and efficiency by supporting and driving improvements within the organization.

Key Responsibilities:

- The individual shall have a comprehensive understanding and experience of using Quality Risk management framework and shall possess excellent investigational report writing skills.
- The individual shall have hands-on experience of using structured RCA (Root Cause Analysis) methodologies such as impact assessment, Fish bone diagram, 5 whys, Timeline & process mapping for investigation of deviations.
- Experience in handling Investigations and Deviations related to Process (Upstream / Downstream), Product & Equipment
- Understanding of core manufacturing unit operations such as sampling, monitoring, and continuous process support.
- The individual shall have broad experience working in GxP environment and handling procedural requirements for HA audits.
- The individual will also be responsible for offering technical and scientific expertise to address process-specific matters, ensuring compliance with cGMPs, SOPs (Standard Operating Procedures), and relevant guidelines and functional standards (such as HSE (Health, Safety and Environment) and NOSSCE).
- Prior experience of handling internal and health authority audits and inspections is preferred
- Ensure overall inspection readiness for area of responsibility -Reporting of technical complaints / adverse events / special case scenarios related to Novartis products within 24 hours of receipt -Distribution of marketing samples (where applicable).
- Certification in investigation handling - Root cause analysis (RCA) is preferred.

Essential Requirements:

- Bachelor's degree in pharmacy, Pharmaceutical Technology, Chemical Engineering, Biomedical engineering, Biotechnology, Chemistry, or equivalent science streams. Desirable MSc/MS. or equivalent experience.
- Min 5 years of experience in MS&T, Quality Assurance, Regulatory or in the manufacturing of pharmaceutical drug substance or drug products in Sterile/Large Molecules platform/facility
- Minimum of 5 years of pharmaceutical process validation and cleaning validation.
- Should be familiar and able to perform basic statistical evaluations using Minitab or other statistical analysis tools.
- Proficient knowledge on deviation handling, incident investigations, root cause analysis, and CAPA management.
- Knowledge of risk assessment and risk management programs.
- Should be familiar with regulatory guidance on validation, product filing and post approval changes.
- Basic knowledge of statistical analysis, results interpretation, and usage of statistical tools (Example: Minitab, Statistica etc.).
- Good communication, presentation and interpersonal skills.

Desirable Requirements:

- Bachelor's degree in pharmacy, Pharmaceutical Technology, Chemical Engineering, Biomedical engineering, Biotechnology, Chemistry, or equivalent science streams. Desirable MSc/MS. or equivalent experience.

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>

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Role Requirements

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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Division

Operations
Business Unit
Innovative Medicines
Location
India
Site
Hyderabad (Office)
Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited
Functional Area
Technical Operations
Job Type
Full time
Employment Type
Regular
Shift Work
No
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