

QA Compliance Expert – Reg CMC Facilitator

Job ID REQ-10013378 Sep 03, 2024 India

Resumen

Supporting product maintenance, and activities throughout the product life-cycle using regulatory strategies and documents related to CMC (Chemistry, Manufacturing & Control). This applies to sector-specific (global and local) products and is intended to ensure timely market supply in compliance with regulatory requirements. Supporting change - and inspection management within the QA Compliance Team.

About the Role

QA Compliance Expert – Reg CMC Facilitator

Location - Hyderabad

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Key Responsibilities:

- Maintaining close cooperation with RA CMC to discuss regulatory requirements, strategies and knowledge of global product dossiers to stay up-to-date.
- Conducting training to ensure appropriate knowledge and regulatory compliance.
- Supporting the area in effective change control. Examination of reg. relevance and pre-evaluation amendments to Novartis products and customer products.
- Contact person for regulatory matters and intermediary between RA CMC and production unit at strategy decisions and in the product life cycle.
- Support of timely reviews of CMC documents for defined products; Support with and Identification of challenges in the course of regulatory compliance audits.
- Implementation and overview of initiatives to improve (regulatory) compliance.
- Coordination, guidance, and support in the preparation of CMC responses to health authorities for specific products.

Essential Requirements:

- Advanced University or academic degree in chemistry, biology, pharmacy, engineering or equivalent.
- Fluent English (German desired).
- More than 3 years of experience in an operational GxP area, in Manufacturing, Development or QA or $\frac{1}{3}$

Regulatory Affairs; with a thorough knowledge of biologic drug substance manufacturing processes for recombinant proteins and/or nucleic acids.

Ability to speak up and to take Quality decisions during challenging situations.

Desirable Requirements:

- Expertise in organization dynamics and culture, ability to gain trust and confidence at all levels in the organization, leadership, and project management experience.
- Ability to work independently and effectively in international, complex, and multifaceted environments.

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

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Ubicación

India

Sitio

Hyderabad (Office)

Company / Legal Entity

IN10 (FCRS = IN010) Novartis Healthcare Private Limited

Functional Area

Quality

Job Type

Full time

Employment Type

Regular

Shift Work

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.

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