NOVARTIS

# ESO Regulatory CMC facilitator

Job ID
REQ-10091301
Ноя. 24, 2026
Индия

### Сводка

#LI-Hybrid

Location: Hyderabad, India

Relocation Support: This role is based in Hyderabad, India. Novartis is unable to offer relocation support: please only apply if accessible.

Your regulatory expertise can help ensure medicines reach patients around the world without interruption. As an ESO Regulatory CMC Facilitator, you will play a key role in shaping and coordinating regulatory Chemistry, Manufacturing and Controls activities across External Supply, supporting product changes, transfers and lifecycle activities.

Working closely with cross-functional and global partners, you will help translate evolving regulatory requirements into practical solutions, maintain strong compliance and enable the timely supply of medicines to markets worldwide. This is an opportunity to bring your expertise, collaborative mindset and regulatory perspective to a role where your contribution can make a meaningful difference for patients.

### About the Role

**Key Responsibilities**

- Manage regulatory Chemistry, Manufacturing and Controls activities arising from product changes, transfers, divestments and remediation initiatives.
- Assess change requests and determine their regulatory impact as part of the External Supply change-control process.
- Coordinate complete, consistent source documentation for timely preparation of regulatory Chemistry, Manufacturing and Controls submissions.
- Facilitate high-quality regulatory documentation that supports efficient product registrations, lifecycle maintenance and ongoing compliance.
- Coordinate responses to health authority questions, ensuring accurate and timely regulatory information for External Supply products.
- Monitor regulatory commitments, required reports and supporting documentation to maintain compliance with applicable health authority requirements.
- Support product launches, transfers and changes by reviewing dossier compliance and relevant technical documentation.
- Build regulatory capabilities across External Supply by sharing regulatory intelligence, lessons learned and change-management expertise.

**Essential Requirements**

- University degree in Pharmacy, Chemistry, Biology or another relevant scientific discipline.
- Professional experience in a regulatory Chemistry, Manufacturing and Controls environment.
- Experience within a manufacturing, quality assurance, laboratory or comparable technical environment.
- Working knowledge of regulatory requirements and submission processes supporting product registration and lifecycle management.
- Experience collaborating across interdisciplinary or matrix teams while coordinating activities across multiple projects.
- Fluent English, with strong written and verbal communication skills.

**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

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

**Benefits and Rewards:** Learn about all the ways we'll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

Дивизион
Operations

Business Unit
Quality
Место
Индия
Сайт
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

### 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.

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