

Medical Lead, CML

Job ID
REQ-10043406

Mar 06, 2025

South Korea

Summary

- In line with overall product strategy, the Medical Lead is responsible for supporting the design, implementation and execution of Medical Affairs plans for assigned Therapy Area, providing scientific information, helping design & organise clinical studies, building educational dialogue with MEs and regulatory stakeholders and work collaboratively within the team and with cross functions
- #LI-Hybrid

About the Role

Major accountabilities:

- Ensure medical enquiries are responded to in a high quality, timely manner, and in accordance with applicable standards; establish standard response documents as appropriate, for frequently asked questions.

- Provide medical/scientific input into the development and execution of clinical trial or clinical research related activities, including initiation and oversight of clinical studies / clinical research within the respective therapeutic area.
- Support country strategy for Non Interventional Studies/Investigator Initiated Trial activities.
- Coordinate review and approval of medical materials and locally developed promotional materials; ensure medical materials provided from global or region for stakeholder engagement and events are tailored to local needs, and reviewed/approved per local/P3 guidelines.
- Ensure medical insights are provided to cross functional groups, including, but not restricted to: Pharmacovigilance, Regulatory affairs, Market Access, QA, Commercial teams, Brand team and others.
- Responsible for risk identification and assessment, mitigation planning as well as implementation and monitoring of appropriate internal controls within the area of responsibilities.
- Reporting of technical complaints / adverse events / special case scenarios related to Novartis products within 24 hours of receipt
- Distribution of marketing samples (where applicable)

Minimum Requirements:

- Experience: At least 3 years in pharmaceutical companies, including medical department or clinical practice roles.
- Clinical Design and Research: Expertise in creating clinical designs and analyzing English papers. Proactively updates and shares information.
- Innovative Environment Creation: Fosters innovation and new product launches while collaborating with related departments.
- Ethical Communication: Communicates clearly and flexibly with internal and external customers, based on ethical awareness.
- Team Collaboration and Independence: Respects team diversity, collaborates effectively, and works independently in various situations.
- Languages: English & Korean

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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do not discriminate in recruitment, hiring, training, promotion or other employment practices for reasons of race, color, religion, gender, national origin, age, sexual orientation, gender identity or expression, marital or veteran status, disability, or any other legally protected status. We are committed to building diverse teams, representative of the patients and communities we serve, and we strive to create an inclusive workplace that cultivates bold innovation through collaboration and empowers our people to unleash their full potential.

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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部門

International

部門

Innovative Medicines

国

South Korea

勤務地

Seoul

Company / Legal Entity

KR01 (FCRS = KR001) Novartis Korea Limited

Functional Area
Research & Development

職種
Full time

雇用形態
Regular

Shift Work
No

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