Medical Advisor, HEMA

Job ID
REQ-10020243
Oct 18, 2024
China

Resumen

About this role:
In line with overall product strategy, the Medical Advisor is responsible for supporting the design, implementation and execution of Medical Affairs plans for assigned Therapy Area, providing scientific information, helping design and organize clinical studies, building educational dialogue with KOLs and regulatory stakeholders

About the Role

Your key responsibilities:
Your responsibilities include, but are not limited to:
• Support country medical affairs strategy in line with the global strategy, country insights and market conditions, & secure implementation of planned Medical Affairs activities within the designated therapy area(s).
• Coordinate scientific meeting, symposia, congresses, Continuous Medical Education (CME) and other medical / scientific exchange and engagement activities which could bring additional value to the relevant therapy area; develop strategic engagement plan(s) for country customer-facing medical activities and events, and ensure timely execution of planned medical affairs activities in an efficient and compliant way.
• 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.

What you'll bring to the role:
•Education & Qualifications: Masters' degree or above, medical background
•Languages: English& Chinese
•Collaborating across boundaries
•Medical advisor and new launch experience preferred
•Operations Management and Execution

•Project Management

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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Novartis is committed to building an outstanding, inclusive work environment and diverse team's representative of the patients and communities we serve.

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.china@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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Benefits and Rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: https://www.novartis.com/careers/benefits-rewards

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Job Type
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