Clinical Research Medical Advisor - Oncology

Job ID
393316BR
Mai 03, 2024
Espagne

Résumé

Clinical Research Medical Advisor - Oncology, Spain or UK, #LI-Remote. The Clinical Research Medical Advisor is a Global Clinical Development role with medical responsibility for the clinical trial process. It is a bridge between Study Site Operations (SSO) clinical trials and Medical Affairs, aligning technical, operations & strategy. Based remotely in Spain or alternatively hybrid onsite in Barcelona, we are ideally searching for a Medical Doctor (MD) Oncologist with Clinical Development experience but will consider MD, PhD or Scientist level applications from other therapy areas

About the Role

Key Responsibilities:
• 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.
• 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.
• 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 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

Essential Requirements:
• An MD (oncologist ideally) or a PhD or Scientist with experience of being a Medical Advisor on clinical trials
• Experience working on Global Clinical Trials including Clinical Trial Design, Data & Reporting
• Knowledge of both SSO Operational Clinical Trials and Medical Affairs
• Operations Management and Execution Project Management collaborating across boundaries
• Strong people and strategic planning skills to pull several different functions together on projects
• Strong communication and negotiation skills pus the ability to speak, read and write in English (essential)
• A background as a Medical Science Liaison, Medical Affair Specialist or Study Lead/coordinator from a CRO as a Medical Advisor on a clinical trial would be considered

Desirable Requirements:
• Previous clinical experience with patients

• Oncology experience is preferred but other therapy areas will be considered

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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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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.

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Functional Area
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