Senior Clinical Development Medical Director* (80-100%**) " >

Job ID
325251BR
Aug 11, 2021
Switzerland

Job Description

Senior Clinical Development Medical Director*, Gastrointestinal cancers

22! the number of oncology drugs we have on the market and more than 25 new molecular entities in development targeting key molecular pathways in cancer biology. And while we're proud of that, in this world of digital and technological transformation, we must also ask ourselves this: how can we continue to improve and extend even more people's lives?

We believe the answers are found when curious, courageous and collaborative people like you are empowered to ask new questions, make bolder decisions and take smarter risks. In Clinical Development & Analytics, our aim is to design innovative, patient friendly clinical development plans to rapidly bring outstanding treatments to patients, caregivers and healthcare systems. To achieve this goal we rely on a dedicated team of experts, including the Clinical Development Medical Director.

The Senior Clinical Development Medical Director (Senior CDMD) is responsible for leading the strategic planning and management of the assigned clinical program(s) from an end-to-end clinical development perspective. As Senior CDMD, you will have oversight of the clinical development for the assigned programs and drive execution of the clinical development plan. In addition, you will enable an empowered organization, which can navigate in a matrix environment and adjust quickly to business needs.

Your responsibilities will include, but are not limited to:
• Providing clinical leadership and strategic medical input for all clinical deliverables in the assigned project or section of a clinical program
• Leading development of clinical sections of trial and program level regulatory documents
• Driving execution the assigned clinical program and/or clinical trial in partnership with global line functions, assigned Global Trial Directors (GTDs), and regional/country medical associates, where applicable
• Supporting (Senior) Global Program Clinical Head (GPCH) in ensuring overall safety of the molecule for the assigned section, and may act as a core member of the Safety Management Team (SMT), supporting overall program safety reporting in collaboration with Patient Safety colleagues
• Supporting the Clinical Development Head (CDH) by providing medical input into Clinical Development Plan (CDP), Integrated Development Plan (IDP) and Clinical Trial Protocol (CTP) reviews, and contributing to/driving development of disease clinical standards for new disease areas
• As a medical expert, supporting the (Sr.) GPCH or CDH in interactions with external and internal stakeholders and decision boards
• May work with NIBR (Novartis Institute of Biomedical Research/ Translational Medical Sciences) to drive transition of pre-PoC (Proof of Concept) projects to DDP (Development Decision Point) and with BD&L (Business Development & Licensing) including target identification and due diligences together with other

medical matters, as needed.

Minimum Requirements

What you bring to the role:
• MD or equivalent medical degree is required in addition to advanced knowledge and clinical training in medical/scientific area; Clinical practice experience 4 years (including residency) and board certification or eligibility in oncology or hematology preferred
• Minimum of 7 years of experience in clinical research or drug development
• Experience in an academic clinical research or industry environment spanning clinical activities in Phases I through IV required. • 2 years of contribution to and accomplishment in all aspects of conducting clinical trials (e.g., planning, executing, reporting and publishing) in a global/matrix environment in pharmaceutical industry required.
• Working knowledge of Oncology is required, with proven ability to interpret, discuss and present efficacy and safety data relating to clinical trial(s) and proven ability to understand and interpret basic and clinical scientific research reports
• Demonstrated ability to establish effective scientific partnerships with key stakeholders
• Working knowledge of GCP, clinical trial design, statistics, and regulatory and clinical development processes
• Previous global people management experience is preferred, though this may include management in a matrix environment.

*Final job title and associated responsibilities will be commensurate with the successful candidates' level of expertise.

**Some restrictions to flexible working models may apply and will be discussed at interview if applicable

Why consider Novartis?

769 million. That's how many lives our products touched in 2020. And while we're proud of that fact, in this world of digital and technological transformation, we must also ask ourselves this: how can we continue to improve and extend even more people's lives?

We believe the answers are found when curious, courageous and collaborative people like you are brought together in an inspiring environment. Where you're given opportunities to explore the power of digital and data. Where you're empowered to risk failure by taking smart risks, and where you're surrounded by people who share your determination to tackle the world's toughest medical challenges.

Imagine what you could do at Novartis!

Commitment to Diversity & Inclusion:
Novartis embraces diversity, equal opportunity and inclusion. 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.

Join our Novartis Network: If this role is not suitable to your experience or career goals but you wish to stay connected to learn more about Novartis and our career opportunities, join the Novartis Network here: https://talentnetwork.novartis.com/network

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