Clinical Program Leader

Job ID
REQ-10013760
Aoû 28, 2024
Suisse

Résumé

#LI-Hybrid

6000! That's the number of associates in the BioMedical Research (BR). This division is the innovation engine of Novartis, focusing on powerful new technologies that have the potential to help produce therapeutic breakthroughs for patients!

Based on our understanding of cancer at the molecular, cellular and organismal levels, Translational Clinical Oncology or TCO will design and execute innovative exploratory clinical trials in order to develop the next generation of highly effective therapeutics that transform the lives of patients with cancer. This role is to provide strategic medical guidance and lead the development of experimental oncology agents in the TCO portfolio, from the Phase of preclinical development, continuing through clinical First in Human and Phase 1b/2 studies.

About the Role

Your responsibilities will include but are not limited to:
• Acts as a clinical leader responsible for assigned global clinical program(s) -driving medical strategy implementation and operational deliverables for investigational products in a defined therapeutic area
• Ensure effective cross-functional communications to align with global strategy and leads the development of clinical sections of trial and program level regulatory documents

• Acts as the medical expert, engages interactions with external stakeholders (e.g., regulatory authorities, key opinion leaders, data monitoring committees, advisory boards etc.) and internal NVS stakeholders
• Contributes to medical/scientific training of relevant Novartis stakeholders. May serve as speaker for medical/scientific training -May serve on or lead global initiatives (e.g., process improvement, training, SOP development, other Clinical Development line function initiatives)
• Experience leading early and/or late phase Oncology clinical programs from the pharma/biotech industry plus credible experience from an academic medical center. In case of no industry experience then significant academic experience with credible PI/co-PI clinical study leadership
• Track record of significant contributions to your field over time, creating new concepts and seeking out new clinical discovery opportunities / approaches

Role Requirements:

Medical degree, oncology board certified preferred, and PhD level basic Science or equivalent expertise
· ≥ 3 years technical, operational and managerial experience in planning, executing, reporting and design of

clinical trials studies in a pharmaceutical company, academic centre or contract research organisation
· Good knowledge of Good Clinical Practice, clinical trial design, statistics, regulatory processes, and global clinical development process
· Good knowledge of oncology and experience in early clinical development preferred.
· Good communication, writing and organization and skills, fluent in English written and spoken

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

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

Join our Novartis Network: Not the right Novartis role for you? Sign up to our talent community to stay connected and learn about suitable career opportunities as soon as they come up: https://talentnetwork.novartis.com/network

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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Site
Basel (City)
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C028 (FCRS = CH028) Novartis Pharma AG
Functional Area
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Job Type
Full time
Employment Type
Regular
Shift Work
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