NOVARTIS

Clinical Sciences Trial Leader/Senior Clinical Sciences Trial Leader, Translational Medicine (Multiple Listings) 80-100%

Job ID
REQ-10079860
Jun 07, 2026
LOC_CH

About the Role

Key Responsibilities:

- Operationally and scientifically lead complex clinical studies
- Drive and deliver aspects of global clinical trial execution from study design, set-up and recruitment to final reporting
- Lead the matrix-management of robust global clinical trial teams, collaborating across the organization and externally with service providers/investigator sites
- Drive the development of clinical study protocols, clinical operations execution plans and other required documents
- Ensure quality conduct and document standards are applied across the clinical trial lifecycle
- Lead clinical data review (Sr. Clinical Sciences Trial Leader)
- Support clinical data review (Clinical Sciences Trial Leader)
- Contribute to project/program level plans (Sr. Clinical Sciences Trial Leader)
- May mentor junior team members
- Support optimization of standard methodologies in clinical trial operations and adopt an open learning and sharing environment
- Unleash the value of data and digital within clinical studies Ultimately, YOU will help bring new technologies and therapies closer to our patients

Essential Requirements:

- A minimum of BSc in life sciences is required. A MSc, PharmD or PhD is desired.
- Clinical Sciences Trial Leader: 2-4 years' experience, in clinical trial management/operations, coupled with a broad knowledge of the drug development field; preferably within the pharma industry.
- Sr. Clinical Sciences Trial Leader 4+ years' experience in clinical trial management/operations, coupled with a broad knowledge of the drug development field; preferably within the pharma industry.
- Clinical Sciences Trial Leader level: Leadership potential and well-developed interpersonal skills.
- Sr. Clinical Sciences Trial Leader level: Demonstrated leadership experience and well-developed interpersonal skills.
- A track record of collaborating with and influencing a wide range of people, and of building strong partnerships
- Strong project management experience; excellent planning, prioritization and organizational skills; used to managing multiple priorities concurrently
- High change agility, thriving in an open and dynamic environment.
- Able to learn proactively, tackle issues and take accountability
- Clear written and verbal expression of ideas; an active communicator

Desirable Requirements:

- Experience in study design and protocol development/writing is highly desirable

This is a dual posting. The final level & title of the offer role would be determined by the hiring team based on the skills, experience & capabilities required to perform the role at the level the role has been offered.

Accessibility and accommodation:
Novartis is committed to working with and providing reasonable accommodation to all individuals. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to receive more detailed information about the essential functions of a position, please send an e-mail to inclusion.switzerland@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message

Role Requirements

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

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally.
Read our handbook (PDF 30 MB)

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Site
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