

Senior Clinical Operations Manager

Job ID
REQ-10076583
апр 27, 2026
Индия

Сводка

Provide or supervise operational and logistical support to clinical trials in Biomedical Research (BR) with focus on high complexity and/or priority status. Oversee junior colleagues' activities related to study operations. Acts as a subject matter expert and/or cross-functional liaison.

About the Role

Major accountabilities:

- Provides operational and logistical support to clinical trials in Biomedical Research (BR) with focus on high complexity and/or priority status, in compliance with Novartis processes and Good Clinical Practice (GCP).
- Perform or supervise defined activities to support the Clinical Trial Team (CTT) throughout the study lifecycle, via study assignment and/or on-demand support.
- Maintain, share, and actively disseminate up-to-date knowledge of ICH-GCP, external regulations, and internal procedures. Continuously strengthen expertise through training and practical application of Novartis Standard Operating Procedures (SOPs) and internal policies.
- Lead finalization and management of clinical, regulatory and study-related documents in scope of role such as study protocols, patient-facing documents, etc.
- Responsible for ownership and leading the preparation of in-scope study documents, e.g. Clinical Study Report (CSR) appendices, patient-facing documents, protocol training material, etc.
- Lead or oversee interactions and communications with relevant functions including Novartis country organizations to prepare, collect, and/or compile relevant documents, and timely follow-up on pending actions as necessary.
- Lead or oversee external communication, such as newsletter development, external meeting organization.

Key performance indicators:

- Timely, efficient and quality execution of trial related activities in compliance with Novartis processes and GCP.
- Proactive operational planning with effective contingency and risk mitigation plans.
- High quality contributions to study or submission documents (i.e. study protocols, patient-facing documents)
- Strong leadership skills to be able to support management in team competency building, lead/contribute to local/global initiatives and best practice sharing across programs and/or departments -Clearly demonstrates Novartis Values and Behaviors (i.e. Innovation, Quality, Collaboration, Performance, Courage and Integrity).

Minimum Requirements:

- At least 3 to 4 years' experience in clinical trial /development
- Project management experience evidenced by the ability to organize effectively and deliver results. Adept organizational skills and quality mindset with attention to detail.
- Effective communication (verbal and written); proactive communication skills.
- Well-developed interpersonal skills with the ability to build rapport, manage, and coach others. Solid presentation skills and ability to mentor / train small groups.
- Ability to successfully interact with a wide range of people, including global teams, different cultures, diverse experience backgrounds, etc.

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>

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Дивизион
Biomedical Research
Business Unit
Research
Место
Индия
Сайт
Hyderabad (Office)
Company / Legal Entity
IN10 (FCRS = IN010) Novartis Healthcare Private Limited
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No

Accessibility and accommodation

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