Global Feasibility Lead / Clinical Operations Program Associate Director

Job ID
REQ-10057885

7月 24, 2025

United Kingdom

摘要

#LI-Hybrid (3 days per week on-site)
Location: London (The Westworks)

This role is based in London, UK. Novartis is unable to offer relocation support for this role: please only apply if this location is accessible for you.

Are you passionate about driving innovation in clinical development? At Novartis, we ' re seeking a strategic and analytical leader to join our Global Clinical Operations team as a Clinical Operations Program Associate Director. In this pivotal role, you will oversee and coordinate early modeling viability assessments, global feasibility evaluations, recruitment projections, and site allocation strategies across assigned trials and programs.

You ' ll be at the forefront of strategic planning, leveraging data-driven analysis and global-local insights to shape the success of our clinical trials–making a meaningful impact on global health outcomes.

About the Role

Major accountabilities:

- **Analyse diverse data sources** such as clinical trial data, historical performance, scientific publications, epidemiology, and the broader clinical and commercial landscape to inform strategic decisions.
- **Deliver modeling and viability analyses** to support early recruitment projections and guide clinical trial planning.
- **Lead end-to-end feasibility assessments** in close collaboration with country feasibility managers, clinical trial teams, and medical experts.
- **Recommend optimal country and site allocations**, identifying associated risks and opportunities in alignment with the global program strategy.
- **Provide strategic input on study design** based on data insights and feasibility findings.

Minimum requirements:

- A degree in Life Sciences or a related scientific discipline.
- A proven background in pharmaceutical clinical drug development.
- A solid understanding of all aspects of clinical development, with a particular focus on clinical trial design and execution.
- Demonstrated expertise in data-driven feasibility for global clinical trials.
- The ability to collaborate effectively in a dynamic, matrixed environment.

Desirable requirements:

- A proactive approach to risk management, with a track record of identifying, flagging, and resolving potential issues, along with experience in strategic scenario planning.
- Strong communication skills, with the ability to engage effectively across local, regional, and global matrixed teams.
- Excellent project management capabilities, with a knack for problem-solving and conflict resolution.
- A data-savvy mindset, with the ability to interpret analytical insights and clearly communicate the rationale behind decisions.
- A solid understanding of the international drug development process, including GCP/ICH standards and health authority regulations.
- Proficiency in Microsoft Office tools, particularly MS Teams, Excel, and Word.

Commitment to Diversity and Inclusion / EEO paragraph

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

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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

部门
Development

Business Unit
Universal Hierarchy Node

地点
United Kingdom

站点
London (The Westworks)

Company / Legal Entity
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.

Functional Area
Research & Development

Job Type
Full time

Employment Type

Regular

Shift Work
No

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