Clinical Research Associate

Job ID
REQ-10055247

6月 27, 2025

United Kingdom

摘要

Location: Field Based

Relocation Support: This role is field based. Novartis is unable to offer relocation support: please only apply if accessible.

Step into a role where your work truly matters. As a Clinical Research Associate (CRA) at Novartis, you ' ll help advance innovative treatments by ensuring excellence in clinical trial execution. Working within Study & Site Operations in Global Drug Development (GDD), you ' ll manage site relationships and oversee on-site and remote monitoring from initiation through completion of Phase I-IV trials. You ' ll ensure compliance with ICH/GCP, local regulations, and SOPs, while proactively managing site performance, recruitment, and quality. As the main point of contact for sites, you ' ll identify needs early and resolve issues efficiently. Join a passionate team, support breakthrough science, and help deliver life-changing therapies to patients faster.

About the Role

Key Responsibilities:

- Manage study sites to ensure compliance with protocols, regulations, and Novartis procedures
- Conduct site initiation visits and deliver tailored training to site personnel
- Perform ongoing site monitoring to ensure data quality and patient safety
- Implement site management strategies to address compliance and operational issues
- Maintain accurate and timely documentation of all monitoring activities
- Ensure continuous updates to global and local electronic systems
- Identify, resolve, and escalate site-related issues as appropriate
- Maintain up-to-date Trial Master Files and collect essential site documents
- Support audit and inspection readiness, ensuring timely corrective actions
- Collaborate with internal teams to support recruitment and contingency planning

Essential Requirements:

- Degree in a scientific or health-related discipline; advanced degree preferred
- Fluent in written and spoken English
- Up to two years of experience in clinical research, with focus on monitoring and trial execution
- Strong knowledge of international standards including GCP and ICH
- Proven ability to manage sites independently and resolve issues proactively
- Willingness and ability to travel domestically and internationally as needed
- Full UK driving license

Desirable Requirements:

- Experience working in a global pharmaceutical or CRO environment

Commitment to Diversity & Inclusion:

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

部门
Development

Business Unit
Universal Hierarchy Node

地点
United Kingdom

站点
Field Force (England / Wales)

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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