Clinical Research Medical Advisor

Job ID
REQ-10056412

8月 08, 2025

Brazil

摘要

Supervisa la ejecuci ó n e interpretaci ó n de investigaciones de ensayos cl í nicos, actividades de recopilaci ó n de datos y operaciones cl í nicas. Establece y aprueba m é todos cient í ficos para el dise ñ o e implementaci ó n de protocolos cl í nicos, sistemas de recopilaci ó n de datos e informes finales. Ayuda en investigaciones cl í nicas nuevas y constantes y en ensayos cl í nicos y asegura la eficiencia y el oportuno procesamiento de acuerdos de confidencialidad y acuerdos cl í nicos. Superv. el cumplim. de los protoc. y determ. la terminaci ó n de los estudios. Gestiona archivos cl í nicos y reglamentarios y mantiene el inventario cl í nico previsto para la distribuci ó n a sitios de investigaci ó n. Puede interactuar con sitios de investigaci ó n, consultores cl í nicos, Organizaciones de Investigaci ó n de Contratos y otros proveedores. Selecciona, desarrolla y eval ú a personal para asegurar la operaci ó n eficiente de la funci ó n.

About the Role

Key Responsibilities

- Validate study designs and assess trial feasibility based on clinical practice and competitive analysis.
- Drive fast and high-quality trial site start-up through expert input during planning phases.
- Provide clinical expertise for IRB/EC interactions and informed consent content.
- Develop trial plans that address recruitment challenges and ensure data quality.
- Deliver robust training on indications, compounds, and protocols to internal and external stakeholders.
- Lead clinical recruitment strategies using physician insights and patient engagement.
- Support regulatory inspections and audits with scientific and clinical expertise.
- Ensure adherence to safety standards and provide medical input on adverse events.

Essential Requirements

- Advanced scientific degree (M.D. highly preferred; Ph.D. or Pharm.D. also considered).
- Strong understanding of clinical development processes and ICH/GCP guidelines.
- Minimum 3 years of experience in clinical development or clinical practice.
- Proven ability to lead cross-functional teams and resolve complex clinical issues.
- Excellent communication skills in English and the local language.
- Ability to deliver high-quality presentations and adapt across therapeutic areas.

Desirable Requirements

- Subspecialty training or experience with Real World Evidence (RWE).
- Familiarity with innovative study designs and data sources such as registries or electronic health records.

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部门

Development

Business Unit
Innovative Medicines

地点
Brazil

站点
Santo Amaro

Company / Legal Entity
BR03 (FCRS = BR003) NOVARTIS BIOCIENCIAS S.A

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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