NOVARTIS

Clinical Development Medical Manager

Job ID
REQ-10090055

十月 26, 2026

China

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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

Major Accountabilities

-Supervisa programas cl í nicos en todas las indicaciones que ejecutan la estrategia m é dica para desarrollo y productos comercializados en un á rea terap é utica definida.
-Es responsable de asegurar la comunicaci ó n alineada con Asociados en Investigaciones Cl í nicas del Pa í s/la Agrupaci ó n, Gerentes y otros interesados clave en la ejecuci ó n y el progreso de estudios cl í nicos.
-Identificar nuevos sitios para ensayos cl í nicos; analizar la capacidad y hacer recomendaciones para la inclusi ó n de ensayos.
-Facilitar la preparaci ó n y recopilaci ó n de documentos a nivel sitio; resolver problemas seg ú n la necesidad.
-Puede ejecutar iniciaci ó n y capacit. del sitio.
-Implem. la gesti ó n total del sitio incluyendo visitas de supervisi ó n, evaluaci ó n reguladora, gesti ó n de suministro de medicamentos y resoluci ó n de problemas del sitio, para asegurar el cumplim.
-Rastrear puntos clave en la ejec. de ensayos; identificar problemas; resolver problemas y escalar como corresponda.
-Puede gestion. el reclutam. y ejec. planes de conting., como sea necesario.
-Completar la preparac./generaci ó n de informes de supervisi ó n de estudios.
-Puede dirigir y presidir reuniones locales del equipo del estudio, asistir y partic. en reunion. de equ. de ensayos cl í nic. glob.
-Notificaci ó n de quejas t é cnicas / eventos adversos / escenarios de casos especiales relacionados con los productos Novartis dentro de las 24 horas posteriores a la recepci ó n
-Distribuci ó n de muestras de comercializaci ó n (cuando proceda)

Key Performance Indicators

-Entregar resultados de satisfacci ó n del cliente para clientes internos y externos
-Entrega de ensayos cl í nicos a est á ndares de calidad y plazos acordados
-Cumplimiento de la pol í tica y directrices de Novartis y de las normativas externas.

Work Experience

Colaboraci ó n internacional
Amplitud funcional
Gesti ó n de crisis

Skills

Monitoreo Cl í nico
Investigaci ó n Cl í nica
Ensayos cl í nicos
Desarrollo de f á rmacos
Ciencias de la Salud
Protocolo de Ensayo Cl í nico
Ciencias biol ó gicas
Cumplimiento normativo
Habilidades para la toma de decisiones

Language

Ingl é s

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)

部门
Development

Business Unit
Development

地点
China

站点
Shanghai (Shanghai)

Company / Legal Entity
CN14 (FCRS = CN014) China Novartis Institutes for BioMedical Research Co., Ltd.

Functional Area
Research & Development

Job Type
Full time

Employment Type

Regular

Shift Work
No

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