

Evidence Generation Manager

Job ID
REQ-10041377
Febr. 25, 2025
Kolumbien

Zusammenfassung

Contribuye, con una supervisión adecuada, a todos los aspectos de los ensayos clínicos globales para ofrecer resultados de estudio dentro de los estándares de programación, presupuesto, calidad/cumplimiento y rendimiento. Puede liderar aspectos específicos del proceso de ensayo clínico global. Contribuye a la excelencia operativa a través de la mejora de procesos y el intercambio de conocimientos.

About the Role

Major accountabilities

Evidence Generation: - To support the coordination, development, and implementation of innovative integrated evidence plans. - To provide expertise on evidence generation methodology and innovative approaches. - To identify opportunities for evidence generation aligned with medical and brand priorities, through other cross-functional activities including digital engagement. - To anticipate and evaluate future needs and trends for RWE information within specified health care environments.

RWE Study Development, Execution and Oversight: - Gather and cross-functionally share relevant insights in the framework of the evidence integrated plans development (advisory boards, scientific trends etc), to leverage the local RWE opportunities. - Accountable for RWE integrated evidence plans for priority TAs, in close collaboration with Medical Leads, as well as Market Access, to meet increasing external RWE demands from payers, HCPs, policymakers and patients.

- Accountable for RWE projects' timely delivery and study management, including development of concept sheets for global approvals, protocols, external approvals, study report/publication and scientific disclosure. - Leading and/ or contributing with technical expertise and research methodological understanding for observational/epidemiologic research, ensuring projects are conducted in line with legal and compliance framework.

Strategic and Business Mindset: - Patient and customer-centric thinking in co-creating impactful integrated evidence plans with understanding of business and client needs, in line with the local CPO priorities. - Support target patient population outcomes as a metric of medical impact. - Broad understanding of across all disease areas and drug development.

- Building and managing strategic partnership with HCO/Third Party Organization to access external data source / analysis, ensuring information governance and data protection. - Maintain understanding and engagement with local health data and research landscape.

Leadership & Culture: - Ensure and drive cross-enterprise capability building in the area of expertise. - Lead the development of effective evidence generation team building through enterprise mindset & cross-functional

excellence. - Effectively operate in cross-functional teams, including early launch, BE&E (data & insights). - Role model for our culture, values & behaviors, consistently demonstrating the highest ethics and integrity-based standards. - Demonstrate enterprise perspective and delivery of medical evidence to address priority business challenges.

Other:

- Track deviations and support implementation/resolution of local CAPA plans.
- Follow up / support to audits and self assessments processes.

Ideal Background

Languages: English

Experience

Clinical Research Phases

Clinical Trial Design, Data Review & Reporting

Collaborating across boundaries

Critical Negotiations

Financial Management

Operations Management and Execution

Competencies

Breakthrough Analysis

Digital & Technology Savvy

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>

Commitment to Diversity and 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>

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Abteilung

International

Business Unit

Innovative Medicines

Ort

Kolumbien

Website

Bogota (Pharmaceuticals / GDD / NTO / CTS)

Company / Legal Entity
CO01 (FCRS = CO001) Novartis de Colombia S.A
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No
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