

Clinical Research Medical Advisor

Job ID REQ-10059570

8月 06, 2025

Canada

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:

- Provide clinical development and indication expertise specific to the country/cluster.
- Validate study designs, contribute to the start-up phase, provide robust training to clinical operations teams and support recruitment and trial awareness.
- Offer clinical/medical expertise for IRB/EC interactions and informed consent forms.
- Interact with external experts, regulatory authorities, and patient advocacy groups.
- Leverage innovation in clinical trial planning and recruitment strategies.
- Support internal stakeholders and decision boards regarding clinical trials.
- Review and resolve trial-related scientific/clinical/medical issues and support protocol deviations.

Essential Requirements:

- Scientific degree M.D., Ph.D., or Pharm.D. (M.D. highly desirable) Subspecialty training desirable
- 3 years of clinical development experience in the pharmaceutical industry or clinical practice.
- Ability to manage a study from the scientific/medical/clinical perspective, and a demonstrated capability to problem solve and mediate complex scientific/clinical/medical/operational issues.
- Ability to lead effectively by communicating well, motivating a cross-functional team, and handling and delegating responsibilities.
- Agility to move quickly across different therapeutic areas and indications.
- Demonstrated problem-solving skills and comfort with complexity.
- Ability to prepare and deliver high quality presentations.
- Speaks and writes English and French

Desirable Requirements:

• Sound understanding of the overall clinical development process, and ICH/GCP principles.

At Novartis Canada, we are determined to be a valued partner and advocate, with a deep understanding of patient needs along the entire care journey - from drug development, to diagnosis, to access and beyond. Part of the way we are doing this is by leveraging data, technology, and partnerships.

Research & Development: we focus on four core therapeutic areas: Cardiovascular, Renal & Metabolic, Immunology, Neuroscience and Oncology. We also develop and deliver treatments through other promoted and established brands, which today are helping millions of patients. Over the last three years, our average annual research and development investment in Canada was over \$30 million, and we conduct clinical trial research in every region throughout Canada.

Commitment to Diversity and Inclusion: Novartis is committed to building outstanding, inclusive work environment and diverse team 's representatives of the patients and communities we serve.

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部门 Development

Business Unit Innovative Medicines

地点 Canada

站点 Montreal

Company / Legal Entity CA04 (FCRS = CA004) NOVARTIS PHARMA CANADA INC.

Functional Area Research & Development

Job Type Full time

Employment Type

Regular

Shift Work No

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