Associate Clinical Development Director (Neurosciences)

Job ID
REQ-10055875

6月 27, 2025

United Kingdom

摘要

As our Clinical Development Director in Neurosciences you will be responsible for the scientific and clinical strategy of assigned clinical trials, scientific monitoring, and reporting of quality data. In addition, you may be responsible for the clinical and scientific strategy of assigned sections of a clinical development program, depending on the size and complexity.

About the Role

our responsibilities include, but are not limited to:

Provide clinical leadership, medical and scientific strategic input, and contribute to development of trial related documents (e.g., CTPs, informed consent form, case report forms, data monitoring committee charters, data analysis plan, reports, publications) for assigned clinical trial(s) consistent with Integrated Development Plan (IDP); develop materials for trial-related advisory boards, data monitoring committees, investigators meetings, and protocol training meetings for Novartis local

medical organizations.

- Provide clinical and scientific input and contribute to clinical sections of trial and program level regulatory documents (e.g., Investigator's Brochures, Health Authority briefing books, safety updates, submission dossiers, and responses to Health Authorities).

- In collaboration with appropriate Clinical Trial Team (CTT) members: Ensure clinical support of trials as needed; conduct ongoing medical and scientific review of clinical trial data with Clinical Scientific Expert(s) with appropriate oversight from Medical Lead; manage patient safety reports on trial data to safety and clinical boards (e.g., Safety Management Team (SMT), GCT, GPT) with appropriate oversight from Medical Lead; provide input into final analyses and interpretation including the development of the Clinical Study Report(s) (CSRs), publications and internal/external presentations.

- Support Therapeutic Area Head (TAH) with contributing to peer-review of IDPs, CTPs, and other clinical documents across various indications and programs, and support development of TA strategies, as needed.

- May contribute to the medical and scientific evaluation for Business Development & Licensing (BD&L) opportunities.

- Contribute to talent and career development of CD associates through on-boarding, coaching, and/or mentoring support; develop and foster CD culture; and may contribute to the performance evaluation of CTT members.

- Contribute to medical/scientific training of relevant Novartis stakeholders on the disease area and compound/molecule. May serve as speaker for franchise medical/scientific training.

- Contribute to global initiatives (e.g., process improvement, training, SOP development, other Clinical Development line function initiatives).

What you'll bring to the role:

- Advanced degree in life sciences/ healthcare (or clinically relevant degree) is required.

- 5+ years of involvement in clinical research or drug development in an academic or industry environment spanning clinical activities in Phases I through IV; 2 or more years of contribution to and accomplishment in all aspects of conducting clinical trials (e.g. planning, executing, reporting and publishing) in a global/matrix environment in pharmaceutical industry.

- Proven ability to interpret, discuss and present efficacy and safety data relating to clinical trial(s) or program level.

- Working knowledge of the disease area is desired with proven ability to interpret, discuss and present efficacy and safety data relating to clinical trial(s) or program level

- Working knowledge of GCP, clinical trial design, statistics, and regulatory and clinical development processes.

This hybrid role can be based in London, Dublin or Basel

Read our handbook to learn about all the ways we'll help you thrive personally and professionally: Novartis Life Handbook

Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.

Accessibility and accommodation
Novartis is committed to working with and providing reasonable accommodation to all individuals. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to receive more detailed information about the essential functions of a position, please send an e-mail to diversity.inclusionch@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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
London (The Westworks)

Company / Legal Entity
GB16 (FCRS = GB016) Novartis Pharmaceuticals UK Ltd.

Alternative Location 1
Basel (City), Switzerland

Alternative Location 2
Dublin (NOCC), Ireland

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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