

Associate Clinical Development Director - CV

Job ID
REQ-10055661

6月 25, 2025

Switzerland

摘要

The Clinical Development Director (CDD) in the Cardio Renal & Metabolic (CRM) Development Unit is responsible for leading the strategic planning and management of the assigned clinical program(s) from an end-to-end clinical development perspective. As a CDD in the CV TA, you will have oversight of the clinical development for the assigned programs and drive the execution of the clinical development plan. In addition, you will enable an empowered organization, which can navigate in a matrix environment and adjust quickly to business needs.

About the Role

Major accountabilities:

- Providing clinical leadership and strategic input for all clinical deliverables in the assigned project or section of a clinical program. Clinical deliverables may include clinical sections of individual protocols or sub studies consistent with the Integrated Development Plans (IDP), clinical data review, program

specific standards, clinical components of regulatory documents/registration dossiers, and publications

- Leading development of clinical sections of trial and program level regulatory documents
- Driving execution of the section of the clinical program in partnership with global line functions, assigned Global Trial Directors (GTDs), and regional/country medical associates, if applicable
- Overseeing/conducting ongoing medical and scientific review of clinical trial data with Clinical Scientific Expert(s) with appropriate oversight from Medical Lead
- Supporting (Sr.) GPCH in ensuring overall safety of the molecule for the assigned section and may be a core member of the Safety Management Team, supporting overall program safety reporting in collaboration with Patient Safety
- As a clinical expert, supporting the (Sr.) GPCH or CDH in interactions with external and internal stakeholders and decision boards
- Contributing to scientific training of relevant Novartis stakeholders on the disease area and compound/molecule. May serve as speaker for franchise medical/scientific training and may be the Program Manager of other associates

Minimum Requirements:

- Advanced degree in life sciences / healthcare (or clinically relevant degree) is required. PharmD or PhD is strongly preferred***
- Minimum of 7 years experience in clinical research or drug development
- Working knowledge of the assigned disease area is desired with proven ability to interpret, discuss and present efficacy and safety data relating to clinical trial(s) or program level
- Demonstrated ability to establish effective working relationship with key investigators
- Working knowledge of GCP, clinical trial design, statistics, and regulatory and clinical development processes
- Strong communication skills with the ability to work in a cross functional and global organization

Why Novartis: Our purpose is to reimagine medicine to improve and extend people ' s lives and our vision is to become the most valued and trusted medicines company in the world. How can we achieve this? With our people. It is our associates that drive us each day to reach our ambitions. Be a part of this mission and join us! Learn more here: <https://www.novartis.com/about/strategy/people-and-culture>

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部门
Development

Business Unit
Universal Hierarchy Node

地点
Switzerland

站点
Basel (City)

Company / Legal Entity
C028 (FCRS = CH028) Novartis Pharma AG

Alternative Location 1
Dublin (NOCC), Ireland

Alternative Location 2

London (The Westworks), United Kingdom

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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