

Clinical Development Director (CDD) - Cardio, Renal, Metabolism

Job ID
REQ-10004658
Mai 21, 2024
Etats-Unis

Résumé

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 Renal 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***
- Fluent oral and written English
- Minimum 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: 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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