

Clinical Development Medical Director

Job ID REQ-10050940

6月 14, 2025

USA

摘要

Onsite
#LI-Onsite
East Hanover, New Jersey

About the role:

As the Clinical Development Medical Director (CDMD) you will lead the planning and management of the assigned clinical program(s) from an end-to-end clinical development perspective and drive execution of the plan, enable an empowered organization, which can navigate in a matrix environment and adjust quickly to business needs. In Clinical Development Oncology, our aim is to design innovative, patient friendly clinical development plans to rapidly bring outstanding treatments to patients, caregivers and healthcare systems. We are striving to develop treatments for Lung, Breast & Prostate Cancers, MDS & AML, CML and sickle-cell disease, and are pushing the boundaries of innovation with CAR-T and Radioligand therapies.

About the Role

Your Key Responsibilities:

- Providing clinical leadership and strategic medical input for all clinical deliverables in the assigned project or section of a clinical program
- Lead development of clinical sections of trial and program level regulatory documents
- Drive execution of the program and/or clinical trial in partnership with global line functions, assigned Global Trial Directors, and regional/country medical associates
- Support the Global Program Clinical Head in ensuring overall safety of the molecule for the assigned section, and may act as a core member of the Safety Management Team, supporting overall program safety reporting in collaboration with Patient Safety colleagues
- Support the Clinical Development Head by providing medical input into the Clinical Development Plan, Integrated Development Plan and Clinical Trial Protocol reviews. and contributing to development of disease clinical standards for new disease areas
- As a medical specialist, supporting the GPCH or CDH in interactions with external and internal partners and decision boards
- May work with the Novartis Institute of Biomedical Research/ Translational Medical Sciences
 to drive transition of pre-PoC projects to DDP and with BD&L including target identification
 and due diligences together with other medical matters, as needed.

Video Link https://www.youtube.com/watch?v=ggbnzRY9z8w

The ideal location for this role is the East Hanover site but remote work may be possible (there may be some restrictions based on legal entity). Please note that this role would not provide relocation as a result. If associate is remote, all home office expenses and any travel/lodging to East Hanover for periodic live meetings will be at the employee's expense. The expectation of working hours and travel (domestic and/or international) will be defined by the hiring manager. This position will require 1% travel.

Role Requirements:

Essential Requirements:

- MD or equivalent medical degree is required, in addition to extensive knowledge and clinical training in medical/scientific areas.
- 3+ years minimum in clinical research or drug development.
- Working knowledge of Hematology/Oncology with a proven track record to interpret, discuss and present efficacy & safety data relating to clinical trials.
- Solid understanding of GCP, clinical trial design, statistics, and regulatory and clinical development processes.
- Demonstrated ability to establish effective scientific partnerships with key partners.

Desired Requirements:

- Previous global people management experience is preferred, though this may include management in a matrix environment.
- Clinical practice experience 4+years (including residency) preferred.

Novartis Compensation and Benefit Summary:

The salary for this position is expected to range between \$236,600 and \$439,400 per year. The final salary offered is determined based on factors like, but not limited to, relevant skills and experience, and upon joining Novartis will be reviewed periodically. Novartis may change the published salary range based on company and market factors. Your compensation will include a performance-based cash incentive and, depending on the level of the role, eligibility to be considered for annual equity awards.

US-based eligible employees will receive a comprehensive benefits package that includes health, life and disability benefits, a 401(k) with company contribution and match, and a variety of other benefits. In addition, employees are eligible for a generous time off package including vacation, personal days, holidays and other leaves.

To learn more about the culture, rewards and benefits we offer our people click here.

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

You'll receive:

You can find everything you need to know about our benefits and rewards in the Novartis Life Handbook.

https://www.novartis.com/careers/benefits-rewards

Accessibility and Reasonable Accommodations: The Novartis Group of Companies are committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the application process, or in order to perform the essential functions of a position, please send an e-mail to tas.nacomms@novartis.com call +1(877)395-2339 and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

Join our Novartis Network: If this role is not suitable to your experience or career goals but you wish to stay connected to learn more about Novartis and our career opportunities, join the Novartis Network here: https://talentnetwork.novartis.com/network

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

EEO Statement:

The Novartis Group of Companies are Equal Opportunity Employers. We do not discriminate in recruitment, hiring, training, promotion or other employment practices for reasons of race, color, religion, sex, national origin, age, sexual orientation, gender identity or expression, marital or veteran status, disability, or any other legally protected status.

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The Novartis Group of Companies are committed to working with and providing reasonable accommodation to individuals with disabilities. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the application process, or to perform the essential functions of a position, please send an e-mail to us.reasonableaccommodations@novartis.com or call +1(877)395-2339 and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

部门 Development

Business Unit Universal Hierarchy Node

地点 USA



站点

East Hanover

Company / Legal Entity U014 (FCRS = US014) Novartis Pharmaceuticals Corporation

Functional Area Research & Development

Job Type Full time

Employment Type Regular

Shift Work No

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