Associate Clinical Development Medical Director

Job ID
REQ-10057431

7月 08, 2025

USA

摘要

The Associate Clinical Development Medical Director (Assoc. CDMD) is responsible for the scientific and medical strategy of assigned clinical trial(s), medical and scientific monitoring, and reporting of quality data. May be responsible for the scientific and medical strategy of assigned sections of a clinical development program, depending on the size and complexity.

About the Role

MAJOR ACCOUNTABILITIES:

- Is a global clinical manager or country / cluster leader responsible for clinical program(s) across indications executing medical strategy for development and marketed products in a defined therapeutic area -Responsible for the scientific and medical strategy of assigned clinical trial(s), medical and scientific monitoring.
- May be responsible for the scientific and medical strategy of assigned sections of a clinical

development program.
- Contributes to the development of clinical sections of trial and program level regulatory documents -Contributes to the execution of the section of the clinical program in partnership with global line functions.
- Contributes to ensuring overall safety of the compound for assigned trial(s) in collaboration with Patient Safety.
- Supports by contributing medical input into IDP and CTP reviews and contributing/driving development of disease clinical standards for new disease areas -Contributes to medical/scientific training of relevant Novartis stakeholders.
- May serve as speaker for franchise medical/scientific training -Contributes to the global initiatives (e.g., process improvement, training, SOP development, other Clinical Development line function initiatives) -Contributes to talent and career development of CD associates through on-boarding, coaching, and/or mentoring support; -Reporting of technical complaints / adverse events / special case scenarios related to Novartis products within 24 hours of receipt -Distribution of marketing samples (where applicable)

REQUIREMENTS

- MD or equivalent medical degree required. Advanced knowledge and clinical training in a medical/scientific area (e.g., internal medicine or sub-specialty) required, with Medical Board certification preferred; Clinical practice experience $\geq$ 4 years (including residency) preferred
- $\geq$ 3 years of involvement in clinical research or drug development in an academic or industry environment spanning clinical activities in Phases I through IV. $\geq$ 2 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
- 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, written and oral
- Strong interpersonal skills
- Strong negotiation and conflict resolution skills
- Proven ability to work independently or in a cross-functional team setting

The salary for this position is expected to range between $204,600 and $379,400 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.

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.

Accessibility & 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 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

状态
Massachusetts

站点
Cambridge (USA)

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

Alternative Location 1
East Hanover, New Jersey, USA

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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REQ-10057431

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