

Translational Medicine Director/Associate*- Cardiovascular & Metabolic

Job ID

REQ-10087155

Sep 03, 2026

LOC_CH

About the Role

LI- hybrid

TME Director Cardiovascular Metabolism

Oversees the planning, execution, and interpretation of clinical trials research, data collection activities and clinical operations, serving a leadership role in the matrixed trial team. Establishes and approves scientific methods for design and implementation of clinical protocols, data collection systems and final reports. Support new and ongoing clinical research and clinical trials and ensure efficient and timely processing of confidentiality agreements and clinical agreements

Key Responsibilities:

- Lead global project teams through PoC phase to drive implementation of the PoC strategy; participate in project teams through program life cycle from NTRC to registration
- Responsible for clinical portions of the Integrated Development Plan (IDP) through PoC.
- Acts as medical monitor for one or more clinical studies.
- Communicate clinical team matters to project teams and relevant decision boards (such as the Disease Area Decision Board).
- Convene relevant (internal and external) leaders to consider proposed approach to PoC.
- Evaluate clinical centers and foster communication with crucial collaborating investigators.
- Oversee publication and external presentation of PoC clinical study results.
- Accountable for compound related biomarker strategies; work closely with biomarker experts in implementation.
- In collaboration with research scientists identify, develop and implement strategy for preclinical support of program related objectives; this includes assessment of medical need, proposal of clinical development pathways, and review of preclinical data for clinical implications and other relevant activities
- Participate in team presentations to Health Authorities as TM Expert

Essential Requirements:

- Doctoral degree, MD required with clinical subspecialty training preferred in cardiovascular disease or endocrinology/metabolic disease; Additional PhD/post-doctoral equivalent research preferred.
- At least 1-2 years' experience in a pharmaceutical/biotech company, CRO, or academic medical center, or related experience; 5 plus years to be considered for Director level.
- Recognized for medical expertise.
- Recognized for scientific expertise and passion for science: respected by colleagues internally and externally, having made significant contributions to the field and created / established new concepts; record of high quality publications in international scientific journals.
- Excellent written and oral communication/presentation skills.
- Independence: Able to work independently on study planning and execution, and with supervision on project planning.
- Leadership of Collaborations: Able to lead multidisciplinary teams in a matrix organization.
- Innovation: Seeks out new clinical discovery opportunities and PoC approaches.
- Additional consideration for candidates with advanced research experience in data science, genetics/genomics, artificial intelligence applied to clinical research, or digital/wearable health science.
- Fluent oral and written English

****This is a dual level posting. The final level & title of the offer role would be determined by the hiring team based on the skills, experience & capabilities required to perform the role at the level the role has been offered.**

The base salary offered is determined based on gender-neutral objectives, such as relevant skills, competencies and experience in accordance with the Novartis pay setting policy and upon joining Novartis will be reviewed periodically.

Associate Director: 122,500.00 - 175,000.00 - 227,500.00 CHF Annual

Director: 150,500.00 - 215,000.00 - 279,500.00 CHF Annual

In addition to your base salary, you may be eligible for a performance-based bonus depending on certain performance parameters.

The rewards of being part of our team go far beyond base pay and incentives. We also offer a variety of competitive benefits in kind to help you thrive personally and professionally, such as insurance plans, retirement plans, wellbeing resources and global recognition programs. In addition, we provide flexible and hybrid working options, where possible, and minimum 14 weeks paid parental leave.

Pay equity is a fundamental principle of our employment policy and reflects our commitment to create a diverse, equitable and inclusive environment that treats all employees with dignity and respect, as outlined in our Code of Ethics.

Read our brochure to learn more about our global total rewards offering:

https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf

Note: Benefits and compensation may vary by country and are subject to local legal requirements, including provisions of collective bargaining agreements where applicable. A full overview of your compensation package, including any relevant collective bargaining agreement details applicable to your role based on your employment location and Novartis employer entity, will be communicated separately to you during the application process.

Role Requirements

Primary location salary range

CHF150,500.00 - CHF279,500.00

Division

DIV_RE

Business Unit

Research

Location

LOC_CH

Site

Basel (City)

Company / Legal Entity

C028 (FCRS = CH028) Novartis Pharma AG

Functional Area

FCT_RD

Job Type

Full time

Employment Type

Regular

Shift Work

No

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