

TA Lead

Job ID
REQ-10060086

9月 16, 2025

China

摘要

-Designs and provides oversight of clinical research programs. Builds relationships with key opinion leaders and applies their input to enhance study design and protocols. Serves as medical/scientific consultant to marketing or research project teams and government regulatory agencies. Establishes the criterion essential for determining the safety, efficacy, and medical utilities. Interprets results of Phase I-III investigations in preparation for new-drug or medical device application. May serve as safety expert for individual clinical projects. May be responsible for post marketing studies

About the Role

Major accountabilities:

- Is a global clinical leader responsible for clinical program(s) across indications or a large regional clinical leader driving medical strategy for development and marketed products in a defined therapeutic area -Owns the risk benefit assesment for the program(s) -Is accountable

for the design, implementation, and execution of a clinical development program(s) to support decision milestones, regulatory requirements and market access -Leads execution on broad strategic direction and contributes to development of disease/therapeutic area strategy -Leads the creation of clinical components of key documents e.g., Clinical Trial Protocols (CTPs), Investigator ' s Brouchures, Clinical Study Reports (CSRs), regulatory documents etc. with high levels of quality and consistency -Acts as the medical expert, leads interactions with external stakeholders (e.g., regulatory authorities, key opinion leaders, data monitoring committees, advisory boards etc.) and internal NVS stakeholders -Collaborates across functions to ensures continuous evaluation of drug safety profile -Drives scenario development for Clinical Development to support decision analysis and optimal resource allocation in program -Ensures career development of program reports through active participation in the performance management -Distribution of marketing samples (where applicable)

Key performance indicators:

- Achievement of unit objectives -Delivery of Clinical Trials to quality standards and agreed timelines -Adherence to Novartis policy and guidelines and external regulations.

Minimum Requirements:

Work Experience:

- Strategy Development.
- Functional Breadth.
- People Challenges.
- People Leadership.
- Representing the organization.

Skills:

- Clinical Decision Making.
- Clinical Research.
- Clinical Trials.
- Disease Area Knowledge.
- Drug Development.
- Leadership.
- People Management.
- Risk Management.
- Strategy Development.

Languages :

- English.

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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部门
Development

Business Unit
Innovative Medicines

地点
China

站点
Shanghai (Shanghai)

Company / Legal Entity
CN06 (FCRS = CN006) Beijing Novartis Pharma Co., Ltd

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work

No

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Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.



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