

Director Group lead Immunology, PK Sciences (80-100%)

Job ID
REQ-10063995

10月 20, 2025

Switzerland

摘要

Welcome to where we thrive together! Are you ready to join a community where you can make a real impact on the world by supporting the global pipeline of the Immunology TA across all phases of development? At Novartis, we believe in creating a positive and inclusive work environment where we can solve the toughest healthcare challenges together.

About the Role

Location: Basel, Switzerland

This role is based in Basel, Switzerland. Novartis is unable to offer relocation support for this role: please only apply if this location is accessible for you.

Key responsibilities:

- Accountable for development and implementation of the PKS strategy for a portfolio of program(s) and coordinates pharmacokinetic / drug metabolism/ modelling and simulation related elements for pre-clinical and/or clinical projects. Identifies potential project hurdles, contributes to dose selections and provides solutions and establishes contingency plans. Represents PKS on IMM project teams(s).
- Responsible for supervising, training, and mentoring PKS PTM, and to ensure line function excellence with respect to efficiency, and scientific / technical project support.
- Participates and contributes internal strategy Boards and review committees dealing with projects assigned to direct reports
- Responsible for resource planning, management, and administration (FTEs and direct costs)
- Reviews study protocols, internal strategy documents and HA related documents for projects assigned to direct reports
- Proactively contributes expert pharmacokinetic /drug metabolism, pre-clinical and clinical pharmacology input into key pre-clinical and clinical, and regulatory documents including clinical study protocols, clinical study reports, investigator brochures, IND / IMPDs and NDA/BLAs within agreed timelines and addresses all regulatory requirements with minimal supervision. Provides input to key NIBR, GDD, milestones (DC, IDPA, ISA, TDP etc).
- Responsible for the PK, PK/PD, IG and M&S component of study protocols, reports, project summaries and development plans within agreed timelines and which meet regulatory requirements. Prepare appropriate responses to Health Authority questions (globally).
- Oversees or performs PK and PK/PD analyses using a variety of tools and approaches. Integrates, interprets and reports data to project teams and other customers. Support project teams with modeling- and simulation-based input to clinical trial design, evaluations, integrated analysis, and reporting.
- Monitors study timelines, objectives and budgets, ensure accuracy of project progress in company tracking tools. Assures rapid and effective communication of data to project teams.
- Leads PKS sub-teams. Coordinates with appropriate sub-team members and ensures implantation of project strategy and/or prepares summary documentation.

Essential Requirements:

- PhD / Pharm.D. level scientist with relevant experience in drug metabolism and pharmacokinetics or related biologic background.
- Relevant TA/DA experience and demonstrated track record on coaching and mentoring of associates would be preferred
- Minimum 8-10 years in drug development functions (e.g. DMPK, Clinical Pharmacology) with a minimum of 5 years ' experience within a relevant LF in industry, academia or regulatory agency
- Extensive and in-depth knowledge of drug metabolism and pharmacokinetics including pharmacology PK and PK/PD evaluation, immunogenicity assessments and regulatory requirements.
- Demonstrated success leading and influencing in a multi-disciplinary environment including global project teams. Aware of the most recent developments in drug development sciences and the implications for current projects.
- Strong coaching, mentoring and people management skills

Desirable Requirements:

- Proven record as team player/leader with excellent influencing & negotiation skills.
- Excellent organizational and project skills

Commitment to Diversity and Inclusion / EEO paragraph:

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

Accessibility and Accommodation:

Novartis is committed to working with and providing reasonable accommodation to all individuals. If, because of a medical condition or disability, you need a reasonable accommodation for any part of the recruitment process, or in order to receive more detailed information about the essential functions of a position, please send an e-mail to inclusion.switzerland@novartis.com and let us know the nature of your request and your contact information. Please include the job requisition number in your message.

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

Biomedical Research

Business Unit

Pharma Research

地点

Switzerland

站点

Basel (City)

Company / Legal Entity

C028 (FCRS = CH028) Novartis Pharma AG

Functional Area

Research & Development

Job Type

Full time

Employment Type

Regular

Shift Work

No

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