

Senior Clinical Supply Project Lead

Job ID
REQ-10054079

6月 10, 2025

Austria

摘要

The GCS Project Lead (PL) leads, represents, manages and supports GCS project team and operates as single point of contact for clinical and technical teams across GDD on clinical supply strategy. The PL ensures complete project oversight in GCS and retains accountability for project deliverables.

The CSPL has operational end to end responsibility for assigned activity. Leads and manages all project and local network activities and participates in cross-functional teams.

About the Role

Key Responsibilities:

- Represent GCS/Technical Research & Development (TRD) in the Global Clinical Trial Team (GCT) and act as business partner to support and influence the decision-making process on the supply strategy and packaging design to be adopted in clinical trials. Attends TRD sub-

team meeting providing insights and guidance from clinical studies perspective.

- Assess clinical development plan scenarios, converts them in demand forecast to support long term supply and capacity planning for TRD/Novartis Operations (NO). As business partner to the clinical teams, drive GCS assessment to support unplanned clinical study requirements.
- Leads overall clinical supply strategy in alignment with clinical and technical requirements and constraints. Assess risks & opportunities and define strategies to ensure supply continuity and increase supply flexibility and responsiveness in case of new clinical study initiatives.
- Oversees the entire End to End (E2E) supply strategy (from Drug Substance to clinical sites) and reviews the planning assumption adopted in the supply plans and forecasts. Participates in the decision-making process in GCS to select the most appropriate supply model.
- Operates as first level of escalation and provide clear overview on issue, supply impact and mitigation plan to GCS management in case of supply risk / issue. Leads communication and manages stakeholder 's expectations in case of critical issue.
- Understands and proactively manages the interactions of project, network and/or platform related activities within and outside of GCS. Acts as ambassador for GCS in TRD and clinical environment.
- Ensures overall budget adherence of the financial resources allocated to the project in GCS. Acts as point of contact for GCS Finance department, manages the budget allocated to the project and discuss variation that could require additional financial resources. Leads the cost assessment of packaging, distribution, booklet, and comparator activities in case of new clinical study initiatives.
- Inspire/coach/lead team members. Provide coaching and technical training as subject matter expert or recognized technical expert. Act as a mentor for junior and senior associates globally.
- Proactively communicate key issues and any critical topic in a timely manner to the appropriate management level and to/or any other relevant project team members. Consolidate data evaluation, propose solutions, and contribute to risk mitigation plans.
- In close cooperation with the Unit Head, drive the unit long term strategic plan and its implementation. Ensure current and future needs are fully met, unit projects are assigned, adequately resources, delivered on time and in full compliance.

Essential Requirements:

- Advanced Degree in science, engineering, or relevant discipline (Ph.D., MBA or equivalent)
- >8 years of practical experience in chemical / pharmaceutical industry or > 4 years of experience in field of expertise
- Thorough knowledge of Drug Development processes and Clinical Supply processes
- Comprehensive knowledge about project management, excellent organization, and planning skills
- Strong knowledge of relevant regulations (e.g., Good Manufacturing Practice (GMP), Health, Safety & Environment (HSE) etc.).
- Demonstrates cross-functional problem-solving and idea generation skills
- Strong communication, presentation, and advanced coaching skills.
- Excellent negotiation, proven leadership, interpersonal skills and Ability to work in interdisciplinary teams.

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In addition to a market-competitive base salary, we offer an attractive incentive program, a modern company pension scheme, childcare facilities, learning and development opportunities as well as worldwide career possibilities within the Novartis group. Level 5: In accordance with Austrian law, we are obliged to disclose the minimum salary as stated in the collective bargaining agreement. For this position the minimum salary is €77 543,90 /year (on a full-time basis). The actual salary will be significantly higher, as we strive to maintain a competitive position in the market and consider your previous experience, qualifications and individual competencies.

If because of a medical condition, physical disability or a neurodiverse condition you require an adjustment during the recruitment process, please reach out to disabilities.austria@novartis.com and let us know the nature of your request as well as your contact information. The support which we can provide will include advice on suitable positions as well as guidance at all stages of the application process. Austrian law provides candidates the opportunity to involve the local disability representative, Behindertenvertrauensperson (BVP), in the application process. If you would like to request this, please let us know in advance as a note on your CV.

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部门
Development

Business Unit
Sandoz

地点
Austria

站点
Schaftenau

Company / Legal Entity
AT33 (FCRS = AT033) Novartis Pharmaceutical Manufacturing GmbH

Functional Area
Research & Development

Job Type
Full time

Employment Type
Regular

Shift Work
No

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Adjustments for Applicants with Disabilities

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