

Supply Chain Manager - Global Clinical Supply

Job ID
REQ-10075881
апр 16, 2026
Австрия

Сводка

Location: Schafteuau, Austria #onsite

Role Purpose:

The Supply Chain Manager (SCM) is responsible for Demand and Supply Planning from Clinical Finished Good (CFG) to Drug Substance (DS) and Booklet labels, ensuring demand fulfillment for assigned projects. The SCM acts as key contributor to the Clinical Supply & Operations Planning (CS&OP) process in TRD/GCS and provides transparency on supply constraints and manages related aspects accordingly within TRD.

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

Major accountabilities:

- Harmonizes the supply strategy within GCS and contributes to the supply strategy of CHAD/PHAD/Biologics.
- Participates in the GPMM along with the CSPL and CTSM ensuring alignment between demand and supply.
- Ensures demand fulfillment and coverage of supply and regulatory aspects by contributing to GCS agenda at TRD Sub team CMC meeting. Represent GCS at TRD Sub-team on supply chain aspects.
- Actively contributes to the portfolio manufacturing schedule alignment (from DS to CFG) Defines most cost-efficient ordering levels from CFG to DS, minimizing waste and allowing flexibility to accommodate demand variability.
- Drives Long term demand and capacity planning (LTDCP) coordinating with the CSPL, DPPL, DSPL and TPL. Adheres to SCM KPIs including the one being part of the SPE for project and unit.
- Proactively manages and adheres to functional performance indicators with a focus on supply planning excellence.
- Data and Digital savviness in SC domain. Manages Ordering and master data requirements in SAP within the scope of the role.
- Adapt and implement **Rapid Response (Maestro)** for portfolio supply& demand planning, network design and scenario building.
- TRAFFIC – Establish the Supply chain design in alliance with Funds Flow, Customs & Trade Compliance and TRD sub-team for portfolio.
- Drive the Change control strategy for clinical supplies from GCS perspective.
- Provides impact assessment on clinical supplies and contribute to the regulatory submission strategy.
- Integrates Comparator supply strategy into the TRD procurement, blinding & release planning.

Minimum Requirements:

Work Experience:

- Degree in science, engineering or equivalent.
- Fluent English
- >5 years of practical experience in chemical / pharmaceutical industry or > 3 years of experience in field of expertise
- Good expertise in related field.
- Good knowledge about the Drug Development process
- Basic project management, good organization and planning skills
- Knowledge of relevant regulations (e.g., GMP, HSE etc.) and Novartis specific standards.
- Demonstrates problem-solving and idea generation skills.
- Good presentation skills
- Intermediate Leadership skills
- Very good communication, negotiation and interpersonal skills. Ability to work in interdisciplinary teams.

Austria Adjustments for Applicants with Disabilities: 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.

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 €65,504.54/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

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>

Benefits and Rewards: Learn about all the ways we'll help you thrive personally and professionally. [Read our handbook \(PDF 30 MB\)](#)

Дивизион
Development
Business Unit

Development
Место
Австрия
Сайт
Schafftenau
Company / Legal Entity
AT33 (FCRS = AT033) Novartis Pharmaceutical Manufacturing GmbH
Functional Area
Research & Development
Job Type
Full time
Employment Type
Regular
Shift Work
No

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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