NOVARTIS

Supply Chain Manager - Global Clinical Supplies

Job ID
REQ-10077014
May 07, 2026
LOC_CH

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.

Commitment to Diversity and Inclusion: 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.

Role Requirements

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)

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Development
Location
LOC_CH
Site
Basel (City)
Company / Legal Entity
C028 (FCRS = CH028) Novartis Pharma AG
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Employment Type
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Shift Work
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