

# Supply Chain Manager - Global Clinical Supplies

Job ID  
REQ-10077014  
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Швейцария

## Сводка

Location: Basel, Switzerland #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.

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](mailto: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.

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C028 (FCRS = CH028) Novartis Pharma AG  
Functional Area  
Research & Development  
Job Type  
Full time  
Employment Type  
Regular  
Shift Work  
No

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