NOVARTIS

Supply Chain Manager

Job ID
REQ-10031133
Dec 03, 2024
Austria

Summary

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

About the Role

Key Responsibilities:

- Creates and maintains the end-to-end supply plan from CFG to DS
- Harmonizes the supply strategy within GCS and contributes to the supply strategy of Chemical and Analytical Development/ Pharmaceutical Development (CHAD/PHAD)/Biologics
- Leads the Clinical Demand Planning Meeting (CDPM) ensuring alignment between demand and supply
- Ensures demand fulfilment and coverage of supply and regulatory aspects by contributing to GCS agenda at TRD Sub team Chemistry/Manufacturing/Controls (CMC) meeting. Represent GCS at TRD Sub-team on supply chain aspects.
- Optimizes the inventory strategy at PP and CFG level together with CTSM
- 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 Clinical Supply Project Lead (CSPL), Drug Product Project Leader (DPPL), Drug Substance Project Lead (DSPL) and Technical Project Leader (TPL).
- Adheres to SCM KPI for project and unit
- Data and Digital savviness in SC domain. Manages Ordering and master data requirements in SAP within the scope of the role
- 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

Essential Requirements:

- Degree in science, engineering or equivalent.
- >5 years of practical experience in chemical / pharmaceutical industry or > 3 years of experience in field of expertise
- Experience in SAP

- Experience in Kinaxis RapidResponse
- 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 and fundamental Leadership skills.
- Very good communication, negotiation, and interpersonal skills. Ability to work in interdisciplinary teams.

Desirable Requirements:

- Experience of running projects independently

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Benefits and rewards: Read our handbook to learn about all the ways we'll help you thrive personally and professionally: https://www.novartis.com/careers/benefits-rewards

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 4: 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 €60,212.18/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.

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.

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Division
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Business Unit
Sandoz
Location
Austria
Site
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
Apply to Job

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