Head of Manufacturing and Supply Chain for Lymphatica Medtech

Lymphatica Medtech SA

Lymphatica Medtech is a pioneer in implantable systems for lymphedema treatment. Spin out of EPFL and CHUV in 2017, the growing company has received national and international recognitions for its breakthrough approach.

We work in a competent and motivated team, implementing an ambitious project to have a significant impact for lymphedema patients and society as a whole.

Job Summary

The primary purpose of the Head of Manufacturing and Supply Chain is to lead efforts in consolidating the supply and manufacturing chain for the production of a Class III implantable medical device, in relation to both clinical trial and market entry phase.

The Head of Manufacturing and Supply Chain will be part of the Management Team, and will act as a strategic thought partner to the CEO.

The job will be based in Milano (Italy), and it will combine the need for frequent travelling to suppliers and production partners, with the possibility of flexible working hours and remote working.

Responsibilities

- Consolidating the company's supply chain and manufacturing strategy
- Managing suppliers and contract negotiations
- Managing and conducting supplier onsite visits and regulatory / quality audits
- Supervising design transfer activities
- Developing and executing strategy to improve product margin and lifetime customer value e.g. through design input and manufacturing and supply chain efficiencies
- Managing inventory of raw goods and finished materials
- Collaborating with Regulatory and Quality Affairs to ensure compliance of manufacturing operations with regulatory requirements for medical devices
- Reviewing, updating and maintaining Standard Operating Procedures (SOP) and Processes (PCS) where related to production, incoming inspection, supplier handling and storage/inventory.

- Acting as a thought partner with the broader team including Quality and Regulatory Affairs, R&D and Management teams.

Qualifications & Skills

- Degree in mechanical engineering or equivalent
- Proven working experience in leading engineering positions and or quality/regulatory positions for medical device manufacturing.
- Certificate of internal/external auditor training according to ISO 13485 or ISO 9001 is a plus.
- Proven experience in supplier management and contract negotiations.
- Result-driven and persistent with a desire to overcome challenges with a positive mindset.
- Ability to adapt and learn quickly in a changing business environment.
- Excellent relationship management skills, with the ability to work collaboratively with internal and external teams.
- Adept at giving and receiving feedback and mentoring team members.
- Proficient written and verbal communication in English and Italian. French is a plus.

How to Apply

Please submit your CV and cover letter to contact@lymphatica.ch. We look forward to reviewing your application!