

## Production Manager

Contract: CDI, permanent

TISSIUM is a fast-paced medical device company, dedicated to the development of innovative tissue reconstruction solutions. We are leveraging our technology platforms to develop novel solutions to disrupt the field of surgery and positively impact the life of patients.

We are actively looking to recruit a **Production Manager** to be part of TISSIUM's Development Factory. This person will lead and manage all stages of the production activities, including management of the team, production time tracking, process improvement.

### Duties & Responsibilities

- ✓ Define, implement, and monitor all the aseptic processes and dedicated quality control aspects of the company's polymer production according to GMP, ISO 13485 and ISO 13408 requirements;
- ✓ Establish and monitor budgets and timelines relating to all bulk polymer processing activities;
- ✓ Support the development and implementation of the company's bulk polymer processing strategy especially all related aseptic processes;
- ✓ Lead and support the manufacturing processes, including scale up, process validation, process improvement either internally or in collaboration with the company's contract manufacturers;
- ✓ Proactively advise and support the product development team at all stages of the product development cycle to yield highly manufacturable products and minimize manufacturability, sourcing, and scalability risks;
- ✓ Coordinate supply activities in collaboration with the Supply Manager and ensure adequate processed polymer supply to all internal and external customers;
- ✓ Manage the production team;  
Implement, improve and maintain a lean manufacturing mindset (production timelines, 5S, performance management tools, KPI's implementation).

### Job Environment/Interactions

- ✓ Assist in the conduction of all validation activities and in the preparation of the Design Dossier documentation by working closely with the company's product development team;
- ✓ Support the QA department in the conduction and management of the company's CAPA activities.

### Qualifications & Skills

- ✓ Engineering degree, pharmacist or equivalent;
- ✓ A minimum of 5 to 10 years of experience in manufacturing of medical products (pharmaceuticals or medical devices);
- ✓ Strong experience in aseptic production and associated processes;
- ✓ Strong experience in production's activities management;
- ✓ Excellent understanding of quality requirements for the production of medical devices or pharmaceutical products and specifically the requirements of ISO 13485, ISO 13408 and GMP standards and practices;

# TISSIUM

- ✓ Strong know-how in manufacturing process design, technology transfer, scale-up operations, analytical development and process validation;
- ✓ Good communication skills (written and verbal) in English;

## **Job Competencies & Personal Attributes**

- ✓ Strong leadership skills and attention to details;
- ✓ Strong ability in driving third parties to adhere to timelines and budgets;
- ✓ Strong planning, problem solving and negotiation skills;
- ✓ Excellent team work abilities;
- ✓ Capacity to understand complicated processes and associated contract manufacturers' interactions;
- ✓ Creativity in approaching and solving technical challenges;
- ✓ Autonomy and good organization skills;

The Production Manager will report to the Site Director. The position is based in Roncq (59), France. To apply to this position please email your CV to [jobs@tissium.com](mailto:jobs@tissium.com)