KiOmed Pharma recruits :

"Product Development Manager – Aesthetic"

Innovative Medical Devices based on KiOmedine®,
an exclusive highly pure non-animal polymer

Possibilité d'embauche CDI / Herstal (Belgium).

Capitalizing on a history of innovation and expertise in exclusive chitosan chemistry, KiOmed Pharma, a spinout from KitoZyme, develops a unique pipeline of medical devices that address unmet medical needs in high impact pathologies and major social burdens such as invalidating osteoarthritis, aesthetic medicine and ophthalmology.

Located in Belgium, Liège and composed of a fast-growing team, KiOmed Pharma has launched on the market its first class III CE marked medical device, KioMedine^{VS}one® indicated in the treatment of symptomatic patients suffering from knee osteoarthritis. A R&D pre-market pipeline of medical devices is currently under development, more specifically in dermo-aesthetic medicine and ophthalmologic diseases.

KiOmed Pharma has the ambition and the resources to grow to a worldwide class and aim at succeeding the commercial launches and growth on its markets.

Responsabilities:

- Develops project plans and establishes a schedule of project tasks and milestones including the registration steps according to MDR 2017/745.
- Plans, organizes and coordinates the execution of tasks with the different stakeholders and partners involved in the project,
- Identifies, qualifies and negotiates with external partners and subcontractors and ensures the implementation of appropriate contractual framework with external partners.
- Manages project execution with a focus on meeting milestones, budget and deadlines. Provides critical analyses of results and their interpretation.
- Reports to the manager any element of non-compliance or unplanned event that may potentially or actually harm the design and/or business activities.
- Proposes and implements fallback plans to manage unexpected events.
- Effectively and periodically monitors project progress and budget and prepare project status
- Ensures compliance with safety instructions and the maintenance and improvement of the quality system in force for design activities.

- Contributes to the preparation of the technical dossier for submission and its follow-up after CE marking
- Archives the documentation linked to the technical dossier
- Contributes to scientific monitoring and the securing of intellectual property as well as to the company's inventive step

Profile:

- Scientific background, PharmD, Scientist (MS or PhD).
- Experience of at least 5 years in the development of implantable medical devices
- In-depth knowledge of European regulations and of design of medical devices.
- Effective project management and leadership skill.
- Strong analytical and problem solving skill. Objective oriented
- Ability to work in rapidly changing environment
- Excellent communication skills in English speaking environment
- Ability to write document
- Strong interest in medical technologies.

Offer:

A position within a fast-growing and innovative Belgian biotech company with a proprietary and unique technology platform targeting unmet medical needs in multibillion dollar markets.

An attractive package in line with the responsibilities and your experience.

If you are interested in this position, please send, in either French or English, your CV and details of current remuneration package to Catherine.Philippart@kiomedpharma.com