Telix Pharmaceuticals Limited
ACN 616 620 369
Suite 401, 55 Flemington Road
North Melbourne
Victoria, 3051
Australia

Job Description: Process Engineer
Type: Full Time
Location: EMEA (Location – Liège)
Immediate

About Telix Pharmaceuticals Limited

Telix Pharmaceuticals Limited ("Telix", the "Company") is an Australian public Company (ASX:TLX) headquartered in Melbourne with operations in Europe, the US and Japan. Our mission is to be a leading, global biopharmaceutical Company in the field of "theranostic" radiopharmaceuticals and the Company is currently developing a mid-late stage pipeline of products in oncology and rare diseases.

About the Role

You contribute to the achievement of the Group's strategic goals by the implementation, ***improvement and maintenance*** of a robust supply chain flow and operational structure for the delivery of Telix products. You manage the ***introduction of new products*** in operations on site. The Process Engineer will collaborate internally with other departments and externally with his counterparts in other Telix juridisctions to ***ensure that the Supply Chain delivers products*** across all Telix regions with the same standards of quality. The role is mainly located in Liège, Belgium.

Your ***key responsabilities*** will be as follows:

- Responsible for ***writing and maintaining Standard Operating Procedures***, work instructions, Master Batch Records and protocols for the Supply Chain department
- ***Management of Supply Chain related deviation***, Change control, Corrective and Preventive Actions (CAPA), Supplier/subcontractor Change Requests, products and processes validation
- Responsible for ***installation, qualification and maintenance*** of equipment on Liège's site.
- Responsible for the *implementation of new products* on site, ***from Design output to product realization***.
- Management of ***GMP infrastructure***

About You

You have at least a ***minimum of 5 to 10 years experience in a Supply Chain environment***, preferably in a pharmaceutical/biotechnology level. In addition, you are manufacturing oriented and have perfect ***technical writer skills***. Furthermore, you have a proven experience in pharmaceutical drug supply environment with ***strong GMP/GDP background***. Moreover, you have experience working with an ERP system (preferably ***SAP B1***) and with a ***QMS*** (preferably Master Control).

Specific skills required

- Commitment to the vision and mission of Telix
- Ability to ***work under pressure***
- Ability to ***prioritise*** competive priorities/ objectives
- Ability and willingness to ***work collaboratively*** with local and global team members across multiple timezones
- Demonstrated ability to function well in a collaborative team environment
- Ability to ***plan, create, strategize and drive execution*** of multiple projects under tight timelines,
- Ability to ***prioritize and manage time*** effectively
- Strong ***organizational and interpersonal skills***
- Demonstrated proficiency in ***Microsoft Office***
- Excellent oral and written ***communication skills***
- Strong ethics in ***decision making***
- A conscientious and rigourous attitude
- Perfectly ***Bilingual English-French***

Why work at Telix?

We are a dynamic, fast-growing biopharmaceutical company where employees have a shared purpose: to help people with cancer and rare diseases live longer, better quality lives. This is an exciting time for Telix and we are looking for like-minded, passionate professionals to join us on the journey.

No agency submissions will be considered for this role.