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

TLX-JD
Job Description : Director - Clinical Operations
Type : Full-time
Location : Melbourne, Australia
Date : 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-to-late stage pipeline of products in prostate, kidney and brain cancer.

Description

The Director of Clinical Operations is a key member of the Company's senior leadership team and as such is involved in setting the Company's long-term vision and short-term goals and objectives.

The position will be responsible for the leadership and oversight of clinical study operations by organising and coordinating the planning, implementation, management, execution and completion of clinical programs according to applicable regulations and guidance, laws and Standard Operating Procedures (SOPs). This position will also be responsible for the development of clinical operations systems/processes and effective communication within and outside the company related to the implementation of processes and successful completion of studies.

This position reports to the Chief Medical Officer (CMO).

Responsibilities

- Setting clinical and strategic goals in partnership with the CMO.
- Oversight responsibility for clinical operations, CRO engagement and responsible for representing the clinical operations activities of the company to external stakeholders including clinicians, collaborators and regulatory authorities.
- Reporting of clinical trial progress as part of board and management team reporting functions.
- Business partnership functions appropriate for a senior clinical role, including competitive analysis, market / product intelligence, support in partnering and collaboration discussions and treatment landscape analysis.
- Participation in the review and business case development for new pipeline opportunities, production indication expansion, etc.
- As a member of the executive team, shared responsibility for the performance of management in meeting agreed goals and objectives and ensuring that the necessary service provider and human resources are in place to enable the Company to meet those goals and objectives; and
- Ensuring the identification and management of risks are robust and appropriate.

Qualifications, Skills and Experience

- Degree in Life Sciences
- At least 10 years pharmaceutical/biotechnology industry clinical trial operations experience including in drug development (Phase 1-3) and with regulatory submissions – ideally key US and EU filings (BLA, NDA, MAA)

- Demonstrated ability to manage a clinical research project according to international standards through all stages from clinical study concept development to final study report.
- Knowledge of global (US and EU) drug development regulations and guidelines; experience in interactions with regulatory authorities
- Well-developed leadership and project management skills, including demonstrated cross-functional communication, interpersonal and influence management skills with a proven track record to interact with multiple employees across multiple locations throughout the globe

Contact:

For further details and information pertaining to compensation for the role, as well as expressions of interest, please contact Telix human resources at employment@telixpharma.com or visit our careers page at www.telixpharma.com .