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

Job Description: Director of Clinical Operations
Type: Full Time
Location: USA
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-late stage pipeline of products in oncology and rare diseases.

Description

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

Reporting to the *Group Chief Medical Officer*, the successful candidate will be responsible for the leadership and oversight of US clinical study and contributing to the function's strategy in partnership with the CMO, Global Director of Clinical Operations and other members of the clinical and medical teams. The individual will actively manage operations by organising and coordinating the planning, implementation, management, execution, and completion of clinical programs according to applicable laws and regulations and SOPs. This position will also be responsible for the development of clinical operations systems and processes and communicating said processes and the successful completion of studies to internal and external stakeholders.

In addition to having overview of US Clinical Operations, the successful candidate will act as a representative for Telix's clinical operation activities and collaborate accordingly with external stakeholders such as clinicians and partners.

To be considered for this role, candidates must hold a degree in Life Sciences and demonstrate progressive experience in a similar environment. Specifically, experience in drug development (Phase 1-3) and with regulatory submissions – ideally key US and EU filings – will be highly regarded. Candidates will have a track record in managing a clinical research project in accordance with international standards through all stages: from concept development to final study report.

Agency support is not required for this role and agency submissions will not be accepted.