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

Job Description: Director or VP of Tumour Microenvironment Research
Type: Full Time
Location: Australia, Europe or United States
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

Telix is expanding again and we are on the hunt for a Director or VP of Tumour Microenvironment (TME) Research. The incumbent will be Telix's expert and lead in building new research and innovation expertise and capability in specialized oncology research topics. As a growth area of research for Telix, the initial incumbent will have the opportunity to determine the strategic direction of the company's research into the field by developing, planning and initiating a research strategy to capitalize on agents targeting the Tumour Microenvironment. The successful candidate will report to the Chief Scientist but will also work closely with the other Telix's internal R&D members, preclinical manager, clinical and project teams across multiple disciplines.

Key accountabilities for the role will include:

- Leading the TME research strategy for Telix by developing and implementing a research plan that will identify synergies between current and new products and add value to the pipeline
- Prioritising and driving research projects to evaluate, optimise and ultimately deliver meaningful research outcomes
- Fostering collaborations & partnerships with leaders & strategic partners in the field
- In conjunction with the Chief Scientist, ensuring that the aims and progress of the Research Theme are communicated effectively and shared both internally and externally
- Supporting colleagues to enhance the development of clinical-stage products by providing technical input on the design and interpretation of clinical studies
- Maintaining an awareness of the current literature, developments in the field
- Effectively managing program timelines, resources & budgets

Title and location are negotiable however to be considered, candidates must have:

- PhD in a quantitative discipline (e.g. biology, biochemistry, radiobiology, pharmacology…) or equivalent practical experience.
- Relevant applied research experience working in oncology, tumour biology and, ideally tumour microenvironment projects.
- R&D experience in the commercial environment with a solid understanding of the drug development process including preclinical studies, clinical studies and regulatory processes.
- Demonstrated knowledge of the TME research field including relevant topics of tumour biology, biomarkers of disease, immunology, histopathology and, ideally radiobiology.

- Proven prior experience managing complex technical research programs and successful 3rd party academic or commercial collaborations.
- Recognised commercial acumen to guide corporate strategy in this area and prioritise key commercial projects.

No agency submissions will be considered for this role.