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**Job Description: Director or VP of Targeted Alpha Research**

**Type: Full Time**

**Location: Australia, Europe or United States**

**Date: Immediate**

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**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 and searching for a Director or VP of Targeted Alpha Research to contribute to the achievement of the Group’s strategic goals by leading research projects in the field of Targeted Alpha Therapy (TAT) within the Telix portfolio. The incumbent will report to the Chief Scientist and will also work closely with other internal R&D members, the preclinical manager and clinical and project teams across multiple disciplines. As a growth area of research for Telix, the initial incumbent will have the opportunity to determine the strategic direction of the company’s targeted alpha strategy by developing, planning and initiating a research theme around alpha-emitting radioisotopes & related technologies.

Key accountabilities for the role will include:

- Leading the TAT research strategy in the company by developing and implementing a long-term research strategy will create platform capabilities & add value to Telix’s proprietary products and pipeline
- Prioritising and driving research projects with a commercial mindset to evaluate, optimise and ultimately deliver meaningful research outcomes
- Leading collaborations and partnerships with strategic partners relating to the production, radiolabelling processes, chelator chemistry and in vivo evaluation of relevant alpha-emitting radioisotopes
- Ensuring the delivery of ongoing and future projects by building capability through various channels
- Supporting colleagues by providing technical input on the design and interpretation of clinical studies with TAT agents.
- Maintaining an awareness of the current literature, developments in the field and encouraging strategic partnerships with companies or academic groups to identify best industry practice
- Managing program timelines, resources & budgets.

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

- PhD in a quantitative discipline (e.g. chemistry) or equivalent practical experience
- Relevant applied research experience working in radiopharmaceuticals, molecular-targeted radiation and, ideally with alpha-emitting radioisotopes
- R&D experience in the commercial environment with a solid understanding of the drug development process including preclinical studies, cGMP principles, clinical studies and regulatory processes
- Demonstrated knowledge of the TAT research field including relevant topics of radiochemistry, isotopes (and their supply chain), chelators, targeting agents and radiobiology

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

No agency submissions will be considered for this role.