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

TLX-JDXXX

Job Description:	**Medical Affairs Director**
Type:	**Full-time**
Location:	**Flexible – East Coast USA (preferably Indianapolis) or Australia (preferably Melbourne or Brisbane)**
Date:	**Immediate**

About Telix Pharmaceuticals Limited

Telix Pharmaceuticals Limited ("Telix", the "Company") is a 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 prostate, kidney and brain (GBM) cancer.

Description

The role involves supporting the development and commercialisation of a range of Telix's products globally – specifically working as part of a multi-disciplinary team to provide input from the Medical affairs perspective .

The Medical Affairs Director is responsible for providing expert contribution to the clinical development and marketing capabilities of Telix Pharmaceuticals, to support research, development and commercialisation of current and new radiopharmaceuticals.

The role can be based in either East Coast USA (preferably Indianapolis), Australia (Melbourne, Sydney or Brisbane), or Belgium (preferably Liege). The Medical Affairs Director reports to the Chief Medical Officer but is expected to have extensive interaction with key staff in the USA and other international locations.

Key Objectives: To contribute to the achievement of the company's strategic goals through the leadership, management and development of the Medical Affairs function in order to establish the medical/scientific platform for the company's portfolio. To ensure that all Medical Affairs activities are conducted within required timelines and budget, to the highest standards of Good Laboratory Practice (GLP), ICH guidelines, Good Clinical Practice (GCP) and according to regulatory guidelines, laws and Standard Operating Procedures (SOPs).

Key Accountabilities:

1. Manage the Global Medical Affairs (MA) function and ensure delivery of objectives according to agreed timelines and budget.
2. Develop and implement MA plans to support establishment of the medical/scientific platform for the company's portfolio.
3. Liaise with USA, Europe and APAC teams to ensure alignment of MA plans and implementation across the business to optimally support regional needs.
4. Design, scope and deliver late-phase (Phase IIIb/IV) clinical trials and other clinical engagement activities, both sponsored activities by the Company and investigator-initiated.
5. Manage clinical/product advisory boards and KOL engagement.

6. Develop and implement medical communications (including but not limited to abstracts, publications and conference materials) for the company's products in development and on-market.
7. 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.
8. Support and input to pharmacoeconomic modelling and strategy development, market access and pricing activities.
9. Provide medical and operational input into clinical development plans as well as other company functions as needed.
10. Participate in the review and business case development for new pipeline opportunities, production indication expansion, etc.
11. Ensure the identification and management of risks are robust and appropriate.
12. Management reporting.
13. Personal development – maintain medical/clinical knowledge and make recommendations for professional development and training.

Education and Experience:

- Medical or PhD qualifications
- 7-10 years of Medical Affairs experience at global pharmaceutical/biotechnology level
- Team leadership
- Experience in radiopharmaceuticals preferred

Competencies:

- 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 (including Outlook, Word, PowerPoint, Excel)
- Excellent oral and written communication skills

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.