# POSITION DESCRIPTION

**Job Description: Head of Pharmacovigilance (PV)**
**Type: Full Time**
**Location: Indianapolis, Liege or Melbourne**
**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

Reporting to the Chief Medical Officer, this role is responsible for overseeing the pharmacovigilance and medical information functions of Telix Pharmaceuticals and to support research, development and commercialisation of current and new radiopharmaceuticals. The Head of PV will also be expected to contribute to the achievement of the company's strategic goals through establishing and maintaining systems and processes for drug safety and drug information worldwide for the company's portfolio of products in development and in commercialisation. There will also be responsibility for ensuring that the Company's obligations to safety reporting are met worldwide and 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).

The Head of PV will be responsible for:

- Developing, implementing and maintaining systems and processes for Pharmacovigilance for the Company's development and commercialized products including urgent and periodic safety reporting and risk monitoring
- Developing and implementing systems and processes for medical information provision in collaboration with the regional medical team
- Overseeing all pharmacovigilance activities with 3rd party providers
- Providing pharmacovigilance input into clinical trial design
- Tracking, planning and preparing annual safety reporting
- Overseeing company's PVMF for product submissions
- Acting as QPPV
- ** In due course, building an internal pharmacovigilance team.

We are open to candidates in any of our three key centres: Indianapolis, Liege or Melbourne.

To be considered, candidates must have:

- A scientific background with University degree
- Pharmacovigilance experience in biotech or CRO
- Detailed knowledge of global pharmacovigilance regulations

For further details and information pertaining to compensation for the role, as well as expressions of interest, please contact Telix People and Culture at: employment@telixpharma.com