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

Job Description : Quality Assurance Specialist
Type : Full-time
Location : Indianapolis
Date : Immediate

About Us

Telix Pharmaceuticals Limited (Telix) is an Australian public Company (ASX: TLX) headquartered in Melbourne with operations in Europe, the United States and Japan. Telix's mission is to be the leading, global radiopharmaceutical company in the field of "theranostic" medicine and the Company is currently developing an extensive early through to commercial stage pipeline of products in prostate, kidney and brain cancer.

About the Role

Reporting to the US Quality Manager, the *Quality Assurance Specialist* will support the delivery of approved manufacturing, clinical and regulatory program plans for the Group's lead assets through the creation, review and approval of key documents. This role will also coordinate and assist in the active management and implementation of the Telix program/s as directed. More specifically, key accountabilities will include:

- Preparing, reviewing & approving documents within Telix Quality Management System (QMS)
- Maintaining quality requirements, quality system registers and training on eQMS
- Writing and implementing deviations, investigations, CAPA and change control reports
- Collaborating with production and manufacturing teams to monitor all work according to quality system requirements and reviewing and evaluating all standard operating procedures, batch records and quality control results to ensure compliance
- Implementing quality risk management measures to ensure compliance commensurate with stage of product lifecycle
- Providing investigational support associated with quality and technical complaints
- Providing support to cross-functional team.

About You

You hold a Bachelors degree in a related field and have some relevant and demonstrable experience within pharmaceutical/biotechnology. You're likely in the early stages of your career however you are committed to continuing professional development and training. You can write SOPs and have experience reviewing and releasing manufactured batches. You also have experience with sterility assurance principles and ideally have knowledge of applicable regulatory regulations within the relevant jurisdictions.

Why work at Telix?

We are a dynamic, fast-growing biopharmaceutical company where employees have a shared purpose: to help people with cancer and rare diseases live longer, better quality lives. This is an exciting time for Telix and we are looking for like-minded, passionate professionals to join us on the journey.

Agency support is not required for this role and no submissions will be considered.