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

Job Description: Asia Pacific QA Associate
Type: Full Time
Location: Melbourne, Australia
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

The Asia Pacific Quality Assurance Associate is a newly created role, established to keep pace with the organisation's rapid growth. Reporting to the Global Director of Quality, the QA Associate's primary remit will be to support the Asia Pacific (AP) Telix business in particular, the China Grand Pharma Partnership. The successful individual will contribute to the achievement of the Group's strategic quality goals by supporting the effective delivery of approved manufacturing, clinical and regulatory program plans for the AP Group's lead asset/s through the creation, review and approval of key documents.

Other elements of the role include:

- Preparation, review and approval of documents within the Telix Quality Management System (QMS) and eQMS (Master Control)
- Maintaining quality requirements, quality system registers and training in the eQMS.
- Participation in writing, implementing and approving Deviations, Investigations, CAPA Change Control and complaint reports.
- Participation in the preparation of the manufacturing and quality sections for regulatory submissions.
- Involvement in the management and completion of internal quality audits
- Engagement and effective management of CMOs (vendors involved in manufacture, analysis and release of drug) and other external suppliers
- Collaboration with production / manufacturing teams to monitor all work according to requirements under the Telix quality system.
- Review and evaluation of all SOPs, batch records and quality control results to ensure compliance with Telix quality and regulatory requirements

To be considered, you will require:

- A Bachelor's degree in relevant discipline
- Experience in quality assurance from a regulated industry
- Knowledge of applicable regulations within Australia, Japan, China and other Asia Pac countries is highly desirable.

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

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