



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

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**Job Description: Clinical Quality Assurance Associate**

**Type: Full Time**

**Location: Australia or USA**

**Date: Immediate**

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**About Telix Pharmaceuticals Limited**

Telix is an Australian public company (ASX: TLX) headquartered in Melbourne with international operations in Europe, the US and Japan. Our mission is to be a leading, global biopharmaceutical company that delivers on the promise of precision medicine through targeted radiation, and we are currently developing a portfolio of clinical-stage products that address significant unmet medical need in oncology and rare diseases.

**About The Role**

Demand for capability within our Global Quality Team continues to grow! The Clinical Quality Associate will be responsible for Clinical QA oversight and support for all clinical development and manufacturing activities. This individual will be a valued partner and will act as the single point of accountability for one or more clinical development projects/programs. Key accountabilities will include:

- Delivering the QA audit programme for clinical development projects/programmes
- Assisting and managing QA Plans for all programmes where audit activities are performed
- Coordinating, managing and/or leading investigations for quality issues, supporting root cause analysis activities, and ensuring subsequent corrective and preventative actions provide necessary remediation
- Generating compliance statements for the Clinical study report where audit activities have been performed

This individual can be based in Australia or US and must be comfortable working a flexible and hybrid working environment.

**About You**

You hold a Bachelor's degree in Sciences. You have demonstrated experience within auditing and clinical trials and a thorough understand of Good Clinical Practice standards. Previous QA processes/requirements and cGMP manufacturing and testing knowledge is highly preferred. You're also a forward-thinking and a proactive team player with a global mindset and have an aptitude to learn, understand and contribute.

Preferred Qualifications:

- Two or more years' experience in Good Clinical Practice
- Experience in the working of a Quality Management System
- Demonstrated
- CIPM Certified

**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. There is opportunity for individual growth and development and, an exciting pipeline of work. This is a genuine opportunity to be part of an organisation which is entering its next phase of maturity.

Application deadline is Friday 29 April 2022

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