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

Job Description: Clinical Project Manager
Type: Full Time
Location: Japan (Kyoto or Tokyo)
Date: Immediate

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 vision is to be a leading, global biopharmaceutical company in the field of “theranostic” radiopharmaceuticals and we are currently developing a portfolio of clinical-stage products that address significant unmet medical needs in oncology and rare diseases.

Description

Telix is growing again! We are currently looking to hire a Clinical Project Manager to join our Japanese team, ideally based out of Kyoto or Tokyo. The CPM will be responsible for managing all aspects of the quality delivery of clinical research studies (phase 1 to 4) from the planning phase through to study closeout. The roles coordinate work streams and cross-functional project teams and will be ultimately responsible for ensuring that Telix clinical trials within Japan (and other parts of Asia Pacific) are conducted to high global quality and regulatory standards, within the corporate timelines and budget. Key accountabilities include:

- Proactively managing all aspects of the trial process including vendor selection, site feasibility and selection, trial timelines, budgets, resources and vendor relationships
- Developing integrated study management plans with the core project team
- Developing and managing all study-related documents and implementing quality standards
- Monitoring all work performed by the external partners
- Ensuring effective project plans are in place and operational for each trial within trial appropriate SOPs
- Managing risks proactively and leading problem solving and resolution efforts

To be considered, candidates must have a minimum of a Masters in life sciences and progressive experience in a similar role for a pharmaceutical/biotechnology company and/or clinical research organization. Candidates must also demonstrate an ability to pivot study strategy quickly and lead teams in the right direction. Other desirable attributes include experience in conflict resolution, project management practice, risk management and an understanding of project financials.

CPMs need to allow for flexibility to work hours that cater for international time zones across the USA, Asia-Pacific and EMEA and have some availability to travel.

We are a team of passionate individuals who love what we do. We just need more help to do it. Come join us on our mission to help patients with cancer live longer, better quality lives!

No agency submissions will be considered for this role.