



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

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**Job Description: Regulatory Affairs Specialist**

**Type: Full Time**

**Location: EMEA (preferably  
Liège – Belgium)**

**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.

**About the Role**

*Provide support for the implementation and execution of biologic regulatory strategy for Telix pipeline.*

Working closely with the Regulatory Affairs Head in Europe, the RA associate will ensure that all aspects of the European Strategy are observed and implemented. Regulatory affairs associates are responsible for knowing all the regulations that apply to the industry they work in and ensuring that they're followed. As part of their duties, they will investigate the processes in place, review data and documents and talk to staff members.

**About You**

You live every day as a new challenge and you feel comfortable being in a fast-changing environment, so we are looking for flexible, resilient, and self-motivated individuals to join the team. You will ideally have a Master’s degree in the medical or paramedical sector or equivalent by experience. You have a strong understanding of the European drug biotechnology regulatory framework. Furthermore, you have acquired experience in interacting with regulatory authorities! You will also be confident in building relationships and collaborating with medical experts. You are problem-solving oriented and have a rigorous attitude towards the business.

English is critical however please highlight other languages of fluency in your application. The role will be in Europe (preferably in Liège, Belgium) and you will report to the local Regulatory Affairs Head in Europe. The successful candidate will be expected to interact with key staff in the USA and Australia

**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.

No agency submissions will be considered for this role.