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

Completion of Phase I Enrolment of Japanese Renal Cancer Study

Melbourne (Australia) – 27th October 2020. Telix Pharmaceuticals Limited (ASX: TLX, 'Telix', the 'Company') today announces the completion of Phase I enrolment of the Company's Phase I/II ZIRDAC-JP study of Telix's renal cancer diagnostic imaging product TLX250-CDx (^{89}Zr-girentuximab) in Japan.

The ZIRDAC-JP study (Zirconium Dosing and Comparison in Japan) is a Japanese multi-centre Phase I/II study that will recruit approximately 40 patients in total. The objective of the Phase I component of the ZIRDAC-JP study is to comparatively evaluate the safety, tolerability, radiation dosimetry and pharmacokinetics / pharmacodynamics of TLX250-CDx in Japanese patients.[1]

The patient population for the ZIRDAC-JP trial has been selected to be identical to Telix's global Phase III ZIRCON trial, which is presently recruiting patients at 33 sites internationally and is expected to complete recruitment in Q1 2021. The ZIRDAC-JP study has been designed in consultation with the Japanese Pharmaceuticals and Medical Devices Agency (PMDA) to collect the necessary data to potentially bridge to the ZIRCON study, by confirming that dosing and pharmacology of TLX250-CDx in Japanese patients is equivalent to non-Japanese patients. Following data review and consultation with the PMDA, Telix expects to commence the Phase II component of the ZIRDAC-JP study in early 2021.

Telix Pharmaceuticals Japan K.K. President, Dr. Shintaro Nishimura stated, "The efficient completion of enrolment into the Phase I part of the ZIRDAC-JP study is a testament to the commitment of Dr Nakaigawa, the principal investigator and his team at Yokohama City University Hospital, and most importantly the patients who participated in this trial under COVID-19 conditions. This represents a significant milestone for Telix Japan and the broader Japanese nuclear medicine community. We now look forward commencing the Phase II part of the ZIRDAC-JP study and ultimately bringing this breakthrough kidney cancer imaging product to market in Japan in coordination with the global development for this asset."

About Telix Pharmaceuticals Limited

Telix is a clinical-stage biopharmaceutical company focused on the development of diagnostic and therapeutic products using Molecularly Targeted Radiation (MTR). Telix is headquartered in Melbourne, Australia with international operations in Belgium, Japan and the United States. Telix is developing a portfolio of clinical-stage oncology products that address significant unmet medical needs in prostate, kidney and brain cancer. Telix is listed on the Australian Securities Exchange (ASX: TLX). For more information visit www.telixpharma.com.

Telix Corporate Contact

Dr Christian Behrenbruch
Telix Pharmaceuticals Limited
CEO
Email: chris@telixpharma.com

Telix Investor Relations

Dr David N. Cade
Telix Pharmaceuticals Limited
CBO and Head of Investor Relations
Email: david.cade@telixpharma.com

[1] ClinicalTrials.Gov Identifier: NCT04496089

Important Information

This announcement does not constitute an offer to sell, or a solicitation of an offer to buy, securities in the United States, or in any other jurisdiction in which such an offer would be illegal. The securities referred to herein have not been and will not be registered under the United States Securities Act of 1933 (the "U.S. Securities Act"), or under the securities laws of any state or other jurisdiction of the United States and may not be offered or sold within the United States, unless the securities have been registered under the U.S. Securities Act or an exemption from the registration requirements of the U.S. Securities Act is available. None of the technologies or products described in this document have received a marketing authorisation in any jurisdiction. This announcement has been authorised for release by Dr Christian Behrenbruch, Managing Director and Chief Executive Officer.