UNITED STATES
SECURITIES AND EXCHANGE COMMISSION
Washington, D.C. 20549

FORM 6-K

REPORT OF FOREIGN PRIVATE ISSUER
PURSUANT TO RULE 13a-16 OR 15d-16 OF
THE SECURITIES EXCHANGE ACT OF 1934

For the month of April, 2026

Commission File Number: **001-42128**

Telix Pharmaceuticals Limited

(Translation of registrant's name into English)

55 Flemington Road
North Melbourne, Victoria 3051, Australia
(Address of principal executive offices)

Indicate by check mark whether the registrant files or will file annual reports under cover of Form 20-F or Form 40-F.

Form 20-F ☒ Form 40-F ☐

INFORMATION CONTAINED IN THIS FORM 6-K REPORT

On April 14, 2026 (Melbourne, Australia), Telix Pharmaceuticals Limited (the "Company") filed announcements with the Australian Securities Exchange titled "Telix Refinances Convertible Bonds," a copy of which is attached to this Form 6-K as Exhibit 99.1 and "Investor Presentation," a copy of which is attached to this Form 6-K as Exhibit 99.2.

99.1	Press release – April 14, 2026
99.2	Presentation – April 14, 2026

SIGNATURES

Pursuant to the requirements of the Securities Exchange Act of 1934, the registrant has duly caused this report to be signed on its behalf by the undersigned, thereunto duly authorized.

Telix Pharmaceuticals Limited

Date: April 14, 2026

By: /s/ Christian Krautkramer
Name: Christian Krautkramer
Title: Group General Counsel

Exhibit 99.1

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

ASX ANNOUNCEMENT

Telix Refinances Convertible Bonds

Melbourne (Australia) and Indianapolis, IN (U.S.) – April 14, 2026. Telix Pharmaceuticals Limited (ASX: TLX, NASDAQ: TLX) ("**Telix**") today launches an offering of US$550 million convertible notes due 2031 to be issued by its wholly-owned subsidiary, Telix Pharmaceuticals (Investments) Inc. (the "**Issuer**"), and guaranteed by Telix and Telix Pharmaceuticals (US) Inc. (the "**Offering**"). The convertible notes, also referred to as "convertible bonds" ("**Convertible Bonds**"), are convertible into fully paid ordinary shares in Telix ("**Ordinary Shares**").

Managing Director and Group CEO, Dr. Christian Behrenbruch, said: "The refinance of the existing Convertible Bonds represents our proactive approach to capital management. The new Convertible Bonds will continue to provide the business with cost effective financing."

The new Convertible Bonds represent attractive, low-cost financing to Telix and are non-dilutive until any potential future conversions occur. The initial conversion price will be at a premium to Telix's current share price.

After deduction of commissions, professional fees and other administrative expenses, it is intended that the net proceeds will be used to repurchase the existing convertible bonds due 2029 ("**Existing Convertible Bonds**"). Any funds raised above that required for the repurchase of the Existing Convertible Bonds will be applied to general corporate purposes.

Convertible Bonds Offering

- It is intended that the Convertible Bonds will be listed on the Official List of Singapore Exchange Securities Trading Limited ("**SGX-ST**")
- The Offering is being marketed to eligible investors with the final terms of the Convertible Bonds to be determined via a bookbuild process expected to be completed prior to market open on Wednesday, 15 April 2026
- Concurrent with the Offering, a delta placement of Ordinary Shares will be executed to facilitate hedging activity by investors in relation to the Convertible Bonds. The clearing price per Ordinary Share under the delta placement will be used as the reference share price for the Convertible Bonds
- More details on the key terms of the Convertible Bonds are provided in the table below

Stock Borrow Facility

To assist the implementation of the Convertible Bonds Offering, Elk River Holdings Pty Ltd as the trustee for The Behrenbruch Family Trust[1] ("**Stock Lender**") intends to enter into a stock lending agreement with an affiliate of J.P. Morgan ("**Stock Borrower**") pursuant to which the Stock Lender will lend a certain number of Ordinary Shares to the Stock Borrower, and the Stock Borrower will be required to return the borrowed Ordinary Shares to the Stock Lender pursuant to the terms of the agreement ("**Stock Borrow Facility**").

[1] Dr Behrenbruch holds an indirect interest.

Concurrent Repurchase

Additionally, Telix is conducting a reverse bookbuilding process to receive indications of interest from holders of the Existing Convertible Bonds on a repurchase of the outstanding Existing Convertible Bonds ("**Concurrent Repurchase**"). The number of Existing Convertible Bonds to be repurchased and the purchase price will be determined by the reverse bookbuilding process.

Adviser

J.P. Morgan Securities plc ("**J.P. Morgan**") is Sole Bookrunner on the Offering and Sole Dealer Manager on the Concurrent Repurchase.

Key terms of the Convertible Bonds

Issuer	Telix Pharmaceuticals (Investments) Inc.
Guarantors	Telix Pharmaceuticals Limited and Telix Pharmaceuticals (US) Inc.
Expected Issue Size	US$550 million
Ranking	Direct, unconditional, unsubordinated and unsecured obligations of the Issuer and Guarantors
Maturity Date	5 years
Investor Put Option	At the end of year 3
Coupon / Yield	1.50 – 1.75%
Conversion Premium	35.0 – 37.5%
Reference Share Price	The clearing price of the Delta Placement – see below
Delta Placement	J.P. Morgan will run a bookbuilding process to facilitate some or all of the hedging activity that may be executed by investors in the Convertible Bonds The clearing price of the Delta Placement will be used as the reference share price to determine the initial conversion price of the Convertible Bonds The manner of conducting the Delta Placement will be determined by J.P. Morgan in consultation with Telix
Conversion Price Adjustment	Standard anti-dilutive adjustments including conversion price adjustment for all dividends paid by Telix
Listing	SGX-ST
Selling Restrictions	Reg S (Cat 2) only

About Telix Pharmaceuticals Limited

Telix is a global biopharmaceutical company focused on the development and commercialization of radiopharmaceuticals with the goal of addressing significant unmet medical need in oncology and rare diseases. Telix is headquartered in Melbourne (Australia) with international operations in the United States, United Kingdom, Brazil, Canada, Europe (Belgium and Switzerland), and Japan, Telix is listed on the Australian Securities Exchange (ASX: TLX) and the Nasdaq Global Select Market (NASDAQ: TLX).

Visit www.telixpharma.com for further information about Telix, including details of the latest share price, ASX and U.S. Securities and Exchange Commission (SEC) filings, investor and analyst presentations, news releases, event details and other publications that may be of interest. You can also follow Telix on LinkedIn, X and Facebook.

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SVP Investor Relations and
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Telix Investor Relations (U.S.)
Ms. Annie Kasparian
Director Investor Relations and
Corporate Communications
annie.kasparian@telixpharma.com

This announcement has been authorized for release by the Telix Pharmaceuticals Limited Board of Directors.

Legal Notices

Cautionary Statement Regarding Forward-Looking Statements.

You should read this announcement together with our risk factors, as disclosed in our most recently filed reports with the Australian Securities Exchange (ASX), U.S. Securities and Exchange Commission (SEC), including our Annual Report on Form 20-F filed with the SEC, or on our website.

The information contained in this announcement is not intended to be an offer for subscription, invitation or recommendation with respect to securities of Telix Pharmaceuticals Limited (Telix) in any jurisdiction, including Australia, Singapore, and the United States. The information and opinions contained in this announcement are subject to change without notification. To the maximum extent permitted by law, Telix disclaims any obligation or undertaking to update or revise any information or opinions contained in this announcement, including any forward-looking statements (as referred to below), whether as a result of new information, future developments, a change in expectations or assumptions, or otherwise. No representation or warranty, express or implied, is made in relation to the accuracy or completeness of the information contained or opinions expressed in the course of this announcement.

This announcement may contain forward-looking statements, including within the meaning of the U.S. Private Securities Litigation Reform Act of 1995, that relate to anticipated future events, financial performance, plans, strategies or business developments. Forward-looking statements can generally be identified by the use of words such as "may", "expect", "intend", "plan", "estimate", "anticipate", "believe", "outlook", "forecast" and "guidance", or the negative of these words or other similar terms or expressions. Forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause our actual results, levels of activity, performance or achievements to differ materially from any future results, levels of activity, performance or achievements expressed or implied by these forward-looking statements. Forward-looking statements are based on Telix's good-faith assumptions as to the financial, market, regulatory and other risks and considerations that exist and affect Telix's business and operations in the future and there can be no assurance that any of the assumptions will prove to be correct. In the context of Telix's business, forward-looking statements may include, but are not limited to, statements about: the initiation, timing, progress, completion and results of Telix's preclinical and clinical trials, and Telix's research and development programs; Telix's ability to advance product candidates into, enroll and successfully complete, clinical studies, including multi-national clinical trials; the timing or likelihood of regulatory filings and approvals for Telix's product candidates, including TLX101-Px and TLX250-Px, manufacturing activities and product marketing activities; Telix's sales, marketing and distribution and manufacturing capabilities and strategies; the commercialization of Telix's product candidates, if or when they have been approved; Telix's ability to obtain an adequate supply of raw materials at reasonable costs for its products and product candidates; estimates of Telix's expenses, future revenues and capital requirements; Telix's financial performance; developments relating to Telix's competitors and industry; the anticipated impact of U.S. and foreign tariffs and other macroeconomic conditions on Telix's business, including as a result of war or other geopolitical conflicts; and the pricing and reimbursement of Telix's product candidates, if and after they have been approved. Telix's actual results, performance or achievements may be materially different from those which may be expressed or implied by such statements, and the differences may be adverse. Accordingly, you should not place undue reliance on these forward-looking statements.

Neither this announcement nor any copy hereof may be taken into or distributed in the United States.

*The information contained in this announcement is not for distribution, directly or indirectly, in or into the United States. The Convertible Bonds, the guarantees and the Ordinary Shares to be issued upon conversion of the Convertible Bonds have not been, and will not be, registered under the U.S. Securities Act of 1933, as amended (the "**Securities Act**") or the securities laws of any state or other jurisdiction of the United States and they may not be offered or sold, resold, transferred or delivered, directly or indirectly, within the United States or to, or for the account or benefit of U.S. persons (as defined in Regulation S under the Securities Act) except pursuant to an exemption from, or in a transaction not subject to, the registration requirements of the Securities Act and applicable state or local securities laws. The Convertible Bonds and the guarantees are being offered and sold solely outside the United States in an "offshore transaction" as defined in, and in reliance on Regulation S under the Securities Act.*

Nothing in this announcement or anything attached to it shall form the basis of any contract or commitment.

The Concurrent Repurchase is not being made and will not be made, directly or indirectly, in or into the United States. This includes, but is not limited to, facsimile transmission, electronic mail, telex, telephone, the internet and other forms of electronic communication. The Existing Convertible Bonds may not be tendered in the Concurrent Repurchase by any such use, means, instrumentality or facility from or within the United States or by persons located or resident in the United States as defined in Regulation S of the Securities Act. Any purported tender of Existing Convertible Bonds made by a person located in the United States will not be accepted.

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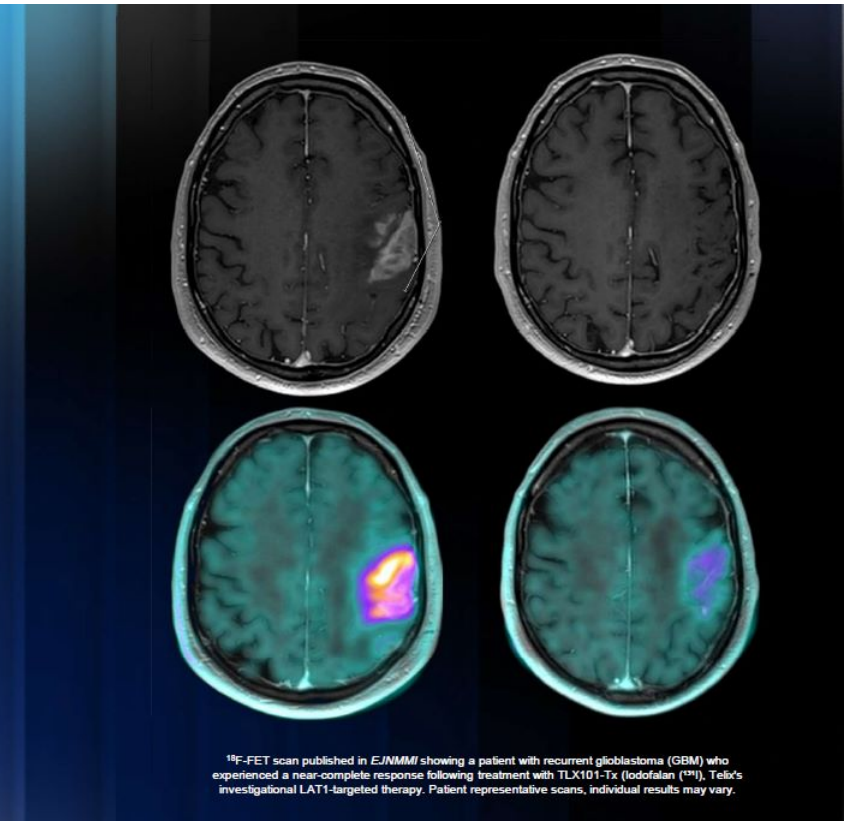
Telix

Investor presentation

April 14, 2026

ASX: TLX | NASDAQ: TLX



^{18}F-FET scan published in *EJNMMI* showing a patient with recurrent glioblastoma (GBM) who experienced a near-complete response following treatment with TLX101-Tx (Iodofalan (^{131}I)), Telix's investigational LAT1-targeted therapy. Patient representative scans, individual results may vary.

Forward looking statements

You should read this presentation together with our risk factors, as disclosed in our most recently filed reports with the Australian Securities Exchange (ASX), U.S. Securities and Exchange Commission (SEC), including our Annual Report on Form 20-F filed with the SEC, or on our website.

The information contained in this presentation is not intended to be an offer for subscription, invitation or recommendation with respect to securities of Telix Pharmaceuticals Limited (Telix) in any jurisdiction, including Australia, Singapore, and the United States. The information and opinions contained in this presentation are subject to change without notification. To the maximum extent permitted by law, Telix disclaims any obligation or undertaking to update or revise any information or opinions contained in this presentation, including any forward-looking statements (as referred to below), whether as a result of new information, future developments, a change in expectations or assumptions, or otherwise. No representation or warranty, express or implied, is made in relation to the accuracy or completeness of the information contained or opinions expressed in the course of this presentation.

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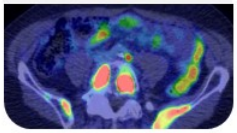
Telix

Five pillars underpinning our global leadership in radiopharma

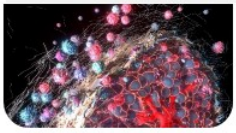
Telix

Integrated Theranostic Approach
See It. Treat It.

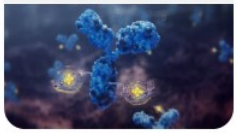
High value therapeutics pipeline



R&D platform for new molecular entities



Precision Medicine portfolio



Specialist commercial organization

Vertically integrated manufacturing and supply chain

Our performance in 2025[1]

Enabling investments that will drive value creation

- **Strong top-line growth, underlying profitability and cash generation**
- **Group revenue $804M up 56% YoY, meeting increased guidance**

- Precision Medicine revenue up 22% YoY to $622M
- RLS Radiopharmacies adds diversified revenue stream – $170M reported revenue[2]
- Group EBITDA[3] $40M, positive cash balance of $142M

Over $500M investment across R&D, commercial infrastructure and strategic investments has delivered:

Pipeline: Four assets in pivotal / Ph3 trials[4]

ProstACT™ Global · IPAX BrIGHT · LUTEON · BiPASS

High-value clinical research

Commercial: Increased market share driven by our multi-product strategy[5]

Gozellix® (kit for the preparation of gallium Ga 68 gozetotide injection) · illuccix® (kit for the preparation of gallium Ga 68 gozetotide injection)

Platform for continued growth

Infrastructure and global supply chain

RLS Radiopharmacies · IsoTherapeutics · ARTMS

Enhancing our commercial offering

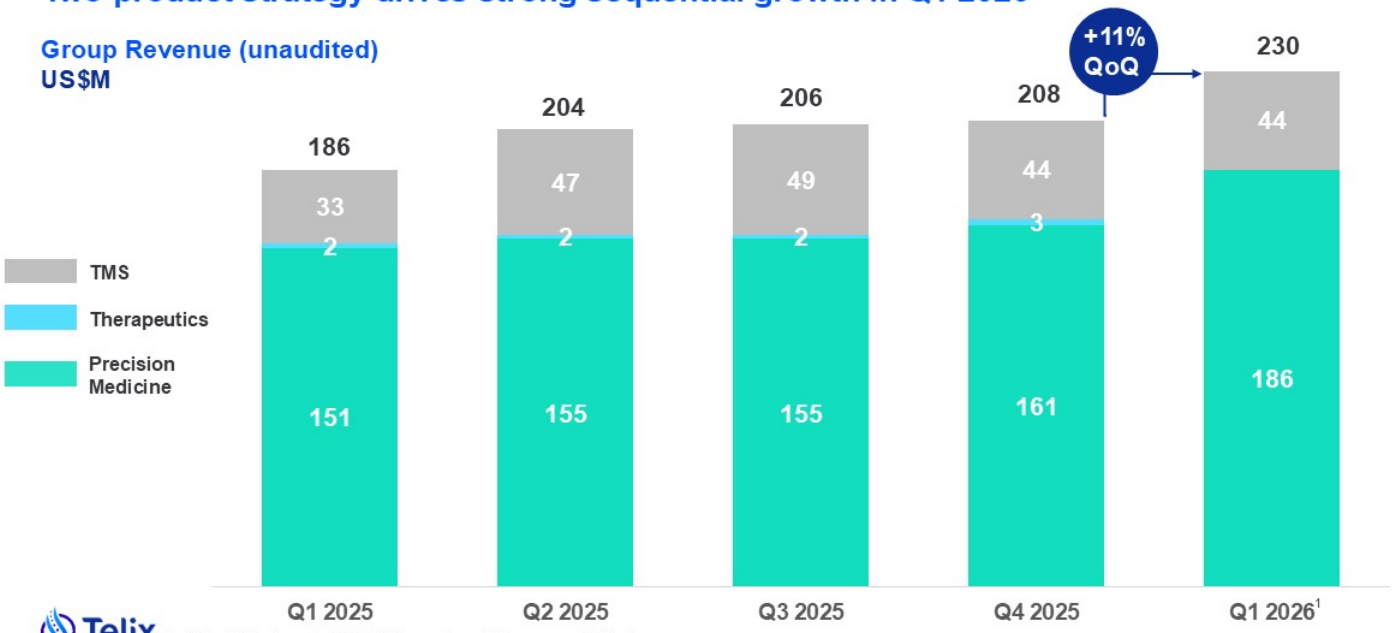
Telix

1. All figures throughout presentation provided in USD.
2. Since acquisition close as of January 28, 2025.
3. Earnings before interest, tax, depreciation and amortization.
4. ProstACT Global, NCT06520345; IPAX BrIGHT, NCT07100730; LUTEON, NCT07197580; BiPASS, NCT07052214.
5. Gozellix currently only approved and available in the U.S.

Q1 2026: Group revenue of US$230M, up 11% QoQ

Two-product strategy drives strong sequential growth in Q1 2026

Group Revenue (unaudited)
US$M



1. Telix ASX disclosure April 7, 2026. Revenue is provided on an unaudited basis.

Delivering on targeted milestones in early 2026

Demonstrating continued commercial momentum and pipeline advancement

Pathway to launch new Precision Medicine products

- **Pixclara®[1] (TLX101-Px, Floretyrosine F 18 or ^{18}F-FET):** PDUFA[2] goal date of September 11, 2026[3]
- **Pixlumi®[1] (TLX101-Px):** European MAA[4] submitted
- **Illuccix® (TLX591-Px) New Drug Application** accepted in China

Therapeutic pipeline advancement

- **ProstACT Global Phase 3 Study of TLX591-Tx in prostate cancer:** Part 1 objectives achieved, acceptable safety and tolerability[5]
- **Two additional pivotal therapy trials recruiting: IPAX BrIGHT (TLX101-Tx, brain)** first patient enrolled, **LUTEON (TLX250-Tx, kidney)** open for recruitment

Strategic collaboration with Regeneron

- Focused on next-generation therapies, endorsement of Telix's radiopharma development and manufacturing platform[6]

Telix

1. Brand name subject to final regulatory approval.
2. Prescription Drug User Fee Act.
3. Telix ASX disclosure April 10, 2026.
4. Marketing Authorization Application.
5. Telix ASX disclosure March 10, 2026.
6. Telix ASX disclosure April 13, 2026.

Telix

Precision Medicine

Precision Medicine portfolio

Telix

Precision Medicine near-term growth strategy

PSMA portfolio provides platform for growth, potential upside from new product launches

Gozellix® (kit for the preparation of gallium Ga 68 gozetotide injection)

- **Successful launch (U.S.)**
- **BiPASS™,** Phase 3 study launched and recruiting rapidly – will support significant market expansion for Illuccix and Gozellix

Illuccix® (kit for the preparation of gallium Ga 68 gozetotide Injection)

- **Commercially available in 22 countries**[1]
- **Pursuing further geographic expansion in China** (NDA accepted)[2] **and Japan** (Phase 3 study dosing patients)[3]

Zircaix[7] (zirconium Zr 89 girentuximab senvedoxam-crdi)

- Successful Type A meetings completed to **align with FDA** on key resubmission items
- **ZIRCON-X** study showed TLX250-Px imaging has meaningful impact on clinical decisions[4]

Pixclara[7] (floretyrosine F 18)

- **MAA filed for TLX101-Px in Europe** and **NDA accepted by U.S. FDA,** assigned **PDUFA goal date** September 11, 2026[5]
- In a survey of 100 physicians (U.S.), **~70% indicated they are ready to prescribe Pixclara** upon FDA approval[6]

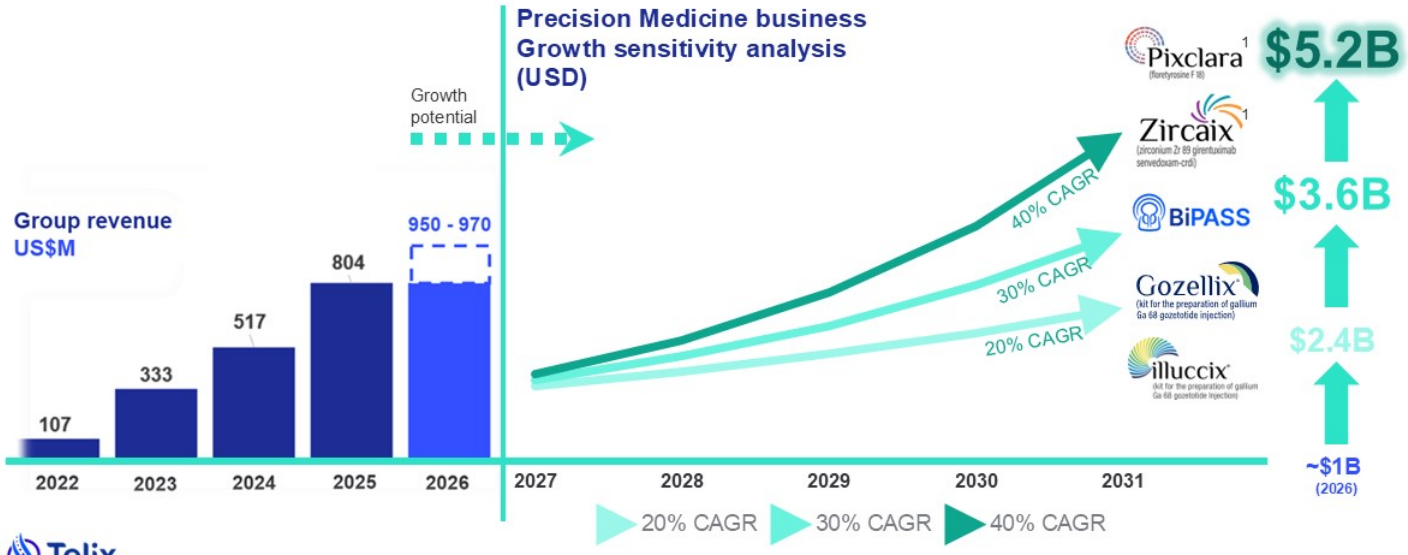
Regulatory submission focus on Pixclara and Zircaix in the U.S. to pave way for future launches

Telix Precision Medicine

1. Austria, Czech Republic, Italy, Slovakia, Switzerland, UK, France, Germany, Spain, Portugal, Belgium, Luxembourg, Netherlands, Denmark, Sweden, Finland, Norway, Australia, NewZealand, Brazil, U.S. and Canada.
2. Chinese National Medical Products Administration (NMPA) Center for Drug Evaluation (CDE) accepted the filing of a NDA, Telix ASX disclosure January 20, 2026.
3. Telix ASX disclosure January 26, 2026.
4. Telix media release November 20, 2025.
5. Telix ASX release April 10, 2026.
6. Shoreline Research, Awareness trial and utilization report, Jan, 2026
7. Brand names subject to final regulatory approval.

Precision medicine: Five-year outlook

Building on the current commercial portfolio with BiPASS, Zircaix and Pixclara[1]



Telix

Precision Medicine

Not intended as a forecast or guidance, subject to change due to market conditions and regulatory approvals.

1. Brand names subject to final regulatory approval.

Telix

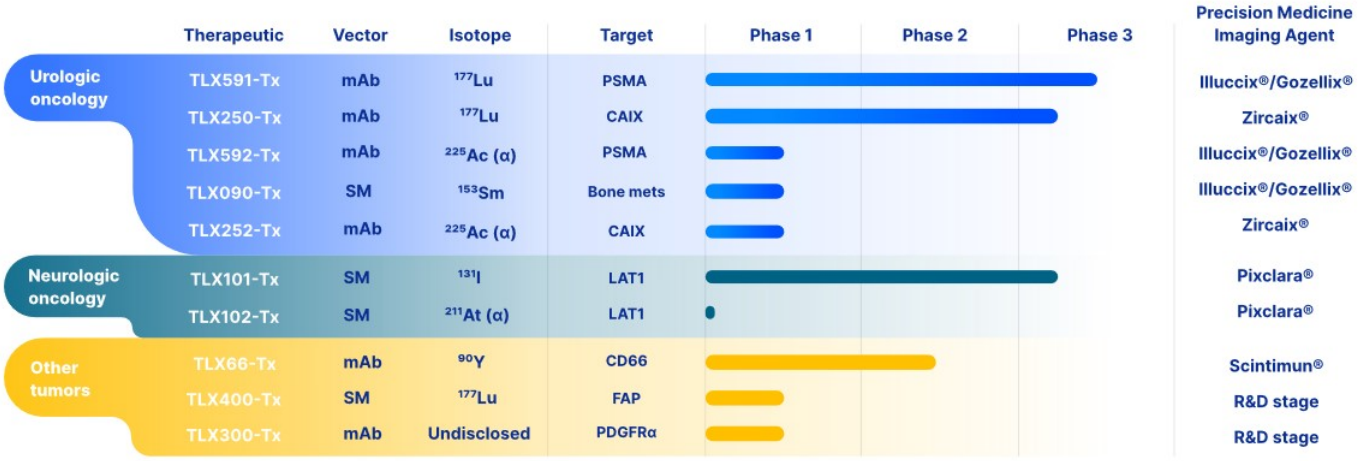
Therapeutics

Therapeutics portfolio

Telix

Therapeutics pipeline: Late-stage and “next-generation” assets

Building a leadership position in urologic and neurologic oncology



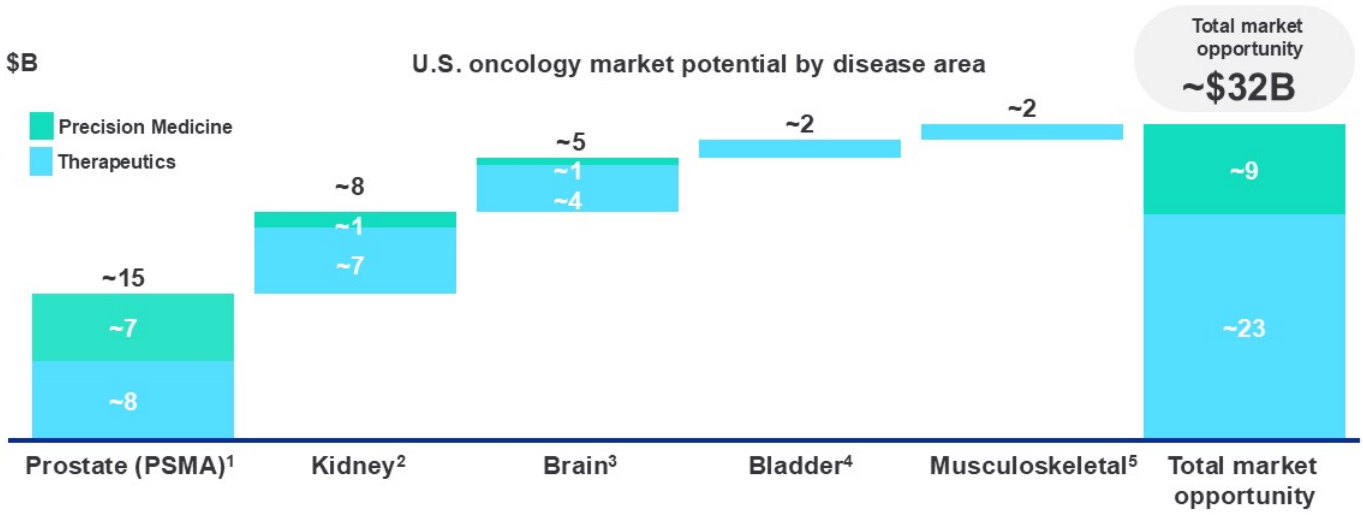
Telix Therapeutics

PSMA: Prostate-specific membrane antigen.
CAIX: Carbonic anhydrase IX.
LAT1: L-Type amino acid transporter 1.
CD66: Cluster of differentiation 66.

PDGFRα: Platelet-derived growth factor receptor alpha.
mAb: Monoclonal antibody.
SM: Small molecule.
FAP: Fibroblast activation protein.

Potential $32B market opportunity

Long-term growth potential across our Precision Medicine and Therapeutics pipeline



1. Datamonitor Cancer Patient-Based Forecast and Management Internal Estimates. Prostate-specific membrane antigen.
2. Datamonitor Renal Cell Carcinoma patient-based forecast model and Management Internal Estimates.
3. Datamonitor Glioblastoma patient-based forecast model, and Management Internal Estimates. Leptomeningeal disease (Brain): Nguyen, A.; Nguyen, A.;Dada, O.T.; Desai, P.D.; Ricci, J.C.;Godbole, N.B.; Pierre, K.;Lucke-Wold, B. Leptomeningeal Metastasis: A Review of the Pathophysiology, Diagnostic, and Therapeutic Landscape. Curr. Oncol. 2023.
4. Datamonitor Bladder Cancer 2024.
5. Lowery, Caitlin D., et al. "Olaratumab Exerts Antitumor Activity in Preclinical Models of Pediatric Bone and Soft Tissue Tumors." Clinical Cancer Research, vol. 24, no. 4, Feb. 2018, pp. 847-857. American Association for Cancer Research. Estimates updated Dec, 2025.

Late-stage assets with upcoming clinical inflection points

Phase 3	Phase 2/3	Phase 2/3	Phase 1
ProstACT™ Global	IPAX BrIGHT	LUTEON	Solace
TLX591-Tx, first-in-class rADC for mCRPC **Part 1 lead in** safety, dosimetry, data readout[1] – primary objectives met **Part 2** (randomized treatment expansion), **currently dosing patients (ex-U.S.)**[2]	**TLX101-Tx, potential first radiotherapy in recurrent GBM** **Part 1** (dose optimization), enrolling patients[3] in Australia and Europe **Part 2**, primary endpoint: OS **ODD** in U.S. and Europe **IPAX-2**: Phase 1 trial, newly diagnosed patients – data in 2026	**TLX250-Tx, first-in-class rADC for advanced or metastatic ccRCC** **Part 1** (dose optimization), open for recruitment in Australia. Primary endpoints: safety, RP3D[4] **IND and CTA submissions in 2026 (US/EU)** **Part 2** primary endpoint: mPFS	**TLX090-Tx, novel treatment for bone pain in patients with osteoblastic lesions** **Part 1** (dose escalation), currently dosing patients (U.S.) Primary endpoints: safety, dosimetry[5] **Part 2** (dose selection). Primary endpoint: optimal dose (safety, pain score)

rADC = radio antibody-drug conjugate, mCRPC = metastatic Castration-Resistant Prostate Cancer, GBM = Glioblastoma, OS = Overall survival, ODD = Orphan drug designation, EAP = Expanded access program, ccRCC = clear cell Renal Cell Carcinoma, RP3D = recommended phase 3 dose, IND = Investigational newdrug, CTA = Clinical trial application, mPFS = median progression free survival, SoC = Standard of care.

Telix Therapeutics

1. Telix ASX disclosure March 10, 2026.
2. Part 2 is enrolling in Australia, NewZealand, and Canada, and has also received regulatory approval to commence in China, Singapore, South Korea, Türkiye, and the United Kingdom. Japanese regulator Pharmaceuticals and Medical Devices Agency (PMDA) has granted approval for a Japan-specific Part 1 in nine patients, prior to commencing Part 2.
3. ClinicalTrials.gov ID: NCT07100730.
4. ClinicalTrials.gov ID: NCT05663710.
5. ClinicalTrials.gov ID: NCT07197645.

ProstACT Global Phase 3 (Part 1 Lead-in): Key findings

Primary and secondary endpoints: Safety and dosimetry

- **Study objectives met:** Confirmed safety, pharmacokinetics, dosimetry across cohorts
- **No new safety signals:** Hematologic events transient and manageable
- **Tolerability profile** supported by dosimetry and low-grade non-hematologic events
- Lesion dosimetry indicates no difference in **absorbed dose profile** across cohorts
- **No adverse drug-drug interactions** observed in TLX591-Tx combinations

Demonstrates feasibility of integrating TLX591-Tx with contemporary, global standards of care

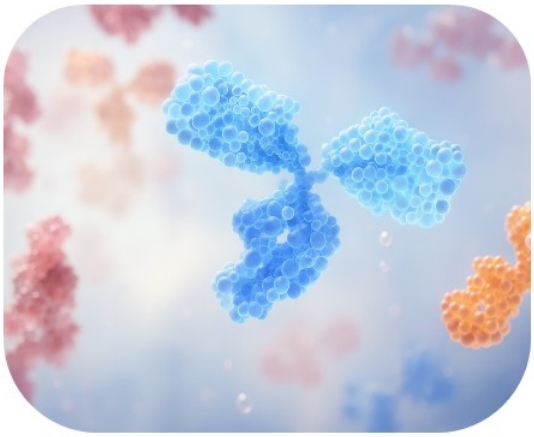
Telix Therapeutics

Source: ProstACT Global Part 1 data on file.

Strategic collaboration with Regeneron (NASDAQ: REGN)

Highly complementary capabilities present a unique opportunity

- Collaboration to **jointly develop and commercialize** next generation radiopharmaceutical therapies, including **targeted alpha therapies**[1]
- Telix to receive **US$40 million cash upfront** with option to co-fund through commercialization and profit share or **earn up to US$2.1 billion** in development and commercial milestone payments **plus low double-digit royalties**
- Leverages Regeneron's **extensive biologics expertise**, with Telix's radiopharma development platform and **global manufacturing and supply chain infrastructure**
- Spans **multiple solid tumor targets**, from Regeneron's portfolio of proprietary, clinically validated antibodies with initial focus on **lung cancer**



1. Telix ASX disclosure April 13, 2026.

2026 strategic priorities

Telix

FY 2026 guidance

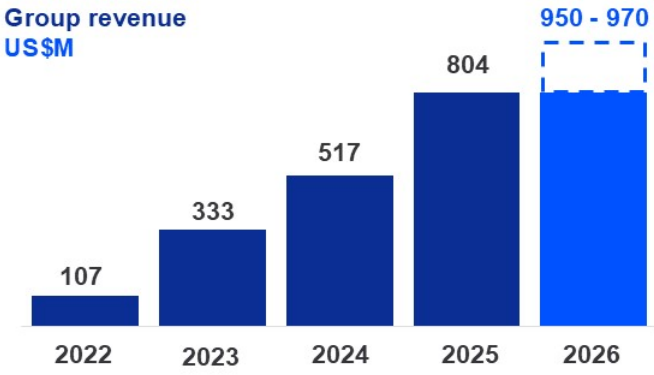
Continued growth trajectory

Revenue guidance range: $950M to $970M

- Based on approved products and geographies
- ~25% growth in Precision Medicine revenue
- Full year of RLS revenue

R&D guidance range: $200M to $240M

- Increased allocation to therapeutics pipeline
- R&D investment will be guided by clinical data outcomes and milestones
- Self-funded through commercial performance



We will continue reinvesting our earnings to position the Company for long-term growth

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Based on expected global and domestic economic conditions and subject to known and unknown risks, uncertainties and other factors that may cause our actual results to differ materially. See Disclaimer for more information.

A catalyst rich 2026

Select milestones for Therapeutics candidates

- **Strategic collaboration with Regeneron**
- **TLX591-Tx** for mCRPC, ProstACT Global
  - **Part 1 data readout**
  - Part 2 international site expansion, interim analysis[1]
- **TLX250-Tx** for ccRCC, LUTEON, open for recruitment
- **TLX101-Tx** for recurrent GBM
  - IPAX BrIGHT, first patient enrolled
  - IPAX 2- data readout MTD (Max tolerated dose)
- **TLX090-Tx** for metastatic bone pain, SOLACE, enrollment completion
- **TLX592-Tx** for mCRPC, AlphaPRO, patient dosing
- **TLX102-Tx** for recurrent GBM and leptomeningeal disease, trial commencement
- **TLX252-Tx** for ccRCC and other CAIX-expressing tumors, trial commencement
- **TLX400-Tx** recommencement of clinical activity

Select milestones for Precision Medicine candidates

- **Pixclara[2] NDA resubmission (U.S.) accepted,** PDUFA goal date September 11, 2026
- **Pixlumi[2] MAA acceptance (Europe)**
- **Zircaix[2] BLA resubmission (U.S.)**
- **Illuccix, Gozellix** BiPASS™ enrollment completion
- **Illuccix** Japan trial, enrollment completion
- **Illuccix** China, regulatory approval/launch
- **TLX593-Px (AlFluor™)** trial commencement

Select milestones for Telix Manufacturing Solutions

- **Key RLS sites:** commence **cyclotron** installations EU (Brussels) and Japan (Yokohama) cyclotrons in production
- **50 ARTMS QIS installations** globally

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BLA = Biologics license application, QIS = QUANTM irradiation system, a cyclotron-based isotope production system.

1. Protocol ^{177}Lu-TLX591-203.
2. Brand name subject to final regulatory approval.

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Q&A

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Illustration of TLX591-Tx
Lutetium (^{177}Lu) rosopatamab tetraxetan