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PRESENTATION OF INFORMATION

All currency amounts in this presentation are in Australian dollars unless otherwise stated.



PRESENTING



Will Knox

CEO

Will comes from a long history of medical and biological technology development. Prior to Tetratherix, Will has worked for small and large businesses in Australia and offshore such as Cochlear, Medtronic and LifeHealthcare. Will has also established his own company in regenerative medicine that was acquired in 2017 by Australian private equity. Combining both the clinical and commercial worlds, Will brings a unique and well respected combination of skillsets to our business.



Dr Ali Fathi

FOUNDER & CTO

Ali is a world-renowned researcher and inventor of Tetramatrix and has continued the development of the technology using his technical and entrepreneurial vision. Ali is a globally respected chemical engineer and is one of Australia's most published young researchers. Using his unique approach to engineering science-based solutions for the real world, Ali has created a technology that has been deliberately considered from start to finish.



Cherie Beach

CFO

Cherie is the epitome of a strategic CFO and brings many years of experience within healthcare and medical technology. Cherie was most recently a senior finance leader at Cochlear and Johnson&Johnson, and has a long history of strategic financial stewardship in our industry. Cherie has an unparalleled combination of leading global financial process with fast-growing innovative thinking - a combination that allows us to grow rapidly while ensuring commercial stability & reliability.



Terence Abrams

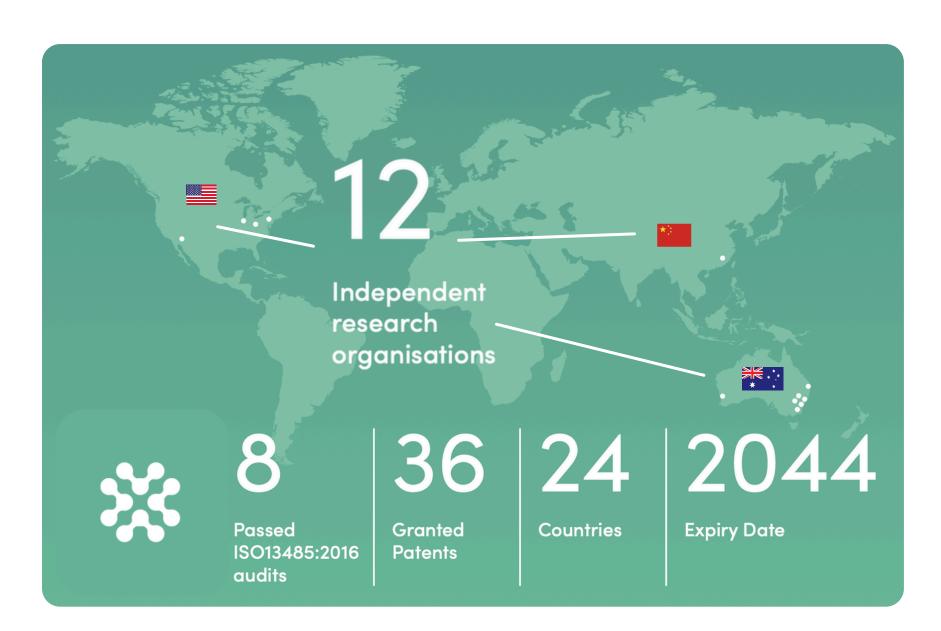
FOUNDER & COO

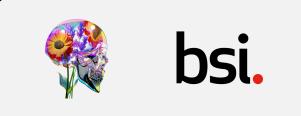
Terence (Tez) is an extremely experienced chemical engineer who has over a decade of experience in bespoke compounding, polymer production & infrastructure design for complex biomaterials. Tez is the engine that drives the unique differentiation behind our advanced manufacturing. Adopting critical thinking and first principles, Tez is a pragmatic problem solver who combines chemistry, quality systems and commercial objectives - a skillset that is standalone in our industry.

The executive leadership team are supported by a deep bench of scientific & commercial advisors who are highly regarded leaders in their respective fields



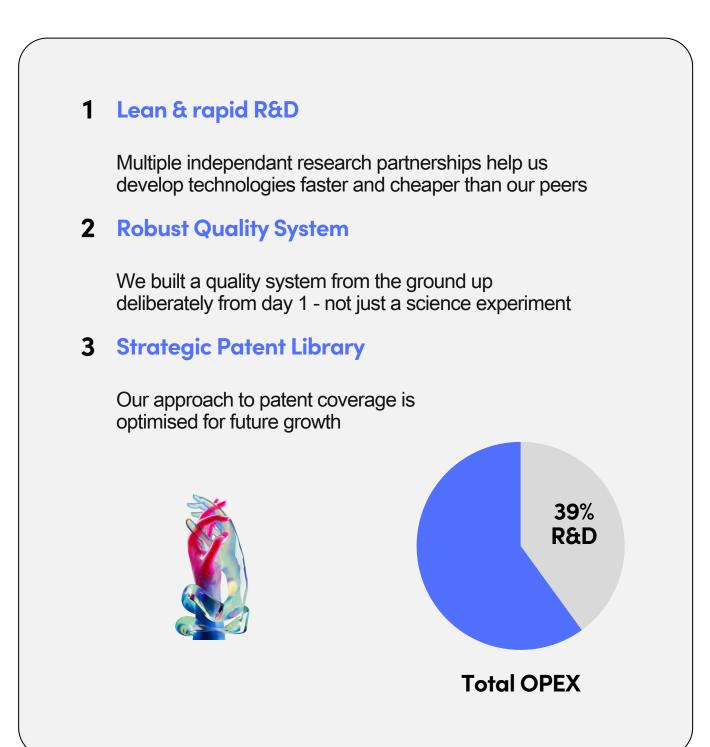
An operational moat with IP Leadership built on a global R&D infrastructure. This sets us apart.





Supported by complete IP coverage

Patent stack with 36 granted patents from 9 families of patent, extending to 2044 & beyond and fully owned by Tetratherix





TetramatrixTM. A proven material platform that provides differentiation across multiple indications

New chemistry from a single Platform



PNPHO



PPHO



PH2NO

3

Technical and translational offering



Minimally Invasive



Clinically Modular



5

Products under development

- Three clinical trials
- 3 confirmed partnerships



An entirely new range of chemical entities that are built on a platform with deliberate conviction

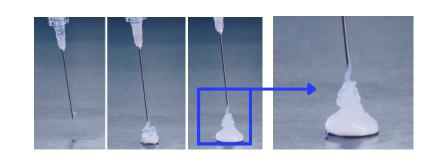


TetramatrixTM platform technology is the world's first biostealth fluid matrix

Intelligent



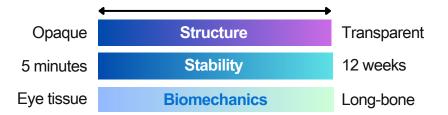
A decade of polymer programming has led to an ideal consistency & physical performance. The material is an injectable fluid to avoid causing damage to the body during its application. Using physiological temperature, material transitions into a 3D matrix.



Modular



A biomaterial platform built with unique polymer programming akin to "medical Lego" to form implantable products to solve a wide range of clinical problems. The platform has built products for 7 anatomical sites with 3 chemical entities derived from the core technology.

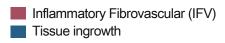


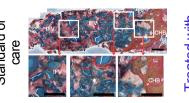
Biomimetic

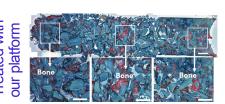


The matrix has similar water content and mechanical properties to natural tissue, and therefore is impervious to the body, helping heal injuries or physically manipulating the body during surgical interventions.

Human Histological Evaluation





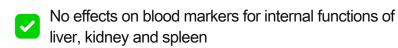


Safe



Due to the unique polymer programming of the material, once the matrix serves its purpose, the material gradually and safely bioresorbs in the body with no impact locally or systemically. Biocompatibility data from multiple clinical trials show no impact to internal organs or blood markers.

Preclinical studies (equivalent to 5 litre product administration)



Fully resorbed and excreted from the body

1



Why does the world need Tetratherix?

The evolving dynamics of the global healthcare system is demanding innovative and cost-effective biomaterials

Trends in healthcare delivery

How does Tetramatrix address the problem?



Rising patient expectations

Patients are demanding increasingly higher quality of care, with a particular focus on reducing recovery times and lowering risk of complications (e.g. infection, blood loss, pain) - which is also a driver of increasing healthcare costs for patients and payers.



Minimally invasive delivery Water-based solution injected through fine gauge needle



Safe, biocompatible and bioresorbable No foreign body reaction upon application



Need for cost-effective, decentralised care

Increasing global healthcare spending and demand for healthcare services is necessitating investment in cost-effective tools and treatments, including those that can be delivered outside a traditional hospital setting, to minimise burden on the healthcare system



Low cost, scalable production

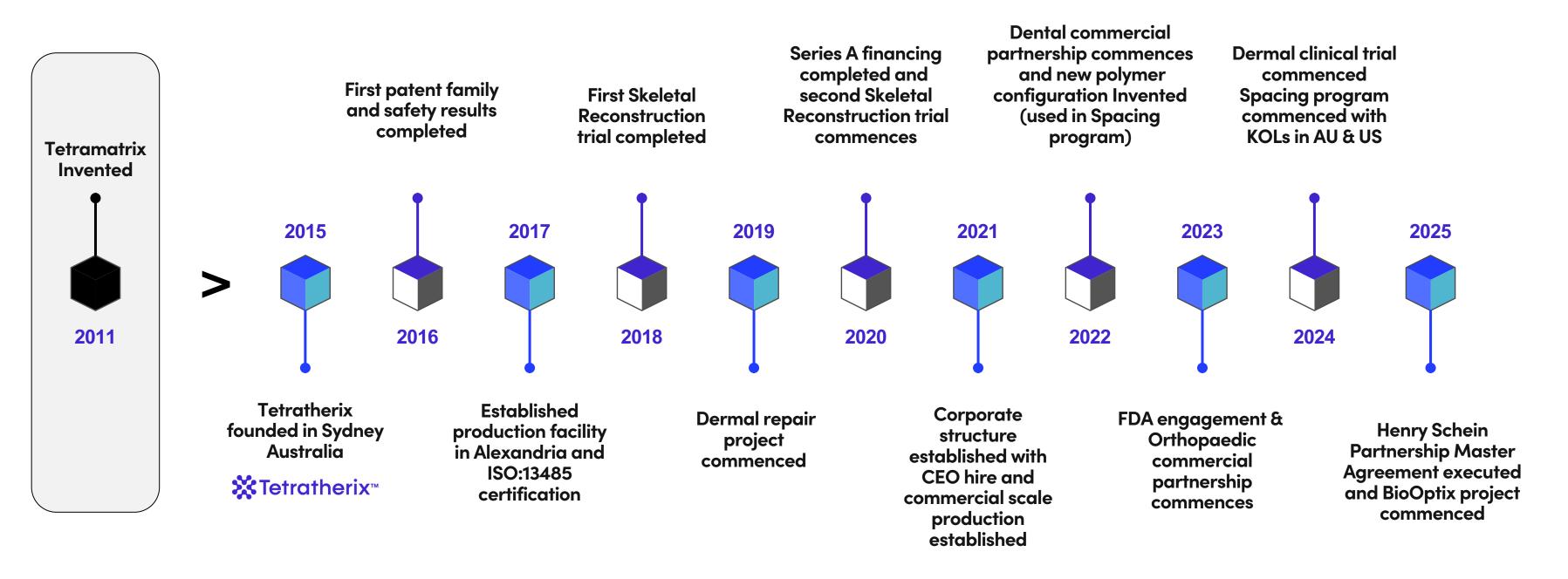
Non-labour intensive, low cost and readily available materials



Seamlessly integrated into existing workflows Delivered via needle with no additional equipment required

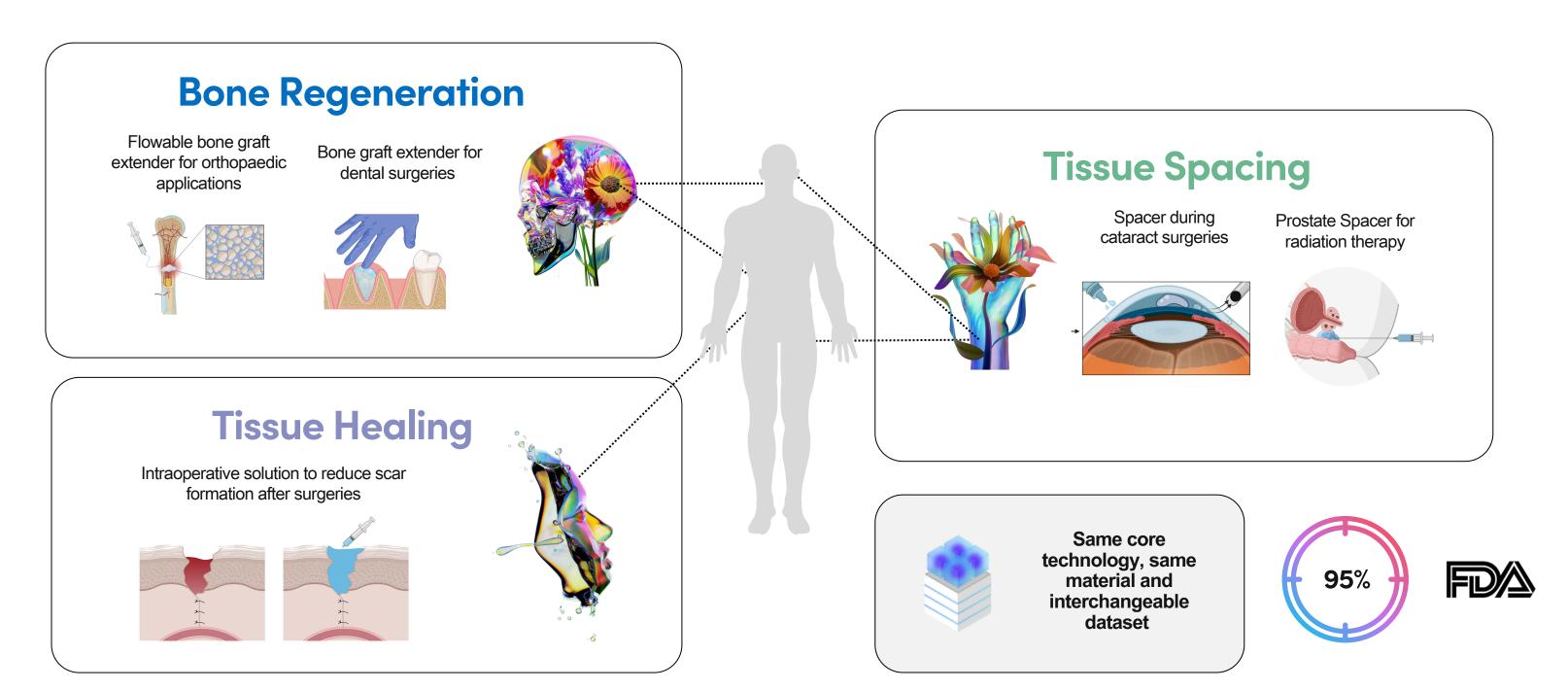


Company History





Derived products from the TetramatrixTM platform technology span 3 large franchises and significant near-term commercial opportunities



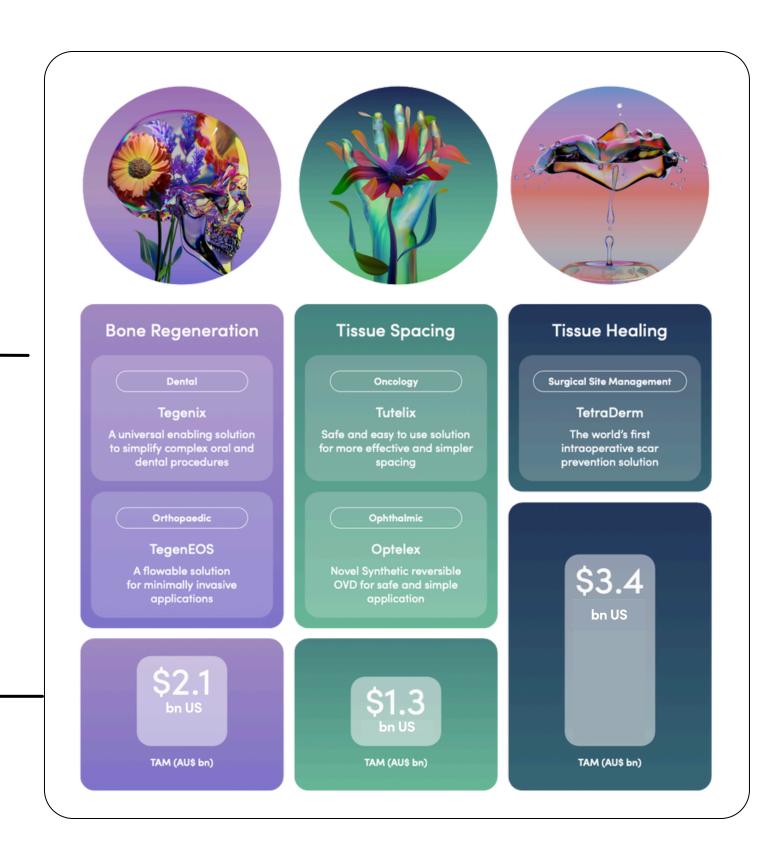


15 years and 14 programs have helped define 5 initial products on the Tetramatrix[™] platform technology



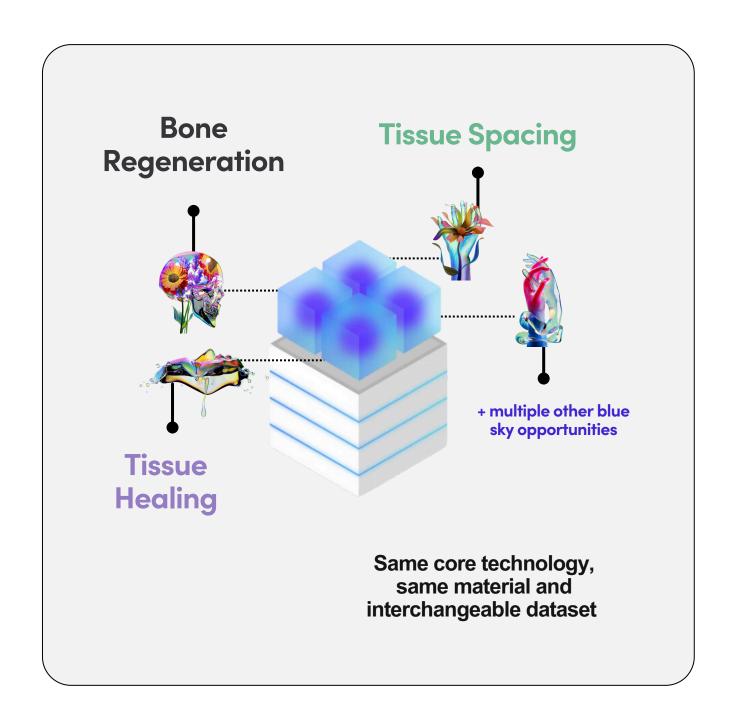
An initial addressable combined market of US\$6.7B







Different clinical needs. Different market segments. Single TetramatrixTM platform technology.

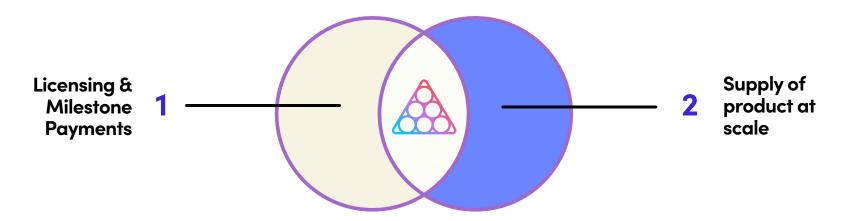


Think of us like a software platform business...

The Tetramatrix biomaterial is our **core 'platform' technology**. We have completed the foundational safety, efficacy and manufacturing work which is transferable across different clinical applications of the platform.

We **license the IP** in a specific field to a leader in the segment. These partners complete the "final 20%" to turn the application into a clinical product. We then manufacture the product and supply at attractive unit economics to our industry leading partners who distribute it though their sales channels.

We enjoy periodic milestone payments as well as a steady flow of income from product supply - a dual revenue source brings the best of both worlds.





A capital light model with compounding revenue

While we build our own production and R&D infrastructure, we avoid committing capital on establishing multiple sales & distribution networks by utilising the capabilities of our partners' established teams.

Importantly, we operate in unison with our partners. Our high margin revenue compounds over time by adding new partnerships and launching new products.

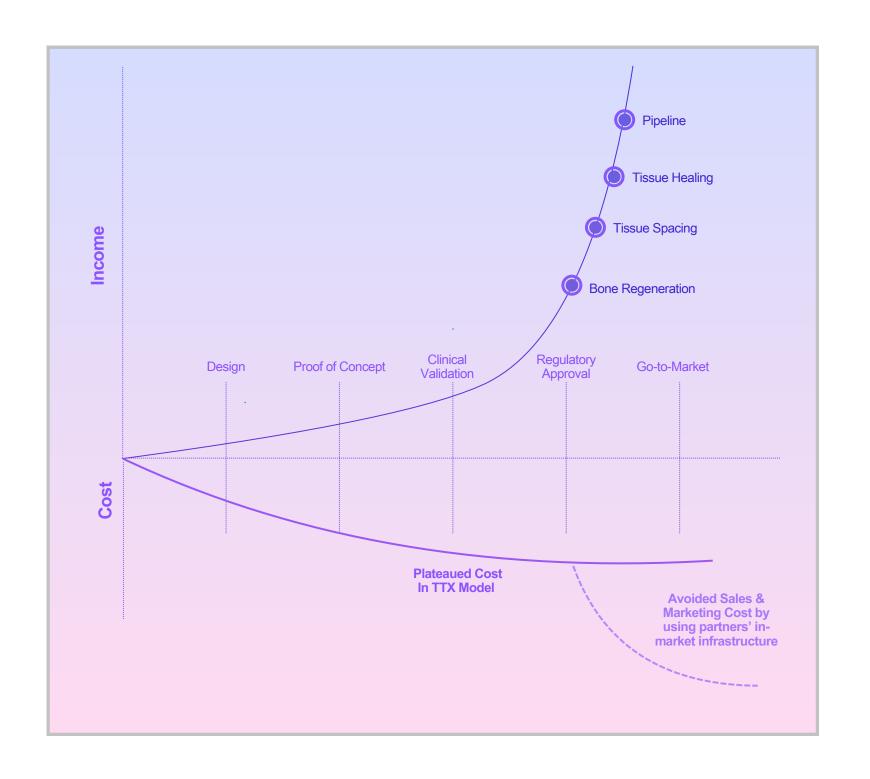
Similar to a 'build-to-buy' model, we partner differently to, and at earlier stages than, traditional medtech companies. Therefore, our return profile is different and not built on expectations of an eventual acquisition.

"Tetratherix is software platform thinking applied to the vast opportunity for smart medicines. I'm excited about the many applications that can be delivered over time and the impact they will have to millions of people."

Rod Drury
Xero Founder & Tetratherix Investor









A partnership model built on long term trust & deliberate pathway to market

Our Partnership model



X Tetratherix™

Tetratherix uses
foundational Tetramatrix™
data to conduct proof-ofconcept and pilot trials,
aligning endpoints and
assessments with partners
throughout the process

STAGE 2 Transition to Market



PARTNER

Tetratherix and its partner co-develop regulatory, reimbursement, and launch strategies.
Alignment on product form, branding, and short-term manufacturing underpins regulatory submission and market entry

STAGE 3 In Market

PARTNER

With medtech launch experience, the partner designs a roll-out strategy focused on key regions and clinical marketing to ensure a steady, sustainable market entry

A true 'win-win' for TTX and its partners

Staged partnership built on tight collaboration that limits risk and builds long-term sustainability

Reducing Risk with Diversified Franchises



By having multiple shots on goal across a range of market and partner opportunities, we reduce the regulatory, technical and commercial risk inherent to our peers in the medical technology industry.

Faster Path to Free Cash Flow for TTX



We deliberately utilise our partners' global infrastructure for sales and distribution. This creates a capital-light cost structure upon which sits rapidly compounding revenue.

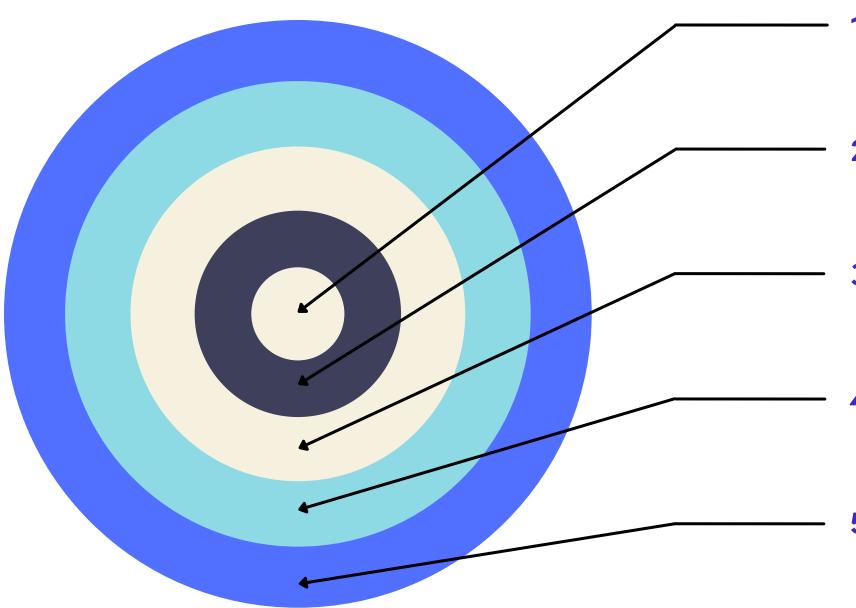
Enabling Cost-Effective R&D for Partners



Early integration of the TetramatrixTM platform technology into partners' internal development programs reduces their R&D inertia and reliance on late-stage M&A to meet their customers' needs.



Strategic Commercialisation built on decades of global launches. It's in our DNA.



1 Commercial Viability Assessment as a first step

We determine commercial reality of the program including unit economics, reimbursement, ASC vs Hospital program as well as gross margin splits for all stakeholders. Not simply a great science experiment.

Partnership Confirmation after many years of diligence

We ensure our partner of choice is motivated, aligned and the right choice on the ground. Focus of strategic portfolio not only now but over the next 5+ years. This can take time but its worth it.

3 US Seeding Program following 510k clearance

Our products have to introduced into the market effectively to ensure sustainable and effective long term growth. This includes careful selection of a small number of US tier 1 KOLs who provide clinical feedback and controlled usage in early stages of US launch.

4 US Market Expansion (P2P)

Following a 1-2 year seeding period, we work with our selected partner to expand into tier 2 KOLs via careful peer-to-peer education & marketing with inclusion of messaging tips and advice from seeding KOLs. Organic US growth follows.

Geographical Expansion on the back of refining innovation uptake

With US tier 1 and tier 2 usage under our belts, we utilise this loyal and leading group of physicians to collect US data for CE mark, Asia and Sth American expansion - with these markets adopting our seeding approach where suitable.



Not simply a science experiment. TetramatrixTM provides real world economics.

	Bone Regeneration	Tissue Spacing	Tissue Healing
ТАМ	US\$3.4bn	US\$1.3bn	US\$2.1bn
Payer	Patient out- of-pocket	Reimbursed (US)	Patient out- of-pocket
Expected GM¹ for Partner	70-80%	~80-90%	~70-80%
Expected GM for TTX	60-70%	~75-85%	~60-70%



We verify the final end user pricing and the gross margin targets for our strategic partners before progressing

1. Weighted Blended Gross Margin



Advance manufacturing and operations

We have established advanced manufacturing in Sydney. To meet growing demand, a new facility is planned to be commissioned in 2026, with a modular design to allow us to quickly scale as needed.

Our manufacturing process is:

Highly Scalable with supply chain security

- Our manufacturing process is design in a 'POD' framework whereby every POD can operate independently
- The POD has a foot print of 200 sqm, and can be replicated multiple times without our facility floorspace
- Allows multiple X increases in the production capacity in a fast and controlled manner

De-risked



- We have previously increased our production yield per batch from laboratory scale (10 g per batch) to a commercial scale (1.2kg) through multiple cycles of process optimisation and scale up
- The production process is not labour-intensive and requires only off-the-shelf equipment
- All raw materials are catalogue products and easily accessible from multiple suppliers, de-risking any supply chain risk
- A new, larger site in Alexandria, Sydney has been identified with negotiations ongoing for a long-term lease

Externally Validated



- Production blueprint validated for further upscale with an external European CMO
- Our process has been governed by an EN ISO13485 certified quality management system since 2017 successfully completing three re-certification audits

Manufacturing POD







Relentless execution with Regular newsflow in first 12 months post-listing

Bone Regeneration	DENTAL		Henry Schein Suppl Licensing agreem execution		m the FDA Submission	FDA Clearance
	ORTHOPAEDIC		Master Partr Agreeme execution	ent agreement ex	ELIA SUBMICCIÓN	FDA Clearance
Tissue Spacing	ONCOLOGY	FDA Pre-submission/ Regulatory Pathway Defined	Tutelix First In Human Cohort 1	Quality Agreement execution	US/AU Tutelix Prima Capital Raise Point for Co	
Tissue Healing	OPHTHALMIC	BioOptix Licensing Agreement	Strategic global partnership	US/ AU Capital Raise	Preclinical Data Read-out	FDA Pre-submission & Regulatory Pathway
	SURGICAL SITE MGMT	Cohort 2 Commenced	Year 1 Follow up for Cohort 1 results published	Primary End-poi Cohort 2	nt FDA Pre-submission	TetraDerm Cohort 3 Initiation (major surgeries)

FY2026 FY2027

