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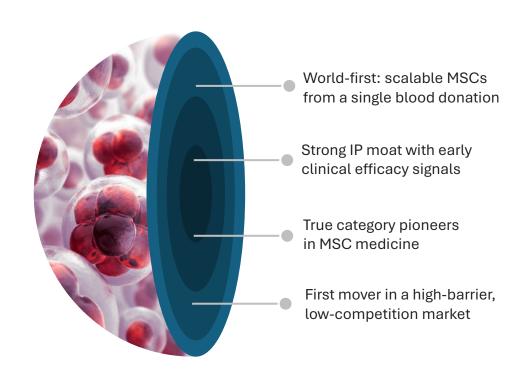
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Cynata Therapeutics

Next-generation Mesenchymal Stromal Cells (MSCs)¹

- Commercialising Cymerus™, a novel platform producing MSCs from a *single* blood donation once, at scale
- MSCs are powerful immune-modulating and tissue-repairing cells, but naturally scarce in the human body
- Traditional manufacturing relies on continuously finding donors and faces inconsistency, potency loss, and scaling limits
- Cynata solves the MSC production bottleneck, unlocking full therapeutic and commercial potential
- Four clinical programs underway, including advanced **Phase 2 & 3 trials** across major indications

Front-Runner Status



Capital Structure

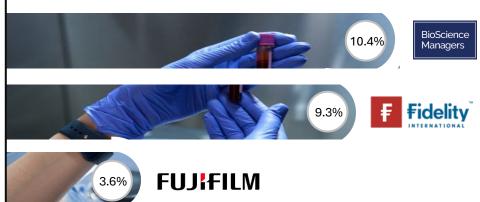
Strong balance sheet. Positioned for upside.

- **Backed by institutions:** Fidelity, BioScience Managers, and Fujifilm among top holders
- **Funded through milestones**: Cash runway secured to mid-2026 through landmark clinical readouts
- **Attractive valuation:** Current market cap does not reflect value of advanced clinical pipeline and near-term catalysts
- Tightly held register: Top 20 own ~47%

Financial Information

Share price (6 August 2025)	A\$0.175
Shares on issue	~225m
Market capitalisation	~A\$39.5m
Cash ¹	~ A\$5m

Top Holders



MSCs: Nature's Repair Cells

Powerful.

Poised for clinical impact.

What are MSCs?

- Naturally found in small quantities in the body
- Regulate and support immune system and reduce inflammation¹
- Support tissue repair¹

Why They Matter

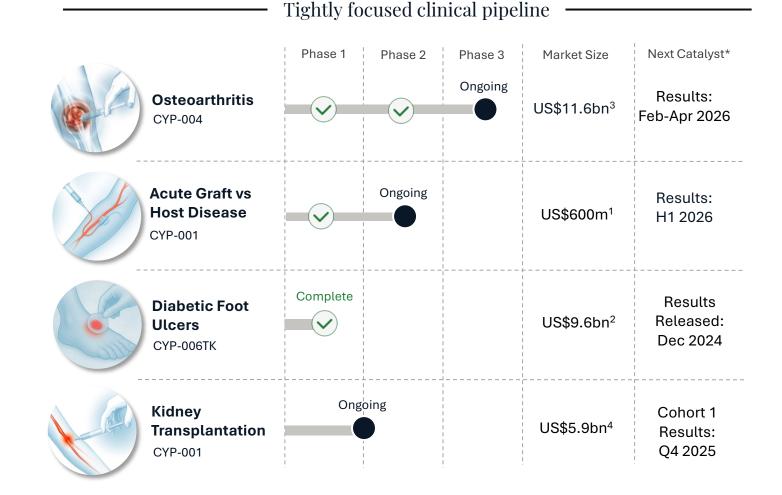
- Target root causes of many diseases, not just symptoms
- Safe and well-tolerated in human trials
- Immune-privileged no donor matching needed

Where They're Going

- 1,800+ clinical trials initiated globally
- Addressing high-burden diseases with limited treatment options
- Entering a phase of clinical maturity & market readiness

Cynata Is At An Inflection Point

- Advancing four clinical programs to address critical treatment gaps
- Company- (and industry-) defining data readouts this financial year
- Manufacturing process established and ready to scale
- Positioned well for global licensing and joint venture deals



^{1.} Global Graft versus Host Disease Market 2019-2029 (Reflects forecast market in 2026)

^{2.} Zion Market Research, 2019 (represents global treatment market in 2025)

^{3.} Persistence Market Research 2018 research report: "Osteoarthritis Treatment Market: Global Industry Analysis (2012-2016) and Forecast (2017-2025) (Reflect OA market by 2025);

[.] Organ Transplant Immunosuppressant Drugs Market in 2026, Grand View Research, Inc., 2019

^{*} Timing of events is approximate, based on the Company's information as at the date of this presentation, and subject to change. Timing refers to calendar years.

Osteoarthritis

There's no cure. Only symptom relief or invasive surgery.



- Degenerative joint disease
- Causes pain, stiffness, and reduced mobility
- \$486 billion economic burden in the U.S. alone¹



- Affects 600 million globally with no curative treatment²
- Current options don't address cartilage degradation
- Surgery is costly, invasive, and not suitable for all



- Early phase 3rd party studies with traditional MSCs have shown:
 - Strong safety
 - Pain relief
 - Improved joint function
- Third party MSC data in OA, and Cymerus™ MSC track record supported our direct phase 3 entry



- ~320-patient study using Cymerus™ MSCs
- Investigating change in pain and cartilage thickness
- Final results anticipated February April 2026

Managed by
University of Sydney

Funded by NHMRC

cvnata.com

Acute Graft vs Host Disease (aGvHD)

When life-saving transplants become life-threatening.









- Life threatening disease post-bone marrow transplant
- Transplanted cells attack recipient's tissues as "foreign"
- Affects up to 50% of patients and damages organs^{1,2}

- Steroids fail in about 50% of cases (first line treatment)³
- <20% 2-year survival rate in patients who fail to respond to steroids³
- Current drugs have poor efficacy and safety

87% Improved by at least 1 grade

53% Showed no signs of aGvHD

60% Alive after two years

Serious adverse events related to CYP-001

- Global trial underway (60 patients)
- Investigating overall response rate at day 28
- Results expected H1 CY26

Regulatory Support

FDA FOR Orphan Drug Control Designation F

FDA Cleared IND EU IMPD Cleared

Cynata's Broader Disease Pipeline

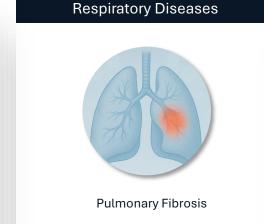
Current Additional Clinical Programs

Strong Pre-Clinical Data



Kidney Transplantation (Phase 1/2 – Ongoing)

- Cynata's Phase 1/2 trial underway (Netherlands) with Cymerus[™] MSCs
- Investigating reduced reliance on immunosuppressants
- Results from Cohort 1 expected H2 2025







Diabetic Foot Ulcers (DFU) (Phase 1 – Successful)

- Phase 1 trial of Cymerus[™] MSCs showed 83.6% wound reduction vs. 47.8% with standard care
- · Safe, well tolerated
- Exploring next development steps

Cardiovascular Diseases



Artery Disease



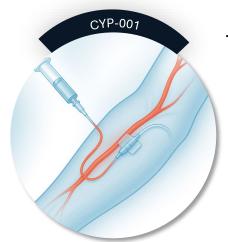
Cynata Products Deliver Promise

Built to meet critical needs.

Osteoarthritis

- Delay or avoid knee replacement surgery
- Improve mobility, daily function & quality of life
- Reduce long-term economic healthcare burden





Acute GvHD

- Safer, more effective therapy that improves survival rates
- Reduce reliance on steroids and immunosuppressants
- · Support faster recovery

Diabetic Food Ulcers

- Faster wound healing
- Reduce risk of amputation
- Improve independence and outcomes





Kidney Transplantation

- Reduce long-term drug toxicity
- Improves graft survival and function
- Enhance quality of life post-transplant



Conventional MSC Manufacturing

Incredible science. Unrealised potential due to manufacturing limitations.



Steps Repeat For Each MSC Batch

MSCs sourced from adult tissue (e.g. bone marrow)

Drawbacks:

- Many donors needed
- Donor variability
- Expensive, complex logistics

MSCs must be separated from other cells

Drawbacks:

- Low yield
- Inconsistent activity, varying from donor to donor

Cells grown in lab to reach treatment scale

Drawbacks:

- Potency loss with over-expansion
- Functional decline

Expanded MSCs used to treat small number of patients

Drawbacks:

· Quality inconsistency

CymerusTM Solves the MSC Bottleneck

One donor. One time. A platform built to scale.



We don't need to repeat steps 1&2 ever again



Blood sample from one healthy adult - just once

Advantages:

- No ongoing donor sourcing
- Simplifies supply chain
- Cuts cost and eliminates donor variability

Blood cells reprogrammed into Induced Pluripotent Stem Cells (iPSCs), then stored in a master cell bank.

Advantages:

- Permanent source
- Consistent starting material
- Unlimited expansion

CYP's iPSC cell bank is large enough for commercialisation of multiple products at global scale iPSCs directed into MSCs via novel Cymerus™ process

Advantages:

- Consistent, potent MSCs
- Scalable and reproducible
- Enables off-the-shelf global therapies at scale

Step 3 is endlessly repeatable

What Our Manufacturing Breakthrough Unlocks

From scientific first to leadership in MSC medicine.

- Only company able to produce clinicalgrade MSCs from iPSCs at scale
- Eliminates decades-old barrier to scalable MSC commercialisation

Positions Cynata as the global standard for MSC manufacturing

Fit for mainstream use in multibillion-dollar markets

Enables licensing and JV deals for off-the-shelf therapies

Sustainable competitive edge with strong IP moat

The Next 9-12 Months Will Define Cynata

Upcoming Landmark Readouts ———

Osteoarthritis
CYP-004

Market Size

US\$11.6bn²

Phase 3 Results:
Feb-Apr 2026
(Efficacy & safety)

Phase 2 Results:
H1 2026
(Efficacy & safety)

Upcoming Preliminary Readout



Kidney TransplantationCYP-001

US\$5.9bn³

Cohort 1 Phase 1/2 Results: Q4 2025 Later stage results could become a strategic trigger

Licensing / Partnering

- Positive Phase 2 & 3 results can trigger regional or indication-specific deals
- All our indications are attractive licensing targets with clear market needs
- Partnership revenue can help fund future trials without equity dilution

M&A Potential

- Compelling data + scalable platform = highly strategic assets
- Strong potential for synergies with other MSC technologies in the market

Joint Development / Pharma Alliances

- Shared risk models appeal to global pharma with aligned pipelines
- Allows Cynata to enter new markets with global reach and local execution

All commercial activities referenced are potential future options only. No agreements have been made.

- 1. Global Graft versus Host Disease Market 2019-2029 (Reflects forecast market in 2026)
- 2. Persistence Market Research 2018 research report: "Osteoarthritis Treatment Market: Global Industry Analysis (2012-2016) and Forecast (2017-2025) (Reflect OA market by 2025);
- 3. Organ Transplant Immunosuppressant Drugs Market in 2026, Grand View Research, Inc., 2019

Go-To-Market Strategy

Lean. Scalable. Capital-Efficient.

- Billions already spent by MSC peers validating market and use-cases
- 2 Cynata to benefit from proof of concepts delivered by peers, reducing capital burn
- Cynata could leapfrog with higherperforming, scalable MSC products

- Recognised global manufacturing partner with established capacity
- Capital-light model with limited infostructure investment
- Revenue via licensing, JVs, and deals focus remains on science and pipeline

The Investment Case

Why Cynata? Why Now?

Solves the MSC manufacturing bottleneck with singledonor scalability



Lead asset backed by FDA Orphan Drug designation and solid IP protection



Fully funded through major clinical milestones



