

Investor Webinar

Melbourne, Australia; 22 June 2026: Cynata Therapeutics Limited (ASX: “CYP”, “Cynata”, or the “Company”), a clinical-stage biotechnology company specialising in cell therapeutics, reminds shareholders that an investor webinar will be held today, Monday 22 June 2026, at 8:30am AEST, hosted by CEO and Managing Director, Dr Kilian Kelly, and Chair, Dr Geoff Brooke. Please visit the following link to register: <https://cynata.com/webinars/rvJo2r-cynata-investor-webinar>

A copy of the presentation to be delivered during the webinar is attached to this announcement.

After the webinar is complete, a recording will be available on the Company’s [InvestorHub](#) portal.

-ENDS-

Authorised for release by Dr Kilian Kelly, CEO & Managing Director

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About Cynata Therapeutics (ASX: CYP)

Cynata Therapeutics Limited (ASX: CYP) is an Australian clinical-stage stem cell and regenerative medicine company focused on the development of therapies based on Cymerus™, a proprietary therapeutic stem cell platform technology. Cymerus™ overcomes the challenges and limitations of conventional MSC production by using induced pluripotent stem cells (iPSCs) to achieve economic manufacture of cell therapy products, including mesenchymal stem cells (MSCs), at commercial scale without the necessity to obtain tissue from multiple donors on an ongoing basis, and without the complexity and product inconsistency resulting from conventional methods.

Cynata Therapeutics encourages all current investors to go paperless by registering their details with the designated registry service provider, [Automic Group](#).



Investor Webinar: Results Update

June 2026

(ASX:CYP)

Important Information

Summary information

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Two Pivotal Readouts:

A surprising and disappointing outcome

- This has been a very difficult few days for the Company
- Cynata reported top-line results from two pivotal trials
 - the Phase 2 trial of CYP-001 in acute Graft versus Host Disease (aGvHD)
 - and the Phase 3 SCULpTOR trial of CYP-004 in knee osteoarthritis (OA).
- Neither trial met its primary endpoint. There were no statistically significant differences between the active and control groups on the primary or key secondary measures in either study.
- The aGvHD outcome was particularly unexpected. Our Phase 1 trial had delivered strongly positive efficacy signals, giving us firm grounds for confidence that were not reproduced at Phase 2.
- In OA, the active group achieved the pain reduction, but an unusually strong placebo response prevented a treatment effect from being demonstrated.
- Both products were again confirmed to be safe and well tolerated, with adverse event profiles comparable to control.



OA: Phase 3 Results

Key findings:

- There were no safety concerns identified, with a similar adverse event profile between groups.
- There were no statistically significant differences between the active and control groups in either of the co-primary endpoints:
 - The proportion of participants reaching the patient-acceptable symptom state (PASS) threshold for knee pain at 24 months was 51.7% in the active group compared to 48.1% in the control group ($p=0.5907$).
 - There was an average loss of central medial femorotibial (cMFT) cartilage thickness of 0.27mm in the active group compared to 0.21mm in the control group ($p=0.1453$).
- There was a substantial and durable reduction in knee pain intensity at all timepoints from 3-24 months compared to baseline (secondary endpoint), with an average reduction in pain score of 26.8/100 in the active group and 25.3/100 in the control group after 24 months. However, there were no significant differences between groups at any timepoint.
- Analysis of additional outcome measures is ongoing and will be reported in due course



aGvHD: Phase 2 Results

Key Findings:

- There were no safety concerns identified, with a similar adverse event profile between groups.
- There were no significant differences between groups in the primary or key secondary endpoints:

Primary endpoint	Active	Control	P Value ¹
Day 28 Overall Response Rate (ORR)	57.7%	54.8%	>0.9999
Secondary endpoints			
Day 28 Complete Response Rate (CRR)	38.5%	35.5%	>0.9999
Day 100 ORR	23.1%	32.3%	0.5581
Day 100 CRR	19.2%	29.0%	0.5392
Durable ORR (patients with OR at each of Day 28, Day 60 and Day 100)	19.2%	32.3%	0.3681
Day 100 overall survival rate	88.1%	82.3%	0.5137





Thank You.

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