



10 November 2025

Sydney, Australia

Investor Presentation and CEO AGM Presentation

Nyrada Inc (ASX:NYR), a clinical-stage biotechnology company developing Transient Receptor Potential Canonical (TRPC) ion channel inhibitors to treat a range of medical conditions provides the enclosed presentation slides in connection to investor meetings to be held in the week commencing Monday 10 November 2025 and for its 12 November 2025 Annual General Meeting CEO presentation.

– ENDS –

About Nyrada Inc.

Nyrada Inc. is a clinical-stage biotechnology company focused on the discovery and development of innovative small-molecule therapies, specifically targeting Transient Receptor Potential Canonical (TRPC) ion channels. The company's lead candidate, Xolatryp™, has shown efficacy in both cardioprotection and neuroprotection, and has just completed a first-in-human Phase I clinical trial. Nyrada Inc. (ARBN 625 401 818) is incorporated in Delaware, US, with limited liability for its stockholders.

www.nyrada.com

Authorised by John Moore, Non-Executive Chair, on behalf of the Board.

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AGM Presentation

James Bonnar
Managing Director and CEO

12 November 2025 | Sydney, Australia

Improving Lives, Offering Hope

ASX:NYR

Authorised by Mr. John Moore, Non-Executive Chair,
on behalf of the Board.



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Nyrada Overview

- › Clinical-stage biotech company developing TRPC ion channel inhibitors to treat a range of medical conditions
- › Lead drug candidate – Xolatryp (formerly NYR-BI03).
 - › Potent 3rd generation TRPC 3, 6 and 7 channel inhibitor
 - › Novel and well-understood mechanism of action
 - › Solid scientific foundation
 - › Preclinical efficacy across multiple therapeutic areas
- › Progressing Clinical Development
 - › Phase I completed (safety, tolerability, PK)
 - › Progressing into Phase IIa - acute myocardial infarction targeting ischemia reperfusion injury



Company Snapshot

Company Structure

ASX:NYR Overview at 6 November 2025

CDIs on Issue (m)	239.5
Share Price at close 5 Nov. 2025 (AU\$)	\$0.60
Market Capitalisation (AU\$m)	\$215.5
Options and Performance Rights (m)*	49.4
Board and Staff Holding (%)	10.3

* Excludes options to be considered at 12 November 2025 Annual General Meeting

Leadership

Nyrada Leadership

Non-Executive Chair	Mr. John Moore
Managing Director and Chief Executive Officer	Mr. James Bonnar
Chief Scientific Officer	Dr. Benny Evison
Chief Financial Officer	Mr. Cameron Jones
Scientific Advisory Board Chair	Prof. Garry Housley

Share Register – 6 November 25

Position	Holder Name	Holding	% IC
1	MR MARK AZZI	39,666,809	16.56%
2	MATT CORP WA PTY LTD	11,408,580	4.76%
3	ALTNIA HOLDINGS PTY LTD	9,921,725	4.14%
4	KYRIACO BARBER PTY LTD	7,503,106	3.13%
5	MR JOHN MOORE	7,072,756	2.95%
6	CELTIC FINANCE CORP PTY LTD	4,700,000	1.96%
7	HARLUND INVESTMENTS PTY LTD	4,150,000	1.73%
8	CELTIC CAPITAL PTE LTD	4,000,000	1.67%
9	CITICORP NOMINEES PTY LIMITED	3,592,561	1.50%
10	MS ROCHELLE SEMAAN	3,109,684	1.30%
	Total	95,125,221	39.72%
	Total issued capital - selected security class(es)	239,497,037	100.00%

Price Chart



Origins of TRPC Channel Inhibition

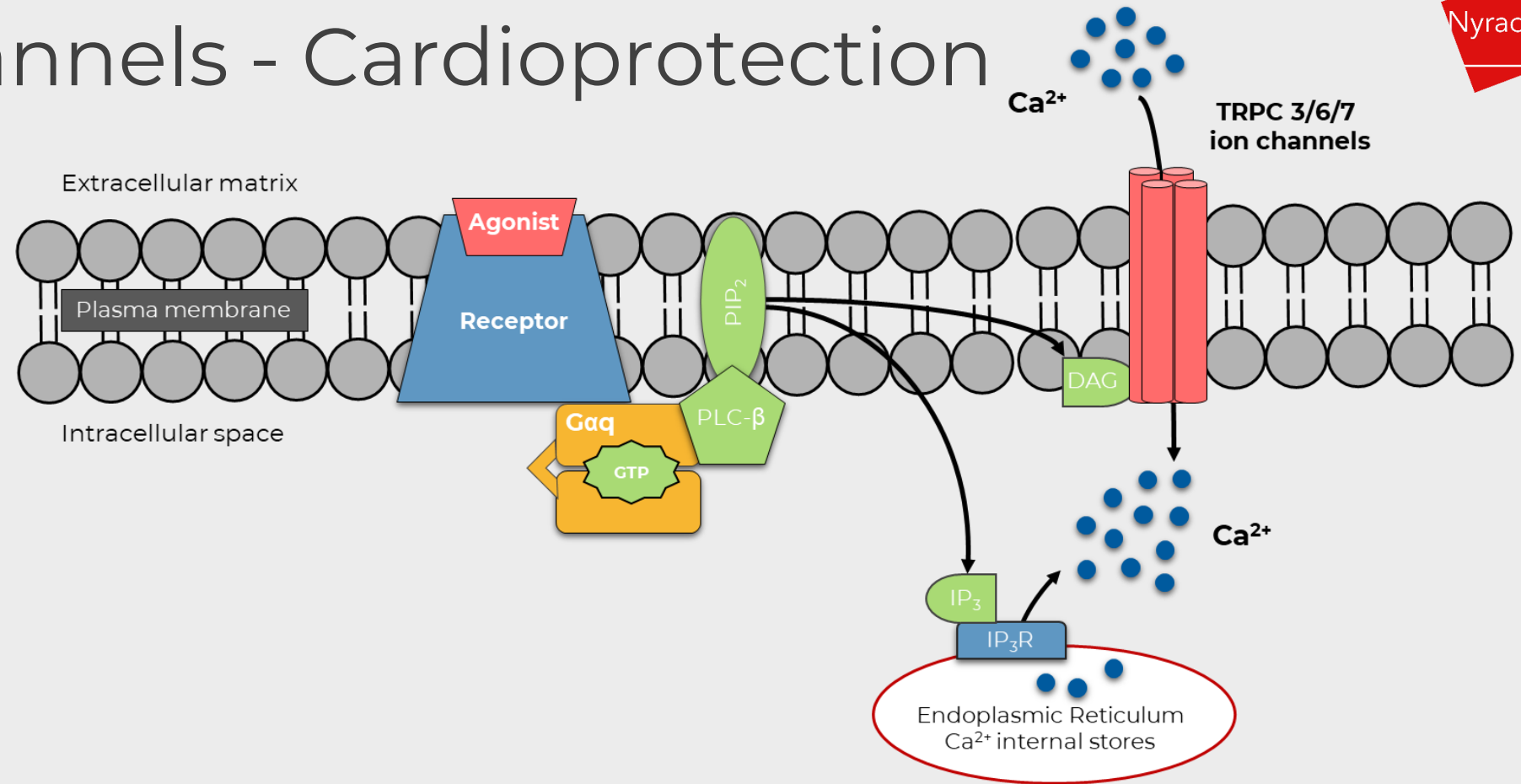
Lutz Birnbaumer



Birnbaumer's Mice



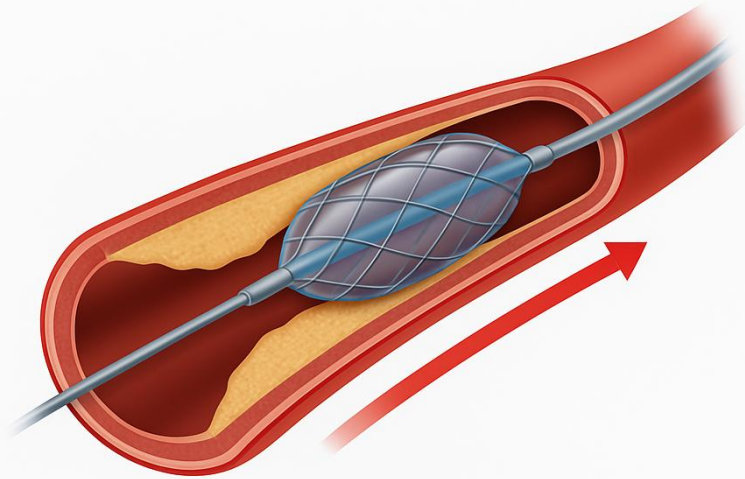
TRPC Channels - Cardioprotection



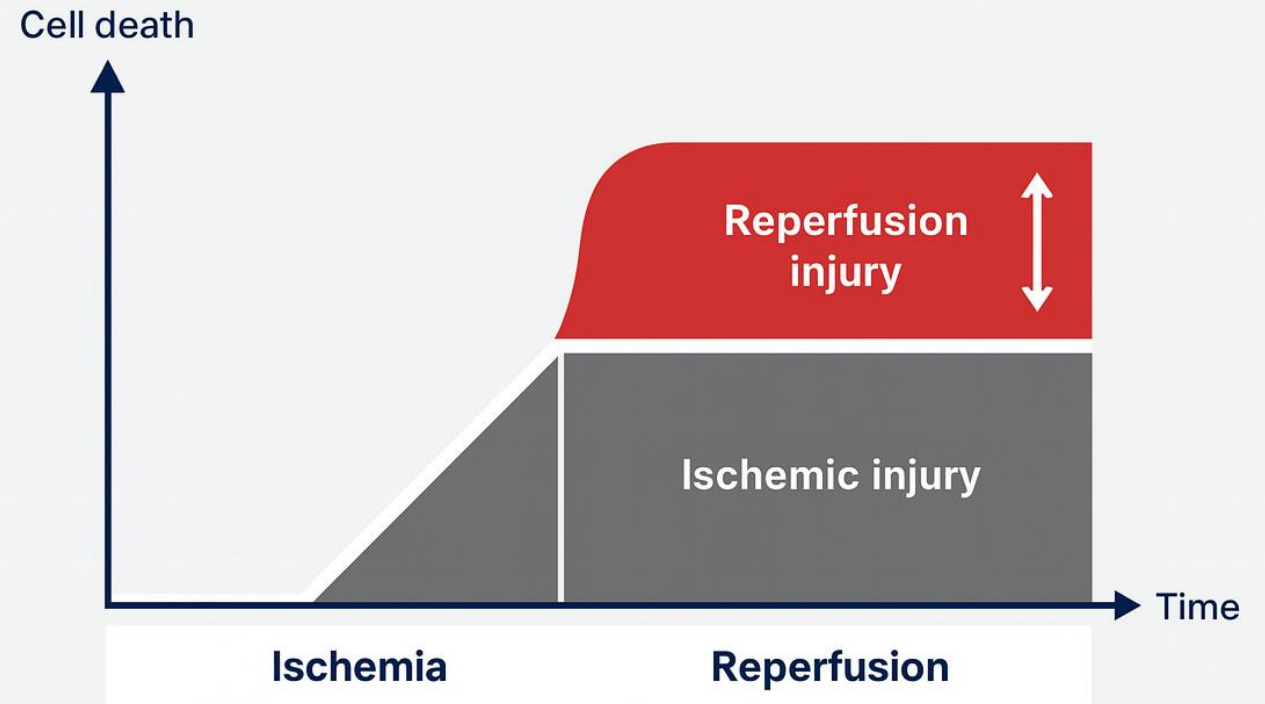
Cell Type	TRPC Activation Effect	Xolatryp Inhibition Effect
Cardiomyocytes	Calcium overload → cell death	Prevents damage, preserves function and contractility
Endothelial	Impaired repair → poor blood flow	Enhances vessel repair, reduces inflammation
Fibroblasts	Excessive scarring → stiff heart	Limits scarring, preserves flexibility

Ischemia Reperfusion – Last Cardiac Frontier

Coronary PCI (angioplasty)



Blood flow restoration



PCI (Balloon Angioplasty):

- First procedure in 1977, stents in 1994,
- Standard of care for STEMI by early 2000s

Large Market Opportunity – Myocardial Ischemia Reperfusion Injury

Globally:

~15-20 million
people suffer heart
attack annually

~15%
mortality within 30
days

No current FDA approved treatments

Effective treatment will improve patient outcomes and reduce high costs associated with long-term care of brain injury survivors.

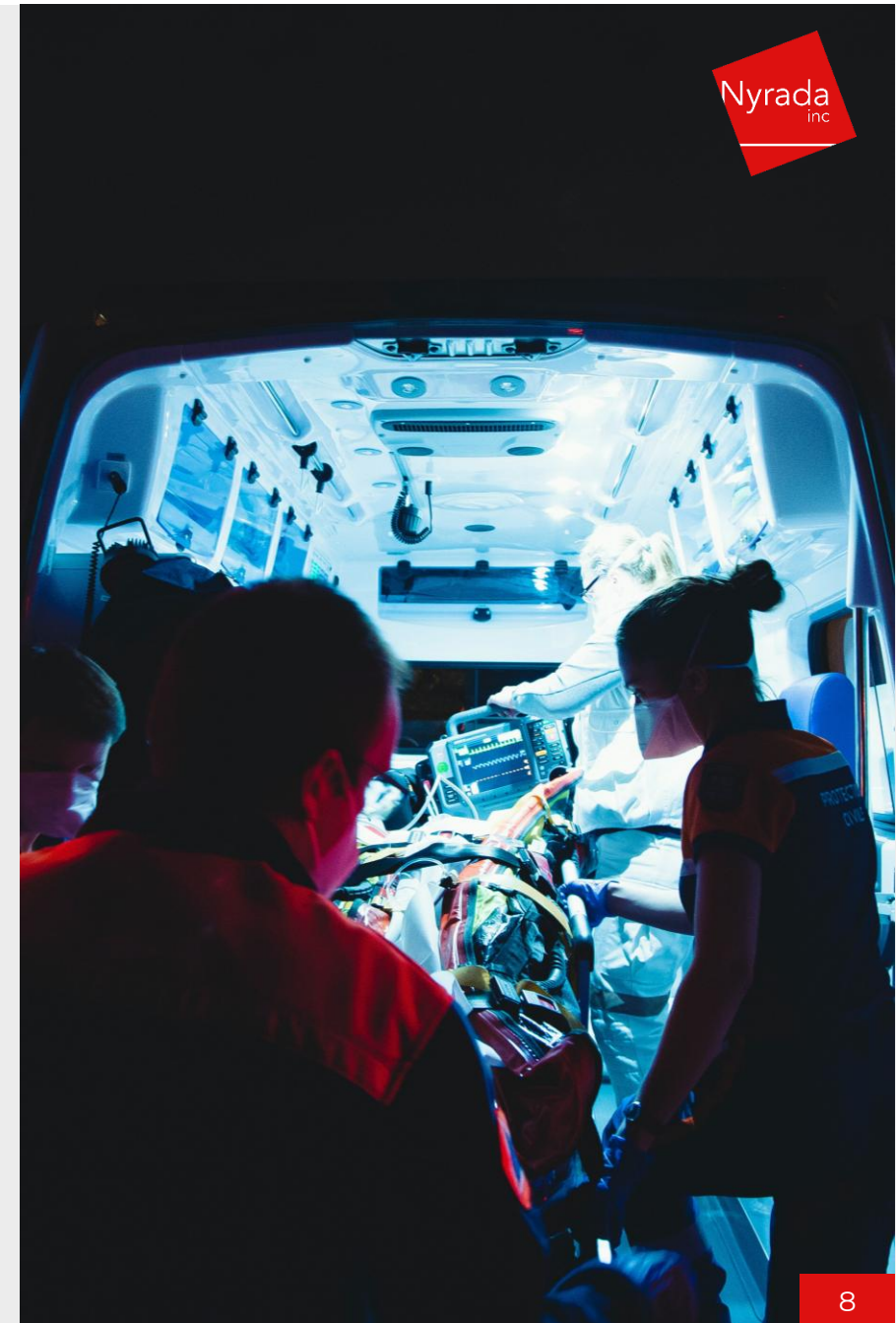
Large and growing treatment market*:

Currently
~US\$1.7 billion

Growing
~7.7% CAGR

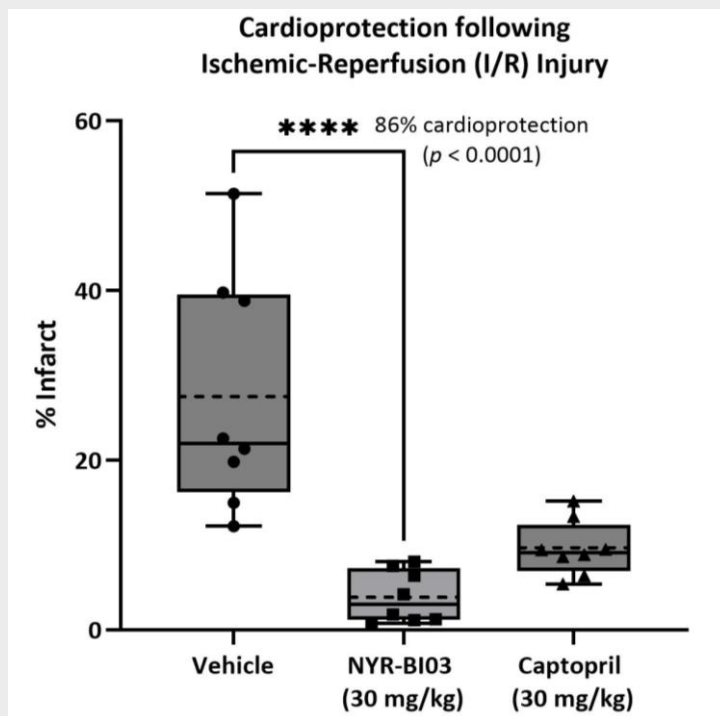
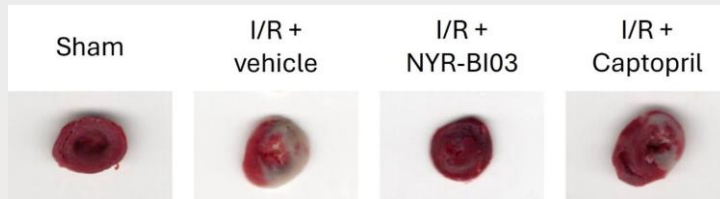
Forecast
**~US\$2.3 billion by
2029**

* Research and Markets - www.researchandmarkets.com/reports/6089883/ischemia-reperfusion-injury-market-report



Preclinical Study 1

Key Preclinical Results:



Xolatryp showed strong efficacy limiting cardiovascular damage resulting from myocardial ischemia-reperfusion (IR) injury

- **86%** Cardioprotection
- **43%** increase in left ventricular ejection fraction
- **50%** increase in fractional shortening

Key blood biomarker markers assessed

- **42%** decrease in AST levels
- **45%** decrease in LDH levels
- **32%** decrease in Troponin I

Superior efficacy compared to FDA-approved, Captopril



Preclinical Study 2

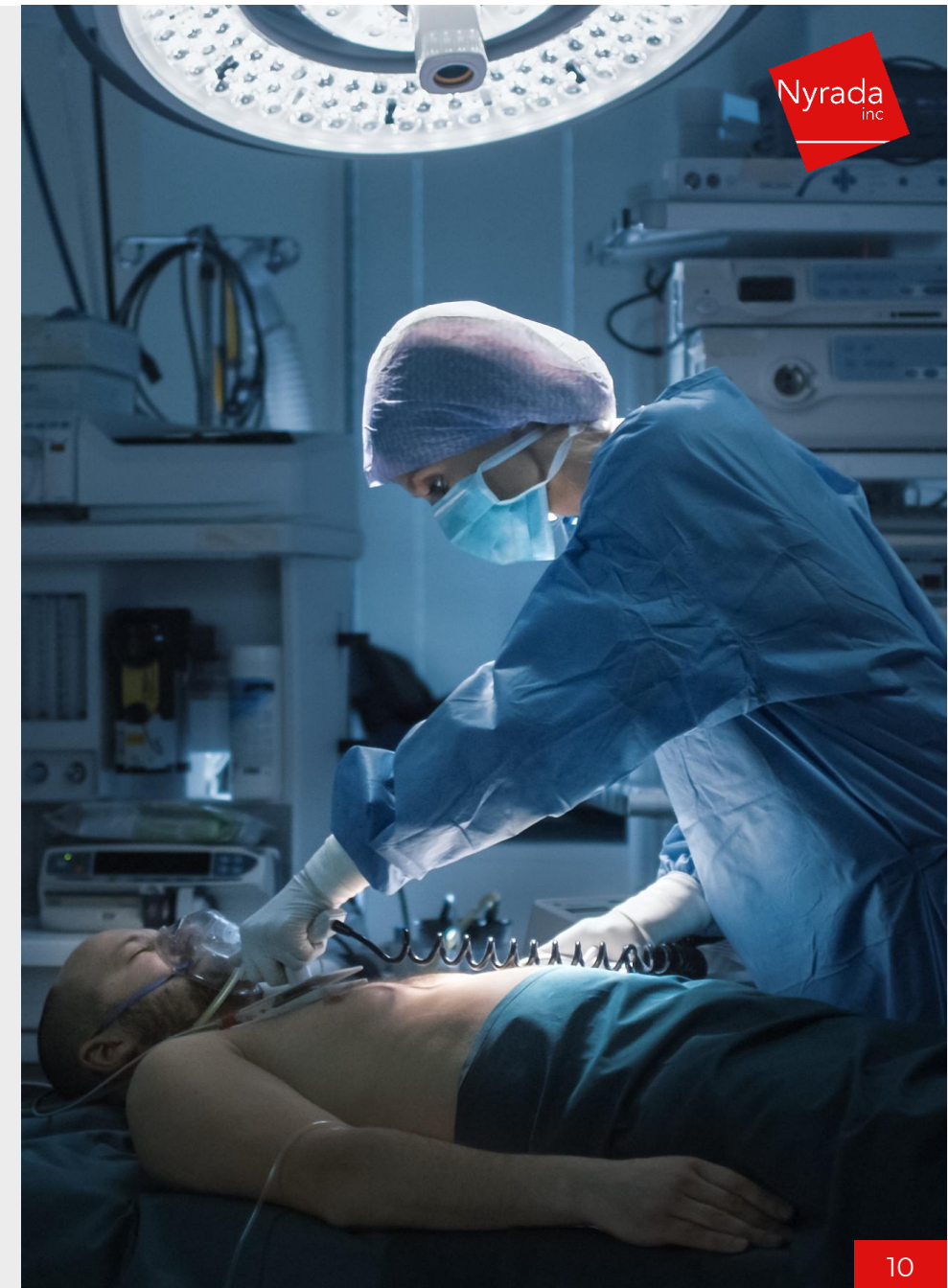
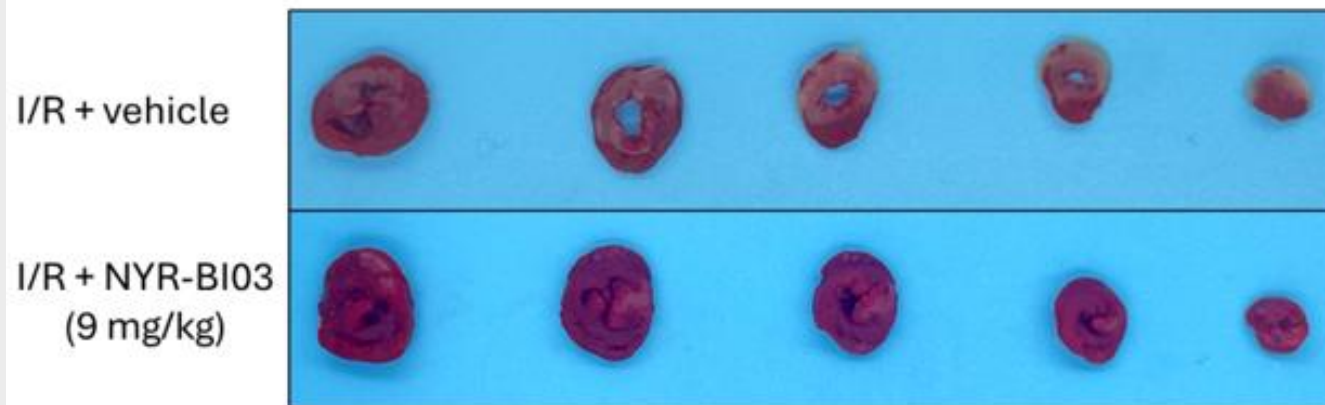
Key Preclinical Results:

Xolatryp showed strong efficacy limiting cardiovascular damage resulting from myocardial ischemia-reperfusion injury when administered as a short-duration intravenous infusion

- **42%** Cardioprotection
- **88%** decrease in arrhythmias at 1 hour
- **90%** decrease in arrhythmias at 3 hours

Key blood biomarker markers assessed

- **32%** decrease in Troponin I
- **21%** decrease in ALT levels

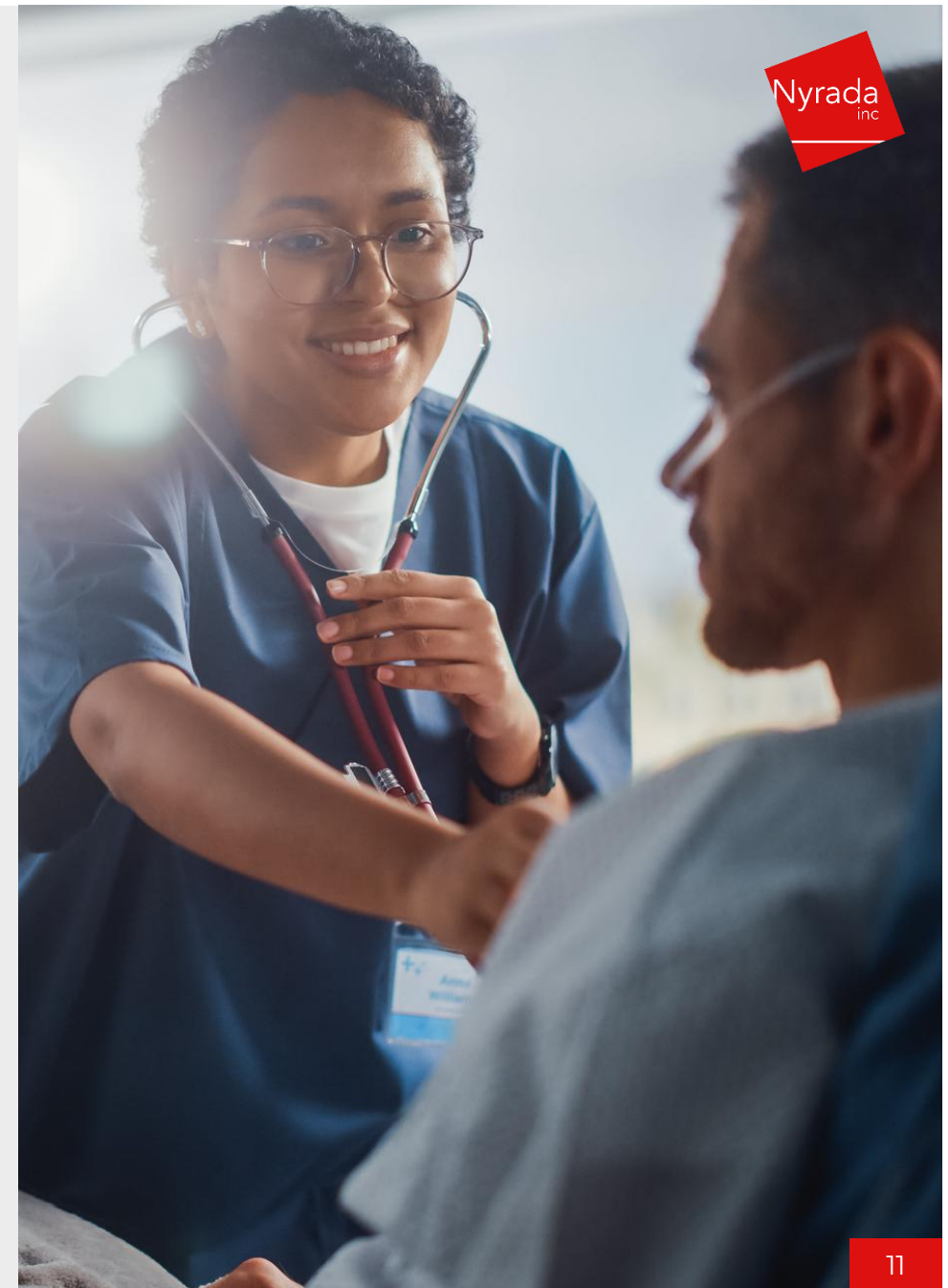
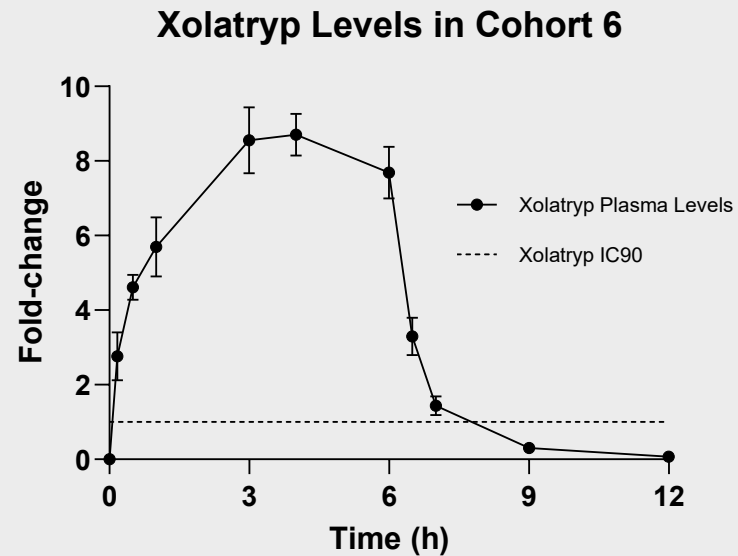


Phase I Clinical Trial

Key Results:

Xolatryp met its Primary Endpoint with all doses safe and well tolerated, with no dose-limiting, or dose-related safety issues.

- **48 healthy participants** (36 received drug, 12 received placebo).
- **Nil SAEs**
- **10 AEs**
 - All mild or moderate
 - 5 *not related* to Xolatryp, 1 *unlikely related* to Xolatryp.
 - Most frequently reported AE was headache.
- **Pharmacokinetics**
predictable with linear blood concentrations over time.
- **10 minutes** to reach therapeutic levels



Phase IIa Clinical Trial

Key Features:

Xolatryp to be assessed for safety and preliminary efficacy in patients with ST-Elevation Myocardial Infarction (STEMI) who undergo primary Percutaneous Coronary Intervention (PCI)

Primary End Point – Safety

Further End Points – Exploratory efficacy potentially including:

- Cardiac function
- Cardiac injury size
- Biomarkers including Troponin I levels
- Incidence of arrhythmias of interest

Scope (subject to change)

- 200 patients (up to approximately; placebo and drug – 1:1)
- 9 to 18 months (indicative)
- 6 sites (initially)

Key operational risk – recruitment rate impacting duration and number of sites



R&D Activities

Evolving Pipeline:

Xolatryp demonstrated preclinical efficacy:

- Myocardial ischemia reperfusion injury
- Ischemic stroke
- Moderate to severe TBI

Literature suggests **Xolatryp** may have efficacy in:

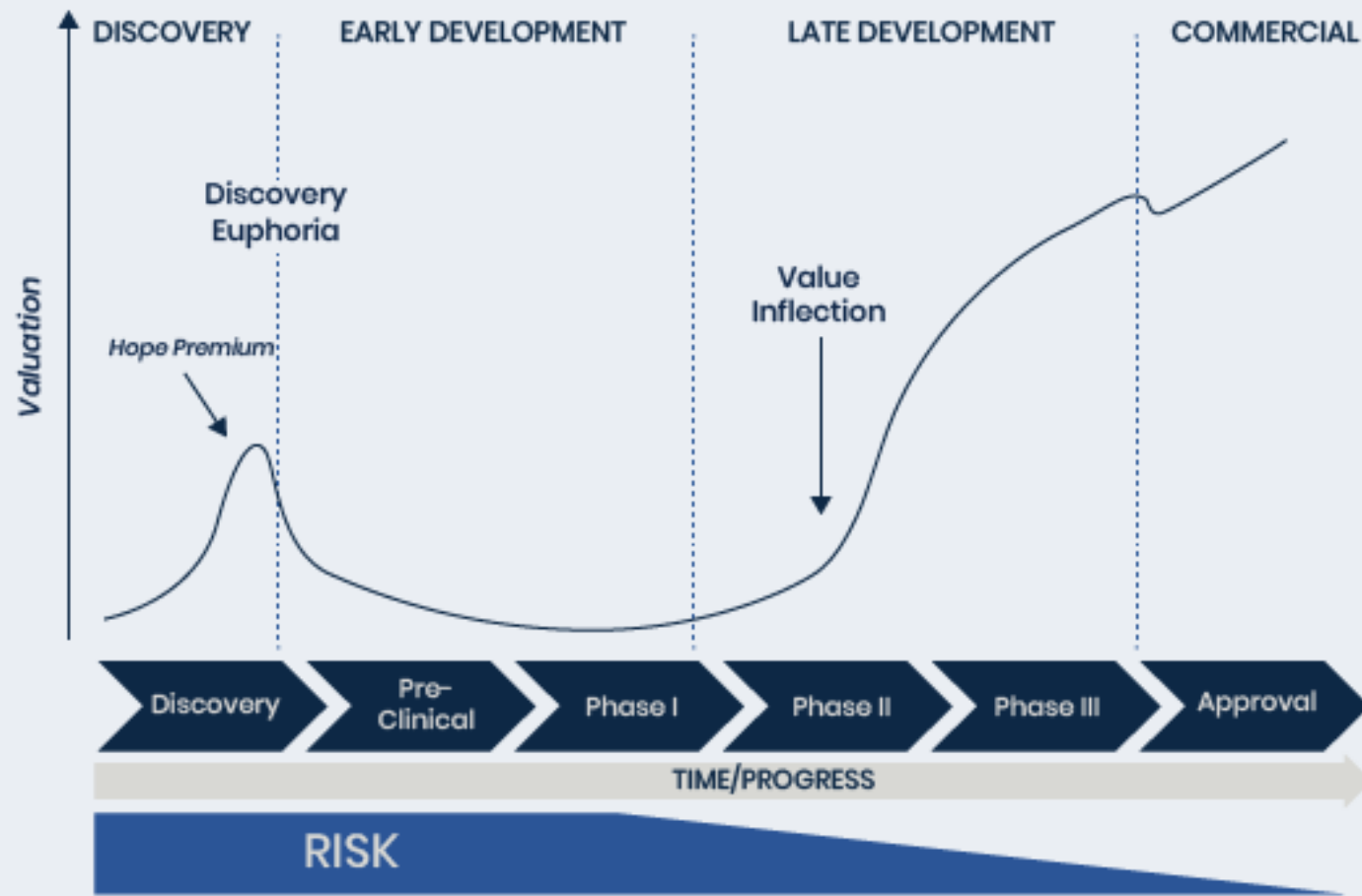
- Protection from cardiotoxicity (induced by anti-cancer drugs)
- Oncology
- Ischemic reperfusion injury – other organs
- Cardiac hypertrophy and fibrosis
- Pain management
- Epilepsy

Xolatryp formulation:

- Current delivery is via intravenous infusion
- Work planned for an oral dose form



Poised for Value Inflection



Source: Karst Peak Rx - www.kp-rx.com



Financial Overview and Outlook



FY2025 Highlights and FY2026 Outlook

› Resources

- › Cash balance of AU\$7.92 million at 30 September 2025
- › Expected R&D rebate of AU\$2.16 million (subject to Government Agency Review)
- › AU\$8.25 million (before costs) in new equity capital raised in August 2025.

› Programs

- › Progressing Xolatryp into Phase IIa clinical trial
 - › Targeted HREC submission November 2025
 - › Targeted first patient dosing March 2026

Operating Results Summary

	FY2025 (AU\$)	FY2023 (AU\$)
R&D Costs	4,376,215	2,030,502
Corporate and admin expenses	1,061,480	577,842
Share-based payment expense	177,218	358,074
Professional services expense	381,618	477,948
Employment benefits expense	1,225,077	1,127,500

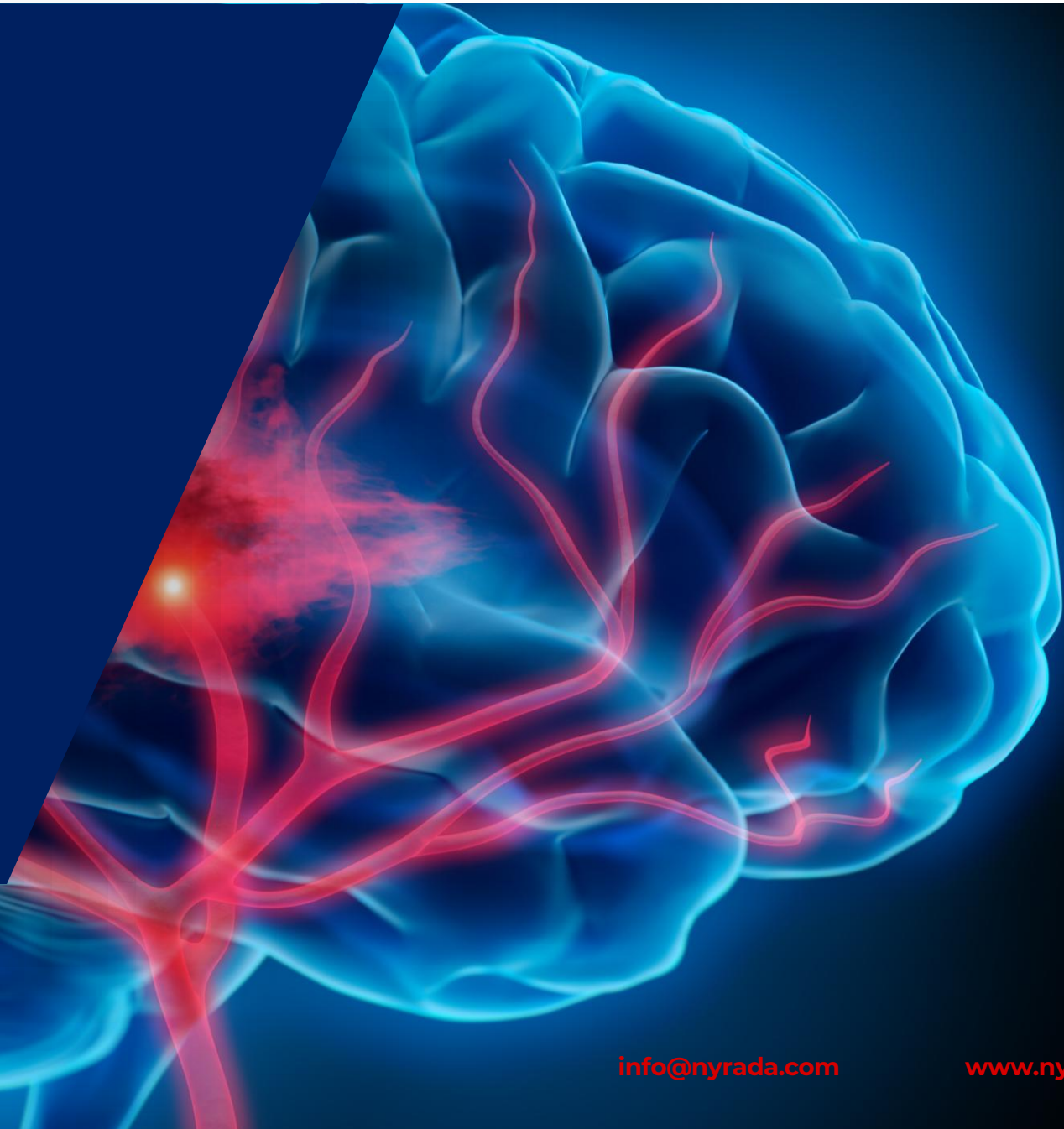
Conclusion

› **Nyrada – the company**

- › Pioneering TRPC channel inhibition therapies to treat a range of medical conditions
- › AU\$7.92 million cash position at end Sep 2025
- › AU\$8.25 million capital raised in Aug 2025
- › AU\$2.16 million R&D rebated expected in Dec 2025

› **Xolatryp – the lead drug asset**

- › Solid scientific foundations and well understood mechanism of action
- › Composition of matter patent pending
- › Preclinical efficacy demonstrated in AMI, ischemic stroke, and traumatic brain injury (TBI)
- › Phase I (safety, tolerability, PK) clinical trial completed
- › Phase IIa (safety and efficacy) clinical trial pending



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