

Investor Presentation

October 2024

Improving Lives, Offering Hope

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

ASX:NYR



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About Nyrada

- Drug discovery and development company specialising in rational design of novel small molecule therapeutics.
- > Nyrada's lead drug candidate NYR-BI03:
 - demonstrated strong preclinical efficacy protecting the brain from secondary injury following stroke.
 - demonstrated strong preclinical efficacy in protecting heart following acute myocardialischemic reperfusion injury.
 - preclinical TBI efficacy study with Walter Reed Army Institute of Research and UNSW in progress.
 - in advanced stages of Good Laboratory Practice safety and tolerability testing.
- Exploratory works for other indications and opportunities.
- > Commercially focused business model and expert team.



Investment Proposition

- Pioneering transient receptor potential canonical (TRPC) channel blocking therapies.
- > First-in-class neuroprotection therapy with novel mode of action.
- Cardioprotection therapy with superior performance to Captopril, an FDA approved ACE inhibitor.
- One drug asset targeting two significant therapeutic areas - neuroprotection and cardioprotection.
- Collaborations with world leading institutions: Walter Reed Army Institute of Research (WRAIR) and UNSW Sydney. Strategic partnership with Rebion.
- Proven and globally experienced board and team.
- Cash position of AU\$4.77M at 30 June 2024.
 AU\$0.99M R&D rebate expected around December 2024.



One Drug

))) NYR-BIO3 (((

GLP data package nearing completion

Approaching Phase I clinical trial in 4QCY2024

Two Applications



Neuroprotection



Cardioprotection

Three Markets

STROKE

~US\$52.2 billion by 20322

TRAUMATIC BRAIN INJURY

~US\$5.5 billion by 20344

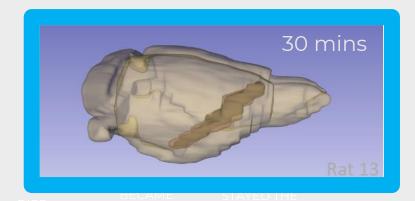
MYOCARDIAL INFARCTION

~US\$3.7 billion by 20325

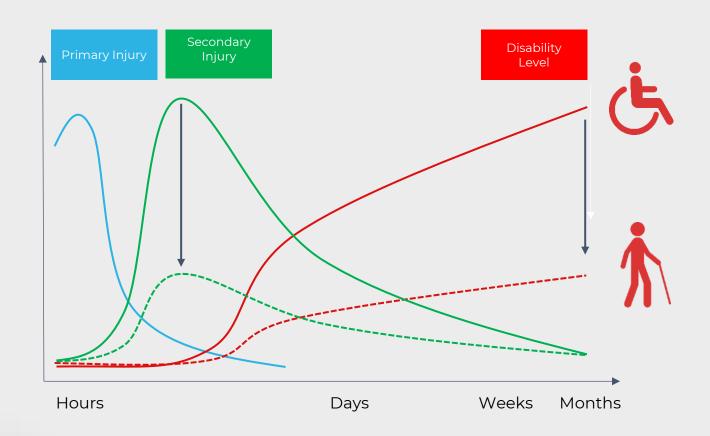
Neuroprotection – Stroke and TBI



Serial reconstruction from MRI







Nyrada drug NYR-BI03 An acute 3-day intravenous treatment

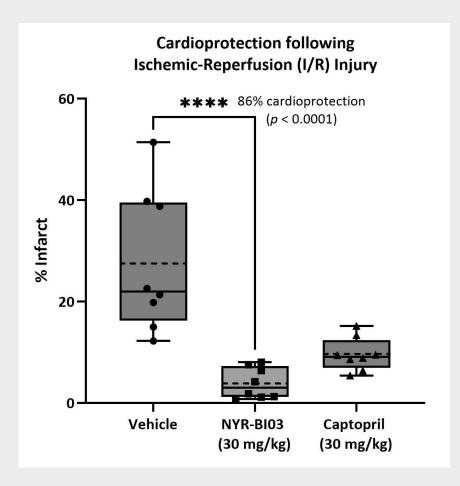


Reduce secondary injury resulting from stroke or TBI

- Improve survivability, limit disability
- Improve quality of life

Cardioprotection

Key Preclinical Results:



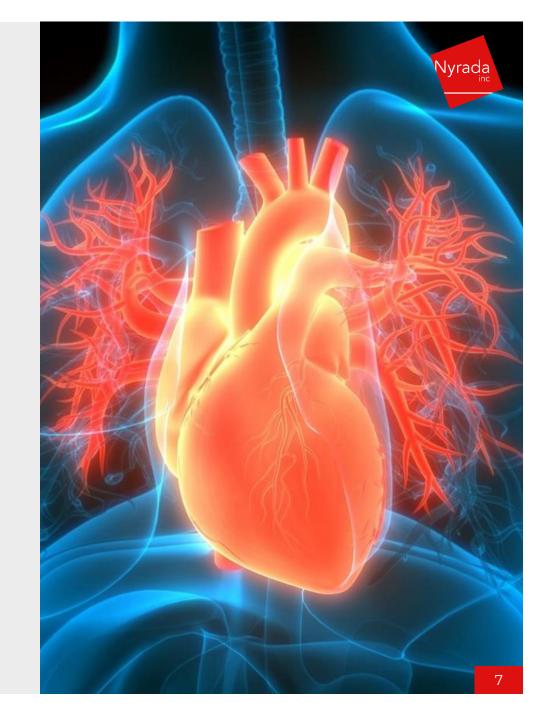
NYR-BI03 showed strong efficacy limiting cardiovascular damage associated with coronary heart disease

86%

Cardioprotection following myocardial ischemicreperfusion injury

Superior efficacy

Compared to FDA-approved, Captopril



GLP Safety Studies Approaching Conclusion

GLP Study	Reported
AMES	16 July 2024
hERG	16 July 2024
Rat CNS	06 August 2024
Rat Respiratory	20 August 2024
Dog cardiovascular safety	09 September 2024
14-day dog toxicology	27 September 2024
14-day rat toxicology	
In vitro micronucleus	
<i>In vivo</i> micronucleus	



Near Term Catalysts





Early to Mid 4QCY2024 NYR-BI03 GLP study updates

Mid 4QCY2024

NYR-BI03 Human Research Ethics application submission

Late 4QCY2024

NYR-BI03 Phase I clinical trial

Early 1QCY2025

WRAIR TBI study update

Appendices



Indicative Phase I Study Design



OBJECTIVES

To assess the safety, tolerability, and pharmacokinetics of NYR-BI03

DESIGN

- Randomised, double-blind placebo –controlled, dose escalation design
- 5 cohorts; 8 participants each cohort; 6:2 active and placebo treatments
- 3 cohorts will be single ascending doses
- 2 cohorts will be given continuous infusion doses

PARTICIPANTS

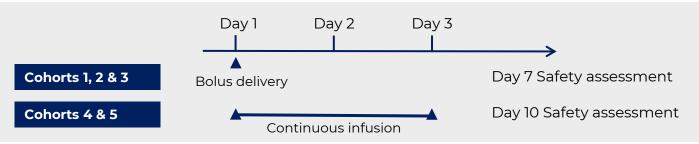
- Male and female healthy volunteers
- 18 50 years age



Cohort number	Dose administered
1	Low dose single bolus
2	Medium dose single bolus
3	High dose
4	Low dose continuous infusion (72 hrs)
5	High dose continuous infusion (72 hrs)

LOCATION & DURATION

- Study will be conducted at a clinical trial centre in Australia expected to commence 4QCY2024
- Study duration will vary between 1 4 days



*trial design subject to ethics approval

Large Market Opportunity – Stroke

Globally:

~15 million people suffer strokes annually¹

~5 millionleft permanently disabled¹

One approved drug class for stroke suitable for <15% of patients (tPA - tissue plasminogen activator).

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

Large and growing treatment market:

Currently ~US\$30.3 billion²

Growing ~7.5% CAGR²

Forecast ~US\$52.2 billion by 2030²



Large Market Opportunity – Traumatic Brain Injury (TBI)

Globally:

~5.5 million people suffer severe TBA annually³

~55 million
living with effects of

living with effects of medically treated TBI³

No current FDA approved treatments

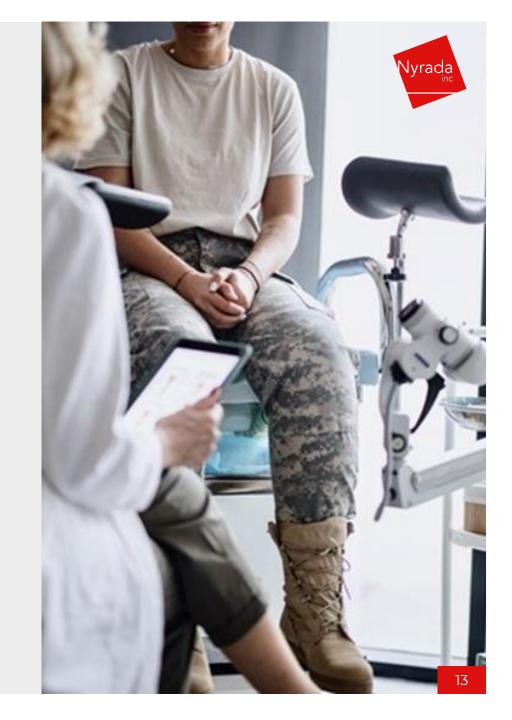
Effective treatment will improve patient outcomes and reduce high costs associated with long-term care of brain injury survivors.

Large and growing treatment market:

Currently ~US\$3.5 billion⁴

Growing ~6.2% CAGR⁴

Forecast ~US\$5.5 billion by 2030⁴



Nyrada's Lead Drug Candidate NYR-BI03





First-in-Class with Novel Mechanism of Action

- NYR-BI03 is a first-in-class therapy.
- Novel mechanism of action.
- Australian developed innovation.



Significant Unmet Clinical Need and Market Opportunity

- Targeting multiple indications.
- Stroke, TBI and Cardiac heart disease are leading causes of death and disability.
- No current FDA approved drugs to treat secondary brain injury.

Management Team with Proven Industry Experience





James Bonnar - CEO

- Business executive with 25 years of experience in healthcare companies in the UK, China, New Zealand, and Australia
- Experience in drug manufacture, preclinical development, clinical operations, regulatory affairs, and quality assurance
- Biotech experience spanning various therapeutic areas including cardiometabolic disease, neurodevelopment disorders, and brain injury



Cameron Jones - CFO

- Finance executive with experience as CFO and Company Secretary of ASX Listed and VC investee healthcare companies
- Supported several healthcare companies through IPOs, capital raisings and M&A transactions
- Managing Director of Bio101, financial services firm
- Chartered Accountant, Member of the Governance Institute of Australia and Registered Tax Agent



Dr Benny Evison - CSO

- More than 20 years of experience in the discovery and development of small molecule inhibitors as therapies for various cancers, cardiovascular diseases and neurodegenerative diseases
- Obtained a PhD at La Trobe
 University (Melbourne, Australia)
 in biochemistry and molecular
 biology, and a postdoctoral
 fellowship in chemical biology at
 St Jude Children's Research
 Hospital, (Memphis TN)

Supported by specialist advisers:

- Prof Gary Housley Brain injury program
- Prof Junichi Nabekura Brain injury program
- Dr Jim Palmer
 Medicinal Chemist
- Dr Mike Bickerdike Toxicologist

High Calibre Board with international experience

- Nyrada operates under the direction of a governing board of international calibre
- Strong track record in realising the value of biotech companies
- Experience in dealmaking, US/EU/AU capital markets, and relevant therapeutic area experience





John Moore
Non-Executive Chair



Marcus Frampton
Non-Executive Director



Dr lan Dixon
Non-Executive Director



Dr Rüdiger Weseloh Non-Executive Director



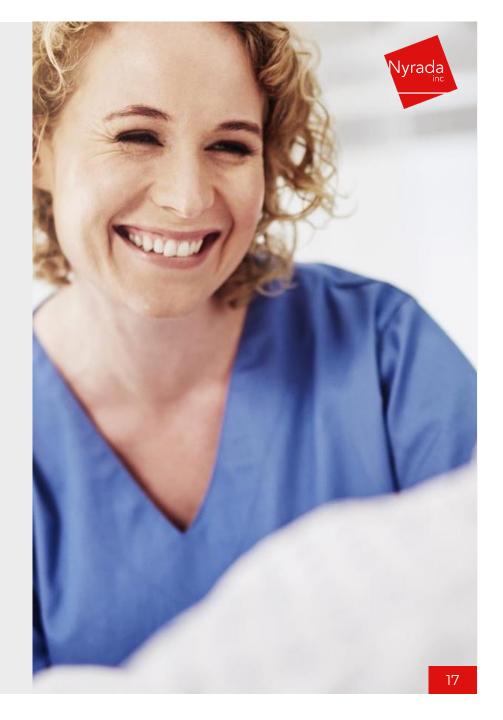
Dr Gisela Mautner Non-Executive Director



Christopher Cox
Non-Executive Director

References

- 1 World Health Organization https://www.emro.who.int/health-topics/stroke-cerebrovascular-accident/index.html#:~:text=Annually%2C%2015%20million%20people%20worldwide,cause%20is%20high%20blood%20pressure
- 2 Databridge Market Research https://www.databridgemarketresearch.com/reports/global-stroke-market .
- 3 National Academy of Sciences https://nap.nationalacademies.org/catalog/25394/traumatic-brain-injury-a-roadmap-for-accelerating-progress
- 4 Databridge Market Research https://www.databridgemarketresearch.com/reports/global-traumatic-brain-injuries-treatment-market
- 5 Spherical Insights https://www.globenewswire.com/en/news-release/2023/05/30/2678779/0/en/Global-Myocardial-Infarction-Market-Size-To-Grow-USD-3-7-Billion-By-2032-CAGR-of-6-8.html



Brain Injury Solution Animation



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