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### **About Nyrada**



- Nyrada is a US incorporated, Australia based drug discovery and development company specialising in novel small molecule therapeutics.
- Commercially focused business model and expert team.
- Experienced Board of Directors and Scientific Advisory Board with impressive track record in biotech.
- Nyrada's lead Brain Injury Program drug candidate NYR-BI03 demonstrated strong efficacy protecting the brain from secondary injury in recently announced preclinical study. Program.
- Low-cost exploratory works ongoing in Cholesterol Lowering Program.



#### **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 and Internationally Experienced Governing Board



- Nyrada operates under the direction of a governing board of international calibre
- Strong track record in finding and 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

#### **Nyrada Investment Proposition**



- Globally experienced and proven team.
- > First-in-class therapy with novel mode of action.
- Significant target market and unmet clinical demand.
- Collaborations with world leading institutions, Walter Reed Army Institute of Research (WRAIR) and UNSW Sydney.
- > Cash position at 31 March 2024 was A\$5.60M.



## March 2024 Capital Raise



#### Nyrada has concluded a placement for A\$1,755,000 before costs

## New Capital

- A placement of approximately A\$1.755M of new equity capital, comprising 23,400,000 Chess Depositary Interests (CDIs), to professional and sophisticated investors.
- The Placement issue price is A\$0.075 per CDI, representing a 16.7% discount to the last traded price and 26.1% discount to the 15-day VWAP.

## Director Participation

- Nyrada Directors will subscribe for an additional \$0.210M of CDIs on the same terms as the placement CDIs.
- This allocation to Directors is subject to shareholder approval with an Extraordinary General Meeting (EGM) provisionally scheduled for 16 May 2024.

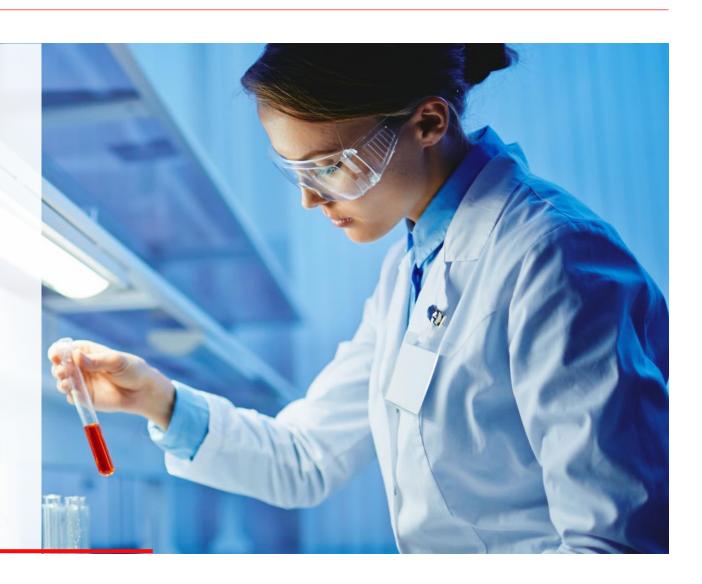
## Use of Funds

- Remaining preclinical work and a Phase I in-human clinical trial of NYR-BI03
- Prepare and submit Investigational New Drug (IND) applications with the US FDA for NYR-BI03's use in both traumatic brain injury (TBI) and stroke
- Planning and preparation for a Phase II in-human trial of NYR-BI03's effectiveness in TBI and stroke.

### **Key 3QFY2024 Financials**



- A\$5.60 million cash at 31 March 2024
- A\$4.65 million cash at 31 December 2023
- 3QFY2024 cash operating outflows A\$0.73 million
- 2QFY2024 cash operating outflows A\$1.21 million
- Fully subscribed placement raised A\$1.76 million (before costs)
- Additional A\$0.21 million (before costs) expected following 16 May 2024 EGM



### Nyrada's Lead Brain Injury 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 and TBI are leading causes of death and disability.
- No current FDA approved drugs to treat secondary brain injury.

## **Statistically Significant Neuroprotection Achieved**



Recent preclinical study demonstrated strong efficacy and good tolerability

42%

## of brain injury in penumbra region rescued on average

MRI brain imaging showed a statistically significant neuroprotection achieved when animals received treatment (*p* value 0.021)<sup>1</sup>

41%

## mean level of biomarkers for NYR-BI03 animals compared to control vehicle

NfL biomarker levels likely reflect neuroprotection. The mean level for the NYR-BI03 animals was 41% lower than the control vehicle (*p* value 0.068)<sup>2</sup>

## Supporting NYR-BI03's favourable safety profile, the current study had no drug-related adverse effects.

#### **Preclinical Study Design**

- The study involved inducing a focal ischemic stroke using a photothrombotic model.
- 16 test animals were treated with either NYR-BI03 or vehicle 30 mins following the induced brain injury. Treatment was conducted for 72 hours via continuous intravenous infusion.

<sup>1.</sup> p value 0.021, Mann-Whitney Rank Sum Test.

<sup>2.</sup> *p* value 0.068, t-test.

#### **Large Market Opportunity – Stroke**



Globally:

~15 million
people suffer
strokes annually<sup>1</sup>

~5 million left permanently disabled<sup>1</sup>

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 billion<sup>2</sup>

Growing ~7.5% CAGR<sup>2</sup>

Forecast ~US\$52.2 billion by 2030<sup>2</sup>



### **Large Market Opportunity – Traumatic Brain Injury (TBI)**



Globally:

**~5.5 million**people suffer severe
TBA annually<sup>3</sup>

~55 million
living with effects of medically treated TBI<sup>3</sup>

No current FDA approved treatments

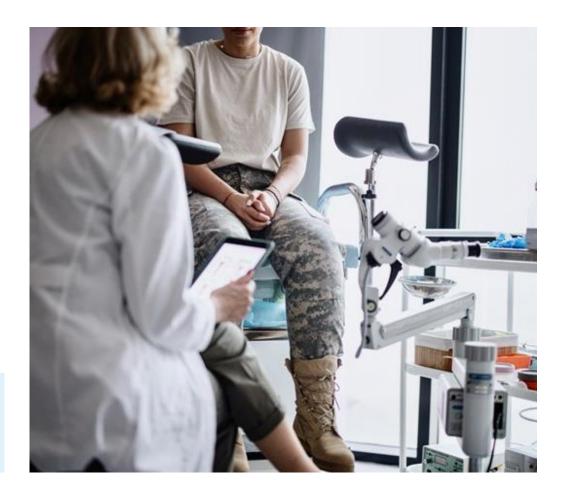
Effective treatment will improve patient outcomes and reduce high costs associated with long-term care of brain injury survivors.

#### **Large and growing treatment market:**

Currently ~US\$3.5 billion<sup>4</sup>

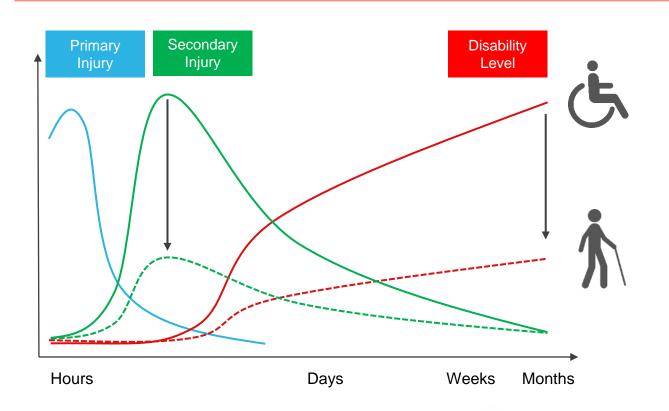
Growing ~6.2% CAGR<sup>4</sup>

Forecast ~US\$5.5 billion by 2030<sup>4</sup>

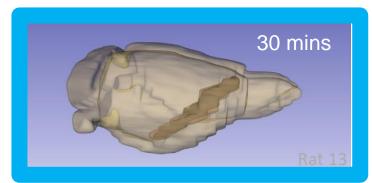


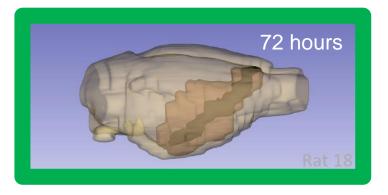
#### **Brain Injury Trajectory, Patient Outcomes, Treatment Aims**





#### Serial reconstruction from MRI





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



Reduce secondary injury resulting from stroke or TBI

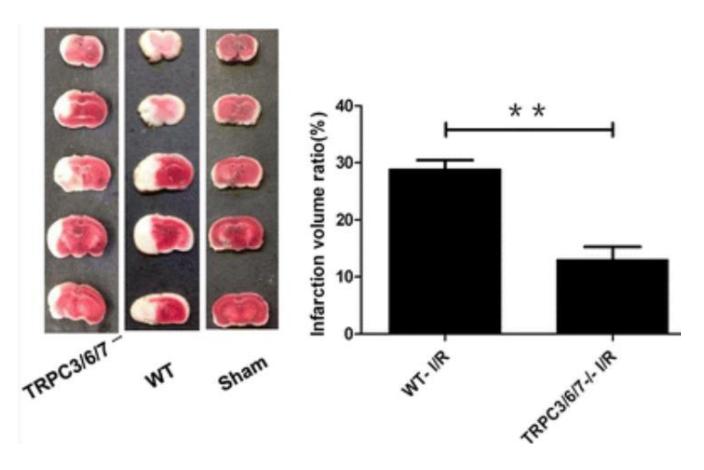
- Improve survivability, limit disability
- Improve quality of life

### **Proof-of-Concept:**

#### TRPC 3/6/7 Knockout Mouse Model



#### Middle cerebral artery occlusion (MCAO) stroke model shows neuroprotection in TRPC3,6,7 triple knockout mouse

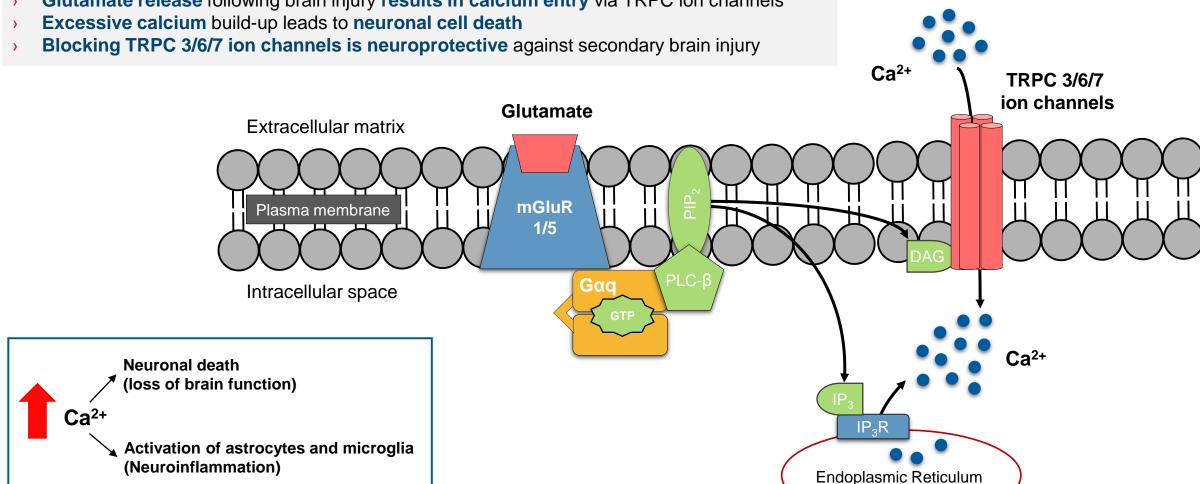


- TRPC 3/6/7 KO mice have significantly smaller lesion size compared to WT in a mouse middle cerebral artery occlusion model (Chen et al. Mol. Neurobiol. 2017)
- NYR-BI03 blocks TRPC3/6/7 channels

#### TRPC 3/6/7 Ion Channels as a Therapeutic Target



Glutamate release following brain injury results in calcium entry via TRPC ion channels

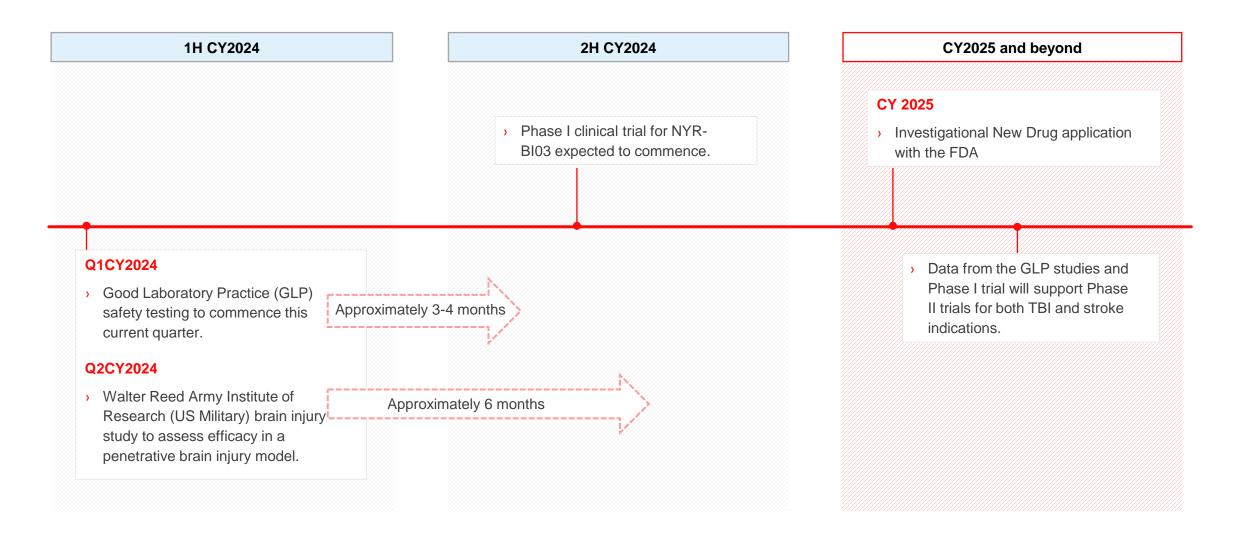


NYRADA INC (ASX:NYR) — CORPORATE PRESENTATION

Ca<sup>2+</sup> internal stores

## **Brain Injury Program Milestones Ahead**





#### **Brain Injury Program - Indicative Phase I Study Design**



#### Commencing in 2HCY2024

**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**

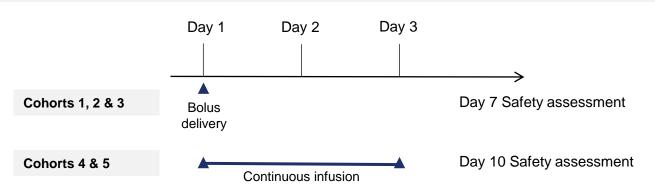
- Male and female healthy volunteers
- 18 50 years age



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 2H CY2024
- The study duration will vary between 1 4 days



\*trial design subject to ethics approval

## **Summary**



# Nyrada is advancing a novel drug candidate to reduce the secondary brain injury caused by stroke or traumatic brain injury.





NYR-BI03 protects the brain from secondary injury in newly announced preclinical study

## First-in-class therapy



With a novel mechanism of action

## Large patient markets



Significant market opportunity in stroke and traumatic brain injury markets

## Near term milestones



Final safety tests commencing in coming weeks and first-in-human trials in 2HCY2024

#### **Nyrada's Vision and Strategy**





To become a high growth pharmaceutical discovery and development company specialising in early-stage drug development of novel treatments.



To develop treatments for diseases where there is an unmet clinical need, or where current treatments are suboptimal, and to monetise the value of these treatments through advancing highly optimised drug candidates towards out-licensing.

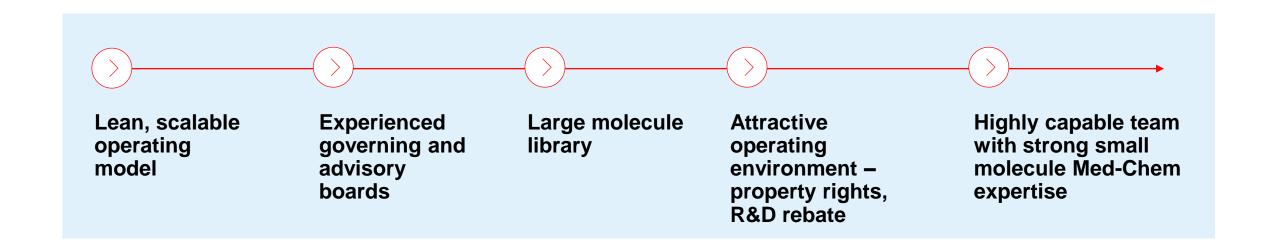


Delivering upon our vision and strategy through our brain injury program via a TRPC ion channel blocker.

#### **Commercially Focused Business Model**



We aim to create value for patients and shareholders by advancing highly optimised drug candidates towards out-licensing.



#### References



- 1 World Health Organization <a href="https://www.emro.who.int/health-topics/stroke-cerebrovascular-accident/index.html#:~:text=Annually%2C%2015%20million%20people%20worldwide,cause%20is%20high%20blood%20pressure">https://www.emro.who.int/health-topics/stroke-cerebrovascular-accident/index.html#:~:text=Annually%2C%2015%20million%20people%20worldwide,cause%20is%20high%20blood%20pressure</a>
- 2 Databridge Market Research <a href="https://www.databridgemarketresearch.com/reports/global-stroke-market">https://www.databridgemarketresearch.com/reports/global-stroke-market</a> .
- 3 National Academy of Sciences <a href="https://nap.nationalacademies.org/catalog/25394/traumatic-brain-injury-a-roadmap-for-accelerating-progress">https://nap.nationalacademies.org/catalog/25394/traumatic-brain-injury-a-roadmap-for-accelerating-progress</a>
- 4 Databridge Market Research <a href="https://www.databridgemarketresearch.com/reports/global-traumatic-brain-injuries-treatment-market">https://www.databridgemarketresearch.com/reports/global-traumatic-brain-injuries-treatment-market</a>







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



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