



Nyrada Quarterly Activities Report & Appendix 4C

Highlights:

- Nyrada remains on track for 2QFY2025 commencement of first in-human Phase I clinical trial for its Brain Injury drug candidate NYR-BI03.
- Good Laboratory Practice (GLP) safety testing studies of NYR-BIO3 continuing with results to be progressively disclosed from 1QFY2025.
- Strategic partnership agreement with Rebion to advance research and development of brain injury therapies and outcomes.
- Nyrada Chair and Non-Executive Director CDI subscription completed, raising A\$0.21 million (before costs).
- Cash position of A\$4.77 million as at 30 June 2024.

Nyrada Inc (ASX:NYR) ("Nyrada or "Company"), a drug development company specialising in novel small molecule therapeutics announces today its Quarterly Activities Report and Appendix 4C for the three-month period concluding 30 June 2024.

Brain Injury Program

Nyrada is developing NYR-BIO3, a first in class neuroprotection treatment for both traumatic brain injury (TBI) and stroke. In February 2024, the Company disclosed preclinical stroke study results showing NYR-BIO3 provided a statistically significant level of neuroprotection, rescuing 42% of brain injury in the penumbra region in treated animals.

Good Laboratory Practice (GLP) Studies

Following the demonstration of brain injury neuroprotection efficacy in a preclinical stroke study, in late 3QFY2024, Nyrada commenced Good Laboratory Practice (GLP) studies to assess the safety of NYR-BIO3 in two animal species. The successful completion of a comprehensive set of GLP studies is a necessary precondition to undertake a first-in-human clinical Phase I study, scheduled to commence in 2QFY2025.



The studies that comprise the GLP safety assessment of NYR-BIO3 include:

- hERG (Human Ether-a-go-go-related Gene);
- AMES (Bacterial Reverse Mutation Test);
- CNS (Central Nervous System), cardiovascular and respiratory safety;
- In vitro and In vivo micronucleus; and
- 14-day toxicity studies in two animal species.

Nyrada anticipates the hERG and AMES study results will be reported in the coming weeks, and CNS and respiratory study results soon after. The remaining study results are expected to be disclosed thereafter with all studies expected to be reported in early 2QFY2025.

Walter Reed Traumatic Brain Injury (TBI) Study

Nyrada's collaborative TBI study with the <u>Walter Reed Army Institute of Research</u> (WRAIR) commenced in early 4QFY2024. This study is assessing the efficacy of NYR-BIO3 in a rodent model of penetrating traumatic brain injury (PTBI). This PTBI model is proprietary to WRAIR and seeks to mimic the serious head injuries suffered by military service members.

As part of this study, the degree to which NYR-BIO3 provides neuroprotection following a PTBI will be assessed and measured.

Results from this study are expected to be reported before the conclusion of 1QFY2025.

Rebion Strategic Partnership

In late June 2024, Nyrada signed a <u>Strategic Partnership Agreement</u> with Boston-based medical device development company Rebion. Rebion uses Neural Performance Scanning technology to identify and monitor functional impairments in the brain stemming from disease or injury.

Nyrada and Rebion will collaborate to advance therapies and outcomes for TBI sufferers. This Strategic Partnership Agreement spans joint research, conference presentations, and applications for non-dilutive funding grants.

Phase I Human Clinical Trial

Subject to the successful completion of GLP studies, Nyrada intends to commence a first-in-human Phase I clinical trial for NYR-BIO3 in 2QFY2025. This Phase I trial will assess the safety of NYR-BIO3 in healthy human volunteers. The study will be conducted in Australia and will confirm the safe dose range to take forward into subsequent clinical trials.



The data from the stroke study, WRAIR TBI study, GLP studies and Phase I trial will support a Phase II efficacy trial of NYR-BIO3 for both stroke and TBI indications in humans.

Corporate and Financial Update

As at 30 June 2024, Nyrada had a cash position of A\$4.77 million (A\$5.60 million as at 31 March 2024).

The Company's cash balance was enhanced during the June quarter by A\$0.21 million (before costs) from the issue of CDIs to Nyrada's Chair and Non-Executive Directors on the same terms as those investors who participated in the March 2024 capital raise (A\$0.075 per CDI).

Total cash operating outflows for the June 2024 quarter were approximately A\$1.05 million, offset by approximately \$35,000 interest income received. Total cash operating outflows for the March 2024 quarter were approximately A\$1.02 million.

As per prior years, the Company intends to lodge an application under the Australian Government's R&D Tax incentive program for research and development expenditure undertaken in the 2024 financial year. An estimate of the quantum of the refund will be provided in Nyrada's audited financial results to be reported in August 2024.

In accordance with Listing Rule 4.7C, payments made to related parties and their associates included in item 6.1 of the Appendix 4C were approximately A\$102,000 and included Director fees (approximately A\$77,000 for the quarter ending 30 March 2024).

-ENDS-



About Nyrada Inc

Nyrada is a drug discovery and development company specialising in novel small molecule therapies. The Company has two main programs, each targeting market sectors of significant size and considerable unmet clinical need. These are a drug to treat brain injury, specifically traumatic brain injury and stroke, and a cholesterol lowering drug. Nyrada Inc. ARBN 625 401 818 is a company incorporated in the state of Delaware, US, and the liability of its stockholders is limited.

www.nyrada.com

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

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Forward-Looking Statements

This announcement may contain forward-looking statements. You can identify these statements by the fact they use words such as "aim", "anticipate", "assume", "believe", "continue", "could", "estimate", "expect", "intend", "may", "plan", "predict", "project", "plan", "should", "target", "will" or "would" or the negative of such terms or other similar expressions. Forward-looking statements are based on estimates, projections, and assumptions made by Nyrada about circumstances and events that have not yet taken place. Although Nyrada believes the forward-looking statements to be reasonable, they are not certain. Forward-looking statements involve known and unknown risks, uncertainties and other factors that are in some cases beyond the Company's control that could cause the actual results, performance, or achievements to differ materially from those expressed or implied by the forward-looking statement.

Appendix 4C

Quarterly cash flow report for entities subject to Listing Rule 4.7B

Name of entity

Nyrada Inc.		
ABN	Quarter ended ("current quarter")	
54 625 401 818	30 June 2024	

Con	solidated statement of cash flows	Current quarter \$A'000	Year to date (12 months) \$A'000
1.	Cash flows from operating activities		
1.1	Receipts from customers	-	-
1.2	Payments for		
	(a) research and development	(525)	(1,982)
	(b) product manufacturing and operating costs	-	-
	(c) advertising and marketing	-	-
	(d) leased assets	-	-
	(e) staff costs	(282)	(1,118)
	(f) administration and corporate costs	(247)	(1,359)
1.3	Dividends received (see note 3)	-	-
1.4	Interest received	35	136
1.5	Interest and other costs of finance paid	-	-
1.6	Income taxes paid	-	-
1.7	Government grants and tax incentives	-	3,557
1.8	Other (provide details if material)	-	-
1.9	Net cash from / (used in) operating activities	(1,019)	(766)

2.	Cas	sh flows from investing activities	
2.1	Pay	ments to acquire or for:	
	(a)	entities	-
	(b)	businesses	-
	(c)	property, plant and equipment	-
	(d)	investments	-
	(e)	intellectual property	-
	(f)	other non-current assets	-

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Con	solidated statement of cash flows	Current quarter \$A'000	Year to date (12 months) \$A'000
2.2	Proceeds from disposal of:		
	(a) entities	-	-
	(b) businesses	-	-
	(c) property, plant and equipment	-	-
	(d) investments	-	-
	(e) intellectual property	-	-
	(f) other non-current assets	-	-
2.3	Cash flows from loans to other entities	-	-
2.4	Dividends received (see note 3)	-	-
2.5	Other (provide details if material)	-	-
2.6	Net cash from / (used in) investing activities	-	-

3.	Cash flows from financing activities		
3.1	Proceeds from issues of equity securities (excluding convertible debt securities)	210	1,965
3.2	Proceeds from issue of convertible debt securities	-	-
3.3	Proceeds from exercise of options	-	-
3.4	Transaction costs related to issues of equity securities or convertible debt securities	(25)	(139)
3.5	Proceeds from borrowings	-	-
3.6	Repayment of borrowings	-	-
3.7	Transaction costs related to loans and borrowings	-	-
3.8	Dividends paid	-	-
3.9	Other (provide details if material)	-	-
3.10	Net cash from / (used in) financing activities	185	1,826

4.	Net increase / (decrease) in cash and cash equivalents for the period		
4.1	Cash and cash equivalents at beginning of period	5,603	3,709
4.2	Net cash from / (used in) operating activities (item 1.9 above)	(1,019)	(766)
4.3	Net cash from / (used in) investing activities (item 2.6 above)	-	-

Con	solidated statement of cash flows	Current quarter \$A'000	Year to date (12 months) \$A'000
4.4	Net cash from / (used in) financing activities (item 3.10 above)	185	1,826
4.5	Effect of movement in exchange rates on cash held	-	-
4.6	Cash and cash equivalents at end of period	4,769	4,769

5.	Reconciliation of cash and cash equivalents at the end of the quarter (as shown in the consolidated statement of cash flows) to the related items in the accounts	Current quarter \$A'000	Previous quarter \$A'000
5.1	Bank balances	2,269	5,603
5.2	Call deposits	2,500	-
5.3	Bank overdrafts	-	-
5.4	Other (provide details)	-	-
5.5	Cash and cash equivalents at end of quarter (should equal item 4.6 above)	4,769	5,603

6.	Payments to related parties of the entity and their associates	Current quarter \$A'000
6.1	Aggregate amount of payments to related parties and their associates included in item 1	102
6.2	Aggregate amount of payments to related parties and their associates included in item 2	-
	if any amounts are shown in items 6.1 or 6.2, your quarterly activity report must includ nation for, such payments.	e a description of, and an

The amount at 6.1 includes Director fees and salary (including superannuation) and consulting fees for directors and related parties.

7.	Financing facilities Note: the term "facility" includes all forms of financing arrangements available to the entity. Add notes as necessary for an understanding of the sources of finance available to the entity.	Total facility amount at quarter end \$A'000	Amount drawn at quarter end \$A'000
7.1	Loan facilities	-	-
7.2	Credit standby arrangements	-	-
7.3	Other (please specify)	-	-
7.4	Total financing facilities	-	-
7.5	Unused financing facilities available at qu	uarter end	-
7.6	Include in the box below a description of each rate, maturity date and whether it is secured facilities have been entered into or are proposinclude a note providing details of those facilities.	or unsecured. If any add osed to be entered into af	tional financing
	include a note providing details of those facil	lities as well.	

8.	Estimated cash available for future operating activities	\$A'000
8.1	Net cash from / (used in) operating activities (item 1.9)	(1,019)
8.2	Cash and cash equivalents at quarter end (item 4.6)	4,769
8.3	Unused finance facilities available at quarter end (item 7.5)	-
8.4	Total available funding (item 8.2 + item 8.3)	4,769
8.5	Estimated quarters of funding available (item 8.4 divided by item 8.1)	4.7
	Note: if the entity has reported positive net operating cash flows in item 1.9, answer item figure for the estimated quarters of funding available must be included in item 8.5.	8.5 as "N/A". Otherwise, a

8.6 If item 8.5 is less than 2 quarters, please provide answers to the following questions:

8.6.1 Does the entity expect that it will continue to have the current level of net operating cash flows for the time being and, if not, why not?

Answer:

8.6.2 Has the entity taken any steps, or does it propose to take any steps, to raise further cash to fund its operations and, if so, what are those steps and how likely does it believe that they will be successful?

Answer:

8.6.3 Does the entity expect to be able to continue its operations and to meet its business objectives and, if so, on what basis?

Answer:

Note: where item 8.5 is less than 2 quarters, all of questions 8.6.1, 8.6.2 and 8.6.3 above must be answered.

Compliance statement

- This statement has been prepared in accordance with accounting standards and policies which comply with Listing Rule 19.11A.
- 2 This statement gives a true and fair view of the matters disclosed.

	9 July 2024
Date:	
	By Order of the Board
Authorised by:	(Name of body or officer authorising release – see note 4)

Notes

- This quarterly cash flow report and the accompanying activity report provide a basis for informing the market about the
 entity's activities for the past quarter, how they have been financed and the effect this has had on its cash position. An
 entity that wishes to disclose additional information over and above the minimum required under the Listing Rules is
 encouraged to do so.
- 2. If this quarterly cash flow report has been prepared in accordance with Australian Accounting Standards, the definitions in, and provisions of, AASB 107: Statement of Cash Flows apply to this report. If this quarterly cash flow report has been prepared in accordance with other accounting standards agreed by ASX pursuant to Listing Rule 19.11A, the corresponding equivalent standard applies to this report.
- 3. Dividends received may be classified either as cash flows from operating activities or cash flows from investing activities, depending on the accounting policy of the entity.
- 4. If this report has been authorised for release to the market by your board of directors, you can insert here: "By the board". If it has been authorised for release to the market by a committee of your board of directors, you can insert here: "By the [name of board committee eg Audit and Risk Committee]". If it has been authorised for release to the market by a disclosure committee, you can insert here: "By the Disclosure Committee".
- 5. If this report has been authorised for release to the market by your board of directors and you wish to hold yourself out as complying with recommendation 4.2 of the ASX Corporate Governance Council's *Corporate Governance Principles* and *Recommendations*, the board should have received a declaration from its CEO and CFO that, in their opinion, the financial records of the entity have been properly maintained, that this report complies with the appropriate accounting standards and gives a true and fair view of the cash flows of the entity, and that their opinion has been formed on the basis of a sound system of risk management and internal control which is operating effectively.