29 OCTOBER 2025

ASX RELEASE

QUARTERLY ACTIVITIES AND CASH FLOW REPORTS – SEPTEMBER 2025

Key Highlights from the Quarter

- **Further promising developments from the ongoing ACCENT trial of narmafotinib in metastatic pancreatic cancer were presented, including additional confirmed responses and an interim median Progression Free Survival figure of 7.6 months. This latter result is a 2 month improvement on historical data for chemotherapy alone**
- **Ethics approval was obtained in Australia for the new trial of narmafotinib, investigating a combination with the chemotherapy FOLFIRINOX, and the first patients in this trial have been dosed**
- **A key patent protecting narmafotinib has been allowed by the US Patent and Trademarks Office**
- **Completion of a capital raise of A$27.5 million (before costs) funding the company into 2027**

Melbourne, Australia: Amplia Therapeutics Limited (ASX: ATX), ("Amplia" or the "Company"), a company developing new approaches for the treatment for cancer and fibrosis, is pleased to announce further progress across its small molecule, focal adhesion kinase (FAK) inhibitor program and the release of its Appendix 4C Cash Flow Report (attached) for the quarter ending 30 September 2025.

Operations Update

The Company continues to make significant strides in the clinical development of its best-in-class FAK inhibitor, narmafotinib, for the treatment of metastatic pancreatic cancer. The Company has reported encouraging results from ongoing trials, regulatory milestones, and patent advancements, reinforcing its commitment to improving outcomes for patients with this challenging disease.

Clinical Trial Updates

In July the Company announced that the ACCENT clinical trial, which investigates narmafotinib in combination with standard chemotherapies gemcitabine and Abraxane®, recorded additional confirmed partial responses (PRs) bringing the total at that time to 17. A confirmed partial response (PR) is defined as tumour shrinkage exceeding 30%, sustained for at least two months and without the appearance of new cancerous lesions. The objective response rate (ORR) in the trial increased from 29% (16 out of 55 patients) to 31% (17 out of 55 patients), surpassing the 23% response rate reported for chemotherapy

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alone[1]. These data include a patient where a complete response (CR) was recorded and an additional patient who, after surgical intervention, was reported as a pathological CR, both of which are very rare occurrences in the treatment of metastatic pancreatic cancer. More recently, the Company updated the ORR to 33% (18 out of 55 patients) after an additional patient recorded a confirmed PR[2].

Interim data in the ACCENT trial up to 20 July 2025 was disclosed on 6 August 2025 and indicated a median progression-free survival (PFS) of 7.6 months, notably higher than the 5.5 months observed with chemotherapy alone and exceeding results from other similar trials. For this metastatic patient population where the cancer is highly aggressive, a two-month improvement in PFS is considered a notable improvement, and indeed the last drug approved in pancreatic cancer (NALIRIFOX) was based on a two-month improvement in overall survival over standard-of-care treatment.

Importantly, narmafotinib continues to show good tolerability in patients. The adverse event profile for the narmafotinib combination closely mirrors that of the chemotherapy regimen alone, with negligible additional patient burden for most participants. Durability of response is also promising, with seven patients remaining on trial for more than 12 months. The mean duration on trial for patients determined at 20 July 2025 was 202 days, compared to 117 days in the MPACT study. This has improved to 219 days from an analysis at the end of the quarter[2].

Expansion into FOLFIRINOX Combination

Last year the Company reported that its planned trial in the US, under an Investigational New Drug (IND) application, had been cleared by the US Food and Drug Administration (FDA)[3]. In July this year, Amplia received ethics approval in Australia for the new Phase 1b/2a clinical trial combining narmafotinib with FOLFIRINOX chemotherapy, following on from US ethics approval received the month prior. FOLFIRINOX, a more aggressive chemotherapy treatment, is commonly used in the treatment of advanced pancreatic cancer in the US and Western Europe. This open-label trial aims to identify the optimal daily dose of narmafotinib, given orally, when paired with FOLFIRINOX, which is administered intravenously every two weeks. The trial is designed following the Project Optimus guidance from the FDA[4], and will proceed in two stages: dose exploration with up to 27 patients and a two dose comparison across 40 patients. Patient recruitment has commenced at sites in Melbourne and Sydney, with several US sites to begin recruitment later this year. The first participants, of a cohort of up to nine patients, have already started dosing, marking an important milestone for the program.

Regulatory and Commercial Progress

On the regulatory front, the United States Adopted Names (USAN) Council has approved "narmafotinib" as the generic drug name for Amplia's FAK inhibitor, following earlier approval by the World Health Organization (WHO) for global use. This formal recognition is crucial for the drug's commercial development, particularly in the US market.

Amplia has also announced the allowance of a key patent by the US Patent and Trademark Office, covering the specific salt and crystal form of narmafotinib used in clinical trials. The patent, which secures protection until at least 2040 in multiple jurisdictions including Australia, Japan, Europe, India, Korea, Singapore, and New Zealand, ensures the company's intellectual property remains safeguarded and commercially valuable.

Future Outlook

As trials mature and further data become available, Amplia remains optimistic about improving outcomes for pancreatic cancer patients through innovative combination therapies. Business development and partnering discussions have advanced in recent months with a number of potential partners and collaborators showing interest in the developing dataset.

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[1] *New England Journal of Medicine* 2013; 369: 1691 – 703
[2] ASX Announcement 9 October 2025
[3] ASX Announcement 18 January 2024
[4] https://www.fda.gov/about-fda/oncology-center-excellence/project-optimus (accessed October 2025)

Capital Raise

On 23 July 2025 the Company announced a capital raise of $27.5 million (before costs) to support the ongoing clinical activities and additional planned activities, funding the Company into 2027. The capital raise comprised a successful institutional placement raising A$25.0 million (before costs) and a Share Purchase Plan seeking to raise an additional $2.5 million. The Placement was strongly supported by existing and new institutional and sophisticated investors in Australia and offshore.

Financial update

Amplia finished the September 2025 quarter with a cash position of $29.2 million (June 2025: $7.0 million). The Company also received its Research and Development Tax Incentive refund for the year ended 31 March 2025 of $3.8m in October 2025, giving Amplia a $33.0 million proforma cash balance on 30 September 2025.

During the quarter, the Company had net operating cash outflows of $3.8 million in relation to operating activities (June 2025: $3.8 million). Operating cashflows included:

- Outflows of $1.1 million for staff and administration/corporate costs; and
- Outflows of $2.6 million for research and development costs, which primarily related to trial costs, Contract Research Organisation (CRO), manufacturing and other CMC related costs incurred in relation to the ACCENT Phase 2 clinical trial for narmafotinib with gemcitabine and Abraxane® and initiation costs with its FOLFIRINOX clinical trial for narmafotinib combined with FOLFIRINOX.

Payments to Related Entities

In accordance with Listing Rule 4.7C, payments made to related parties and their associates included in item 6.1 of the Appendix 4C incorporates directors' fees, salaries and superannuation. Total payments made for the quarter equals $172,500 and relate to payments to the CEO/Managing Director in line with employment contracts and payments to the Non-Executive Directors.

- End -

For Further Information

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This ASX announcement was approved and authorised for release by the Board of Amplia Therapeutics.

About Amplia Therapeutics Limited

Amplia Therapeutics Limited is an Australian pharmaceutical company advancing a pipeline of Focal Adhesion Kinase (FAK) inhibitors for cancer and fibrosis. FAK is an increasingly important target in the field of cancer and Amplia has a particular development focus in fibrotic cancers such as pancreatic and ovarian cancer. FAK also plays a significant role in a number of chronic diseases, such as idiopathic pulmonary fibrosis (IPF). For more information visit www.ampliatx.com and follow Amplia on X (@ampliatx) and LinkedIn.

About Narmafotinib

Narmafotinib (AMP945) is the company's best-in-class inhibitor of the protein FAK, a protein over-expressed in pancreatic cancer and a drug target gaining increasing attention for its role in solid tumours. The drug, which is a highly potent and selective inhibitor of FAK, has shown promising data in a range of preclinical cancer studies. Narmafotinib is currently undergoing a clinical trial (the ACCENT trial) where it is dosed in combination with the chemotherapies gemcitabine and Abraxane® in first-line patients with advanced pancreatic cancer. The trial has already achieved its desired outcome in achieving a response rate of 33%, superior to chemotherapy alone and an interim mPFS of 7.6 months has been reported. A second trial – AMPLICITY – has recently opened and is being run under an IND at sites in Australia and the US, investigating the combination of narmafotinib with the chemotherapy FOLFIRINOX in advanced pancreatic cancer patients.

Appendix 4C

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

Name of entity

AMPLIA THERAPEUTICS LIMITED

ABN	**Quarter ended ("current quarter")**
16 165 160 841	30 September 2025

Consolidated statement of cash flows		**Current quarter $A'000**	**Year to date (6 months) $A'000**
1.	**Cash flows from operating activities**		
1.1	Receipts from customers	-	-
1.2	Payments for		
	(a) research and development	(2,600)	(5,588)
	(b) product manufacturing and operating costs	-	-
	(c) advertising and marketing	-	-
	(d) leased assets	-	-
	(e) staff costs	(373)	(912)
	(f) administration and corporate costs	(760)	(1,116)
1.3	Dividends received (see note 3)	-	-
1.4	Interest received	68	156
1.5	Interest and other costs of finance paid	-	-
1.6	Income taxes paid	-	-
1.7	Government grants and tax incentives	-	-
1.8	Other (payment of GST)	(169)	(183)
1.9	**Net cash from / (used in) operating activities**	**(3,834)**	**(7,643)**

2.	**Cash flows from investing activities**		
2.1	Payments to acquire or for:		
	(a) entities	-	-
	(b) businesses	-	-
	(c) property, plant and equipment	(21)	(26)
	(d) investments	-	-
	(e) intellectual property	-	-
	(f) other non-current assets	-	-

Consolidated statement of cash flows		Current quarter $A'000	Year to date (6 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 (bank guarantee and security deposit)	-	(64)
2.6	**Net cash from / (used in) investing activities**	**(21)**	**(90)**

3.	**Cash flows from financing activities**		
3.1	Proceeds from issues of equity securities (excluding convertible debt securities)	27,647	27,647
3.2	Proceeds from issue of convertible debt securities	-	-
3.3	Proceeds from exercise of options	238	238
3.4	Transaction costs related to issues of equity securities or convertible debt securities	(1,809)	(1,809)
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 (repayment of lease liability)	(18)	(32)
3.10	**Net cash from / (used in) financing activities**	**26,058**	**26,044**

4.	**Net increase / (decrease) in cash and cash equivalents for the period**		
4.1	Cash and cash equivalents at beginning of period	6,963	10,863
4.2	Net cash from / (used in) operating activities (item 1.9 above)	(3,834)	(7,643)
4.3	Net cash from / (used in) investing activities (item 2.6 above)	(21)	(90)

Consolidated statement of cash flows		Current quarter $A'000	Year to date (6 months) $A'000
4.4	Net cash from / (used in) financing activities (item 3.10 above)	26,058	26,044
4.5	Effect of movement in exchange rates on cash held	(10)	(18)
4.6	**Cash and cash equivalents at end of period**	**29,156**	**29,156**

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	2,428	1,338
5.2	Call deposits	26,728	5,625
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)**	**29,156**	**6,963**

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	173
6.2	Aggregate amount of payments to related parties and their associates included in item 2	-

Note: if any amounts are shown in items 6.1 or 6.2, your quarterly activity report must include a description of, and an explanation for, such payments.

The amount at 6.1 includes Director fees and salary (including superannuation) for the CEO and Managing Director and Non-Executive Directors.

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 quarter end**	-

7.6 Include in the box below a description of each facility above, including the lender, interest rate, maturity date and whether it is secured or unsecured. If any additional financing facilities have been entered into or are proposed to be entered into after quarter end, include a note providing details of those facilities as well.

N/A

8.	**Estimated cash available for future operating activities**	**$A'000**
8.1	Net cash from / (used in) operating activities (item 1.9)	(3,834)
8.2	Cash and cash equivalents at quarter end (item 4.6)	29,156
8.3	Unused finance facilities available at quarter end (item 7.5)	-
8.4	Total available funding (item 8.2 + item 8.3)	29,156
8.5	**Estimated quarters of funding available (item 8.4 divided by item 8.1)**	**7.6**

Note: if the entity has reported positive net operating cash flows in item 1.9, answer item 8.5 as "N/A". Otherwise, a figure for the estimated quarters of funding available must be included in item 8.5.

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: N/A

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: N/A

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: N/A

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

1 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.

29 October 2025

Date: .....

The Board of Directors

Authorised by: .....
(Name of body or officer authorising release – see note 4)

Notes

1. 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.