

ASX RELEASE 20 December 2022

LETTER TO SHAREHOLDERS: END OF YEAR SUMMARY AND PLANS FOR 2023

Dear Shareholder,

As 2022 draws to a close, I would like to take a moment to extend my most sincere gratitude to each and every one of Amplia Therapeutics' (ASX:ATX) supporters. We are backed by an enviable network of investors, corporate leaders, clinical and scientific experts, and other stakeholders who are truly invested in our journey.

Over the last 12 months, the Amplia team has executed our clinical and business development program with purpose and enthusiasm, bringing us closer to changing the treatment landscape for people diagnosed with cancer and fibrotic diseases.

As a long-standing member of the Board, and recently appointed CEO and MD, it has been extremely encouraging to witness this energy in action, and I am confident that we are on track for a successful and transformative year ahead.

Investor presentation

We have prepared an Investor Presentation showcasing some of the highlights of the last 12 months, and outlining our plans for 2023, appended here and available on our website (https://www.ampliatx.com/site/investor-information/current-presentations)

Key achievements in 2022 include:

- Following ethics clearance, the Company <u>commenced recruitment into the ACCENT trial</u> of our lead drug candidate, AMP945 in first-line pancreatic cancer patients. The response to the trial has been extremely positive, and <u>recruitment for the first cohort is now complete</u>.
- Presented the design and rationale for the ACCENT trial to a global audience at the <u>American Association for Cancer Research (AACR) Special Conference on Pancreatic Cancer³</u>
- Released <u>new preclinical data showing comparable efficacy of AMP945 to current standard of care in an animal model of lung fibrosis</u>⁴, providing compelling encouragement for potential clinical trials of AMP945 in idiopathic pulmonary fibrosis.
- Progressed the clinical development of our second FAK inhibitor, AMP886, including the <u>report</u> of preclinical data demonstrating its potential as a treatment for acute myeloid leukemia⁵.
- Received a \$1.8m R&D tax refund⁶ as part of the Australian Government's Research and Development Tax Incentive Refund scheme.

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¹ ASX Announcement – First Patient Recruited to ACCENT Trial in Pancreatic Cancer, 2 August 2022

² ASX Announcement – ACCENT Trial Recruitment Progress, 29 November 2022

³ ASX Announcement – Presentation at Major Pancreatic Cancer Conference, 25 July 2022

⁴ ASX Announcement – AMP945 Shows Efficacy in Model of Lung Fibrosis, 2 June 2022

⁵ ASX Announcement – AMP886 Activity in Acute Myeloid Leukemia (AML), 3 October 2022

⁶ ASX Announcement – Amplia Receives \$1.8m R&D Tax Incentive

Amplia in the News

The progress of our ACCENT Clinical Trial has consistently captured the attention of top tier and industry press this year. A definite highlight was a story filmed with Channel 9 News and the Garvan Institute, showcasing the scientific potential of AMP945.

Click here to watch the Channel 9 News story (https://www.youtube.com/watch?v=sZnOG7KgLLM)

I anticipate plenty of positive news to share with our shareholders as we continue to progress the ACCENT trial, and move through our planned clinical and corporate milestones. I thank you for your support as we continue this journey.

I wish you and your loved ones a safe and restful break over the festive season.

With thanks,

Dr Chris Burns

Amplia CEO and MD

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

- End -

For Further Information

info@ampliatx.com www.ampliatx.com

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 cancer and Amplia has a particular development focus in fibrotic tumours such as pancreatic and ovarian cancers. FAK also plays a significant role in a number of chronic diseases, such as idiopathic pulmonary fibrosis (IPF) and the Company is also developing its FAK inhibitors in these indications.



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2022 Year in Review and Growth Plans for 2023

December 2022



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Milestones acomplished in 2022

- Commenced recruitment of patients for ACCENT trial in Pancreatic Cancer
- Recruitment for first cohort of patients completed
- Demonstrated increased survival in preclinical Pancreatic Cancer model
- Completed 90 day toxicology study to support
 3 month clinical dosing
- Positive results in comparison with market leader in pre-clinical pulmonary fibrosis model
- Positive results of second FAK inhibitor,
 AMP886, in preclinical model of acute myeloid leukemia

Pancreatic Cancer



ACCENT Clinical Trial

- Apr: Received HREC approval
- May: Pre-IND meeting completed
- Aug: First patient recruited
- Sep: ACCENT trial presented at AACR
- Nov: First dosing cohort recruitment complete

Pre-clinical Studies

• Feb: Increased survival demonstrated in human pancreatic cancer model

Chemistry, Manufacturing, Controls

- Jul: Completed manufacture of 5kg batch of AMP945 demonstrating that scale up could be achieved
- Feb: Completed manufacture of first batch of capsules
- May: Completed manufacture of remaining capsules required for trial

IPF Idiopathic Pulmonary Fibrosis



Toxicology Studies

 Aug: Completed 90 day toxicology studies that support chronic dosing studies for up to 3 months

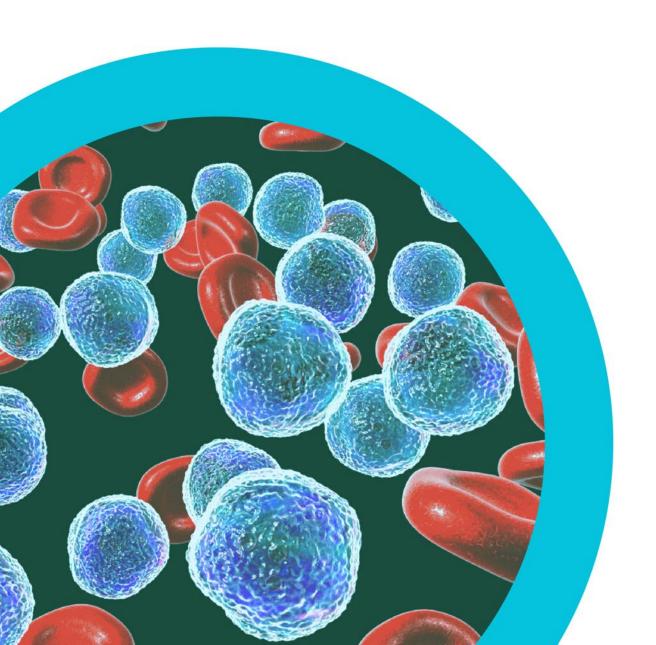
Pre-clinical Studies

• **Jul**: AMP945 shown to be as effective as current market leader Ofev® (nintedanib) in industry-standard model of IPF

Chemistry, Manufacturing, Controls

 Aug: Manufactured sufficient capsules for a 3 month clinical trial

AML Acute Myeloid Leukaemia



Pre-clinical Studies

 Oct: AMP886, a second FAK inhibitor, was shown to inhibit acute myeloid leukemia (AML) in an industry-standard model of disease

Chemistry, Manufacturing, Controls

Aug: Sufficient AMP886 has been manufactured for pre-clinical studies

New Team Members

We have continued to grow the Amplia Team to increase our expertise across the breadth of pre-clinical and clinical studies, and manufacturing



Chris Burns
CEO and Managing
Director

November 2022



Anthony Bishop

Principal Development

Manager

February 2022



Terrie-Anne Cock
Head of Translational
Biology

July 2022



May 2022

CMC Project Manager



Growth Plans for 2023

- Complete dose selection for ACCENT Trial
- Commence Part B of ACCENT Trial
- Commence process improvement for manufacturing
- Submit IND to US FDA
- Progress towards a second clinical trial with AMP945
- Continue to progress the development of AMP886



Thank You.

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