

ASX RELEASE

27 March 2025

ENCOURAGING UPDATED DATA FROM PANCREATIC CANCER TRIAL

HIGHLIGHTS

- *Encouraging updated data from the ongoing ACCENT trial, evaluating narmafotinib with chemotherapy for advanced pancreatic cancer*
- *13 Confirmed partial responses have been observed to date*
- *Narmafotinib in combination with chemotherapy demonstrates good tolerability, with adverse event profiles similar to chemotherapy alone*
- *Additional ACCENT trial data has been accepted for presentation at the American Association of Cancer Research Annual Meeting in April*

Melbourne, Australia: Amplia Therapeutics Limited (ASX: ATX), (“Amplia” or the “Company”), is pleased to report updated data from the Company’s ongoing Phase 2a clinical trial. The [ACCENT trial](#) is investigating the Company’s best-in-class FAK inhibitor narmafotinib in combination with standard-of-care chemotherapy in the treatment of advanced pancreatic cancer. This latest data analysis includes all reported data up to 12 March 2025 and builds on the previous interim data release¹ which covered data up to 6 December 2024.

Recruitment for the Phase 2a trial began in January 2024 and was completed ahead of schedule in January this year. A total of 55 advanced pancreatic patients have enrolled in the study at the trial sites in Australia and Korea. The trial is fully enrolled and as of 12 March 2025, 29 patients remain on study.

The primary endpoint for the study is the objective response rate where data for the narmafotinib + chemotherapy combination is compared against historical data for chemotherapy treatment alone. At this time, 13 confirmed partial responses (PRs) have been observed (defined as a >30% reduction in tumour size sustained over two months with no new lesions apparent), noting that tumour shrinkage assessments for the majority of patients enrolled since December are yet to be undertaken. An outcome of 15 PR’s or more, by the end of the study, would indicate the narmafotinib combination performs better than chemotherapy alone. The drug continues to be well tolerated by patients with the rate and type of adverse events for the narmafotinib combination being very similar to that reported for chemotherapy alone.

Further data from the ACCENT trial has been accepted for presentation at the annual meeting of American Association of Cancer Research to be held next month in Chicago. The poster, *Narmafotinib (AMP945) in combination with gemcitabine and nab-paclitaxel in first-line patients with advanced pancreatic cancer (ACCENT trial) a Phase 1b 2a study: Interim analysis Part B*, will be presented on Monday 28 April at 2pm (US CDT), and will be released to the market on that date.

Amplia CEO and MD Dr Chris Burns commented: “Data from the ACCENT trial continues to be extremely positive, with promising efficacy and good tolerability observed to date, and with tumour responses still being measured for the patients remaining on study. Our scheduled presentation of additional data at AACR, one of the world’s premier cancer meetings, allows us to engage with

¹ ASX Release 11 December 2024

clinicians and scientists about the considerable potential narmafotinib is demonstrating in the treatment of pancreatic cancer.”

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

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.

About the ACCENT Trial

The ACCENT trial is entitled ‘*A Phase 1b/2a, Multicentre, Open Label Study of the Pharmacokinetics, Safety and Efficacy of AMP945 in Combination with Nab-paclitaxel and Gemcitabine in Pancreatic Cancer Patients*’.

The trial is a single-arm open label study conducted in two stages. The first stage (Phase 1b), completed in November 2023, determined an optimal dose of narmafotinib (AMP945) by assessing the safety, tolerability, pharmacokinetics and preliminary efficacy when dosed in combination with gemcitabine and Abraxane in first-line patients with advanced pancreatic cancer.

The second stage (Phase 2a) of the trial is designed to assess efficacy in combination with gemcitabine and Abraxane. The primary endpoints are Objective Response Rate (ORR) and Duration on Trial (DOT) with secondary endpoints being Progression Free Survival (PFS) and Overall Survival (OS). Safety and tolerability will continue to be assessed.

The trial is being conducted at seven sites in Australia and five sites in South Korea.

More information about the ACCENT trial can be found via the ACCENT trial [site](#), the Amplia Therapeutics [website](#) and at ClinicalTrials.gov under the identifier [NCT05355298](#).

The Company will provide further updates on the trial as data is accrued.

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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 [Twitter](#) (@ampliatx) and [LinkedIn](#).