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ASX RELEASE

AMPLIA AND ANZGOG FINALISE PROTOCOL FOR OVARIAN CANCER CLINICAL TRIAL

HIGHLIGHTS

- *The PRROSE trial will investigate the combination of narmafotinib with standard of care chemotherapy in patients with advanced ovarian cancer*
- *Following close collaboration between ANZGOG and Amplia the clinical trial protocol has been finalised*
- *Ethics and site governance approvals will be submitted with the goal of opening trial sites before the end of 2026*

Melbourne, Australia: Amplia Therapeutics Limited (ASX:ATX; OTCQB:INNMF), ("Amplia" or the "Company"), and the Australia New Zealand Gynaecological Oncology Group (ANZGOG) are pleased to announce that the clinical trial protocol for the planned PRROSE Phase 2 ovarian cancer study has been finalised, marking an important milestone toward study commencement.

The ANZGOG and Amplia teams have worked collaboratively over the last three months to finalise the PRROSE trial protocol. With this completed, submission for ethics and site governance approvals will be undertaken, with site activation and patient recruitment planned before the end of 2026.

The PRROSE trial will explore the combination of Amplia's FAK inhibitor, narmafotinib, in combination with standard-of-care chemotherapy, in women with high-grade serous ovarian cancer (HGSOC). The study will focus on patients whose tumours demonstrate a poor response to chemotherapy prior to planned interval debulking surgery, a group with a significant unmet medical need.

The study will be led by Dr Gwo Yaw Ho of Monash Health and Monash University and will be sponsored and coordinated by ANZGOG. Approximately 26 patients across selected clinical sites in Australia and New Zealand are expected to be enrolled into the trial.

Dr Chris Burns, CEO and Managing Director of Amplia, commented: "This is an important study for a patient population who currently have limited treatment options. We are grateful to the ANZGOG team, working closely with our clinical team at Amplia, for their diligent efforts to complete the trial protocol. We now look forward to finalising ethics submissions with the goal of opening trial sites before the end of the year."

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Dr Gwo Yaw Ho, Lead Investigator, said: "Completion of the study protocol enables us to move towards evaluating this promising therapeutic approach in patients with high-grade serous ovarian cancer who are at increased risk of poor outcomes. We look forward to initiating the study and exploring the potential clinical benefit of combining narmafotinib with chemotherapy."

The PRROSE study was first announced in May 2026[1] and is designed to explore the potential role of FAK inhibition in improving outcomes for patients with ovarian cancer.

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

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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 X (@ampliatx) and LinkedIn.

About the Australia New Zealand Gynaecological Oncology Group (ANZGOG)

ANZGOG is the peak national gynaecological cancer research organisation in Australia and New Zealand. It's more than 1650 members work in hospitals, research institutions, universities, government, the non-profit sector, and industry. Members include surgeons, physicians, radiation oncologists, psychologists, social workers, nurses, trial coordinators, pure researchers, allied health professionals and cancer consumers who are based in all states of Australia and regions in New Zealand. ANZGOG's vision is ***Advancing research, saving lives*** and its mission is to improve outcomes and quality of life for everyone with a lived experience of gynaecological cancer by conducting and promoting clinical trials and multidisciplinary research. For more information, visit www.anzgog.org.au and follow ANZGOG on LinkedIn

About the PRROSE trial

The PRROSE trial will evaluate the safety of narmafotinib in combination with standard-of-care chemotherapy (carboplatin and paclitaxel) in HGSOC patients who have demonstrated poor response to up-front standard-of-care platinum-based chemotherapy prior to planned interval debulking surgery. Approximately one in five ovarian cancer patients do not respond adequately to initial chemotherapy,

[1] ASX Announcement May 8 2026

limiting their ability to undergo surgery and contributing to poor clinical outcomes. This study is designed to address this significant unmet medical need. The study will therefore also explore whether the addition of narmafotinib can increase the proportion of patients eligible for successful surgical resection.

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 achieved its desired outcome in achieving an objective response rate of 36%, superior to chemotherapy alone, and a mOS of 11.1 months has been reported.