

## Appendix 4C quarterly webinar – 11:00am AEST Friday 1 August

Syntara Limited (ASX:SNT), a clinical-stage drug development company, is pleased to announce that it will hold an investor webinar following the announcement of the Company's Appendix 4C and quarterly update.

CEO Gary Phillips will deliver an update and investor presentation as part of the webinar, to be held at 11:00am AEST Friday 1 August 2025.

Shareholders, investors and interested parties are encouraged to register to attend the presentation at the following link:

[https://us02web.zoom.us/webinar/register/WN\\_9SKnUn9mQ3OTn0YYIN12gQ](https://us02web.zoom.us/webinar/register/WN_9SKnUn9mQ3OTn0YYIN12gQ)

After registering, you will receive a confirmation email containing information about joining the webinar as well as dial-in details for those that wish to join by phone.

Questions can be submitted on the day or sent in advance to [benl@nwrcommunications.com.au](mailto:benl@nwrcommunications.com.au)

Please note a replay of the webinar will be available at the above-mentioned link shortly following the conclusion of the live session.

Syntara notes that its securities will be suspended from quotation today as it did not lodge its periodic report for the period ended 30 June 2025 by the relevant due date. While Syntara's Quarterly Shareholder Report was filed on 30 July 2025, the associated Appendix 4C Quarterly Cash Flow Report was not lodged at the same time due to an administrative error. The Company has since lodged the Appendix 4C on 31 July 2025 and expects the suspension of securities to be lifted on Monday 4 August 2025.

#ENDS#

**SOURCE:**

Syntara Limited (ASX: SNT),  
Sydney, Australia  
(ABN: 75 082 811 630)

**AUTHORISED FOR RELEASE TO ASX BY:**

Syntara Limited Disclosure Committee.

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## About Syntara

Syntara Limited (ABN: 75 082 811 630) is a clinical stage drug development company targeting extracellular matrix dysfunction with its world-leading expertise in amine oxidase chemistry and other technologies to develop novel medicines for blood cancers and conditions linked to inflammation and fibrosis.

Lead candidate amsulostat (also known as SNT-5505 and previously as PXS-5505) is for the bone marrow cancer myelofibrosis which causes a build-up of scar tissue that leads to loss of red and white blood cells and platelets. Amsulostat has recently been granted Fast Track Designation, having already achieved FDA Orphan Drug Designation and clearance under an Investigational New Drug Application for development in myelofibrosis. After encouraging phase 2a trial results when used as a monotherapy in myelofibrosis, amsulostat is now being studied with a JAK inhibitor in a suboptimal response setting. Protocols for another two Phase 1c/2 studies with amsulostat in patients with a blood cancer called myelodysplastic syndrome are in development and expected to commence recruitment by H1 2025.

Syntara is also advancing topical pan-LOX inhibitors with SNT-9465 in a Phase 1a/b study of hypertrophic scars and continuing the ongoing collaboration with Professor Fiona Wood and the University of Western Australia studying SNT-6302 in keloid scars. SNT-4728 is being studied in collaboration with Parkinson's UK as a best-in-class SSAO/MAO-B inhibitor to treat sleep disorders and slow progression of neurodegenerative diseases like Parkinson's by reducing neuroinflammation. Other Syntara drug candidates target fibrotic and inflammatory diseases such as kidney fibrosis, MASH, pulmonary fibrosis and cardiac fibrosis.

Syntara developed two respiratory products available in world markets (Bronchitol® for cystic fibrosis and Aridol®- a lung function test), which it sold in October 2023.

Syntara is listed on the Australian Securities Exchange, code SNT. The company's management and scientific discovery team are based in Sydney, Australia. [www.syntaraTX.com.au](http://www.syntaraTX.com.au).

## Forward-Looking Statements

Forward-looking statements in this media release include statements regarding our expectations, beliefs, hopes, goals, intentions, initiatives or strategies, including statements regarding the potential of products and drug candidates. All forward-looking statements included in this media release are based upon information available to us as of the date hereof. Actual results, performance or achievements could be significantly different from those expressed in, or implied by, these forward-looking statements. These forward-looking statements are not guarantees or predictions of future results, levels of performance, and involve known and unknown risks, uncertainties and other factors, many of which are beyond our control, and which may cause actual results to differ materially from those expressed in the statements contained in this document. For example, despite our efforts there is no certainty that we will be successful in partnering any of the products in our pipeline on commercially acceptable terms, in a timely fashion or at all. Except as required by law we undertake no obligation to update these forward-looking statements as a result of new information, future events or otherwise.