Media Release

2 February 2024

Syntara share purchase plan

Clinical stage drug development company Syntara Limited (ASX: **SNT**) today advised that the Syntara Limited Share Purchase Plan (SPP) announced on 19th December 2023 closed on 30th January 2024 with eligible applications for $303,000.

Syntara further advises that it will accept all eligible applications and issue 13,771,861 fully paid shares that will rank equally with existing shares from allotment on 6 February 2024.

The new shares issued under the SPP will be issued at $0.022 per share, being the same price that was paid by sophisticated and institutional investors under the placement announced on 19th December 2023.

Gary Phillips, Chief Executive Officer commented, "The total of $10.3[1] million raised from the SPP and placement significantly strengthens the Syntara balance sheet. When combined with the expense reductions flowing from exiting the mannitol business unit, the Company now has a cash runway through to results from several clinical trials in mid-2025."

1. Before costs of the offer

#ENDS#

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

AUTHORISED FOR RELEASE TO ASX BY:
David McGarvey, Chief Financial Officer, and Company Secretary:

+61 2 9454 7200,
david.mcgarvey@syntaraTX.com.au

CONTACT:

Syntara Media:
Felicity Moffatt:
+61 418 677 701
felicity.moffatt@syntaraTX.com.au

Syntara Investor relations:
Rudi Michelson
(Monsoon Communications)
+61 411 402 737
rudim@monsoon.com.au

JOIN THE SYNTARA MAILING LIST HERE

About Syntara

Syntara Limited (ABN: 75 082 811 630) is a clinical stage drug development company with a focus on blood-related cancers. The company's highly productive drug discovery engine is driven by its expertise in amine oxidase inhibitors.

Syntara is managing three phase 2 clinical studies in diseases of high unmet need with a further two potential phase 1c/2 studies being evaluated for 2024. Lead candidate SNT-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. SNT-5505 has 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, PXS-5505 is now being studied with a JAK inhibitor in a further phase 2 myelofibrosis study with interim data by Q4 2024.

Syntara is also advancing both oral and topical pan-LOX inhibitors in scar prevention and scar modification programs as part of an ongoing collaboration with Professor Fiona Wood and the University of Western Australia. SNT-4728 is being studied in collaboration with Parkinson's UK as a best-in-class SSAO/MAOB 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, NASH, pulmonary fibrosis and cardiac fibrosis.

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

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.

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.