PTX Strongly Supported Placement Raises $7 million to Fund Expanded Pipeline

HIGHLIGHTS:

- **Strongly supported $7 million placement follows oversubscribed $6.5 million SPP bringing total funds raised to $13.5 million**
- **Placement and SPP bring pro forma cash on hand to over $20.3 million[1]**
- **PTX now appropriately funded to vigorously advance its expanded pipeline of cell therapy and CAR-T, in addition to targeted therapies PTX-100 and PTX-200**

MELBOURNE Australia, 26 August 2020 – Prescient Therapeutics (ASX: PTX) ("**Prescient**" or the "**Company**"), a biotechnology company developing personalised medicines for cancer, is pleased to announce that it has received firm commitments to raise $7,046,084 (before costs) by way of a share placement (**Placement**) to professional and sophisticated investors of 128,110,611 fully paid ordinary shares (**Placement Shares**) at $0.055 (5.5 cents) per share, being the same issue price as the recently completed Share Purchase Plan (**SPP**).

Demand for the Placement was secured last week, immediately following completion of the SPP, and significantly exceeded the Company's placement capacity. The SPP and Placement have raised a combined $13.5 million before costs. Following settlement of the Placement, Prescient will have over $20.3 million[1] cash on hand.

Prescient CEO and Managing Director Steven Yatomi-Clarke said, "We are delighted by the strong response to the Placement, which in part reflected the excess demand for the SPP. Both existing and new shareholders have shown that they share Prescient's enthusiasm for its expanded pipeline of innovative cancer treatments."

"In addition to progressing targeted therapies and Cell Therapy Enhancement programs, these funds will also be applied towards the development of OmniCAR, a next generation CAR-T platform that aims to significantly broaden CAR-T's addressable market, while overcoming many limitations of existing CAR-T therapies. These represent tremendous market opportunities. We welcome new shareholders as they join us on our journey to develop innovative and personalised cancer therapies."

Details of Placement

Prescient will issue 128,110,611 shares utilising the Company's placement capacity under Listing Rule 7.1 (76,866,367 shares) and 7.1A (51,244,245 shares). The Company is not utilising the temporary ASX facility of additional 10% capacity under Listing Rule 7.1.

[1] *Based on unaudited current cash balance including proceeds from Placement and SPP (excluding issue costs).*

The Placement shares will be issued on or around Friday, 28 August 2020 subject to receipt of funds on Thursday, 27 August 2020. The Placement shares will rank equally with existing fully paid shares of the Company. The Placement was managed by the Company.

Use of Funds

The funds raised from the Placement and SPP will be utilised for development of targeted therapies; Cell Therapy Enhancements; the development of Prescient's next-generation CAR-T platform, OmniCAR; working capital; and costs of the offer.

To stay updated with the latest PTX news click here:
https://prescienttherapeutics.investorportal.com.au/register

– Ends –

About Prescient Therapeutics Limited (Prescient)

Prescient Therapeutics is a clinical stage oncology company developing personalised medicine approaches to cancer, including targeted and cellular therapies.

Cell Therapies

OmniCAR: is a universal immune receptor platform enabling controllable T-cell activity and multi-antigen targeting with a single cell product. OmniCAR's modular CAR system decouples antigen recognition from the T-cell signalling domain. It is the first universal immune receptor allowing post-translational covalent loading of binders to T-cells. OmniCAR is based on technology licensed from Penn; the SpyTag/SpyCatcher binding system licensed from Oxford University; and other assets.
The targeting ligand can be administered separately to CAR-T cells, creating on-demand T-cell activity post infusion and enables the CAR-T to be directed to an array of different tumour antigens.
OmniCAR provides a method for single-vector, single cell product targeting of multiple antigens simultaneous or sequentially, whilst allowing continual re-arming to generate, regulate and diversify a sustained T-cell response over time.

Cell Therapy: Prescient has several other initiatives underway to develop new cell therapy approaches.

Targeted Therapies

PTX-100 is a first in class compound with the ability to block an important cancer growth enzyme known as geranylgeranyl transferase-1 (GGT-1). It disrupts oncogenic Ras pathways by inhibiting the activation of Rho, Rac and Ral circuits in cancer cells, leading to apoptosis (death) of cancer cells. PTX-100 is believed to be the only RhoA inhibitor in the world in clinical development. PTX-100 is currently in a PK/PD basket study of hematological and solid malignancies, focusing on cancers with Ras and RhoA mutations. In a previous Phase 1 trial in advanced solid tumours, PTX-100 was well tolerated and achieved stable disease.

PTX-200 is a novel PH domain inhibitor that inhibits an important tumour survival pathway known as Akt, which plays a key role in the development of many cancers, including breast and ovarian cancer, as well as leukemia. Unlike other drug candidates that target Akt inhibition which are non-specific kinase inhibitors that have toxicity problems, PTX-200 has a novel mechanism of action that specifically inhibits Akt whilst being comparatively safer. This highly promising compound has encouraging Phase 2a data in HER2-negative breast cancer; Phase 1b/2 in relapsed and refractory AML and Phase 1b in recurrent or persistent platinum resistant ovarian cancer:

COVID-19 Therapies

Two assets are being assessed by the Doherty Institute for antiviral activity against SARS-CoV-2, the virus that causes COVID-19 disease.

Find out more at https://prescienttherapeutics.investorportal.com.au/, or connect with us via Twitter @PTX_AUS and LinkedIn.

The Board of Prescient Therapeutics Limited has approved the release of this announcement.

For more information please contact:

Steven Yatomi-Clarke	**Investor enquiries:**	**Media enquiries:**
CEO & Managing Director	Warrick Lace – Reach Markets	Andrew Geddes – CityPR
Prescient Therapeutics	+61 404 656 408	+61 2 9267 4511
steven@ptxtherapeutics.com	warrick.lace@reachmarkets.com.au	ageddes@citypublicrelations.com.au

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Certain statements made in this document are forward-looking statements within the meaning of the safe harbor provisions of the United States Private Securities Litigation Reform Act of 1995. These forward-looking statements are not historical facts but rather are based on the current expectations of Prescient Therapeutics Limited ("Prescient" or the "Company"), their estimates, assumptions, and projections about the industry in which Prescient operates. Material referred to in this document that use the words 'estimate', 'project', 'intend', 'expect', 'plan', 'believe', 'guidance', and similar expressions are intended to identify forward-looking statements and should be considered an at-risk statement. These forward-looking statements are not a guarantee of future performance and involve known and unknown risks and uncertainties, some of which are beyond the control of Prescient or which are difficult to predict, which could cause the actual results, performance, or achievements of Prescient to be materially different from those which may be expressed or implied by these statements. These statements are based on our management's current expectations and are subject to a number of uncertainties and risks that could change the results described in the forward-looking statements. Risks and uncertainties include, but are not limited to, general industry conditions and competition, general economic factors, global pandemics and related disruptions, the impact of pharmaceutical industry development and health care legislation in the United States and internationally, and challenges inherent in new product development. In particular, there are substantial risks in drug development including risks that studies fail to achieve an acceptable level of safety and/or efficacy. Investors should be aware that there are no assurances that results will not differ from those projected and Prescient cautions shareholders and prospective shareholders not to place undue reliance on these forward-looking statements, which reflect the view of Prescient only as of the date of this announcement. Prescient is not under a duty to update any forward-looking statement as a result of new information, future events or otherwise, except as required by law or by any appropriate regulatory authority.

Certain statements contained in this document, including, without limitation, statements containing the words "believes," "plans," "expects," "anticipates," and words of similar import, constitute "forward-looking statements." Such forward-looking statements involve known and unknown risks, uncertainties and other factors that may cause the actual results, performance or achievements of Prescient to be materially different from any future results, performance or achievements expressed or implied by such forward-looking statements. Such factors include, among others, the following: the risk that our clinical trials will be delayed and not completed on a timely basis; the risk that the results from the clinical trials are not as favourable as we anticipate; the risk that our clinical trials will be more costly than anticipated; and the risk that applicable regulatory authorities may ask for additional data, information or studies to be completed or provided prior to their approval of our products. Given these uncertainties, undue reliance should not be placed on such forward-looking statements. The Company disclaims any obligation to update any such factors or to publicly announce the results of any revisions to any of the forward-looking statements contained herein to reflect future events or developments except as required by law.

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Supplemental COVID-19 Risk Factors

Please see our website : Supplemental COVID-19 Risk Factors