

ASX Announcement | 29 March 2021 Noxopharm Limited (ASX:NOX)

Presentation for Switzer Small and Micro Cap Virtual Investor Day

Sydney 29 March 2021: Australian clinical-stage drug development company Noxopharm Limited (ASX:NOX) provides the corporate presentation for the Switzer Small and Micro Cap Virtual Investor Day to be held Tuesday 30 March 2021.

The presentation will focus on the Company's proprietary drug technology platform that is behind the Company's aim in developing a pipeline of drugs to meet the major unmet needs of cancer and septic shock. The link between these two important community ailments is a disturbance in the immune and inflammatory systems, with the Noxopharm technology platform able to address that imbalance.

The presentation reviews the significant progress made in establishing Veyonda[®] (NOX66) as a highly sought-after, next generation immuno-oncology drug, to be used to boost the anti-cancer effectiveness of most standard anti-cancer therapies including radiotherapy, chemotherapy and immune checkpoint inhibitors.

Also reviewed is the progress of the potential role in using Veyonda to block progression into septic shock, believed responsible for much long-term disability and death in COVID-19 patients.

Dr Kelly will review the Company's progress in generating the underlying proof-of-concept data towards it goal of raising both the value and international profile of Veyonda.

Dr Kelly's presentation titled, "Switzer Small & Micro Cap Virtual Investor Day Presentation March 2021" is attached.

Registration for the virtual investor day can be found by visiting: <u>https://info.switzer.com.au/2021-switzer-small-micro-cap-investor-day</u>

-ENDS-

Graham Kelly, CEO and Managing Director of Noxopharm, has approved the release of this document to the market on behalf of the Board of Directors.

About Noxopharm

Noxopharm Limited (ASX:NOX) is an Australian clinical-stage drug development company focused on the treatment of cancer and cytokine release syndrome/septic shock.



Veyonda is the Company's first pipe-line drug candidate currently in Phase 2 clinical trialing. Veyonda has three main drug actions – highly selective inhibition of sphingosine kinase, STING signaling and autophagy. Sphingosine kinase inhibition contributes to its dual-acting oncotoxic and immuno-oncology functions designed to enhance the effectiveness and safety of standard oncology treatments, i.e., chemotherapies and immune checkpoint inhibitors; STING signaling inhibition provides an anti-inflammatory effect, contributing to an anti-cancer action, but also potentially blocking sepsis; autophagy inhibition is believed to augment the immunotherapy effect of radiotherapy, in particular the triggering of an abscopal response.

Noxopharm also is the major shareholder of US biotechnology company Nyrada Inc (ASX:NYR), and wholly owns Pharmorage, a private drug development company focused on drug development in the areas of sepsis and autoimmunity.

To learn more visit: https://www.noxopharm.com/

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SWITZER SMALL & MICRO CAP VIRTUAL INVESTOR DAY March 2021



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Noxopharm is building a pipeline of drugs based on a unique and proprietary <u>technology platform</u> for the treatment of <u>cancer</u> and <u>septic shock</u>

Of est. 55 million deaths globally from disease:

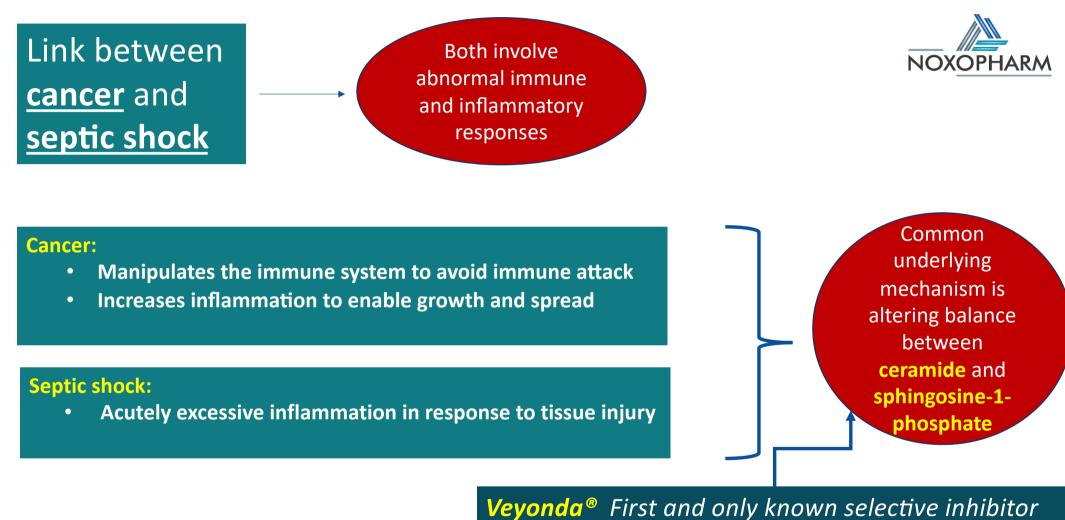
- ~ 10 million are from cancer*
- ~ 10 million are from septic shock*



* World Health Organization (2020)

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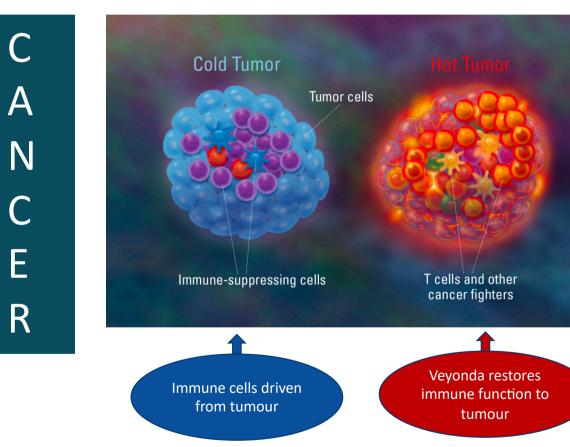




Veyonda[®] First and only known selective inhibito of sphingosine kinase 1 to enter the clinic

Veyonda[®] breakthrough <u>COLD to HOT</u> effect





Majority of human tumours are COLD (lacking immune cells) Converting **COLD** tumors to **HOT** means that standard anti-cancer treatments should work better

Veyonda being developed as a combination drug to boost effectiveness of common cancer treatments

Source: Dana-Faber Cancer Institute

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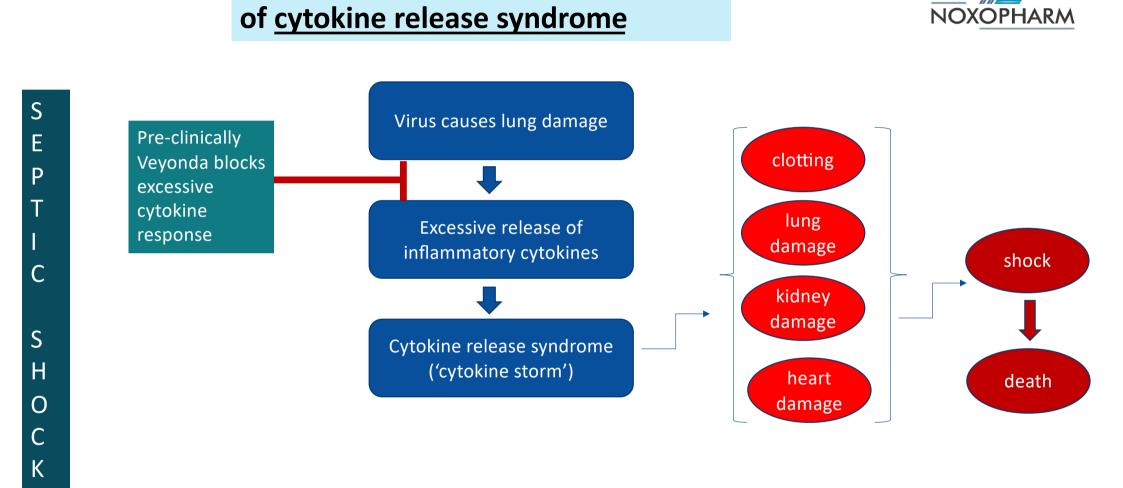
Veyonda[®] Two clinical studies confirm anti-cancer boost. 3rd key study now underway



C A	DARRT program	 Veyonda + (low-dose) external beam radiotherapy (EBRT) Phase I study / 25 men / end-stage prostate cancer Combination delivers major boost to anti-cancer effect of EBRT¹
N C F	LuPIN program	 Veyonda + ¹⁷⁷Lu-PSMA-617 (Novartis) Phase I/II study / 56 men / end-stage prostate cancer Combination delivers major boost to reports of anti-cancer effect of Novartis drug alone²
R	IONIC program	 Veyonda + Opdivo[®] (Bristol Myers Squibb) Phase I/II study / 30 patients / all forms of solid cancer Study started³

1. ASCO ASM June 2020; 2. ASCO Genitourinary Cancer Conference Feb 2021; 3. ASX announcement 19 March 2021

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Breakthrough potential treatment

Veyonda[®]

Veyonda[®] NOXCOVID-1 clinical study ongoing



NOXCOVID program

- 40 patients with moderate COVID-19 disease
- Requiring hospitalization and low-volume oxygen supplementation
- Dose-escalation arm (400-1800 mg) complete
- Dose-expansion arm (1800 mg) ongoing

Objective: to demonstrate that Veyonda

- Reduces the cytokine release syndrome (CRS)
- Reduces disease progression requiring ICU care and mechanical ventilation
- Reduces incidence of CRS-related disability and death
- Has the potential to be a cost-effective treatment for septic shock generally

SUMMARY



Both DARRT and LuPIN have provided clear evidence of proof-of-concept

IONIC trial

The objective of this trial is to use Veyonda to overcome resistance of most cancers to anti-cancer drug Opdivo[®] (Bristol Myers Squibb) currently with annual sales of USD8 billion

Veyonda is well tolerated as a combination treatment

Robust IP strategy based on PCT patents around use claims

Our commercial end-point for Veyonda

Noxopharm prepared to take Veyonda all the way to market. However, opportunity as a general booster of most forms of cancer therapy seen as being so large that a partnership potentially is more likely

Currently working with Novartis (USD 200 B) and Bristol Myers Squibb (USD 140 B) drugs in combination therapy

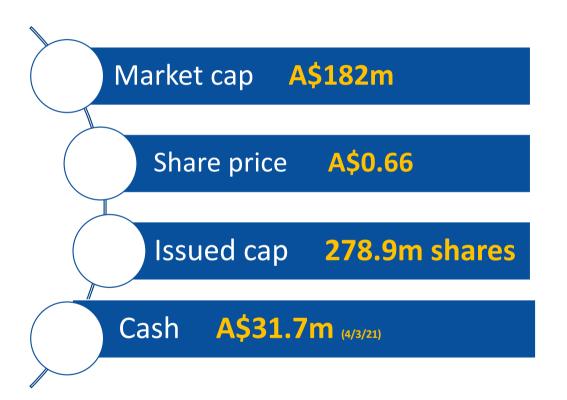
Commercial transaction seen as potentially likely during current Phase 2 trialling





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Key metrics





News Flow (next 6 months)

- IONIC-1 and DARRT-2 start patient recruitment
- COVID-19 clinical trial completion
- Growing first-in-class drug pipeline
- Pharmorage (subsidiary) progressing novel drug development for sepsis and autoimmunity

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For further information

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