

ASX Announcement |14 August 2020 Noxopharm Limited ASX:NOX

Noxopharm August 2020 EGM Corporate Presentation

Australian clinical stage drug development company Noxopharm (ASX:NOX) is pleased to provide to shareholders and the market generally the corporate presentation for today's Extraordinary General Meeting (EGM).

Following voting formalities, the Company will take the opportunity to present this corporate presentation and an update on recent progress in its R&D activities and plans for upcoming clinical trials that were the subject of the recent capital raise. Noxopharm's CEO and Founder (Dr Graham Kelly) and CMO (Dr Gisela Mautner) will present the update virtually.

This presentation can be found at: https://www.noxopharm.com/site/investors/presentations

About Noxopharm

Noxopharm Limited (ASX:NOX) is an Australian clinical-stage drug development company focused on treating cancer with Veyonda[®], its lead drug candidate.

Veyonda[®] is a dual-acting oncotoxic and immuno-oncology drug designed to enhance the effectiveness and safety of standard oncology treatments, i.e., chemotherapy, radiotherapy and immuno-oncology drugs. The drug acts by harnessing the body's immune system to inflict damage on cancer cells in the body and has shown promise in treating a broad spectrum of cancers.

Veyonda[®] also is to undergo evaluation as a treatment for septic shock, starting with a Phase 1 study in patients with moderate COVID-19 disease.

Noxopharm also has an active research and development (R&D) program for additional drug candidates and is the major shareholder of US biotechnology company Nyrada Inc. (ASX:NYR).

To learn more please visit: <u>https://www.noxopharm.com/</u>

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

Forward Looking Statements

This announcement may contain forward-looking statements. You can identify these statements by the fact they use words such as "aim", "anticipate", "assume", "believe", "continue", "could", "estimate", "expect", "intend", "may", "plan", "predict", "project", "plan", "should", "target", "will" or "would" or the negative of such terms or other similar expressions. Forward-looking statements are based on estimates, projections and assumptions made by Noxopharm about circumstances and events that have not yet taken place. Although Noxopharm believes the forward-looking statements to be reasonable, they are not certain. Forward-looking statements involve known and unknown risks, uncertainties and other factors that are in some cases beyond the Company's control that could cause the actual results, performance or achievements to differ materially from those expressed or implied by the forward-looking statement.



Noxopharm EGM – August 2020 Presentation Noxopharm Limited (ASX:NOX)

Graham Kelly - Executive Chairman and CEO

Veyonda[®]



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Immuno-oncology (I-O) drugs have taken the oncology world by storm and are the future of cancer therapy

> Current market for I-O drugs = ~<u>A\$30 billion p.a. (2019)</u>

I-O drugs currently limited in use Potential market predicted to be <a>A\$150 billion p.a.

NOX believes that **Veyonda[®]** will be one of the main drugs driving that A\$120 billion increase in I-O use and value



This presentation

1. Who We Are

2. Veyonda[®] Explained

3. Our Business

4. Investment Case



01 Who we are

Who We Are

Australian clinical-stage drug development company

ASX: NOX [listed Aug 2016]

Focus on large markets of unmet need – cancer and septic shock

Commercial goal of out-licensing at Phase 2 stage

Veyonda[®] - a major commercial opportunity with 'blockbuster' potential

A disruptive technology transforming the management of cancer



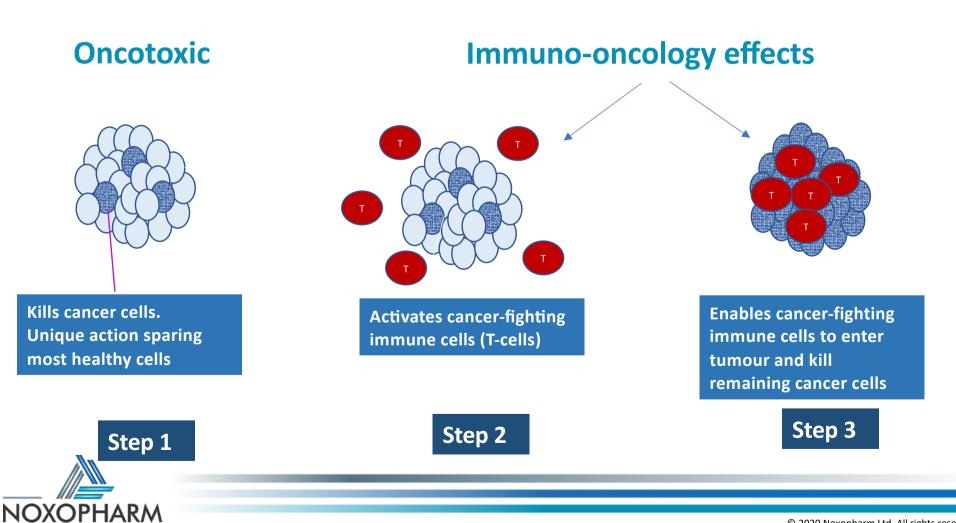
First-in-class

Veyonda[®] explained

Dual oncotoxic and immuno-oncology drug

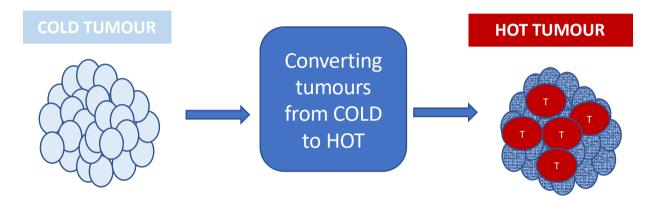
Veyonda[®] Explained

First-in-class anti-cancer drug with unique 3-step action



Veyonda[®] Explained

Increasing I-O therapy effectiveness depends on boosting the number of cancerfighting cells inside tumours. Referred to as converting tumours from COLD to HOT



Majority of human tumours are COLD:

- No active immune cells present
- Believed to be a major cause of limited effectiveness of current cancer therapies

Converting human tumours to HOT:

- Increased immune function
- Increased effectiveness of chemotherapy, radiotherapy and immuno-oncology therapy

COLD to <u>HOT</u> conversion current major goal of pharma industry Limited success to date <u>Noxopharm believes Veyonda[®] will be the 'breakthrough' drug</u>



ONCOLOGY



SEPTIC SHOCK

Clinical Objective

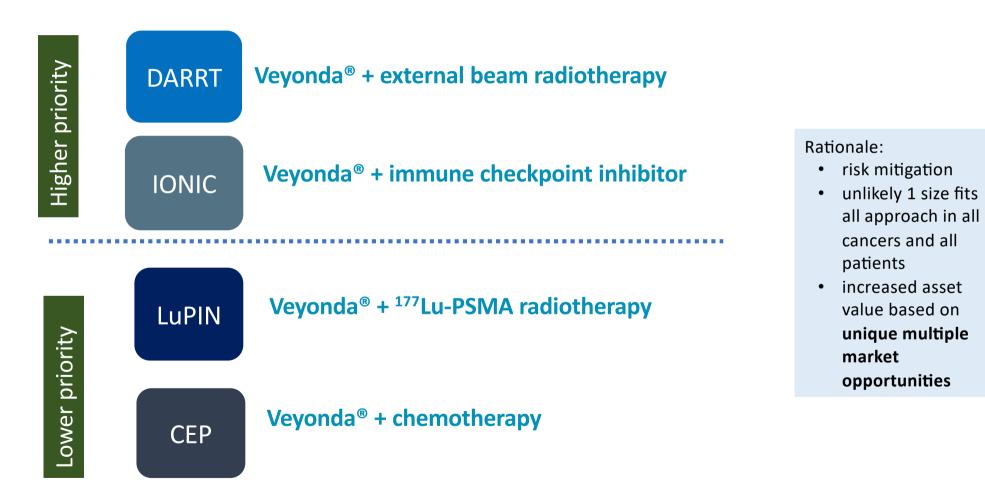
To see Veyonda[®] come to market as a well-tolerated, costeffective, immuno-oncology (I-O) drug assisting immune checkpoint inhibitors, radiotherapy and chemotherapy to work more effectively across most forms of cancer

Commercial Objective

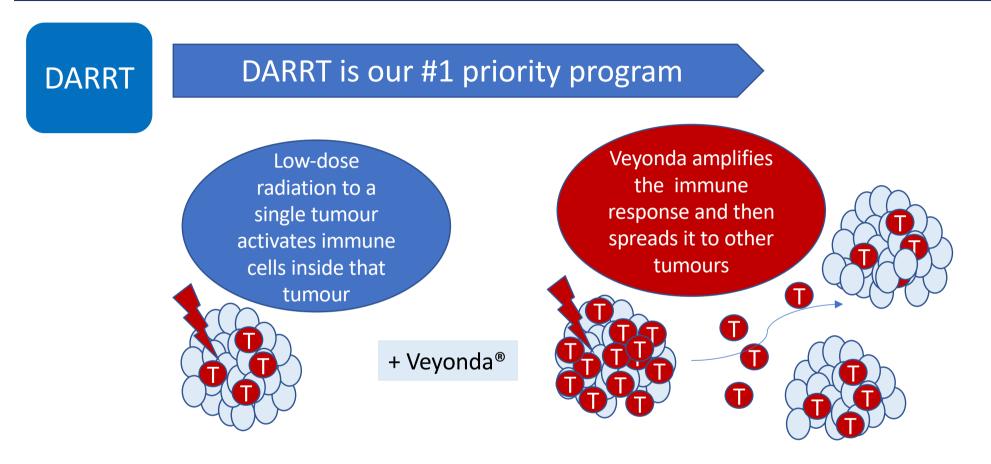
To provide comprehensive pre-clinical and clinical data packages that are compelling for 'blockbuster' trade deals



Aiming to show an I-O benefit four different ways



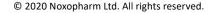




COLD → HOT effect

- Known as an 'abscopal response'. Normally an extremely rare event
- DARRT aiming to make abscopal responses commonplace





DARRT

DARRT is our #1 priority program

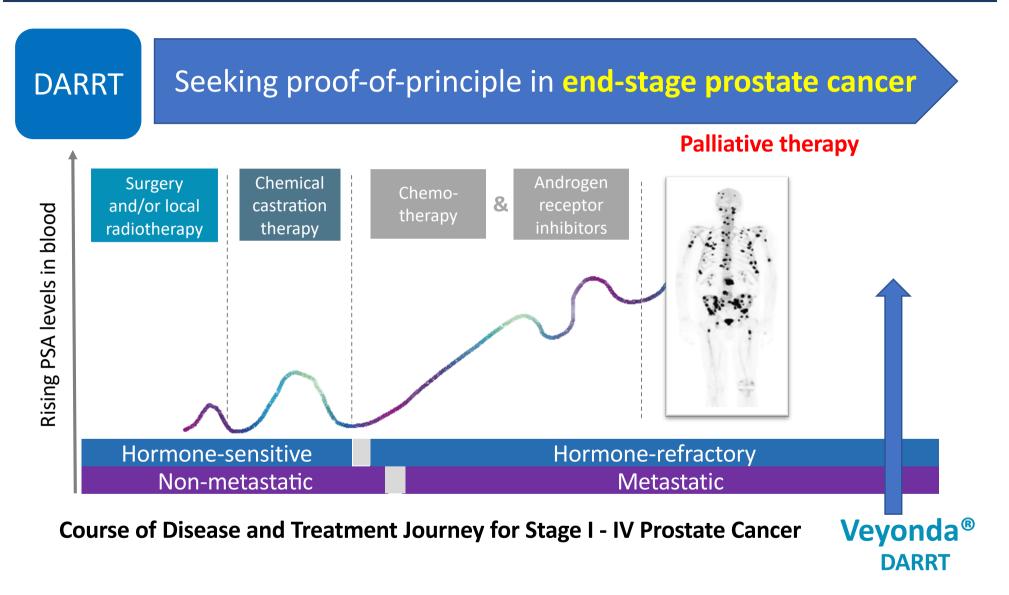
Aimed at the largest sector in the oncology market

- End-stage cancer where treatment is limited to palliative care
- Little competition
- Multi-billion dollar market opportunity

Attractive form of anti-cancer therapy

- Well-tolerated, short-course of therapy in out-patient clinic
- Most common form of radiotherapy (= low cost, ready availability)







DARRT

DARRT-1 Phase 1b study

25 men with end-stage prostate cancer who had stopped responding to treatment, with metastatic and progressive disease, and were considered to have limited life-spans

- Clear evidence of an I-O effect
- 16 men completed the 14 days of treatment and the 6month follow up period
- In 10 men, tumours had stopped growing or were reduced in size
- Abscopal responses confirmed in 4 men*
- Treatment well tolerated

 First known demonstration of abscopal responses in prostate cancer in more than isolated cases



IONIC

IONIC study is our #2 priority program

Immune checkpoint inhibitors

- Keytruda (Merck); Opdivo & Yervoy (BMS)
- Minority of cancers are responsive cancers (eg. melanoma, lung, kidney, bladder)
- Majority of cancers non-responsive (eg. breast, prostate, ovarian, bowel)
- Nevertheless, 2019 global sales ~<u>A\$30 billion</u>
- Achieving responses in majority of cancers estimated to yield market worth >A\$150 billion p.a.

Objective = by converting tumours from COLD to HOT, to use Veyonda to boost the response rates to immune checkpoint inhibitors and make Veyonda a key contributor to the ~A\$120 billion missing market opportunity

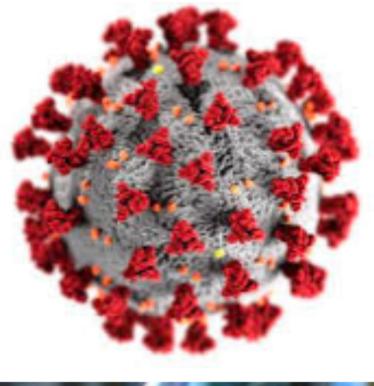


And then came SARS-CoV-2

and the pandemic

and with it, an unexpected opportunity.....

the possibility of having an effective therapy for SEPTIC SHOCK





Our Business – Veyonda[®] and septic shock

SEPTIC SHOCK

SEPTIC SHOCK: ONE OF THE SINGLE LARGEST CAUSES OF DEATH IN THE WORLD

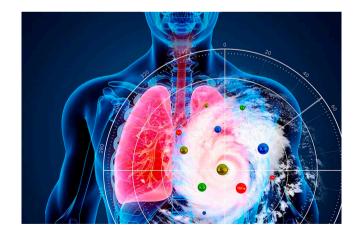
- An estimated 10 million people die each year in the world from septic shock. That is about <u>20% of all human deaths</u>
- An estimated minimum of <u>5000</u> people die from SEPTIC SHOCK each year in Australia
- SEPTIC SHOCK occurs when the body produces an excessively high immune/inflammatory response to infection (viral, bacterial) or trauma (accidents, major surgery)
- That hyper-response produces something called a <u>CYTOKINE STORM</u> resulting in multiple organ failure. This is known as **SEPTIC SHOCK**
- **SEPTIC SHOCK** is invariably fatal because we lack effective therapies
- Up to this year, the most common viral cause of SEPTIC SHOCK in Australia was seasonal influenza with up to about 3,000 deaths p.a.
- In 2020, a new viral cause has emerged SARS-CoV-2 virus



Our Business – Veyonda[®] and septic shock

NOXCOVID

<u>First</u> clinical trial of a <u>STING</u> <u>blocker</u> in COVID-19 patients



Cytokine Storm

- Septic shock is triggered by a CYTOKINE STORM
- About 5% of COVID-19 patients develop a CYTOKINE STORM
- CYTOKINE STORM results in clotting and organ failure
- **CYTOKINE STORM** is associated with excessive STING signaling
- Veyonda only known <u>inhibitor of STING signaling in the</u> clinic

NOXCOVID study

- Phase 1 (pilot) study in patients with moderate to severe illness
- **30-40** patients
- Objectives: safety, biomarkers (cytokines), clinical responses



Our Business – Veyonda[®] and septic shock

NOXCOVID

Q. You have explained that Veyonda is working as an anti-cancer drug by promoting the immune system. If septic shock is driven by an immune system that is already over-active, wouldn't Veyonda just make things worse?

SEPTIC SHOCK A. Yes it could, if it wasn't for the fact that one of the special and unique features of Veyonda is that it only works on systems behaving abnormally, restoring their function to normal

- In cancer, the abnormality is a suppressed immune system.
 Veyonda restores this function to normal by activating it
- In septic shock, the abnormality is an over-active immune system.
 Veyonda restores this function to normal by reducing it

Veyonda is the only drug that blocks the main source of a CYTOKINE STORM that is already in the clinic and proven to be well-tolerated



Large commercial opportunities

Areas of major M&A activity

Strong news flow

Focused commercial strategy

Experienced executive and Board



Investment case

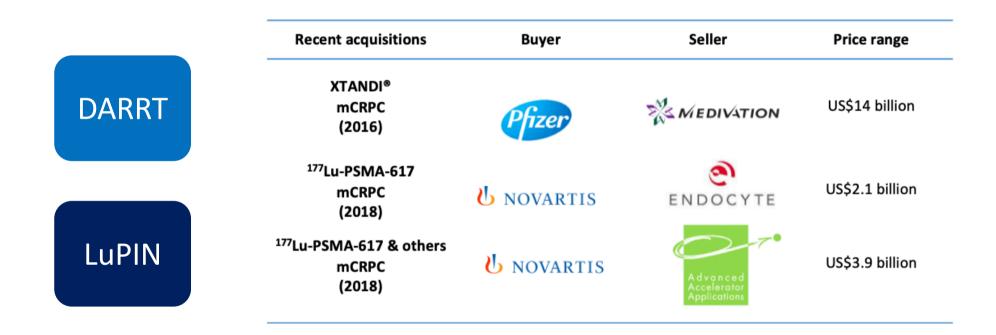
Major market opportunities

DARRT	 Focus on end-stage prostate cancer Estimated 300,000 deaths globally; 33,000 in the U.S. Aiming to make Veyonda standard of care with radiotherapy in end-stage prostate cancer
IONIC	 Checkpoint inhibitor 2019 sales = A\$30 billion Potential market estimated >A\$150 billion p.a. Aiming to make Veyonda the go-to drug to increase response rates to immune checkpoint inhibitors
SEPTIC SHOCK	 Septic shock is major cause of death globally; responsible for estimated 20% of all human deaths Associated with viral and bacterial infections; trauma No effective treatment. Value of market unknown, but expected to be \$multibillion Aiming to develop a treatment, based on Veyonda, that will reduce death and long-term disability in COVID-19 patients



Investment case

Prostate cancer major area of M&A activity





Multiple programs = multiple catalysts Q3 20 - Q1 21

DARRT	 Phase 2 multi-national study in ~200 men; DARRT-2 Anticipated start Q1 2021 Pending appointment of CRO
LuPIN	 Imminent - last patient final treatment Imminent - publication of data from first 32 men Anticipated conference updates on remaining 24 men
IONIC	 Logistical planning current Anticipated start Q1 2021
NOXCOVID	 Aiming to enrol first patient Q3 2020 Aiming to complete enrolment Q4 2020, with results expected in Q1 2021
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Investment Case

Focused Commercial Strategy

Attract potential partners by continuing to undertake clinical trials to develop Veyonda[®] as standard of care in treatment of end-stage prostate cancer

Position Veyonda[®] as a major opportunity to increase response rates to immune checkpoint inhibitors

Position Veyonda[®] as a potential COVID-19 treatment preventing progression of patients onto requiring ICU care and mechanical ventilation



DARRT-2 Phase 2 data LuPIN Phase 2 data

IONIC study

NOXCOVID Phase 1 study



Investment Case

Executive Team



Dr Graham Kelly -Executive Chairman and CEO



Jeanette Bell Chief Operating Officer



Dr Gisela Mautner Chief Medical Officer



Shawn Van Boheemen Chief Financial Officer



Key metrics

Number of Shares	213.2 million shares outstanding
Board shareholding	19.8%
SP (13 Aug 2020)	A\$0.32
Listing date	9 August 2016
Market cap (13 Aug 2020)	A\$67.16 M
Cash position	• AU\$ 7.1M (30 June 2020)





Learn more about Noxopharm Limited (ASX:NOX)

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