



HC Wainwright BIOCONNECT Virtual Conference Jan 10-13 2022

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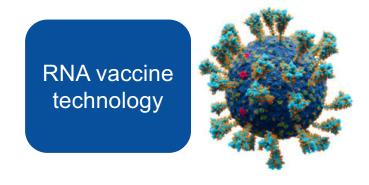


Our fields of interest











Our points of distinction

Small molecule drug platform

A unique family of drugs marked by:

 multiple biological functions based on inhibition of protein folding (thiol-disulfide interchange) and disruption of plasma membrane function (transmembrane electron potential)

a first-in-class means of addressing multiple dysfunctions within the tumour micro-environment

a first-in-class means of addressing selectively **key inflammatory signaling pathways including cGAS-STING and TBK1**



Our points of distinction

A unique platform of oligonucleotides designed to minimize the unwanted triggering of inflammatory receptors

Oligonucleotide drug platform

Oligos inhibiting cGAS-STING signaling:

as treatments for autoimmune diseases

Oligos inhibiting Toll-like receptors (TLRs):

- as treatments for TLRdriven autoimmune diseases
- to reduce inflammatory side-effects of mRNA vaccines



Oncology programs - Veyonda®



Suppository dosage form of idronoxil (IDX)

(3-(4-hydroxyphenyl)-2H-chromen-7-ol)

Selective inhibitor of:

- external membrane NADH oxidases
- sphingosine kinase 1
- TBK1

Biological functions:

- cytostatic/cytotoxic to most forms of cancer cells
- inhibits DNA repair and autophagy
- activates CD4+ and CD8+ T-cells
- reverses S-1-P gradient restores T-cell populations in tumours



Oncology programs - Veyonda®

Program	Combination	Indication	Phase 1	Phase 2
DARRT	Veyonda + EBRT	Prostate, lung, breast	DARRT-1 Completed	DARRT-2 Active
IONIC	Veyonda + nivolumab	Multiple	IONIC-1 Active	
LuPIN	Veyonda + 177LuPSMA-617	Prostate	LuPIN Completed	
CEP	Veyonda + chemotherapy	Soft Tissue Sarcomas	CEP-1 Completed	
			CEP-2 in Start Up	



DARRT-2 Trial

Oncology programs - Veyonda®



Rationale: abscopal response to EBRT via autophagy inhibition + T-cell activation + tumor COLD to HOT conversion

- Multi-national Phase 2 trial. ~100 patients
- IND received from FDA
- Active recruitment in two U.S. sites
- More sites in coming months in AUS and Europe
- Study is in 2 parts:
 - Dose escalation: 1200 mg to 2400 mg; any solid tumour
 - Dose expansion: final dose; focus on prostate cancer, breast and lung cancer







IONIC Trial

Oncology programs - Veyonda®



Rationale: IDX converts tumors from COLD to HOT by reversal of intra-tumoral S-1-P gradient and activation of CD4+ and CD8+ T-cells

- Phase 1 trial of Veyonda® + nivolumab (Opdivo®; BMS)
- Investigator initiated study; Australian only sites
- First 2 patients enrolled
- Cohort 1: Non-responding tumours to PD-1 inhibitors
- Cohort 2: PD-1 inhibitor naive tumours





CEP-2 Trial

Oncology programs - Veyonda®



Rationale: IDX chemo-enhancement of alkylating agents

- Phase 1 trial of Veyonda® + doxorubicin IND from FDA received
- Soft tissue sarcomas
- Trial to be conducted in the U.S. due to strong interest
- Contract negotiations with clinical sites ongoing
- Enrolment expected shortly



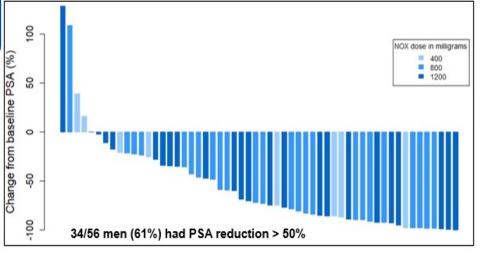
LuPIN Trial

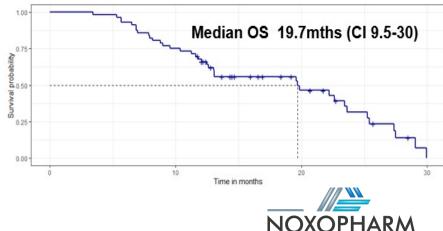
Oncology programs - Veyonda®



Rationale: IDX radio-enhancement of 177 lutetium-PSMA

- Promising results have been published in peer-reviewed medical journals:
 - Any PSA reduction in 86% of patients
 - PSA fall of >50% in 61% of patients
 - Median Overall Survival of 19.7 months
- Discussions regarding potential trial in Europe currently underway





Oncology - Pipeline



A pipeline of 3 exciting new anti-cancer drugs with novel actions

Drug 1: Glial brain cancers (GBM, DIPG)

Drug 2: Pancreatic cancer

G-protein coupled receptor inhibitors

Targeting highly aggressive cancers dependent on cancer-derived stromal cells for growth signals

Purpose:

- Stop aggressive cancer growth
- Prevent killing of healthy brain tissue to make room for cancer growth

Purpose:

- Stop aggressive cancer growth
- Enhance cancer-killing effect of chemotherapy
- Remove scar tissue blocking access to cancer by chemo drugs

Drug 3: Multiple cancer types

Second generation IDX

Most active form of IDX in the body

Purpose:

- Potential for greater potency, reduced dosage, stronger IP
- Potential oral dosage form



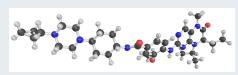
Acute Inflammation - Sepsis



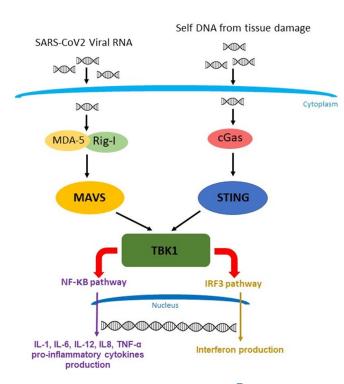
Acute inflammation - sepsis

Drug discovery program

Unique family of small molecule inhibitors of Tank-Binding Kinase 1 (TBK1)



Development of drugs to block the cytokine release syndrome associated with viral and bacterial infections





Autoimmunity



Drug discovery program

Oligonucleotides targeting cGAS-STING



Developing drugs for various chronic inflammatory/ autoimmune diseases





RNA Vaccines

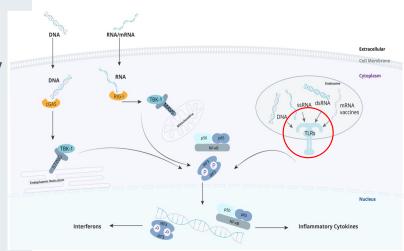


Oligonucleotides targeting Toll-like receptors (TLRs)

Therapeutic aim = to reduce the pro-inflammatory side-effects of RNA drugs and vaccines and treat TLR-driven inflammatory diseases

In the case of RNA vaccines, to:

- Improve their safety
- Permit higher dosages of viral antigen to be used
- Improve manufacturing efficiencies







Key Metrics

(at 7 January 2022)

Market cap	A\$124m
Share price	A\$0.425c
Issued shares	~292.2m
Cash (at 30/9/21)	A\$23.6 m
\$5.9M cash rebate rec'e	d Jan 2022

Anticipated News Flow (next 6 months)

Progress in:

- IONIC-1, DARRT-2 & CEP-2 clinical program
- Oncology drug pipeline
- Sepsis and autoimmunity disease drug discovery programs
- mRNA vaccine technology program



