

PYC Therapeutics

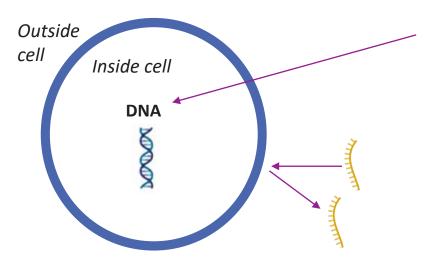
Investor Relations Presentation September 2019



The delivery challenge: getting big drugs inside cells



The promise of breakthrough medicine can only be realised once the 'delivery challenge' has been solved



The Opportunity

The *highest value drug targets* (DNA, RNA, protein) exist *inside* cells

The Challenge

But... The cell membrane has evolved over hundreds of millions of years to **keep foreign substances** (**like drugs**) **out**

The Implication: Many breakthrough therapeutics fail due to an inability to reach their target



"If the CRISPR gene editing system is to live up to its disease-curing potential, researchers must devise a plan to deliver it into the body"



"Merck's Alan Sachs, on RNAi's Big Challenge: Delivery, Delivery, Delivery"



"For the most common inherited monogenic disorders ... effective gene therapy is likely to remain a delivery challenge"

^{1 &#}x27;CRISPR's Breakthrough problem', Chemical & Engineering News, Feb 2017;

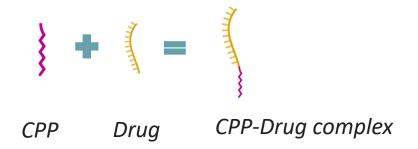
^{2 &#}x27;Merck's Alan Sachs, on RNAi's Big Challenge: Delivery, Delivery, Delivery', Exome by Xconomy, Jan 2010

PYC Therapeutics solves the delivery challenge with our Cell-Penetrating Peptide (CPP) technology

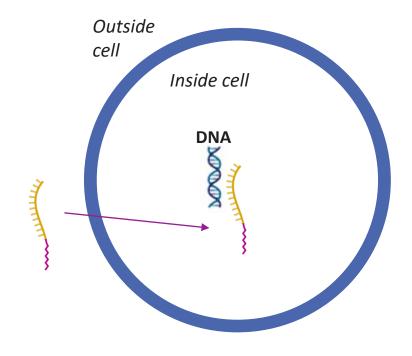


PYC Therapeutics' solution

Cell Penetrating Peptides (CPPs) are *able to cross the cell membrane* – CPPs can be joined to a drug cargo to deliver it inside the cell



Precision medicine is now a reality

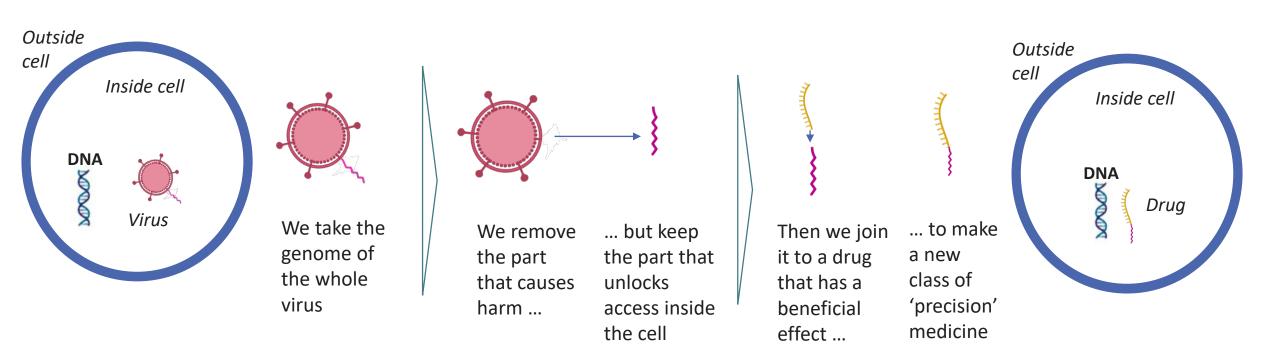


"By taking advantage of the unique cell-translocation property of these short peptides, various payloads of proteins, nucleic acids, or even nanoparticle-based carriers were **delivered into all cell types with unparalleled efficiency**" 1

PYC's delivery platform uses Nature's solution to a complex problem



Nature has worked out how to cross the cell membrane – viruses and other micro-organisms can enter human cells

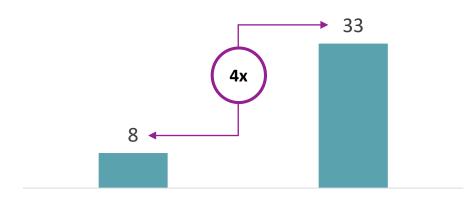


PYC has a clear competitive advantage in the amount of drug cargo that we can deliver



Our delivery technology delivers 4x as much drug cargo inside cells than our nearest competitor's

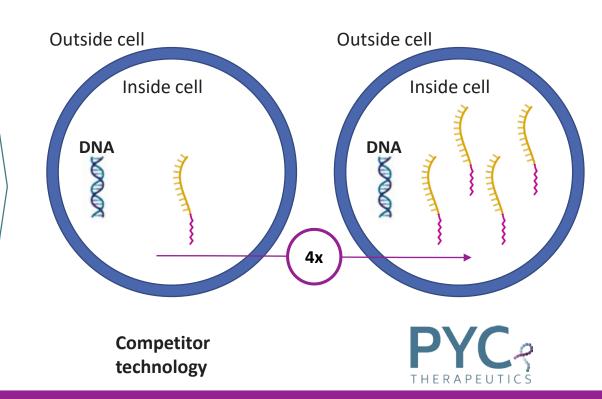
% of cells with drug successfully delivered¹ in animals



Competitor technology



Getting enough drug cargo inside the cell is the ratelimiting step in the development of precision drugs



PYC's competitive advantage has been proven in both animals and human cells

PYC drives shareholder returns through two commercial applications of our delivery 'platform'





Development of our own pipeline of drugs



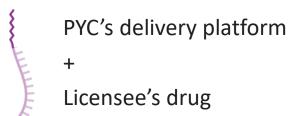
PYC's delivery platform

+

PYC's drug

- Develop our own cargoes for our area of focus, starting with Inherited Retinal Disease
- PYC owns the 'whole molecule'

Licensing our delivery platform to other Pharma Co



- Leverage our delivery platform to generate up-front revenue and gain a trailing interest in third party molecules
- Use the out-licensing model to scale across multiple disease indications across diverse target tissues

Developing our drugs: PYC is targeting significant markets of unmet need



Illustrative drug value proposition – Inherited Retinal Disease example

Development stage

Discovery

Preclinical

Plase 1

Phase 2

Phase 3

Marketing

Current progress



Potential by 2025 2+ Clinical drugs 1 Marketed drug

We have a strong base to enable success in reaching and capturing the full opportunity in the Inherited Retinal Disease market



Success in animal models

Both efficacy and toxicology



Success in human cells

Proof of concept established



3D human retina model

Organoid or "retina in a dish" model coming in the immediate future



Success from similar drugs
ASOs in other inherited retinal
diseases are clinically validated



Serving unmet need

Opportunity to combine phases 2/3 in clinical trials



Accessible patient population *Disease registries assist distribution*

PYC has a clear set of milestones to deliver our lead drug

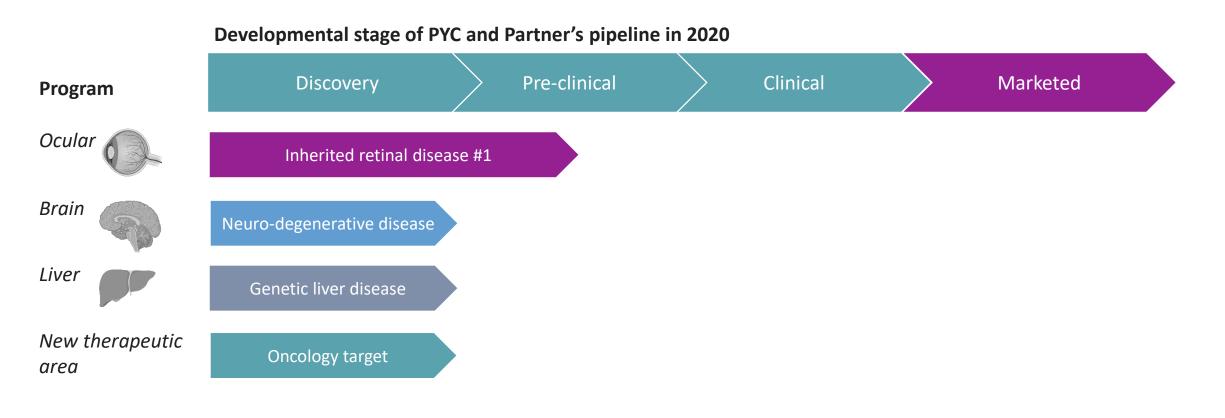


Milestone impacting valuation and progress to market

TH	PYC	2								
Pre-clinical		> IND-enabling		IND	Phase 1		Phase 2		Phase 3	NDA
 Proof of concept animal efficacy Proof of concept human cell data Proof of concept "3D model of the retina" data (all informs if the drug will work in humans) 	•	Drug molecule finalised Large animal toxicology results (informs if it's safe in humans)	•	Investigational New Drug (IND) approval from US FDA	First dosing of Human Patients with the Rare Inherited Retinal Disease Phase 1 results including primarily a) is the drug safe in humans, and secondarily b) does the drug work in humans	•	First dosing in 2/3 trail Phase 2/3 resuprimarily a) dowork in humadose Continued reaphase 1 cohor	ults in oes th ons at o	cluding e drug safely our intended s from the	New Drug Application (NDA) approval from US FDA to market the drug First commercial patient dosing Applications in other major markets including Europe and Japan

2 Licencing out PYC's CPP delivery platform





"Haven't heard of RNA Therapeutics yet? You will" 1

PYC will be operating in a competitive landscape



Antisense Oligo landscape examples

	PYC? THERAPEUTICS	KODIAK	Apellis	ProQR THERAPEUTICS	STOKE
Geographic base	Australia	US	US	Netherlands	US
Platform or asset	Platform	Platform	Asset	Asset	Platform
Development stage	Pre-clinical	Clinical (Phase 1)	Clinical (Phase 2)	Clinical (Phase 1b)	Pre-clinical
Lead indication	Ocular rare disease	Wet AMD	Ocular immunotherapy	Ocular rare disease	Neurological rare disease
Cash reserves (AUD) as at 30 June 2019	~\$10M	~\$120M	~\$260M	~\$140M	~\$350M
Market Cap (AUD) as at 26 August 2019	~\$90M	~\$580M	~\$2,500M	~\$415M	~\$1,600M

Eye disease landscape examples

The scientific team



Scientific Advisory Board



Shohei Koide B.S. M.S. Ph.D **Professor of Biologics** Design at **New York University** World-leader in the design of biologics



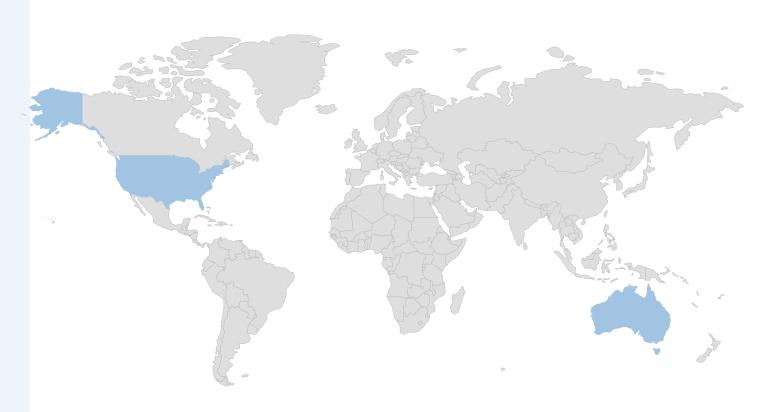
Stephen Doberstein B.S.Ch.E, Ph.D Chief Research & Development Officer at **Nektar Therapeutics** 17 years experience in biotechnology



Judy Lieberman MD, Ph.D Professor of Pediatrics at **Harvard Medical School** First-class University board representation



Rakesh Veedu Ph.D Head of precision nucleic acid therapeutics research at the Centre for Comparative Genomics Expert in antisense oligos



Operational Team



Rohan Hockings MBBS (Hons.), JD GDLP Experience across both clinical and commercial roles



Kaggen Ausma LLB, B.Econs (Hons.) Previous roles in McKinsey & Co and CLSA Asia-Pacific



Katrin Hoffmann, Ph.D 20 years experience in biomedical research



Science Team 23 Scientists based at the Harry Perkins Institute of Medical Research

Key collaborators



Centre for Molecular Medicine and Innovative Murdoch Therapeutics



Inventors of (USFDA approved) ASO drug Eteplirsen

Clinical expertise in the eye (initial target tissue of interest)

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