

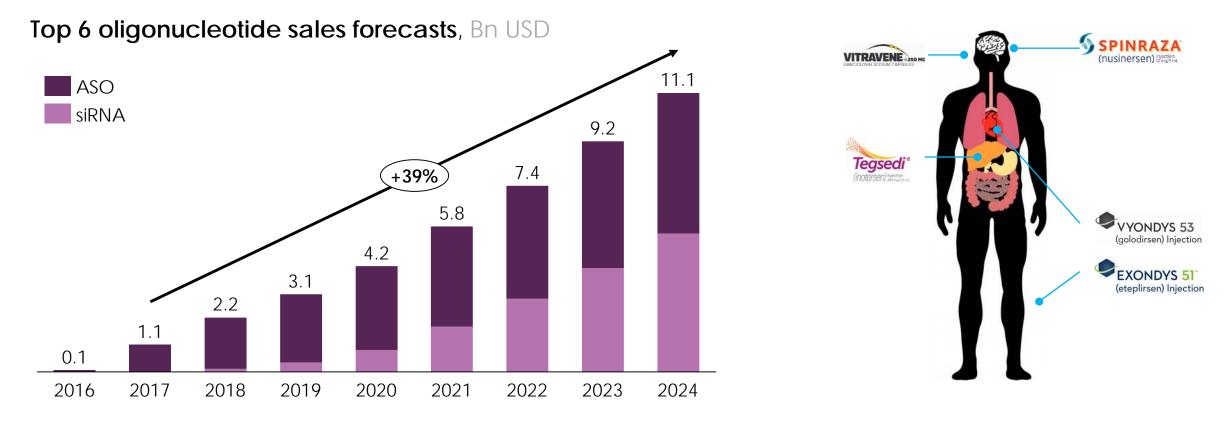


# Investor Relations Presentation February 2020

Where we are competing: Oligonucleotides are set to become the third major class of drug



# "In the long run, oligonucleotides are likely to become a major class of therapeutics, on par with small molecules and biologics"



# Why we can win: PYC now has the capabilities to progress opportunities from discovery through development



### PYC's competitive advantages extend throughout the entire drug development value chain

2. Drug design

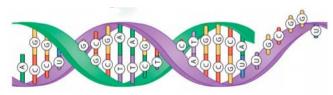
1. Selection of target disease

- Clinicians guide the choice of diseases that we target
- PYC is developing treatments for the greatest patient needs



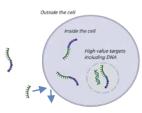
 PYC's drugs are designed by world-leading experts

 PYC's drugs maximise potency and precision



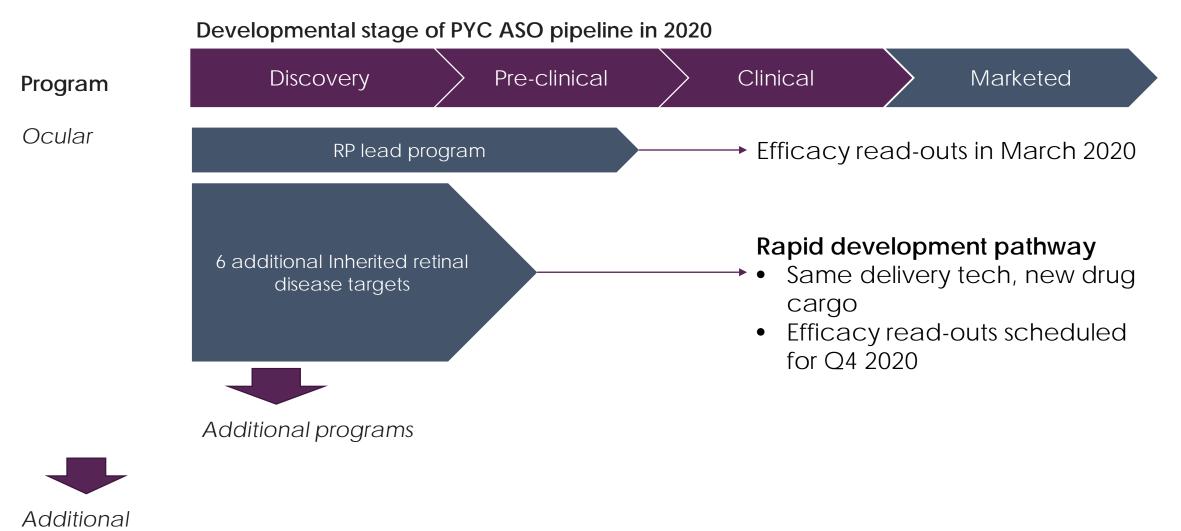
## 3. Drug delivery

- PYC has a defensible competitive advantage
- PYC's delivery technology ensures more drug reaches its target



PYC's partnership with the Lions Eye Institute ensures that we start with the patient in mind PYC's R&D team is responsible for multiple FDA-approved products PYC's delivery technology is proven to outperform our competitors How we are executing: PYC has multiple assets in development and is actively expanding





How we are executing: PYC is specifically focused on diseases with attractive target markets



Rare ocular disease pipeline assessments

Rare unmet need

Strong reimbursement cases

Attractive eligibility for FDA concessions

1,000-8,000 patients

100-500k USD pricing

Orphan designation Priority designation Fast track designation

Attractive, sustainable end markets with a rapid path to entry

For prevalence see Sullivan et al, 'Prevalence of disease-causing mutations in families with autosomal dominant retinitis pigmentosa: a screen of known genes in 200 families' Invest Ophthalmol Vis Sci. 2006 Jul;47(7):3052-64.

For pricing see Exondys 51, Spinraza, Luxturna,

For concession see https://www.fda.gov/patients/learn-about-drug-and-device-approvals/fast-track-breakthrough-therapy-accelerated-approval-priority-review





PYC's lead program is fully funded through to market



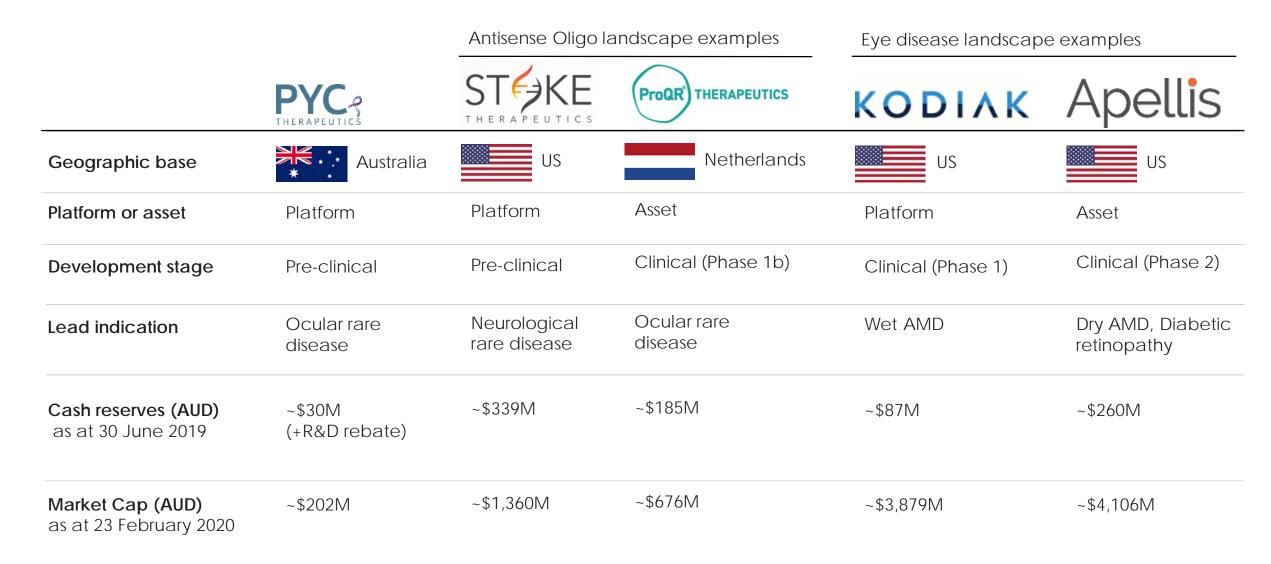
PYC is on the ground in the US alongside the regulator, specialist life sciences investors, and industry partners



PYC has multiple shots on goal at the critical IND milestone this year

# How we are executing: We must build the credibility of our technology to close the valuation gap to peers





## PYC Corporate snapshot



#### Financial Information (23 February 2020)

Share price	\$0.069
Number of shares	2,930M
Market Capitalisation	\$202M
Cash (1-Jan-20)	\$30M
Debt (1-Jan-20)	Nil
Enterprise Value	\$172M

Source: IRESS

Note:

1 Excludes 10m unlisted options exercisable at A\$0.06 before 30 May 2020



Alan Tribe – Chairman	Ala
Doug Huey – Executive Director	Dr
Dr Rohan Hockings – Executive Director	Sie
Dr Bernard Hockings – Non-Executive Director	Ar

Top shareholders	%
Alan Tribe	27.1%
Dr Bernard Hockings	12.0%
Sietsma Holdings	10.0%
Anthony Barton	5.0%

#### Share price performance (1 year)

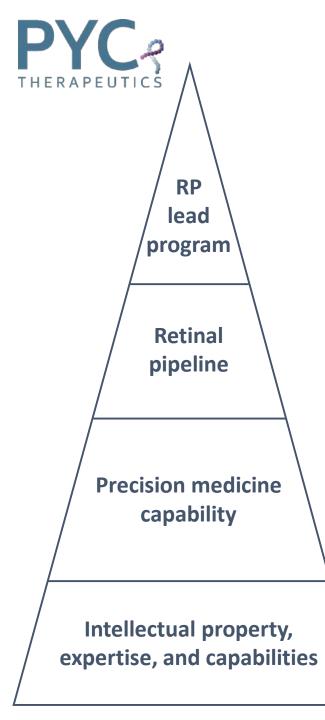


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### **Overview of PYC**

- **High-performing precision medicine** lead drug combines the advantages of Cell Penetrating Peptides (CPP) and Anti Sense Oligonucleotides (ASO) to treat a form of Retinitis Pigmentosa
- Favorable clinical path and market targets a severe unmet need within a >US\$1bn market
- **Pre-clinical validation** completing final assessments to finalise the pre-clinical data pack, with upcoming Investigational New Drug (IND)-enabling stage-gate in Q2 2020
- Established development pipeline six opportunities being developed with a subset expected to reach the IND-enabling stage-gate by end of 2020
- **Distinctive capabilities** Vision Pharma partnership combines PYC's RNA therapeutic expertise with the Lions Eye Institute's world leading retinal tissue and clinical expertise
- **Competitive advantages** enhanced performance and safety of CPP-ASO's v. other ASO and broader RNA/DNA therapeutics
- **Favorable pre-clinical timelines** highly efficient process to design and validate RNA therapeutics once targets are identified
- Applicable to broad set of tissues currently evaluating performance in multiple tissues
- **Proprietary peptide libraries** diversity of libraries, quality of screening assays, richness of data, and advanced analytics capabilities allow for broad applications
- **Capabilities** Internal expertise in precision medicine coupled with partnerships with clinicians. Commercial excellence and global reach through Australian and US offices