

ASX: ZLD OTCQB: ZLDAF WWW.ZELIRATX.COM

ZELIRA THERAPEUTICS A LEADING GLOBAL MEDICINAL CANNABIS COMPANY





DISCLAIMER & IMPORTANT NOTICE

Disclaimer

This presentation has been prepared by Zelda Therapeutics Ltd ACN 103 782 378 ("Company"). It does not purport to contain all the information that a prospective investor may require in connection with any potential investment in the Company. You should not treat the contents of this presentation, or any information provided in connection with it, as financial advice, financial product advice or advice relating to legal, taxation or investment matters.

No representation or warranty (whether express or implied) is made by the Company or any of its officers, advisers, agents or employees as to the accuracy, completeness or reasonableness of the information, statements, opinions or matters (express or implied) arising out of, contained in or derived from this presentation or provided in connection with it, or any omission from this presentation, nor as to the attainability of any estimates, forecasts or projections set out in this presentation.

This presentation is provided expressly on the basis that you will carry out your own independent inquiries into the matters contained in the presentation and make your own independent decisions about the affairs, financial position or prospects of the Company. The Company reserves the right to update, amend or supplement the information at any time in its absolute discretion (without incurring any obligation to do so).

Neither the Company, nor its related bodies corporate, officers, their advisers, agents and employees accept any responsibility or liability to you or to any other person or entity arising out of this presentation including pursuant to the general law (whether for negligence, under statute or otherwise), or under the Australian Securities and Investments Commission Act 2001, Corporations Act 2001, Competition and Consumer Act 2010 or any corresponding provision of any Australian state or territory legislation (or the law of any similar legislation in any other jurisdiction), or similar provision under any applicable law. Any such responsibility or liability is, to the maximum extent permitted by law, expressly disclaimed and excluded.

Nothing in this material should be construed as either an offer to sell or a solicitation of an offer to buy or sell securities. It does not include all available information and should not be used in isolation as a basis to invest in the Company.

Future matters

This presentation contains reference to certain intentions, expectations, future plans, strategy and prospects of the Company.

Those intentions, expectations, future plans, strategy and prospects may or may not be achieved. They are based on certain assumptions, which may not be met or on which views may differ and may be affected by known and unknown risks. The performance and operations of the Company may be influenced by a number of factors, many of which are outside the control of the Company. No representation or warranty, express or implied, is made by the Company, or any of its directors, officers, employees, advisers or agents that any intentions, expectations or plans will be achieved either totally or partially or that any particular rate of return will be achieved.

Given the risks and uncertainties that may cause the Company's actual future results, performance or achievements to be materially different from those expected, planned or intended, recipients should not place undue reliance on these intentions, expectations, future plans, strategy and prospects. The Company does not warrant or represent that the actual results, performance or achievements will be as expected, planned or intended.



2

ZELIRA THERAPEUTICS

- pain and autism.
- Disruptive 'Launch, Learn, & Develop' model for rapid commercialization.
- Revenues from US licensing deals commenced in Q1, 2020

• Globally integrated company developing, and marketing clinically validated medicinal cannabis products

• US, Australia and EU footprint to rapidly access the largest, most profitable & fastest growing cannabis markets

Leading pipeline of products in clinical development targeting large addressable markets for insomnia, chronic

• Multiple branded products targeting consumer and pharmaceutical markets set-to launch globally in 1H 2020.







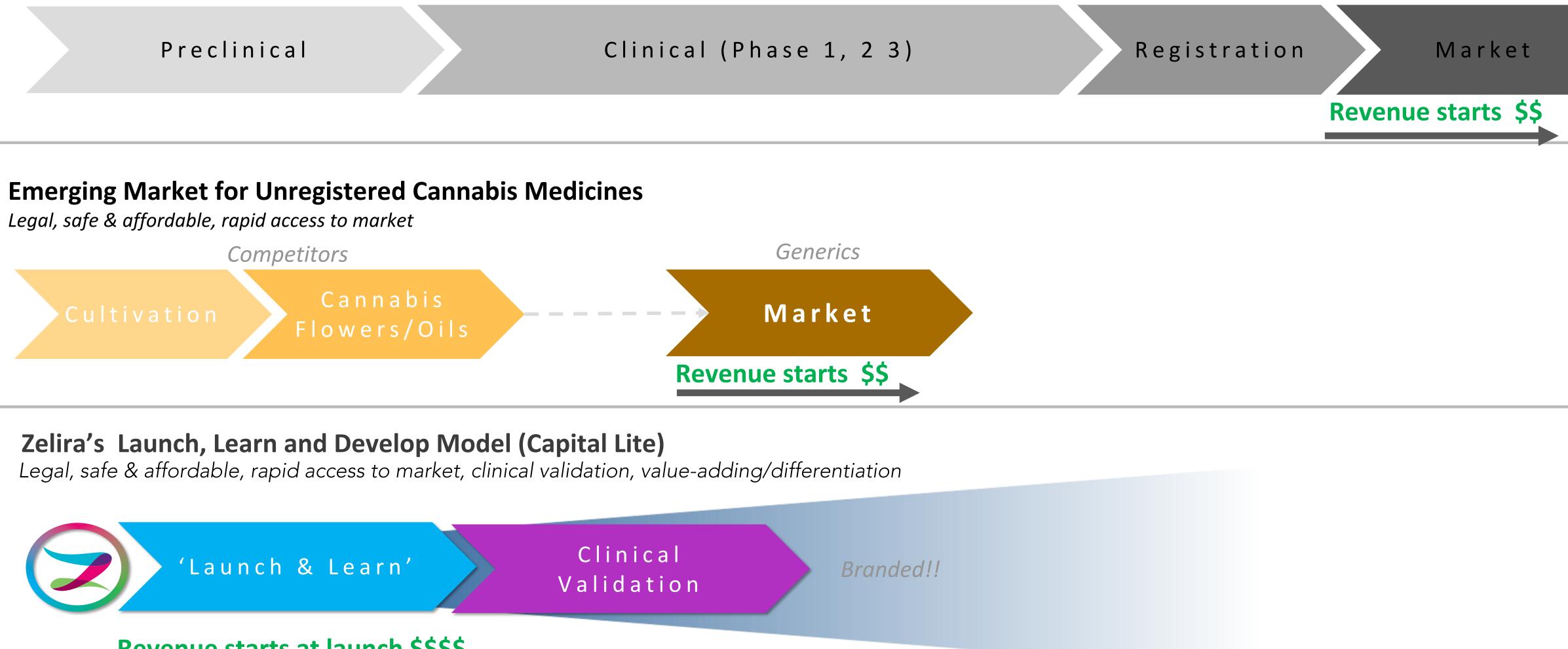
COMMERCIALIZATION STRATEGY

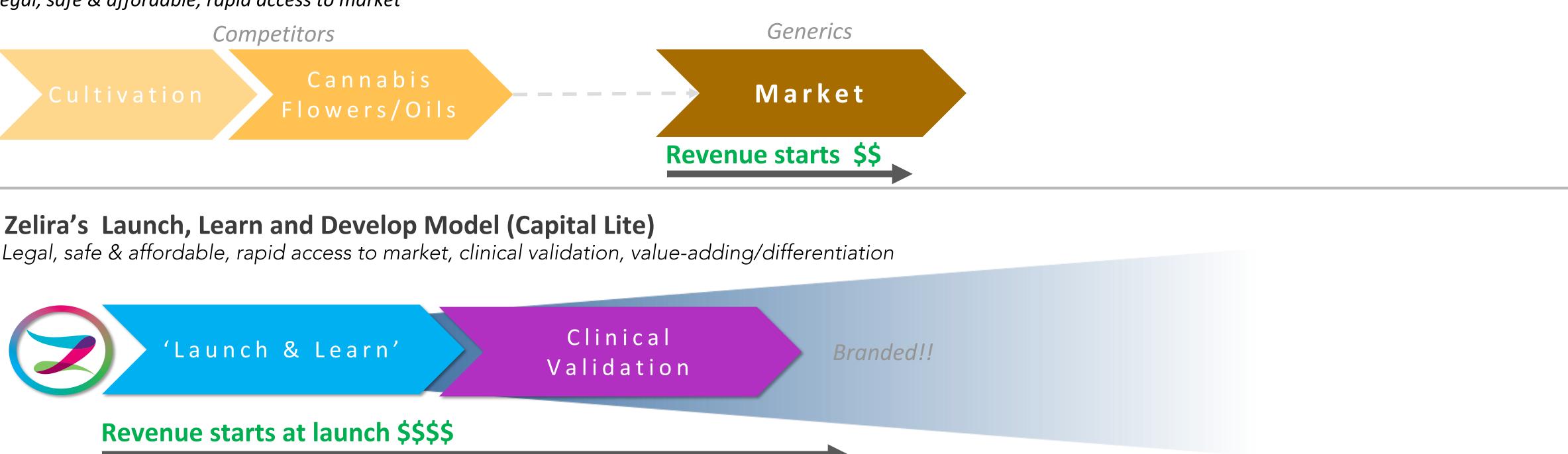


MEDICINAL CANNABIS: ACCELERATED PATH TO MARKET

Conventional Pharmaceutical Development Path

10 years to market, \$1 Billion-expensive medicine







DIFFERENTIATED COMMERCIALIZATION MODEL



Value Adding

- Clinically validated
- **Proprietary formulations**
- GMP grade
- **IP** protection
- Branded

Competitors



Branding Attracts **Price Premium**

Revenues from Global Distribution/Licensing



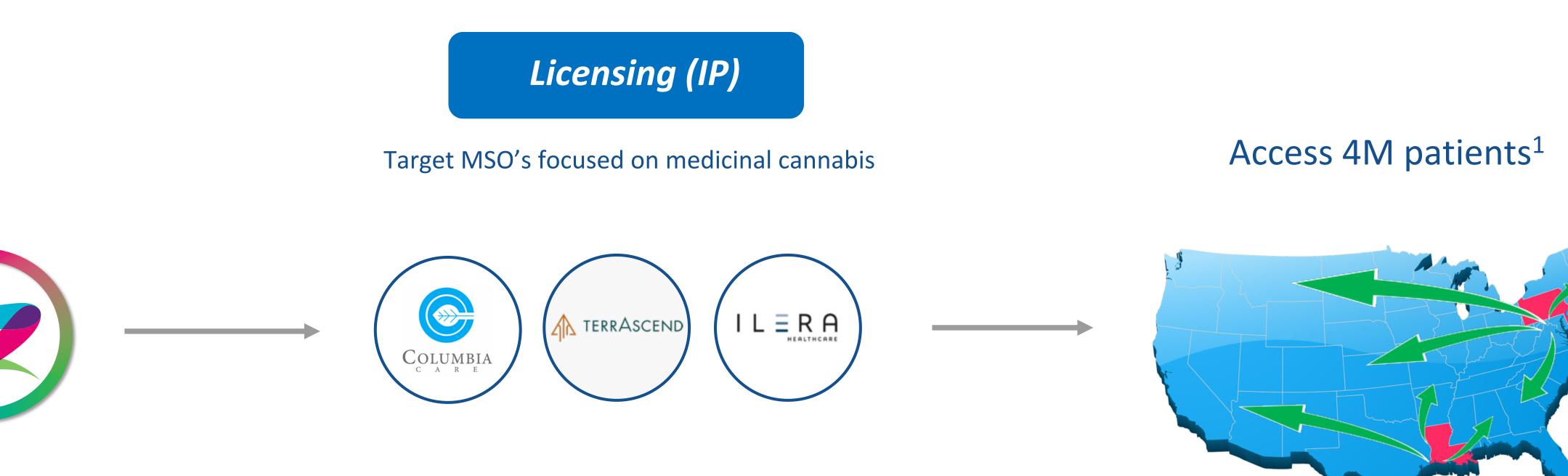
- Generics
- **Downward pressure on price to compete**







ZELIRA'S COMMERCIALIZATION STRATEGY - USA



- US\$10BB market is >100X larger than Australia
- Novel virtual distribution strategy to accelerate access to entire US market
- License products, IP, technical and clinical dossiers
- Access to premium validated products drives deal-flow

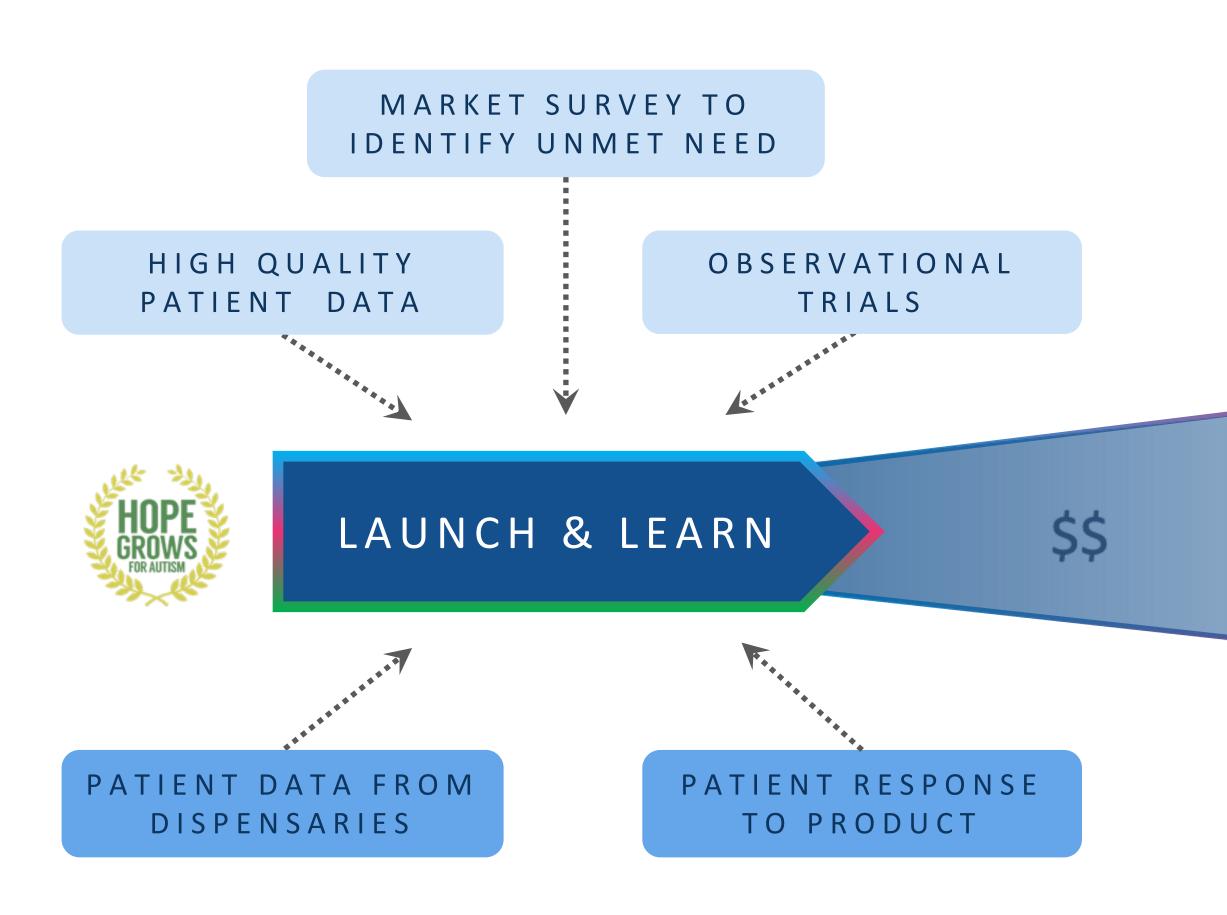




HOW ARE WE DIFFERENT?



ZELIRA'S "LAUNCH, LEARN AND DEVELOP" MODEL



CLINICAL TRIALS

Clinically Validated Products

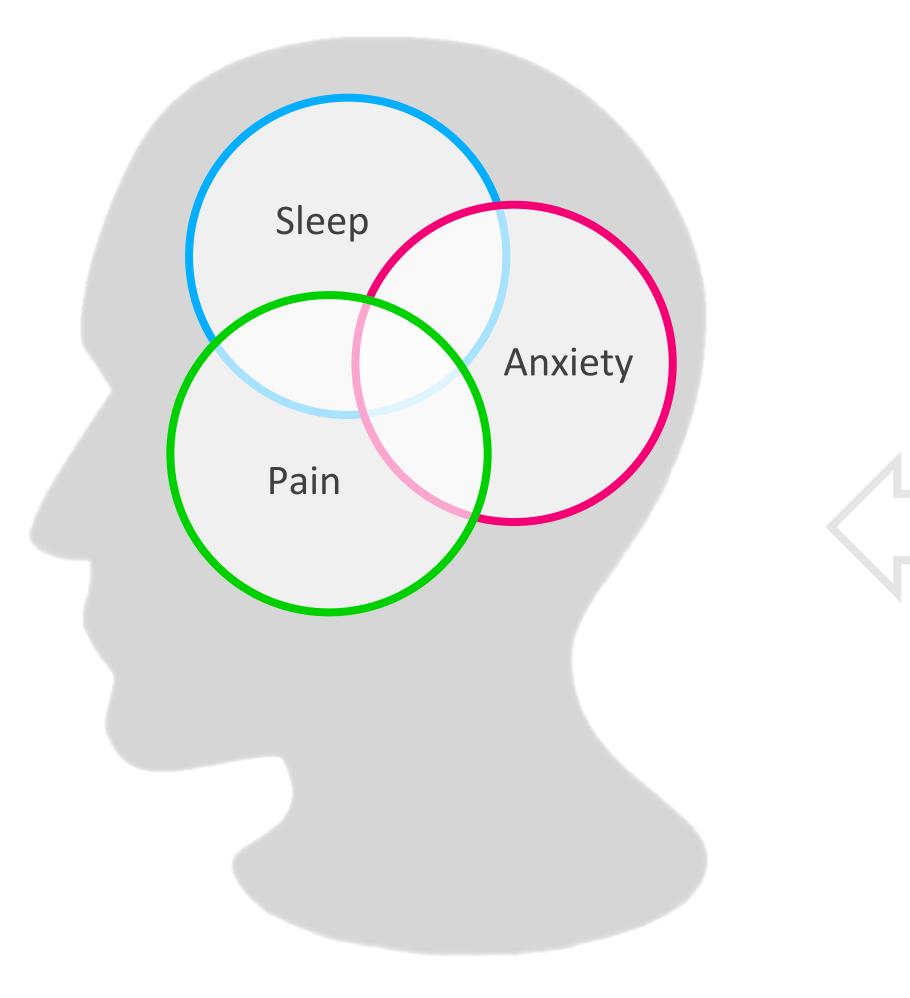
\$\$\$\$





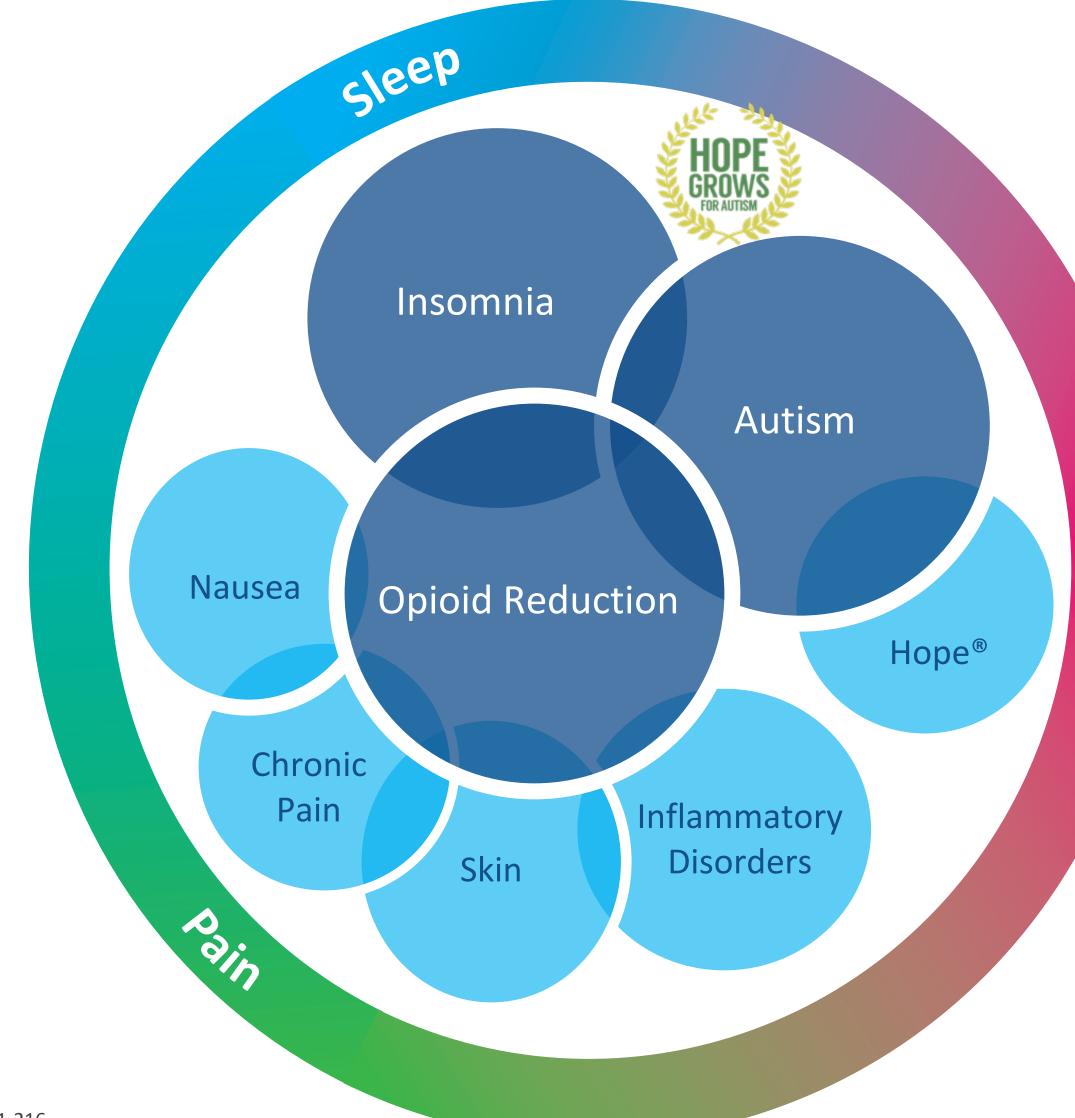


FOCUS ON CURRENT PATIENT NEEDS = LARGE ADDRESSABLE MARKET



Most common reasons patients use medicinal cannabis¹.

1. Lintzeris, N et al., Medical Cannabis in Australia, 2016: the Cannabis as medicine Survey (CAM-16). The Medical Journal of Australia, 209(05), 211-216.



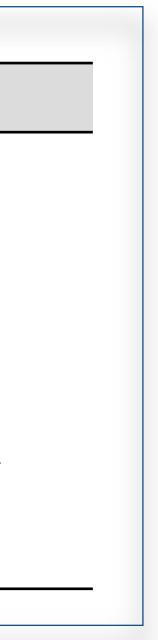




STRONG INTELLECTUAL PROPERTY PORTFOLIO

- Seven patent families filed covering:
 - Cancer*
 - Skin Disease (topical)*
 - Sleep Disorders*
 - Cancer diagnostics*
 - Autism
 - Anxiety
 - Opioid Reduction
- Patents (*granted) provide broad composition of matter protection until 2036.
- Generating novel IP is core to Zelira's commercialization strategy.
- Strong and defensible IP will be essential as patent landscape becomes increasingly litigious.

Patent	Priority Date	Filing
Cancer	August 2016	WO2018/023166
Skin Disease	August 2016	WO2018/023164
Sleep Disorders	August 2016	WO 2018/023163
Insomnia	November 2017	AU2017904818
Cancer Prognosis	October 2016	WO 2018/071986 A
Autism	September 2017	AU2017903766





STRATEGIC FOCUS ON DISRUPTING TRADITIONAL PHARMA MARKETS

~US\$330B

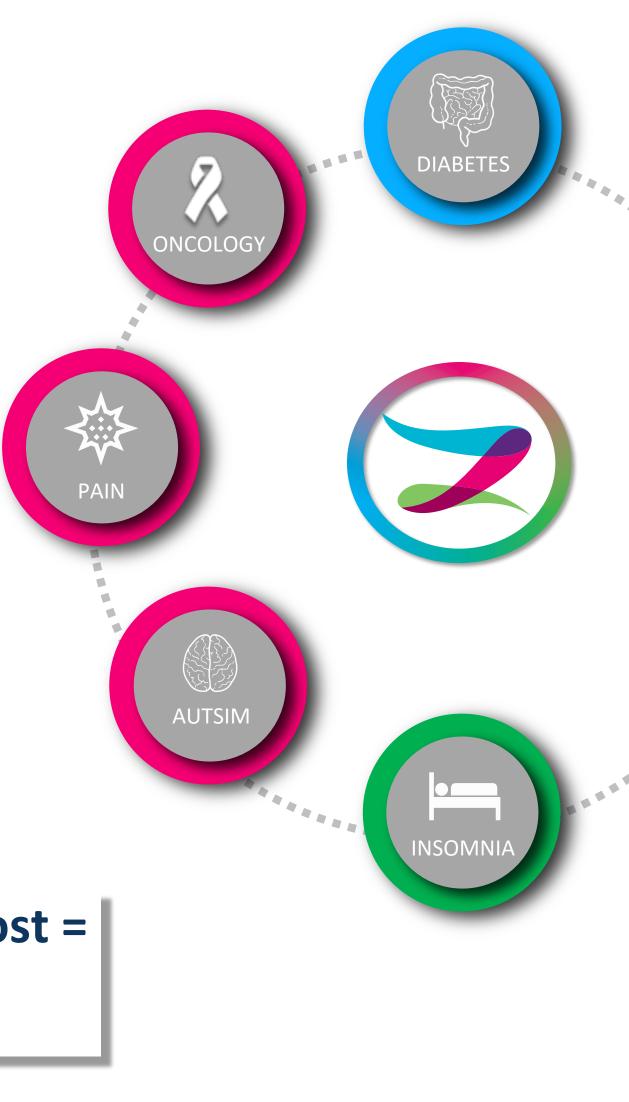
TOTAL DRUG SPEND IN USA

\$2 B

Source: https://www.statista.com/statistics/238698/us-health-spending-leading-areas/ https://blog.marketresearch.com/top-6-things-to-know-about-the-28-billion-sleep-market DIABETES

\$52 B

- \$46 B ONCOLOGY
- \$32 B CHRONIC PAIN
- \$17 B MENTAL HEALTH
 - INSOMNIA

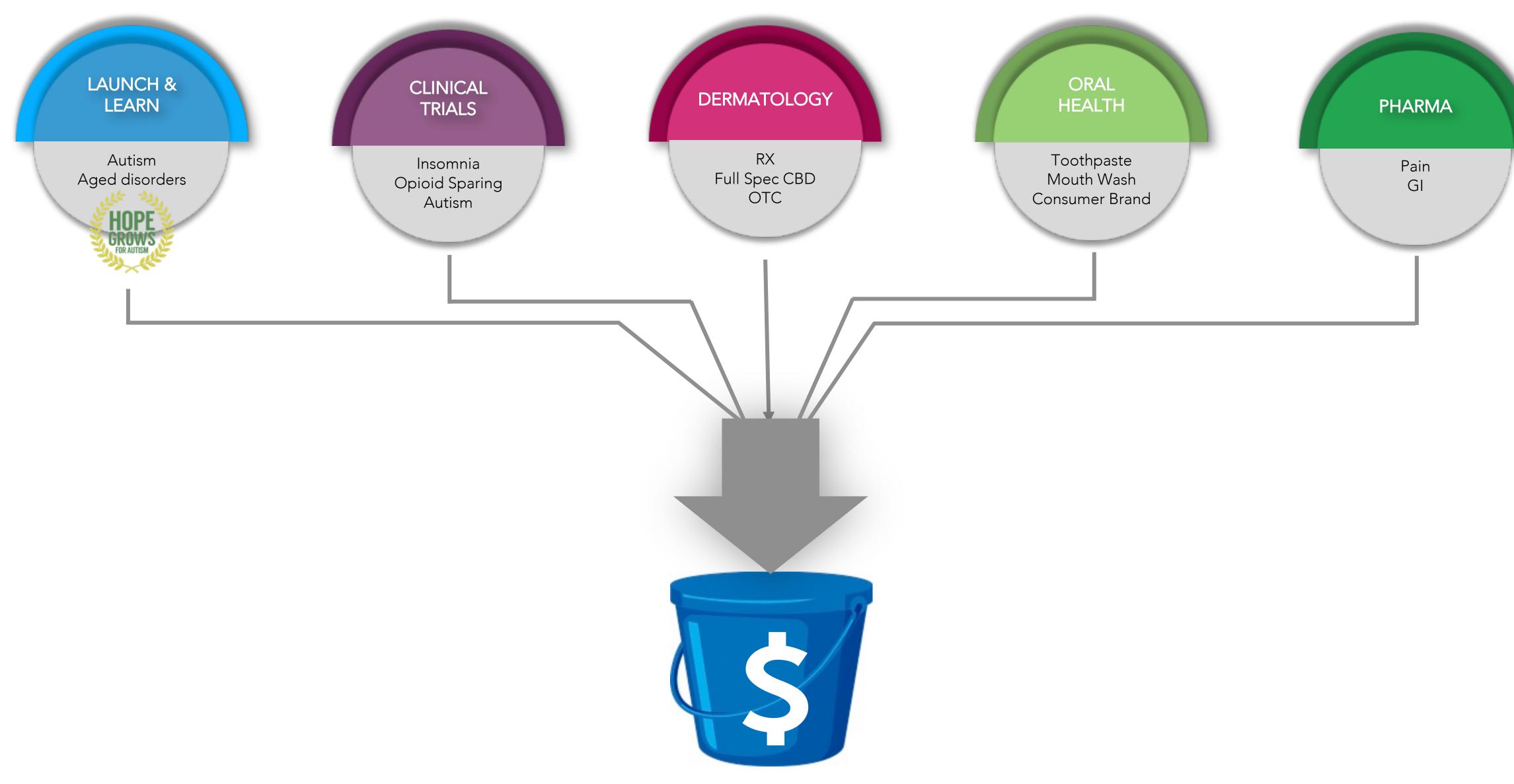


Estimated annual cannabis substitution cost = \$4.86 billion in 2019

(New Frontier Data¹)



ZELIRA'S DIVERSIFIED REVENUE MODEL





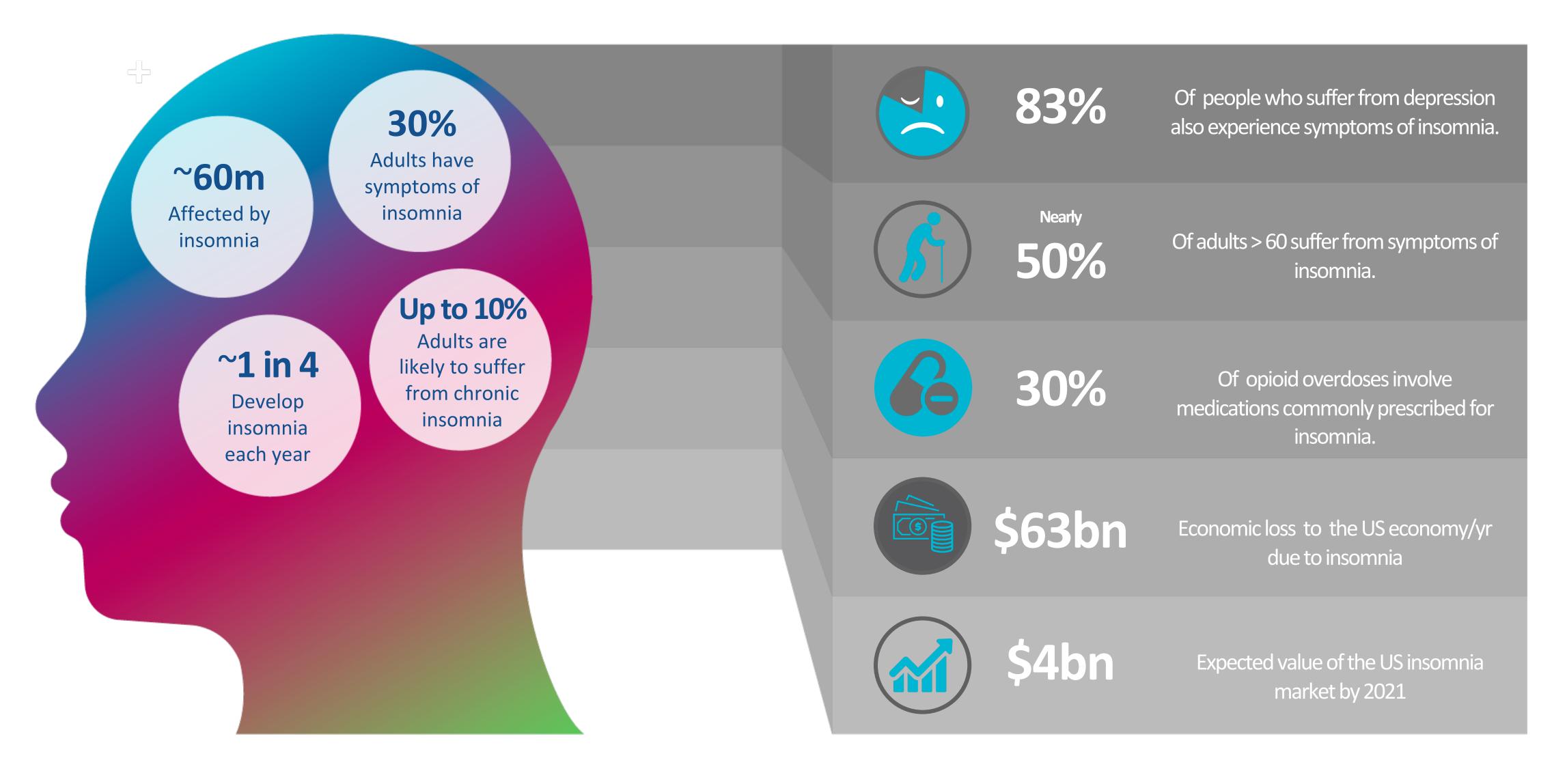


WORLD'S FIRST PHASE 1A/2B CLINICAL TRIAL TARGETING **SLEEP DISORDERS**

EFFICACY OF ZELIRA'S MEDICINAL CANNABIS FORMULATION (ZTL-101) FOR TREATING CHRONIC INSOMNIA



KEY INSOMNIA STATS-USA MARKET



Insomnia is the most common sleep disorder in America



ZELIRA: INSOMNIA PHASE 1A/2B TRIAL

A Study to Evaluate the Efficacy of Sublingual Cannabinoid Based Medicine Extract (ZTL-101) **Compared with Placebo for the Treatment of Sleep Disorders Due to Insomnia**

Primary Outcomes:

- To evaluate safety and tolerability of increasing doses o the sublingual cannabinoid extract ZTL-101.
- To evaluate the efficacy of the sublingual cannabinoid extract ZTL-101 containing THC for improving insomnia symptoms in people with chronic insomnia.

- Investigational Team: PI Prof Peter Eastwood, Clinical Prof David Hillman, Ass Prof Nigel McArdle, Dr Melissa Ree, Dr Jennifer Walsh
 - **Study design:** Randomised double-blind, placebo-controlled, crossover
 - **ANZCTR:** ACTRN12618000078257

	Secondary Outcomes:
of	 To evaluate the efficacy of ZTL-101 as compared to placebo for improving <u>objective</u> and <u>subjective</u> sleep quantity and quality in people with insomnia. To evaluate <u>quality of life</u> improvements in people with insomnia when using ZTL-101 as compared with placebo.



ZTL-101 MET PRIMARY AND SECONDARY ENDPOINTS

Primary Endpoints

SAFE

- No serious adverse events
- Adverse events mild and transient
- Maximal dose well tolerated

EFFICACIOUS



- 36% reduction in Insomnia Severity Index (ISI)
- Treatment was dose responsive
- High dose patients became subclinical to insomnia
- Statistically significant reduction in ISI scores at all doses

Secondary Endpoints

Improved Objective and Subjective Measures of Insomnia



Statistically significant improvement in:

- Time spent asleep
- Time taken to return to sleep after waking
- Sleep quality
- Feeling rested

Improved Quality of Life

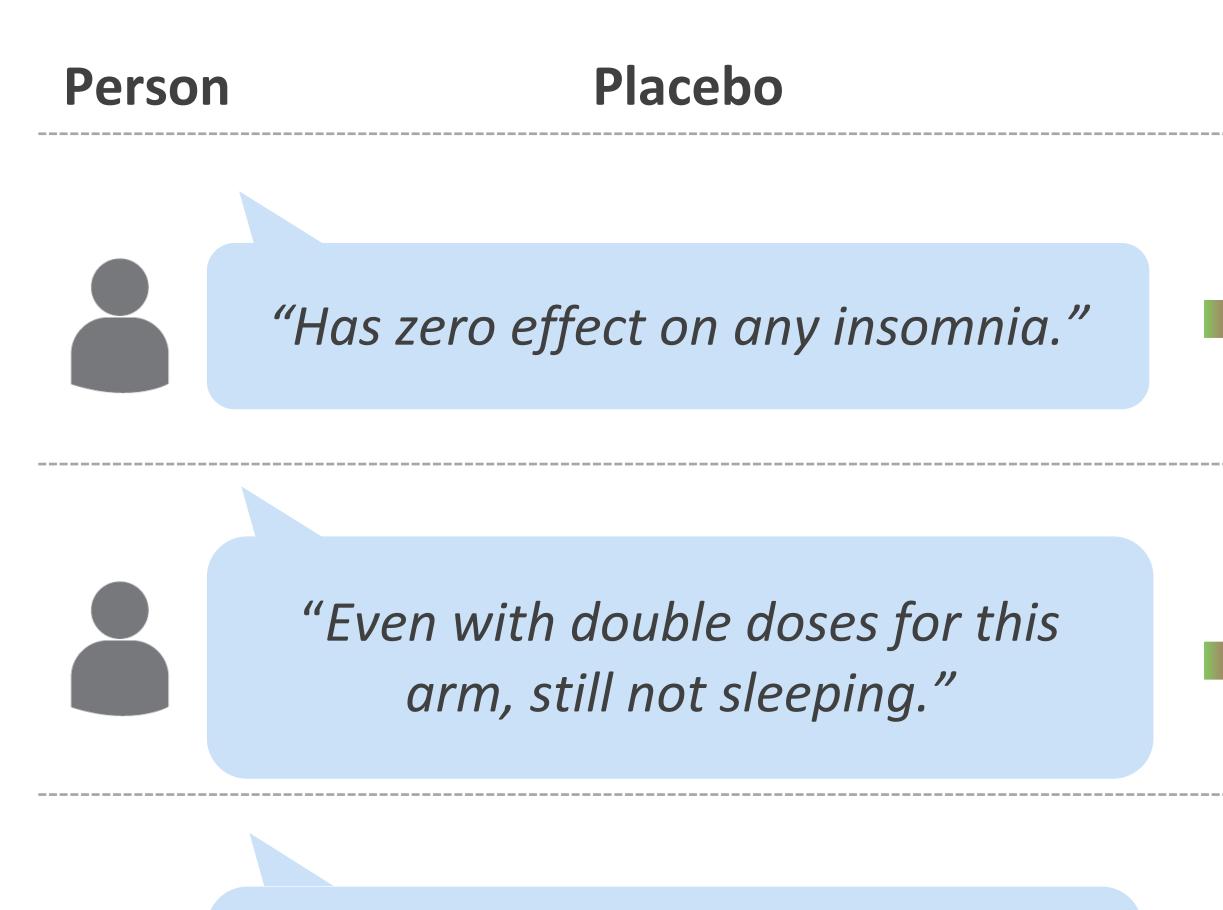


Statistically significant:

- Reduction in levels of fatigue
- Improvement in functioning



PATIENT COMMENTS*



"Stayed awake for hours trying to sleep most nights, same as prior to taking any medication..."

* Responses from participants after spending night in sleep lab following 2 weeks of treatment course with either placebo or ZTL-101

ZTL-101

"Fell asleep and did not wake up in middle of night...more than in past 22 years."

"I haven't slept this well in a very long time."

"Slept well, night passed quickly."









ZTL-101 COMPARABLE TO CURRENT SLEEP MEDICATIONS

Criteria	ZTL-101	Temazepam (Temtabs®) Aspen	Zolpidem (Stilnox [®] /Ambian [®]) Sanofi	Suvorexant (Belsomra®) Merck	Melatonir
Rapid Sleep Onset	+/-	\checkmark	\checkmark	Х	X
Improved Sleep Maintenance	\checkmark	+/-	X	\checkmark	X
Increase Total Sleep	\checkmark	\checkmark	\checkmark	\checkmark	Х
Improve 'Quality of Sleep'	\checkmark	\checkmark	\checkmark	X	X
Feel Rested Upon Waking	\checkmark				
No Serious Side-Effects	\checkmark	+/-	+/-	+/-	X
Maintain effect long-term	TBD	X	X	\checkmark	X
No potential for addiction	TBD	X	X	\checkmark	X







- First clinically validated medicinal cannabis drug targeting chronic insomnia
- Phase 1b/2b clinical trial successfully achieved all primary and secondary endpoints
- **ZTL-101:** safe and tolerable
- **ZLT-101** is efficacious: Patients slept longer, slept sooner after waking, spent more time asleep and reported feeling rested, less stressed and fatigued the next day
- **ZTL-101** addresses unmet need for insomnia medications that improve quality of life
- On-track to launch into global markets by Q3 2020

SUMMARY FOR ZTL-101



CORPORATE



CORPORATE SNAPSHOT

FINANCIALS (as at 30 April 20	20)	SHAR
		\$0.09
		\$0.08 \$0.07
	AUD\$	\$0.06
Share Price	\$0.046	\$0.05 \$0.04
 52w Range 	\$0.022 - 0.091	\$0.03
	γ····	Ma 10M
 Current shares on issue 	966m	
 Market Capitalization 	\$44m	CAPI
• Cash (Mar 2020)	\$4.02m	Struct
 Cash Burn (2019 FY) 	\$2.6m	Direct
	γ = · · · · ·	Top 20
		Emplo

RE PRICE (as at 30 April 2020)



PITAL STRUCTURE (Fully Diluted²)

ture

- ctors Holdings: 37% 20 Shareholders: 72%
- Employee Options: 95m

Major Shareholders

- Ilera Investors 48%
- Jason Peterson 3.9%
- Harry Karelis 3.6%
- Merchant Fund 1.7%



COMPARATIVE ANALYSIS

	Company Name	Market Cap (USD)	Revenue (2019)	Market (to Reven Ratio
pharmaceuticals	GW Pharmaceuticals	\$3.0BB	\$311M	9.1x
	Tilray	\$723MM	\$167M	4.3x
Trulieve	Trulieve	\$1.4BB	\$163M	8.6x
botanix PHARMACEUTICALS	Botanix Pharmaceuticals	\$26MM	\$0	n/a
Zelira THERAPEUTICS	Zelira Therapeutics	\$37MM	\$0	n/a
TETRA BIO-PHARMA	Tetra Bio Pharma	\$72MM	\$0	n/a

Cap nue o	Product & Development Status
κ	Approved for Sativex (MS spasticity) and Epidolex [®] (childhood epilepsy), multiple Phase 1-3 trials. Synthetic THC/CBD
<	Cannabis THC/CBD Cultivation, Process, Distribution
<	Cannabis THC/CBD Cultivation, Process, Distribution

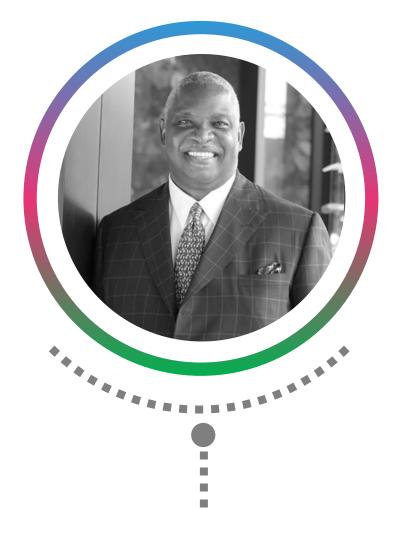
Phase 1-2 Trials. CBD only (Acne, Plaque Psoriasis, Atopic Dermatitis)

Phase 2 (insomnia), Phase 1 (Opioid Reduction), HOPE[®] launched in US.

Developing cannabinoid derived medicines targeting pain (Phase 2), ophthalmology (Phase 1) and oncology (Phase 1)







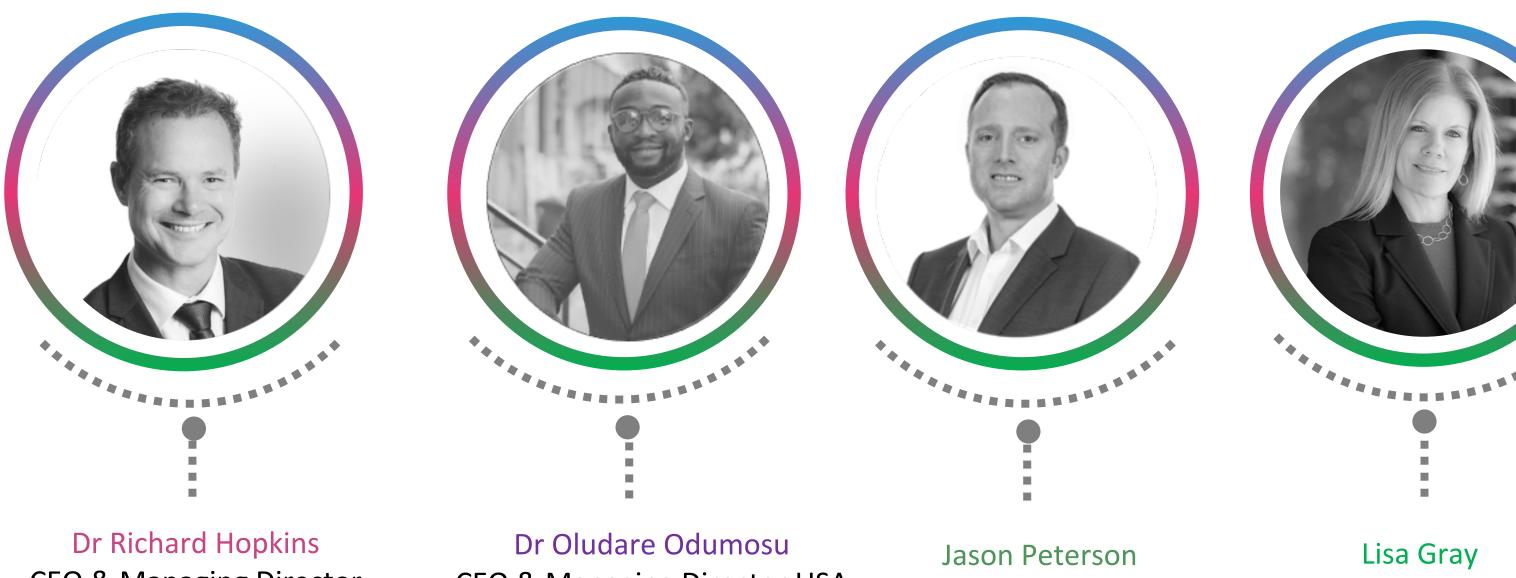
Osagie Imasogie Chairman of the Board

- Over 30 years in the field of law, finance, business management, healthcare and the pharmaceutical industry.
- Co-founder and the Senior Managing Partner of PIPV Capital, a Private Equity Firm focused on the Life Sciences vertical.
- Chairman and Founder of Ilera Healthcare, iCeutica, Inc., Churchill Pharma, Ception Therapeutics Inc. and Trigenesis Therapeutics Inc.



Harry Karelis Vice Chairman of the board

- Co-founder corporate advisory & investment firm, Gemelli Group
- Founding Director/Shareholder of numerous ASX-listed companies including in the global medicinal cannabis sector.
- 25 years experience in the financial services sector, specialising in med-tech private equity investing.



CEO & Managing Director EX-US

- Experienced bio- pharmaceutical executive.
- 19 years in corporate leadership roles. CEO/MD of 3 biotechnology companies.
- Involved in multiple pharma licensing deals generating >\$12m in revenues
- Co-founder of Phylogica.

BOARD OF DIRECTORS

- **CEO & Managing Director USA**
- Global life sciences/pharmaceutical innovation & development
- Post-clinical development of Iroko Pharmaceutical's Zorvolex® Tivorbex[®] and Vivlodex[®] through FDA approvals and successful US market commercialization.
- Founding COO of Ilera Healthcare. Ilera healthcare was acquired by TerrAscend for up to \$225M in 2019
- CSO/EVP of llera Therapeutics

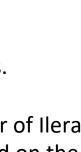
Director

- Founder, Director and Head of Corporate at Stock Broking and Corporate Advisory firm, CPS Capital.
- Founding Director/Shareholder of numerous ASX-listed companies including in the global medicinal cannabis sector.
- +25 years of experience in the financial advisory sector.

Director

- Served as COO for GlaxoSmithKline ("GSK") Pharmaceuticals Ventures.
- Co-Founder and Vice Chair of Ilera Healthcare, and was a lead on the sale of this business to TerrAscend (TER.CN).
- Vice Chair for Ilera Holistic Healthcare and Ilera Therapeutics.
- Co-Founder and Managing Partner of PIPV Capital.

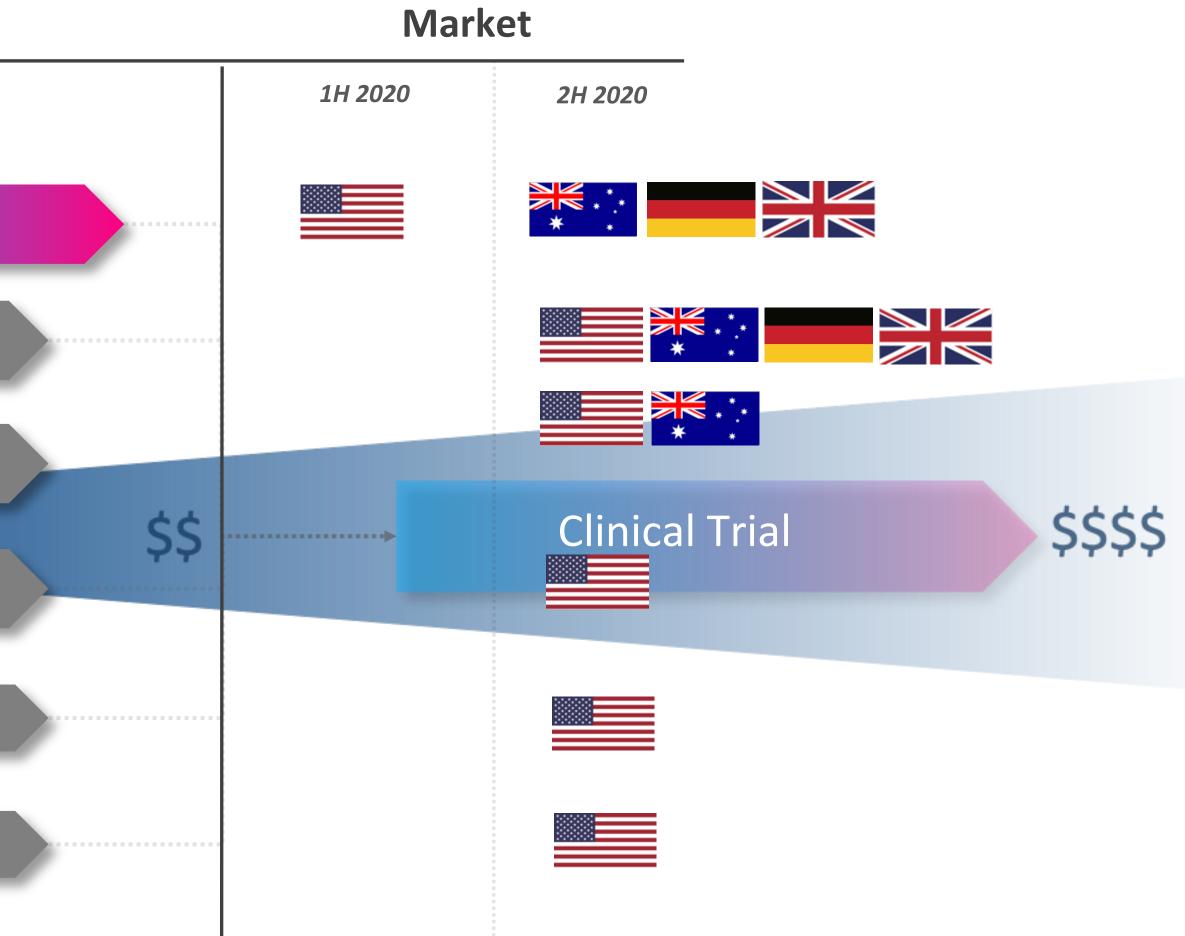




RICH PRODUCT PIPELINE LAUNCHING IN 2020



JULLER JAN		
GROWS FOR AUTISM	Autism	Launched
	Insomnia (ZTL-101)	(2020)
	Aged Disorder	(2020)
	Oral Health	(2020)
	Dermatology	2021
	Targeted Pain/GI	(2021)







UPCOMING MILESTONES

• Clinical Trial Results

- Phase II Insomnia Trial Successfully achieved primary and secondary endpoints.
- Phase Ib Opioid Sparing Trial (Reports 1H 2020). Fully recruited.

International launch of multiple proprietary products

- HOPE[™] Grows for Autism[®] (Launched in US, Q3 2020 in Australia)
- Aged Disorder (2H 2020 in US)
- Insomnia (Q3 2020 in Australia)
- Oral Healthcare (2020)
- Dermatology (2021)

• New Clinical Trials

• Autism clinical trial (2020)

• Trials targeting established pharmaceutical markets for pain and gastrointestinal therapies (2020)



ZELIRA THERAPEUTICS OFFERS INVESTORS EXPOSURE TO A RAPIDLY

EMERGING GLOBAL INDUSTRY AT VERY ATTRACTIVE VALUATIONS AND

SIGNIFICANT VALUE DRIVERS OVER THE NEXT 3-18 MONTHS

