



Investor Presentation

February 2024

Forward Looking Statements

This presentation contains forward-looking statements made within the meaning of the Private Securities Litigation Reform Act of 1995 by Anavex® Life Sciences Corp. and its representatives. These statements can be identified by introductory words such as “expects,” “plans,” “intends,” “believes,” “will,” “estimates,” “forecasts,” “projects,” or words of similar meaning, and by the fact that they do not relate strictly to historical or current facts. Forward-looking statements frequently are used in discussing potential product applications, potential collaborations, product development activities, clinical studies, regulatory submissions and approvals, and similar operating matters. Many factors may cause actual results to differ from forward-looking statements, including inaccurate assumptions and a broad variety of risks and uncertainties, some of which are known and others of which are not. Known risks and uncertainties include those identified from time to time in reports filed by Anavex Life Sciences Corp. with the Securities and Exchange Commission, which should be considered together with any forward-looking statement. No forward-looking statement is a guarantee of future results or events, and one should avoid placing undue reliance on such statements. Anavex Life Sciences Corp. undertakes no obligation to update publicly any forward-looking statements, whether as a result of new information, future events or otherwise. Anavex Life Sciences Corp. cannot be sure when or if it will be permitted by regulatory agencies to undertake clinical trials or to commence any particular phase of any clinical trials. Because of this, statements regarding the expected timing of clinical trials cannot be regarded as actual predictions of when Anavex Life Sciences Corp. will obtain regulatory approval for any “phase” of clinical trials. We also cannot be sure of the clinical outcome for efficacy or safety of our compounds. Potential investors should refer to the risk factors in our reports filed on Edgar.

Anavex Overview

NASDAQ

AVXL

HEADQUARTERS

NYC

TEAMMATES

42

CASH & CASH EQUIVALENTS¹

\$143.8M

CNS DISORDERS
MARKET OPPORTUNITY²

\$232.2B

PATIENTS WITH CNS
DISORDERS WORLDWIDE³

67.5M+

**World-wide patent protection for all
listed product candidates**



Anavex is dedicated to therapeutic discovery and development of novel small molecule treatments for central nervous system (CNS) diseases including Alzheimer's Disease, Parkinson's Disease (Dementia), Schizophrenia and Rett Syndrome, among other rare diseases



Company's precision platform, SIGMACEPTOR™, has enabled the development of drugs that unlock the body's own defenses to treat, and in some cases potentially reverse, CNS conditions



Aiming to bring the Company's treatments to market to benefit patients across the globe

Our Values



Precision Therapeutics

Every person is unique: aspire to develop therapeutic treatments tailored to suit each individual's unique medical condition



Community-based Insights

Value and rely on community-based partnerships to inform our decision making and strategy development processes



Scientific Innovation

Therapeutic development model is designed to identify and innovate novel therapeutic solutions that are rooted in science



Inclusion & Equitable Access

Global institutional need for change: committed to serving the diverse human population and to implementing inclusive and equitably accessible therapeutic programs

1. As of Dec 31, 2023.

2. Source: <https://www.alliedmarketresearch.com/central-nervous-disorders-therapeutics-market-A13121>

3. Sources available on slide 24 of this presentation.

Well-Positioned to Expand Transformative Precision Medicine Platform & Capitalize on Significant Market Opportunities



Precision medicine platform and novel central nervous system mechanism improve chance of clinical success



Several promising differentiated therapeutics for challenging CNS diseases with unmet medical needs



Achieved multiple successful clinical milestones and progressing from a research focus to a commercial stage company to bring therapies to patients around the world



Positioned for future expansion with worldwide commercial rights and strong IP foundation



Sufficient cash runway due to disciplined operations and non-dilutive cash sources, such as Michael J. Fox Foundation, International Rett Syndrome Foundation and The Australian Government

Anavex Precision Platform Enables a Novel Approach

Targeting CNS conditions with precision and restoring neuronal homeostasis via SIGMAR1 activation

Proprietary SIGMACEPTOR™ Discovery Platform produces small molecule therapeutic candidates for targeting the SIGMAR1 receptor

Age- and chronic related Changes

Chronic CNS pathologies, including progressive chronic Alzheimer's, cause exhaustion of the body's own SIGMAR1 activators, impairing the body's response to chronic cellular stress

Progressive CNS Pathology (e.g., AD, PD)

Impaired body-own compensatory SIGMAR1 response to chronic cellular stress

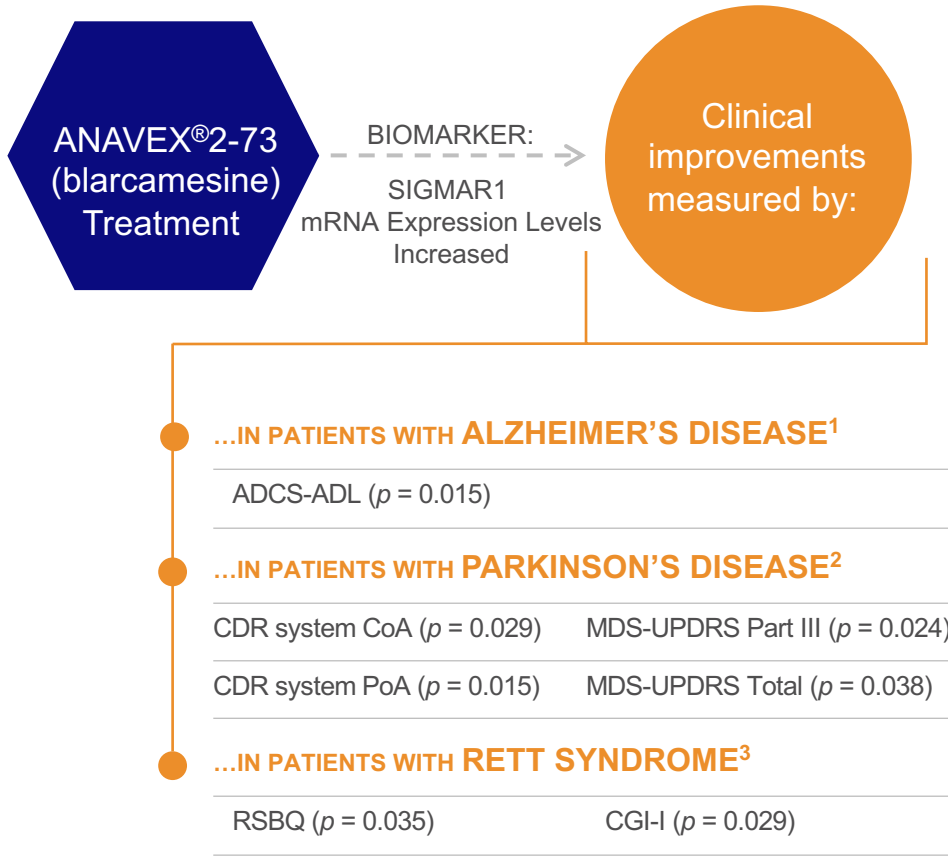
ANAVEX®2-73 (blarcamesine)

ANAVEX®2-73 (blarcamesine) re-establishes the body's own SIGMAR1 response and restores SIGMAR1 levels

Beneficial therapeutic effect for patients

SIGMAR1 target binding affinity is so specific that even when patients carry a variant receptor, still powerful effects observed. All patients regardless of genotype stand to benefit

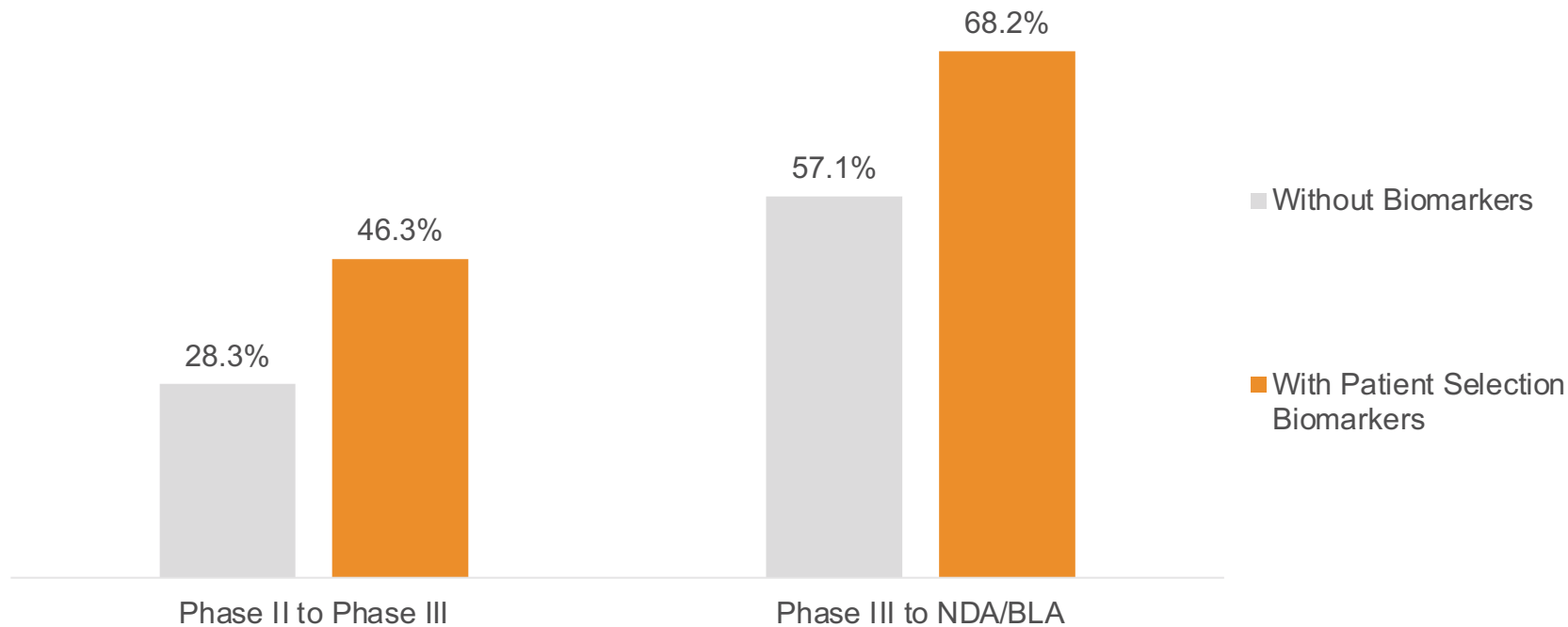
ANAVEX®2-73 Small Molecule Treatment Correlates with Efficacy Biomarkers in Alzheimer's, Parkinson's and Rett Syndrome Patients



Confirmed with independent SIGMAR1 variant analysis

1. Hampel H et al. Alzheimer's Dement. 2020;00:1–14
2. Results from ANAVEX®2-73-PDD-001 Parkinson's disease dementia study
3. Results from U.S. ANAVEX®2-73-RS-001 Rett syndrome study

Precision Medicine Increases Likelihood of Clinical Success Through Efficacy of Drug Targeting



**Biomarkers
increase
probability of
success**

Precision Platform Shows Promise for Superior Care

Anavex is positioned to target significant unmet medical needs across CNS conditions



Achieved



Near Term



Long Term

SIGMAR1 activation established as a new platform class

- ☒ **ANAVEX®2-73 (blarcamesine)**
Clinical study results in broad CNS indications confirm SIGMAR1 technology
- ☒ **Parkinson's disease dementia:** Complete data ANAVEX®2-73 Phase 2 study
- ☒ **Rett syndrome:** Complete data ANAVEX®2-73 U.S. adult Phase 2 study
- ☒ **Rett syndrome:** Top-line data AVATAR: Potentially pivotal Phase 3 adult ANAVEX®2-73 clinical trial
- ☒ **Rett syndrome:** EXCELLENCE exceeded enrollment target: Potentially pivotal Phase 2/3 pediatric clinical trial
- ☒ **Top-line data Phase 1 ANAVEX®3-71 clinical trial**
- ☒ **Alzheimer's disease:** Top-line data ANAVEX®2-73-AD-004: Potentially pivotal Phase 2b/3 clinical trial

SIGMAR1 technology to succeed traditional modalities

- ☐ **Alzheimer's disease:** Data from the blarcamesine Phase 2b/3 ANAVEX®2-73-AD-004 trial to be published in an upcoming peer-reviewed journal
- ☒ **Top-line data Rett syndrome:** EXCELLENCE Phase 2/3 pediatric clinical trial: 2H2023
- ☒ **Parkinson's disease dementia:** Data of 48-week OLE Phase 2 study
- ☐ **Parkinson's disease:** Initiation of ANAVEX®2-73 imaging-focused trial and Phase 2b/3 6 months trial
- ☐ **Fragile X:** Initiation of potentially pivotal ANAVEX®2-73 Phase 2/3 clinical trial
- ☐ **Schizophrenia:** Initiation of ANAVEX®3-71 Phase 2 clinical trial
- ☐ **New Rare disease:** Initiation of potentially pivotal ANAVEX®2-73 Phase 2/3 clinical trial
- ☐ **Publications:** Continued clinical publications involving ANAVEX®2-73 and ANAVEX®3-71

SIGMAR1 to open up new opportunities beyond the horizon

- Expanded CNS indications
- Regenerative medicine¹
- Disease prevention²

1. K. Ruscher, T. Wieloch, *The involvement of the sigma-1 receptor in neurodegeneration and neurorestoration*, *Journal of Pharmacological Sciences*, Volume 127, Issue 1, 2015, Pages 30-35, ISSN 1347-8613, <https://doi.org/10.1016/j.jpshs.2014.11.011>.

2. L. Nguyen et al., *Role of sigma-1 receptors in neurodegenerative diseases*, *Journal of Pharmacological Sciences*, Volume 127, Issue 1, 2015, Pages 17-29, ISSN 1347-8613, <https://doi.org/10.1016/j.jpshs.2014.12.005>.

Well-Positioned to Expand Transformative Precision Medicine Platform & Capitalize on Significant Market Opportunities



Precision medicine platform and novel central nervous system mechanism improve chance of clinical success



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Transformative Precision Platform Drives Impactful Therapies

Novel approach and transformative technology will likely continue to produce a variety of CNS treatments

Proprietary SIGMACEPTOR™ Discovery Platform	SIGMAR1 mRNA and SIGMAR1 variant biomarkers with genomic precision	Targeted treatment
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Late-Stage Pipeline has Achieved Clinical Milestones for Ongoing Development

ANAVEX®2-73 (blarcamesine)

- Alzheimer’s Disease
- Parkinson’s Disease Dementia
- Parkinson’s Disease
- Rett Syndrome

ANAVEX®3-71 (AF710B)

- Schizophrenia
- Frontotemporal Dementia (FTD)
- Alzheimer’s Disease

Promising Early-Stage Pipeline

ANAVEX®2-73 (blarcamesine)

- Fragile X Syndrome
- Infantile Spasms
- Angelman Syndrome
- Undisclosed Rare Disease

ANAVEX®1-41

- Depression
- Stroke
- Neurodegenerative Diseases

ANAVEX®1066

- Visceral Pain
- Acute & Neuropathic Pain

Treating Alzheimer’s Disease (AD)






Progressive, neurological disease and the most common cause of dementia¹

- Progressive development; slowly destroys memory and thinking skills
- Impacts nearly every aspect of a person’s life as it progresses: short-term memory loss and confusion, difficulty learning new things, delusions and disorientation, inability to recognize common things
- Current annual cost of dementia is estimated at \$1T, a figure set to double by 2030



~35M
people worldwide living with AD²

~6.5M
people in the U.S. living with AD²

CANDIDATE	STAGE 1	STAGE 2	STAGE 3
ANAVEX®2-73 (blarcamesine)	 ANAVEX®2-73-001	 AD ANAVEX®2-73-002/3	 AD ANAVEX®2-73-AD-004
ANAVEX®3-71	 ANAVEX®3-71-001	 (Planned Trial)	

1. Source: www.alz.org/alzheimers-dementia/what-is-dementia/types-of-dementia/parkinson-s-disease-dementia

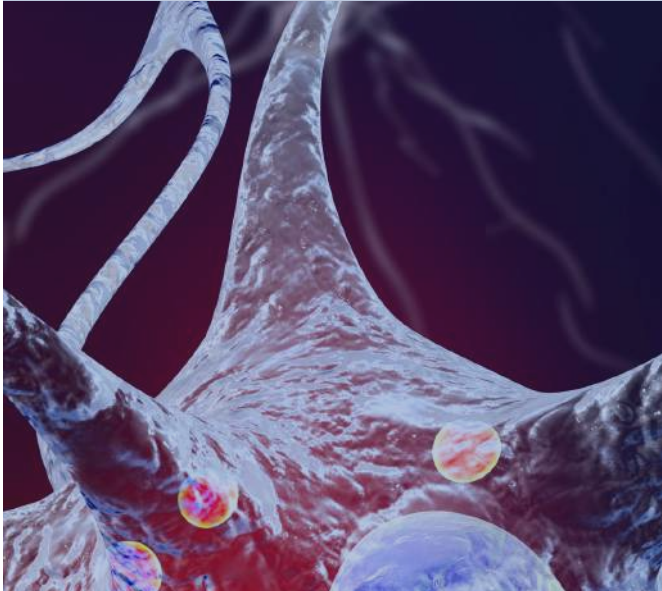
2. Sources available on slide 24 of this presentation.



Treating Parkinson's Disease (PD) & Parkinson's Disease Dementia (PDD)







Motor disorder in which patients suffer from tremors in their extremities and head, stiff limbs and inability to relax muscles during episodes

- Up to 80% of those with Parkinson's disease (PD) eventually experience Parkinson's disease dementia
- Progressive development; can cause numerous cognitive and behavioral deficits
- Parkinson's disease is a fairly common neurological disorder in older adults, estimated to affect nearly 2% of those over the age of 65



>1%
of the world population has PD

>1.5M
Americans affected by PD today







	CANDIDATE	STAGE 1	STAGE 2	STAGE 3
PARKINSON'S DISEASE	ANAVEX®2-73 (blarcamesine)	 ANAVEX®2-73-001	 (Planned Trial)	 (Planned Trial)
PARKINSON'S DISEASE DEMENTIA	ANAVEX®2-73 (blarcamesine)	 ANAVEX®2-73-001	 PDD ANAVEX®2-73-PDD-001	 (Planned Trial)

Source: Aarsland D, Creese B, Politis M, Chaudhuri KR, Ffytche DH, Weintraub D, Ballard C. Cognitive decline in Parkinson disease. Nat Rev Neurol. 2017 Apr;13(4):217-231. doi: 10.1038/nrneurol.2017.27. Epub 2017 Mar 3. PMID: 28257128; PMCID: PMC5643027; www.alz.org/alzheimers-dementia/what-is-dementia/types-of-dementia/parkinson-s-disease-dementia

Treating Rett Syndrome

Neuro-developmental disease in girls with both movement impairment and cognitive impairment¹

- Cognitive and motor delays begin to manifest along with slower head growth following relatively normal infancy (i.e., around 1.5 to 3 years of age, loss of spoken language and hand skills begin to develop)
- Associated with four key symptoms: loss of expressive language, loss of fine motor skills, impaired ability to walk and repetitive hand movement
- Second most common cause of severe intellectual disability in females

CANDIDATE	STAGE 1	STAGE 2	STAGE 3
ANAVEX®2-73 - Pediatrics (blarcamesine)	 ANAVEX®2-73-001		 EXCELLENCE ANAVEX®2-73-RS-003
ANAVEX®2-73 - Adults (blarcamesine)	 ANAVEX®2-73-001	 U.S. ANAVEX®2-73-RS-001	
ANAVEX®2-73 - Adults (blarcamesine)	 ANAVEX®2-73-001		 AVATAR ANAVEX®2-73-RS-002

~350,000
patients diagnosed with Rett
Syndrome worldwide¹

~11,000
patients diagnosed with Rett
Syndrome in the U.S.²

1. Source: <https://www.rettssyndrome.org/about-rett-syndrome>.

2. Sources available on slide 24 of this presentation.

Real World Evidence (RWE): Patients with Rett Syndrome Positive Feedback with Blarcamesine

Voice of the Patients: Real World Evidence (RWE)

- Brigitte: “*We did get a surprise once with her mobility. We heard a noise from our family room, and next we looked, and Madeline had climbed twelve steps upstairs to her bedroom by herself.*”
- Jayne: “*Within a week of starting the Anavex open label extension, she only had one seizure and then she went three months without a seizure.*”
- See related link for more video comments from parents at [RSAA/parent stories](#).
- >91% of patients completing the EXCELLENCE trial continued into a 48-week open-label extension study (OLE)
- To date, of the pediatric patients who completed the OLE, 93% have joined the Compassionate Use Program
- Compassionate Use level for adult patients from AVATAR trial after 48-week OLE is >96%
- As of today, some patients with Rett syndrome have been **on blarcamesine-treatment for >4 years**, combined OLE and Compassionate Use Program

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Multiple Clinical Milestones Achieved and Promising Ongoing Development Driving Progress towards Commercialization

CANDIDATE	PRECLINICAL	PHASE 1	PHASE 2	PHASE 3
ANAVEX®2-73 blarcamesine       	ALZHEIMER'S DISEASE			AD ANAVEX®2-73-AD-004
	PARKINSON'S DISEASE DEMENTIA		ANAVEX®2-73-PDD-001	
	PARKINSON'S DISEASE	ANAVEX®2-73-PET-001		ANAVEX®2-73-PD-001
	*RETT SYNDROME			EXCELLENCE ANAVEX®2-73-RS-003
	*RETT SYNDROME		AVATAR ANAVEX®2-73-RS-001	Fast Track, Rare Pediatric, Orphan Drug (U.S./EU)
	*RETT SYNDROME			AVATAR ANAVEX®2-73-RS-002
	*INFANTILE SPASMS			
	*FRAGILE X			
	ANGELMAN'S			
ANAVEX®3-71 AF710B	UNDISCLOSED RARE DISEASE			
	SCHIZOPHRENIA	ANAVEX®3-71-001		
	*FRONTOTEMPORAL DEMENTIA (FTD)	ANAVEX®3-71-001		
ANAVEX®1-41	ALZHEIMER'S DISEASE	ANAVEX®3-71-001		
	DEPRESSION			
	STROKE			
ANAVEX®1066	NEURODEGENERATIVE DISEASES			
	VISCERAL PAIN			
	ACUTE & NEUROPATHIC PAIN			

Legend

Solid bar = completed trials
Dashed lines = planned trials

* Orphan Drug Designation by FDA

ANAVEX[®]2-73-AD-004 Phase 2b/3 Alzheimer's Disease Trial

Global, multicenter, randomized, double-blind, placebo-controlled, parallel group, 48-week trial evaluating ANAVEX[®]2-73 (blarcamesine) once daily oral

Solid trial design produced reliable and meaningful data

N=508

Early AD patient population

- Confirmed AD pathology
- Patients aged 60 to 85 years
- MMSE score 20-28

RANDOMIZATION 1:1:1

Blarcamesine
50 mg

Blarcamesine
30 mg

Placebo

Co-Primary and Secondary Endpoints

- ADAS-Cog₁₃¹
- ADCS-ADL²
- CDR-SB³

Other Pre-specified Analyses

- Genetic variant SIGMAR1
- Structural and functional MRI
- Biomarkers: Abeta₄₀/Abeta₄₂, T-tau, P-tau, NFL, YKL-40, neurogranin, BACE1, DNA and RNA sequencing

1. AD Assessment Scale-Cognitive subscale. ADAS-Cog is the most common cognitive assessment instrument used in AD clinical trials all over the world.
2. AD Cooperative Study-Activities of Daily Living Scale. ADCS-ADL is the most common functional assessment instrument used in AD clinical trials all over the world.
3. Clinical Dementia Rating-Sum of Boxes.

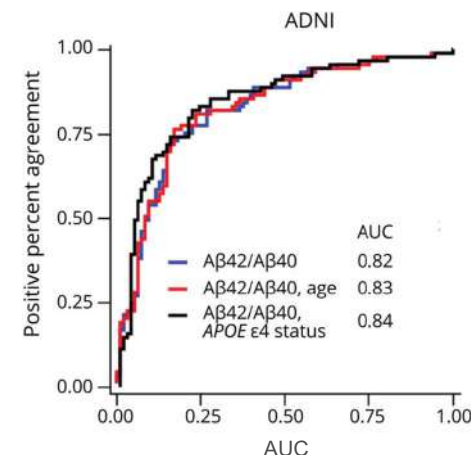
ANAVEX[®]2-73 (blarcamesine) Shows Robust Clinical Efficacy and Slows Neurodegeneration in Early Alzheimer's Disease

- Change from baseline to 48 weeks between the blarcamesine and placebo groups in least-squares mean (LSM) using mixed model for repeated measures (MMRM)¹:
 - ✓ ADAS-Cog₁₃: -1.783 [95% CI, -3.314 to -0.251]; (p = 0.0226)
 - ✓ CDR-SB: -0.456 [95% CI, -0.831 to -0.080]; (p = 0.0175)
- Validated biomarkers of amyloid beta pathology, plasma Aβ₄₂/40 ratio increased significantly (P = 0.048)
- Significant reduction in brain volume loss, including whole brain (P = 0.0005), measured through MRI
- ANAVEX[®]2-73 (blarcamesine) was generally safe and well tolerated

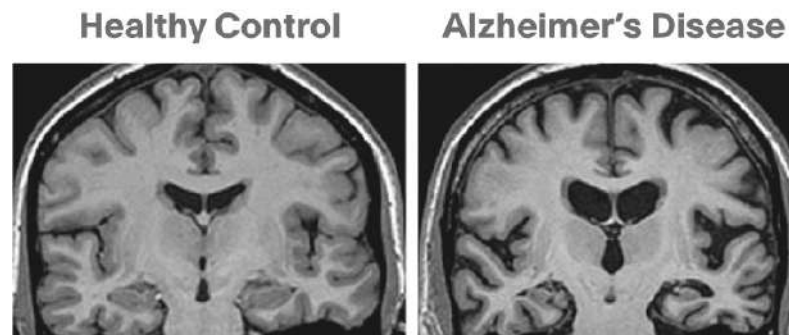
Next Steps

- Initiated regulatory submission of oral blarcamesine for Alzheimer's disease to European Medicines Agency (EMA)
- Starting to explore possible commercial activities and examining innovative strategies to effectively engage patients, providers and payers
- Data from the ANAVEX[®]2-73-AD-004 trial to be published in an upcoming peer-reviewed journal
- Continue open-label extension study ATTENTION-AD to follow participants over 96 weeks

Plasma Aβ₄₂/40 alone predicted amyloid PET status in Alzheimer's disease²



Brain volume loss (atrophy) in Alzheimer's disease³

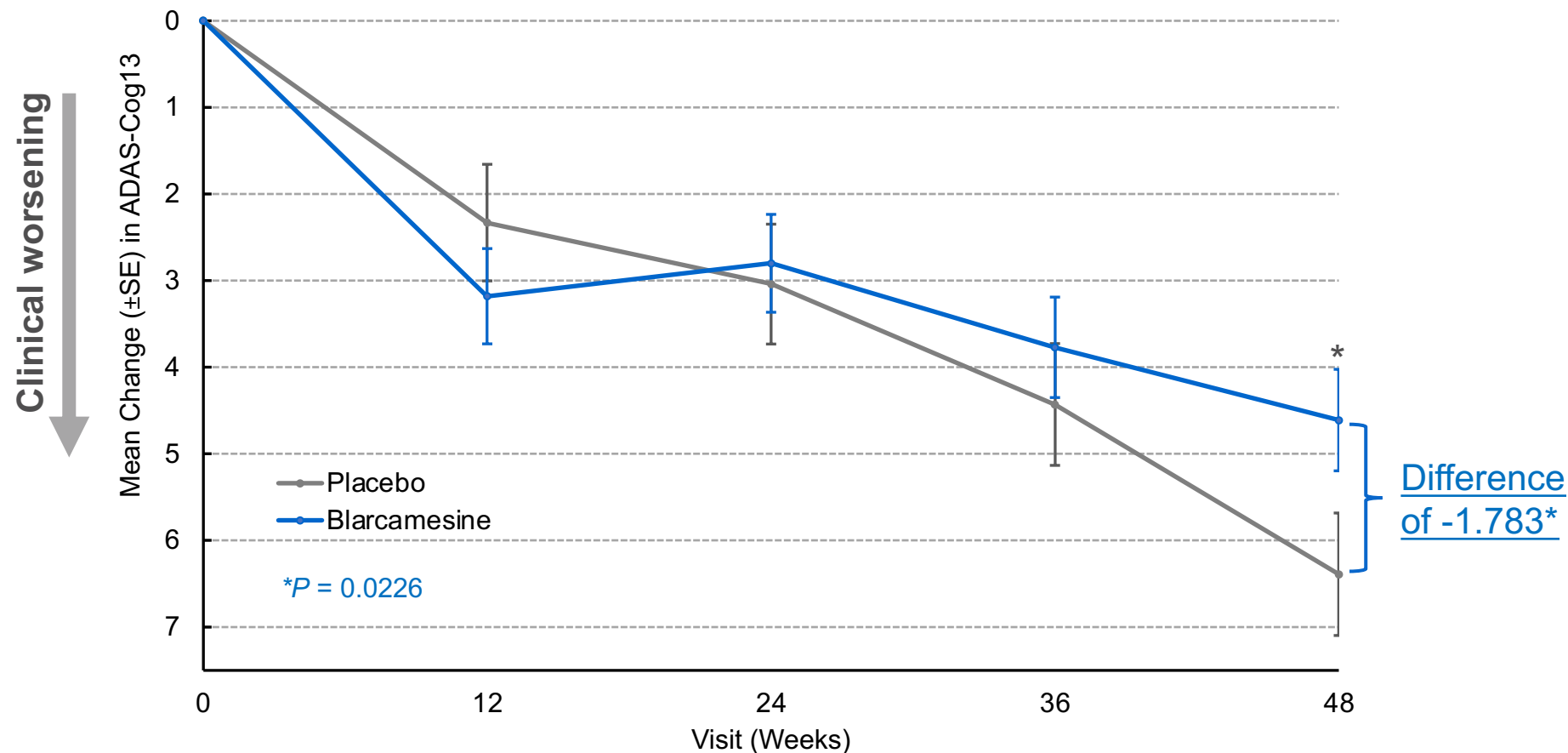


1. Gatekeeping approach: If both co-primary endpoints significant at an alpha level of 0.05, then secondary endpoint evaluated at the same alpha level. If only one primary endpoint significant at an alpha level of 0.025, then the secondary endpoint evaluated at the same level of 0.025.
2. Exemplified by Li, Yan et al. "Validation of Plasma Amyloid-β 42/40 for Detecting Alzheimer Disease Amyloid Plaques." *Neurology* vol. 98,7 (2022): e688-e699.
3. Exemplified by <https://www.omegapds.com/mri-can-help-diagnose-alzheimers-disease/>

Key Cognitive Endpoint ADAS-Cog₁₃

Blarcamesine vs Placebo at 48 Weeks

Change of -1.783 points, $P = 0.0226$; 27.9% slowing of decline at 48 Weeks



Blarcamesine at Week 48 has **stronger response** than donanemab at 76 Weeks (ADAS-Cog₁₃ difference of -1.35; 19.2% slowing: TRAILBLAZER-ALZ 2)¹

No known amyloid-related imaging abnormalities of edema or effusion observed with blarcamesine compared to donanemab

Blarcamesine **orally once daily** versus challenges of biologic mAb-based intravenous drug

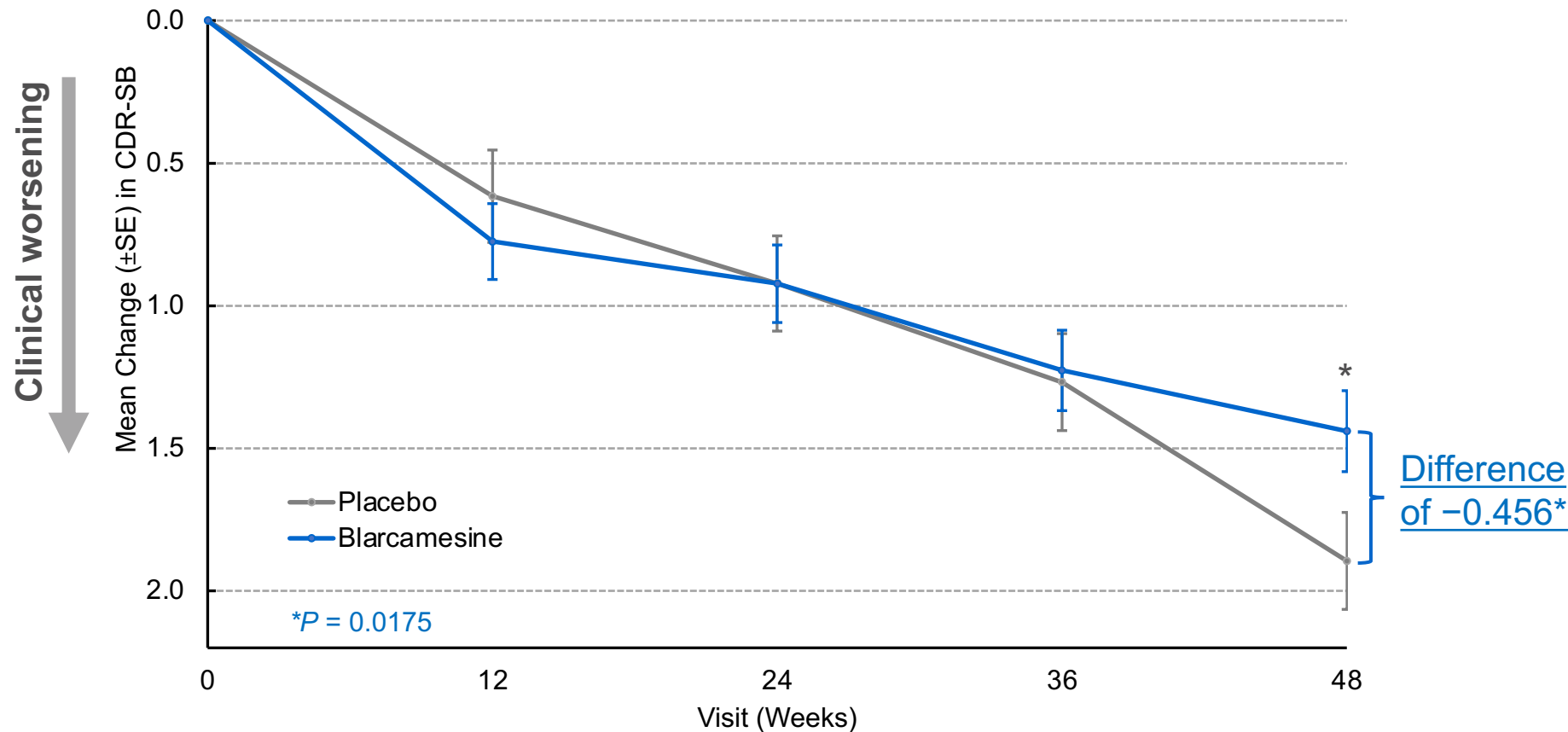
Note: Least-squares mean (LSM) using mixed model for repeated measures (MMRM); CDR-SB, Clinical Dementia Rating-Sum-of-Boxes; SE, standard error.

1. Sims et al. (2023) Donanemab in Early Symptomatic Alzheimer Disease: The TRAILBLAZER-ALZ 2 Randomized Clinical Trial. JAMA 330(6)

Key Functional and Cognitive Endpoint CDR-SB

Blarcamesine vs Placebo at 48 Weeks

Change of -0.456 points, $P = 0.0175$; 24% slowing of decline at 48 Weeks



Blarcamesine at Week 48 has **faster response** than Leqembi (lecanemab) at Week 72¹

Blarcamesine attains similar response **24 weeks earlier** despite more advanced patients

Blarcamesine **orally once daily** versus challenges of biologic mAb-based intravenous drug

Note: Least-squares mean (LSM) using mixed model for repeated measures (MMRM); CDR-SB, Clinical Dementia Rating-Sum-of-Boxes; SE, standard error.

1. Lecanemab CDR-SB -0.451 difference compared to placebo at 72 weeks; Source: https://www.eisai.com/ir/library/presentations/pdf/e4523_221130.pdf

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Worldwide Dementia Cases Projected to Grow to Over 130M by 2050

Positioned to capitalize on a significant and growing market opportunity to treat CNS diseases

>\$20T

Cumulative costs of Alzheimer's and dementia care from 2015 to 2050

1 in 3

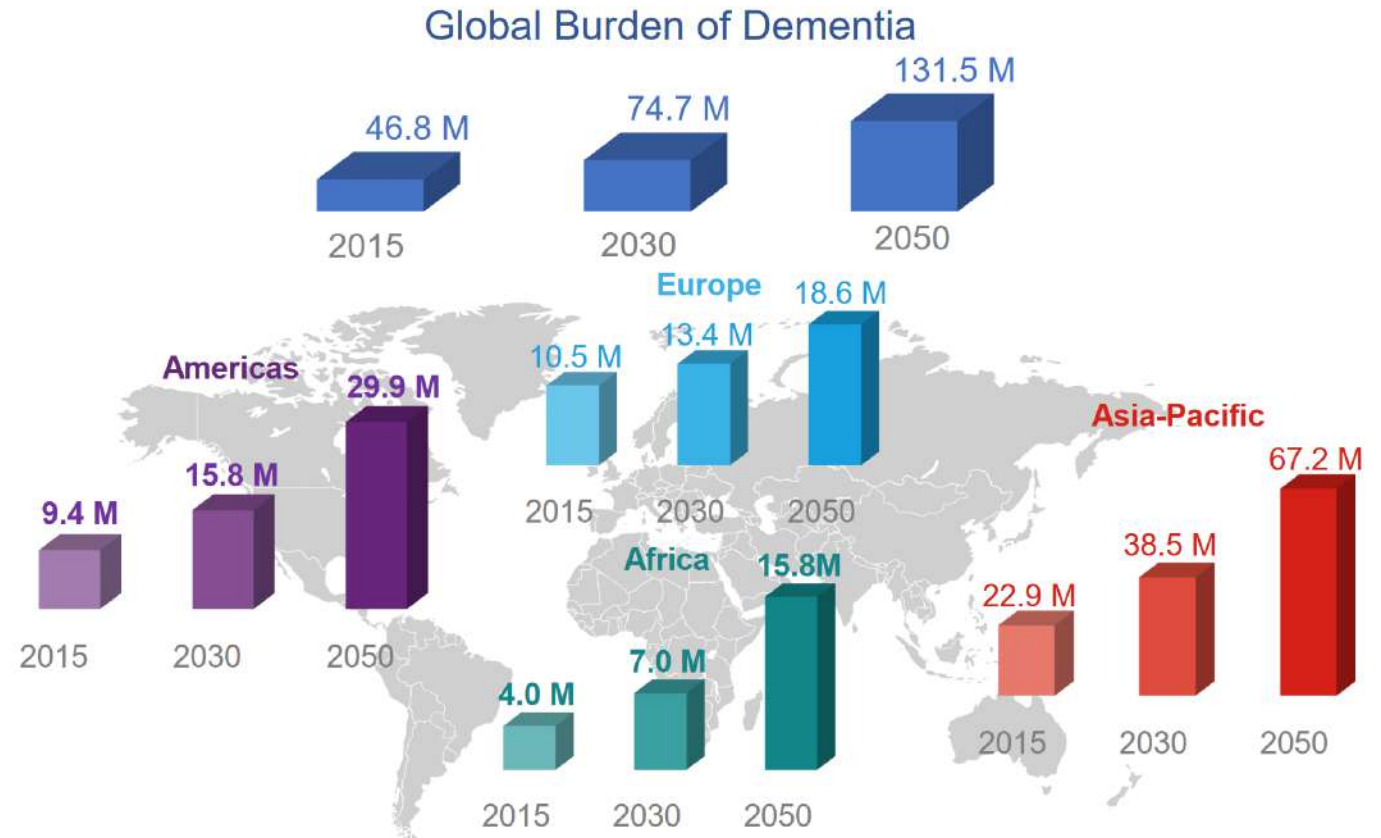
Medicare dollars will be spent on people living with Alzheimer's and other dementias in 2050

>11M

The number of Americans providing unpaid care for people with Alzheimer's or other dementias

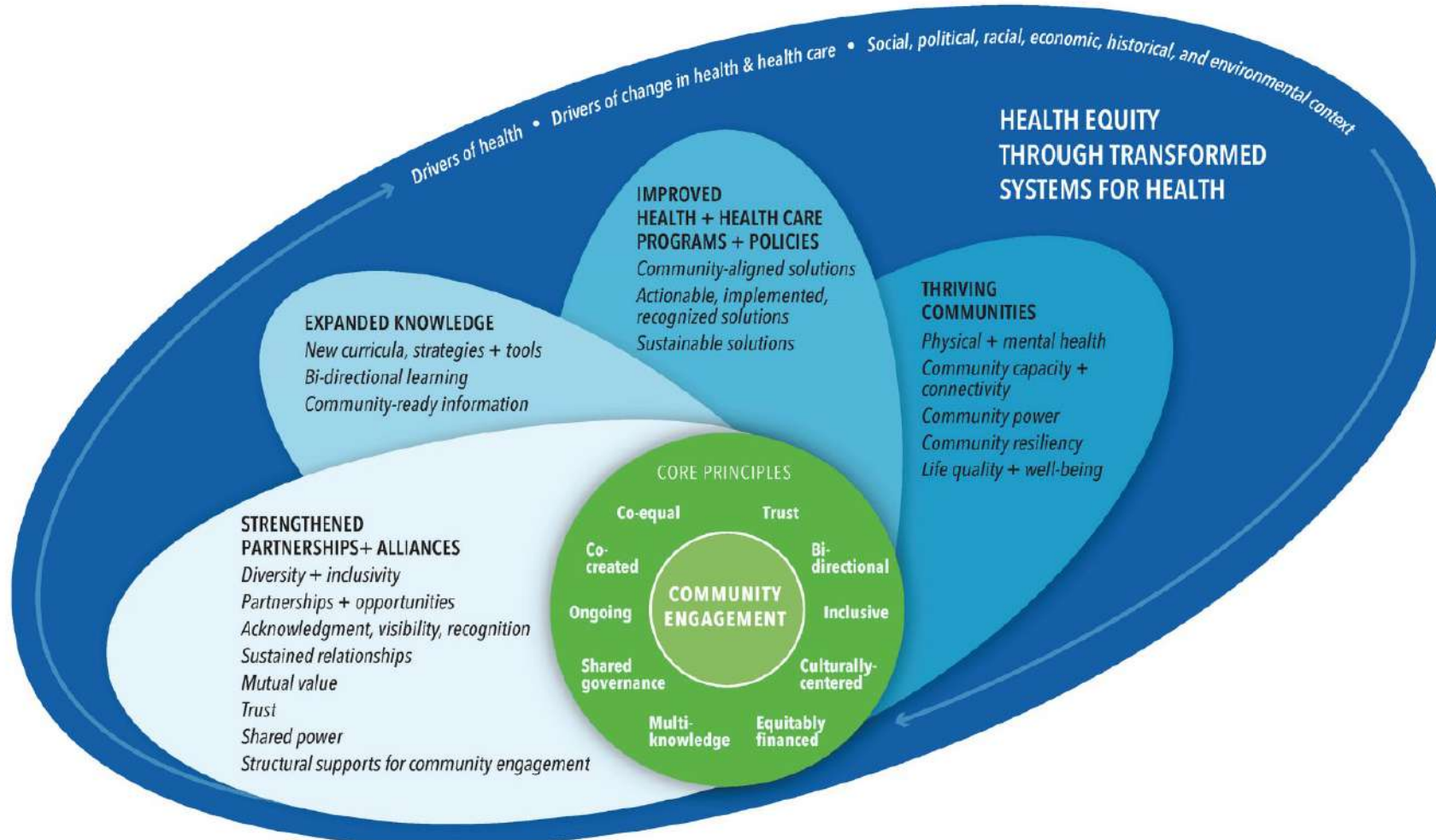
Targeting these markets using a differentiated and transformative precision platform

PEOPLE LIVING WITH DEMENTIA AROUND THE WORLD



Starting to Explore Possible Commercial Activities

Examining innovative strategies to effectively engage patients, providers and payers



High demand from Alzheimer's disease patients and families for easy access and scalable treatment options

Intended to reduce the need for complex procedures for the treatment of people with Alzheimer's disease

Blarcamesine **orally once daily** versus challenges of biologic mAb-based intravenous drug

Addressable CNS Diseases Globally with Therapeutic Disruption Potential

U.S. AND GLOBAL PATIENT NUMBERS

INDICATION	USA	EUROPE	ASIA	GLOBAL
Alzheimer's Disease (AD) ^{1,2}	~6,500,000	~7,800,000	~23,000,000	~35,000,000
Parkinson's Disease (PD) ^{3,4}	~1,000,000	~1,400,000	~3,000,000	~10,000,000
Schizophrenia ^{5,6*}	~1,600,000	~3,000,000	~9,000,000	~24,000,000
Frontotemporal Dementia (FTD) ⁷	~60,000	~65,000	~500,000	~800,000
Rett Syndrome (RTT) ^{8*}	~11,000	~13,000	~37,000	~350,000
Fragile X Syndrome (FXS) ^{9,10*}	~62,500	~150,000	~900,000	~1,400,000

1. Alzheimer's Association. 2022 Alzheimer's Disease Facts and Figures. *Alzheimers Dement* 2022;18

2. *Dementia in the Asia Pacific Region. Alzheimer's Disease International* 2014; 10

3. Marras C et al 2018. *npj Parkinson's Disease* volume 4, Article number: 21

4. GBD 2016 Parkinson's Disease Collaborators. *The Lancet* 2018 Volume 17, Issue 11, P3939-953

5. National Alliance on Mental Illness, 2019; Schizophrenia. World Health Organization. Accessed January 2024. <https://www.who.int/news-room/fact-sheets/detail/schizophrenia>

6. Fasseh et al., 2018. *Eur J Public Health*. 2018 Dec 1;28(6):1043-1049

7. Knopman & Roberts 2011. *J Mol Neurosci* 2011;45(3):330-335

8. *Rettsyndrome.org*, 2016

9. National Fragile X Foundation, 2022

10. Hunter et al., 2014. *Am J Med Genet A*. 2014 Jul;164A(7):1648-5

* Patient estimates derived from the published prevalence estimate range for the regional population

Worldwide Commercial Rights to Capitalize on Valuable Pipeline and Global Opportunity

Aiming to bring lead therapies to patients in Europe, Asia-Pacific, and the U.S. following regulatory discussion



Foundation for More Cost Effective & Safer Treatments for CNS Conditions

Oral Solid ANAVEX®2-73 (blarcamesine)	Oral Liquid ANAVEX®2-73 (blarcamesine)	Oral Solid ANAVEX®3-71 (AF710B)
<ul style="list-style-type: none">Alzheimer’s DiseaseParkinson’s DiseaseParkinson’s Disease Dementia	<ul style="list-style-type: none">Rett SyndromeFragile X SyndromeInfantile SpasmsAngelman Syndrome	<ul style="list-style-type: none">SchizophreniaFrontotemporal Dementia (FTD)Alzheimer’s Disease

Orally-administered candidates offer immense potential for clinical benefit relative to costly and logistically challenging biologic mAb-based drugs, which also often present additional safety challenges

~60%
of established small-molecule drug products available commercially are administered orally¹

~90%
of the global market share of all pharmaceutical formulations for humans are oral¹

~84%
of the best-selling pharmaceutical products are orally administered¹

1. Prasad V., De Jesús K., Mailankody S. (2017). The high price of anticancer drugs: origins, implications, barriers, solutions. Nat. Rev. Clin. Oncol. 14 (6), 381. 10.1038/nrclinonc.2017.31

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Anavex's Strong Financial Profile Supports Operations and Clinical Programs are Funded for 4 Years

Strong balance sheet supported by non-dilutive funding sources



\$143.8M

Cash and cash equivalents¹



~82M

Shares outstanding¹

**Non-dilutive
funding sources**



Disciplined approach to operational expenditures



27.8M

Fiscal year 2023 cash utilization¹

Sufficient cash runway



4

Est. Years of Runway

Sustainable cash runway due to disciplined operations and non-dilutive cash sources

Values-Driven Team with Track Record and Expertise Capable of Advancing Anavex's Cutting-Edge Precision Platform

Christopher U. Missling, PhD

President & CEO

20+ years of experience in the healthcare industry within large pharmaceutical companies, the biotech industry and investment banking



Edward R. Hammond, MD, MPH, PhD

Chief Medical Officer

15+ years of exceptional expertise in clinical drug development, including the approval of medicines and beyond



Walter E. Kaufmann, MD

Chief Scientific Officer

20+ years of experience in clinical studies with focus on developing novel therapies for genetic disorders associated with intellectual disability



David Goldberger, RPh, MLS

SVP of Regulatory Affairs

40+ years pharmacy practice and pharmaceutical industry experience



Kun Jin, PhD

VP Head of Biostatistics

27+ years of experience with US Food and Drug Administration (FDA)



Daniel Klammer, PhD

VP of Business Development & Scientific Strategy

15+ years of experience in neuroscience and the orphan disease space, with acquisition, partnering and R&D experience in Europe and the USA



Purpose-Built Scientific Advisory Board

Diverse skillset tailored
to Anavex's portfolio



CNS Drug development



Trial design and analysis



Academic and research
thought leadership



Clinical expertise in
treating CNS diseases

Andrew Cole, MD



Corinne Lasmezas, PhD



Dag Aarsland, MD, PhD



Daniel Weintraub, MD



Jacqueline French, MD



Jeffrey Cummings, MD



Norman Relkin, MD, PhD



Ottavio Arancio, MD, PhD



Paul Aisen, MD



Tangui Maurice, PhD



Timo Grimmer, MD



Marwan Sabbagh, MD



Well-Positioned to Expand Transformative Precision Medicine Platform & Capitalize on Significant Market Opportunities

\$232.2B

CNS DISORDERS
MARKET OPPORTUNITY¹

67.5M+

PATIENTS WITH CNS
DISORDERS WORLDWIDE²

1. Source: <https://www.alliedmarketresearch.com/central-nervous-disorders-therapeutics-market-A13121>

2. Sources available on slide 24 of this presentation.



Precision Medicine Platform and Novel Central Nervous System Mechanism Improve Chance of Clinical Success

Targeting CNS conditions with genomic precision and restoring neuronal homeostasis via SIGMAR1 activation



Several promising Differentiated Therapeutics for Challenging CNS Diseases with Unmet Medical Needs

Novel approach and transformative technology will likely continue to produce a variety of CNS treatments



Achieved multiple successful Clinical Milestones and Progressing from a Research focus to a Commercial stage Company to bring Therapies to Patients around the World

ANAVEX®2-73 (blarcamesine) robust clinical efficacy and slows neurodegeneration in early Alzheimer's disease



Positioned for Future Expansion with Worldwide Commercial Rights and Strong IP Foundation

Positioned to capitalize on significant market opportunities around the globe with orally-administered candidates

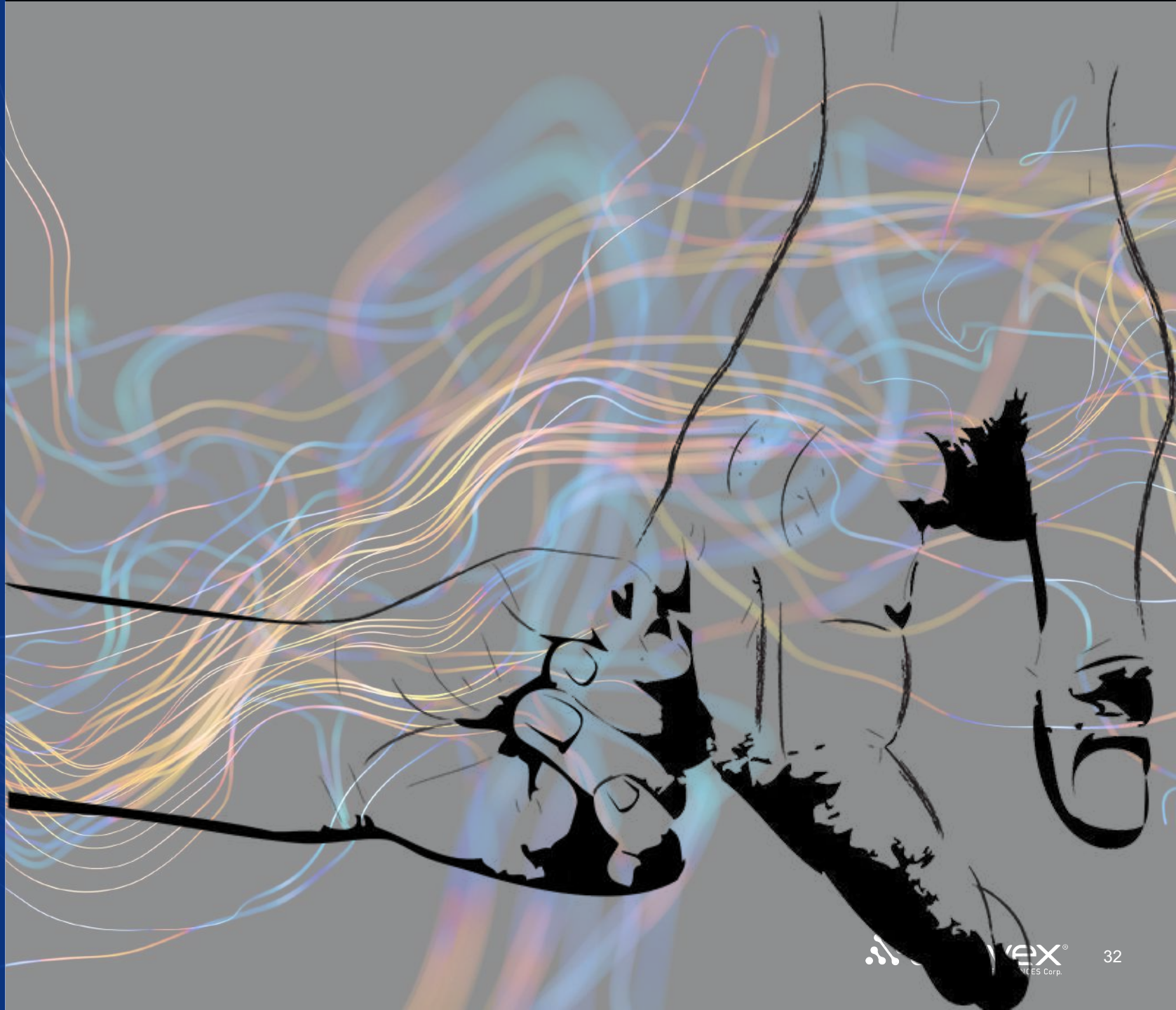


Sufficient Cash Runway Due to Disciplined Operations and Non-Dilutive Cash Sources, such as Michael J. Fox Foundation, International Rett Syndrome Foundation and the Australian government

Estimated that operations and clinical programs are funded for 4 years

Anavex's Inherent Advantage is Platform Scalability

Equitable and Accessible for Diverse Populations, and Maintaining Sustainability within Global Healthcare Systems



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