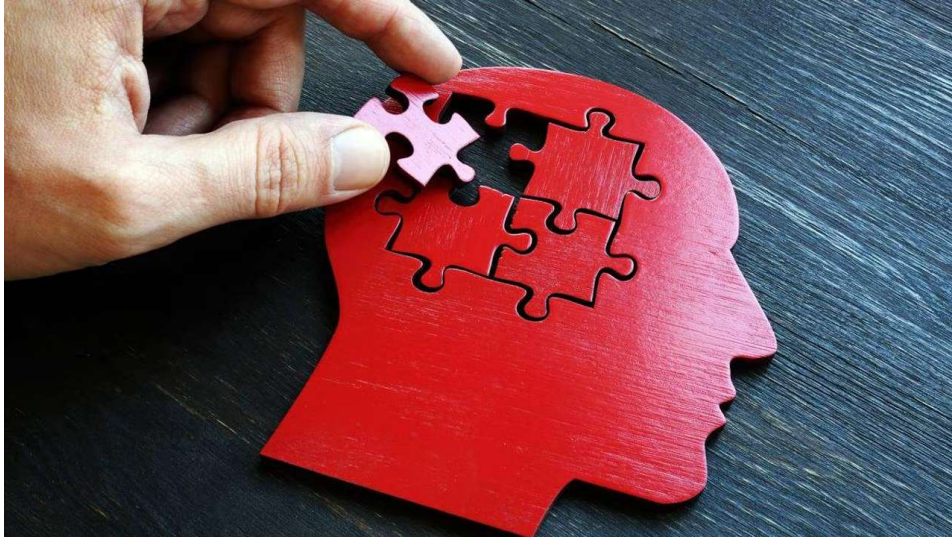


Business **Stockhead**

The dementia challenge: Which ASX



The search for a dementia breakthrough is gathering pace in Australia and around the world. Pic: Getty Images

Dementia is the leading cause of death in Australia while ranking among major causes of disability and dependency in older people



dependency among older people.
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Morning Feed: What's cooking on the ASX?
In Australia, [dementia is the leading cause of death](#), while globally it ranks among the major causes of disability and dependency in older people. Market action and updates from selected ASX-listed companies.

That research effort includes Australia, which is marking Dementia Action Week. To coincide with the week, seven Australian startups were announced as recipients of a combined \$15.6 million in funding to accelerate the



developing new therapies capable of slowing or changing its progression.

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Everything you need to know before the ASX opens global Alzheimer's drug development pipeline, with 192 clinical trials testing 158 drugs in 2026, according to an annual report led by Alzheimer's researcher Dr Jeffrey Cummings.

For ASX investors, the opportunity extends from cognitive assessments in Alzheimer's trials to potential treatments and therapies targeting the shared



focused on tackling the disease.

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Cogstate at forefront of Alzheimer's clinical trials

work to develop potential treatments for Alzheimer's disease, which is the most common cause of dementia, including US-based Eli Lilly and Japanese



(brand name Kisunla).

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Inmate's never before seen execution demand

Donanemab is a monoclonal antibody administered by intravenous (IV) infusion and designed to target and remove amyloid plaques from the brain.

The phase III trial assessed whether removing the plaques could slow cognitive and functional decline in people with early Alzheimer's disease.

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[Follow](#) In July 2026, Australia's Therapeutic Goods Administration (TGA) approved a blood test that can help identify amyloid pathology associated with Alzheimer's disease.

Roche's Elecsys Phospho-Tau (217P) Plasma (pTau217) test was developed in collaboration with Eli Lilly and was also cleared in August by the FDA.

"Eli Lilly's and Eisai's phase III trials are expected to conclude in 2027 and if they work Alzheimer's has changed forever because we now start treating people at the pre-symptomatic stage," O'Connor said.

CGS also has a Cognigram Digital Cognitive Assessment System, used by physicians to measure cognitive function and detect changes over time.

It assesses areas including processing speed, attention, visual learning, working memory, visual motor function and executive function.

The test can be used to assess cognition at a single point in time or track cognitive change through periodic assessments.

"There is a potential future where a simple blood test can identify the formation of plaques in the brain," O'Connor said.

"Cogstate digital assessments can then identify any functional impairment which will provide insight into the potential rate of decline.

"Then, through drugs approved for use in this earlier stage of disease, we can remove plaques before the horrendous memory impairment takes hold. It is super exciting."

Countdown to Actinogen's phase III Alzheimer's trial readout

[Actinogen Medical \(ASX:ACW\)](#) is the only ASX-listed healthcare company with an advanced program focused on Alzheimer's disease.

ACW's lead candidate Xanamem is in a phase 2b/3 trial dubbed XanaMIA, with topline results expected in November.

Xanamem is described by the company as the "first and only potent oral inhibitor of 11 β -HSD1", an enzyme involved in regulating cortisol in brain regions linked to memory and mood.

The oral drug candidate is designed to reduce excess cortisol production in the brain while preserving normal cortisol levels elsewhere in the body, with the aim of slowing cognitive decline.

"As national conversations around brain health take centre stage for Dementia Action Week, our team is staying sharply focused on our November trial readout," ACW chief financial officer Will Souter told Stockhead.

"We are rapidly approaching a pivotal moment that we hope will deliver a meaningful new treatment option for patients and their families."

"Dementia Action Week is a timely reminder of why we do what we do every day at Actinogen.

With trial sites fully enrolled and our upcoming November results on track, we are working hard to advance Xanamem toward becoming a potential breakthrough in Alzheimer's care."

Alterity eyes broader dementia potential for lead drug

[Alterity Therapeutics \(ASX:ATH\)](#) is developing its lead asset ATH434 for multiple system atrophy (MSA), a rare and aggressive Parkinsonian disorder.

The neurodegenerative disease, which affects up to 50,000 patients in the US, is characterised by the build-up of excess iron and aggregation of a protein called alpha-synuclein in the central nervous system (CNS).

ATH said the underlying biology is relevant across a broader group of neurodegenerative diseases.

Alpha-synuclein is a shared hallmark of MSA, Parkinson's disease, and dementia with Lewy bodies, one of the most common forms of dementia after Alzheimer's disease. Alzheimer's itself is driven by different proteins, amyloid-beta and tau.

The company describes ATH434 as an "iron chaperone", an orally administered small molecule designed to redistribute excess reactive iron within brain cells.

"ATH434 targets excess reactive iron in the brain, a key driver of alpha-synuclein misfolding," CEO Dr David Stamler told Stockhead.

"Right now our focus is on Multiple System Atrophy, a rapidly progressing disease with no approved therapy.

"The alpha-synuclein protein misfolds in a number of other neurodegenerative diseases, including dementia with Lewy bodies and in Parkinson's disease, where many patients develop dementia."

In preclinical models of MSA and Parkinson's disease, ATH434 reduced alpha-synuclein aggregation, preserved neurons, and improved motor function.

ATH is preparing for a phase III trial of ATH434 in MSA after reaching agreement with the US Food and Drug Administration (FDA) at an End-of-Phase 2 meeting on key elements of the study design, including the dose and primary endpoint. ATH434 has FDA Fast Track designation and orphan drug designation in the US and EU.

While MSA is ATH's immediate focus, the company sees potential for ATH434 to be explored in other neurodegenerative diseases, including dementia with Lewy Bodies.

"The biology we are targeting is shared across these conditions, which is why the mechanism is relevant well beyond MSA," Stamler said.

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