

# Promising Alzheimer's 'drug' halts memory loss

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A new class of experimental drug-like small molecules is showing great promise in targeting a brain enzyme to prevent early memory loss in Alzheimer's disease, according to Northwestern Medicine research.

Developed in the laboratory of D. Martin Watterson, the molecules halted memory loss and fixed damaged communication among brain cells in a mouse model of Alzheimer's.

"This is the starting point for the development of a new class of drugs," said Watterson, lead author of a paper on the study and the John G. Searle Professor of Molecular Biology and Biochemistry at Northwestern University Feinberg School of Medicine. "It's possible someday this class of drugs could be given early on to people to arrest certain aspects of Alzheimer's."

Changes in the brain start to occur ten to 15 years before serious [memory problems](#) become apparent in Alzheimer's.

"This class of drugs could be beneficial when the [nerve cells](#) are just beginning to become impaired," said Linda Van Eldik, a senior author of the paper and director of the University of Kentucky Sanders-Brown Center on Aging.

The study is a collaboration between Northwestern's Feinberg School, Columbia University Medical Center and the University of Kentucky. It will be published June 26 in the journal *PLOS ONE*.

The [novel drug](#)-like molecule, called MW108, reduces the activity of an enzyme that is over-activated during Alzheimer's and is considered a contributor to [brain inflammation](#) and impaired neuron function. Strong communication between neurons in the brain is an essential process for [memory formation](#).

"I'm not aware of any other drug that has this effect on the [central nervous system](#)," Watterson said.

"These exciting results provide new hope for developing drugs against an important molecular target in the brain," said Roderick Corriveau, program director at the National Institute of Neurological Disorders and Stroke, which helped support the research. "They also provide a promising strategy for identifying small molecule drugs designed to treat Alzheimer's disease and other neurological disorders."

Watterson and his collaborators have a new National Institutes of Health (NIH) award to further refine the compound so it is metabolically stable and safe for use in humans and develop it to the point of starting a phase 1 clinical trial.

Other senior authors on the paper are Wayne Anderson, professor of molecular pharmacology and biological chemistry at Feinberg, and Ottavio Arancio, M.D., associate professor of pathology and cell biology at Columbia.

## **Compound Strikes New Bull's Eye**

The compound strikes at a new, single target that has long flown under the radar in Alzheimer's drug development. The target is a stress-related protein kinase, p38alpha MAPK.

"We think this protein kinase target is one of the key players in the early

to mid-stage of several diseases of the central nervous system and cancer," Watterson said. Recent neuroscience research has shown that the target is activated in neurological disorders such as Alzheimer's.

These other diseases include amyotrophic lateral sclerosis (ALS or Lou Gehrig's disease), Parkinson's and frontotemporal dementia. In cancer, the protein helps malignant tumors evade the immune system as they metastasize and alter their microenvironment in a stealth type of expansion. Scientists at the University of California at San Diego currently are studying MW108 in preclinical cancer immunology research.

## **"Drug" Prevents Alzheimer's-like Symptoms in Mice**

In a key memory experiment in the study, mice brains were injected with beta-amyloid, whose increase is one hallmark of Alzheimer's in humans. One group of mice was then administered MW108 and another group was administered a placebo.

Next, each group of mice was taught environmental cues to learn how to swim through a water maze to find a resting platform. Then the mice were placed in a different arm of the maze and tested on their ability to remember the location of the platform based on the environmental cues.

The mice administered MW108 found the resting platform in the water maze as quickly as a control group of mice. The mice given the placebo made more mistakes and took longer to find the platform. They also had difficulty learning the location of the resting platform during the teaching phase.

"The results show the compound prevented the cognitive impairment," Van Eldik said.

## Can You Hear Me Now?

Another experiment in the study directly tested the compound's effect on the synaptic signal, the strength of the electrical connection between brain cells. Like a signal on your cell phone, you can't hear what the caller is saying if you don't have a strong signal. The same goes for brain cell communication.

"That's why we think people with Alzheimer's disease have trouble learning," said Columbia's Arancio, who performed the synaptic signal experiment. "If you can't transmit information from one cell to another, you can't learn."

Arancio's team took slices of a normal mouse brain and incubated them with beta-amyloid or a combination of beta-amyloid and the MW108 compound. Then they pulsed electrical current—similar to the current in a brain – through each slice and measured the strength of the electrical signal in the synapses. The mouse brain treated with beta-amyloid had a lower signal that decayed more rapidly, indicating the synapse is impaired and not communicating well. The mouse brain treated with beta-amyloid and the MW108 had a strong signal that fired normally. The compound prevented the impairment of the synapse.

## Damaging the Wiring

The over-activated protein kinase p38 MAPK damages the wiring of the communication network within the brain. Neurons communicate with other neurons at structures called synapses. Over activation of p38 MAPK damages the synapses and alters the normal functioning of neurons, impairing communication.

The p38 MAPK is also present in glial cells, which are critical to the

brain's health and comprise 90 percent of [brain cells](#). They control the strength and duration of the synaptic signal. Too much activation of p38 MAPK in glial cells impairs their supportive function and can result in the release of neurotoxic molecules further harming the synapse.

MW108 protects the brain in two different but complementary ways. By inhibiting p38 MAPK, it prevents both inflammation in the glial cells and disruption of the neuronal messages at the synapse. The result is a robust signal between neurons and within the larger communication network, which protects memory formation.

## **Designing the Right Puzzle Piece To Disable the Target**

The new drug-like probe strikes a single bull's eye that selectively disables the lone protein kinase, whose over activation is a major contributor to the [brain](#) network dysfunction. The single target approach of MW108 is novel. The prevailing view has been that multiple kinases in a network had to be disabled in order to restore normal function. There are an estimated 400-500 kinases in the human genome, with literally dozens being activated during disease processes. The research of Watterson and collaborators shows that striking the right one can be highly effective.

Watterson envisions that an array of drugs eventually will be used to treat Alzheimer's and other complex neurological diseases. He said MW108 was designed to minimize drug interaction so it could be used in combination with other drugs.

Northwestern scientists were able to rapidly design the novel small molecule MW108 to selectively disable the protein kinase because of Northwestern's premier structural genomics programs, Watterson said.

Wayne Anderson is one of the leaders of Northwestern's structural genomics programs, which use state-of-the-art expertise and technology to map the three-dimensional atomic structure of proteins involved in human disease. Anderson and Valerie Grum-Tokars, a junior structural biologist on the team, developed a three-dimensional structure of human p38 MAPK, enabling the chemists to design and synthesize novel drug-like small molecules that would disable it.

"We found the piece that fits precisely into the binding site of the protein kinase and prevents its operation," Anderson said.

Provided by Northwestern University

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