

Calcium is key to age-related memory loss

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Research at the University of Leicester is offering new clues into how and why cognitive functions such as memory and learning become impaired with age. A paper published recently in a specialist neuroscience journal shows that a crucial factor is calcium levels in specific cells in the brain.

As we get older, our memory starts to fail and it becomes harder to learn new things. It would not be unreasonable to assume that this is caused by [brain cells](#) gradually dying off but that doesn't happen. So what causes age-related cognitive impairment?

The answer lies in synapses, the electro-chemical connections between neurons that use neurotransmitter molecules to create the web of functions within the central nervous system. Professor Nick Hartell from the University of Leicester's Department of Neuroscience, Psychology and Behaviour looked at whether calcium levels in the hippocampus, part of the brain necessary for learning and memory, might play a part.

Most research in this area has concentrated on post-synaptic cells—the ones which receive neurotransmitters—simply because measuring calcium levels in pre-synaptic cells is very difficult. Nick and his colleagues stepped up to the challenge, by developing a special strain of mice which express a calcium-sensing [fluorescent protein](#) within the pre-synaptic parts of their hippocampus.

The research used mazes and object recognition tests to study the cognitive functions of mice at ages of 6, 12, 18 and 24 months, and found a clear correlation between cognitive ability and pre-synaptic [calcium levels](#). In older mice, which perform less well in the tests, the homeostatic processes that should keep intracellular calcium within limits start to falter, creating a build-up of calcium in pre-synaptic cells within the hippocampus.

Experimentally raising the level of intracellular pre-synaptic calcium in the brains of young mice altered the synaptic properties so that they behaved like those from the older mice. Most fascinating of all the results is that the reverse is also true: lowering intracellular [calcium](#) in old mouse brains rejuvenates their synapses—which obviously has enormous potential significance for age-related health issues in humans.

More information: Daniel Pereda et al. Changes in presynaptic calcium signalling accompany age-related deficits in hippocampal LTP and cognitive impairment, *Aging Cell* (2019). [DOI: 10.1111/acer.13008](https://doi.org/10.1111/acer.13008)

Provided by University of Leicester

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