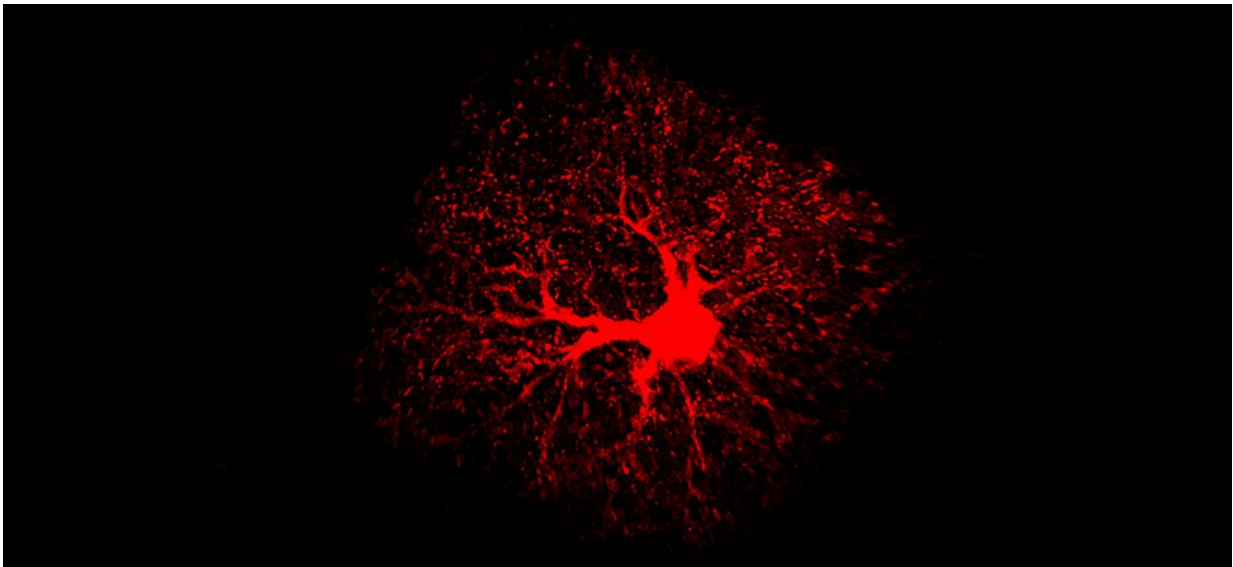


Adult astrocytes are key to learning and memory

April 24 2020, by Graciela Gutierrez



An image of an astrocyte of the adult mouse brain labeled with tdTomato red fluorescent protein. Credit: The Deneen lab

Researchers at Baylor College of Medicine reveal that astrocytes, the most abundant cells in the brain, play a direct role in the regulation of neuronal circuits involved in learning and memory. The findings appear in the journal *Neuron*.

"It has become increasingly clear that astrocytes are much more than supportive cells in the healthy adult [brain](#). They play a direct role in a

wide variety of complex and essential functions, including neuronal communication through synapses and regulation of neural circuit functions," said corresponding author Dr. Benjamin Deneen, professor of neurosurgery and a member of the Center for Stem Cell and Regenerative Medicine at Baylor. "In this study, we show a new role of astrocytes in normal brain function."

Previous work showed that astrocytes comprise diverse populations with unique cellular, molecular and functional properties. They occupy distinct brain regions, indicating regional specialization. There is evidence suggesting that [transcription factors](#)—proteins involved in controlling [gene expression](#)—regulate astrocyte diversity. Deneen and his colleagues looked to get a better understanding of the role transcription factor NFIA, a known regulator of astrocyte development, played in adult mouse brain functions.

The researchers worked with a mouse model they had genetically engineered to lack the NFIA gene specifically in adult astrocytes in the entire brain. They analyzed several brain regions, looking for alterations in astrocyte morphology, physiology and gene expression signatures.

"We found that NFIA-deficient astrocytes presented defective shapes and altered functions," said Deneen, who holds the Dr. Russell J. and Marian K. Blattner Chair and is a member of the Dan L Duncan Comprehensive Cancer Center at Baylor. "Surprisingly, although the NFIA gene was eliminated in all [brain regions](#), only the astrocytes in the hippocampus were severely altered. Other regions, such as the cortex and the brain stem, were not affected."

Astrocytes in the hippocampus also had less calcium activity—calcium is an indicator of [astrocyte](#) function—as well as a reduced ability to detect neurotransmitters released from neurons. NFIA-deficient astrocytes also were not as closely associated with neurons as normal astrocytes.

Importantly, all these morphological and functional alterations were linked to defects in the animals' ability to learn and remember, providing the first evidence that astrocytes are to some extent controlling the neuronal [circuits](#) that mediate learning and memory.

"Astrocytes in the brain are physically close to and communicate with neurons. Neurons release molecules that astrocytes can detect and respond to," Deneen said. "We propose that NFIA-deficient astrocytes are not able to 'listen' to [neurons](#) as well as normal astrocytes, and, therefore, they cannot respond appropriately by providing the support needed for efficient memory circuit function and neuronal transmission. Consequently, the circuit is disrupted, leading to impaired learning and memory."

More information: Anna Yu-Szu Huang et al, Region-Specific Transcriptional Control of Astrocyte Function Oversees Local Circuit Activities, *Neuron* (2020). [DOI: 10.1016/j.neuron.2020.03.025](https://doi.org/10.1016/j.neuron.2020.03.025)

Provided by Baylor College of Medicine

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