New class of molecules may protect brain from stroke, neurodegenerative diseases

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Credit: Louisiana State University

Research led by Nicolas Bazan, MD, PhD, Boyd Professor and Director of the Neuroscience Center of Excellence at LSU Health New Orleans, has discovered a new class of molecules in the brain that synchronize cell-to-cell communication and neuroinflammation/immune activity in
response to injury or diseases. Elovanoids (ELVs) are bioactive chemical messengers made from omega-3 very long chain polyunsaturated fatty acids (VLC-PUFAs,n-3). They are released on demand when cells are damaged or stressed.

"Although we knew about messengers from omega-3 fatty acids such as neuroprotectin D1 (22 carbons) before, the novelty of the present discovery is that elovanoids are made of 32 to 34 carbon atoms in length," notes Nicolas Bazan, MD, PhD, Boyd Professor and Director of the Neuroscience Center of Excellence at LSU Health New Orleans. "We expect that these structures will profoundly increase our understanding of cellular cross talk to sustain neuronal circuitry and particularly to restore cell equilibrium after pathological insults."

Working in neuronal cell cultures from the cerebral cortex and from the hippocampus and a model of ischemic stroke, the researchers found that elovanoids not only protected neuronal cells and promoted their survival, but helped maintain their integrity and stability. The work is published in Science Advances.

"Our findings represent a breakthrough in the understanding of how the complexity and resiliency of the brain are sustained when confronted with adversities such as stroke, Parkinson's or Alzheimer's and neuroprotection signaling needs to be activated," says Dr. Bazan. "A key factor is how neurons communicate among themselves. These novel molecules participate in communicating messages to overall synaptic organization to ensure an accurate flow of information through neuronal circuits. We know how neurons make synaptic connections with other neurons, however these connections have to be malleable to change strength appropriately. Elovanoids might play a central role as synaptic organizers, especially important in conditions resulting from synaptic dysfunction such as autism or amyotropic lateral sclerosis, for which we have no therapeutic answers."
Although the occurrence of very long chain polyunsaturated fatty acids has been well documented, what has not been known is their significance and potential to be converted into biochemical triggers to resolve injury, inflammation and other threats to neuronal communication and cell survival.
The researchers discovered the structure and characteristics of two eloovanoids - ELV-N32 and ELV-N34 - in the brain. Starting with neuron cell cultures and then an experimental model of stroke, they found that eloovanoids were activated when cells underwent either oxygen/glucose deprivation or excitotoxicity - early events associated with stroke, epilepsy, Parkinson's, traumatic brain injury and other neurodegenerative diseases. They determined the concentrations and therapeutic windows at which eloovanoids conferred neuroprotection. The team found that eloovanoids overcame the damaging effects and toxicity of these early events. In the stroke model, eloovanoids reduced the size of the damaged brain area, initiated repair mechanisms and improved neurological/behavioral recovery.

"Our findings provide a major conceptual advance of broad relevance for neuronal cell survival and brain function, particularly in regards to ischemic stroke," adds Bazan. "In the near future, we hope to apply this knowledge to prevent traumatic brain injury, chronic traumatic encephalopathy, stroke and neurodegenerative diseases. Elovanoids may offer an answer to be tested as a potential therapy."

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