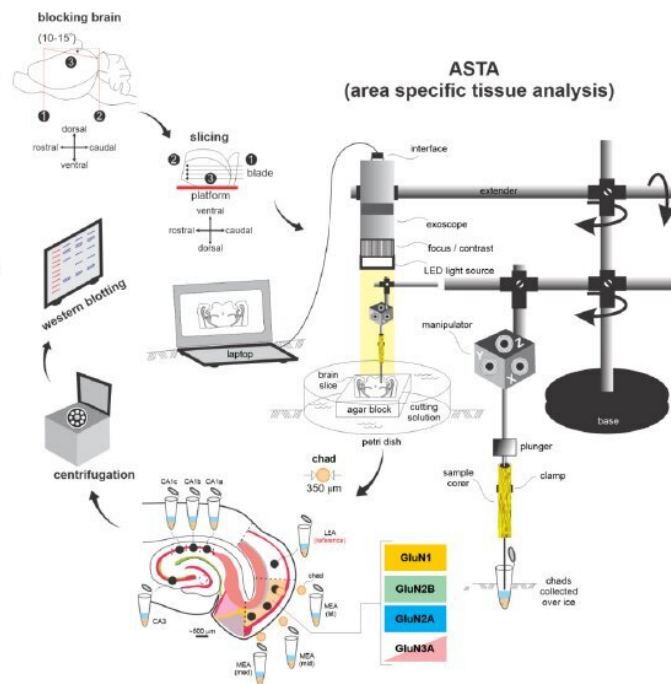


# Link between specific brain protein and vulnerability to temporal lobe epilepsy neurodegeneration discovered

July 18 2022, by Audrey Post

## GluN3 Subunit Expression Correlates with Increased Vulnerability of Hippocampus and Entorhinal Cortex to Neurodegeneration in a Model of Temporal Lobe Epilepsy

- Harvesting of *brain chads* using ASTA (area specific tissue analysis) for assaying region-specific expression of NMDAR subunits in the entorhinal cortex and the hippocampus in the context of Temporal Lobe Epilepsy (TLE).
- Our findings reveal that GluN3 subunit expression is correlated with the region-specific pattern of cell loss and neurodegeneration, observed in these regions in TLE.
- The expression profile of this subunit sheds light on the spatial extent of the pathology observed in these regions and implicates the GluN3 subunit of NMDARs in hippocampal and entorhinal cortical pathology underlying TLE.



Graphical abstract. Credit: *Journal of Neurophysiology* (2022). DOI: 10.1152/jn.00070.2022

A team of Florida State University College of Medicine researchers has

found a link between a specific protein in the brain and increased vulnerability to neurodegeneration for individuals with temporal lobe epilepsy (TLE).

Their findings are published in the *Journal of Neurophysiology*.

TLE is the most common form of epilepsy in adults and is often resistant to medication. Professor of Biomedical Sciences Sanjay Kumar, who led the study, said the team used a [novel technique](#) that made it possible to study small amounts of tissue from hard-to-reach regions within the brain. Kumar, FSU researcher Stephen Beesley and former doctoral student Thomas Sullenberger focused on a [chemical messenger](#) called [glutamate](#) and one of its receptors, N-methyl-D-aspartate (NMDA).

Glutamate plays a major role in learning and memory, and it must be present in the right concentration at the right time for the brain to function properly. It is also the body's most abundant amino acid, a building block of protein.

The team discovered that although two proteins commonly associated with NMDA—GluN1 and GluN2—were evenly distributed in a critical hippocampal region of the brain, a third one—GluN3—was distributed on a gradient. A pattern of neuron loss in the hippocampal and para-hippocampal regions of the brain is a hallmark feature of TLE.

"The relationship between GluN3 and [cell loss](#) was not known until this research," Kumar said. "This advance in [cellular biology](#) is an important step for developing therapies to help patients."

Because GluN3 makes neurons more susceptible to calcium-induced cellular damage, the discovery helps researchers narrow the focus to identify exactly where neurons are dying and in how large an area.

Kumar has applied to patent the novel technique, known as area-specific tissue analysis (ASTA), that he developed. ASTA's added precision created an improved method of testing for both the presence and volume of specific proteins linked to TLE.

Ultimately, Kumar said, discovering the gradient distribution of GluN3 will allow researchers to access more relevant tissue samples.

"This research shows how area-specific tissue analysis can be a useful tool," he said. "I'm excited to explore what further research with this technique can uncover."

The Kumar Lab focuses on deciphering the basic mechanisms underlying TLE, as well as identifying and isolating vulnerable cells and circuits within the hippocampal region to promote the discovery of more effective therapies and interventions.

**More information:** Stephen Beesley et al, GluN3 subunit expression correlates with increased vulnerability of hippocampus and entorhinal cortex to neurodegeneration in a model of temporal lobe epilepsy, *Journal of Neurophysiology* (2022). [DOI: 10.1152/jn.00070.2022](https://doi.org/10.1152/jn.00070.2022)

Provided by Florida State University

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