

## The surprising twist behind a rare neurological disorder

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Stoichiometry of transport and locations of EA6 mutations in the hEAAT1 structure. (A) The transport of each substrate molecule (glutamate or aspartate)



is coupled to the cotransport of 3 sodium ions (Na+) and 1 proton (H+), and the countertransport of a potassium ion (K+); the transport process also activates a thermodynamically uncoupled chloride channel. (B and C) Locations of EA6-related mutations in the structure of hEAAT1 (PDB: 5LLU) (B) and on a topology schematic (C) are shown. Cartoon representation of transmembrane domains (TMDs) 1 and 2 from the scaffold domain are omitted for clarity, while the whole scaffold domain is shown in transparent white surface representation. TMDs in B are colored as per C. Figure was made using ChimeraX. Credit: *Journal of Clinical Investigation* (2022). DOI: 10.1172/JCI154891

A fruit fly study, by an international group of researchers, has pinpointed how a genetic mutation causes one of the important molecular machines in our cells to malfunction. The findings provide a biological mechanism behind episodic ataxia 6, a rare neurological disorder.

Episodic ataxia 6 is a debilitating illness that severely impacts balance and coordination and causes vertigo. Patients with episodic ataxia 6 are known to have mutations in the gene SLC1A3, that codes for <u>glutamate</u> <u>transporters</u>. Glutamate transporters are molecular machines on the surface of our cells that control chemical signals and make sure our cells seamlessly talk to one another, in particular our neurons, and are vital for normal brain function.

The importance of glutamate transporters in nerve signaling and memory is widely acknowledged, but even what they look like in our cells has remined a mystery until recently.

A previous University of Sydney study found the glutamate transporter not only looks like a "twisted elevator," but is also an efficient multitasker, transporting glutamate, water and <u>chloride ions</u> at the same time.

Until now, the exact impact of the SLC1A3 mutations had on glutamate



transporters was unknown.

In this study, researchers found the SLC1A3 mutation triggered a "fault" in the "twisted elevator" mechanism, in particular the ability of the glutamate transporter to move <u>chloride</u> ions.

The mutation didn't just increase or increase chloride flow, the error went both ways.

The researchers found <u>fruit flies</u> with the genetic mutation that impacted chloride flow walked round in circles, and their movements were unsteady.

But in an unexpected twist to the story, the researchers found that the mutations only impacted chloride ion transport. There was no change to the ability to move glutamate.

The results could help our understanding of the condition and in the future develop treatments for patients suffering from episodic ataxia 6 and similar disorders.

"A key step towards improved treatment for episodic ataxia 6 is knowing whether the disease stems from an issue in glutamate transport, or the chloride channel, or even both," said senior author Professor Renae Ryan from the School of Medical Sciences, Faculty of Medicine and Health.

"Our results strongly support that the mutations impact the chloride channel."

The discovery, published in the *Journal of Clinical Investigation*, was coled by researchers from the University of Sydney, and the Research Institute of the McGill University Health Centre, in collaboration with



University of Illinois computational scientists.

## Mapping the mutations in fruit flies

"Studying the role of glutamate transporter protein mutations in episodic ataxia 6 is not straightforward. This protein has a dual function as both a glutamate transporter and as a chloride channel." Said Professor Don van Meyel, senior scientist in Brain Repair and Integrative Neuroscience Program at McGill University.

"The chloride channel function is activated by the glutamate transport function, making it difficult to disentangle one function from the other."

The research team carefully analyzed the impact of five sub types of episodic ataxia 6 related mutations.

"We studied the impacts of these mutations in frog oocytes (eggs), then using the fruit fly model, we demonstrated that an increase or decrease in glutamate transporter chloride channel activity can affect <u>motor</u> <u>function</u>," said first author Qianyi Wu, Ph.D., who was a Ph.D. student with Professor Ryan at the time of this study.

"We were able to show that several <u>mutations</u> in glutamate transporter impacted the <u>chloride channel</u> while the glutamate transporter function was unaffected."

"This suggests that episodic ataxia type 6 may be directly linked to how chloride levels are balanced in the brain, and not to glutamate transport," adds co- author Azman Akhter who was a Masters of Science student with Professor van Meyel.

"Since the glutamate transporter is highly concentrated in <u>glial cells</u> in the <u>human brain</u>, we believe they contribute to chloride homeostasis, an



idea supported by our fruit fly model."

"Additionally, we were able to identify a mechanism by which one severe mutation in the SLC1A3 gene also causes an unusual leak of sodium ions through the glutamate," explains Dr. Shashank Pant, who conducted computer modeling of the <u>glutamate</u> transporter structure in the lab of collaborator Dr. Emad Tajkhorshid at the University of Illinois.

## Science in action

This study is an example of how <u>scientific knowledge</u> about the inner workings of our cells are built up over years of research. Professor Ryan says each study along the way has been crucial to gradually building their understanding.

"This is a really nice translational study moving from basic protein characterization and computer modeling to examining its role and what happens during disease in an animal model."

**More information:** Qianyi Wu et al, Ataxia-linked SLC1A3 mutations alter EAAT1 chloride channel activity and glial regulation of CNS function, *Journal of Clinical Investigation* (2022). DOI: 10.1172/JCI154891

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