How excitatory/inhibitory balance is maintained in the brain

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Just as a thermostat is used to maintain a balanced temperature in a home, different biological processes maintain the balance of almost everything in our bodies, from temperature and oxygen to hormone and blood sugar levels. In our brains, maintaining the balance—or homeostasis—between excitation and inhibition within neural circuits is important throughout our lives, and now, researchers at the RIKEN Brain Science Institute and Nagoya University in Japan, and École Normale Supérieure in France have discovered how disturbed inhibitory connections are restored.

Published in *Cell Reports*, the work shows how inhibitory synapses are stabilized when the neurotransmitter glutamate triggers stored calcium to be released from the endoplasmic reticulum in neurons.

"Imbalances in excitation and inhibition in the brain has been linked to several disorders," explains lead author Hiroko Bannai. "In particular, forms of epilepsy and even autism appear to be related to dysfunction in inhibitory connections."

One of the key molecules that regulates excitation/inhibition balance in the brain is the inhibitory neurotransmitter GABA. When GABA binds to GABAA receptors on the outside of a neuron, it prevents that neuron from sending signals to other neurons. The strength of the inhibition can change depending on how these receptors are spaced in the neuron's membrane.

While GABAA receptors are normally clustered together, continual neural activation of NMDA receptors by the neurotransmitter glutamate—as occurs naturally during learning and memory, or in epilepsy—leads to an excess of incoming calcium, which ultimately causes the receptors to become more spread out, reducing how much the neuron can be inhibited by GABA.

To combat this effect, the receptors are somehow continually re-clustered, which maintains the proper excitatory/inhibitory balance in the brain. To understand how this is accomplished, the team focused on another signaling pathway that also begins with glutamate, and is known to be important for brain development and the control of neuronal growth.

In this pathway glutamate binds to the mGluR receptor and leads to the release of calcium from internal storage into the neuron's internal environment. Using quantum dot-single particle tracking, the team was able to show that after release, this calcium interacts with protein kinase C to promote clustering of GABAA receptors at the postsynaptic membrane—the place on a neuron that receives incoming neurotransmitters from connecting neurons.
Note that both pathways are triggered by glutamate and require calcium, although the calcium comes from different, non-overlapping sources. These opposing pathways act to keep excitation and inhibition balanced in the brain by adjusting the location of GABA receptors.

Credit: RIKEN

These findings show that glutamate activates distinct receptors and patterns of calcium signaling for opposing control of inhibitory GABA synapses.

Notes Bannai, "it was surprising that the same neurotransmitter that triggers GABA receptor dispersion from the synapse, also plays a completely opposite role in stabilizing GABA receptors, and that the processes use different calcium signaling pathways. This shows how complex our bodies are, achieving multiple functions by maximizing a limited number of biological molecules.

Pre-activation of the cluster-forming pathway completely prevented the dispersion of GABA receptors that normally results from massive excitatory input, as occurs in status epilepticus—a condition in which epileptic seizures follow one another without recover of consciousness. Bannai explains, "further study of the molecular mechanisms underlying the process we have uncovered could help develop treatments or preventative medication for pathological excitation-inhibition imbalances in the brain.

"The next step in understanding how balance is maintained in the brain is to investigate what controls which pathway is activated by glutamate. Most types of cells use calcium signals to achieve biological functions. On a more basic level, we believe that decoding these signals will help us understand a fundamental biological question: why and how are calcium signals involved in such a variety of biological phenomena?"


Provided by RIKEN