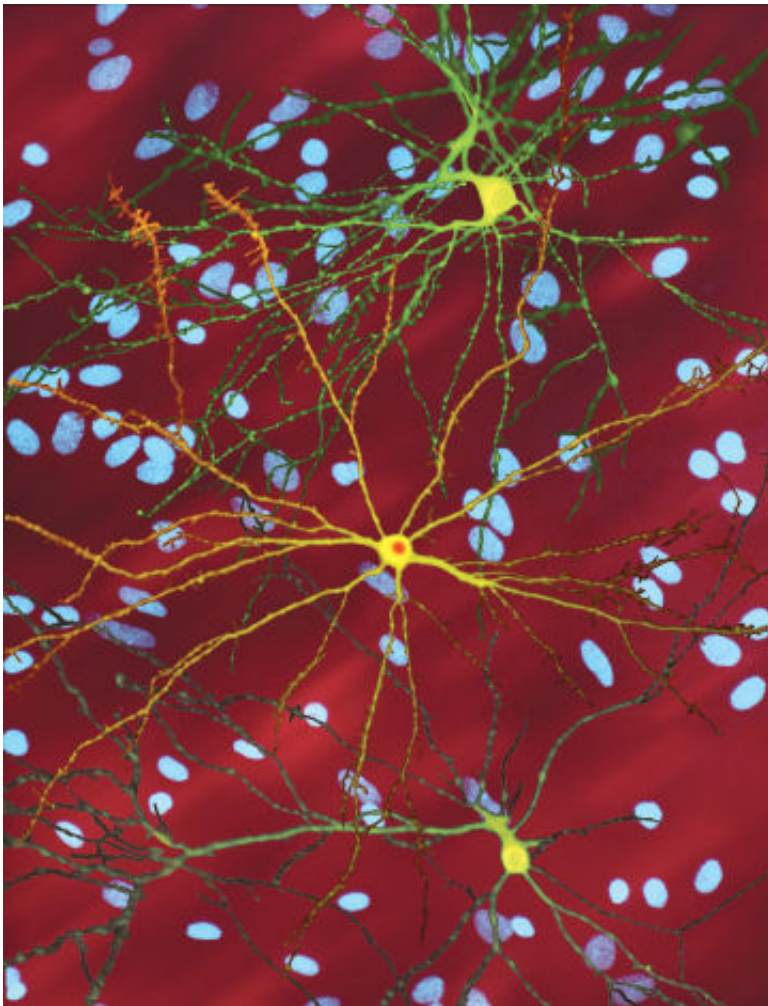


Huntington's disease linked to dysfunction of brain structure

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A montage of three images of single striatal neurons transfected with a disease-associated version of huntingtin, the protein that causes Huntington's disease. Nuclei of untransfected neurons are seen in the background (blue). The neuron in the center (yellow) contains an abnormal intracellular accumulation of huntingtin called an inclusion body (orange). Credit: Wikipedia/ Creative

Northwestern Medicine scientists identified a link between Huntington's disease and dysfunction of the subthalamic nucleus, a component of the basal ganglia, a group of brain structures critical for movement and impulse control.

Huntington's [disease](#) is characterized by the progressive loss of [nerve cells](#) in the [brain](#) and affects approximately 1 in 10,000 people. This fatal disorder is caused by a hereditary defect in a single gene.

"Although the genetic basis of the disease is well established, why the mutation leads to the expression of symptoms and loss of brain tissue remains poorly understood," explained senior author Mark Bevan, professor of physiology at Northwestern University Feinberg School of Medicine.

The study was published Dec. 20 in the journal *eLife*.

The debilitating symptoms of Huntington's disease typically manifest in adulthood and involve loss of motor and cognitive function, depression and personality changes. From the point of onset, symptoms develop and intensify over the following 10 to 25 years until death, typically due to complications associated with the disease.

"While research into Huntington's disease has focused on other parts of the [basal ganglia](#), the subthalamic [nucleus](#) has been largely overlooked," said Bevan. "This is surprising because patients with Huntington's disease have fewer nerve cells in the subthalamic nucleus. People who have suffered damage to the subthalamic nucleus exhibit excessive movement and impulsive behavior, similar to patients with Huntington's

disease."

Using mice genetically engineered to carry the Huntington's disease gene, scientists discovered the electrical activity of the subthalamic nucleus was lost. Furthermore, impaired subthalamic activity was caused by anomalous receptor signaling, leading to defective energy metabolism and accumulation of damaging oxidants. The authors also found abnormalities in the subthalamic nucleus occur earlier than in other brain regions, and that subthalamic nucleus nerve cells progressively degenerate as the mice age, mirroring the human pathology of Huntington's disease.

"Our findings suggest early problems in the [subthalamic nucleus](#) not only contribute to the symptoms of Huntington's disease, but are also likely to impair the processing capacity and health of other brain structures, more traditionally associated with the disease," Bevan said.

Currently, there is no cure for Huntington's disease; treatment can only alleviate some of the symptoms. A better understanding of aberrant brain receptor signaling that leads to nerve cell dysfunction could reveal a target for therapy, according to the authors.

Provided by Northwestern University

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