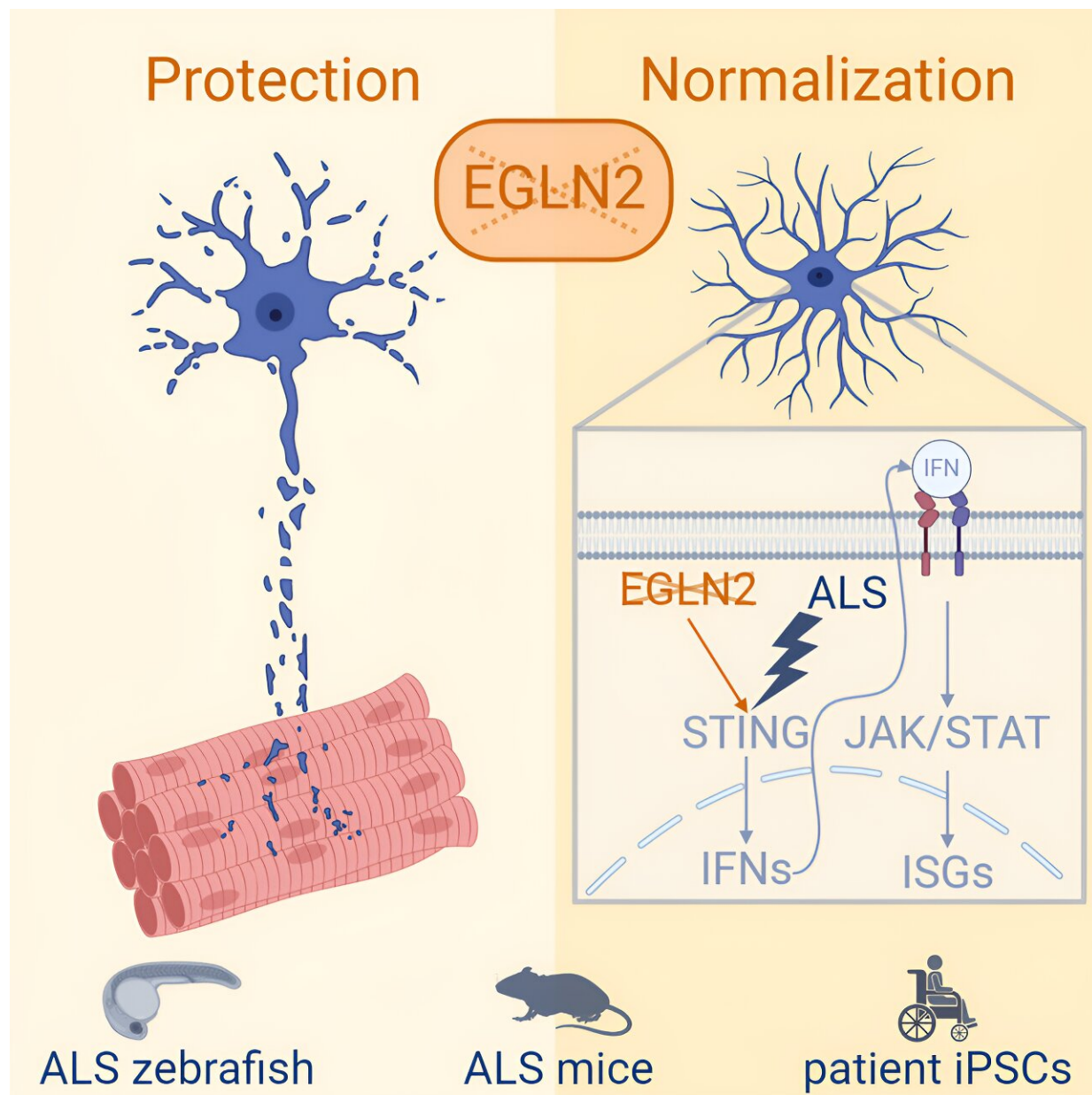


New research identifies potential therapeutic target for amyotrophic lateral sclerosis

September 10 2024



Credit: *Cell Reports* (2024). DOI: 10.1016/j.celrep.2024.114719

Amyotrophic lateral sclerosis (ALS), also known as Lou Gehrig's disease, is the most common degenerative motor neuron disease in adults. It is characterized by a selective loss of motor neurons, resulting in progressive muscle weakness and paralysis, as well as swallowing and speech difficulties. Patients usually succumb to the disease within 2 to 5 years after diagnosis.

Despite extensive research, there are currently no effective treatments to halt or reverse the progression of the disease.

From cancer to ALS research

Oxidative stress, metabolic dysfunction, and neuroinflammation are well-known hallmarks of ALS, which gave the researchers a target for their quest to understand the disease: a group of enzymes called EGLNs—metabolic sensors that regulate both cellular inflammation and metabolism.

Previously, the lab of Prof. Peter Carmeliet from the VIB-KU Leuven Center for Cancer Biology explored the role of the EGLN2 enzyme in cells experiencing low oxygen levels. The research demonstrated that inhibiting EGLN2 protected [muscle cells](#), liver cells, and cortical neurons against [oxidative stress](#).

This prompted further study into its potential protective effects on [motor neurons](#). Building on this work and in collaboration with the lab of Prof. Bart De Strooper, the Van Den Bosch team at the VIB-KU Leuven Center for Brain & Disease Research investigated the enzyme's relevance to ALS.

A potential therapeutic target

The researchers discovered that downregulating EGLN2 protects motor neurons and can mitigate the ALS phenotype in both zebrafish and mouse models. They also found that EGLN2 influences the pro-inflammatory cascade in astrocytes, a type of brain cell that supports motor neurons.

These findings suggest that targeting EGLN2 could potentially reduce inflammation and slow down the progression of ALS. The research is [published](#) in the journal *Cell Reports*.

"We used a multi-model approach to investigate the effects of EGLN2 in different cell types," says Christine Germeys, first author of the study. "We included zebrafish, mice, and induced [pluripotent stem cells](#) (iPSCs) from an ALS patient and used single-nuclei RNA sequencing to understand the underlying processes behind EGLN2 regulation."

This approach, a collaboration with the Nucleomics and Single Cell expertise units, allowed them to examine changes in [gene expression](#) at the level of individual cells, providing a detailed view of how EGLN2 influences disease progression.

"Targeting EGLN2 could represent a promising therapeutic strategy for ALS," says Prof. Ludo Van Den Bosch. "While further research is needed, this discovery brings us closer to understanding how to slow or prevent this devastating disease."

More information: Christine Germeys et al, Targeting EGLN2/PHD1 protects motor neurons and normalizes the astrocytic interferon response, *Cell Reports* (2024). [DOI: 10.1016/j.celrep.2024.114719](https://doi.org/10.1016/j.celrep.2024.114719)

Provided by VIB (the Flanders Institute for Biotechnology)

Citation: New research identifies potential therapeutic target for amyotrophic lateral sclerosis (2024, September 10) retrieved 10 September 2024 from

<https://medicalxpress.com/news/2024-09-potential-therapeutic-amyotrophic-lateral-sclerosis.html>

<p>This document is subject to copyright. Apart from any fair dealing for the purpose of private study or research, no part may be reproduced without the written permission. The content is provided for information purposes only.</p>
--