

Neuroscientists find molecular clue in ALS, suggesting potential new drug target

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Recent study from Houston Methodist Research Institute uncovers a link between motor neurons' inability to repair oxidative genome damage in ALS and suggests that DNA ligase-targeted therapies may prevent or slowdown disease progression. Credit: Hegde Lab

Houston Methodist researchers have discovered a repair defect in nerve cells that send messages to the brain, opening the door to a possible therapy for preventing or slowing down ALS. Also known as Lou



Gehrig's disease, ALS is a terminal illness marked by rapidly progressive muscle weakness that ultimately affects a person's ability to swallow and breathe.

Led by Muralidhar Hegde, Ph.D., and Haibo Wang, Ph.D., the team described their findings in an article titled "Mutant FUS causes DNA ligation defects to inhibit oxidative damage repair in Amyotrophic Lateral Sclerosis" in the Sept. 11 issue of *Nature Communications*. The new pathway they uncovered suggests that a DNA Ligase-targeted therapy may prevent or slowdown disease progression.

The research was funded by the National Institute of Health, the Muscular Dystrophy Association, ALS Association and Houston Methodist Research Institute.

Other researchers collaborating with Hegde and Wang on this paper were Joy Mitra, Pavana Hegde, Bradley Eckelmann and Sankar Mitra from Houston Methodist; Wenting Guo, Tijs Vandoorne and Ludo Van Den Bosch from Leuvan Brain Institute, Belgium; and Alan Tomkinson from University of New Mexico.

More information: Haibo Wang et al, Mutant FUS causes DNA ligation defects to inhibit oxidative damage repair in Amyotrophic Lateral Sclerosis, *Nature Communications* (2018). DOI: 10.1038/s41467-018-06111-6

Provided by Houston Methodist

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