New way to study swallowing could one day lead to improved treatments for ALS

November 14 2018

There is no cure for amyotrophic lateral sclerosis (ALS), also known as Lou Gehrig's Disease, but new findings from the University of Missouri School of Medicine and the University of Missouri College of
Veterinary Medicine are deepening researchers' understanding of a common ALS symptom: swallowing problems. Researchers have discovered how to mimic human swallowing problems in a rodent model of ALS, which will allow more targeted study on how to preserve and restore swallowing function—a major factor in quality of life. The research could one day lead to new treatments to slow the disease and improve the well-being of individuals with ALS.

"ALS is a neurodegenerative disease that makes it hard to study one aspect, like swallowing function, when the entire body is wasting away," said study co-author Teresa Lever, Ph.D., associate professor of otolaryngology at the MU School of Medicine. "We created a rodent model that allows us to study only the tongue, which is an essential muscle for several upper airway functions, including swallowing, breathing and speaking. By targeting a muscle we know is affected by ALS, we hope to better understand how we might eventually slow ALS' ability to compromise multiple upper airway functions."

ALS affects swallowing by killing the hypoglossal motor neurons that send impulses from the brain along the hypoglossal nerve to the muscles that control the tongue. As more motor neurons die, the tongue begins to weaken, and swallowing, breathing and speaking become more difficult.

Lever's co-author, Nicole Nichols, Ph.D., assistant professor of Biomedical Sciences at the MU College of Veterinary Medicine and investigator at the MU Dalton Cardiovascular Research Center, examined the impact of motor neuron death on hypoglossal nerve output, which controls tongue function.

"We have developed a unique rodent model that has reproducible hypoglossal motor neuron death and only tongue dysfunction, which will allow for further study to preserve upper airway function," Nichols said. "Specifically, we'll be able to evaluate the adaptability of the surviving
motor neurons in response to the loss of others. What we want to do next is try to make the surviving neurons work a little harder to show that we can improve tongue function both in the short and long term."


Provided by University of Missouri-Columbia

Citation: New way to study swallowing could one day lead to improved treatments for ALS (2018, November 14) retrieved 8 October 2023 from [https://medicalxpress.com/news/2018-11-swallowing-day-treatments-als.html](https://medicalxpress.com/news/2018-11-swallowing-day-treatments-als.html)

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