



Get In. We're Ending ALS

What We're Up To

On Friday, March 8, Amylyx Pharmaceuticals announced that their Phase III PHOENIX Trial for AMX0035 (RELYVRIO) did not meet prespecified primary or secondary endpoints. We share the ALS community's heartbreak and frustration over this outcome, but we appreciate Amylyx's transparency. You can read I AM ALS' full statement here:

<https://www.iamals.org/updates/phoenix-amx0035-trial-results/>

ALS Community Awards

Announcing a call for nominations for the inaugural Community Awards! These awards recognize people for their incredible leadership and contributions to furthering the movement to end ALS. Nominations will be open through April 5th.

**SUBMIT A
NOMINATION!**

COMMUNITY UPDATES



Clinical Trials Team

ALS TDI's Nadia Sethi came to the Clinical Trials Team and presented the ALS Trial Navigator. Check out this powerful tool here:

<https://www.als.net/als-trial-navigator/>

Legislative Affairs Team



The team's incredibly important and time sensitive work to secure \$75M for the ACT for ALS' Expanded Access Programs and doubling the CDMRP to \$80M for 2025 continues. We need constituents in **CA, NV, NJ, NY, OK, & TX** to support these efforts. Please click here to become a constituent and share this link far and wide to help us recruit:

<https://www.iamals.org/action/help-the-legislative-team/>



Lou Gehrig Day Team

This team is happy to announce that they will host **Jonathan Eig**, author of *Luckiest Man: The Life and Death of Lou Gehrig*, for a virtual author talk on May 1st at 7pm EST! We can't think of a better way to kick off ALS Awareness Month and get everyone energized and excited as we gear up for Lou Gehrig Day on June 2nd! Sign up to attend:

<https://www.iamals.org/action/the-luckiest-man-jonathan-eig-talks-lou-gehrig/>



Veterans Team

The Veterans Team is working hard to get the Elizabeth Dole Home Care Act passed in Congress. Help them push this bill across the finish line: use this action to contact your Senators and encourage them to support the bill. <https://www.iamals.org/action/elizabeth-dole-home-care-h-r-542-s-141/>



Youth Lou Gehrig Day Team

Here's our pitch: step up to the plate and hit a home run by joining our team! We are recruiting youth leagues to raise ALS awareness and learn about the great Lou Gehrig for the 4th annual Lou Gehrig Day. Get your youth league involved or join the team:

<https://www.iamals.org/action/youth-lou-gehrig-day/>



We've covered a lot of topics in our [Ask Me Anything ALS](#)

series, from tracheostomies and feeding tubes to finding meaning. What would you like to learn about and discuss with other people impacted by ALS? Fill out this form to help us choose future AMA topics!

https://secure.everyaction.com/T17ra3_la0e3i9tsi5i6vQ2



In Memoriam: Stacie Votaw

We are incredibly saddened to announce the passing of Stacie Votaw. Stacie was a fierce advocate for herself and for others, as well as an accomplished writer and artist. Her work was a focal point of I AM ALS' virtual art gallery in 2023. She will be deeply missed. Read Stacie's story in her own words [here](#), and check out two of her written and artistic pieces: "[Transforming Oxygen](#)" and "[Can You Speak ALS?](#)"

WHAT'S COMING UP?

March Social Hour - March 28th, 3pm ET

This month's I AM ALS Social Hour is coming up on Tuesday, March 28th! Join your fellow ALS advocates for a peaceful hour of low-key social time and conversation.

[Register](#)

2024 ALS Outlook - March 20th, 7pm ET

This webinar from Everything ALS will be hosted by Dr. Merit Cudkowicz as part of their Expert Talks Series. It will focus on the outlook for ALS in 2024.

[Register](#)

The Role of Multidisciplinary Care in ALS - March 28th, 1pm ET

This webinar from Les Turner ALS Foundation is part of their ALS Learning Series. In this webinar, team members and people living with ALS will explore the crucial role of collaboration in delivering holistic care and how multidisciplinary clinics can help people living with ALS to live

longer and with a higher quality of life.

[Register](#)

SEE ALL EVENTS



I AM ALS is a patient-led community that provides critical support and resources to patients, caregivers and loved ones. It empowers advocates to raise awareness and lead the revolution against ALS in driving the development of cures. Learn more at iamals.org.

I AM ALS



1200 Pennsylvania Ave NW, #14135 Washington, DC 20044