



Get In. We're Ending ALS

What We're Up To:

Later in March, an advisory panel will gather to understand more about AMX0035 and share its input to the FDA. This is another step in the FDA's review of Amylyx' treatment for ALS. I AM ALS is hosting a Community Conversation where we will share more about Amylyx' path to this date as well as what this advisory panel meeting will entail. Join us this Tuesday, March 1, 2022 at 2 p.m. ET by signing up [here](#).

Sign up for the Community Conversation

Join I AM ALS to learn more about the FDA's review of AMX0035.

[RSVP HERE](#)

COMMUNITY UPDATES



Clinical Trials Team

This team continues to diligently update ALS Signal with new studies and clinical trials around the world. Take a look at the team's handiwork [here](#).



Legislative Affairs Team

ACT for ALS has passed, but now we need to ensure the funding is there to do what the law permits. Want to help get ACT for ALS fully funded? Please fill out [this form](#). I AM ALS' Legislative Affairs Community Team will then be in touch with you soon!

Many Shades of ALS Team



This team is taking over I AM ALS' Twitter account this Monday, February 28 to celebrate the team's first birthday! To join in on the celebration and get all the updates, follow I AM ALS on Twitter [here](#).



Veterans Team

This team established the ALS Veterans Champion Award to recognize individuals and organizations that distinguish themselves by going above and beyond to address the needs of veterans living with ALS and their loved ones. If you know someone that fits the criteria, nominate them [here](#).



Community Spotlight: Matt Toole

Matt joined I AM ALS just recently, but he's been an ALS advocate for more than 10 years. He is an ALS Research Ambassador through NEALS and has participated in NEALS PEACe committee meetings, patient advisory boards, ALS support group meetings and many more! Read more of Matt's story [here](#).

WHAT'S COMING UP?

Ask Me Anything ALS: Tracheostomy

You got questions? We got answers! We know the decision to get a tracheostomy is complex and can be scary. Thankfully ALS community members are here to listen and answer your questions as best they can and connect you to amazing resources this coming Wednesday, March 2 at 5:30 p.m. ET.

[Register](#)

Brewfest for ALS

Join ALS TDI on Saturday, March 19 for Brewfest for ALS in Vista, CA.

Guests will be able to sample beers from local Ales for ALS brewers and enjoy food trucks and live music.

[Register](#)

Ask Me Anything ALS: Raising Kids

Supporting children while your family navigates ALS can present unique challenges. There are community members experiencing these challenges now – parents living with ALS, caregivers, ALS gene-carriers and child development professionals. They are here to listen and answer your questions as best they can and connect you to resources on April 29 at 11:30 a.m. ET.

[Register](#)



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



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