



GET IN. WE'RE ENDING ALS

"Hope is a four letter word. But not one to be afraid of." - Brian Wallach, I AM ALS Co-Founder

What We're Up To

While we seek cures, we stand committed to people living with ALS today having a chance to access treatments that could slow disease progression and give them more time. We firmly believe that even one more person with access to an ALS treatment that might work for them is progress. [Take a look at a recent statement we shared with the community about patient access to therapies.](#)

Join Us

Lou Gehrig Day in Major League Baseball is coming up, want to be in on the action?

[SIGN UP HERE](#)

COMMUNITY UPDATES



Community Outreach Team

The team members continue their campaign to diversify the image of ALS by collecting stories of women diagnosed with ALS under the age of 35. [Click here](#) to become part of the movement.



Familial ALS Team

This team is reaching out to people impacted by hereditary

diseases to get them to submit proposals to their upcoming conference, Our DNA Does Not Define Us. Interested in submitting a proposal? [Click here.](#)



Legislative Affairs Team

Community team members have contacted 519 of the 535 members of the 117th Congress. Last week alone, they contacted more than 183 congressional leaders!!! This team is powerful! [Join them!](#)



Veterans Affairs Team

The team is busy researching and creating additional resources to help improve the health and wellbeing of veterans living with ALS. [Click here to view I AM ALS current resources for veterans.](#)



Meet Seth Palmer: [Seth](#) is an incredible ALS advocate from North Carolina who lost his Granny to ALS, is currently living with ALS and is on the I AM ALS Legislative Affairs Team. Not only does he build relationships with legislators, but he's also creating legislative training sessions for advocates! Be on the lookout for his incredible trainings!

WHAT'S COMING UP?

Reflections on Clinical Trial Selections

Unite Genomics will host Sandy Morris, Marc Litt, John Russo and Tim Lowrey on April 14th to discuss how as people living with ALS they decided what clinical trials to participate in and what they take into consideration related to clinical trial design.

[Register](#)

Project ALS' Women & the Brain

Join author & sex educator Emily Nagoski, PhD for an open and invigorating discussion about the brain, sex and intimacy.

[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