



GET IN. WE'RE ENDING ALS.

"When the lights go out for the last time all I want - all I think any of us want - is to have a smile on my face knowing I did everything I could to leave this world a better place." - Brian Wallach, I AM ALS Co-Founder

What We're Up To

Swing batter batter! We're still vibing from the great news that MLB has joined the revolution and will help spread ALS awareness on Lou Gehrig Day. Can you count the ways baseball has brought hope to you in ALS? Tell us [here](#).

Make Hope Real

Join us on our mission to give the community access to critical treatments.

SIGN THE PETITION

COMMUNITY UPDATES



Clinical Trials Team

This incredible team updates [ALS Signal: Clinical Research Dashboard](#) regularly to ensure the information remains current. Give it a gander.



Community Outreach Team

These advocates want you to [join them!](#) They even made an [incredible video](#) to show you how much they want you on their team.



Legislative Affairs Team

The team really needs your help! We need more advocates to put their name on the [Every Community Map](#) to show Congress ALS impacts every single congressional district. So, what are you waiting for?



Meet [Gary Kurtis](#): Gary lost his father to ALS and since then has dedicated himself to ending ALS. And that he has! On our Legislative Affairs Team he moves ALS legislation forward by building relationships with senators and representatives in Pennsylvania, Massachusetts and New Jersey and getting them to become ALS champions.

WHAT'S COMING UP?

Becca

Target ALS is hosting [ÒBecca,Ó](#) a one-night-only concert fundraiser in memory of Rebecca Luker. Affectionately titled [ÒBecca,Ó](#) the show will feature stories and songs from Luker's career, performed by Broadway's best-known talent, and will raise much-needed funds for ALS research.

[Click here for more details](#)

Brigance Brigade Caregiver Club

Caregiver Club is a bi-monthly virtual meetup of caregivers of people living with ALS offered free of charge. The purpose of this group is to cultivate a community of ALS caregivers who can share knowledge and practice self-care.

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



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