



GET IN. WE'RE ENDING ALS

"The power to change this fight didn't come from one person or org.; instead, it was in you! Every patient, caregiver, loved one and those who lost someone to ALS. All we had to do was listen to and empower you, and you would change the world." - Brian Wallach, I AM ALS Co-Founder

What We're Up To

Each person we lose to ALS leaves behind a legacy. This week we launched a legacy wall to honor them. Take a moment to [read their stories](#) and remember the incredible souls we've lost too soon.

Celebrate

I AM ALS just turned two!
Thank you for all of your hard work and dedication!

BLOW OUT
THE CANDLES

COMMUNITY UPDATES



Community Outreach Team

The Community Outreach Team is collaborating on a content plan for I AM ALS's [LinkedIn](#). Make sure you don't miss it!



Legislative Affairs Team

The Legislative Affairs Team set an aggressive agenda for the first 100 days of the new administration. Check out their

commitments [here](#).



Veterans Affairs Team

Veterans Affairs Team member [Jeremy Van Tress](#) joined the Veterans Health Administration (VHA) ALS Executive Committee as an ex officio member. This team is relentlessly ensuring people with ALS are at the table.



Shout out to Jeremy, a member of the Familial ALS Team for spreading awareness of ALS through sharing [his story](#).

WHAT'S COMING UP? JANUARY 31 - FEBRUARY 6

The Hermstad Legacy: Advances in Treatments for ALS

Hosted by Project ALS, join special guest and ALS research advocate Ben Stiller, ALS clinicians, researchers and dedicated patient advocates to learn about the development of jacifusen D a novel gene therapy for ALS.

[Register](#)

HEALEY ALS Platform Trial Webinar

Join the webinar, CNM-Au8, Mechanism of Action and Science with representatives from Clene Nanomedicine, who will discuss CNM-Au8, which is Regimen C in the HEALEY ALS Platform Trial.

[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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