IAM ALS



GET IN. WE'RE CURING ALS

"Strength (noun): humility mixed with empathy mixed with competence mixed with just the right amount of confidence to know that none of us can do this alone. " - Brian Wallach, Co-Founder of I AM ALS

What's Up With Us

Happy (belated) Thanksgiving!!! We are thankful for you, an incredibly amazing community that <u>brings us so much</u> <u>hope</u>.

You may be wondering: Who is that Bitmoji? It's Michael, I AM ALS' community lead organizer!

Join the Party! Hey! Let's hang out. Join the I AM ALS Social Hour.

SIGN UP HERE

COMMUNITY UPDATES



Legislative Affairs Team

The HouseÕs ACT for ALS bill now has <u>193 cosponsors</u>! The Team is keeping that momentum, while also turning its attention to the <u>Senate companion bill</u>. Help this team get more ACT for ALS cosponsors in the House and Senate by adding your name to the <u>Every Community Map</u>! LetÕs show Congress that ALS impacts every congressional district.



Veterans Affairs Team

Just try keeping up with this team! The Veterans Affairs Team began its next project by outlining how to conduct a gap analysis of veteransÕ ALS health care. To find resources on veterans care, <u>click here</u>.



Meet <u>Shelly</u>: Not only is Shelly Hoover an amazing advocate on our Veterans Affairs Team and Legislative Affairs Team, but sheÕs also a talented author! Check out ShellyÕs new book <u>In RubyÕs Shoes.</u>

WHAT'S COMING UP? DECEMBER 6-13

International ALS MND Symposium

Next week will be filled with discussions on the latest advances in research and clinical management at the annual ALS MND virtual symposium. Our incredible <u>ALS Signal</u> and <u>PaCTD Rating system</u> leaders also will be hosting live poster sessions. Learn more

Clinical Trials 101

New to learning about clinical trials? YouÕre not alone. Join The Les Turner ALS FoundationÕs virtual learning session next week. <u>Register</u>

SEE ALL EVENTS

ALS IS RELEATLESS SO ARE WE



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 <u>iamals.org</u>.



1200 Pennslyvania Ave NW, #14135 Washington, DC 20044