



# GET IN. WE'RE CURING ALS

"When all is said and done, all I want is to live with purpose and change the world in some way for the better. So, let's do this." - Brian Wallach, I AM ALS Co-Founder

## What We're Up To

Join the thousands of advocates who have told their representative and senators that the ALS community deserves to have a congressional hearing with the FDA about ALS, ALS therapy development, and the FDA's progress implementing their 2019 guidelines, [Developing Drugs for Treatment Guidance for Industry](#). [Click here](#) to join the movement!

## Play Ball!

Want to help organize community events for Lou Gehrig Day?

[SIGN UP HERE](#)

## COMMUNITY UPDATES



### Clinical Trials Team

Team members frequently update ALS Signal: Clinical Trial Dashboard. Spread the word of this incredible resource by posting and sharing [this poster](#).



### Community Outreach Team

Team members are planning another panel of people living with

and impacted by ALS to educate nursing students. To request the Community Outreach Team present at your organization, click [here](#).



#### Familial ALS Team

This team is busy recruiting community members, researchers and clinicians of hereditary disease communities to submit ideas for panels for their conference, Our DNA Does Not Define Us. Help them out by sharing their [call for proposals](#).



#### Legislative Affairs Team

The team successfully contacted all US representatives and senators to put ALS on their minds and agendas. Congratulations, team!



Meet Teresa Turtle: Teresa is an avid baseball fan (Dodgers) who is so pumped for Lou Gehrig day. She is also a tireless ALS advocate, who contributes to I AM ALS Legislative Affairs, Veterans and Community Outreach Teams. Phew! That's a lot!

## WHAT'S COMING UP?

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### Celebrate Becca

Join a concert honoring the life of Broadway star Rebecca Luker, featuring stories and songs performed by some of Broadway's best-known talent.

[Register](#)

## ALS Drug Development: Ethical Considerations

Join the third installment of the NEALS webinar series on ALS drug development moderated by Dr. Richard Bedlack from the Duke ALS Clinic. This series will include talks on the preclinical development of potential therapies, the regulatory pathway, the science of trials, ethical considerations, the clinical trial site level overview, and the importance of patient engagement.

[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](http://iamals.org).

I AM ALS



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