

I AM ALS



Get In. We're Ending ALS

What We're Up To:

The second AdCom regarding AMX0035 is on September 7th. We need the FDA AdCom to know that AMX0035 is safe and effective. Help us spread the word by retweeting I AM ALS tweets about AMX0035 and joining the conversation with all the reasons you believe AMX0035 should be approved. Check out (and share!) some of these tweets [here](#) and [here](#), and keep an eye out for more. You can also read our joint written testimony, crafted with Les Turner ALS Foundation, Answer ALS Foundation, and Team Gleason [here](#).

Men living with ALS - we need you!

Our Peer Support Initiative matches people seeking guidance and advice with mentors who have a similar connection to ALS. Right now, we need more men living with ALS to serve as mentors. Join us!

[APPLY HERE](#)

COMMUNITY UPDATES



Clinical Trials Team

This team has three incredible new co-chairs! Team members Bob Hebron, Diane Hoey, and Daniel McIntyre have joined Sandy Morris to lead this team through their next chapter, and we're so excited to have them. Learn more about them and what the Clinical Trials Team is up to [here](#).

Community Outreach Team



This team would love your help spreading the word about the ALS Support Team's [Online Resource Center](#)! Please share this link on social media and on ALS forums to help more people get the information and resources they need.



Familial ALS Team

This team met with ALS4, a group dedicated to raising awareness and advocating for people with a specific form of genetic ALS, to see how these groups can collaborate moving forward. Learn more about ALS4 [here](#).



Many Shades of ALS Team

This team is busy preparing for Hispanic Heritage Month! They are working on two amazing events, and are partnering with the ALS Association Golden West chapter. They have some amazing Spanish-language resources available on their website – check them out [here](#) and share them with anyone who may need them!



Veterans Team

This team wants you to check out the Justice for ALS Veterans Act and encourage your elected officials to support this important legislation. Learn more about the bill [here](#), and send a message to your congresspeople [here](#).



The [Online Resource Center](#) is an incredible hub of information and resources for everyone living with ALS, along with their caregivers and loved ones. It includes centralized [information](#) from a variety of trusted sources, [answers to your essential questions](#) from others impacted by ALS, [checklists](#) with questions to help you prepare for care conversations, and an interactive list of [resources in your community](#).

[Community Spotlight: Chanel Hobbs](#)



Meet Chanel! Chanel is a member of the Many Shades of ALS team. She lives in Virginia with her high school sweetheart and has two adult daughters who look up to her. She shared her story with us recently about her tracheostomy, and we are so grateful for her honesty and vulnerability. Read Chanel's story [here](#), and check out all of our tracheostomy-related resources [here](#).

WHAT'S COMING UP?

FDA Advisory Committee (AdCom) Meeting Regarding AMX0035

On September 7th, an Advisory Committee will convene to give recommendations and/or answer questions posed by the FDA in regard to potential approval of AMX0035, a promising treatment for ALS. They will meet and review available evidence, written testimony, and hear oral testimony from impacted parties. This meeting will be live-streamed and available to the public – we will share that link as soon as it becomes available, so keep an eye on our social media channels!

[Learn More](#)

Home Automation for People Living with ALS

This webinar from Les Turner ALS Foundation, in partnership with Team Gleason, will discuss what home automation is and provide suggestions on how to implement current available technology. They will also discuss ways that Team Gleason can help with the home automation process. A Q&A will follow the presentation.

[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.

IAM ALS



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