Sponsorship and Involvement for the ALS Community

This month’s Connect for a Cure conversation is all about ways to get people involved before, during and after the spring and summer months. Join us on June 15th at 10:00 AM, June 16th at 10:00 AM, and June 29th at 10:00 AM. To join this month’s Zoom conversation with other members of our Walk community, just email jenna@alsmn.org.

Volunteer Leadership Certificate from the MN Association of Voluntary Nonprofits

Jenna Van Proosdy has started as our new Volunteer Coordinator! Jenna has extensive experience in volunteer management. She has her Master’s in Human Development; and her Bachelor in Business Administration with a focus in Marketing.

Help Wanted!

Volunteers are needed in the Minneapolis/St. Paul Metro Area to serve as Family Support Volunteers. Please contact Jenna at (612) 455-6052 or jenna@alsmn.org for more information.

Youth Mentoring Program

If you have a volunteer request available, or the application process, please call 612-455-6052 or email volunteer@alsmn.org.

Warm Summer Night, A Moment the Vickers Family Will Never Forget

Many believe that all it takes is one song to bring back 1,000 memories, moments we hold dear and relive in our minds every time we listen to it. No one knows this better than the Vickers family, sharing their beautiful story with us. Check out our Chapter’s Facebook page to see these segments and more from our Lou Gehrig Day.

Making ALS a Livable Disease by 2030

For decades, ALS has been a fatal disease without any cures and no significant life-extending treatments. Making ALS a livable disease means significantly longer lives for people with ALS, greater quality of life, and access to local care resources. This is achievable.

Share Your Perspective in the New ALS Focus Survey

Calling all people living with ALS and caregivers, your perspectives are urgently needed! Your responses on the current survey focuses on your experiences with telehealth. The data collected will be open and free to the entire ALS community. The current survey focuses on your experiences with telehealth. The data collected will be open and free to the entire ALS community. It only takes 10 minutes to participate after registering and all information is fully anonymous. FDA, as well as the more than 150 others from the community who have done so online. We continue to hear that people with ALS want effective treatments now, even if they offer modest benefit, and are willing to pay for them.

Please Welcome Our New Volunteer Coordinator!

We are thrilled to announce that Jenna Van Proosdy has started as our new Volunteer Coordinator! Jenna has extensive experience in volunteer management. She has her Master’s in Human Development; and her Bachelor in Business Administration with a focus in Marketing.

Help Wanted!

Volunteers are needed in the Minneapolis/St. Paul Metro Area to serve as Family Support Volunteers. Please contact Jenna at (612) 455-6052 or jenna@alsmn.org for more information.

Warm Summer Night, A Moment the Vickers Family Will Never Forget

Many believe that all it takes is one song to bring back 1,000 memories, moments we hold dear and relive in our minds every time we listen to it. No one knows this better than the Vickers family, sharing their beautiful story with us. Check out our Chapter’s Facebook page to see these segments and more from our Lou Gehrig Day.

Making ALS a Livable Disease by 2030

For decades, ALS has been a fatal disease without any cures and no significant life-extending treatments. Making ALS a livable disease means significantly longer lives for people with ALS, greater quality of life, and access to local care resources. This is achievable.

Share Your Perspective in the New ALS Focus Survey

Calling all people living with ALS and caregivers, your perspectives are urgently needed! Your responses on the current survey focuses on your experiences with telehealth. The data collected will be open and free to the entire ALS community. The current survey focuses on your experiences with telehealth. The data collected will be open and free to the entire ALS community. It only takes 10 minutes to participate after registering and all information is fully anonymous. FDA, as well as the more than 150 others from the community who have done so online. We continue to hear that people with ALS want effective treatments now, even if they offer modest benefit, and are willing to pay for them.

Please Welcome Our New Volunteer Coordinator!

We are thrilled to announce that Jenna Van Proosdy has started as our new Volunteer Coordinator! Jenna has extensive experience in volunteer management. She has her Master’s in Human Development; and her Bachelor in Business Administration with a focus in Marketing.

Help Wanted!

Volunteers are needed in the Minneapolis/St. Paul Metro Area to serve as Family Support Volunteers. Please contact Jenna at (612) 455-6052 or jenna@alsmn.org for more information.