



“What do you do when you are diagnosed with a rare, terminal, neurodegenerative disorder with no cure? All the internet searches were so disheartening, and most doctors have no experience with MSA which can delay diagnosis and symptom management for years. I found [Mission MSA], and there were other people like me all over the world! This international group advocates for research, fundraising, and provides support for patients and caregivers. This organization has given me hope.”

- BELINDA F., MSA PATIENT



“[Mission MSA] has been a lifeline for my family since 2013 when my husband was diagnosed with MSA. This organization has shared so much valuable information and has had a positive impact on our MSA journey. The volunteers are incredibly supportive. We rely on this group so much.

- OLGA C., MSA CARE PARTNER



“I want to thank [Mission MSA] for the helpful information they supplied me. With their help, I was able to put together an information package for staff at the long-term care facility in which my patient resides and present an in-service educating the staff about this horrific and rare disease.”

- JENNIFER C., R.N., B.N., CHPCN(C)



MISSION MSA™

Leading the Charge to Cure
Multiple System Atrophy

Connect with the MSA Community!



Contact

Support Hotline: 866.737.5999

Office Line: 866.737.4999

Email: info@missionmsa.org

Mailing Address:

1660 International Drive, Suite 600
McLean, VA 22102, United States of America



Visit us online at
www.missionmsa.org



MISSION MSA™

Serving the MSA community since 1989.

Enhancing quality of life,
and building hope for people
affected by Multiple System Atrophy



About Mission MSA

At Mission MSA, we are dedicated to improving the quality of life and building hope for people affected by multiple system atrophy through support services, educational resources, research funding, and community engagement.

Starting out as a small support group in 1989, Mission MSA has grown into a sizable 501(c)(3) charitable organization with international impact. With over 30 years of experience in MSA advocacy, we have made significant strides in supporting patients and their families and will continue to provide these services until we have reached a future without MSA.



Support

Facilitating Services for Patients & Care Partners

Patient Support Line: 866-737-5999

Mission MSA is here to provide guidance, information, and community resources to address any questions or concerns you may have.

Virtual & In-Person Support Groups

Join support groups online or in person across the U.S. and abroad. Check out the interactive map of regional support groups on our website.

Community

Offering Opportunities to Connect With Others on Their MSA Journey

MSA Connect

MSA Connect is a dedicated online platform just for the MSA community where you can ask questions, share experiences, and join discussions with others who are going through a similar journey.

Education

Developing & Sharing Reliable Resources

Monthly Webinars

Mission MSA offers monthly webinar programming on a variety of topics related to multiple system atrophy including coping with symptoms, financial planning, finding your care team, and more.

Annual Patient & Family Conferences

Mission MSA's annual conference is the largest MSA conference and features panels and presentations by medical professionals, scientists, patients, and care partners.

Research

Funding Medical Professionals in Search of Better Treatments and Faster Diagnosis

Centers of Excellence

The MSA Centers of Excellence program was created to provide access to the best possible and easily accessible multi-disciplinary clinical care and support services for individuals affected by MSA and their families. Find a full list of these centers on our website.