Looking for a doctor, support group or treatment summary?

Email us with your request!
info@defeatmsa.org

Email us about setting up a named legacy fund or fundraiser!

info@defeatmsa.org

Support Research and those suffering with MSA now!

We are registered in all US states for fundraising!

This brochure is for promotional purposes only and should not be used for diagnosis or treatment. Anyone with questions about MSA should consult with their doctor or other health care professional for diagnosis or treatment options.

Make Checks Payable:

Defeat MSA Alliance
29924 Jefferson Avenue
St. Clair Shores, Michigan
48082 USA

Visit Our Partners:

MSA Awareness Shoe
www.msashoe.org

MSA United Research
www.msaunited.org

Defeat MSA Awareness Shoe Charity - USA 501(c)(3)
29924 Jefferson Avenue
Saint Clair Shores, Michigan 48082
USA
EIN: 47-2700163
MSA Shoe, Trademark Pending TM

We move today, in service of tomorrow.

DEFEAT MSA ALLIANCE (USA)

Call Our Toll Free Help Line:

1(855) KICK-MSA
1(855) 542-5672

www.DefeatMSA.org

501(c)(3); EIN: 46-0661655
A NonProfit Public Charity

What is MSA?

Multiple System Atrophy (MSA) is a rare, rapidly progressing neurodegenerative disorder. MSA impairs the systems that regulate blood pressure, heart rate and the bladder – many of the basic bodily functions that people take for granted every day. People with MSA suffer from dangerously low blood pressure, speech and swallowing difficulties, sleep disturbances, breathing problems, rigidity and tremors. The life expectancy for those with MSA is typically 5 to 10 years. Patients with advanced MSA often become bed-bound, unable to speak and immobile. At present, there is no cure for MSA, no genetic tests to detect it, no therapy to slow it and very few treatments to manage its debilitating effects.

Our Mission

Defeat MSA Alliance is an inclusive US based 501(c)(3) public charity that aspires to balance efforts to support patients, educate medical professionals, raise public awareness, nurture promising research and advocate for the MSA community.

Awareness. Alleviation. Advancement.

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*Volunteer experts provide additional help reviewing research proposals.

Research Registry

People with MSA face challenges learning about research. The International Research Information Registry is a way for MSA community members to provide their contact information and other details, if they wish to be contacted about participating in experimental drug trials and research studies.

Defeat MSA Alliance (USA) and its MSA United Consortium partners continue to strongly advocate for more drug trials and a more equitable implementation of them worldwide. This informational registry is hosted by Defeat MSA Alliance (USA) but available to anyone diagnosed with MSA (or any related disease) globally. It is completely voluntary and strictly confidential.

defeatmsa.org/research-registry/

Interested in new drugs?
Visit our pipeline page!

defeatmsa.org/msa-research/pipelines/

March is MSA Awareness Month

February 28th is Rare Disease Day

May is our Fundraiser Month