International Support & 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.

MSA United Research Consortium and its partners continue to strongly advocate for more drug trials and a more equitable implementation of them worldwide. This information registry is completely voluntary and strictly confidential.

msaunited.org/research-registry/

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 treatments.

Please help those suffering with MSA!

Make Checks Payable:
MSA United Research Consortium - 501c3
29924 Jefferson Avenue
Saint Clair Shores, Michigan 48082 USA

Email: info@msaunited.org

March is MSA Awareness Month
February 28th is Rare Disease Day
May is our Fundraiser Month

Visit Our Members:

Defeat Multiple System Atrophy – Vaincre L'Atrophie Multisystématisée Canada
www.msacanada.ca

Defeat MSA Alliance (USA)
www.defeatmsa.org

Defeat MSA New Zealand
www.defeatmsa.org.nz

Combattiamo l'Atrofia Multisistemica
www.msa-italia.org

MSA Partnership
www.msaireland.org

Landsforeningen Multipel System Atrofi
www.msa-danmark.dk

National MSA Research
nationalmsaresearch.org.au
(Australia)

Atrofia Multisistémica España
www.amsespaña.org

Many Lives,One Hope

MSA UNITED RESEARCH CONSORTIUM

Call Our Toll Free Help Line:

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

www.MSAunited.org

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EIN: 99-1017213

What is MSA?

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

MSA United Research is a global consortium of charitable associations united in their dedication to fight the disease of Multiple System Atrophy. The collective purpose of MSA United aspires to balance: community support, medical education and advocacy, public awareness and scientific research. All members in MSA United Research International are staffed entirely by volunteers.

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*Volunteer experts provide additional help reviewing research proposals.

**RESEARCH. EDUCATION. SUPPORT.
AWARENESS. ADVOCACY.**

Regional Leaders

**Rosemary Arbuthnot
(Northern Ireland)**

**Brent Evans
(New Zealand)**

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***"A seed hidden in the heart of an apple
is an orchard invisible."***
(Kahlil Gibran)