Education and Support Programs

Empower your Patients by Sharing SRNA's Programs

About the Siegel Rare Neuroimmune Association

The Siegel Rare Neuroimmune Association (SRNA) is a not-for-profit international organization dedicated to the support of children, adolescents, and adults with rare neuroimmune disorders including: Acute Disseminated Encephalomyelitis (ADEM), Acute Flaccid Myelitis (AFM), MOG Antibody Disease (MOGAD), Neuromyelitis Optica Spectrum Disorder (NMOSD), Optic Neuritis (ON) and Transverse Myelitis (TM). We support individuals living with these disorders and their families, promote awareness to empower patients, families, clinicians and scientists, build a collaborative and dedicated clinical care network, and help advance scientific understanding and research. Learn more about our programs and how to get involved at **srna.ngo/mp-resource-center**

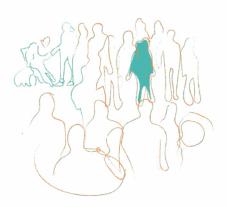




SRNA Through the years connect. care. cure.

Empowerment Through Education and Support

SRNA offers a variety of support and educational programs for individuals with ADEM, AFM, MOGAD, NMOSD, ON, and TM, as well as for caregivers, care partners, and families. Learn more about these programs below.





Support Group Network

srna.ngo/sgn

Patients and caregivers can meet with others and discuss how they are feeling, how they manage their symptoms, ask questions, and offer solutions to problems they have experienced.



srna.ngo/peer-connect

Participants are paired with a trained volunteer who has the same or similar diagnosis and background to offer emotional support, share similar experiences, and recommend resources.



Online Learning Micro-Courses

srna.ngo/courses

SRNA developed the first-ever online micro-course in the field of rare disorders for patients, "Understanding Rare Neuroimmune Disorders," to enhance their understanding of their condition and empower them to make informed decisions about their health and treatment.





srna.ngo/helpline

The Myelitis Helpline is a free online resource that allows those who have recently been diagnosed, or anyone at any point after the onset of a rare neuroimmune disorder, to get answers and support about living with their disorder.







