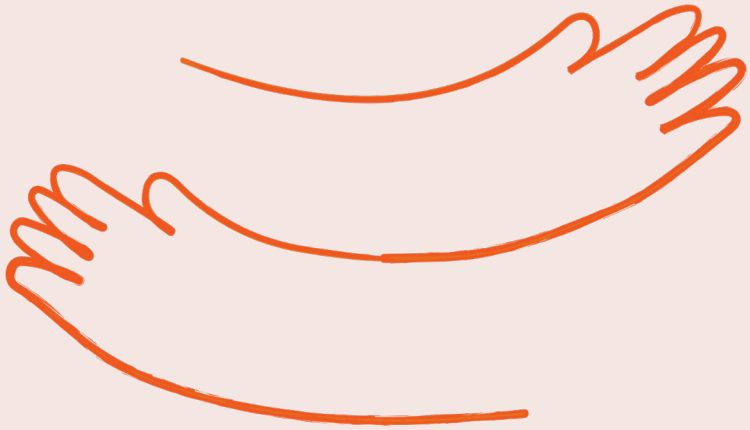
SRNA
connect. care. cure.™

Siegel
Rare Neuroimmune
Association

TOGETHER WE ARE STRONGER

30 years of connection, care, and cure for rare neuroimmune disorders



ADEM | AFM | MOGAD | NMOSD | ON | TM

FROM ISOLATION TO CONNECTION

Rare neuroimmune disorders can strike without warning and change lives forever. For many, these diagnoses bring uncertainty, fear, and isolation. Too often, there are no specialists nearby and no roadmap for care.

That's where SRNA comes in. For 30 years, we've been the bridge from confusion to clarity—connecting families with experts, advancing research, and ensuring that no one faces these disorders alone.

Because while the challenges remain great, so does our commitment: to connect, to care, and to find a cure.

WHY YOUR SUPPORT
MATTERS

WE ADVOCATE for a spectrum of rare neuroimmune disorders that share underlying mechanisms and challenges because what connects these conditions also connects the people living with them, and progress in one drives discovery, treatment, and hope for all.

WE CONNECT families across 121 countries because no one should have to face their diagnosis alone.

WE INVEST deeply in education through online courses, podcasts, webinars, and guides because informed families and clinicians drive better care.

WE TRAIN future clinicians and researchers through the James T. Lubin Fellowship because more specialists in more places mean families can find expert care closer to home.

WE SUPPORT Centers of Excellence dedicated to rare neuroimmune disorders because specialized care and collaboration improve outcomes for everyone.

WE FUND research that uncovers causes, improves diagnoses, and develops better treatments because every discovery moves us closer to answers, better care, and a cure.

IMPACT IN ACTION

CONNECT

15,000+
individuals and families connected across 121 countries

255+
families attended our Quality of Life Family Camps

38+
Walk·Run·N·Roll events hosted across the U.S.

3,000+
individuals connected through support groups worldwide

5,500+
support requests answered through the Myelitis Helpline

CARE

28
symposia hosted, uniting 5,300+ community members and experts

900+
educational resources and online learning tools developed

261
healthcare professionals listed in our global network

785
community members joined the SRNA Registry

CURE

18
Centers of Excellence established

$1,700,000+
allocated to research and training future clinician-scientists

10
fellows trained through the James T. Lubin Fellowship

13
research studies funded, including one that led to an NIH grant of $1.25 million

I TOLD MY WIFE, WITH TEARS IN MY EYES, THAT I FINALLY FELT AT HOME, SURROUNDED BY PEOPLE WHO TRULY UNDERSTAND WHAT IT'S LIKE TO LIVE WITH TRANSVERSE MYELITIS. CONNECTING WITH OTHERS WHO TRULY 'GET IT' WAS BOTH HEALING AND EMPOWERING.

— Don, diagnosed with transverse myelitis

I WOULD BE A RUDDERLESS SHIP WITHOUT SRNA. SRNA PROVIDES ME ACCESS TO DOCTORS, THERAPISTS, RESEARCHERS, SUPPORT GROUPS, CAMPS, SYMPOSIUMS, AND OTHER RESOURCES THAT HAVE GUIDED ME IN MY RECOVERY AND ENHANCED MY QUALITY OF LIFE.

— Ashley, diagnosed with acute disseminated encephalomyelitis in 2007

GOING TO CAMP FOR MANY YEARS HELPED TO BUILD MY CONFIDENCE AND DEMONSTRATE ALL THAT IS POSSIBLE WHILE LIVING WITH A RARE NEUROIMMUNE DISORDER. CAMP GAVE ME MENTORS TO LOOK UP TO, MENTORS THAT I STILL CONNECT WITH TODAY IN MY PERSONAL AND PROFESSIONAL LIFE.

— Abby, initially diagnosed with transverse myelitis in 2006 and later re-diagnosed with acute flaccid myelitis in 2019

WITHOUT SRNA, WE WOULD NOT BE WHERE WE ARE TODAY IN TERMS OF HAVING A DIAGNOSTIC TEST AND A TEAM OF RESEARCHERS COMMITTED TO UNDERSTANDING MOGAD AND ITS IMPACT ON PATIENTS. SRNA IS FOCUSED ON IMPROVING CARE FOR MANY RARE DISEASES THAT ARE OFTEN OVERLOOKED BY THE GENERAL MEDICAL COMMUNITY.

— Becca, diagnosed with MOG antibody disease in 2016

Together, we can reach more families and drive new discoveries. Give today at **srna.ngo/donate** or scan the QR code