

Alie's Fund for Children with MS supports children, teens and young adults with MS between the ages of 2-22, by offsetting the cost of the following:

- Educational Needs
 - Elementary
 - Primary
 - Secondary
- Durable Medical Equipment
- Insurance Co-Pays
- Medical Needs
- Personal and Emotional Needs
- Physician Visits
- Referrals
- Respite Services
- Testing
- Transportation



JOIN THE MOVEMENT to prove you're stronger than it.

Programs, Services and Advocacy

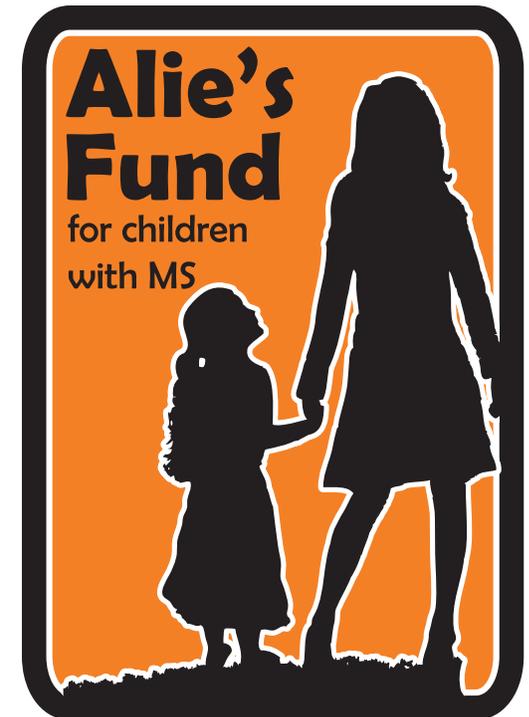
Until a cure is found, the National Multiple Sclerosis Society helps people cope with the disease and maintain independence. To accomplish this, the Society offers educational programs, wellness programs, financial assistance, support groups, counseling and referrals. The Society serves as the premier advocate for individuals with MS and their families nationwide.

There is Hope

Early and ongoing treatment with an FDA-approved therapy can make a difference for people with multiple sclerosis. Learn more about options by talking to your health care professional and contacting the National MS Society at www.nationalmssociety.org.



**National
Multiple Sclerosis
Society**



*"Helping to ensure a
brighter future
for kids with MS."*

Alie's Story

Alie was diagnosed with multiple sclerosis at the age of 14. She is currently living her life to the fullest, thanks to her medical team's care and passion. She is grateful for the medications currently available to slow the progression of her disease.



The National Multiple Sclerosis Society served as a critical source of information and support to Alie and her family. The Society referred doctors with expertise in MS care, worked with her schools to allow extra time for exams and guided her college advisors about the intricacies of her life with MS.

Alie is continuously inspired by the efforts of so many people who raise funds to find solutions, so people with MS can live their best lives. Stories from children affected by financial hardship motivated her to start a fund for children with MS.

An estimated 8,000 - 10,000 children have multiple sclerosis in the United States. Children with MS can experience weakness, fatigue, numbness and tingling, vision problems, loss of balance, difficulty concentrating or remembering, seizures and mental status changes.



"It is not possible to measure the impact this fund will have on the lives of those with MS, but it feels incredibly good to be able to help other children with MS."

Alie's Fund for Children with MS provides funds to support children, teens and young adults ages 2-22 who live with MS. These funds will help defer medical and educational costs, counseling and other needed services to ensure a brighter future for those impacted by MS.

Alie's Fund for Children with MS Form for Support

If you are interested in applying for assistance for Alie's Fund for Children with MS, please complete the form and return to:

**3201 West Commercial Blvd., Ste. 127
Fort Lauderdale, FL 33309**

For more information call 954-731-4224 or visit www.nmssflls.org.

Name _____

____ / ____ / ____
Date of Birth

Age of Diagnosis* _____

Address _____

City, State, Zip Code

(____) _____

Telephone _____

Parent/Guardian Name

(____) _____

Telephone (if different) _____

Preferred way to receive application

Email _____

Fax _____

Mail _____

*Documentation of your MS diagnosis and verification of age is required.