



Who is United MSD Foundation?

Founded in 2016, United MSD Foundation is a registered 501(c)(3) nonprofit serving an international community of Multiple Sulfatase Deficiency families, researchers, and care providers. We exist to bring awareness to MSD, fund research toward treatment, and support families through education, resources, and community.

Bringing Everyone Together

The United MSD Foundation has created a community of MSD families, researchers, healthcare providers, and experts to advance our mission to cure MSD. We work diligently to ensure that these groups are connected and working together effectively.

Family Resources

Everyone's journey with MSD is unique. Rare diseases often leave individuals and families feeling isolated, but please know you are not alone. There is an MSD community waiting to embrace and support you. We support our MSD families in a variety of ways and continue to expand our offerings

Our programs include:

- A Comprehensive Patient Resource Guide
- Clinical Care Guidelines
- Monthly Family Support Meetings
- Family Mentor Program
- Access to a Private MSD Family Support Facebook Group
- Opportunities to Participate in Research
- Biennial MSD Scientific and Family Conference
- Interpreters Available

Contact us TODAY to learn more!