## **Alternate Resources for Patients and Caregivers**

At NS Support, we're a committed partner to families coping with a devastating disease. We understand the challenges you're facing and stand ready to provide support and resources throughout your journey.

Here are some other organizations that may be able to assist you—from foundations working to make treatment more affordable to advocacy groups dedicated to finding a cure.

		ONS

Accessia // Health

Accessia Health provides the financial safety net, products, services, and assistance to patients and families living with rare and chronic

diseases and disabilities

Phone: 800-366-7741 Website: www.accessiahealth.org Contact: assistance@accessiahealth.org

The Assistance Fund

Helps patients and families facing high medical out-of-pocket costs by providing financial assistance for their co-pays, co-insurance,

deductibles, and other health-related expenses through a dedicated fund for patients with Duchenne

Phone: 855-730-5877 Website: https://tafcares.org Contact: https://tafcares.org/about-us/contact/

**DUCHENNE** 

Provides assistance to families for expenses related to the care of a child with Duchenne, and offers help and support to families who need access to recommended treatments, equipment, clinical care, and social services

Phone: 833-428-3301 Website: www.duchennefap.org Contact: info@duchennefap.org



Provides resources to help make treatments affordable for patients with specific, life-altering conditions

Phone: 877-968-7233 Website: www.mygooddays.org Contact: info@mygooddays.org

**HEALTHWELL** FOUNDATION®

HealthWell offers a financial lifeline to underinsured Americans, who require critical medical treatments which they cannot fully afford,

by assisting with their cost-sharing obligations

Phone: 800-675-8416 Website: www.HealthWellFoundation.org Contact: grants@healthwellfoundation.org



Provides assistance to patients and families struggling to obtain life-saving or life-sustaining treatment and care including medication, financial assistance with insurance premiums and co-pays, diagnostic testing, caregiver respite grants, and travel assistance to

treatment sites and clinical trials; offers a dedicated fund for patients with Duchenne

Website: https://rarediseases.org Phone: 800-999-6673 Contact: https://rarediseases.org/contact-us Provides direct financial assistance with co-payments, co-insurance, and deductibles required by the patient's insurer for medications

prescribed to treat and manage their disease

Phone: 866-512-3861 Website: https://copays.org/ Contact: https://copays.org/contact-us/

**PAN** Foundation

**CO-PAY RELIEF** 

Helps underinsured patients with life-threatening, chronic, and rare diseases get the medications and treatments they need by

assisting with out-of-pocket costs

Phone: 866-316-7263 Website: www.panfoundation.org Contact: www.panfoundation.org/contact/

## PATIENT ADVOCACY GROUPS



Akari Foundation educates families on rare diseases and DMD, advocates for the rights and needs of the Hispanic immigrant community within the United States, helps with resources, translations, and interpretations, and are that connection and bond between families and other organizations Website: https://theakarifoundation.org/ Contact: Info@Theakarifoundation.org

Phone: 210-630-5451



Dedicated to finding and funding a cure for Duchenne by breaking the traditional charitable mold through an innovative venture philanthropy model that funds groundbreaking research, early diagnosis, and community education

Phone: 949-872-2552 Website: www.cureduchenne.org

Contact: info@cureduchenne.org

Contact: info@littleherculesfoundation.org



Partners with individuals and families through empowering educational programming, transformational summer camp experiences, financial support for emergencies and accessibility equipment, and by accelerating development of life-changing treatments Phone: 781-585-5566 Website: www.jettfoundation.org Contact: info@jettfoundation.org

Focused on improving the lives of those diagnosed with Duchenne muscular dystrophy through four main pillars: Advocacy, Awareness, Family Assistance, and Funding Research

Phone: 833-428-3301 Website: https://littleherculesfoundation.org

Committed to transforming the lives of people affected by muscular dystrophy

Phone: 800-572-1717 Website: www.mda.org Contact: resourcecenter@mdausa.org

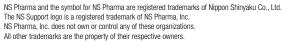
Project Iuscular Dystrophy

🛂 Team Joseph

Accelerate research, impact policy, demand optimal care for every family, and strive to ensure access to approved therapies Phone: 800-714-5437 Website: www.parentprojectmd.org Contact: info@parentprojectmd.org

Fund cutting-edge research to find a treatment or cure for Duchenne muscular dystrophy, and improve the lives of patients and families through advocacy and Family Assistance

> Website: www.teamjoseph.org Contact: info@teamjoseph.org



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