



SPONSORSHIP OPPORTUNITIES



2025



Since 1958, the Muscular Dystrophy Family Foundation has provided financial assistance, quality programs, and other services for people across the state of Indiana facing the challenge of living with a neuromuscular disease. Our goal is to increase the quality of life for members of our community through advocacy, education, and by providing life-changing resources that support greater mobility and independence.

Muscular dystrophy causes a person's muscles that control movement to degenerate. Genetic mutations interfere with the production of muscle proteins necessary to build and maintain healthy muscles. This rare, complicated disease impacts around 50,000 Americans and 1,500 Hoosiers. It can appear as early as infancy or as late as middle-age—or even later. Each type differs in the muscles affected, age of onset, and rate of progression.

There is no cure for muscular dystrophy, but medications and therapies can help people manage their symptoms and slow the course of the disease. Many people with muscular dystrophy lose the ability to walk and end up homebound. Some may have trouble breathing or swallowing. Others are born healthy and are athletic in their youth then diagnosed later in life. They go from leading active lives to being fully-dependent on assistive devices and caregivers, making the emotional challenges just as difficult as the physical changes to which they have to adapt.

When a family reaches out to the MDFF for help, we're often their last hope. Insurance companies, Medicaid, Medicare, and other agencies continue to make drastic cuts to funding. Those with muscular dystrophy need adaptive equipment and services needed just to **experience quality of life, more independence, and opportunities to build meaningful connections**. MDFF provides essential items like wheelchairs, hospital beds, van lifts, ramps for homes, bathroom modifications, and shower chairs.

We have done a lot of good, but we're only getting started. Thanks to generous support from donors, grant writers, partners, medical professionals, staff, and volunteers, our MDFF family continues to help individuals with neuromuscular diseases get what they need to experience a higher quality of life, live more independently, and build meaningful connections.

We would love to partner with you to create a better future for individuals with muscular dystrophy and other neuromuscular diseases.

Thank you,
The MDFF Board of Directors



MDFF Mobility Gala

Date: September 19, 2025
Location: Biltwell Event Center

A change to MDFF's annual September event is coming Fall of 2025. This fundraiser helps provide MDFF with the funds needed to continue our vital programs, including the Adaptive Services Program, Partial Van Assistance Program, Family Events and Accessible Van Giveaway.

Sponsor benefits are subject to change as event details are finalized.

MDFF Gala Fundraiser	All-In \$10,000	Royal \$5,000	Ace \$2,500
Table(s) of 8 attendees	2	1	1
Company Logo on printed program, signage, and mentions during dinner presentation	x	x	x
Company Logo on event webpage & bidding platform	x	x	x
Company recognition in e-newsletter and social media	x	x	
Company recognition at drink stations	x	x	
(3) free beads per person for heads or tails game	x		

**Printed material, e-news, and social media benefits are subject to deadlines*