

Wings Over Wall Street

Imagine a world without ALS. This vision drives the lifesaving mission of the Muscular Dystrophy Association's ("MDA") Wings Over Wall Street event.

For more than 21 years, Wings Over Wall Street has raised over \$12.7 million in support of ALS research. The importance of our work – and your support – will only grow as we continue to be a catalyst in the research community.

Your generosity, through a sponsorship, journal advertisement, or donation enables MDA to fund groundbreaking research breakthroughs and unprecedented progress to advance more clinical trials and develop new standards of ALS treatment and care. Join MDA and over 300 leaders from the finance, insurance, airline, technology, pharmaceutical, and healthcare industries on Thursday, June 9, 2022, at Guastavinos, NYC.

MDA & Amyotrophic Lateral Sclerosis (ALS | Lou Gehrig's Disease): At MDA, we are the leading authority on, and resource for, neuromuscular disease. We are at the center of a broad network of researchers, scientists, clinicians, and caregivers who are focused on cracking the code around ALS.

Since 1950, MDA has invested more than \$170 million in ALS research and care. For over 70 years we have been committed to ALS and the ALS community, serving at 48 designated MDA ALS Care Centers across the country. Today, we have 45 active ALS grants, and the advent of the first-of-its-kind MOVR data technology hub. MDA is helping enable a pipeline of young scientists with the financial and technological resources they need to provide promising research that aims to deliver scientific advancements.

You are the driving force that enables MDA's Wings Over Wall Street to impact families living with ALS. At the heart of the Wings Over Wall Street community are people like you, who make it possible for MDA to lead the way toward a better future through funding research.

Your sponsorship/donation will make an outsized impact! Together, we can Unlock the Cure. When you support Wings Over Wall Street you are empowering people and families living with ALS.

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