TOP 10 ACP BREAKTHROUGHS in People-Powered MS Research



A DECADE OF PROGRESS

Celebrating 10 years of iConquerMS, with 8,500 members reshaping MS research and care. Together, we are ensuring that research addresses the real needs of those living with MS.

RESEARCH POWERED BY YOU

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Ensuring research reflects real MS challenges by engaging actual people, iConquerMS members. This year, our members contributed their vital insights to over a dozen research engagements.

The ACP Repository has been instrumental in advancing our research on MS...we discovered aggregated antibodies that are toxic to neurons – a potential hallmark of MS disease activity. ACP has truly accelerated our progress in understanding MS mechanisms and developing potential therapeutic strategies.

Xiaoli Yu, PhD, Associate Professor Department of Neurosurgery, University of Colorado Anschutz Medical Campus

UNLOCKING MS MYSTERIES



Supporting researchers across the globe with essential biospecimens and data through the ACP Repository. Our new partnership with DNAnexus allows experts to analyze our unparalleled dataset, potentially revealing crucial insights that could unlock new avenues for treatment and lead to cures.

FIGHTING FATIGUE



Launching the largest-ever study on digital treatments for fatigue, thanks to a \$4.5M federal grant. We will bring hope and practical solutions to individuals suffering from MS fatigue, a symptom affecting 70% of those with MS.



REMOTE RESEARCH REVOLUTION

Enabling participants to contribute blood samples and data remotely through our COVER-MS study, **accelerating research timelines and broadening research participation** beyond traditional MS clinic locations.



Donate now! Please visit our

website to learn more about our work and to donate.



AWARD-WINNING ADVOCACY

Earning the 2024 Made with Patients award for advancing patient-centered research through the RIDE Council – showcasing our commitment to inclusive research opportunities.

DIVERSE VOICES, BETTER RESEARCH



Amplifying voices from under-represented communities through our RIDE Council to drive inclusive MS research. This crucial research enriches our understanding of how MS affects diverse populations to produce more effective solutions.

I wanted to see a change. And the only way that I could do that was by sharing my story, aiding in the fight to empower other Latinos with MS. The RIDE Council allowed us to de-stigmatize myths, address fears, encourage, support, and empower. I do not know of any other organization that is putting us, those living with MS, first.

Shawn F., person living with MS

CHAMPIONING YOUNG VOICES

Partnering with kids, teens, and young adults with pediatric MS to create the first-ever patient-driven research agenda for young people with MS. Shaping future research and developing effective treatments and support systems for their unique needs.



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CARING FOR CAREGIVERS

Recognizing the often-overlooked heroes, MS caregivers. We've launched the MS Care Partner Connection. A onestop resource hub, designed by caregivers for caregivers.



EMPOWERING WITH KNOWLEDGE

Translating research into clear, actionable insights for MS patients and doctors to make informed care decisions together, armed with the latest evidence-based information.



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