Advancing Research & Treatment Innovation

Funding LMS-specific studies and global precision-medicine initiatives that accelerate clinical trial development and expand treatment options. Our Research in Review series keeps patients and families informed about emerging research and clinical trials.

International Leiomyosarcoma Research Roundtable

NLMSF - SPAGN International LMS Research Roundtable. Annual meetings since 2018, with over 130 research members. 5 Working groups within MultiOmics, Clinical Trials, PDX Models/ Cell Lines Strategic Imaging, Gynecologic Oncology Published Concensus Papers for Treatment and Management of this disease.



NATIONAL LEIOMYOSARCOMA FOUNDATION

IMPACT IN ACTION

NLMSF.ORG

Patient, Caregiver & Family Community Roundtable

A collaborative think tank network focused on identifying and addressing the common challenges faced by patients, caregivers, and families. *Caregiving Counts* is a bi-monthly series for Caregivers to share information, tips, and support.

Clinical Trials Patient-Caregiver Perspectives

A Community Network for patients sand caregivers sharing information and identifying common challenges (unmet needs) in the clinical trial setting. Survivorship Care Planning Clinic offers what patients and families should know to maintain/enhance quality of life during and after treatment.

Community Empowerment Initiatives

Education, Support & Connection. Awareness programs, education forums, resource support, research information and opportunities, live support network. Bereavement Circle offers small group discussions for families who have lost a loved one to LMS/Sarcoma.

Annual Early Career Awards

Recognizing and encouraging the next generation of physician-scientists dedicated to LMS research. *Early Career Award* and the *Brian Van Tine, MD Faculty Award* for Excellence in LMS Research.

Questions? Contact Annie Achee, President, NLMSF at +1 303 808 3437 / annieachee@aol.com / www.NLMSF.org