NATIONAL LEIOMYOSARCOMA FOUNDATION

IMPACT IN ACTION

NLMSF.ORG

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 Funding of LMS Research Grants

Annual funding of LMS research projects Grant proposals collected, vetted, evaluated, selected - March - September. Gateway Website created for the Research community: lmsresearch.net

International Leiomyosarcoma Research Roundtable

LMSF - 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.

Advancing Research & Treatment Innovation

Accelerating and advancing global research initiatives focused on precision clinical trial development to expand treatment options for patients with leiomyosarcoma (LMS). *LMS Research in Review Series* - research experts share updates on the latest research and treatment updates with patients and families.

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 quarterly network meeting of new patients, patients current in trials, and those that have completed a trial to discuss experiences and common challenges. Survivorship Care Planning Clinic offers what patients and families should know to maintain/enhance quality of life during and after treatment.

Annual Early Career Awards

Recognizing and encouraging the next generation of physician-scientists dedicated to LMS research.

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