LeiomyoSarcoma
SUPPORT & DIRECT RESEARCH FOUNDATION
~~WISHING~~ WORKING FOR THE CURE

www.lmsdr.org

info@LMSDR.org

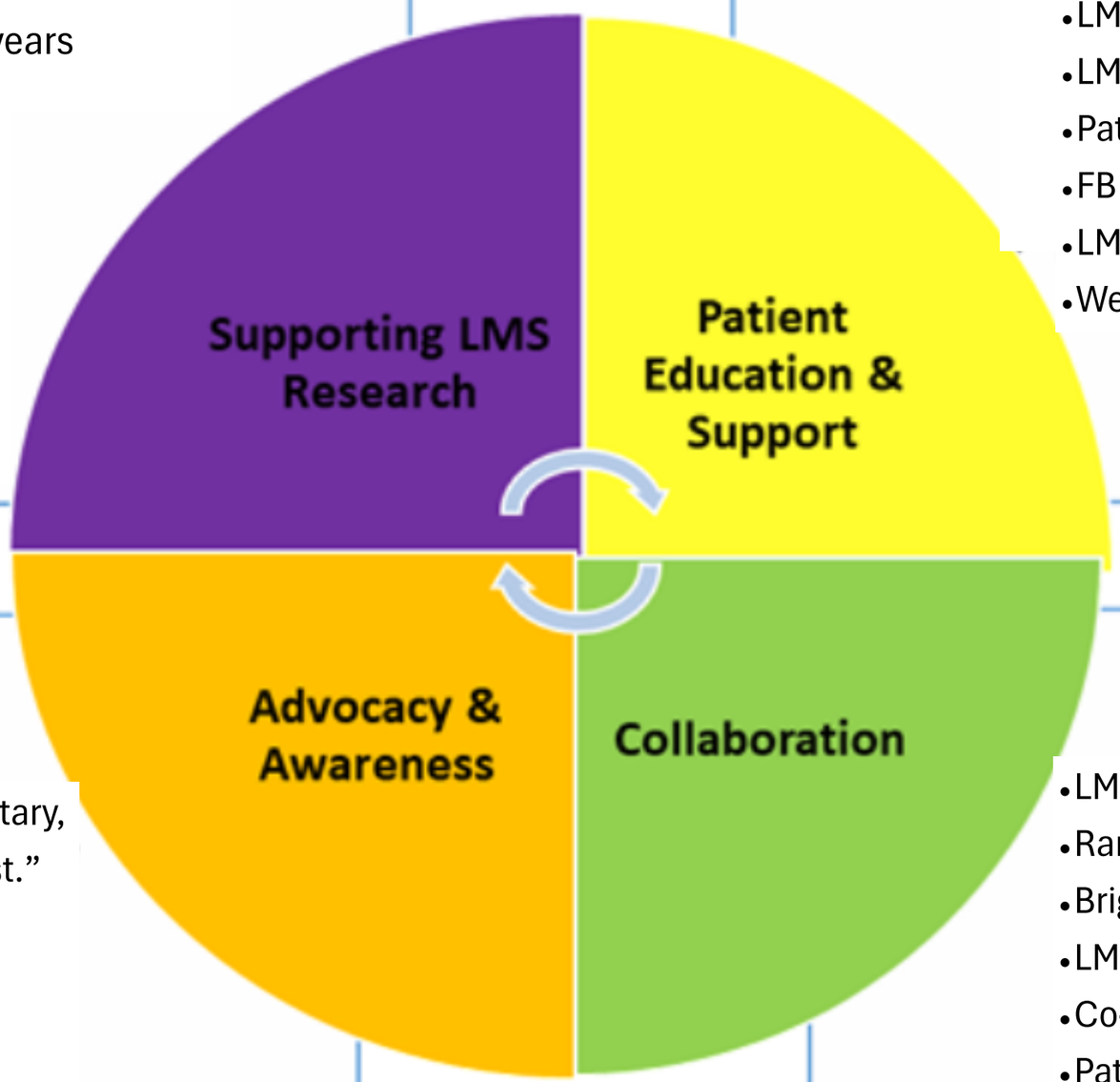
- Originating in 2006, the Leiomyosarcoma Support & Direct Research (LMSDR) foundation is an all-volunteer staff of patients and caregivers, who are passionate about supporting others and helping researchers to find treatments that will work.

What We Do

- Awarded over $5M for LMS Research in the last 20 years

LMSDR Created:

- LMS Patient Registry
- LMS Tissue Bank
- LMS Cell Lines



- LMSDR Website
- LMSeAlerts Newsletter
- LMS Boot Camp videos
- Patient Conferences
- FB Support Group
- LMS Community Blog
- Webinars

- Co-producer of documentary, "Kicking the Hornets' Nest."
- Sarcoma Conferences
- Testified at FDA hearing against morcellation
- LMS Awareness Day

- LMS Project/Broad Institute
- Rare Cancer Foundation
- Briger Foundation
- LMS Research UK
- Co-funding research
- Patient led fundraisers
- Sponsorships

Who Are We?

Board of Trustees



Sharon Anderson (President)



Janine Di Tullio (Exec Director)



Hao Lam (Treasurer)



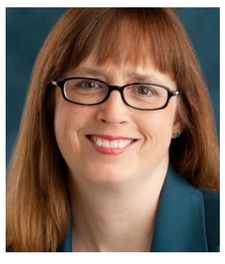
Dondee Murray (Secretary)



Cherry Kim

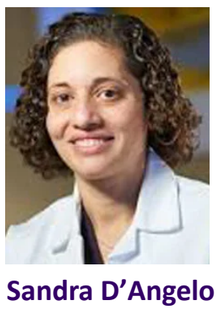


Michael Rhine



Dr Hope K. Haefner

Science Advisory Board



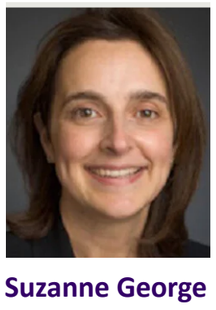
Sandra D'Angelo (Memorial Sloan Kettering)



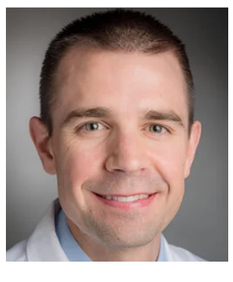
Gina D'Amato (Sylvester, Miami)



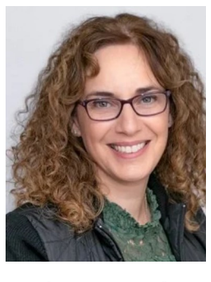
Jerad Gardner (Geisinger, Pensylvania)



Suzanne George (Dana Faber/ Harvard)



Matt Hemming (Dana Faber)



Isis Kanevsky (Pfizer)



Sara Watson (UCL London)



Alex Lazar (MD Anderson)



Matt van de Rijn (Stanford)