

How Can You Get Involved?

Join Our Community

Connect with us through our support groups, online forums, and social media channels.

Raise Awareness

Share information about LMS and LMSR UK with your network, helping to increase understanding and support.

Fundraise or Donate

Donate to our JustGiving page or organise fundraising events to support us.

<https://www.justgiving.com/crowdfunding/LMSRUK>

Volunteer

Offer your time and skills to help us achieve our mission.

Together, we will find a cure.

Thank you!



Meet the LMSR UK Team



Colin Grandson
Treasurer



Maisie England
Chair



Julia Casimo
Secretary



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LEIOMYOSARCOMA RESEARCH UK

Together, we will find a cure.



LMSR UK

WHO WE ARE
AND WHAT WE
DO

Registered Charity 1210455

What is Leiomyosarcoma?

Leiomyosarcoma (LMS) is a rare and aggressive form of cancer that affects the smooth muscles. It presents unique challenges due to its rarity and aggressive nature, often leading to a poor prognosis. Research into LMS is currently underfunded and fragmented, resulting in limited treatment options and a lack of access to specialist care for many patients.

Who are LMSR UK?

We are a newly established organisation founded by LMS patients and their loved ones. Driven by the urgent need for better treatments and outcomes, we are dedicated to making a real difference in the lives of those affected by LMS. Our mission is to support research, provide education, and foster a strong community for LMS patients and their families across the UK.



Our Aims

Establish a UK Network of LMS Experts

We aim to create a network of researchers and experts to coordinate and focus research efforts, encourage collaboration, and share best practices, ultimately leading to improved outcomes for LMS patients.

Educate and Inform

We provide up-to-date, accessible information about LMS through leaflets, our website, social media, and patient information sessions.

Provide Support

We offer a safe and supportive community for those affected by LMS through local and online support groups and forums.

Raise Awareness

We actively work to increase awareness of LMS within the community and among medical professionals, advocating for better diagnosis and treatment options.

Sponsor Research

We are dedicated to funding research into the prevention, diagnosis, and treatment of LMS, with the goal of improving patient outcomes and developing the next generation of medical experts and researchers.



What Does This Mean for Patients?

Reliable Information

Access to clear and up-to-date information about LMS and the latest research developments.

Community and Connection

Opportunities to connect with other LMS patients and families through support groups and online forums, fostering a sense of belonging and shared experience.

Knowledge

Resources to help you advocate for yourself and make informed decisions about your treatment and care.

Hope for the Future

By funding research and fostering collaboration, we are actively working towards better treatments and improved outcomes for LMS patients.