

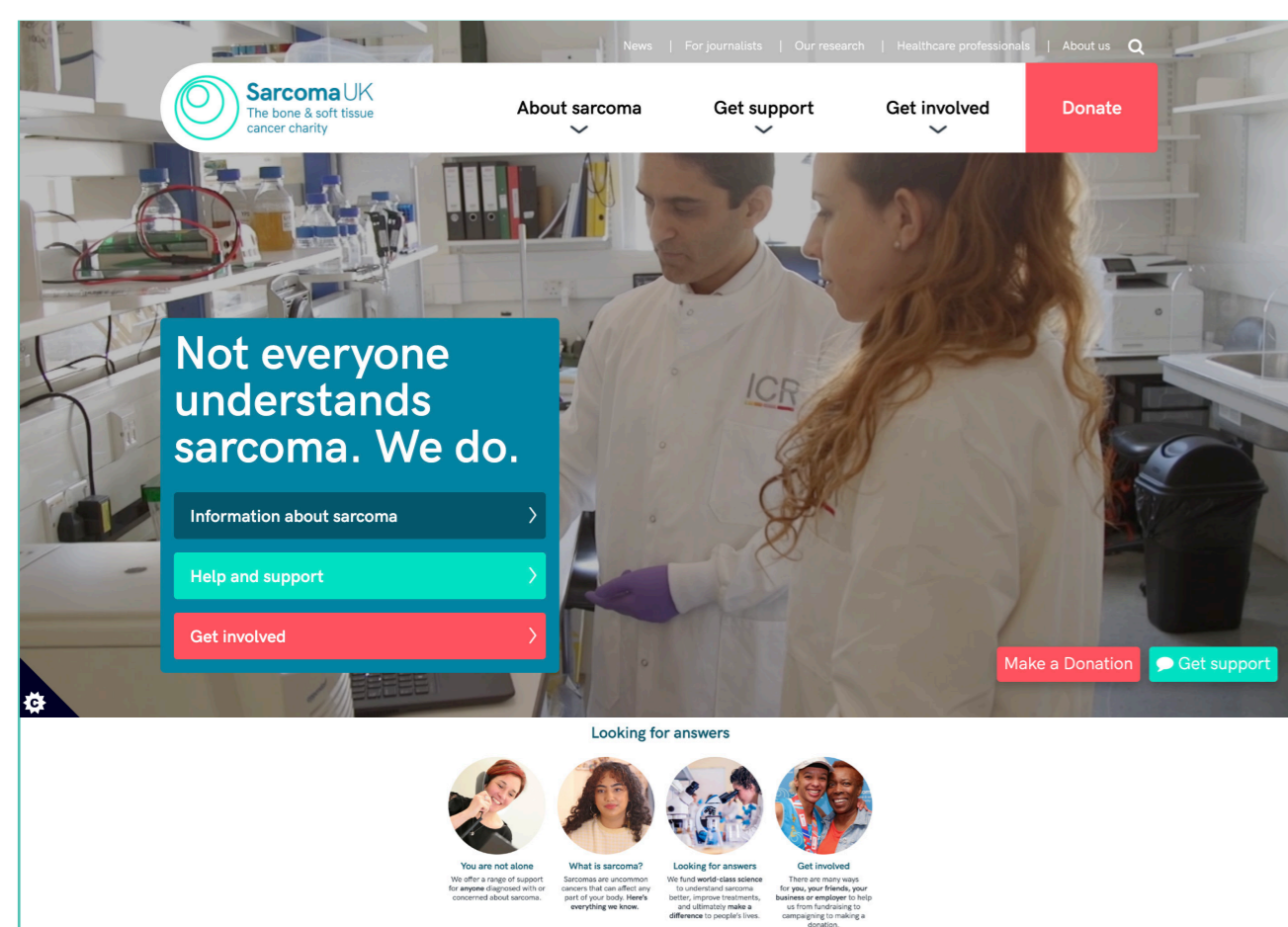
Sarcoma UK

Our mission is to ensure everyone affected by sarcoma receives the best treatment, care, information and support available and to create the treatments of the future

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Who we are

- Sarcoma UK was founded by Roger Wilson CBE, a sarcoma patient.
- We are the only cancer charity in the UK focusing on all types of sarcoma.



What we do

Guided by our strategy *Tackling Sarcoma Together*, we:

- Find answers through funding sarcoma **research**.
- Drive **awareness** of sarcoma cancer.
- Provide **information** and **support** to anyone affected by sarcoma.
- **Campaign** for better treatments and to improve standards of care.
- **Fundraise** to help us continue our work



Patient Involvement

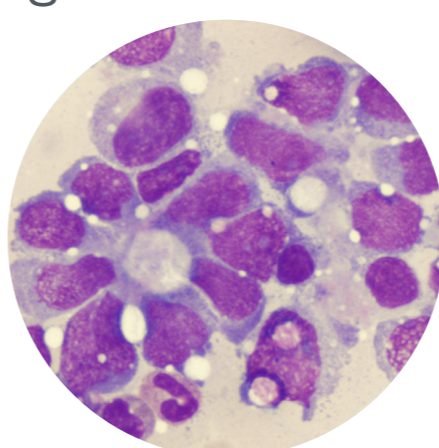
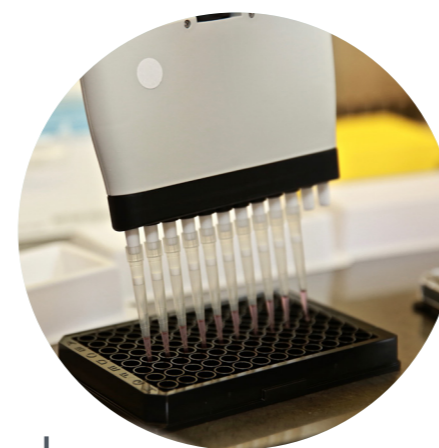


- We want to make sure the voice of people affected by sarcoma is heard throughout the organisation and within the Sarcoma community.
- Our Patient Involvement Network is made up of people affected by sarcoma who want to shape our work
- The Network now has over **120 members**
- Opportunities offered include:
 - ▶ Reviewing research proposals
 - ▶ Joining recruitment interview panels
 - ▶ Reviewing information resources
 - ▶ Joining focus groups

Research



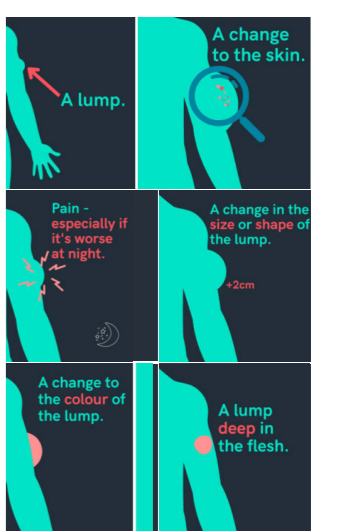
- Since 2009, we've invested over **£5 million** into research across the UK
- We fund research into:
 - ▶ Cause
 - ▶ Diagnosis
 - ▶ Treatment
 - ▶ Quality of life
- We offer small, large grants and PhD Studentships
- We prioritise collaboration and fund research in partnership with LifeArc, GIST Cancer UK, the Bone Cancer Research Trust and the Grace Kelly Childhood Cancer Trust
- We target our research funding into areas of patient priority including:
 - ▶ Genomics
 - ▶ Diagnosis
 - ▶ Ultra-rare subtypes
- Our research funding has led to over **140 publications and presentations**, and over £5 million of follow on funding for sarcoma researchers



Policy



- Our Policy Priorities:
 - ▶ Earlier and more accurate diagnosis
 - ▶ Access to the best possible care
 - ▶ Access to appropriate psychological support
- Our 2020 report, *Delays Cost Lives* calls on policy makers to improve early diagnosis of sarcoma
- We have launched a new sarcoma education module for GPs and new Signs and Symptoms resources for people concerned they have sarcoma



Support

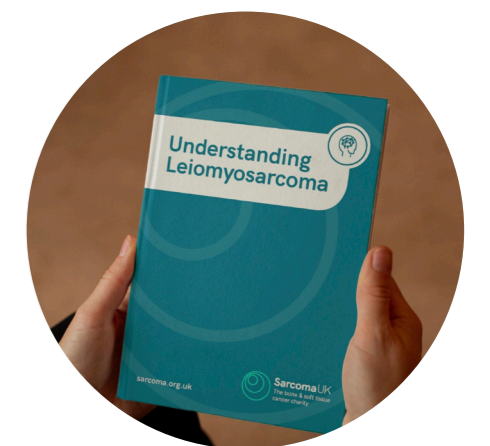


- The Support Line is staffed by healthcare professionals with expertise in sarcoma
- The team can be contacted by phone, email and text
- We've been in contact with almost 5,000 individuals

Top 3 reasons to contact the Support Line:

1. Emotional support
2. To discuss treatments
3. Questions pre-diagnosis

Information



- We publish digital and print information for people affected by sarcoma and healthcare professionals
- In the past 12 months we've sent out **18,256 print information resources**
- We've published a soft tissue sarcoma data hub to help people to see where their subtype fits in
- We have a clinical trials hub to allow people to search for and find trials for their sarcoma subtype.

