



The Future of Sarcoidosis Research is in

OUR Hands

"I consider it a privilege to be part of the FSR Patient Registry. As a sarcoidosis patient, there is so much that is out of my control. Being part of the registry lets me feel that I am contributing towards a greater good for all of us with this nasty disease."
– Jim K., FSR Advocate

"I joined the FSR Patient Registry because finding better treatment for sarcoidosis begins with research and that research begins with patient involvement. I believe all sarcoidosis patients should take the time to join the registry to help accelerate research."
– Jessica R., FSR Advocate

LEAVE YOUR MARK ON SARC!

Every patient has a unique experience with sarcoidosis. By sharing your personal experiences with this disease in the FSR Patient Registry, you have the opportunity to help accelerate research quickly and conveniently!

REASONS YOU SHOULD JOIN THE FSR PATIENT REGISTRY

- Assist in the better understanding of the variation and progression of sarcoidosis.
- Contribute to clinical research and new drug development.
- Provide the medical community information to help develop clinical guidelines.
- Provide reports on patient outcomes to help improve patient care.
- Get connected to IRB-approved clinical research in your area.
- Receive notifications when publications using the registry data are released.

YOU ARE ELIGIBLE TO JOIN IF:

- You have been diagnosed with sarcoidosis.
- You are a caretaker for someone with sarcoidosis.

**Patients under 18 must register with a parent or legal guardian*

JOIN THE REGISTRY IN TWO SIMPLE STEPS!

1

Go to this link or scan the QR code:
<https://fsr-sarc.patientcrossroads.org/sign-up.html>

2

Complete the 45–60-minute survey



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