

Summit 2022 FAQ

Q: What if I can't make both days, should I still register?

A: We totally understand that not everyone will be able to attend all of both days, or even at all during the weekend! Which is exactly why the entire summit platform, sessions, connections with other attendees, and exhibit booths chalked full of valuable information, will be available until September 30th. This gives everyone plenty of time to see all the sessions they wanted to attend, or ones they missed. We'd love to see you at the summit!

Q: What sarcoidosis manifestations will be covered?

A: The FSR Global Virtual Sarcoidosis Patient Summit: Unveiling Possibilities will have sessions on various manifestations.

For example, if you are interested in Neurosarcoidosis, we have a Neurosarcoidosis 101 session that is meant for folks new to the disease, and a separate session on Chronic Neurosarcoidosis for folks who have been managing this disease for a while. In addition to this, we will have sessions with Patient Advocates, many of whom suffer from neurosarcoidosis or systemic sarcoidosis. We will also have a chat room dedicated to neurosarcoidosis. Finally, the platform connects patients to one another. People who register for a neurosarc session will be suggested as contacts for others who registered for the same session.

The summit has sessions on pulmonary sarcoidosis, skin (cutaneous sarcoidosis), ocular sarcoidosis, neurosarcoidosis, cardiac sarcoidosis, and gastric sarcoidosis.

You can review the entire agenda for both days on the Summit webpage to learn more.

Q: I am interested in hearing more about sarcoidosis research, is this the conference for me?

A: Yes, this the right conference for you. We have a number of sessions specifically dedicated to advancements in sarcoidosis research and research updates will be discussed by global sarcoidosis experts throughout.

You can review the entire agenda for both days on the **Summit webpage** to learn more.

Q: I am a caregiver of someone living with sarcoidosis. Am I welcome to attend?

A: We encourage caregivers to join us for this summit. There is lots of helpful information for you and your loved one impacted by sarcoidosis. In addition, this year's summit has two session dedicated to caregivers and an exclusive chat board for caregivers.

You can review the entire agenda for both days on the Summit webpage to learn more.

Q: I am interested in learning about how others manage their disease and cope with pain and fatigue. Do you have any sessions for this?

A: This summit is full of sessions on living with sarcoidosis and managing symptoms. Join sessions on living with fatigue, managing pain and how diet and nutrition may help to improve symptoms. Join us for a meaningful discussion on "life hacks" those living with sarcoidosis use to improve their every day lives.

You can review the entire agenda for both days on the <u>Summit webpage</u> to learn more.

Q: Is there any way for me to learn about clinicians that are treating those living with sarcoidosis?

A: This year, FSR is excited to be able to feature 14 clinic partners who are members of the FSR Global Sarcoidosis Clinic Alliance through their booths in the exhibit hall. Visit their booths to learn more about the clinics and to view resources to help you live with sarcoidosis.

Q: After I register, will I receive and email with information about when I am able to access the platform?

A. Once you complete registration you will receive a confirmation email. If you do not receive a confirmation email within 15 minutes of completing your registration, please reach out to Mindy Buchanan, Director of Patient Programs, mindy@stopsarcoidosis.org Your confirmation will not have the link to join the summit, the email with detailed instructions with how to join the Summit will come separately on July 27th from moreply@swapcard.com. It is a good idea to add this email address to your known senders to ensure you get the information you need to join. If you do not receive this email by the end of the day on July 27th, please reach out to Mindy.