

TAKE PART IN CDRC'S SARCOMA PATIENT PARTNERED RESEARCH



Share your experience living with Sarcoma

Sarcoma subtypes will be evaluated for a repurposing drugs initiative to advance treatment potential and options. We need you to help find better treatments and to learn from your experiences. We want to find out if there are any drugs, supplements, or natural products not prescribed for your sarcoma that make your life better or that might cause harm. We also want to help crowdsource treatment options that can be shared publicly with the community. If you want to know more about this project or have any questions, please send an email to cdrc@c-path.org.

WHY WE NEED YOU!

Our goal is to better understand the current therapies being used for rare and ultra-rare sarcomas. Your medical records and patient information is the best source to find out the drugs you have taken and the treatments you have tried. By collecting and sharing all this information, we will help patients make better decisions about treatment and identify more treatment options. In the future, we hope to study even more types of sarcomas.

xcures.com/cdrc



HOW DO I PARTICIPATE?

- Medical Records are helpful, but don't tell the whole story. We need your help! Your story can inform more treatment options for sarcoma patients.
- Share your treatments that you have received
- Sign up for xCures to help consolidate your medical records with your permission. Review and amend your records to share your story.
 - xCures will gather and provide access to all of your medical records, produce a visual summary of your care history and, when appropriate, provide information regarding clinical trials.

FOR MORE INFORMATION

- Check out the CDRC FAQ page
- Visit xcures.com/cdrc
- Email cdrc@c-path.org for more information

Goals:



Collecting
deidentified data on
lived experiences



Collaborating more
effectively to ask the
right questions



Sharing insights gained
to help identify
treatments and to make
better treatment
decisions

**PLEASE SHARE THIS INFORMATION WITH OTHER FAMILIES LIVING WITH
ANGIOSARCOMA, EHE, OR PECOMA.
THE MORE PEOPLE PARTICIPATE, THE MORE WE WILL ALL LEARN!**

PARTNERS & COLLABORATORS

