



SYNOVIAL SARCOMA REGISTRY AND BIOSPECIMEN REPOSITORY

WHAT IS THE PURPOSE OF THIS RESEARCH STUDY?

To establish a registry and biorepository for synovial sarcoma patients by collecting and storing data and biospecimens, such as blood, saliva and/or buccal (cheek) swabs, and tumor tissue. To use this collected data to advance research and improve outcomes for patients in the future.

WHO MAY PARTICIPATE?

Anyone who has a diagnosis of synovial sarcoma. There is no age restriction. This study is open to all ages. Families of deceased patients are also welcome to contribute.

WHAT WILL BE ASKED OF YOU?

As a participant in the research, you will:

- Provide data on your cancer treatment and agree to let the study team review your health records.
- Provide saliva and/or buccal (cheek) swab samples.
- Provide blood samples at the time of a clinical blood draw.
- Agree to have your leftover tumor tissue stored in a repository at CHOP.
- Have genetic testing done on your provided biospecimens.

Participation in the research study is voluntary and will not affect any services that you or your child may be receiving at CHOP or your local health care institution. You are free to withdraw from the program at any time.

I WANT TO HELP! WHO DO I CALL?

If you are interested in participating in the study or want to learn more about it please contact:

Study Team
Dr. Theodore Laetsch, MD
Lauren Gutstein, Project Manager
(267) 827-8145
SynovialSarcomaRegistry@chop.edu

Complete interest form at
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