NIH NATIONAL CANCER INSTITUTE

CHILDHOOD CANCER
DATA INITIATIVE

PEDIATRIC, ADOLESCENT, & YOUNG ADULT

RARE CANCER STUDY

HELP ADVANCE RARE CHILDHOOD CANCER RESEARCH

The CCDI Pediatric, Adolescent, and Young Adult Rare Cancer Study is a longitudinal, observational study that will collect different types of health and genomic data from people with very rare childhood cancers.

Data collected will also go into a database that all researchers can use to better understand how or why very rare childhood cancers develop and how they behave over time, so that they can find effective ways to treat people with these cancers. Personal information will be removed from the data.

WHO CAN JOIN?

ANYONE INTERESTED MUST:

- Be a child, teen, or young adult (age 39 or younger)
- Have been diagnosed with a rare solid tumor
- Have been diagnosed less than one year ago
- Be willing to provide tumor and saliva or cheek swab samples for testing

There will be no experimental treatments or in-person appointments—just surveys, and data and sample collection. Compensation for participating will be provided.

Specific cancer treatment is not part of this study. Any treatment you or your child are receiving will continue as prescribed by your provider at the institution where you are being treated.

INTERESTED IN JOINING?

Contact us:
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Visit for more information:
cancer.gov/ccdi-rare-cancer

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