

Family members of children with cancer may also be at risk

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When a child is diagnosed with cancer, one of the first questions the parents ask is "Will my other children get cancer?" A new study from Huntsman Cancer Institute (HCI) at the University of Utah suggests the answer to that question depends on whether a family history of cancer exists. The research results were published online in the *International Journal of Cancer* and will appear in the November 15 print issue.

The study, led by Joshua Schiffman, M.D., medical director of HCI's High Risk Pediatric Cancer Clinic and a pediatric hematologist/oncologist in the Department of Pediatrics at the University of Utah, examined the family [medical history](#) of 4,482 [children](#) diagnosed with [cancer](#) over a 43-year period to determine the [cancer risk](#) in their relatives.

The research team found that when children were diagnosed with any kind of cancer at age 18 or younger, the risk to their parents, siblings, or children for childhood cancer doubled compared to families with no childhood [cancer patients](#). If the [cancer diagnosis](#) came when the child was age 4 or less, the risk to close relatives for childhood cancer increased almost four times.

"No one had previously studied the question, so we simply told parents there was no evidence of increased risk to the other children," said Schiffman. "Now we can give an evidence-based answer—the risk depends on your family history of cancer."

This is the first study that uses the Utah Population Database (UPDB) to broadly examine the risk of all [types of cancer](#) in relatives of children with cancer. This unique resource at the University of Utah links genealogies and cancer registry data from Utah to medical records and vital records, including Utah [death certificates](#).

"Because our data came from the UPDB, the

assessment of family history in our study does not rely on self- or family-reported medical history," said lead author Karen Curtin, Ph.D., a genetic epidemiologist and UPDB assistant director. "Self-reporting of family medical history depends on an individual's memory, while our data comes from the statewide Utah Cancer Registry that records nearly all cancer cases, which reduces possible errors in reporting family cancers."

The team also assessed known inherited genetic syndromes in adult relatives of pediatric cancer patients. They found cancers associated with Li-Fraumeni Syndrome (LFS) seemed to be driving the increased risk to relatives in families with a history of cancer.

"Not all children's cancers are hereditary," said Schiffman. "But the numbers in this study suggest that the proportion of hereditary childhood cancers may be significantly higher than the 5-10% generally cited in adult hereditary cancers, and likely even more than 20%."

"LFS is one of the most devastating cancer syndromes," said Schiffman. "It causes a variety of cancers in both children and adults. For people with LFS, the lifetime risk of getting cancer is 80% to 90%, but with increased and early screening for tumors, there's early indication of a very high survival rate, perhaps even approaching 100%. In a previous study, LFS patients who did not receive early screening only had a 20% survival rate."

Although [childhood cancer](#) rarely occurs in the population, based on their findings, the authors recommended collection of three generations of family medical history for all newly diagnosed pediatric cancer patients and referral of families with a history of early-onset cancers in children or adults for genetic counseling. In addition, parents of children diagnosed with cancer before age five with a family history of cancer should be advised of the potential for increased risk to other children in the

family.

"We want to encourage the taking of a family history as part of routine care. With all cancers, but perhaps especially with childhood cancers, so many other questions seem so urgent, a [family history](#) may not seem to be the most pressing issue," said co-author Wendy Kohlmann, director of HCI's Genetic Counseling Program. "When families are referred into genetic counseling, we can provide them with more information about the risks and actions they can take."

"For families with previously unidentified LFS, following these recommendations could actually save the lives of multiple family members if at risk individuals are identified and early cancer surveillance programs implemented," Schiffman said.

The data analyzed in this study indicated that outcomes for [pediatric cancer](#) patients are worse in families with a history of cancer. Schiffman said that further studies are planned to learn the clinical implications of this observation.

Provided by University of Utah Health Sciences

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