

Study reveals properties of cells fated to relapse in acute lymphoblastic leukemia

March 3 2020



Cancer cell during cell division. Credit: National Institutes of Health

Scientists at St. Jude Children's Research Hospital and Princess Margaret Cancer Centre in Toronto, Canada, have reported that subpopulations of leukemic cells present at diagnosis can cause relapse in children with

acute lymphoblastic leukemia. The findings have implications for current and future therapy. The work recently appeared as an advance online publication in *Cancer Discovery*.

While there have been significant advances in the treatment of ALL, the disease recurs in 15-20% of pediatric and 40-75% of [adult patients](#). Prognosis for patients who relapse is poor. Treatment for ALL eliminates some leukemia [cells](#), while others survive and progress to relapse; but why this happens has remained a key question in [cancer research](#).

There are distinct populations of ALL cells called subclones, whose behavior and properties differ from their fellow leukemia cells in part based on their underlying genetics. Researchers identified and isolated subclones, present at diagnosis, that somehow survive therapy and eventually initiate relapse. The scientists called these cells diagnosis relapse initiating clones, or dRI.

"We didn't just want to do a [genetic analysis](#); we wanted to understand what the precise clonal hierarchy looks like to get a better understanding of when relapse arises and how it can be affected by treatment," said co-senior author Charles Mullighan, M.D., MBBS, of the St. Jude Department of Pathology. "This work helped us show that treatment-resistant clones may be present at diagnosis, before treatment commences, and to determine how these subclones at diagnosis behave during therapy."

The researchers conducted advanced genomic and transcriptomic analysis on the cells. Results showed that dRI clones have increased tolerance to standard chemotherapy. Findings also show that dRI clones have alterations in epigenetic, metabolic and pro-survival pathways that provide new avenues for overcoming resistance.

Advanced genomics coupled with sophisticated modeling

In addition to advanced genomic and transcriptomic analysis, the researchers relied on sophisticated modeling with patient-derived xenografts to conduct this study. The scientists created the models using differing doses of leukemia cells in mice. This increased their ability to capture rare subclones in the models. This step was instrumental in showing the existence of subclones at diagnosis that harbor latent or relapse-specific genetic variants.

"Xenografting added considerable new insight into the evolutionary fates and patterns of subclones obtained from diagnosis samples," said co-senior author John Dick, Ph.D., senior scientist at Princess Margaret Cancer Centre, professor of molecular genetics at the University of Toronto, and scientific co-leader of the Acute Leukemia Translational Research Initiative at the Ontario Institute for Cancer Research. "We were able to gather extensive information about the genetics of the subclones from our models, which helped us describe the trajectories of each subclone and the order in which they acquired mutations. With such subclones isolated by xenografting, we could go on to examine the reasons why these subclones uniquely could both survive therapy and regenerate relapse disease."

The research described in the *Cancer Discovery* paper is a companion study to work published in *Blood Cancer Discovery* in January 2020. The *Blood Cancer Discovery* paper described the genomic landscape, patterns and mechanisms of clonal evolution from [diagnosis](#) to relapse in 92 patients on the Total Therapy protocols for ALL treatment at St. Jude. Together, these papers provide an integrated genomic and functional approach to describing the underlying genetics and mechanisms of relapse for ALL.

Improving the care and treatment of ALL

These findings have important implications for the clinical care of patients with ALL. When clinicians know that a patient is at risk of relapse, they can increase treatment up-front to try to prevent relapse. Additionally, knowing which subpopulation of cells is going to drive the [relapse](#) provides a clue for developing new treatment approaches.

More information: Stephanie M. Dobson et al. Relapse fated latent diagnosis subclones in acute B lineage leukaemia are drug tolerant and possess distinct metabolic programs, *Cancer Discovery* (2020). [DOI: 10.1158/2159-8290.CD-19-1059](https://doi.org/10.1158/2159-8290.CD-19-1059)

Esmé Waanders et al. Mutational Landscape and Patterns of Clonal Evolution in Relapsed Pediatric Acute Lymphoblastic Leukemia, *Blood Cancer Discovery* (2020). [DOI: 10.1158/0008-5472.BCD-19-0041](https://doi.org/10.1158/0008-5472.BCD-19-0041)

Provided by St. Jude Children's Research Hospital

Citation: Study reveals properties of cells fated to relapse in acute lymphoblastic leukemia (2020, March 3) retrieved 20 April 2023 from <https://medicalxpress.com/news/2020-03-reveals-properties-cells-fated-relapse.html>

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