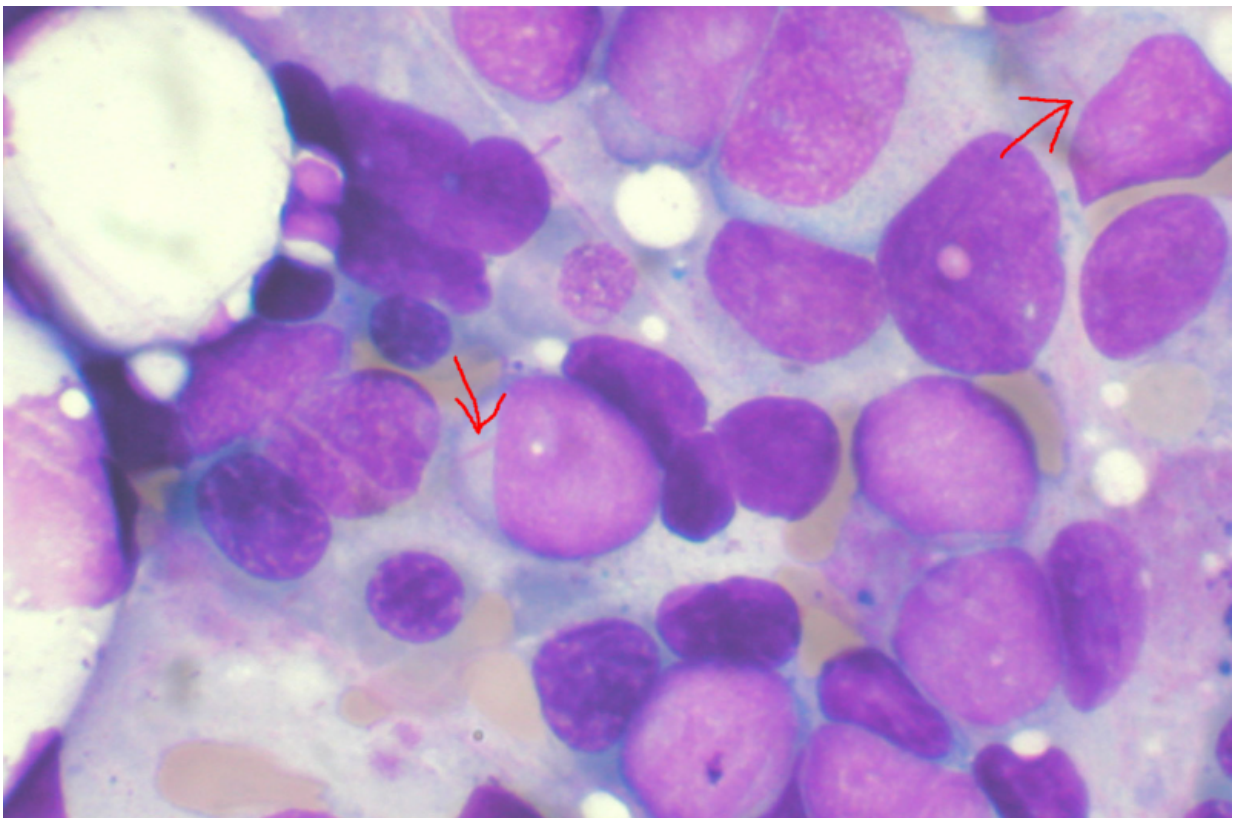


# Researchers discover drug resistance mechanism in leukemia, identify treatment strategy

October 6 2020

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Bone marrow aspirate showing acute myeloid leukemia. Several blasts have Auer rods. Credit: Wikipedia

Like packaging designed to create a protective environment around a

product, the mix of cells and fluids immediately surrounding human bone marrow provides critical protective and nourishing conditions for hematopoietic—or bone marrow-originating—stem cells. Immune cells and other specialized components native to this small-scale "microenvironment" ensure that newly emerged hematopoietic stem cells are healthy and functional.

Nonetheless, the [bone marrow](#) microenvironment is vulnerable to manipulation, notably in the case of [leukemia](#). Leukemia, a cancer of blood-forming tissues, modulates the [bone](#) marrow niche to protect and promote the survival of tumor cells. Now, in a highly collaborative research effort, scientists at the Lewis Katz School of Medicine at Temple University (LKSOM) show that this protective environment also gives leukemia cells with insufficient expression of BRCA1 and BRCA2 genes the ability to resist [anticancer drugs](#) known as PARP inhibitors. This resistance, the researchers discovered, hinges on overexpression of a molecule known as transforming growth factor beta receptor (TGFβR) [kinase](#), which is located on the leukemia cell surface.

The new study, published online October 6 in the journal *Cell Reports* and involving researchers at multiple institutions in the United States and abroad, is the first to show that resistance to PARP inhibitors in leukemia can be overcome by combining PARP inhibition with blockade of TGFβR kinase activation.

"Leukemia cells usually reside in two environments in the body, the bone marrow and the blood," explained Tomasz Skorski, MD, Ph.D., DSc, Professor of Microbiology and Immunology, Associate Professor at the Fels Institute for Cancer Research and Molecular Biology at LKSOM, and senior investigator on the study. "Previous research has shown that leukemia cells displaying deficiency of BRCA1 and BRCA2 proteins are sensitive to PARP inhibition while circulating in the blood. We discovered that the same leukemia cells are resistant to the inhibitors in

the bone marrow microenvironment."

PARP inhibitors trigger a phenomenon known as "synthetic lethality" in cancer cells. They kill malignant cells by shutting down a specific DNA repair mechanism and are effective especially against cells with BRCA1 and BRCA2 gene mutations, in which the homologous recombination mechanism of DNA repair has already been disabled.

Dr. Skorski and colleagues investigated the mechanistic impact of the bone marrow microenvironment on PARP inhibitor resistance in BRCA1/2-deficient leukemia cells derived from human patients and leukemia-bearing mice. The cells were grown in incubation conditions that mimicked the bone marrow microenvironment, enabling the malignant cells to establish drug resistance.

Examination of factors in the replicated bone marrow microenvironment identified TGF- $\beta$ 1 as an important player in resistance. TGF- $\beta$ 1, a protein generated by stromal cells in the bone marrow niche, activates TGF $\beta$ R kinase. Leukemia cells in the bone marrow microenvironment were found to be highly responsive to TGF- $\beta$ 1, owing to low oxygen levels that induce overexpression of TGF $\beta$ R kinase on their surface.

With TGF $\beta$ R kinase levels noticeably elevated in leukemia cells, the researchers decided to test the effects of TGF $\beta$ R kinase inhibition. They observed that treatment with molecules that blocked TGF $\beta$ R kinase activation by TGF- $\beta$ 1 not only halted signaling along the TGF- $\beta$ 1-TGF $\beta$ R kinase axis but also rendered [cells](#) sensitive to PARP inhibitors.

Those observations carried through in animals, where targeting of the TGF $\beta$ R kinase restored leukemia cell sensitivity to PARP inhibitors. Moreover, leukemia-bearing mice treated with a PARP inhibitor plus a TGF $\beta$ R kinase inhibitor survived longer than mice treated only with

PARP inhibition.

"We've now discovered a central and constitutive mechanism underlying PARP drug resistance in leukemia," Dr. Skorski said. "And we went a step further, showing that resistance can be overcome through a therapeutic strategy that combines inhibitors targeting PARP and TGF $\beta$ R kinase."

Dr. Skorski and colleagues plan next to investigate this strategy clinically.

"The drugs we experimented with in our latest research are already approved for use in patients," he noted. "We look forward to future collaborations that will allow us to translate our findings on combined PARP and TGF $\beta$ R kinase inhibitor therapy to the clinic."

Provided by Temple University

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