

# Misregulated protein breakdown promotes leukemias and brain cancer

November 9 2017

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An enzyme that is responsible for the breakdown of specific amino acids in food plays a key role in the development of leukemias and brain cancer, according to scientists from the German Cancer Research Center (DKFZ) in Heidelberg. They have reported their findings in *Nature*. The researchers have discovered a surprising link between energy metabolism and the so-called epigenetic code. These labels in the DNA of cancer stem cells determine the activity of genes, and thus, many cellular functions. The authors think that blocking this enzyme is a promising approach to combat cancer.

Acute myeloid leukemia (AML) is an aggressive type of blood [cancer](#) that very often relapses in the wake of successful initial treatment. Stem cells that are resistant to therapy are believed to be responsible. Scientists working with Andreas Trumpp from the German Cancer Research Center (Deutsches Krebsforschungszentrum, DKFZ) and HI-STEM wanted to understand the molecular background to this resistance. To this end, they examined patient samples by comparing the composition of proteins of AML stem cells and leukemia cells without stem cells properties.

The investigators found suspiciously high levels of an enzyme called BCAT1 in the stem cells. These levels rose even higher during a cancer recurrence. The researchers considered this to be a clue that BCAT1 might be linked to therapy resistance. Cancer researchers have suspected for some time that the BCAT1 enzyme, which is responsible for the breakdown of specific proteins in food, plays a role in the development

of malignant tumors. A team led by Bernhard Radlwimmer from DKFZ had recently found that an overproduction of BCAT1 increases the aggressiveness of [malignant brain tumors](#) and breast cancer.

The two research groups have now joined forces in order to find out how BCAT1 influences the therapy resistance of [leukemia stem cells](#). They discovered a previously unknown process in which a key molecule of [energy metabolism](#) plays a crucial role. BCAT1 reduces the levels of this key molecule and this leads to increased levels of chemical labels in the DNA.

"These tiny methyl groups that are attached to DNA determine whether particular genes are active or silent and, thus, have an immense impact on all [cellular functions](#)," said Simon Raffel, one of the study's first authors.

This result was notable, because in another 30 percent of AML cases, defects in other enzymes lead to the same consequence: They increase cancer-promoting methylation of DNA. AML is known for an extremely heterogeneous pattern of genetic alterations. However, misregulated methylation with its drastic consequences for the whole cell appears to be a common characteristic of this malignant disease.

The finding that BCAT1 drives cancer-promoting methylation in AML stem cells and other cancer [stem cells](#) opens up new options for therapy: "A blockade of the enzyme using a targeted agent might normalize DNA methylation and thereby reduce cancer spread and therapy resistance," Trumpp said.

**More information:** Simon Raffel et al, BCAT1 restricts  $\alpha$ KG levels in AML stem cells leading to IDHmut-like DNA hypermethylation, *Nature* (2017). [DOI: 10.1038/nature24294](https://doi.org/10.1038/nature24294)

Provided by German Cancer Research Center

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