

Ready to start planning your care? Call us at [800-525-2225](tel:800-525-2225) to make an appointment.

×



Memorial Sloan Kettering
Cancer Center

[Make an Appointment](#)

[Press Releases](#)

[Learn About Our Center & Treatment](#)

[Refer a Patient](#)

ABOUT US

[Our mission, vision & core values](#)

[Leadership](#)

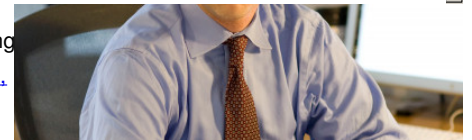
[History](#)

[Equality, diversity & inclusion](#)

[Annual report](#)

[Give to MSK](#)

The discovery, published in the July issue of *Cancer Cell*, provides a key insight into what first goes wrong in the development of many leukemias. The finding was made by a research group led by [Ross L. Levine, MD](#), a member of the Human Oncology Pathogenesis Program and the Leukemia Service at Memorial Sloan Kettering, and Iannis Aifantis, PhD, a member of the NYU Cancer Institute.



Ross Levine

Central to the discovery was the development of a mouse model lacking *TET2* function, which will serve as a valuable research tool. “We now have a model that will allow us to look for therapeutic targets that might be effective against leukemias caused by the *TET2* mutation,” said Dr. Levine. “After proving that *TET2* loss confers a new capacity on these stem cells, we can start investigating whether existing or novel therapies might block that effect.”

Researchers have known that mutations in the *TET2* gene are common in many blood cancers, but other gene mutations are associated with leukemias as well, so the role of *TET2* in leukemia development was unclear. Drs. Levine and Aifantis created the first *TET2*-deficient mouse model to answer this question.

This study is a chapter in a story that is evolving very rapidly. Many other research groups are studying the basic biology of epigenetic regulators like *TET2* in parallel to us, so what we and others learn about the mechanism is laying the groundwork for the development of novel therapies for leukemia patients.

Ross L. Levine, MD, a member of the Human Oncology Pathogenesis Program and the Leukemia Service at Memorial Sloan Kettering

Because *TET2* appeared to be relevant to blood cancers, the mice were engineered to carry a *TET2* mutation in blood cells only. The loss of *TET2* had two dramatic effects: 1) it increased the function of blood stem cells, which in turn allowed these mutant cells to accumulate in the bone marrow and outnumber normal stem cells, and 2) over the next six months, the mice went on to develop myeloid leukemia.

“For the first time, we have definitive proof for what a *TET2* mutation by itself does to the blood cells,” said Dr. Levine, noting that while this mutation alone may not always lead to leukemia — unknown mutations in other genes may need to occur as well — the results of the experiment prove that *TET2* plays a critical role.

The study also provides a glimpse into how mutations in genes known as epigenetic regulators contribute to leukemia development. These types of

genes, which include *TET2*, function by modifying how other genes are expressed without altering their DNA sequence. Instead, epigenetic regulators modify the structure of molecules that surround DNA. Dr. Levine and his colleagues are now trying to further define exactly how *TET2* deficiency changes gene expression in the mutant stem cells, spurring them to outperform the normal stem cells.

Although the mice in the study specifically developed myeloid leukemia, the researchers say it is likely that *TET2* deficiency plays a role in many blood cancers and may contribute to other types of cancer as well. Because mutations in *TET2* frequently occur with mutations in other genes that are linked to cancer, Dr. Levine is working on creating mouse models that carry multiple mutations.

“This study is a chapter in a story that is evolving very rapidly,” Dr. Levine said. “Many other research groups are studying the basic biology of epigenetic regulators like *TET2* in parallel to us, so what we and others learn about the mechanism is laying the groundwork for the development of novel therapies for leukemia patients.”

Researchers from Weill Cornell Medical College, the University of Chicago, and the MD Anderson Cancer Center also contributed to this work. The research was supported by the National Institutes of Health, the [Starr Cancer Consortium](#), and the Howard Hughes Medical Institute.

- Connect
 - Contact us
 - Locations
- APPOINTMENTS
 - 800-525-2225
 -
 -
 -
 -
 -
- About MSK
 - About us
 - Careers ■
 - Giving ■
- Cancer Care
 - Adult cancer types
 - Child & teen cancer types
 - Integrative medicine
 - Nutrition & cancer
 - Find a doctor
- Research & Education
 - Sloan Kettering Institute
 - Gerstner Sloan Kettering Graduate School ■
 - Graduate medical education
 - MSK Library ■

[Communication preferences](#)

[Cookie preferences](#)

[Legal disclaimer](#)

[Accessibility statement](#)

[Privacy policy](#)

[Price transparency](#)

[Public notices](#)

© 2024 Memorial Sloan Kettering Cancer Center