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FOR THE MEDIA



This X-ray image shows metastatic bone tumors in a mouse with kidney cancer. The metastatic tumors appear as hollow or dark areas in the bones.

Summary

Scientists have identified genes and biological mechanisms that one day could be targeted with drugs to stop kidney cancer from spreading to the bone, brain, or other organs.

[Metastasis](#), the process by which some tumors spread from their site of origin to other body parts, accounts for more than nine out of ten cancer-related deaths. But scientists still know relatively little about the genes and biological processes that cause metastasis to occur — information that potentially could translate into new ways to stop cancers from reaching advanced or terminal stages.

Now a Memorial Sloan Kettering research team has shed light on the mechanisms by which [kidney cancer](#) metastasizes to distant organs, including the lungs, bone, and brain. [Published in the journal *Nature Medicine* in December](#) , the findings point to potential therapeutic strategies to control the spread of the disease. The study also offers scientific insights that could advance metastasis research in other cancer types.

Focusing on Kidney Cancer

The research was led by cancer biologist [Joan Massagué](#) , Chair of the [Sloan Kettering Institute's Cancer Biology and Genetics Program](#) and Director of the [Metastasis Research Center](#) . In recent years, his laboratory has uncovered basic causes of metastasis in a number of cancer types, including [breast](#) and [lung cancers](#) .

"In this study, we focused on clear cell renal cell carcinoma, the most common subtype of kidney cancer, for which there is an urgent need for more-effective therapies," says postdoctoral fellow [Sakari Vanharanta](#) , the first author of the *Nature Medicine* report. "The metastatic form of this disease is almost always incurable."

In the vast majority of patients, clear cell renal cell carcinoma (ccRCC) tumors carry DNA changes in a gene called *VHL*. These mutations have been shown to cause the formation of primary kidney tumors, but they do not necessarily lead to metastasis. Until recently, researchers did not know what makes some renal cell carcinoma cells capable of forming secondary tumors in distant organs.

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New Potential Drug Targets

In the recent study, the team addressed this question by performing experiments in mouse models and cell lines, and by analyzing biological and clinical data from more than 700 patients with ccRCC, whose tumors had been analyzed in large-scale cancer genomics projects.

They discovered that two genes called *CYTIP* and *CXCR4* are activated in metastatic tumor cells but inactive in non-metastatic cells. Their experiments suggest that the activation of the two genes might be essential for the spread of kidney cancer.

"*CXCR4* has been linked to metastasis before in this and other tumor types, including [breast cancer](#) ," Dr. Vanharanta says. "Now, our study shows that blocking *CXCR4* function with a drug called plerixafor can reduce kidney cancer metastasis in mice." Plerixafor (Mozobil TM) is currently used to stimulate blood stem cells in some cancer patients treated by bone marrow transplantation.

The researchers plan to investigate further whether *CXCR4* and *CYTIP*, and other genes identified in the study, might offer new targets for the development of more-effective drugs for kidney cancer.

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Exploring the Epigenetics of Metastasis

In addition, the investigators explored the mechanisms by which the *CXCR4* and *CYTIP* genes are switched on in kidney cancer cells to incite metastasis. Their study revealed that the genes undergo a series of epigenetic changes — modifications in the proteins that package a cell's DNA and regulate genes.

Unlike gene mutations, which alter a cell's genetic code, epigenetic changes leave the DNA sequence unaffected. Nevertheless, such changes can influence a cell's behavior by switching individual genes on or off.

Epigenetic modifications are commonly seen in many types of cancer and have recently been associated with more-advanced disease. However, little is known about the specific genes and mechanisms by which tumor cells may reconfigure their epigenetic makeup, causing a person's disease to progress and establish itself in new organs.

"Our study has demonstrated with clear examples how epigenetic alterations can lead to the activation of metastasis-inducing genes," Dr. Vanharanta notes. "This is a conceptual advancement that is likely to help us understand how metastasis occurs in kidney cancer as well as in other cancer types."

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