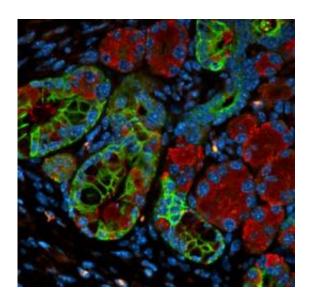


Gene linked to pancreatic cancer growth, study finds

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When Kras is turned off, early stage precancerous pancreatic lesions transform into normal pancreas cells. Credit: University of Michigan Comprehensive Cancer Center

A mutant protein found in nearly all pancreatic cancers plays a role not only in the cancer's development but in its continued growth, according to a new study from University of Michigan Comprehensive Cancer Center researchers. The finding suggests a possible target for developing new ways to treat this deadly disease.

Researchers have known that mutations in the Kras gene are what cause pancreatic cancer to develop. These mutations are frequently seen in



common precancerous lesions, suggesting it has an early role in pancreatic cancer.

The new study, published in the February <u>Journal of Clinical</u> <u>Investigation</u>, finds that in mice, mutant Kras also keeps the tumor growing and helps precancerous tumors grow into invasive cancer. When the researchers turned off Kras, the tumors disappeared and showed no signs of recurring.

The researchers were able to manipulate Kras in a mouse model that they designed to look at Kras at various points in pancreatic <u>cancer</u> <u>development</u>. In the <u>precancerous lesions</u>, turning off Kras eliminated the tumors in mice and the pancreas tissue returned to normal, with no signs of the cancer returning. With <u>invasive cancer</u>, inactivating Kras killed off the cancer but left the pancreas with fibrous areas similar to <u>scar tissue</u>. Tumors did not recur.

Researchers hope this finding provides the basis for future drug development.

"Right now no drugs specifically target Kras, but there are drugs that target the cellular processes downstream of Kras. We next need to figure out which of these downstream effectors of Kras are important in pancreatic cancer," says study author Marina Pasca di Magliano, Ph.D., assistant professor of surgery and of cell and developmental biology at the U-M Medical School.

Kras is also known to play a role in lung and <u>colon cancer</u>. But it is likely the biggest player in pancreatic cancer, where more than 90 percent of all tumors have mutated Kras. Pancreatic cancer is one of the most deadly types of cancer: about 4 percent of patients are alive five years after their diagnosis. The disease is often diagnosed when surgery is not an option and it tends to be resistant to available chemotherapies.



"There is a dire need for new therapies for pancreas cancer based on a better understanding of the biology of this disease. My lab is now looking at the downstream inhibitors of Kras to try to find the best target," Pasca di Magliano says.

More information: *Journal of Clinical Investigation*, Vol. 122, No. 2, February 2012

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