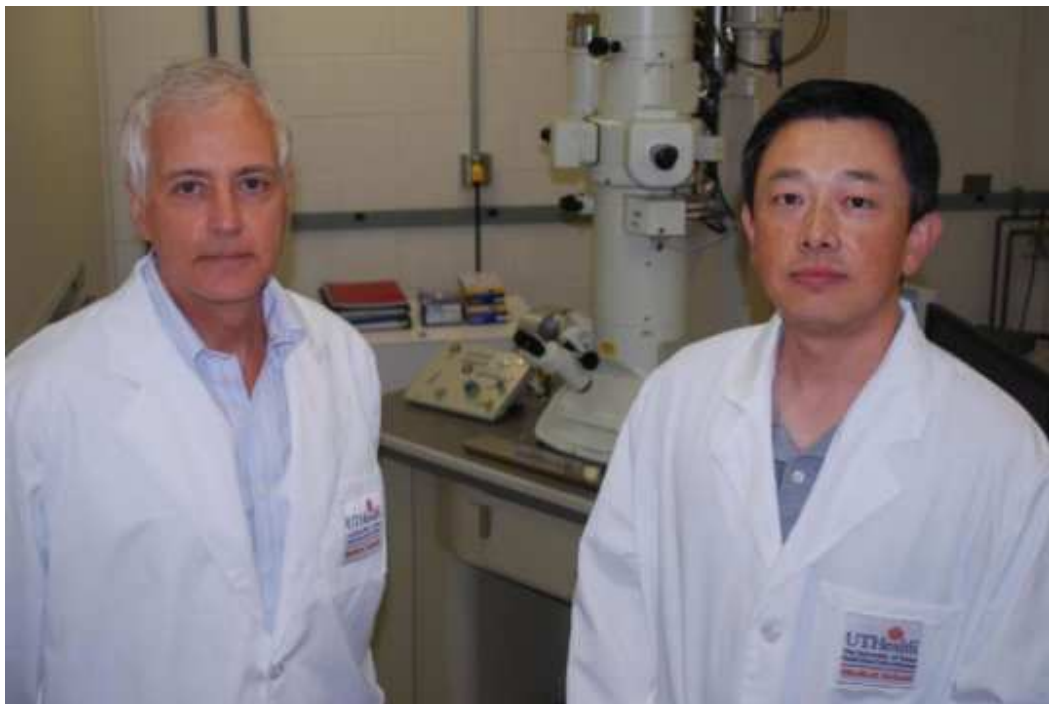


## Scientists discover electrical control of cancer cell growth

Date: August 24, 2015

Source: University of Texas Health Science Center at Houston

Summary: The molecular switches regulating human cell growth do a great job of replacing cells that die during the course of a lifetime. But when they misfire, life-threatening cancers can occur. Research has now revealed a new electrical mechanism that can control these switches.



UTHealth's John Hancock, M.B., B.Chir, Ph.D., ScD, left, and Yong Zhou, Ph.D., are studying what causes cancer at the molecular level.

*Credit: The University of Texas Health Science Center at Houston (UTHealth)*

The molecular switches regulating human cell growth do a great job of replacing cells that die during the course of a lifetime. But when they misfire, life-threatening cancers can occur. Research led by scientists at The University of Texas Health Science Center at Houston (UTHealth) has revealed a new electrical mechanism that can control these switches.

This information is seen as critical in developing treatments for some of the most lethal types of cancer including pancreatic, colon and lung, which are characterized by uncontrolled cell growth caused by breakdowns in cell signaling cascades.

The research focused on a molecular switch called K-Ras. Mutated versions of K-Ras are found in about 20 percent of all human cancers in the United States and these mutations lock the K-Ras switch in the on position.

"When K-Ras is locked in the on position, it drives cell division, which leads to the production of a cancer," said John Hancock, M.B., B.Chir, Ph.D., ScD, the study's senior author and chairman of the Department of Integrative Biology and Pharmacology at UTHealth Medical School. "We have identified a completely new molecular mechanism that further enhances the activity of K-Ras."

Findings appear in *Science*, a journal of the American Association for the Advancement of Science.

The study focused on the tiny electrical charges that all cells carry across their limiting (plasma) membrane. "What we have shown is that the electrical potential (charge) that a cell carries is inversely proportional to the strength of a K-Ras signal," Hancock said.

With the aid of a high-powered electron microscope, the investigators observed that certain lipid molecules in the plasma membrane respond to an electrical charge, which in turn amplifies the output of the Ras signaling circuit. This is exactly like a transistor in an electronic circuit board.

Yong Zhou, Ph.D., first author and assistant professor of integrative biology and pharmacology at UTHealth Medical School, said, "Our results may finally account for a long-standing but unexplained observation that many cancer cells actively try to reduce their electrical charge."

Initial work was done with human and animal cells and findings were subsequently confirmed in a fruit fly model on membrane organization.

"This has huge implications for biology," Hancock said. "Beyond the immediate relevance to K-Ras in cancer, it is a completely new way that cells can use electrical charge to control a multitude of signaling pathways, which may be particularly relevant to the nervous system."

Hancock's co-authors at UTHealth include Ching-On Wong Ph.D., Kwang-Jin Cho, Ph.D., Dharini van der Hoeven, Ph.D., Hong Liang, M.D., Dhananjay Thakur, Jialie Luo, Ph.D., Michael Zhu, Ph.D., Hongzhen Hu, Ph.D., and Kartik Venkatachalam, Ph.D.

Co-authors from The University of Arizona include Milos Babic, Ph.D., and Konrad Zinsmaier Ph.D.

At UTHealth Medical School, Hancock is the vice dean for basic research, executive director of the Brown Foundation Institute of Molecular Medicine for the Prevention of Human Diseases and holder of the John S. Dunn Distinguished University Chair in Physiology and Medicine.

Hancock and Venkatachalam are on the faculty of The University of Texas Graduate School of Biomedical Sciences at Houston.

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### Story Source:

The above post is reprinted from [materials](#) provided by [University of Texas Health Science Center at Houston](#). (<https://www.uth.edu/>) *Note: Materials may be edited for content and length.*

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