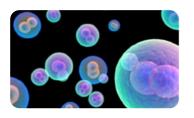


## Stopping cell migration may help block fibrosis and the spread of cancer

Discoveries by a Yale-led team of scientists could lead the way for development of new therapies for treating fibrosis and tumor metastasis. The researchers have both uncovered a signaling pathway that promotes cell migration in certain forms of pulmonary fibrosis, a deadly lung disease, and developed a drug treatment that ma block the cancer cell migration. The study appears in the Advance Online Publication of Nature Cell Biology.

By <u>Helen Dodson</u><sup>21</sup> May 2012



Pulmonary fibrosis is the development of excessive connective tissue in the lungs. It can develop without a known cause, a produces permanent scar tissue in the lungs. There is no known cure or treatment to slow down its progress.

Cell migration is the biological process by which cells move around the body, often contributing to the development or spre of diseases such as fibrosis or metastatic cancer.

Fibroblast connective tissue cells and cancer cells migrate in a way that requires sustained activation of signaling pathway But until now the regulation of these cellular functions has been poorly understood. The Yale team studied the workings of stimulant of cell migration known as lysophosphatidic acid (LPA). They identified the chemical reactions that ultimately induce cell migration, but more importantly, discovered a way to block the pathway via inhibition of the proteins responsibl for promoting the migration.

Senior author Dianqing (Dan) Wu, professor of pharmacology and vascular biology at Yale School of Medicine and meml of Yale Cancer Center, explained, "Our ability to block the pathway provides a potential therapeutic target for treating pulmonary fibrosis, a very serious disease that lacks effective treatments, and other types of fibrosis. Because cancer cel particularly melanoma and lung cancer cells containing activated BRAF genetic mutations, can use this signaling pathway migrate, blocking this pathway could also prevent metastasis of these cancers."

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