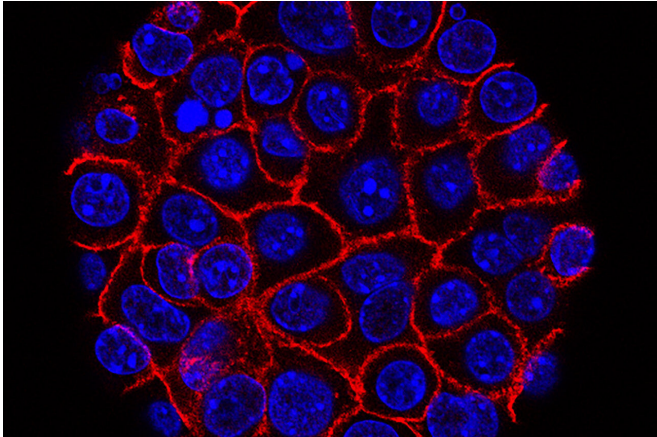


Researchers discover cancer-associated fibroblasts induce drug sensitivity

August 16 2022



Credit: Min Yu (Eli and Edythe Broad Center for Regenerative Medicine and Stem Cell Research at USC), USC Norris Comprehensive Cancer Center

Cells in the surrounding tumor microenvironment have a significant impact on cancer growth, survival and responses to therapy. Cancer-associated fibroblasts in the tumor environment have typically been associated with tumor progression and resistance to therapy, despite some studies suggesting that these fibroblasts may also sensitize cancer cells to therapy. In a new article published in *Science Signaling*, Moffitt Cancer Center researchers shed light on these conflicting studies and demonstrate that cancer-associated fibroblasts can promote or inhibit drug sensitivity based on the type of tumor cell and the drug used for treatment.

Through a series of laboratory experiments, the research team determined the impact of cancer-associated fibroblasts on drug responses among different non-small cell lung cancer cell lines. They discovered that the presence of cancer-associated fibroblasts had varying effects on [tumor cells](#) based on both the type of non-small cell lung cancer and the drug used for treatment. For

example, the presence of cancer-associated fibroblasts induced resistance to two different MEK inhibitors in non-small cell lung cancer cell lines with a mutant KRAS protein. However, cancer-associated fibroblasts sensitized non-small cell lung cancer cell lines with a mutant EGFR protein to EGFR inhibitors. Interestingly, normal lung-associated fibroblasts never sensitized cells to drug treatment, suggesting that cancer-associated fibroblasts secrete a factor that causes differential responses to drug treatment in a cell-context manner.

The researchers compared cancer-associated fibroblasts to normal fibroblasts to identify factors that would produce these disparate effects. They found that cancer-associated fibroblasts had alterations in the levels of secreted proteins that are part of the insulin-like growth factor (IGF) signaling pathway, which is involved in cell growth, death and migration. Specifically, cancer-associated fibroblasts secreted higher levels of proteins called IGF binding proteins (IGFBPs), which inhibit IGF signaling; and lower levels of IGFs, which activate IGF signaling. In combination, these alterations result in inhibitory effects on the IGF signaling pathway.

In further analyses, the researchers found that IGFBPs sensitized lung cancer cell lines to EGFR inhibitor treatment, while IGF proteins induced resistance to EGFR inhibitor treatment. They identified that survival signaling in response to EGFR inhibitor treatment was dependent on the proteins IGF1R and FAK, which are both part of the IGFBP signaling pathway. Importantly, they discovered that drugs that blocked the activity of IGF1R and FAK sensitized mutant EGFR lung [cancer cells](#) to EGFR inhibitors, suggesting that this combination approach may be effective in the clinic.

"These results highlight tumor-suppressive effects competing with otherwise tumor-promoting effects of cancer-associated fibroblasts and add to the

growing evidence that eliminating cancer-associated [fibroblasts](#) in an undifferentiated way may be detrimental to cancer therapy," said lead study author Lily Remsing Rix, Ph.D., a research scientist at Moffitt.

"We show that mechanistic understanding not just of cancer-associated fibroblast-mediated resistance, but also of their tumor-suppressive pathways, can lead to rational design of improved therapeutic approaches that mimic these effects and may delay the onset of drug resistance," added Uwe Rix, Ph.D., associate member of the Department of Drug Discovery at Moffitt and principal investigator of the study.

More information: Lily L. Remsing Rix et al, IGF-binding proteins secreted by cancer-associated fibroblasts induce context-dependent drug sensitization of lung cancer cells, *Science Signaling* (2022). DOI: [10.1126/scisignal.abj5879](https://doi.org/10.1126/scisignal.abj5879)

Provided by H. Lee Moffitt Cancer Center & Research Institute

APA citation: Researchers discover cancer-associated fibroblasts induce drug sensitivity (2022, August 16) retrieved 8 December 2022 from <https://medicalxpress.com/news/2022-08-cancer-associated-fibroblasts-drug-sensitivity.html>

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