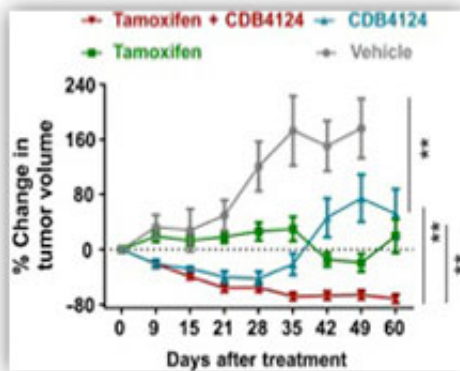


Steroid receptor crosstalk informs therapies for breast cancer treatment

June 28 2016



Tumor cells in the placebo-treated mice grew rapidly. Tamoxifen prevented the tumors from growing. CDB4124 initially caused the tumors to regress, but after 35 days, the tumors began growing again. The combination of tamoxifen and CDB4124 caused tumors to shrink 70 percent by day 60. Credit: University of Chicago Medical Center

One of the first clues pathologists look for in tissue from a newly diagnosed breast cancer patient is the estrogen receptor, a nuclear protein that converts hormonal messages in the bloodstream into instructions for the cell about how to behave. They also look for the presence of progesterone receptors, primarily to confirm that the estrogen receptor is active.

In the June 24 issue of *Science Advances*, however, researchers radically

upgrade the significance of the progesterone receptor. They show that when exposed to estrogens and progestins, these [receptor proteins](#) interact with different sets of binding sites in the cell's chromosomes, with the progesterone receptor dramatically altering how estrogen receptors interact with the cell's DNA.

"In the last year or two, researchers have tuned into the extensive and previously unrecognized cross talk between the progesterone and estrogen receptors," said study author Geoffrey Greene, PhD, Co-Director of the Ludwig Center for Metastasis Research and Chairman of the Ben May Department for Cancer Research at the University of Chicago. "We now know that this relationship can be better understood and potentially exploited."

"This observation is important," he added, "because more than two-thirds of breast cancers contain both estrogen and [progesterone receptors](#)."

Previous studies from many labs, including a group at Cambridge University, showed that when exposed to the primary female sex hormone, estradiol, the [estrogen receptor](#) activates genes that encourage tumor cells to grow and divide. "In this setting, the cancer cells grow faster," Greene said. "They divide faster and they live longer, which helps the breast cancer progress to a more advanced state."

"Our study, as well as that of the Cambridge group, showed that when progesterone or a progestin is added, it changes the landscape in the nucleus," Greene said. "It opens up a whole new set of binding sites for the estrogen receptor that now work in conjunction with the progesterone receptor." This combination inhibits cellular proliferation, cell survival and pathways involved in metastasis.

"Our data further suggest that, despite the historical bias toward the

effects of estrogen on the estrogen receptor, it's the progesterone receptor that dominantly controls estrogen receptor activity when both receptors are present and activated."

Inhibiting both receptors makes tumors shrink

In their *Science Advances* paper, Greene, lead author Hari Singhal, now a postdoctoral scholar at Harvard University, and colleagues took a step further. Building on the well-established use of tamoxifen, an estrogen antagonist, to treat women with estrogen receptor-positive breast cancers, they decided to inhibit the activities of both receptors with antagonists to see if there would be an added benefit of combining two receptor-selective drugs. Using a well-established ER+/PR+ human breast cancer model (T47D) implanted in immune-compromised mice, they tested their theory in four groups of mice.

One group received a placebo. A second group received tamoxifen. The third group was treated with a progesterone receptor antagonist, an experimental drug known as CDB4124 (Telapristone). The fourth group was treated with tamoxifen plus CDB4124 to simultaneously block both the estrogen and progesterone receptors.

Tumor cells in the placebo-treated mice grew rapidly, to about 200 percent their original size in seven weeks. Tamoxifen, the estrogen antagonist, prevented the tumors from growing but did not cause them to shrink. CDB4124, the progestin antagonist, initially caused the tumors to regress, but after 35 days, the tumors began growing again; by seven weeks they were about 50 percent larger than their original size.

Significantly, the combination of tamoxifen and CDB4124 caused tumors to shrink. "Virtually full regression was observed in a subset of mice," Singhal said. By day 60, the average tumor volume in mice treated with tamoxifen plus CDB4124 was 70 percent lower than the

original tumors.

"These findings," the authors note, "emphasize the clinical value of assessing both progesterone receptor and estrogen receptor expression in [breast cancer](#) samples." Not only is the progesterone receptor an "essential modulator of estrogen-receptor-regulated genes," but it also significantly contributes to the "prognostic value of estrogen receptors in ER+/PR+ breast cancers." In addition, it might serve as an important combinatorial target in these breast cancers.

Provided by University of Chicago Medical Center

Citation: Steroid receptor crosstalk informs therapies for breast cancer treatment (2016, June 28) retrieved 5 May 2024 from

<https://medicalxpress.com/news/2016-06-steroid-receptor-crosstalk-therapies-breast.html>

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