

Drug zeroes in on mutated nuclear receptors found in cancer, leaves normal proteins alone

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Killer T cells surround a cancer cell. Credit: NIH

Researchers at Sanford Burnham Prebys Medical Discovery Institute (SBP) have published a study in *Nature Communications* shedding new light on how K-80003 (TX803), an anti-cancer agent discovered at the



Institute, prevents activation of the PI3K pathway, resulting in inhibition of cancer cell growth. Because the PI3K pathway is common to many cancers, K-80003 could have broad therapeutic applications. Tarrex Biopharma, Inc. has licensed the compound and announced they will soon begin Phase 1 clinical trials at the Dana Farber Cancer Center for patients with colorectal cancer.

"K-80003 binds specifically to a truncated form of the retinoid X receptor-alpha (tRXR α) protein that promotes tumors," says Xaio-kun Zhang, Ph.D., adjunct professor and senior author of the paper. "When it binds tRXR α , it freezes the protein into an inactive (tetrameric) configuration that prevents it from stimulating the PI3K pathway.

"A major goal of this study was to dig deeper and find out why this compound is so effective, and appears to have so few side effects," adds Zhang. "We wanted to visualize how tRXR α and K-80003 physically interact—at atomic resolution—in a way that makes this such a promising cancer drug."

Working with the lab of Robert Liddington, Ph.D., professor at SBP, the research team used X-ray crystallography to find that six molecules of K-80003 bound to the tetramer interfaces.

"Our findings show that K-80003 creates a configuration that locks the tetramer into a tight ball, hiding all of its binding sites so it can't trigger the PI3K pathway," explains Liddington.

"We know tRXR α protein is specifically produced by tumor cells, while normal cells only produce full-length RXR α protein. Taken together, our findings explain the drug's potency and specificity for <u>cancer</u> cells and why the compound has such low toxicity," adds Zhang.

Because RXR α is a nuclear receptor, it has the ability to control gene



expression. The mutation that creates tRXR α puts the protein into overdrive, activating PI3K in the cytoplasm and accelerating tumor growth. K-80003 acts like a Denver Boot, clamping down on tRXR α and eliminating these hyperactive growth signals. In addition, K-80003 often brings tRXR α back to the nucleus, which may further impair its function and enhance the drug's therapeutic impact.

But that was not the end of the discovery. The team also found an LxxLL motif (a common sequence of amino acids) in the p85 α protein, which acts as an intermediary between RXR α and PI3K. Identifying LxxLL sheds new light on how tRXR α binds to the cytoplasmic p85 α protein and activates PI3K. When tRXR α becomes a tetramer, p85 α can no longer bind to it.

These findings also help illustrate how nuclear receptors can function outside the nucleus, and how changes in both conformation and location can dramatically alter their function. In the long-term, understanding these mechanisms could lead to a variety of new therapies. But right now, this research shows why K-80003 has so much potential.

"The tRXR α protein is produced in almost all cancers," said Zhang, "so the compound could be very effective for many different types, such as breast, liver and colon."

More information: Lluís Mañosa et al, Inverse barocaloric effect in the giant magnetocaloric La–Fe–Si–Co compound, *Nature Communications* (2011). DOI: 10.1038/ncomms16066

Provided by Sanford-Burnham Prebys Medical Discovery Institute

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