

Study links normal stem cells to aggressive prostate cancer

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A study that revealed new findings about prostate cells may point to future strategies for treating aggressive and therapy-resistant forms of prostate cancer.

The study proved that the prostate basal cell layer contains adult stem cells which possess a unique gene expression profile resembling the deadliest form of prostate cancer. The research was led by The University of Texas MD Anderson Cancer Center with findings published in the Feb. 29, 2016 online issue of *Nature Communications*.

"It has become very controversial as to whether the human prostate contains adult stem cells or not and where they are located within the basal or luminal cell compartments," said Dean Tang, M.D., Ph.D., professor of Epigenetics and Molecular Carcinogenesis. "Our study provided definitive evidence that the prostate basal cell layer harbors self-renewing adult stem cells that are enriched in stem-cell genes."

Tang and his team headed by Dingxiao Zhang, Ph.D., an instructor in Tang's lab, show that the findings point to a "theoretical rationale for combining Pol-I and MYC inhibitors to treat highly aggressive forms of prostate cancer which are resistant to endocrine therapy." Pol-I is an enzyme involved with DNA replication and MYC is a regulator gene that plays a role in cell death and transformation.

The prostate gland contains basal and luminal cells, both of which have been identified as "cells-of-origin" for prostate cancer in recent mouse

studies. However, the question of whether and where stem cells were present in the human prostate has been largely a medical mystery and a constant debate until now.

Tang's team completed a genome-wide analysis of human benign prostate basal and luminal cells using RNA sequencing and found that they expressed genes differently and that some [basal cells](#) represented self-renewing [adult stem cells](#).

"Strikingly, we found that basal stem cells also expressed a large cohort of 'proneural' genes that are normally involved in regulating the nervous system development," said Tang. "These proneural genes seem to play important functions in conferring stem cell-like properties upon some basal cells."

This finding is important because a subset of [prostate cancers](#) (less than 5 percent) are highly aggressive and do not respond to current anti-prostate cancer treatments such as endocrine therapy.

"Surprisingly, these hard-to-treat cancers also express a gene signature that overlaps with our normal basal stem cell gene expression profile, suggesting that basal stem cells may represent the cell-of-origin for these prostate cancers," said Tang. "Of significance, the basal stem cell [gene expression profile](#) is also linked to endocrine therapy-resistant cancer which is lethal to virtually all advanced prostate cancer patients."

Tang's team also found that basal [stem cells](#) are enriched in a genetic component that is partially regulated by MYC which offers hope that the deadliest of prostate cancers and therapy-resistant prostate cancers may have a new therapeutic option.

"Our studies establish that therapy that combines Pol-I and MYC inhibitors may be a potential new line of treatment for highly metastatic

and endocrine therapy-resistant [prostate](#) cancer," said Tang.

More information: Dingxiao Zhang et al. Stem cell and neurogenic gene-expression profiles link prostate basal cells to aggressive prostate cancer, *Nature Communications* (2016). [DOI: 10.1038/ncomms10798](https://doi.org/10.1038/ncomms10798)

Provided by University of Texas M. D. Anderson Cancer Center

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