

Understanding why men get more cancer than women

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Peter Mac-led research has shed new light on why men are more likely than women to get cancer, uncovering a new role for the X chromosome in affecting cancer risk and mortality.

While men are known to have a higher risk of developing, and dying from, cancer compared to women—even when matched for other known [cancer risk](#) factors like age, race, smoking history and cancer stage—the drivers of this are poorly understood.

To address this apparent sex-disparity in cancer, Dr. Sue Haupt and team studied the DNA of men and women diagnosed with 12 different types of non-reproductive cancers. They focused on the gene TP53—which produces a natural protector of our genome—looking for differences in the mutation of this gene, and irregular behavior of its partner proteins.

"TP53 is the most commonly mutated gene in human cancer, with more than half of all human cancers harboring a genetic alteration that interferes with the function of p53 protein," says Dr. Haupt, co-lead author on the study.

The scientists showed, for the first time, that TP53 mutations are more common in males than females, which could account in part for the greater cancer risk in men.

They then focused their search on the X chromosome, as males have only one X chromosome (XY) putting them at higher risk than females (XX) of developing diseases if genes on their X chromosome become dysfunctional.

"By looking specifically on the X chromosome we were able to find—for the first time—a group of p53-regulating genes that we thought could, in part, explain the sex disparity in cancer," Dr. Haupt says.

Using a new computational approach developed by Franco Caramia, co-lead author and a bioinformatician Ph.D. student in the Haupt lab, they found a cluster of genes located on the X chromosome that can affect

the activity of p53 protein, even if the cancers don't have a mutation in TP53 itself.

They found while women have a higher incidence of mutations on the X chromosome, these mutant genes are often not expressed to make proteins. This was particularly true for the p53-regulating genes. In contrast, men commonly express the mutant forms of X-linked p53-regulating genes.

"These findings come together to unveil a fascinating story of safeguarding of women from p53-induced cancers across three intricate and complex layers of biological protection," says Dr. Haupt.

"First, women are less likely to possess mutations in TP53. Second, the presence of p53-regulating genes on the X chromosome means men are particularly vulnerable to defects in these genes. And lastly, there appears to be a barrier in [women](#) that prevents the expression of mutant X-linked p53-regulating [genes](#)."

Prof Ygal Haupt, the study's senior author, said the findings suggest genetically-assigned sex should be taken into consideration when it comes to risk assessment and potentially even treatment of cancers.

"For non-reproductive cancers, clinical decision-making rarely takes sex into account. Our results suggest a person's chromosomal makeup could directly impact whether they will respond to a specific treatment. This can be particularly important for the use of new drugs aimed at reactivating p53, which are currently in [clinical trials](#)," says Prof Haupt.

This Peter Mac-driven research was done in collaboration with scientists at the Walter and Eliza Hall Institute of Medical Research, MD Anderson Cancer Centre in U.S. and the Karolinska Institute in Sweden.

This research is contained in the paper "Identification of [cancer](#) sex-disparity in the functional integrity of p53 and its X chromosome network" published in the journal *Nature Communications*.

More information: Gennady Korotkevich et al. Fast gene set enrichment analysis, (2016). [DOI: 10.1101/060012](https://doi.org/10.1101/060012)

Provided by Peter MacCallum Cancer Centre

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