

Aggressive prostate cancer secrets revealed in landmark study

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Stylized image of prostate cancer from a man with a BRCA2 mutation. Credit: Monash University

A landmark study, led by Monash University's Biomedicine Discovery Institute with the involvement of the Peter MacCallum Cancer Centre, has revealed the reason why men with a family history of prostate cancer who also carry the BRCA2 gene fault have a more aggressive form of prostate cancer.

The study, published today in *Nature Communications*, involving a consortium of Melbourne and Toronto researchers and clinicians, reports the molecular profile of the prostate cancers in men with the BRCA2 fault is similar to the profile seen in patients with advanced cancer: explaining why - right from diagnosis - BRCA2 patients have a poor



outcome.

The international team of scientists, led by Professor Gail Risbridger and Dr Renea Taylor from the Monash Biomedicine Discovery Institute (BDI) and Dr Rob Bristow from the Princess Margaret Cancer Centre in Toronto, Canada, in collaboration with clinicians from the Peter MacCallum Cancer Centre, kConFab, Austin Health and the Olivia Newton-John Cancer Centre, worked to unlock the secrets of why prostate cancer in BRCA2 men behaves aggressively.

This study, part of a larger Victorian Cancer Agency funded program of translational research, had previously reported that men who carried the BRCA2 gene fault were at a higher risk of having a more aggressive form of prostate cancer if a cell pathology known as IDCP (intraductal carcinoma of the prostate) was present; the IDCP cell pathology predicted these men were much more likely to have a poor clinical outcome.

The reason why this happens is now clear. Following surgery to remove the cancerous prostate, the patient's specimen was made available to the laboratory for genomic analyses. Surprisingly, the study showed that these early, untreated, prostate cancers were genetically similar to cancers that are usually seen in men with more advanced cancer that has spread to other parts of the body. This was in contrast to the cancers seen in men who don't carry a BRCA2 gene fault and who rarely have cancer spread at diagnosis. This was confirmed when compared to data from a companion study, published at the same time in Nature and led by the Toronto group, which looked at prostate cancer tissue samples from more than 320 patients with prostate cancer who don't carry a BRCA2 gene fault.





Professor Gail Risbridger and Dr Renea Taylor, Monash Biomedicine Discovery Institute. Credit: Steven Morton

Put together, these studies identified why the presence of the BRCA2 gene fault led to markedly different clinical outcomes, with the disease progressing rapidly in this group of men.

Director of Monash Partners Comprehensive Cancer Consortium and Monash BDI lead author, Professor Gail Risbridger, explained why these findings are important.

"This study shows how different these tumours are from 'regular'



tumours and emphasises the importance of men knowing if they have a family history of prostate, breast or ovarian cancer in their family and may carry the BRCA2 gene fault," Professor Risbridger said.

Fellow Monash BDI lead author, Dr Renea Taylor, highlighted that these findings prompt the need to think about whether these men should be managed differently.

"As the tumours in men with the BRCA2 gene fault are so different from the 'get-go', our findings raise the question about whether these patients should be managed differently at diagnosis," Dr Taylor said.

Director of Genitourinary Oncology at the Peter MacCallum Cancer Centre and study author, Associate Professor Declan Murphy, spoke about possible therapeutic implications."We now know that the BRCA2 fault is seen in many more men presenting with advanced prostate cancer than previously realised. Also, as prostate cancer progresses, the BRCA2 fault begins to develop in prostate cancer secondaries, and drives the aggressive behaviour of the cancer. Therefore these new findings detailing the genomic instability of BRCA2 prostate cancer are important as we may be able to target this with new therapies," Associate Professor Murphy said.

More information: Renea A. Taylor et al, Germline BRCA2 mutations drive prostate cancers with distinct evolutionary trajectories, *Nature Communications* (2017). DOI: 10.1038/ncomms13671

Provided by Monash University

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