

A Case-Based Clinical Approach to the Investigation, Management and Screening of Families with BRCA2 Related Prostate Cancer

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Abstract: *BRCA2* is the most commonly implicated DNA damage repair gene associated with inherited prostate cancer. *BRCA2* deficient prostate cancer typically presents at a younger age, is more poorly differentiated, and is associated with worse survival outcomes than non-*BRCA2* associated prostate cancer. Despite these unfavourable prognostic implications, poly-ADP ribose polymerase inhibitors and platinum-based chemotherapy have been identified as potent targeted therapeutic agents towards *BRCA1/2* deficient cancer cells. This review article explores the literature surrounding *BRCA2*-related prostate cancer through a familial clinical scenario. The investigation, diagnosis and management of *BRCA2* deficient prostate cancer will be explored, alongside the implications of the identification of a germline pathogenic *BRCA2* variant within a family, cascade screening and prostate cancer surveillance in unaffected male *BRCA2* carriers. A greater understanding of the molecular pathogenesis of DNA damage repair gene deficient prostate cancer, coupled with new treatment paradigms and widened access to both somatic and germline genetic analysis for prostate cancer patients and their families will hopefully enable the robust implementation of high quality evidence-based clinical pathways for both the management and identification of *BRCA2* deficient prostate cancer and improved screening, early detection and prevention strategies for individuals at increased genetic risk of prostate cancer.

Keywords: prostate cancer, BRCA, genomics, PARP inhibitors, clinical management

Introduction

Prostate cancer (PC) is the most common cancer in males in the UK and remains one of the leading causes of morbidity and mortality within the cancer patient population, with approximately 1.4 million new diagnoses and around 375,000 associated deaths globally in 2020.¹ The majority of PC occurs sporadically in males over the age of 65. These cancers are driven by tumour-specific genetic changes that drive oncogenesis (somatic pathogenic variants). Twin studies in Scandinavia have suggested that just over half of predisposition to PC may be due to genetic factors.² Currently, the majority of these are common variants contributing to polygenic inheritance, where multiple genetic loci associated with a low-moderate risk increase an individual's lifetime risk of developing PC. A smaller proportion of cases are due to single moderate-high risk genetic variants.³ These are primarily pathogenic variants in DNA damage repair (DDR) genes.

Common variants predisposing to PC are predominantly identified through large-scale Genome Wide Association Studies (GWAS), and over 160 single

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nucleotide polymorphisms (SNPs) associated with PC have been identified.⁴ Whilst individually these SNPs are low risk (relative risk ~1.1–1.3), their use in combined polygenic risk scores is likely to facilitate future identification of more men at risk of PC in clinical practice and they have given significant insights into the biological pathways and networks underlying oncogenesis.^{4,5}

DNA sequencing of germline DNA in patients with PC is the mainstay of identification of rarer moderate-high penetrance monogenic variants predisposing to PC. In the main, these genes act within DNA damage repair pathways and include *BRCA1*, *BRCA2*, *ATM*, *PALB2*, *HOXB13* as well as mismatch repair genes indicated in Lynch Syndrome, particularly *MSH2*.^{6–10} Germline variants in *BRCA2* have been implicated in the pathogenesis of 1–2% of PCs and have been reported to confer an increased relative risk of approximately 2.4–8.6-fold in men below the age of 65 years, making it the most commonly implicated DDR gene associated with inherited PC.^{3,11–13} *BRCA2* germline variants confer a 20% lifetime risk of developing PC and are associated with not only early-onset disease, but also a worse prognosis than non-carriers of *BRCA2* germline variants.^{3,6,7,11,14–18}

The *BRCA2* gene, located on chromosome 13q12.3, is tumour suppressor gene that follows an autosomal dominant pattern of inheritance. *BRCA2* encodes for a protein which helps to repair double-stranded DNA (dsDNA) breaks through homologous recombination (HR) repair. Most *BRCA2* pathogenic variants are protein truncating variants leading to the formation of a truncated, non-functional protein. *BRCA2* deficient PC in most cases, occurs due to biallelic loss of the *BRCA2* gene within the tumour (loss of both copies of the *BRCA2* gene). This can occur in the tumour alone or can be due to the inheritance of a germline monoallelic pathogenic variant, with loss of the second allele within the tumour in accordance with Knudson's two-hit hypothesis. Therefore, men with inherited *BRCA2* variants have a higher propensity for developing PC if the one remaining unaffected *BRCA2* allele develops a second single somatic pathogenic variant leading to oncogenesis.^{7,19} There are rare exceptions, where the mutant allele only is lost in a tumour, leaving the normal copy implying that in some cases a haplo-insufficiency mechanism leads to tumour development.²⁰

In this review article, we will consider *BRCA2*-related PC, considering a hypothetical family (Figure 1) and discuss the investigation, diagnosis and management of

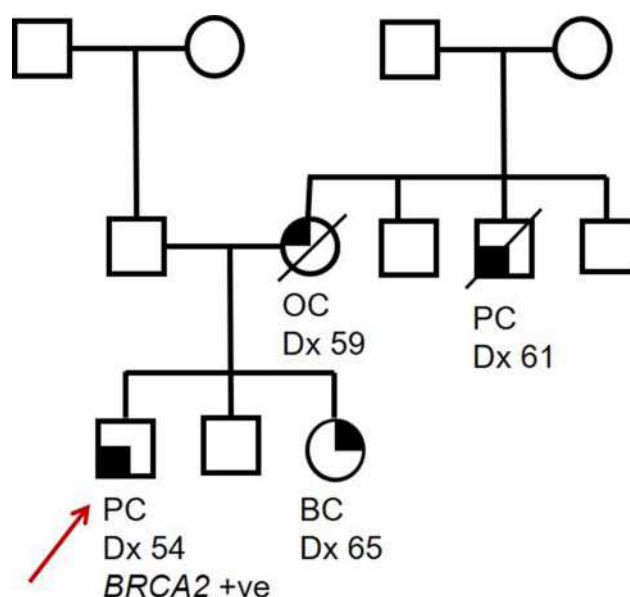


Figure 1 Pedigree of a hypothetical family carrying a pathogenic *BRCA2* germline variant. Proband (indicated by red arrow) was diagnosed with PC at 54 years, proband's sister was diagnosed with BC at 41 years and proband's mother (deceased) was diagnosed with OC at 59 years. A maternal uncle (deceased) was diagnosed with PC at 61 years. Circles indicate females, squares indicate males. A line through a figure indicates that individual is deceased.

Abbreviations: PC, prostate cancer; BC, breast cancer; OC, ovarian cancer; Dx, diagnosis.

BRCA2 deficient PC, alongside the implications of the identification of a germline pathogenic variant in the *BRCA2* gene within a family and PC surveillance in unaffected male *BRCA2* carriers. This clinical case-based approach is intended to summarise the implications of a diagnosis of *BRCA2* deficient prostate cancer for both the proband and wider family and is complementary to more comprehensive systematic reviews^{21–27} providing more in depth scientific review on the diagnosis and management of DDR gene deficient prostate cancer.

Identification and Management of *BRCA2* Deficient Prostate Cancer

The proband in Figure 1 has just been diagnosed with PC at a relatively young age. There are fortunately a number of different treatment options for PC which range from conservative management with active surveillance to radical medical and/or surgical options. The selection of an appropriate clinical treatment plan is dependent on a number of factors which include (but are not limited to) pre-existing patient risk factors, tumour grading and staging, life-expectancy and patient preference. Mainstay treatment options for low-high risk localised PC include active surveillance, radical prostatectomy, radiotherapy (external

beam or brachytherapy) and androgen-deprivation therapy, and high intensity focused ultrasound.^{11,28,29}

Compared to *BRCA* proficient PC (*BRCA2*-ve), *BRCA2* deficient PC (*BRCA2*+ve) typically presents at a younger age, is more poorly differentiated and aggressive, and is associated with worse survival outcomes than non-*BRCA2* associated PC. Evidence for the more adverse clinical presentation comes from multiple studies. Carter et al³⁰ provided evidence that germline *BRCA2* carriers are more likely to be reclassified from Gleason score (GS) 3+3 at diagnosis to GS $\geq 4+3$ (4.1% (*BRCA2*+ve) vs 0.7% (*BRCA2*-ve), ($p=0.01$)) versus GS 3+4 (2.1% (*BRCA2*+ve) vs 0.6% (*BRCA2*-ve), ($p=0.03$)). They indicated that *BRCA2* carriers have more unfavourable prognostic indicators and a greater chance of progression from localised PC to metastatic castration-resistant PC (mCRPC) (which has a median survival rate 18–36 months).

A systematic review by Gleicher et al³¹ compared PC characteristics in both carriers of the *BRCA2* germline variant (*BRCA2*+ve) and non-*BRCA2* carriers (*BRCA2*-ve). It was found that *BRCA2* carriers had higher prostate-specific antigen (PSA) values at diagnosis ($p<0.001$), a higher proportion of PCs with a Gleason score >7 (64% (*BRCA2*+ve) vs 49% (*BRCA2*-ve), ($p<0.001$)). In addition, a higher proportion of *BRCA2*+ve tumours have T3/T4 staging (41% (*BRCA2*+ve) vs 29% (*BRCA2*-ve), ($p<0.001$)) and higher rates of metastatic disease at diagnosis (26% (*BRCA2*+ve) vs 8% (*BRCA2*-ve), ($p<0.001$)). A study by Wokolorczyk et al³² found that PC with Gleason scores >8 were more likely in men carrying germline *BRCA2* variants than in non-carriers (75% (*BRCA2*+ve) vs 22% (*BRCA2*-ve), ($p<0.05$)). This group stipulated that *BRCA2* carriers had a lower mean age than non-*BRCA2* carriers at the time of their PC diagnosis (57 years (*BRCA2*+ve) vs 61 years (*BRCA2*-ve), $p=0.3$). In a study by Na et al,³³ 6.07% of men with lethal PC were later identified as carriers of pathogenic variants in either *BRCA1/2* or *ATM* compared to 1.44% of men with localised PC who were carriers of these variants ($p=0.0007$).

The treatment pathway is therefore more challenging in these patients. More potent therapeutic agents are often considered rather than more conservative measures with active surveillance.^{7,8,11,28,34–36}

Therapeutic Options in *BRCA2* Deficient Prostate Cancer

Whilst a diagnosis of *BRCA2* deficient PC has unfavourable prognostic implications, the understanding of the

underlying molecular defect leading to oncogenesis enables a precision medicine approach targeting the DDR deficient (DDRd) cancer cells specifically. Poly-ADP ribose polymerase (PARP) inhibitors and platinum-based chemotherapeutic agents have been identified as particularly valuable treatment options for cancers brought about from either somatic or germline variants in DDR genes such as *BRCA2* and may become standard of care for *BRCA2* deficient PC patients in the near future.^{3,17,19,37–39}

PARP Inhibitors

One of the main driving forces for oncogenesis in *BRCA1/2* deficient cells relates to the disruption of the HR DNA repair pathway as a result of biallelic *BRCA*-loss. In cells lacking functional *BRCA* proteins, HR is impaired, forcing the cell to utilise alternative DDR mechanisms such as (i) non-homologous end joining (NHEJ) which is less accurate than HR and thus susceptible to errors and (ii) single strand (ss) DNA break repair, which is mediated by the PARP enzymes, PARP1 and PARP2. Increased dependency on PARP-mediated DDR pathways makes *BRCA1/2* deficient cells suitable targets for PARP inhibitor therapy, which would disable PARP1 and PARP2. This forces the *BRCA* deficient cells to try to mend DNA damage in the double strand (ds) DNA pathway, which they are unable to do due to the *BRCA* deficiency. Accumulation of these ds-DNA breaks increases cell stress, leading to cell cycle arrest and cell death via apoptosis. This process is known as synthetic lethality.^{40–43}

PARP inhibitors were first investigated as a novel treatment approach for *BRCA* deficient tumours in 2005 and have since proven to be of clinical benefit, with pharmaceuticals such as rucaparib and olaparib being approved for mCRPC treatment in the US following the results of clinical trials such as TOPARP-A, TOPARP-B, PROfound and TRITON (Table 1).^{17,41,44–49} Whilst PARP inhibitor use has been documented as a beneficial treatment for *BRCA* deficient mCRPC patients, resistance to these drugs is not uncommon and may occur via the acquisition of somatic *BRCA1/2* variants in the germline allele which restore the reading frame (ie *BRCA* reversion variants) and thus the normal functioning of the *BRCA* protein.^{50–56} Case reports by Carneiro et al⁵⁷ and Ma et al⁵⁸ have identified multiple reversion variants in *BRCA2* deficient mCRPC patients who developed resistance to olaparib treatment. Ma et al⁵⁸ postulated that the efficacy of PARP inhibitor therapy may be reliant on the proportion of DDR variants in relation to other variants present within

Table 1 Summary of Mentioned Poly-ADP Ribose Polymerase (PARP) Inhibitor Clinical Trials Involving Metastatic Castration-Resistant Prostate Cancer Patients

Clinical Trial (Trial No.)	Trial Description	Patient Characteristics	Treatment Given	Primary Trial Endpoint	Main Findings
TOPARP-A (NCT01682772)	Phase II trial	50 mCRPC patients with unknown carrier status for pathogenic DDR gene variants	Olaparib 400mg twice daily	Treatment response rate	NGS identified 16 patients with deleterious DDR gene variants. 14/16 (88%) of this subgroup had a positive treatment response, including all 7 patients with BRCA2-deficient tumours.
TOPARP-B (NCT01682772)	Randomised phase II trial	98 progressing mCRPC patients carrying with pathogenic DDR gene variants	Olaparib 400mg or 300mg twice daily	Treatment response rate	A positive treatment response observed in 54.3% of patients randomised to receive 400mg olaparib compared to 39.1% of patients randomised to receive 300mg olaparib. The most pronounced treatment effect was observed in the BRCA1/2 carrier subgroup, with an overall response rate of 83%.
PROfound (NCT02987543)	Randomised Phase III trial	387 mCRPC patients carrying germline pathogenic variants in BRCA1/2 or ATM (cohort A) or 12 other known DDR genes (cohort B)	Olaparib 300mg twice daily or SOC treatment (enzalutamide or abiraterone) in a 2:1 ratio	Imaging-based PFS	In both study cohorts overall, imaging-based PFS was significantly extended in those on olaparib treatment compared to those receiving SOC (5.8 months vs 3.5 months; hazard ratio for progression or death, 0.49; 95% CI 0.38–0.63; $p<0.001$). There was a more pronounced treatment effect observed in cohort A (7.4 months vs 3.6 months, hazard ratio, 0.34; 95% CI 0.25–0.47; $p<0.001$), with hazard ratios of 0.41 and 0.21 calculated for BRCA1 and BRCA2 patients respectively.
TRITON2 (NCT02952534)	Ongoing phase II trial	115 mCRPC patients carrying pathogenic DDR gene variants whose tumours had progressed on 1–2 lines of AR-directed therapy and 1 line of taxane-based chemotherapy	Rucaparib 600mg twice daily	Objective response rate, PSA response rate	Improvements in radiographical and PSA results were demonstrated in the BRCA-deficient mCRPC patient subgroup. Response rates to rucaparib therapy were similar in both patients with somatic BRCA variants and those with germline BRCA variants.
TRITON3 (NCT02975934)	Ongoing phase III trial	mCRPC patients carrying pathogenic variants in BRCA1/2 or ATM whose tumours progressed on one prior AR-directed therapy	Rucaparib 600mg twice daily vs SOC treatment (physician's choice of second-line AR-directed therapy or docetaxel in chemotherapy-naïve patients)	Radiographic PFS	

Abbreviations: mCRPC metastatic castration-resistant prostate cancer; NGS, next generation sequencing; DDR, DNA damage repair; PFS, progression free survival; SOC, standard of care; PSA, prostate specific antigen; AR, androgen receptor.

the mutational landscape of the tumour. These reports support the utility of circulating tumour DNA (ctDNA) analysis monitoring to predict PARP inhibitor resistance in mCRPC patients.

Platinum-Based Chemotherapy

In recent years platinum-based chemotherapy has shown efficacy in the management of mCRPC associated with pathogenic DDR gene variants including *BRCA2*.^{37–39} Platinum-based chemotherapeutic agents work by generating cross-links between purine bases in DNA which disrupts DNA repair, causing cell death via apoptosis.⁵⁹

Platinum-based chemotherapy is not routinely used to treat mCRPC based on evidence from trials which did not show overall survival benefit from standard of care.⁶⁰ However, in studies specifically selecting mCRPC patients with pathogenic variants in recognised DDR genes,^{37–39,61,62} platinum-based chemotherapy was found to ameliorate patient quality of life by decelerating tumour progression (Table 2). Reports by Mota et al⁶³ and Simmons et al⁶⁴ (Table 2) highlight the challenge of tumour resistance which is especially important in *BRCA*-associated PCs which have already been recognised as difficult to treat.

Identification of *BRCA2*-Deficient Prostate Cancer

For our patient in Figure 1, it is important to diagnose that he has *BRCA2* deficient PC not only for his immediate therapeutic management, but also to consider any possible heritable predisposition to cancer which may require wider cascade screening in the family. Evidence suggesting a *BRCA2* deficient PC can be acquired from multiple sources.

Tumour Phenotype and Somatic Pathogenic Variants

As previously discussed, *BRCA2* deficient PC often presents with poorer prognostic indicators than *BRCA* proficient PC, which may trigger suspicion in the treating clinician.^{11,18} Identification of specific biallelic pathogenic variants in *BRCA2* can be undertaken by next-generation sequencing (NGS) of tumour-derived DNA. From February 2021, the UK National Genomic Test Directory⁶⁵ specified that somatic gene sequencing of *BRCA1* and *BRCA2* via NGS panel approaches should be undertaken for all PCs, with the addition of *ATM/CDK12* for metastatic resistant PC cases. This standard of care pathway should now be implemented

by all seven Genomic Laboratory Hubs (GLHs) in the UK, reducing inequities in testing pathways which have previously existed in the UK, and leading to the identification of more *BRCA2* deficient PC. The identification of a somatic pathogenic variant in the *BRCA1/2* genes in PC should then trigger germline testing for inherited susceptibility according to the European Society of Medical Oncology (ESMO) guidelines.⁶⁶ Somatic small panel testing is likely to be expanded to a wider number of DDR-related genes through gene panel testing, with likely expansion of tumour testing to whole genomes in the future. Whole genome somatic NGS analysis enables identification of driver mutations driving oncogenesis, but in addition enables the identification of mutational signatures, and tumour mutational burden, which can provide evidence for DDR driving oncogenesis and precision medicine approaches.⁶⁷

Identification of *BRCA2*-Related Germline Susceptibility to Prostate Cancer

Family history is a widely established PC risk factor, with some families displaying significant aggregation of PC cases; a man with a first-degree relative affected by the disease has at least twice the chance of developing this condition compared with the general population. Current access to *BRCA* germline genetic testing for PC patients in the UK relies on them meeting National Genomic Test Directory eligibility criteria R208 (Hereditary Breast and Ovarian Cancer) in which family history is an essential component.⁶⁸ Given the primary focus on breast and ovarian cancer in the eligibility criteria and the fact that systematic family history assessments for those diagnosed with PC is not well embedded within clinical pathways, men presenting with PC are likely to be underserved with respect to germline genetic testing. More robust educational and training is required in secondary care to ensure documentation and collection of family history in men with PC to ensure eligible men, such as our proband from Figure 1, are identified as having likely genetic susceptibility requiring germline testing.

Historically, germline genetic testing of PC patients typically involves predictive testing of a known gene in patients with a previously identified familial variant increasing cancer risk, or, diagnostic genetic testing of *BRCA1/2* in the context of a significant family history of breast, ovarian and prostate cancer. In our family in Figure 1, diagnostic *BRCA* testing in such pedigrees has often been instigated by

Table 2 Summary of Mentioned Platinum-Based Chemotherapy Reports Involving Metastatic Castration-Resistant Prostate Cancer Patients

Publication Authors	Publication Type	Patient Characteristics	Treatment Given	Primary Study Endpoint	Main Findings
Pomerantz et al ³⁹	Retrospective cohort study	141 mCRPC patients who were subdivided into carriers and non-carriers of pathogenic germline <i>BRCA2</i> variants	≥ 2 doses of carboplatin and docetaxel	Treatment response rate	6/8 (75%) of <i>BRCA2</i> carriers exhibited a positive treatment response compared with 23/133 (17%) of non-carriers ($p < 0.001$).
Zafeiriou et al ³⁸	Case report	3 metastatic PC patients, 1 of whom was identified as a germline carrier of a pathogenic <i>BRCA2</i> variant	<i>BRCA2</i> carrier received carboplatin and docetaxel following tumour progression on other therapies	Treatment response	44% PSA reduction was observed along with signs of disease regression on MRI following 6 cycles of treatment in the <i>BRCA2</i> carrier.
Cheng et al ³⁷	Case report	3 mCRPC patients with biallelic <i>BRCA2</i> loss who had progressed on previous therapy	Carboplatin and docetaxel (patient 1), carboplatin and doxorubicin (patient 2), carboplatin and docetaxel (patient 3)	Treatment response	PSA reductions were observed in all three patients. A positive clinical response was noted in patient 2 and significant radiographical response noted in patient 3.
Slootbeek et al ⁶¹	Retrospective cohort study	71 mCRPC patients, of which 14 carried pathogenic <i>DDR</i> gene variants (including 7 <i>BRCA2</i> carriers)	Carboplatin	PSA response, radiographical response	≥ 50% PSA reduction and objective radiographical response observed in all patients with <i>BRCA2</i> deficient tumours. Median overall survival was significantly longer in <i>BRCA2</i> deficient patients compared with patients carrying <i>BRCA2</i> wild type (21 months vs 7 months, $p = 0.041$).
Schmid et al ⁶²	Retrospective cohort study	508 mCRPC patients: 80 <i>DDR</i> gene variant carriers (cohort 1), 98 non-carriers of <i>DDR</i> gene variants (cohort 2) and 300 patients of unknown <i>DDR</i> gene variant carrier status (cohort 3)	Platinum monotherapy (carboplatin/cisplatin) or platinum combination therapy (carboplatin/cisplatin/oxaliplatin + docetaxel/etoposide/paclitaxel)	PSA response, radiographical response	The most profound treatment response was observed in the <i>BRCA2</i> carrier patient subgroup, of which 23/44 (63.9%) had ≥ 50% reduction in PSA value after treatment and soft tissue responses were noted in 17/44 (38.6%).
Mota et al ⁶³	Prospective cohort study	109 mCRPC patients	Carboplatin/cisplatin or carboplatin/cisplatin + taxane chemotherapy/etoposide	Association between <i>DDR</i> variants and response to platinum chemotherapy, response to platinum chemotherapy in patients who had progressed on PARP inhibitor therapy	Patients with <i>DDR</i> variants were more likely to have a ≥ 50% reduction in PSA following platinum-based chemotherapy (odds ratio, 7.0; 95% CI 1.9–29.2). Treatment response to platinum-based chemotherapy was only seen in 37% <i>DDR</i> gene variant carriers who had progressed on previous PARP inhibitor therapy.

Simmons et al ⁶⁴	Case report	1 mCRPC patient with a germline <i>BRCA2</i> variant	Sequential treatment with carboplatin followed by rucaparib	NGS results following provision of treatment	12 different somatic reversion variants (thought to confer resistance against carboplatin by restoring the C-terminal end of <i>BRCA2</i>) were discovered using NGS after 4 months on carboplatin chemotherapy. As a result of these reversion mutations, rucaparib had little benefit in slowing tumour progression.
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Abbreviations: mCRPC, metastatic castration-resistant prostate cancer; PC, prostate cancer; PSA, prostate specific antigen; MRI, magnetic resonance imaging; DDR, DNA damage repair; PARP, poly-ADP ribose polymerase; NGS, next generation sequencing.

a female presenting with breast or ovarian cancer rather than a male presenting with PC. To widen our ability to identify germline susceptibility to PC, diagnostic gene panel testing in PC patients of PC-associated DDR genes (cancer susceptibility genes) where a family history suggestive of inherited predisposition is absent will detect additional germline gene carriers.¹⁰

The detection of germline variants can (i) inform choice of treatment, (ii) facilitate genetic testing of at-risk family members (cascade screening), and (iii) help to determine the risk of developing other cancer types associated with the genes where variants were identified.^{9,69} It is therefore important that novel strategies to improve identification of carriers are being implemented.

In our family in Figure 1, the brother of our proband has a 50% (1 in 2) chance of also carrying the germline *BRCA2* pathogenic variant identified in his brother. He should be offered a predictive genetic test via clinical genetics services to establish his own risk. In this case, predictive genetic testing identified that the brother also carries the *BRCA2* pathogenic variant. This should now facilitate entry of the unaffected brother into Screening, Prevention and Early Detection (SPED) pathways. The sister of the proband with breast cancer is likely to also carry the *BRCA2* pathogenic variant but should also be offered germline predictive testing to clarify. If shown to carry the variant she will need discussions about considering risk reducing bilateral salpingo-oophorectomy to reduce her risk of ovarian cancer. The *BRCA2* pathogenic variant is likely to have been inherited from the deceased mother who died of ovarian cancer, but cascade testing to the maternal uncles and cousins should be initiated to identify all at risk individuals. The father could be offered predictive testing if no maternal relative is identified with the *BRCA2* pathogenic variant to ensure he is not a carrier, as occasionally, phenocopies of cancer diagnoses can mask the side of the family from which the variant was inherited.

Barriers to access to SPED for unaffected male *BRCA2* carriers exist across healthcare systems. There is a lack of awareness and understanding among male *BRCA*-carriers about their increased risk of developing *BRCA*-associated cancers, which include PC, pancreatic cancer and male breast cancer, and this can have a detrimental effect on health outcomes.⁷⁰ Contributing factors to misinformation among male *BRCA*-carriers include (i) limited availability of relevant information and resources pertaining to genetic

risk of *BRCA*-associated cancers such as PC and male breast cancer,^{71–73} and (ii) lack of awareness among healthcare personnel about men's inherited cancer predisposition and available treatment options.⁷⁴ *BRCA1/2* carriers are labelled as persons with “Hereditary Breast and Ovarian Cancer syndrome”⁷⁵ which is misleading because *BRCA1/2* variants increase the risk of other cancers like PC, pancreatic cancer and melanoma,^{76,77} but it also feminizes the condition^{73,78} leading to stigma within the male population.

Improved education and training around identification of genetic susceptibility to PC, more streamlined access to family history assessment, somatic and germline testing for PC patients will all increase identification of individuals with inherited susceptibility to PC. *BRCA*-risk information specifically adapted for the male population will also aid the promotion of more proactive health behaviours which in turn enable earlier detection of *BRCA*-associated disease in male *BRCA2* carriers.⁷⁰

Prostate Cancer Screening in Unaffected *BRCA2* Mutation Carriers

Consider now the brother of our proband from Figure 1. We have identified he has an increased likelihood of developing *BRCA2*-related PC, and he will want to know how this risk may be managed.

Whilst measuring serum concentrations of PSA as a means of screening for PC has become common practice,⁷⁹ there are no universally accepted PC screening guidelines for men.⁸⁰ Screening the general population for PC using PSA testing is not currently recommended based on the results of previous research,^{81–85} which highlights the low specificity of the test.

Data from IMPACT study,^{80,86} an international multi-centre PC screening study investigating the utilisation of PSA testing in *BRCA1/2* carrier males, have suggested that PSA testing in these men (who are at increased risk of PC) may be of greater clinical benefit than screening the general population. Following the completion of the first IMPACT study screening round,⁸⁰ data showed that in *BRCA2* carriers, the positive predictive value (PPV) of biopsy using a PSA threshold of 3.0ng/mL was 48%, compared with a lower 33.3% PPV seen in the control group. In addition, there was a higher frequency of intermediate-high risk PC detected in *BRCA2* carriers with PSA values above the 3.0ng/mL threshold for further

investigation with biopsy. It was also noted the males diagnosed with PC below the age of 50 years were all *BRCA1/2* carriers, and *BRCA2* carriers reaching the 3.0ng/mL PSA threshold were significantly younger than their counterparts with elevated PSA values. After 3 more years of follow-up (4 screening rounds), data analysis reiterated the utility of PSA screening in *BRCA1/2* carriers described previously. When compared with non-*BRCA2* carriers, *BRCA2* carriers had a significantly higher PPV of PSA >3.0 ng/mL (31% vs 18%; $p=0.025$), a higher PC incidence rate per 1000 person years (19.4 vs 12.0; $p=0.03$), a significantly higher frequency of intermediate-high risk PC ($p=0.011$) and were significantly younger at the time of PC diagnosis ($p=0.044$), with the youngest age of onset being recorded in a 41-year-old *BRCA2* carrier. Overall, the results so far from the ongoing IMPACT study further substantiate the earlier age of onset and more aggressive PC phenotype seen in *BRCA2* carriers which has been widely reported throughout the literature,^{11,31–34,36,87} and also highlights the efficacy of using PSA testing in this patient subgroup.

In another prospective PC screening study, Segal et al⁸⁸ sought to investigate the use of PSA testing and multi-parametric magnetic resonance imaging (mpMRI) in 188 male *BRCA1/2* carriers. Whilst MRI is widely used for breast cancer screening in female *BRCA1/2* carriers,⁸⁹ this study was the first to investigate the use of this imaging modality for PC screening. Prostate biopsies were offered to patients with either an elevated PSA value (PSA >3 ng/mL or elevated age-stratified PSA) and/or a suspicious prostatic lesion on mpMRI (PI-RADS ≥ 3). Of these patients, 92 men (85%) underwent a prostate biopsy and PC was detected in 16 men (8.5%). Amongst the 92 men having a prostate biopsy, 5 different PC screening strategies making use of either (i) PSA testing alone, (ii) mpMRI alone or (iii) both PSA testing and mpMRI were compared using decision curve analysis. Overall, using mpMRI in isolation proved to be the best screening tool, picking up 15/16 cancers. Interestingly, in *BRCA1/2* carriers under the age of 55 years, using mpMRI irrespective of PSA value had the most benefit whilst PSA screening on its own had zero net benefit. In men older than 55 years, triaging with PSA testing before offering mpMRI had the most benefit.

Based on the results of the IMPACT study^{80,86} and those reported by Segal et al,⁸⁸ national and/or international guidelines for unaffected male *BRCA2* carriers, such as the brother of our proband in Figure 1, are likely to be

published in the near future. This is likely to specify PSA testing for male *BRCA2* carriers from age 40, with subsequent investigations required for individuals with PSA ≥ 3 ng/mL.

Conclusion

In this review article, we have discussed the investigation and management of *BRCA2* deficient PC using a case study of a family with a pathogenic germline *BRCA2* variant. We have outlined how *BRCA2* deficient PC in the proband might present, and management options which may have specific relevance for *BRCA2* deficient PC, such as PARP inhibitors and platinum-based chemotherapy. We have also highlighted the need to distinguish between somatic and germline *BRCA2* pathogenic variants which may have relevance for cancer predisposition in other family members, and the screening which can be offered to unaffected male *BRCA2* carriers. The inclusion of all these facets to *BRCA2* deficient PC investigation and management in our review provides a comprehensive overview of the clinical approach to a family in which *BRCA2* deficient PC is diagnosed. The rapid advances in our understanding of the molecular pathogenesis of DDRd PC and new treatment paradigms, alongside the widened access to both somatic and germline genetic analysis for PC patients and their families will hopefully enable the robust implementation of high-quality evidence based clinical pathways for both the management and identification of *BRCA* deficient PC and improved screening, early detection and prevention strategies for individuals at increased genetic risk of PC.

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