Germline testing is becoming increasingly relevant in prostate cancer (PCa) screening, prognosis, and management. A subset of patients with PCa harbor pathogenic/likely pathogenic variants (P/LPVs) in genes mediating DNA-repair processes, and these P/LPVs have implications for cancer screening, treatment, and cascade testing. As a result, it is recommended that all men with high-risk localized and metastatic PCa undergo routine germline testing. As more PCa patients undergo germline testing, it is important that clinicians and genetics experts recognize current disparities in germline testing rates among racial/ethnic minorities in the United States. The reasons for these disparities are multiple and require similarly manifold consideration to close the germline testing gap and reduce inequities in PCa screening, management, and treatment.

INTRODUCTION
Prostate cancer (PCa) is the most common cancer among men and one of the leading causes of deaths worldwide [1]. In 2021, it is estimated that 248,530 men in the United States (US) will be diagnosed with PCa and 34,130 individuals will die secondary to this disease [2]. While age, race/ethnicity, and family history are established risk factors for PCa, it is now recognized that a proportion of PCa susceptibility is attributed to genetic predisposition. Advances in molecular sequencing technologies have identified several PCa susceptibility genes, many related to known hereditary cancer syndromes, including hereditary breast and ovarian cancer (HBOC) syndrome (BRCA1, BRCA2, ATM, CHEK2, and PALB2) and Lynch syndrome (MLH1, MSH2, MSH6, and PMS2) [3]. As a result of these findings, recommendations for germline testing based on clinical features and family history have expanded. The identification of pathogenic/likely pathogenic variants (P/LPVs) in PCa predisposition genes may help inform cancer screening strategies for patients and family members, treatment options in the metastatic setting, and clinical trial enrollment.

As germline testing becomes more clinically relevant and widely available, it is important to recognize the risk of exacerbating health disparities among racial/ethnic minorities with PCa and develop systematic strategies to bridge disparities in germline testing. Reasons for these disparities are multifaceted and include patient, clinician, and system factors. Additionally, current PCa clinical trials and genetic studies do not reflect the diverse populations of individuals at-risk or suffering from this disease. In this review, we discuss the indications for germline testing in men with PCa, barriers to germline testing in diverse populations, and potential strategies to bridge the disparities gap with the expansion of germline testing for men with PCa.

DISPARITIES IN OUTCOMES OF MEN WITH PCA
There are documented disparities in the incidence, treatment, and mortality of PCa between Black and non-Black men [2, 4–6]. Notably, Black men are diagnosed with PCa at nearly twice the rate of non-Hispanic white (NHW) men [2], and Black men with local/regional PCa have been found to be less likely to receive treatment with curative intent than NHW men [7]. Further, the PCa mortality rate is twice as high in Black men compared to NHW men [2]. The National Cancer Institute estimates that Black men have a 4.72% lifetime risk of dying of PCa compared to a 2.86% risk among NHW men [5]. Although biological differences may account for a portion of the disparity in overall PCa survival, it has been suggested that improved access to care, including screening, follow-up, and therapy may be effective in reducing this disparity [8]. It is important to note that one limitation of studies of PCa incidence and mortality is that most data on Black men does not stratify them by country/region of origin—Black men are not a homogenous group and there may be differences in PCa incidence and mortality for Caribbean, African, and African-American men [9].

Similarly, despite the genetic and cultural diversity of Hispanic men in the US, individual subgroups are typically combined. Notably, significant heterogeneity has been observed among Hispanic men with PCa [10]. Overall, PCa occurs less often in Hispanic men than in NHW men [1]. However, Mexican-American men have been found to have more advanced stage PCa at diagnosis [11] and are significantly more likely to have aggressive PCa following radical prostatectomy [12]. While prostate cancerspecific mortality (PCSM) is comparable between Hispanic and NHW men, Puerto Rican men have been shown to have significantly higher PCSM than NHW men, Black men, and all...
other Hispanic subgroups [10]. Ultimately, the dearth of PCa studies examining individual Hispanic subgroups makes it difficult to compare them to NHW men.

Despite these documented disparities, African-American/Canadian, Asian/Pacific Islander, and Hispanic populations are typically underrepresented in germline testing, clinical trials (Table 1), and study cohorts [13]. One study analyzed 72 global phase III and IV prevention, screening, and treatment PCa clinical trials between 1987 and 2016: 59 trials reported race/ethnicity data, and 96% of patients enrolled in these studies were NHW men. African and Caribbean medical centers were particularly underrepresented in these trials [14]. Concordant studies have shown that the majority of PCa patients receiving germline testing are NHW men [15, 16], with as high as 95% being English-speaking men [16]. Underrepresentation of racial/ethnic minorities in germline testing is not unique to PCa and exists among patients with various other malignancies [17–19].

Racial/ethnic minority populations in the US are expected to grow rapidly over the coming decades, underscoring the need to address and resolve these disparities. It is projected that by 2045, NHW people will make up <50% of the total US population. While NHW populations are expected to decline, all other racial/ethnic minority populations are expected to grow; in particular, Hispanic populations are the fastest growing demographic and are expected to comprise 24.6% of the US population in 2045, up from 18.7% in 2020 [20].

Table 1. Representation of diverse racial/ethnic groups in prostate cancer clinical trials involving germline testing.

| Trial          | Race  | Ethnicity (Hispanic or Latino) |
|---------------|-------|-------------------------------|
|               | N     | White | Black | Asian | Other/Unknown | No | Yes | Unknown |
| METASTATIC HORMONE SENSITIVE PROSTATE CANCER |       |       |       |       |               |    |     |         |
| CHARTR2—Docetaxel [72] | 790   | 674   | 76    | NR    | 40           | NR | NR | NR     |
| STAMPEDE—Docetaxel [73] | 2962  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| LATITUDE—Abiraterone [74] | 1199  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| STAMPEDE—Abiraterone [75] | 1917  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| TITAN—Alapatreumide [76] | 1052  | 719   | 19    | 229   | 85           | NR | NR | NR     |
| ARCHES—Enzalutamide [77] | 1150  | 926   | 16    | 155   | 53           | NR | NR | NR     |
| ENZAMET—Enzalutamide [78] | 1125  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| MCRPC         |       |       |       |       |               |    |     |         |
| TAX 327—Docetaxel [79] | 1006  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| TROPIC—Cabazitaxel [80] | 755   | 631   | 40    | 58    | 26           | NR | NR | NR     |
| IMPACT—Sipuleucel-T [81] | 512   | 461   | 30    | NR    | 21           | NR | NR | NR     |
| ALSYMPCA—Radium-223 [82] | 921   | 865   | NR    | NR    | NR           | NR | NR | NR     |
| COU-AA-301—Abiraterone post-chemo [83] | 1195  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| COU-AA-302—Abiraterone pre-chemo [84] | 1088  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| AFFIRM—Enzalutamide post-chemo [85] | 1199  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| PREVAIL—Enzalutamide pre-chemo [86] | 1717  | 1324  | 34    | 167   | 192          | 1527| 38  | 152    |
| NMCPRC        |       |       |       |       |               |    |     |         |
| SPARTAN—Paladatamide [87] | 1200  | 800   | 68    | 140   | 192          | NR | NR | NR     |
| PROSPER—Enzalutamide [88] | 1401  | NR    | NR    | NR    | NR           | NR | NR | NR     |
| ARAMIS—Darolutamide [89] | 1509  | NR    | NR    | NR    | NR           | NR | NR | NR     |

MCRPC metastatic castration-resistant prostate cancer, NMCPRC non-metastatic castration-resistant prostate cancer, NR not reported.

GENOMICS OF NON-WHITE MEN WITH PCA

P/LPVs have been found to be prevalent in men with PCa. A 2016 multi-institutional study found that the incidence of germline mutations in genes mediating DNA-repair processes (including but not limited to BRCA2, ATM, CHEK2, BRCA1, RAD51D, and PALB2) among 692 men with metastatic PCa was 11.8% [3]. For patients with localized disease, the prevalence of P/LPVs ranged from 2 to 6%, with increased prevalence in men with higher Gleason scores and higher-risk PCa. Notably, S76 (83%) of the men in this study were NHW men. Additionally, a 2021 study found that 9.5% of PCa patients with high-risk localized disease had P/LPVs, most frequently in BRCA2 and ATM [21].

In order to better understand the genomic landscape of racial/ethnic minorities, there is a need to more extensively examine P/LPV rates in non-white men with PCa. More recent studies have found that the prevalence of P/LPVs varies across racial/ethnic groups. When compared to NHW men, Hispanic men with PCa have been found to have similar rates of P/LPVs in the BRCA2 gene than their NHW counterparts [23]. Among patients with metastatic PCa, mutations in DNA-repair genes have been found to occur more often in Black men than in NHW men [24]. However, these studies were limited by small sample size.

The lack of diversity in germline testing cohorts is thought to be a contributor to higher rates of variants of uncertain significance (VUS) in racial/ethnic minorities [15, 25]. Notably, African-American, Hispanic, and Asian/Pacific Islander PCa patients have been found to be more likely to have a VUS than those with European ancestry [23, 26]. In one study of PCa patients referred to Color Genomics for germline testing, VUS rates in HBOC and Lynch syndrome genes were 21% in NHW men, while 26.6% and 33.3% in African-American/Canadian and Asian/Pacific Islander men, respectively [15]. Increasing the proportion of underrepresented groups in germline testing cohorts is predicted to result in the reclassification of VUS, which will assist in cancer risk stratification and targeted therapy strategies [15].
INDICATIONS FOR GERMLINE TESTING AND IMPLICATIONS OF TESTING RESULTS

Recent studies on the incidence of P/LPVs among men with PCa have resulted in updated guidance regarding which patients should receive germline testing. The most recent iteration of the National Comprehensive Cancer Network guidelines for PCa recommends germline testing for all men with high-risk localized and metastatic PCa, Ashkenazi Jewish ancestry, a family history of high-risk germline mutations, or a positive family history of cancer [27, 28]. Given emerging data on the association between intraductal/cribriform and ductal histologies and P/LPVs, testing is considered for men with these histologic subtypes [28]. Other professional societies and expert panels have also provided recommendations for germline testing for men with PCa, largely based on evidence synthesis, consensus agreement, and expert opinion (Table 2).

Expanding germline testing uptake may help clinicians predict outcomes in men with PCa by detecting ethnicity-dependent biomarkers and mutations that drive aggressive tumor biology [29]. Germline mutations in DNA-repair genes, particularly BRCA1/2 and ATM, are associated with aggressive PCa and significantly shorter survival time: mutation carriers have been found to have a higher proportion of Gleason Score ≥7 (71%) than noncarriers (31%) and mutation frequency has been found to be significantly higher in patients that have died of PCa than in localized PCa patients [30].

Germline testing also has implications regarding candidacy of select treatments, including platinum chemotherapy, poly(ADP-ribose) polymerase (PARP) inhibitors, and checkpoint inhibition for patients with metastatic castration-resistant prostate cancer (mCRPC) [31]. In 2020, the PARP inhibitor Olaparib was FDA approved for the treatment of adult patients with deleterious/suspected deleterious germline or somatic homologous recombination repair (HRR) gene-mutated mCRPC who have progressed following prior treatment with enzalutamide or abiraterone. Olaparib is FDA approved for a panel of 14 genes, including BRCA1/2. Of the patients in the PROfound trial, 69% were white, 29% were Asian, and 1% were Black [32]. Another therapy, Rucaparib, was FDA approved in 2020 for the treatment of adult patients with a deleterious BRCA1/2 mutation (germline and/or somatic)-associated mCRPC who have been treated with androgen receptor-directed therapy and a taxane-based chemotherapy. For the 115 patients enrolled in the TRITON2 study, the majority were white (73%) and 10% were Black; other racial/ethnic groups were not specified [33]. Pembrolizumab has also been FDA approved for patients with refractory metastatic cancers with MSI-high or MMR deficiency (dMMR) status based on tumor assessment that had progressed following prior treatments [34]. Pembrolizumab has shown antitumor activity with an acceptable safety profile in an unselected subset of patients with mCRPC [35].

Another indication for germline testing is cascade testing, which refers to germline testing among relatives of patients with cancer-associated P/LPVs; it has historically had decreased uptake in the community at around 30% or less. Being a PCa patient with a germline P/LPV in a DNA-repair (HRR) gene-mutated mCRPC who have progressed following prior treatments [34]. Pembrolizumab has shown antitumor activity with an acceptable safety profile in an unselected subset of patients with mCRPC [35].

Table 2. Guidelines on germline testing in prostate cancer.

| Source | Regional (N1)/metastatic prostate cancer | NCCN very high and high-risk localized prostate cancer | NCCN intermediate/low/very low risk localized prostate cancer |
|--------|-----------------------------------------|-----------------------------------------------------|-------------------------------------------------------------|
| National Comprehensive Cancer Network Version 1.2022 [27, 28] | Recommend | Recommend | Recommend if a family history of: \* Ashkenazi Jewish ancestry \* High-risk germline mutations (e.g., BRCA1/2, Lynch mutation) \* PCa in brother/father/multiple family members diagnosed with PCa (not GG1) at ≤60 years of age or who died from PCa \* ≥3 cancers on same side of family, especially diagnoses ≤50 years of age: bile duct, breast, colorectal, endometrial, gastric, kidney, melanoma, ovarian, pancreatic, prostate (not GG1), small bowel, or urothelial cancer Consider for: Intraductal/cribriform histology |
| Philadelphia Prostate Cancer Consensus Meeting Publication 2019 [90] | Recommend | • Consider for T3a or higher. • Consider for intraductal/ductal pathology. • Consider for Gleason 4 (Gleason & sum) or above. • Consider for Ashkenazi Jewish ancestry. • Consider for family history of two or more cancers in HBOC/Lynch spectrum in any relatives on the same side of the family (especially if diagnosed at age <50 years). • Recommend for family history of one brother/father/two or more male relatives with one of the following: • PCa at age <60 years • Died of PCa. • Metastatic PCa. |
| AUA/ASTRO/SUO 2017 and 2021 [91–93] | Recommend | Recommend if a strong family history of specific cancers (e.g., breast, ovarian, pancreatic, other gastrointestinal tumors, and lymphoma). Not recommended |

PCa Prostate Cancer, HBOC Hereditary Breast and Ovarian Cancer Syndrome, NCCN National Comprehensive Network, AUA American Urological Association, ASTRO American Society of Radiation Oncology, SUO Society of Urologic Oncology, GG Grade Group.
Table 3. Challenges and solutions.

| Challenges                                      | Solutions                                                                 |
|------------------------------------------------|---------------------------------------------------------------------------|
| Shortage of CGCs and limitations of current GC models | • Increase clinical training capacity for CGCs  
|                                                  | • Offer pretest GC and select posttest GC via alternative methods (telemedicine, group GC)  
|                                                  | • Automate risk assessment                                                  |
| Differences in the quality of care for minority patients | • Increase access to genetic services, contain costs, and address provider implicit bias  |
| Medical mistrust                                   | • System wide interventions to address gaps in healthcare delivery  
|                                                  | • Increase representation of minorities in healthcare  
|                                                  | • Community outreach                                                      |
| Lack of knowledge regarding testing                | • Increase genetics education among patients and community health providers  
|                                                  | • Culturally tailored genetic counseling                                   |
| Prohibitive cost and lack of insurance coverage for germline testing | • Increase payer coverage  
|                                                  | • Low-cost testing and government subsidies                                 |
| Understudied link between PCa and breast/ovarian cancers | • Address similar disparities in germline testing among women with HBOC syndrome  
|                                                  | • Physician and patient directed education regarding genetic link between PCa and HBOC syndrome |

CGC Clinical Genetic Counselor, GC Genetic Counseling, PCa Prostate Cancer, HBOC syndrome Hereditary Breast and Ovarian Cancer Syndrome.

REASONS FOR THE DISPARITIES IN GERMLINE TESTING AMONG RACIAL/ETHNIC MINORITIES WITH PCA AND POTENTIAL STRATEGIES TO BRIDGE THE GAP

Although germline testing is now routinely recommended for high-risk localized, locally advanced, and metastatic PCa patients, there is a disparity in the proportion of white vs. non-white PCa patients receiving germline testing [15, 16, 26]. We propose several reasons and potential solutions for this disparity, including the (1) nationwide shortage of genetic counselors to facilitate germline testing within current genetic counseling models, (2) differences in access to quality healthcare between white and non-white patients, (3) healthcare system mistrust among non-white men leading to unfavorable attitudes towards research and reluctance to seek care, (4) lack of knowledge or education about germline testing, (5) prohibitive cost of germline testing, and (6) understudied link between PCa and breast/ovarian cancer (Table 3).

Challenge and solution: shortage of genetic counselors and limitations of current genetic counseling models

Certified Genetic Counselors (CGCs) and physicians work cooperatively to facilitate germline testing and provide counseling, risk assessment, and result interpretation to PCa patients. The shortage of clinical cancer CGCs engaged in direct patient care creates an unmet need for genetic services that disproportionately affects socioeconomically disadvantaged, rural, and racial/ethnic minority patients. As the demand for germline testing grows, CGC workforce growth limitations will need to be addressed. One such limitation is clinical training capacity. Proposed solutions include novel clinical training techniques, such as nonclinical or extra-disciplinary training placements, rural clinical placements, peer supervision/assisted learning, role-emerging placements, clinical audit, and patient simulation. Perhaps most important is the need to recruit, train, and retain clinical supervisors by providing dedicated support personnel and professional development opportunities [37].

In addition to a nationwide shortage of CGCs engaged in direct patient care, existing genetic counseling models are becoming increasingly inadequate given the number of PCa patients referred for germline testing. The current time-intensive model of assessing family histories for genetic risk, providing pretest and posttest counseling, ordering appropriate testing, and interpreting test results over multiple in-person sessions is increasingly less feasible. Increased genetic literacy among medical oncologists, urologists, and radiation oncologists, including knowledge of patient risk factors and family history, genetics and genetic conditions, and available genetic services, may alleviate bottlenecks at the genetic counseling level [38].

Modifications to existing workflows within oncology practices may expand genetic resources for patients. Automating the risk assessment would be one such modification, whereby patient-completed family history questionnaires facilitate referral and testing processes: automated electronic medical record features can trigger genetic counseling referrals or alert clinical teams to patients with elevated cancer risks or who meet guidelines for germline testing. This would allow CGCs to prioritize posttest visits, especially those involving complex counseling or abnormal results [39]. Other practical strategies focus on increasing CGC efficiency and patient volumes, including group genetic counseling sessions. Additionally, establishing support roles, such as genetic counseling assistants, can alleviate administrative burdens [8]. Likewise, patient advocates and language interpreters in the genetic counseling setting can provide resources and translation services for non-English-speaking patients, which would further alleviate burdens on monolingual English-speaking CGCs and reduce patient miscommunication.

Rural patients are particularly disadvantaged by current genetic counseling models, given the scarcity of CGCs in more rural counties and among populations with a low median household income [40]. Teledicine, which has been adopted by many clinics in the COVID-19 era [41], can help bridge this gap: video genetics education and genetic counseling may be as effective as traditional genetic counseling and has resulted in a similar uptake of germline testing without compromising the tenants of informed consent [42]. Teledicine models do, however, need to adapt to potential challenges, including limited internet access, scheduling issues, billing questions, and state licensure regulations [41].

Challenge and solution: differences in the quality of care between white and non-white patients

There is overwhelming evidence that there are disparities in the quality of healthcare between white and non-white patients, even when insurance status, income, age, and severity of conditions are comparable [43]. Significant disparities have been noted for definitive therapy for PCa [6], with Black men being particularly underrepresented in PCa research, including validation studies of new clinical tools like genomic testing [15, 44]. One explanation...
for this disparity is that minority-serving physicians have been found to be significantly less likely to have ever referred a patient for germline testing or counseling, specialty services, or clinical trials [45]. This may be the result of many underlying issues, including access and cost.

Strategies to integrate genetic services into minority community health settings will be critical in ensuring the accessibility of germline testing. Because most CGCs are concentrated within large academic medical centers and hospital systems, the incorporation of satellite campuses and clinics into medically underserved communities would greatly expand access. In tandem, minority community health programs can practice evidence-based medicine through the implementation of clinical pathways to ensure that all patients are receiving the minimum standard of care. This will require expanding physician knowledge and awareness of current PCa clinical practice guidelines, as well as integration of these guidelines into existing workflows (Fig. 1).

Despite the expansion of germline testing guidelines for PCa patients, germline testing is not routinely covered by insurance. Coverage policies for germline testing in PCa patients are nonspecific and nonuniform across insurance companies, and physicians may not recommend genetic services for patients who cannot afford the out-of-pocket costs [46]. Expanding insurance coverage to include PCa patients that meet recommendations for germline testing may alleviate cost barriers. Additionally, in the absence of genetic services in medically underserved communities, expanded insurance coverage for transportation costs may benefit those who cannot access such services due to geographic barriers and for whom in-person counseling may help overcome hesitation due to unfamiliarity with telemedicine and/or lack of trust in the healthcare system.

**Challenge and solution: medical mistrust leading to unfavorable attitudes towards research and medicine**

Healthcare disparities among racial/ethnic minorities are thought to contribute to long-standing generational mistrust in healthcare-providing entities in the US. Medical mistrust has been shown to lower utilization of routine checkups and preventive care services [47–49], including referrals for genetic counseling and testing. Delays in these services may prevent a substantial number of men from obtaining recommended services until an advanced stage of illness [47]. This mistrust becomes a barrier to an emphasis on prostate health [50] and precludes racial/ethnic minorities from seeking PCa screening, germline testing, and treatment.

The lack of representation within medical institutions, as well as subsequent language barriers, may be a contributor to medical mistrust. In 2019, 5.0% and 5.8% of physicians identified as Black and Hispanic, respectively [51]. Further, 10.0% of CGCs in the US identified as non-white in 2021 [52]. Representation improves patient-clinician communication and rapport: when provided a doctor of the same race, Black men have been found to be more likely consent to invasive services, such as blood draws and biopsies, and discuss personal matters or health issues [53]. Hispanic men, in contrast, may face language barriers with clinicians: monolingual English-speaking clinicians may have limited communication with patients or rely on interpreters or translated materials, which may convey confusing or even contradictory information [54]. Issues of representation can be addressed by actively recruiting racial/ethnic minorities to the healthcare workforce and creating student training programs targeting these populations [55]. Language challenges can be addressed by employing multilingual, culturally cognizant interpreters in clinics where the need exists [56].

Medical mistrust may also stem from implicit bias, which refers to the unconscious and unintentional attitudes and stereotypes attributed towards a group of people. Implicit bias may contribute to health disparities by shaping physician behavior and producing differences in treatment along the lines of race, ethnicity, and gender. Healthcare professionals can combat implicit bias by individualizing, which involves a conscious focus on specific information about a patient instead of their race, ethnicity, or gender [57]. Addressing implicit bias early is essential: genetic counseling and nursing programs, medical schools, and healthcare professional training programs can expand and emphasize coursework in racial sensitivity and implicit bias. Additionally, addressing implicit bias in continuing medical education may help minimize biases.

Medical mistrust may also result from a lack of trust regarding the use of genetic information. Despite the passage of the Genetic Information Nondiscrimination Act in 2008, which was designed to protect Americans against discrimination in health insurance and employment based on their genetic information [58], utilization of genetic services among racial/ethnic minorities is disproportionately low [59]. In response, providers need to anticipate and dispel patient fears about germline testing. Patients may believe that their results are not confidential or that positive results will leave them susceptible to discrimination, reduced access to care, or insurance coverage loss [44]. Patients may also conflrate germline testing ordered by a clinician with direct-to-consumer DNA testing provided by companies that have faced controversy for sharing customers’ data with law enforcement and pharmaceutical companies.

Outreach and community support may help combat medical mistrust. Distributing medical literature directly to underserved populations has been shown to have positive results; however, personalized interactions between clinicians and racial/ethnic minority communities may further build trust and assuage fears about genetic services in order to encourage participation in germline testing and clinical trials [59]. Outreach and educational efforts within community institutions (such as churches) that involve partners and spouses, as well as cancer survivors within the community, may play a pivotal role [60].

**Challenge and solution: lack of knowledge regarding testing**

A lack of knowledge regarding germline testing and its implications for PCa screening, diagnostics, and treatment may present further barriers [38]. The availability of reliable, easy-to-understand information regarding the effects of P/LPVs on disease, as well as the importance of personal or family history of disease, is crucial [56].

Access to clear, concise tools about genetics is important because the complexity of such tools may compromise their effectiveness in identifying individuals at-risk for PCa. Genetics education among the general public is also important because individuals who are aware of and ask for specialized genetic services are the most likely to receive them [25]. Clear, simple, prescriptive education on genetics needs to be widely available to all PCa patients, and physicians will need to communicate the
advantages of genetic counseling and germline testing when they encounter high-risk localized and metastatic PCa patients who may benefit from it [40].

A lack of cross-cultural communication may also prevent racial/ethnic minorities from seeking or consenting to germline testing. The cultural impact of cancer can have an effect on patients’ attitudes towards germline testing: some South Asian and African-American communities have been found to take on a fatalistic view of cancer, associating the diagnosis with death; they may not wish to pursue testing if they believe nothing can be done to prevent or treat it [61]. Culture can also have an effect on the acceptance of test results: patients who receive germline testing may fear that their results will ostracize them from their family or community. Additionally, patients may not understand what a positive, negative, or inconclusive result means in the context of their own health and their family’s health. One solution could be culturally tailored genetic counseling (CTGC) and testing programs, which have been developed and evaluated to improve access to risk assessment services, subsequently enhancing the quality of care among patients from racial/ethnic minority groups. CTGC consists of education about risk factors for hereditary disease, personalized risk information, and discussions about the benefits, limitations, and risks of germline testing [56].

**Challenge and Solution: prohibitive cost and lack of insurance coverage for germline testing**

Access to germline testing is often limited by access to quality, affordable health insurance, which varies by race/ethnicity: NHW people are more likely to have health insurance than racial/ethnic minorities [62]. Additionally, NHW people are more likely to have private health coverage as opposed to public health coverage, such as Medicare and Medicaid [62]. Patients who are uninsured, underinsured, or insured by government programs may face significant barriers to obtaining care—for example, they may be denied care by private physicians, leading them to seek care in emergency departments, public hospital systems, or local health departments which may not offer the same referrals or specialty and preventative services as private practices [63]. Overall, having health insurance is strongly associated with undergoing PCa screening, lower stage of cancer at diagnosis, treatment for local/regional disease, prostatectomy, PCa survival, and quality of life [5]. Without health insurance, the cost of germline testing is often prohibitive. And, even when germline testing is covered by insurance, there may be prohibitive out-of-pocket costs, including deductibles and copayments. Additionally, not all insurers cover germline testing for PCa [46], including some private insurers and public options such as Medicare and Medicaid [64].

There are programs that increase PCa patient access to genetic services by offering free or reduced cost germline testing. Color Genomics offers a relatively low-cost risk analysis of several genes associated with PCa, as well as access to CGC and physician services [65]. Additionally, Invitae offers free germline testing and counseling for hereditary PCa through their Detect Hereditary Prostate Cancer program, in which eligible patients work with a genetic counselor or physician to order testing [66]. These, along with research studies such as the patient-driven PROMISE registry, which offers medical Color Genomics germline testing by mail to men with any stage of PCa [67], can reduce cost and access barriers.

Additional strategies for increasing payer coverage for germline testing and reducing test costs are necessary in order to create equitable access for all PCa patients. Strategies to increase insurance coverage may include clarifying and expanding Current Procedural Terminology codes to allow coverage for more specific tests and adding genetic specialists to insurance company staff to address shortages of genetic expertise. To address the costs of genetic tests, government subsidy programs and cost caps may be helpful in mitigating the cost to both the patient and insurers. Ultimately, there is a need for healthcare coverage reform in the US. In the absence of publicly funded healthcare, improving provider discussions about out-of-pocket costs is critical for ensuring informed patient testing and treatment decisions [64].

**Challenge and solution: understudied link between PCa and breast/ovarian cancers**

P/LPVs of BRCA1/2 have been found to increase the risk of multiple cancers, including those of the breast, ovary, and prostate. These P/LPVs have clinical implications for PCa patients as well as their families. BRCA1/2 associated HBOC should be suspected in individuals with family history of PCa and other cancers associated with HBOC syndrome [68], and likewise, PCa risk should be considered in individuals with a personal history of male breast cancer and/or family history suggestive of HBOC syndrome [69].

As with PCa, there are disparities in germline testing rates among racial/ethnic minority women with HBOC syndrome. These disparities are thought to be a result of multiple factors, including medical mistrust and fears of discrimination on the basis of genetic information [70]. Such disparities are thought to contribute to concordant disparities among racial/ethnic minority men with PCa. Therefore, addressing germline testing disparities among women with HBOC syndrome may aid in identifying male family members at-risk for developing PCa, as well as those already diagnosed with PCa who would benefit from germline testing’s impact on treatment options.

One particular challenge for PCa patients is that the names of certain PCa predisposition genes and familial risk factors (e.g., BRCA1/2 and HBOC syndrome) do not make obvious their link to PCa. As a result, it may not be clear to PCa patients that hereditary mutations in these genes affect them, as their names only indicate a link to breast and ovarian cancers. To address this misconception, one proposal is to change the name of HBOC syndrome to remove the sex specificity of the name [71], which may reduce confusion about its relevance to men.

**CONCLUSION**

It is widely accepted that a subset of PCa susceptibility is attributed to inherited predisposition. Because the identification of alterations in PCa predisposition genes may help inform screening strategies for patients and family members, treatment options in the metastatic setting, and clinical trial enrollment, it will become increasingly important to bridge the gap for PCa patients who are underserved with regard to germline testing. Issues to be addressed include a shortage of genetics professionals, disparities in care, medical mistrust, misinformation, and misunderstanding regarding germline testing, costs, and the understudied link between PCa and breast/ovarian cancer.

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Concept and design: RRM. Acquisition, analysis or interpretation of data: NW, JS. Drafting of paper: NW, RRM. Critical revisions of paper for important intellectual content: NW, JS, JJD, HHC, LM, and RRM. Approval of final paper: NW, JS, JJD, HHC, LM, and RRM. Supervision: RRM.

COMPETING INTERESTS

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ADDITIONAL INFORMATION

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