



# What Do Men with Metastatic Prostate Cancer Consider When Making Treatment Decisions? A Mixed-methods Study

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**Objective:** Metastatic prostate cancer (mPCa) patients often make complicated treatment decisions, yet decision aids to facilitate shared decision-making for mPCa are uncommon. To inform the development of patient-centered mPCa decision aids, we examined what mPCa survivors considered most important when making treatment decisions.

**Methods:** Using an exploratory sequential approach, we conducted three focus groups with 14 advanced prostate cancer survivors (n=5, n=3, n=6 in each group) to identify considerations for making treatment decisions. Focus groups were audio-recorded and transcribed, and we identified qualitative themes. We then developed a quantitative survey to assess the importance of each theme and administered the survey to mPCa survivors (N=100). We used relative frequencies to determine the most strongly endorsed items and chi-squared and Fisher's exact tests to assess associations with participant characteristics.

**Results:** Focus groups yielded 11 themes, and the resulting survey included 20 items. The most strongly endorsed mPCa treatment considerations were: relying on physician's treatment recommendations (79% strongly agree); wanting to feel well enough to spend quality time with loved ones (72% strongly agree); the importance of dying in a manner consistent with one's wishes (70% strongly agree); hoping to eliminate cancer completely (68% strongly agree); and optimizing treatment efficacy (65% strongly agree). Age, race, marital status, employment status, and self-reported health were related to how strongly men endorsed various considerations for mPCa treatment decision-making.

**Conclusion:** We identified multiple considerations that mPCa survivors appraised when making treatment decisions. These data may inform the development of patient-centered decision aids for mPCa.

**Keywords:** decision-making, focus groups, metastasis, prostate cancer, quality of life

Prostate cancer (PCa) is the most common cancer among men in the US, accounting for more than 20% of annual male cancer diagnoses.<sup>1</sup> Although localized PCa is highly treatable, metastatic PCa (mPCa) is incurable and the second leading cause of cancer death in the US (five-year survival of 31%, 33,330 deaths expected in 2020).<sup>1</sup> Men with mPCa face choices between multiple noncurative systemic treatment options with various side effect profiles, financial obligations, and modes of delivery, but all treatments result in similar survival benefits.<sup>2</sup> There is often not a "best" mPCa treatment option in terms of expected efficacy, and the optimal sequence of treatments to optimize efficacy remains an active area of investigation.<sup>3-7</sup>

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Due to the complexity of treatment options, professional societies (eg, American Urological Association) recommend shared decision-making as an ideal approach to prostate cancer treatment decision-making.<sup>8</sup> In shared decision-making, patients and providers collaboratively make treatment decisions based on clinical information (ie, available treatments, risks, benefits, costs) and patients' values and preferences.<sup>9,10</sup> However, shared decision-making is complex, and it can be difficult to accomplish in the limited time available for clinical consultation and care. Thus, decision aids can facilitate shared decision-making by increasing patient knowledge about treatment options.<sup>11</sup> For prostate cancer, decision aids have largely been studied in the context of localized disease and prostate-specific antigen (PSA) screening.

Decision aids range from basic informational pamphlets to more sophisticated and interactive technology and internet-based tools. Despite considerable variability in the quality of decision aids and their adherence to criteria outlined in the International Patient Decision Aid Standards,<sup>12,13</sup> decision aids for localized prostate cancer and PSA screening are associated with improved decisional outcomes (eg, reduced decisional conflict and regret, improved knowledge, satisfaction, and perceptions of control).<sup>12,14–16</sup> However, few empirically supported decision aids have been developed for mPCa, leaving men with mPCa with little evidence-based guidance for facilitating shared decision-making. This is a significant oversight, particularly as the landscape of mPCa treatments continues to increase in complexity. Because patients' values and preferences are a critical component of shared decision-making, an important first step in the development of empirically supported patient-centered decision aids is to better understand mPCa patients' considerations when making treatment decisions.

To address this need, we identified and characterized the considerations that affect how mPCa patients make treatment decisions using a mixed-methods approach. First, we conducted focus groups to identify treatment decision-making considerations. Then, using the resulting information, we developed and administered a survey to mPCa survivors to determine which treatment decision-making considerations they most strongly endorsed. Finally, we explored associations between patient characteristics and treatment decision-making considerations.

## Methods

### Participants and Procedures

We performed an exploratory sequential mixed-methods study including an initial qualitative phase of data collection and analysis, followed by a quantitative data collection and analysis phase. This study was approved by the Institutional Review Board at Vanderbilt University Medical Center (VUMC; approval #140871).

#### Qualitative Phase: Focus Groups

From August to September 2014, men with advanced incurable prostate cancer were recruited from urology and medical oncology clinics at VUMC to participate in focus groups (k=3) as a qualitative exploration of treatment decision-making considerations for men with incurable prostate cancer. All participants provided informed consent, including consent to publish anonymized responses. Focus groups were completed in person and required patients to report to a facility that was distinct from the clinical setting and to come at a time outside of scheduled appointments. To improve feasibility of recruitment and attendance at focus groups, both men with biochemical recurrence and with mPCa were included in the focus groups if their prostate cancer was considered incurable. A standard moderator's guide was used for each group that included questions about relevant topic areas generated from the literature and clinical experience. Broad topic areas identified a priori were queried related to what is important to men with incurable PCa as they make treatment decisions for their PCa. These topics were explored to identify specific patient-developed themes, and focus groups were performed until thematic saturation was reached. Focus groups were audio-recorded, transcribed verbatim, de-identified, and analyzed systematically to identify qualitative themes. Initial themes were developed using the first focus group transcript, and themes were added and revised to capture the content of discussions. The coding system was reviewed and revised by the principal investigator AKM, and the final coding system was applied to all three focus group transcripts. Participants also reported sociodemographic and clinical information via a demographics survey.

#### Quantitative Phase: Survey

For the quantitative phase of the study, we developed a survey based on the themes identified in the focus groups to determine which of the treatment considerations mPCa survivors most strongly endorsed as being important to

them. Items were developed to probe aspects of each theme identified in the qualitative phase of the study. From October 2015 to November 2016, mPCa survivors who had not participated in the focus groups were recruited from urology and medical oncology clinics at VUMC and online from ZEROcancer.com, a nonprofit organization that provides patients and caregivers with prostate cancer education and advocacy activities. The survey component of this study was largely hypothesis generating, thus a recruitment goal of 100 participants was deemed acceptable. After providing informed consent, including consent to publish anonymized responses, participants completed surveys describing their sociodemographic and clinical characteristics and rated their perceived health (ie, “How would you describe your overall health right now?”) with the following response options: excellent, good, fair, and poor. Participants rated their agreement with statements about treatment decision-making on a Likert-scale from strongly agree (1) to strongly disagree (5). Participants recruited from ZEROcancer.com completed the survey online, whereas participants recruited from VUMC clinics completed the survey in clinic using paper forms. The data that support these findings are available from the corresponding author upon reasonable request.

## Data Analysis

Qualitative data were divided into short quotes containing a single thought or idea and loaded into Microsoft Excel to facilitate coding by two coders. Quantitative data were analyzed using IBM SPSS Statistics for Windows, Version 25.0 (IBM Corporation, Armonk, NY, USA). We used descriptive statistics to characterize the participants. For focus group data, an initial codebook was developed and two independent coders assessed the transcripts for overarching themes.<sup>17</sup> Disagreements were resolved by consensus. Analysis continued until thematic saturation was reached (ie, the point at which no further themes were identified).<sup>18</sup>

For survey data, we used relative frequencies to determine the most strongly endorsed items (ie, proportion of participants who responded strongly agree). Item responses were dichotomized in this way to allow us to identify the treatment decision-making considerations that participants endorsed most strongly. We used Fisher’s exact test to explore associations between participant characteristics and likelihood of strongly endorsing each item (ie, strongly agree vs other response) across the following

participant characteristics: age (median split <68 vs ≥68 years old), race (non-white vs white), marital status (married/partnered vs not married/partnered), employment status (working vs retired) and self-reported health (poor/fair vs good/excellent). Significance was determined by  $p < 0.05$ .

## Results

### Participant Characteristics

Table 1 describes the characteristics of the focus group participants (N=14 total across k=3 focus groups) and survey participants (N=100). Most focus group participants had mPCa (79%) and were median ages of 71, 62, and 72 years old in each of the focus groups, respectively. Participants were primarily white (93%) and married/partnered (79%).

Survey participants all had mPCa and were a median age of 68 years old. Participants were primarily white (88%), married/partnered (80%), retired (63%), and insured (99%). There were no characteristic differences between participants recruited from VUMC (n=77) and ZEROcancer.com (n=23), with the exception that participants recruited from ZEROcancer.com were more likely to be retired ( $\chi^2(1)=6.02$ ,  $p=0.01$ ).

### Qualitative Themes of Treatment Decision-making Considerations

Focus groups yielded 11 themes related to treatment decision-making. Here, we briefly describe the themes and present select representative quotes. Additional quotes are included in the [Supplemental Table](#).

#### Confidence in Physician’s Expertise

Most participants reported relying on their physician’s recommendations for planning their treatment approach. As treatments failed and new treatments were identified, participants relied heavily on their physician to identify next steps.

I just listen to what the doctor says and follow her advice.  
I’m content with that.

#### Eliminating Cancer

The possibility of completely eliminating cancer was a salient consideration among participants. Despite the cognitive acceptance that advanced prostate cancer is incurable, many participants expressed hope that the next treatment or discovery could result in cure.

I think we are all just hoping for that miracle drug.

**Table 1** Sociodemographic and Clinical Characteristics of Men Who Participated in the Focus Groups and Survey

Variables	Focus Group 1 (n=5)	Focus Group 2 (n=3)	Focus Group 3 (n=6)	Survey (n=100)
Age, years; median	71	62	72	68
Race/ethnicity; n (%)				
White	5 (100)	3 (100)	5 (83)	88 (88)
Black/African American	0	0	0	10 (10)
Other	0	0	1 (17)	2 (2)
Marital status; n (%)				
Married or partnered	5 (100)	2 (67)	4 (67)	80 (80)
Not married or partnered	0	1 (33)	2 (33)	19 (19)
Employment status; n (%)				
Working				35 (35)
Retired				63 (63)
Health insurance status; n (%)				
Insured				99 (99)
Not insured				1 (1)
Self-reported health; n (%)				
Excellent or good				68 (68)
Fair or poor				30 (30)
Disease status; n (%)				
Biomedical recurrence	1 (20)	1 (33)	1 (17)	0
Metastatic	4 (80)	2 (67)	5 (83)	100 (100)
Most recent PSA, ng/mL; M (SD)				12.33 (23.66)
Site of metastasis; n (%)				
Bone				79 (79)
Lymph node				27 (27)
Liver				2 (2)
Lung				3 (3)
Other				3 (3)
Do not know				7 (7)
Total number of metastases; n (%)				
1				77 (77)
2				11 (11)
3				5 (5)
Do not know				7 (7)

**Notes:** Only brief sociodemographic information was collected from focus group participants. Among survey participants, variables with missing data are age (n=96), marital status (n=99), employment status (n=98), self-reported health (n=98), and most recent PSA (n=97).

**Abbreviations:** M, mean; PSA, prostate-specific antigen.

### Prolonging Survival

Participants prioritized treatments that would help them prolong survival. For many, family and marital relationships influenced this factor.

My wife says, “Just do what you can to stick around.”

### Quality of Life (QOL)

Many participants felt that prolonging survival must be balanced with QOL, defined as continuing to live in

a reasonably normal way, carrying on basic activities of daily living, and maintaining quality interpersonal relationships. Some participants expressed a high value of QOL, whereas others were willing to sacrifice QOL for longer survival.

For me it’s really a pretty easy decision. I’m willing to live in discomfort. I’ll do whatever.

I would go for the quality of life over the quantity over life, I think every time.

### Side Effects

Participants described many treatment side effects including memory loss, trouble thinking, pain, bruising, fatigue, erectile dysfunction, and loss of libido. They discussed how side effects can interfere with daily activities, including work and hobbies. While participants were aware of potential treatment-related side effects, they often chose treatments to prolong survival or suppress the cancer despite these side effects.

Chemo has pretty serious side effects of its own, so you know there's that trade off all the time.

### Pain Relief

Pain is a common and intractable side effect of advanced prostate cancer, and pain relief was identified as a significant consideration. Some participants reported willingness to use powerful narcotics to manage cancer-related pain. Other participants expressed unwillingness to undergo further surgery because of the anticipated pain and potential disability.

Radiation won't cure [the cancer] but it will relieve the pain.

### Treatment Efficacy

Optimizing treatment efficacy is a concern when treating advanced prostate cancer. Several participants discussed the importance of optimal sequencing of treatments, so as not to eliminate future treatment options. Others commented that having prostate cancer creates complications for treating other, unrelated health issues.

If I had gone on that drug trial, then I wouldn't have been able to do the second.

### Financial Toxicity

Although paying for treatment was often not the primary concern, participants considered the cost of treatments and whether treatments would be covered by insurance. Participants voiced complaints about high out-of-pocket costs, insurance companies refusing to cover treatments, and delays in treatment due to extended insurance review.

[The insurance company] called me and said, "You're approved." Then they called back saying, "Well, the good news is that you are approved, but it's going to cost you \$10,000." I said, "That's not good news."

### Treatment Accessibility/Convenience

For some, having to travel for treatment was a barrier to care. They described the burden of some treatments being

unavailable in their local area. Although this was not a primary obstacle for most participants, it was discussed by some as a consideration for how they determined next steps.

They don't have [that treatment option] around here, so you're gone for 6 months . . . [if you decide to pursue that treatment].

### Fastest Treatment

Many participants described a sense of time urgency and considered the ability to be treated quickly of high importance. In practice, this can affect treatment choices requiring prior authorization (eg, infusion treatments) or complicated treatment planning requiring multiple specialties (eg, radiopharmaceuticals, immunotherapy). This can also be a barrier for patients who must apply for financial assistance to offset costs of care (eg, via co-payment relief funding sources).

The idea in the beginning was that I had to [get treated] right now. I've got to do everything today.

### Decision Regret

Some participants felt they had made the best treatment choices they could, whereas others expressed regrets and wished they had chosen a different initial treatment approach.

[Looking back] we should've said "No, we're not doing this [treatment]."

## Survey Analysis of Treatment Decision-making Considerations

We developed a 20-item survey to assess treatment decision-making considerations based on the 11 qualitative themes. Table 2 shows the proportion of participants who responded "strongly agree" to each item. The largest proportion of participants strongly endorsed items reflecting reliance on their doctor for making treatment decisions (79% strongly agree), wanting to feel well enough to spend quality time with loved ones is (72% strongly agree), the importance of dying in a manner consistent with one's wishes (70% strongly agree), hoping to eliminate cancer completely (68% strongly agree), and treatment sequencing to maintain future treatment options (65% strongly agree). By contrast, the smallest proportion of participants strongly endorsed items reflecting less time urgency (20% strongly agree) and pain relief (25% strongly agree).

**Table 2** Proportion of Participants Who Strongly Agreed with Statements About Treatment Decision-making Considerations

Treatment Decision-making Consideration	Survey Item	Percent of Participants Responding "Strongly Agree"
Confidence in physician's expertise		
	• I rely on my doctor to tell me how to treat my prostate cancer.	79
Eliminating cancer		
	• When I choose a cancer treatment, I am hoping for one that will eliminate the cancer completely.	68
Prolonging survival		
	• Above all, living as long as I can is my most important goal.	53
	• I need to live as long as possible, no matter how I feel, so I can continue to take care of my loved ones.	32
Quality of life		
	• I am more interested in having a good quality of life rather than the longest life possible.	59
	• If I am going to die from cancer, the most important thing for me is to die with dignity.	70
	• It is very important to me that I do not end my life so sick I am stuck in bed.	63
	• I want a treatment that lets me keep working or doing the things I enjoy even if it means I will not live as long.	51
	• Feeling well enough to spend as much quality time as possible with my loved ones is the most important thing.	72
	• I would choose to die sooner rather than be a burden to my loved ones.	52
Side effects		
	• I am willing to put up with uncomfortable side effects as long as the treatment is slowing the growth of my cancer.	39
	• I would put up with severe side effects to avoid needing to be in a wheelchair.	34
	• I would put up with severe side effects to make sure I have time to organize my affairs for my loved ones.	56
Pain relief		
	• The most important thing is finding a treatment that minimizes pain.	25
Treatment efficacy		
	• I try to choose treatments that will not close any doors to future treatments.	65
Financial toxicity		
	• Avoiding financial trouble due to treatment of my prostate cancer is very important to me.	54

(Continued)



**Table 2** (Continued).

Treatment Decision-making Consideration	Survey Item	Percent of Participants Responding “Strongly Agree”
Treatment accessibility		
	• I am willing to travel as far as necessary to get the best treatment.	53
Fastest treatment		
	• I am not in a hurry to treat my cancer, I would rather take my time and make the best choices.	39
	• I would choose a treatment that is available now rather than wait for a treatment that might work better, but requires waiting.	20
Decision regret		
	• I choose my treatment so I will not look back with regret.	62

Exploratory Fisher’s exact test revealed associations between participant characteristics and the likelihood of strongly endorsing individual survey items (Table 3).

### Age

Relative to younger participants, a larger proportion of older participants strongly endorsed the importance of dying with in a manner consistent with one’s wishes (56% vs 78%, respectively; OR=2.80, 95% CI: 1.15–6.82,  $p=0.027$ ) and valuing QOL over prolonging survival (32% vs 58%, respectively; OR=3.00, 95% CI: 1.28–3.00,  $p=0.013$ ). In addition, relative to younger participants, a marginally larger proportion of older participants strongly endorsed the importance of avoiding financial toxicity (42% vs 62%, respectively; OR=2.29, 95% CI: 1.00–5.22,  $p=0.063$ ).

### Race

Relative to white participants, a larger proportion of non-white participants strongly endorsed the importance of prolonging survival to care for loved ones (26% vs 75%, respectively; OR=0.12, 95% CI: 0.03–0.47,  $p=0.001$ ), and a marginally larger proportion of non-white participants strongly endorsed the importance of minimizing pain (22% vs 50%, respectively; OR=0.28, 95% CI: 0.08–0.95,  $p=0.068$ ).

### Marital Status

Relative to married/partnered participants, a larger proportion of single participants strongly endorsed willingness to endure severe side effects to avoid needing a wheelchair (29% vs 55%, respectively; OR=0.33, 95% CI: 0.12–0.90,  $p=0.036$ ), and a marginally smaller proportion of single participants strongly endorsed the importance of feeling

well enough to spend quality time with loved ones (76% vs 55%, respectively; OR=2.63, 95% CI: 0.95–7.29,  $p=0.092$ ).

### Employment Status

Relative to retired participants, a marginally larger proportion of working participants strongly endorsed hope for eliminating cancer completely (62% vs 80%, respectively; OR=2.50, 95% CI: 0.95–6.58,  $p=0.074$ ).

### Self-reported Health

Relative to participants with self-reported good/excellent health, a larger proportion of participants with self-reported poor/fair health strongly endorsed hope for eliminating cancer completely (43% vs 78%, respectively; OR=0.22, 95% CI: 0.09–0.54,  $p=0.001$ ), willingness to endure uncomfortable side effects to slow cancer growth (37% vs 63%, respectively; OR=0.34, 95% CI: 0.14–0.82,  $p=0.017$ ), and willingness to endure severe side effects to avoid needing a wheelchair (17% vs 41%, respectively; OR=0.29, 95% CI: 0.10–0.84,  $p=0.021$ ). Moreover, relative to participants with self-reported good/excellent health, a marginally larger proportion of participants with self-reported poor/fair health strongly endorsed the importance of sequencing cancer treatments (50% vs 71%, respectively; OR=0.47, 95% CI: 0.17–1.01,  $p=0.067$ ) and avoiding decision regret (47% vs 68%, respectively; OR=0.42, 95% CI: 0.17–1.01,  $p=0.071$ ).

## Discussion

In this hypothesis-generating study, we identified what advanced prostate cancer survivors considered when

**Table 3** P-values Indicating the Significance of Fisher's Exact Test Assessing the Relationships Between Participants' Sociodemographic Characteristics and Likelihood of Strongly Endorsing Each Survey Item

Survey Item	Age	Race	Marital Status	Employment Status	Health Status
I rely on my doctor to tell me how to treat my prostate cancer	1.000	0.712	0.356	0.799	0.431
When I choose a cancer treatment, I am hoping for one that will eliminate the cancer completely	1.000	0.747	0.285	0.074*	0.001***
Above all, living as long as I can is my most important goal	0.540	0.130	0.317	0.675	0.279
I need to live as long as possible, no matter how I feel, so I can continue to take care of my loved ones	0.272	0.001***	1.000	0.823	0.489
I am more interested in having a good quality of life rather than the longest life possible	0.013**	0.760	1.000	0.213	0.118
If I am going to die from cancer, the most important thing for me is to die with dignity	0.027**	0.749	0.287	0.117	0.477
It is very important to me that I do not end my life so sick I am stuck in bed	0.292	1.000	0.445	0.393	1.000
I want a treatment that lets me keep working or doing the things I enjoy even if it means I will not live as long	0.675	0.543	0.617	0.671	0.825
Feeling well enough to spend as much quality time as possible with my loved ones is the most important thing	0.656	1.000	0.092*	0.817	0.628
I would choose to die sooner rather than be a burden to my loved ones	0.303	0.223	0.132	1.000	0.827
I am willing to put up with uncomfortable side effects as long as the treatment is slowing the growth of my cancer	0.683	0.219	0.210	1.000	0.017**
I would put up with severe side effects to avoid needing to be in a wheelchair	1.000	0.329	0.036**	1.000	0.021**
I would put up with severe side effects to make sure I have time to organize my affairs for my loved ones	0.834	0.530	1.000	0.198	0.120
The most important thing is finding a treatment that minimizes pain	0.345	0.068*	0.576	0.473	0.801
I try to choose treatments that will not close any doors to future treatments	0.515	0.533	0.794	0.512	0.067*
Avoiding financial trouble due to treatment of my prostate cancer is very important to me	0.063*	0.137	0.803	0.293	0.666
I am willing to travel as far as necessary to get the best treatment	0.148	0.368	0.618	0.675	0.129
I am not in a hurry to treat my cancer, I would rather take my time and make the best choices	0.217	1.000	0.114	1.000	0.108
I would choose a treatment that is available now rather than wait for a treatment that might work better, but requires waiting	0.201	1.000	0.127	0.832	0.268
I choose my treatment so I will not look back with regret	1.000	0.527	0.607	1.000	0.071*

**Notes:** Sociodemographic variables were dichotomized as follows: age, median split <68 vs ≥68 years old; race, nonwhite vs white; marital status, married/partnered vs not married/partnered; employment status, working vs retired; self-reported health, poor/fair vs good/excellent. \* $p < 0.10$ , \*\* $p < 0.05$ , \*\*\* $p < 0.01$ .

making treatment decisions, and we investigated the relative importance of treatment considerations among mPCa survivors. The three most strongly endorsed treatment decision-making considerations were related to relying on the physician's recommendations and maintaining QOL (ie, to spend time with loved ones and dying in a manner consistent with one's wishes). Many participants also strongly endorsed the importance of treatment efficacy (ie, treatment sequencing), but this was only the fifth most strongly endorsed consideration. Subgroup analyses showed that mPCa survivors may have different priorities when making treatment decisions depending on demographic characteristics. For example, a larger proportion of older mPCa survivors strongly endorsed survey items

related to maintaining QOL and dying with in a manner consistent with one's wishes than younger mPCa survivors, whereas a larger proportion of non-white mPCa survivors strongly endorsed wanting to prolong survival to care for loved ones than white mPCa survivors. These findings highlight the importance of tailoring discussions about mPCa treatment decision-making to individual patient characteristics and values.

Our findings are mostly aligned with past studies of treatment considerations for prostate cancer. One study in localized prostate cancer found that patients considered whether potential treatments could eliminate their cancer, extend their survival, delay disease progression, and preserve their QOL in the context of treatment side effects.<sup>19</sup>



We identified many similar treatment considerations among men with mPCa, highlighting significant overlap in what patients consider while making treatment decisions for prostate cancer across the disease continuum.

Similar to a study in localized prostate cancer<sup>19</sup> and a study of patients with advanced prostate cancer,<sup>20</sup> we found that most patients relied heavily on their physician's recommendations. However, another study in localized disease found that patients relied more on the stories and experiences of others, particularly men who have been previously treated for localized prostate cancer.<sup>21</sup> It is possible that as prostate cancer progresses to advanced disease, men may have fewer acquaintances with similar disease histories and experiences, making acquaintances less helpful for informing treatment decision-making. Thus, patients may increasingly rely on their physician's expertise as prostate cancer progresses to advanced disease. Men with mPCa could benefit from programs that provide opportunities for peer support, such as group-based psychosocial and educational interventions.<sup>22</sup> Group-based programs such as these inherently increase opportunities to learn from the experiences of other men with mPCa. In addition, when designing decision aids for mPCa, researchers should carefully consider the role of patient narratives.<sup>23,24</sup> Notably, past work has shown that in clinical practice, the treatment that men receive for localized prostate cancer is more aligned with physician recommendations than with patient preferences.<sup>25</sup> It is unclear whether this finding generalizes to men with mPCa. Nevertheless, participants in our study of mPCa clearly described the factors they consider when making treatment decisions, with priorities differing by key socio-demographic characteristics. Future studies should explore strategies for ensuring that patient preferences are not lost in the treatment decision-making process.

We found that financial toxicity, or problems related to the out-of-pocket costs of treatment (eg, co-payments, deductibles, co-insurance),<sup>26</sup> was not a major consideration for mPCa treatment decision-making. This is in contrast with recent assessments of decision-making in advanced prostate cancer, in which patients were not willing to accept significant out-of-pocket expenses for treatments.<sup>27</sup> Our sample was mostly retired and adequately insured, which could influence our results. The experience of financial toxicity likely varies by health literacy, socioeconomic status, and other factors, and the influence of financial toxicity warrants further investigation among mPCa survivors.<sup>28</sup>

Finally, a recent discrete choice experiment of prostate cancer patients and oncologists identified that treatment efficacy and pain control were similarly important when considering treatment choices.<sup>29</sup> In our study, treatment efficacy emerged as more important than pain control for mPCa survivors, with treatment efficacy identified as the fifth most strongly endorsed consideration. This underscores the importance of treatment efficacy and sequencing among mPCa patients.

## Study Limitations

Our sample sizes were small, and there was little socioeconomic diversity in our focus group and survey participants. Most participants were between 60 and 75 years old, white, married, retired, and insured, which may limit the generalizability of our findings to younger men, racial/ethnic minorities, and men who continue to work through cancer treatment. In particular, our finding that financial toxicity did not play a large role in treatment decision-making for mPCa could be due to the characteristics of our sample and may not reflect the values of the broader population of mPCa survivors. We did not include caregivers, partners, or other loved ones in our analyses; inclusion of these important cancer care partners in future studies may enhance our understanding of treatment considerations, as spouses are frequently involved in the process of treatment planning.<sup>30–33</sup> Finally, this study was cross-sectional and we could not assess potential variations in how treatment considerations change over time. Longitudinal studies of treatment decision-making considerations for mPCa will be a valuable addition to this field.

## Clinical Implications

Understanding what men with mPCa consider when making treatment decisions can inform how clinicians approach challenging conversations about treatment options in practice. mPCa survivors clearly endorsed the importance of their relationship and trust in their physician, and open communication about patients' values are important for optimizing the doctor–patient relationship.<sup>34</sup> Studies should continue to elucidate strategies to enable patients to participate in the decision-making process and communicate their treatment considerations and preferences. This may ultimately improve the quality of treatment decisions and outcomes for mPCa.<sup>20,30,35</sup>

## Conclusions

We identified multiple considerations that men with mPCa report are important for treatment decision-making. Several aspects of QOL were consistently identified, including feeling well enough to spend time with loved ones and dying in a manner consistent with one's wishes. These values should be considered as critical components of any treatment decision for this population. Efforts to understand aspects of treatment decisions in more diverse populations remain necessary and are ongoing.

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## Disclosure

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