

# CANCER SUPPORT COMMUNITY

A Global Network of Education and Hope

# Background

Treatment by a multidisciplinary health care team can improve patient outcomes among people living with cancer, but the extent to which prostate cancer (PC) patients in the community access specialty providers remains unclear.

# Aims

This study explored, in a national sample of prostate cancer patients, 1) associations between number of providers and treatments discussed/received, and 2) predictors of informed treatment decision-making and treatment regret.

## Methods

Of 311 PC patients enrolled in the Cancer Support Community's online Cancer Experience Registry, 181 indicated seeing one or more physician providers during treatment (primary care provider, urologist, radiation oncologist, medical oncologist, pain specialist). Informed decision-making and treatment regret were measured using the Prostate Cancer Outcomes Measures; each scale was calculated as the sum of 5 individual items items (1=not at all to 5=very much) and standardized to range between 0 and 100. In logistic regression analysis, we modeled high levels of informed decisionmaking (>=75 out of 100) and treatment regret (<25 out of 100), adjusting for stage at diagnosis and number of treatments discussed.

## **Participants**

	Total Sample N = 181	1 Provider n = 72	2 Providers n = 53	3-5 Providers n = 56	Chi-square p-value	
	%	%	%	%		
Age >65 years	59	69	57	48	.06	
Non-Hispanic White	87	92	87	82	.27	
College Graduate	70	72	69	69	.89	
Geographic region						
Rural	23	20	19	30		
Suburban	35	39	28	37	.28	
Urban	42	41	53	33		
Time since diagnosis						
< 2 years	36	35	37	38		
2-5 years	31	25	37	32	.50	
5+ years	33	41	27	30		
Stage at diagnosis						
<i>O, I, II</i>	37	43	40	27		
III, IV	34	15	36	55	<.001	
Don't know	29	42	25	18		
Ever Metastatic	30	17	28	48	.001	
Advanced prostate cancer patients were more likely to be treated by a team of providers than a sinale provider.						

## **Acknowledgments**

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# Health Care Team and Treatment Decision-Making Experiences **Among Prostate Cancer Patients**

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## Results



Patients reporting a single provider for treatment primarily consulted a urologist or radiation oncologist. Patients with 3-5 providers typically included a urologist and oncology specialists. Half of patients with multiple providers saw a PCP, and fewer than 1 in 5 consulted a pain specialist.

\*Chi-square test, p<.05

# **Prostate Cancer Treatment(s) Discussed with Provider** Single Provider 2 Providers 3-5 Providers 50

Patients seeing multiple providers were more likely to discuss hormone therapy and cancer clinical trials but less likely to discuss surgery and active surveillance.

\*Chi-square test, p<.05



\*Chi-square test, p<.05

## References

<sup>1</sup>Clark, J. A., Bokhour, B. G., Inui, T. S., Silliman, R. A., & Talcott, J. A. (2003). Measuring patients' perceptions of the outcomes of treatment for early prostate cancer. Medical Care, 923-936.

<sup>2</sup> Cegala, D. J., Bahnson, R. R., Clinton, S. K., David, P., Chi Gong, M., Monk III, J. P., & Pohar, K. S. (2008). Information seeking and satisfaction with physician–patient communication among prostate cancer survivors. *Health Communication*, *23*(1), 62-69.



Learn more or join the Registry at www.CancerExperienceRegistry.org

## Logistic Regression Analysis Predicting High Levels of Informed Decision-Making and Low Levels of Treatment Regret



\*p=.98, \*\*p=.48; Hosmer and Lemeshow's goodness of fit test. % is predicted probability holding other variables constant in the model. Interaction between stage at diagnosis and number of providers was not significant in either model. Models adjust for stage at diagnosis and total number of treatments discussed.

- Conclusions
- oncology specialist.
- contrary to our hypothesis.
- of prostate cancer.

cancer survivors and their caregivers.

High Informed Decision Making* (n = 172)			Low Treatment Regret** (n = 172)			
%	Odds ratio (95% CI)	<i>p</i> -value	%	Odds ratio (95% CI)	<i>p</i> -value	
63.1	1.00 (Reference)		82.7	1.00 (Reference)		
50.5	0.59 (0.28, 1.29)	.19	66.7	0.42 (0.17, 1.01)	.053	
40.2	0.39 (0.18, 0.88)	.023	55.1	0.26 (0.11, 0.62)	.003	

After accounting for stage at diagnosis and total number of treatments discussed:

Having multiple providers was associated with lower informed decision-making:

• 63% of patients with a single provider were classified as high informed decision-making • 51% of patient with 2 providers were classified as high informed decision-making • 40% of patients with 3-5 providers were classified as high informed decision making

Having fewer providers was associated with low treatment regret:

• 83% of patients with a single provider were classified as having low treatment regret • 67% of patients with 2 providers were classified as having low treatment regret • 55% of patients with 3-5 providers were classified as having low regret

Advanced prostate cancer patients are more likely than those with early stage disease at diagnosis to be treated by both a urologist and at least one

Patients seeing multiple providers were more likely to discuss cancer clinical trials and more likely to receive non-surgical treatment(s).

Even after accounting for stage at diagnosis and number of treatment(s) discussed with provider(s), having multiple providers was associated with low levels of informed decision-making and high levels of treatment regret,

While treatment by multiple medical providers may expand available treatment options, patients can still experience treatment regret, and access to multiple providers alone does not ensure care coordination.

Integrated care and shared decision-making between patients, families, and providers remains ever important in the evolving treatment landscape

The Cancer Experience Registry is an online research initiative that captures the immediate and ongoing or changing social and emotional experiences of

• The Registry is for all cancer survivors and caregivers, and also includes 13 disease-specific surveys. • Findings contribute toward advancing research, health care and policy.

• Over 14,000 cancer survivors and caregivers are registered in the Cancer Experience Registry.