

Perceptions About Cancer Clinical Trials Among Prostate Cancer Survivors

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Background

- Clinical trials are an important mechanism to advance innovation and quality patient care, yet patient participation in clinical trials remains low in the United States.
- Recruitment into prostate cancer clinical trials (CCTs) remains challenging despite efforts to enhance patient understanding of and access to CCTs.

Aims

The objective of this study was to examine perceptions of CCTs among prostate cancer (PC) survivors in the Cancer Support Community's Cancer Experience Registry sample.

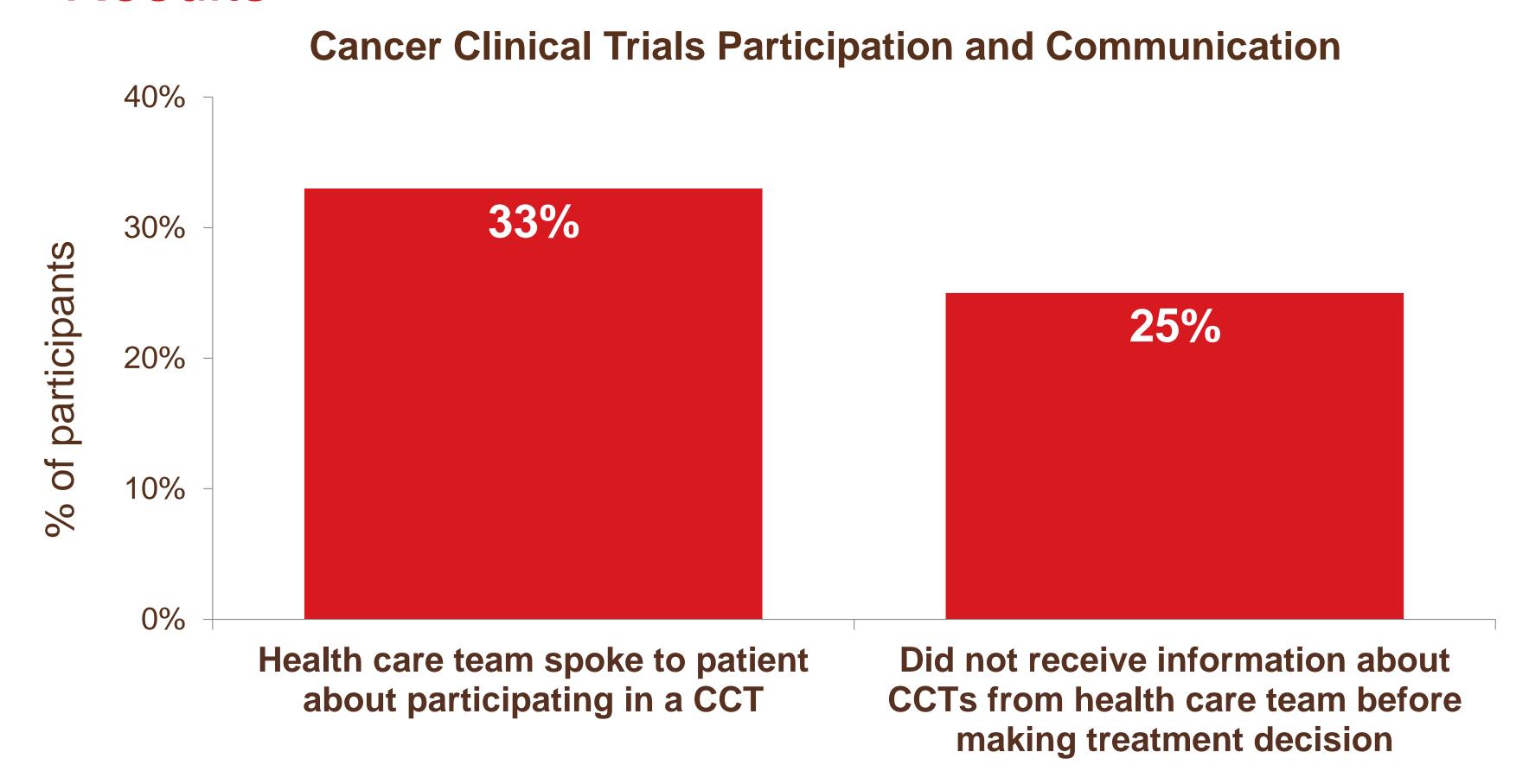
Methods

- 86 PC survivors enrolled in the Cancer Support Community's Cancer Experience Registry, provided demographic and clinical history, and rated their agreement (0 = strongly disagree to 4 = strongly agree) with eight statements about cancer clinical trials (CCTs).
- Bivariate associations between CCT counseling and individual factors (sociodemographic information, where treatment received, history of CCT counseling and CCT participation) were examined via Spearman's rank correlation and chi-square tests.

Participants

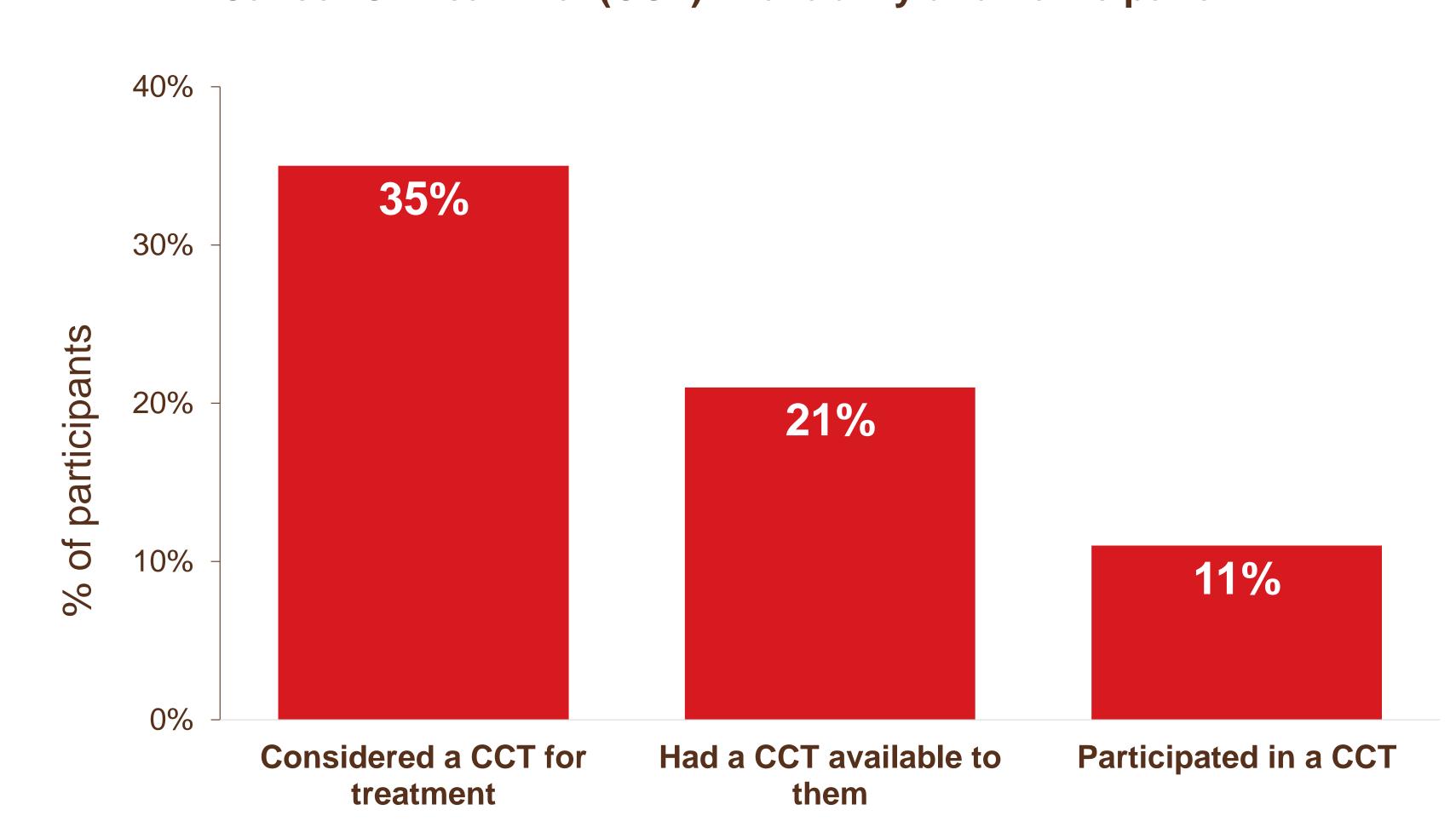
<i>N</i> = 86	Mean/ n	SD / %
Age (years)	65	7
	Range: 50 – 85	
White	82	95%
Time since diagnosis (years)	4	4
Treatment history		
Surgery	21	24%
Radiation	29	34%
Both surgery and radiation	17	20%
Hormone therapy (current)	14	16%
Hormone therapy (past)	7	8%
Disease status		
Diagnosed 5+ years ago	27	32%
Recurrence	19	22%
Metastatic disease	26	31%

Results



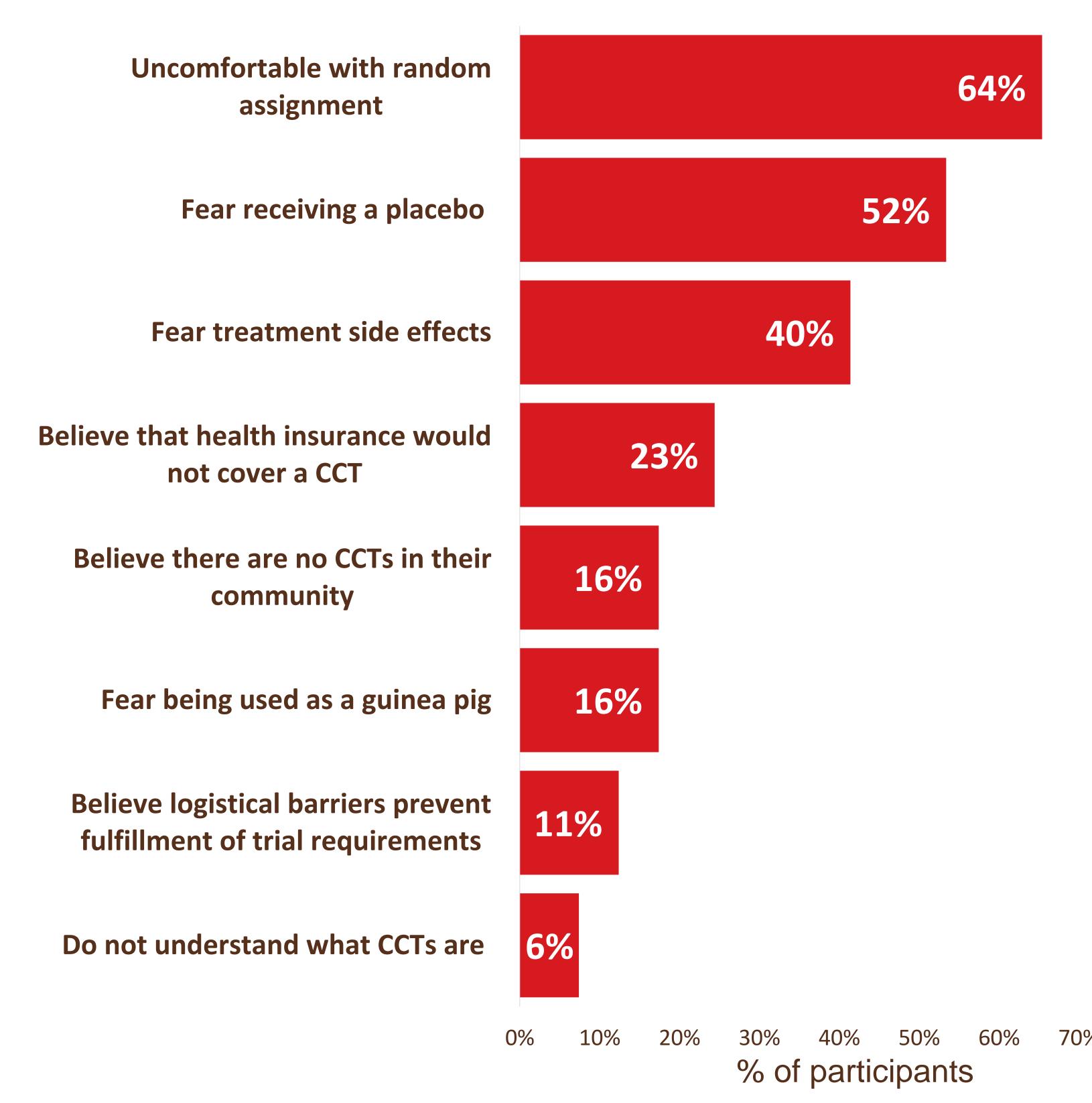
- One-third of respondents reported that their health care team spoke to them about participating in a CCT.
- There was a non-significant trend between being counseled about participating in a CCT and where cancer care was received; 47% reported that their health care team spoke to them about participating in a CCT if all or part of care was received at an academic or comprehensive cancer center or private oncology practice vs. 24% if care was received at a community hospital/cancer center (χ^2 =3.03, p=.082).

Cancer Clinical Trial (CCT) Availability and Participation



- Among all prostate cancer survivors participating in this study, about 1 in 5 reported that they had a clinical trial available to them.
- More than a third of prostate cancer participants considered a clinical trial for their treatment, and about 1 in 10 participated in clinical trial.

Beliefs and Perceptions about Cancer Clinical Trials



- Almost two thirds of participants reported feeling uncomfortable with random assignment.
- Over half of participants feared receiving a placebo; 4 in 10 feared treatment side effects.

Implications and Conclusions

- Many prostate cancer survivors are uncomfortable with random assignment to treatment in a CCT and fear receiving a potentially ineffective placebo.
- These findings also underscore that the majority of prostate cancer survivors do not recall having conversations with their providers about CCTs.
- There is a need to support comprehensive discussions between patients and providers about all prostate cancer treatment options, including CCTs, as well as innovative efforts to raise awareness and education about CCTs.
- Multivariate analysis will enhance understanding of factors contributing to clinical trial perceptions among prostate cancer survivors.

Acknowledgments

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References

- Jimenez, R., Zhang, B., Joffe, S., Nilsson, M., Rivera, L., Mutchler, J., . . . Prigerson, H. G. (2013). Clinical trial participation among ethnic/racial minority and majority patients with advanced cancer: what factors most influence enrollment? J Palliat Med, 16(3), 256-262.
- Byrne, M. M., Tannenbaum, S. L., Glück, S., Hurley, J., & Antoni, M. H. (2014). Participation in cancer clinical trials: why are patients not participating? Med Decis Making, 34(1), 116-126.



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

- The Registry is for all cancer survivors and caregivers, but also includes 10 disease-specific surveys.
- Findings contribute toward advancing research, health care and policy.
- Over 12,000 cancer survivors and caregivers are registered in the Cancer Experience Registry.

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