

Cancer-Related Distress and Cost of Care Concerns Among a Community-Based Sample of Individuals with Cancer

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Background

 Cancer survivors are at risk for financial burden due to substantial cost of care (CoC), which may impact quality of life

Aims

 The objective of this study was to examine the relationship between patients' cost of care experiences and psychosocial distress

Methods

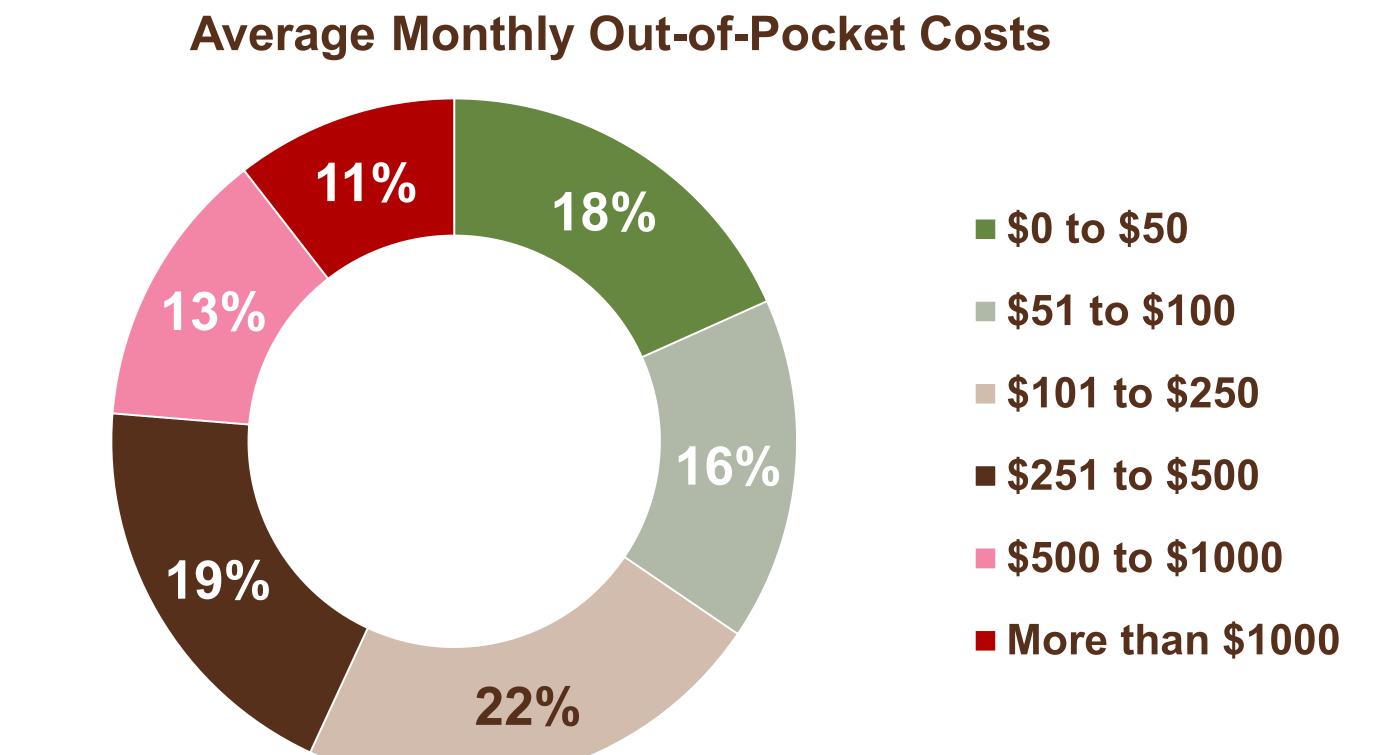
- 3,318 individuals diagnosed with cancer enrolled in Cancer Support Community's Cancer Experience Registry®, provided demographic, clinical, and cost of care information, and completed CancerSupportSource®, a 25-item distress screening tool that includes 2-item depression risk and 2-item anxiety risk subscales
- We used logistic regression to examine the role of cost of care in predicting odds of anxiety and depression risk, controlling for demographic and clinical variables significant in bivariate analyses

Participants

	M/n	SD/%
Age (years)	58	12
		Range: 18 – 83
18-44 years old	391	13%
45-64 years old	1,765	57%
≥65 years old	919	30%
Employed full-time	1,051	33%
Employed full-time ages 18-64 years (N=2,144)	909	43%
Female	2,480	75%
Non-Hispanic White	2,748	85%
Currently receiving treatment	1,828	61%
Ever experienced recurrence	782	29%
Currently in remission	1,084	49%
Time since diagnosis ≤5 <i>years</i>	2,043	69%
Ever diagnosed with metastatic cancer	583	26%
Cancer type		
Breast Cancer	1386	42%
Chronic Myeloid Leukemia	314	10%
Multiple Myeloma	313	9%
Prostate Cancer	180	5%
Lung Cancer	154	5%
Chronic Lymphocytic Leukemia	122	4%
Colorectal Cancer	103	3%
Ovarian Cancer	90	3%
* Additional cancer diagnoses include Non-Hodgki	in's Lymphoma (2%) Head and

* Additional cancer diagnoses include Non-Hodgkin's Lymphoma (2%), Head and Neck Cancer (2%), Melanoma (2%), and Endometrial Cancer (1%), among others

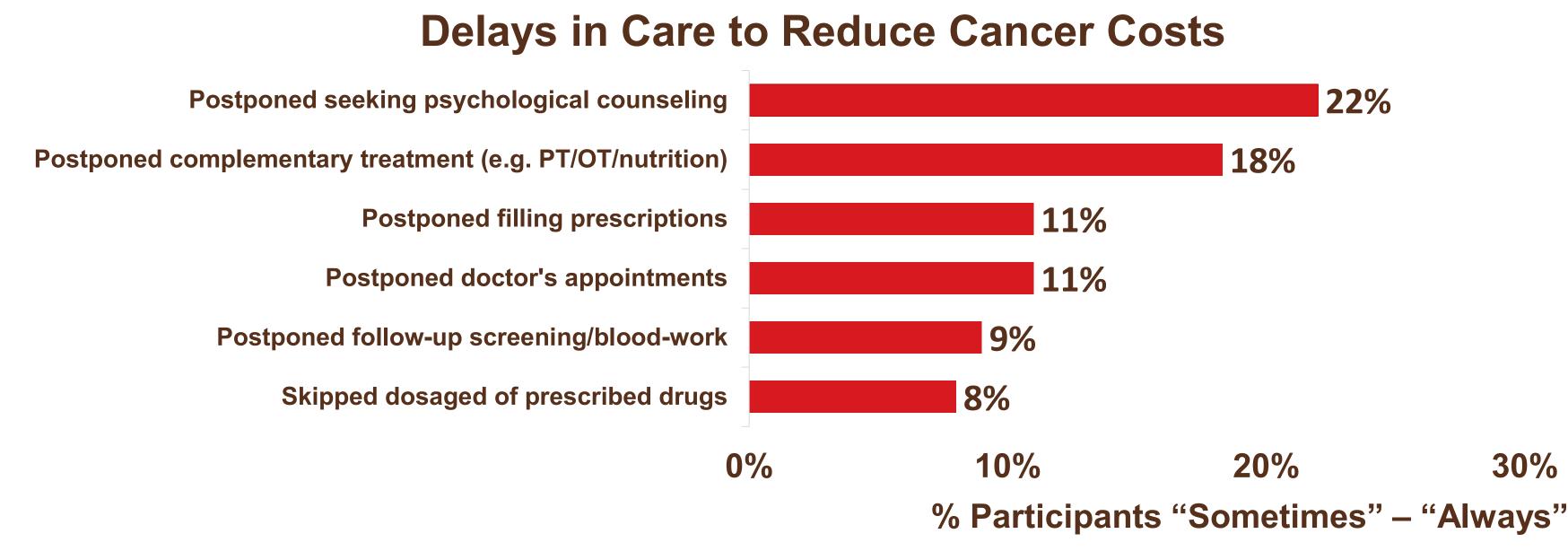
Results



 Nearly one-fourth (24%) of respondents spent more than \$500 per month out-of-pocket for costs such as prescriptions, insurance co-pays, transportation and parking, meals while at the hospital, etc.

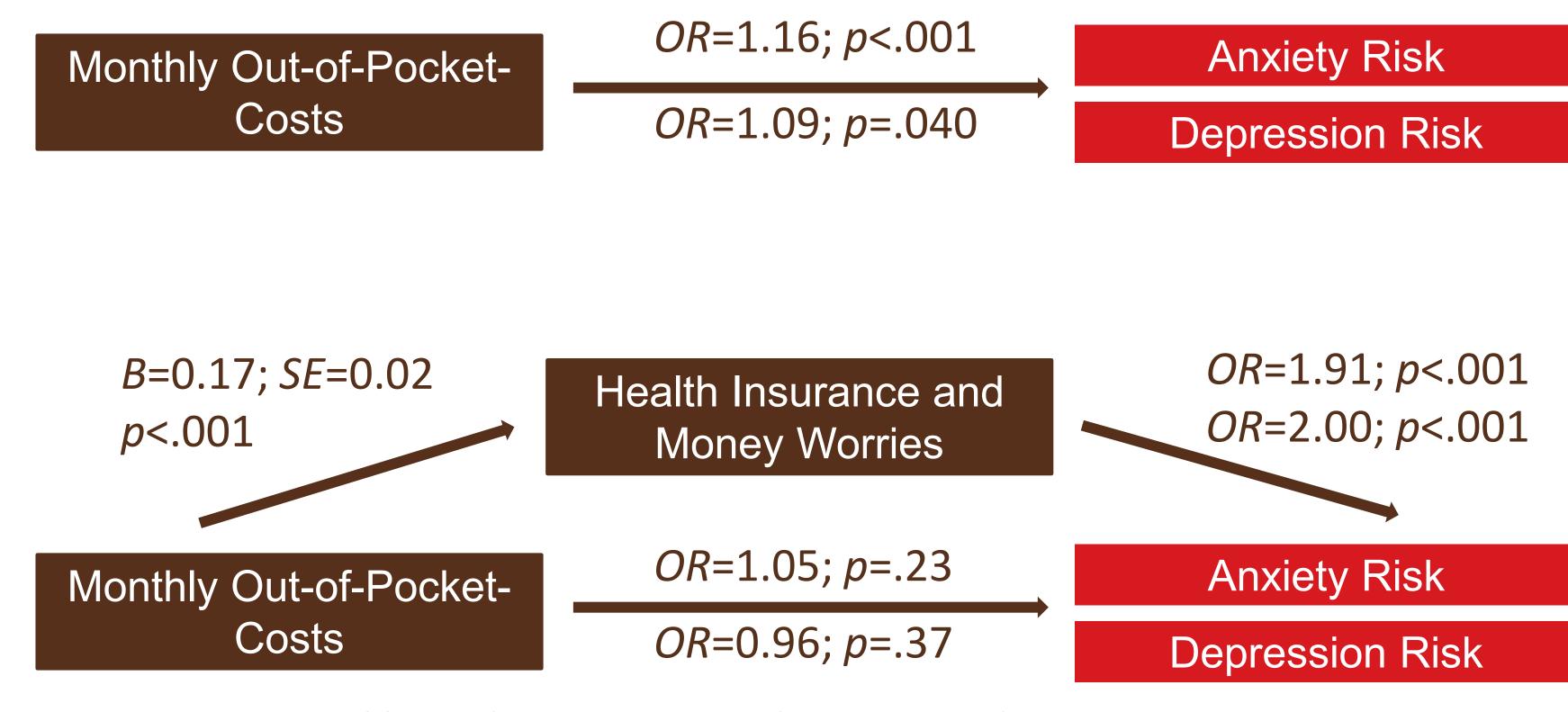
Financial Efforts to Offset Cost 32% **Depleted savings** Borrowed against or used money from retirement plan Liquidated assests Used pharmaceutical assistance programs Applied or used public assistance

• Due to the cost of cancer, 32% of respondents depleted their savings, 18% borrowed against or used retirement savings, and 12% liquidated assets



- About 1 in 5 sometimes to always postponed psychological counseling (22%) or complementary therapy (18%) to reduce cost of care
- About 1 in 10 postponed filling prescriptions (11%) or skipped doses (8%)

Financial Worries Mediate Associations Between Out-of-Pocket Costs and Anxiety and Depression Risk



* Controlling for demographic, clinical, and financial variables significant in bivariate analysis

- In multivariate logistic regression, the associations between higher out-of-pocket costs and greater odds of anxiety risk and odds of depression risk were significantly mediated by financial worries (ps<.001)
- Health insurance and money worries was associated with greater odds of anxiety risk (OR = 1.91, p < .001) and depression risk (OR = 2.00, p < .001)
- Postponing psychological counseling was associated with greater odds of anxiety risk (OR=1.31, p<.001) and depression risk (OR=1.39, p<.001)
- Having a member of the HCT discuss cost of care was associated with lower odds of depression risk (OR = 0.71, p < .05)

Implications and Conclusions

- Higher out-of-pocket costs predict increased concern about health insurance and money worries, which in turn predicts odds of anxiety risk and depression risk
- Additionally, postponing psychological counseling to reduce cost of care predicts anxiety and depression risk odds
- These findings demonstrate the need for doctor-patient cost of care communication and financial counseling to mitigate the impact of financial burden on quality-of-life, treatment, and health outcomes

Acknowledgments

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- CANCER EXPERIENCE REGISTRY A PROGRAM of the CANCER SUPPORT COMMUNITY

% Participants Endorsing "Yes"

- 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 and also includes 12 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.

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