

Lung Cancer Symptom Burden and Quality of Life: Findings from the Cancer Experience Registry®

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Background

- Given earlier diagnosis and personalized treatment, lung cancer survival rates are increasing.
- Lung cancer can be associated with significant emotional distress, which affects health-seeking behaviors and health care utilization.

Aims

 The study examines lung cancer patients' quality of life compared to other US population groups and cancer-related correlates of depression and anxiety.

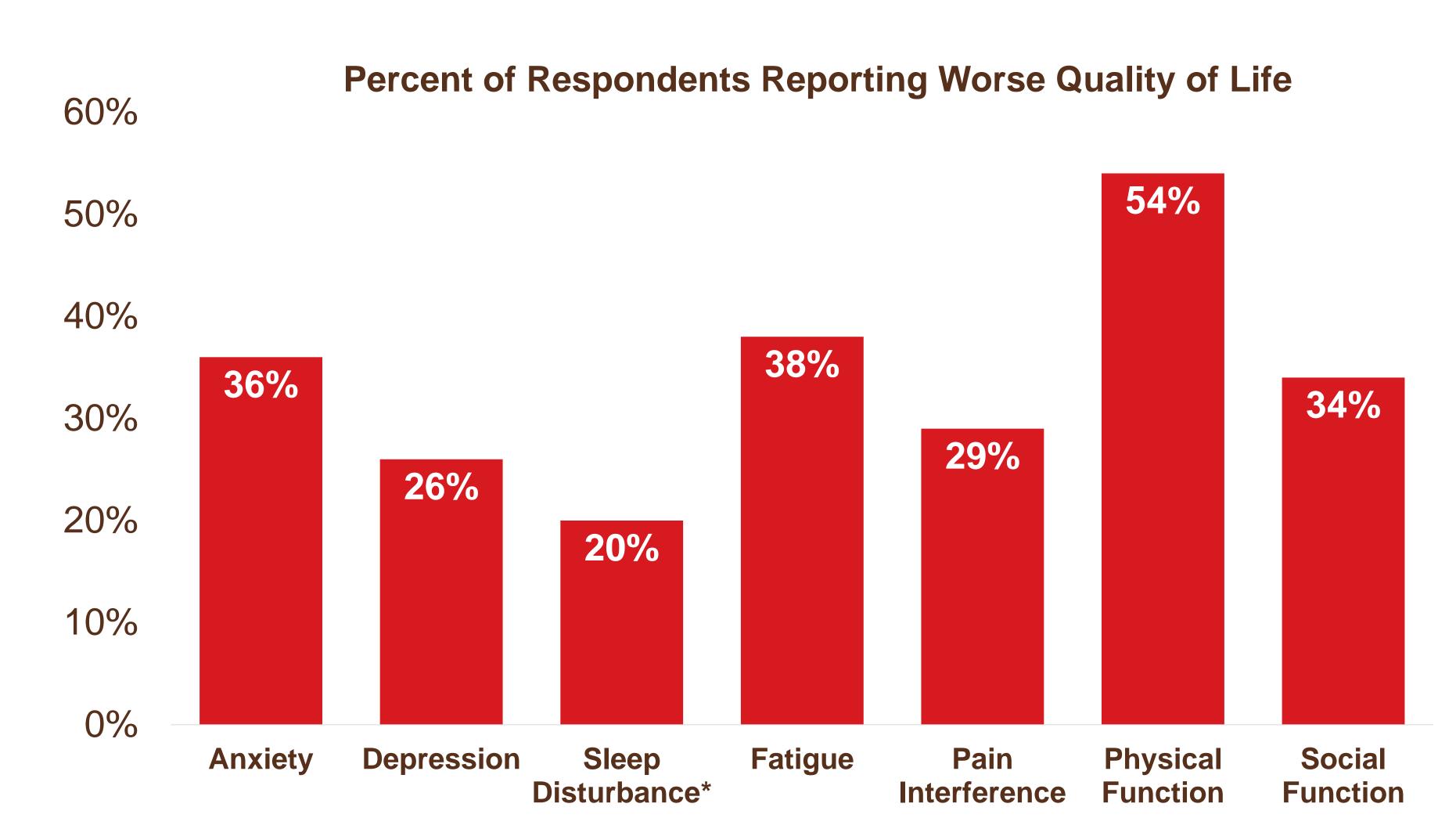
Methods

- 149 individuals with lung cancer enrolled in the Cancer Support Community's online Cancer Experience Registry and completed surveys including the Patient-Reported Outcomes Measurement Information System (PROMIS-29 v2.0).
- Using multiple regression analysis, we identify predictors of PROMIS T-scores for depression and anxiety.
- Independent variables included sociodemographic factors (age, gender, education, race), clinical history (time since diagnosis, relapse, lung cancer type, line of therapy, currently receiving treatment, type of therapies received), and symptom burden (number of comorbidities, physical functioning, fatigue, pain interference).
- Variables significant in bivariate analysis (p<.05) were included in multivariate models.

Participants

<i>N</i> = 149	M/n	SD/%	
Age	62 years	9	
	Range: 36	Range: 36 – 84 years	
Non-Hispanic White	126	86%	
Female	98	66%	
Time Since Diagnosis	3 years	5	
	Range: 0 – 49 years		
Type of Lung Cancer			
Small Cell	20	14%	
Non-Small Cell	119	86%	
Treatment Type			
Chemotherapy (ever / current)	112 / 57	76% / 38%	
Immunotherapy (ever / current)	30 / 22	20% / 15%	
Currently Receiving Treatment	79	53%	
Experienced a Relapse	34	25%	
Ever Metastatic	62	45%	

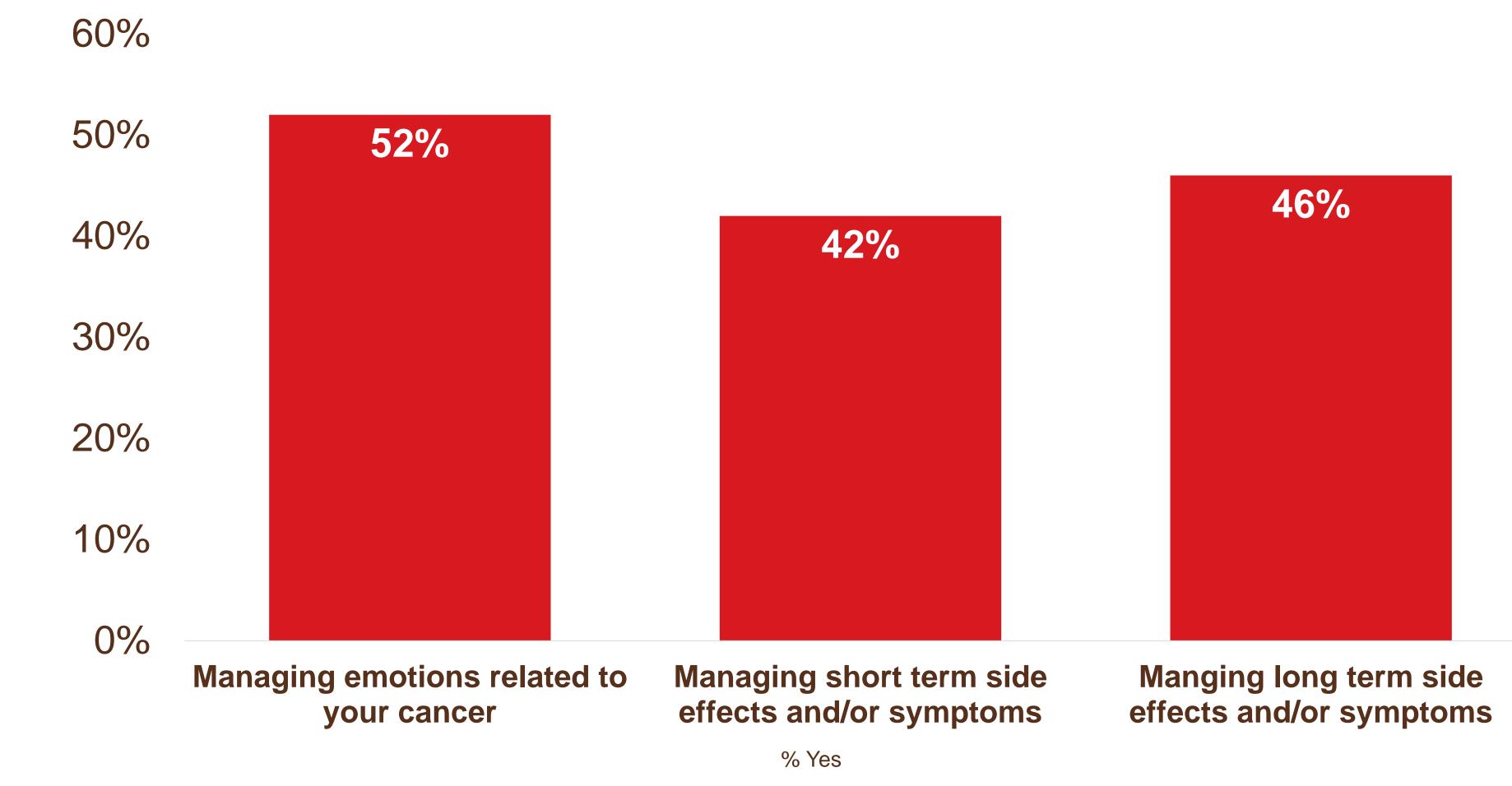
Results



Note: Relative to PROMIS-29 US population norms, except where * denotes US population group balanced to include more people with chronic illness; worse denoted as +1SD for symptom scales and -1SD for function scale

 PROMIS scores for lung cancer patients were worse than national averages (95% CI) for anxiety (54.2-57.6), fatigue (55.4-58.8), sleep disturbance (51.6-54.6), physical functioning (39.3-41.9), and social functioning (43.1-46.2).

Physical and Emotional Unmet Needs



- More than half of participants report wishing that they had received more help managing the emotions related to their cancer.
- Almost half of participants report wishing they had received help managing the short and long term side effects and/or symptoms of their cancer.

Predictors of Elevated Levels of Anxiety

	Semipartial <i>r</i>	p
Anxiety ($R^2 = .51$, $F(8,121) = 15.64$, $p<.001$)		
Female gender	.18	<.01
Time since diagnosis	19	<.01
Self-reported health	17	<.01
Fatigue	.25	<.001
Pain interference	.23	<.01

*Significant associations only are displayed

• Eight predictors explained 51% of the variance in anxiety (R^2 =.51, F(8,121)=15.64 p<.001); and, greater anxiety was associated with female gender, lesser time since diagnosis, worse self-reported health, greater fatigue, and greater pain interference.

Predictors of Elevated Levels of Depression

	Semipartial <i>r</i>	p
Depression ($R^2 = .51$, $F(9,119) = 13.74$, $p<.001$)		
Female gender	.17	<.05
Immunotherapy	18	<.01
Self-reported health	14	<.05
Fatigue	.25	<.001
Pain interference	.23	<.01
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*Significant associations only are displayed

• Nine predictors explained 51% of the variance in depression ($R^2 = .51$, F(9,119)=13.74, p<.001); and, greater depression was associated with female gender, not being treated with immunotherapy, poorer self-reported health, greater fatigue, and greater pain interference.

Implications and Conclusions

- Substantial proportions of lung cancer survivors experience worse healthrelated quality of life compared to the general U.S. population
- Symptom burden, especially fatigue and pain interference, was associated with anxiety and depression
- These results suggest the need for more comprehensive symptom management efforts throughout survivorship care, including increased access to palliative and supportive care services.

Acknowledgments

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- 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