JU Insight



Amy N. Luckenbaugh^(D), Christopher J. D. Wallis, Li-Ching Huang et al.

Correspondence: Amy N. Luckenbaugh (telephone: 615-322-2101; FAX: 615-322-8990; email: <u>amy.n.luckenbaugh@vumc.org</u>). Full-length article available at www.auajournals.org/10.1097/JU.00000000002370.

Study Need and Importance: The absolute differences in cancer-specific and overall mortality between treatment modalities (active surveillance, radical prostatectomy and radiotherapy with or without androgen deprivation therapy) for localized prostate cancer are small, and thus, treatment-related morbidity is carefully considered when making treatment decisions. Mental health outcomes in these patients have been poorly explored; therefore, we evaluated the association between prostate cancer treatment type and patientreported depression and emotional well-being over time using previously validated Centers for Epidemiologic Studies Depression (CES-D) and Medical Outcomes Study 36-item Short Form survey (SF-36) scores from the prospective population-based CEA-SAR (Comparative Effectiveness Analysis of Surgery and Radiation) study.

HE JOURNAL

ww.auajournals.org/journal/juro

What We Found: We found no effect of treatment modality on depressive symptoms (see figure). However, we identified a number of factors associated with declines in mental health regardless of treatment type, including older age, poorer health, being unmarried and having lower baseline CES-D scores.

Limitations: This is an observational study, and thus treatment choice is nonrandom, which can lead to confounding. Additionally, the CES-D and SF-36 have been validated in a general population, but not in a prostate cancer-specific population and thus may fail to detect small differences between treatment groups. Finally, using active surveillance as the referent group may contribute to the limited impact of treatment modality on mental health outcomes, as it is possible that there is a large mental health burden for those on active surveillance with prostate cancer.

Interpretation for Patient Care: Although we did not find clinically important differences in mental health outcomes for men with localized prostate cancer based on treatment received, we did identify several patient characteristics associated with poorer mental health outcomes. These characteristics may allow for early identification of patients who are most at risk for adverse mental health outcomes following prostate cancer treatment.

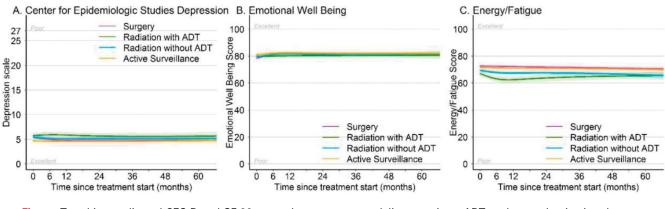


Figure. Trend in unadjusted CES-D and SF-36 scores by treatment modality over time. ADT, androgen deprivation therapy.

THE JOURNAL OF UROLOGY[®] © 2022 by American Urological Association Education and Research, Inc. https://doi.org/10.1097/JU.000000000002370 Vol. 207, 1029-1037, May 2022 Printed in U.S.A.

www.auajournals.org/jurology **1029**





Association between Treatment for Localized Prostate Cancer and Mental Health Outcomes

Amy N. Luckenbaugh[®],^{1,*} Christopher J. D. Wallis,¹ Li-Ching Huang,² Daniela Wittmann,³ Zachary Klaassen,^{4,5} Zighuo Zhao,² Tatsuki Koyama,² Aaron A. Laviana,⁶ Ralph Conwill,⁷ Michael Goodman,⁸ Ann S. Hamilton,⁹ Xiao-Cheng Wu,¹⁰ Lisa E. Paddock,¹¹ Antoinette Stroup,¹¹ Matthew R. Cooperberg,¹² Mia Hashibe,¹³ Brock B. O'Neil,¹⁴ Sherrie H. Kaplan,¹⁵ Sheldon Greenfield,¹⁵ Karen E. Hoffman,¹⁶ David F. Penson¹ and Daniel A. Barocas¹

¹Department of Urology, Vanderbilt University Medical Center, Nashville, Tennessee

² Department of Biostatistics, Vanderbilt University Medical Center, Nashville, Tennessee

³Department of Urology, University of Michigan Medical Center, Ann Arbor, Michigan

⁴Division of Urology, Medical College of Georgia at Augusta University, Augusta, Georgia

⁵Georgia Cancer Center, Augusta, Georgia

⁶Department of Surgery and Perioperative Care, Dell Medical School, Austin, Texas

⁷Office of Patient and Community Education, Patient Advocacy Program, Vanderbilt Ingram Cancer Center, Vanderbilt University Medical Center, Nashville, Tennessee ⁸Department of Epidemiology, Emory University Rollins School of Public Health, Atlanta, Georgia

⁹Department of Preventive Medicine, Keck School of Medicine at the University of Southern California, Los Angeles, California

¹⁰Department of Epidemiology, Louisiana State University Health Sciences Center New Orleans School of Public Health, New Orleans, Louisiana

¹¹Department of Epidemiology, Cancer Institute of New Jersey, Rutgers Health, New Brunswick, New Jersey

¹²Department of Urology, University of California, San Francisco, California

¹³Department of Family and Preventative Medicine, University of Utah School of Medicine, Salt Lake City, Utah

¹⁴Department of Urology, University of Utah Health, Salt Lake City, Utah

¹⁵Department of Medicine, University of California Irvine, Irvine, California

¹⁶Department of Radiation Oncology, The University of Texas MD Anderson Cancer Center, Houston, Texas

Abbreviations and Acronyms

ADT = androgen deprivation therapy

 $CaPSURE^{TM} = Cancer of the$ Prostate Strategic Urologic Research Endeavor

CEASAR = Comparative Effectiveness Analysis of Surgery and Radiation

CES-D = Centers for Epidemiologic Studies Depression

CES-D10 = 10-question version of CES-D

EPIC = Expanded Prostate Cancer Index Composite

PSA = prostate specific antigen

SF-36 = Medical Outcomes Study 36-item Short Form survey **Purpose**: We aimed to compare patient-reported mental health outcomes for men undergoing treatment for localized prostate cancer longitudinally over 5 years.

Materials and Methods: We conducted a prospective population-based analysis using the Comparative Effectiveness Analysis of Surgery and Radiation (CEA-SAR) study. Patient-reported depressive symptoms (Centers for Epidemiologic Studies Depression [CES-D]) and domains of the Medical Outcomes Study 36item Short Form survey evaluating emotional well-being and energy/fatigue were assessed through 5 years after treatment with surgery, radiotherapy (with or without androgen deprivation therapy) and active surveillance. Regression models were adjusted for outcome-specific baseline function, demographic and clinicopathological characteristics, and treatment approach.

Author Contributions: Conceptualization of the research: ANL, CJW, DW, ZK, ZZ, TK, AAL, RC, MG, ASH, XCW, LEP, AS, MRC, MH, BBO, SHK, SG, KEH, DFP, DAB; Data curation: LCH, ZZ, TK; Formal Analysis: ANL, CJW, LCH, ZK, ZZ, TK, RC, DFP, DAB; Funding acquisition: DFP, DAB; Methodology: ANL, CJW, LCH, ZZ, TK, DAB; Writing—original draft: ANL, CJW, DAB; Writing—review and editing: ANL, CJW, LCH, DW, ZK, ZZ, TK, AAL, RC, MG, ASH, XCW, LEP, AS, MRC, MH, BBO, SHK, SG, KEH, DFP, DAB: Disclaimers: There are no relevant disclaimers.

Submission History: This work was presented at the Society of Urologic Oncology meeting in December 2019 as a poster presentation. * Correspondence: Department of Urology, Vanderbilt University Medical Center, A1302 Medical Center North, Nashville, Tennessee 37232-2765 (telephone: 615-322-2101; FAX: 615-322-8990; email: amy.n.luckenbaugh@vumc.org).

Editor's Note: This article is the second of 5 published in this issue for which category 1 CME credits can be earned. Instructions for obtaining credits are given with the questions on pages 1167 and 1168.

THE JOURNAL OF UROLOGY[®] © 2022 by American Urological Association Education and Research, Inc. https://doi.org/10.1097/JU.00000000002370 Vol. 207, 1029-1037, May 2022 Printed in U.S.A.

Copyright © 2022 American Urological Association Education and Research, Inc. Unauthorized reproduction of this article is prohibited.

Accepted for publication November 22, 2021.

Support: This work was supported by the Agency for Healthcare Research and Quality (1R01HS019356, 1R01HS022640, R01CA230352); the Patient-Centered Outcomes Research Institute (CE-12-11-4667); and the National Cancer Institute (NIH/NCI grant 5T32CA106183). Data management was facilitated by Vanderbilt University's Research Electronic Data Capture (REDCap) system, which is supported by the Vanderbilt Institute for Clinical and Translational Research grant (UL1TR000011 from NCATS/NIH). The funders had no role in the study design, data collection, and analysis, decision to publish, or preparation of the manuscript.

Conflict of Interest: The authors have no disclosures relevant to this work.

Results: A total of 2,742 men (median [quartiles] age 64 [59–70]) met inclusion criteria. Baseline depressive symptoms, as measured by the CES-D, were low (median 4, quartiles 1–8) without differences between groups. We found no effect of treatment modality on depressive symptoms (p=0.78), though older age, poorer health, being unmarried and baseline CES-D score were associated with declines in mental health. There was no clinically meaningful association between treatment modality and scores for either emotional well-being (p=0.81) or energy/fatigue (p=0.054).

Conclusions: This prospective, population-based cohort study of men with localized prostate cancer showed no clinically important differences in mental health outcomes including depressive symptoms, emotional wellbeing, and energy/fatigue according to the treatment received (surgery, radiotherapy, or surveillance). However, we identified a number of characteristics associated with worse mental health outcomes including: older age, poorer health, being unmarried, and baseline CES-D score which may allow for early identification of patients most at risk of these outcomes following treatment.

Key Words: prostatic neoplasms, mental health, quality of life

For patients diagnosed with localized prostate cancer, guideline-recommended treatment options include active surveillance, radical prostatectomy and radiotherapy, based on risk-stratification. Absolute differences in cancer-specific and overall mortality between these approaches are small,^{1,2} thus treatment-related morbidity is paramount in treatment decision making. While health-related quality of life outcomes following prostate cancer treatments such as urinary symptoms, erectile dysfunction and bowel symptoms are well reported³⁻⁶ and the burden of other interventions to manage treatment-related complications is increasingly recognized,⁷⁻¹⁰ less attention has been paid to the association between treatment for localized prostate cancer and mental health outcomes.

While mental health outcomes of treatment have been poorly explored among patients with prostate cancer, one small (368) cross-sectional study of Black men with prostate cancer found a relatively high prevalence of major depressive symptoms (33%), with an increased likelihood among those who underwent radiotherapy (odds ratio 2.38, 95% confidence interval 1.02-5.51). However, these data are limited by the lack of generalizability, difficulties with causation with a cross-sectional study design and the need for reproducibility.¹¹ In contrast, the association between treatment and mental health outcomes is better established in both breast cancer and colorectal cancer,^{12,13} in which sexual dysfunction and body image concerns contribute to emotional distress and worsening psychosocial function over time.

In this context, we evaluated the association between prostate cancer treatment and patient-reported depression and emotional well-being over time applying previously validated Centers for Epidemiologic Studies Depression (CES-D) and Medical Outcomes Study 36-item Short Form survey (SF-36) scores to data from the prospective population-based Comparative Effectiveness Analysis of Surgery and Radiation (CEASAR) study.

MATERIALS AND METHODS

Study Population

The CEASAR study enrolled men with clinically localized prostate cancer (cT1–cT2, prostate specific antigen [PSA] <50 ng/dL) from 5 population-based SEER (Surveillance, Epidemiology, and End Results) program registries and the observational Cancer of the Prostate Strategic Urologic Research Endeavor (CaPSURETM)prostate cancer registry from 2011–2012, as previously described.^{14,15} Institutional Review Board approval was obtained from Vanderbilt University Medical Center (coordinating center, IRB No. 110299) and from each participating site.

Participants completed surveys at baseline, 6 months and 1, 3 and 5 years following enrollment (with the final survey being completed in September 2017). Data regarding tumor characteristics, treatment choice and treatment dates were obtained from medical chart abstraction 1 year after enrollment.^{14,15} Any treatment received after 1 year was based on patient report. Survival was determined from vital status followup data obtained from the SEER and CaPSURE registries.

Exposure

The exposure of interest was the primary treatment modality, categorized as active surveillance, surgery, radiation with androgen deprivation therapy (ADT) and radiation without ADT. Treatment modality was determined primarily based on 1-year chart abstraction, supplemented by patient report.

Outcomes

Depressive symptoms were assessed by using a 10-question version of the previously validated CES-D scale (CES-D10).^{16,17} The CES-D10 has been validated across diverse populations.^{18,19} To reduce respondent burden, the CES-D was modified to a 9-question version. We adjusted the overall CES-D score to reflect this difference (from a standard score of 30 points to 27 points.) Domain scores ranged from 0–27, where 27 indicated more severe depressive symptoms. Notably, to our knowledge, a clinically meaningful difference on this scale has not been described though scores above 19 have strong specificity and positive predictive value for major depression.

The validated SF-36 was used to evaluate emotional well-being and energy and fatigue.^{20,21} For the SF-36,

domain scores range from 0-100 with 100 indicating the best function. The results of these domain scores were analyzed over time for each treatment modality. We interpreted results based on previously determined minimally clinically important differences of 6 and 9 for emotional well-being and energy and fatigue, respectively.^{22,23}

Statistical Analysis

Clinical and sociodemographic characteristics were evaluated by treatment modality, categorized as active surveillance, surgery, radiation with ADT and radiation without ADT. Treatment group differences were assessed using Kruskal-Wallis tests and χ^2 tests for continuous and categorical variables, respectively.

The study endpoints of CES-D score and SF-36 domain scores for emotional well-being and energy and vitality were reported as adjusted mean score differences (with 95% confidence intervals). To further evaluate the associations between treatments and measures of mental health over time, using the longitudinal survey data, we fit multivariable longitudinal linear regression models for CES-D, emotional well-being and energy and vitality adjusting for time since treatment (continuous, restricted cubic splines using 4 knots), age at diagnosis (continuous, restricted cubic splines using 3 knots), race (White, Black, Hispanic, Asian, other), education (less than high school, high school graduate, some college, college graduate, graduate/professional school), marital status (not married, married), comorbidity as measured with total burden index for prostate cancer-TIBI-CaP²⁴ (categorical: 0-2, 3-4, 5 or more), income (less than \$30,000, \$30,001-\$50,000, \$50,001-\$100,000, more than \$100,000), insurance status (Medicare; private or health maintenance organization; and Veterans Administration, military, Medicaid, other or uninsured), D'Amico risk category (low, intermediate, high), site (Utah, Atlanta, Los Angeles, Louisiana, New Jersey, CaPSURE), baseline physical functioning (continuous, linear), baseline general health (continuous, linear), baseline social support (continuous, linear), baseline participatory decision making scale (continuous, linear), baseline sexual function score (continuous, linear), time-varying sexual function scores (at 6 months, 1, 3 and 5 years, continuous, linear) and corresponding baseline value of the outcome (continuous, restricted cubic splines using 3 knots). Covariates were obtained from patient-reported surveys and chart abstraction, as appropriate.

Comparing between treatment modalities, we utilized active surveillance as the referent. To allow for variable estimation of treatment effect at different time points, we included the interaction terms between treatment and time since treatment in the models. In all models, to account for the correlation due to repeated measures obtained on the same subjects from multiple time points, the Huber-White method was implemented by the robcov function in the rms R package to estimate the variancecovariance matrices.^{25,26} The results of models were reported as mean differences between treatment groups and the associated 95% confidence intervals. All missing covariate values were imputed 10 times using the MICE (multiple imputation using chained equations) implemented by the aregImpute function in the rms R package. The Expanded Prostate Cancer Index Composite

(EPIC)-26 sexual function score was missing at 4%, 9%, 10%, 20% and 28% at each time point, respectively. Income was missing in 11%; however, all other variables had less than 5% missing. To graphically represent the trends in CES-D and SF-36 scores, we fit simpler models that included time since treatment start and treatment modality, along with their interaction terms. Statistical significance was considered for all 2-sided p values <5%. All analyses were conducted using R version 4.0.2 (R Foundation for Statistical Computing, Vienna, Austria).

RESULTS

Among 2,742 patients included in the analysis, 372 (13.7%) underwent active surveillance, 1,419 (51.8\%) underwent surgery, 630 (23.0%) underwent radiation therapy without ADT and 321 (11.7%) underwent radiation with ADT (fig. 1). Differences in baseline characteristics across the treatment groups are in keeping with prior reports: men who underwent surgery as their primary treatment type were younger with fewer comorbidities while men undergoing active surveillance were more likely to have low risk disease characteristics, including lower PSA, clinical stage T1 and a low D'Amico risk category. Men with features of high risk disease (Gleason 8, 9, 10, T2, high D'Amico risk category) were more likely to undergo radiation therapy with ADT (table 1). Baseline urinary function, urinary incontinence and bowel function domain scores were similar across all treatment groups. In contrast, baseline sexual function domain scores were higher for men treated with radical prostatectomy compared to those treated with radiation or active surveillance (table 1).

At baseline, the median CES-D score in this cohort was 4 (interquartile range 1-8), indicating a low prevalence of depressive symptoms (table 1). We found no evidence of a clinically meaningful treatmentrelated effect on longitudinal assessments of depressive symptoms measured with the CES-D score, whether assessed continuously or dichotomized at 9 (tables 2 and 3). In addition to higher baseline CES-D score (p < 0.001), on multivariable analysis, significant predictors of decline in CES-D score were older age (p=0.001), higher comorbidity (p < 0.001), poorer overall health (p=0.001) and physical function (p=0.008), being unmarried (p=0.02), lower income (p=0.002) and lower baseline participatory decision making score (p=0.003). Interestingly, social support (p=0.39) and education (p=0.12) were not associated with worsening CES-D scores, nor were race (p=0.38), insurance status (p=0.95), D'Amico risk group (p=0.99) or registry site (p=0.11).

Assessing emotional well-being and energy/fatigue domains of the SF-36, we found that while baseline scores were overall quite high, lower scores were reported among those undergoing radiotherapy (with or without ADT), a difference which persisted over

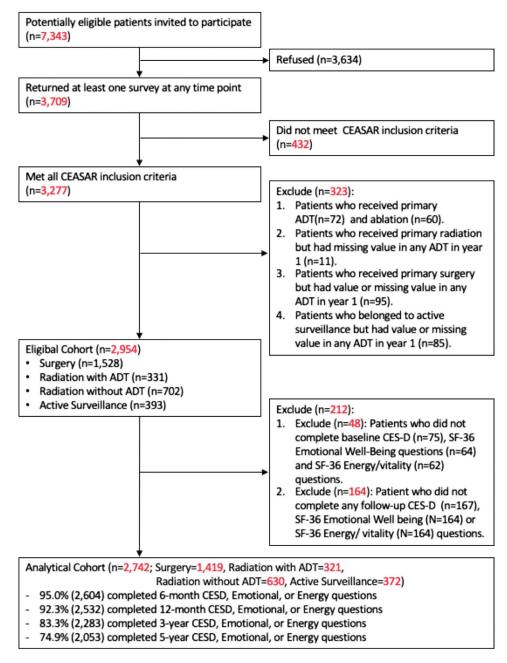


Figure 1. Flow chart demonstrating inclusion criteria for the study.

time (fig. 2). However, on adjusted and unadjusted analysis treatment type was not associated with a clinically significant difference in longitudinal assessments of SF-36 emotional well-being or energy/fatigue scores at 6 months or 1, 3 or 5 years following treatment initiation, despite statistically significant differences (table 2 and supplementary table 1, <u>https://www. jurology.com</u>). Minor declines in SF-36 energy/fatigue scores at 6 and 12 months among those treated with radiation and ADT (fig. 2) are unlikely to be clinically meaningful and were not statistically significant (table 2). Notably, there was a statistically significant difference in the SF-36 energy/fatigue domain between patients who underwent surgery compared to radiotherapy, though this failed to meet the threshold for a clinically important difference and also diminished with time. Further pairwise testing between surgery and radiotherapy, and between radiotherapy with and without ADT demonstrated no clinically meaningful differences in the captured mental health related outcomes (supplementary table 2, <u>https://www.jurology.</u> <u>com</u>). In addition to baseline SF-36 emotional wellbeing (p <0.001), significant predictors of decline in SF-36 emotion well-being scores were older age (p <0.001), higher comorbidity (p <0.001), lower income (p=0.004), general health (p=0.02) and physical



Table 1. Baseline demographic, socioeconomic and disease characteristics by treatment type

	Active Surveillance		Surgery		Radiation with ADT		Radiation without ADT		p Value
No. pts	372	(00 70)	1,419	(57	321	(05 7 5)	630	(00 70)	0.00
Median yrs age at diagnosis (IQR) No. race (%):	67	(62—72)	62	(57—66)	70	(65—74)	67	(62—72)	<0.001 0.001
White	296	(80)	1,069	(76)	220	(69)	481	(76)	0.001
Black	39	(11)	165	(12)	56	(18)	103	(16)	
Hispanic	22	(6)	113	(8)	25	(8)	27	(4)	
Asian	9	(2)	40	(3)	13	(4)	10	(2)	
Other No. TIBI-CaP category (%):	5	(1)	21	(1)	4	(1)	8	(1)	<0.001
0-2	95	(26)	452	(34)	49	(16)	133	(22)	<0.001
3-4	142	(39)	586	(44)	114	(37)	253	(43)	
5 or more	126	(35)	307	(23)	147	(47)	206	(35)	
No. \$ income (%):									< 0.001
<30,000	63	(18)	211	(17)	101	(36)	124	(22)	
30,001-50,000	79 105	(23)	213	(17)	67	(24)	129	(23)	
50,001—100,000 >100,000	105 97	(31) (28)	422 422	(33) (33)	67 47	(24) (17)	174 126	(31) (23)	
No. education (%):	57	(20)	722	(00)	17	(17)	120	(20)	<0.001
Less than high school	25	(7)	113	(8)	52	(17)	59	(10)	
High school graduate	66	(18)	271	(20)	67	(22)	121	(21)	
Some college	77	(21)	298	(22)	68	(22)	145	(25)	
College graduate	91 104	(25)	324	(24)	57	(19)	135	(23)	
Graduate/professional school No. marital status (%):	104	(29)	332	(25)	62	(20)	129	(22)	<0.001
Not married	67	(19)	216	(16)	70	(23)	153	(26)	<0.001
Married	294	(81)	1,120	(84)	235	(77)	437	(74)	
No. employment status (%):									< 0.001
Full time	160	(43)	876	(62)	80	(25)	250	(40)	
Retired/part time/unemployed	210	(57)	530	(38)	235	(75)	371	(60)	<0.001
No. insurance status (%): Medicare	214	(58)	464	(33)	221	(69)	388	(62)	<0.001
Private/health maintenance organization	142	(38)	884	(62)	78	(24)	211	(34)	
Veterans Administration/military/Medicaid/none	16	(4)	68	(5)	22	(7)	29	(5)	
Median PSA at diagnosis, corrected (IQR)	5.2	(3.9—7.0)	5.1	(4.2-6.8)	7.0	(4.9—11.3)	5.5	(4.3-7.3)	< 0.001
No. clinical tumor stage (%):	000	(0.4)	1 000	(75)	04.0	(00)	404	(70)	<0.001
T1 T2	309 57	(84) (16)	1,069 347	(75) (25)	218 103	(68) (32)	491 138	(78) (22)	
No. biopsy Gleason score (%):	57	(10)	347	(23)	103	(32)	130	(22)	<0.001
6 or less	330	(89)	717	(51)	44	(14)	334	(53)	0.001
3+4	33	(9)	437	(31)	113	(35)	197	(32)	
4+3	6	(2)	147	(10)	57	(18)	58	(9)	
8, 9, 10	1	(0)	114	(8)	107	(33)	36	(6)	0.001
No. D'Amico risk category (%): Low risk	293	(79)	614	(43)	28	(9)	305	(49)	<0.001
Intermediate risk	67	(18)	598	(43)	144	(45)	261	(43)	
High risk	10	(3)	205	(14)	149	(46)	60	(10)	
No. site (%):									< 0.001
Utah	56	(15)	119	(8)	24	(7)	52	(8)	
Atlanta	47	(13)	189	(13)	25	(8)	188	(30)	
Los Angeles Louisiana	116 93	(31) (25)	409 356	(29) (25)	76 141	(24) (44)	100 143	(16) (23)	
New Jersey	28	(23)	241	(23)	36	(44)	143	(20)	
CaPSURE	32	(9)	105	(7)	19	(6)	19	(20)	
Median SF-36 Physical Function (IQR)	95	(80—100)	100	(85-100)	90	(65-100)	90	(75-100)	< 0.001
Median SF-36 Emotional Well-Being (IQR)	88	(72-92)	84	(68-92)	88	(72-92)	84	(72-92)	800.0
Median SF-36 Energy & Fatigue (IQR)	75	(60-85)	75	(60-85)	70	(55-85)	75	(58-85)	< 0.001
Median depression (CES-D10) score (IQR) Median social support, median (IQR)	3 95	(1—6) (75—100)	4 95	(1—8) (75—100)	4 95	(2—8) (75—100)	4 95	(1—8) (70—100)	0.05 0.046
Median participatory decision making (IQR)	95 86	(68-96)	95 86	(75-100) (71-93)	95 79	(64-89)	95 79	(68-92)	<0.040
Median prostate cancer-specific burden, baseline (IQR)		(0.0-37.1)		(8.6-45.7)		(5.7-42.9)		5.7-42.9	< 0.001
Median EPIC-26 sexual function, baseline (IQR)	75	(42-89)	80	(38—95)	50	(12-80)	65	(32—85)	< 0.001
Median EPIC-26 urinary incontinence, baseline (IQR)	100	(85—100)	100	(79—100)	100	(79—100)	100	(85—100)	0.2
Median EPIC-26 urinary irritative, baseline (IQR)	88	(75-94)	88	(75-100)	88	(75-94)	88	(75-100)	0.011
Median EPIC-26 bowel function, baseline (IQR) Median EPIC-26 hormonal domain score at baseline (IQR)	100	(92-100)	100	(96—100)	100	(92-100)	100	(92-100)	0.012
weulan Eric-zo normonal uomain score at baseine (IUR)	95	(85—100)	95	(85—100)	90	(75—95)	95	(85—100)	< 0.001

function (p <0.001), social support (p=0.001) and baseline participatory decision making scores (p=0.002). Notably, unlike for changes in CES-D

score, marital status was not associated with changes in emotional well-being measured by SF-36 (p=0.17), nor was race (p=0.26), insurance

	Surgery vs Active Surveillance			Radiation	(+ADT) vs Active	Surveillance	Radiation (no ADT) vs Active Surveillance		
Time (yrs)	Effect 95% CI		p Value	Effect	95% CI	p Value	Effect	95% CI	p Value
				CES-D10)				
0.5	-0.1	(-0.8, 0.5)	0.669	0.2	(-0.5, 1.0)	0.5	0.4	(-0.3, 1.0)	0.264
1	-0.5	(-0.9, -0.1)	0.024	-0.3	(-0.9, 0.3)	0.351	0	(-0.5, 0.4)	0.881
3	-0.3	(-0.7, 0.2)	0.316	0.1	(-0.6, 0.9)	0.713	0	(-0.6, 0.5)	0.873
5	-0.5	(-1.0, 0.0)	0.067	-0.1	(-0.9, 0.7)	0.78	0	(-0.6, 0.6)	0.959
				Emotional Wel	II-Being				
0.5	1.9	(-0.1, 4.0)	0.065	0.6	(-1.7, 3.0)	0.598	0.4	(-1.7, 2.5)	0.721
1	2.4	(1.0, 3.8)	< 0.001	1.8	(-0.3, 3.9)	0.094	0.6	(-0.9, 2.2)	0.432
3	1.7	(0.0, 3.3)	0.044	0	(-2.4, 2.4)	0.988	0.8	(-1.0, 2.6)	0.376
5	1.2	(-0.6, 3.0)	0.187	0.4	(-2.4, 3.2)	0.769	0.6	(-1.4, 2.6)	0.562
				Energy/Fati	que				
0.5	4.7	(2.2, 7.2)	< 0.001	-1	(-4.0, 2.0)	0.528	0.3	(-2.3, 2.9)	0.84
1	3	(1.4, 4.6)	< 0.001	-0.9	(-3.3, 1.5)	0.455	-1.5	(-3.3, 0.2)	0.085
3	2.1	(0.3, 3.9)	0.023	-2.2	(-5.0, 0.5)	0.113	-0.2	(-2.2, 1.8)	0.845
5	1.5	(-0.4, 3.4)	0.134	-1.1	(-4.0, 1.8)	0.455	-1.8	(-4.0, 0.5)	0.128

Table 2. Effect of treatment modality on longitudinal assessment of CES-D and SF-36 emotional well-being and energy/fatigue scores at 6 months and 1, 3 and 5 years following index, adjusted for the effect of patient demographic, tumor and baseline functional characteristics

status (p=0.88), D'Amico risk group (p=0.62) or registry site (p=0.24).

DISCUSSION

In this population-based, prospective cohort study of men with localized prostate cancer, we found no clinically meaningful association between treatment approach (including active surveillance, radical prostatectomy and radiotherapy) and measures of mental health including depressive symptoms (captured using the validated CES-D) and emotional well-being and energy/fatigue (captured as domains of the validated SF-36). These findings were consistent with our hypothesis that treatment type would not impact overall mental health outcomes in men with localized prostate cancer. However, on multivariable analysis we did identify baseline characteristics associated with declining emotional well-being following prostate cancer treatment including older age, poor overall health, unmarried status, and worse baseline depression and emotional well-being symptoms. These characteristics may allow clinicians to identify patients most at risk of declines in mental health following prostate cancer diagnosis and treatment to target interventions to address these issues.

To our knowledge, this is the first prospective evaluation of mental health outcomes in men with

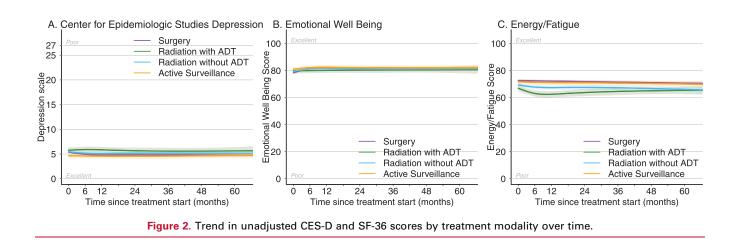
localized prostate cancer. While previous work has demonstrated an increased utilization of antidepressants following diagnosis for patients who received surgery or radiotherapy, but not active surveillance, this relied on administrative records and prescriptions as a proxy for symptoms.²⁷ In contrast, and in keeping with our findings, there does not appear to be an increased risk of suicide amongst patients diagnosed with prostate cancer, whereas there are increased risks of suicide amongst patients with other malignancies.²⁸

Although we did not demonstrate a relationship between treatment type and mental health outcomes in men with localized prostate cancer, we did find several factors predictive of declining mental health outcomes. Consistent with research in other malignancies, older, unmarried patients had declining emotional well-being in our analysis. In the bladder cancer population, prior research has demonstrated those at risk for suicidal death were typically elderly, unmarried men.²⁹ An additional vulnerable population we identified was those with poor depression and poor emotional well-being symptoms prior to treatment. It is imperative that urologists seize available opportunities to identify and intervene in patients with mental health concerns both at the time of diagnosis and during followup. In addition to the importance of addressing patient distress and morbidity

Table 3.

Time (yrs)	Surgery vs Active Surveillance			Radiation (+A	ADT) vs Active S	urveillance	Radiation (no ADT) vs Active Surveillance		
	Odds Ratio	95% CI	p Value	Odds Ratio	95% CI	p Value	Odds Ratio	95% CI	p Value
				CES-D >9 (bina	ary)				
0.5	0.9	(0.6, 1.6)	0.829	1.3	(0.7, 2.2)	0.448	1.4	(0.8, 2.5)	0.181
1	0.9	(0.6, 1.3)	0.556	1	(0.6, 1.5)	0.858	1.1	(0.8, 1.6)	0.528
3	0.8	(0.6, 1.3)	0.404	1.4	(0.8, 2.3)	0.236	0.9	(0.5, 1.3)	0.484
5	0.8	(0.6, 1.2)	0.364	1	(0.6, 1.8)	0.983	1.3	(0.8, 2.0)	0.323

Copyright © 2022 American Urological Association Education and Research, Inc. Unauthorized reproduction of this article is prohibited.



associated with these symptoms, prior work has demonstrated that significant mental health care utilization is independently associated with worse cancerspecific and all-cause mortality.³⁰ While psychological interventions are beyond the scope of most clinicians treating prostate cancer, we ought to appropriately screen and subsequently refer those at risk.

Our findings should be considered in the context of several limitations. First, as this is an observational study, treatment choice is nonrandom and thus there is the potential for confounding by indication. However, given the baseline similarities between groups, and the longitudinal nature of assessment and control for clinical factors, it is unlikely that this explains the findings. Second, due to the nature of the survey employed, we utilized a modified CES-D 10 with patients completing only 9 questions rather than 10 questions on the recommendation of our psychometrician in order to reduce respondent burden as other included instruments captured overlapping concepts. We adjusted the overall CES-D score to reflect this difference (from a standard score of 30 points to 27 points.) Similarly, the SF-36 and CES-D have been validated in a general population, but may not detect minute differences in our population of overall healthy men, and should not be used in isolation for diagnosing depression or mental health changes. Third, in the context of a finding of no significant differences, we must consider the potential for type II error. However, given the small differences noted which did not meet established threshold for clinically meaningful differences, increases in sample size are unlikely to change study conclusions. Additionally, many patients with low risk disease in the CEASAR study received active intervention which, while common at the time, is not reflective of current practice patterns which now favor active surveillance in this cohort. Finally, we must not underestimate

the mental health burden of being diagnosed with prostate cancer including for those who choose active surveillance as their primary treatment strategy. Thus, using the active surveillance group as the referent group may contribute to the limited impact of treatment modality on mental health outcomes. Perhaps a more appropriate referent group, and consideration for future work, would be a group of healthy age-matched men without a cancer diagnosis.

These limitations notwithstanding, in this population-based, prospective cohort study of men with localized prostate cancer we found no clinically meaningful association between treatment approach (including active surveillance, radical prostatectomy and radiotherapy) and measures of mental health including depressive symptoms (captured using the validated CES-D) and emotional well-being and energy/fatigue (captured as domains of the validated SF-36). We further identified characteristics of patients with a higher likelihood of declining mental health following prostate cancer diagnosis, independent of treatment approach, including older age, being unmarried, worse overall health and worse baseline mental health.

CONCLUSIONS

Careful evaluation of patients at risk for adverse mental health outcomes is warranted among all treatment groups, and appropriate psychiatric assistance should be provided to these patients to optimize the comprehensive care we provide to prostate cancer patients.

ACKNOWLEDGMENT

This article was written on behalf of all CEASAR investigators.

REFERENCES

- Hamdy FC, Donovan JL, Lane JA et al: 10-Year outcomes after monitoring, surgery, or radiotherapy for localized prostate cancer. N Engl J Med 2016; **375**: 1415.
- Wallis CJ, Saskin R, Choo R et al: Surgery versus radiotherapy for clinically-localized prostate cancer: a systematic review and meta-analysis. Eur Urol 2016; **70:** 21.
- Wallis CJD, Glaser A, Hu JC et al: Survival and complications following surgery and radiation for localized prostate cancer: an international collaborative review. Eur Urol 2018; 73: 11.
- Resnick MJ, Koyama T, Fan KH et al: Long-term functional outcomes after treatment for localized prostate cancer. N Engl J Med 2013; 368: 436.
- Barocas DA, Alvarez J, Resnick MJ et al: Association between radiation therapy, surgery, or observation for localized prostate cancer and patient-reported outcomes after 3 years. JAMA 2017; **317**: 1126.
- Hoffman KE, Penson DF, Zhao Z et al: Patientreported outcomes through 5 years for active surveillance, surgery, brachytherapy, or external beam radiation with or without androgen deprivation therapy for localized prostate cancer. JAMA 2020; **323**: 149.
- Wallis CJ, Herschorn S, Saskin R et al: Complications after radical prostatectomy or radiotherapy for prostate cancer: results of a population-based, propensity score-matched analysis. Urology 2015; 85: 621.
- Wallis CJ, Cheung P, Herschorn S et al: Complications following surgery with or without radiotherapy or radiotherapy alone for prostate cancer. Br J Cancer 2015; **112**: 977.
- Wallis CJ, Mahar A, Cheung P et al: New rates of interventions to manage complications of modern prostate cancer treatment in older men. Eur Urol 2015; 69: 933.
- Wallis CJ, Mahar AL, Cheung P et al: Hospitalizations to manage complications of modern prostate cancer treatment in older men. Urology 2016; 96: 142.
- Kinlock BL, Parker LJ, Howard DL et al: Prevalence and correlates of major depressive symptoms among Black men with prostate cancer. Ethn Dis 2017; 27: 429.

- Congard A, Christophe V, Duprez C et al: The self-reported perceptions of the repercussions of the disease and its treatments on daily life for young women with breast cancer and their partners. J Psychosoc Oncol 2019; **37**: 50.
- Reese JB, Handorf E and Haythornthwaite JA: Sexual quality of life, body image distress, and psychosocial outcomes in colorectal cancer: a longitudinal study. Support Care Cancer 2018; 26: 3431.
- Barocas DA, Chen V, Cooperberg M et al: Using a population-based observational cohort study to address difficult comparative effectiveness research questions: the CEASAR study. J Comp Eff Res 2013; 2: 445.
- Cooperberg MR, Broering JM, Litwin MS et al: The contemporary management of prostate cancer in the United States: lessons from the Cancer of the Prostate Strategic Urologic Research Endeavor (CapSURE), a national disease registry. J Urol 2004; **171:** 1393.
- Irwin M, Artin KH and Oxman MN: Screening for depression in the older adult: criterion validity of the 10-item Center for Epidemiological Studies Depression scale (CES-D). Arch Intern Med 1999; 159: 1701.
- Smarr KL and Keefer AL: Measures of depression and depressive symptoms: Beck Depression Inventory-II (BDI-II), Center for Epidemiologic Studies Depression scale (CES-D), Geriatric Depression Scale (GDS), Hospital Anxiety and Depression scale (HADS), and Patient Health Questionnaire-9 (PHQ-9). Arthritis Care Res (Hoboken), suppl., 2011; 63: S454.
- Zhang W, O'Brien N, Forrest JI et al: Validating a shortened depression scale (10 item CES-D) among HIV-positive people in British Columbia, Canada. PLoS One 2012; 7: e40793.
- Baron EC, Davies T and Lund C: Validation of the 10-item Centre for Epidemiological Studies Depression scale (CES-D-10) in Zulu, Xhosa and Afrikaans populations in South Africa. BMC Psychiatry 2017; 17: 6.
- McHorney CA, Ware JE Jr and Raczek AE: The MOS 36-Item Short-Form Health Survey (SF-36): II. Psychometric and clinical tests of validity in measuring physical and mental health constructs. Med Care 1993; **31**: 247.

- Ware JE Jr and Sherbourne CD: The MOS 36item Short-Form Health Survey (SF-36). I. Conceptual framework and item selection. Med Care 1992; **30**: 473.
- Skolarus TA, Dunn RL, Sanda MG et al: Minimally important difference for the Expanded Prostate Cancer Index Composite short form. Urology 2015; 85: 101.
- Jayadevappa R, Malkowicz SB, Wittink M et al: Comparison of distribution- and anchor-based approaches to infer changes in health-related quality of life of prostate cancer survivors. Health Serv Res 2012; 47: 1902.
- Litwin MS, Greenfield S, Elkin EP et al: Assessment of prognosis with the total illness burden index for prostate cancer: aiding clinicians in treatment choice. Cancer 2007; 109: 1777.
- White H: A heteroskedasticity-consistent covariance matrix estimator and a direct test for heteroskedasticity. Econometrica 1980; 48: 817.
- Huber PJ: The behavior of maximum likelihood estimates under nonstandard conditions. In: Proceedings of the Fifth Berkeley Symposium on Mathematical Statistics and Probability. Edited by LM Le Cam and J Neyman. University of California 1967; vol 5.1, pp 221–233.
- Matta R, Wallis CJD, Goldenberg MG et al: Variation and trends in antidepressant prescribing for men undergoing treatment for nonmetastatic prostate cancer: a population-based cohort study. Eur Urol 2019; **75:** 3.
- Klaassen Z, Wallis CJD, Chandrasekar T et al: Cancer diagnosis and risk of suicide after accounting for prediagnosis psychiatric care: a matched-cohort study of patients with incident solid-organ malignancies. Cancer 2019; 125: 2886.
- Klaassen Z, Goldberg H, Chandrasekar T et al: Changing trends for suicidal death in patients with bladder cancer: a 40+ year populationlevel analysis. Clin Genitourin Cancer 2018; 16: 206.
- Klaassen Z, Wallis CJD, Goldberg H et al: The impact of psychiatric utilisation prior to cancer diagnosis on survival of solid organ malignancies. Br J Cancer 2019; **120**: 840.