Associations between body mass index and bladder cancer survival: Is the obesity paradox short-lived?

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Abstract

Introduction: We investigated the associations of pre-surgical body mass index (BMI) with bladder cancer outcomes in patients treated with radical cystectomy.

Methods: We retrospectively analyzed data from 488 bladder cancer patients treated with radical cystectomy between 1994 and 2007 and followed up until 2016. Cox regression with step function (time-segment analysis) was conducted for overall survival because the proportional hazard assumption was violated.

Results: Of 488 bladder cancer patients, 155 (31.8%) were normal weight, 186 (38.1%) were overweight, and 147 (30.1%) were obese. During the median followup of 59.5 months, 363 (74.4%) patients died, including 197 (40.4%) from bladder cancer. In adjusted Cox regression analyses, BMI was not significantly associated with bladder cancer-specific survival for overweight (hazard ratio [HR] 0.79, 95% confidence interval [CI] 0.57–1.10, p=0.16) or obese (HR 0.76, 95% CI 0.52–1.09, p=0.13) patients. In the Cox regression with step function for overall survival, the time interaction was significant overall (p=0.020) and specifically for overweight patients (p=0.006). In the time-segment model, the HR for overweight during the first 63 months was 0.66 (95% CI 0.49–0.90, p=0.008), whereas it was 1.41 (95% CI 0.89–2.23, p=0.14) after 63 months. Although not statistically significant, a similar pattern was observed for obese patients.

Conclusions: Our findings suggest that overweight and obese bladder cancer patients had better outcomes within the first five years after radical cystectomy; however, there were no differences in longer-term survival. These data suggest that the obesity paradox in bladder cancer patients treated with radical cystectomy may be short-lived.

Introduction

Bladder cancer is the fifth most common cancer in Canada¹ and the tenth most common cancer worldwide.² Radical

cystectomy is the main treatment for muscle-invasive bladder cancer and high-risk non-muscle-invasive bladder cancer.³ Although bladder cancer survival has increased slightly,⁴ bladder cancer remains the eighth most common cause of cancer mortality in Canada.¹ The 90-day mortality rate after radical cystectomy ranges from 3.2–7.5%^{5,6} and the five-year relative survival rate ranges from 16–55% depending on the tumor stage.⁷

Overweight and obesity are potential risk factors for developing bladder cancer because fat mass is related to inflammatory processes, alterations in sex hormone metabolism, abnormal levels of insulin and insulin-like growth factor, adipokine pathways, and microenvironment perturbations that contribute to tumor cell growth and proliferation.^{8,9} The association of obesity with bladder cancer survival, however, is much less clear. Some studies have reported that higher body mass index (BMI) is associated with a worse prognosis,^{10,11} whereas other studies have found no association.^{12,13} Conversely, some studies have reported that higher BMI is associated with better survival in bladder cancer patients,¹⁴⁻¹⁶ a phenomenon known as the obesity paradox.

Obesity may be linked to cancer treatment-related symptoms, treatment tolerance and response, recurrence, and long-term survival. Understanding the association between obesity and survival in bladder cancer patients after radical cystectomy may improve prognosis and guide bladder cancer management. The purpose of the present study was to investigate the association of pre-surgical BMI with bladder cancer outcomes in patients treated with radical cystectomy.

Methods

After institutional review board approval (Health Research Ethics Board of Alberta (HREBA) – Cancer Committee #20-0294), we analyzed an existing database of bladder cancer patients undergoing radical cystectomy from the Northern Alberta Urology Centre in Edmonton, Canada. The data set was collected retrospectively and included bladder cancer patients undergoing radical cystectomy between 1994 and 2007 and followed up for mortality outcomes until 2016. The primary exposure of interest was BMI, defined as weight

in kilograms divided by the square of height in meters (kg/m²). BMI was measured objectively during a clinical visit prior to radical cystectomy and was recorded in the patient electronic medical record. Eligible patients had to have a measure of BMI recorded in the data set.

The primary outcomes were length of hospitalization, 90-day mortality, bladder cancer-specific survival (BCSS), and overall survival (OS). Length of hospitalization was the number of days hospitalized from the time of radical cystectomy until discharge. Ninety-day mortality was assessed as death from any cause within 90 days of the radical cystectomy. BCSS was defined as the time from radical cystectomy to the date of death from bladder cancer or last followup if the patient had not died of bladder cancer. If the patient died of other causes, the event was censored at the time of death. OS was determined as the time from radical cystectomy to the date of death from any cause. Patients who were still alive were censored at the date of last followup.

Patients were categorized as normal weight (18.5–24.9 kg/m²), overweight (25–29.9 kg/m²), or obese (≥30 kg/m²). Descriptive statistics were used to summarize the data overall and by BMI category. Missing values for covariates (<3%) were replaced with the mean for continuous variables and the mode for categorical variables. Univariable associations between BMI and clinicopathological variables were assessed using Pearson Chi-squared for categorical variables and ANOVA for continuous variables. Life tables were used to obtain median survival time overall and by BMI category. Kaplan-Meier curves were used to describe survival probabilities of OS by BMI category. Log-rank tests were used to assess differences in unadjusted survival probabilities by BMI category.

Proportional hazard assumption was verified graphically using the log minus log plot and considered valid if the curves run in parallel. Cox proportional model was used to estimate the hazard ratio (HR) and 95% confidence intervals (CI) for 90-day mortality and BCSS. We observed evidence of non-proportionality for OS, indicating that the effect of a fixed baseline covariate (i.e., BMI) may change over time.¹⁷ Therefore, a Cox regression with a step function (time-segment analysis) was applied to explore the time varying coefficient.^{17,18} Two separate analyses were conducted based on a split of the median survival time.¹⁸ All the survival analyses were adjusted for established prognostic factors, including age, sex, cancer subtype (urothelial vs non-uro-thelial), lymphovascular invasion, tumor grade, pathological T stage, pathological N stage, and adjuvant chemotherapy. All tests were two-sided and statistical significance was set at p<0.05. All analyses were performed using SPSS (SPSS Inc., Chicago, IL, U.S.).

Results

A total of 508 patients were initially available in the data set. Twenty patients were excluded from the analyses due to missing data on weight or height (n=6), implausible BMI (n=4), and being underweight (n=10). The final sample included 488 bladder cancer patients who underwent radical cystectomy and had a valid measure of BMI before surgery.

Table 1 summarizes the baseline demographic and disease characteristics of the bladder cancer patients overall and by BMI category. Mean age at diagnosis was 65.8 years, BMI was 27.7 kg/m², and 78.5% were males. Most patients were diagnosed with urothelial carcinoma (89.8%), highgrade tumors (88.5%), and \geq T2 stage (71.1%). Overall, 155 patients (31.8%) were normal weight, 186 (38.1%) were overweight, and 147 (30.1%) were obese. Sex differed among the BMI categories, with the overweight and obese groups having more men (p<0.05). Regarding disease profile, overweight and obese patients generally had a more favorable prognostic profile compared to normal weight patients, including a significantly lower pathological T stage (p<0.05).

Table 2 summarizes the treatment and survival outcomes of bladder cancer patients overall and by BMI category.

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		Body mass index (kg/m²)				
Variables	Overall	Normal weight 18.5–24.9	Overweight 25–29.9	Obese ≥30	р	
	(n=488)	(n=155)	(n=186)	(n=147)		
Body mass index (kg/m²), M±SD	27.7±4.8	22.6±1.6	27.3±1.4	33.4±3.3	-	
Weight (kg), M±SD	81.8±17.1	65.8±9.5	81.4±9.6	99.3±13.5	-	
Sex (male), n (%)	383 (78.5)	107 (69.0)	154 (82.8)ª	122 (83.0)ª	0.002	
Age (years), M±SD	65.8±10.0	65.6±10.9	66.7±9.9	64.7±9.1	0.18	
Urothelial carcinoma, n (%)	438 (89.8)	138 (89.0)	164 (88.2)	136 (92.5)	0.41	
Lymphovascular invasion, n (%)	153 (31.4)	59 (38.1)	55 (29.6)	39 (26.5)ª	0.08	
High-grade, n (%)	432 (88.5)	143 (92.3)	159 (85.5)	130 (88.4)	0.15	
Pathologic T stage (≥T2), n (%)	347 (71.1)	124 (80.0)	130 (69.9)ª	93 (63.3)ª	0.005	
Pathologic N stage (N+), n (%)	104 (21.3)	41 (26.5)	37 (19.9)	26 (17.7)	0.15	

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Variables	Body mass index (kg/m²)					
	Overall	Normal weight 18.5–24.9	Overweight 25–29.9	Obese ≥30	р	
	(n=488)	(n=155)	(n=186)	(n=147)		
Neoadjuvant chemotherapy, n (%)	5 (1.0)	2 (1.3)	2 (1.1)	1 (0.7)		
Surgical procedure, n (%)						
Anterior exenteration (women)	105 (100)	48 (100)	32 (100)	25 (100)		
Radical cystoprostate (men)	376 (98.2)	103 (96.3)	152 (98.7)	121 (99.2)		
Positive surgical margin, n (%)	60 (12.3)	18 (11.6)	23 (12.4)	19 (12.9)	0.94	
Lymph node dissection, n (%)						
None/limited	37 (7.6)	18 (11.6)	12 (6.5)	7 (4.8)	0.63	
Standard	416 (85.2)	124 (80.0)	161 (86.6)	131 (89.1)		
Extended	35 (7.2)	13 (8.4)	13 (7.0)	9 (6.1)		
Diversion, n (%)						
lleal conduit	379 (77.7)	121 (78.1)	147 (79.0)	111 (75.5)	0.62	
Neobladder	99 (20.3)	29 (18.7)	36 (19.4)	34 (23.1)		
Other	10 (2.0)	5 (3.2)	3 (1.6)	2 (1.4)		
Estimated blood loss (ml), M±SD	1.3±0.8	1.3±0.9	1.3±0.8	1.3±0.8	0.74	
Hospital stay (days), M±SD	14.7±16.8	13.7±14.5	13.2±8.4	17.5±24.9 ^b	0.050	
Adjuvant chemotherapy, n (%)	79 (16.2)	30 (19.4)	26 (14.0)	23 (15.6)	0.39	
30-day mortality, n (%)	15 (3.1)	7 (4.5)	1 (0.5)ª	7 (4.8) ^b	0.039	
90-day mortality, n (%)	31 (6.4)	19 (12.3)	2 (1.1)ª	10 (6.8) ^b	<0.001	
Bladder cancer death, n (%)	197 (40.4)	74 (47.7)	74 (39.8)	49 (33.3)ª	0.038	
Overall death, n (%)	363 (74.4)	117 (75.5)	138 (74.2)	108 (73.5)	0.92	

Table 2. Treatment and survival outcomes of bladder cancer patients, overall and by body mass index

Only 1% of patients underwent neoadjuvant chemotherapy. Surgical procedures did not vary among BMI categories; all women in the study underwent anterior exenteration and 98.2% of men had a radical cystoprostatectomy; 85.2% of patients had a standard lymph node dissection; and 77.7% had ileal conduit diversion. In the 30-day mortality analyses, overweight patients had a lower proportion of deaths compared to normal and obese patients (p=0.039) but the overall number of deaths was small (n=15; 3.1%).

Regarding our primary outcomes of interest, obese patients had a prolonged hospital stay compared to overweight patients (17.5 ± 1.4 vs. 13.2 ± 1.3 days, p<0.05). After adjusting for prognostic factors, hospitalization remained significantly longer for obese vs. overweight patients (17.8 ± 1.4 vs. 13.0 ± 1.2 days, p=0.010) and became significantly longer for obese vs. normal weight patients (17.8 ± 1.4 vs. 14.0 ± 1.4 days, p=0.036). In the 90-day mortality analyses, overweight patients had a lower proportion of deaths compared to normal and obese patients (p<0.001). The results remained statistically significant in the multivariable Cox regression analyses, where overweight patients (HR 0.09, 95% CI 0.02–0.38, p=0.001) were less likely to die within 90-days after radical cystectomy (Table 3).

During the median followup of 59.5 months (interquartile range 13–144), 363 (74.4%) patients died, including 197 (40.4%) from bladder cancer. Univariable analyses for bladder cancer-specific mortality showed that being obese was associated with improved BCSS (p=0.011). After adjusting for established prognostic factors in the multivariable Cox regression analyses (Table 3), the difference in BCSS was not significant for either overweight (HR 0.79, 95% CI 0.57–1.10, p=0.16) or obese (HR 0.76, 95% CI 0.52–1.09, p=0.13) patients. The supplementary table provides the multivariate analysis, including all the covariates for the association of BMI with 90-day mortality, bladder cancer-specific mortality, and all-cause mortality.

Median OS for normal weight, overweight, and obese patients was 34.4, 88.5, and 70.8 months, respectively. The OS probability was not significantly different by BMI category (Figure 1). When assessing the proportional hazard assumption using the graphical model log-log plot, we observed evidence of non-proportional hazards among the BMI categories for OS. The violation of the proportional hazard assumption indicates that the hazard is transient over time (i.e., time-varying coefficient) and may lead to biased effect estimates when followup is long. One method to address the non-proportionality is dividing the Cox model into time intervals, therefore, we conducted two separate analyses split at the median survival of 63 months. In this analysis, BMI was included as a time-varying coefficient and the overall time interaction was significant (p=0.020). More specifically, the interaction was significant for over-

Table 3. Cox proportional HR and 95% CI for the				
association of body mass index with 90-day mortality,				
bladder cancer-specific mortality, and all-cause mortality				

	Overall (n=488)						
Body mass index (kg/m²)	No. of deaths/ no. of person- years	HR (95% CI)	р				
90-day mortality							
Normal weight (18.5–24.9)	19/985	Referent					
Overweight (25–29.9)	2/1417	0.09 (0.02–0.38)	0.001				
Obese (≥30)	10/1071	0.63 (0.29–1.39)	0.25				
Per 1-unit increase		0.93 (0.85–1.02)	0.11				
Bladder cancer-specific mortality							
Normal weight (18.5–24.9)	74/985	Referent					
Overweight (25–29.9)	74/1417	0.79 (0.57–1.10)	0.16				
Obese (≥30)	49/1071	0.76 (0.52–1.09)	0.13				
Per 1-unit increase		0.98 (0.95–1.01)	0.21				
All-cause mortality							
Normal weight (18.5–24.9)	117/985	Referent					
Overweight (25–29.9)	138/1417	0.84 (0.65–1.09)	0.18				
Obese (≥30)	108/1071	1.00 (0.77–1.31)	1.00				
Per 1-unit increase		1.01 (0.98–1.03)	0.70				
All-cause mortality with time-varying covariate			0.020*				
≤63 months							
Normal weight (18.5–24.9)	88/136	Referent					
Overweight (25–29.9)	82/166	0.66 (0.49–0.90)	0.008				
Obese (≥30)	72/141	0.88 (0.64–1.20)	0.42				
>63 months							
Normal weight (18.5–24.9)	29/841	Referent					
Overweight (25–29.9)	56/1244	1.41 (0.89–2.23)	0.14				
Obese (≥30)	36/920	1.38 (0.84–2.28)	0.21				
*p-value for time-interaction adjusted for age, sex, cancer subtype, lymphoyascular							

*p-value for time-interaction adjusted for age, sex, cancer subtype, lymphovascular invasion, tumor grade, pathological T stage, pathological N stage, and adjuvant chemotherapy. Cl: confidence interval; HR: hazard ratio.

weight (HR 2.13, 95% Cl 1.24–3.66, p=0.006) but not obese patients, although the interaction was in the same direction (HR 1.57, 95% Cl, 0.87–2.82, p=0.13). In the time-segment model, the estimated HR for overweight during the first 63 months was 0.66 (95% Cl 0.49–0.90, p=0.008), whereas it was 1.41 (95% Cl 0.89–2.23, p=0.14) after 63 months. The estimated HR for obese during the first 63 months was 0.88 (95% Cl 0.64–1.20, p=0.42), whereas it was 1.38 (95% Cl 0.84–2.28, p=0.21) after 63 months (Table 3). We conducted a sensitivity analysis for all survival analyses that excluded non-urothelial cancers and found no meaningful differences compared to the overall analyses (data not shown).

Discussion

In this retrospective analysis of a previously established database, overweight and obese patients had a lower 90-day mortality rate and a better OS within the first five years of radical cystectomy. After five years, however, overweight and obese patients had an increased risk of all-cause mortality, resulting in no difference in longer-term OS. These data suggest that the obesity paradox in bladder cancer patients treated with radical cystectomy may be driven by a lower 90-day mortality rate and fewer early bladder cancer-specific deaths and, therefore, short-lived. Our findings suggest a more nuanced interpretation of the obesity paradox for bladder cancer survival.

The role of BMI as a predictor of survival outcomes in bladder cancer patients varies considerably among studies.^{19,20} A recent systematic review and meta-analysis of BMI and survival in urothelial cancer patients treated with radical surgery reported that overweight was associated with better BCSS (HR 0.793, 95% CI 0.706-0.891, p<0.001) but worse OS (HR 1.125, 95% CI 1.018-1.244, p=0.021). Conversely, obesity was significantly associated with both worse BCSS (HR 1.138, 95% CI 1.028-1.261, p=0.013) and OS (HR 1.308, 95% CI 1.192-1.436, p<0.001).²⁰ This review noted substantial heterogeneity among the studies, although the source of the variance was not identified. One possible source of heterogeneity is that the review included Asian and Western-based studies with varying definitions of the BMI categories. Moreover, the studies had relatively short median followup times (from 21-64.1 months), making it difficult to compare the short-term and long-term survival experience of overweight and obese bladder cancer patients.

The obesity paradox appears to only hold during the first five years after radical cystectomy, when most deaths are due to surgical complications or bladder cancer, and then dissipates during further followup, when most deaths are due to other causes. After five years, obesity appears to exert its usual negative association with survival. This unique finding in our study was prompted by a violation of the proportional hazard assumption in our data set, which indicated BMI as a time-varying coefficient for OS.²¹ Assessment of the proportionality of hazards is a fundamental step for survival analyses using the Cox model. When the assumption is violated, a significant effect in the early (or late) followup period may be missed.¹⁸ The verification of this assumption, however, is not commonly reported in survival analysis of BMI in oncology, including investigations in bladder cancer.^{11-15,22}

It is possible that our observation may provide another explanation of the inconsistency in the obesity paradox across different cancers. Petrelli et al observed that the association between obesity and cancer survival may differ based on cancer type.²³ More specifically, these authors reported evidence of the obesity paradox in lung, renal, and

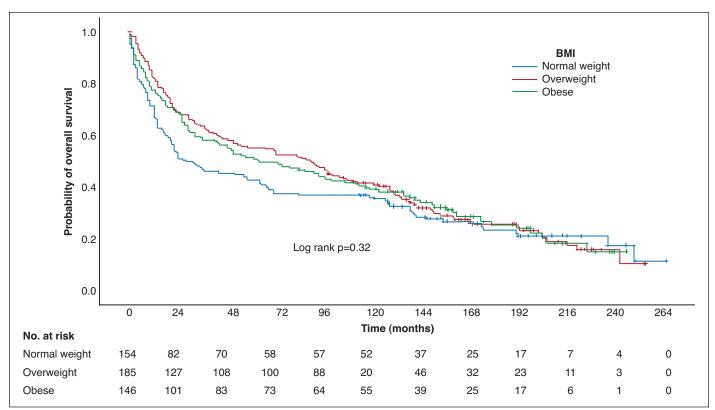


Figure 1. Kaplan-Meier curves for overall survival stratified by body mass index category. BMI: body mass index.

metastatic melanoma cancers — all cancers that generally have higher rates of surgical complications and more cancer deaths that occur early in the cancer trajectory. Conversely, they noted that the traditional negative association of BMI with survival seemed to hold in patients with breast, colon, and uterine cancers — all cancers that generally have lower rates of surgical complications and fewer early cancer-specific deaths, where patients are more likely to die of other causes during longer-term followup.

Nevertheless, the obesity paradox, whether short-lived or not, should be interpreted with caution. Explanations in the literature include methodological biases, such as unmeasured confounding.²⁴ For instance, smoking status was not assessed in our study and it may confound the association between BMI and survival in bladder cancer patients. Other studies have suggested clinical and biological differences among BMI categories, including body composition and tumor expression.²⁴

For example, in patients with clear-cell renal cell carcinoma, another malignancy of the genitourinary tract, differences in gene expression pathways within the tumor microenvironment might be associated with the obesity paradox.^{25,26} Sanchez et al reported that obese patients had significant tumor upregulation angiogenic pathways in comparison to normal weight patients; therefore, they were more likely to increase local drug delivery and benefit from therapy.²⁵ Hakimi et al identified that fatty acid synthase (*FASN*) gene expression, an indicator of tumor aggressiveness and prognosis, differed by BMI categories.²⁶ *FASN* was significantly upregulated in the normal BMI group and down-regulated in the obese group, suggesting that the tumors of obese patients may be less aggressive than normal weight patients with renal cell carcinoma.²⁶ In our sample, overweight and obese patients had a more favorable prognostic profile compared to normal weight patients. The literature has conflicting findings, however, regarding the associations of BMI with tumor characteristics in bladder cancer patients and further investigation is needed.^{10,22}

BMI as a measure of obesity is also questionable since it fails to distinguish body composition compartments among individuals in the same BMI range.²⁷ Psutka et al explored the association of BMI and fat mass index with OS in patients treated with radical cystectomy and urinary diversion.¹⁶ Increased BMI was associated with improved five-year OS, however, obesity based on fat-mass index had no association. In addition, after adjusting for the presence of low skeletal muscle mass, the association disappeared.¹⁶ Skeletal muscle mass is an important prognostic factor not assessed by BMI. Bladder cancer patients with lower skeletal muscle mass are at increased risk of early complications, cancer-specific mortality, and all-cause mortality.^{28,29} Bladder cancer patients have a high prevalence of low skeletal muscle mass at diagnosis and post-

treatment.³⁰ It is plausible that patients in the normal weight range might have lower muscle mass and, therefore, lower survival rates compared to overweight and obese patients.

Another possible explanation for better survival rates among overweight patients relates to the amount of subcutaneous fat. Cancer patients with higher subcutaneous adipose tissue experience longer survival than those with low subcutaneous fat.³¹ The moderate amount of fat allows patients to survive weight losses that occur with treatment and may provide nutritional benefits.^{24,32} These explanations are only speculations, however, and further research is warranted.

Strengths and limitations

The strengths of our study include a sufficient sample size, the balanced distribution across the BMI categories, the relatively long followup time, and the verification of the proportional hazard assumption for the survival analysis.

The limitations of our study include the exploratory nature of the analysis, the retrospective design of the study, the lack of data on potential confounders (e.g., smoking status, comorbidities), the use of BMI as the measure of obesity, and the measurement of BMI at only one time point. In addition, treatment has evolved since the completion of data collection (1994-2007), which may alter the association of obesity with survival. For example, our 90-day mortality rate of 12.3% is higher than that reported in the current literature, possibly due to changes in treatments and/or the likely higher rate of normal weight patients in our more dated sample. Finally, the generalizability of our results is limited only to patients treated with radical cystectomy. Future research should collect more sophisticated measures of obesity at multiple time points and check the proportional hazard assumption for a potential time-varying interaction with obesity. Moreover, other measures of health-related fitness besides body composition (e.g., cardiovascular fitness, muscular strength, physical functioning) may also be important predictors of clinical outcomes in bladder cancer patients, as they have been in breast cancer patients.³³

Conclusions

This exploratory analysis suggests that the obesity paradox in bladder cancer patients treated with radical cystectomy may only apply to postsurgical complications and bladder cancer deaths that typically occur early in the survival trajectory. With sufficient followup, the benefits of obesity on bladder cancer survival are reversed by an increased risk of other deaths, resulting in no long-term survival advantage for overweight/obese bladder cancer patients. Whether these associations were due to unmeasured confounding, limitations of BMI, or different tumor expression among the BMI groups is unknown. Well-designed, prospective studies that include a direct measure of body composition and tumor expression pathways are warranted to understand if and how body composition is associated with survival outcomes in bladder cancer.

Competing interests: The authors do not report any competing personal or financial interests related to this work.

This paper has been peer-reviewed.

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