

Prognostic significance of combined pretreatment body mass index (BMI) and BMI loss in patients with esophageal cancer

This article was published in the following Dove Press journal:
Cancer Management and Research

Wen-Shen Gu^{1,2,*}
Wei-Zhen Fang^{1,2,*}
Chun-Yue Liu^{1,2}
Kun-Yi Pan^{1,2}
Rui Ding^{1,2}
Xiao-Hui Li³
Chao-Hui Duan^{1,2}

¹Guangdong Provincial Key Laboratory of Malignant Tumor Epigenetics and Gene Regulation, Sun Yat-Sen Memorial Hospital, Sun Yat-Sen University, Guangzhou 510120, People's Republic of China; ²Department of Laboratory Medicine, Sun Yat-Sen Memorial Hospital, Guangzhou 510120, People's Republic of China; ³Key Laboratory of Oncology in South China, Collaborative Innovation Center for Cancer Medicine, Department of Clinical Laboratory Medicine, Sun Yat-Sen University Cancer Center, Guangzhou 510060, People's Republic of China

*These authors contributed equally to this work

Background: Body mass index (BMI) has been associated with a risk of esophageal cancer. However, the influence of BMI and BMI loss on people with esophageal cancer that were treated with different therapies has not been described in China.

Methods: In total, 615 consecutive patients that underwent esophagectomy and/or chemotherapy/radiotherapy were classified according to the Asian-specific BMI (kg/m²) cutoff values. The impact of BMI and BMI loss on long-term overall survival (OS) was estimated using the Kaplan–Meier method and Cox proportional hazard models.

Results: Multivariate analysis showed that overweight and obese patients had a more favorable survival than normal weight and underweight patients ($p=0.017$). Patients with a low BMI and high BMI loss before therapy had worse OS than others ($p=0.001$). Subgroup analysis showed that patients with a high BMI were more likely to suffer hypertension ($p<0.001$) and receive only surgery ($p<0.001$), and they were less likely to be smokers ($p=0.007$) and anemic ($p<0.001$). Conversely, patients with high BMI loss were more likely to be anemic ($p=0.001$), to have advanced pathological stage ($p=0.012$), and to receive chemotherapy and radiotherapy ($p=0.001$). Moreover, the mortality rate was higher when patients had a high BMI loss. There is no survival benefit of higher BMI in the non-esophageal squamous cell carcinoma (ESCC) group.

Conclusion: Pretreatment BMI was an independent prognostic factor for long-term survival in esophageal cancer patients treated with different treatments. The overall survival was increased in esophageal cancer patients with a high pretreatment BMI and no BMI loss. There is no survival benefit of higher BMI in the non-ESCC group.

Keywords: BMI, BMI loss, esophageal cancer, long-term survival

Introduction

Esophageal cancer is one of the most common digestive system malignancies in the world. It is especially prevalent in some Asian countries, and about half of the cases occur in China.^{1–3} With the advancements in surgical techniques and therapy, the outcomes of esophageal cancer have improved, but the long-term survival rate is still unsatisfactory.^{1,3,4} A common presenting symptom of esophageal cancer is weight loss.^{5,6} This is because most patients will experience dysphagia, which leads to reduced food intake.⁷ Moreover, increased energy consumption is induced by the tumor, which can also contribute to weight loss.⁸ Considerable weight loss in cancer patients is a risk factor for long-term survival. Many studies have indicated that excessive postoperative or post-diagnostic weight loss leads to worse long-term outcomes in many cancers,^{9–11} including esophageal cancer.^{12,13} Furthermore, several studies have demonstrated that

Correspondence: Chao-Hui Duan
Guangdong Provincial Key Laboratory of Malignant Tumor Epigenetics and Gene Regulation, Sun Yat-Sen Memorial Hospital, Sun Yat-Sen University, No. 107 Yanjiang West Road, Guangzhou, People's Republic of China
Tel +86 208 133 2072
Fax +86 208 133 2072
Email 1725012289@qq.com

a high body mass index (BMI) is a well-recognized risk factor for esophageal cancer.^{12,14–16} However, these studies were primarily concerned with the western population and patients who had undergone surgical tumor resection. Thus far, the impact of both BMI and a reduction in BMI prior to therapy on the long-term survival in Chinese esophageal cancer patients treated with adjuvant therapy, including chemotherapy and/or radiotherapy, has not been explored. Therefore, the aim of this study was to evaluate the impact of BMI and BMI loss prior to therapy on long-term survival in esophageal cancer patients treated with various types of therapy.

Patients and methods

Patients

We identified consecutive patients with esophageal cancer at the Sun Yat-sen University Cancer Center. Based on our previous study,¹⁷ patients with esophageal squamous cell carcinoma (ESCC) were identified between May 2007 and December 2008. Patients with non-ESCC were identified between March 2006 and September 2016 because of low incidence (Table S1). Follow-up time was calculated from the date of surgery to the event or date of the last contact. Follow-up continued until July 2018. The endpoint was overall survival (OS). OS was calculated from the time of surgery to the time of death from any causes or the date of the last follow-up, if the patients were still alive. Patients were excluded if their BMI was unknown or if they had a history of other cancers. Patient characteristics were collected from retrospective medical record reviews using a standardized data collection form. The most common surgical approaches included the left transthoracic procedure, the Ivor-Lewis approach, and the cervicothoracoabdominal procedure. Pathologic stage was determined according to the 7th edition AJCC staging system.¹⁸ The study was approved by the Ethics Committee of the Sun Yat-sen University Cancer Center and patient consent to review their medical records was not required for the research project did not involve personal privacy and commercial interests. Patient information would be kept strictly confidential and in compliance with the Helsinki declaration.

BMI value and BMI loss group

BMI (kg/m^2) was calculated based on a direct measurement of height and weight at diagnosis. Patients were classified according to the Asian-specific BMI cutoff value of the Chinese population.¹⁹ The three categories were as follows: underweight ($<18.5 \text{ kg}/\text{m}^2$), normal weight ($18.5\text{--}22.9 \text{ kg}/\text{m}^2$), and

overweight and obese ($\geq 23.0 \text{ kg}/\text{m}^2$). BMI loss was calculated based on height and weight loss compared to their weight at diagnosis. BMI loss was divided into the following three groups: group 1 (no BMI loss), group 2 (BMI loss ≤ 2), group 3 (BMI loss > 2) refer to some researches.^{13,20,21}

Statistical analysis

Statistical analysis was performed using the SPSS 19.0 for Windows software system (SPSS Inc., Chicago, IL, USA). Differences between the three groups were tested by the Kruskal–Wallis test. The associations between BMI categories and clinicopathologic parameters were analyzed by the χ^2 -square test or Fisher's exact test. Survival curves were calculated by the Kaplan–Meier method and analyzed by the log-rank test. Multivariable analysis was performed using Cox's proportional hazards regression model using the enter procedure (the entry and removal probabilities were 0.05 and 0.10, respectively). *p*-values < 0.05 were regarded as statistically significant.

Results

Baseline characteristics by BMI and BMI loss

There were 615 patients pathologically confirmed with esophageal cancer enrolled in this study. Of them, 474 (77.1%) were men, and 141 (22.9%) were women. All patients were between 23–82 years of age, with a median age of 59 years. The characteristics and parameters of the patients are listed in Tables 1 and 2. Patients with a high BMI were more likely to be diagnosed with hypertension ($p < 0.001$) and less likely to be smokers ($p = 0.007$) or anemic ($p < 0.001$). Furthermore, overweight and obese patients were less likely to require adjuvant therapy in addition to their operation ($p < 0.001$). On the contrary, patients with high BMI loss were more likely to receive chemotherapy and radiotherapy ($p = 0.001$). Moreover, BMI and BMI loss demonstrated a positive correlation with the cancer stage. Patients with high BMI loss and a low BMI had higher values for T ($p = 0.007$) classification and lymph node metastasis ($p < 0.001$) (Table S2). There were no significant differences in histopathology, gender, age, alcohol history, diabetes mellitus, family history, and cancer location.

Univariate and multivariate analysis

The median OS of the entire cohort was 31.0 months, and it ranged from 1.0–144.0 months. At the end of the observation period, 61.1% (376/615) of patients had died due to disease-related or unrelated factors (Table 3). To evaluate the prognostic

Table I Relationship between the BMI and the clinical characteristics in 615 patients with esophageal cancer

Characteristics	Overall (n=615)	BMI			p-value
		<18.5(%) (n=88)	18.5–22.9(%) (n=307)	≥23.0(%) (n=220)	
Histopathology					
ESCC	446(72.5)	65(73.9)	225(73.3)	156(70.9)	0.828
Non-ESCC	169(27.5)	23(26.1)	82(26.7)	64(29.1)	
Small cell carcinoma	63(37.3)	5(21.7)	31(37.8)	27(42.2)	0.638
Adenocarcinoma	46(27.2)	2(8.7)	21(25.6)	23(35.9)	
Gender					
Male	474(77.1)	62(70.5)	240(78.2)	172(78.2)	0.280
Female	141(22.9)	26(29.5)	67(21.8)	48(21.8)	
Age (years)					
≤59 years	314(51.1)	54(61.4)	147(47.9)	113(51.4)	0.083
>59 years	301(48.9)	34(38.6)	160(52.1)	107(48.6)	
Alcohol					
Never	375(62.0)	50(56.8)	177(57.7)	148(67.3)	0.057
Ever	240(38.0)	38(43.2)	130(42.3)	72(32.7)	
Smoking					
Never	217(35.3)	24(27.3)	98(31.9)	95(43.2)	0.007
Ever	398(64.7)	64(72.7)	209(68.1)	125(56.8)	
Diabetes mellitus					
No	587(95.4)	85(96.6)	293(95.4)	209(95.0)	0.833
Yes	28(4.6)	3(3.4)	14(4.6)	11(5.0)	
Hypertension					
No	453(73.7)	81(92.0)	235(76.5)	137(62.3)	0.000
Yes	162(26.3)	7(8.0)	72(23.5)	83(37.7)	
Anemia					
No	548(89.1)	68(77.3)	273(88.9)	207(94.1)	0.000
Yes	67(11.9)	20(22.7)	34(11.1)	13(5.9)	
Family history					
No	499(81.1)	72(81.8)	259(84.4)	168(76.4)	0.067
Yes	116(19.9)	16(18.2)	48(15.6)	52(23.6)	
Location					
Up	56(9.1)	7(8.0)	31(10.1)	18(8.2)	0.938
Middle	380(61.8)	55(62.5)	189(61.6)	136(61.8)	
Low	179(29.1)	26(29.5)	87(28.3)	66(30.0)	
Stage					
I and II	320(52.0)	39(44.3)	161(52.4)	120(54.5)	0.262
III and IV	295(48.0)	49(55.7)	146(47.6)	100(45.5)	
Treatment					
CHO+RAD	36(5.8)	3(3.4)	20(6.5)	13(5.9)	0.003
Surgery only	324(52.7)	38(43.2)	149(48.5)	137(62.3)	
SUR+CHO+RAD	255(41.5)	47(53.4)	138(45.0)	70(31.8)	

Note: Bold values indicate significant differences ($p < 0.05$).

Abbreviations: BMI, body mass index; CHO+RAD, chemotherapy and radiotherapy; SUR+CHO+RAD, surgery and chemotherapy and radiotherapy; ESCC, esophageal squamous cell carcinoma.

Table 2 Relationship between the BMI loss and the clinical characteristics in 615 patients with esophageal cancer

Characteristics	Overall (n=615)	BMI loss			p-value
		0%(n=365)	≤2%(n=207)	> 2%(n=43)	
Histopathology					
ESCC	446(72.5)	266(72.9)	151(72.9)	29(67.4)	0.741
Non-ESCC	169(27.5)	99(27.1)	56(27.1)	14(32.6)	
Small cell carcinoma	63(37.3)	36(36.4)	24(42.9)	3(21.4)	0.689
Adenocarcinoma	46(27.2)	30(30.3)	14(25.0)	2(14.3)	
Gender					
Male	474(77.1)	279(76.4)	162(78.3)	33(76.7)	0.882
Female	141(22.9)	86(23.6)	45(21.7)	10(23.3)	
Age					
≤59 years	314(51.1)	170(47.8)	112(54.1)	23(50.3)	0.317
>59 years	301(48.9)	186(52.2)	95(45.9)	20(49.7)	
Alcohol					
Never	375(62.0)	230(63.0)	124(59.9)	21(61.0)	0.183
Ever	240(38.0)	135(37.0)	83(40.1)	22(39.0)	
Smoking					
Never	217(35.3)	136(37.3)	67(32.4)	14(35.3)	0.464
Ever	398(64.7)	229(62.7)	140(67.6)	29(64.7)	
Diabetes mellitus					
No	587(95.4)	348(95.3)	197(95.2)	42(95.4)	0.765
Yes	28(4.6)	17(4.7)	10(4.8)	1(4.6)	
Hypertension					
No	453(73.7)	266(72.9)	151(72.9)	36(83.7)	0.299
Yes	162(26.3)	99(27.1)	56(27.1)	7(16.3)	
Anemia					
No	548(89.1)	333(91.2)	184(88.9)	31(72.1)	0.001
Yes	67(11.9)	32(8.8)	23(11.1)	12(27.9)	
Family history					
No	499(81.1)	291(79.7)	169(81.6)	39(90.7)	0.215
Yes	116(19.9)	74(20.3)	38(18.4)	4(9.3)	
Location					
Up	56(9.1)	32(8.8)	21(10.1)	3(7.0)	0.477
Middle	380(61.8)	235(64.4)	121(58.5)	24(55.8)	
Low	179(29.1)	98(26.8)	65(31.4)	16(37.2)	
Stage					
I and II	320(52.0)	208(57.0)	92(44.4)	20(52.0)	0.012
III and IV	295(48.0)	157(43.0)	115(55.6)	23(48.0)	
Treatment					
CHO+RAD	36(5.8)	16(1.7)	15(7.2)	5(11.6)	0.001
Surgery only	324(52.7)	200(56.3)	99(47.8)	25(58.1)	
SUR+CHO+RAD	255(41.5)	149(42.0)	93(44.9)	13(30.2)	

Note: Bold values indicate significant differences ($p < 0.05$).

Abbreviations: BMI, body mass index; CHO+RAD, chemotherapy and radiotherapy; SUR+CHO+RAD, surgery and chemotherapy and radiotherapy; ESCC, esophageal squamous cell carcinoma.

Table 3 Overall survival analysis in 615 patients with esophageal cancer

Variables	Overall survival (OS)			
	Mean	Median	5 year OS(%)	10 year OS(%)
Histopathology				
ESCC	68.1	47.0	46.0	39.2
Non-ESCC	52.9	21.0	43.8	39.1
Small cell carcinoma	32.7	17.0	36.5	34.9
Adenocarcinoma	59.3	33.0	47.8	43.5
Gender				
Male	69.3	39.0	45.8	40.5
Female	63.0	28.0	39.7	34.8
Age (years)				
≤59	70.0	37.0	44.9	41.7
>59	60.8	37.0	43.9	36.5
TNM stage[#]				
I-II	89.0	60.0	59.4	54.1
III-IV	44.4	19.0	28.1	23.1
Alcohol				
Never	71.2	43.0	46.4	41.9
Ever	58.5	31.0	41.3	35.0
Smoking				
Never	1.7	44.0	46.5	42.4
Ever	65.7	34.0	43.3	37.4
Diabetes mellitus				
No	67.5	37.0	44.0	39.0
Yes	67.1	69.0	53.6	42.9
Hypertension				
No	66.6	37.0	43.9	38.9
Yes	66.5	37.0	45.7	40.1
Anemia				
No	69.6	38.0	45.3	40.7
Yes	50.0	27.0	37.3	26.9
BMI loss				
No	74.4	50.0	50.1	43.6
Yes	54.3	28.0	36.0	32.8
Treatment				
CHO+RAD	46.3	19.0	38.9	38.9
Surgery only	69.4	39.0	44.8	41.0
SUR+CHO+RAD	6.6	39.0	44.7	36.9
BMI status				
<18.5 kg/m ²	52.5	26.0	33.0	29.5
18.5–22.9 kg/m ²	65.0	33.0	43.0	36.2
≥23.0 kg/m ²	77.3	51.0	50.5	47.3

Notes: [#]TNM denotes tumor-node-metastasis.

Abbreviations: CHO+RAD, chemotherapy and radiotherapy; SUR+CHO+RAD, Surgery and chemotherapy and radiotherapy; ESCC, esophageal squamous cell carcinoma.

value of BMI and BMI loss in esophageal cancer patients, the demographic data, clinicopathological features, life style, BMI, and BMI loss were evaluated using univariate and multivariate Cox regression models (Table 4). The univariate analysis revealed that histopathology, TNM stage, anemia, BMI, and BMI loss were significantly associated with the survival of esophageal cancer patients. However, gender, age, alcohol history, smoking history, diabetes mellitus, high blood pressure, and treatment were not significantly associated. Thus, the significant variables were selected for multivariate analysis using the enter method. The multivariate analysis showed that histopathology (HR =1.545; 95%CI: 1.227–1.946; $p<0.001$), TNM stage (HR =2.533; 95%CI: 2.050–3.131; $p<0.001$), and BMI (HR =0.829; 95%CI: 0.711–0.966; $p=0.017$) were independent prognostic indicators of esophageal cancer survival. What's more, when we model BMI and BMI loss as continuous predictors, it shows that for every 1 unit increased in BMI and BMI loss, the HR decreased to 0.952(0.921–0.984) ($p=0.004$) and increased to 1.039(1.004–1.076) ($p=0.030$). Thus, these findings suggest that BMI prior to therapy may serve as a novel independent prognostic factor for esophageal cancer.

Table 4 Multivariate Cox hazards analysis for OS in 615 patients with esophageal cancer

Variables	Multivariate survival analysis	
	HR(95% CI)	p-value*
Histopathology		
ESCC	1.523(1.209–1.918)	0.000
Non-ESCC		
TNM stage#		
I-II	2.533(2.050–3.131)	0.000
III-IV		
Anemia		
No	1.262(0.929–1.713)	0.137
Yes		
BMI loss		
No	1.194(0.970–1.469)	0.095
Yes		
As continuous variables	1.039(1.004–1.076)	0.030
BMI status		
<18.5 kg/m ²	0.829(0.711–0.966)	0.017
18.5–22.9 kg/m ²		
≥23.0 kg/m ²		
As continuous variables	0.952(0.921–0.984)	0.004

Notes: Bold values indicate significant differences ($p<0.05$). *Cox hazard regression model. #TNM denotes tumor-node-metastasis.

Abbreviations: BMI, body mass index; ESCC, esophageal squamous cell carcinoma; OS, overall survival.

Prognostic significance of BMI

To further explore the prognostic significance of BMI in esophageal cancer patients, Kaplan-Meier survival curves were generated, and the groups were compared using the log-rank test. The patients were categorized into the following three groups: the underweight group (<18.5 kg/m²), the normal weight group (18.5–22.9 kg/m²), and the overweight and obese group (≥23.0 kg/m²). In the entire cohort, overweight and obese patients had significantly better 10 year OS than underweight and normal weight patients (Figure 1). The cumulative 10 year survival rate in the overweight and obese group was 47.3%, whereas it was only 29.5% and 36.2% in the underweight and normal weight groups, respectively. Additionally, patients were categorized into either an ESCC group or a non-ESCC group to evaluate the 10 year OS differences according to the BMI index. This is because the 10 year survival rates in these two groups were different ($p<0.001$, Figure 2A). The 10 year OS in patients with a high BMI was significantly longer than patients with a low BMI in both the whole cohort ($p=0.004$, Figure 1) and the ESCC cohort ($p=0.003$, Figure 2B). However, there was no significant difference in 10 year OS among the three BMI groups in the non-ESCC group ($P=0.667$, Figure 2C).

Combining the prognostic significance of BMI and BMI loss

To further explore the prognostic significance of combining BMI and BMI loss, we generated Kaplan-Meier survival curves and performed the log-rank test for the whole cohort. The prognostic significance is shown in Figure S1. Patients were divided into six groups: group 1 (BMI <18.5 kg/m² and BMI loss =0), group 2 (18.5≤ BMI <23.0 kg/m² and BMI loss =0), group 3 (BMI ≥23.0 kg/m² and BMI loss =0), group 4 (BMI <18.5 kg/m² and BMI loss >0), group 5 (18.5≤ BMI <23.0 kg/m² and BMI loss >0), group 6 (BMI ≥23.0 kg/m² and BMI loss >0). The 10 year OS rates of the six groups were significantly different ($p=0.001$). Patients with a high BMI and no BMI loss (group 3) had a significantly higher survival probability than the other five groups. Conversely, the survival rate of underweight patients with BMI loss (group 4) was the lowest among the six groups. The order of the survival rates of the six groups from high to low was as follows: group 3 (51.0%) > group 6 (40.0%) > group 2 (39.2%) > group 1 (36.4%) > group 5 (32.1%) > group 4 (22.7%) (Figure S1B). Although BMI loss was not an independent prognostic factor for esophageal cancer (HR =1.194, 95% CI: 0.970–1.469, $P=0.095$), there was

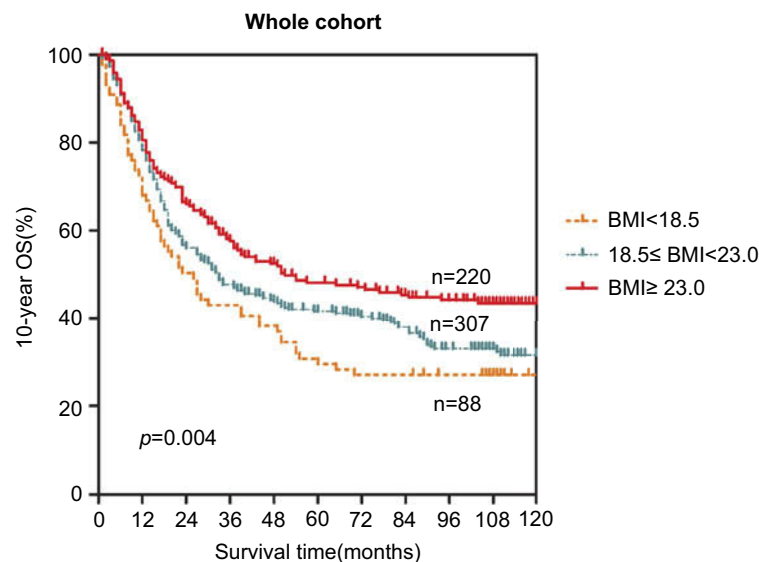


Figure 1 Prognostic significance of BMI in 615 esophageal cancer patients. The 10 year overall survival rate was calculated using the Kaplan-Meier method and analyzed with the log-rank test. The patients were categorized into a low BMI group, normal BMI group and a high BMI group, according to the Asian-specific BMI cutoff. A high BMI level was a favorable prognostic factor in whole esophageal cancer patients cohort.

Abbreviation: BMI, body mass index.

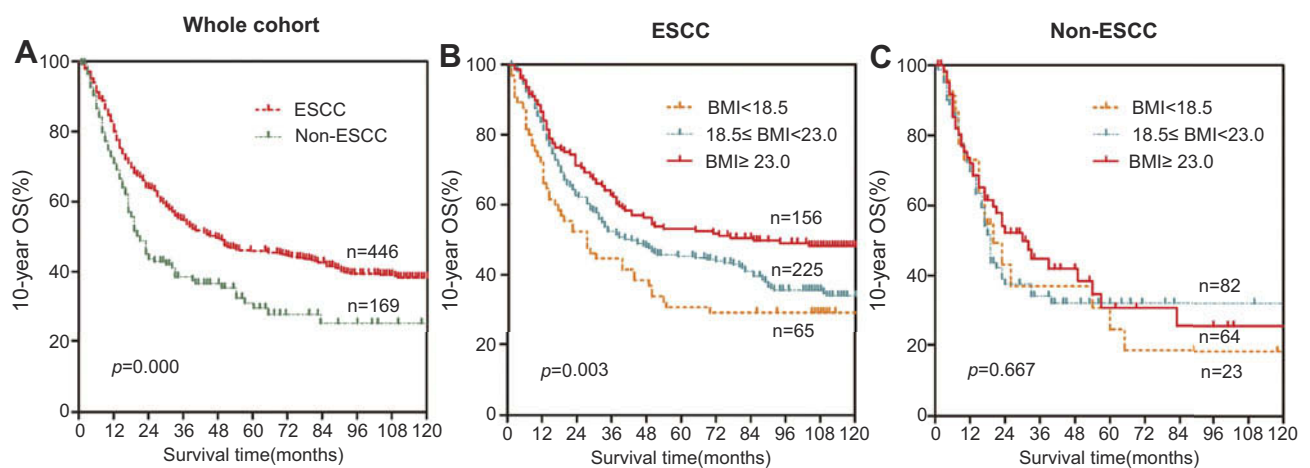


Figure 2 Prognostic significance of BMI in ESCC and non-ESCC groups. The overall survival rate was calculated using the Kaplan-Meier method and analyzed with the log-rank test. The patients were categorized into either the ESCC group or the non-ESCC group. **(A)** The 10 year overall survival rate was different in these two groups. **(B)** A high BMI was a favorable prognostic factor in the ESCC group. **(C)** There were no statistically significant differences in the non-ESCC group.

Abbreviations: BMI, body mass index; ESCC, esophageal squamous cell carcinoma.

a significant difference ($p < 0.001$) in the 10 year OS among the three BMI loss groups in the whole cohort (Figure S1A). To further explore the changes related to BMI loss, we calculated the mortality rate (Figure S1C). The results indicate that the mortality rate increased as BMI loss increased.

Discussion

Our study demonstrates that pretreatment BMI could be an independent prognostic factor for long-term OS in esophageal cancer patients. We found that the 10 year OS of patients with

a high BMI were significantly longer than patients with a low BMI, which was true for both the entire cohort and the ESCC group. This result was consistent with previous studies in both benign tumors²²⁻²⁵ and malignant tumors.^{12,26-33} Although there have been studies that investigated the relationship between esophageal cancer and BMI,^{12,16,34-39} these studies primarily focused on operative squamous cell carcinoma, esophageal adenocarcinoma, or patients only treated with adjuvant therapy. Moreover, a population study of esophageal cancer patients treated with different therapies, including surgery and

adjuvant therapy, has not been explored. In other words, there are no reports of clinical research on the relationship between pretreatment BMI and esophageal cancer patients treated with adjuvant therapy in the Chinese population.

Although BMI levels were different in different studies, the results of this study were consistent with other studies on the prognosis of esophageal cancer.^{12,37,40} The reason why overweight and obese patients might have improved survival is not clear. Previous studies have indicated that the prognostic advantage for overweight and obese patients might be attributed to the fact that overweight and obese patients were more likely to never have smoked, never have consumed alcohol, and not have lost weight before surgery.^{16,41} In our study, patients with a high BMI were less likely to be smokers ($p=0.007$). Moreover, it was previously reported that pre-operative malnutrition was associated with poor survival in cancer patients.⁴² Overweight and obese patients might have a better nutritional reserve to resist cancer.⁴³ In this study, patients with a high BMI were less likely to be anemic ($p<0.001$), and this may imply better physical conditions.

As a calculation that indicates body fat, BMI is different from other unchangeable factors, such as age, race, and gender. It is potentially more modifiable. The factor that affects body weight may be a factor that influences the relationship between BMI and the prognosis of esophageal cancer patients. Weight loss is a common presenting symptom of digestive system neoplasms, especially in oral cancer, esophageal cancer, and gastric carcinoma. This is because most patients will experience dysphagia and odynophagia, which lead to reduced food intake. Also contributing to weight loss and malnutrition is the increased energy consumption and altered nutrient metabolism that is associated with systemic inflammation, which can be induced by the tumor.^{5,6,44} This is especially true for esophageal cancer patients after surgery.^{45,46} It is known that significant weight loss in cancer patients is a risk factor for long-term survival. Several studies have reported that pre-operative weight loss $>10\%$ of body weight can increase the risk of mortality after surgery by more than 10% in esophageal cancer patients.^{13,20,21} In this study, we chose BMI loss instead of weight loss to evaluate the influence of weight change on survival by taking patient height into consideration. The Hazard Ratios (HRs) of BMI loss suggest that pretreatment BMI loss increased

mortality, although this was not statistically significant (HR =1.194; 95%CI: 0.970–1.469; $p=0.095$). However, the mortality rate of patients in the present cohort increased as BMI loss increased. BMI loss was closely related to the cancer stage; a higher BMI loss had higher T ($p=0.007$) classifications.

Nevertheless, we should acknowledge that the association between BMI and survival might be influenced by certain significant confounding factors. Studies have shown that a higher household income was associated with a 33–38% decrease in risk of death in esophageal cancer.⁴⁰ Thus, improving the general socioeconomic status is a promising approach for reducing the tremendous burden of upper gastrointestinal cancers.³⁸ In other words, patients with good socioeconomic status, income status, and education were more likely to receive better treatment because of better health consciousness and financial support.

However, this is a single-center retrospective analysis, and we need to validate the key findings and conclusions with other commonly used validation methods, including an independent cohort, in the future.

Conclusion

Our study provides more concrete evidence that a high BMI is associated with favorable long-term survival in esophageal cancer patients treated with different therapies in China. Patients with a low BMI and high BMI loss before therapy had a worse outcome than other patients with esophageal cancer. These results emphasize that the nutritional status in malnourished patients with esophageal cancer should be improved before therapy for a better prognosis.

Acknowledgments

We thank the staff at the Director of Clinical Laboratories, Sun Yat-Sen Memorial Hospital, and Sun Yat-sen University Cancer Center for providing support on research conditions in this study. This work was supported by the National Natural Science Foundation of China (NSFC, No. 31170147), the Guangdong Natural Science Foundation (Grant Numbers: S2013010014007 and 2014A030313070) and the Guangdong Province Science and Technology Project Plan and Social Development Foundation (2010A030400006).

Disclosure

The authors report no conflicts of interest in this work.

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Supplementary materials

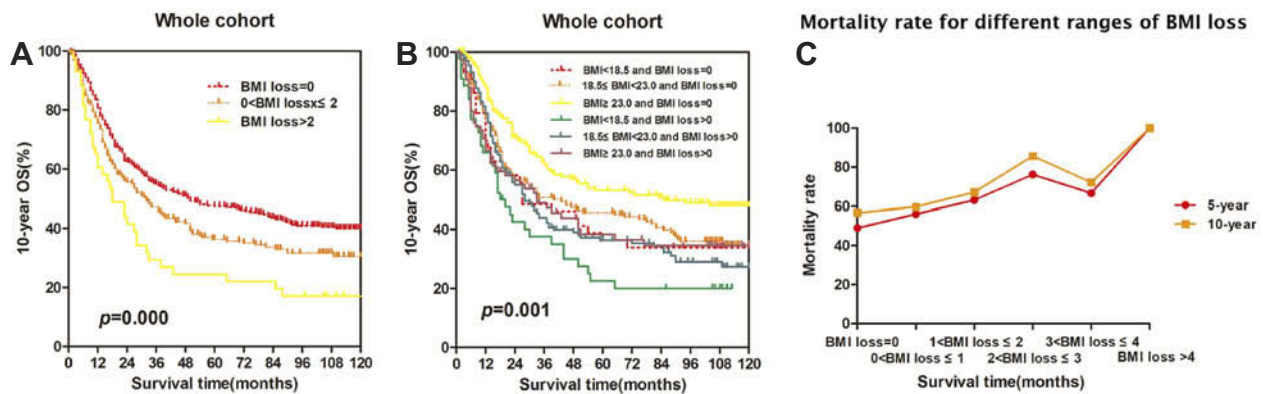


Figure S1 Combining the prognostic significance of BMI and BMI loss. The overall survival rate was calculated using the Kaplan-Meier method and analyzed with the log-rank test. The patients were categorized into three groups: group 1 (no BMI loss), group 2 (BMI loss ≤ 2), and group 3 (BMI loss > 2). **(A)** Patients with no BMI loss had a higher 10 year overall survival rate. **(B)** Patients with a high BMI and no BMI loss had a significantly higher survival probability than the other groups. **(C)** The mortality rate increased as BMI loss increased.

Table S1 Main cancer type in 169 patients with non-ESCC

Histopathology	Number (%)
Small cell carcinoma	63(37.3)
Adenocarcinoma + adenosquamous carcinoma	46(27.2)
Sarcomatoid carcinoma + carcinosarcoma	31(18.3)
Neuroendocrine carcinoma	12(7.1)
Malignant melanoma	8(4.7)
Mesenchymoma	5(3.0)
Signet-ring cell carcinoma	1(0.6)
Rhabdomyosarcoma	1(0.6)
Neurinoma	1(0.6)
Mucoepidermoid carcinoma	1(0.6)

Abbreviation: ESCC, esophageal squamous cell carcinoma.

Table S2 Relationship between the BMI or BMI loss and the TNM stage in 615 patients

Characteristics	Overall (n=615)	BMI			p-value	BMI loss				p-value	
		<18.5 (%) (n=88)	18.5–22.9 (%) (n=307)	≥23.0 (%) (n=220)		0% (n=365)	1% (n=82)	2% (n=125)	≥3% (n=43)		
T classification											
1–2	196	20(22.7)	104(33.9)	72(32.7)	0.133	132(36.2)	28(34.1)	25(20.0)	11(25.6)		
3–4	419	68(77.3)	203(66.1)	148(67.3)		233(63.8)	54(65.9)	100(80.0)	32(74.4)		
N classification											
No	283	33(37.5)	147(71.0)	103(46.8)	0.000	183(50.1)	34(41.5)	50(40.0)	16(37.2)		0.096
Yes	332	55(62.5)	160(29.0)	117(53.2)		182(49.9)	48(58.5)	75(60.0)	27(62.8)		
M classification											
No	585	84(95.5)	289(94.1)	212(96.4)	0.498	346(94.8)	77(93.9)	121(96.8)	41(95.3)		0.775
Yes	30	4(4.5)	18(5.9)	8(3.6)		19(5.2)	5(6.1)	4(3.2)	2(4.7)		

Notes: Bold value indicates significant differences (p<0.05).

Abbreviation: BMI, body mass index.

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