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# Loss of skeletal muscle mass during neoadjuvant treatments correlates with worse prognosis in esophageal cancer: a retrospective cohort study

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## Abstract

**Background:** Nutritional deficits, cachexia, and sarcopenia are extremely common in esophageal cancer. The aim of this article was to assess the effect of loss of skeletal muscle mass during neoadjuvant treatment on the prognosis of esophageal cancer patients.

**Methods:** Esophageal cancer patients ( $N = 115$ ) undergoing neoadjuvant therapy and surgery between 2010 and 2014 were identified from our surgery database and retrospectively analyzed. Computed tomography imaging of the total cross-sectional muscle tissue measured at the third lumbar level defined the skeletal muscle index, which defined sarcopenia ( $\text{SMI} < 52.4 \text{ cm}^2/\text{m}^2$  for men and  $< 38.5 \text{ cm}^2/\text{m}^2$  for women). Images were collected before and after neoadjuvant treatments.

**Results:** Sarcopenia in preoperative imaging was prevalent in 92 patients (80%). Median overall survival was 900 days (interquartile range 334–1447) with no difference between sarcopenic (median = 900) and non-sarcopenic (median = 914) groups ( $p = 0.872$ ). Complication rates did not differ (26.1% vs 32.6%,  $p = 0.725$ ). A 2.98% decrease in skeletal muscle index during neoadjuvant treatment correlated with poor 2-year survival (log-rank  $p = 0.04$ ).

**Conclusion:** Loss of skeletal muscle tissue during neoadjuvant treatment correlates with worse overall survival.

**Keywords:** Esophageal neoplasms, Thoracic surgery, Sarcopenia, Body composition, Malnutrition

## Background

Esophageal cancer (EC) is intimately related to weight changes and poor nutritional status, since the most common symptoms of EC are dysphagia and weight loss [1]. In locally advanced EC, baseline nutritional status has been linked to survival after definitive chemoradiotherapy [2]. Preoperative weight loss has also been linked to worse outcomes [3]. A high body mass index (BMI) has not been found to have a significant effect on survival in EC [4, 5].

In the recent years, there has been an increasing amount of studies on frailty and especially sarcopenia as

prognostic factors in cancers. Sarcopenia is defined as the progressive loss of muscle related to aging or disease [6]. Sarcopenia has been associated with worse outcomes in many types of cancers such as hepatocellular carcinoma, colorectal cancer, and small cell lung cancer [7–9]. For esophageal and gastroesophageal junction cancers, there are conflicting reports. Worse long-term outcomes have been reported in resected esophageal or gastroesophageal junctional cancers [10–12]. Sarcopenia has been linked to increased pulmonary and other complication rates [13, 14]. Decreased skeletal muscle area during neoadjuvant therapy has also been associated with poorer outcomes and risk of positive clinical resection margin [15, 16]. There are also reports of sarcopenia not being an independent prognostic risk factor for mortality, morbidity, or poor outcomes in EC after neoadjuvant chemoradiotherapy or chemotherapy [14, 17–19]. One

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study found a significant correlation with lean psoas mass and survival in patients not undergoing neoadjuvant treatment, but no such effect on patients receiving neoadjuvant treatment [20].

The aim of this trial was to assess the effect of sarcopenia and loss of skeletal muscle index during neoadjuvant treatments in patients undergoing esophagectomy for EC. The primary end-point is overall survival, and secondary end points are recurrence-free survival and complication rates.

## Methods

### Patients

Patients who underwent surgical resection and neoadjuvant therapy for EC between 2010 and 2014 were identified in retrospect from our surgery database using type of surgery and diagnosis of esophageal or junctional cancer as identifiers ( $N = 118$ ). Patients who had no eligible imaging for analysis of the skeletal muscle index were excluded ( $N = 3$ ), leaving 115 patients. Collected data included patient characteristics, weight, and weight-loss data before and during the treatment and follow up, primary tumor characteristics and staging, specifics of neoadjuvant, endoscopic, surgical and adjuvant treatments, post-operative and long-term complications, and overall survival. Weight loss here is defined as unintentional deviation from healthy weight (weight 6 months before diagnosis). Complications were collected as recommended by an international consensus statement [21]. 30- and 90-day overall survival rates and 2-year overall survival and recurrence-free survival rates were also collected.

CT (computed tomography) scans were collected from time of initial staging, post-neoadjuvant, 6-, and 18-month follow-up visits. CT scans were excluded if there was impaired visibility at third lumbar vertebra or over 1 month of time interval between the CT scan and associated event (start of neoadjuvant therapy, operation, 6-month follow-up visit, or 18-month follow-up visit).

### Preoperative staging

All patients underwent gastroscopies with biopsies that confirmed the diagnosis of EC. All patients underwent CT scans of the thorax, abdomen, and pelvis and a routine total body PET-CT scan. Endoscopic ultrasound was done unless prevented by tumor obstruction or stent insertion and assessed the invasion depth of the tumor and identified regional enlarged lymph nodes. PET/CT scans were routinely repeated after neoadjuvant treatments for preoperative planning and to assess the radiologic response of the tumor.

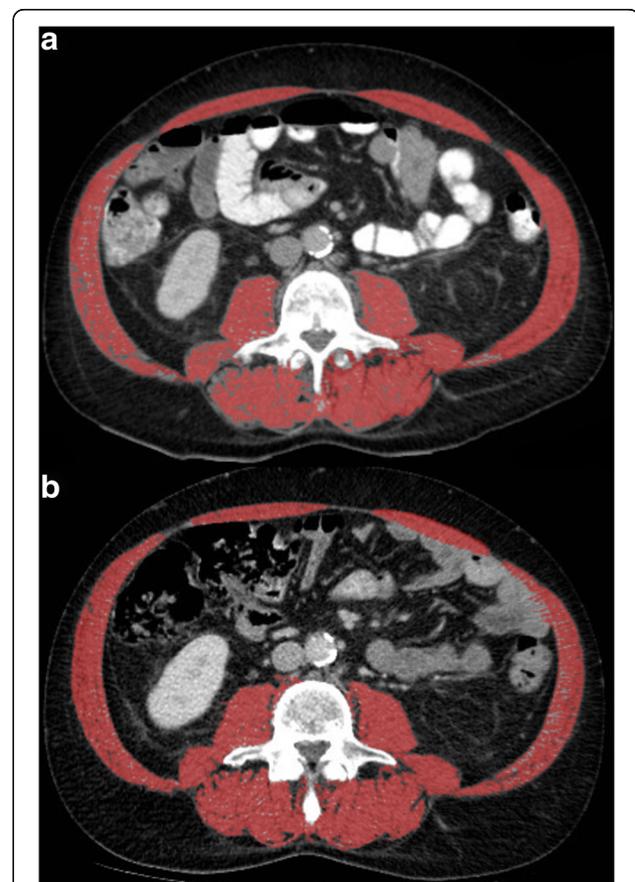
### Neoadjuvant treatment protocols

Treatment strategies were discussed and decided together with oncologists. Patients with nodal disease spread (cN+)

or transmural tumor invasion ( $cT \geq 3$ ) underwent neoadjuvant treatment, unless contraindicated. Neoadjuvant therapy was epirubicin–oxaliplatin–capecitabine neoadjuvant chemotherapy as per MAGIC (Medical Research Council Adjuvant Gastric Infusional Chemotherapy) protocol for esophageal adenocarcinoma [22]. Squamous cell carcinoma was treated with neoadjuvant chemoradiotherapy consisting of 2 cycles of platin- and 5-fluorouracil-based therapy over 5–6 weeks. Chemosensitization was followed by a 45 Gy total dose of radiation to the tumor and regional nodes, in 1.8 Gy daily fraction.

### Preoperative endoscopic procedures

Endoscopic mucosal resection (EMR) with or without radiofrequency ablation (RFA) was discussed as an option for patients with mucosal tumors or high-grade dysplasia. The initial decision whether to proceed with endoscopic treatments or to do an esophagectomy was a shared decision between the surgeon and the patient. If endoscopic mucosal resection showed submucosal spread or there was cancer recurrence, treatment proceeded to esophagectomy. One patient included in this



**Fig. 1** An example of SMI delineating. **a** Shows a male patient without sarcopenia (SMI = 57.6 cm<sup>2</sup>/m<sup>2</sup>), whereas **b** shows the same patient with sarcopenia (SMI = 47.9 cm<sup>2</sup>/m<sup>2</sup>) at follow-up

**Table 1** Baseline characteristics of the study patient population

	Overall	Pre-operative sarcopenia		p
		No	Yes	
Number of patients (%)	115	23	92	
Sex (%)				
Female	29 (25.2)	9 (39.1)	20 (21.7)	0.147
Male	86 (74.8)	14 (60.9)	72 (78.3)	
ECOG (%)				
0	40 (35.1)	11 (47.8)	29 (31.9)	0.140
1	69 (60.5)	10 (43.5)	59 (64.8)	
2	5 (4.4)	2 (8.7)	3 (3.3)	
Smoking status (%)				
Current smoker	32 (27.8)	11 (47.8)	21 (22.8)	0.018
Ex-smoker	40 (34.8)	3 (13.0)	37 (40.2)	
Non-smoker	43 (37.4)	9 (39.1)	34 (37.0)	
T stage (%)				
1	2 (1.8)	0 (0.0)	2 (2.2)	0.894
2	10 (8.9)	2 (8.7)	8 (9.0)	
3	86 (76.8)	17 (73.9)	69 (77.5)	
4	14 (12.5)	4 (17.4)	10 (11.2)	
N stage (%)				
0	33 (29.5)	6 (26.1)	27 (30.3)	0.258
1	74 (66.1)	15 (65.2)	59 (66.3)	
2	4 (3.6)	1 (4.3)	3 (3.4)	
3	1 (0.9)	1 (4.3)	0 (0.0)	
Cancer type (%)				
Adenocarcinoma	88 (76.5)	21 (91.3)	67 (72.8)	0.061
SCC*	27 (23.5)	2 (8.7)	25 (27.2)	
Pathological grade** (%)				
1	9 (16.1)	2 (28.6)	7 (14.3)	0.723
2	22 (39.3)	3 (42.9)	19 (38.8)	
3	25 (44.7)	2 (28.6)	23 (46.9)	
Tumor location (%)				
Lower third	95 (82.6)	20 (87.0)	75 (81.5)	0.646
Middle third	17 (14.8)	3 (13.0)	14 (15.2)	
Upper third	3 (2.6)	0 (0.0)	3 (3.3)	
Neoadjuvant treatment (%)				
Chemoradiation	28 (24.3)	2 (8.7)	26 (28.3)	0.051
Chemotherapy	87 (75.7)	21 (91.3)	66 (71.7)	
Operation type (%)				
MIE†	78 (67.8)	18 (78.3)	60 (65.2)	0.464
Thoracotomy	26 (22.6)	2 (8.7)	24 (26.1)	
Hybrid-laparoscopy	5 (4.3)	1 (4.3)	4 (4.3)	
Hybrid-VATS‡	3 (2.6)	1 (4.3)	2 (2.2)	
Transhiatal	3 (2.6)	1 (4.3)	2 (2.2)	
Preop. endo. treatment (%)				
Any	40 (34.8)	4 (17.4)	36 (39.1)	0.087

**Table 1** Baseline characteristics of the study patient population (Continued)

	Overall	Pre-operative sarcopenia		p
		No	Yes	
Stent	35 (30.4)	3 (13.0)	32 (34.8)	0.076
PEG§	4 (3.5)	1 (4.3)	3 (3.3)	1.000
EMR¶	1 (0.9)	1 (4.3)	0 (0.0)	0.451
Age, years (mean [SD])	63 [9]	59 [8]	64 [9]	0.015
Height, cm (median [IQR])	174 [166,179]	171 [160,174]	175 [167,180]	0.009
Preop. weight, kg (mean [SD])	74 [16]	82 [18]	73 [15]	0.017
Weight loss, kg (median [IQR])	7 [0, 13]	6 [0, 12]	8 [2, 12]	0.509
FEV1% (mean [SD])	90 [18]	90 [21]	91 [18]	0.914
Creatinine, umol/l (mean [SD])	71 [19]	72 [18]	71 [20]	0.810
CCI (median [IQR])	5 [4, 6]	5 [4, 6]	5 [5, 6]	0.098

\*Squamous cell carcinoma

\*\*Grade not reported on all pathological reports

†Minimally invasive esophagectomy

‡Video-assisted thoracoscopic surgery

§Percutaneous endoscopic gastrostomy

¶Endoscopic mucosal resection

study had a prior EMR. Patients with obstructing tumor growth and marked dysphagia preventing the ingestion of solid foods were treated with insertion of a self-expandable metallic stent (SEMS, N = 35) or a percutaneous endoscopic gastrostomy (PEG, N = 4) tube.

**Surgical treatment and follow-up**

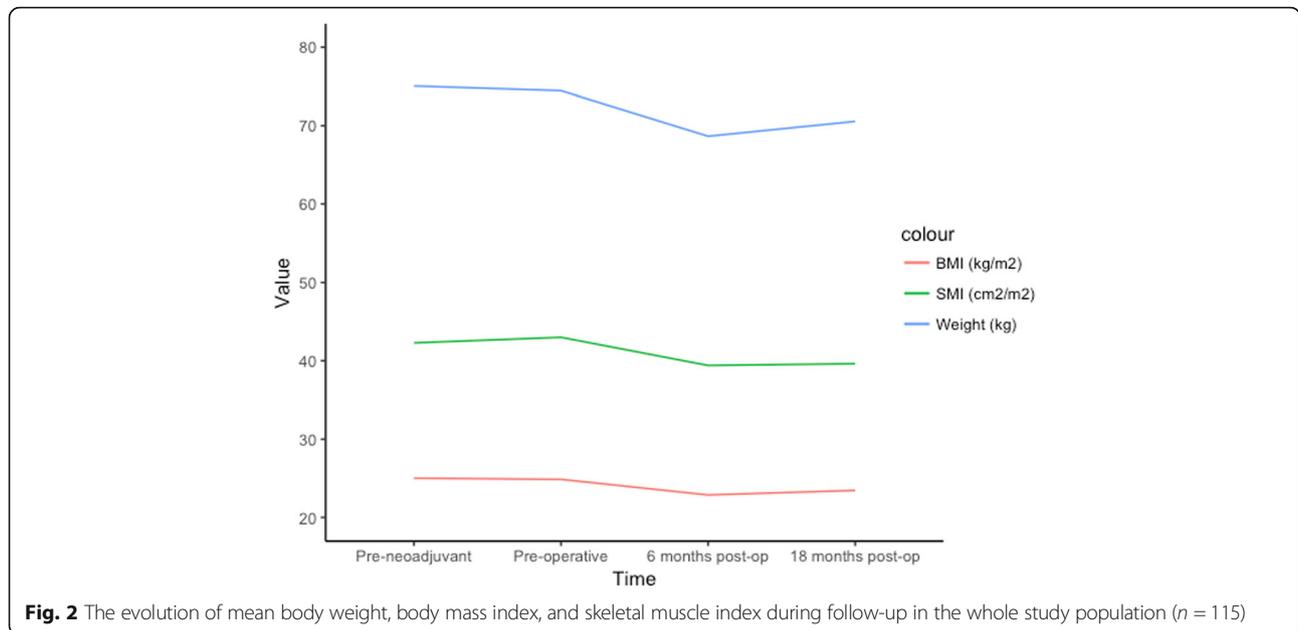
Surgical techniques included minimally invasive esophagectomy, hybrid minimally invasive esophagectomy with either laparoscopy or video-assisted thoracoscopic surgery, Ivor-Lewis esophagectomy, 3-part esophagectomy, and transhiatal esophagectomy. These techniques are described elsewhere [23]. Post-operative stage of the tumor was reported according to eighth edition AJCC/IAUC staging [24]. Amount of positive lymph nodes and total lymph node count was recorded.

Patients were followed until death or January 2017, yielding a follow-up period of at least 24 months. Patients were met at an outpatient clinic 1 month after surgery to assess the recovery from surgery. Gastroscopies were done every 6 months for 2 years after surgery and annually for up to 5 years. CT scans were taken 6 months after surgery, 18 months after surgery, and annually up to 5 years.

All the treatments discussed here are standard approaches in our institution.

**Measurement of muscle parameters and sarcopenia definition**

Scans were coded in order to blind the researcher from outcome. Images were imported to Osirix® Version 3.3



**Table 2** Analysis of complication rates by preoperative sarcopenia

Level	Preoperative Sarcopenia		p
	No (N = 23)	Yes (N = 92)	
30-day mortality (%)	0 (0.0)	3 (3.3)	0.884
90-day mortality (%)	1 (4.3)	6 (6.5)	1.000
Any complication (%)	17 (73.9)	62 (67.4)	0.725
Clavien-Dindo score (%)			0.886
0	6 (26.1)	30 (32.6)	
1	0 (0.0)	3 (3.3)	
2	4 (17.4)	10 (10.9)	
3a	3 (13.0)	13 (14.1)	
3b	7 (30.4)	20 (21.7)	
4a	2 (8.7)	11 (12.0)	
4b	0 (0.0)	2 (2.2)	
5	1 (4.3)	3 (3.3)	
Chyle leak (%)	1 (4.3)	8 (8.7)	0.795
Anastomotic leak (%)	2 (8.7)	13 (14.1)	0.729
Conduit necrosis (%)	1 (4.3)	2 (2.2)	1.000
Recurrent nerve palsy (%)	1 (4.3)	7 (7.6)	0.927
Intraoperative Complications <sup>†</sup> (%)	3 (13.0)	9 (9.8)	0.939
Pulmonary complications <sup>‡</sup> (%)	6 (26.1)	26 (28.3)	1.000
Reoperation rate (%)	2 (8.7)	8 (8.7)	1.000
Operative bleeding, ml (median [IQR])	200 [150, 400]	400 [150, 700]	0.091
ICU stay, days (median [IQR])	3 [1, 5]	2 [1, 4]	0.535
Tracheostomy rate (%)	1 (4.3)	8 (8.7)	0.795

<sup>†</sup>Intraoperative vessel, conduit or airway injury or conversion to open esophagectomy

<sup>‡</sup>Contains ARDS (acute respiratory distress syndrome), pneumonia, atelectasis requiring intervention, pleural effusion or pneumothorax requiring intervention, pulmonary embolism, and acute aspiration

(32-bit Pixmeo, Sarl, Switzerland). We selected a single image on the level of L3, with both transverse processes and delineated abdominal muscles by use of a semi-automated selection of region of interest. Psoas, quadratus lumborum, paraspinal, transverse abdominal, external oblique, internal oblique, and rectus abdominis muscles were included. The Hounsfield unit threshold range for skeletal muscle was -29 to +150. The images were manually corrected, if needed, by the propulsion and brush tools in Osirix®. The cross-sectional total muscle area at the level of L3 (cm<sup>2</sup>) was divided by the square of height (m<sup>2</sup>), which produced the skeletal muscle index (SMI). This method is suggested as the preferred method of measuring the muscle mass of cancer patients [25]. SMI limit for sarcopenia was <52.4 cm<sup>2</sup>/m<sup>2</sup> for men and <38.5 cm<sup>2</sup>/m<sup>2</sup> for women, based on a previous study by Prado et al. [26] For the survival and complication analyses, the preoperative SMI values were used, unless stated otherwise.

The process of delineating the abdominal muscle mass is shown in Fig. 1.

**Statistical analysis**

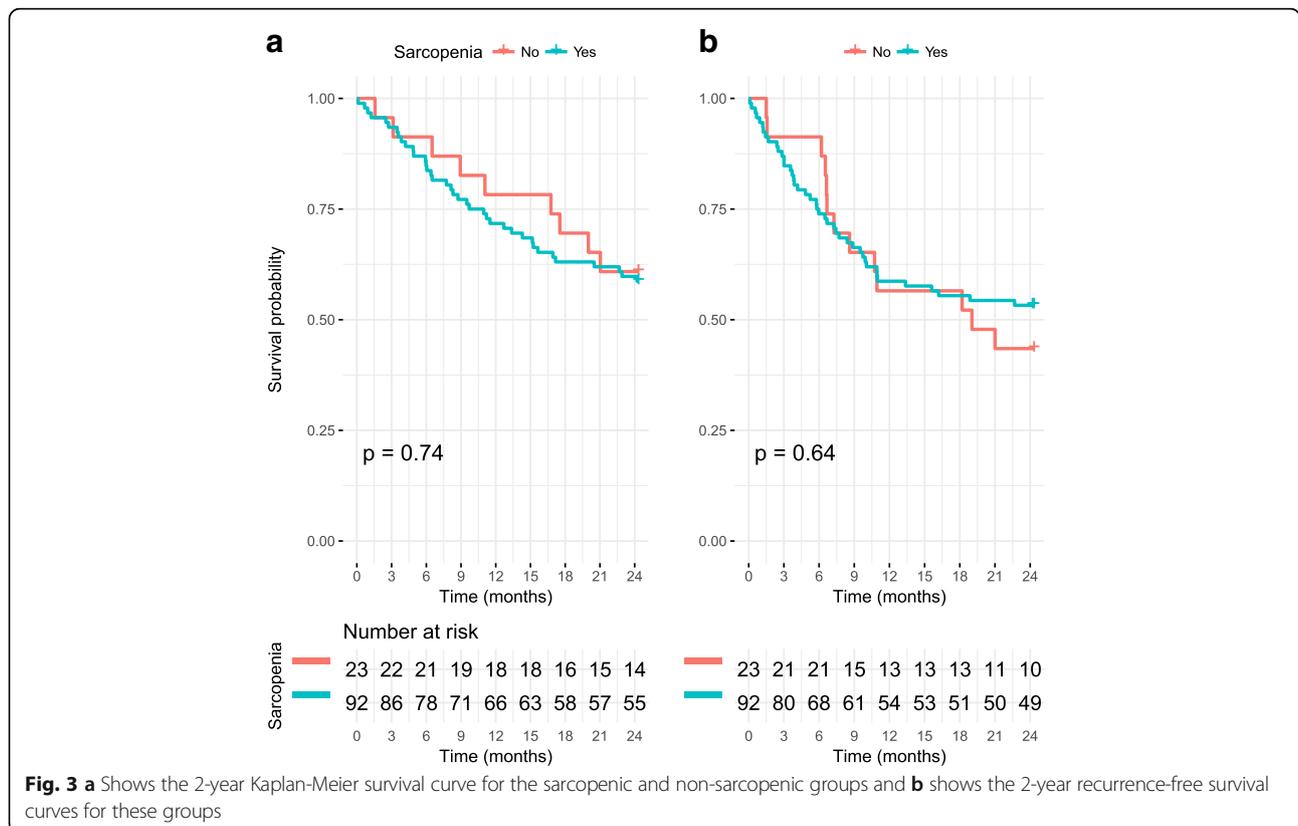
All statistical analysis was done with R Project. (R Core Team, 2016). R: A language and environment for statistical computing. R Foundation for Statistical Computing, Vienna, Austria. URL: <https://www.R-project.org/>).

Continuous parameters were tested for normality using Shapiro-Wilk’s test and deemed normally distributed if  $p > 0.05$ . Normal continuous data is presented as mean and standard deviation (SD) whereas non-normal data is described with median and interquartile range (IQR). For comparing normal scalar variables between two groups, independent samples Student’s *t* test was used and for non-normal variables Mann-Whitney *U* test was used. The 2-tailed  $\chi^2$  test served for categorical variables. Kaplan-Meier survival analysis and log-rank test demonstrated the possible difference of survival between groups.

**Results**

Baseline characteristics of the study population are displayed in Table 1. Included patients numbered 115, of which sarcopenia was found in 92 (80%). Median overall survival was 900 days (interquartile range 334–1447). Patients with and without preoperative sarcopenia are compared in Table 1. The sarcopenic patients were statistically significantly older, taller, weighed less, and had smaller prevalence of current smokers.

The progression of mean weight- and body composition-related parameters during neoadjuvant treatments and follow-up is displayed in Fig. 2. Ninety-one (79.1%) patients had sarcopenia before neoadjuvant treatments, 92 (80%) were sarcopenic before esophagectomy, 82 of 99 (82.8%)



**Table 3** Characteristics by SMI change groups

	SMI change		<i>p</i>
	< - 2.98%	> - 2.98%	
Number of patients (%)	57	58	
Sex (%)			
Female	19 (33.3)	10 (17.2)	0.076
Male	38 (66.7)	48 (82.8)	
ECOG (%)			
0	15 (26.3)	25 (43.9)	0.108
1	40 (70.2)	29 (50.9)	
2	2 (3.5)	3 (5.3)	
Smoking status (%)			
Current smoker	20 (35.1)	12 (20.7)	0.086
Ex-smoker	21 (36.8)	19 (32.8)	
Non-smoker	16 (28.1)	27 (46.6)	
T stage			
1	2 (3.8)	0 (0.0)	0.549
2	5 (9.3)	5 (8.6)	
3	42 (77.8)	44 (75.9)	
4	5 (9.3)	9 (15.5)	
N stage			
0	14 (25.9)	19 (32.8)	0.445
1	38 (70.4)	36 (62.1)	
2	1 (1.9)	3 (5.2)	
3	1 (1.9)	0 (0.0)	
Cancer type (%)			
Adenocarcinoma	40 (70.2)	48 (82.8)	0.111
SCC*	17 (29.8)	10 (17.2)	
Pathological grade**			
1	3 (13.0)	6 (18.2)	0.790
2	9 (39.1)	13 (39.4)	
3	12 (47.9)	14 (42.4)	
Tumor location (%)			
Lower third	45 (78.9)	50 (86.2)	0.357
Middle third	11 (19.3)	6 (10.3)	
Upper third	1 (1.8)	2 (3.4)	
Neoadjuvant treatment (%)			
Chemoradiation	19 (33.3)	11 (19.0)	0.079
Chemotherapy	38 (66.7)	47 (81.0)	
Operation type (%)			
MIE†	39 (68.4)	39 (67.2)	0.931
Thoracotomy	13 (22.8)	13 (22.4)	
Hybrid-laparoscopy	2 (3.5)	3 (5.2)	
Hybrid-VATS‡	2 (3.5)	1 (1.7)	
Transhiatal	1 (1.8)	2 (3.4)	

**Table 3** Characteristics by SMI change groups (Continued)

	SMI change		<i>p</i>
	< - 2.98%	> - 2.98%	
Preop. endo. treatment (%)			
Any	26 (45.6)	14 (24.1)	0.026
Stent	23 (40.4)	12 (20.7)	0.037
PEG§	2 (3.5)	2 (3.4)	1.000
EMR¶	1 (1.8)	0 (0.0)	0.993
Age, years (mean [SD])	63 [9]	63 [9]	0.817
Height, cm (median [IQR])	173 [164,179]	174 [168,179]	0.350
Preop. weight, kg (mean [SD])	74 [17]	75 [16]	0.840
Weight loss, kg (median [IQR])	10 [0, 15]	6 [2, 10]	0.131
FEV1% (mean [SD])	89 [21]	92 [16]	0.527
Creatinine, umol/l (mean [SD])	70 [22]	72 [16]	0.484
CCI (median [IQR])	5 [5, 6]	5 [4, 6]	0.397

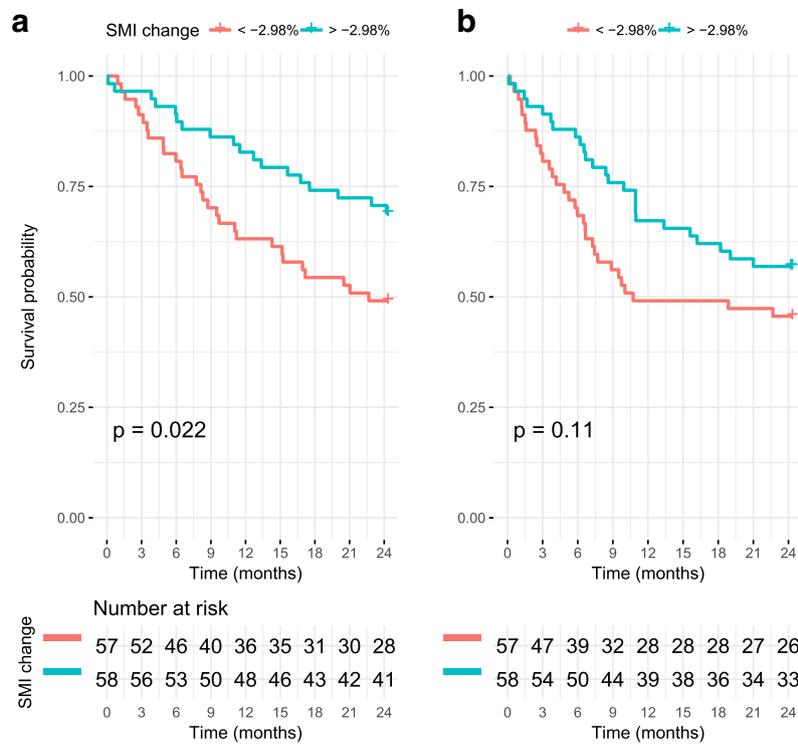
\*Squamous cell carcinoma  
 \*\*Grade not reported on all pathological reports  
 †Minimally invasive esophagectomy  
 ‡Video-assisted thoracoscopic surgery  
 §Percutaneous endoscopic gastrostomy  
 ¶Endoscopic mucosal resection

were sarcopenic 6 months post-operatively, and 67 of 78 (85.9%) 18 months post-operatively.

Table 2 shows the relevant complication rates and Clavien-Dindo scores of patients. The amount of intra-operative bleeding and length of initial intensive care unit stay are also displayed. There was no statistically significant difference between the sarcopenic or non-sarcopenic groups in any complication groups. An additional table shows a more exhaustive table of complication rates [see Additional file 1].

There was no statistical difference in 2-year overall survival or recurrence-free survival between the preoperative sarcopenic and non-sarcopenic groups (*p* = 0.74 and *p* = 0.64, respectively). The Kaplan-Meier curves are shown in Fig. 3a, b. Neither preneoadjuvant SMI nor preoperative SMI had an effect on OS (*p* = 0.6023 and *p* = 0.3843) or RFS (*p* = 0.3241 and *p* = 0.9273).

Patients were divided into groups based on the change between preneoadjuvant measurement and preoperative measurement of SMI. Median percentual change of SMI (- 2.98%) was used as the cut-off value, as has been done in a previous study [15]. The baseline characteristics are displayed in Table 3. The group with more muscle loss had more preoperative stents inserted (*p* = 0.037). SMI change below the median correlated with 2-year overall survival (*p* = 0.022) but not 2-year RFS (*p* = 0.11), as shown in Fig. 4a–b. The change of SMI during neoadjuvant treatments was significantly different between 2-year survivors and non-survivors with mean changes of - 0.69 and - 6.20%, respectively (*p* = 0.01259).



**Fig. 4** Kaplan-Meier survival analyses of the patients divided into groups by skeletal muscle index change by cut-off of  $-5\%$ . **a** Shows the 2-year overall survival rates between the groups and **b** shows the 2-year recurrence free survival

Finally, a multivariate Cox regression analysis was done using the backwards elimination method with a  $p$  value limit of 0.2. The model constructed is shown in Table 4. The model showed that the patients who had percentual SMI change over the median ( $-2.98\%$ ) had better survival than those whose SMI had decreased more ( $p = 0.049$ ; HR  $-0.609$ ; HR 95% CI  $0.297-0.997$ ). No other covariates reached significance (N stage, T stage, or CCI). There was 11.75 events per variable (47 events, 4 variables), which is over the suggested limit of 10 (Table 4) [27].

**Discussion**

Our findings suggest that loss of skeletal muscle mass during neoadjuvant treatment of EC is a marker of poor prognosis. Sarcopenia itself was not correlated with poorer

oncological outcomes; however, its prevalence is high in this population and increases post-esophagectomy.

The amount of skeletal muscle lost during neoadjuvant treatment seems to predict a poorer prognosis. A median cut-off of  $-2.98\%$  produced significantly different 2-year overall survival rates as seen in Fig. 4a. To the authors’ best knowledge, this finding has not been previously reported in published literature, although similar findings have been reported in squamous cell carcinoma patients [15]. A previous study failed to show this correlation, but concluded that the amount of skeletal muscle lost during neoadjuvant therapy differed significantly between survivors and non-survivors, which was confirmed by our study [16]. Skeletal muscle wasting post-operatively has been associated with worse outcomes in thoracic esophageal cancer [28].

**Table 4** Multivariate Cox regression analysis of the covariates affecting OS

	HR	95% CI	$p$
$\Delta$ SMI%* during neoadj. (>median vs. <median)	-0.609	0.297-0.997	0.049
N stage (per level)	0.486	0.940-2.801	0.082
T stage (per level)	0.496	0.993-2.892	0.086
CCI** (per level)	0.175	0.973-1.459	0.091

\*The percentual change in skeletal muscle index during neoadjuvant treatments  
 \*\*Charlson comorbidity index

The progression of sarcopenia is not stopped by resection of the tumor according to our data as shown in Fig. 2. The prevalence of sarcopenia increases with time in follow-up. Similar findings have been reported in the literature [14]. Esophagectomy seems to affect the body composition of many patients. An interventional randomized study did not see statistically significant change in the weight of patients' post-esophagectomy at 7 days, regardless of the method of nutritional support [29]. A previous study has shown that malnutrition and weight loss is common even years after EC surgery [30]. Whether this change in body mass post-esophagectomy contributes to the morbidity or mortality of operatively treated EC patients is unknown.

Sarcopenia was not correlated with worse overall survival or recurrence-free survival in our study. This finding contradicts many previous studies and is supported by some previous studies [10–18]. Complication rates in any of the complication subgroups did not differ significantly between sarcopenic and non-sarcopenic groups. There was a statistically non-significant increase in operative bleeding with sarcopenic patients.

Our study has a number of limitations. The study is retrospective in nature without randomization. This increases the risk for systemic errors and selection bias. The number of patients included in the study is of adequate size taking into account the incidence of this disease, but statistical power is of concern especially in regard to complication rates and specific complication where the number of events is low or non-existent.

## Conclusions

In conclusion, our study found an interesting correlation between the loss of skeletal muscle during neoadjuvant treatment and worse oncological outcomes in surgically treated EC patients. This finding should guide further investigation into the interventions for nutritional support in esophageal cancer and into the significance of indirect measurements of body composition in the prognosis of EC patients.

## Additional file

**Additional file 1:** Complication rate analysis by preoperative sarcopenia. (XLSX 34 kb)

## Abbreviations

AC: Adenocarcinoma; BMI: Body mass index; CCI: Charlson Comorbidity Index; CI: Confidence interval; CRT: Chemoradiotherapy; CT: Computed tomography; EC: Esophageal cancer; ECOG: The Eastern Cooperative Oncology Group; EMR: Endoscopic mucosal resection; EOX: Epirubicin–oxaliplatin–capecitabine; EUS: Endoscopic ultrasound; HU: Hounsfield unit; ICU: Intensive care unit; IQR: Interquartile range; MIE: Minimally invasive esophagectomy; OS: Overall survival; PEG: Percutaneous gastrostomy; PET: Positron emission tomography; RFA: Radiofrequency ablation; RFS: Recurrence-free survival; ROI: Region of interest; SCC: Squamous cell

carcinoma; SD: Standard deviation; SEMs: Self-expandable metallic stent; SMI: Skeletal muscle index; TMA: Total muscle area; VATS: Video assisted thoroscopic surgery

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## Availability of data and materials

The datasets used and/or analyzed during the current study are available from the corresponding author on reasonable request.

## Authors' contributions

TJ contributed to the study design, collected, analyzed, and interpreted the data, wrote the manuscript, and contributed to the editing the manuscript. II contributed to the study design, analysis and interpretation of the data, and editing of the manuscript. JK and JS contributed to the study design and editing of the manuscript. JR was a major contributor to the study design, interpretation of the results, and the editing of the manuscript. All authors read and approved the final manuscript.

## Ethics approval and consent to participate

This study was approved by local medical ethical committees and was done according to the revised version of the Declaration of Helsinki (October 2008, Seoul).

## Consent for publication

Not applicable.

## Competing interests

The authors declare that they have no competing interests.

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## References

- Daly JM, Fry WA, Little AG, Winchester DP, McKee RF, Stewart AK, et al. Esophageal cancer: results of an American College of Surgeons Patient Care Evaluation Study. *J Am Coll Surg*. 2000;190(5):562–72. discussion 72–3
- Di Fiore F, Leclaire S, Pop D, Rigal O, Hamidou H, Paillot B, et al. Baseline nutritional status is predictive of response to treatment and survival in patients treated by definitive chemoradiotherapy for a locally advanced esophageal cancer. *Am J Gastroenterol*. 2007;102(11):2557–63.
- van der Schaaf MK, Tilanus HW, van Lanschot JJB, Johar AM, Lagergren P, Lagergren J, et al. The influence of preoperative weight loss on the postoperative course after esophageal cancer resection. *J Thorac Cardiovasc Surg*. 2014;147(1):490–5.
- Grotenhuis BA, Wijnhoven BP, Hotte GJ, van der Stok EP, Tilanus HW, van Lanschot JJ. Prognostic value of body mass index on short-term and long-term outcome after resection of esophageal cancer. *World J Surg*. 2010; 34(11):2621–7.
- Blom R, Lagarde S, Klinkenbijl J, Busch O, MIV H. A high body mass index in esophageal cancer patients does not influence postoperative outcome or long-term survival. *Ann Surg Oncol*. 2012;19(3):766–71.
- Cruz-Jentoft AJ, Baeyens JP, Bauer JM, Boirie Y, Cederholm T, Landi F, et al. Sarcopenia: European consensus on definition and diagnosis: report of the European working group on sarcopenia in older people. *Age Ageing*. 2010; 39(4):412–23.
- Harimoto N, Shirabe K, Yamashita YI, Ikegami T, Yoshizumi T, Szejima Y, et al. Sarcopenia as a predictor of prognosis in patients following hepatectomy for hepatocellular carcinoma. *Brit J Surg*. 2013;100(11): 1523–30.

8. Miyamoto Y, Baba Y, Sakamoto Y, Ohuchi M, Tokunaga R, Kurashige J, et al. Sarcopenia is a negative prognostic factor after curative resection of colorectal cancer. *Ann Surg Oncol*. 2015;22(8):2663–8.
9. Go SI, Park MJ, Song HN, Kang MH, Park HJ, Jeon KN, et al. Sarcopenia and inflammation are independent predictors of survival in male patients newly diagnosed with small cell lung cancer. *Support Care Cancer*. 2016;24(5):2075–84.
10. Tamandl D, Paireder M, Asari R, Baltzer PA, Schoppmann SF, Ba-Ssalamah A. Markers of sarcopenia quantified by computed tomography predict adverse long-term outcome in patients with resected oesophageal or gastro-oesophageal junction cancer. *Eur Radiol*. 2016;26(5):1359–67.
11. Kudou K, Saeki H, Nakashima Y, Edahiro K, Korehisa S, Taniguchi D, et al. Prognostic significance of sarcopenia in patients with Esophagogastric junction cancer or upper gastric cancer. *Ann Surg Oncol*. 2017;24(7):1804–10.
12. Paireder M, Asari R, Kristo I, Rieder E, Tamandl D, Ba-Ssalamah A, et al. Impact of sarcopenia on outcome in patients with esophageal resection following neoadjuvant chemotherapy for esophageal cancer. *Eur J Surg Oncol*. 2017;43(2):478–84.
13. Nishigori T, Okabe H, Tanaka E, Tsunoda S, Hisamori S, Sakai Y. Sarcopenia as a predictor of pulmonary complications after Esophagectomy for thoracic esophageal cancer. *J Surg Oncol*. 2016;113(6):678–84.
14. Elliott JA, Doyle SL, Murphy CF, King S, Guinan EM, Beddy P, et al. Sarcopenia: prevalence, and impact on operative and oncologic outcomes in the multimodal management of locally advanced esophageal cancer. *Ann Surg*. 2017;266(5):822–30.
15. Liu JJ, Motoyama S, Sato Y, Wakita A, Kawakita Y, Saito H, et al. Decreased skeletal muscle mass after neoadjuvant therapy correlates with poor prognosis in patients with esophageal cancer. *Anticancer Res*. 2016;36(12):6677–85.
16. Reisinger KW, Bosmans JWAM, Uittenbogaart M, Alsoumaili A, Poeze M, Sosef MN, et al. Loss of skeletal muscle mass during Neoadjuvant chemoradiotherapy predicts postoperative mortality in esophageal cancer surgery. *Ann Surg Oncol*. 2015;22(13):4445–52.
17. Grotenhuis BA, Shapiro J, van Adrichem S, de Vries M, Koek M, Wijnhoven BP, et al. Sarcopenia/muscle mass is not a prognostic factor for short- and long-term outcome after esophagectomy for cancer. *World J Surg*. 2016;40(11):2698–704.
18. Awad S, Tan BH, Cui H, Bhalla A, Fearon KCH, Parsons SL, et al. Marked changes in body composition following neoadjuvant chemotherapy for oesophagogastric cancer. *Clin Nutr*. 2012;31(1):74–7.
19. Yip C, Goh V, Davies A, Gossage J, Mitchell-Hay R, Hynes O, et al. Assessment of sarcopenia and changes in body composition after neoadjuvant chemotherapy and associations with clinical outcomes in oesophageal cancer. *Eur Radiol*. 2014;24(5):998–1005.
20. Sheetz KH, Zhao L, Holcombe SA, Wang SC, Reddy RM, Lin J, et al. Decreased core muscle size is associated with worse patient survival following esophagectomy for cancer. *Dis Esophagus*. 2013;26(7):716–22.
21. Low DE, Alderson D, Ceconello I, Chang AC, Darling GE, D'Journo XB, et al. International consensus on standardization of data collection for complications associated with esophagectomy esophagectomy complications consensus group (ECCG). *Ann Surg*. 2015;262(2):286–94.
22. Cunningham D, Allum WH, Stenning SP, Thompson JN, Van de Velde CJH, Nicolson M, et al. Perioperative chemotherapy versus surgery alone for resectable gastroesophageal cancer. *New Engl J Med*. 2006;355(1):11–20.
23. Stiles BM, Altorki NK. Traditional techniques of esophagectomy. *Surg Clin North Am*. 2012;92(5):1249–63.
24. Rice TW, Ishwaran H, Hofstetter WL, Kelsen DP, Apperson-Hansen C, Blackstone EH, et al. Recommendations for pathologic staging (pTNM) of cancer of the esophagus and esophagogastric junction for the 8th edition AJCC/UICC staging manuals. *Dis Esophagus*. 2016;29(8):897–905.
25. Fearon K, Strasser F, Anker SD, Bosaeus I, Bruera E, Fainsinger RL, et al. Definition and classification of cancer cachexia: an international consensus. *Lancet Oncol*. 2011;12(5):489–95.
26. Prado CMM, Liefers JR, McCargar LJ, Reiman T, Sawyer MB, Martin L, et al. Prevalence and clinical implications of sarcopenic obesity in patients with solid tumours of the respiratory and gastrointestinal tracts: a population-based study. *Lancet Oncol*. 2008;9(7):629–35.
27. Vittinghoff E, Mc Culloch CE. Relaxing the rule of ten events per variable in logistic and cox regression. *Am J Epidemiol*. 2007;165(6):710–8.
28. Mayanagi S, Tsubosa Y, Omae K, Niihara M, Uchida T, Tsushima T, et al. Negative impact of skeletal muscle wasting after neoadjuvant chemotherapy followed by surgery on survival for patients with thoracic esophageal cancer. *Ann Surg Oncol*. 2017;24(12):3741–7.
29. Page RD, Oo AY, Russell GN, Pennefather SH. Intravenous hydration versus naso-jejunal enteral feeding after esophagectomy: a randomised study. *Eur J Cardio-Thorac*. 2002;22(5):666–72.
30. Martin L, Lagergren J, Lindblad M, Rouvelas I, Lagergren P. Malnutrition after oesophageal cancer surgery in Sweden. *Br J Surg*. 2007;94(12):1496–500.

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