



# Nomogram for the prediction of individualized overall survival of patients diagnosed with small cell esophageal carcinoma

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**Background:** A nomogram was developed for the estimation of individualized overall survival (OS) of patients diagnosed with small cell esophageal carcinoma (SCEC).

**Methods:** From the SEER dataset, 427 patients diagnosed with SCEC during the period from 2004 to 2015 were selected as training sets. For the establishment of a nomogram capable of estimating the OS possibility of patients diagnosed with SCEC, a group of independent prognostic factors were identified and incorporated. The effectiveness of the nomogram was then both externally and internally verified among 159 patients from Fudan University Shanghai Cancer Center (FUSCC) who were diagnosed with SCEC between 2006 and 2015. The predictive accuracy and discriminative ability of the nomogram were measured by concordance index (C-index). Comparisons between nomogram and the AJCC staging systems (6th and 7th) were performed with calibration plots and area under the curves (AUC) values.

**Results:** We identified age, gender, primary site, SEER stage, surgery, radiotherapy, and chemotherapy as seven independent risk factors which were then used to set up the nomogram. Calibration curves indicated that the prediction of the nomogram was consistent with real observations for the possibilities of 1-, 3-, and 5-year OS, and applying the nomogram to the cohort for validation led to reproducible results. Moreover, the C-indices and AUC values were higher in the nomogram than those in the AJCC staging system AJCC which is also aimed at the prediction of OS.

**Conclusions:** This study resulted in the establishment of the first nomogram for the prediction of individualized OS of patients diagnosed with SCEC. The accuracy rate of prediction of this model may be higher than previously established staging systems.

**Keywords:** Small cell esophageal carcinoma (SCEC); overall survival (OS); prognostic model

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## Introduction

Neuroendocrine carcinoma (NEC) is an esophageal tumour with low incidence, constituting only 3% of all esophageal cancers (1,2). Of tumours classified by the WHO

originating in the digestive system, NECs are considered as poorly-differentiated carcinomas, whose mitotic count is higher than 20 per 10 high power field, and whose index of proliferation is higher than 20 percent. Both small

and large cell carcinomas exist in NECs (3,4). Primary small cell esophageal carcinoma (PSCEC) is the most common extrapulmonary small cell carcinomas (5,6) and its prevalence is increasing overtime (7,8). Most patients with SCEC have a poor prognosis, with a median survival time of is only 8–13 months following diagnosis (5).

Due to the rarity and tumour heterogeneity of SCEC, a specific staging system, clinical trials, or standard treatment have not as yet been established. Additionally, as SCEC is a highly aggressive malignancy with dismal survival, considerable controversy regarding its prognostic factors exists, with the few studies investigating this being retrospective, with small sample size, and producing inconsistent results (9-13).

A nomogram is a statistical model based on a set of vital factors and aimed to estimate the possible result of a clinical event, such as the recurrence rate of a disease or the possibility of death (14). It is widely regarded as a tool that can reliably predict the outcomes of clinical treatments of numerous cancers, and in some (15,16), has been verified as producing higher accuracy than previously developed staging systems. While nomograms are now used as alternative tools and are even seen as a novel criterion to manage cancer patients, none have been established to estimate the OS of patients diagnosed with SCEC. Therefore, it is imperative that investigations should be made in which SCEC OS is estimated on the basis of a large-scale cohort of patients.

For a precise estimation of the clinical treatment effect of patients diagnosed with SCEC, a nomogram was set up to investigate the contribution of each clinical variable bearing significance, as well as the relevant SEER data. In addition, an independent group of SCEC patients from Fudan University Shanghai Cancer Center (FUSCC) were employed to externally validate the model. We present the following article in accordance with the TRIPOD reporting checklist (available at <https://dx.doi.org/10.21037/atm-21-3900>).

## Methods

### *Study population*

This retrospective research was approved by the Institutional Review Board of FUSCC. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013). With the support of the National Cancer Institute, information was sourced and obtained from 18 cancer registries in the SEER dataset which cover around one third of the total population of the United States. By referring to the 2008 ICD-0-3/WHO between 2004 and 2015, all patients

diagnosed with SCEC were selected. Several covariates were also investigated incorporating race, gender, age when the patient was diagnosed, primary tumour location, registration region, marital status, SEER summary stage, tumour-node-metastasis (TNM) stage (6th and 7th edition), surgery type, and the use of chemotherapy and radiation. On the other hand, SCEC patients treated in the FUSCC between 2006 and 2015 were also selected as a validation cohort, and only those diagnosed with SCEC in the FUSCC for the first time were chosen, while those treated with anti-cancer therapies such as radiotherapy and chemotherapy were not considered. In addition to the aforementioned variables, performance status score (PS), smoking status, alcohol use, food habit, vascular invasion, perineural involvement, and pathology (pure or mixed SCEC) were also sought and included, while patients whose information was incomplete were excluded.

### *Statistical analyses*

Statistical analysis was performed using R-3.5.2 for Windows. OS was defined as the duration from the time when a patient was initially diagnosed with SCEC to the time when they died or attended their last follow-up, and was the primary endpoint of the research. Continuous data were analyzed using Student's t test and categorical data using Chi-square test. Log-rank tests and Kaplan-Meier survival curves were employed to compare OS between different subgroups of SCEC patients and to delineate features influencing OS. Univariate and multivariable Cox regression analyses were applied for survival analysis. Using the methods in our previous study, a novel nomogram was built based on prognostic parameters derived from the SEER data (17). To evaluate whether the survival predicted by the nomogram was consistent with the observed survival, we referred to its calibration curves, which were validated by 1,000 resamples. We investigated the prognostic performance of the nomogram by using both calibration plots and time-dependent receiver operating characteristic (tdROC) curve analyses, and further compared its performance with that of the AJCC staging system through calculating C-indices and area under the curves (AUC) values. A two-sided P value smaller than 0.05 suggested statistical significance.

## Results

### *Patient baseline characteristics*

We retrospectively enrolled 427 patients diagnosed with SCEC in the SEER database between 2004 and 2015. To

verify the effectiveness of the nomogram, 159 patients were gathered as a cohort from the FUSCC through referring to the set inclusion and exclusion standards, and their baseline clinical pathologic features are summarized in *Table 1*. Most patients, both in the training group and in the verification group, were male, married, had lymph node metastasis, and received chemotherapy. Lower thoracic SCEC was the most common primary tumor site (44%) in the SEER database, while middle thoracic SCEC (64.8%) was the most common site in the SEER database. The SEER cohort were relatively older (mean 67.4±11.4 years-of-age) at diagnosis than the validation cohort (mean 61.0±7.6 years-of-age), were associated with higher probability of distant metastasis (49.4%), and were more likely to have undergone radiotherapy (52.5%). By contrast in the validation cohort, as a regional stage predominated (45.3%), most patients received surgery (60.4%), while radiotherapy only involved 40.9% of patients. As listed in *Table 1*, we retrospectively collected information on PS, pathology (pure or mixed SCEC), smoking status, alcohol use, food habits, vascular invasion, perineural involvement, and surgical type in the validation group.

#### ***Independent prognostic factors in the training cohort***

A univariate analysis was performed in the training cohort to clarify the clinical parameters having vital linkages with the OS of SCEC patients. *Figure 1A* shows that race, gender, age, marital status, CHSDA region, primary tumour site, surgery, SEER stage, and clinical therapies such as chemotherapy and radiotherapy display remarkable associations with OS. However, multivariate analysis showed that race, marital status, and CHSDA region lost their vital roles, while the remaining variables remained significant (*Figure 1B*). The 3-year OS rates for patients with SEER localized, regional, and distant stages were 27.7%, 20.5%, and 2.4%, respectively. The Kaplan-Meier survival curves for different SEER stages are shown in *Figure 2A*. Of note, neither the 6th or 7th AJCC TNM staging system was an independent prognostic factor in the Cox regression model, and the Kaplan-Meier survival curves were overlapped among AJCC stages I–IV (*Figure 2B,2C*). Better survival was found among the patients treated with chemotherapy than those never receiving it, and radiotherapy and surgery were also found to be associated with favourable outcomes.

#### ***Prognostic nomogram for OS***

A nomogram incorporating all variables of significance

identified by using the covariates mentioned above was developed, and as shown in *Figure 3*, the nomogram predicting 1-, 3-, and 5-year OS was built on the basis of chosen parameters that had a hazard ratio. The nomogram showed the greatest contributing factor to prognosis was the SEER distant stage, followed by chemotherapy, unknown SEER stage and regional stage, radiotherapy, upper primary tumour site, and male gender. *Table S1* shows the exact values of all predictive parameters of the nomogram. Through adding all these values and placing them into the total-point scale, the survival probabilities were calculated.

#### ***Validation of the predictive accuracy of the nomogram for OS***

The C-index of the nomogram, which was employed to indicate OS, was 0.71 (95% CI, 0.67–0.75) in the training set by using 1,000× bootstrap resampling. The predicted OS was further verified externally in the separate group of 159 patients from the FUSCC, who had been diagnosed with SCEC. The calibration plots presented an optimal agreement between the outcomes predicted by the nomogram and those actually observed in terms of the 1-, 3-, and 5-year OS (*Figure 4A-4C*). Internal and external validation results both suggested appreciable reliability of the nomogram.

#### ***Comparison of the nomogram with AJCC staging systems***

We did comparisons between the SCEC nomogram with the 6th and 7th AJCC TNM staging systems by calculating C-indices and area under the curves (AUC) values to access the prediction accuracy. The nomogram yielded a significantly larger C-index for predicting OS in the group aimed at verification compared with that of the AJCC stages [nomogram: 0.71 (95% CI, 0.67–0.75); TNM 6th: 0.63 (95% CI, 0.59–0.67); TNM 7th: 0.64 (95% CI, 0.60–0.68);  $P < 0.001$ ]. *Figure 4D-4F* and *Table S2* show the AUC value of the nomogram was slightly higher than that of the TNM staging system (nomogram *vs.* TNM 6th *vs.* 7th, 1-year OS rate, 0.92 *vs.* 0.59 *vs.* 0.81; 3-year OS rate, 0.89 *vs.* 0.67 *vs.* 0.65; 5-year OS rate, 0.83 *vs.* 0.61 *vs.* 0.66). These results implied both the short-term OS and the long-term OS were predicted more accurately by the nomogram than the AJCC staging systems.

## **Discussion**

SCEC is an exceptionally rare diagnosis, with its prognostic

**Table 1** Basic demographic and patient characteristics in the training and validation cohort of SCEC patients

Variables	FUSCC (n=159)	SEER (n=427)	P
Race (%)			<0.001
NR	159 (100.0)	0 (0.0)	
Others	0 (0.0)	94 (22.0)	
White	0 (0.0)	333 (78.0)	
Gender (%)			0.001
Female	38 (23.9)	168 (39.3)	
Male	121 (76.1)	259 (60.7)	
CHSDA region (%)			<0.001
East	0 (0.0)	153 (35.8)	
Northern Plains	0 (0.0)	84 (19.7)	
NR	159 (100.0)	0 (0.0)	
Pacific Coast	0 (0.0)	167 (39.1)	
Southwest	0 (0.0)	23 (5.4)	
Primary site (%)			<0.001
Lower	35 (22.0)	188 (44.0)	
Middle	103 (64.8)	105 (24.6)	
Others	7 (4.4)	101 (23.7)	
Upper	14 (8.8)	33 (7.7)	
TNM 7th (%)			<0.001
I	2 (1.3)	0 (0.0)	
IA	12 (7.5)	5 (1.2)	
IB	14 (8.8)	24 (5.6)	
IIA	4 (2.5)	24 (5.6)	
IIB	31 (19.5)	19 (4.4)	
III	1 (0.6)	0 (0.0)	
IIIA	17 (10.7)	28 (6.6)	
IIIB	7 (4.4)	0 (0.0)	
IIIC	20 (12.6)	21 (4.9)	
IV	51 (32.1)	168 (39.3)	
NR	0 (0.0)	138 (32.3)	

**Table 1** (continued)**Table 1** (continued)

Variables	FUSCC (n=159)	SEER (n=427)	P
TNM 6th (%)			<0.001
I	14 (8.8)	28 (6.6)	
IIA	18 (11.3)	28 (6.6)	
IIB	37 (23.3)	14 (3.3)	
III	37 (23.3)	46 (10.8)	
IIIA	1 (0.6)	0 (0.0)	
IVA	8 (5.0)	8 (1.9)	
IVB	44 (27.7)	144 (33.7)	
IV NOS	0 (0.0)	20 (4.6)	
NR	0 (0.0)	139 (32.6)	
T stage 7th (%)			<0.001
T1	8 (5.0)	0 (0.0)	
T1a	0 (0.0)	51 (11.9)	
T1b	23 (14.5)	8 (1.9)	
T2	50 (31.4)	41 (9.6)	
T3	51 (32.1)	50 (11.7)	
T4	3 (1.9)	0 (0.0)	
T4a	10 (6.3)	38 (8.9)	
T4b	11 (6.9)	16 (3.7)	
TX	3 (1.9)	223 (52.2)	
N stage 7th (%)			<0.001
N0	38 (23.9)	114 (26.7)	
N1	58 (36.5)	133 (31.1)	
N2	44 (27.7)	7 (1.6)	
N3	19 (11.9)	0 (0.0)	
NX	0 (0.0)	173 (40.5)	
M stage 7th (%)			<0.001
M0	108 (67.9)	143 (33.5)	
M1	51 (32.1)	168 (39.3)	
MX	0 (0.0)	116 (27.2)	
T stage 6th (%)			<0.001
T1	31 (19.5)	59 (13.8)	
T2	50 (31.4)	41 (9.6)	
T3	51 (32.1)	50 (11.7)	
T4	24 (15.1)	54 (12.6)	
TX	3 (1.9)	223 (52.2)	

**Table 1** (continued)

Table 1 (continued)

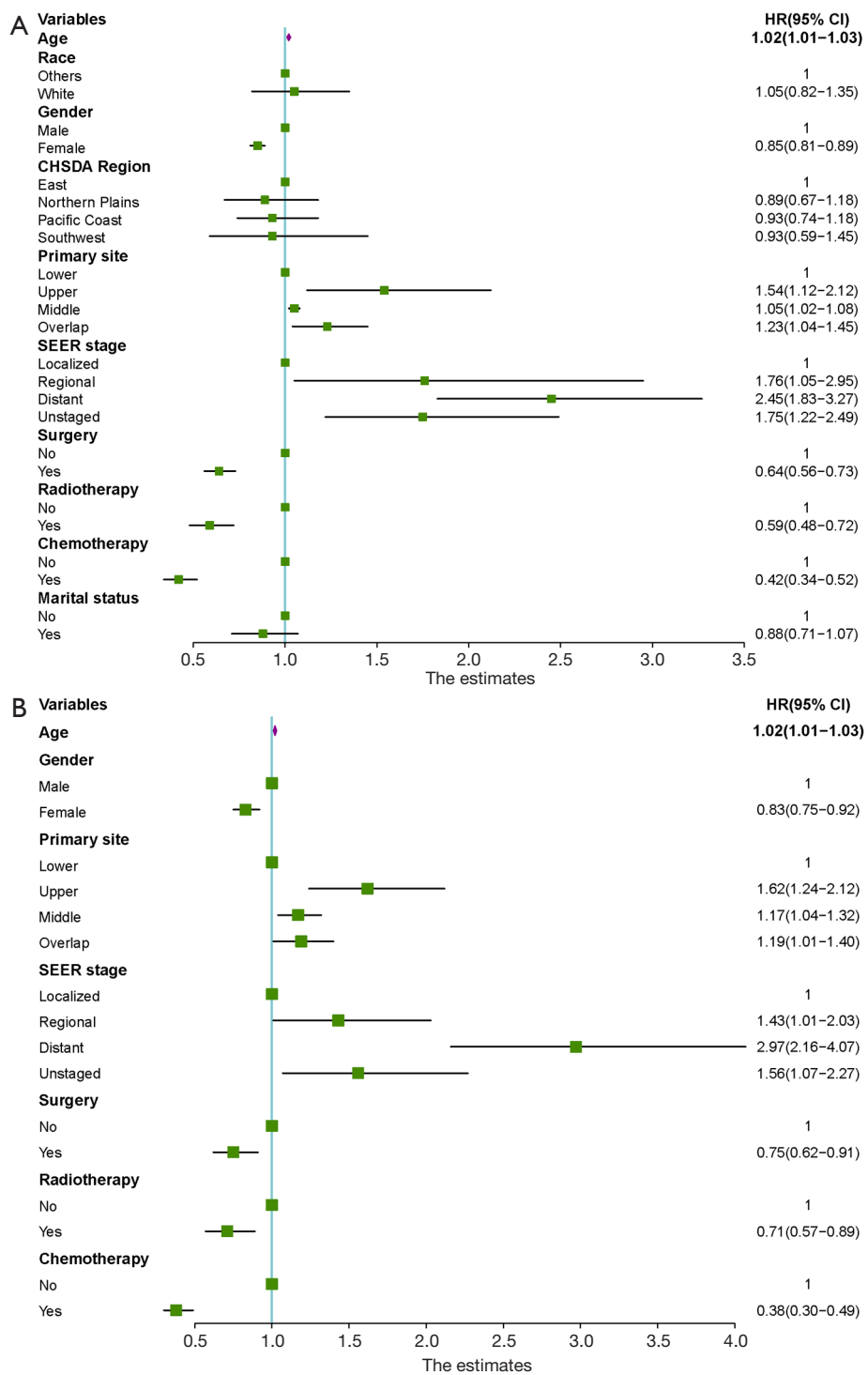
Variables	FUSCC (n=159)	SEER (n=427)	P
N stage 6th (%)			<0.001
N0	40 (25.2)	117 (27.4)	
N1	119 (74.8)	136 (31.9)	
NX	0 (0.0)	174 (40.7)	
M stage 6th (%)			<0.001
M0	108 (67.9)	139 (32.6)	
M1	1 (0.6)	20 (4.7)	
M1a	8 (5.0)	8 (1.9)	
M1b	42 (26.4)	144 (33.7)	
MX	0 (0.0)	116 (27.2)	
SEER stage (%)			<0.001
Localized	37 (23.3)	77 (18.0)	
Regional	72 (45.3)	76 (17.8)	
Distant	50 (31.4)	211 (49.4)	
Unstaged	0 (0.0)	63 (14.8)	
VALSG stage			
Limited	115 (72.3)	NR	
Extensive	44 (27.7)	NR	
Surgery (%)			<0.001
No	63 (39.6)	388 (90.9)	
Yes	96 (60.4)	39 (9.1)	
Radiotherapy (%)			0.016
No	94 (59.1)	203 (47.5)	
Yes	65 (40.9)	224 (52.5)	
Chemotherapy (%)			0.006
No	29 (18.2)	128 (30.0)	
Yes	130 (81.8)	299 (70.0)	
Age [mean (SD)]	60.99 (7.64)	67.37 (11.37)	<0.001
Marital status (%)			<0.001
No	0 (0.0)	192 (45.0)	
Yes	159 (100.0)	235 (55.0)	
Performance status (%)			<0.001
0	61 (38.4)	0 (0.0)	
1	94 (59.1)	0 (0.0)	
2	4 (2.5)	0 (0.0)	
NR	0 (0.0)	427 (100.0)	

Table 1 (continued)

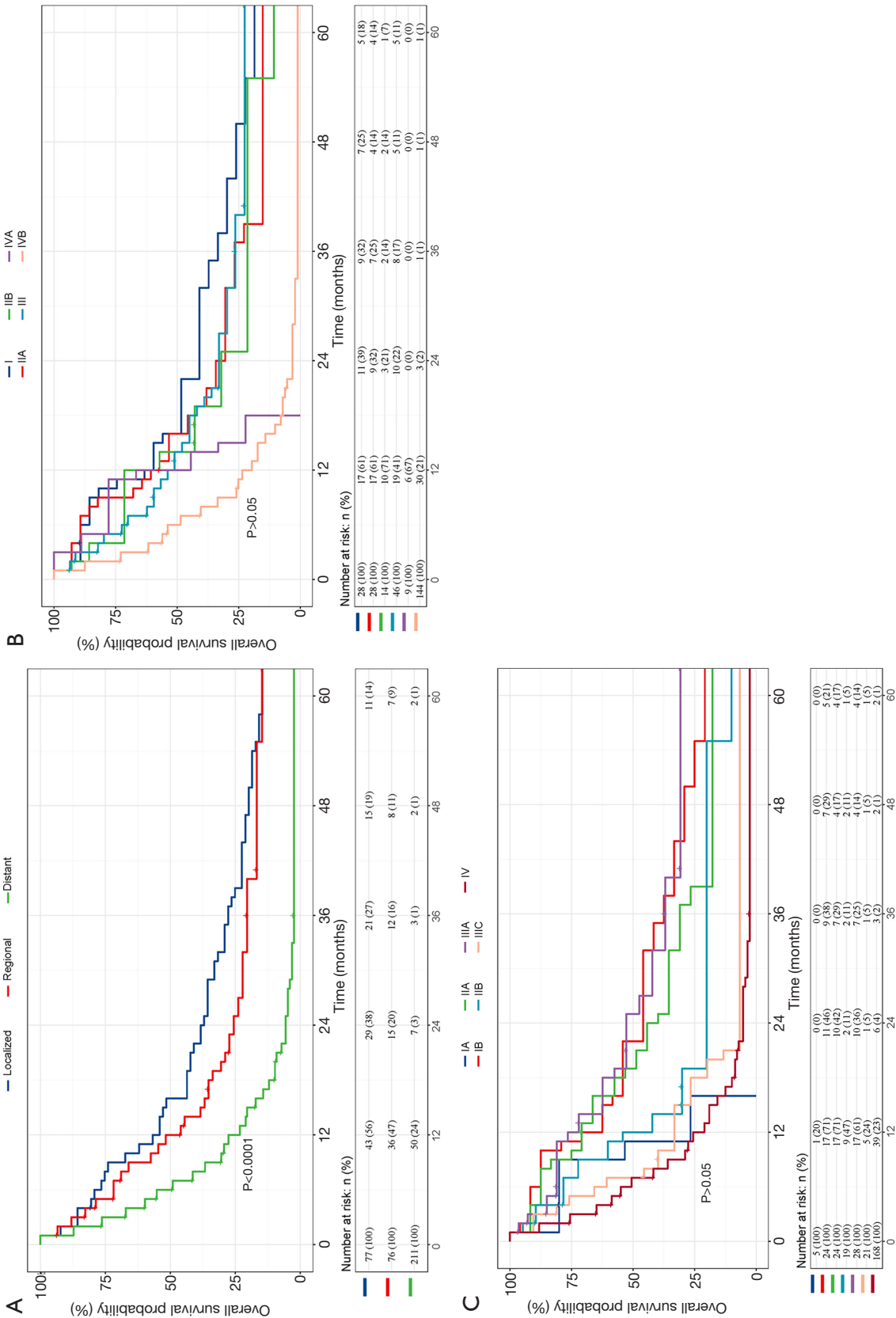
Table 1 (continued)

Variables	FUSCC (n=159)	SEER (n=427)	P
Pathology			
Pure SCEC	131 (82.4)	NR	
Mixed SCEC	28 (17.6)	NR	
Smoking status (%)			<0.001
N	68 (42.8)	0 (0.0)	
NR	0 (0.0)	427 (100.0)	
Y	91 (57.2)	0 (0.0)	
Alcohol status (%)			<0.001
N	91 (57.2)	0 (0.0)	
NR	0 (0.0)	427 (100.0)	
Y	68 (42.8)	0 (0.0)	
Food habit (%)			<0.001
N	76 (47.8)	0 (0.0)	
NR	19 (11.9)	427 (100.0)	
Y	64 (40.3)	0 (0.0)	
Vascular invasion (%)			<0.001
NR	70 (44.0)	427 (100.0)	
Y	36 (22.6)	0 (0.0)	
N	73 (45.9)	0 (0.0)	
Perineural involvement (%)			<0.001
NR	70 (44.0)	427 (100.0)	
Y	16 (10.1)	0 (0.0)	
Ivor-Lewis	62 (39.0)	0 (0.0)	
Surgical type (%)			<0.001
McKeown	8 (5.0)	0 (0.0)	
No	70 (44.0)	0 (0.0)	
NR	0 (0.0)	427 (100.0)	
Palliative resection	3 (1.9)	0 (0.0)	
Sweet's	3 (1.9)	0 (0.0)	
Three field	13 (8.2)	0 (0.0)	

FUSCC, Fudan University Shanghai Cancer Center; SEER, Surveillance, Epidemiology, and End Results database; NR, not reported; CHSDA, Contract Health Service Delivery Areas; NOS, not of specific; VALSG, Veterans Administration Lung Study Group; SD, standard deviation; SCEC, small cell oesophageal carcinoma.

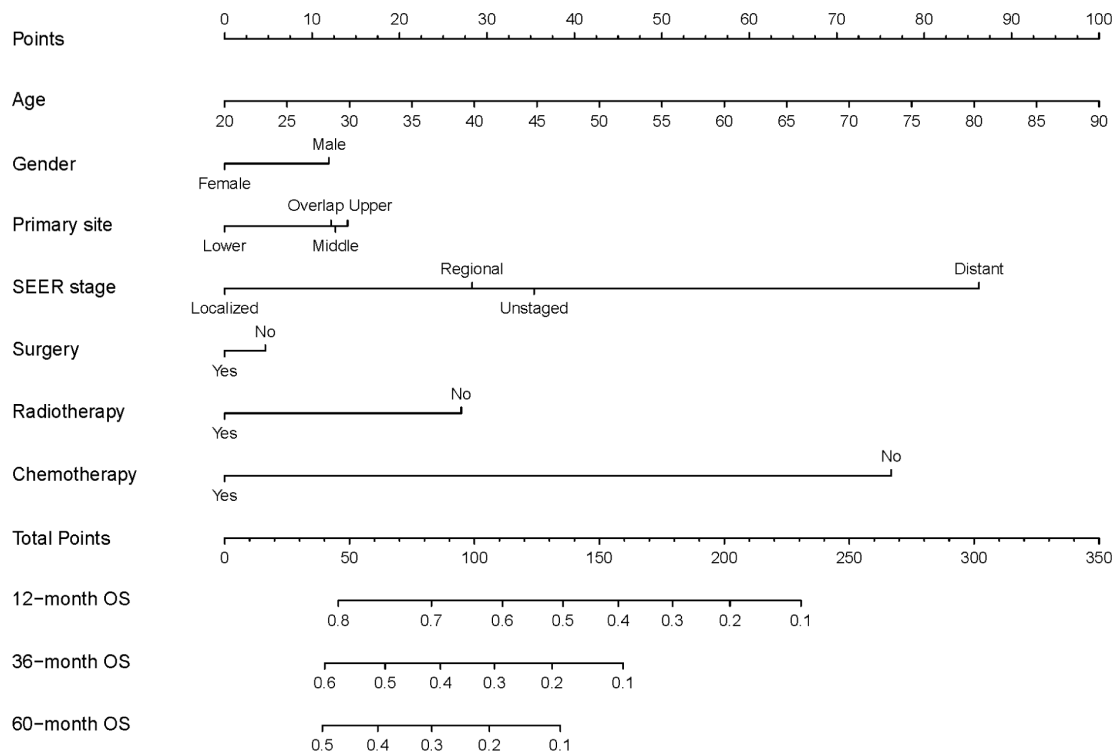


**Figure 1** Cox regression analysis for OS in SCEC patients from the training cohort. Univariate (A) and multivariate (B) Cox regression in the SEER cohort. SEER, Surveillance, Epidemiology, and End Results database; SCEC, small cell oesophageal carcinoma.



**Figure 2** Overall survival in the training cohort for SCEC patients based on different staging system. (A) SEER stage, (B) AJCC 6thstage, (C) AJCC 7th stage. SEER, Surveillance, Epidemiology, and End Results database; SCEC, small cell oesophageal carcinoma; VALSG, Veterans Administration Lung Study Group; AJCC, American Joint Committee on Cancer.



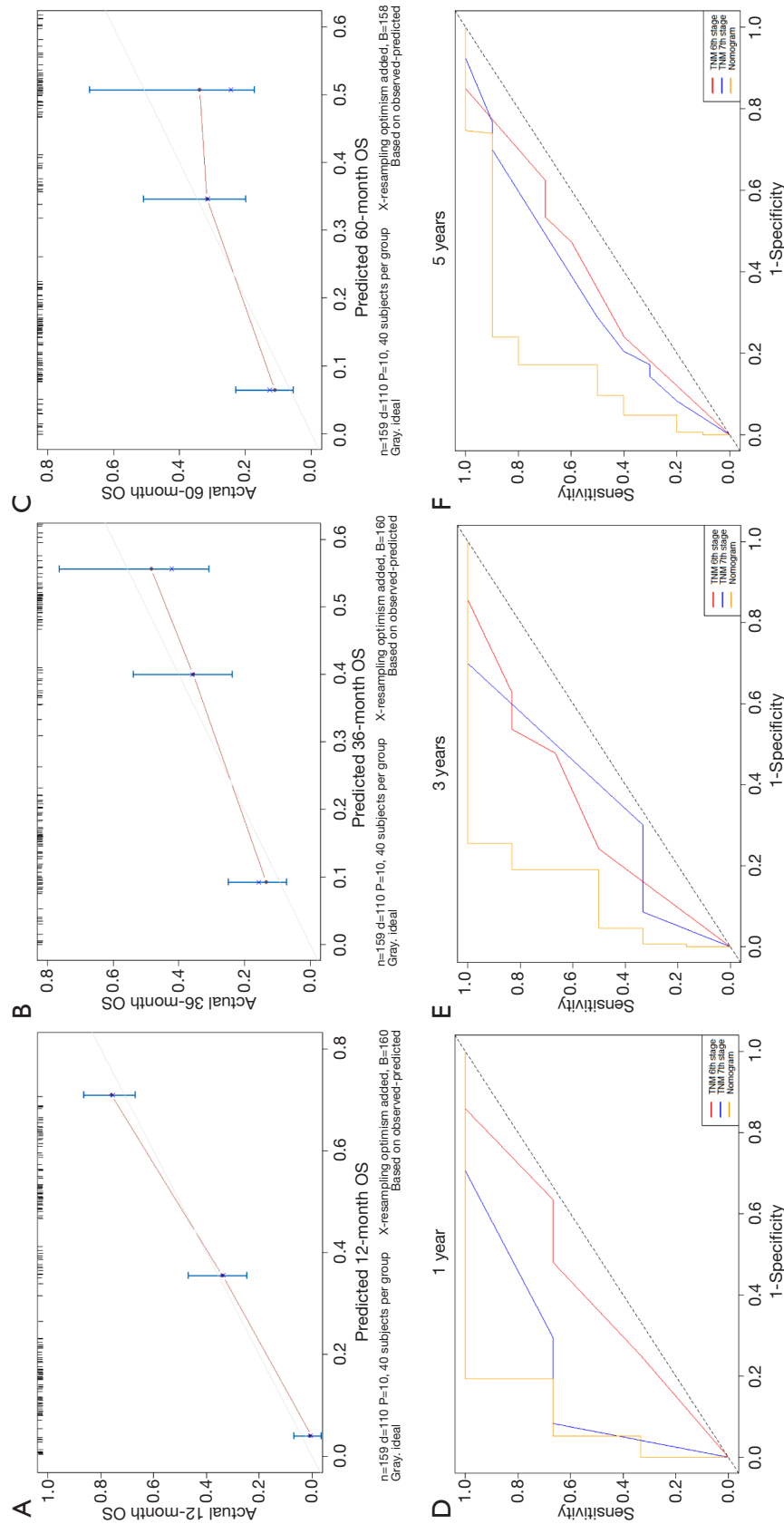


**Figure 3** Nomogram estimating 1-, 3-, and 5-year overall survival for SCEC patients based on the SEER cohort. SCEC, small cell oesophageal carcinoma; SEER, Surveillance, Epidemiology, and End Results database.

factors and treatment strategy poorly understood. As existing knowledge on these factors has largely been derived from small series or case reports, there is an urgent need to more accurately determine the survival characteristics of SCEC patients. In this study, the data of many SCEC patients were obtained from the SEER dataset, and a comprehensive nomogram was established for the predictive calculation of their OS. The model showed satisfactory agreement in its application to both training and validation groups of SCEC patients, and obtained superiority over AJCC staging systems, with more accurate individual prediction of survival probability, demonstrating its clinical application. The extremely low incidence of SCEC has led to there being no specific staging system for the disease, and to date, the TNM-based AJCC staging systems have been adopted to assess the survival of patients. However, our results showed the predictive value of the 6th and 7th AJCC staging systems for the estimation of OS in SCEC patients to be poor, and through AJCC stages I–IV, the survival curves were obviously overlapped. Instead of TNM stages, SEER stages were incorporated into the multivariate COX analysis and showed none of T, N,

or M classifications were significant prognostic factors. The standardization and simplification of SEER staging ensures over-time consistency among various definitions that exert an impact on OS in our nomogram, and accords with previously released statistics (18). However, as there is a wider application of AJCC staging systems in clinical practice, whether TNM stages correlate with the survival of SCEC patients remains controversial. One study of 64 patients suggested that T stage was an independent prognostic factor (10), with other data suggesting the N stage was associated with survival (11,19). Limitations notably exist in all previous studies, including a small sample size in the former and the majority of SCEC patients being non-metastatic. The present research shares the limitation of other SEER-data-based studies in that its interpretation is restricted. As we also failed to investigate tumour invasion, distant metastases, as well as lymph node metastases of 27.2–52.2% of patients, there were 138 (32.3%) and 139 (32.6%) patients with unknown 7th and 6th TNM stage. However, both the AJCC and SEER categorizations of parameters only incorporate tumour-related variables, and the clinical characteristics of patients





**Figure 4** Validation of the predictive accuracy of the nomogram for overall survival in the FUSCC cohort. Calibration curves (A-C) and time-dependent ROC curves (D-F) at 1-, 3-, and 5-year. ROC, receiver operating characteristic; FUSCC, Fudan University Shanghai Cancer Center.

and data on previous therapies is not taken into account, resulting in an inaccurate estimation of OS as several studies in other cancer types have revealed (15,16,20,21). Future studies are required to establish a more precise and individual staging system specified for SCEC.

In the current study, besides the SEER stage, we showed that age, male gender, tumor location in the upper-third of the esophagus, and insufficient or no radiotherapy, chemotherapy, and/or surgery were closely linked to mortality in SCEC patients. Our data for age (9,13,18,22), male gender (9,18,22), and primary tumour site (9,22) as prognostic factors accords with previously published literature on SCEC. Zhang and colleagues (23) built a competing-risk model to assess the cause-specific survival (CSS) in patients with esophageal NEC which demonstrated that primary tumour site, distant metastases, and clinical treatment were closely linked to CSS, which supports our results.

Zhang's study also concluded surgery was a protective factor that lowered the possibility of cause-specific mortality in SCEC patients by 76%, followed by radiotherapy and chemotherapy, whereas our results indicated that the death risk of patients treated with chemotherapy decreased by 62% (HR =0.38, 95% CI, 0.30–0.49), followed by radiotherapy (HR =0.71, 95% CI, 0.57–0.89), and surgery (HR =0.75, 95% CI, 0.62–0.91). This difference may be because Zhang's nomogram enrolled 162 esophageal NEC patients from 1998 to 2004 making the data relatively old, and may have included patients with large cell carcinomas. Moreover, Zhang's nomogram did not include any staging system, which is a major limitation acknowledged by the authors in their paper and its editorial letter (24). In addition, they did not verify their nomogram externally. An optimal treatment strategy has not been provided in national guidelines for SCEC due to a lack of sufficient data. SCEC is widely acknowledged as a complicated disease, and metastatic disease has occurred in 40–60% of patients at the time of diagnosis (5). While chemotherapy was regarded as the foundation of multidisciplinary as well as local treatment for SCEC, the recurrence rate has remained high and the prognosis far from ideal (5,11,25), and while surgery continues to play a predominant role, its use is limited to early stage and/or resectable SCEC patients (6,19,26,27). In the current study, we enrolled 427 SCEC patients from the SEER database and 49.4% patients were distant SEER stage, which is a figure consistent with the literature and may indicate our results are close to the real world. On this basis, we propose chemotherapy-based

systemic therapy for the local treatment of SCEC.

However, there are several limitations to this study. Above all, the information relevant to tumour site, TNM and VALSG stage, lifestyle, performance status, pure/mixed SCEC, surgical margins, and other detailed clinical information could not be ascertained from SEER, which restricted the reliability as well as precision of our prognostic analyses. Secondly, our study developed a nomogram to estimate individualized OS, which did not take competing risks into consideration, and more investigation into cause-specific mortality in SCEC patients is warranted. Last but not least, the retrospective nature and limited sample size of our analysis inevitably produced substantial bias, and prospective and multicentre studies are required to validate these factors.

## Conclusions

Despite its limitations, as well as the rarity of SCEC, the nomogram in our research was developed and verified on the basis of a large population of SCEC patients and on the basis of combining non-anatomic factors with anatomic ones, incorporating both clinical characteristics and therapy methods. While the estimations of the nomogram were relatively accurate when predicting the prognosis of SCEC patients, more research based on larger-scale samples is required to validate its applicability and update our findings.

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## Footnote

*Reporting Checklist:* The authors have completed the TRIPOD reporting checklist. Available at <https://dx.doi.org/10.21037/atm-21-3900>

*Conflicts of Interest:* All authors have completed the ICMJE uniform disclosure form (available at <https://dx.doi.org/10.21037/atm-21-3900>). The authors have no conflicts of interest to declare.

*Ethical Statement:* The authors are accountable for all

aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. This retrospective study was approved by the institutional review board and independent ethics committee of FUSCC. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013).

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**Table S1** Detailed scores of all predictors in the nomogram

Variables	Nomogram points
Age	
20	0
25	7
30	14
35	21
40	29
45	36
50	43
55	50
60	57
65	64
70	71
75	78
80	86
85	93
90	100
Gender	
Female	0
Male	12
Location	
Low	0
Overlap	12
Middle	13
Upper	14
Stage	
Localized	0
Regional	28
Distant	86
Unstaged	35
Surgery	
Yes	0
No	5
Radiotherapy	
Yes	0
No	27
Chemotherapy	
Yes	0
No	76

**Table S2** AUC at 1-, 3-, and 5- years in the FUSCC cohort

	1 year		3 years		5 years	
	AUC	95% CI	AUC	95% CI	AUC	95% CI
TNM 6th	0.588	0.293-0.882	0.672	0.478-0.866	0.609	0.442-0.776
TNM 7th	0.806	0.661-0.983	0.653	0.474-0.831	0.655	0.496-0.815
Nomogram	0.919	0.899-0.939	0.886	0.798-0.973	0.831	0.698-0.963

AUC, area under the curves; FUSCC, Fudan University Shanghai Cancer Center; CI, confidence interval.