



Risk stratification model for patients with stage I invasive lung adenocarcinoma based on clinical and pathological predictors

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Background: The aim of this study was to propose a new kind of pathological classification and further establish a prognostic model for resected stage I invasive adenocarcinoma (IADC).

Methods: Clinicopathological data were collected from 2 hospitals. The new proposed pathological reclassification was defined according to certain subtype instead of a predominant one. Survival curves were plotted by Kaplan-Meier analysis. Cox regressions were analyzed for recurrence-free survival (RFS) and overall survival (OS), through which prognostic scores and stratification models were established. The comparison between risk models and the eighth edition of tumor, node, metastasis (TNM) classification was conducted through receiver operating characteristic curves (ROC), as identified by the area under the curve (AUC) and z test.

Results: In all, 1,196 patients were enrolled. At multivariable analysis, solid and micropapillary of the new pathological reclassification, along with stage IA3 and IB were independent predictors for poorer RFS. Stage IB and smoking status significantly indicated worse OS. After normalization and standardization of log-hazard ratio (HR), personalized scores were calculated and the risk stratifications with 3 risk groups were generated. Compared with TNM classification, the risk model of RFS showed advantage over early-recurrence prediction (1-year: 0.653 vs. 0.556, P=0.033; 3-year: 0.663 vs. 0.076, P=0.008). No marked difference was observed in long-term RFS or OS.

Conclusions: Considering the harboring of certain patterns may be a new concept in adenocarcinoma classification. The risk stratification model based on this pathological classification and the eighth TNM classification showed remarkable superiority over TNM alone in predicting early recurrence of stage I adenocarcinoma. However, TNM classification remained valuable for long-term recurrence and survival prediction.

Keywords: Pathological stage I lung invasive adenocarcinoma; new pathological classification; risk stratification model; the eighth edition TNM classification

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Introduction

Non-small-cell lung cancers (NSCLC) have become a severe public health issues across all countries of the world, and especially in China, which currently has an all-time high level of NSCLC-related morbidity and mortality (1,2). As one of the major subtypes of NSCLC, lung adenocarcinoma (ADC) accounts for approximately 70% of NSCLC cases (3). Although lobectomy has been recommended as the first choice for early-staged NSCLC, with sublobar resection as an appropriate substitution to well-selected patients (4-6), the 5-year survival rate still remains dissatisfactory (7).

With the widespread use of low-dose computed tomography (LDCT), there has been a significant boost in the detection of early-stage NSCLC, particularly small-sized ADC (8). The tumor, node, metastasis (TNM) classification system has been considered the gold standard for lung cancer survival outcome for decades; the eighth TNM edition of lung cancer indicates a 5-year survival of 90%, 85%, 80%, and 73% for stage IA1, IA2, IA3, and IB cases, respectively (9). However, despite this reliable prognostic survival, the application of the TNM system alone might not be able to indicate the risk of tumor recurrence after complete resection (7).

The International Association for the Study of Lung Cancer (IASLC)/American Thoracic Society (ATS)/European Respiratory Society (ERS) have recognized the predominant histologic subtype as the standard classification to resected ADC. The subtypes with predominant patterns include lepidic-predominant ADC (LEP), acinar-predominant ADC (ACN), papillary-predominant ADC (PAP), solid-predominant ADC (SOL), micropapillary-predominant ADC (MIP) and other variants (10,11). However, ADC displays the most heterogeneity with the most mixed histological patterns compared with other non-ADC lung cancers, different subtype of ADC with varying potential influences on recurrence and the postsurgical adjuvant therapy (12,13). For instance, the poor prognosis of ADC patients existed even when the SOL/MIP components were not in predominance and postoperative therapy should be carefully considered (14). Besides, invasive adenocarcinoma (IADC) patients harboring LEP

component were clearly correlated with relatively better prognosis compared with those without LEP component according to our previous studies (15,16). Hence, we hypothesized that the specific subtype of stage I ADC rather than the predominant type would truly be indicative of the recurrence and survival outcome.

In this research, we tried to establish a risk stratification model combining other independent prognostic factors with the eighth edition TNM classification and extend the content of the TNM classification system. Therefore, we defined a proposed pathological classification with respect to the specific subtype of ADC and further developed a clinicopathological risk stratification model for the personalized prediction of pathological stage I IADC based on the data pool from Shanghai Chest Hospital and Fudan University Shanghai Cancer Center.

We present the following article in accordance with the TRIPOD reporting checklist (available at <http://dx.doi.org/10.21037/tlcr-21-393>).

Methods

Patients

All procedures performed in this study involving human participants were in accordance with the Declaration of Helsinki (as revised in 2013). The study was approved by ethics board of Shanghai Chest Hospital (KS1992) and informed consent was not needed. A total of 3,368 cases with resected pulmonary nodules ≤ 3 cm admitted from January 2007 to December 2011 to the Shanghai Chest Hospital and Fudan University Shanghai Cancer Center were screened for eligibility for investigation. According to the inclusion and exclusion criteria (*Figure 1*), 1,196 cases with pathological stage I IADC were enrolled. Non-ADC lesions such as squamous carcinoma, large-cell and small-cell lung cancer; noninvasive ADC including atypical adenomatous hyperplasia (AAH), ADC *in situ* (AIS), and minimally invasive ADC (MIA) identified by final pathology; multiple lesions indicating multiple primary ADC confirmed by examination and pathology; patients who received neoadjuvant therapy or palliative treatment; the local extension indicating advanced staging such as

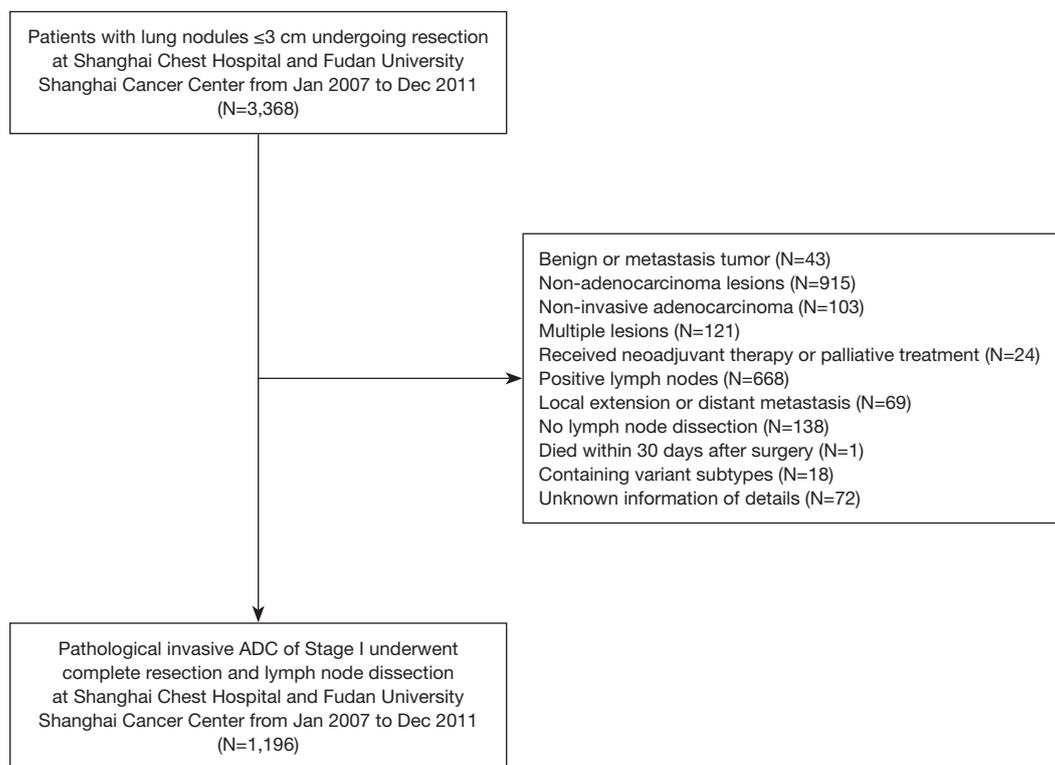


Figure 1 Study cohort flowchart. A total of 3,368 patients with pulmonary nodules ≤ 3 cm who underwent surgical resection at Shanghai Chest Hospital and Fudan University Shanghai Cancer Center were identified between January 2007 and December 2011. After exclusion, 1,196 patients were included in this study.

T3 and T4 with respect to the eighth TNM classification; patients who died within 30 days of surgery; cases with positive lymph nodes; and cases with no dissection of lymph nodes in the hilum or mediastinum. In addition, patients with variant subtypes or those with missing data were also excluded.

Outcomes variables and patient follow-up

The primary endpoint was recurrence-free survival (RFS), defined as survival from the date of surgical operation to the date of recurrence or last follow-up. The secondary endpoint was overall survival (OS), defined as the survival between surgery and death or last follow-up. All patients completed the preoperative routine examinations including enhanced chest computed tomography (CT) scans, head CT scans or magnetic resonance imaging (MRI), abdominal ultrasound, and bone scintigraphy. Positron emission tomography (PET) was performed when required. After the surgery, our routine follow-up time was every 3 months for

the first postoperative year; CT scans, neck and abdominal ultrasound, head MRI, and bone scintigraphy were conducted at postoperative month 6 and repeated every year thereafter. Patients would be contacted via telephone calls prior to their follow-up with appointments fixed in the outpatient department.

Definition and identification of the proposed new risk pathological classification of IADC

In this study, we proposed a new pathological classification system for ADC. To be specific, any subtype of IADC harboring MIP was classified under the new classification of MIP (nMIP). The subtype containing SOL not coexisting with MIP was classified as the new classification of SOL (nSOL). Histological subtypes containing PAP except for MIP/SOL were therefore classified as nPAP. The proposed new classification of nACN was defined as ADC containing ACN instead of MIP/SOL/PAP. The remaining pathological subtypes were divided into nLEP (containing

LEP without MIP/SOL/PAP/ACN) and others. For a more accurate and unbiased assessment, hematoxylin and eosin (HE)-stained slides of the pathological samples were all rechecked by 2 experienced pathologists from each hospital.

Statistical analysis

The association between the categorical variables was calculated by Pearson χ^2 or Fisher's exact test. Univariable and multivariable Cox regressions were conducted to investigate the potential independent prognostic factors of RFS and OS as estimated by hazard ratio (HR) and 95% confidence interval (95% CI). The results of the HR obtained above were registered as the status of log-HR, from which the personalized risk scores were generated (17,18). Then, normalization and standardization of these scores were divided by the smallest value of the obtained risk coefficients and the accumulation of personalized scores for patients were derived (18). Thus, the risk score stratification model (including low-, intermediate-, and high-risk classes) for patients' clinical outcomes could be presented and the cutoffs, which were decided upon by the trisection of risk values (low risk class with score ≥ 0 and < 1 , intermediate risk class with scores ≥ 1 and < 2 , and high risk class with scores ≥ 2). Kaplan-Meier survival curves were plotted for comparison between risk model and the eighth TNM classification, and identified by the log-rank test. The curves of receiver operating characteristics (ROC) were performed to analyze the sensitivity and specificity between our risk model and the eighth TNM classification through the estimation of the area under the curve (AUC), the difference of which was verified by z test.

Results

The baseline characteristics of pathological stage I IADC

The clinicopathological characteristics of 1,196 patients are listed in *Table 1*. The predominant patterns were classified according to the 2015 World Health Organization classification, and included lepidic-predominant ADC (LEP), acinar-predominant ADC (ACN), papillary-predominant ADC (PAP), solid-predominant ADC (SOL), micropapillary-predominant ADC (MIP), and other variants (7). The proposed pathological classification was classified according to our previous definition. The relationship between the predominant patterns and our proposed classification can

been seen in *Table S1*.

The primary cohort mainly consisted of females (61%), and more than half (54%) of our candidates were aged ≤ 60 years. With respect to smoking status, the majority (almost 64%) of patients were never-smokers, while the smoking patients (including cessation) accounted for less than 36% of the total. Therefore, the never-smokers constituted the majority of the candidates. As for the location of ADC, the most common site of all the resected stage I IADC patients was still the right upper lobe (RUL), accounting for 32.6% of cases, followed by the left upper lobe at 25%, the right lower lobe (RLL) at 18.9%, the left lower lobe (LLL) at 14.1%, and the right middle lobe (RML) at 9.4%. In this study, patients without adjuvant chemotherapy were still in the majority, accounting for 76.5% of patients; meanwhile, patients with adjuvant chemotherapy (including standard adjuvant chemotherapy and incomplete one) constituted only 23.5% of patients. As for the tumor size of the resected stage I ADC cases, the majority of our primary cohort consisted of tumor sizes of 1 to 2 cm and 2 to 3 cm, at 47.4% and 43.2%, respectively. Meanwhile, sizes below 1 cm accounted for fewer than 9.4% of cases, probably owing to the lower aggressiveness of smaller-sized tumors. Lobectomy is still recommended as the gold stand approach worldwide. Hence, nearly 96.7% of patients underwent lobectomy in our primary cohort, while only 3.3% underwent anatomical sublobar resections. And all the patients received systemic lymph dissection or sampling. For pathological subtypes, ACN and PAP were the two most common subtypes among stage I ADC patients, accounting for 46.7% and 36% of cases, respectively. These subtypes were followed by SOL, MIP, and IMA, all of which had lower than 5% predominance. Specifically, MIP accounted for less than 0.5% of all subtype, which indicated a potential relationship between the subtype and high invasiveness and metastatic ability. In reference to our proposed new pathological classification, nPAP accounted for the majority of our primary cohort (54.2%), followed by the classification of nACN (23.1%), nSOL (11.4%), and nLEP (5.3%). nMIP and others accounted for only 3%, respectively. We selected the eighth TNM classification as our standard classification in this study and classified our resected stage I IADC patients into stage IA1, IA2, IA3, and IB accordingly. The stage IA2 IADC patients accounted for the majority (35.4%) of patients, followed by stage IB (29.9%), stage IA3 (26%), and stage IA1

Table 1 Baseline characteristics of the primary cohort with stage I invasive adenocarcinoma who underwent complete resection and lymph node dissection

Variables	No. of patients	(%)
Sex		
Male	468	39.1
Female	728	60.9
Age		
≤60 years	645	53.9
>60 years	551	46.1
Smoking history		
No	764	63.9
Yes	432	36.1
Location		
RUL	390	32.6
RML	113	9.4
RLL	226	18.9
LUL	299	25
LLL	168	14.1
Adjuvant chemotherapy		
No	915	76.5
Yes	281	23.5
Size		
≤1 cm	113	9.4
≤2 cm	566	47.4
≤3 cm	517	43.2
Surgery		
Sublob	40	3.3
Lob	1,156	96.7
Pathology		
LEP	103	8.6
ACN	558	46.7
PAP	430	36
SOL	52	4.3
MIP	6	0.5
Others	47	3.9

Table 1 (continued)

Table 1 (continued)

Variables	No. of patients	(%)
Proposed new pathology		
nLEP	63	5.3
nACN	276	23.1
nPAP	648	54.2
nSOL	137	11.4
nMIP	36	3.0
Others	36	3.0
8 th TNM classification		
IA1	104	8.7
IA2	423	35.4
IA3	311	26
IB	358	29.9
LN		
≤10	803	67.1
>10	393	32.9
PI		
No	838	70.1
Yes	358	29.9
LVI		
No	1,128	94.3
Yes	68	5.7

RUL, right upper lobe; RML, right middle lobe; RLL, right lower lobe; LUL, left upper lobe; LLL, left lower lobe; sublob, sublobectomy; lob, lobectomy; LEP, lepidic-predominant ADC; ACN, acinar-predominant ADC; PAP, papillary-predominant ADC; SOL, solid-predominant ADC; MIP, micropapillary-predominant ADC; nLEP, new classification of LEP; nACN, new classification of ACN; nPAP, new classification of PAP; nSOL, new classification of SOL; nMIP, new classification of MIP; TNM, tumor, node, metastasis; LN, harvested lymph nodes; PI, pleural invasion; LVI, lymphovascular invasion.

(8.7%). For lymph nodes, we selected 10 as the cutoff value for the harvested lymph nodes, and patients with fewer than 10 lymph nodes still constituted the majority (67.1%), with only 32.9% of patients having more than 10 harvested lymph nodes. Moreover, 29.9% patients had pleural invasion (PI) in stage IB, indicating that stage IB

Table 2 Univariable Cox proportional hazards regressions for patients with stage I invasive adenocarcinoma who underwent complete resection and lymph node dissection

Variables	RFS			OS		
	HR	95% CI	P	HR	95% CI	P
Sex	0.674	0.497–0.914	0.011	0.545	0.352–0.843	0.006
Age	1.214	0.895–1.645	0.213	1.166	0.754–1.803	0.491
Smoking history	1.375	1.012–1.869	0.042	2.039	1.316–3.157	0.001
Location	0.947	0.854–1.051	0.306	1.106	0.875–1.180	0.833
Adjuvant chemotherapy	1.888	1.370–2.601	<0.001	1.744	1.098–2.769	0.018
Size	1.561	1.210–2.014	0.001	1.551	1.079–2.230	0.018
Surgery	0.883	0.391–1.995	0.765	0.827	0.261–2.624	0.748
Pathology	1.058	0.913–1.227	0.455	1.145	0.928–1.411	0.206
Proposed new pathology	1.393	1.211–1.603	<0.001	1.298	1.043–1.614	0.019
8 th TNM Classification	1.576	1.333–1.862	<0.001	1.483	1.172–1.876	0.001
LN	2.337	0.851–2.998	0.064	1.180	0.739–1.885	0.488
PI	2.024	1.490–2.748	<0.001	1.739	1.118–2.703	0.014
LVI	1.976	1.180–3.309	0.010	1.961	0.945–4.071	0.071

RFS, recurrence-free survival; OS, overall survival; HR, hazard ratio; CI, confidence interval; TNM, tumor, node, metastasis; LN, harvested lymph nodes; PI, pleural invasion; LVI, lymphovascular invasion.

or PI could be included in later multivariable analysis. Patients with lymphovascular invasion (LVI) accounted for only 5.7% of cases, which was likely related to the relatively restrained invasiveness of early-staged IADC in lymphatic vessels and surrounding blood vessels.

Recurrence and survival analysis for pathological stage I IADC

The median follow-up time was 51.51 months (range, 1–77 months), during which there were 166 (13.9%) patients who relapsed (including local recurrence and distant metastasis) and 81 (6.8%) patients who died (63 due to cancer-specific death and 18 deaths due to other causes). The 5-year RFS and OS were 86.3% and 94.1%, respectively (Figure S1).

Univariable and multivariable cox regressions for pathological stage I IADC

In the analysis of univariable cox regression, we enrolled all the related baseline clinicopathological data of resected stage I IADC patients into the Cox model and analyzed

the potential risk factors that impacted the recurrence and survival outcome of pathological stage I IADC patients (Table 2). With respect to RFS, females demonstrated much better RFS compared with males (HR 0.674, 95% CI, 0.497–0.914, $P=0.011$), which indicated female sex as the potential protective factor of recurrence to pathological stage I IADC. As for smoking history, patients with a smoking history demonstrated worse recurrence as compared to never-smokers (HR 1.375, 95% CI, 1.012–1.869, $P=0.042$), and thus smoking history was indicated to be potential risk factor for postoperative recurrence. Patients who received adjuvant chemotherapy also demonstrated a worse recurrence rate compared to patients who did not receive adjuvant chemotherapy (HR 1.888, 95% CI, 1.370–2.601, $P<0.001$), a finding which requires further investigation. Tumor size as well as the eighth TNM classification was also associated with RFS rate (tumor size: HR 1.561, 95% CI, 1.210–2.014, $P=0.001$; TNM: HR 1.576, 95% CI, 1.333–1.862, $P<0.001$). Furthermore, PI and LVI also showed themselves as potential recurrence factors for resected stage I IADC (PI: HR 2.024, 95% CI, 1.490–2.748, $P<0.001$; LVI: HR 1.976, 95% CI, 1.180–3.309, $P=0.010$). In contrast to the traditional

predominant subtypes, our proposed new classification of pathological stage I IADC was significantly associated with the recurrence outcome (HR 1.393, 95% CI, 1.211–1.603, $P < 0.001$), while predominant subtypes were not ($P = 0.455$). Furthermore, patient age ($P = 0.213$), tumor location ($P = 0.306$), surgical approach ($P = 0.765$), and harvested lymph nodes ($P = 0.064$) also showed no statistical difference, indicating that they have no impact on the recurrence for pathological stage I IADC.

As for the clinical survival outcomes, female sex was still a potential protective factor for pathological stage I IADC (HR 0.545, 95% CI, 0.352–0.843, $P = 0.006$), while smoking history was a potential risk factor for OS (HR 2.039, 95% CI, 1.316–3.157, $P = 0.001$). However, those patients receiving postoperative chemotherapy still demonstrated potentially worse OS (HR 2.039, 95% CI, 1.316–3.157, $P = 0.001$). Tumor size as well as the eighth TNM classification was also potential risk factors for OS (tumor size: HR 1.551, 95% CI, 1.079–2.230, $P = 0.018$; TNM classification: HR 1.483, 95% CI, 1.172–1.876, $P = 0.001$). Furthermore, PI was also revealed to be a potential risk factor for OS (HR 1.739, 95% CI, 1.118–2.703, $P = 0.014$). In the univariable analysis of OS, our proposed new classification of pathological stage I IADC strongly indicated the potential ability for OS prognosis, with the similar results to RFS (HR 1.298, 95% CI, 1.043–1.614, $P = 0.019$). Meanwhile, age ($P = 0.491$), tumor location ($P = 0.833$), surgical approach ($P = 0.748$), traditional predominant subtypes ($P = 0.206$), harvested lymph node ($P = 0.488$), and LVI ($P = 0.071$) showed no statistical difference, indicating these have no prognostic ability in OS for resected stage I IADC.

After univariable analysis, we enrolled all the potential risk factors related to RFS and OS into the multivariable cox regression models, respectively (Table 3). Two independent risk factors were identified as being associated with cancer recurrence in our proposed pathological classification and TNM classification: nSOL (HR 5.428, 95% CI, 1.277–23.074, $P = 0.022$), nMIP (HR 10.784, 95% CI, 2.415–48.154, $P = 0.002$), stage IA3 (HR 3.231, 95% CI, 1.144–9.127, $P = 0.027$), and stage IB (HR 4.202, 95% CI, 1.504–11.736, $P = 0.006$). Interestingly, the predominant patterns were not associated with postsurgical relapse.

With respect to OS, our proposed pathological classification, similar to the conventional predominant patterns, did not show any statistical influence, while stage IB and smoking history strongly corresponded with worse

survival outcome (Tables 2,3).

The RFS-related risk stratification model for pathological stage I IADC

Accordingly, the results of RFS multivariable analysis were applied and log-HR was calculated and standardized, from which the personalized scores were derived in turn (Table 4). The highest scores were 2 points, and the lowest one was 0. Among all the independent recurrent risk factors of resected stage I IADC, nMIP received the most points (2 points), followed by nSOL (1.4 points), stage IB (1.2 points), and stage IA3 (1 point), with nLEP, nACN, nPAP, etc., as well as stage IA1 and IA2, all receiving 0 points.

The risk stratification model based on our proposed new pathological classification and the eighth TNM classification was established according to the statistical analysis above (Figure 2A), and the corresponding 5-year RFS of low-risk group, intermediate-risk group and high-risk group in the risk stratification model was 93.0%, 81.6% and 64.4%, respectively. Furthermore, the survival curve of the eighth TNM classification was also plotted (Figure 2B), indicating a 5-year RFS of stage IA1, stage IA2, stage IA3 and stage IB was 96.0%, 89.4%, 83.1% and 76.1%, respectively.

In order to compare the prognostic capability between those 2 above approaches, ROCs were plotted by means of AUC with the SPSS software. After these 2 approaches were compared in terms of AUCs of different postoperative times (1-, 3-, 5-year, and total postoperative time), we found that there was a difference in recurrence prediction between the risk stratification model and the eight TNM classification across the different postoperative times.

For 1-year recurrence prediction of resected stage I IADC, the risk stratification model demonstrated superiority, as it could much more precisely predict the prognosis of 1-year recurrence after surgery (1-year: model *vs.* TNM =0.653 *vs.* 0.556; $P = 0.033$).

For 3-year prediction, our risk stratification model still showed better performance compared to the traditional TNM classification (3-year: model *vs.* TNM =0.663 *vs.* 0.607; $P = 0.008$). All the ROC curves and AUCs for these performances are shown in Figure 3A,B, respectively.

For 5-year and total RFS prediction, however, no remarkable difference in prognostic prediction was found, which indicated that our risk stratification model had no prognostic advantage for long-term recurrence compared

Table 3 Multivariable Cox proportional hazards regressions for patients with stage I invasive adenocarcinoma who underwent complete resection and lymph node dissection

Variables	RFS			OS		
	HR	95% CI	P	HR	95% CI	P
Sex						
Male	(Ref =1)			(Ref =1)		
Female	-	-	0.067	-	-	0.349
Smoking history						
No	(Ref =1)			(Ref =1)		
Yes	-	-	0.187	1.990	1.284–3.083	0.002
Adjuvant chemotherapy						
No	(Ref =1)			(Ref =1)		
Yes	-	-	0.059	-	-	0.294
Proposed new pathology						
nLEP	(Ref =1)		<0.001	(Ref =1)		
nACN	2.948	0.700–12.422	0.141	-	-	0.548
nPAP	2.376	0.573–9.855	0.233	-	-	0.780
nSOL	5.428	1.277–23.074	0.022	-	-	0.085
nMIP	10.784	2.415–48.154	0.002	-	-	0.176
Others	2.856	0.517–15.784	0.229	-	-	0.394
8 th TNM Classification						
IA1	(Ref=1)		<0.001	(Ref=1)		
IA2	2.081	0.736–5.885	0.167	2.609	0.611–11.136	0.195
IA3	3.231	1.144–9.127	0.027	4.125	0.974–17.475	0.054
IB	4.202	1.504–11.736	0.006	5.176	1.243–21.557	0.024
LVI						
No	(Ref=1)			-	-	-
Yes	-	-	0.168	-	-	-

RFS, recurrence-free survival; OS, overall survival; Ref, reference; HR, hazard ratio; CI, confidence interval; nLEP, new classification of LEP; nACN, new classification of ACN; nPAP, new classification of PAP; nSOL, new classification of SOL; nMIP, new classification of MIP; TNM, tumor, node, metastasis; PI, pleural invasion; LVI, lymphovascular invasion.

Table 4 Standardized risk score coefficients based on recurrence-free survival

Recurrence-free Survival	Scores				
	0	1	1.2	1.4	2
Proposed new pathology	nLEP/nACN/nPAP/Others			nSOL	nMIP
8 th TNM classification	IA1 and IA2		IA3	IB	

nLEP, new classification of LEP; nACN, new classification of ACN; nPAP, new classification of PAP; nSOL, new classification of SOL; nMIP, new classification of MIP; TNM, tumor, node, metastasis.

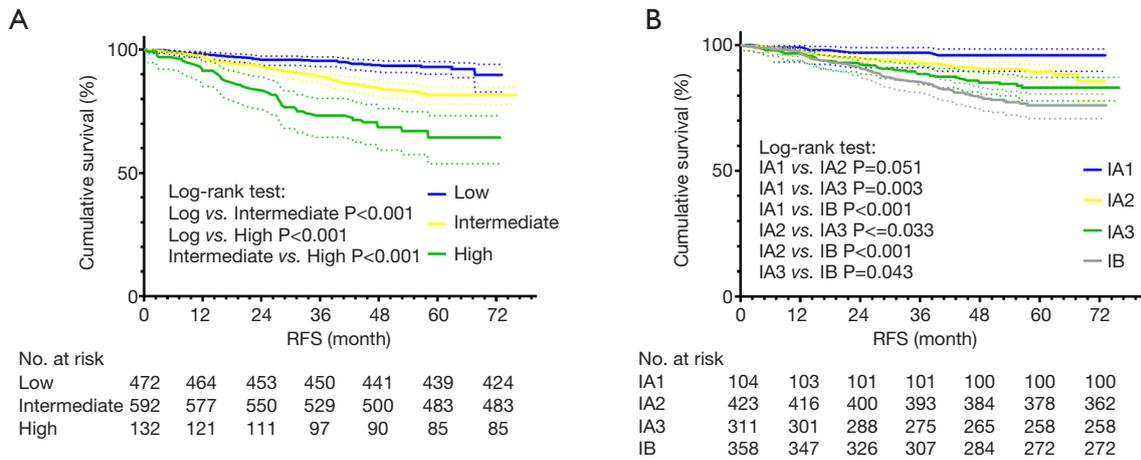


Figure 2 Recurrence-free survival according to the risk stratification model (A) and the eighth TNM classification (B).

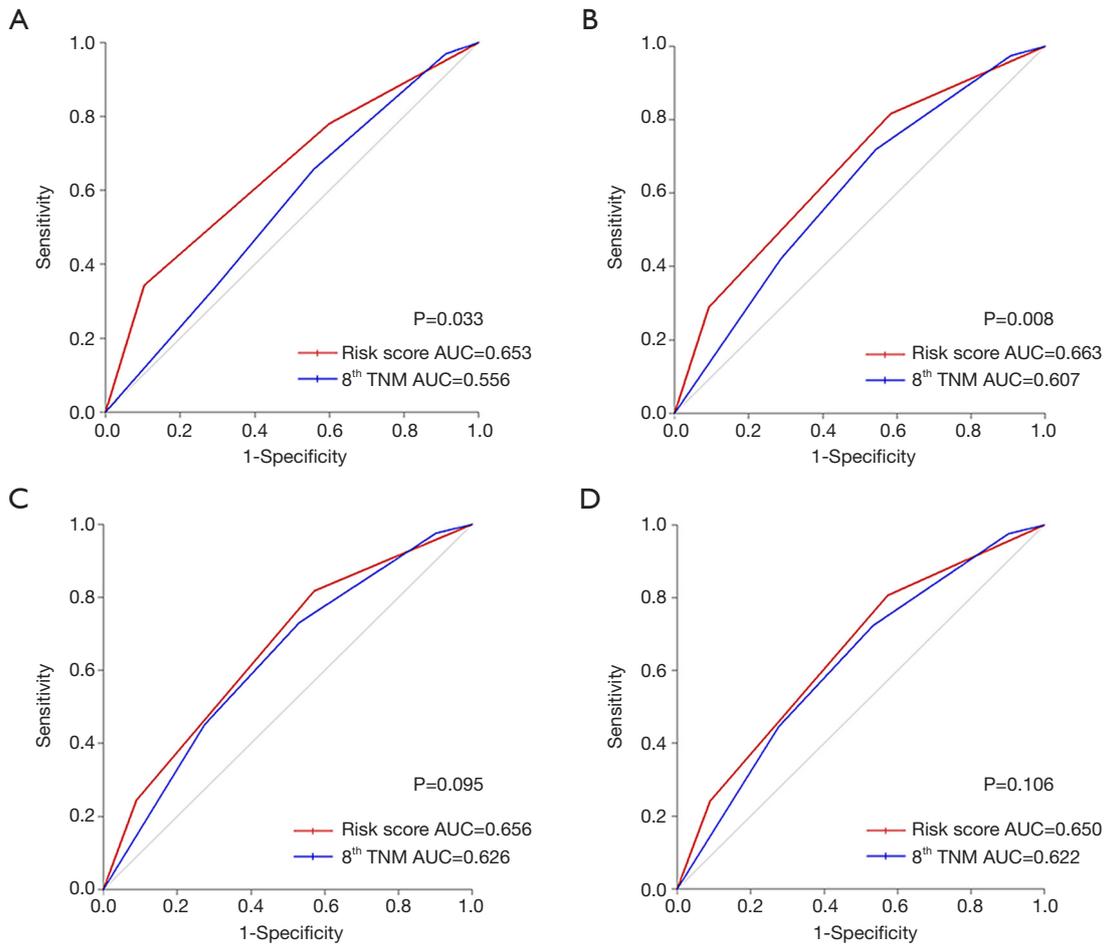


Figure 3 Prognostic performance comparison between risk stratification model and the eighth TNM classification for recurrence-free survival. AUC comparison between the risk model and TNM classification in 1-year RFS (A), 3-year RFS (B), 5-year RFS (C), and total RFS (D).

with TNM classification (5-year: model *vs.* TNM =0.656 *vs.* 0.626, P=0.095; total: model *vs.* TNM =0.650 *vs.* 0.622, P=0.106). The ROC curves and the related AUCs for these performances were shown in *Figure 3C,D*, respectively.

The OS-related risk stratification model for pathological stage I IADC

As for the risk model of OS, individualized scores were also established in accordance with previous methods (*Table S2*). Among all the independent risk scores for OS, the highest score was 2.4 points, and the lowest one score was 0 points. Among all the independent risk factors, stage IB had 2.4 points, followed by smoking history with 1 point. Meanwhile, never-smokers, stage IA1, stage IA2, and stage IA3, all received 0 points.

We thus established the risk stratification model for OS prediction using different risk scores of different patients; the cutoffs for the low-risk, intermediate-risk, and high-risk group are mentioned in the Methods. Furthermore, the stratified survival curves of the risk model and TNM system are presented in *Figure S2*.

To compare the sensitivity and specificity of OS prediction between the risk stratification model and the eighth TNM classification, we created ROC curves and AUCs using SPSS software. However, in contrast to the results mentioned above, the risk model did not demonstrate significant predictive ability in short-term survival prognosis compared with TNM system, meaning our risk stratification model was not superior to the traditional TNM classification for short-term postoperative survival (1-year: model *vs.* TNM =0.396 *vs.* 0.431, P=0.822; 3-year: model *vs.* TNM =0.608 *vs.* 0.592, P=0.648). Additionally, the risk stratification model and TNM classification system showed no difference in prognostic ability for the long-term survival outcome of resected stage I IADC (5-year: model *vs.* TNM =0.599 *vs.* 0.594, P=0.836; total: model *vs.* TNM =0.610 *vs.* 0.602, P=0.740). All the related ROC curves and AUCs are plotted in the *Figure S3A,B,C,D*.

Discussion

With the discovery of the activation of epidermal growth factor receptor (EGFR) mutations in 2004, it was expected that the application of lung cancer gene-specific diagnosis and therapy would bring about a revolution in the clinical management of lung cancer (19). In addition to *EGFR*, several genes including *ALK* and *ROS1*, along with their

related targeted therapy were also confirmed as representing promising treatment options for advanced and stage IV NSCLC (20,21). Eventually, there were more protocols and clinical practice management guidelines created for the suggested targeted treatments (22,23). As genes for as routine practice among resected early-staged ADC have not been tested and their prognostic significance is still not confirmed, it is still of great importance for clinicians to evaluate and predict precise recurrence and survival outcome for early-staged ADC patients using the clinical and pathological characteristics.

According to the novel ADC pathological classification published by the IASLC/ATS/ERS in 2011, any histological pattern should be recorded with an assessment quantified by 5% increments with a single predominant pattern being recommended to represent the certain tumor (11). While approximately 8–90% of IADCs have been recognized to be the mixed patterns, focusing on a specific predominant pattern while ignoring the coexisting ones might lead to incorrect prognostic assessment (12). To date, large numbers of previous studies have proven that SOL/MIP are associated with worse clinical outcome (24,25); MIP is particularly correlated with a higher degree of aggressiveness, LVI, and lymph node metastasis (26,27). Interestingly, MIP still show a consistently prognostic impact, even when the presence is $\geq 5\%$ or $\geq 1\%$ (12,14,16,28,29). Thus, it was reasonable to speculate that an MIP component could become a representative classification of certain ADC types even when not predominant. Compared with MIP, SOL displayed limited prognostic impact when not coexistent with MIP (30,31). Therefore, the reclassification of nSOL was proposed. Previous studies have uniformly classified ACN/PAP into the same group for the relatively similar prognostic outcome (12). Nevertheless, we tried to separate these 2 patterns in our proposed classification. Interestingly, in contrast to the eighth TNM classification system, our proposed system demonstrated marked superiority in early-stage ADC recurrence prediction. From this we could recognize that harboring MIP could definitively determine the prevalence of disease relapse regardless of whether a predominant pattern was present. This “all or none” effect was also present in nSOL, indicating that the higher risk of recurrence was mostly an influence of the SOL component. Therefore, the classification of IADC could be reclassified according to certain type of pathology instead of predominant one. Conversely, nPAP/nACN showed no statistical difference in RFS or OS, probably

due to the prevalence of LEP shows a tendency of better prognosis (14), thus prompting for further investigation of the latter's prognostic effect.

Pignon *et al.* and Chansky *et al.* have pointed out that almost 50% of early-staged NSCLC suffer from cancer relapse (32,33). Although, the current TNM system has, for decades, maintained its significance in static and universal applicability, its prognostic ability has been constantly scrutinized and modified (9,34). We noticed that this conventional staging system was becoming less useful, especially among early-staged ADC when the T category became the only variable for prognosis analysis. In this research, our risk stratification model for RFS was constructed with the combination of two well-known and easy-to-extract clinicopathological system, our proposed pathological classification and the eighth TNM system, all of which were verified and accepted through multivariable Cox regressions. After being stratified into 3 risk levels by personalized scores, the risk model showed higher prognostic sensitivity and specificity compared with the traditional TNM classification in early recurrence prediction. Furthermore, the high-risk group could also be a better indicator for adjuvant therapy, owing to the MIP and SOL were contained in this group which was considered as a high risk for cancer recurrence (12,30). A clear statistical difference was evident between our risk model and TNM classification for 5-year and long-term recurrence prediction, implying an indispensable role of the traditional TNM classification in long-term prognosis of ADC recurrence. As for OS, our proposed pathological classification did not perform significantly better, strongly suggesting relatively low importance of histological subtypes on clinical survival outcome, which was consistent with a previous study (12). With consideration to these findings, the TNM classification can still be said to be unparalleled in its ability in predicting survival prognosis. Furthermore, smoking history also indicated poor OS, as previously suggested (35). Unfortunately, the risk model based on these 2 independent factors did not demonstrate significant superiority over conventional TNM classification in predicting OS.

In addition to this risk stratification model for early recurrence prediction, several risk factors, including tumor budding, tumor spread through air spaces (STAS), and even liquid biopsy, have been recently verified as reliable independent predictors of postoperative recurrence for resected ADC patients (36,37). It is therefore necessary to reconsider and recombine these potential risk factors into

our risk stratification model for future work.

There were several limitations in this research. First, this study retrospectively collected patients from 2 hospitals. The nature of this retrospective analysis inevitably introduced biases, in that our sample population likely lacks universality and representativeness. Second, although HE slides were all double-checked by 2 experienced pathologists in each hospital, the subjective nature of this examination might have led to discrepancies between pathologists. Third, we determined that any pathological pattern should be recorded when it showed $\geq 5\%$ predominance. For example, nMIP was noted when MIP was $\geq 5\%$. On the other hand, we did not consider the component when it was $< 5\%$. The consequences of using this cutoff are unknown and require further investigation.

Conclusions

Harboring a certain pattern of a histological subtype, such as MIP/SOL, instead of merely considering the predominant subtype, could be recommended as a new kind of representative classification, and this classification was clearly correlated with worse RFS. Furthermore, the risk stratification model based on this proposed pathological classification showed clear superiority for predicting early recurrence of resected stage I ADC when compared to the eighth TNM classification, suggesting that the influence of certain pathological subtypes should not be neglected when estimating potential recurrence risk.

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Footnote

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[org/10.21037/tlcr-21-393](https://doi.org/10.21037/tlcr-21-393)). 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. All procedures performed in this study involving human participants were in accordance with the Declaration of Helsinki (as revised in 2013). The study was approved by ethics board of Shanghai Chest Hospital (KS1992) and informed consent was not needed.

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Supplementary

Table S1 The relationship between the predominant patterns of adenocarcinoma and the proposed new pathological classification of IADC.

New pathology	LEP	ACN	PAP	SOL	MIP	Others	Total
nLEP	61	0	0	0	0	2	63
nACN	25	247	0	0	0	4	276
nPAP	15	233	396	0	0	4	648
nSOL	2	66	17	52	0	0	137
nMIP	0	12	17	0	6	1	36
Others	0	0	0	0	0	36	36
Total	103	558	430	52	6	47	1196

Abbreviations: LEP, lepidic-predominant ADC; ACN, acinar-predominant ADC; PAP, papillary-predominant ADC; SOL, solid-predominant ADC; MIP, micropapillary-predominant ADC; nLEP, new classification of LEP; nACN, new classification of ACN; nPAP, new classification of PAP; nSOL, new classification of SOL; nMIP, new classification of MIP.

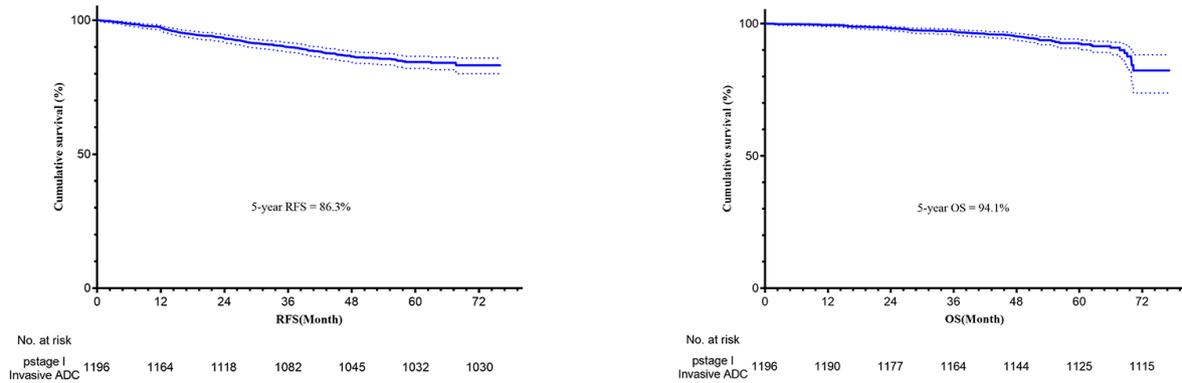


Figure S1 Recurrence-free survival and overall survival in the primary cohort. Kaplan-Meier curve for relapse-free survival and overall survival in 1,196 patients with resected stage I IADC.

Table S2 Standardized risk score coefficients based on overall survival.

Overall survival	Scores		
	0	1	2.4
Smoking history	No	Yes	
8 th TNM classification	IA1/IA2/IA3		IB

Abbreviations: TNM, tumor, node, metastasis.

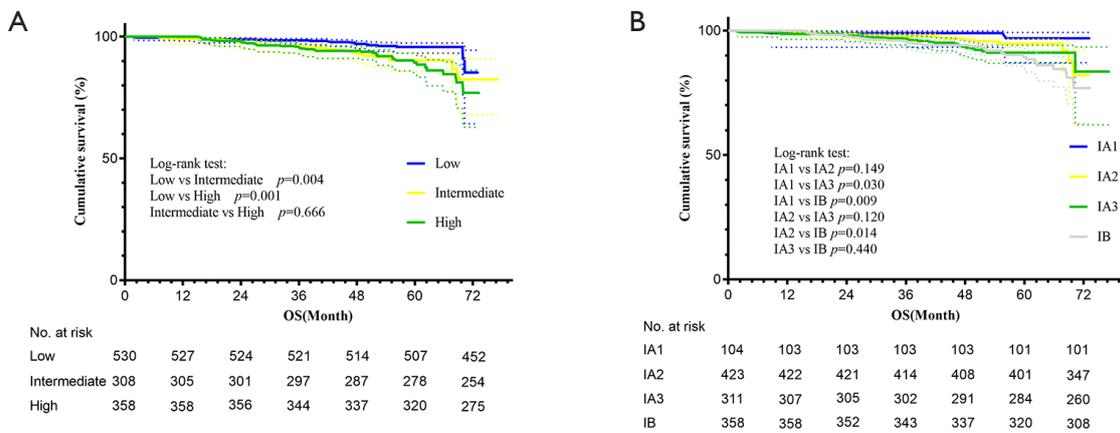


Figure S2 Overall survival according to the risk stratification model (A) and the eighth TNM classification (B). Kaplan-Meier curves for overall survival in 1,196 patients with resected stage I ADC, plotted by risk stratification model and the eighth TNM Classification, respectively.

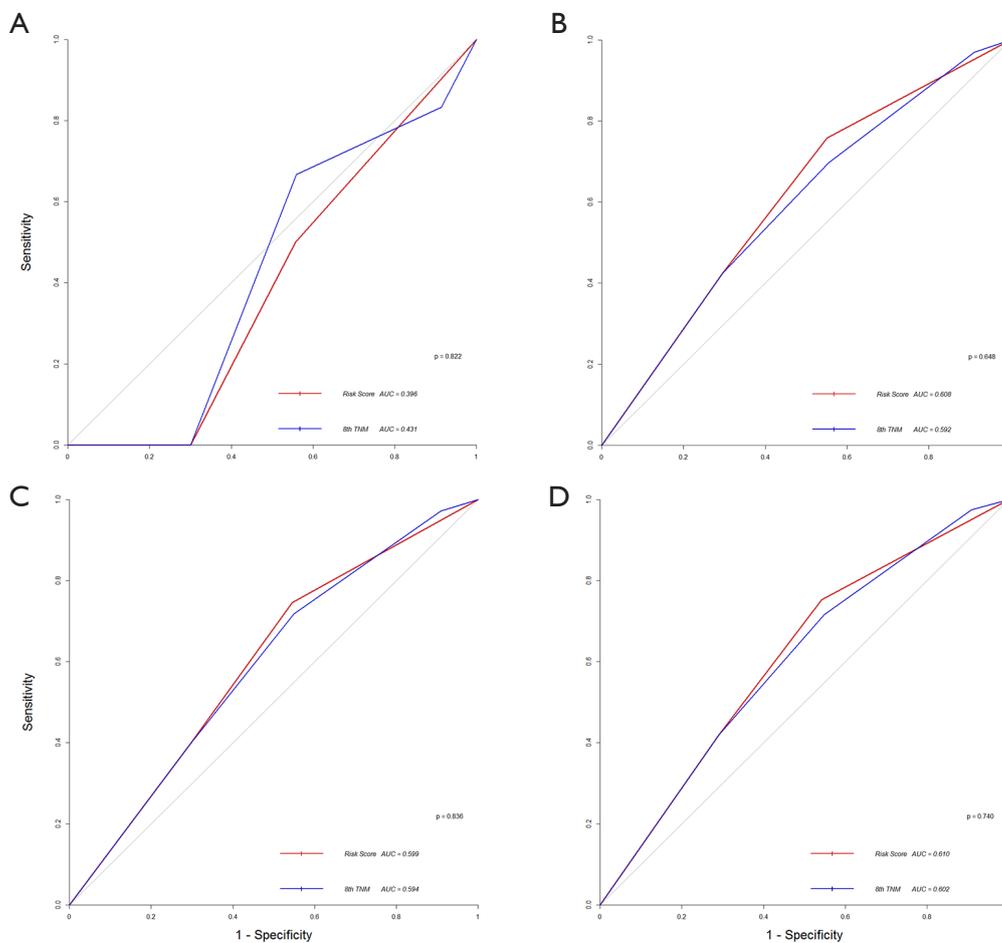


Figure S3 Prognostic performance comparison between risk stratification model and the eighth TNM classification for overall survival. AUC comparison between the risk model and TNM classification in 1-year OS (A), 3-year OS (B), 5-year OS (C), and total OS (D).