ORIGINAL RESEARCH

Platelet Count is Associated with the Rate of Lymph Node Metastasis in Lung Adenocarcinoma

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Purpose: Emerging studies have revealed that platelets are involved in tumor metastasis in lung adenocarcinoma (ADC). The solid pathological subtype of lung ADC is associated with metastasis, recurrence, and poor prognosis. However, there is no study exploring the relationship between platelets and different lung pathological subtypes.

Patients and Methods: The association between platelet counts and lymph node metastasis was analyzed in 852 patients with lung ADC who underwent surgery and lymph node dissection. Multivariate logistic analysis was conducted to identify the risk factors of lymph node metastasis. Then, lymph node metastasis and other factors were analyzed to determine their correlation with platelet count and histological subtype.

Results: We found that the platelet count was associated with lymph node metastasis (P = 0.01) in multivariable analysis, independent of tumor size, predominant subtype, visceral pleural invasion, and microvessel invasion. In patients with a platelet count ≥300 × 109/L, the rate of lymph node metastasis was 38.5%, almost twice as high as that in patients with a platelet count <300 × 109/L (23.2%). Additionally, elevated platelet counts, even those within the normal range, were significantly associated with a higher rate of lymph node metastasis. The mean platelet count in patients with solid-predominant histology (269.70 ± 69.38 × 109/L) was significantly higher than that in patients with other histologies (P < 0.001).

Conclusion: Elevated platelet counts are significantly associated with a higher rate of lymph node metastasis, even if the platelet counts are within the reference range. Platelet counts were significantly higher in patients with solid-predominant histology than in patients with other histologies. In addition, VEGF-C may play an important role in lymphatic metastasis in patients with lung ADC. We hypothesize that antiplatelet therapy may reduce lymph node metastasis in lung ADC patients.

Keywords: lung adenocarcinoma, platelets, lymph node, solid-predominant, pathological subtype

Introduction

Lung cancer is the most commonly diagnosed cancer (11.6% of all cancer diagnoses) and the leading cause of cancer-related death (18.4% of all cancer-related deaths) in both men and women.[1] Lung adenocarcinoma (ADC) is an independent pathologic subtype of lung cancer that has been widely studied. In 2011, the International Association for the Study of Lung Cancer, American Thoracic Society, and European Respiratory Society (IASLC/ATS/ERS) proposed a new histologic classification of lung ADC.[2] Invasive lung ADC was divided into six major pathological subtypes: lepidic, acinar, papillary, micropapillary, solid, and invasive mucinous adenocarcinoma. Several studies have reported that

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the micropapillary and solid subtypes of lung ADC are associated with metastasis, recurrence, and poor prognosis.[3–7] However, the reason for this phenomenon is unclear.

Emerging research suggests that platelets are functional players in many steps of the metastatic process.[8–11] It is clear that cancer cells can induce abnormalities in platelet number and function. In turn, platelets can promote tumor metastasis.[12,13] Tumor-educated platelets (TEPs) may be an ideal ally of tumor progression.[14] Conversely, platelets can be continuously activated and augment cancer-associated thrombosis.[15] Elevated platelet counts have been observed in patients with advanced cancer affecting the breast, colon, gastric, lung, ovary, and esophagus.[16–22] Further, elevated platelet counts are sometimes related to poor prognosis.[16–22] Although there is still some controversy in breast cancer, most preclinical studies indicate an association between elevated platelets and metastasis,[23] as well as with lymph node metastasis, and poor prognosis in lung cancer.[24,25]

Lung ADC metastasis is mainly classified into hematogenous and lymphatic metastases. Previous studies have mainly reported that preoperative elevated platelet counts have a role in promoting hematogenous metastasis and predicting a poor prognosis in non-small cell lung cancer patients,[23,24] the relationships between platelets, lymph node metastasis, and different ADC subtypes have not been widely studied. We conducted this study to explore the relationship of platelets, lymph node metastasis, and different ADC subtypes.

Patients and Methods
Ethical Statement and Patients
This retrospective study has been approved bythe institutional review board of the Qilu Hospital of Shandong University (KYLL-2016-097). All patients provided informed consent for the use of their clinical information. One thousand four hundred and seventy-two patients underwent surgical resection for pathologically diagnosed invasive lung ADC between January 2013 and December 2016 in the Qilu Hospital. All cases were collected through the medical record information database of the Qilu Hospital of Shandong University. The inclusion criteria were as follows: (1) patients underwent surgery for pathologically confirmed lung invasive ADC, and the pathological subtypes were recorded according to the 2011 IASLC/ATS/ERS standard;[2] (2) patients underwent

computed tomography, magnetic resonance imaging, bone scans, and positron emission tomography in accordance with the recommendations of the Chinese Society of Clinical Oncology guidelines within 14 days before surgery; (3) no distant metastasis or other malignancies; (4) no neoadjuvant chemotherapy or radiotherapy; (5) no hematological diseases before surgery; and (6) no antiplatelet drugs such as aspirin. After screening, a total of 852 patients were eligible for inclusion.

Blood Samples
All patients underwent routine blood testing in our hospital before surgery. Blood was collected in a blood collection tube containing EDTA for anticoagulation between 6:00 and 8:00 in the morning when patients were in the fasting state, and blood was sent for examination within 2 hours.

Pathological Examination
After surgery, all excised specimens were fixed in formalin and immediately stained with hematoxylin and eosin, with at least two cut surfaces of lymph nodes and three cut surfaces of tumor tissue being stained. Microscopic evaluation was performed by two pathologists, and if there were differences in the results, no less than two additional pathologists discussed the case to reach an agreement. Pathologists adopted the newly announced IASLC/ATS/ERS classification system for lung ADC. Since most lung ADCs fall into the mixed subtype, comprehensive histologic subtyping has been proposed to make a semiquantitative assessment of the percentages of the various histologic components and to classify tumors according to the predominant histologic subtype.[2] The percentage of each histological component was recorded in 5% increments. The predominant subtype was defined as the subtype with the largest percentage, even if it accounted for less than 50% of the tumor. Minor components referred to subtypes that occupied more than 5% of the tumor but were not the predominant subtype. The presence of visceral pleural invasion (VPI),[26] microvessel invasion (MVI),[27,28] and spread through air spaces (STAS) were also determined.[29,30]

Statistical Analysis
Clinical and pathological characteristics, such as gender, age, smoking status, platelet count, platelet groups, tumor location, surgery, predominant subtype, tumor diameter, T stage, STAS, VPI, and MVI were analyzed to determine their correlation with lymph node metastasis. Continuous random variables, such as platelet count, age, and tumor

diameter, were expressed as mean ± standard deviation (SD) or mean ± standard error of mean (SEM) and analyzed using the independent samples *t*-test. On univariate analysis, the effect of each factor on the probability of lymph node metastasis and MVI was assessed based on the logistic regression model. All variables with P < 0.1 in the univariate analysis were entered into the multivariate analysis. On multivariate analysis, the stepwise logistic regression analysis was conducted by utilizing these factors. Odds ratios (ORs) and 95% confidence intervals (CIs) were reported for both univariate and multivariate analyses. Lymph node metastasis, STAS, VPI, and MVI were analyzed to determine the correlation with predominant subtypes, the solid-predominant/minor components subtype, and the platelet count. P values were calculated using Pearson's Chi-square test or Fisher's exact test, and the significance level was set at 0.05. The data were analyzed using IBM SPSS Statistics, version 25.0 (IBM, USA).

Results

The characteristics of all 852 patients are listed in Table 1. We divided patients based on N stage: 635 (74.5%) had pathologically validated N0 disease and 217 (25.5%) had validated N1 or N2 disease. No significant differences were observed in terms of gender, age, smoking history, or STAS between the two groups. Platelet count, tumor location, surgical method, predominant subtype, pathological T stage, VPI, and MVI were significantly different between the two groups.

Risk Factors of Lymph Node Metastasis

We first performed univariate regression analysis of variables predicting lymph node metastasis (Table 2). In univariate analysis, platelet count, tumor location, surgical method, tumor size, predominant subtype, VPI, and MVI were significantly associated with lymph node metastasis (P < 0.05). In multivariate analysis, we found that platelet count (aOR = 1.004, 95% CI = 1.001–1.007; P = 0.010), tumor size (aOR = 1.699, 95% CI = 1.471–1.963; P < 0.001), VPI (aOR = 1.635, 95% CI = 1.053–22.332; P = 0.029), and MVI (aOR = 3.074, 95% CI = 1.773–5.330; P < 0.001) remained independent risk factors for lymph node metastasis. The predominant subtype was also a significant independent risk factor of lymph node metastasis. Compared to patients with the lepidic subtype, patients with the acinar subtype had an aOR of 10.430 (95% CI = 3.693–29.457, P < 0.001), patients with

Table 1 Clinicopathological Characteristics of Different Lymph Node Status in 852 Patients with Lung Adenocarcinoma

	Total (n=852)	N0 (n=635)	N1/2 (n=217)	P value
Gender				0.148[a]
Male	392(46)	283(44.6)	109(50.2)	
Female	460(54)	352(55.4)	108(49.8)	
Age (year)		59.77 ±9.376	59.53 ±9.870	0.745
Smoking				0.211
Current or ex	277(32.5)	199(31.3)	78(35.9)	
Never	575(67.5)	436(68.7)	139(64.1)	
PLT (×10^9/L)		235.57 ±58.390	258.59 ±62.733	0.000[a]
Tumor location				0.016[a]
LUL	200(23.5)	143(22.5)	57(26.3)	
LLL	140(16.4)	98(15.4)	42(19.4)	
RUL	278(32.6)	226(35.6)	52(24)	
RML	57(6.7)	45(7.1)	12(5.5)	
RLL	123(19.4)	54(24.9)	177(20.8)	
Surgery				0.012[a, b]
Lobectomy	792(93)	593(93.4)	199(91.7)	
Limited resection	54(6.3)	41(6.5)	13(6.0)	
Pneumonectomy	6(0.7)	1(0.2)	5(2.3)	
Predominant subtype				0.000[a]
L	148(17.4)	144(22.7)	4(1.8)	
A	426(50)	317(49.9)	109(50.2)	
P	129(15.1)	100(15.7)	29(13.4)	
MP	23(2.7)	6(0.9)	17(7.8)	
S	86(10.1)	39(6.1)	47(21.7)	
IMA	40(4.7)	29(4.6)	11(5.1)	
Tumor size (cm)		2.267 ±1.118	3.514 ±1.785	0.000[a]
T stage				0.000[a]
T1	663(77.8)	546(86.0)	117(53.9)	
T2	147(17.3)	76(12)	71(32.7)	
T3/T4	42(4.9)	29(2.0)	13(13.4)	
STAS				0.967
Absent	765(89.8)	570(89.8)	195(89.9)	
Present	87(10.2)	65(10.2)	22(10.1)	
VPI				0.000[a]
Absent	708(83.1)	547(86.1)	161(74.2)	
Present	144(16.9)	88(13.9)	56(25.8)	
MVI				0.000[a]
Absent	773(90.7)	603(95)	170(78.3)	
Present	79(9.3)	32(5)	47(21.7)	

Notes: Data are presented as n (%) or means ± SD. P value, and OR were calculated using the Chi-square test and Fisher's exact test. [a] Statistically significant. [b] Fisher's exact test. **Abbreviations:** OR, odds ratio; LUL, left upper lobe; LLL, left lower lobe; RUL, right upper lobe; RML, right middle lobe; RLL, right lower lobe; L, lepidic; A, acinar; P, papillary; MP, micropapillary; S, solid; IMA, invasive mucinous adenocarcinoma; STAS, spread through air spaces; VPI, visceral pleural invasion; MVI, microscopic vessel invasion.

Table 2 Univariate and Multivariate Logistic Analyses of the Risk Factors Associated with Lymph Nodes Metastasis

Variables	Univariate Analysis			Multivariate Analysis		
	OR	95% CI	P value	aOR	95% CI	P value
Gender						
Male	1					
Female	0.797	0.585–1.085	0.149			
Age (year)	0.997	0.981–1.014	0.745			
Smoking						
Current or ex	1					
Never	0.813	0.588–1.125	0.211			
PLT (×10^9/L)	1.006	1.004–1.009	0.000[a]	1.004	1.001–1.007	0.010[a]
Tumor location						
LUL	1					
LLL	1.075	0.669–1.728	0.764			
RUL	0.577	0.375–0.888	0.012[a]			
RML	0.669	0.330–1.357	0.265			
RLL	1.101	0.707–1.716	0.669			
Surgery						
Limited resection	1					
Lobectomy	1.058	0.556–2.016	0.863			
Pneumonectomy	15.769	1.686–147.509	0.016[a]			
Tumor size (cm)	1.921	1.678–2.200	0.000[a]	1.699	1.471–1.963	0.000[a]
Predominant subtype						
L	1			1		
A	12.379	4.477–34.227	0.000[a]	10.430	3.693–29.457	0.000[a]
P	10.440	3.559–30.622	0.000[a]	6.779	2.228–20.633	0.001[a]
MP	102.000	26.144–397.946	0.000[a]	68.016	16.734–276.455	0.000[a]
S	43.385	14.727–127.805	0.000[a]	19.235	6.262–59.082	0.000[a]
IMA	13.655	4.064–45.880	0.000[a]	5.952	1.587–22.332	0.008[a]
STAS						
Absent	1					
Present	0.989	0.594–1.648	0.967			
VPI						
Absent	1			1		
Present	2.162	1.481–3.156	0.000[a]	1.635	1.053–22.332	0.029[a]
MVI						
Absent	1			1		
Present	5.210	3.223–8.422	0.000[a]	3.074	1.773–5.330	0.000[a]

Notes: P value, OR, and 95% CI were calculated using the Chi-square test and Fisher's exact test. aOR, adjusted odds ratio. [a] Statistically significant.
Abbreviation: CI, confidence interval.

the papillary subtype had an aOR of 6.779 (95% CI = 2.228–20.633, P = 0.001), patients with the micropapillary subtype had an aOR of 68.016 (95% CI = 16.734–276.455, P < 0.001), patients with the solid subtype had an aOR of 19.235 (95% CI = 6.262–59.082, P < 0.001), and patients with invasive mucinous ADC had an aOR of 5.952 (95% CI = 1.587–22.332, P = 0.008) (Table 2).

The Influence of Platelet Count on Tumor Behavior

To determine the effect of platelet count on tumor behavior, we divided the patients into the reference range platelet count group (group α: platelet count 100–300 × 10^9/L) and the thrombocytosis group (group β: platelet count ≥300 × 10^9/L). There were

Table 3 The Influence Between Normal Range Platelet Count (Group α) and Thrombocytosis (Group β) on Microscopic Vessel Invasion (MVI) and Lymph Nodes Metastasis

PLT Group	α (720)	β (130)	P value
T stage, n (%)			0.019[a]
T1	570(79.2)	91(70)	
T2	120(16.7)	27(20.8)	
T3/T4	30(4.2)	12(9.2)	
Lymph node status, n (%)			0.000[a]
N0	553(76.8)	80(61.5)	
N1/2	167(23.2)	50(38.5)	
STAS, n (%)			0.156
Absent	651(90.4)	112(86.2)	
Present	69(9.6)	18(13.8)	
VPI, n (%)			0.525
Absent	601(83.5)	119(80.8)	
Present	105(80.8)	25(19.2)	
MVI, n (%)			0.001[a]
Absent	663(92.1)	108(83.1)	
Present	57(7.9)	22(16.9)	

Notes: Data are presented as n (%). Group a: platelet<300×10^9/L, group b: platelet ≥ 300×10^9/L (two patients of which the platelet < 100×10^9/L were excluded). [a] Statistically significant.

significant differences between group α and group β in T stage (P = 0.019), lymph node positivity (P < 0.001), and the incidence of MVI (P < 0.001). In group α, the

rate of lymph node metastasis was 23.2%, and in group β, it was 38.5% (P < 0.001). (Table 3)

Previous studies have only analyzed these indicators between patients in the platelet reference range and those with thrombocytosis; no previous study has examined clinicopathological factors in patients with platelet counts within the reference range. Therefore, to further investigate the relationship between platelet count and tumor behavior, we divided the patients into six groups based on the platelet count (group 1: platelet count $<150 \times 10^9$/L, group 2: platelet count 150–200×10^9/L, group 3: platelet count 200–250×10^9/L, group 4: platelet count 250–300×10^9/L, group 5: platelet count 300–400×10^9/L, and group 6: platelet count $\geq 400 \times 10^9$/L). Our results showed that as the platelet count increased, the lymph node-positive rate (P < 0.001) and the incidence of MVI (P < 0.001) significantly increased. In addition, the rate of lymph node metastasis increased from group 1 to group 4, despite these patients all having platelet counts within the reference range. In group 6, the rate of lymph node metastasis reached 53.3%. However, no significant difference was observed between the platelet count groups in T stage (P = 0.158) or the incidence of STAS (P = 0.184) (Table 4).

Table 4 The Influence of Different Platelet Counts on Microscopic Vessel Invasion (MVI) and Lymph Nodes Metastasis

PLT Group	1(n=33)	2(n=170)	3(n=313)	4(n=206)	5(n=115)	6(n=15)	P value
T stage, n (%)							0.131
T1	25(75.8)	139(81.8)	252(80.5)	156(75.7)	83(72.2)	8(53.3)	
T2	7(21.2)	26(15.3)	50(16.0)	37(18.0)	22(19.1)	5(33.3)	
T3/T4	1(3.0)	5(2.9)	11(3.5)	13(6.3)	10(8.7)	2(13.3)	
Lymph node status, n (%)							0.000[a]
N0	29(87.9)	140(82.4)	242(77.3)	144(69.9)	73(63.5)	7(46.7)	
N1/2	4(12.1)	30(17.6)	71(32.7)	62(30.1)	42(36.5)	8(53.3)	
STAS, n (%)							0.265
Absent	33	154(90.6)	281(89.8)	185(89.8)	100(87)	12(80)	
Present	0	16(9.4)	32(10.2)	21(10.2)	15(13)	3(20)	
VPI, n (%)							0.012[a]
Absent	30(90.9)	150(88.2)	267(85.3)	156(75.7)	92(80)	13(86.7)	
Present	3(9.1)	20(11.8)	46(31.9)	50(24.3)	23(20)	2(13.3)	
MVI, n (%)							0.000[a]
Absent	33	163(95.9)	291(93.0)	178(86.4)	95(82.6)	13(86.7)	
Present	0	7(4.1)	22(7.0)	28(13.6)	20(17.4)	2(13.3)	

Notes: Data are presented as n (%). [a]Statistically significant. Group 1: platelet<150×10^9/L, group 2: 150≤ platelet<200×10^9/L, group 3: 200≤ platelet<250×10^9/L, group 4: 250≤ platelet<300×10^9/L, group 5: 300≤ platelet<400×10^9/L, group 6: platelet ≥400×10^9/L.

The Influence of Different Subtypes on Platelet Count

As the predominant subtype of ADC influences tumor behavior, such as lymph node metastasis, and poor prognosis, we investigated the relationships between subtype and platelet counts. We found that the platelet count in those with the solid-predominant subtype ($269.70 \pm 69.38 \times 10^9$/L) was significantly higher than those in patients with other subtypes (lepidic: $235.86 \pm 4.611 \times 10^9$/L, acinar: $240.36 \pm 2.749 \times 10^9$/L, papillary: $234.52 \pm 5.883 \times 10^9$/L, micropapillary: $231.17 \pm 10.451 \times 10^9$/L, and invasive mucinous adenocarcinoma: $242.95 \pm 10.761 \times 10^9$/L; $P < 0.001$). The solid-predominant subtype also had the second-highest lymph node-positive rate (54.7%; $P < 0.001$) and the highest incidence of MVI (23.3%; $P < 0.001$) (Table 5). We further compared the solid-minor components group (patients with more than 5% solid subtype but with a predominant subtype other than solid; n = 52) and the solid-predominant group (n = 86) (Table 6). The platelet count in the solid-predominant group was significantly higher than that in the solid-minor components group ($269.70 \pm 69.378 \times 10^9$/L vs $238.35 \pm 58.473 \times 10^9$/L; $P < 0.001$). We observed similar results in the lymph node-positive rate (54.7% vs 22.2%; $P < 0.001$) and the incidence of MVI (23.3% vs 7.7%; $P < 0.001$).

Discussion

Here, we investigated the possible correlation between platelet counts and lymph node metastasis in 852 lung invasive ADC cases. Thrombocytosis was significantly related to lymph node metastasis in lung ADC patients. In addition, higher platelet counts within the reference range were also associated with increased lymph node metastasis. Further, patients with the solid-predominant subtype had significantly higher platelet counts than patients with other pathological subtypes. We speculate that this may be one of the causes of the higher rate of lymph node metastasis in patients with the solid-predominant subtype.

Table 6 The Influence of Different Contents of Solid Subtypes on Platelet Counts

Subtypes	Solid-Minor Components (n=52)	Solid-Predominant (n=86)	*P* value
PLT (×10^9/L)	238.35±58.473	269.70±69.378	0.000[a]
Lymph node status, n (%)			0.000[a]
N0	596(77.8)	39(45.3)	
N1/2	170(22.2)	47(54.7)	
STAS, n (%)			0.405
Absent	690(90.1)	75(87.2)	
Present	76(9.9)	11(12.8)	
VPI, n (%)			0.097[a]
Absent	642(83.8)	66(76.7)	
Present	124(16.2)	20(23.3)	
MVI, n (%)			0.000[a, b]
Absent	707(92.3)	66(76.7)	
Present	59(7.7)	20(23.3)	

Notes: Data are presented as n (%) or means ± SD. [a] Statistically significant. [b] Fisher's exact test.

Table 5 The Influence of Different Predominant Subtype on Platelet Count

Subtype	L (n=148)	A (n=420)	P (n=128)	MP (n=23)	S (n=84)	IMA (n=39)	P value
PLT (×10^9/L)	235.86±4.611	240.36±2.749	234.52±5.883	231.17±10.451	269.70±7.570	242.95±10.761	0.000
Lymph node status							0.000
N0, n (%)	144(97.3)	317(74.4)	100(77.5)	6(26.1)	39(45.3)	29(72.5)	
N1/2, n (%)	4(2.7)	109(25.6)	29(22.5)	17(73.9)	47(54.7)	11(27.5)	
STAS							0.114[a]
Absent	139(93.9)	387(90.8)	109(84.5)	20(87)	75(87.2)	35(87.5)	
Present	9(6.1)	39(9.2)	20(15.5)	3(13)	11(12.8)	5(12.5)	
VPI							0.023[a]
Absent	133(89.9)	360(84.5)	99(76.7)	19(82.6)	66(76.7)	31(77.5)	
Present	15(10.1)	66(15.5)	30(23.3)	4(17.4)	20(23.3)	9(22.5)	
MVI							0.000[a]
Absent	145(98)	397(93.2)	110(85.3)	18(78.3)	66(76.7)	37(92.5)	
Present	3(2)	29(6.8)	19(14.7)	5(21.7)	20(23.3)	3(7.5)	

Notes: Data are presented as n (%)or means ±SEM. [a]Statistically significant.

There are complex interactions between tumor cells and platelets. However, the specific mechanism of their interaction is still under study. Malignancies may promote the production of platelets in a variety of ways,[31–33] and platelets can affect tumor behavior in return. Platelets can promote tumor cell migration and invasion into the surrounding microenvironment, as well as facilitating tumor arrest at the endothelium, extravasation, and seeding.[34–36] They can also protect tumor cells from NK cell-mediated lysis.[37,38] Several studies have reported that platelet depletion results in reduced average tumor weight in patients receiving chemotherapy. Conversely, platelet transfusion increases average aggregate tumor weight.[39–41] Recent evidence suggest that increases in platelets and megakaryocytes can also inhibit metastasis in a site-specific manner.[23] However, these findings were only in preclinical studies in breast cancer. Based on our results, we can conclude that an increase in the platelet count, even within the reference range, is associated with lymph node metastasis in lung ADC patients.

Higher platelet levels indicate lymph node metastasis in patients with invasive ADC.[32,33] We examined platelet levels based on histological subtype and found that the platelet count and the incidence of MVI in patients with solid-predominant tumors were significantly higher than in patients with other histologies. Further, the rate of lymph node metastasis increased with increased platelet count, even if the platelet count was within the reference range. In group α (platelet count: $100–300 \times 10^9$/L), the rate of lymph node metastasis was 23.2%, whereas it was 38.5% in group β (platelet count: $>300 \times 10^9$/L; $P < 0.001$). In patients with a platelet count $>400 \times 10^9$/L, the rate of lymph node metastasis was 53.3%. The rates of lymph node metastasis in the four groups with platelet counts above 200×10^9/L were almost twice as high as the rates in the two groups with platelet counts below 200×10^9/L. Groups α and β also differed significantly in terms of T stage (Table 3). However, there was no significant difference in T stage among groups 1, 2, 3, and 4 (all within the reference range; Table 4). However, there was a significant difference in N stage. As the two main parameters for tumor growth, T stage and N stage are the most likely to be directly related to elevated platelets counts. Therefore, we can conclude that platelet count may have a direct relationship with lymph node metastasis, at least in patients with platelet counts in the reference range.

The relationship between elevated platelet counts and lymph node metastasis is not fully understood. Platelets are known to contain many angiogenic factors, such as vascular endothelial growth factor (VEGF) and platelet-derived growth factor (PDGF). VEGFs are present in resting platelets, which are then released on activation,[42,43] and cancer cells can stimulate the secretion of VEGF-A and other pro-angiogenic factors from platelets.[44] VEGF-C and VEGF-D facilitate tumor metastasis via modulating the lymphatic vessels,[45–47] and VEGF-C can induce sentinel node lymphatic angiogenesis and promote lymphatic metastasis even before the tumor has metastasized.[47,48] In addition, PDGF can stimulate the production of VEGFs by fibroblasts, which can expand the lymphatic vasculature.[47,49,50] We hypothesize that platelets may promote lymphatic metastasis of lung ADC tumor cells in a similar fashion. Some preclinical studies have shown that the exit routes provide by lymph node blood vessels may play role in metastatic tumor cell dissemination.[51] Further, VEGF-C and VEGFR-3 are related to lymphangiogenesis and the development of non–small cell lung cancer.[52] However, as VEGF-C can influence megakaryopoiesis, the correlation of platelet counts and lymph node metastasis may be indirect.[53]

We found that patients with the solid-predominant subtype had the highest platelet counts. Therefore, we speculate that the increased platelets in the solid-predominant subtype may facilitate the increased lymphatic metastasis we observed. Tumor-induced lymphangiogenesis via VEGF-C/VEGFR-3 is the major mechanism through which metastasis to the lymph nodes is promoted.[54] It is therefore very likely that the VEGF-C level is increased locally or systemically in patients with solid-predominant subtypes. However, this has not been assessed. Those with the solid-predominant subtype have higher levels of laminin-5, ezrin, VEGF, and CD204-TAMs than other subtypes.[55] Studies have proven that increases in these factors are significantly related to lymphatic metastasis and poor prognosis in those with resected non-small cell lung cancer.[55,56] However, the effects of platelets in different lung ADC subtypes have not been studied. This is the first study to examine the relationship between platelets and subtypes in lung ADC.

There is some evidence to suggest that aspirin can decrease levels of PDGF and VEGF in the platelet releasate.[57–60] However, previous studies focused on how antiplatelet therapy inhibits platelet-mediated angiogenesis rather than platelet-mediated lymphatic spread. Based on our results, we hypothesize that antiplatelet therapy may reduce lymph node metastasis in all lung ADC patients. Because lymphatic system metastasis is a common cause

of treatment failure and death in postoperative patients with non-small cell lung cancer,[61] targeted therapy against lymph node metastasis can play an important postoperative role. Therefore, antiplatelet therapy such as low-dose aspirin may be a potent direction for lung ADC treatment. Long-term follow-up studies have shown that regular use of aspirin reduces the long-term risk of several cancers, as well as the risk of distant metastasis.[62–64] For lung cancer, the benefit of aspirin was only confirmed in ADC.[25] Anticoagulants are effective for patients with venous thromboembolism. However, antiplatelet therapy may be effective in the lymphatic system because of its effect on the platelet releasate.[44] In a large clinical trial, small doses of aspirin reduced the risk of distant metastases in many ADCs and the overall risk of fatal ADC in the trial population.[62] Another study showed that aspirin can reduce mortality in older patients with several types of ADC.[63] In 2016, the US Preventive Services Task Force recommended regular use of low-dose aspirin in some people with risk factors for colorectal cancer.[65] A large retrospective study including more than 10 million people in South Korea confirmed that long-term use of low-dose aspirin was associated with a decreased risk of lung cancer.[66] Recognition of the tumor-promoting effects of platelets emphasizes the role of anti-platelet therapy in the prevention and treatment of cancer. Nevertheless, identifying cancer patients who will benefit from antiplatelet therapy is still an important issue. The potential risks of antiplatelet therapy should be fully evaluated and investigated.

There are some limitations to this study. First, all patients had invasive lung ADC and had undergone resection. Therefore, patients who were unable to tolerate surgery were excluded. Second, the classification of lung ADC subtypes is still being studied, and non-dominant subtypes might still play a role in tumor biology. Third, pathological inspection of biopsy specimens cannot fully represent the true content of pathological subtypes.[67] Fourth, as a retrospective study, selection bias and performance bias are inevitable. Future multicenter clinical studies with larger samples may validate our results.

Conclusions

Elevated platelet counts are significantly associated with a higher rate of lymph node metastasis, even if the platelet counts are within the reference range. Platelet counts were significantly higher in patients with solid-predominant histology than in patients with other histologies. In addition, VEGF-C may play an important role in lymphatic

metastasis in patients with lung ADC. We hypothesize that antiplatelet therapy may reduce lymph node metastasis in lung ADC patients.

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Disclosure

The authors report no conflicts of interest in this work.

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