Differential clinical manifestations and clinical outcome of cancer-related pulmonary embolism

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Background/Aims: Although acute pulmonary embolism (PE) adversely impacts survival and should be treated regardless of cancer, the treatment rate of cancer-related PE is relatively low. We aimed to compare clinical characteristics and long term prognosis of PE in patients with or without cancer.

Methods: From March 2010 to December 2013, patients with newly diagnosed PE were analyzed. Baseline demographics, comorbidities, cancer status and clinical manifestations of PE were recorded. We defined primary composite outcome as recurrent venous thromboembolism (VTE) and death from PE.

Results: Among a total of 976 patients with PE, the 703 (72.0%) had cancer-related PE. Cancer-related PE group was more frequently asymptomatic (54.5% vs. 13.2%, \( p < 0.001 \)), less extensive (involvement of bilateral pulmonary arteries: 42.8% vs. 51.3%, \( p = 0.017 \); lung infarction: 5.3% vs. 10.3%, \( p = 0.005 \)) and less likely to accompany right ventricular dysfunction (10.3% vs. 27.2%, \( p < 0.001 \)) compared with the non-cancer PE group. Anticoagulation was less frequently undergone in patients with cancer-related PE than those without cancer (62.0% vs. 81.7%, \( p < 0.001 \)). A composite of recurrent VTE and death from PE was significantly higher in the cancer-related PE group (14.4% vs. 6.6%, \( p = 0.001 \)).

Conclusions: Although PE in cancer patients were seem to be less aggressive initially, compared to those without cancer, they had significantly poor prognosis. Given a high rate of recurrent VTE and relatively similar risk of anticoagulation associated bleeding events in cancer patients, more active treatment of PE is warranted in cancer patients.

Keywords: Pulmonary embolism; Venous thromboembolism; Neoplasms

INTRODUCTION

The prevalence of venous thromboembolism (VTE) in patients with active cancer has been reported as four to seven times higher than those without cancer and is increasing with cancer status [1,2]. This is because (1) cancer patients have increased thrombotic risk due to the malignancy itself or chemotherapy, (2) cancer patients frequently undergo serial computed tomography (CT) scanning, which enables early detection of pulmonary embolism (PE). With the increased use of various anticancer drugs that are potentially thrombogenic, the incidence of VTE continues to increase [3-6]. Because the pathogenesis of VTE in cancer patients is quite different from VTE of other cause, such as thrombophilia, trauma, or immobilization, the clinical progress of cancer-related VTE is also different. However, available data regarding the differential clinical manifestations of
cancer-related VTE are limited.

Although current guidelines for PE management uniformly recommend that active cancer patients should receive anticoagulation regardless of the location and extent of PE [7,8], the treatment rate of cancer-related PE in real-world practice is quite low [9]. Concerns about high bleeding tendency or potential interactions between anticoagulants and anticancer drugs in patients with advanced stage cancer cause physicians to neglect to treat PE. However, given that the expected survival increasing in patients with active cancer, it is important to treat combined morbidities such as deep vein thrombosis (DVT) and PE. In the present study, we identified differential clinical manifestations and prognosis of cancer-related PE compared to PE of other causes.

METHODS

Study population and design
The study cohort consisted of consecutive patients diagnosed with PE between October 2010 and December 2013 in our clinic. Among 1,164 eligible patients, patients with known PE before visiting our clinic (n = 58), diagnosed with PE by abdomen CT scan that partially covered the pulmonary artery (n = 55), with filling defect in pulmonary artery stump site after cancer operation (n = 21), who were lost to follow-up (n = 20), and who were finally diagnosed with tumor emboli (n = 19) were excluded. After additional exclusion of 15 patients with chronic PE, 976 patients were included in final analysis (Fig. 1).

Clinical, laboratory, echocardiography, CT findings of PE, and underlying cancer status were collected from our PE registry. The local Institutional Review Board of Samsung Medical Center (2017-12-073) approved this study and waived the requirement for written informed consent.

Provoking factors causing PE were evaluated and, classified as active cancer, major surgery within three months of index event that required general anesthesia lasting 30 minutes, immobilization defined as absolute bed rest for 5 consecutive days and airplane flight longer than six hours, pregnancy or puerperium at the time of PE, usage of oral contraceptives or hormonal replacement therapy, or trauma. PE was considered ‘unprovoked’ in the absence of the above factors. Unprovoked PE was classified as hereditary thrombophilia, antiphospholipid syndrome, or idiopathic event. Patients were classified as having cancer-related PE when they had recurrent or progressive cancer or any malignancy that required curative or palliative treatment at the time of PE detection. Both solid and hematologic malignancies were eligible.

On CT imaging, PE was defined as the existence of contrast filling defects in one or more branches of the pulmonary artery. Location of PE and associated lung infarction was assessed. Proximal PE was defined as PE involving main or lobar level of pulmonary arteries on CT imaging. Distal PE was defined as PE involving segmental or subsegmental level of the pulmonary arteries. Lower extremity DVT was confirmed using Doppler ultrasonography or CT. Proximal DVT was defined as occurring in the popliteal vein and above. Presence of right ventricular (RV) dysfunction was assessed using two-dimensional echocardiography.

Clinical outcomes and definitions
The primary outcome of this study was a composite of recurrent DVT/PE and PE-related death. Secondary end points included all cause of death, which was considered to be a cardiac cause unless a definite non-cardiac cause was established. Clinically significant major and minor bleeding events during anticoagulation were also assessed. Major bleeding was defined according to the In-
ternational Society on Thrombosis and Hemostasis criteria as fatal bleeding or clinically overt bleeding together with a decrease in hemoglobin level of at least 2 g/dL or transfusion of at least 2 units of packed red blood cells occurring at a critical site (intracranial, intraspinal, intraocular, retroperitoneal, pericardial, intraarticular, and intramuscular with compartment syndrome) [10]. All non-major bleeding was considered minor bleeding.

Statistical analyses
Baseline characteristics are summarized with continuous variables and expressed as mean ± standard deviation or median with interquartile range (IQR). Categorical data are presented as a percentage and the number of events. Continuous variables were analyzed using an independent t test or Wilcoxon rank sum test as appropriate, and categorical variables were analyzed using a chi-square test or Fisher’s exact test.

Event-free survival curves were constructed using the Kaplan-Meier estimator and compared using log-rank tests. Cox proportional-hazards regression with backward elimination was used to estimate the hazard ratio as a measure of the relative risk of PE among cancer patients compared with non-cancer patients adjusting for baseline characteristics and comorbidity. All analysis was conducted with SPSS software version 23.0 (IBM Co., Armonk, NY, USA). Statistical significance was concluded at a 2-sided significance level of 0.05 for all analyses.

RESULTS
Among 976 patients who were finally analyzed (age: 65 years [IQR, 54 to 73]; 53.2% male), 703 (72.0%) were classified to the cancer-related PE group. Lung cancer (44.8%) was the most prevalent, followed by colon cancer (8.7%), gynecological malignancy (7.3%), and lymphoma (5.8%). Most patients had advanced stage cancer with distant metastasis (68.4%). Among 273 patients (28.0%) with non-cancer-related PE, 192 patients had provoked PE, which was commonly caused by immobilization (33.0%) or surgery (23.1%). Table 1 shows the baseline characteristics. Patients with cancer-related PE were more frequently male, less commonly smokers, and younger than those with non-cancer PE (male: 56.5% vs. 44.7%, \( p = 0.001 \); smoker: 6.7% vs. 12.1%, \( p = 0.006 \); age: 63 years [IQR, 55 to 71] vs. 69 years [IQR, 54 to 77], \( p < 0.001 \)). There were no significant differences in pre-existing diabetes, hypertension, or DVT between two groups except the presence of chronic kidney disease (5.5% vs. 9.2%, \( p = 0.041 \)).

Table 2 shows the initial clinical manifestation of PE in both groups. Cancer-related PE group was more frequently asymptomatic compared to non-cancer-related PE (76.3% vs. 41.5%, \( p < 0.001 \)). Critical vital status at the time of PE diagnosis, including hypoxia, shock, and tachycardia, was significantly less severe in patients with cancer-related PE than in those without cancer (all \( p < 0.05 \)). PE-related cardiac distress was also less frequent in patients with cancer-related PE (RV dysfunction:...
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https://doi.org/10.3904/kjim.2018.267

10.3% vs. 27.2%, \( p < 0.001 \); N-terminal pro-blood natriuretic peptide: 436.0 pg/mL [IQR, 183.0 to 1,630.0] vs. 900.0 pg/mL [IQR, 144.0 to 3,625.0], \( p = 0.028 \). Cardiac enzymes also tended to be lower in patients with cancer-related PE. On chest CT, involvement of bilateral pulmonary arteries (42.8% vs. 51.3%, \( p = 0.017 \)) and combined lung infarction (53.3% vs. 10.3%, \( p = 0.005 \)) were less prevalent in patients with cancer-related PE than those with non-cancer PE. There was no significant difference in the incidence of coexisting DVT.

Anticoagulation was less frequent in patients with cancer-related PE than those without cancer (62.0% vs.

Table 2. Clinical manifestation of PE

| Characteristic                  | PE without cancer (n = 273) | PE with cancer (n = 703) | Total (n = 976) | \( p \) value |
|--------------------------------|----------------------------|--------------------------|-----------------|--------------|
| Initial symptom/sign           |                            |                          |                 |              |
| No symptom                     | 36 (13.2)                  | 383 (54.5)               | 419 (42.9)      | < 0.001      |
| Dyspnea                        | 139 (50.9)                 | 201 (28.6)               | 340 (34.8)      | < 0.001      |
| Chest pain                     | 41 (15.0)                  | 32 (4.6)                 | 73 (7.5)        | < 0.001      |
| Leg swelling                    | 47 (17.2)                  | 65 (9.2)                 | 112 (11.5)      | < 0.001      |
| Fever                          | 50 (18.3)                  | 40 (5.7)                 | 90 (9.2)        | < 0.001      |
| Hypoxia                        | 134 (50.0)                 | 151 (22.3)               | 285 (30.2)      | < 0.001      |
| Tachycardia                    | 114 (42.5)                 | 233 (34.4)               | 347 (36.7)      | 0.020        |
| Shock                          | 35 (12.1)                  | 36 (5.3)                 | 71 (7.5)        | < 0.001      |
| Unexpected PE\(^a\)            | 40 (14.7)                  | 474 (67.4)               | 514 (52.7)      | < 0.001      |
| Laboratory finding             |                            |                          |                 |              |
| D-dimer                        | 5.9 (2.8–15.4)             | 9.5 (4.7–19.1)           | 8.0 (3.7–18.0)  | < 0.001      |
| Hemoglobin                     | 12.1 (10.0–13.4)           | 11.2 (10.1–12.6)         | 11.4 (10.0–13.0) | 0.001      |
| Platelet                       | 197 (152–250)              | 203 (141–270)            | 200 (144–264)   | 0.751        |
| NT-proBNP                      | 900 (144–3,625)            | 436 (183–1,630)          | 586 (167–2,502) | 0.028        |
| CK-MB                          | 1.97 (0.78–4.75)           | 1.56 (0.57–3.32)         | 1.83 (0.60–3.91) | 0.024        |
| Troponin I                     | 0.029 (0.006–0.132)        | 0.019 (0.006–0.115)      | 0.022 (0.006–0.118) | 0.145        |
| Echocardiographic finding      |                            |                          |                 |              |
| RV dysfunction                 | 41 (27.2)                  | 17 (10.3)                | 58 (18.4)       | < 0.001      |
| D-shaped LV                    | 32 (21.2)                  | 14 (8.5)                 | 46 (14.6)       | 0.001        |
| CT finding                     |                            |                          |                 |              |
| Proximal\(^b\)                 | 170 (62.3)                 | 415 (59.0)               | 585 (59.9)      | 0.354        |
| Bilateral involvement          | 140 (51.3)                 | 301 (42.8)               | 441 (45.2)      | 0.017        |
| Lung infarction                | 28 (10.3)                  | 37 (5.3)                 | 65 (6.7)        | 0.005        |
| Coexisting DVT\(^c\)           | 139 (60.2)                 | 160 (57.6)               | 299 (58.7)      | 0.550        |
| Distal DVT                     | 45 (19.5)                  | 52 (18.7)                | 97 (19.1)       | 0.836        |
| Proximal DVT                   | 94 (40.7)                  | 108 (38.8)               | 202 (39.7)      | 0.836        |

Values are presented as number (%) or median (interquartile range).

PE, pulmonary embolism; NT-proBNP, N-terminal pro-blood natriuretic peptide; CK-MB, creatine kinase myocardial band; RV, right ventricle; LV, left ventricle; CT, computed tomography; DVT, deep vein thrombosis.

\(^a\)Unexpected PE was defined as incidentally detected PE on computed tomographic examination without PE-related symptom/ sign.

\(^b\)Proximal PE includes filling defects in the main, central, and lobar pulmonary artery.

\(^c\)DVT is confirmed using Doppler ultrasonography or computed tomography. Proximal DVT was defined as occurring in the popliteal vein and above.
Patients with cancer-related PE had more frequently use of vitamin K antagonist as anticoagulation than those with non-cancer PE. On the other hand, in the cancer-related PE group, the frequency of low molecular weighted heparin was relatively higher compared to non-cancer PE group. Details on PE treatment are provided in Table 3. Thrombolysis and surgical thrombectomy were rarely performed in patients with cancer-related PE. Among patients treated with anticoagulants, clinically relevant major or minor bleeding events were not different between the two groups (major bleeding: 3.8% vs. 3.3%, p = 0.850; minor bleeding: 1.7% vs. 2.2%, p = 0.601) (Table 4). During follow-up periods, a composite of recurrent VTE and death from PE was significantly higher in the cancer-related PE group (14.4% vs. 6.6%, p = 0.001) (Fig. 2). In particular, recurrent VTE events occurred more frequently in patient with cancer-related PE (11.1% vs. 4.0%, p = 0.001). No anticoagulation and current smoking were independently associated with poor prognosis in patients with non-cancer PE, while no anticoagulation was the only prognostic factor in those with cancer-related PE (Table 5).

After excluding patients who presented with unexpected incidentally detected PE, subgroup analysis was also performed in those with symptomatic PE (229 patients with cancer-related PE and 233 patients with non-cancer PE). Baseline characteristics and clinical manifestations of PE in this subgroup are shown in Supplementary Tables 1 and 2. Patients with suspected cancer-related PE consistently had a higher risk of recurrent VTE com-
pared to those with suspected non-cancer PE (Fig. 3). In multivariate analysis, the presence of cancer itself was the strongest poor prognostic factor for PE irrespective of suspected PE or incidental PE (adjusted hazard ratio, 9.75; 95% confidence interval, 2.4 to 39.2; \( p = 0.001 \)) (Supplementary Table 3).

**DISCUSSION**

In this observational study, we investigated the clinical features and prognosis of PE according to causes that were cancer-related or not. Compared to patients with non-cancer PE, those with cancer-related PE tended to be more asymptomatic and had a lower rate of combined cardiac dysfunction and smaller burden of thromboembolism. However, despite less severe clinical manifestations, PE-related adverse events including recurrent VTE and death from PE were significantly higher in patients with cancer-related PE. The rate of anticoagulation-related bleeding events was similar in the two groups. Our data show the differential clinical manifesta-
tations of cancer-related PE and subsequent PE-related clinical outcomes from our consecutive PE registry data.

Although currently recommended treatment strategies for cancer-related PE are not different than those for non-cancer PE, in terms of real world clinical practice, the treatment rate of PE is relatively low in patients who present with cancer-related PE compared to patients without cancer [9]. In our data, the treatment rate of cancer-related PE was only 62.0%, which is significantly lower than those diagnosed with other causes of PE. There are several possible explanations for our findings. First, as shown in the results, the clinical features of PE were far less severe with cancer-related PE than with non-cancer PE, despite uncontrolled hypercoagulability related to malignancy and the poor functional status of cancer patients. Second, concern regarding potential interactions between anticoagulants and anticancer drugs or relatively short expected survival may cause physicians to waver in treating PE in patients with cancer. Moreover, the high bleeding risk associated with bone marrow suppression hampers the use of anticoagulants in patients with cancer. The relatively mild clinical manifestations at PE diagnosis and concerns about anticoagulation-related complications are likely the main causes for neglecting to provide adequate treatment of thromboembolism in patients with cancer.

Incidentally detected PE is more frequent in patients with cancer than in those without cancer. However, there is a conflict regarding the efficacy of anticoagulation in incidentally detected PE. While some researchers report that cancer patients with incidental PE have a similar risk for mortality and PE recurrence compared to those with suspected PE, some studies demonstrated that the risk of progressive or recurrent PE without anticoagulation is low in patients with incidental PE [11-13]. For this reason, we also analyzed a subgroup diagnosed with suspected PE. Even in patients with suspected PE, patients with cancer consistently had mild initial clinical features, a low rate of anticoagulation, and higher VTE recurrence than patients without cancer.

Although physicians may feel burdened by anticoagulation therapy for patients with cancer because of concerns about bleeding complications, malignancy does not raise the risk of bleeding in practice. At the time of the study, when before the effectiveness and safety of new oral anticoagulant (NOAC) was confirmed in cancer-related PE patients, the use of NOAC was relatively low in this study. However, only 36 patients treated with anticoagulants experienced major bleeding events in our study. Furthermore, there were no statistically significant differences in bleeding complications between cancer-related PE and non-cancer PE. These observa-
tions support the current guideline recommendations for treatment with anticoagulation, even in patients with cancer-related PE.

Although the association between smoking and VTE recurrence remains controversial, several studies showed that some factors induced by smoking may promote formation of a clot and subsequent VTE. In our study, smoking was a poor prognostic factor in patients with non-cancer PE. However, with cancer-related PE, there was no significant association between smoking and VTE recurrence. This is because the presence of airflow limitation did not increase the risk of VTE recurrence in cancer patients with PE [14].

In contrast to previous studies, cancers associated with especially high rates of PE are not consistent with other research [15-17]. Several western studies showed that pancreatic cancer is associated with a high VTE risk [18]. In this study, however, lung cancer was the most common cancer in patients with cancer-related PE. This may result from differences in the incidence of cancer among races. Lung cancer is the most common cancer in all of Asia [19]. In particular, it is the second most common cancer in Korea. Similarly, a report from China described lung cancer as the most common malignancy diagnosed in patients with PE [20]. With advances in anticancer treatment, especially in patients with lung cancer, expected survival is significantly increasing. Therefore, it is important to treat comorbidities in addition to treating cancer itself. Given the higher rate of recurrent VTE and relatively similar risk of anticoagulation-associated bleeding events in patients with cancer, more active PE treatment is warranted.

Study limitation

This study has some limitations. Its retrospective design may enhance information bias. To minimize this bias, a prespecified and standardized protocol was used to thoroughly review all medical charts of included patients. Cancer patients may have undergone heightened surveillance for VTE, leading to surveillance bias. Although the risk of VTE varies with different types of cancer [18], this study included all stages of cancer. Further research is needed to describe the characteristics and outcomes of PE according to cancer stage. Median follow-up duration was 10.4 months. The follow-up duration was shorter than expected, which is largely attributed to cancer-related deaths.

In conclusion, the treatment rate of PE was 62.0% in patients with current cancer. In patients with cancer, PE tended to have less aggressive initial clinical manifestations and thrombus burden compared to PE in patients without cancer. However, given the high rate of recurrent VTE, more active PE treatment is warranted.

KEY MESSAGE

1. Pulmonary embolism (PE) with cancer was less aggressive initially than those without cancer.
2. PE related adverse events were higher in patients with cancer.
3. In terms of bleeding complication, there were no statistically significant differences between cancer-related PE and non-cancer PE.

Conflict of interest

No potential conflict of interest relevant to this article was reported.

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### Supplementary Table 1. Baseline characteristics in patients with symptomatic PE

| Variable                  | Suspected PE | Total (n = 462) | p value |
|---------------------------|--------------|-----------------|---------|
|                           | Without cancer (n = 233) | With cancer (n = 229) |         |
| Age, yr                   | 70 (54–77)   | 64 (56–72)      | 66 (54–75) | 0.011 |
| Male sex                  | 101 (46.1)   | 118 (51.5)      | 219 (47.4) | 0.078 |
| BMI, kg/m²                | 24.1 (21.4–27.1) | 23.3 (21.2–25.4) | 23.7 (21.4–26.1) | 0.014 |
| Current smoker            | 28 (12.2)    | 11 (4.8)        | 39 (8.5)  | 0.005 |
| Comorbidities             |              |                 |         |
| Diabetes                  | 31 (13.3)    | 41 (17.9)       | 72 (15.6) | 0.173 |
| Hypertension              | 89 (38.2)    | 78 (34.1)       | 167 (36.1) | 0.355 |
| Chronic kidney disease    | 21 (9.0)     | 18 (7.9)        | 39 (8.4)  | 0.656 |
| Previous DVT              | 11 (4.7)     | 16 (7.0)        | 27 (5.8)  | 0.299 |
| Previous cancer           | 26 (11.2)    | 9 (1.9)         | 35 (7.6)  | 0.003 |

Values are presented as median (interquartile range) or number (%).
PE, pulmonary embolism; BMI, body mass index; DVT, deep vein thrombosis.
## Supplementary Table 2. Clinical manifestation in patients with symptomatic PE

| Variable                  | Suspected PE | Total (n = 462) | p value |
|---------------------------|--------------|-----------------|---------|
|                           | PE without cancer (n = 223) | PE with cancer (n = 229) |         |
| Initial symptom/sign      |              |                 |         |
| No symptom                | 9 (3.9)      | 10 (4.4)        | 19 (4.1) | 0.785   |
| Dyspnea                   | 136 (58.4)   | 148 (64.6)      | 284 (61.5) | 0.167   |
| Chest pain                | 39 (16.7)    | 24 (10.5)       | 63 (13.6) | 0.050   |
| Leg swelling              | 46 (19.7)    | 46 (20.1)       | 92 (19.9) | 0.926   |
| Fever                     | 50 (21.5)    | 34 (14.8)       | 84 (18.2) | 0.065   |
| Hypoxia                   | 131 (57.0)   | 135 (59.5)      | 266 (58.2) | 0.586   |
| Tachycardia               | 108 (47.0)   | 132 (58.1)      | 240 (52.5) | 0.017   |
| Shock                     | 35 (15.2)    | 31 (13.7)       | 66 (14.4) | 0.635   |
| Laboratory finding        |              |                 |         |
| D-dimer                   | 7.1 (3.3–16.9) | 12.6 (6.4–23.9) | 7.9 (2.7–19.1) | 0.001   |
| Hemoglobin                | 12.4 (9.8–13.7) | 11.0 (9.8–12.5) | 11.4 (9.8–13.1) | < 0.001 |
| Platelet                  | 198 (147–241) | 168 (117–224)   | 191 (136–247) | 0.058   |
| NT-proBNP                 | 1,235 (210–3,847) | 636 (229–2,110) | 741 (204–2,920) | 0.095   |
| CK-MB                     | 2.29 (0.95–5.48) | 1.89 (0.69–3.67) | 1.89 (0.73–4.20) | 0.064   |
| Troponin I                | 0.043 (0.009–0.246) | 0.024 (0.008–0.125) | 0.029 (0.007–0.134) | 0.207   |
| Echocardiographic finding |              |                 |         |
| RV dysfunction            | 41 (30.1)    | 17 (18.7)       | 58 (25.6) | 0.052   |
| D-shaped LV               | 32 (23.5)    | 12 (13.2)       | 44 (19.4) | 0.053   |
| CT finding                |              |                 |         |
| Proximal                  | 149 (65.9)   | 142 (62.0)      | 291 (65.6) | 0.666   |
| Bilateral involvement     | 131 (56.2)   | 123 (53.7)      | 254 (55.6) | 0.587   |
| Lung infarction           | 27 (11.6)    | 15 (6.6)        | 42 (9.1)  | 0.060   |
| Coexisting DVT            |              |                 |         |
| Distal DVT                | 40 (19.4)    | 34 (19.2)       | 74 (19.3) | 0.979   |
| Proximal DVT              | 84 (40.8)    | 74 (41.8)       | 158 (41.3) | 0.979   |

Values are presented as number (%) or median (interquartile range).

PE, pulmonary embolism; NT-proBNP, N-terminal pro-blood natriuretic peptide; CK-MB, creatine kinase myocardial band; RV, right ventricle, LV, left ventricle; CT, computed tomography; DVT, deep vein thrombosis.

aProximal PE includes filling defects in the main, central and lobar pulmonary artery.

bDVT is confirmed using Doppler ultrasonography or computed tomography. Proximal DVT was defined as occurring in the popliteal vein and above.
**Supplementary Table 3. Predictive factors for recurrent VTE and death from PE**

| Variable                      | Adjusted HR | 95% CI     | p value |
|-------------------------------|-------------|------------|---------|
| All patients with PE          |             |            |         |
| Cancer                        | 9.75        | 2.4–39.2   | 0.001   |
| No anticoagulation            | 3.95        | 1.1–11.8   | 0.048   |
| Current smoking               | 5.3         | 1.6–17.3   | 0.006   |
| Bilateral lung involvement    | 3.46        | 1.0–11.8   | 0.048   |
| Incidental PE                 | 0.1         | 0.0–0.95   | 0.045   |
| Suspected PE                  |             |            |         |
| Cancer                        | 8.70        | 2.0–37.4   | 0.004   |
| No anticoagulation            | 7.96        | 2.1–30.5   | 0.002   |
| Current smoking               | 5.94        | 1.8–20.1   | 0.004   |

Adjusted for age, gender, body mass index, cancer, no anticoagulation, duration of treatment, proximal located PE, bilateral lung involvement, shock, tachycardia, hypoxia, smoking, right ventricular dysfunction, unexpected PE, N-terminal pro-blood natriuretic peptide, creatine kinase myocardial band, hemoglobin, and D-dimer.

VTE, venous thromboembolism; PE, pulmonary embolism; HR, hazard ratio; CI, confidence interval.