



By

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Introduction

- **Pulmonary embolism (PE) is a major international health problem.**
- The diagnosis is often difficult to obtain and is frequently missed.
- Mortality in untreated PE is approx. 30%, but with adequate treatment, it can be reduced to 2-8%.
- The prevalence of PE at autopsy (12-15%) in hospitalized patient has not changed over the last 3 decades.

Classification

For clinical purposes PE can be classified into two main groups: massive and non-massive.

- Massive PE consists of shock and/or hypotension (defined as a systolic blood pressure <90 mmHg or a pressure drop of 40 mmHg for >15 min if not caused by new-onset arrhythmia, hypovolemia or sepsis).
- Otherwise *non-massive PE* can be diagnosed.
- A subgroup of patients with non-massive PE may be identified by echocardiographic signs of right ventricular hypokinesis. This subgroup is called *submassive*, because there is growing evidence that the prognosis of this patient group may be different from those with non-massive PE and normal right ventricular function.

Diagnosis of PE

- **PE** has a wide range of clinical presentation.
- A reasonable clinical suspicion is required to avoid missing the diagnosis of PE.
- First line diagnostic tests, such as ECG, chest X-ray and blood-gas analysis are indicated to assess clinical probability of PE and general condition of the patient.

Diagnosis of PE

- Clinical evaluation is accurate to discriminate a subgroup of patients with a low likelihood of PE.
- Clinical probability may be estimated empirically or explicitly by a prediction rule.
- Patients with a low clinical probability of PE, no lower limb deep vein thrombosis and a nondiagnostic lung scan have a very low risk of PE.

Risk factors of VIE

- ◆Age >40 yr
- **L**ong-haul air travel
- History of venous thromboembolism
- Surgery requiring >30 min of anesthesia
- Prolonged immobilization
- *Cerebrovascular accident
- **Congestive heart failure**
- Cancer
- Fracture of pelvis, femur, or tibia

Risk factors of VIE

- Obesity
- **Pregnancy or recent delivery**
- **Estrogen therapy**
- Inflammatory bowel disease
- Genetic or acquired thrombophilia
 - >Factor V Leiden
 - >Antithrombin III deficiency
 - >Protein C deficiency
 - >Protein S deficiency
 - >Prothrombin G20210A mutation
 - >Anticardiolipin antibody syndrome
 - >Lupus anticoagulant

Symptoms of PE

	PE	No PE
yspnea	80 %	59 %
hest pain pleuritic	52 %	43 %
hest pain substernal	12 %	8 %
Cough	20 %	25 %
Iemoptysis	11 %	7 %
yncope	19 %	11 %

Signs of PE

	PE	No PE
achypnoea (20/min)	70%	68%
achycardia (> 100/min)	26 %	23 %
gns of DVT	15 %	10 %
ever > 38.5	7 %	17 %
anosis	11 %	9 %

Routine Investigation

	PE	No PE
est X-ray		
electasis or infiltrate	49%	45%
eural effusion	46%	33%
eural-based opacity (infarction)	23%	10%
evated diaphragm	36%	25%
creased pulmonary vascularity	36%	6%
nputation of hilar artery*	36%	1%
poxemia	75%	81%
		100/

Diagnostic Prediction

Wells et. al.:

>7Clinical Data

• DVT	$(3\cdot 0)$	points)	
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- Alternative diagnosis is less likely than PE (3.0 points)
- Heart rate above 100/min
 (1.5 points)
- Immobilisation or surgery in the previous 4 w (1.5 points)
- Previous DVT or PE
 (1.5 points)
- Haemoptysis (1 point)
- Cancer, being treated currently orwithin the previous 6 months or palliative (1 point)

> Clinical Probability:

- Low < 2
- Intermediate 2-6

Diagnostic Prediction

British Thoracic Society

- A standard assessment of pre-test clinical probability might include the following.
 - A. The patient has clinical features compatible with PE (raised respiratory rate, which may be accompanied by haemoptysis, pleuritic chest pain, or both)
- Plus two other factors:
 - 1. The absence of another reasonable clinical explanation
 - 2. The presence of a major risk factor

> Assessment

- A. PLUS 1. AND 2.: high pre-test clinical probability
- A PIJIS 1 OP 2 intermediate protect clinical probability

Modified pretest probability for DVT

nical feature	Score
derness along entire deep vein system	1
lling of the entire leg	1
ater than 3 cm difference in calf circumference	1
ing oedema	1
ateral superficial veins	1
t factors present:	
Active cancer	1
Prolonged immobility or paralysis	1
Recent surgery or major me <mark>dical illne</mark> ss	1
native diagnosis likely (ruptured Baker's cyst in rheumatoid ritis, superficial thrombophlebitis, or infective cellulitis)	-2

- Most cases of deep vein thrombosis (DVT) about 90%) start in the calf solated DVT of the calf rarely causes:
- ➤ Leg symptoms (80% of the cases of symptomatic DYT involve the proximal veins)
- >Clinically important pulmonary embolism.
- About 1/4 of untreated cases of DVT in the calf will extend to involve the proximal veins and do so within a week of

Most patients with symptomatic proximal DVT and without chest symptoms have lung scan evidence of oulmonary embolism (about 40% have 'high-probability'' lung scans). These abnormalities are often misdiagnosed as new pulmonary embolism during reatment.

- About 75% of all patients who are diagnosed with PE have DVT; about 2/3 of these cases nvolve the proximal veins
- >Patients with less extensive PE are less likely to have proximal DVI'
- >Up to 25% of patients with symptomatic PE have clinical evidence of DYT.
- Without treatment, about 1/2 of patients with symptomatic proximal DVT or PE are expected to have recurrent VTE within 3

After PE, as compared with DVT, at east within the first 3 months, a high proportion of recurrent episodes of VTE are fatal PE (case fatality rate over 2-fold higher) 10% of symptomatic PE cases are estimated to be fatal within an hour of irst symptoms:

> 5%-10% of patients with PR have shock

- ➤ About 50% of patients who are diagnosed with PE have echocardiographic evidence of RV dysfunction at presentation, a finding that is associated with an elevated short-term mortality.
- With treatment of PE, about 50% resolution of perfusion defects is expected after 2–4 weeks. Eventually, complete resolution of PE embolism is expected to occur in about 2/3 of

Natural History & Prognosis

imary

- Iortality in untreated PE is 25-30%
- ntreated VTE has a high risk of (fatal or on-fatal) recurrence.
- Inticoagulant therapy reduces the mortality patients with PE by 75%.
- The prognosis of treated, non-massive VTE is ainly dependent on co-existing illnesses, such a malignancy or CV diseases

Take home message

PE is a common medical emergency that aces nearly all medical specialties The diagnosis is often difficult and frequently nissed Mortality of untreated PE about 30% that can be reduced to 2-8 % with adequate reatment So, use of clinical prediction rules and liagnostic algorithms is mandatory to avoid his fatal clinical amargancy aspecially in

