

Utilità degli score prognostici cardiovascolari per il medico di pronto soccorso



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DEFINITION

Classification systems may help to quantify symptoms and stages of disease, specific scores enable risk stratification and may facilitate decision-making in various cardiac disorders

The number of points, goals, runs, etc. achieved in a game or by a team or an individual.



THE CHALLENGE: ACUTE CHEST PAIN

Cardiac	Pulmonary	Vascular	Gastro-intestinal	Orthopaedic	Other
Myopericarditis Cardiomyopathies ^a	Pulmonary embolism	Aortic dissection	Oesophagitis, reflux or spasm	Musculoskeletal disorders	Anxiety disorders
Tachyarrhythmias	(Tension)-Pneumothorax	Symptomatic aortic aneurysm	Peptic ulcer, gastritis	Chest trauma	Herpes zoster
Acute heart failure	Bronchitis, pneumonia	Stroke	Pancreatitis	Muscle injury/ inflammation	Anaemia
Hypertensive emergencies	Pleuritis		Cholecystitis	Costochondritis	
Aortic valve stenosis				Cervical spine pathologies	
Tako-Tsubo cardiomyopathy					
Coronary spasm					
Cardiac trauma					

Hospital attendances and admissions for acute chest pain present a substantial burden, accounting for 5% of all emergency department attendances, and 40% of acute medical admissions



Acute Chest Pain → AMI

1st History

2nd 12-lead ECG

3rd Troponin

Diagnosis



No ST-Elevation

ST-Elevation



Other

Unstable Angina

NSTEMI

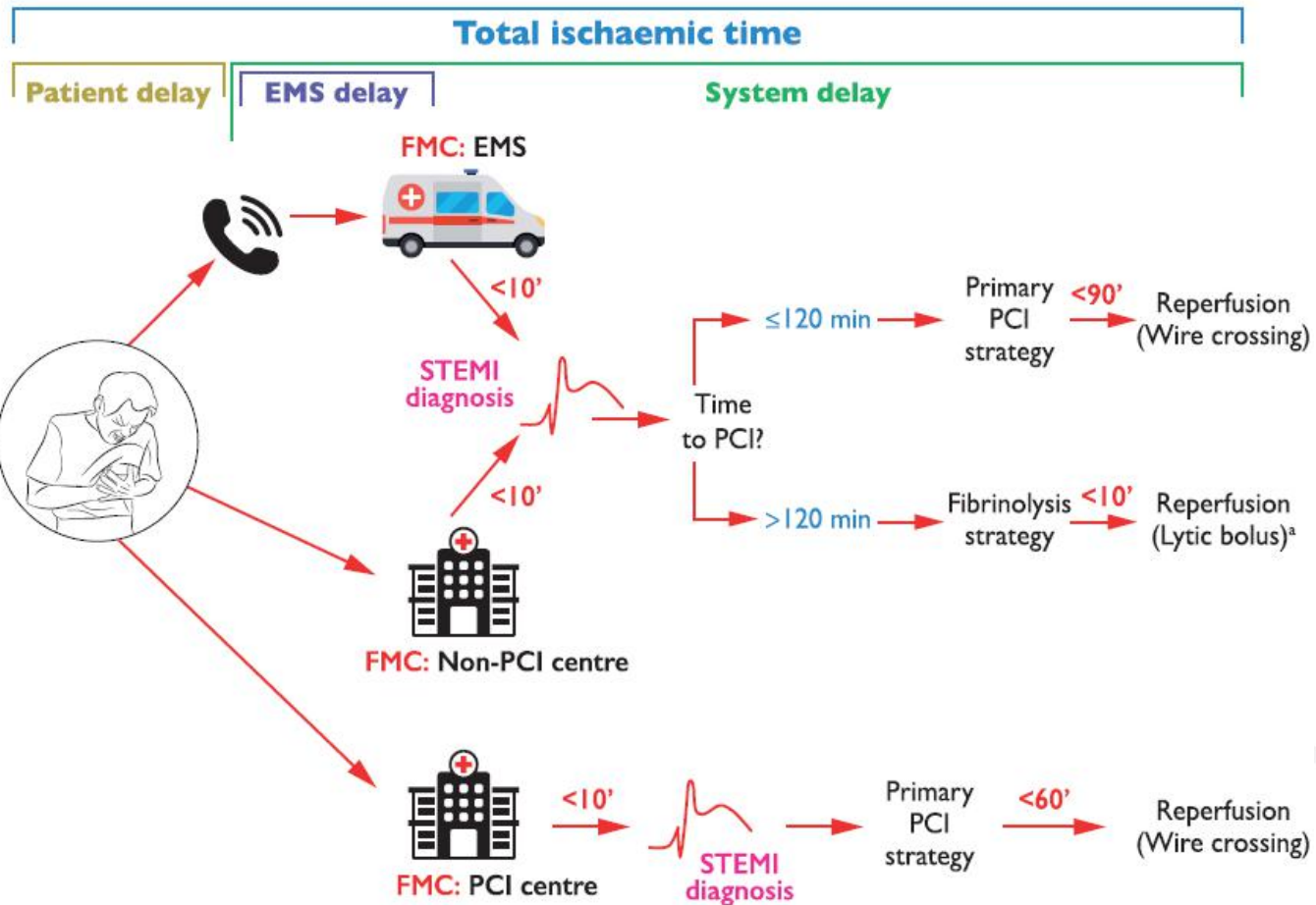
STEMI

AMI



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The ECG

The Diagnostic Key?

- The standard ECG is the single best test to identify patients with an AMI upon E.D. presentation
- But sensitivity is still far from ideal
 - ST elevation in 50% of **AMI's**
 - 1-5% of **AMI's** have a normal initial ECG
 - 4 - 23% of pts. with unstable angina have a normal ECG



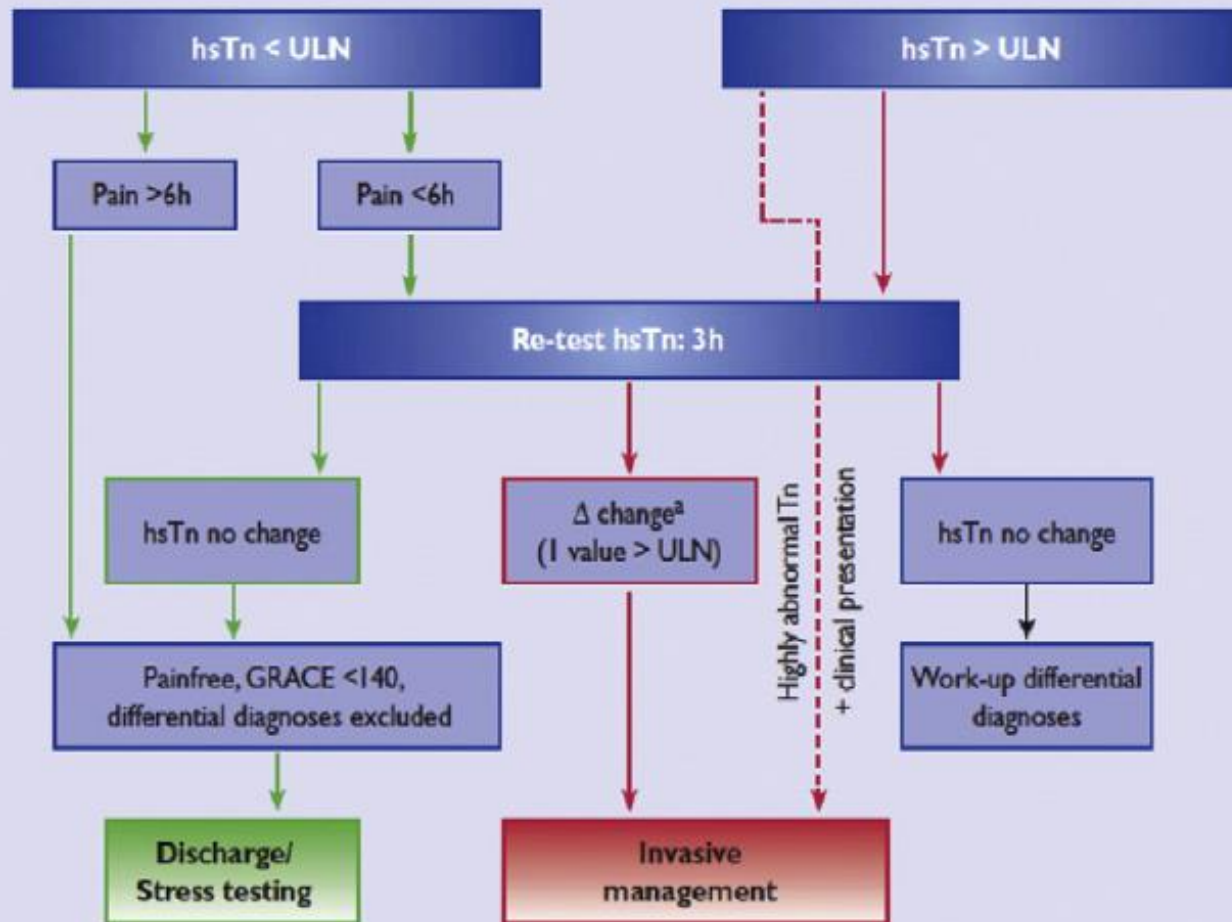
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Acute Chest Pain



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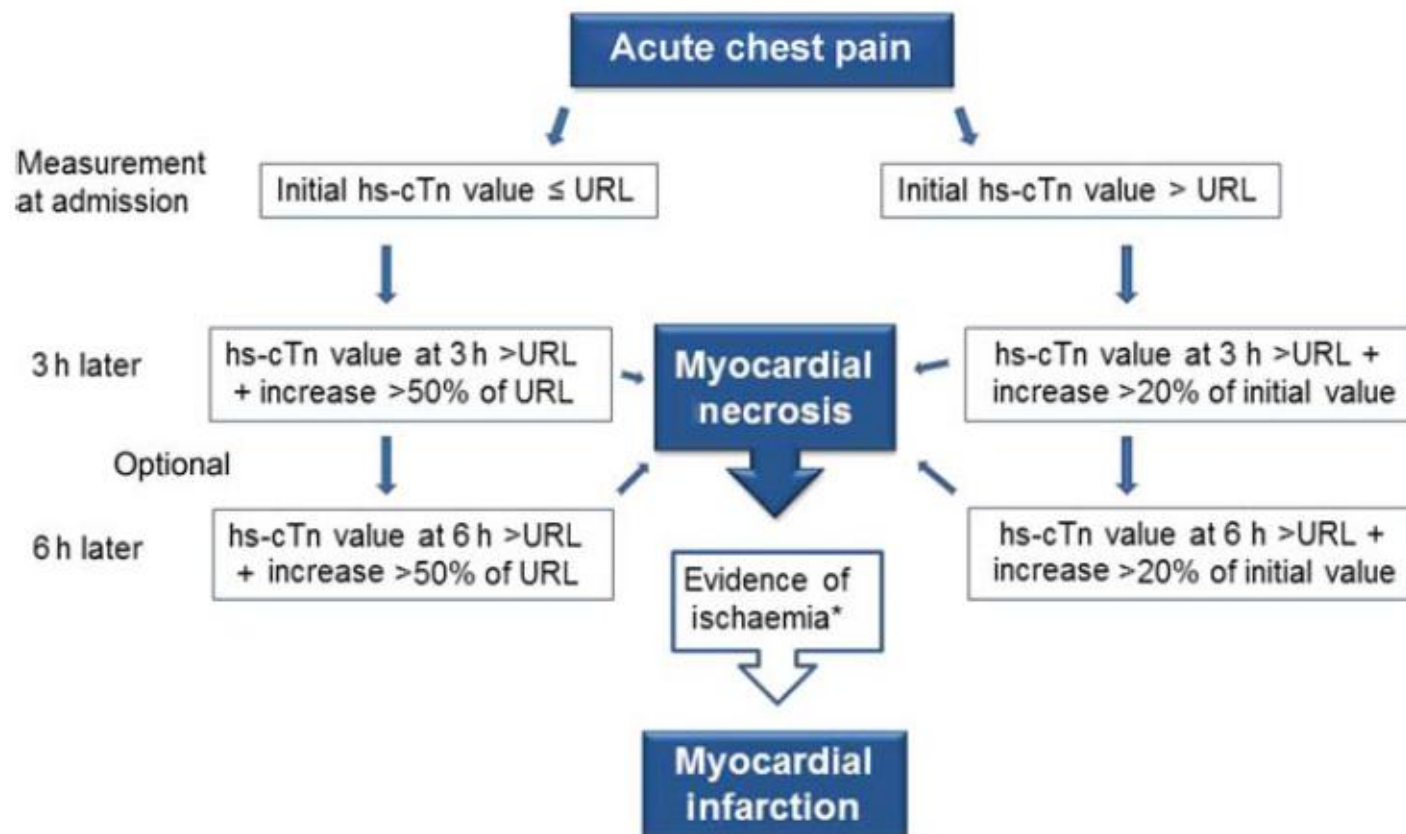
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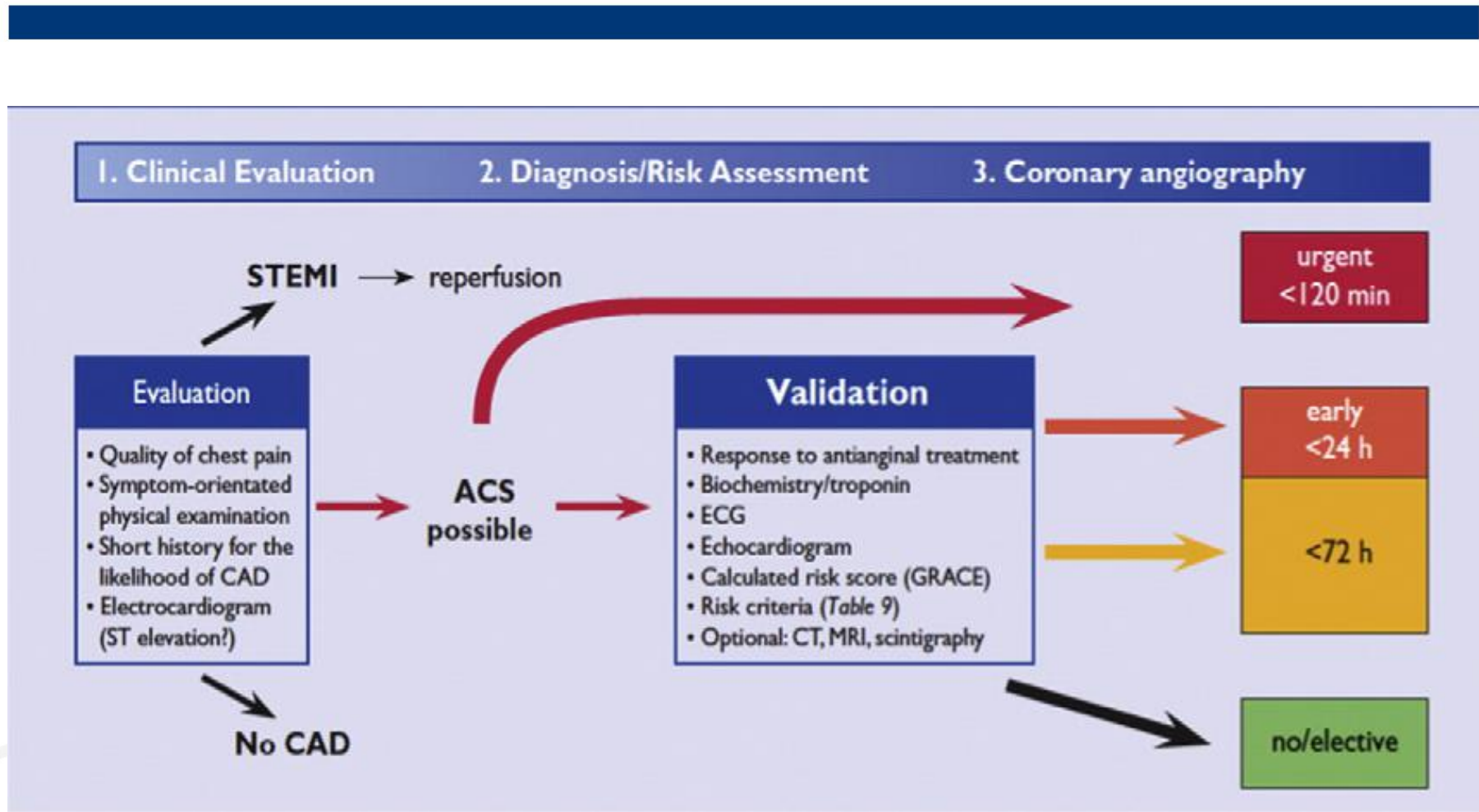
ESC rapid rule-out of ACS with hs-cTn algorithm.



Rapid early rule-in of AMI with high-sensitivity cardiac troponin



The current ESC guideline on the use of hs-cTn recommends that for patients with a negative initial hs-cTn or values close to the upper reference limit URL, a 50% delta value or an absolute increase of 7 ng/L at 3 h can be used to diagnose MI. In patients with elevated hs-cTn at presentation, a delta value of 20% at 3 h can be used.

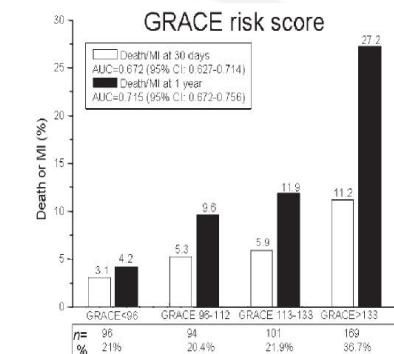
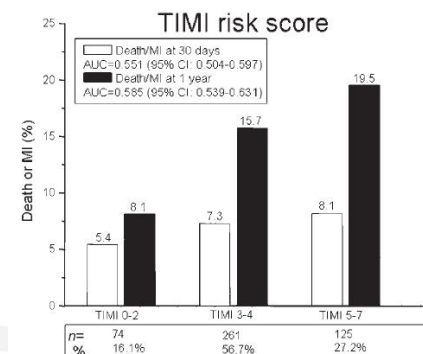
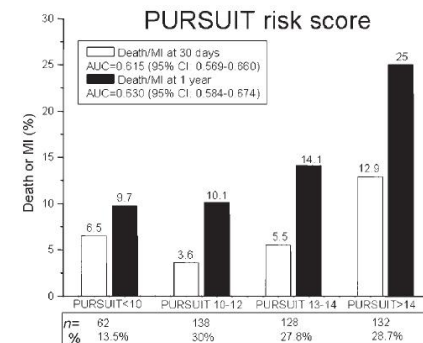


TIMI, PURSUIT, and GRACE risk scores: sustained prognostic value and interaction with revascularization in NSTEMI-ACS

PURSUIT (0-18)	Age, separate points for enrolment diagnosis	Decade [UA (MI)]
	50	8 (11)
	60	9 (12)
	70	11 (13)
	80	12 (14)
	Sex	
	Male	1
	Female	0
	Worst CCS-class in previous 6 weeks	
	No angina or CCS I/II	0
	CCS III/IV	2
	Signs of heart failure	2
	ST-depression on presenting ECG	1

TIMI (0-7)	Age ≥ 65 years	1
	≥ 3 risk factors for CAD	1
	Use of ASA (last 7 days)	1
	Known CAD (stenosis $\geq 50\%$)	1
	>1 episode rest angina in <24 h	1
	ST-segment deviation	1
	Elevated cardiac markers	1

GRACE (0-258)	Age (years)	
	<40	0
	40-49	18
	50-59	36
	60-69	55
	70-79	73
	≥ 80	91
	Heart rate (bpm)	
	<70	0
	70-89	7
	90-109	13
	110-149	23
	150-199	36
	>200	46
	Systolic BP (mmHg)	
	<80	63
	80-99	58
	100-119	47
	120-139	37
	140-159	26
	160-199	11
	>200	0
	Creatinine (mg/dL)	
	0-0.39	2
	0.4-0.79	5
	0.8-1.19	8
	1.2-1.59	11
	1.6-1.99	14
	2-3.99	23
	>4	31
	Killip class	
	Class I	0
	Class II	21
	Class III	43
	Class IV	64
	Cardiac arrest at admission	43
	Elevated cardiac markers	15
	ST-segment deviation	30



MINI-GRACE SCORE



Calculator

1. INPUT DATA > 2. DEATH / DEATH MI RESULTS

Age (years)	<input type="text"/>	ST-segment deviation	<input type="checkbox"/>
Heart rate (bpm)	<input type="text"/>	Cardiac arrest at admission	<input type="checkbox"/>
Systolic blood pressure (mmHg)	<input type="text"/>	Elevated troponin*	<input type="checkbox"/>
CHF (Killip class)	<input type="text"/>	* Or other necrosis cardiac biomarkers	
Diuretic usage	<input type="checkbox"/>		
Creatinine (mg dL ⁻¹ / μmol L ⁻¹)	<input type="text"/>		
Renal failure	<input type="checkbox"/>		
<input type="button" value="RESET"/>		<input type="button" value="CALCULATE"/>	

The MG risk score for 6 month mortality from hospitalization with AMI comprised six of the eight GRACE variables:

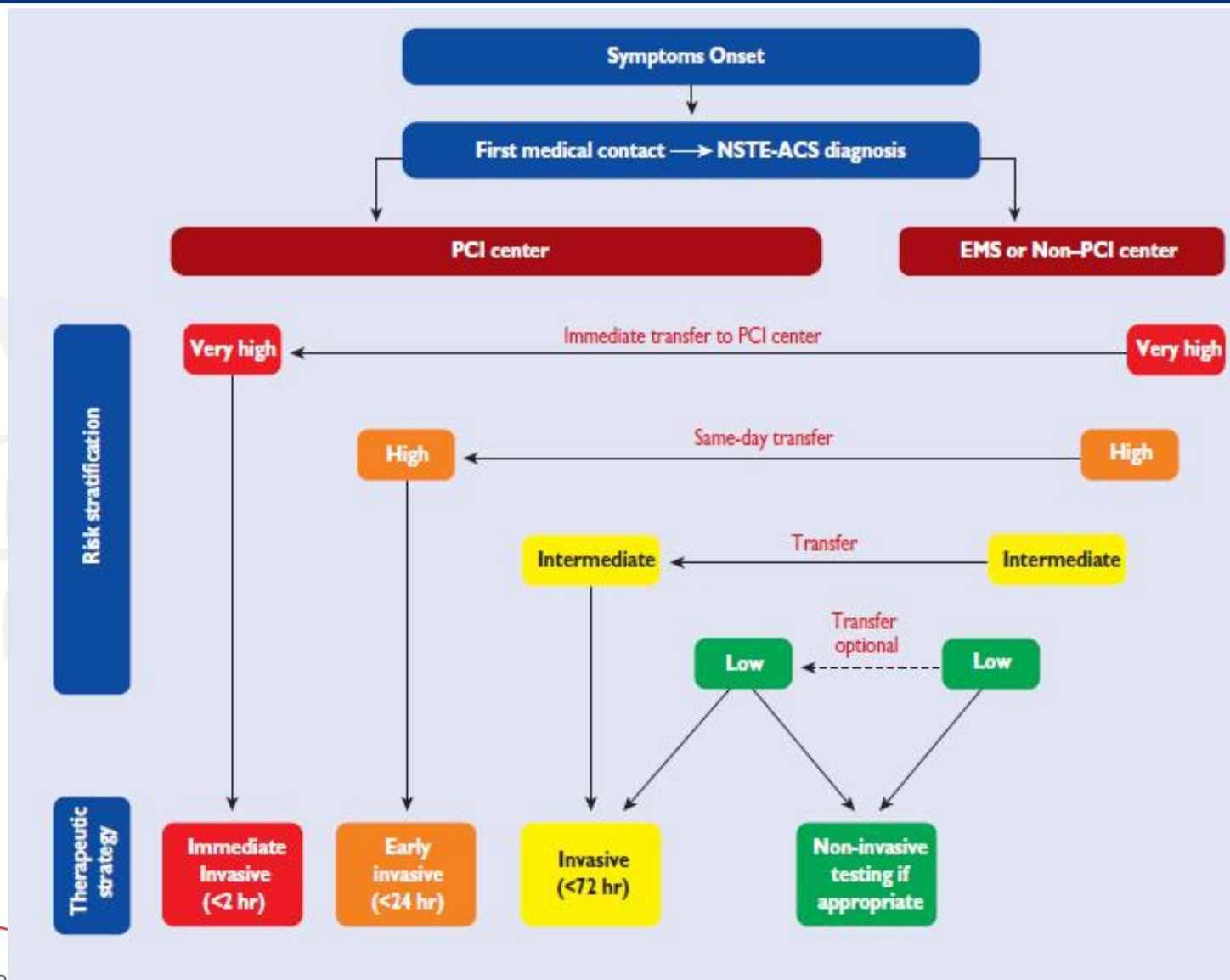
- age
- admission systolic blood pressure
- heart rate
- electrocardiographic ST segment deviation
- cardiac arrest
- elevated cardiac enzymes (defined as a cardiac troponin concentration >0.05 ng/ml).

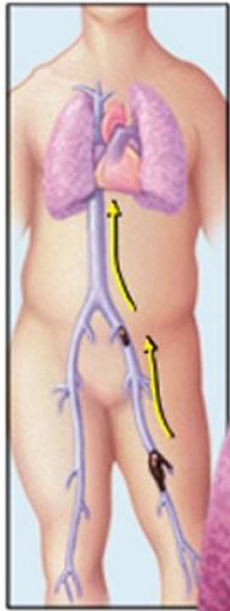


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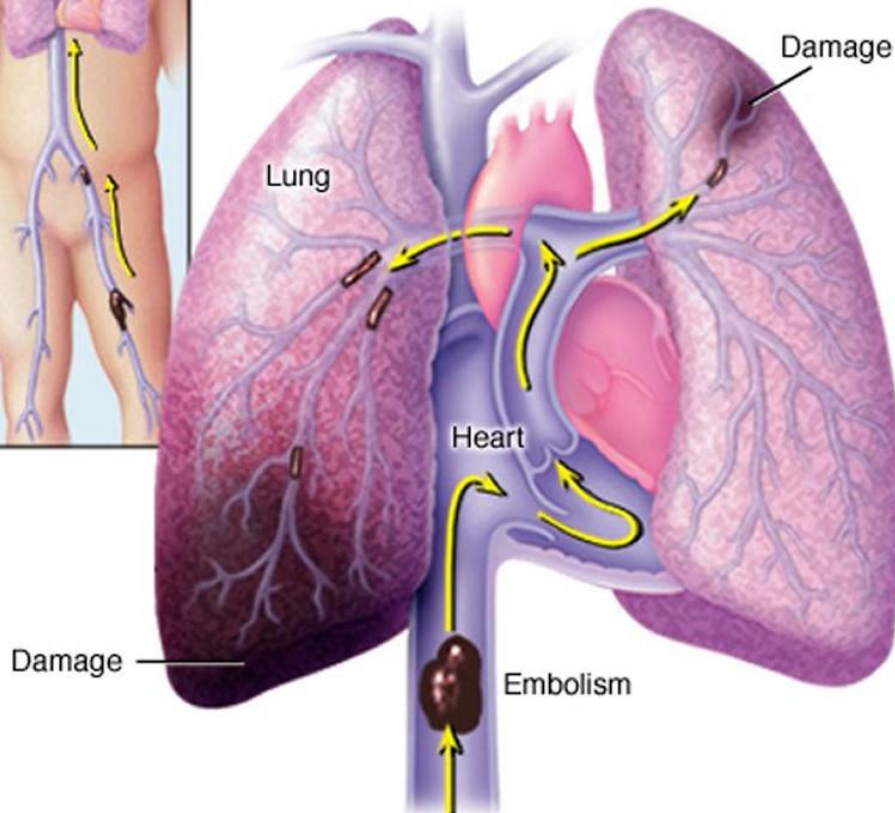
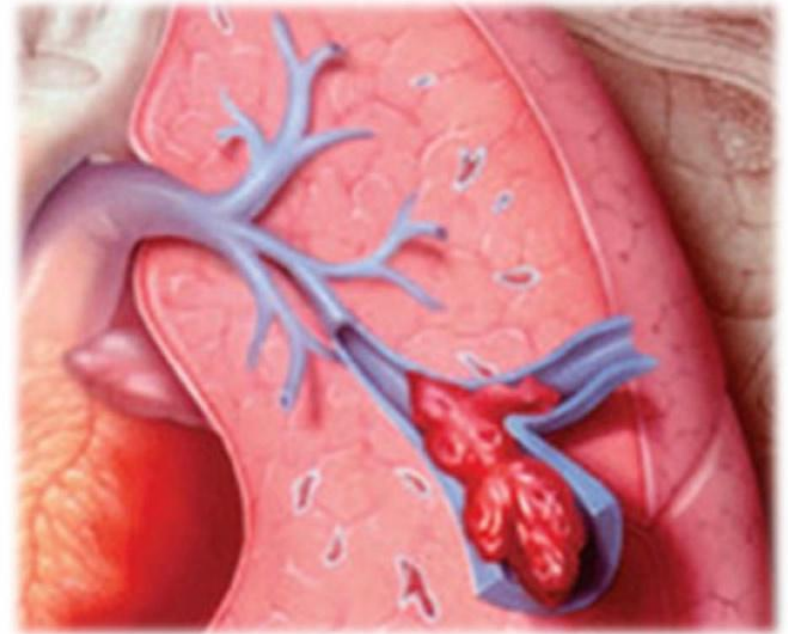
Simms AD et al. Heart 2013





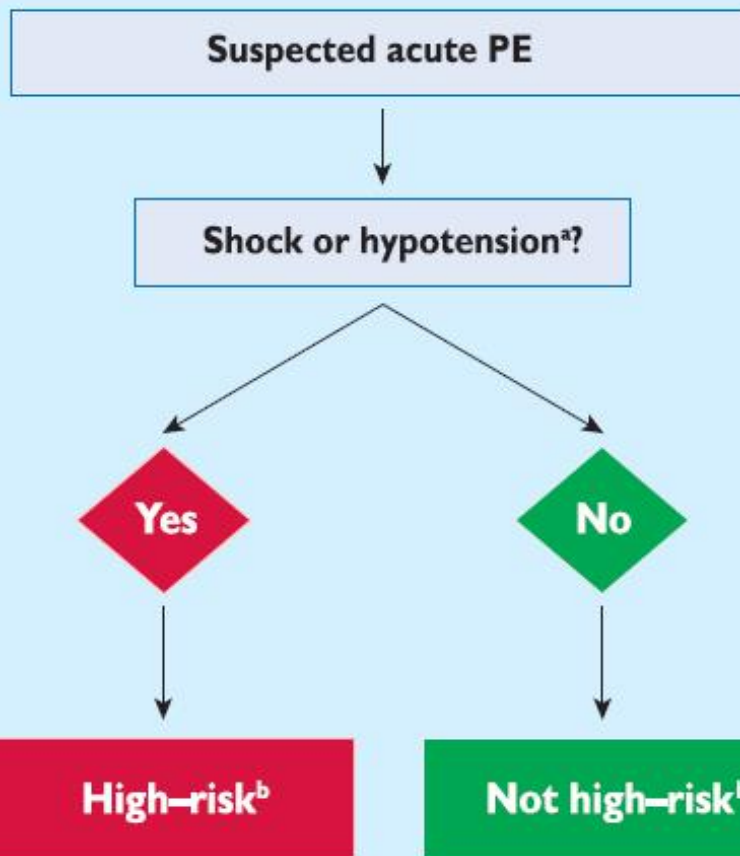


Pulmonary Embolism



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PE = pulmonary embolism.

^aDefined as systolic blood pressure <90 mm Hg, or a systolic pressure drop by ≥ 40 mm Hg, for >15 minutes, if not caused by new-onset arrhythmia, hypovolaemia, or sepsis.

^bBased on the estimated PE-related in-hospital or 30-day mortality.



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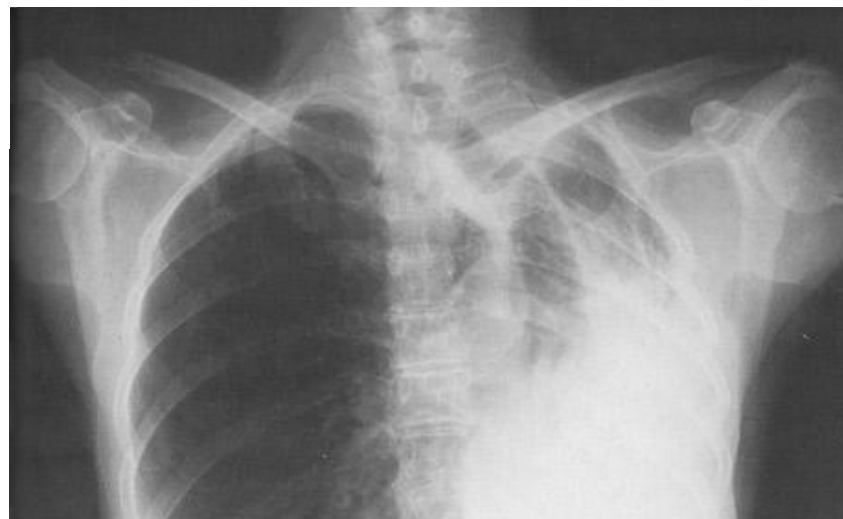
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Early mortality risk		Risk parameters and scores			
		Shock or hypotension	PESI class III-V or sPESI $\geq 1^a$	Signs of RV dysfunction on an imaging test ^b	Cardiac laboratory biomarkers ^c
High		+	(+) ^d	+	(+) ^d
Intermediate	Intermediate-high	-	+	Both positive	
	Intermediate-low	-	+	Either one (or none) positive ^e	
Low		-	-	Assessment optional; if assessed, both negative ^e	

Recommendations	Class ^a	Level ^b
Initial risk stratification of suspected or confirmed PE—based on the presence of shock or persistent hypotension—is recommended to identify patients at high risk of early mortality.	I	B
In patients not at high risk, use of a validated clinical risk prediction score, preferably the PESI or sPESI, should be considered to distinguish between low- and intermediate-risk PE.	IIa	B
In patients at intermediate risk, assessment of the right ventricle with echocardiography or CT, and of myocardial injury using a laboratory biomarker, should be considered for further risk stratification.	IIa	B



Clinical presentation



Feature	PE confirmed (n = 1880)	PE not confirmed (n = 528)
Dyspnoea	50%	51%
Pleuritic chest pain	39%	28%
Cough	23%	23%
Substernal chest pain	15%	17%
Fever	10%	10%
Haemoptysis	8%	4%
Syncope	6%	6%
Unilateral leg pain	6%	5%
Signs of DVT (unilateral extremity swelling)	24%	18%

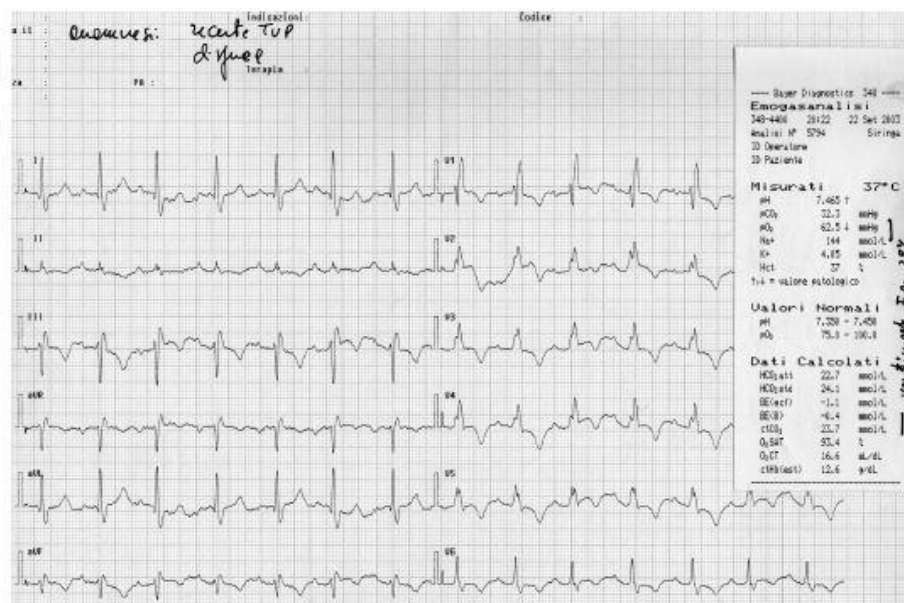


Figura 1



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ESC Guidelines

Items	Clinical decision rule points	
	Original version ⁹³	Simplified version ¹⁶⁷
Wells rule		
Previous PE or DVT	1.5	1
Heart rate ≥ 100 b.p.m.	1.5	1
Surgery or immobilization within the past four weeks	1.5	1
Haemoptysis	1	1
Active cancer	1	1
Clinical signs of DVT	3	1
Alternative diagnosis less likely than PE	3	1
Clinical probability		
Three-level score		
Low	0–1	N/A
Intermediate	2–6	N/A
High	≥ 7	N/A
Two-level score		
PE unlikely	0–4	0–1
PE likely	≥ 5	≥ 2

Revised Geneva score	Original version ⁹³	Simplified version ¹⁶⁸
Previous PE or DVT	3	1
Heart rate 75–94 b.p.m. ≥ 95 b.p.m.	3 5	1 2
Surgery or fracture within the past month	2	1
Haemoptysis	2	1
Active cancer	2	1
Unilateral lower limb pain	3	1
Pain on lower limb deep venous palpation and unilateral oedema	4	1
Age > 65 years	1	1
Clinical probability		
Three-level score		
Low	0–3	0–1
Intermediate	4–10	2–4
High	≥ 11	≥ 5
Two-level score		
PE unlikely	0–5	0–2
PE likely	≥ 6	≥ 3

USE OF THE D-DIMER FOR DETECTING PULMONARY EMBOLISM IN THE EMERGENCY DEPARTMENT

Table 4. Imaging Results of Patients With False-Negative D-dimer

Imaging Results
Small segmental PEs
Acute subsegmental PE
Solitary small PE
Segmental to subsegmental PE
Small segmental PE
A new nonocclusive thrombus in the right distal main pulmonary artery extending into several segmental branches
Segmental to subsegmental pulmonary thromboembolic disease in the right upper and right middle lobes
Possible bilateral nonocclusive segmental/subsegmental pulmonary thromboembolism
High probability VQ scan

Table 3. Bivariate Characteristics of Patients With False-Negative D-dimer

Characteristic	n (%)
Male	7 (78)
Cancer	0 (0)
History of PE/DVT	4 (44)
Pregnant	0 (0)
COPD	1 (11)
Recent surgery	1 (11)
Exogenous estrogen	0 (0)
Shortness of breath	5 (56)
Cough	4 (44)
Hemoptysis	1 (11)
Syncope	1 (11)
Fever > 100.4°F	0 (0)
Unilateral leg swelling	0 (0)
Current smoker	2 (22)
Positive DVT US	0 (0)



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Glober N et al. J Emerg Med. 2018

Suspected PE with shock or hypotension

CT angiography immediately available

No^a

Yes

Echocardiography

RV overload^b

No

Yes

CT angiography
available
and
patient stabilized

CT angiography

positive

negative

Search for other causes
of haemodynamic instability

PE-specific treatment:
primary reperfusion^c

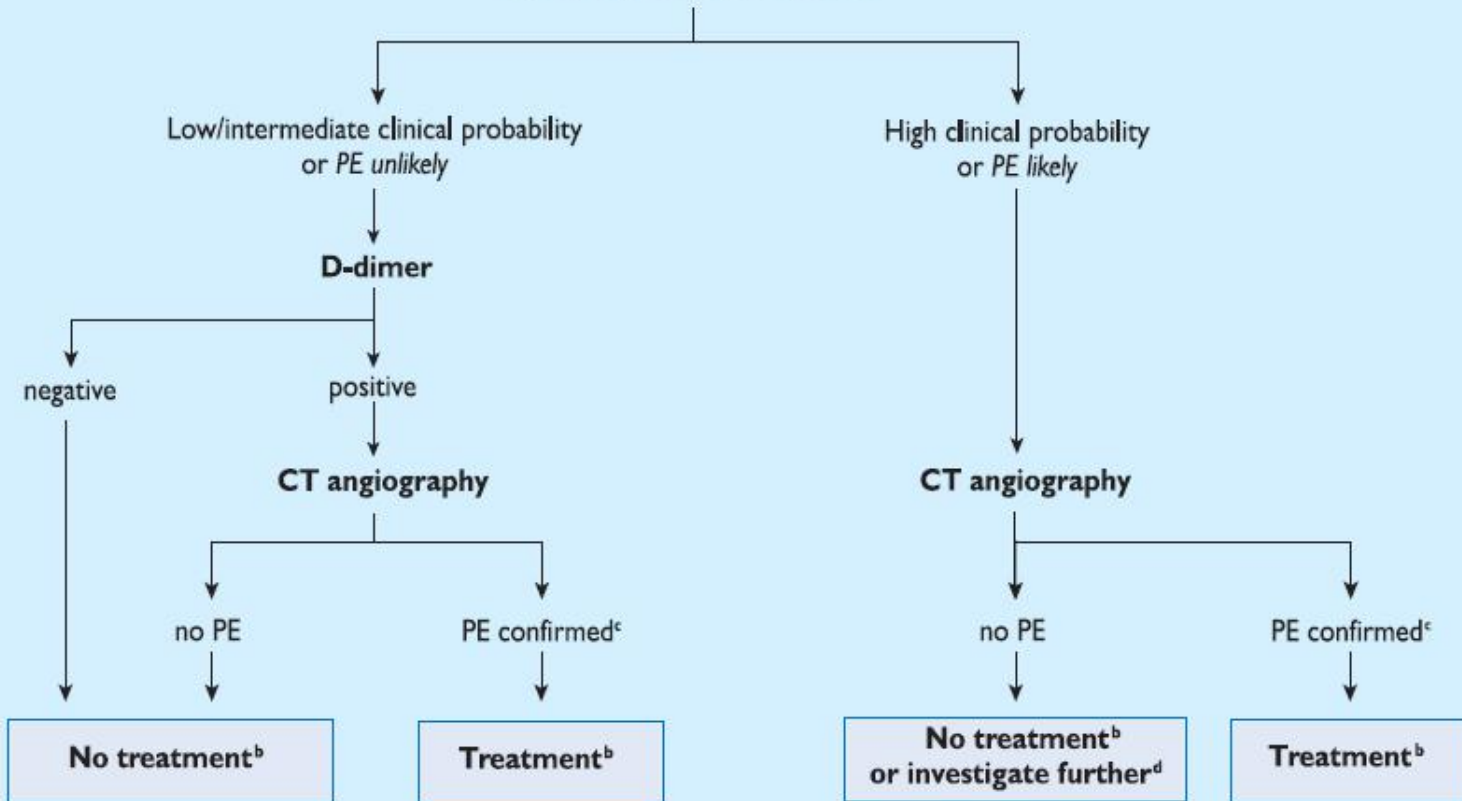
Search for other causes
of haemodynamic instability

No other test available^b
or patient unstable

Suspected PE without shock or hypotension

Assess clinical probability of PE

Clinical judgment or prediction rule^a



Pulmonary embolism severity index (PESI)

Parameter	Original version ²¹⁴	Simplified version ²¹⁸
Age	Age in years	1 point (if age >80 years)
Male sex	+10 points	–
Cancer	+30 points	1 point
Chronic heart failure	+10 points	1 point
Chronic pulmonary disease	+10 points	
Pulse rate ≥ 110 b.p.m.	+20 points	1 point
Systolic blood pressure <100 mm Hg	+30 points	1 point
Respiratory rate >30 breaths per minute	+20 points	–
Temperature <36 °C	+20 points	–
Altered mental status	+60 points	–
Arterial oxyhaemoglobin saturation <90%	+20 points	1 point
Risk strata^a		
	Class I: ≤ 65 points very low 30-day mortality risk (0–1.6%) Class II: 66–85 points low mortality risk (1.7–3.5%) Class III: 86–105 points moderate mortality risk (3.2–7.1%) Class IV: 106–125 points high mortality risk (4.0–11.4%) Class V: >125 points very high mortality risk (10.0–24.5%)	0 points = 30-day mortality risk 1.0% (95% CI 0.0%–2.1%) ≥ 1 point(s) = 30-day mortality risk 10.9% (95% CI 8.5%–13.2%)



Il PESI è lo score più ampiamente convalidato fino ad oggi nella valutazione prognostica dei pz con EP acuta

Clinical suspicion of PE

Shock / hypotension?

Yes

No

**Diagnostic algorithm
as in Figure 3**

**Diagnostic algorithm
as in Figure 4**

PE confirmed

**Assess clinical risk
(PESI or sPESI)**

PESI class III-IV
or sPESI ≥ 1

PESI class I-II
or sPESI = 0

Intermediate risk

Consider further
risk stratification

**RV function (echo or CT)^a
Laboratory testing^b**

Both positive

One positive
or both negative

High risk

Intermediate-high risk

Intermediate-low risk

Low risk^c

Primary reperfusion

**A/C; monitoring;
consider rescue
reperfusion^d**

A/C; hospitalization^e

**A/C; consider early
discharge and home
treatment, if feasible^f**

Pulmonary embolism rule-out criteria (PERC) rule in European patients with low implicit clinical probability (PERCEPIC): a multicentre, prospective, observational study

Pulmonary Embolism Rule-Out Criteria

Variable

Age <50 years

Pulse <100 beats per minute

SaO₂ ≥95% on room air

No hemoptysis

No exogenous estrogen use

No prior venous thromboembolism

No surgery or trauma requiring hospitalization within the past 4 weeks

No unilateral leg swelling



To limit the use of unnecessary, costly, time-consuming, and potentially harmful investigations in patients suspected of pulmonary embolism, Kline and colleagues developed the pulmonary embolism rule-out criteria (PERC), a clinical decision rule, based on eight parameters easily available at initial emergency department assessment.



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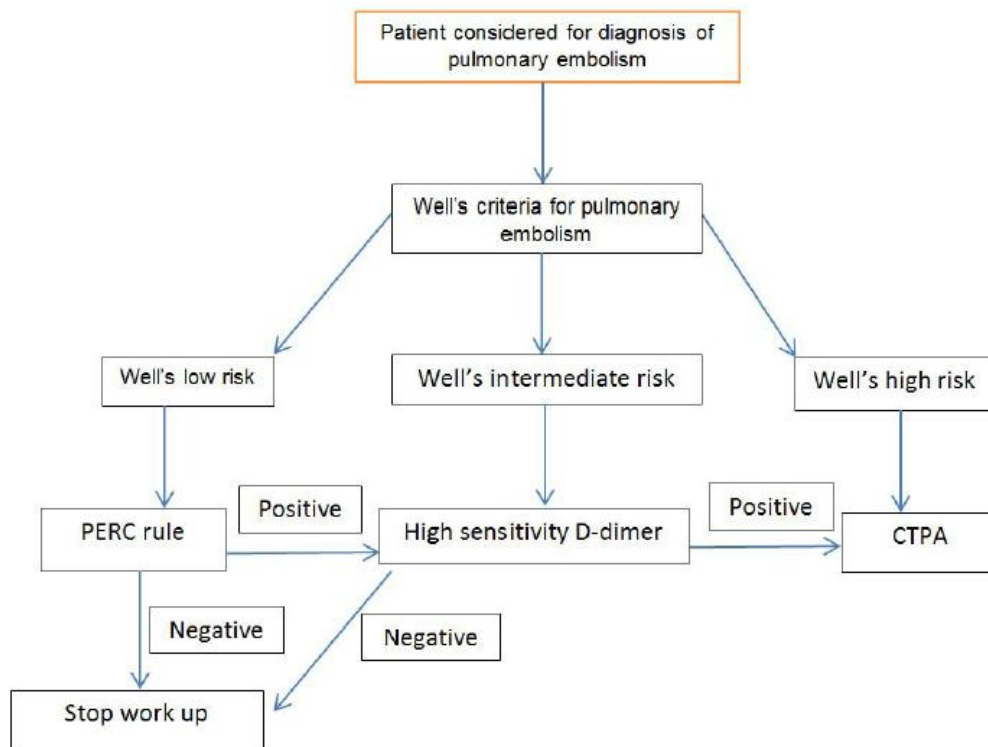
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Lancet Haematol 2017;



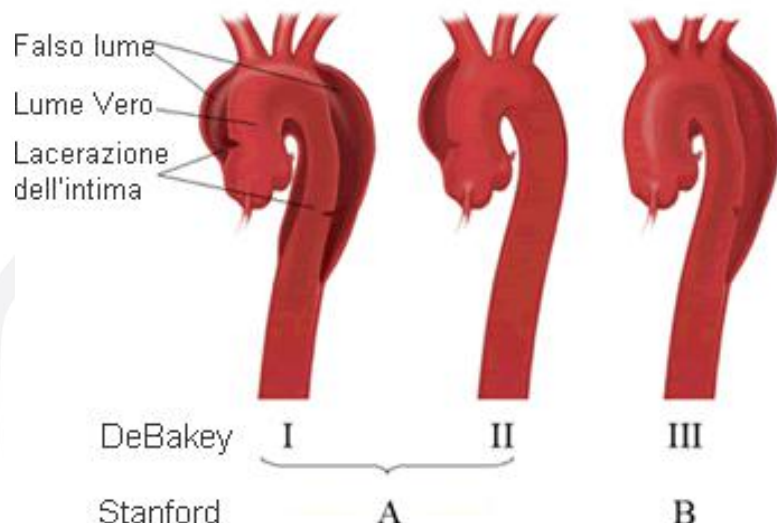
Knowledge Translation of the PERC Rule for Suspected Pulmonary Embolism: A Blueprint for Reducing the Number of CT Pulmonary Angiograms



Results: CTPA declined from 1,033 scans for 98,028 annual visits (10.53 per 1,000 patient visits (95% CI [9.9-11.2])) to 892 scans for 101,172 annual visits (8.81 per 1,000 patient visits (95% CI [8.3-9.4]) $p < 0.001$. The absolute reduction in PACT ordered was 1.72 per 1,000 visits (a 16% reduction). Patient characteristics were similar for both periods.

AORTIC DISSECTION

Anatomia e Classificazione della Dissezione Aortica



	STANFORD A	STANFORD B
Dolore toracico	80%	70%
Dolore dorsale	40%	70%
Dolore migrante	85%	85%
Insufficienza aortica moderato-severa	40 – 75%	-----
Tamponamento cardiaco	10 – 20%	-----
Sindrome coronarica acuta	10 – 15%	10%
Scompenso cardiaco	10%	5%
Versamento pleurico	15%	20%
Sincope	15%	5%
Deficit neurologici	10%	5%
Insufficienza renale acuta	20%	10%
Ischemia arti inferiori	10%	10%

Dati clinici utili a stabilire la probabilità "a priori" di SAA

Condizioni ad alto rischio	Tipo di dolore ad alto rischio	Caratteristiche cliniche ad alto rischio
<ul style="list-style-type: none"> ✓ S. Marfan (o altra connettivopatia) ✓ Anamnesi familiare positiva per malattia aortica ✓ Malattia aortica nota ✓ Aneurisma aorta toracica noto ✓ Pregressa procedura invasiva coinvolgente l'aorta (inclusa chirurgia cardiaca) 	<p>Dolore <u>toracico/addominale/dorsale</u> con una delle seguenti caratteristiche:</p> <ul style="list-style-type: none"> - Inizio improvviso - Intensità severa - Di tipo "squarciante" 	<p>Evidenza di deficit <u>perfusivo</u> per:</p> <ul style="list-style-type: none"> - <u>Iposfigmia</u> - Differenza di PA sistolica in diversi punti di misurazione - Deficit focale neurologico - Soffio diastolico aortico di nuova insorgenza - Ipotensione/shock



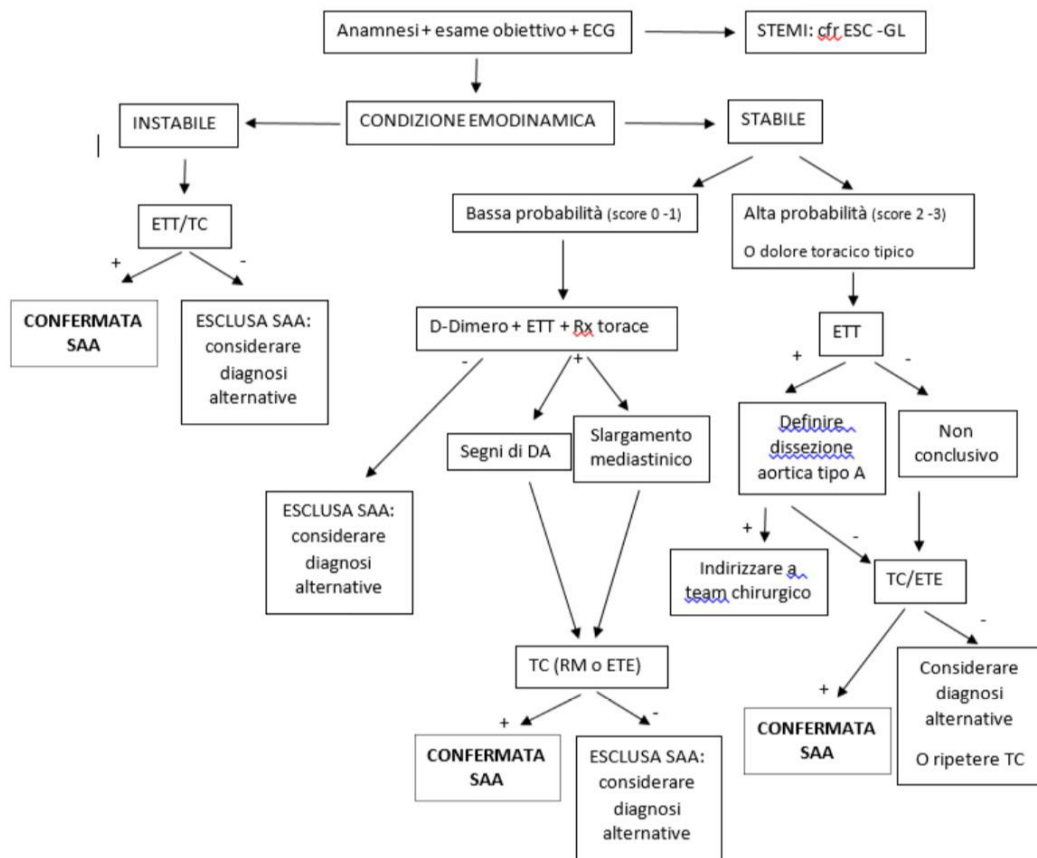
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DOLORE TORACICO ACUTO



Raccomandazione	Classe	Livello
Valutazione anamnestica e clinica		
Accurata raccolta dell'anamnesi e delle caratteristiche clinico-sembiotologiche	I	B
Test di laboratorio		
L'interpretazione dei biomarkers non deve essere disgiunta dalla valutazione di probabilità clinica pre-test	IIa	C
In caso di bassa probabilità clinica, la negatività del DDimero deve essere considerata diagnostica nella esclusione della SAA	IIa	B
In caso di probabilità clinica intermedia e positività del DDimero deve approfondito l'iter diagnostico con test di imaging	IIa	B
In caso di elevata probabilità di DA la valutazione del DDimero non è indicata	III	C
Imaging		
ETT è raccomandato come valutazione iniziale	I	C
In caso di pz instabile con sospetta SAA, è raccomandato eseguire (in relazione a disponibilità della metodica ed esperienza dell'operatore):		
ETE	I	C
TC	I	C
In caso di pz stabile con sospetta SAA le seguenti metodiche sono raccomandate (o dovrebbero essere considerate) in relazione alla disponibilità della metodica ed all'esperienza dell'operatore		
TC	I	C
RM	I	C
ETE	IIa	C
In caso di iniziale negatività dei test di imaging e persistenza di sospetto clinico di SAA, è raccomandato ripetere CT o RM	I	C
RX torace può essere considerato in caso di bassa probabilità di SAA	IIb	C
In caso di DA di tipo B non complicata, in terapia medica, è raccomandato ripetere TC o RM nei primi giorni	I	C



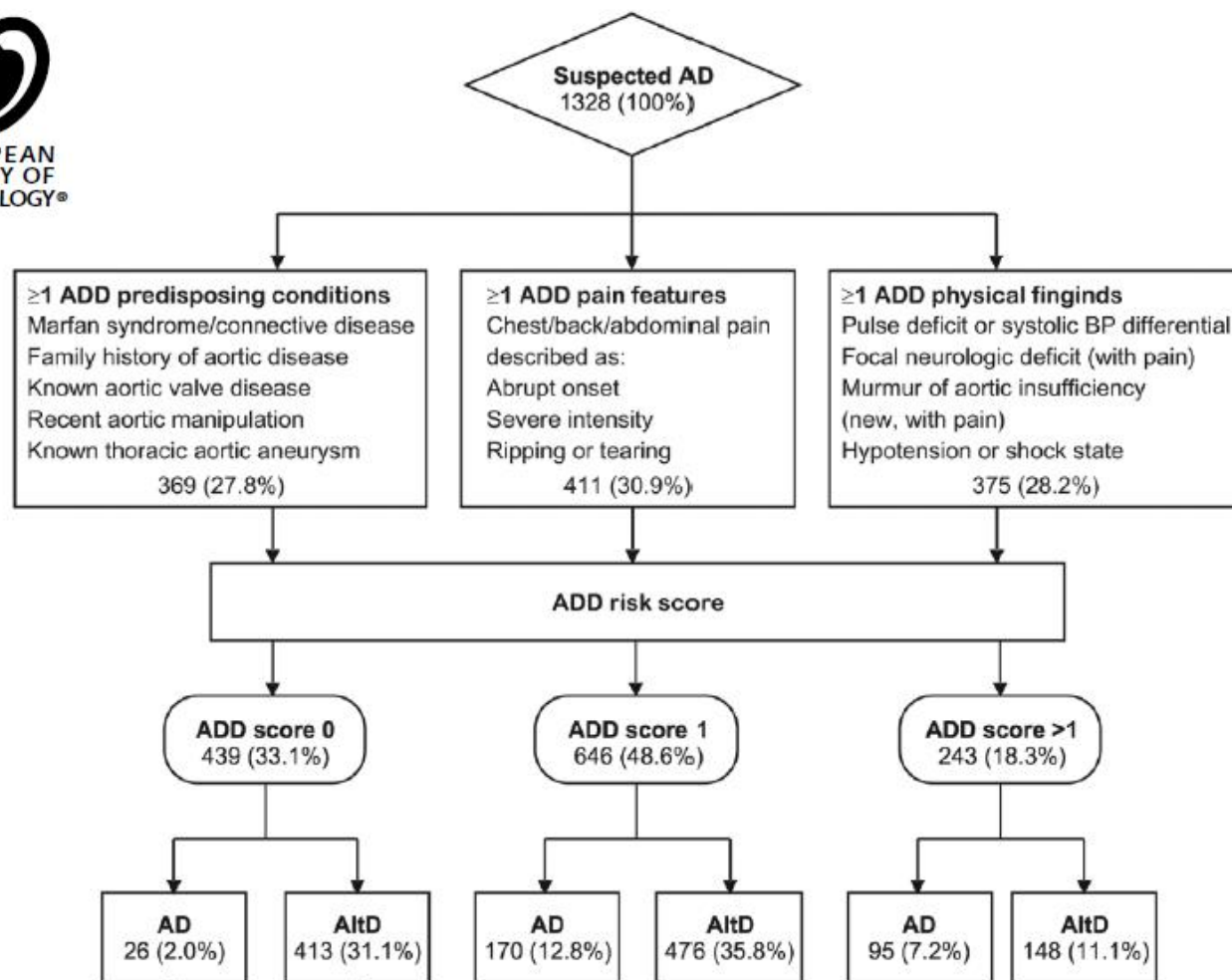
Aortic pain with immediate onset, a tearing or ripping character, or both; mediastinal widening, aortic widening, or both on chest radiography; and pulse differentials, blood pressure differentials, or both ($P < .001$ for all) were identified as independent predictors of acute aortic dissection. Probability of dissection was low with absence of all 3 variables (7%), intermediate with isolated findings of aortic pain or mediastinal widening (31% and 39%, respectively), and high with isolated pulse or blood pressure differentials or any combination of the 3 variables ($\geq 83\%$).

Table 3. Risk for Types A and B Acute Aortic Dissection According to 3 Clinical Predictors

Variable	No. (%) of Patients		Probability of Dissection, %
	Dissection (n = 128)	No Dissection (n = 122)	
No sign present	5 (4)	65 (53)	7
Aortic pain alone	13 (10)	29 (24)	31
Mediastinal widening, aortic widening, or both alone	11 (9)	17 (14)	39
Aortic pain + mediastinal widening, aortic widening, or both	50 (39)	10 (8)	83
Pulse differentials, blood pressure differentials, or both alone	2 (2)	0	100
Aortic pain + pulse differentials, blood pressure differentials, or both	11 (9)	1 (1)	92
Mediastinal widening + pulse differentials, blood pressure differentials, or both	2 (2)	0	100
Aortic pain + mediastinal widening, aortic widening, or both + pulse differentials, blood pressure differentials, or both	34 (27)	0	100



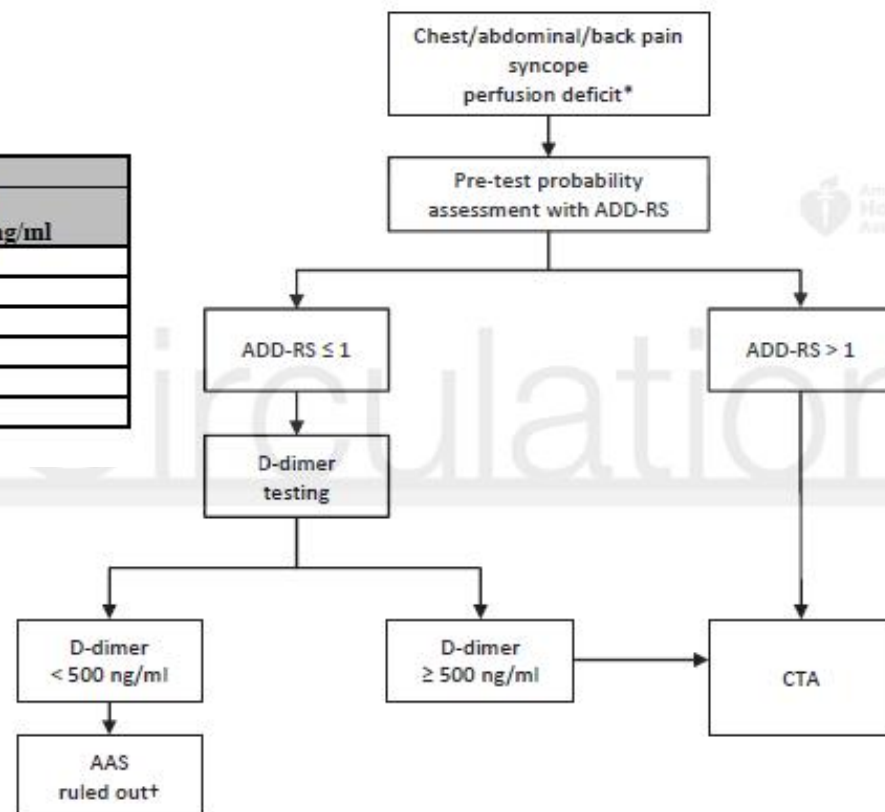
Diagnostic performance of the aortic dissection detection risk score in patients with suspected acute aortic dissection



Diagnostic Accuracy of the Aortic Dissection Detection Risk Score Plus D-Dimer for Acute Aortic Syndromes: The ADvISED Prospective Multicenter Study

Diagnostic variables	Diagnostic strategy	
	ADD risk score = 0 plus D-dimer < 500 ng/ml	ADD risk score ≤ 1 plus D-dimer < 500 ng/ml
Sensitivity	99.6% (97.7-100%)	98.8% (96.4-99.7%)
Specificity	18.2% (16.4-20.2%)	57.3% (54.9-59.7%)
PPV	15.4% (13.7-17.3%)	25.8% (23-28.7%)
LR+	1.22 (1.19-1.25)	2.31 (2.18-2.45)
NPV	99.7% (98.1-100%)	99.7% (99.1-99.9%)
LR-	0.02 (0.003-0.16)	0.02 (0.01-0.07)

Variables are presented as percent and 95% confidence interval (in brackets).



CONCLUSIONI

Emerging evidence worthy of mention illustrates that a physician Gestalt may perform better than sole reliance on clinical scoring systems. This new body of research illustrates the German concept of Gestalt theory, a philosophical and psychiatric principle in which the process is taken into consideration versus the content—in other words, the whole is not the sum of its parts, but greater than the sum of its parts. A physician's clinical judgment should not be replaced by clinical scoring systems, but should instead be used in conjunction with evidence-based validated systems when deciding the most likely diagnosis for a patient.





- Dottore, comincio a dubitare delle sue diagnosi...

Grazie



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