

E-OUAL EMERGENCY OUALITY NETWORK

Clinical Policy: Suspected Acute Venous Thromboembolic Disease





Presenters



Stephen J. Wolf, MD



American College of Emergency Physicians[®]

Clinical Policy: Critical Issues in the Evaluation and Management of Adult Patients Presenting to the Emergency Department With Suspected Acute Venous Thromboembolic Disease

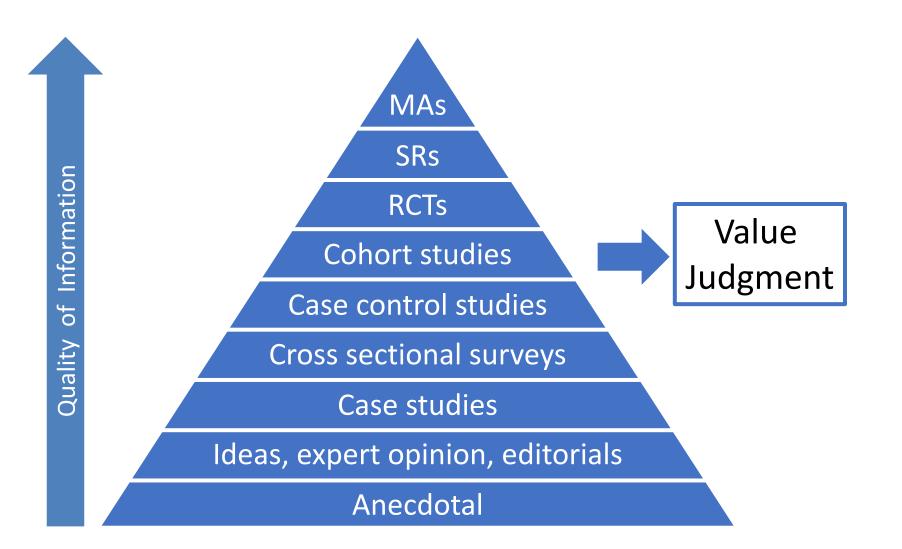


From the American College of Emergency Physicians Clinical Policies Subcommittee (Writing Committee) on Thromboembolic Disease:
Stephen J. Wolf, MD (Subcommittee Chair; Committee Co-Chair)
Sigrid A. Hahn, MD, MPH
Lauren M. Nentwich, MD
Ali S. Raja, MD, MBA, MPH
Scott M. Silvers, MD
Michael D. Brown, MD, MSc (Committee Co-Chair)

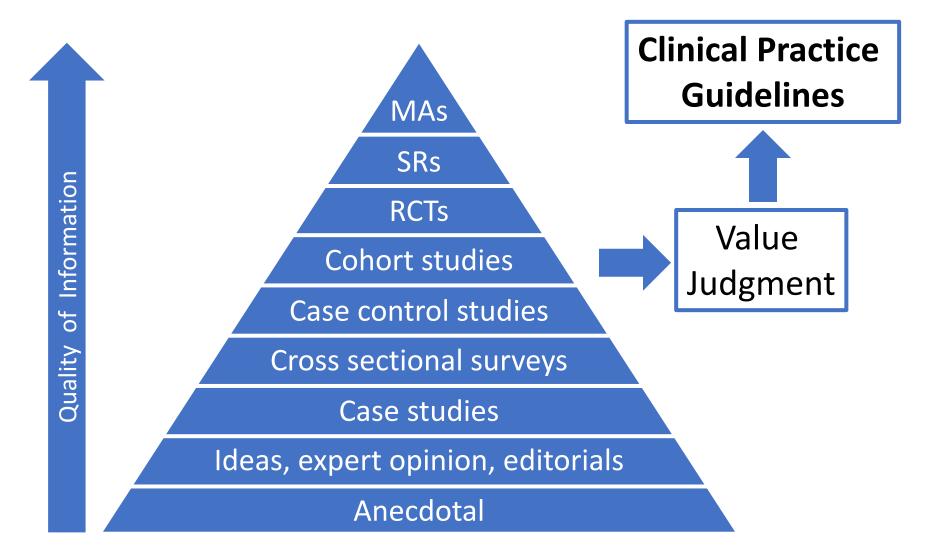
> Stephen J Wolf, MD, FACEP Department of Emergency Medicine Denver Health Medical Center University of Colorado School of Medicine



Hierarchy of Evidence



Value-Based Evidence



Clinical Policies Committee



IOM Standards for Trustworthiness

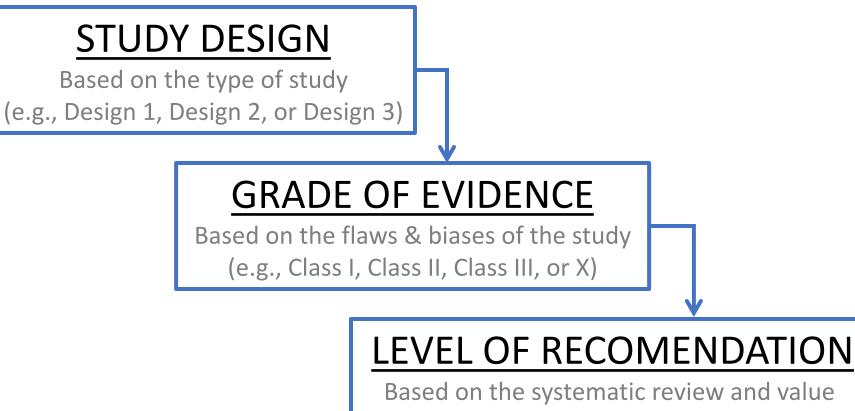
Establishing Transparency Management of Conflicts of Interest Group Composition Systematic Review Intersection **Evidence Foundations for and Rating Strength of Recommendations Articulation of Recommendations External Review** Updating

ACEP's Process

Topic selection Subcommittee appointed Critical questions developed Literature search & grading Subcommittee writing **Oversight committee input** Expert review & open comment **Board approval & dissemination**

١

Getting from Point A to B



judgments (e.g., Level A, Level B, or Level C)

How trustworthy are ACEP Clinical Policies for imaging recommendations? Clinical Policy: Critical Issues in the Evaluation and Management of Adult Patients Presenting to the Emergency Department With Suspected Acute Venous Thromboembolic Disease



From the American College of Emergency Physicians Clinical Policies Subcommittee (Writing Committee) on Thromboembolic Disease:
Stephen J. Wolf, MD (Subcommittee Chair; Committee Co-Chair)
Sigrid A. Hahn, MD, MPH
Lauren M. Nentwich, MD
Ali S. Raja, MD, MBA, MPH
Scott M. Silvers, MD
Michael D. Brown, MD, MSc (Committee Co-Chair)

> Acep.org/ClinicalPolicies @ACEPNation

Critical Questions: VTE

Diagnostic Questions

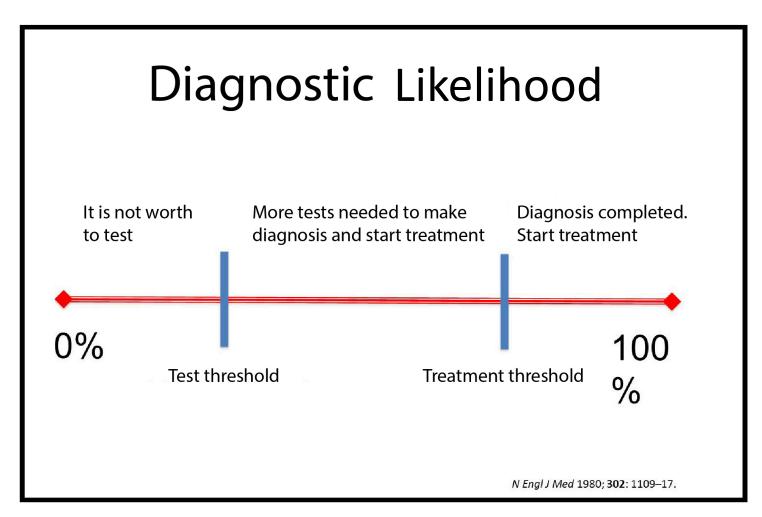
- In adult patients with suspected acute PE, can a clinical prediction rule be used to identify a group of patients at very low risk for the diagnosis of PE for whom no additional diagnostic workup is required?
- In adult patients with low to intermediate pretest probability for acute PE, does a negative age adjusted D-dimer result identify a group of patients at very low risk for the diagnosis of PE for whom no additional diagnostic workup is required?

Critical Questions: VTE

Management Questions

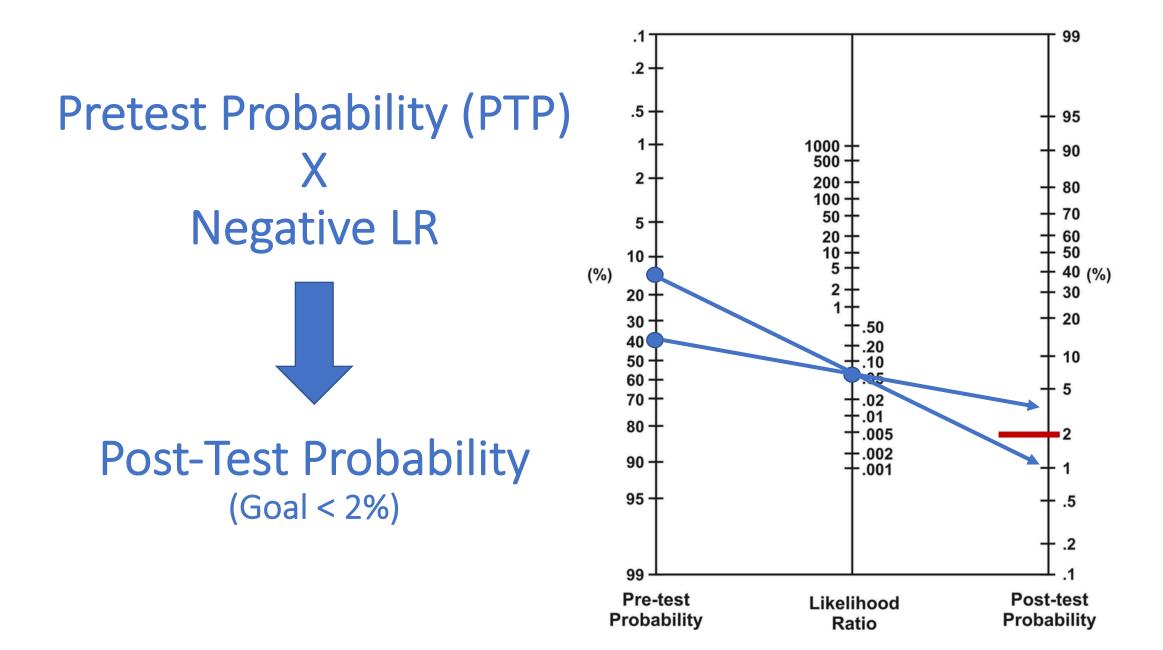
- In adult patients with subsegmental PE, is it safe to withhold anticoagulation?
- In adult patients diagnosed with acute PE, is initiation of anticoagulation and discharge from the ED safe?
- In adult patients diagnosed with acute lower-extremity DVT who are discharged from the ED, is treatment with a NOAC safe and effective compared with treatment with LMWH and VKA?

EMB & Stewardship



Testing Threshold for VTE

In consideration of the cost of evaluation, the risk of false positives, and the risk of complications related to testing, studies have supported using a predefined posttest probability threshold of less than 2.0% to exclude the diagnosis of VTE. Why 2%? My colleagues always say they want to miss the bad stuff less than 1% of the time



Critical Question

In adult patients with suspected acute PE, can a **clinical prediction rule** be used to identify a group of patients at very low risk for the diagnosis of PE for whom **no additional diagnostic workup** is required?

> 47 identified > 19 graded 4 Class II, 4 Class III, 11 Class X

Pulmonary Embolism Rule-out Criteria (PERC)

- 1. Age < 50 year
- 2. Pulse Rate < 100 beats/min
- 3. SaO2 > 94% (at sea level)
- 4. No Recent Trauma or Surgery
- 5. No Unilateral Leg Swelling
- 6. No Previous PE or DVT
- 7. No Hormone Use
- 8. No Hemoptysis

PERC Performance

Clinical Policy

Table 1. PERC performance.

							PERC Performance				
Study Cohorts	Class	Pretest Probability	N	PE (%)	PERC Determination	Sensitivity (95% CI), %	Specificity (95% CI), %	Negative LR (95% CI)	Posttest VTE (%) (95% Cl)		
Low-Risk Cohorts	;										
Kline et al ²⁴	II	Low	1,427	114 (8)	Prospective	96 (90-99)	27 (25-30)	0.16 (0.07-0.38)	1.4 (0.4-3.2)		
Kline et al ²⁶	II	Low	5,425	163 (3)	Prospective	97 (96-99)	22 (21-23)	0.12 (0.07-1.19)	1.3 (0.8-1.9)		
Hugli et al ²⁷	II	Low	587	57 (10)	Retrospective	79 (67-88)	33 (29-37)	0.63 (0.04-1.06)	6.4 (3.7-6.8)		
Wolf et al ³¹	III	Low	60	1 (2)	Retrospective	100 (25-100)	22 (12-35)	0 (*)	0 (0-24.7)		
Penaloza et al ³⁰	III	Low	399	26 (7)	Retrospective	100 (99-100)	9 (6-11)	0 (*)	0 (0-5)		
Undifferentiated-F	Risk Coh	orts									
Kline et al ²⁶	II	All	8,138	561 (7)	Prospective	96 (94-97)	25 (24-26)	0.17 (0.11-0.25)	1.0 (0.6-1.6)		
Hugli et al ²⁷	II	All	1,675	357 (21)	Retrospective	97 (94-98)	16 (14-18)	0.21 (0.12-0.37)	5.4 (3.1-9.3)		
Wolf et al ³¹	III	All	120	16 (12)	Retrospective	100 (79-100)	16 (10-24)	0 (*)	0 (0-17.6)		
Crichlow et al ²⁹	III	All	152	18 (12)	Prospective	100 (78-100)	10 (6-17)	0 (*)	0 (0-23.2)		
Penaloza et al ³⁰	III	All	959	286 (30)	Retrospective	99 (97-100)	10 (8-13)	0.13 (0.05-0.36)	5.4 (1.7-12.5)		
Bozarth et al ²⁸	Ш	All	719	32 (5)	Retrospective	97 (94-100)	12 (10-15)	0.26 (0.04-1.82)	1.2 (0-6.5)		

CI, confidence interval; *LR*, likelihood ratio; *PE*, pulmonary embolism; *PERC*, pulmonary embolism rule-out criteria; *VTE*, venous thromboembolism; *Undefined given 100% sensitivity

Critical Question

In adult patients with suspected acute PE, can a clinical prediction rule be used to identify a group of patients at very low risk for the diagnosis of PE for whom no additional diagnostic workup is required?

Level B Recommendation

For patients who are at low risk for acute PE, use the PERC to exclude the diagnosis without further diagnostic testing.

Clinical prediction rules are easy to misapply – where do you see cracks in the evidence translation?

Critical Question

In adult patients with low to intermediate pretest probability for acute PE, does a negative **age-adjusted D-dimer** result identify a group of patients at very low risk for the diagnosis of PE for whom no additional diagnostic workup is required?

> 59 identified > 42 graded 3 Class II, 7 Class III, 32 Class X

Age-Adjusted D-Dimer Goal

Improve diagnostic efficiency Reduce unnecessary testing Reduce test-related complications Steward health care resources

Age-Adjusted D-Dimer

Important note

D-dimer assays are reported as either the concentration of DDU or as FEU, depending on the calibration for the assay. The 2 numeric values are easily convertible because the mass of one FEU equals approximately half of one DDU (ie,1 FEU=2DDU).

Strategies:

Fixed age-adjusted cutoff Incremental age-adjusted cutoff

Study	Class	CPR	PTP	AADD cutoff (µg/L)	CDD Sensitivity (%; 95% Cl)	AADD Sensitivity (%; 95% Cl)	CDD Miss Rate (%; 95% Cl)	AADD Miss Rate (%; 95% Cl)	% Cohort With Negative CDD (95% CI)	% Cohort With Negative AADD (95% CI)
Righini et al ⁴³ *	II	sRGS or Wells	Non-high or unlikely	$Age \times 10^{\dagger}$	NR	NR	1/810 (0.1; 0-0.7)	2/1,141 (0.2; 0-0.6)	28 (27-30)	40 (38-42)
Flores et al ⁴⁵	П	Wells	Non-high	$Age \times 10^{\dagger}$	100 (94-100)	100 (94-100)	0/92 (0; 0-3.9)	0/121 (0; 0-3.0)	28 (23-33)	37 (32-42)
van Es et al ⁴⁴	II	Wells	Unlikely	$Age \times 10^{+}$	99 (99-100)	99 (98-99)	13/2,035 (0.7; 0.4-1.1)	22/2,369 (0.9; 0.6-1.5)	28 (21-37)	33 (25-42)
van Es et al ⁴⁷ *	III	Wells	Unlikely	$Age \times 10^{+}$	NR	NR	1/60 (1.7; 0-8.9)	2/92 (2.2; 0-7.6)	15 (11-18)	22 (18-26)
Gupta et al ⁴⁹	Ш	NR	Any	$Age \times 10^{+}$	100 (94-100)	97 (90-100)	0/72 (0; 0-5.0)	2/165 (1.2; 0.1-4.3)	7 (7-9)	16 (14-19)
Friz et al ⁵⁰	Ш	NR	Any	$Age \times 10^{+}$	100 (97-100)	98 (94-100)	0/8 (0; 0-36.9)	2/28 (7.1; 0.9-23.5)	2 (1-3)	6 (4-8)
Jaconelli et al ⁵²	III	Wells	Unlikely	Age×5 [‡]	95 (86-99)	95 (86-99)	3/859 (0.3; 0.1-1.0)	3/989 (0.3; 0.1-0.9)	65 (62-68)	75 (72-77)
Sharp et al ⁴⁸	III	NR	Any	$Age \times 10^{+}$	98 (96-99)	93 (90-95)	10/16,660 (0.1; 0-0.1)	36/19,584 (0.2; 0.1-0.3)	54 (53-54)	63 (62-64)
Douma et al ⁴⁶	III	Wells	Unlikely	$Age \times 10^{+}$	NR	NR	2/983 (0.2; 0.1-0.7)	7/1,093 (0.6; 0.3-1.3)	46 (43-48)	51 (49-53)
Douma et al ⁴⁶	Ш	RGS	Non-high	$Age \times 10^{+}$	NR	NR	0/561 (0; 0.0-0.7)	2/663 (0.3; 0.1-1.1)	34 (32-37)	40 (38-43)
Sharp et al ⁴⁸	III	NR	Any	1,000 [†]	98 (96-99)	84 (81-87)	10/16,660 (0.1; 0.0-0.1)	80/23,146 (0.3; 0.3-0.4)	54 (53-54)	74 (74-75)
Friz et al ⁵⁰	III	NR	Any	1,000 [†]	100 (97-100)	96 (91-99)	0/8 (0; 0-36.9)	4/61 (6.6; 1.8-15.9)	2 (1-3)	13 (10-16)
Kline et al ⁵¹ * [§]	III	sRGS or Wells	Any	1,000 [†]	94 (88-97)	92 (86-96)	8/152 (5.3; 2-10.1)	10/185 (5.4; 2.6-9.7)	22 (19-26)	27 (24-31)

AADD, age-adjusted D-dimer; CDD, conventional D-dimer; CI, confidence interval; CPR, clinical prediction rule; NR, not reported; PTP, pretest probability; RGS, revised Geneva score; sRGS, simplified revised Geneva score. *Multiple CPRs were used; for simplicity, only results for Wells are presented.

[†]D-dimer value reported in FEUs.

[‡]D-dimer value reported in DDUs;

Study	Class	CPR	РТР	AADD cutoff (µg/L)	CDD Sensitivity (%; 95% Cl)	AADD Sensitivity (%; 95% CI)	CDD Miss Rate (%; 95% Cl)	AADD Miss Rate (%; 95% Cl)	% Cohort With Negative CDD (95% Cl)	% Cohort With Negative AADD (95% CI)
Righini et al ⁴³ *	II	sRGS or Wells	Non-high or unlikely	$Age \times 10^{+}$	NR	NR	1/810 (0.1; 0-0.7)	2/1,141 (0.2; 0-0.6)	28 (27-30)	40 (38-42)
Flores et al ⁴⁵	Ш	Wells	Non-high	$Age \times 10^{\dagger}$	100 (94-100)	100 (94-100)	0/92 (0; 0-3.9)	0/121 (0; 0-3.0)	28 (23-33)	37 (32-42)
van Es et al ⁴⁴	II	Wells	Unlikely	$Age \times 10^{+}$	99 (99-100)	99 (98-99)	13/2,035 (0.7; 0.4-1.1)	22/2,369 (0.9; 0.6-1.5)	28 (21-37)	33 (25-42)
van Es et al ⁴⁷ *	III	Wells	Unlikely	$Age \times 10^{+}$	NR	NR	1/60 (1.7; 0-8.9)	2/92 (2.2; 0-7.6)	15 (11-18)	22 (18-26)
Gupta et al ⁴⁹	III	NR	Any	$Age \times 10^{+}$	100 (94-100)	97 (90-100)	0/72 (0; 0-5.0)	2/165 (1.2; 0.1-4.3)	7 (7-9)	16 (14-19)
Friz et al ⁵⁰	III	NR	Any	$Age \times 10^{+}$	100 (97-100)	98 (94-100)	0/8 (0; 0-36.9)	2/28 (7.1; 0.9-23.5)	2 (1-3)	6 (4-8)
Jaconelli et al ⁵²	III	Wells	Unlikely	Age×5 [‡]	95 (86-99)	95 (86-99)	3/859 (0.3; 0.1-1.0)	3/989 (0.3; 0.1-0.9)	65 (62-68)	75 (72-77)
Sharp et al ⁴⁸	III	NR	Any	$Age \times 10^{+}$	98 (96-99)	93 (90-95)	10/16,660 (0.1; 0-0.1)	36/19,584 (0.2; 0.1-0.3)	54 (53-54)	63 (62-64)
Douma et al ⁴⁶		Wells	Unlikely	$Age \times 10^+$	NR	NR	2/983 (0.2; 0.1-0.7)	7/1,093 (0.6; 0.3-1.3)	46 (43-48)	51 (49-53)
Douma et al ⁴⁶	III	RGS	Non-high	$Age imes 10^+$	NR	NR	0/561 (0; 0.0-0.7)	2/663 (0.3; 0.1-1.1)	34 (32-37)	40 (38-43)
Sharp et al ⁴⁸	Ш	NR	Any	1,000 [†]	98 (96-99)	84 (81-87)	10/16,660 (0.1; 0.0-0.1)	80/23,146 (0.3; 0.3-0.4)	54 (53-54)	74 (74-75)
Friz et al ⁵⁰	III	NR	Any	1,000 [†]	100 (97-100)	96 (91-99)	0/8 (0; 0-36.9)	4/61 (6.6; 1.8-15.9)	2 (1-3)	13 (10-16)
Kline et al ⁵¹ * [§]	III	sRGS or Wells	Any	1,000†	94 (88-97)	92 (86-96)	8/152 (5.3; 2-10.1)	10/185 (5.4; 2.6-9.7)	22 (19-26)	27 (24-31)

AADD, age-adjusted D-dimer; CDD, conventional D-dimer; CI, confidence interval; CPR, clinical prediction rule; NR, not reported; PTP, pretest probability; RGS, revised Geneva score; sRGS, simplified revised Geneva score. *Multiple CPRs were used; for simplicity, only results for Wells are presented.

[†]D-dimer value reported in FEUs.

[‡]D-dimer value reported in DDUs;

Study	Class	CPR	РТР	AADD cutoff (µg/L)	CDD Sensitivity (%; 95% Cl)	AADD Sensitivity (%; 95% Cl)	CDD Miss Rate (%; 95% Cl)	AADD Miss Rate (%; 95% Cl)	% Cohort With Negative CDD (95% CI)	% Cohort With Negative AADD (95% CI)
Righini et al ⁴³ *	II	sRGS or Wells	Non-high or unlikely	$Age \times 10^{+}$	NR	NR	1/810 (0.1; 0-0.7)	2/1,141 (0.2; 0-0.6)	28 (27-30)	40 (38-42)
Flores et al ⁴⁵	П	Wells	Non-high	$Age \times 10^{\dagger}$	100 (94-100)	100 (94-100)	0/92 (0; 0-3.9)	0/121 (0; 0-3.0)	28 (23-33)	37 (32-42)
van Es et al ⁴⁴	II	Wells	Unlikely	$Age \times 10^{+}$	99 (99-100)	99 (98-99)	13/2,035 (0.7; 0.4-1.1)	22/2,369 (0.9; 0.6-1.5)	28 (21-37)	33 (25-42)
van Es et al ⁴⁷ *	III	Wells	Unlikely	$Age \times 10^{+}$	NR	NR	1/60 (1.7; 0-8.9)	2/92 (2.2; 0-7.6)	15 (11-18)	22 (18-26)
Gupta et al ⁴⁹	111	NR	Any	Age $\times 10^{+}$	100 (94-100)	97 (90-100)	0/72 (0; 0-5.0)	2/165 (1.2; 0.1-4.3)	7 (7-9)	16 (14-19)
Friz et al ⁵⁰	III	NR	Any	$Age \times 10^{+}$	100 (97-100)	98 (94-100)	0/8 (0; 0-36.9)	2/28 (7.1; 0.9-23.5)	2 (1-3)	6 (4-8)
Jaconelli et al ⁵²	III	Wells	Unlikely	$Age \times 5^{\ddagger}$	95 (86-99)	95 (86-99)	3/859 (0.3; 0.1-1.0)	3/989 (0.3; 0.1-0.9)	65 (62-68)	75 (72-77)
Sharp et al ⁴⁸	III	NR	Any	$Age \times 10^{+}$	98 (96-99)	93 (90-95)	10/16,660 (0.1; 0-0.1)	36/19,584 (0.2; 0.1-0.3)	54 (53-54)	63 (62-64)
Douma et al ⁴⁶	III	Wells	Unlikely	$Age \times 10^{+}$	NR	NR	2/983 (0.2; 0.1-0.7)	7/1,093 (0.6; 0.3-1.3)	46 (43-48)	51 (49-53)
Douma et al ⁴⁶	III	RGS	Non-high	$Age \times 10^{+}$	NR	NR	0/561 (0; 0.0-0.7)	2/663 (0.3; 0.1-1.1)	34 (32-37)	40 (38-43)
Sharp et al ⁴⁸	III	NR	Any	1,000†	98 (96-99)	84 (81-87)	10/16,660 (0.1; 0.0-0.1)	80/23,146 (0.3; 0.3-0.4)	54 (53-54)	74 (74-75)
Friz et al ⁵⁰		NR	Any	1,000 [†]	100 (97-100)	96 (91-99)	0/8 (0; 0-36.9)	4/61 (6.6; 1.8-15.9)	2 (1-3)	13 (10-16)
Kline et al ⁵¹ * [§]		sRGS or Wells	Any	1,000†	94 (88-97)	92 (86-96)	8/152 (5.3; 2-10.1)	10/185 (5.4; 2.6-9.7)	22 (19-26)	27 (24-31)

AADD, age-adjusted D-dimer; CDD, conventional D-dimer; CI, confidence interval; CPR, clinical prediction rule; NR, not reported; PTP, pretest probability; RGS, revised Geneva score; sRGS, simplified revised Geneva score. *Multiple CPRs were used; for simplicity, only results for Wells are presented.

[†]D-dimer value reported in FEUs.

[‡]D-dimer value reported in DDUs;

Study	Class	CPR	РТР	AADD cutoff (µg/L)	CDD Sensitivity (%; 95% CI)	AADD Sensitivity (%; 95% CI)	CDD Miss Rate (%; 95% CI)	AADD Miss Rate (%; 95% Cl)	% Cohort With Negative CDD (95% CI)	% Cohort With Negative AADD (95% CI)
Righini et al ⁴³ *	II	sRGS or Wells	Non-high or unlikely	$Age \times 10^{\dagger}$	NR	NR	1/810 (0.1; 0-0.7)	2/1,141 (0.2; 0-0.6)	28 (27-30)	40 (38-42)
Flores et al ⁴⁵	П	Wells	Non-high	$Age \times 10^{\dagger}$	100 (94-100)	100 (94-100)	0/92 (0; 0-3.9)	0/121 (0; 0-3.0)	28 (23-33)	37 (32-42)
van Es et al ⁴⁴	II	Wells	Unlikely	$Age \times 10^{\dagger}$	99 (99-100)	99 (98-99)	13/2,035 (0.7; 0.4-1.1)	22/2,369 (0.9; 0.6-1.5)	28 (21-37)	33 (25-42)
van Es et al ⁴⁷ *	III	Wells	Unlikely	$Age \times 10^{+}$	NR	NR	1/60 (1.7; 0-8.9)	2/92 (2.2; 0-7.6)	15 (11-18)	22 (18-26)
Gupta et al ⁴⁹	III	NR	Any	$Age \times 10^{+}$	100 (94-100)	97 (90-100)	0/72 (0; 0-5.0)	2/165 (1.2; 0.1-4.3)	7 (7-9)	16 (14-19)
Friz et al ⁵⁰	III	NR	Any	Age $\times 10^{+}$	100 (97-100)	98 (94-100)	0/8 (0; 0-36.9)	2/28 (7.1; 0.9-23.5)	2 (1-3)	6 (4-8)
Jaconelli et al ⁵²	III	Wells	Unlikely	Age $\times 5^{\ddagger}$	95 (86-99)	95 (86-99)	3/859 (0.3; 0.1-1.0)	3/989 (0.3; 0.1-0.9)	65 (62-68)	75 (72-77)
Sharp et al ⁴⁸	III	NR	Any	$Age \times 10^{\dagger}$	98 (96-99)	93 (90-95)	10/16,660 (0.1; 0-0.1)	36/19,584 (0.2; 0.1-0.3)	54 (53-54)	63 (62-64)
Douma et al ⁴⁶	III	Wells	Unlikely	Age $\times 10^{+}$	NR	NR	2/983 (0.2; 0.1-0.7)	7/1,093 (0.6; 0.3-1.3)	46 (43-48)	51 (49-53)
Douma et al ⁴⁶	III	RGS	Non-high	$Age \times 10^{\dagger}$	NR	NR	0/561 (0; 0.0-0.7)	2/663 (0.3; 0.1-1.1)	34 (32-37)	40 (38-43)
Sharp et al ⁴⁸	III	NR	Any	1,000 [†]	98 (96-99)	84 (81-87)	10/16,660 (0.1; 0.0-0.1)	80/23,146 (0.3; 0.3-0.4)	54 (53-54)	74 (74-75)
Friz et al ⁵⁰	III	NR	Any	1,000 [†]	100 (97-100)	96 (91-99)	0/8 (0; 0-36.9)	4/61 (6.6; 1.8-15.9)	2 (1-3)	13 (10-16)
Kline et al ⁵¹ * [§]	III	sRGS or Wells	Any	1,000†	94 (88-97)	92 (86-96)	8/152 (5.3; 2-10.1)	10/185 (5.4; 2.6-9.7)	22 (19-26)	27 (24-31)

AADD, age-adjusted D-dimer; CDD, conventional D-dimer; CI, confidence interval; CPR, clinical prediction rule; NR, not reported; PTP, pretest probability; RGS, revised Geneva score; sRGS, simplified revised Geneva score. *Multiple CPRs were used; for simplicity, only results for Wells are presented.

[†]D-dimer value reported in FEUs.

[‡]D-dimer value reported in DDUs;

Critical Question

In adult patients with low to intermediate pretest probability for acute PE, does a negative age adjusted D-dimer result identify a group of patients at very low risk for the diagnosis of PE for whom no additional diagnostic workup is required?

Level B Recommendation

In patients older than 50 years deemed to be low or intermediate risk for acute PE, clinicians may use a negative age-adjusted D-dimer result to exclude the diagnosis of PE. In your opinion, does the research suggest D-Dimer testing increases or decreases CT imaging use? Clinical Policy: Critical Issues in the Evaluation and Management of Adult Patients Presenting to the Emergency Department With Suspected Acute Venous Thromboembolic Disease



From the American College of Emergency Physicians Clinical Policies Subcommittee (Writing Committee) on Thromboembolic Disease:
Stephen J. Wolf, MD (Subcommittee Chair; Committee Co-Chair)
Sigrid A. Hahn, MD, MPH
Lauren M. Nentwich, MD
Ali S. Raja, MD, MBA, MPH
Scott M. Silvers, MD
Michael D. Brown, MD, MSc (Committee Co-Chair)

> Stephen J Wolf, MD, FACEP Department of Emergency Medicine Denver Health Medical Center University of Colorado School of Medicine





TCPi Transforming Clinical Practices Initiative



For More Information

- E-QUAL Website
 - www.acep.org/equal
 - equal@acep.org

• Contacts:

- Nalani Tarrant: (Director)
- ntarrant@acep.org
- Dhruv Sharma: (Project Manager)
- dsharma@acep.org

