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HEART POCT

Dosaggio di Troponine ad elevavata sensibilità in un
setting preospedaliero

Dott.ssa Margherita Scorpiniti, UOC 118 Arezzo



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IL DOLORE TORACICO

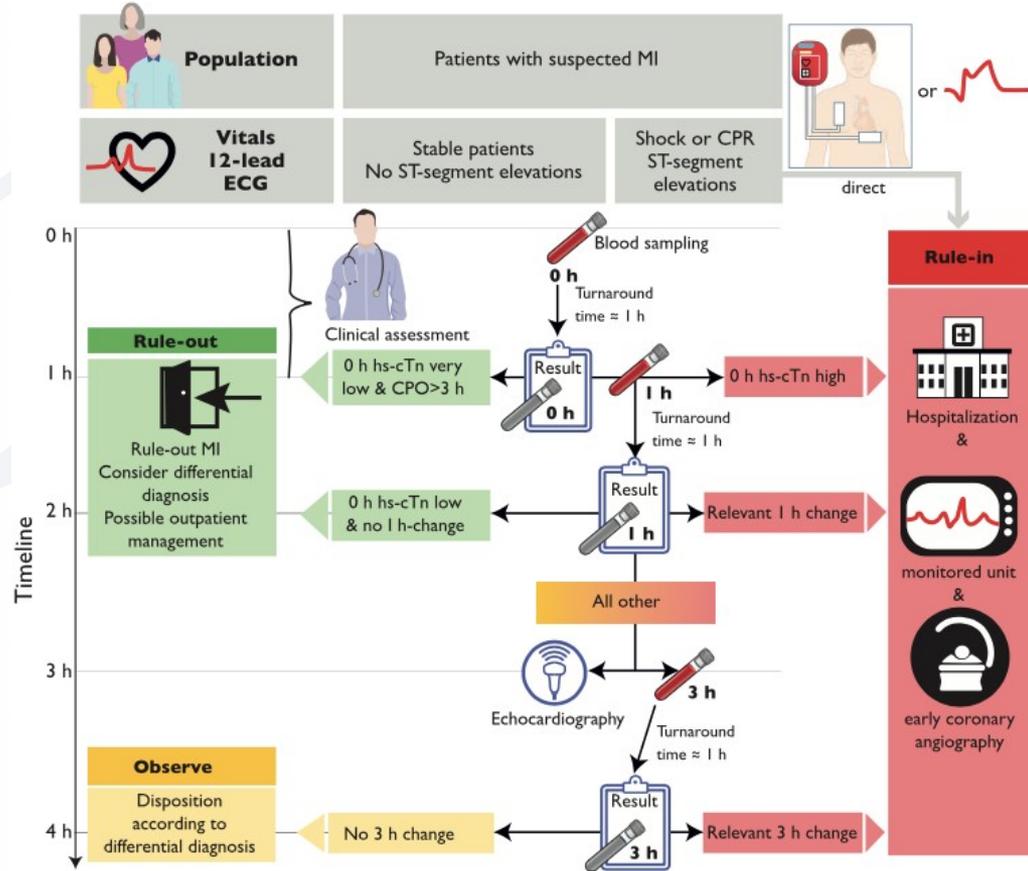


Nella pratica attuale *:

- circa la metà dei pazienti che presentano dolore toracico viene dimesso con diagnosi di causa non cardiaca del dolore
- dei pazienti ricoverati solo il 25% ha una diagnosi finale di SCA
- circa il 2% dei pazienti con dolore toracico con SCA viene erroneamente dimesso.

*Stepinska J, Lettino M, Ahrens I, Bueno H, Garcia-Castrillo L, Khoury A, Lancellotti P, Mueller C, Muenzel T, Oleksiak A, Petrino R, Guimenez MR, Zahger D, Vrints CJ, Halvorsen S, de Maria E, Lip GY, Rossini R, Claeys M, Huber K. Diagnosis and risk stratification of chest pain patients in the emergency department: focus on acute coronary syndromes. A position paper of the Acute Cardiovascular Care Association. Eur Heart J Acute Cardiovasc Care. 2020 Feb;9(1):76-89.

2020 ESC Guidelines for the management of acute coronary syndromes in patients presenting without persistent ST-segment elevation



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Ma se la troponina la misurassimo a casa?



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Letteratura POCT

Practice review



Point-of-care testing with high-sensitivity cardiac troponin assays: the challenges and opportunities

Louise Cullen ,¹ Paul O Collinson ,² Evangelos Giannitsis³

Clinical Chemistry 65:12
1592-1601 (2019)

Point-of-Care Testing

Diagnostic Evaluation of a High-Sensitivity Troponin I Point-of-Care Assay

Nils A. Sörensen,^{1,2*} Johannes T. Neumann,^{1,2} Francisco Ojeda,¹ Evangelos Giannitsis,³ Eberhard Spanuth,⁴ Stefan Blankenberg,^{1,2} Dirk Westermann,^{1,2†} and Tanja Zeller^{1,2†}

- Rapida esecuzione
- Buona fattibilità
- Accuratezza diagnostica (protocollo 0-1h)

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VOL. 75, NO. 10, 2020

Early Diagnosis of Myocardial Infarction With Point-of-Care High-Sensitivity Cardiac Troponin I



Jasper Boeddinghaus, MD,^{a,b,*} Thomas Nestelberger, MD,^{a,b,*} Luca Koechlin, MD,^{a,b,c} Desiree Wussler, MD,^{a,b,d} Pedro Lopez-Ayala, MD,^{a,b} Joan Elias Walter, MD,^{a,b,d} Valentina Troester, MD,^{a,b} Paul David Ratmann, MD,^{a,b} Funda Seidel, MD,^{a,b} Tobias Zimmermann, MD,^{a,b,d} Patrick Badertscher, MD,^{a,b,e} Karin Wildi, MD,^{a,b,f} Maria Rubini Giménez, MD,^{a,b,g} Eliska Potlukova, MD,^{a,b,d} Ivo Strebler, MSc,^{a,b} Michael Freese, RN,^{a,b} Óscar Miró, MD,^{b,h} F. Javier Martin-Sanchez, MD,^{b,i} Damian Kawecki, MD,^{b,j} Dagmar I. Keller, MD,^k Danielle M. Gualandro, MD,^{a,b,l} Michael Christ, MD,^{b,m} Raphael Twerenbold, MD,^{a,b} Christian Mueller, MD,^{a,b} for the APACE Investigators



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Point-of-care testing with high-sensitivity cardiac troponin assays: the challenges and opportunities

Louise Cullen ¹, Paul O Collinson ², Evangelos Giannitsis³



assays may not have a significant impact on ED efficiency. The literature to date illustrates that it is not the provision of rapid cTn results alone that is important but their inclusion within a clinical decision-making pathway.³² Widespread adoption of



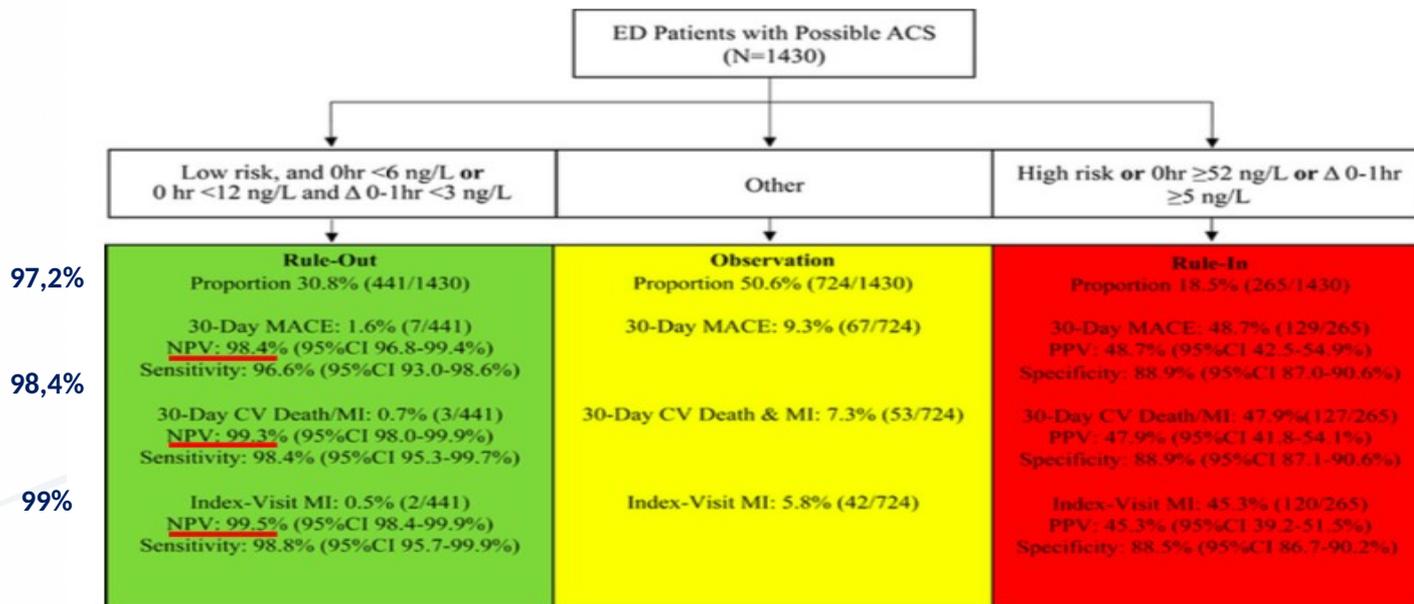
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“High Sensitivity Cardiac Troponin T in a Multisite United States Cohort”

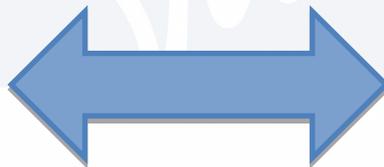
Benjamin M. Gerretsen, MD REBEL EM Medical Category: CIRCULATION, March 2021



1462 pt: The 0/1-h high-sensitivity troponin algorithm combined with the HEART score

HEART POCT

Una proposta di studio clinico



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HEART

HEART score for chest pain patients

H istory (Anamnesis)	Highly suspicious	2	
	Moderately suspicious	1	
	Slightly suspicious	0	
E CG	Significant ST-deviation	2	
	Non-specific repolarisation disturbance / LBBB / PM	1	
	Normal	0	
A ge	≥ 65 years	2	
	45 – 65 years	1	
	≤ 45 years	0	
R isk factors	≥ 3 risk factors <i>or</i> history of atherosclerotic disease	2	
	1 or 2 risk factors	1	
	No risk factors known	0	
T roponin	≥ 3x normal limit	2	
	1-3x normal limit	1	
	≤ normal limit	0	
		Total	

Risk factors for atherosclerotic disease:

Hypercholesterolemia	Cigarette smoking
Hypertension	Positive family history
Diabetes Mellitus	Obesity (BMI>30)



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ATELLICA VTLi

Siemens



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Tipo di Device	Analizzatore	99th Percentile (µg/L unless noted)	% CV at 99th Percentile	Percentuale dei normali misurati ≥ LoD	Tempo per il risultato	Campione
Sistemi palmari	Abbott i-STAT	Overall: 0.08	16.5%	-	10 minuti	vWB/PL 17 µL
	Roche cobas h 232 - TnT	-	-	-	12 minuti	vWB/PL 150 µL
	Siemens ATELLICA VTLi hs-cTnI	Overall: 22.9 ng/L F: 18.5 ng/L M: 27.1 ng/L	6.5% (plasma) 6.1% (WB) at 22.9 ng/L	Overall: 84% F: 80% M: 87%	8 minuti	Capillary/vWB/ PL 30-100 µL
Sistemi da banco	Quidel/Alere TriageTrue hs-cTnI	Overall: 20.5 ng/L F: 14.4 ng/L M: 25.7 ng/L	5.0-5.9% at 21 ng/L (plasma) 5.9-6.5% at 22 ng/L (WB)	Overall: ≥ 50%	20 minuti	vWB/PL 175 µL
	Radiometer AQT90 FLEX TnI	Overall: 0.023	12.3%	-	<19 minuti	vWB/PL 2000 µL
	Radiometer AQT90 FLEX TnT	Overall: 0.017	15.2%	-	<13 minuti	vWB/PL 2000 µL
	PATHFAST hs-cTnI (GEPA)	Overall: 27.9 ng/L F: 20.3 ng/L M: 29.7 ng/L	6.1%	Overall: 66.3% F: 52.8% M: 78.8%	<17 minutes	vWB/Siero/PL 100 µL

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Prospective study sampling 1089 patient

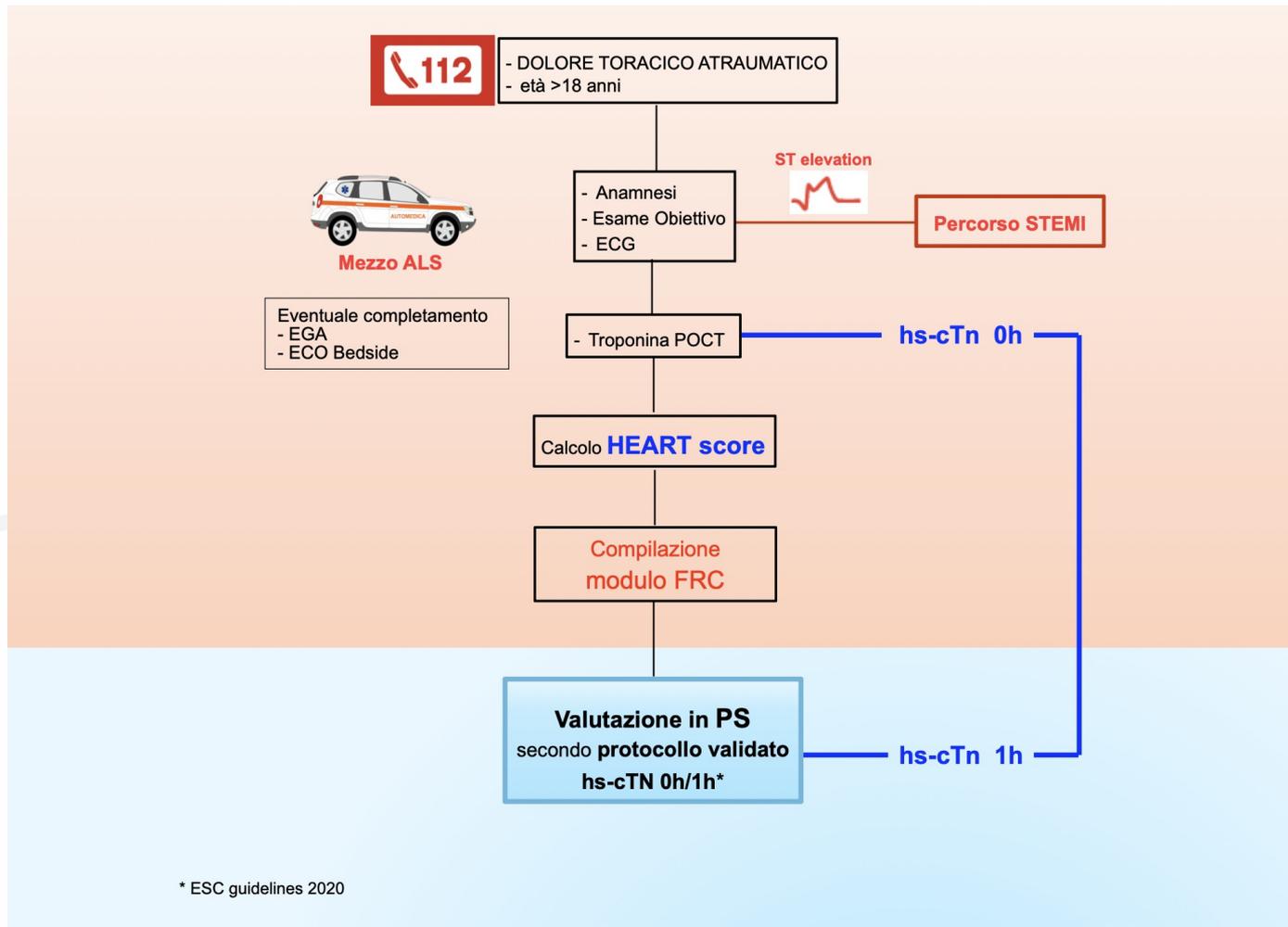
Popolazione	99th URL (ng/L)	Timepoint	Soggetti		Sensibilità (95% CI)	Specificità (95% CI)	PPV (95% CI)	NPV (95% CI)
Generale	22.9	Baseline	998	91	64.8% (54.6–73.9%)	85.7% (83.4–87.7%)	29.2% (25.0–33.8%)	96.4% (95.3–97.2%)
		2 ore	998	91	81.3% (72.1–88.0%)	84.6% (82.2–86.7%)	32.5% (28.7–36.4%)	98.0% (97.0–98.7%)
Uomini	27.1	Baseline	615	56	67.9% (54.8–78.6%)	86.2% (83.2–88.7%)	30.9% (25.5–36.9%)	96.7% (95.3–97.7%)
		2 ore	615	56	80.4% (68.2–88.7%)	84.7% (81.7–87.3%)	32.4% (27.6–37.5%)	97.9% (96.5–98.8%)
Donne	18.5	Baseline	383	35	65.7% (49.2–79.2%)	85.4% (81.5–88.6%)	29.1% (22.6–36.6%)	96.5% (94.5–97.7%)
		2 ore	383	35	82.9% (67.3–91.9%)	84.3% (80.4–87.6%)	32.6% (26.8–38.9%)	98.2% (96.3–99.1%)

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Diagnostic Performance of a Rapid, Novel, Whole Blood, Point of Care High-Sensitivity Cardiac Troponin I Assay for Myocardial Infarction.
Ian L.GunsolusaKarenSchulzbYaderSandovalcStephen W.SmithdeBrittanyLindgrenfBrynnOkesongFred S.Applehl
<https://doi.org/10.1016/j.clinbiochem.2022.04.008>

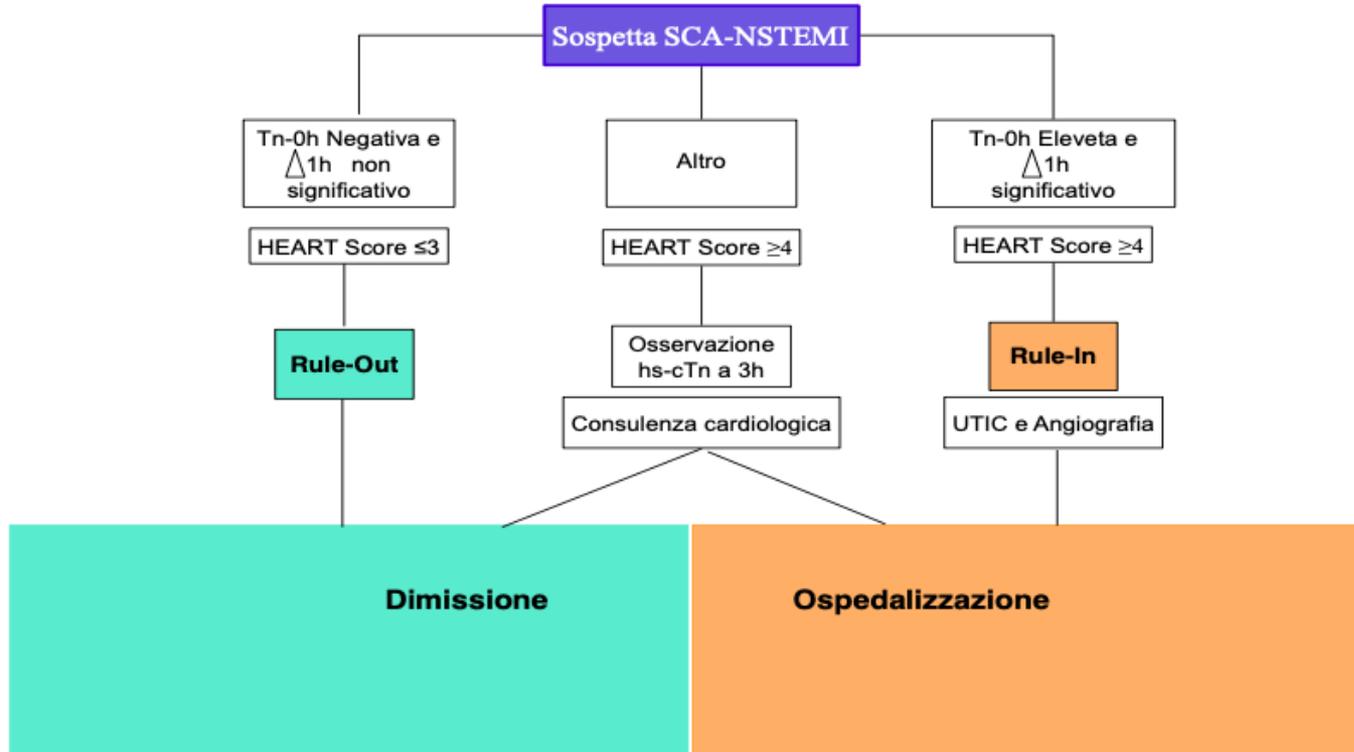
Criteri inclusione

- età >18 anni
- dolore toracico non traumatico
- consenso informato del paziente



* ESC guidelines 2020

PROTOCOLLO hs-cTN 0-1h e HEART Score



HEART POCT

Una proposta di studio clinico

Outcome primario

- Valutare la fattibilità della hs-cTn POCT in ambito pre-ospedaliero.
- Valutare la performance della hs-cTn POCT nel rule-in e rule-out di paziente con sospetta SCA tramite l'utilizzo dell'algoritmo 0/1 ora in combinazione con l'HEART score.

Outcome secondario

- Valutare a posteriori se con l'introduzione della hs-cTn POCT e HEART Score nel pre-ospedaliero, i pazienti identificati a basso rischio potevano essere gestiti a domicilio senza necessità di invio in PS.
- Valutare se, con l'assetto tecnologico e culturale attuale, il medico di AM può individuare cause non cardiache di dolore toracico a domicilio.

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Punti di forza

- Risultato in 8 min
- Accuratezza diagnostica
- Maneggevolezza
- Sangue intero capillare o venoso, plasma
- Hs-cTnI con ridotta influenza dei valori di creatinina

Limiti

- Durante la processazione deve rimanere su un piano fisso
- La cartuccia del reagente deve essere conservata a 2-8°C.
- Deve essere usato entro 15 minuti dall'apertura della busta
- Stoccaggio a temperatura ambiente (18-26°C) fino a 8h



BMJ Open Acute rule-out of non-ST-segment elevation acute coronary syndrome in the (pre)hospital setting by HEART score assessment and a single point-of-care troponin: rationale and design of the ARTICA randomised trial

Goarls W A Aarts ¹, Cyril Camaro ¹, Robert-Jan van Geuns,¹ Etienne Cramer,¹ Roland R J van Kimmenade,¹ P Damman,¹ Pierre M van Grunsven,² Eddy Adang,³ Paul Giesen,⁴ Martijn Rutten,⁴ Olaf Ouwendijk,⁵ Marc E R Gomes,⁶ Niels van Royen¹

Multicentrico: 800 Pz
poct+heart fattibile e accurato
Riduzione dei costi di gestione ospedaliera

ClinicoEconomics and Outcomes Research

Dovepress

open access to scientific and medical research

 Open Access Full Text Article

ORIGINAL RESEARCH

Economic evaluation of point-of-care testing in the remote primary health care setting of Australia's Northern Territory

This article was published in the following Dove Press journal:
ClinicoEconomics and Outcomes Research

Conclusion: This study demonstrated that POCT when used to aid decision making for acutely ill patients delivered significant cost savings for the NT health care system by preventing unnecessary emergency medical retrievals.

Territori remoti e rurali: 200 Pz



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Grazie per l'attenzione



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