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Point-of-Care Guides

Risk Stratification of Patients Presenting with Syncope

SAN FRANCISCO SYNCOPE RULE ³	ROSE RISK SCORE⁵	OESIL RISK SCORE ⁶
Risk factors		
Systolic blood pressure < 90 mm Hg	Brain natriuretic peptide level ≥ 300 pg per mL (300 ng per L)	Age > 65 years History of
Shortness of breath	Bradycardia (≤ 50 beats per	cardiovascular
ECG: Nonsinus rhythm	minute)	disease
or new changes present	Rectal examination shows fecal occult blood	Syncope without a prodrome
History of congestive heart failure	Anemia (hemoglobin level < 9.0	Abnormal ECG
Hematocrit < 30 percent	g per dL [90.0 g per L])	findings
	Chest pain associated with syncope	
	ECG with Q wave (not in lead III)	
	Oxygen saturation ≤ 94 percent on room air	
Risk groups*		
No factors present: 0.3 percent	No factors present: 1.5 percent	0 to 1 factor present (low risk): 0.6 percent
≥ 1 factors present: 15.2 percent	≥ 1 factors present: 16.5 percent	2 to 4 factors present (high risk): 31 percent
Accuracy of score		
98 percent sensitive	87 percent sensitive	97 percent sensitive
56 percent specific	66 percent specific	73 percent specific

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LR+ = 2.9	LR+ = 2.5	LR+ = 3.6

$$LR-=0.03$$
 $LR-=0.2$ $LR-=0.11$

NOTE: Results are shown for the study used to originally develop and validate each risk score.

*—The San Francisco Syncope Rule and ROSE risk score measure risk of serious outcome or death at one month; the OESIL risk score measures risk of all-cause mortality at 12 months.

ECG = electrocardiography; LR- = negative likelihood ratio; LR+ = positive likelihood ratio; OESIL = Osservatorio Epidemiologico sulla Sincope nel Lazio; ROSE = Risk Stratification of Syncope in the Emergency Department.

Information from references 3, 5, and 6.

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