

Prognostic value of stress echocardiography assessed by the ABCDE protocol

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Received 16 March 2021; revised 4 May 2021; editorial decision 22 June 2021; accepted 19 July 2021; online publish-ahead-of-print 27 August 2021

See page 3879 for the editorial comment on this article (doi:10.1093/eurheartj/ehab562)

Aim

The aim of this study was to assess the prognostic value of ABCDE-SE in a prospective, large scale, multicentre, international, effectiveness study. Stress echocardiography (SE) was recently upgraded to the ABCDE protocol: step A, regional wall motion abnormalities; step B, B lines; step C, left ventricular contractile reserve; step D, Doppler-based coronary flow velocity reserve in left anterior descending coronary artery; and step E, electrocardiogram-based heart rate reserve.

Methods and results

From July 2016 to November 2020, we enrolled 3574 all-comers (age 65 ± 11 years, 2070 males, 58%; ejection fraction $60 \pm 10\%$) with known or suspected chronic coronary syndromes referred from 13 certified laboratories. All patients underwent clinically indicated ABCDE-SE. The employed stress modality was exercise ($n = 952$, with semi-supine bike, $n = 887$, or treadmill, $n = 65$ with adenosine for step D) or pharmacological stress ($n = 2622$, with vasodilator, $n = 2151$; or dobutamine, $n = 471$). SE response ranged from score 0 (all steps normal) to score 5 (all steps abnormal). All-cause death was the only endpoint. Rate of abnormal results was 16% for A, 30% for B, 36% for C, 28% for D, and 37% for E steps. During a median follow-up of 21 months (interquartile range: 13–36), 73 deaths occurred. Global χ^2 was 49.5 considering clinical variables, 50.7 after step A only ($P = \text{NS}$ (not significant)) and 80.6 after B–E steps ($P < 0.001$ vs. step A). Annual mortality rate ranged from 0.4% person-year for score 0 up to 2.7% person-year for score 5.

Conclusion

ABCDE-SE allows an effective prediction of survival in patients with chronic coronary syndromes.

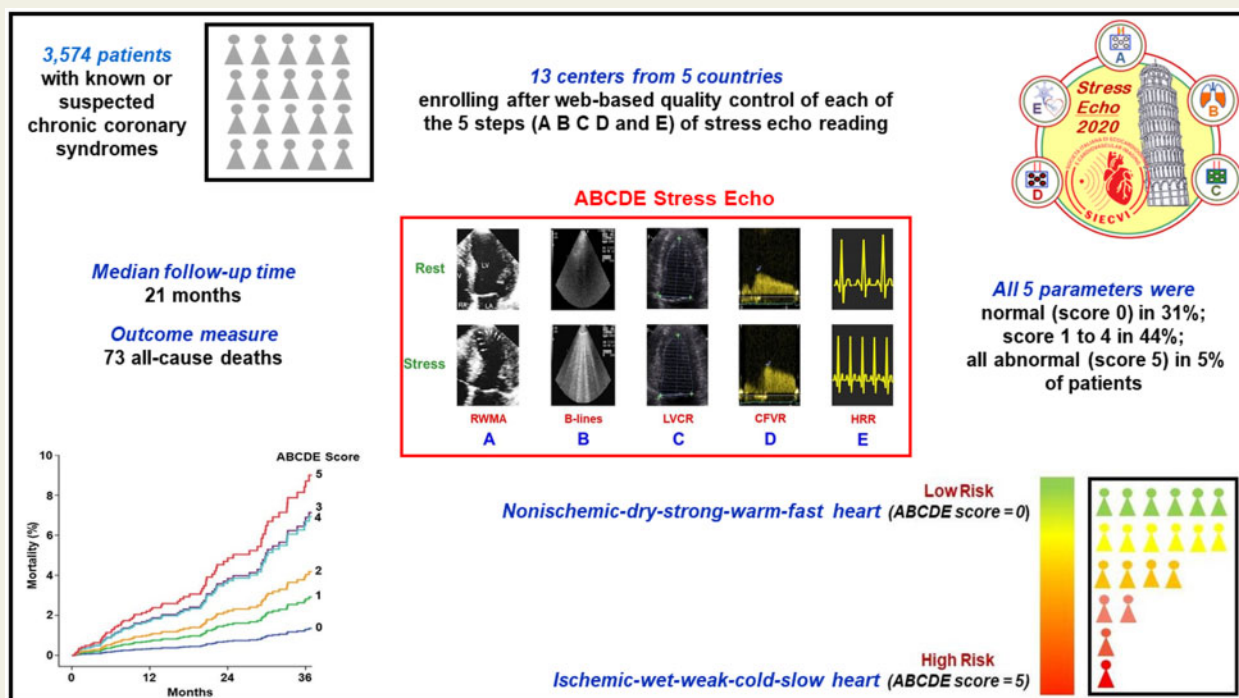
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Graphical Abstract



This study recruited 3,574 patients from 13 stress echo laboratories of 5 countries and shows the prognostic value of the new state-of-the-art cardiac functional testing with stress echo ABCDE protocol: Step A for regional wall motion abnormalities; step B for B-lines obtained with 4-site simplified scan and lung ultrasound; step C for contractile reserve with volumetric echocardiography; step D for Doppler-based assessment of coronary flow reserve in mid-distal left anterior descending coronary artery; step E for EKG-based heart rate reserve.

Keywords

Echocardiography • Outcome • Stress • Vulnerability

Introduction

Stress echocardiography (SE) is an established diagnostic technique based on the detection of inducible myocardial ischaemia as regional wall motion abnormality (RWMA). This approach has a recognized clinical value but it seems unfit to describe the emerging complexity and heterogeneity of the individual patient vulnerability. Therefore, SE was recently upgraded to the ABCDE protocol:¹ step A for RWMA, which is the only sign established in guidelines and recommendations;^{2,3} step B for B lines by lung ultrasound;^{4,5} step C for left ventricular contractile reserve (LVCR);^{6,7} step D for Doppler-based coronary flow velocity reserve (CFVR) in the left anterior descending coronary artery (LAD);⁸ and step E for imaging-independent heart rate reserve (HRR) based on the electrocardiogram (ECG).⁹⁻¹¹ This approach applies to all types of physical or pharmacological stress modalities and allows a more integrated assessment of patient vulnerability above and beyond the detection of anatomic coronary artery disease (CAD). ABCDE-SE allows assessment of inducible myocardial ischaemia (step A), pulmonary congestion (step B), contractile reserve (step C), coronary microvascular dysfunction (step D), and cardiac sympathetic reserve (step E). The ABCDE-SE protocol is

more likely to capture many potential sources of vulnerability of the patient with CAD^{12,13} than a simpler approach based only on RWMA.

The study hypothesis was that each pathophysiological variable may contribute independently and incrementally to the prognostic vulnerability in the individual patient and, therefore, the outcome would progressively worsen with the increasing number of abnormal steps during cardiac functional stress. To test this hypothesis, ABCDE-SE was performed in patients with chronic coronary syndromes, referred for clinically driven SE in accredited laboratories¹⁴ in the network of the international, multicentre, prospective SE 2020 study¹⁵ (Graphical abstract).

Methods

Study population

In this prospective study, we initially screened 4213 patients recruited from July 2016 to November 2020 by 13 certified laboratories of 5 countries (Bulgaria, Italy, Poland, Russian Federation, Serbia). Of these initial 4213, 639 did not complete the full protocol with missing information on step B ($n = 281$), step C ($n = 18$), or step D ($n = 286$). The main reasons

were that the step was performed prior to specific training and certification for steps B and C, or the test was attempted but data were not obtained because of technical failure in step D. An additional 54 patients were lost to follow-up. The final study population included 3574 patients all studied with the full ABCDE protocol with outcome information.

The inclusion criteria were: (i) age >18 years; (ii) referral for known or suspected CAD; (iii) no severe primary valvular or congenital heart disease, or presence of prognosis-limiting comorbidities, such as advanced cancer, reducing life expectancy to <1 year; (iv) echocardiography of acceptable quality at rest and during stress; (v) willingness to give their written informed consent allowing scientific utilization of observational data, respectful of privacy rights; and (vi) recruitment in centres with systematic follow-up program.

All patients underwent SE testing as part of a clinically driven evaluation and according to the referring physician's indications. Indications were in recommendation Class I or IIa according to the European Society of Cardiology² and/or American Society of Echocardiography,³ falling into one of the three broad categories: initial test to diagnose obstructive CAD in patients with preserved ejection fraction, especially in patients with intermediate pre-test probability of disease ($n = 1400$, 39%); inconclusive exercise ECG or unable to exercise ($n = 452$, 13%); and prognosis assessment or re-assessment in stable CAD, with or without global left ventricular dysfunction ($n = 1722$, 48%).

Written informed consent was obtained from all patients before testing. The study protocol was reviewed and approved by the institutional ethics committees as a part of the SE 2020 study (148—Comitato Etico Lazio-1, 16 July 2016; ClinicalTrials.gov Identifier NCT03049995). The study was funded partly by the Italian National Research Council (Ageing project) and with travel grants of the Italian Society of Echocardiography and Cardiovascular Imaging with dedicated sessions during national meetings. No support from industry was received.

Stress echocardiography

We used commercially available ultrasound machines. All patients underwent comprehensive transthoracic echocardiography at rest¹⁶ and underwent physical or pharmacological SE according to the protocol recommended by the European Association of Echocardiography¹⁷ and American Society of Echocardiography.¹⁸ The exercise protocol was semi-supine bike or post-treadmill. We used dipyridamole dose up to 0.86 mg/kg (over 4 or 6 min), dobutamine starting from 5 up to 40 µg/kg/min with atropine co-administration up to 1 mg, and adenosine up to 0.14 µg/kg/min over 6 min. In one centre (Belgrade), ABCE steps were acquired with treadmill exercise, and step D with adenosine (0.14 µg/kg/min in 2 min) in the same patients on the same day, 1 h after the treadmill test.

Criteria for terminating the test were severe chest pain, diagnostic ST-segment shift, excessive blood pressure increase (systolic blood pressure ≥ 240 mmHg, diastolic blood pressure ≥ 120 mmHg), limiting dyspnoea, maximal predicted heart rate, and significant arrhythmias. Echocardiographic imaging was performed from parasternal long- and short-axis views, and apical four- and two-chamber views, using conventional two-dimensional echocardiography. Contrast use was recommended in cases of suboptimal acoustic window, when ≥ 2 segments were not optimally visualized for 50% or more of endocardial length, and its use ranged from 0% to 40% in recruiting centres, depending on local lab policy and reimbursement issues. Anti-anginal drugs were usually not suspended before testing.

Step A included assessment of wall motion abnormalities and was performed in all patients. Wall motion score index (WMSI) was calculated in each patient at baseline and peak stress, in a 4-point score ranging from 1 (normal) to 4 (dyskinetic) in a 17-segment model of the left ventricle.

Step B of protocol included the assessment of B lines with lung ultrasound and the four-site simplified scan, from mid-axillary to mid-clavicular lines on the third intercostal space, each site scored from 0 (normal horizontal A lines) to 10 (white lung with coalescent B lines).⁵

Step C of protocol included the force-based assessment of LVCR as the stress/rest ratio of force, calculated as systolic blood pressure/end-systolic volume.⁷

Coronary flow velocity reserve (step D) was assessed during the standard SE examination using intermittent imaging of wall motion and LAD.⁸ Coronary flow in the mid-distal portion of the LAD was imaged from the low parasternal long-axis view and/or modified apical two-, three-, or four-chamber view under the guidance of colour Doppler flow mapping. All studies were digitally stored to simplify offline reviewing and measurements. At each time point, three optimal profiles of peak diastolic Doppler flow velocities were measured, and the results were averaged.

Heart rate reserve (step E) was calculated as the peak/rest heart rate from 12-lead ECG.¹¹

All steps were performed by the same sonographer/cardiologist with the same transducer for cardiac, lung, and coronary scan—although occasionally a different high-frequency transducer was used for coronary flow. All steps were acquired at rest and peak stress. If needed, steps were repeated after 5 min in the recovery phase (Figure 1).

A detailed visual description of the scanning procedure is also available in a 9-min movie from the consortium (YouTube. ABCDE Stress Echo 2030: How I Make It. More easily done than said. Available at: <https://www.youtube.com/watch?v=O4-5FjSF7ao>, accessed 26 May 2021).

Stress echocardiography positivity criteria

All positivity criteria were determined a priori.

The A criterion was considered positive in the presence of stress-induced RWMA (WMSI stress > rest), when at least two adjacent segments of the same vascular territory of the left ventricle showed an increment of at least one point of segmental score during SE.

The B criterion was considered positive in the presence of stress or rest B lines ≥ 2 units.⁵

The C criterion was considered positive in the presence of force-based LVCR ≤ 2.0 for exercise or dobutamine and ≤ 1.1 for vasodilators.⁷

The D criterion was considered positive in the presence of CFVR ≤ 2.0 .⁸

The E criterion was considered positive in the presence of HRR <1.80 for exercise or dobutamine or <1.22 for dipyridamole or adenosine.^{9–11}

As required by SE 2020 protocol, all readers had passed the quality control for each of the four imaging parameters upstream to starting patient recruitment.¹⁵

Stress echocardiography response was summarized with a score ranging from 0 to 5 as follows: score 0 (all ABCDE markers within normal limits) or scores 1–5, according to the number of abnormal steps (e.g. score 5 indicated that all 5 steps were abnormal).

Data storage and analysis

The results for each test were entered in the data bank at the time of testing by each recruiting centre and sent monthly to the core lab with the electronic case report form with clinical data. After checking for internal consistency by trained technical staff, and double-checking with the centre for data verification on possibly inconsistent input, the data were added to the data bank and locked.

Coronary angiography

Invasive coronary angiography ($n = 1421$) or non-invasive multidetector coronary angiography ($n = 20$) showing no CAD was available in 1441

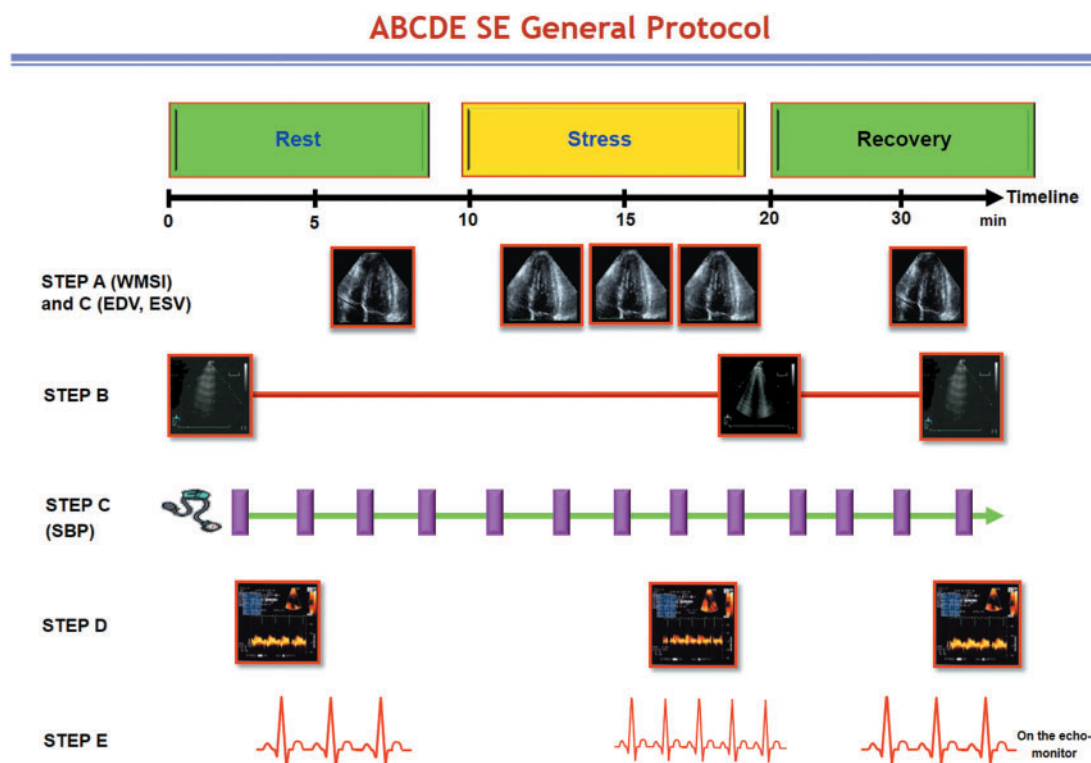


Figure 1 The ABCDE protocol. The completion of the test from preparation phase to written response takes ~30 min. Recovery phase is optional, and advised in the presence of test positivity. Step A and step C require the very same images. Step E requires only one electrocardiogram lead. Step B can be completed in the early (<30 s) recovery phase. SBP, systolic blood pressure; SE, stress echocardiography.

patients. Coronary angiography was indicated by the referring physician based on symptoms, individual clinical characteristics, and non-invasive imaging results. Obstructive significant CAD was defined by a quantitatively assessed coronary diameter reduction $\geq 50\%$ in the view showing the most severe stenosis. Images were read by experienced invasive cardiologists unaware of the SE results.

Outcome data analysis

Deaths were identified from the national health service database. Non-deceased participants were contacted directly. Follow-up data were obtained from review of the patient's hospital record, personal communication with the patient's physician, review of the patient's chart, a telephone interview with the patient or a patient's close relative conducted by trained personnel, and a staff physician visiting the patients at regular intervals in the outpatient clinic. To avoid misclassification of the cause of death, overall death was considered. Assessors were blinded to clinical and SE results.

Statistical analysis

Categorical data are expressed in terms of number of subjects and percentage while continuous data are expressed as mean \pm standard deviation or median (minimum–maximum) depending on variable distribution. For continuous variables, differences among groups were tested with one-way analysis of variance followed by Bonferroni *post hoc* tests for between-group comparisons or Kruskal–Wallis followed by Mann–Whitney test using the Bonferroni correction, as appropriate. χ^2

test or Fisher's exact test were used to compare the distribution of categorical variables among groups.

Survival was estimated using the Kaplan–Meier method and survival curves were compared by means of the log-rank test. Univariate Cox proportional hazards models were used to identify candidate predictors for selected endpoints. All variables with P -value < 0.10 at univariate analysis were considered for inclusion in the multivariate Cox proportional hazards model. Collinearity was verified for all the models. We used the variance inflation factors to check the presence of collinearity and, for the multivariable model containing the positivity components, the overall mean variance inflation factor was 1.16. Moreover, values for all the variables were below 1.4, thus suggesting no evidence of collinearity in the model.

Non-proportionality of hazard was assessed using the Schoenfeld test. The incremental value of ABCDE-SE was evaluated comparing multivariable models with and without individual steps using global X^2 value to evaluate improvement of goodness-of-fit as well as continuous net reclassification index and integrated discrimination index to assess its incremental value. Statistical significance was set at P -value < 0.05 . All analyses were performed using STATA (STATA Corp. Stata statistical software: Release 14. College Station, TX, USA).

Results

The main clinical characteristics of the 3574 study patients are described in Table 1. The employed stress was exercise ($n = 952$,

with semi-supine bicycle exercise: $n = 887$ or treadmill, $n = 65$), or pharmacological ($n = 2622$) with vasodilator ($n = 2151$; dipyridamole, $n = 2112$; adenosine, $n = 39$), or dobutamine ($n = 471$). Patients with more advanced ABCDE score (4 or 5), when compared to patients with lower scores, were more often hypertensives and studied under anti-ischaemic therapy with beta-blockers and nitrates (Table 1).

Stress echocardiography findings

The main SE findings are described in Table 2. Figures 2 and 3 show examples of ABCDE negativity and ABCDE positivity, respectively. Rate of abnormal results was 16% for A, 30% for B, 36% for C, 28% for D, and 37% for E steps.

The positivity rate in various centres differed significantly, possibly due to different referral patterns and also because of the limited sample in some centres that determined a high degree of variability. In details, positivity rate ranged from 0% to 56% for step A, from 0% to 60% for step B, from 16% to 100% for step C, from 13% to 100% for step D, and from 8% to 86% for step E (Supplementary material online, Table S1). Each centre had its preferred stress modality.

Stress modality by centre is shown in Supplementary material online, Table S2.

All five parameters were normal (score 0) in 1114 patients (31%), all five were abnormal (score 5) in 172 patients (5%). The abnormal score was 1 in 1064 (30%), 2 in 610 (17%), 3 in 349 (10%), and 4 in 265 (7%) patients.

In the 1441 patients with coronary angiography available, 848 patients (59%) showed no CAD, 373 patients (26%) showed one-vessel CAD, and 220 patients (15%) showed multivessel CAD. ABCDE score increased progressively with increasing extent of CAD (Figure 4).

Outcome data results

During a median follow-up of 21 months (interquartile range: 13–36), 73 deaths occurred. In the univariable analysis, predictors of all-cause mortality were step B [hazard ratio (HR) 2.621, 95% confidence interval (CI) 1.654–4.152, $P < 0.001$], step D (HR 2.578, 95% CI 1.624–4.093, $P < 0.001$), and step E (HR 2.955, 95% CI 1.848–4.725; $P < 0.001$), but not step A (HR 1.333, 95% CI 0.731–2.430; $P = 0.349$) and step C (HR 1.581, 95% CI 0.997–2.506; $P = 0.051$) (Table 3). In

Table 1 Clinical characteristics of the patients

	All patients ($n = 3574$)	ABCDE score						P-value
		0 ($n = 1114$)	1 ($n = 1064$)	2 ($n = 610$)	3 ($n = 349$)	4 ($n = 265$)	5 ($n = 172$)	
Age (years)	65 ± 11	$62 \pm 11^{a,b,c,d}$	65 ± 11^c	66 ± 11	$68 \pm 11^{d,e}$	66 ± 10	64 ± 9	<0.001
BSA (m^2)	1.87 ± 0.22	1.86 ± 0.24^e	1.87 ± 0.22	1.87 ± 0.24	1.85 ± 0.20	1.89 ± 0.19	1.92 ± 0.21	0.022
Sex (M/F)	2070 (58)	658 (59)	597 (56)	346 (57)	193 (55)	158 (60)	118 (69)	0.302
	1504 (42)	456 (41)	467 (44)	264 (43)	156 (45)	107 (40)	54 (31)	
Type of stress								<0.001
Exercise	952 (27)	113 (10)	241 (23)	171 (28)	101 (29)	178 (67)	148 (86)	
Vasodilator	2151 (60)	959 (86)	648 (61)	291 (48)	158 (45)	72 (27)	22 (13)	
Dobutamine	471 (13)	42 (4)	175 (16)	148 (24)	90 (26)	15 (6)	1 (1)	
Hypertension	2747 (77)	833 (75)	790 (74)	465 (75)	193 (76)	275 (79)	155 (90)	<0.001
Diabetes	846 (24)	212 (19)	244 (23)	167 (27)	111 (32)	67 (25)	45 (26)	<0.001
Previous PCI/CABG	1310 (40)	394 (37)	376 (43)	197 (43)	114 (46)	127 (52)	102 (62)	<0.001
Coronary angiography available	1441 (45)	432 (39)	411 (39)	287 (47)	161 (46)	104 (39)	46 (27)	<0.001
No CAD)	848 (59)	309 (71)	236 (64)	174 (61)	78 (48)	22 (21)	2 (4)	
1 vessel	373 (26)	108 (25)	112 (27)	66 (23)	43 (27)	33 (32)	11 (24)	
Multivessel	220 (15)	15 (4)	36 (9)	47 (16)	70 (25)	49 (47)	33 (72)	
History of myocardial infarction	928 (26)	259 (23)	275 (26)	150 (25)	96 (27)	77 (29)	71 (41)	<0.001
Beta-blockers	2179 (61)	529 (52)	635 (60)	393 (64)	245 (70)	193 (69)	134 (78)	<0.001
ACE-inhibitors/ARB	2343 (66)	651 (58)	678 (64)	415 (68)	256 (73)	198 (75)	145 (84)	<0.001
Nitrates	101 (3)	7 (1)	16 (2)	22 (4)	17 (5)	17 (6)	22 (13)	<0.001
Calcium channel blockers	633 (18)	146 (13)	169 (16)	114 (19)	89 (26)	71 (27)	44 (26)	<0.001
Diuretics	793 (22)	142 (11)	241 (23)	177 (29)	112 (32)	78 (29)	61 (35)	<0.001

Values are expressed as mean \pm standard deviation or n (%).

ACE, angiotensin-converting enzyme; ARB, angiotensin receptor blocker; BSA, body surface area; CABG, coronary artery bypass graft; CAD, coronary artery disease; F, female; M, male; PCI, percutaneous coronary intervention.

^a $P < 0.05$ vs. ABCDE score 1.

^b $P < 0.05$ vs. ABCDE score 2.

^c $P < 0.05$ vs. ABCDE score 3.

^d $P < 0.05$ vs. ABCDE score 4.

^e $P < 0.05$ vs. ABCDE score 5.

Table 2 Haemodynamic, rest, and stress echocardiographic findings

	All patients (n = 3574)	ABCDE score						P-value
		0 (n = 1114)	1 (n = 1064)	2 (n = 610)	3 (n = 349)	4 (n = 265)	5 (n = 172)	
Rest WMSI	1.10 ± 0.23	1.05 ± 0.17 ^{a,b,c,d}	1.07 ± 0.19 ^{a,b,c,d}	1.11 ± 0.24 ^{c,d}	1.15 ± 0.29 ^d	1.18 ± 0.33	1.24 ± 0.36	<0.001
Stress WMSI	1.15 ± 0.31	1.04 ± 0.14 ^{a,b,c,d}	1.06 ± 0.18 ^{a,b,c,d}	1.12 ± 0.25 ^{b,c,d}	1.22 ± 0.30 ^{c,d}	1.53 ± 0.37 ^d	1.82 ± 0.47	<0.001
ΔWMSI (stress–rest)	0.06 ± 0.21	-0.01 ± 0.07 ^{a,b,c,d}	-0.01 ± 0.09 ^{b,c,d}	0.01 ± 0.11 ^{b,c,d}	0.06 ± 0.15 ^{c,d}	0.35 ± 0.36 ^d	0.58 ± 0.31	<0.001
A positivity	570 (16)	0	16 (2)	59 (10)	116 (33)	207 (78)	172 (100)	<0.001
B lines at rest	0.9 (0–35)	0.1 (0–1)	1.0 (0–32)	1.4 (0–30)	1.7 (0–34)	1.6 (0–20)	2.1 (0–35)	<0.001
B lines at peak	1.7 (0–40)	0.1 (0–1) ^{a,b,c,d,e}	1.4 (0–40) ^{a,b,c,d}	2.3 (0–24) ^{b,c,d}	3.1 (0–33) ^{c,d}	4.0 (0–40) ^d	6.3 (0–39)	<0.001
B positivity	1072 (30)	0	299 (28)	264 (43)	185 (53)	152 (57)	172 (100)	<0.001
DBP rest (mmHg)	78 ± 10	78 ± 10 ^d	78 ± 10	79 ± 10	79 ± 9	79 ± 10	80 ± 11	0.011
DBP stress (mmHg)	78 ± 15	75 ± 14 ^{a,b,c,d,e}	78 ± 15 ^{c,d}	78 ± 15 ^{c,d}	79 ± 15 ^d	81 ± 15	83 ± 15	<0.001
SBP rest (mmHg)	132 ± 17	130 ± 15 ^{a,b,c,d,e}	132 ± 17 ^{a,b}	134 ± 18	136 ± 18	134 ± 17	135 ± 20	<0.001
SBP stress (mmHg)	142 ± 34	134 ± 30 ^{a,b,c,d,e}	143 ± 35 ^{c,d}	145 ± 35 ^{c,d}	144 ± 34 ^{c,d}	153 ± 32	156 ± 28	<0.001
EF rest (%)	60 ± 10	61 ± 6 ^{a,b}	61 ± 8 ^b	59 ± 16	58 ± 10	60 ± 11	61 ± 10	<0.001
EF stress (%)	69 ± 11	74 ± 8 ^{a,b,c,d,e}	71 ± 9 ^{a,b,c,d}	69 ± 11 ^{b,c,d}	63 ± 12 ^{c,d}	59 ± 12 ^d	56 ± 11	<0.001
EDV rest (mL)	84 ± 29	77 ± 20 ^{a,b,c,d,e}	83 ± 27 ^{c,d}	87 ± 29 ^d	89 ± 22 ^d	98 ± 39	104 ± 37	<0.001
EDV stress (mL)	80 ± 29	72 ± 21 ^{a,b,c,d,e}	78 ± 26 ^{c,d}	83 ± 29 ^{c,d}	84 ± 32 ^{c,d}	96 ± 37	104 ± 37	<0.001
ESV rest (mL)	34 ± 19	30 ± 10 ^{a,b,c,d,e}	35 ± 14 ^{c,d}	36 ± 19 ^d	38 ± 22	41 ± 29	43 ± 25	<0.001
ESV stress (mL)	26 ± 18	19 ± 8 ^{a,b,c,d,e}	23 ± 13 ^{a,b,c,d}	28 ± 17 ^{b,c,d}	33 ± 21 ^{c,d}	41 ± 28 ^d	48 ± 27	<0.001
Force rest (mmHg/mL)	4.62 ± 1.96	4.86 ± 1.73 ^{c,d}	4.63 ± 1.90 ^{c,d}	4.60 ± 2.23 ^d	4.51 ± 2.21	4.23 ± 1.87	4.04 ± 2.06	<0.001
Force peak (mmHg/mL)	7.39 ± 4.19	8.46 ± 3.93 ^{a,b,c,d}	8.00 ± 4.24 ^{a,b,c,d}	7.07 ± 4.56 ^{b,c,d}	6.04 ± 3.86 ^{c,d}	4.93 ± 2.68	4.27 ± 2.32	<0.001
LVCR	1.61 ± 0.72	1.77 ± 0.72 ^{a,b,c,d}	1.76 ± 0.77 ^{a,b,c,d}	1.56 ± 0.75 ^{b,c,d}	1.35 ± 0.53 ^{c,d}	1.17 ± 0.38	1.08 ± 0.29	<0.001
C positivity	1295 (26)	0	263 (25)	334 (55)	271 (78)	255 (96)	172 (100)	<0.001
LAD rest (cm/s)	26.8 ± 12.4	25.6 ± 7.4 ^{c,d}	24.8 ± 11.0 ^{b,c,d}	25.3 ± 13.3 ^{c,d}	27.2 ± 12.4 ^{c,d}	33.2 ± 17.6 ^d	38.6 ± 21.2	<0.001
LAD stress (cm/s)	58.1 ± 24.0	65.5 ± 17.4 ^{a,b,c,d,e}	59.0 ± 25.1 ^{a,b,c,d}	53.8 ± 27.7	50.1 ± 23.3	49.7 ± 23.9	49.5 ± 25.5	<0.001
CFVR	2.25 ± 0.55	2.54 ± 0.36 ^{a,b,c,d,e}	2.42 ± 0.44 ^{a,b,c,d}	2.16 ± 0.45 ^{b,c,d}	1.89 ± 0.45 ^{c,d}	1.56 ± 0.42 ^d	1.31 ± 0.28	<0.001
D positivity	994 (28)	0	111 (10)	221 (36)	245 (70)	245 (92)	172 (100)	<0.001
Heart rate rest (b.p.m.)	68 ± 11	65 ± 10 ^{a,b,c,d,e}	68 ± 11 ^{a,d}	70 ± 13 ^d	69 ± 11 ^d	70 ± 12 ^d	73 ± 11	<0.001
Heart rate stress (b.p.m.)	103 ± 25	99 ± 21 ^{a,e}	106 ± 28	108 ± 28 ^b	102 ± 26	103 ± 20	102 ± 18	<0.001
HRR	1.53 ± 0.37	1.52 ± 0.31 ^{d,e}	1.56 ± 0.43 ^{b,c,d}	1.56 ± 0.42 ^{b,c}	1.48 ± 0.34	1.48 ± 0.28	1.40 ± 0.19	<0.001
E positivity	1320 (37)	0	375 (35)	342 (56)	230 (66)	201 (76)	172 (100)	<0.001

Values are expressed as mean ± standard deviation, or n (%), except B-lines, expressed as median (range).

CFVR, coronary flow velocity reserve; DBP, diastolic blood pressure; EDV, end-diastolic volume; EF, ejection fraction; ESV, end-systolic volume; HRR, heart rate reserve; LAD, left anterior descending coronary artery; LVCR, left ventricular contractile reserve; SBP, systolic blood pressure; WMSI, wall motion score index.

^aP < 0.05 vs. ABCDE score 2.

^bP < 0.05 vs. ABCDE score 3.

^cP < 0.05 vs. ABCDE score 4.

^dP < 0.05 vs. ABCDE score 5.

^eP < 0.05 vs. ABCDE score 1.

the multivariable analysis, ABCDE-SE was an independent predictor of mortality with scores 3 (HR 3.472, 95% CI 1.483–8.135; $P = 0.004$), 4 (HR 4.045, 95% CI 1.595–10.259; $P = 0.004$), and 5 (HR 5.678, 95% CI 2.106–15.313; $P < 0.001$). Annual mortality rate ranged from 0.4% person-year for score 0 up to 2.7% person-year for score 5. Survival was worse in patients with ABCDE score 5 and best in patients with ABCDE score 0 (Figure 5).

In the incremental analysis, global X^2 of clinical model for the prediction of death increased from 49.5 considering clinical variables positive at univariate analysis to 50.7 after step A only ($P = \text{NS}$), and 80.6 after BCDE steps ($P < 0.001$ vs. step A). The increase over clinical evaluation was not significant after step A (after considering clinical

variables) and step C (after considering step B) and significant for steps B, D, and E (Figure 6).

Adding ABCDE-SE, risk reclassification was significantly improved with net reclassification index: 23.4% (95% CI 0.26–46.5%; $P = 0.048$). Integrated discrimination index was 0.51% (95% CI 0.01–0.94%; $P = 0.020$).

Discussion

A higher ABCDE-SE score indicates a less benign outcome. The risk is extremely low when all biomarkers are normal. In the individual

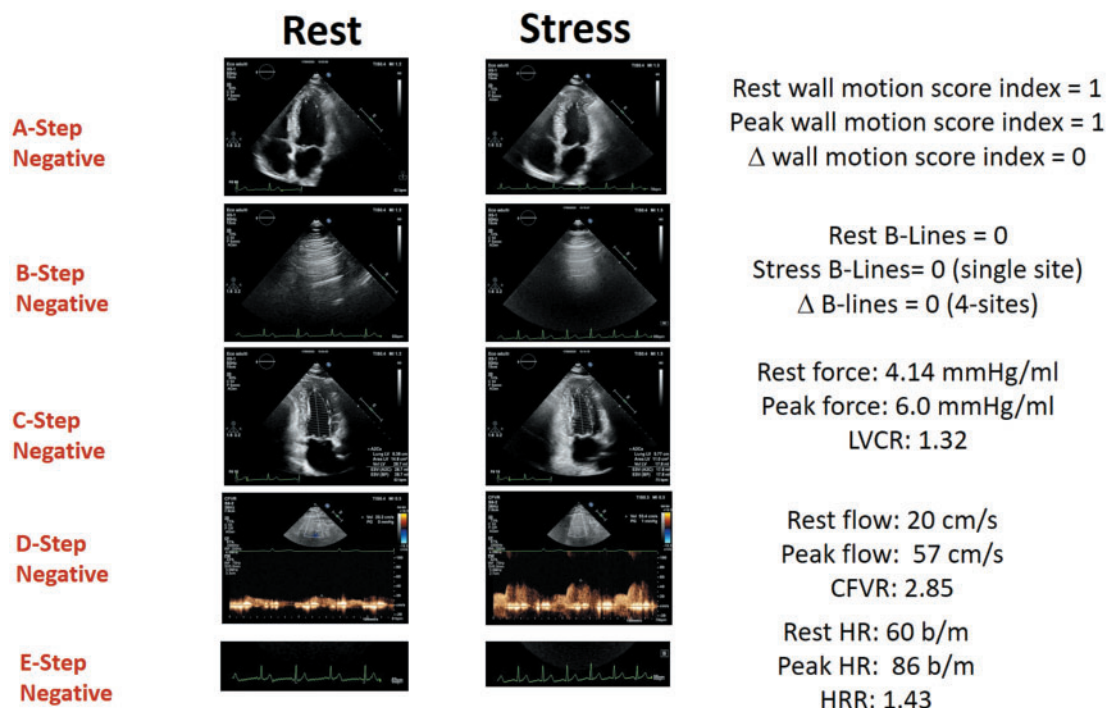


Figure 2 A normal ABCDE study. Left column: rest images. Right column, stress images. From top to bottom: A step: normal wall motion at rest and during stress; B step: normal A lines in lung ultrasound at rest and during stress; C step: reduced end-systolic volume and increased force during stress; D step: normal increase of pulsed-wave Doppler peak diastolic flow: $56/20 = 2.85$; E step: normal heart rate increase on electrocardiogram ($86/60 = 1.43$). Stress modality: dipyridamole. CFVR, coronary flow velocity reserve; HR, hazard ratio; HRR, heart rate reserve; LVCR, left ventricular contractile reserve.

biomarker analysis, the more established step A is also the less useful, outperformed by steps B, C, D and E for predicting all-cause death. Prognostically meaningful indices such as HRR and B lines are also very simple to obtain and analyse. Step B requires basic 2D technology and a minimal training, acquisition, and analysis time. Step E is imaging independent and automatically read in real time on the monitor of the echocardiographic machine. The value of ABCDE approach is coherent with the current understanding of the pathophysiological complexity of CAD. The individual patient with CAD may have multiple vulnerabilities, which are ignored, and missed, by the conventional step A approach. With ABCDE, the focus is shifted from coronary stenoses to the patient as a whole, with a simultaneous assessment of multiple vulnerabilities, from coronary microvascular dysfunction¹⁹ to cardiac autonomic unbalance.²⁰ It is conceivable that an SE-driven tailored therapy might be effective in improving prognosis through targeted efforts against the abnormal steps, which may trigger specific countermeasures. ABCDE-SE changes the binary step A response in a refined stratification of the overall level of risk, with a better identification of the prevailing phenotype coming into play in a specific patient.

Comparison with previous studies

The findings of the present study are consistent with previous reports showing the diagnostic and prognostic values of step B,^{5,21} step C,⁶ step D,^{8,22} and step E^{9–11} separately performed. The low positivity

rate with ischaemia by SE or myocardial perfusion imaging has been observed in recent years, possibly due to a change in population characteristics.²³ The prognostic value of various steps has also been extensively shown to be additive to stress-induced RWMA for step B,⁵ step C,⁶ step D,⁸ and step E.²⁴ Step D offers independent information beyond steps A, C, and E.^{8,24} The results of the present study confirm and extend previous evidence since all steps were simultaneously assessed in the same population allowing a separate analysis for all-cause death.

The test positivity based on RWMA did not predict mortality.⁶ Most patients are now studied under therapy that masks inducible ischaemia. In addition, step A is the only parameter embedded in general cardiology guidelines² and is utilized by referring cardiologists to decide ischaemia-driven revascularization to improve symptoms and possibly, in some subsets, survival.

Clinical implications

Cardiac functional testing with ABCDE-SE allows gaining of a comprehensive insight on patient vulnerability still with an extraordinarily simple and feasible test with low cost, minimal risk, zero radiation, and near-zero environmental impact. Therefore, it has the ideal features of economic, radiological, and environmental sustainabilities that are especially important for health care in the COVID-19 era. The protocol can be applied with all stresses, and this is especially important since exercise can be easily replaced with pharmacological

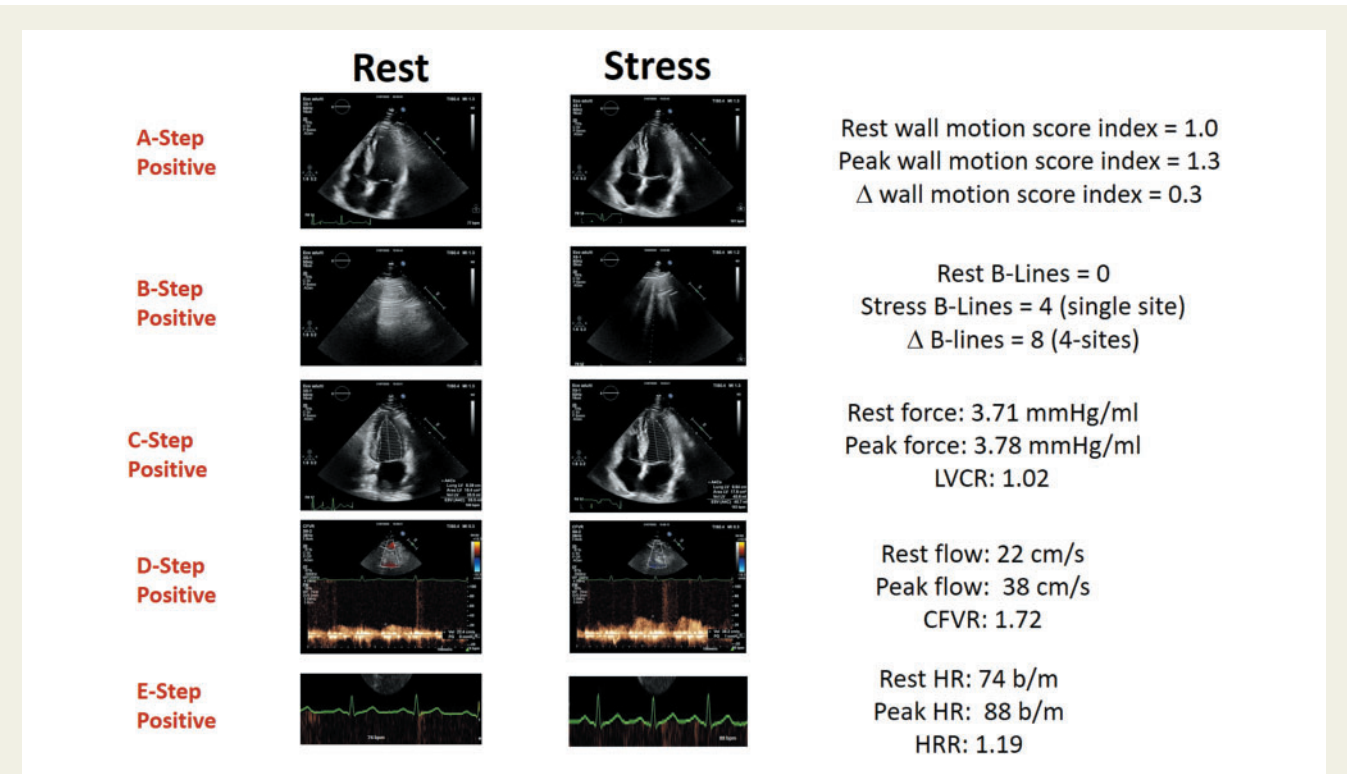
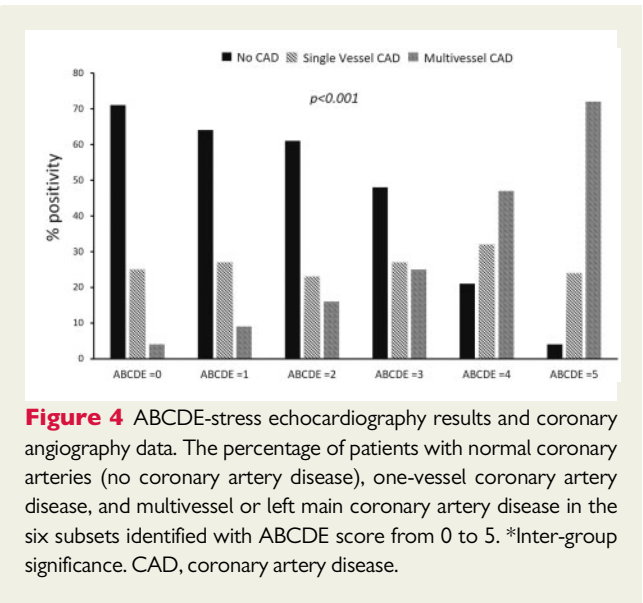


Figure 3 An abnormal ABCDE study. Left column: rest images. Right column, stress images. From top to bottom: A step: normal wall motion at rest and abnormal wall motion of apical region during stress; B step: no B lines at rest, four B lines in lung ultrasound during stress; C step: dilated end-systolic volume and blunted force reserve during stress; D step: reduced increase of pulsed-wave Doppler peak diastolic flow—38/22 = 1.72; E step: blunted heart rate on electrocardiogram (87/75 = 1.16). Stress modality: dipyridamole. CFVR, coronary flow velocity reserve; HR, hazard ratio; HRR, heart rate reserve; LVCR, left ventricular contractile reserve.



test in periods of intense viral epidemiological pressure when pharmacological stress is preferred over exercise because of added safety concerns and needed personal protective equipment during exercise.²⁵

The use of the ABCDE protocol is not mainly intended for the primary detection of CAD, since epicardial artery stenosis is the target of step A. However, the presence of advanced ABCDE score, even with normal step A, points to more extensive anatomic CAD in patients with chronic coronary syndromes. Patients with chest pain and CAD may show pulmonary congestion, reduced LVCR, a reduced CFVR, and blunted HRR in the absence of RWMA. A similar pattern can be found, even more frequently, in patients with chest pain with angiographically normal coronary arteries. The reassuring view of the classical ischaemic cascade with RWMA as the early and often only marker of inducible ischaemia is clearly challenged by these findings.

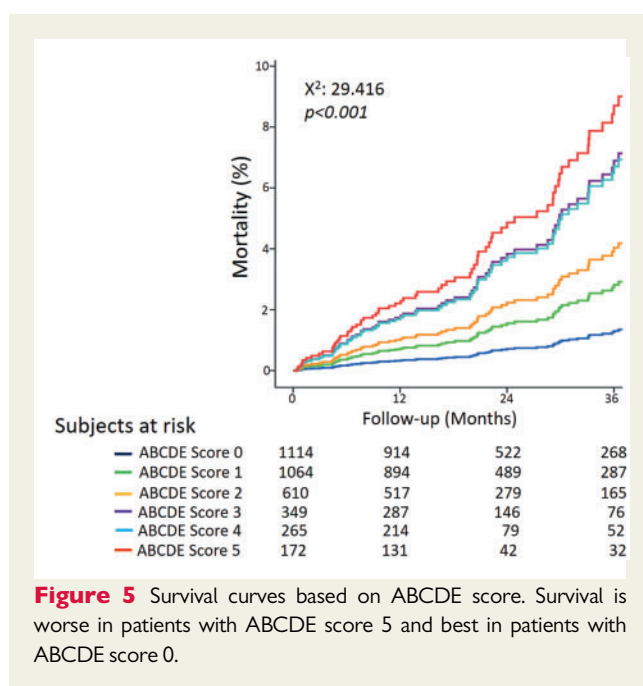
The main applications of ABCDE-SE are the identification of functional mechanisms of disease and symptoms, risk stratification, guide to therapy, or objective assessment of therapy efficacy. Each step identifies a specific phenotype, a biomarker of risk, and a potential selective target of tailored therapy.

An abnormal step A indicates anti-ischaemic therapy with beta-blockers, calcium channel blockers or nitrates, and possibly revascularization.² Diuretic therapy is recommended in patients with pulmonary congestion,² which is optimally identified and quantified with an abnormal step B after stress. Angiotensin-converting enzyme inhibitors are recommended in patients with asymptomatic left ventricular dysfunction after myocardial infarction,² but dysfunction is better identified with contractile reserve in step C. A selectively abnormal

Table 3 Predictors of all-cause mortality

Variables	Univariable Cox regression analysis		Multivariable Cox regression analysis	
	HR (95% CI)	P-value	HR (95% CI)	P-value
Age (years)	1.074 (1.047–1.101)	<0.001	1.065 (1.038–1.092)	<0.001
Male sex	1.968 (1.167–3.321)	0.011	1.964 (1.162–3.320)	0.012
Type of stress	0.759 (0.101–5.703)	0.789		
Exercise (reference)	1			
Dipyridamole	0.859 (0.495–1.491)	0.588		
Dobutamine	0.741 (0.332–1.654)	0.464		
Adenosine	1.317 (0.175–9.890)	0.789		
Hypertension (%)	1.547 (0.849–2.820)	0.154		
Beta-blockers therapy (%)	1.016 (0.631–1.637)	0.948		
Diabetes (%)	2.562 (1.613–4.070)	<0.001	1.946 (1.221–3.103)	0.005
Prior MI	1.280 (0.785–2.087)	0.322		
Reduced EF (<50%)	1.866 (1.041–3.343)	0.03		
A positivity	1.333 (0.731–2.430)	0.349		
B positivity	2.621 (1.654–4.152)	<0.001		
C positivity	1.581 (0.997–2.506)	0.051		
D positivity	2.578 (1.624–4.093)	<0.001		
E positivity	2.955 (1.848–4.725)	<0.001		
ABCDE score 0	1		1	
ABCDE score 1	2.182 (0.987–4.824)	0.054	1.798 (0.811–3.988)	0.149
ABCDE score 2	2.974 (1.301–6.797)	0.010	2.187 (0.934–5.043)	0.066
ABCDE score 3	5.438 (2.352–12.573)	<0.001	3.472 (1.483–8.135)	0.004
ABCDE score 4	5.289 (2.095–13.354)	<0.001	4.045 (1.595–10.259)	0.004
ABCDE score 5	6.923 (2.573–18.625)	<0.001	5.678 (2.106–15.313)	<0.001

CI, confidence interval; EF, ejection fraction; HR, hazard ratio; MI, myocardial infarction.



step D suggests the presence of coronary microvascular disease, and statins are recommended.² Abnormal step E implies a reduced cardiac sympathetic reserve, potentially targetable with a variety of possible therapies reducing the overactive sympathetic nervous system through the blockade of the beta-adrenergic or renin-angiotensin-aldosterone systems, or neuromodulation therapy aimed to restore autonomic balance through a sympatho-modulatory intervention, such as for instance renal denervation.²⁶ The overall global ABCDE risk score may be important to titrate therapy for event prevention.

Study limitations

Results were accessible to the referring physician, but the information beyond step A is unlikely to have affected subsequent decision-making since no evidence was available to support a change in management based on steps B–E.

Stress echocardiography does not tolerate improvisation, and a dedicated training is required to start ABCDE activity. However, the additional training and technology requirements are minimal compared to standard conventional SE, and analysis of wall motion and volumes is rapidly becoming operator independent and objective in the era of artificial intelligence.²⁷ The most difficult technical hurdle is

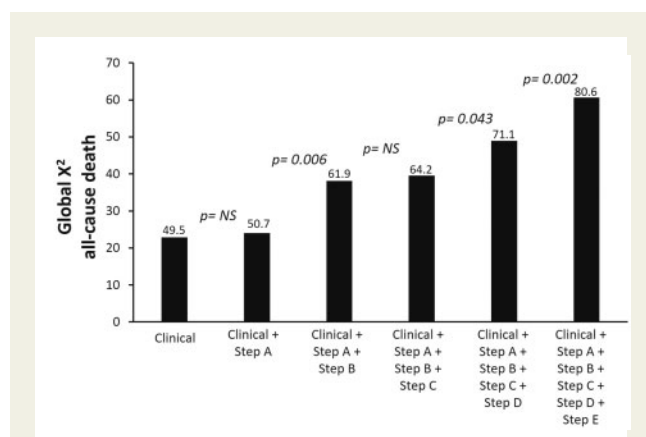


Figure 6 Incremental prognostic value of ABCDE positivity. The incremental value of ABCDE on global X² for predicting all-cause death was non-significant for step A (after considering clinical variables) and step C (after considering step B) and significant for steps B, D, and E. LVEF, left ventricular ejection fraction. NS, ***.

the acquisition of step D for LAD flow. This parameter was recommended with vasodilator stress echo since 2009¹⁷ and is already present in the 2019 European Society of Cardiology guidelines on chronic coronary syndromes for the assessment of coronary microvascular dysfunction in patients with ischaemia and normal coronary arteries.² The next step for those willing to try and start is to put together this information in the clinical arena and to build their own experience in their own patients.

The original statistical plan was to include 5000 patients with 3-year follow-up for mortality analysis, but as already described for many studies, COVID-19 pandemic abruptly slowed the recruitment everywhere at the beginning of 2020,²⁸ which was the planned last year of the SE 2020 study.

The outcome endpoint was all-cause death, which is a methodologically sound and clinically meaningful indicator but does not allow a separate analysis for different causes of death. In practice, the endpoint death is extremely robust, while the adjudication of cardiac or non-cardiac death, and even more sudden or non-sudden cardiac death, can be more problematic.²⁹ More granular endpoints may be preferentially predicted by individual biomarkers: step A predicts unstable angina;^{3,17} step B predicts acute decompensated heart failure;⁵ step C predicts progression of dyspnoea class and left ventricular dilatation;⁶ step D predicts myocardial infarction;⁸ and step E predicts electrical instability and arrhythmias.³⁰

The echocardiographic images were not adjudicated by a core lab but at each recruiting centre. This allowed a substantial sparing of resources and was also a methodological prerequisite for an effectiveness study, mirroring real-life conditions with unselected patients. A mandatory web-based quality control was implemented for conventional and innovative parameters upstream to patient recruitment since the volume of activity is necessary but not sufficient to ensure the quality of reading.¹⁴

The percentage of exercise, vasodilator, and dobutamine is different in the different subgroups, and this may influence at least the extent of positivity, although we found that the rate of positivity was similar with different stresses when stress-specific cut-off values are

used.⁵ The score 0–5 underuses the potential of stratification since every single letter used to build the score, from A to E, is analysed as a binary (positive vs. negative) response but is amenable to further titration of severity, from mild to moderate to severe impairment. These aspects are certainly important and will likely be addressed in the future with larger sample sizes, longer follow-up, and artificial intelligence-based data analysis as planned in the SE 2030 study.

The overall score generated by the ABCDE protocol is useful to identify the level of global risk, with higher scores requiring more aggressive treatment and closer surveillance. On top of this methodological and conceptual framework, further steps can be added in specific patients, from mitral regurgitation (step F) for valvular or ischaemic heart disease to step G (for gradients) in hypertrophic cardiomyopathy up to step R (for right ventricular function) in valvular and some forms of congenital heart disease.

Conclusions

ABCDE-SE allows assessment of multiple different vulnerabilities of the patient above and beyond epicardial artery stenosis. Vulnerabilities include pulmonary congestion, reduced contractile reserve, coronary microvascular dysfunction, and cardiac autonomic unbalance. This spectrum of phenotypes identifies the individual risk more accurately than inducible ischaemia and RWMA. Each step is important in modulating outcome and is a potential selective therapeutic target, paving the way to personalized assessment of risk and tailored therapy approaches.

Supplementary material

Supplementary material is available at *European Heart Journal* online.

Funding

The study was partially funded by CNR-MIUR (National Research Council, Italian Ministry of University and Research) Ageing subproject (Progetto P001328, Progetto di Interesse-Invecchiamento).

Conflict of interest: none declared.

Data availability

The data presented in this study are available on request from the principal investigator (QC). The data are not publicly available due to privacy.

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