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Myocardial Viability and Survival in Ischemic Left Ventricular Dysfunction

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Abstract

BACKGROUND—The assessment of myocardial viability has been used to identify patients with coronary artery disease and left ventricular dysfunction in whom coronary-artery bypass grafting (CABG) will provide a survival benefit. However, the efficacy of this approach is uncertain.

METHODS—In a substudy of patients with coronary artery disease and left ventricular dysfunction who were enrolled in a randomized trial of medical therapy with or without CABG, we used single-photon-emission computed tomography (SPECT), dobutamine echocardiography, or both to assess myocardial viability on the basis of pre-specified thresholds.

RESULTS—Among the 1212 patients enrolled in the randomized trial, 601 underwent assessment of myocardial viability. Of these patients, we randomly assigned 298 to receive medical therapy plus CABG and 303 to receive medical therapy alone. A total of 178 of 487 patients with viable myocardium (37%) and 58 of 114 patients without viable myocardium (51%) died (hazard ratio for death among patients with viable myocardium, 0.64; 95% confidence interval [CI], 0.48 to 0.86; $P = 0.003$). However, after adjustment for other baseline variables, this association with mortality was not significant ($P = 0.21$). There was no significant interaction between viability status and treatment assignment with respect to mortality ($P = 0.53$).

CONCLUSIONS—The presence of viable myocardium was associated with a greater likelihood of survival in patients with coronary artery disease and left ventricular dysfunction, but this relationship was not significant after adjustment for other baseline variables. The assessment of myocardial viability did not identify patients with a differential survival benefit from CABG, as compared with medical therapy alone. (Funded by the National Heart, Lung, and Blood Institute; STICH ClinicalTrials.gov number, NCT00023595.)

Coronary artery disease is an important contributor to the rise in the prevalence of heart failure and in associated mortality and morbidity.¹⁻⁴ It has not been clearly established whether coronary-artery bypass grafting (CABG) has a role in improving the symptoms and

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*A list of the investigators in the Surgical Treatment for Ischemic Heart Failure (STICH) trial is provided in the Supplementary Appendix, available at NEJM.org.

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the rate of survival of patients with coronary artery disease and heart failure. We conducted the multicenter Surgical Treatment for Ischemic Heart Failure (STICH) trial^{5,6} to examine two hypotheses, one of which (hypothesis 1) compared the efficacy of medical therapy alone with that of medical therapy plus CABG in patients with coronary artery disease and left ventricular dysfunction.

Left ventricular dysfunction in patients with coronary artery disease is not always an irreversible process related to previous myocardial infarction, since left ventricular function improves substantially in many patients and may even normalize after CABG.⁷⁻⁹ The assessment of myocardial viability with the use of single-photon-emission computed tomography (SPECT) or low-dose dobutamine echocardiography is commonly performed to predict improvement in left ventricular function after CABG, and numerous studies have suggested that the identification of viable myocardium with the use of such methods also predicts improved survival after CABG.^{8,10-33} However, previous studies that have suggested an association between myocardial viability and outcome have been retrospective in nature, and it is uncertain in most of these studies whether the decision to perform CABG may have been driven by the results of the tests, whether adjustment for key baseline variables was adequate, and whether patients who did not undergo CABG received aggressive medical therapy for heart failure.³⁴ In this substudy, we report the outcome of patients who were randomly assigned to receive medical therapy alone or medical therapy plus CABG in the hypothesis 1 component of STICH study and who also underwent assessment of myocardial viability.

METHODS

STUDY DESIGN

The rationale and design of the STICH trial have been described previously.⁵ We conducted a multicenter, nonblinded, randomized trial that was funded by the National Heart, Lung, and Blood Institute. The hypothesis 1 comparison involved 99 sites in 22 countries. Details of the study design for the hypothesis 1 comparison are reported in the study by Velazquez et al. in this issue of the *Journal*.⁶ The trial protocol, which was approved by the ethics committee at each study center, is available with the full text of this article at NEJM.org. The authors of this report assume responsibility for the completeness and accuracy of the data and the analyses and for the fidelity of the study to the trial protocol.

STUDY PATIENTS

Patients with angiographic documentation of coronary artery disease amenable to surgical revascularization and with left ventricular systolic dysfunction (ejection fraction, $\leq 35\%$) were eligible for enrollment. Exclusion criteria included left main coronary artery stenosis of more than 50%, cardiogenic shock, myocardial infarction within 3 months, and a need for aortic-valve surgery. All patients provided written informed consent.

Patients were randomly assigned to receive medical therapy alone or medical therapy plus CABG. A “risk at randomization” (RAR) score was calculated for each patient with the use of an equation derived in an independent data set from multiple variables with a known power to predict the 5-year risk of death without CABG.³⁵ Details of the RAR score are provided in the Supplementary Appendix, available at NEJM.org.

STUDY PROCEDURES

In the initial design of the STICH trial, viability testing with SPECT was required for the enrollment of patients.⁵ However, this requirement proved to be an impediment to enrollment. Therefore, the protocol was subsequently revised to make viability testing

optional and to allow the use of either SPECT or dobutamine echocardiography for viability testing. Investigators at all study centers were strongly encouraged to perform viability testing in every patient, but the decision to perform the test was left up to the recruiting investigators. Therefore, only a subgroup of patients in the hypothesis 1 component of the trial underwent viability testing. Details regarding the selection of patients for imaging are described in the Supplementary Appendix.

Independent core laboratories that were funded by the National Heart, Lung, and Blood Institute in which investigators were unaware of study-group assignments and the individual characteristics of patients coordinated data collection and analysis for the SPECT and dobutamine echocardiography studies. Details of the imaging protocols for identifying and quantifying the extent of viable myocardium are provided in the Supplementary Appendix. Briefly, thresholds of the extent of viable myocardium were prespecified to classify patients in a binary fashion as either having or not having substantial myocardial viability. For SPECT, patients with viability were defined as those with 11 or more viable segments on the basis of relative tracer activity. For dobutamine echocardiography, patients with viability were defined as those with 5 or more segments with abnormal resting systolic function but manifesting contractile reserve during dobutamine administration. Core laboratory measurements were submitted to the Duke Clinical Research Institute, which performed all statistical analyses.

PATIENT FOLLOW-UP AND OUTCOMES

After trial enrollment, patients were followed every 4 months for the first year and every 6 months thereafter. The primary outcome was death from any cause. Secondary end points included death from cardiovascular causes and a composite of death from any cause or hospitalization for cardiovascular causes. Definitions of the trial end points are provided in the report on the main study by Velazquez et al.⁶ All end points were adjudicated by an independent clinical events committee.⁶ The comparisons of outcomes that were related to treatment were based on intention-to-treat analyses. Analyses that were based on actual treatment received were also performed to account for crossovers.⁶

STATISTICAL ANALYSIS

We used means and standard deviations to summarize the baseline clinical characteristics of patients unless otherwise specified. We used the Wilcoxon rank-sum test, the Chi-square test, or Fisher's exact test, as appropriate, to assess baseline differences in individual variables between patients who underwent viability testing and those who did not undergo such testing and between patients who underwent testing who met the prespecified criteria for viability and those who did not meet such criteria. To further describe and characterize these differences, we developed multivariable propensity models using logistic regression to identify the key baseline clinical characteristics that distinguished patients who underwent viability testing from those who did not undergo such testing. Among patients who underwent viability testing, we also used these models to distinguish patients with viable myocardium from those without viable myocardium.

We used the Cox proportional-hazards model to assess the relationship between the presence of viable myocardium and the end point of death from any cause. We compared the strength of the univariate relationship between viability and the rate of death with the strength of other known prognostic factors, including the left ventricular ejection fraction, the left ventricular end-systolic and end-diastolic volume indexes, and the RAR score. Finally, we used multivariable Cox model analyses to assess the relationship between viability and each end point with adjustment for other known prognostic factors, including age, sex, race or ethnic group, heart failure class at baseline, history of myocardial infarction, previous

revascularization, ejection fraction, number of diseased vessels, chronic renal insufficiency, mitral regurgitation, history of stroke, and history of atrial fibrillation.

To test whether patients in the CABG group had a better outcome than those in the medical-therapy group among patients with viable myocardium, as compared with those without viable myocardium, we used Kaplan–Meier curves to examine the outcomes in each study group, according to viability status. We used the Cox regression model to test for an interaction between treatment and viability status. We used three separate prespecified analyses to assess the association between myocardial viability and outcome according to study-group assignment (for details, see the Supplementary Appendix).

RESULTS

PATIENTS

Of 1212 patients enrolled in the hypothesis 1 comparison, 601 who underwent assessment of myocardial viability were included in the analysis reported here. The numbers of patients undergoing each viability test, the timing of the tests, and baseline characteristics of the patients are detailed in the Supplementary Appendix. Among the 601 patients, 487 were found to have viable myocardium on the basis of the prespecified criteria, and 114 were found not to have viable myocardium. In the subgroup of 487 patients with myocardial viability, 244 were assigned to receive medical therapy plus CABG, and 243 were assigned to receive medical therapy alone. Likewise, in the subgroup of 114 patients without myocardial viability, 54 were assigned to receive medical therapy plus CABG, and 60 were assigned to receive medical therapy alone. The baseline characteristics of patients who were assigned to undergo CABG or to receive medical therapy were similar in each subgroup (Table 1, and Table S6 in the Supplementary Appendix).

OUTCOMES

During a median of 5.1 years of follow-up of 601 patients, there were 236 deaths (39%). These deaths included 58 of 114 patients without myocardial viability (51%) and 178 of 487 patients with myocardial viability (37%). Patients with viable myocardium had lower overall rates of death than those without viable myocardium (hazard ratio among patients with viable myocardium, 0.64; 95% confidence interval [CI], 0.48 to 0.86; $P = 0.003$) (Fig. 1). However, after adjustment for other significant baseline prognostic variables in a multivariable model, the prespecified viability status was no longer significantly associated with the rate of death ($P = 0.21$) (Supplementary Appendix).

Patients with myocardial viability also had lower rates of the secondary end points of death from cardiovascular causes (hazard ratio, 0.61; 95% CI, 0.44 to 0.84; $P = 0.003$) and a composite of death or hospitalization for cardiovascular causes (hazard ratio, 0.59; 95% CI, 0.47 to 0.74; $P < 0.001$) (Fig. S1 in the Supplementary Appendix). The relationship between myocardial viability and death from cardiovascular causes was not significant on multivariable analysis ($P = 0.34$), but the relationship with the composite of death or hospitalization for cardiovascular causes remained significant ($P = 0.003$).

There was no significant interaction between myocardial viability and study-group assignment with respect to death ($P = 0.53$) (Fig. 2), death from cardiovascular causes ($P = 0.70$), or the composite of death or hospitalization for cardiovascular causes ($P = 0.39$) on the basis of Kaplan–Meier and Cox model analyses. Additional prespecified analyses that were based on median viability scores or on a continuous model of viability and risk also revealed no significant interactions. This was the case whether the results were examined for all patients who had undergone viability testing, those with SPECT data alone, or those with dobutamine echocardiography data alone (Table S8 in the Supplementary Appendix).

Analysis of outcomes on the basis of the treatment received rather than that assigned showed similar trends, again reflecting no interaction between viability status and treatment with respect to death from any cause ($P = 0.96$), death from cardiovascular causes ($P = 0.26$), or death or hospitalization for cardiovascular causes ($P = 0.98$) (Fig. S3 and Table S9 in the Supplementary Appendix).

DISCUSSION

In this substudy of the STICH trial, in which all patients were eligible for CABG as well as optimal medical therapy, we analyzed patients for whom data were available with respect to myocardial viability in order to determine whether the presence of viable myocardium had an influence on the outcome. On univariate analysis, there was a significant association between myocardial viability and outcome. However, this association was not significant on multivariable analysis that included other prognostic variables. The findings of this multivariable analysis do not necessarily indicate that myocardial viability does not have pathophysiological importance in patients with coronary artery disease and left ventricular dysfunction. Instead, it is likely that some of the other variables in the analysis (e.g., left ventricular volumes and ejection fraction) are causally determined by the extent of viable myocardium.

A second, and more important, objective of this substudy was to determine whether the presence of substantial myocardial viability influenced the likelihood of benefit from medical therapy plus CABG, as compared with medical therapy alone. We did not find a significant interaction between myocardial viability and medical versus surgical treatment with respect to the rates of death from any cause or from cardiovascular causes or the rate of death or hospitalization for cardiovascular causes. This was true whether patients were grouped according to the assigned treatment (i.e., intention-to-treat analysis) or to the treatment actually received.

Conclusions that can be drawn from our results are limited by a number of factors. First, viability data were not available for all the patients who were enrolled in the STICH hypothesis 1 comparison. The study patients represent slightly less than 50% of the randomized group. Furthermore, viability testing was not performed on a randomly selected subgroup of patients but, rather, was obtained according to test availability and the judgment of the recruiting investigator. The differences in baseline characteristics between patients who underwent viability testing and those who did not undergo such testing suggest that at least some patients may have been selected for testing on the basis of clinical factors (Table S2 in the Supplementary Appendix).

Second, only 114 of 601 patients who underwent assessment of myocardial viability (19%) were deemed not to have viable myocardium on the basis of our prespecified criteria. This small number limited the power of our analysis to detect a differential effect of CABG, as compared with medical therapy, in patients with myocardial viability, as compared with those without myocardial viability, although in additional analyses that were based on median viability scores or that used a continuous model of viability, no effect of viability was detected.

Third, we cannot exclude the possibility that results of viability testing could have influenced subsequent clinical decision making. There was a nonsignificant trend toward higher rates of surgery among patients who underwent viability testing on the day of randomization or on the subsequent day than among those who underwent such testing before randomization. However, the timing of viability testing relative to randomization

does not appear to have influenced the rate of patient crossovers (Table S3 in the Supplementary Appendix).

Fourth, our analysis was based on SPECT and dobutamine echocardiography assessment of myocardial viability. An analysis of outcomes on the basis of a combination of these two tests poses important limitations, given the fundamental differences in the viability information provided by SPECT and dobutamine echocardiography (one related to membrane integrity and the other to contractile reserve) and the differences in analytic approaches between the two methods. However, results were similar whether SPECT and dobutamine echocardiography data were combined or analyzed separately. We also did not incorporate other approaches, such as positron-emission tomography (PET)^{36,37} or contrast-enhanced magnetic resonance imaging (MRI).^{38,39} However, in a meta-analysis and other reviews, SPECT and dobutamine echocardiography have been found to have similar prognostic potential to that of PET,^{8,40,41} and there are limited data regarding outcomes in patients with chronic ischemic left ventricular dysfunction who were studied on MRI.

The lack of a differential survival benefit between CABG and medical therapy in patients with viable versus those with nonviable myocardium in this study differs markedly from results of previous retrospective studies and meta-analyses.^{8,10-33} This finding may reflect, in part, the low rates of death among patients with viable myocardium who were assigned to receive medical therapy (approximately 7% per year) in this study, as compared with previously reported rates (which exceeded 15% per year in many studies). Adherence to guidelines-recommended therapies was high in our trial, whereas data on medical therapy are lacking in many previous retrospective analyses. Medical therapy, like CABG, has the potential to improve left ventricular function in patients with dysfunctional but viable myocardium.⁴²⁻⁴⁴ This finding underscores the prognostic importance of providing evidence-based therapies to high-risk patients with left ventricular dysfunction, as well as the consideration of CABG for those who are candidates for revascularization. The lack of interaction between myocardial-viability status and benefit from CABG in this study indicates that assessment of myocardial viability alone should not be the deciding factor in selecting the best therapy for these patients. These findings also highlight the need for prospectively designed studies to determine the role of cardiac imaging in clinical decision making.⁴⁵

In summary, we conducted a substudy of the STICH trial to determine whether the presence of substantial myocardial viability influenced the likelihood of benefit from medical therapy plus CABG, as compared with medical therapy alone, in patients with coronary artery disease and left ventricular dysfunction. We did not find a significant interaction between myocardial-viability status and medical versus surgical treatment with respect to the rates of death from any cause or from cardiovascular causes or the rate of death or hospitalization for cardiovascular causes.

Supplementary Material

Refer to Web version on PubMed Central for supplementary material.

APPENDIX

Northwestern University Feinberg School of Medicine, Chicago (R.O.B., T.A.H.); Medical University of Vienna, Vienna (G.M.); Duke Clinical Research Institute, Duke University Medical Center, Durham, NC (K.L.L., L.S., E.J.V., R.H.J.); Ohio State University Medical Center, Columbus (P.F.B.); National Heart, Lung, and Blood Institute, Bethesda, MD (P.D.-N.); Medical University of Lodz, Lodz (J.D.), and John Paul II Hospital, Krakow (K.W.) —

both in Poland; Instituto Dante Pazzanese de Cardiologia (P.S.F.) and Instituto do Coração–Hospital das Clínicas da Faculdade de Medicina da Universidade de São Paulo (J.C.N.) — both in São Paulo; Jefferson Medical College, Philadelphia (A.M.F.); Heart Center, University of Leipzig, Leipzig, Germany (T.D.); Montefiore Medical Center/Albert Einstein College of Medicine, New York (R.E.M.); Cedars–Sinai Medical Center, Los Angeles (D.S.B.); Division of Cardiovascular Diseases, Mayo Clinic, Rochester, MN (P.A.P.); Zala County Hospital, Zalaegerszeg, Hungary (N.A.); Washington Hospital Center, Washington, DC (F.M.A., J.A.P.); and University Hospital Favalaro Foundation, Buenos Aires (L.E.F.)

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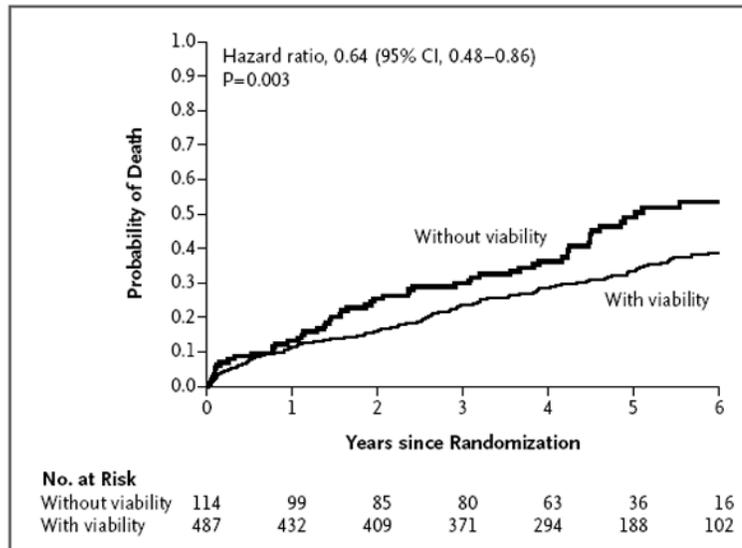


Figure 1. Kaplan–Meier Analysis of the Probability of Death, According to Myocardial Viability Status

The comparison that is shown has not been adjusted for other prognostic baseline variables. After adjustment for such variables on multivariable analysis, the between-group difference was not significant ($P = 0.21$).

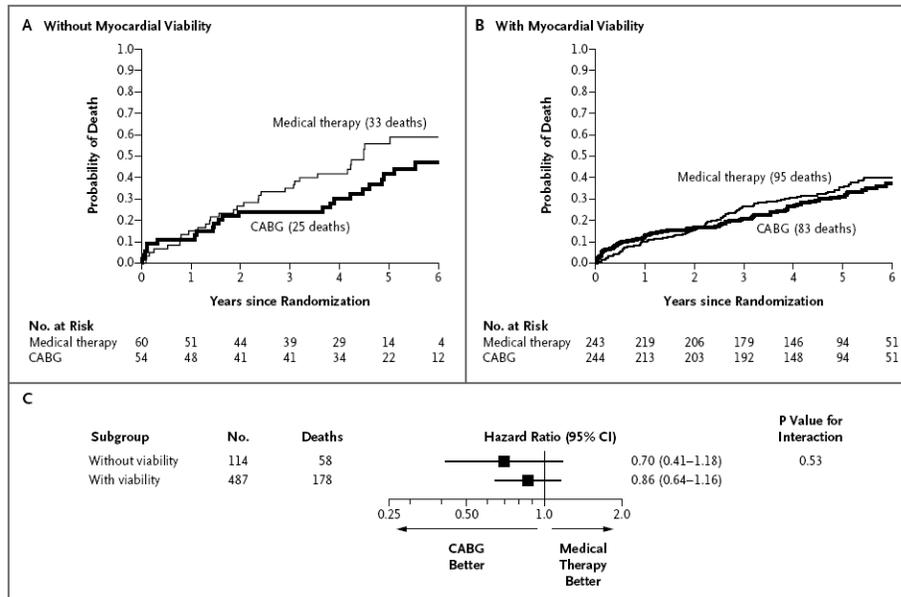


Figure 2. Kaplan–Meier Analysis of the Probability of Death According to Myocardial-Viability Status and Treatment

At 5 years in the intention-to-treat analysis, the rates of death for patients without myocardial viability were 41.5% in the group assigned to undergo coronary-artery bypass grafting (CABG) and 55.8% in the group assigned to receive medical therapy (Panel A). Among patients with myocardial viability, the respective rates were 31.2% and 35.4% (Panel B). There was no significant interaction between viability status and treatment assignment with respect to mortality ($P = 0.53$) (Panel C).

Table 1
 Baseline Characteristics of Patients Who Underwent Assessment of Myocardial Viability.*

Characteristic	All Patients (N = 601)	Patients with Myocardial Viability (N = 487)		Patients without Myocardial Viability (N = 114)		
		Medical Therapy (N = 243)	CABG (N = 244)	Medical Therapy (N = 60)	CABG (N = 54)	
Age — yr	60.7±9.4	60.0±9.7	61.5±9.2	61.6±8.5	60.0±9.2	0.34
Male sex — no. (%)	521 (87)	205 (84)	211 (86)	55 (92)	50 (93)	1.00
Previous myocardial infarction	481 (80)	190 (78)	183 (75)	56 (93)	52 (96)	0.68
Current Canadian Cardiac Society angina class — no. (%)						0.03
0	236 (39)	101 (42)	101 (41)	18 (30)	16 (30)	
I	94 (16)	34 (14)	34 (14)	19 (32)	7 (13)	
II	253 (42)	104 (43)	99 (41)	23 (38)	27 (50)	
III	14 (2)	3 (1)	8 (3)	0	3 (6)	
IV	4 (1)	1 (<1)	2 (1)	0	1 (2)	
Highest New York Heart Association functional class in 3 previous mo — no. (%)						0.25
I	27 (4)	15 (6)	9 (4)	0	3 (6)	
II	212 (35)	94 (39)	88 (36)	14 (23)	16 (30)	
III	275 (46)	100 (41)	111 (45)	36 (60)	28 (52)	
IV	87 (14)	34 (14)	36 (15)	10 (17)	7 (13)	
Risk-at-randomization score [†]	12.5±8.8	11.9±8.4	12.8±9.0	13.7±9.8	12.0±8.8	0.37
Medications at baseline — no. (%)						
Beta-blocker	534 (89)	221 (91)	216 (89)	52 (87)	45 (83)	0.62

Characteristic	All Patients (N = 601)		Patients with Myocardial Viability (N = 487)		Patients without Myocardial Viability (N = 114)	
	Medical Therapy (N = 243)	CABG (N = 244)	P Value	Medical Therapy (N = 60)	CABG (N = 54)	P Value
ACE inhibitor	514 (86)	210 (86)	0.37	54 (90)	48 (89)	0.85
ARB	46 (8)	20 (8)	0.99	3 (5)	3 (6)	1.00
ACE inhibitor or ARB	554 (92)	227 (93)	0.25	57 (95)	51 (94)	1.00
Statin	508 (85)	193 (79)	0.02	56 (93)	47 (87)	0.26
Aspirin	513 (85)	205 (84)	0.54	56 (93)	43 (80)	0.03
Coronary artery disease distribution — no. (%)						
No. of diseased vessels with $\geq 75\%$ stenosis			0.79			0.45
0	12 (2)	6 (2)		2 (3)	1 (2)	
1	152 (25)	62 (26)		17 (28)	11 (20)	
2	221 (37)	87 (36)		18 (30)	24 (44)	
3	215 (36)	88 (36)		23 (38)	18 (33)	
Left ventricular ejection fraction — %	26.7 \pm 8.6	28.1 \pm 8.4	0.30	22.6 \pm 8.5	23.3 \pm 9.1	0.50
Left ventricular end-diastolic volume index — ml/m ² of body-surface area	122.8 \pm 41.9	117.8 \pm 37.9	0.63	152.3 \pm 51.3	140.0 \pm 53.8	0.16
Left ventricular end-systolic volume index — ml/m ² of body-surface area	91.7 \pm 38.9	85.8 \pm 34.3	0.97	120.8 \pm 49.6	111.2 \pm 50.8	0.25

* Plus-minus values are means \pm SD. ACE denotes angiotensin-converting enzyme, ARB angiotensin-receptor blocker, and CABG coronary-artery bypass grafting.

[†]The risk-at-randomization score ranges from 1 to 32, with higher numbers indicating a higher predicted rate of death. Among patients receiving medical therapy, a score of 1 predicts a rate of 18% and a score of 32 predicts a rate of 99% over 5 years.