

Long-term outcomes after coronary artery bypass graft with or without surgical ventricular reconstruction in patients with severe left ventricular dysfunction

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Background: Patients with chronic myocardial infarction (MI) and severe left ventricular (LV) dysfunction have poor clinical outcomes. This study aimed to determine whether coronary artery bypass graft (CABG) with surgical ventricular reconstruction (SVR) leads to further improvement in long-term patient outcomes compared with isolated CABG (I-CABG).

Methods: From April 2010 to June 2013, 140 consecutive patients with chronic MI and severe LV dysfunction who received contrast-enhanced cardiovascular magnetic resonance imaging (CE-CMR) within 1 month before surgery were enrolled in this study. The cardiovascular events (CVEs) and long-term survival of patients who underwent CABG and SVR were compared with those who met the criteria for SVR but received I-CABG.

Results: A total of 140 patients were included in the final analysis, including 70 patients who underwent CABG and SVR and 70 patients who underwent I-CABG. No differences were observed in the baseline characteristics, LV function, and late gadolinium enhancement (LGE) between the two groups. CABG+SVR patients experienced a longer cardiopulmonary bypass (CPB) time (116.0±35.0 vs. 100.2±23.8 minutes, P=0.002) and ventilation time [median (interquartile range): 22.0 (17.0, 37.0) vs. 20.0 (15.0, 24.0) hours, P=0.019] than I-CABG patients. During a mean follow-up of 123.1±12.7 months (range, 102–140 months), the CABG+SVR group had fewer rehospitalizations for congestive heart failure (CHF) (4.3% vs. 19.1%, P=0.007), but no statistical difference in the mortality rate was observed (2.9% vs. 4.4%, P=0.987). The cumulative CVE-free survival rate was significantly higher in CABG+SVR patients (87.0% vs. 67.6%, P=0.007).

Conclusions: Our findings indicated that patients with chronic MI and severe LV dysfunction experienced similar perioperative outcomes after CABG+SVR or I-CABG. However, the CABG+SVR group resulted in fewer rehospitalizations for CHF and a higher cumulative CVE-free survival rate.

Keywords: Coronary artery bypass graft (CABG); contrast-enhanced cardiovascular magnetic resonance imaging (CE-CMR); left ventricular aneurysm; surgical ventricular reconstruction (SVR)

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Introduction

Although patients with chronic myocardial infarction (MI) and severe left ventricular (LV) dysfunction are at higher surgical risk, coronary artery bypass graft (CABG) is still the most widely applied technique to improve symptoms and prognosis in these patients (1-3). However, a small group of patients still develop LV dilatation after a maladaptive response to ischemic injury, leading to a spherically shaped LV and the formation of an aneurysm as well as the clinical syndrome of congestive heart failure (CHF). Surgical ventricular reconstruction (SVR) is a specific procedure developed for the management of CHF with LV remodeling caused by coronary artery disease (CAD) (4). Previous studies have demonstrated that this procedure restores the normal cardiac size and the elliptical shape of the heart. It has also been reported to reduce the LV volume and improve left ventricular ejection fraction (LVEF) and New York Heart Association (NYHA) class in these patients (5).

Although the benefit of SVR has been demonstrated, controversy remains as to whether CABG+SVR is superior to isolated CABG (I-CABG). Prucz et al. reported that SVR together with CABG might improve LV function to a greater degree than I-CABG and result in fewer rehospitalizations for CHF (6). However, the STICH trial concluded that despite CABG+SVR reducing the LV volume, this anatomical change was not associated with a greater improvement in symptoms or better survival (7). Prior et al. further reported that elective addition of SVR in patients undergoing CABG was not associated with a greater improvement in mortality (8). Although many previous studies sought to compare patients who underwent CABG and SVR to those who received I-CABG and different conclusions were drawn, only a few studies selected patients using a consistent imaging modality, and no studies provided a definitive answer about which kind of patients might benefit from SVR.

In recent years, contrast-enhanced cardiovascular magnetic resonance imaging (CE-CMR) has emerged as an accurate and non-invasive modality for the detection and quantification of myocardial scars and is commonly applied in the evaluation of patients with chronic MI and LV dysfunction to select appropriate treatment strategies. Therefore, it is the gold standard imaging modality for the assessment of not only LV function but also the transmurality of the scars (9). Based on the American Heart Association (AHA) 17-segmental model (10), our previous studies have demonstrated that the cardiac function of

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patients with >4 scar segments did not improve after I-CABG, and those with \geq 6 scar segments experienced a higher risk of cardiovascular events (CVEs) post-I-CABG (11,12). However, the question remains as to whether SVR plus CABG can further improve the prognosis of patients with a considerable amount of myocardial scarring. Using this cohort of patients, we aimed to compare the outcomes of patients who underwent CABG and SVR to those who received I-CABG. We present the following article in accordance with the TREND reporting checklist (available at https://jtd.amegroups.com/article/view/10.21037/jtd-22-1214/rc).

Methods

Study population

This study included 140 consecutive patients with chronic MI and severe LV dysfunction clinically referred for CE-CMR [LV function and myocardial late gadolinium enhancement (LGE)] within 1 month before first-time surgery from April 2010 to June 2013. Patients were enrolled based on the following criteria: (I) CAD with >70% stenosis in two or more major vessels scheduled for surgery; (II) dyspnea as the predominant symptom; (III) previous Q-wave MI on electrocardiogram (ECG) and a history of MI \geq 3 months before surgery; and (IV) LVEF of CMR \leq 35% and at least two adjacent segments with wall motion abnormalities at rest, and the presence of anterior akinesia or dyskinesia of the LV on CMR. Patients were excluded if they had any of the following conditions: (I) any prior cardiac surgical interventions or concomitant surgical procedures (mitral/aortic valve repair or replacement); (II) hypertrophic obstructive cardiomyopathy or myocarditis; and (III) contraindications for CE-CMR examination (non-compatible biometallic implants, allergy to contrast agents, claustrophobia, etc.). This study was conducted in accordance with the Declaration of Helsinki (as revised in 2013) and was approved by the institutional review board of Fuwai Hospital (No. 2010-259). All patients provided written informed consent.

An experienced surgical team was blinded to the patient identification and reviewed the CE-CMR of all subjects with LVEF \leq 35% to determine their eligibility for SVR. During surgical exploration, the team was more inclined to perform SVR in cases with the presence of anterior/anteroseptal MI (scar tissue) and dominant anterior/anteroseptal akinesia or dyskinesia of the LV (6,13-15).



Video 1 Coronary artery bypass graft plus surgical ventricular reconstruction for a patient with chronic myocardial infarction and severe left ventricular dysfunction. This patient was a 64-year-old man with six scar segments on CE-CMR and a left ventricular ejection fraction of 28.1%. The patient underwent coronary artery bypass graft plus surgical ventricular reconstruction and survived without cardiovascular events during the follow-up. CE-CMR, contrast-enhanced cardiovascular magnetic resonance imaging.

Consequently, patients were divided into two groups according to the surgical strategy: those who underwent I-CABG (70 cases) and those who underwent CABG and SVR (70 cases). Transthoracic echocardiography (TTE) was performed preoperatively, before discharge, at 3, 6, and 12 months after surgery, and annually thereafter.

Surgical technique

All surgeries were performed by the same surgical team. CABG aimed to obtain complete revascularization, which was technically possible and performed in all patients. The left internal mammary artery (LIMA) and great saphenous vein (GSV) were harvested from each patient and the quality of grafts was assessed intra-operatively using a transittime flow probe (TTFP; Medi-stim Butterfly flowmeter, Oslo, Norway). After surgery, all patients received standard pharmacotherapy for CAD.

In this study, cardioplegia was applied and I-CABG was performed under cardiac arrest. After aortic crossclamping, cardiac arrest was accomplished via the antegrade administering of cold blood cardioplegia (CBC) (4 °C) via the aortic root. Heart topical cooling was applied throughout the entire procedure. The CBC solution was a mixture of whole oxygenated blood and hyperkalemic solution at a 4:1 ratio. The initial induction dose was 15 to 20 mL/kg, and half of the initial dose was administered every 20 to 30 minutes. Additionally, the LIMA was anastomosed to the left anterior descending (LAD) branch, while the GSV was anastomosed to the obtuse marginal (OM) branch or posterior descending artery (PDA) branch.

For patients who underwent CABG and SVR, the SVR component was most commonly performed during a single period of cardioplegic arrest. The surgical indications for SVR included an anterior or anteroseptal MI (myocardial scar tissue), and the presence of dominant anterior akinesia or dyskinesia of the LV. In this procedure, after ventriculotomy is centered in the zone of anterior asynergy, a suture is placed in the interior of the ventricle to encircle the myocardial scar at the boundary between akinetic and viable tissue. The tightening of this suture brings the healthy myocardium together. The ventriculotomy defect was closed with a linear closure unless the incision was more than 2 to 3 cm, in which case a Dacron patch was used (16,17) (Figure S1 and Video 1). In our study, aneurysmectomy with linear repair was performed in 55 patients (78.6%), while circular reconstruction was performed in 15 patients (21.4%). After surgery, all patients received standard pharmacotherapy.

CMR protocol and imaging analysis

CMR was performed using a 1.5 Tesla scanner (Avanto, Siemens AG, Germany) according to a standardized scanning protocol. To evaluate the functional parameters, ECG-gated cine images were acquired with a steadystate free precession (SSFP) sequence in long-axis planes and contiguous short-axis slices from the atrioventricular ring to the apex as previously described (11). Ten to 15 minutes after intravenous injection of 0.2 mmol/kg of gadolinium-diethylenetriamine pentaacetic acid (Gd-DTPA) (gadopentetate dimeglumine, Magnevist, Bayer Healthcare Pharmaceuticals, Wayne, NJ, USA), LGE images were obtained using a phase-sensitive inversionrecovery gradient-echo pulse sequence in identical long-axis and short-axis planes.

The LGE evaluation and post-processing were performed using Argus software (Siemens AG, Munich, Germany). The cardiac function and LGE images were evaluated by two independent experienced radiologists (MJ Lu and SH Zhao) who were blinded to the clinical data using an identical 17-segment model. A five-point scale system was used to describe the transmural extent of LGE in each of the segments (scar score): 0 = no LGE,

1 = 1-25% LGE, 2 = 26-50% LGE, 3 = 51-75% LGE, and 4 = 76-100% LGE. If no agreement on the interpretations was reached, the image was reevaluated by two radiologists until a consensus was achieved. A cut-off value of 50% LGE was the optimal threshold to define segmental viability for predicting recovery of cardiac function (18). Additionally, the extent of scar tissue was quantified using the following definitions (18): (I) spatial extent, the number of affected segments; (II) normal segments, the number of segments with a scar score of 0; (III) viable segments, the number of segments with a scar score of 1 or 2; (IV) scar segments, the number of segments with a scar score of 3 or 4; and (V) total scar score (TSS), summation of the segmental scar scores for each patient. The severity of segmental wall motion was determined on a four-point scale system: 0 = normal, 1 = hypokinesis, 2 = akinesis, and 3 = dyskinesis. The wall motion score (WMS) was the summation of the WMSs for all 17 segments of the heart.

Outcomes and follow-up

The primary endpoint of the present study was a composite of CVEs defined as death from any cause, rehospitalization for CHF (a severe clinical syndrome characterized by the presence of dyspnea or limited exertion due to impaired cardiac ventricular filling or lowered cardiac contraction, leading to rehospitalization instead of staying at home), lifethreatening ventricular arrhythmia (VA), non-fatal MI, and severe angina pectoris (AP) (11).

The subjects were followed up regularly for CVEs from the first day after discharge to the latest follow-up via telephone contact with the patients or their relatives, outpatient visit, or medical records review. All causes of death, CVEs, and functional status of the patients were recorded in detail.

Statistical analysis

Statistical analysis was performed using SPSS version 20.0 (IBM Corp., Armonk, NY, USA) and SAS software (Version 9.4; SAS Institute Inc, Cary, NC). Continuous variables were presented as the mean \pm standard deviation (SD) or median (interquartile range). Categorical variables were reported as absolute numbers and percentages. Between-group comparisons were performed using the Chi-squared test or Fisher's exact test for categorical variables, and the Student's *t*-test or Mann-Whitney U test for continuous variables that were normally and non-normally distributed,

respectively. The improvement of LVEF and LV size as well as the effect of surgery were compared using a paired *t*-test. Survival curves were generated by the Kaplan-Meier method and compared by the log-rank test. All statistical tests were two-tailed, and P<0.05 was considered statistically significant.

Results

Study population

A total of 140 patients were enrolled and included in the final analysis (*Figure 1*). The baseline characteristics of the I-CABG group were well-matched to those of the CABG+SVR cohort, and no significant differences were observed between the two groups (*Table 1*). The mean age of patients in the I-CABG and CABG+SVR groups was 57.7±8.4 and 58.2±7.2 years, respectively (P=0.715). In addition, other baseline characteristics including hypertension, diabetes mellitus, and patients with NYHA class III/IV were similarly distributed between the two groups.

Cardiac surgery

The CABG+SVR group had significantly longer operation time and CPB time than the I-CABG group (P=0.021 and P=0.002, respectively). The duration of aortic crossclamping and the ventilation time were also longer for patients undergoing CABG and SVR (P=0.008 and P=0.019). The intensive care unit (ICU) duration, hospital stay, and number of grafts per patient were not significantly different between the two groups (P=0.064, P=0.340, and P=0.060, respectively). The procedural characteristics and outcomes of surgery are shown in *Table 2*.

In-hospital mortality was 2.9% in the I-CABG group (two patients) compared with 1.4% (one patient) in the CABG+SVR group (P=1.000). In the I-CABG group, there were two in-hospital mortalities: one owing to an exacerbation of CHF and the other owing to multiple organ dysfunction syndrome (MODS). In addition, the only one in-hospital mortality in the CABG+SVR group was attributable to MODS on postoperative day 9. The incidence of postoperative complications, including newonset atrial fibrillation (AF), reoperation for bleeding, perioperative MI, stroke, life-threatening VA, renal failure requiring dialysis, and low cardiac output syndrome (LCOS), were similarly distributed among the two groups (*Table 2*).



Figure 1 Flow diagram illustrating the study design and patient categorization. A total of 140 patients were finally enrolled and analyzed; 70 subjects underwent I-CABG and 70 subjects received CABG and SVR. CABG, coronary artery bypass graft; I-CABG, isolated coronary artery bypass graft; LVEF, left ventricular ejection fraction; SVR, surgical ventricular reconstruction.

CE-CMR results

No significant difference was observed between the I-CABG and CABG+SVR patients in terms of LV function or volume at baseline or LGE (including spatial extent, number of scar segments, viable segments, and TSS at baseline). The CMR results revealed more severe LV adverse remodeling in the CABG+SVR group, as evidenced by larger left ventricular end-diastolic diameter (LVEDD) and left ventricular end-systolic volume index (LVESVI), but this was only marginally statistically significant (P=0.055 and P=0.079, respectively). The detailed CE-CMR results are summarized in *Table 3*. Moreover, the typical CE-CMRs of two patients at baseline and 6 months after I-CABG or CABG+SVR are shown in Figures S2,S3.

Long-term outcomes

The average time for follow-up of all subjects was 123.1 ± 12.7 (range, 102-140) months. Both groups showed a significant improvement in LVEF by TTE. The mean LVEF improved from $37.5\%\pm7.5\%$ to $45.4\%\pm8.5\%$ in the

I-CABG group (P<0.001) compared with $35.9\% \pm 8.4\%$ to $48.1\% \pm 8.9\%$ in the CABG+SVR group (P<0.001) (*Figure 2A*). The change in LVEDD between 6 months and the latest follow-up in I-CABG patients was not significantly different, while that for patients who underwent CABG and SVR was significantly different (P=0.506 vs. P<0.001, respectively, *Figure 2B*). Likewise, the change of LVEF/LVEDD between baseline and follow-up in I-CABG patients was lower than that for CABG+SVR patients (P=0.002 and P<0.001, respectively, *Table 4*). In addition, CABG+SVR patients were more likely to have a $\geq 5\%$ increase in LVEF during follow-up (84.1% vs. 60.3%, P=0.002, *Table 4*).

From baseline to the latest follow-up, the proportion of patients with NYHA class I increased, while that of patients with NYHA class III/IV decreased (*Figure 3A*). Likewise, the proportion of patients without severe AP increased, while that of patients with Canadian Cardiovascular Society (CCS) class III/IV decreased (*Figure 3B*). Moreover, the NYHA class improved significantly from 2.7±0.7 to 1.9±0.9 (P<0.001) in the I-CABG group and from 2.8±0.7 to 1.5±0.7

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Variables	All patients (n=140)	I-CABG (n=70)	CABG+SVR (n=70)	P value
Age, years	58.0±7.8	57.7±8.4	58.2±7.2	0.715
Men	124 (88.6)	61 (87.1)	63 (90.0)	0.595
Hypertension	79 (56.4)	38 (54.3)	41 (58.6)	0.609
Diabetes mellitus	57 (40.7)	29 (41.4)	28 (40.0)	0.863
Hypercholesterolemia	76 (54.3)	40 (57.1)	36 (51.4)	0.497
COPD	23 (16.4)	12 (17.1)	11 (15.7)	0.820
Stroke history	13 (9.3)	5 (7.1)	8 (11.4)	0.560
Current smoker	105 (75.0)	51 (72.9)	54 (77.1)	0.558
Family history of CAD	59 (42.1)	34 (48.6)	25 (35.7)	0.123
Angiographic findings				0.217
Two-vessel lesions	19 (13.6)	7 (10.0)	12 (17.1)	
Three-vessel lesions	121 (86.4)	63 (90.0)	58 (82.9)	
NYHA class III/IV	89 (63.6)	42 (60.0)	47 (67.1)	0.380
LVEF (Echo), %	36.7±8.0	37.5±7.5	35.9±8.4	0.223
Mitral grade				
Mild	63 (30.7)	36 (51.4)	27 (38.6)	0.198
Moderate	20 (14.3)	7 (10.0)	13 (18.6)	0.198
Euroscore	6.0±2.4	5.8±2.2	6.2±2.5	0.238

Table 1 Baseline characteristics of the patients

Values are expressed as mean ± SD or n (%). CABG, coronary artery bypass graft; CAD, coronary artery disease; COPD, chronic obstructive pulmonary disease; Echo, echocardiography; I-CABG, isolated coronary artery bypass graft; LVEF, left ventricular ejection fraction; NYHA, New York Heart Association; SD, standard deviation; SVR, surgical ventricular reconstruction.

in the CABG+SVR group (P<0.001). At the latest followup, patients without angina were distributed similarly between these two groups (63.8% vs. 57.4%, P=0.616, *Figure 3B*). However, the CABG+SVR patients had a higher proportion of NYHA I/II class compared to the I-CABG patients (85.5% vs. 69.1%, P=0.030, *Figure 3A*).

In our analysis, no significant difference between the two groups in preoperative mitral grade was observed. Sixty-three patients had mild mitral valve regurgitation (MR) (36 in the I-CABG group and 27 in the CABG+SVR group), while 20 patients had moderate MR (seven in the I-CABG group and 13 in the CABG+SVR group). In these 83 subjects during follow-up, MR was reduced in 61 (73.5%) patients compared with that before the operation. In addition, the I-CABG patients were more likely to be rehospitalized for CHF compared to the CABG+SVR patients (19.1% *vs.* 4.3%, P=0.007). Three deaths occurred in the I-CABG group, while two deaths occurred in the

CABG+SVR group (4.4% vs. 2.9%, P=0.987). Patients who suffered VA, non-fatal MI, and AP recurrence were distributed similarly between the two groups (*Table 4*). Furthermore, the long-term cumulative CVE-free survival rate was significantly higher in the CABG+SVR patients (87.0% vs. 67.6%, P=0.007, *Figure 4A*), and that for patients overall in our study was 77.4% (*Figure 4B*).

Discussion

The salient findings of the present study were that in patients with a considerable amount of myocardial scarring (as detected by CE-CMR) and severe LV dysfunction, CABG+SVR provided a greater improvement in LVEF and NYHA class, and patients were less likely to be rehospitalized for CHF during follow-up compared to those who underwent I-CABG.

SVR is an effective treatment to improve LV function

Table 2	Perioperative	data and	outcomes	of surgery

Variables	All patients (n=140)	I-CABG (n=70)	CABG+SVR (n=70)	P value
Operation time, min	233.0±49.6	223.4±46.8	242.6±50.7	0.021
CPB time, min	108.1±30.9	100.2±23.8	116.0±35.0	0.002
Cross clamp, min	70.3±19.8	65.9±17.8	74.8±20.8	0.008
Ventilation time, h	20.0 (16.0, 28.7)	20.0 (15.0, 24.0)	22.0 (17.0, 37.0)	0.019
ICU duration, h	71.0 (46.0, 110.0)	45.0 (69.0, 93.0)	82.5 (55.3, 117.8)	0.064
Hospital stay, days	10.5±4.2	10.2±5.3	10.8±2.7	0.340
Grafts per patient	3.3±0.8	3.4±0.8	3.1±0.7	0.060
CABG outcomes				
New-onset AF	12 (8.6)	5 (7.1)	7 (10.0)	0.546
Reoperation for bleeding	5 (3.6)	2 (2.9)	3 (4.3)	1.000
Perioperative MI	7 (5.0)	3 (4.3)	4 (5.7)	1.000
Stroke	3 (2.1)	2 (2.9)	1 (1.4)	1.000
VA	7 (5.0)	3 (4.3)	4 (5.7)	1.000
Renal failure requiring dialysis	7 (5.0)	4 (5.7)	3 (4.3)	1.000
LCOS	12 (8.6)	7 (10.0)	5 (7.1)	0.546
Death	3 (2.1)	2 (2.9)	1 (1.4)	1.000

Values are expressed as mean ± SD, median (interquartile range) or n (%). AF, atrial fibrillation; CABG, coronary artery bypass graft; CPB, cardiopulmonary bypass; I-CABG, isolated coronary artery bypass graft; ICU, intensive care unit; LCOS, low cardiac output syndrome; MI, myocardial infarction; SD, standard deviation; SVR, surgical ventricular reconstruction; VA, ventricular arrhythmia.

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Baseline parameters	All patients (n=140)	I-CABG (n=70)	CABG+SVR (n=70)	P value
LVEF, %	27.5±5.7	28.3±5.6	26.6±5.7	0.077
LVEDD, mm	62.7±7.2	61.5±6.5	63.8±7.7	0.055
LVEDVI, mL/m ²	113.4±31.3	109.4±28.7	117.3±33.4	0.135
LVESVI, mL/m ²	82.3±27.3	78.2±25.3	86.3±28.7	0.079
CI, L/min/m ²	2.2±0.7	2.2±0.6	2.2±0.8	0.998
Dysfunctional segments	15.5±2.0	15.2±2.2	15.8±1.8	0.097
WMS	24.7±4.6	24.0±5.1	25.4±4.0	0.051
Myocardial LGE				
Spatial extent	13.6±2.3	13.6±2.4	13.6±2.2	0.971
Scar segments	5.7±0.8	5.6±0.9	5.8±0.7	0.206
Viable segments	8.2±2.1	8.4±2.0	8.1±2.3	0.429
Total scar score	29.7±5.6	29.2±6.1	30.2±4.9	0.275

Values are expressed as mean ± SD. CABG, coronary artery bypass graft; CI, cardiac index; I-CABG, isolated coronary artery bypass graft; LGE, late gadolinium enhancement; LVEDD, left ventricular end-diastolic diameter; LVEDVI, left ventricular end-diastolic volume index; LVEF, left ventricular ejection fraction; LVESVI, left ventricular end-systolic volume index; SD, standard deviation; SVR, surgical ventricular reconstruction; WMS, wall motion score.



Figure 2 Changes in the LVEF and LVEDD by echocardiography. (A) LVEF improved significantly from baseline to 6 months after surgery (P<0.001) and from 6 months to the latest follow-up in both groups (P<0.001). (B) LVEDD decreased significantly from baseline to 6 months after surgery in both groups (P<0.001). However, the mean reduction in LVEDD between 6 months and the latest follow-up in patients who underwent I-CABG was not statistically significant (P=0.506), while that for patients who underwent CABG and SVR was statistically significant (P<0.001). CABG, coronary artery bypass graft; I-CABG, isolated coronary artery bypass graft; LVEF, left ventricular ejection fraction; LVEDD, left ventricular end-diastolic diameter; SVR, surgical ventricular reconstruction.

in patients with severe CHF and an LV anterior-apical aneurysm (19). Previous studies have shown that SVR not only reduces LVEDV and improves regional myocardial performance in non-ischemic areas, but also eliminates LV desynchrony and has a positive effect on patients' longterm survival (5,20). Yet, it remains controversial whether these benefits are a result of the additional SVR or surgical revascularization. Howlett *et al.* reported that in patients with CHF and LV dysfunction undergoing I-CABG, the rates of cardiovascular hospitalization and CHF hospitalization were 45.2% and 25.6%, respectively (3). Klein *et al.* reported a 1-year survival rate of 90.6% in patients with previous MI and CHF (21). Prucz *et al.* demonstrated an excellent 4-year survival of 75% and 62% in CHF patients with ischemic cardiomyopathy who underwent CABG and SVR and I-CABG, respectively (6). In our patient cohort, the 10-year CVE-free survival rate for patients undergoing CABG and SVR was 87.0%. However, for patients who received I-CABG, this rate was only 67.6%, which was consistent with previous reports. For individuals whom we believed to be qualified for SVR yet did not receive it, *Table 1* showed that these patients were similar to CABG+SVR patients in almost every baseline characteristic. Therefore, the differences in their operative



Figure 3 NYHA heart failure symptoms and CCS angina class at baseline and follow-up. (A) Patents with NYHA class III–IV were distributed similarly between the two groups at baseline. The NYHA class also improved by an average of 1.0 class at follow-up for both groups (P<0.001), with 39.7% NYHA class I in I-CABG patients and 62.3% NYHA class I in CABG+SVR patients. (B) Patients with CCS class III–IV were also distributed similarly between the two groups at baseline. Angina symptoms improved by an average of 0.8 class in both groups (P<0.001). The proportion of patients with no angina increased at follow-up in both groups (P<0.001). CABG, coronary artery bypass graft; CCS, Canadian Cardiovascular Society; I-CABG, isolated coronary artery bypass graft; NYHA, New York Heart Association; SVR, surgical ventricular reconstruction.

Table 4 Outcomes for I-CABG versus CABG+SVR during follow-up

Outcomes	All patients (n=137)	I-CABG (n=68)	CABG+SVR (n=69)	P value
Rehospitalization for CHF	16 (11.7)	13 (19.1)	3 (4.3)	0.007
VA	4 (2.9)	3 (4.4)	1 (1.4)	0.601
Non-fatal MI	3 (2.2)	2 (2.9)	1 (1.4)	0.990
Angina pectoris	3 (2.2)	1 (1.5)	2 (2.9)	1.000
Deaths	5 (3.6)	3 (4.4)	2 (2.9)	0.987
NYHA class I or II	106 (77.4)	47 (69.1)	59 (85.5)	0.030
LVEDD change, %	-6.4±6.4	-4.4±6.3	-8.4±5.8	<0.001
LVEF change, %	9.9±8.3	7.8±8.4	12.0±7.6	0.002
LVEF improvement*	99 (72.3)	41 (60.3)	58 (84.1)	0.002
No or trivial MR	105 (76.6)	47 (69.1)	58 (84.1)	0.115

*, LVEF improvement (Echo) was defined as ≥5% improvement in LVEF at follow-up. NYHA improvement was defined as any improvement in the NYHA class at follow-up. Values are expressed as mean ± SD or n (%). CABG, coronary artery bypass graft; CHF, congestive heart failure; Echo, echocardiography; I-CABG, isolated coronary artery bypass graft; LVEDD, left ventricular end-diastolic diameter; LVEF, left ventricular ejection fraction; MI, myocardial infarction; MR, mitral valve regurgitation; NYHA, New York Heart Association; SD, standard deviation; SVR, surgical ventricular reconstruction; VA, ventricular arrhythmia.



Figure 4 Survival curves of patients who underwent I-CABG *vs.* CABG+SVR. (A) Kaplan-Meier curve shows that the rate of survival without CVEs in patients who underwent CABG and SVR was significantly higher than that for patients who underwent I-CABG (87.0% *vs.* 67.6%, P=0.007, respectively). (B) During a follow-up period of 123.1±12.7 months, the cumulative rate of survival without CVEs for all patients was 77.4%. CABG, coronary artery bypass graft; CVEs, cardiovascular events; I-CABG, isolated coronary artery bypass graft; SVR, surgical ventricular reconstruction.

protocol definitely affected their outcomes.

Numerous studies have demonstrated that preoperative LVESVI predicts survival in patients with ischemic cardiomyopathy (5,8,22). However, SVR may not be appropriate for many patients with LV enlargement and CHF, and we believe that SVR is only appropriate for a highly select group of patients who meet specific eligibility criteria. Current indications that are generally accepted for SVR involve anterior MI or aneurysm and LV akinesia or dyskinesia (23). These factors are not necessarily present in all ischemic cardiomyopathy patients with LV dilatation. Therefore, patients must be appropriately evaluated for a history of MI, LV enlargement, and most significantly, anterior wall nonviability (22). As a result, CE-CMR serves as the preferred modality and is used because it can provide accurate LV function and volume measurements and also determine the scar tissue (24). Castelvecchio et al. indicated that scar location affects long-term survival in CHF patients undergoing SVR (25), while Yamazaki et al. further demonstrated that accurate preoperative assessments of myocardial viability testing using CE-CMR are essential for better stratification of the SVR procedure (22). In the present study, all of the subjects had a considerable amount of scar tissue (≥ 5 scar segments), as evidenced by CE-

CMR. No significant differences were found in the LGE parameters including scar segments, TSS, or LV size-related parameters at baseline between the two groups. The number of grafts in each group was also similar. Therefore, the two groups were comparable enough and we strongly believe that our results accurately reflect the discrepancies in follow-up outcomes between I-CABG and CABG+SVR.

Functional MR caused by ischemia is a common occurrence in individuals with a dilated ventricle. The longitudinal and transverse elongation of the LV causes lateral displacement and papillary muscle dysfunction, hindering leaflet coaptation (26). All individuals undergoing mitral valve procedures at the time of CABG were not included in the present study to ensure that the mitral valve operations were not confounding factors. In our analysis, MR was reduced in 61 (73.5%) of the patients during the follow-up compared with that before surgery, which indicates that both revascularization and SVR improve mitral valve function, and mitral repair or replacement may not be essential in patients with mild to moderate functional MR.

Our data also suggested that SVR can be safely performed in high-risk populations and is a useful strategy to treat chronic MI and severe LV dysfunction because of the differences in long-term prognosis and the percentage

of LVEF increase. The idea that an appropriate SVR must be performed to achieve benefit is also supported by the current data. However, SVR is underutilized in realworld clinical situations, which might be attributable to the primary surgeon not appreciating the potential benefit, being unfamiliar with candidate selection, as well as a lack of training in the surgical technique. However, the number of patients with severe LV dysfunction and large aneurysms has been decreasing in developing countries like China in recent years. The patients in our cohort exhibited significant improvements not only in LVEF and LV sizes but also in NYHA class as well as clinical symptoms. We hope that our findings will increase awareness of SVR so that eligible patients will be referred to surgeons qualified to perform the SVR procedure in the future.

Limitations

Several limitations of this study merit attention. Firstly, despite the relatively long follow-up interval, the major limitation is that this is not a randomized controlled trial (RCT), but a single-center study with a relatively small cohort. Secondly, the classifications were based on a review of the preoperative imaging protocol and were thus inherently subjective. We tried to mitigate this subjectivity by having the same surgical team review all available CE-CMR protocols, TTEs, and ventriculograms. Also, despite having a specific indication of SVR, the final decision on whether to perform SVR is still made by the cardiac surgeon in charge of the procedure. As a result, selectivity bias indeed exists, which might also be a limitation of this study. Finally, CE-CMR was not performed in all patients at follow-up and the LV volume measurements for these patients were not reported. Thus, analyzing the results to detect the degree of revascularization and sufficient myocardial perfusion could not be performed. Nevertheless, all patients underwent TTE at 6 months instead of CE-CMR.

Conclusions

We conclude that in patients with chronic MI and severe LV dysfunction, CABG+SVR provided a greater improvement in terms of LVEF/LVEDD and NYHA class, and patients were less likely to be rehospitalized for CHF compared to those individuals who underwent I-CABG. Based on the 17-segmental model, patients with LV enlargement and a considerable amount of scar tissue (\geq 5 scar segments)

located in the anterior segments will probably benefit from SVR in the long term.

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Footnote

Reporting Checklist: The authors have completed the TREND reporting checklist. Available at https://jtd. amegroups.com/article/view/10.21037/jtd-22-1214/rc

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Conflicts of Interest: All authors have completed the ICMJE uniform disclosure form (available at https://jtd.amegroups.com/article/view/10.21037/jtd-22-1214/coif). The authors have no conflicts of interest to declare.

Ethical Statement: The authors are accountable for all aspects of the work in ensuring that questions related to the accuracy or integrity of any part of the work are appropriately investigated and resolved. The study was conducted in accordance with the Declaration of Helsinki (as revised in 2013) and was approved by the institutional review board of Fuwai Hospital (No. 2010-259). All patients provided written informed consent.

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Figure S1 The procedure of circular reconstruction in a patient who underwent CABG+SVR. During circular reconstruction, after ventriculotomy is centered in the zone of anterior asynergy (A), a suture is placed in the interior of the ventricle to encircle the myocardial scar at the boundary between the akinetic and viable tissue (B). Tightening of this suture brings the healthy myocardium together. Next, a Dacron patch is used to reconstruct the shape of left ventricle (C). Finally, we closed the ventriculotomy with linear closure (D). CABG, coronary artery bypass graft; VA, ventricular arrhythmia.



Figure S2 Case 1. The CE-CMR of a patient at baseline and 6 months after I-CABG. Example of CE-CMR images before and 6 months after I-CABG of a 56-year-old man with a 6-scar segment who suffered from congestive heart failure 9 months after surgery. CE-CMR, contrast-enhanced cardiovascular magnetic resonance imaging; I-CABG, isolated coronary artery bypass graft; LVEDD, left ventricular end-diastolic diameter; LVEDV, left ventricular end-diastolic volume index; LVEF, left ventricular ejection fraction; LVESV, left ventricular end-systolic volume.



Figure S3 Case 2. The CE-CMR of a patient at baseline and 6 months after CABG+SVR. Example of CE-CMR images before and 6 months after CABG+SVR of a 54-year-old man with a 7-scar segments who survived without CVEs during follow-up. CE-CMR, contrastenhanced cardiovascular magnetic resonance imaging; I-CABG, isolated coronary artery bypass graft; CVEs, cardiovascular events; LVEDD, left ventricular end-diastolic diameter; LVEDV, left ventricular end-diastolic volume index; LVEF, left ventricular ejection fraction; LVESV, left ventricular end-systolic volume; SVR, surgical ventricular reconstruction.