Is minimally invasive mitral valve surgery inappropriate for redo-surgery or elderly patients? – Study of 55 consecutive cases following minimally invasive mitral valve replacement via right mini-thoracotomy

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Abstract

Background: Minimally invasive mitral valve surgery has been established as a routine procedure at our institution. In the present study, early and mid-term outcomes of patients who underwent minimally invasive mitral valve replacement (MIMVR), including redo-operations and elderly patients, during the 5-year post-surgery period were analyzed to review short-term morbidity and mortality, and mid-term results. Methods: Preoperative variables, intraoperative findings, and postoperative outcomes of MIMVR patients treated from January 2014 to November 2020 and prospectively stored in a database were reviewed. Survival and freedom from cerebrovascular events were evaluated using life tables and Kaplan-Meier analysis. Results: A total of 445 patients underwent minimally invasive mitral valve surgery during the study period, of whom 55 received mitral valve replacement (MVR), including 18 cases of redo-MVR and 10 elderly ([?]80 years old) patients. Mean age at the time of surgery was 70.7 \pm 11.3 years. The number of patients who underwent conversion to a sternotomy was 0, while 30-day mortality was noted in one (2%). For all MIMVR cases, 1- and 5-year survival was 90.8 \pm 3.9% and 76.5 \pm 7.1%, respectively. Furthermore, freedom from cerebrovascular events and anticoagulation-related complications was 94.3 \pm 3.2% and 84.2 \pm 6.3% at 1 and 5 years, respectively. In univariate analysis, independent predictors of hospital mortality and prolonged hospital stay included infectious endocarditis, while previous cardiac surgery and elderly status were not significant factors. Conclusions: MIMVR can be performed safely and effectively for redo-MVR and in elderly patients with very few perioperative complications. Early and mid-term outcomes in the present cohort were acceptable. Is minimally invasive mitral valve surgery inappropriate for redo-surgery or elderly patients? – Study of 55 consecutive cases following minimally invasive mitral valve replacement via right mini-thoracotomy

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Methods: Preoperative variables, intraoperative findings, and postoperative outcomes of MIMVR patients treated from January 2014 to November 2020 and prospectively stored in a database were reviewed. Survival and freedom from cerebrovascular events were evaluated using life tables and Kaplan-Meier analysis.

Results: A total of 445 patients underwent minimally invasive mitral valve surgery during the study period, of whom 55 received mitral valve replacement (MVR), including 18 cases of redo-MVR and 10 elderly (\geq 80 years old) patients. Mean age at the time of surgery was 70.7±11.3 years. The number of patients who underwent conversion to a sternotomy was 0, while 30-day mortality was noted in one (2%). For all MIMVR cases, 1- and 5-year survival was 90.8±3.9 % and 76.5±7.1%, respectively. Furthermore, freedom from cerebrovascular events and anticoagulation-related complications was 94.3±3.2% and 84.2±6.3% at 1 and 5 years, respectively. In univariate analysis, independent predictors of hospital mortality and prolonged hospital stay included infectious endocarditis, while previous cardiac surgery and elderly status were not significant factors.

Conclusions: MIMVR can be performed safely and effectively for redo-MVR and in elderly patients with very few perioperative complications. Early and mid-term outcomes in the present cohort were acceptable.

Introduction

The feasibility and safety of minimally invasive mitral valve surgery (MIMVS) with the primary goals of improved cosmetic factors and reduced postoperative discomfort, while maintaining the same level of safety and efficacy as with conventional surgery, have been demonstrated in several reports [1]. Most analyses have shown that patients who undergo MIMVS are at low risk and young, and have fewer comorbidities [1]. However, questions remain regarding whether these methods should be limited to young patients or primary cases. On the other hand, few studies have specifically investigated minimally invasive mitral valve replacement (MIMVR) in detail. Regardless of the approach utilized for mitral valve stenosis, the significant usefulness of mitral valve replacement (MVR) has been shown for conditions such as advanced infectious endocarditis or mitral valve insufficiency with low cardiac function, as well as for patients with a previous failed mitral valve repair.

A mitral valve re-operation can be demanding for patients with a patent coronary artery bypass graft or who have undergone a previous aortic valve replacement procedure, as well as those affected by complications following a previous operation (abscess, perivalvular leakage, thrombosis). In addition, risk of graft injury or hemorrhage, presence of dense adhesions, and/or complex valve exposure can make a redo-valve operation through a median sternotomy challenging [2]. In such cases, a minimally invasive surgical approach through a right-sided mini-thoracotomy is considered to be a valid alternative instead of a repeated conventional median sternotomy [3].

In 2014, use of an MIVMR procedure was established at our department, with that via a right mini-thoracotomy (RT approach) now the standard used for treatment of mitral valve disease. The aim of the present study was to analyze early and mid-term outcomes of 55 consecutive patients who underwent MIMVR via an RT approach during a recent 5-year period. In addition, to determine the effectiveness of redo-MIMVR and the procedure in elderly (≥80 years old) patients, we compared the results of primary and redo-MIMVR cases, and also examined adaptation for elderly cases.

Methods

The study complies with the Declaration of Helsinki. The present analysis was approved by the audit board at the Chiba-Nishi general hospital. As this analysis came under clinical audit/quality of care assessment and all data were anonymized following the governance criteria of the NHS, the IRB agreed informed consent was not required.

Patient selection and data collection

A total of 445 patients underwent minimally invasive mitral valve surgery via an RT approach at Chiba-Nishi General Hospital between January 2014 and November 2020. Of the patients treated during the study period, 380 underwent mitral valve repair as the primary procedure, while 65 underwent MIMVR via an RT approach. After excluding 10 of the MIMVR patients who received concomitant aortic valve replacement, 55, including 10 classified as elderly, who underwent that as first-time (n=38) or redo (n=17) procedure were enrolled, each of whom also had late follow-up results available.

The main outcomes investigated were early and late mortality, perioperative complications, and freedom from cerebrovascular events and reoperation. Early mortality was defined as death from any cause occurring within 30 days of the operation or before hospital discharge. Stroke was defined as any new focal or global neurological deficit lasting more than 24 hours,.

All patients were examined postoperatively from 1 to 4 weeks, and again at 12 and 24 weeks, then were contacted thereafter for follow-up information. Follow-up examinations were performed every 24 weeks in the late phase. The median follow-up

period was 34.2±24.6 months (interquartile range 13–52 months) and follow-up examinations were performed in 100% of the cases.

Statistical analysis

Continuous data are expressed as the mean ± standard deviation or median with interquartile range (IQR), while categorical data are shown as percentages. Survival and time-to-event analyses for rates of cerebrovascular events and anticoagulation-related complications were assessed using the Kaplan-Meier method. All reported P values are two-sided and those <0.05 were considered to indicate statistical significance. All statistical analyses of recorded data were performed using the Excel statistical software package (Ekuseru-Toukei 2010; Social Survey Research Information Co., Ltd., Tokyo, Japan).

Results

Baseline characteristics and risk factor prevalence are listed in Table 1. The mean age of the enrolled patients was 70.7 ± 11.3 years and 36~(65%) were female. Eighteen (33%) had previously undergone a cardiac operation, including mitral valve repair in 8, mitral valve replacement in 1, aortic valve replacement in 3, total arch replacement in 2, and

coronary artery bypass grafting (CABG) in 1. The mean preoperative left ventricular ejection fraction was $60.6\pm11.7\%$. Preoperative chronic atrial fibrillation was noted in 20 (36%) patients. There were no statistically significant differences between first-time and redo-MIMVR patients for mean age (71.8±10.1 vs. $68.4\pm14.6\%$, *p*=0.31). The redo population had a significantly higher prevalence for EuroSCORE II (4.37 ± 4.7 vs. 7.79±6.2, *p*=0.03) and left ventricular ejection fraction dysfunction (62.8 ± 11.5 vs. 56.2±11.5%, *p*=0.05), whereas there was no statistically significant difference between the elderly and younger (age <80 years) patients for EuroSCORE II (7.95 ± 6.7 vs. 4.94 ± 5.0 , *p*=0.11). On the other hand, the elderly group had a significantly higher prevalence for diabetes (insulin user) [4 (40%) vs. 6 (13%), *p*=0.048] and pulmonary hypertension (>45 mmHg) [7 (70%) vs. 16 (36), *p*=0.046]. The most predominant pathological factor was rheumatic disease (n=29, 52%), followed by degenerative disease (n=10, 18%) and endocarditis (n=8, 15%), with other such factors noted in 8 (15%).

The operative procedural details are presented in Table 2. All 55 patients underwent a successful MIMVR via an RT. Seven received a mechanical valve and 48 a bioprosthetic valve, and all had an uneventful recovery. There were no cases of intraoperative conversion to a sternotomy. The mean operative time was 215±46.7 minutes, while aortic cross-clamp and cardiopulmonary (CPB) times were 98±26.8 and 131±28.6 minutes, respectively, for the MIMVR. There were no statistically significant differences between first-time and redo-MIMVR patients in regard to CPB (129±28.1

vs. 137±30.7 minutes, p=0.36) or aortic cross-clamp (100±24.4 vs. 96±32.4 minutes, p=0.62) times, though redo-MIMVR patients had a longer operation time (207±46.5 vs. 233±44.2 minutes, p=0.046). There was no statistically significant difference between the elderly and younger patients for operation time. Concomitant procedures were performed in 24 patients (44%), including atrial fibrillation (AF) ablation in 17 (31%), tricuspid valve surgery in 11 (20%), patent foramen ovale/atrial septal defect closure in 2 (4%), and oversewing or use of a closed devise for the left atrial appendage in 15 (27%). Redo-MIMVR patients were less likely to undergo a concomitant procedure [21 (57%) vs. 3 (17%), p=0.005].

Early outcomes

Operative results are presented in Table 3. There was 1 in-hospital mortality (2%), which occurred on postoperative day 92 and was due to acute respiratory distress syndrome in an active IE patient. Median hospital and ICU stays were 11 (IQR 8.8-18) and 3 (IQR 2-4) days, respectively. There was no statistically significant difference between elderly and younger patients in regard to intensive care unit (ICU) length of stay [3 (2-4) vs. 3 (2-4) days, p=0.61], though the elderly group had longer hospital length of stay [16.5 (13-33) vs. 10 (8-15.8) days, p=0.01] and ventilation time [16 (12.5-18) vs. 8 (6-12) hours, p=0.003]. There were no statistically significant difference differences between first-time and redo-MIMVR cases in regard to ICU or hospital stay duration. A blood transfusion during hospitalization was required in 27 (49%) patients,

with those who underwent redo-MIMVR [14 (38%) vs. 13 (72%), p=0.017] or with elderly status [8 (80%) vs. 19 (42%), p=0.033] more likely to undergo a transfusion. Of 35 patients admitted for sinus rhythm, new-onset atrial fibrillation occurred in 7 (13%). No patients had a cerebral vascular accident, while 2 (4%) had an acute renal failure incident that required dialysis, of whom 1 (2%) needed a reoperation for bleeding. Following surgery, all patients underwent a transthoracic echocardiography examination prior to discharge, with no device-related complications or paravalvular leakage observed.

Midterm outcomes

Clinical follow-up examinations were performed in 55 patients (100%) over a mean period of 34.2±24.6 months (interquartile range 13–52 months). None demonstrated prosthetic valve dysfunction or underwent a mitral prosthetic valve reoperation.

Survival

Overall, the 1-, 3-, and 5-year survival rates were 90.8±3.9%, 81.3±5.9%, and

76.5 \pm 7.1%, respectively, while those for the elderly group were 80.0 \pm 12.7%,

64.0±17.5%, and 64.0±17.5%, respectively, and for the younger group were 93.3±3.8%,

84.3±6.1%, and 78.6±7.8%, respectively (Fig. 1A, B). A total of 10 patients (18%), 3

(30%) elderly and 7 (16%) younger, died during the follow-up period, of whom 4

(10.8%) underwent first-time MIMVR and 6 (33%) redo-MIMVR. The cause of death

could only be determined in 8, which was cardiac-related in 3 (3, 14, 30 months), pneumonia in 2 (3, 5 months), a cerebral vascular event in 1 (48 months), an abdominal aortic aneurysm rupture in 1 (19 months), septic shock in 1 (2 months), and unknown in 2 (3, 13 months).

Cerebrovascular events and anticoagulation-related complications

Following mitral valve replacement, the rates for freedom from cerebrovascular events and anticoagulation-related complications after 1, 3, and 5 years were $94.3\pm3.2\%$, $88.4\pm5.1\%$, and $84.2\pm6.3\%$, respectively, while those in elderly patients were 100%, 100%, and 100%, respectively, and in the younger group were $93.2\pm3.8\%$, $86.5\pm5.8\%$, and $81.7\pm7.2\%$, respectively (Fig. 2A, B).

Predictors of hospital mortality and prolonged hospital stay shown by univariate analysis

Following MIMVR surgery, there was 1 in-hospital mortality (2%) and 7 patients had a prolonged (>30-days) hospital stay (12.7%), for a total of 8 (14.5%) cases. Variables assessed as possible predictors of mortality in univariate analysis are shown in Table 4. The only significant (p=0.05) univariate predictor was infectious endocarditis, while previous cardiac surgery, concomitant operation, and elderly status were not found to be significant.

Discussion

Although several reports have described excellent early and long-term results of minimally invasive mitral valve repair [4], few studies have analyzed MIMVR cases in detail. The present analysis of a consecutive series of patients who underwent an MIMVR procedure found it to be safe and associated with excellent postoperative outcomes, while good procedural times, shorter hospital and ICU stays, and acceptable mid-term results were also revealed. As demonstrated by complete echocardiographic follow-up results, no paravalvular leakage developed in any of the patients, including during the late phase. Specifically, overall in-hospital mortality and postoperative neurological events were nearly 0%, with the latter lower than the range of rates for mortality and neurological events recently presented by Modi and colleagues [5] in a meta-analysis of MIMVR cases (4.9% in 5 cohorts with a total of 979 patients, 2.3% in 3 cohorts with a total of 778 patients).

In the present study, since there were no cerebral vascular accident incidents, preoperative contrast CT results were analyzed to detect aorta condition, i.e., the presence of arteriosclerotic disease or parietal thrombus, and determine the perfusion program for each case. Details of the perfusion protocol were previously reported by Nakamura et al. [6]. When significant arteriosclerotic disease was found in the entire aorta and iliac artery, right axillary artery cannulation was employed to establish antegrade perfusion for CPB.

Although early good results and quick recovery following MIMVR have been noted in prior reports [7], there are concerns regarding its long-term efficacy, which has not been adequately elucidated. We found 2 reports that included long-term follow-up findings after MIMVR procedures performed at high-volume centers. The longest functional and echocardiographic follow-up study was conducted by Glauber and colleagues, who analyzed 476 patients who underwent MIMVR via an RT approach between 2003 and 2013 (mean age 67±11 years, incision length 5-7 cm, in-hospital mortality 3.2%, neurological complications 2.6%, reoperation for bleeding 6.4%) [8]. In those cases, survival after replacement at 1, 5, and 10 years was found to be 91.0 \pm 1.4%, 81.3 \pm 2.5%, and 76.2 \pm 3.4% respectively, while the rates for freedom from reoperation after replacement at 1, 5, and 10 years were 98.6±0.6%, 94.5±1.6%, and 83.9±5.5%, respectively. In 2003, Casselman et al. reported early and long-term results in a review of a series of 80 patients who underwent endoscopic MVR with an RT procedure (incision length 4 cm, mean follow-up period 19.6±17.3 months, completed in all) [9]. Although 2 (2.5%) of their patients experienced new onset endocarditis during the follow-up period and 1 underwent a reoperation with a median sternotomy, they noted excellent freedom from late phase death (4 years) in 92.0±3.6% as well as freedom from anticoagulation-related complications (4 years) in 97.0±1.9%. In the present cohort who underwent MIMVR (mean age 70.7±11.3 years, mean incision length 5-6 cm), 5-year survival, freedom from a mitral valve-related reoperation, and freedom from cerebrovascular events and anticoagulation-related complications showed favorable rates of 76.5%, 100%, and 81.7%, respectively. In addition, those rates were also good

after first-time MIMVR at 83%, 100%, and 93%, respectively. As compared to those two other recent reports, our mid-term survival results were relatively better, while the freedom from reoperation was 0% at the 5-year follow-up examination, and there were no device-related complications or cases of paravalvular leakage.

Reoperative mitral valve surgery through a median sternotomy can be particularly challenging due to dense adhesions and is known to carry a substantial risk of injury to vascular structures, which have been found to occur in 7–9% of cases and are associated with increased mortality [10]. A valid alternative that could be employed to avoid risks associated with a redo median sternotomy approach is a right anterolateral minithoracotomy approach [11]. Unfortunately, only 5 patients who underwent a conventional sternotomy were available as a control group for the present study. Our strategy consists of selecting MIMVR whenever possible even for redo-mitral valve procedure cases, as that procedure has been our the first-line choice since 2014. However, that introduces a critical selection bias including cases of redo-mitral valve surgery between MIMVR and conventional sternotomy groups for analysis of procedures performed at our institution. Thus, we sought to compare the present results with those previously reported for cases of redo-MIMVR or conventional full sternotomy redo-mitral valve surgery. Studies regarding the feasibility of redo-MIMVR via an RT procedure have been presented by Ricci et al. [12], Sharony et al. [13], and Thomson et al. [14], with excellent results reported in each (Table 5). The results obtained in the present cases are favorable as compared with those, though remaining

issues related to our method include longer prolonged CPB and cross-clamp times as compared to those prior findings. On the other hand, there was only one in-hospital mortality among the present cohort. Furthermore, the rates for freedom from stroke events and reoperation for bleeding were nearly 0%. In addition, 3 other retrospective observational studies all demonstrated the superiority of an RT approach as compared to a median sternotomy for reoperative mitral valve surgery, with excellent results noted in those cases (Table 5) [15-17]. Pooled analysis showed a mini-thoracotomy mitral valve surgery procedure as a safe alternative to a standard sternotomy, with reduced mortality rates, length of hospital stay, and reoperations for bleeding, along with a comparable risk of stroke.

There is a paucity of literature showing outcomes of MIMVS in older patients. Iribarne et al. demonstrated that it can be performed safely in patients at least 75 years old [18]. Furthermore, they noted that even though a minimally invasive approach was associated with slightly longer CPB and cross-clamp times than a conventional sternotomy, there were no significant differences regarding postoperative morbidity or mortality. Importantly, the mean and median durations of hospitalization were 3 and 1 day shorter, respectively, for patients who underwent an MIMVS procedure, findings with important implications for resource use. Our strategy consists of selecting MIMVR whenever possible even for elderly cases, thus we were not able to compare with a conventional sternotomy approach. In the present cohort, elderly patients (≤80 years old) who underwent MIMVR showed favorable rates for 5-year survival, freedom from a mitral

valve-related reoperation, and freedom from cerebrovascular events and anticoagulation-related complications of 64%, 100%, and 100%, respectively. On the other hand, the cause of cardiac-related death could only be determined in 1 of 3 cases (septic shock after 2 months, fatal arrythmia after 3 months, unknown after 13 months). Nevertheless, the present results demonstrated the superiority of MIMVR in elderly patients the same as with the young group.

Factors with potential effects on the outcome of patients who undergo MIMVR are numerous and many confounding variables exist, though only dependent variables related to in-hospital mortality and prolonged hospital stay (>30 days) were examined in the present study. Univariate analysis indicated infectious endocarditis as a predictor, which might be related to the influence of post-operative antibiotic administration. Interestingly, previous cardiac surgery, concomitant operation, and elderly status were not significant. A multivariate logistic regression model of 409,904 valve procedures performed between 1994 and 2003 and cataloged in the Society of Thoracic Surgeons database demonstrated that the third most important preoperative variable influencing operative mortality is a reoperation (OR 1.61, P < 0.001) [19]. However, the present data suggest that an RT approach after a previous median sternotomy is not an independent predictor of mortality (P=0.344). This important finding indicates that this technique, in least in consideration of early outcome, should be the first choice for reoperative mitral valve surgery in patients who do not need concomitant aortic valve replacement or coronary revascularization. That is in concordance with previously reported data

demonstrating equivalent or lower mortality rates, and less morbidity associated with an RT approach as compared to a reoperative sternotomy [15, 20, 21]. Nevertheless, findings regarding mid- and long-term outcomes are needed.

For treating mitral valve disease or degenerated bioprosthetic valve failure, transcatheter mitral valve replacement (TMVR) therapy has emerged as an encouraging option. Early reports have noted utilization of TAVR devices for degenerated surgical bioprosthetic valves (transcatheter mitral valve-in-valve implantation with balloon-expandable valves) (22) or annuloplasty rings (transcatheter mitral valve-in-ring implantation with balloon-expandable valves) (23), with reasonable results shown. Additionally, utilization of a TAVR valve in the mitral position has been described for treating patients with significant mitral annular calcification (MAC), though early results have demonstrated high rates of mortality and significant complications (24). Overall, survival (30-day all-cause mortality) after these various procedures has been greatest in patients who underwent a mitral valve-in-valve procedure (6.2%), followed by valve-inring (9.9%), with the worst results reported for valve-in-MAC (34.5%) (25), and those findings raise questions regarding whether TMVR is an entirely appropriate alternative for surgical mitral valve surgery. Results presented thus far show that MIMVR is more effective. Additional long-term studies with larger numbers of enrolled patients will be necessary to better assess the efficacy of TMVR and MIMVR.

This study has some limitations. It was conducted in a retrospective manner and lacked a control group for appropriate comparisons. Thus, it was not possible to compare the present case series with a control group, since MIMVR has been our standard approach since 2014 and patients typically demand less invasive procedures. Even though this was a follow-up study conducted over 5 years, the number of patients was small and only 12 were found to be at risk at the 60-month follow-up examination. A well-designed study with an appropriate sample size will be required in the future to validate the advantages of MIMVR.

To conclude, we found that MIMVR can be safely performed with encouraging short and mid-term outcomes, including elderly patients, with very low rates of conversion and mortality as compared to a conventional sternotomy procedure. In addition, a mitral valve re-operation can be safely and effectively performed through a smaller right thoracotomy. Finally, MIMVR was found useful for elderly patients. Based on these findings, we consider MIMVR to be an attractive alternative to conventional MVR.

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Disclosure: The authors have no conflicts of interest to declare.

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Figure legends

Fig. 1A. 1B. Freedom from late death for any reason.

Fig. 2A. 2B. Freedom from cerebrovascular events and anticoagulation-related complications.

Table legends

Table 1. Preoperative clinical characteristics

COPD: chronic obstructive pulmonary disease, LVDd: left ventricular dimension diastole, LVDs: left ventricular dimension systole, LVEF: left ventricular ejection fraction, MVP: mitral valve plasty, OMC: open mitral commissurotomy, AVR: aortic valve replacement, TAR: total arch replacement, CABG: coronary artery bypass graft

Table 2. Procedural details

IQR: interquartile range

Table 3. Early outcomes

Table 4. Univariate analysis for hospital mortality and prolonged hospital stay

Table 5. Principal data and outcomes of interest from individual studies

Data are expressed as absolute number, percentage, or mean.

MT: mini-thoracotomy, ST: median sternotomy, NA: not available, RFB: reoperation for bleeding, USA: United States of America, UK: United Kingdom

Table 1

Variable	MVR (n=55)	First-time MVR (n=37)	Redo MVR (n=18)	P value	Age ≥80 (n=10)	Age <80 (n=45)	P value
Mean age (years ± SD)	70.7±11.3	71.8±10.1	68.4±14.6	0.31	82.4±2.2	68.1±11.4	< 0.01
Female, n (%)	36 (65)	27 (73)	9 (50)	0.09	8 (80)	28 (62)	0.28
Hypertension, n (%)	26 (47)	16 (43)	10 (56)	0.39	7 (70)	19 (42)	0.11
Congestive heart failure, n (%)	15 (27)	10 (27)	5 (31)	0.95	5 (50)	10 (22)	0.07
NYHA class III or IV	18 (33)	11 (30)	7 (39)	0.50	5 (50)	13 (29)	0.20
Diabetes (insulin user), n (%)	10 (18)	9 (24)	1 (6)	0.09	4 (40)	6 (13)	0.048

COPD, n (%)	3 (5)	2 (5)	1 (6)	0.98	1 (10)	2 (5)	0.48
Atrial fibrillation, n (%)	20 (36)	13 (35)	7 (39)	0.79	4 (40)	16 (36)	0.18
Chronic renal failure, n (%)	13 (24)	9 (24)	4 (22)	0.86	4 (40)	9 (20)	0.17
Dialysis-dependent	1 (2)	1 (3)	0		0	1 (2)	
EuroSCORE II, % (mean ± SD)	5.49±5.4	4.37±4.7	7.79±6.2	0.03	7.95±6.7	4.94±5.0	0.11
LVEDD, mm (mean ± SD)	50.7±8.4	50.2±9.1	51.5±7.1	0.60	50.5±8.0	50.7±8.7	0.97
LVESD, mm (mean ± SD)	34.2±8.6	33.1±8.8	36.4±8.4	0.19	32.9±6.3	34.5±9.2	0.60
LVEF, % (mean ± SD)	60.6±11.7	62.8±11.5	56.2±11.5	0.05	63.4±9.2	60.0±12.4	0.41

LVEF <40%, n (%)	3 (5)	1 (3)	2 (11)	0.20	0	3 (7)	0.40
Pulmonary hypertension >45 mmHg (%)	23 (42)	15 (41)	8 (44)	0.78	7 (70)	16 (36)	0.046
Cause							
Degenerative, n (%)	10 (18)	2 (5)	8 (44)		2 (20)	8 (18)	
Rheumatic, n (%)	29 (52)	27 (73)	2 (11)		3 (30)	26 (58)	
Infective, n (%)	8 (15)	7 (19)	1 (6)		3 (30)	5 (11)	
Other, n (%)	8 (15)	1 (3)	7 (39)		2 (20)	6 (13)	

Previous cardiac surgery, n (%)

18 (33)

Mitral valve (MVP/MVR/OMC)	(8/1/2)
Aortic valve (AVR)	3
Arch (TAR)	2
Coronary (CABG)	1
Other	2
Time to re-operation, years ± SD	13.9±12.7

Variable	MVR (n=55)	First-time MVR (n=37)	Redo MVR (n=18)	P value	Age ≥80 (n=10)	Age <80 (n=45)	P value
Operative time, min (mean ± SD)	215±46.7	207±46.5	233±44.2	0.046	226±64.2	213±43.0	0.44
Cardiopulmonary bypass time, min (mean ± SD)	131±28.6	129±28.1	137±30.7	0.36	127±26.2	132±29.6	0.63
Cross-clamp time, min (mean ± SD)	98±26.8	100±24.4	96±32.4	0.62	92.1±12.1	99.7±29.2	0.42
Mechanical valve/bioprosthetic valve, n (%)/n (%)	7/48	3/34	4/14		0/10	7/38	
Concomitant procedures	24 (44)	21 (57)	3 (17)	0.005	5(50)	19(42)	0.65

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Tricuspid valve annuloplasty, n (%)	11 (20)	8 (22)	3 (17)	 2(20)	9(20)
Atrial fibrillation surgery, n (%)	17 (31)	17 (46)	0	 3(30)	14(31)
ASD closure, n (%)	2 (4)	1 (3)	1 (6)	1(10)	1(2)
LAA closure, n (%)	15 (27)	14 (38)	1 (6)	3(30)	12(27)
Conversion to sternotomy, n (%)	0	0	0	 0	0
Arterial cannulation Femoral/Rt. axillary artery, n (%)/n (%)		26 (70)/ 11(30)	7 (39)/ 11 (61)	5 (50)/ 5 (50)	32 (71)/ 13 (29)
Venous cannulation Femoral/femoral+SVC, n (%)/n (%)		34 (92)/ 3 (8)	14 (78)/ 4 (22)	 7 (70)/ 3 (30)	41 (91)/ 4 (9)

Variable	MVR (n=55)	First-time MVR (n=37)	Redo MVR (n=18)	P value	Age ≥80 (n=10)	Age <80 (n=45)	P value
In-hospital mortality, n (%)	1 (2)	0	1 (6)	0.48	0	1 (2)	0.63
Intensive care unit stay, days (median, IQR)	3 (2-4)	3 (2-4)	3 (2-4.8)	0.66	3 (2.3-4)	3 (2-4)	0.61
Hospital stay, days (median, IQR)	11 (8.8-18)	10 (8-18)	11 (10-16)	0.69	16.5 (13-33)	10 (8-15.8)	0.01
Ventilation duration, hours (median ± SD)	9±15.8	8.3±5.2	11.5±5.6	0.25	16 (12.5-18)	8 (6-12)	0.003
Blood transfusion (intra or post op), n (%)	27 (49)	14 (38)	13 (72)	0.017	8 (80)	19 (42)	0.03
Left ventricle free wall rupture, n (%)	0	0	0		0	0	
Low cardiac output syndrome, n (%)	0	0	0		0	0	
Re-exploration, n (%)	1 (2)	0	1 (6)	0.48	0	1 (2)	0.63
Respiratory insufficiency, n (%)	2 (4)	1 (3)	1(6)	0.60	2 (20)	0	0.002
Gastrointestinal bleeding, n (%)	0	0	0		0	0	

Temporary renal replacement therapy, n (%)	2 (4)	0	2 (11)	0.039	0	2 (4)	0.49
Cerebral vascular accident, n (%)	0	0	0		0	0	
Wound complication, n (%)	2 (4)	1 (3)	1 (6)	0.60	0	2 (4)	0.49
Complete atrioventricular block, n (%)	0	0	0		0	0	
New onset postoperative atrial fibrillation, n (%)	7 (13)	6 (16)	1 (6)	0.27	2 (20)	5 (11)	0.45



Variable	Hospital mortality or prolonged hospital stay >30 days (n=8)	P value
	Yes (n=8) No (n=47)	
Mean age, years ± SD	64.5±20.7 71.8±9.5	0.11

Age >80 years, n (%)	3 (63)	7 (15)	0.12
Female, n (%)	3 (38)	33 (70)	0.07
NYHA III or IV	4 (50)	12 (16)	0.16
Congestive heart failure, n (%)	4 (50)	11 (23)	0.12
Chronic kidney disease, n (%)	4 (50)	9 (19)	0.07

Atrial fibrillation, n (%)	1 (13)	19 (40)	0.13
Pre-operative LV EF, % (mean ± SD)	64.5±9.3	60.0±12.2	0.33
Pre-operative LV EF <40%, n (%)	0	3 (6)	0.46
EuroSCORE II, % (mean ± SD)	7.92±4.3	5.08±5.5	0.17
Infective endocarditis, n (%)	4 (50)	4 (9)	0.002

Previous cardiac surgery, n (%)	4 (50)	14 (30)	0.26
Concomitant procedure, n (%)	4(50)	21 (45)	0.78
Tricuspid valve procedure, n (%)	2 (25)	9 (19)	0.70
Mechanical valve replacement, n (%)	2 (25)	5 (11)	0.26

Table 5

Study	Country	Study period	Group (MT/ST)	Participants (male/ female)	Age (years)	MVP/MVR (%)	Mortality events (%)	Stroke events (%)		CPB time (min)	Clamping time (min)	Conversion to sternotomy (no. of patients)
Bolotin et al. [15]	USA	January 1996– June 2003	MT	38 (-/-)	68±2	42/58	2 (5)	NA	NA	160±65	NA	0
			ST	33 (-/-)	63±2	9/91	2(6)	NA	NA	157±54	NA	

Hiraoka et al. [16]	Japan	January 2006– September 2011	MT	10 (5/5)	68±15	0/100	0 (0)	1 (10)	0 (0)	145±25	90±7 (VF-time)	NA
			ST	27 (18/9)	63±15	0/100	1 (4)	1 (4)	2 (7)	135±28	84±19	
Losenno et al. [17]	Canada	September 2000–August 2014	MT	40 (28/12)	68±14	20/80	2 (5)	2 (5)	1 (3)	201±63	123±37 (VF-time)	0
			ST	92 (38/54)	62±13	9/91	10 (11)	2 (2)	6 (7)	180±75	105±46	
Ricci et al. (12)	Italy	October 1997- January 2007	MT	241 (134/107)	65±11	23/66 (other <u>11%)</u>	12 (4.9)	14 (5.8)	12 (4.9)	117±46	71±31]
Sharony et al. (13)	USA	1995-2002	MT	100 (-/-)	NA	31/69	5 (5)	4 (2.5)	8 (5.0)	122±2.9	80±2.3	0
Thomson et al. (14)	UK	1985-2001	MT	125 (33/92)	63	0/100	8 (6.4)	2 (1.6)	4 (3.2)	83.6± 43.1	NA	NA
Present study	Japan	May 2014- November 2020	MT	16 (8/8)	69.5±14	0/100	1 (6)	0 (0)	0 (0)	137±31.7	94±33.1	0
			ST	5 (4/1)	68.6±7.8	0/100	0 (0)	0 (0)	0 (0)	244±94.2	176±77.6	

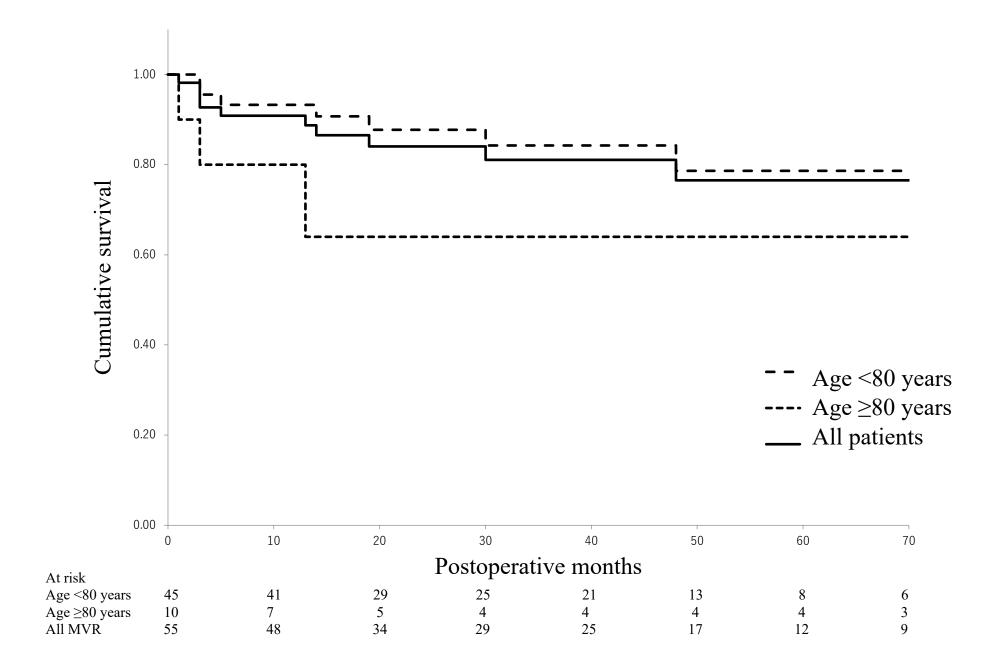


Fig. 1A

