Right ventricular mechanics and contractility after aortic valve replacement surgery: a randomised study comparing minimally invasive versus conventional approach

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ABSTRACT

Objective Minimally invasive aortic valve replacement surgery (MIAVR) is an alternative surgical technique to conventional aortic valve replacement surgery (AVR) in selected patients. The randomised study Cardiac Function after Minimally Invasive Aortic Valve Implantation (CMILE) showed that right ventricular (RV) longitudinal function was reduced after both MIAVR and AVR, but the reduction was more pronounced following AVR. However, postoperative global RV function was equally impaired in both groups. The purpose of this study was to explore alterations in RV mechanics and contractility following MIAVR as compared with AVR.

Methods A predefined post hoc analysis of CMILE consisting of 40 patients with severe aortic valve stenosis who were eligible for isolated surgical aortic valve replacement were randomised to MIAVR or AVR. RV function was assessed by echocardiography prior to surgery and 40 days post-surgery.

Results Comparing preoperative to postoperative values, RV longitudinal strain rate was preserved following MIAVR (−1.5±0.5 vs −1.5±0.4 1/s, p=0.84) but declined following AVR (−1.7±0.3 vs −1.4±0.3 1/s, p<0.01). RV longitudinal strain reduced following AVR (−27.4±2.9% vs −18.8±4.7%, p<0.001) and MIAVR (−26.5±5.3% vs −20.7±4.5%, p<0.01). Peak systolic velocity of the lateral tricuspid annulus reduced by 36.6% in the AVR group (9.3±2.1 vs 5.9±1.5 cm/s, p<0.01) and 18.8% in the MIAVR group (10.1±2.9 vs 8.2±1.4 cm/s, p<0.01) when comparing preoperative values with postoperative values.

Conclusions RV contractility was preserved following MIAVR but was deteriorated following AVR. RV longitudinal function reduced substantially following AVR. A decline in RV longitudinal function was also observed following MIAVR, however, to a much lesser extent.

INTRODUCTION

Minimally invasive aortic valve replacement (MIAVR) surgery has in selected patients become an alternative approach to conventional full sternotomy aortic valve replacement (AVR) surgery. A number of studies have compared the efficacy, long-term results, clinical outcomes and patient satisfaction between AVR and MIAVR in the treatment of severe aortic valve stenosis.3–4 However, less has been explored regarding the impact of
these surgical techniques on right ventricular (RV) function. Evaluation of RV function is of eminent clinical importance in the management of patients with a variety of cardiac diseases. It has been shown that RV function is one of the major determinants of symptoms and survival in patients with heart failure. Furthermore, it has been reported that perioperative RV dysfunction is a predictor of morbidity and mortality following AVR.

Previous studies have consistently reported a reduction in RV longitudinal function assessed by tricuspid annulus peak systolic excursion (TAPSE) and peak systolic velocity of lateral tricuspid annulus (RVS) following AVR. However, conclusions drawn from these findings regarding global RV function have been divergent. While some have proposed a reduction in global RV function as assessed by TAPSE and RVS, others have reported a preserved global RV function despite a reduction in TAPSE and RVS. These conflicting conclusions are reflecting several aspects of: (i) the complex geometry of the RV, making evaluation of global RV function complicated; (ii) the complex pathophysiology leading to decreased RV longitudinal function following AVR; and (iii) the commonly used measurements in previous studies (eg, TAPSE, RVS and fractional area change (RV-FAC)) are not pure measures of myocardial contractility but rather strongly dependent on the loading conditions and mechanical influences from the loss of pericardial support and adherences. While TAPSE and RVS are measures of RV mechanics, it has been shown that strain rate (RV longitudinal strain rate (RV-LSR)) is the best measure of myocardial contractility.

The inclusion criteria were (1) age ≥18 years; (2) severe AS: defined as peak velocity ≥4 m/s, mean gradient ≥40 mm Hg, aortic valve area (AVA) <1.0 cm² or indexed AVA <0.6 cm² by Doppler echocardiography in combination with two-dimensional echocardiographic morphology of severe valvular stenosis; (3) referred for surgical replacement of severe AS in adherence to current guidelines; and (4) sinus rhythm.

Patients were excluded if they had (1) reduced left ventricular ejection fraction (LVEF) <45%; (2) previous cardiac surgery; (3) concomitant other severe valvular heart disease; (4) coronary artery disease (CAD) requiring surgical intervention; and/or (5) urgent or emergent surgery.

Preoperatively, all patients were investigated by coronary angiography for evaluation of coexisting CAD. Echocardiography was performed within 1 week before surgery and 40 days post-surgery. Patients were analysed as treated (the patients who crossed over from one group to the other group were analysed in the new group).

**SURGICAL TECHNIQUES**

Minimally invasive aortic valve replacement surgery

MIAVR was performed via upper partial ministernotomy approach. Approximately a 6 cm vertical skin incision over the upper part of the sternum was applied. A partial J-shaped incision was extended into the right third intercostal space. A small vertical pericardial incision was applied anterior to the ascending aorta and subsequently aortic valve was revealed. CE-marked and FDA-approved mechanical and bioprosthetic aortic valves (conventional stented or sutureless bioprosthesis) were implanted. The pericardial incision was closed at the end of the procedure.

Cardiopulmonary bypass was established through a central arterial and peripheral venous cannulation. Antegrade crystalloid cardioplegia solution was used.

Conventional aortic valve replacement surgery

AVR was performed via full median sternotomy. A complete pericardial incision was performed. Myocardial

**MATERIALS AND METHODS**

**Study design**

The CMILE study was designed as a randomised, single-centre, open-label clinical trial. The methods and design of CMILE study has been previously described elsewhere. In brief, patients assigned to aortic valve replacement surgery at Karolinska University Hospital in Stockholm, Sweden, were eligible for inclusion into the study. Forty patients were randomised either to AVR or MIAVR in a 1:1 ratio, between October 2013 and July 2015.

All patients provided a written informed consent. The study is “Post-results” of CMILE study, registered at clinical trials.gov (NCT01972555).

**Patient population**

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Valvular heart disease

Echocardiographic assessment

Comprehensive transthoracic echocardiography examination was performed according to the recent guidelines. All examinations were performed on a Vivid E9 (GE, Healthcare, Horten, Norway). The images were stored digitally for subsequent analysis. An experienced echocardiographer, who was blinded to all clinical data, assessed all analysis off-line using commercially available software EchoPAC V.201 (GE Vingmed Ultrasound AS, Horten, Norway).

Right heart

RV size and function was quantified in accordance with American Society of Echocardiography Guidelines. RV-LS and RV-LSR were obtained by Speckle Tracking Echocardiography (STE) from RV-focused four-chamber view. The gain setting and the depth of cine loops were optimised in order to achieve a frame rate of at least 50/s. Subsequent analysis was conducted with dedicated, available commercial software for 2D strain analysis on EchoPAC V.201. The region of interest (ROI) was placed over the RV free wall. The width of the ROI was optimised in order to limit it to the myocardium. End systole was predefined by the software used for strain analysis, in which an automatic estimate of aortic valve closure was used. The software automatically calculated segmental and the global RV-LS and RV-LSR. RV stroke volume was calculated according to the existing guidelines.

RVS was measured as the maximal systolic velocity of the RV free wall displacement at the level of tricuspid annulus using cardiac state diagram. Right atrial (RA) volume was assessed with the same method as RA volume.

Automated analysis of the right ventricle

The myocardial traces from the tissue Doppler imaging (TDI) recordings were imported to a software called GHLab (Gripping Heart AB, Stockholm, Sweden), where the phases of the cardiac cycle and velocities were automatically identified by the software. The cardiac mechanical time events are defined according to the Dynamic Adaptive Piston Pump principle that describes the heart as a mechanical pump controlled by its inflow. The movement of the atrioventricular plane initiates the mechanical functioning of the heart and, therefore, the atrial contraction is considered as the starting point of the cardiac cycle. The different phases in the cardiac cycle are defined by shifts in myocardial mechanical work rather than by the opening or closure of the valves. The terms pre-ejection and post-ejection are used instead of isovolumic contraction and relaxation. ROI is chosen at the level of tricuspid annulus of RV free wall. The six phases of the cardiac cycle were identified as: atrial contraction, pre-ejection period, ventricular ejection time, post-ejection period, rapid filling/early diastole and slow filling/diastasis.

The duration of each phase, peak myocardial velocities during right ventricular ejection (RVS), tricuspid annular early diastolic velocity (RVE’) and tricuspid annular late
diastolic velocity (RVA′) were obtained from the software (GHLab) as shown in figure 1. Subsequently right ventricular index of myocardial performance (RIMP) and ratio between E′/A′ were calculated.

Statistical methods
The primary endpoints of this study have previously been reported in the CMILE study. 16 Statistical analyses were performed using SPSS (V.22) and MATLAB. Continuous data are reported as means±SD. Categorical data are presented as frequencies or percentages. Predata and postdata within each group were compared by paired Student’s t-test or Wilcoxon signed-rank test whenever suitable. Predata and postdata between AVR and MIAVR were compared using independent samples t-test or non-parametric independent sample whenever suitable. Categorical data were compared using χ². All tests were performed as two-sided tests with a significance level of 0.05.

RESULTS
A total of 40 patients were enrolled in the study, of which 20 patients were randomised to AVR and 20 patients to MIAVR. One patient who was randomised to MIAVR was converted to AVR intraoperatively due to difficulties in cardioplegia. Two patients who were randomised to AVR died after surgery (1) at 2 days post-surgery due to acute pancreatitis and multiorgan disease; and (2) at 4 days postsurgery due to undiagnosed liver cirrhosis and acute hepatic failure. These patients were excluded from the analysis. Finally, 19 patients were analysed in each group.

Baseline characteristics are presented in (table 1). Mean age was slightly but not significantly lower in the patients undergoing MIAVR compared with patients undergoing AVR. The number of patients with diabetes and previous stroke was slightly higher in the AVR arm but without reaching the statistical significance level. However, the proportion of patients suffering from renal impairment, and hyperlipidaemia was higher in the MIAVR group but did not reach statistical significance. There was no significant difference in patients’ medication between AVR and MIAVR groups, neither before surgery nor after surgery (online supplementary appendix 1).

In the AVR arm, 14 patients received bioprostheses: Edwards Perimount (n=10), St Jude Trifecta (n=2) and Medtronic Hancock II (n=2), and five patients received a mechanical prosthesis: St Jude Regent (n=4) and Sorin Slimline (n=1). In the patients undergoing MIAVR, 14 bioprostheses were also implanted, where 50% were sutureless: Sorin Perseval S (n=6) and Edwards Intuity (n=1), and five mechanical prostheses were implanted: Sorin Bicarbon Slimline (n=5). The mean valve size was 23 mm in both AVR and MIAVR groups. The cardiopulmonary bypass time was longer in the MIAVR group compared with AVR group (113.4±36 vs 87.4±28.2 min, p=0.04). Aortic cross clamp time was similar in the MIAVR and AVR groups (80±21.8 vs 82.9±26.9 min, p=0.72). The 30-day mortality was 10% in the AVR group and none in the MIAVR group. The perioperative/postoperative data are presented in supplementary appendix 2.

| Table 1 | Comparison of baseline clinical characteristics between patients undergoing AVR and patients undergoing MIAVR |
|---------|----------------------------------------------------------------------------------------------------------------|
| Characteristics               | AVR (n=19)                                      | MIAVR (n=19)                                      | P values | Total (n=38)  |
| Male                             | 11 (57.9%)                                      | 12 (63.2%)                                      | 0.74     | 23 (60.5%)    |
| Age (years)                      | 70.8±8                                          | 67.3±9                                          | 0.22     | 70±9          |
| Height (cm)                      | 170.9±9                                         | 172.8±9                                         | 0.55     | 172±9         |
| Weight (kg)                      | 75.5±12                                         | 84.1±19                                         | 0.15     | 81±16         |
| Hypertension                     | 13 (68.4%)                                      | 12 (63.2%)                                      | 0.73     | 25 (65.8%)    |
| Diabetes                         | 5 (26.3%)                                       | 4 (21.1%)                                       | 0.70     | 9 (23.7%)     |
| Previous stroke                  | 2 (10.5%)                                       | 0                                               | 0.15     | 2 (5.3%)      |
| Hyperlipidaemia                  | 6 (31.6%)                                       | 8 (42.1%)                                       | 0.50     | 14 (36.8%)    |
| Previous PCI                     | 0                                               | 1 (5.3%)                                        | 0.31     | 1 (2.9%)      |
| Renal failure                    | 0                                               | 2 (10.5%)                                       | 0.15     | 2 (5.3%)      |
| Systolic blood pressure (mm Hg)  | 132±13                                          | 134±20                                          | 0.70     | 133±16        |
| Diastolic blood pressure (mm Hg) | 72±10                                           | 73±8                                            | 0.89     | 73±9          |
| ECG                              |                                                  |                                                 |          |               |
| LBBB                             | 2 (10.5%)                                       | 0                                               | 0.16     | 2 (5.3%)      |
| RBBB                             | 1 (4.8)                                         | 1 (4.8)                                         | 0.97     | 2 (5.3%)      |
| QRS duration (ms)                | 95.7±16.8                                       | 93.4±10.3                                       | 0.64     | 94.6±13.8     |
| Heart rate (beats/min)           | 66±13                                           | 69±9                                            | 0.60     | 67±11         |

AVR, aortic valve replacement surgery; LBBB, left bundle branch block; MIAVR, minimally invasive aortic valve replacement surgery; PCI, percutaneous coronary intervention; QRS, the second wave in ECG; RBBB, right bundle branch block.
RV size and function

The end-diastolic RV transversal inflow diameter and proximal RV outflow diameter was similar in both groups before and after surgery (table 2).

Prior to surgery, there was no difference in RV-LS and RV-LSR between the patients undergoing AVR or MIAVR. Following surgery, RV-LS was deteriorated in both groups, although without any significant difference between them. RV-LSR was deteriorated in patients treated with AVR comparing preoperative values to values post-surgery (1.7±0.3 vs −1.4±0.3 1/s, p<0.01). However, RV-LSR was preserved in patients treated with MIAVR when comparing preoperative values with postoperative values (−1.5±0.5 vs −1.5±0.4 1/s, p=0.84) (figure 2).

At baseline, there was no difference in RVS between the patients undergoing AVR and patients undergoing MIAVR. After surgery, RVS declined in both groups, however, to a much lesser extent in the MIAVR group compared with the AVR group (18.8% vs 36.6%, p<0.001).

There was no significant difference in RV performance as assessed by RIMP between patients undergoing AVR and MIAVR at baseline. The RIMP was prolonged in both groups following surgery, although without any significant difference between the groups. The postoperative alterations in different phases of one cardiac cycle from baseline in one patient from each group is shown in figure 3.

Table 2: Preoperative baseline and postoperative echocardiographic RV measurements in the two surgical groups

| Variables               | Pre aortic valve surgery | Post aortic valve surgery | Between the groups (preoperative)† | Between the groups (postoperative)† |
|-------------------------|--------------------------|---------------------------|-----------------------------------|-------------------------------------|
| RV longitudinal strain (%) | AVR: −27.4±2.9           | MIAVR: −26.5±5.3         | 0.56                              | 0.25                                |
|                         |                          |                           | P values                          |                                     |
| RV strain rate (1/s)    | AVR: −1.7±0.3            | MIAVR: −1.5±0.5           | 0.32                              | 0.53                                |
|                         |                          |                           | P values                          |                                     |
| RVS (cm/s)              | AVR: 9±2.1               | MIAVR: 10.1±2.9           | 0.39                              | <0.001                              |
|                         |                          |                           | P values                          |                                     |
| RVE’ (cm/s)             | AVR: 8.4±2.9             | MIAVR: 7.1±2.1            | 0.14                              | 0.99                                |
|                         |                          |                           | P values                          |                                     |
| RVA’ (cm/s)             | AVR: 12.2±3.5            | MIAVR: 12.2±2.9           | 0.97                              | <0.001                              |
|                         |                          |                           | P values                          |                                     |
| RV E’/A’               | AVR: 0.84±0.6            | MIAVR: 0.59±0.2           | 0.16                              | 0.04                                |
|                         |                          |                           | P values                          |                                     |
| RV stroke volume (mL)   | AVR: 62.5±18.2           | MIAVR: 63.4±13.1          | 0.64                              | 0.014                               |
|                         |                          |                           | P values                          |                                     |
| RIMP                   | AVR: 0.75±0.2            | MIAVR: 0.68±0.2           | 0.26                              | 0.25                                |
|                        |                          |                           | P values                          |                                     |
| PEP (ms)               | AVR: 123±62              | MIAVR: 104±29             | 0.25                              | 0.69                                |
|                        |                          |                           | P values                          |                                     |
| ET (ms)                | AVR: 300.5±40            | MIAVR: 308.7±37           | 0.54                              | 0.56                                |
|                         |                          |                           | P values                          |                                     |
| POP (ms)               | AVR: 100.8±27            | MIAVR: 103.7±25           | 0.74                              | 0.19                                |
|                         |                          |                           | P values                          |                                     |
| RVOT prox (mm)          | AVR: 30.1±3.2            | MIAVR: 30.2±3.9           | 0.84                              | 0.79                                |
|                         |                          |                           | P values                          |                                     |
| RVD1 (mm)              | AVR: 31.1±4.0            | MIAVR: 28.1±4.4           | 0.75                              | 0.39                                |
|                         |                          |                           | P values                          |                                     |
| RA volume (mL)          | AVR: 40.3±16.7           | MIAVR: 42.6±16.1          | 0.16                              | 0.40                                |
|                         |                          |                           | P values                          |                                     |
| TR Vmax (m/s)           | AVR: 2.7±0.4             | MIAVR: 2.5±0.87           | 0.54                              | 0.67                                |
|                         |                          |                           | P values                          |                                     |
| SPAP (mm Hg)            | AVR: 37.5±9.1            | MIAVR: 31.7±13.7          | 0.76                              | 0.62                                |
|                         |                          |                           | P values                          |                                     |
| HR (min⁻¹)             | AVR: 69.6±12.3           | MIAVR: 63.4±9.1           | 0.22                              | 0.75                                |

Paired t test versus baseline.

*P<0.05; **p<0.01; ***p<0.001.

†Independent samples t-test to compare degree of changes seen following AVR with that seen following MIAVR.

RV, right ventricular; A’, late diastolic myocardial velocity; AVR, aortic valve replacement surgery; E’, early diastolic myocardial velocity; ET, ejection time; HR, heart rate; MIAVR, minimally invasive aortic valve replacement surgery; PEP, pre-ejection period; POP, post-ejection period; RA, right atrial; RIMP, right ventricular index of myocardial performance; RVD1, RV basal linear dimension; RVOT prox, proximal RV outflow diameter; S, peak systolic myocardial velocity; SPAP, systolic pulmonary artery pressure; TR, tricuspid regurgitation.
LV size and function
LV linear dimensions (LVEDD and LVESD) and LV volumes (LVEDV or LVESV) did not differ between the groups at baseline. LVEF was normal and without any difference when comparing patients undergoing AVR to patients undergoing MIAVR, both prior to surgery and postsurgery.

LV global longitudinal strain was within the normal range in both groups and remained within the normal range when comparing values and from baseline to post-surgery.

There was no difference in diastolic function between the AVR group and MIAVR group at baseline or after surgery (table 3).

**DISCUSSION**
The randomised, prospective CMILE study evaluated the impact of MIAVR and AVR on the regional and global RV function. In the present predefined post hoc analysis of the CMILE study, the principal findings were that MIAVR did not significantly affect RV intrinsic myocardial contractility assessed by RV-LSR, while both MIAVR and AVR induced decreased RV function when using methods that are more load dependent and dependent on extrinsic mechanical influence. RV contractile function is of significant clinical importance. It has been shown that RV contractility is one of the most important parameters for prediction of RV failure in patients treated with LV assist device and development of RV failure following

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*Figure 2* Changes in the parameters of right ventricular function in patients undergoing aortic valve replacement surgery (AVR) and minimally invasive aortic valve replacement surgery (MIAVR) comparing preoperative to postoperative values. RIMP, right ventricular index of myocardial performance; RV-FAC, right ventricular fractional area change; RV-LS, right ventricular longitudinal strain; RV-LSR, right ventricular longitudinal strain rate.
transplantation in patients with precapillary pulmonary hypertension. Furthermore, previous studies indicate that impaired RV contractile performance implicates poor outcomes in patients with valvular heart disease. Including more novel and conventional quantitative parameters of RV mechanics and contractility, we observed a significant reduction in regional and global RV function following AVR and MIAVR. Following surgery, RVS was reduced in both groups. However, the magnitude of reduction in RVS was significantly higher following AVR as compared with MIAVR, despite longer cardiopulmonary bypass time in the MIAVR group. Global RV function assessed by RV-LS and RIMP declined in both AVR and MIAVR without any significant difference between the groups. However, RV contractility assessed by RV-LSR was preserved following MIAVR while it was deteriorated following AVR. Previously, it has been acknowledged that load-dependent measures of RV function can change without changes in RV myocardial contractility.  

Figure 3  Example of right ventricular cardiac state diagram (CSD) displaying postoperative alterations in different phases of one cardiac cycle. AVR, aortic valve replacement surgery; MIAVR, minimally invasive aortic valve replacement surgery.

Table 3  Preoperative baseline and postoperative echocardiographic LV measurements in the two surgical groups

| Variables          | Pre aortic valve surgery | Post aortic valve surgery | P values       |
|--------------------|--------------------------|----------------------------|----------------|
|                    | AVR          | MIAVR             | AVR          | MIAVR             | Between the groups (preoperative)† | Between the groups (postoperative)† |
| LVEDD (mm)         | 44.8±5.1     | 46.3±5.3          | 43.7±4.9*    | 47.1±5.6          | 0.67                          | 0.03                          |
| LVESD (mm)         | 26.9±6.2     | 28.7±6.9          | 26.7±5.1     | 30.6±9.9          | 0.40                          | 0.03                          |
| LVEDV (mL)         | 85.6±19.9    | 102±30            | 82±17.2      | 100.5±29.7        | 0.14                          | 0.03                          |
| LVESV (mL)         | 34.9±10.4    | 42.3±13.3         | 36.1±10.4    | 45.5±14.9         | 0.11                          | 0.04                          |
| LVEF (%)           | 60.2±5.7     | 57.8±6.3          | 56.5±5.4**   | 54.6±6.5          | 0.33                          | 0.37                          |
| LVGLS (%)          | −16.5±8.1    | −17.4±2.1         | −15.7±2.7    | −16.9±2.1         | 0.64                          | 0.21                          |
| LVMPI              | 0.81±0.3     | 0.75±0.2          | 0.91±0.2     | 0.89±0.2**        | 0.37                          | 0.63                          |
| Mitral E (m/s)     | 0.81±0.3     | 0.80±0.3          | 0.84±0.2     | 0.89±0.3          | 0.72                          | 0.71                          |
| Mitral A (m/s)     | 0.98±0.4     | 0.81±0.4          | 0.89±0.3*    | 0.79±0.3          | 0.14                          | 0.61                          |
| Mitral E/A         | 0.84±0.2     | 1.2±0.9           | 0.98±0.3     | 1.2±0.4           | 0.14                          | 0.46                          |
| Dec T (ms)         | 228.8±61     | 201.1±58.4        | 200±50       | 203.7±49.7        | 0.30                          | 0.59                          |
| LA volume (mL)     | 83.2±23.3    | 79.6±20.5         | 75.2±22.1    | 79.8±23.7         | 0.53                          | 0.58                          |
| HR (min⁻¹)         | 69.6±12.0    | 63.4±9.1          | 72.5±13.7*   | 71.2±13.0*        | 0.22                          | 0.75                          |

Paired t test versus baseline.
*P<0.05; **P<0.01; ***P<0.001.
LV, left ventricular; A, late diastolic flow velocity; AVR, conventional aortic valve replacement surgery; Dec T, declaration time; E, early diastolic flow velocity; EDD, end-diastolic diameter; EDV, end-diastolic volume; EF, ejection fraction; ESD, end-systolic diameter; ESV, end-systolic volume; GLS, global longitudinal strain; HR, heart rate; LA, left atrium; MIAVR, minimally invasive aortic valve replacement surgery; Mitral E/A, ratio between early diastolic flow velocity and late diastolic flow velocity; MPI, myocardial performance index.
Our findings could provide an explanation for the discrepancies in earlier studies. The inconsistencies could thus be understood better from alterations in postoperative loading conditions and/or varying degrees of negative mechanical influence from the thoracotomy and pericardial manipulation. Incision of the pericardium due to altered pericardial constraint has been proposed to be the main factor responsible for deterioration of RV function. This hypothesis is further strengthened by our results revealing a post-surgical reduction in RVS and RV-LS following both AVR and MIAVR, despite the fact that during MIAVR only limited parts of the pericardium was incised, which was closed at the end of surgery. It can be speculated that RV is dependent on full pericardial support for maintaining its mechanical function, but this would not necessarily be due to the loss of myocardial contractility. Hence, opening even a small part of the pericardium may result in loss of that support. Furthermore, incision of the pericardium may result in geometrical changes of the RV, which is not restored even after closing the pericardium. Although RV longitudinal function was more preserved in MIAVR as compared with AVR, no difference in global RV function was observed between the groups. Although RVS correlates well with global RV function in normal hearts, it does not correlate well with global RV function after open heart surgery. Increasing PASP has also been proposed to contribute to deterioration of RV function following AVR. In our study, we observed a reduction in PASP following aortic valve surgery regardless of surgical approach.

The effect of AVR on RV longitudinal function in our study is consistent with the results of previous studies. Currently, few data are available regarding the effect of MIAVR on RV function. In contrast to our study, Unsworth et al have reported preserved RV longitudinal function following MIAVR. This discrepancy could be explained by the fact that there were only eight patients included in that study. Furthermore, as the type of surgical approach for MIAVR has not been reported, it is difficult to compare the results of that study to ours, since the location and size of the pericardial incision has been suggested to affect the magnitude of post-surgical modifications in RV longitudinal contraction.

To our knowledge, there are only a few studies available that used novel echocardiographic parameters comparing the impact of AVR and MIAVR on RV function in the context of RV mechanics and contractility. Although MIAVR was associated with a better preserved RVS, there was no significant difference between MIAVR and AVR in global RV function as assessed by RV-LS and RIMP. The state of postsurgical RV function can be translated into clinical outcomes. Dalen et al did not show any difference between AVR or MIAVR in 30-day mortality or 2-year survival.

Our findings indicate a more complex picture, where myocardial contractility might be more preserved from MIAVR compared with AVR, but both MIAVR as well as AVR seem to negatively impact RV mechanical function despite preserved contractility in the MIAVR group. A composite of echo-derived parameters should be incorporated in assessment of RV function. Future studies are warranted for investigating long-term results of AVR and MIAVR on RV function and its clinical implications.

Limitations
The main limitation of this study is the relatively small number of patients included in the study. Also, the follow-up time of 40 days in our study was relatively short. Future studies with longer follow-up are required to assess long-term impact of MIAVR and AVR on RV function. Since our study is primarily an echocardiographic study, we would not be able to translate the echocardiographic differences between the groups into clinical outcomes, for example, exercise capacity.

CONCLUSION
Both MIAVR and AVR influence RV mechanical function, but MIAVR does not result in a decline in RV contractility. A composite of conventional and novel echocardiographic parameters should be incorporated in assessment for RV function since many estimates of RV function are load dependent.

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