Heart team approach in treatment of mitral regurgitation: patient selection and outcome

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ABSTRACT

Objective A multidisciplinary heart valve team is recommended for the evaluation of treatment in patients with valvular heart disease, but evidence supporting this concept is lacking. In patients with severe mitral regurgitation, we thought to analyse the patient selection process by the heart team for different treatment options and the outcome after treatment.

Methods In this single-centre cohort study, all patients treated for mitral regurgitation between July 2013 and September 2018 were included. Primary end points during follow-up were all-cause mortality and a combined end point, consisting of all-cause mortality, cardiovascular rehospitalisation and mitral valve reintervention.

Results 179 patients (44.8%) were treated using Mitraclip, 185 (46.2%) by surgical repair and 36 (9.0%) by surgical replacement. The mortality risk according to EuroScore II differed significantly between treatment groups (6.6%±5.6%, 1.7%±1.5% and 3.6%±2.7% for Mitraclip, surgical repair and replacement, respectively, p<0.001). In-hospital mortality for the 3 groups were 3.4%, 1.6% and 8.3%, respectively (p=0.091). Overall, surgical repair patients had higher 4-year survival (HR 0.40 (95% CI 0.26 to 0.63), p<0.001) and fewer combined end points (HR 0.51 (95% CI 0.32 to 0.80), p<0.001) compared with surgical replacement and Mitraclip patients. However, patients undergoing Mitraclip for isolated, primary mitral regurgitation achieved very good long-term survival.

Conclusion The multidisciplinary heart team assigned only low-risk patients with favourable anatomy to surgical repair, while high-risk patients underwent Mitraclip or surgical replacement. This strategy was associated with lower than expected in-hospital mortality for Mitraclip patients and high 4-year survival rates for patients undergoing surgical or percutaneous repair of isolated primary mitral regurgitation.

KEY QUESTIONS

What is already known about this subject?
► According to current guidelines, a multidisciplinary heart valve team is recommended for the evaluation of treatment in patients with severe valvular heart disease; however, there is a lack of data supporting this approach, and the decision-making process is not well defined.

What does this study add?
► In this single-centre cohort study including 400 patients, we thought to analyse the patient selection process by the heart team for different treatment options patients with mitral valve regurgitation and the outcome after treatment.
► The multidisciplinary heart team assigned only low-risk patients with favourable anatomy to surgical repair, while high-risk patients underwent Mitraclip or surgical replacement.
► This strategy was associated with lower than expected in-hospital mortality for Mitraclip patients and high 4-year survival rates for patients undergoing surgical or percutaneous repair of isolated primary mitral regurgitation.

How might this impact on clinical practice?
► A consequent selection process by the heart team requires reduction of factors often biasing treatment assignment.
► In view of future treatment options including percutaneous mitral valve replacement, the heart team selection process becomes even more important and may influence patient outcome.

INTRODUCTION

Patients undergoing surgical treatment for mitral regurgitation (MR) have been studied for decades. In selected patient populations (relatively young patients, primary MR), surgical mitral valve repair (MVR) leads to outstanding repair rates and excellent long-term survival.1 2 However, in patients with secondary MR, and in particular in ischaemic heart disease, it is an ongoing debate whether surgical MVR repair or mitral valve replacement (MVR) is the treatment of choice.3 In addition, for elderly patients outcomes after mitral valve (MV) surgery are far less favourable.4 Percutaneous MV repair using the Mitraclip (MC) procedure...
Figure 1  Decision-making algorithm of the heart team compared with estimated and experienced in-hospital mortality. MDT, multidisciplinary heart team; MR, mitral regurgitation; MV, mitral valve; MVR, surgical MV replacement.

Methods

Patient population
In this retrospective cohort analysis, all patients treated for MR at the Heart Clinic Zurich between July 2013 and September 2018 were included. Three patients treated with percutaneous valve-in-valve/ring replacement were excluded as they have been reported elsewhere.9 Surgical treatment included isolated or combined MV surgery (MV surgery plus coronary bypass and/or additional valve surgery). In the percutaneous group, interventions counted as combined instead of isolated if the MC procedure was part of a beforehand planned series of interventions, including percutaneous coronary and/or other valve interventions. Clinical data, including follow-up data, were extracted from patient charts and by telephone interview. Twelve patients were lost to follow-up. Echocardiographic parameters were quantified according to current guidelines.10 11

MDT decision

The MDT of the Heart Clinic Zurich consists of cardiac surgeons (two dedicated MV surgeons with >600 MV repair operations each), interventional cardiologists (three experienced in MC procedure, main operator with >500 MC interventions), imaging and heart failure specialists as well as cardiac anaesthetists. The MDT meets weekly to discuss all heart valve cases and works as an organisationally and financially independent unit. In cases of disagreement between the team members regarding optimal treatment, decision is taken according to the majority principle of present team members. According to current guidelines, the MDT assigned all patients to one of the three treatment strategies (MC, surgical MV repair or primary MV replacement (I° MVR)).8 12 The primary MDT’s treatment intention was surgical MV repair. Anatomical amenability for MV repair was analysed using three-dimensional transoesophageal echocardiography.13 For patients deemed high risk for surgery but with suitable MV anatomy, percutaneous MV repair using MC was performed.14 Patients were considered ‘high surgical risk’ based on a combination of factors, including estimated mortality (EuroScore II) >4%, age >80 years and additional clinical risk factors not

* Additional criteria: Body mass index (BMI) <30 kg/m²; Diabetes mellitus, severe degenerative back disease, liver cirrhosis
## Table 1  Baseline characteristics of the study population (n=400)

| Clinical findings                          | MC (n=179) | MVrepair (n=185) | I° MVR (n=36) | P value |
|--------------------------------------------|------------|------------------|---------------|---------|
| **Age (years)**                            | 80.3 (8.1) | 65.8 (11.6)      | 70.4 (11.1)   | <0.001  |
| **Male sex, n (%)**                        | 105 (58.7) | 127 (68.6)       | 13 (36.1)     | 0.001   |
| **EuroScore II, (%)**                      | 6.6 (5.6)  | 1.7 (1.5)        | 3.6 (2.7)     | <0.001  |
| **STS score mortality, (%)**               | 4.6 (4.0)  | 1.0 (1.0)        | 3.2 (2.1)     | <0.001  |
| **Body mass index (kg/cm²)**               | 24.6 (4.5) | 24.7 (3.4)       | 25.7 (4.6)    | 0.354   |
| **Systolic blood pressure (mm Hg)**        | 122.4 (22.5) | 133.2 (17.5)   | 130.4 (22.4)  | <0.001  |
| **Heart rate (bpm)**                       | 77.8 (17.5) | 73.2 (15.7)      | 75.8 (15.4)   | 0.040   |
| **Atrial fibrillation, n (%)**             | 62 (36.9)  | 31 (17.1)        | 9 (26.5)      | <0.001  |
| **NYHA ≥III, n (%)**                       | 160 (92.0) | 45 (27.6)        | 14 (45.2)     | <0.001  |
| **Laboratory findings**                    |            |                  |               |         |
| **Haemoglobin (g/L)**                      | 125.4 (18.9) | 138.8 (13.5)   | 130.7 (16.2)  | <0.001  |
| **GFR (mL/min)**                           | 49.9 (19.4) | 77.7 (14.3)      | 70.6 (20.9)   | <0.001  |
| **NTproBNP (pg/mL)**                       | 10'998.6 (30'986.7) | 677.1 (1'024.1) | 2'203.8 (2'265.8) | 0.014   |
| **History of**                             |            |                  |               |         |
| **Hypertension, n (%)**                    | 139 (77.7) | 78 (42.2)        | 17 (47.2)     | <0.001  |
| **Diabetes mellitus, n (%)**               | 23 (12.8)  | 2 (1.1)          | 3 (8.3)       | <0.001  |
| **Coronary artery disease, n (%)**         | 96 (53.6)  | 40 (21.6)        | 11 (30.6)     | <0.001  |
| **Percutaneous coronary intervention, n (%)** | 68 (38.0) | 15 (8.1)        | 2 (5.6)       | <0.001  |
| **Coronary bypass graft surgery, n (%)**    | 28 (15.6)  | 0 (0.0)          | 2 (5.6)       | <0.001  |
| **Prior mitral valve intervention, n (%)**  | 8 (4.5)    | 2 (1.1)          | 5 (13.9)      | 0.001   |
| **Stroke, n (%)**                          | 16 (8.9)   | 9 (4.9)          | 0 (0.0)       | 0.074   |
| **Chronic obstructive pulmonary disease, n (%)** | 25 (14.0) | 4 (2.2)         | 1 (2.8)       | <0.001  |
| **Mitril regurgitation**                   |            |                  |               |         |
| ≥Moderate mitral regurgitation, n (%)      | 179 (100.0) | 185 (100.0)     | 34 (94.4)     | <0.001  |
| Severe mitral regurgitation, n (%)         | 165 (92.2) | 170 (91.9)       | 31 (86.1)     | 0.478   |
| **Aetiology of mitral regurgitation**      |            |                  |               |         |
| Primary, n (%)                             | 102 (57.0) | 175 (94.6)       | 30 (83.3)     | <0.001  |
| Secondary, n (%)                           | 64 (35.7)  | 9 (4.9)          | 4 (11.1)      | <0.001  |
| Combination, n (%)                         | 13 (7.3)   | 1 (0.5)          | 2 (5.6)       | 0.004   |
| **Further echocardiographic findings**     |            |                  |               |         |
| **LVEF (%)**                               | 50.7 (16.4) | 62.6 (7.4)       | 59.5 (10.3)   | <0.001  |
| **LVEDVI (mL/m²)**                         | 82.4 (39.2) | 75.0 (21.4)      | 73.7 (23.4)   | 0.070   |
| **LAVI (mL/m²)**                           | 71.4 (29.0) | 64.5 (30.3)      | 65.9 (19.0)   | 0.101   |
| **RV function reduced, n (%)**             | 66 (39.3)  | 11 (6.4)         | 4 (12.5)      | <0.001  |
| **RV/RA pressure gradient**                | 38.9 (13.5) | 29.0 (12.7)     | 32.6 (11.3)   | <0.001  |
| ≥Moderate tricuspid regurgitation, n (%)   | 53 (31.0)  | 28 (15.7)        | 7 (20.0)      | 0.003   |
| ≥Moderate aortic stenosis, n (%)           | 6 (3.6)    | 2 (1.1)          | 3 (9.1)       | 0.036   |
| ≥Moderate aortic regurgitation, n (%)      | 14 (8.2)   | 9 (5.0)          | 5 (15.2)      | 0.105   |

GFR, glomerular filtration rate; LAVI, left atrial volume index; LVEDVI, left ventricular end diastolic volume index; LVEF, Left ventricular ejection fraction; MC, Mitraclip; MV, mitral valve; MVR, surgical MV replacement; MVrepair, surgical mitral valve repair; NTproBNP, N-terminanted pro-brain natriuretic peptide; NYHA, New York Heart Association; RV, right ventricular; RV/RA, right ventricular/right atrial; STS, Society of Thoracic Surgeons.
Table 2  Anatomical and peri-interventional factors associated with in-hospital outcome

|                         | MC (n=179) | MVrepair (n=185) | I° MVR (n=36) |
|-------------------------|------------|------------------|---------------|
|                         | Success (n=152) | Failed/Died (n=27) | P value | Success (n=166) | Failed/Died (n=19) | P value | Success (n=33) | Died (n=3) | P value |
| **Mitral valve anatomy**|            |                  |              |            |                  |              |            |            |         |
| Calcification leaflet, n (%) | 6 (3.9) | 3 (11.1) | 0.118 | 5 (3.0) | 3 (15.8) | 0.009 | 17 (51.5) | 1 (33.3) | 0.560 |
| Calcification annulus, n (%) | 41 (27.0) | 9 (33.3) | 0.500 | 20 (12.0) | 10 (52.6) | <0.001 | 14 (42.4) | 1 (33.3) | 0.807 |
| **MV prolapse characteristics** |              |                  |              |            |                  |              |            |            |         |
| Single segment, n (%) | 37 (24.3) | 3 (11.1) | 0.130 | 55 (33.1) | 5 (26.3) | 0.550 | 3 (9.1) | 1 (33.3) | 0.212 |
| Two segments ant. or post., n (%) | 6 (3.9) | 3 (11.1) | 0.118 | 19 (11.4) | 2 (10.5) | 0.905 | 3 (9.1) | 1 (33.3) | 0.212 |
| Advanced only post., n (%) | 11 (7.2) | 2 (7.4) | 0.975 | 15 (9.0) | 2 (10.5) | 0.832 | 3 (9.1) | 0 (0.0) | 0.598 |
| Advanced ant.+post., not Barlow, n (%) | 17 (11.2) | 3 (11.1) | 0.991 | 40 (24.1) | 4 (21.1) | 0.769 | 5 (15.2) | 0 (0.0) | 0.482 |
| Barlow, n (%) | 5 (3.3) | 0 (0.0) | 0.342 | 21 (12.7) | 1 (5.3) | 0.349 | 5 (15.2) | 0 (0.0) | 0.482 |
| Flail leaflet, n (%) | 53 (34.9) | 4 (14.8) | 0.039 | 80 (48.2) | 11 (57.9) | 0.426 | 8 (24.2) | 0 (0.0) | 0.348 |
| **Extent of intervention** |              |                  |              |            |                  |              |            |            |         |
| Isolated mitral valve intervention, n (%) | 112 (73.7) | 18 (66.7) | 0.454 | 129 (77.7) | 9 (47.4) | 0.004 | 21 (63.6) | 0 (0.0) | 0.033 |
| MV+PCI (staged) or bypass surgery, n (%) | 27 (17.8) | 9 (33.3) | 0.063 | 14 (8.4) | 3 (15.8) | 0.296 | 3 (9.1) | 1 (33.3) | 0.212 |
| MV+TAVR (staged) or SAVR, n (%) | 15 (9.9) | 1 (3.7) | 0.304 | 6 (3.6) | 4 (21.1) | <0.001 | 3 (9.1) | 2 (66.6) | 0.005 |
| MV+TV intervention, n (%) | 1 (0.7) | 0 (0.0) | 0.675 | 21 (12.7) | 4 (21.1) | 0.313 | 7 (21.2) | 1 (33.3) | 0.640 |
| Triple valve surgery, n (%) | – | – | – | 2 (1.2) | 1 (5.3) | 0.187 | 1 (3.0) | 0 (0.0) | 0.768 |
| Single clip intervention, n (%) | 37 (24.3) | 5 (18.5) | 0.513 | – | – | – | – | – | – |
| Two clips, n (%) | 98 (64.5) | 13 (48.1) | 0.108 | – | – | – | – | – | – |
| ≥3 clips, n (%) | 17 (11.2) | 9 (33.3) | 0.002 | – | – | – | – | – | – |

ant., anterior; MC, Mitraclip; MV, mitral valve; MVR, surgical MV replacement; MVrepair, surgical mitral valve repair; NTproBNP, N-termined pro-brain natriuretic peptide; PCI, percutaneous coronary intervention; post., posterior; SAVR, surgical aortic valve replacement; TAVR, transcatheter aortic valve replacement; TV, tricuspid valve.
covered by EuroScore II such as frailty, obesity or liver disease. If neither MVrepair nor MC seemed feasible according to anatomical criteria, or the surgical repair effort appeared unreasonably high, primary MV replacement was performed.

**In-hospital outcome**

In-hospital outcome was measured as success of the initial treatment strategy and in terms of mortality. ‘Treatment success’ in surgical patients was defined as survival with residual MR ≤ Mild, no secondary MVR and absence of MV stenosis at discharge. The term ‘secondary’ MVR was used for patients in whom primary MVRepair failed and the surgical team decided intraoperatively to convert to MVR. Absence of mitral stenosis was defined as mean transvalvular gradient < 5 mm Hg, or a gradient ≥ 5 mm Hg when haemodynamically explained by heart rate and/or haemoglobin level. In MC patients, a residual MR ≤ Moderate at discharge with a reduction in MR by ≥ 1 grade from baseline was still considered as success. Pure technical success for MC patients was defined according to the M-VARc criteria. 5,12

**Follow-up outcome**

Primary end points during follow-up were all-cause mortality and a combined endpoint, consisting of all-cause mortality, cardiovascular rehospitalisation and MV reintervention.

Secondary end points during follow-up were severity of MR and dyspnoea according to the New York Heart Association classification (NYHA) after 3 months, 1 year and at the last follow-up examination before January 2019. Secondary end points were analysed separately for the accomplished intervention at discharge (MC, MVRepair or all MVR).

**Statistical analysis**

Continuous data are expressed as mean±SD, and categorical data as number and percentage (%). To compare data, we used the t-test, one-way analysis of variance (ANOVA) or repeated measures ANOVA, as appropriate. The Kaplan-Meier survival curves were constructed to estimate event-free survival for different subgroups and were compared with the log rank (Mantel-Cox) test. Univariate and multivariate Cox-regression model was used to analyse the predictors on combined end points. The conditional average treatment effect was estimated for various subgroups of patients as the difference in probability of a combined end point within 1 year between MVRepair and MC using an augmented inverse probability weighted estimate. Both the treatment and outcome model were fitted with a Random Forest regression and a missForest imputation was used for 5.9% missing covariate values, while the width of the CIs was approximated using Bootstrap. The level of significance was set at a p value of <0.05. All statistical analyses were performed using SPSS software (V.25.0, SPSS, Chicago, Illinois, USA) and R (V.3.4.1).

## RESULTS

**MDT decision and baseline characteristics**

Figure 1 demonstrates the assignment to the initial treatment strategy. Seventy-five per cent of MVRepair and 55% of MVR patients had isolated MV surgery. Baseline characteristics are summarised in table 1. Within the MC population, 57.0% of patients were treated for primary (EuroScore II 5.2%) and 35.7% for secondary MR (EuroScore II 9.0%). Of patients assigned to MVRepair, only 4.9% were treated for secondary MR (EuroScore II 3.1%).

**In-hospital outcome**

Treatment success rates at discharge for the initial treatment strategy MC, MVRepair and MVR were 84.9%, 89.7% and 91.7%, respectively (p=0.284). The treatment success rate for patients who underwent minimally invasive MVRepair for isolated MR was 93.6%. For MC patients, the technical success rate according to the M-VARc criteria was 99.4%.

One MC patient (0.6%) and 12 MVRepair patients (6.5%) needed peri-interventional conversion to MV replacement (secondary MVR) and were counted as failure of initial treatment strategy. Anatomical and
per-interventional factors associated with treatment success (or failure) for the three treatment arms are summarised in table 2. Of note, staged percutaneous interventions did not impact initial MC treatment success, while concomitant surgical procedures, in particular aortic valve replacement, were more frequent in patients in whom the surgical treatment strategy failed. A calcified leaflet and/or annulus was more frequent in patients in whom surgical MV repair failed but had no impact on MC or MVR treatment success.

The median (IQR) duration of hospital stay was 5 (4–7.5) days for MC, 10 (9–12) days for MV repair and 12 (10–19.3) days for MVR. Echocardiographic findings at discharge are summarised in table 3.

In-hospital mortality of MC, MV repair and MVR was 3.4%, 1.6% and 8.3%, respectively, $p=0.091$ (figure 1).
### Table 4: Predictors for combined end points

|                     | MC (n=179) | MVrepair (n=185) |
|---------------------|------------|-----------------|
|                     | P value    | HR 95% CI       | P value    | HR 95% CI       |
| **Univariate**      |            |                 |            |                 |
| Age >75 years       | 0.607      | 0.866           | 0.030      | 2.463           |
| Height <170 cm      | 0.505      | 0.860           | 0.034      | 2.376           |
| Female              | 0.166      | 0.723           | 0.087      | 2.018           |
| NYHA ≥3             | 0.426      | 1.286           | 0.106      | 2.043           |
| EuroScore II >2.5%  | 0.033      | 2.324           | 0.002      | 3.602           |
| Atrial fibrillation | 0.900      | 0.970           | 0.173      | 1.845           |
| Anaemia             | 0.047      | 1.565           | 0.027      | 2.893           |
| GFR <60 mL/min      | 0.093      | 1.551           | 0.147      | 2.093           |
| LVEF <60%           | 0.002      | 2.315           | 0.564      | 0.760           |
| LVEDVi >75 mL/m²     | 0.015      | 1.810           | 0.138      | 0.519           |
| RV function reduced | <0.001     | 2.324           | 0.207      | 2.189           |
| RV/RA gradient >30 mm Hg | 0.929 | 0.975 | 0.125 | 2.024 |
| **Multivariate**    |            |                 |            |                 |
| EuroScore II >2.5%  | 0.116      | 1.966           | 0.007      | 3.759           |
| Anaemia             | 0.043      | 1.631           | 0.662      | 1.312           |
| LVEF <60%           | 0.025      | 2.015           | –          | –               |
| RV function reduced | 0.016      | 1.871           | 1.125      | 3.109           |

GFR, glomerular filtration rate; LVEDVi, left ventricular end diastolic volume index; LVEF, Left ventricular ejection fraction; MC, Mitraclip; MVrepair, surgical mitral valve repair; NYHA, New York Heart Association; RV, right ventricular; RV/RA, right ventricular/right atrial.

### Long-term outcome according to initial treatment strategy

Outcomes of the entire cohort during the mean follow-up time of 32.2±17.6 months are reported in figure 2A,B. Single and multivariate regression analysis of factors predicting combined end points in MC and MVrepair patients are summarised in table 4.

When analysing the long-term impact of treatment success at discharge, failure of initial treatment strategy did not influence all-cause mortality for MC or MVrepair patients but led to more combined end points in MC patients (HR 0.528, 95% CI 0.289 to 0.964, p=0.034).

Concomitant heart treatments as well as MR aetiology were additional factors influencing outcome. Patients with combined surgical or percutaneous interventions had a significantly higher all-cause mortality compared with patients with an isolated MV intervention (surgical MV treatment: HR 3.419, 95% CI 1.300 to 8.993, p=0.008; MC patients: HR 2.753, 95% CI 1.569 to 4.892, p<0.001). Furthermore, MC patients with secondary MR had a worse outcome compared with MC patients with a primary MR (figure 2C,D). And in the subgroups of patients with primary MR with an isolated MV treatment (figure 2E,F), the 4-year mortality rates were particularly low in MC patients. Women older than 75 years with small body size appear to profit from the MC procedure when analysing for combined end points at 1 year (figure 3).

The secondary follow-up end points MR grade and dyspnoea according to NYHA class over time are shown in figure 4.

### DISCUSSION

#### MDT decision process

The concept of an interdisciplinary MDT is a centrepiece in modern structural heart disease treatment. Yet, there is no consensus on how the MDT should decide, and evidence supporting the effectiveness of MDT decisions for patients with severe MR is lacking. In our cohort study, we describe the selection process and outcome of an all-comer population with severe MR treated by a dedicated valvular MDT. Counter to the expectations generated by valvular guidelines, the percentage of MR patients assigned to (gold standard) surgical MVrepair was only 46.2%, while 44.8% received percutaneous repair by MC and 9.0% of patients were eligible for primary MVR. In the only other published cohort of patients with MV disease treated according to the MDT decision, the distribution was similar (23% percutaneous and 62% surgical MV repair, 15% MVR). In our cohort, mainly patients with primary MR, favourable repair anatomy and low surgical risk were offered surgical MVrepair, while patients with elevated risk or secondary MR were treated percutaneously. Such a consequent selection process was achieved as our MDT has reduced factors often biasing treatment assignment such as patient referral to a specific doctor or financial interest of the individual MDT members.

As additional treatment options such as percutaneous MV replacement will be available in the future, selecting the right patient for the right treatment will be even more challenging. Percutaneous MVR by compassionate
Subgroup analysis to evaluate whether MVrepair or Mitraclip may be better for patients with isolated, primary MR. GFR, glomerular filtration rate; LAVI, left atrial volume index; LVEDVi, left ventricular end diastolic volume index; LVEF, left ventricular ejection fraction; MC, Mitraclip; MV, mitral valve; MVrepair, surgical mitral valve repair; RV/RA, right ventricular/right atrial.

Figure 4 Mitral regurgitation grade (MR) and dyspnoea (New York Heart Association (NYHA) I–IV) from baseline to the last follow-up examination according to the treatment performed. MC, Mitraclip.
Effectiveness of the MDT treatment decision and room for improvement

The most important question is whether the MDT decision was the right one for the given patient. According to our data, patients assigned to MC achieved a lower in-hospital mortality than expected with surgical treatment, while patients assigned to surgical repair experienced in-hospital mortality rates that met the expectations according to EuroScore II. This was particularly relevant for patients with secondary MR treated by MC, where the reduction of the expected to the achieved in-hospital mortality was largest. Our consequent selection process thus improved the short-term outcome of high-risk surgical patients by assigning them to the lower risk percutaneous treatment option. Furthermore, the benefit of the less invasive percutaneous treatment seemed particularly relevant for MC patients who had isolated, primary MR. Long-term survival rates of these patients was excellent and better than in most previously published series.21 22 In addition, this study corroborates previous data showing a negative impact of concomitant heart treatments on long-term survival and thus emphasises the importance of a careful selection process by the MDT.23

The best treatment option for short elderly patients remains controversial. This study suggests that 1-year outcome for women with an age >75 years and small height (as well as patients with a calcified annulus) might have been further improved if a percutaneous approach had been chosen. Reduced MV repair rates in women have previously been reported24 and are possibly due to the lack of repair space in shorter elderly women with smaller annular dimensions and limited MV prolapse tissue. Two recent propensity-weighted analyses between surgical and percutaneous MV repair in elderly patients came to opposing conclusions, one favouring surgery and one favouring the percutaneous approach.25 26 The reasons for these discrepant results are unclear and need further analysis by randomised controlled trials.

Limitation

This study has the limitations attributed to its retrospective design and data collection. In addition, the decision-making process was not defined 'a priori' but represents the common understanding of our MDT. We cannot exclude that additional factors may have influenced decisions and outcome.

The small number of patients assigned to MVR did prevent further statistical analyses. It may also account for the higher than expected in-hospital mortality in this group (one expected, three actual deaths), a mortality rate comparable to the 5.4% operative mortality in a large series of patients undergoing MVR.27 It does, however, underscore the need for lower-risk percutaneous MV replacement options for this high-risk surgical group.

A similar limitation has to be acknowledged regarding patients with secondary MR. Our team opted for surgical treatment in only 15 patients (9 MV repair, 4 MVR), as opposed to 64 patients with secondary MR treated by MC. Therefore, this study cannot answer the question which treatment option is best for functional MR.

As we are a tertiary referral centre, patients not eligible for surgical or percutaneous treatment options are sent back to the primary cardiologist for optimal medical care. These (overall few) patients were not included in our study and we do not have follow-up data on them.

Conclusion

This study describes the MDT selection process in assigning patients for MR treatment and relates it to outcome. Only low-risk patients with favourable anatomy were offered MV repair, while high-risk patients underwent MC or MVR. This strategy was associated with lower than expected in-hospital mortality for MC patients and high 4-year survival rates for MV repair and MC patients with isolated primary MR.

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Data availability statement Data are available on reasonable request. All data relevant to the study are included in the article.

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