Influence of Mitroflow bioprosthesis structural valve deterioration on cardiac morbidity

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Abstract

Background: This study investigated the extent and nature of cardiac morbidity and cause of mortality in patients with Mitroflow structural valve deterioration (SVD).

Methods: A retrospective study was performed examining the medical records of patients who had received Mitroflow bioprosthesis between February 2001 and April 2014 and died during this period. A total of 211 patients were identified and included in the analyses. To determine the cause of mortality, cases were divided into three predefined groups: cardiovascular death due to SVD (group 1), cardiovascular death with no SVD (group 2) and non-cardiovascular death without SVD (group 3).

Results: Overall mortality in this study was 7.6% at 1 year, 46.4% at 5 years and 97.2% at 10 years. In group 1, 53 patients (25%) died; in group 2, 59 patients (28%) died; and in group 3, 99 patients (47%) died. Hospitalisation for congestive heart failure was observed in 49.1% in the SVD group vs. 10.2 and 13.1% in the two other groups, \( p < 0.001 \). Hospitalisation for endocarditis was also significantly higher in the SVD group (11.3%) than in the two other groups (6.8 and 0%), \( p < 0.05 \). Hospitalisation due to myocardial infarction, cerebral stroke, arrhythmia or other cardiac-related diseases was not significantly different between groups.

Conclusion: Structural valve deterioration in Mitroflow bioprosthesis was associated with a high prevalence of hospital admissions due to congestive heart failure and endocarditis. Patients with Mitroflow bioprosthesis should be systematically and routinely followed with echocardiography, and reoperation should be considered if SVD has developed.

Keywords: Aortic valve replacement, Structural valve deterioration, Bioprosthesis, Mitroflow, Cardiac morbidity

Introduction

Aortic valve replacement has improved life expectancy in patients with severe aortic valve stenosis. Bioprosthetic as opposed to mechanical valves are mainly preferred in patients >65 years, at which age shorter durability is thought to be counterbalanced by freedom from anticoagulation [1]. The Carpentier-Edwards (CE) Perimount (Edwards Lifesciences, Irvine, CA, USA) and the Mitroflow (LivaNova Group Inc., Vancouver, Canada) pericardial aortic bioprosthesis are commonly used worldwide [1–5]. Previous studies show satisfactory mid- and long-term results for both valves [1–5]. However, recent studies have reported early structural valve deterioration (SVD) in the Mitroflow valve compared with the Carpentier-Edwards (CE) Perimount valve [6]. Some studies show a pattern of SVD already 4 years after Mitroflow implantation [7, 8]. This is supported by a recent registry-based study which observed increased reoperation rates for Mitroflow prostheses size 19 and 21 compared with Carpentier-Edwards (CE) Perimount 19 and 21 mm valves [6].
Previous studies concerning Mitroflow valves have focused on all-cause mortality and showed increased crude mortality in Mitroflow bioprosthesis compared with other valve types. However, cause of death related to known SVD in Mitroflow bioprosthesis and the impact of living with a dysfunctional Mitroflow bioprosthesis on cardiac morbidity remain unclear. Therefore, further investigation is needed to clarify the influence exerted by the Mitroflow valve on cardiac morbidity and cause of mortality.

The aim of the present study was to investigate the extent and nature of cardiac morbidity and cause of mortality in patients with Mitroflow structural valve deterioration, and discuss relevant treatment options.

**Materials and methods**

We conducted a retrospective population-based cohort study that included all patients who died after a Mitroflow pericardial bioprosthesis implantation within the study period. The patients were selected from a single centre, Aarhus University Hospital, Denmark, between February 2001 and April 2014. A total of 440 patients who received aortic valve replacement with the Mitroflow bioprosthesis were identified from the Western Denmark Heart Registry [9]. Since all Danes have a unique identification number, the registry provided a complete follow-up for all of the patients who received a Mitroflow bioprosthesis [10]. A total of 267 patients had died at the end of the inclusion period. We censored the first postoperative 90 days for confounding issues regarding operation complications to be able to clearly relate morbidity and mortality to subsequent SVD. On this basis, a total of 227 patients were left in the study (Fig. 1).

Morbidity was defined as either cardiac or non-cardiac diseases. Cardiac morbidities included congestive heart failure, endocarditis, stroke, myocardial infarction, supraventricular and ventricular arrhythmia; the remaining cardiovascular diseases were called “other cardiac diseases”. Non-cardiac morbidities, viz. infection, respiratory disease and cancer, were defined as “other non-cardiac diseases”. Morbidity data were collected from patient charts. Due to missing patient charts, 13 patients were excluded from the analysis. Due to the small sample sizes of Mitroflow 23 (n = 2) and size 25 (n = 1), these cases were also excluded from the study. Consequently, a total of 211 patients with either Mitroflow valve size 19 or 21 were included in the morbidity and cause of mortality analyses (Fig. 1).

To determine the cause of mortality, the cases were divided into three predefined groups: cardiovascular death due to SVD (group 1), cardiovascular death with no SVD (group 2) and non-cardiovascular death without SVD (group 3).

The study was approved by the Danish National Health Board (ref no: 1–16–02–527-16) and the Danish Data protection Agency (ref. no: 2012-58-006).

**Definition of SVD**

SVD was defined by echocardiography as significant aortic valve prosthesis stenosis when the aortic valve area was below < 1.0 cm², calculated from the continuity equation or from direct planimetry using transoesophageal...
echocardiography. Assessment of prosthesis stenosis was also based on the velocity ratio between the peak velocity in the left ventricle outflow tract and peak velocity across the prosthetic valve. A velocity ratio < 0.25 was considered as significant stenosis. In the presence of colour Doppler, SVD was considered as severe aortic insufficiency (vena contracta > 6 mm) along with volume overload on the left ventricle. Moderate aortic stenosis with an aortic valve area between 1 and 1.5 cm² or a velocity ratio of 0.25–0.3 combined with moderate aortic regurgitation was also considered as significant SVD. The information was gathered from echocardiographic recordings.

Statistical analysis
Descriptive data are expressed as mean ± standard deviation for continuous variables and proportions for categorical variables. Time-to-event was calculated from 90 days after the operation until the date of death. The overall incidences are illustrated in graphs. Due to competing risk, cumulative incidence functions were performed to evaluate the incidence of SVD. Cumulative incidence functions were calculated at year 5 and 10. Hospitalisation due to cardiac co-morbidity was analysed with Fisher’s exact test. Statistical analyses were performed using STATA version 14.2 (StataCorp. 2015). A p-value < 0.05 was considered significant.

Results
Patient characteristics
A total of 211 patients were included. Preoperative patient characteristics are outlined in Table 1. The mean age at operation was 77 ± 2 years (range: 62–94 years) and 155 (73.5%) were females. The mean follow-up time from 90 days after operation until death was 5 ± 2.8 years. A total of 1064.9 patient years of evaluation was calculated. The size of the Mitroflow was distributed as follows: size 19 (30.3%) and 21 (69.7%). The indication for Mitroflow bioprosthesis implantation in 208 (98.6%) patients was aortic stenosis; in the rest (n = 3, 1.4%), it was endocarditis. Pre-operatively, 64% of the patients received diuretics, 71% aspirin, 47% beta-blockers, 29% ACE/ATII-inhibitors, 43% statins, 24% calcium antagonists, 12% vitamin-K antagonists and 3% amiodarone. During Mitroflow implantation, 47% received concomitant coronary artery bypass grafting. The three defined subgroups used for morbidity and cause of mortality analyses had the same average age (77 years). A total of six patients (2.8%) received reoperation during the follow-up time, where three patients had endocarditis and the remaining three had aortic stenosis.

Morbidity
Univariate analysis was performed to identify preoperatively clinical or echocardiographic variables that could predict later SVD development in patients with Mitroflow bioprosthesis. Univariate analysis showed no predicting variables for the development of SVD, p = NS (Table 2).

Of the total study population (n = 211), 53 patients (25.1%) developed SVD (group 1) during the follow-up period (13 years). The SVD group (group 1) had the largest hospitalisation rate for cardiac morbidity. In the pre-defined cardiac morbidity diseases, we observed the following: hospitalisation for congestive heart failure was observed in 49.1% of the patients in the SVD group (group 1) vs. 10.2% in group 2 and 13.1% in group 3, p < 0.001. Hospitalisation for endocarditis was also significantly higher in group 1 (11.3%) than in group 2 (6.8%) and group 3 (0%), p < 0.05 (Fig. 2). Hospitalisation due to myocardial infarction, cerebral stroke, arrhythmia or other cardiac-related diseases was not significantly different between the groups. Hospitalisation for cancer was lower in group 1 (3.8%) than in group 2 (8.5%) and group 3 (36.3%), p<0.001. Hospitalisation for other non-cardiac diseases was significantly higher in group 3 (68.7%) than in group 1 (56.6%) and group 2 (52.6%), p<0.05 (Fig. 3).

Mortality
During the follow-up time (13 years), 211 patients died after Mitroflow bioprosthesis implantation. In group 1, 53 patients (25%) died from SVD, in group 2, 59 patients (28%) died from cardiac disease without SVD, and in group 3, 99 patients (47%) died from non-cardiac disease without SVD. The overall mortality in this study was 7.6% at 1 year, 46.4% at 5 years and 97.2% at 10 years. No significant difference in all-cause mortality was found between groups 1, 2 and 3 (Fig. 4). The cumulative incidences at 5 and 10 years showed no significant difference in cause of mortality between groups 1 and 2. The cumulative incidence for groups 1 and 2 at 5 years was 9.5% (4.8; 14.3) vs. 12.2% (6.9; 17.5), and at 10 years 21.8% (15.1; 28.4) vs. 29.7% (18.5; 40.1), respectively. However, the cause-specific mortality in group 3 at 5 years was 13.6% (95% CI: 5.3; 21.9) and at 10 years 17.3% (95% CI: 3.6%; 31.0%) higher than in group 1, p < 0.05 (Fig. 5).

The cause-specific mortality analysis due to SVD was compared for Mitroflow 19 and 21 at 5 and 8 years. We observed a slightly higher but statistically insignificant risk of mortality among patients with a Mitroflow 19 compared with Mitroflow 21 (p < 0.4).

Discussion
In the present study, we describe for the first time cardiac morbidity and cause of death in a patient population that had received a Mitroflow bioprosthesis implantation. In particular, we identified patients who developed SVD in the Mitroflow bioprosthesis and
compared their cardiac-related morbidity and cause of death with that of patients without signs of SVD.

This study demonstrated that patients with SVD had a significantly higher hospital admission rate due to heart failure and endocarditis than patients who died from cardiovascular events without SVD and patients who had a non-cardiac death without SVD. It is well known that heart failure and endocarditis are serious complex conditions associated with reduced survival, requiring long and often repeated hospital stays and impaired quality of life. A total of 53% of patients with SVD were admitted with congestive heart failure, compared with 10% for group 2 and 13% for group 3. Acute admission with congestive heart failure often requires intensive care with intravenous diuretic and inotropic treatment. In-hospital stays are often extended by close follow-ups in an outpatient clinic, with substantial social and economic costs. Furthermore, our study shows that patients with congestive heart failure are often re-admitted (35%).

We know from other studies that congestive heart failure is associated with poor prognosis [11–13]. Randomised controlled studies with beta-blockers and angiotensin-converting enzyme treatment show increased mortality in congestive heart failure patients if they are not treated [14–17]. Therefore, systematic follow-up after Mitroflow implantation with transthoracic echocardiography is

| Table 1 Patient characteristics |
|--------------------------------|
| Total  | Group 1 | Group 2 | Group 3 | P-value |
|--------|---------|---------|---------|---------|
| **Age at operation (years)** | 76.8 ± 5.3 | 76.9 ± 5.7 | 76.9 ± 5.3 | 76.8 ± 5.2 | 0.992 |
| **Women** | 155 (73.5) | 42 (79.3) | 42 (71.2) | 71 (71.7) | 0.543 |
| **BSA (m²)** | 1.73 ± 0.19 | 1.72 ± 0.16 | 1.76 ± 0.21 | 1.72 ± 0.19 | 0.217 |
| **EuroScore II** | 1.6 ± 1.3 | 1.6 ± 1.1 | 1.6 ± 1.1 | 1.6 ± 1.6 | 0.999 |
| **NYHA** | 2.5 ± 0.7 | 2.5 ± 0.8 | 2.6 ± 0.8 | 2.4 ± 0.7 | 0.552 |
| **CCS** | 1.2 ± 1.3 | 1.2 ± 1.3 | 1.4 ± 1.2 | 1 ± 1.3 | 0.141 |
| **Blood pressure** | | | | | |
| Systolic (mmHg) | 147 ± 27 | 150 ± 28 | 144 ± 25 | 147 ± 27 | 0.574 |
| Diastolic (mmHg) | 79 ± 16 | 80 ± 20 | 77 ± 13 | 79 ± 15 | 0.730 |
| eGFR (mL/min/1.73 m²) | 50 ± 24 | 46 ± 18 | 52 ± 25 | 51 ± 25 | 0.574 |
| ≥50 | 53 (44.9) | 11 (39.3) | 19 (47.5) | 23 (46) | 0.783 |
| < 50 | 65 (55.1) | 17 (60.7) | 21 (52.5) | 27 (54) | 0.783 |
| **Smoker** | | | | | 0.503 |
| Never | 68 (32.5) | 21 (40.4) | 17 (28.8) | 30 (30.6) | 0.574 |
| Active | 95 (45.5) | 19 (36.5) | 31 (52.5) | 45 (45.9) | 0.574 |
| Quit | 46 (22) | 12 (23.1) | 11 (18.6) | 23 (23.5) | 0.574 |
| **Diabetes** | 26 (12.4) | 7 (13.5) | 6 (10.2) | 13 (13.1) | 0.830 |
| **Hypertension** | 115 (55) | 33 (63.5) | 34 (58.6) | 48 (48.5) | 0.173 |
| **Ischaemic heart disease** | 108 (51.9) | 31 (59.6) | 37 (63.8)* | 40 (40.8)* | < 0.001 |
| **Previous PCI** | 9 (4.3) | 4 (7.6) | 3 (5.3) | 2 (2.0) | 0.261 |
| **Previous CABG** | 7 (3.4) | 1 (1.9) | 4 (6.8) | 2 (2.1) | 0.224 |
| **Extracardiac arteriopathy** | 23 (10.9) | 7 (13.2) | 10 (17.0) | 6 (6.1) | 0.086 |
| **Atrial fibrillation** | 41 (19.8) | 15 (28.3) | 12 (20.7) | 14 (14.6) | 0.13 |
| **Apoplexia cerebri** | 22 (10.6) | 7 (13.2) | 6 (10.3) | 9 (9.4) | 0.765 |
| **COPD** | 37 (17.9) | 6 (11.5) | 11 (18.6) | 20 (20.8) | 0.365 |
| **Pre-operative echo** | | | | | |
| EF (%) | 54 ± 13 | 53 ± 13 | 54 ± 12 | 55 ± 13 | 0.782 |
| Aorta area (cm²) | 0.68 ± 0.2 | 0.68 ± 0.2 | 0.68 ± 0.2 | 0.68 ± 0.2 | 0.968 |
| Aorta CW max (mmHg) | 74 ± 28 | 73 ± 28 | 70 ± 23 | 76 ± 32 | 0.486 |

Values are mean ± standard deviation or as number and percent
BSA Body surface area, CABG Coronary artery bypass graft, CCS Canadian Cardiovascular Society, COPD Chronic obstructive pulmonary disease, EF Ejection fraction, eGFR Estimated glomerular filtration rate, PCI Percutaneous coronary intervention, NYHA New York Heart Association
*Significantly different, p < 0.05
necessary to identify SVD at early stages and prevent the development of congestive heart failure. However, this study showed no increased cardiac mortality in the SVD group compared with the non-SVD groups. One explanation could be early detection and treatment of SVD-related complications; another possible explanation could be the small population size, which is a limitation of our study.

Hospitalisation due to endocarditis after Mitroflow bioprosthesis implantation was mostly associated with the SVD group. Prosthetic heart valve endocarditis often requires lengthy intensive care with cardiac monitoring, 4–6 weeks of intravenous antibiotics and, in some cases, reoperation of the aortic valve. Furthermore, heart valve prosthesis endocarditis is frequently associated with severe extracardiac infective manifestations. After protracted hospitalisation, patients require extensive cardiac rehabilitation. From a health and socioeconomic perspective, it can be assumed that both congestive heart failure and endocarditis are associated with substantial burdens for patients and society, which should be taken into account when evaluating SVD.

Contrary to our expectations, this study found no significant difference in all-cause mortality between any of the three groups. When we adjusted for cause of death, we found that at 5 and 10 years, significantly more died in group 3 than in group 1 and group 2. The higher mortality in group 3 might be related to concurrent non-cardiac morbidities, such as cancer, that naturally increased the death rate. Even though patients in group 1 were much more frequently re-admitted postoperatively due to heart failure and endocarditis, these conditions did not seem to translate into a higher mortality rate compared with patients who died from cardiovascular conditions without SVD. However, the preoperative risk profile was similar. The differences in co-morbidities between the three groups complicated comparison between the groups. It would have been more accurate to compare the cause of death between the SVD group and the background population. Another limitation was the small size of the SVD group which also complicated comparison with groups 2 and 3.

Previous studies have demonstrated that age (below 70 years and above 80 years), Mitroflow valve size 19,
ejection fraction below 35% and poor New York Heart Association functional class were independent risk factors for developing SVD [5, 18, 19]. Other studies have not identified small valve size as a risk factor [20]. However, the present study found no clinical variables that could predict the development of SVD. A possible explanation could be that the problem of developing SVD does not reside within the patient but in the structure of the Mitroflow valve itself [21]. In contrast to a previous case series [18], Lus F et al. found that the DLA Mitroflow model with its anticalcification treatment did not emerge as a protective factor against SVD in their study [20]. However, it has been mentioned that the pathologic mechanisms of SVD, concerning the degenerated LA/LXA Mitroflow prostheses shows more often calcification than cusp tear, in accordance with the results reported by Luk et al. [21]. Patients with a degenerated Mitroflow prosthesis developed prosthesis stenosis more often than regurgitation. This fact has important clinical consequences since patients with SVD withstand more easily a pressure than a volume overload.

Fig. 3 Kaplan Meier curve for all-cause mortality. Group 1: Cardiovascular death due to SVD; Group 2: Cardiovascular death with no SVD; Group 3: Non-cardiovascular death without SVD. P = NS

Fig. 4 Cause-specific mortality. Mortality is defined by the three different groups. Group 1: Cardiovascular death due to SVD; Group 2: Cardiovascular death with no SVD; Group 3: Non-cardiovascular death without SVD
This is also in agreement with our long-term observations regarding more hospitalizations for congestive heart failure caused by the pressure overload observed.

We suggest that patients with Mitroflow bioprosthesis are systematically followed postoperatively with trans-thoracic echocardiography since SVD is related to severe morbidity and has financial consequences for patients and hospitals. Early detection of SVD could prevent related morbidities and reduce socioeconomic costs. Since most of the patients are elderly, reoperation of the aortic bioprosthesis is often not recommended due to the high risk of mortality. An alternative treatment is a valve-in-valve implantation. Valve-in-valve treatment has been reserved mainly for bioprosthesis size 21 and above, but several studies have had success with fracturing the ring of a small Mitroflow bioprosthesis by high-pressure balloon predilatation and then allowing for valve-in-valve treatment [22–25].

**Limitations**

We acknowledge the limitation that the present study reflects the experiences of a single centre with a limited number of patients. Another limitation is that this study only assesses the smaller Mitroflow valve sizes 19 and 21, so we cannot conclude anything about larger size valves. However, this study has a great strength as the registries allowed for complete follow-up. Additionally, patients were included consecutively from a single centre, which might minimise differences in pre-, peri- and postoperative care.

**Conclusion**

Structural valve deterioration in Mitroflow bioprosthesis was associated with a high prevalence of hospital admissions due to congestive heart failure and endocarditis. SVD developed in 25% of the total study population. Patients with SVD died from cardiac-related complications. The findings of this study have a number of important implications for future practice. Firstly, patients with Mitroflow bioprosthesis should be systematically and routinely followed with echocardiography; and secondly, reoperation should be considered if SVD has developed. Especially, the possibility of a valve-in-valve strategy with transcatheter aortic valve implantation should be considered.

**Abbreviation**

SVD: Structural valve deterioration

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**Availability of data and materials**

Please contact author for data request.
Authors’ contributions
All authors participated in conception and design of the study. FW and ZK obtained, organized and cleaned the dataset. FW performed data analyses.
All authors provided data interpretation and results interpretation. FW drafted the article and all authors provided critical revisions of the article. All authors provided final approval of the article.

Ethics approval and consent to participate
The study was approved by the Danish National Health Board (ref no: 1–16–02-527-16) and the Danish Data protection Agency (ref. no: 2012–58-006). Consent for publication
Not applicable.

Competing interests
The authors declare that they have no competing interests.

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