Minimally invasive redo mitral valve surgery without aortic crossclamp

Reoperação da valva mitral minimamente invasiva sem pinçamento da aorta

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Abstract

Introduction: Reoperations of the mitral valve have a higher rate of complications when compared with the first surgery. With the field of video-assisted techniques for the first surgery of mitral valve became routine, reoperation cases began to arouse interest for this less invasive procedures.

Objective: To assess the results and the technical difficulties in 10 patients undergoing minimally invasive redo mitral valve surgery.

Method: Cardiopulmonary bypass was installed through a cannula placed in the femoral vessels and right internal jugular vein, conducted in 28 degrees of temperature in ventricular fibrillation. A right lateral thoracotomy with 5 to 6 cm in the third or fourth intercostal space was done, pericardium was displaced only at the point of atriotomy. The aorta was not clamped.

Results: Ten patients with mean age of 56.9 ± 10.5 years, four were in atrial fibrillation rhythm and six in sinusal. Average time between first operation and reoperations was 11 ± 3.43 years. The mean EuroSCORE group was 8.3 ± 1.82. The mean ventricular fibrillation and cardiopulmonary bypass was respectively 70.9 ± 17.66 min and 109.4 ± 25.37 min. The average length of stay was 7.6 ± 1.5 days. There were no deaths in this series.

Conclusion: Mitral valve reoperation can be performed through less invasive techniques with good immediate results, low morbidity and mortality. However, this type of surgery requires a longer duration of cardiopulmonary bypass, especially in cases where the patient already has prosthesis. The presence of a minimal aortic insufficiency also makes this procedure technically more challenging.


Resumo

Introdução: Reoperações da valva mitral apresentam maior índice de complicações quando comparadas com a primeira cirurgia. Com o domínio das técnicas videoassistidas para as primeiras cirurgias da valva mitral, os casos de reoperações passaram a despertar interesse para esses procedimentos menos invasivos.

Objetivo: Analisar os resultados e as dificuldades técnicas da retroca valvar mitral minimamente invasiva em 10 pacientes.

Métodos: A circulação extracorpórea foi instalada por meio de colocação de cânulas femorais e cânula na veia jugular interna direita, conduzida em 28 graus de temperatura e fibrilação ventricular. Realizada toracotomia lateral direita com 5 a 6 cm no terceiro ou quarto espaço intercostal. Péricardio foi descolado apenas na região do átrio esquerdo no ponto da atriotomia. A aorta não foi pinçada.
Objective
The goal of this study is to describe the immediate results of ten patients who underwent video-assisted minimally invasive mitral valve reoperation, under hypothermic ventricular fibrillation and without aortic clamping.

METHODS
Tem patients who had previous mitral valve surgery were submitted to reoperation. As previously described [3], conventional induction of anesthesia was used. Conventional orotracheal cannula was used for mechanical ventilation hence one-lung ventilation was not needed. Patients were placed in dorsal decubitus with right side elevation at 15 degrees. Cardiopulmonary bypass was established with dissection of the left femoral artery and vein, followed by the insertion of arterial cannula designed for peripheral access into the femoral artery and the positioning of long venous cannula in the right atrium close to the superior vena cava, which was inserted via the femoral vein. Another venous cannula was inserted in the right internal jugular vein via puncture.

Thoracotomy was performed in the right fourth intercostal space, beginning right below the nipple and extending 4 to 6 cm laterally. The 6.5 mm 30° optics was inserted into the right pleura through the same space as the main incision, via access located 2 to 3 cm posterolateral to the main access. Cardiopulmonary bypass had begun and lungs had stopped when the intercostal space was opened and optics was inserted.

Pericardium was opened at the level of the left atrium and adhesions in that area were removed. A pair of temporary pacing wires was placed in the right ventricle and connected...
to a fibrillator. Patients were cooled down to 28°C and a catheter with CO₂ at 4 L/min was placed in the cavity. A small incision, less than 1 cm long, was made approximately 3 cm above the main incision in order to place the left atrial retractor. The opening of the left atrium was performed in the same way as in conventional surgeries. Prolene 4-0 sutures were applied to the edges of the left atrium to facilitate exposure. A customized retractor, which was designed specifically for this kind of surgery, was inserted into the left atrium (Figure 1).

The most technically challenging part of the procedure is the removal of previously implanted prosthesis. The removal is done in a similar way to conventional surgeries; however, the distance between the valve and the edge of the incision as well as the equipment used, which is usually quite delicate for this part of the surgery, make it difficult to remove the prosthesis.

The remaining steps of the surgery were performed routinely, with the passage of stitches through the valve ring done in a similar way to conventional surgeries. Both visibility of the ring and lowering of the prosthesis to its position were very good (Figure 2).

A long appropriate instrument is required for placing the knots to affix the prosthesis. Once the prosthetic valve was placed (Figure 3), a left ventricle aspirator was inserted via the left atrium and through the prosthesis and atriorrhaphy was done as usual. Defibrillation was performed externally.

After the cannulas were removed, heparin was completely reversed and the right pleural cavity was drained. The intercostal space used for access was closed with 2-0 ethibond suture and the skin was closed with intradermal suture.

- This study was approved by the Ethics Committee, protocol 0003761/10 on October 17th, 2010.

RESULTS

A total of ten patients, four males and six females, were evaluated. Patient ages ranged from 46 to 76 years-old, the mean being 56.9±10.5 years. Two patients (20%) were diabetic, six (60%) had hypertension, and four (40%) had history of smoking. Four patients (40%) were in functional class II, five (50%) were in functional class III, and one (10%) was in functional class IV. Electrocardiogram at rest showed sinus rhythm in six patients (60%) and atrial fibrillation in four patients (40%). Four patients (40%) had normal ejection fraction, five patients (50%) had moderate left ventricular dysfunction, and one patient (10%) had severe ventricular dysfunction. Five patients (50%) had undergone previous mitral valve repair; the other five (50%), mitral valve
replacement with biological prosthesis. The diagnosis which led to a new surgical procedure was: stenosis in two patients (20%), insufficiency in three patients (30%), and double dysfunction in five patients (50%). Mean length of time between the first and second surgeries was 11±3.43 years, ranging from 7 to 17 years. EuroSCORE ranged from 7 to 12, with a mean of 8.3±1.82.

Mechanical prostheses were implanted in eight patients (80%) and biological prostheses were implanted in the other two patients (20%). Size of the prosthesis ranged from 25 to 29 mm, with seven patients (70%) receiving a 27 mm prosthesis.

Mean time of ventricular fibrillation was 70.9±17.66 minutes, ranging from 45 to 100 minutes. Cardiopulmonary bypass time ranged from 66 to 150 minutes, with a mean of 109.4±25.37 minutes. After surgery, patients were taken to the ICU, where mean time of mechanical ventilation was 6.4±3.2 hours, ranging from zero to 12 hours. Length of stay in the ICU ranged from 2 to 3 nights, the mean being 2.5±0.52 nights. Six patients (60%) had low output syndrome after removal of cardiopulmonary bypass, requiring inotropic support. None of the patients in this series needed the reoperation due to bleeding nor did they have stroke, acute myocardial infarction, or acute renal failure. One patient (10%) developed pneumonia in the postoperative period. There were no deaths in this study.

At hospital discharge, five patients (50%) had sinus rhythm and five patients (50%) had atrial fibrillation. Mean length of hospital stay was 7.6±1.5 days, ranging from 5 to 10 days.

**DISCUSSION**

Since the mid-1990s, the medical community has shown growing interest in mini-incisions for valve replacement [4-8]. The right anterolateral thoracotomy seems to be an interesting access for mitral valve surgeries as it allows for direct vision of the line where left atriotomy is usually performed, good exposure of the valve, and minimum initial discomfort for the surgeon. In addition, aesthetic result of this incision is superior to others, since, in women, the breast hides the incision [9].

In the last decade, the use of video in cardiac surgeries has become commonplace. Some studies [10-15] have demonstrated that, in video-assisted surgeries, incisions can be smaller, visualization of the surgical field is improved, and long-term results are the same as those of conventional mitral valve surgery. Furthermore, there is substantial reduction in trauma, mechanical ventilation as well as hospital stay, so the patient can return to routine activities faster than in conventional surgeries, where the patient cannot drive for 45 days as opposed to 10 to 15 days in minimally invasive surgeries.

Dogan et al. [11] published an article comparing conventional and minimally invasive surgeries. Forty patients were randomly assigned to two groups, Group I being comprised of patients undergoing minimally invasive surgery and Group II of patients undergoing conventional surgery. Cardiopulmonary bypass and aortic clamping times were slightly longer in Group I; however, there were no statistically significant differences.

In Group I, none of the patients needed to be converted to conventional surgery and length of both mechanical ventilation and hospital stay was shorter, but again there were no statistically significant differences. There was also no difference between the two groups in terms of drainage by chest drains and pulmonary function. There were no major complications or deaths reported in this series; however, the authors stated that the biggest challenge in Group I was myocardial protection, as they faced a series of problems with endovascular aortic clamping in addition to difficulties using conventional occlusion forceps. The authors concluded that despite the lack of statistically significant data in favor of minimally invasive surgeries, they proved to be just as effective as conventional surgeries, which justifies performing the less invasive procedure.

In a similar article, Holzhey et al. [16] published a study comparing conventional and minimally invasive surgeries in patients over 70 years-old. Like other studies [17-19], the authors stated that duration of cardiopulmonary bypass, aortic clamping as well as the surgery itself was longer. There were no differences in terms of major complications and mortality in 5 to 8 years of follow-up. In the immediate postoperative period, the conventional surgery group showed greater incidence of arrhythmia and pacemaker implant; once again, the difference was not statistically significant. The conclusions reached were the same as the ones from other authors: minimally invasive surgeries are at the very least equal to conventional surgeries and the biggest challenge lies in myocardial protection methods.

Ventricular fibrillation associated with hypothermia has been used as myocardial protection in coronary surgeries for many years, based on the fact that a decompressed heart in ventricular fibrillation associated with hypothermia at 28°C consumes roughly the same amount of energy as a heart in cardiac arrest induced by cardioplegia [3].

On the other hand, in valve surgeries, this kind of heart protection technique has always caused certain concern related to the incidence of stroke, since the cavities are open and the aorta is not clamped.

In a 2008 article, Umakanthan et al. [20] demonstrated how safe ventricular fibrillation associated with hypothermia and without aortic clamping is during minimally invasive mitral valve surgeries. The authors assessed 195 patients who had undergone minimally invasive mitral valve surgery with this kind of myocardial protection. The results
showed incidence of mortality in 30 days at 3%, low output syndrome at 4%, and stroke at 3%, which are equivalent to the results reported by other authors who used myocardial protection with aortic clamping [10,11,16]. The authors attributed the low incidence of stroke to: arterial pressure being kept above 30 mmHg during cardiopulmonary bypass, insertion of cannula for continuous aspiration into the left ventricle, and continuous CO₂ instillation in the cavity throughout the procedure. The authors concluded this type of cardiac management is safe, easily reproduced, and it makes minimally invasive surgery simple.

After this study, minimally invasive mitral valve surgery, which had been limited to patients who had had no previous surgical intervention, started to be performed in patients undergoing reoperations as well.

In patients who have undergone previous surgeries, the heart tends to have a large amount of adhesions, which are difficult to remove through minimally invasive methods, making the aortic clamping almost impossible to achieve. That, as well as inconsistencies in the occlusion of the aorta via endovascular techniques severely limits the indication of less invasive procedures to patients undergoing reoperations. Aortic clamping is unnecessary when the patient is under ventricular fibrillation; therefore, it is possible to remove only the adhesions on top of the left atrium where the incision is made.

In 2010, Ricci et al. [21] presented a study in which 241 patients who had undergone previous mitral valve or other cardiac surgeries were evaluated. The authors started by describing the risks of performing a resternotomy due to adhesions, which could lead to serious injuries, especially to the right ventricle and innominate vein. In the results, the authors mentioned the need to convert the surgery to sternotomy in two patients, one due to aortic dissection; the other, left ventricular perforation. Average length of stay in intensive care was 24 hours, average time of mechanical ventilation was 12 hours, and average blood drainage in 24 hours was 450 ml. Ventricular fibrillation was used in six of the patients who underwent surgery; in the remaining patients, endovascular clamping was used. Furthermore, the authors reported a stroke rate of 5.8%, none in the fibrillation group, and overall mortality of 4.9%. They concluded that the minimally invasive mitral valve reoperation is safe and its benefits are: very low incidence of surgical wound infection, short length of intensive care unit and hospital stay, good aesthetic results, and faster return to routine activities.

A report was published in 2008 by Fortunato et al. [22], describing the case of a patient who underwent endoscopic mitral valve surgery reoperation. The authors performed a comissurotomy in a valve that had been repaired 12 years earlier. They reached the conclusion that the procedure was feasible and effective.

In the present study, 10 patients who underwent minimally invasive mitral valve reoperation were evaluated. All patients underwent the whole procedure without aortic clamping, under hypothermia and ventricular fibrillation, and with continuous CO₂ instillation. The average EuroSCORE of the group was above 8, indicating a group of high risk patients. Yet, there were neither deaths nor strokes observed in this series. The biggest challenge faced in this study was the removal of the mitral prosthesis that had been implanted in the first surgery, which led to patients in this condition being under ventricular fibrillation for a longer period of time. Six patients developed low output syndrome after removal of cardiopulmonary bypass. Consequently, in the last cases, it took longer to start the reheating in order to avoid exposing a heart under fibrillation to higher temperatures. The effect of doing that was somewhat positive making it easier to remove the CPB. Presence of mild aortic insufficiency hinders the surgical procedure; therefore, in cases where there is slight to mild insufficiency, the procedure should not be performed.

CONCLUSION

Minimally invasive mitral valve reoperations under hypothermia associated with ventricular fibrillation without aortic clamping can be safely performed, with good immediate results, low incidence of complications, and no need for ample dissections of the heart. However, longer length of time on cardiopulmonary bypass is required, especially in patients who have previously implanted prosthesis. This type of procedure should not be performed in this manner if there is significant aortic insufficiency.

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