

Robotic cardiac surgery in Brazil

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Background: Brazil, the largest country and economy in South America, is a major driving force behind the development of new medical technologies in the region. Robotic cardiac surgery (RCS) has been evolving rapidly since 2010, when the first surgery using the DaVinci® robotic system was performed in Latin America. The aim of this article is to evaluate short and mid-term results in patients undergoing robotic cardiac surgery in Brazil.

Methods: From March 2010 to December 2015, 39 consecutive patients underwent robotic cardiac surgery. Twenty-seven patients were male (69.2%), with the mean age of 51.3±17.9 years. Participants had a mean ejection fraction of 62±5%. The procedures included in this study were mitral valve surgery, surgical treatment of atrial fibrillation, atrial septal defect closure, resection of intra-cardiac tumors, totally endoscopic coronary artery bypass and pericardiectomy.

Results: The mean time spent on cardiopulmonary bypass (CPB) during RCS was 154.9±94.2 minutes and the mean aortic cross-clamp time was 114.48±75.66 minutes. Thirty-two patients (82%) were extubated in the operating room immediately after surgery. The median intensive care unit (ICU) length of stay was 1 day (ranging from 0 to 25) and the median hospital length of stay was 5 days (ranging from 3 to 25). For each type of procedure, endpoints were individually reported. There were no conversions to sternotomy and no intra-operative complications. Patient follow-up was complete in 100% of the participants, with two early deaths unrelated to the procedures and no re-operations at mid-term.

Conclusions: Despite the heterogeneity of this series, RCS appears to be feasible, safe and effective when used for the correction of various intra- and extra-cardiac pathologies. Adopting the robotic system has been a challenge in Brazil, where its limited clinical application may be related to the lack of specific training and the high cost of technology.

Keywords: Minimally invasive surgery; robotic surgery; valve surgery; Coronary artery bypass; atrial septal defect



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Introduction

Robotic-assisted totally endoscopic cardiac surgery has been performed for more than fifteen years in Europe and United States of America (1-3). Since the first procedures, there have been extensive discussions regarding its safety, viability and clinical outcomes (4). A growing number of large clinical centers have become involved with use of this new modality, with centers showing particular interest in the potential for reduced surgical trauma. The proposed

benefits of reduced tissue manipulation and trauma include shorter length of hospital stay, earlier return to social and professional activities and greater patient satisfaction when compared with conventional procedures (4).

Recent changes in generalized access to medical information through the internet have increased patient desire for less invasive procedures (5). Increasing numbers of patients are pursuing new technologies and procedures that promise to improve their quality of life, minimize postoperative pain and accelerate recovery following

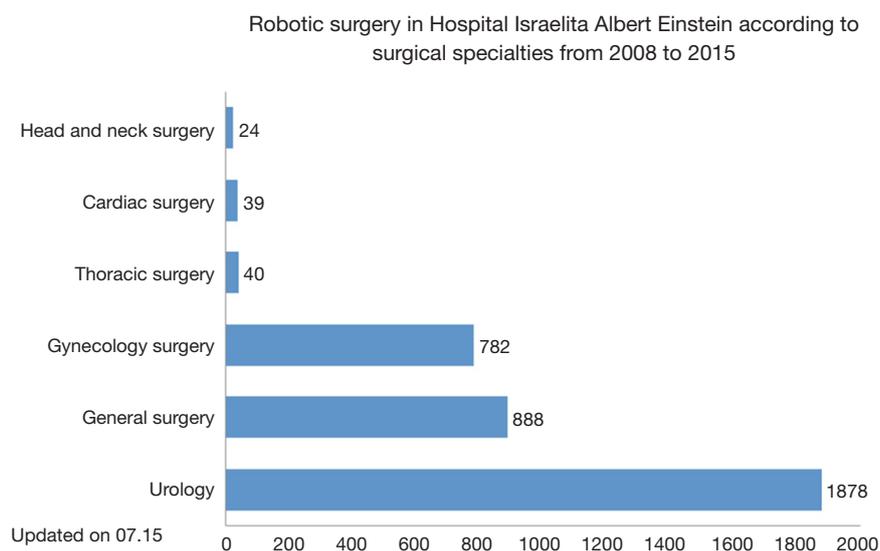


Figure 1 Robotic Surgery in Hospital Israelita Albert Einstein according to surgical specialties from 2008 to 2015.

surgical treatment. Medical institutions worldwide must continue to maintain a high level of excellence and to follow international guidelines of safety and quality of assistance (6).

In 2008, three private hospitals in the state of Sao Paulo incorporated robotic technology into their surgical armamentarium; however, only Hospital Israelita Albert Einstein implemented programs for all surgical specialties, including urology, gynecology, gastrointestinal, head and neck, thoracic and cardiac surgery. From 2008 to 2015, 3,651 patients underwent robotic surgery at Hospital Israelita Albert Einstein (*Figure 1*), corresponding to the majority of cases operated in the whole country. In 2010, the first robotic cardiac surgery in Latin America using the DaVinci® robotic system (Intuitive Surgical, Inc., Sunnyvale, CA, USA) was performed and, since then, 39 surgeries have been performed. Hospital Israelita Albert Einstein remains the only hospital robotic technology to use for cardiac surgery in Brazil.

This article aims to report the results of the first series of cardiac robotic surgeries performed in Brazil. The objectives of this study are to evaluate the short- and mid-term outcomes of patients undergoing robotic cardiac surgery in a general, tertiary-care, and philanthropic hospital.

Methods

This prospective study includes 39 consecutive patients who underwent robotic-assisted cardiac surgery using the DaVinci® robotic system (Intuitive Surgical, Inc.,

Sunnyvale, CA, USA), composed of a set of four robotic arms, an image capture and recording system and a console through which the surgeon controls the robot's movements.

All participants gave written informed consent. Participants were provided with information on the surgical possibilities regarding the performance of the procedure and chose the robotic-assisted approach. The study was approved by the Hospital Israelita Albert Einstein ethics committee under the number CEP/Einstein 11/1501, entitled "Einstein Registry of Robotic-assisted and Minimally Invasive Cardiovascular Surgery".

The procedures were performed between March 2010 and December 2015. Inclusion criteria followed the indications for conventional surgical correction of cardiac pathologies, whether acquired or congenital. Exclusion criteria for the performance of the robotic-assisted procedures included patients with thoracic deformities; severe thoracic trauma; anatomical or pathological abnormalities of the peripheral vascular system; moderate or severe aortic regurgitation; ejection fraction <40%; highly calcified coronary arteries, vessels with a diameter <1.5 mm or multi-vessel disease.

Patients who had mitral valve disease were screened using trans-esophageal echocardiography (TEE). Surgical correction was indicated following the American College of Cardiology (ACC) and the American Heart Association (AHA) guidelines for the management of patients with valve disease (7). Patient follow-up was performed in an outpatient setting, consisting of regular clinical evaluation,

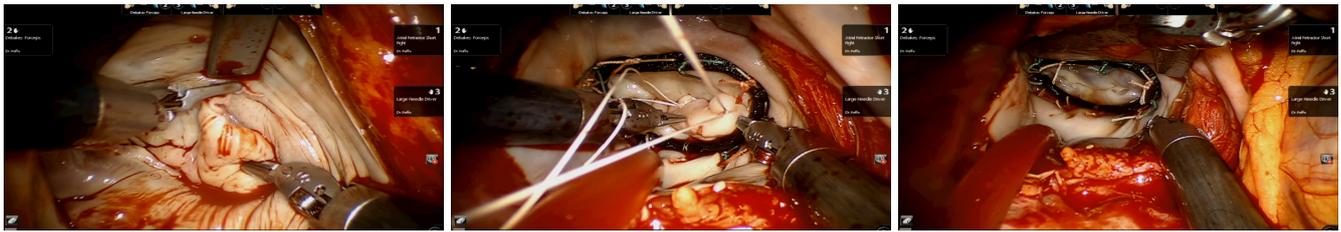


Figure 2 Mitral valve regurgitation repair (P2 prolapse) due to a ruptured chordae using polytetrafluoroethylene neo chords and a complete semi-rigid ring.

New York Heart Association (NYHA) classification and echocardiographic findings.

Surgical technique

Mitral valve operations

Robotic mitral valve repair or replacement was performed using a technique previously described by Chitwood *et al.* (8). Patients were intubated using a Robert Shaw endotracheal tube for selective pulmonary ventilation and positioned with the right hemi-thorax elevated at 20°. Disposable paddles for external cardiac defibrillation were placed in the right scapular and anterolateral region of the left hemi-thorax. A nasopharyngeal thermometer for body temperature control and a three-dimensional transducer for intraoperative trans-esophageal echocardiography were used.

After positioning the patient, markings were made on the thorax surface for trocar introduction. After systemic heparinization, peripheral cannulation of the femoral vein and artery, as well of the right internal jugular vein were performed under trans-esophageal echocardiography guidance. Prior to trocar introduction, the right lung was isolated from ventilation. The trocar for the micro camera was positioned prior to its introduction. The right hemi-thorax was inspected and then the remaining trocars were introduced. The DaVinci® robotic system (Intuitive Surgical, Inc., Sunnyvale, CA, USA) was docked.

Through the second right intercostal space (RICS) in the mid-axillary line (MAL), the Chitwood transthoracic clamp was introduced (Fehling Instruments GMBH & CO. KG, Karlstein, Germany). Cardiopulmonary bypass (CPB) was established and the patient was cooled to 28 °C. With the aid of videothoracoscopy, the ascending aorta was clamped and punctured with a 30-cm metal needle (Geister Medizintechnik, Tuttlingen, Germany) for the administration of antegrade crystalloid cold cardioplegic solution Custodiol HTK (Köhler Chemie GmbH,

Bensheim, Germany). With the aim of reducing the effects of air embolism, carbon dioxide was insufflated from the time of left atrial opening to the time of closure.

The mitral valve was inspected prior to the procedure and then repaired (*Figure 2*) or replaced. Under trans-esophageal echocardiographic monitoring, the heart was thoroughly de-aired, and CPB was discontinued. The femoral and jugular cannulas were removed, and heparin was reverted using protamine at a 1:1 ratio. The right hemi-thorax was drained and the drain was exteriorized through the trocar port of the right robotic arm, directed to the interior of the pericardial sac.

Two patients had atrial fibrillation associated with mitral valve disease, and underwent mitral valve repair. The first patient underwent epicardial ablation with electrical isolation of the pulmonary veins using the Cobra Adhere XL (Estech, San Ramon, CA, USA) system, as described by Bevilacqua *et al.* (9), prior to the establishment of CPB. The second patient underwent endocardial ablation of the left atrium with Cardioblate® Surgical Ablation Monopolar System Medtronic, Inc. (Minneapolis, MN, USA) (10).

Correction of atrial septal defect

Surgical access for ASD repair was similar to the approach used in robotic-assisted mitral valve surgery (11). In ASD repair, the right atrial chamber was accessed after isolation of the superior and inferior venae cavae. The introduction of the micro camera into the right atrium enabled the ASD to be visualized and corrected using a bovine pericardial patch (12). In all seven cases of this series, ASDs were classified as ostium secundum.

Intracardiac tumor resection

Robotic-assisted resection of intra-cardiac tumors also used a similar surgical approach to that described for robotic-assisted mitral valve surgery (13).

For tumours located in the left atrium, left atriotomy

was performed in the region of the right pulmonary veins following cardioplegia. In one patient, the tumor was attached to the left posterior atrial wall, which was broadly resected and replaced by a bovine pericardial patch. In a second patient, the tumor was adherent to the atrioventricular septum, anterior to the anterior cusp of the mitral valve. When the tumor was found in the right atrium, this chamber was opened and, without using cardioplegia, part of the right atrial free wall that the tumor adhered to was resected, and primary suture of the right atrium was performed.

Pericardiectomy

Robotic-assisted pericardial resection was performed by positioning ports in a similar way to that described for mitral valve repair. These procedures were performed off-pump. The pericardium was broadly resected, parallel to the right phrenic nerve, up to the region corresponding to the anterior aspect of the left ventricle.

Treatment of isolated atrial fibrillation

In the treatment of isolated AF, patient preparation, positioning and markings were the same as described for robotic mitral valve surgery. Prior to the introduction of trocars, the right lung was isolated. The micro camera was introduced into the fourth RICS, in the anterior axillary line (AAL). The right thorax was inspected and the remaining trocars were introduced. The DaVinci® robotic system (Intuitive Surgical, Inc., Sunnyvale, CA, USA) was docked.

After opening the pericardial sac, the superior and inferior venae cavae were dissected and a guide was introduced through the transverse sinus, posterior to the left atrium, directly below the pulmonary veins, until emerging in the oblique sinus. The ablation probe Cobra Adhere XL (Estech, San Ramon, CA, USA) was introduced using a guide, and was positioned for ablation. After correctly positioning via endoscopic approach, the epicardial ablation system was activated. After the first ablation, the ablation line was visually checked and, later, electrically checked using CARTO 3DTM (Biosense Webster, Johnson & Johnson, Diamond Bar, California, USA) device for electroanatomical mapping. If a gap was identified by the electroanatomical system, it was corrected by the electrophysiologist.

Totally endoscopic coronary artery bypass

A technique described by Bonatti (14) was used for totally

endoscopic coronary artery bypass (TECAB). TECAB was performed off-pump via a left thoracic approach. After single-lung ventilation had been established with isolation of the left lung, a 12-mm trocar was introduced in the fifth left ICS (LICS), for the introduction of the micro camera. Two trocars for the 8-mm robotic instruments were inserted in the third and seventh LICS in the left AAL. Carbon dioxide was constantly insufflated in the left thorax thus ensuring room for dissection of the left internal thoracic artery (LITA).

After robotic dissection of the LITA, a robotic coronary stabilizer was used, which was introduced in the left hemithorax and connected to the fourth robotic arm. The left anterior descending (LAD) artery segment to be anastomosed was mechanically immobilized and occluded proximally and distally using a tourniquet. After completion of the anastomosis, a TTFM (Medstim, Oslo, Norway) flow-meter was used to check adequate flow through the anastomosis. No patients underwent hybrid approach with additional stenting to other vessels.

Statistical analysis

Continuous variables were expressed as mean, standard deviation (SD), median and range, and categorical data were expressed as count and percentage. The Kaplan-Meier method was used for survival analysis. Data analysis was performed using SPSS software (version 18.0; SPSS, Chicago, IL, USA).

Results

The preoperative diagnoses are demonstrated in *Figure 3*, and the procedures performed are summarized in *Figure 4*.

Twenty-seven patients (69.2%) were male. The mean age was 51.3 ± 17.9 years, ranging from 20 to 83 years and the mean ejection fraction was $62 \pm 5\%$. The New York Heart Association (NYHA) functional class I/II was observed in 69.3% (27/39). The most commonly documented comorbidities included systemic arterial hypertension (43.6%), previous acute myocardial infarction (10.3%) and diabetes (7.7%). A history of cigarette smoking was recorded in 15.4% of patients.

The most common procedure performed was mitral valve surgery (N=20). In echocardiographic assessment, 18 patients showed severe mitral valve regurgitation (MR) due to degenerative change and two had double lesion due to rheumatic disease. Repairs were possible in 15 cases and

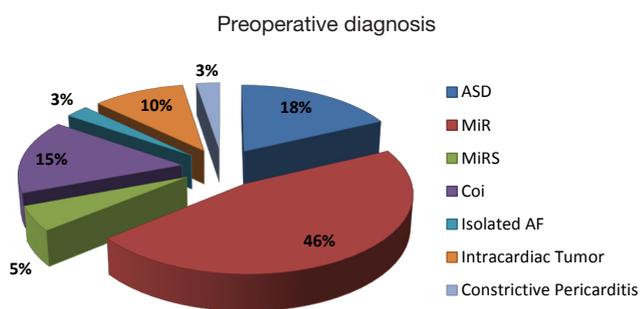


Figure 3 Preoperative diagnosis. ASD, atrial septal defect; MiR, mitral regurgitation; MiRS, mitral regurgitation and stenosis; Coi, coronary insufficiency; AF, atrial fibrillation.

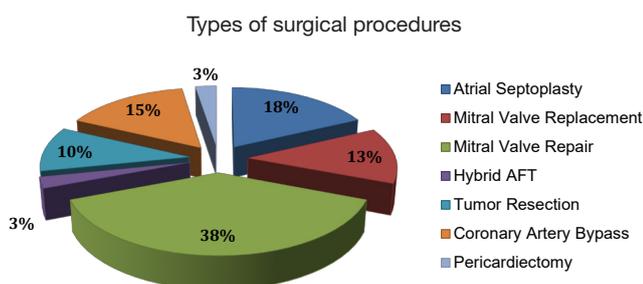


Figure 4 Types of surgical procedures. AFT, atrial fibrillation therapy.

five replacements were performed. Patent foramen ovale was identified in four patients in association with mitral valve disease, and atrial septal suture was performed. Two patients with mitral valve disease had atrial fibrillation and underwent mitral valve repair in addition to surgical ablation.

The remaining cases were assorted as follows: seven patients with ASD (17.9%) were submitted to atrial septoplasty with bovine pericardial patch; six patients (15.3%) underwent TECAB to the LAD due to severe stenosis (>70%) and four had already undergone previous angioplasty and progressed with intracoronary stent stenosis. In all cases, the left internal thoracic artery (LITA) graft was used to revascularize the left anterior descending artery. Four (10.2%) patients underwent tumor resection (three left atrial tumors and one right atrial tumor); one (2.5%) underwent hybrid treatment of atrial fibrillation with epicardial-isolation of the pulmonary veins and endocardial ablation; and one patient underwent pericardiectomy.

All procedures are summarized in *Table 1*.

There were no intra-operative complications. The mean

cardiopulmonary bypass and the mean aortic cross-clamp time are presented in *Table 2*, categorized by the type of procedure. TECAB, pericardiectomy and treatment of isolated atrial fibrillation were performed without the use of cardiopulmonary bypass.

After the procedure, all patients underwent intraoperative TEE showing successful valvuloplasty with absent or trivial mitral regurgitation, prostheses well positioned and functioning normally, no residual shunt after ASD closure and complete tumor resection.

Of the 39 patients, 32 (82,0%) were extubated in the operating room immediately after surgery, and the remaining seven patients were extubated in an ICU with a median tracheal intubation time of 9 hours (ranging from 5.8 to 22 hours). The median ICU length of stay was 1 day (ranging from 0 to 25) and the median hospital length of stay was 5 days (ranging from 3 to 25). Endpoints are shown in *Table 3*, categorized by type of procedure.

All procedures were performed as planned, with none requiring conversion to sternotomy. Post-operative complications included one case of unilateral pulmonary edema (UPE) (2.5%), two cases of atrial fibrillation with rapid ventricular response (5.1%) and three re-explorations for bleeding (7.6%). One patient developed a systemic inflammatory response syndrome (SIRS). Stroke was observed in one patient (2.5%) on the 20th postoperative day, two weeks after hospital discharge. No infections of the surgical sites were reported. One case of femoral pseudoaneurysm was reported following use of peripheral cannulation. However, this thrombosed spontaneously, with preserved arterial perfusion of the limb with no further complications.

In this series, two deaths (5%) were reported in the early post-operative period. One patient had acute respiratory failure due to a bronchopneumonia on the 25th post-operative day. A second patient died from complications of an ischemic stroke after hospital discharge on the 24th post-operative day.

There were no late deaths and all patients are being regularly evaluated on an outpatient basis. Mean follow-up time was $1,126.33 \pm 734.14$ days, ranging from 24 to 2,118 days (*Figure 5*).

Transthoracic echocardiography (TTE) was obtained every 6 months in all non-TECAB patients. In those patients who underwent mitral valve repair, no significant mitral regurgitation was documented. At mid-term, there was no need for re-operation.

On the qualitative analysis, all patients showed a marked

Table 1 Procedures performed

Patient	Gender	Age	Diagnosis	Treatment
1	F	34	ASD	Atrial septoplasty
2	M	39	ASD	Atrial septoplasty
3	M	52	MiRS + PFO	Mitral valve replacement + PFO closure
4	M	53	MiR	Mitral valve repair + annuloplasty
5	F	38	MiRS + PFO	Mitral valve replacement + PFO closure
6	F	22	LA Tumor	LA tumor resection
7	M	74	MiR	Mitral valve repair + annuloplasty
8	M	67	MiR	Mitral valve repair + annuloplasty
9	M	42	MiR	Mitral valve repair + annuloplasty
10	F	60	MiR	Mitral valve repair + annuloplasty
11	F	24	ASD	Atrial septoplasty
12	M	77	MiR + AF	Mitral valve repair + annuloplasty + epicardial ablation
13	M	51	AF	Hybrid AFT treatment
14	F	25	RA Tumor + SVCThrombus	RA tumor resection + SVC thrombectomy
15	M	49	Coi	Coronary artery bypass (LITA – LAD)
16	M	72	Coi	Coronary artery bypass (LITA – LAD)
17	M	32	MiR	Mitral valve repair + annuloplasty
18	M	81	Constrictive pericarditis	Pericardiectomy
19	M	57	Coi	Coronary artery bypass (LITA – LAD)
20	F	44	ASD	Atrial septoplasty
21	F	40	LA Tumor	LA tumor resection
22	M	36	ASD	Atrial septoplasty
23	M	44	MiR	Mitral valve repair + annuloplasty
24	F	60	LA Tumor	LA tumor resection
25	M	20	MiR	Mitral valve replacement
26	F	48	MiR	Mitral valve repair + annuloplasty
27	M	46	MiR	Mitral valve repair + annuloplasty
28	M	48	ASD	Atrial septoplasty
29	M	48	MiR	Mitral valve replacement
30	M	78	MiR	Mitral valve replacement
31	M	55	MiR	Mitral valve repair + annuloplasty + PFO closure
32	F	22	ASD	Atrial septoplasty
33	F	63	MiR	Mitral valve repair + annuloplasty
34	M	83	Coi	Coronary artery bypass (LITA – LAD)
35	M	75	Coi	Coronary artery bypass (LITA – LAD)
36	M	65	MiR	Mitral valve repair + annuloplasty
37	M	47	Coi	Coronary artery bypass (LITA – LAD)
38	M	49	MiR	Mitral valve repair + annuloplasty + PFO closure
39	M	81	MiR	Mitral valve repair + annuloplasty

F, female; ASD, atrial septal defect; M, male; MiR, mitral regurgitation; MiRS, mitral regurgitation and stenosis; PFO, patent foramen ovale; LA, left atrium; AF, atrial fibrillation; RA, right atrium; SVC, superior vena cava; Coi, coronary insufficiency; LITA, left internal thoracic artery; LAD, left anterior descending artery.

Table 2 Intraoperative data

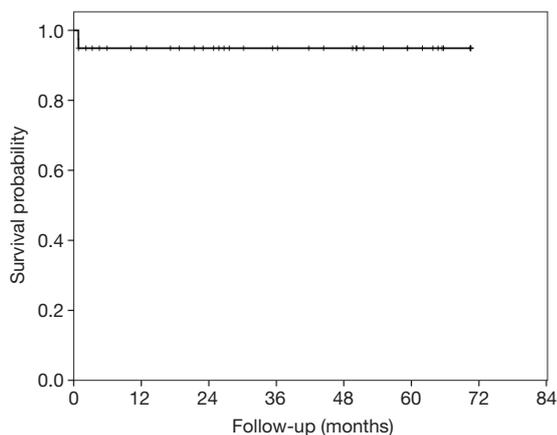
Type of procedure	Cardiopulmonary bypass time (min)		Aortic cross-clamp time (min)	
	Mean \pm SD	Range	Mean \pm SD	Range
Atrial septoplasty	160.06 \pm 86.10	140–240	117.18 \pm 71.71	75–150
Mitral valve repair	151.72 \pm 97.00	120–340	112.36 \pm 78.43	90–260
Mitral valve replacement	156.14 \pm 90.65	170–280	113.57 \pm 75.63	120–210
Tumor resection	128.94 \pm 95.11	120–180	86.94 \pm 73.96	90–105
All	154.92 \pm 94.22	120–340	114.48 \pm 75.66	75–260

*, values expressed in mean \pm standard deviation (SD) and range.

Table 3 Length of stay

Type of procedure	ICU length of stay (days)		Total hospital length of stay (days)	
	Median	Range	Median	Range
Atrial septoplasty	1.5	1–3	5	3–6
Mitral valve repair	1	1–7	5	3–15
Mitral valve replacement	1	1–25	4.5	3–25
Tumor resection	1	1–2	4	4–5
TECAB	5	4–9	5	5–9
All	1	0–25	5	3–25

*, values expressed in median and range. TECAB, totally endoscopic coronary artery bypass.

**Figure 5** Kaplan-Meier survival curve.

improvement in cardiovascular function, with NYHA functional class at pre-operative (I, 25.8%; II, 43.5%; III, 30.7%) and post-operative (I, 84.6%; II, 15.4%; III, 0%) intervals are shown in *Figure 6*.

Discussion

The use of minimally invasive cardiac surgery via mini-thoracotomy was, for many years, limited by the need for direct visualization of the heart and by the use of inappropriate instruments. With the incorporation of video-assisted techniques, it has been shown to be safe and effective in different surgical situations (15-19). However, the acquisition of new abilities such as manipulation

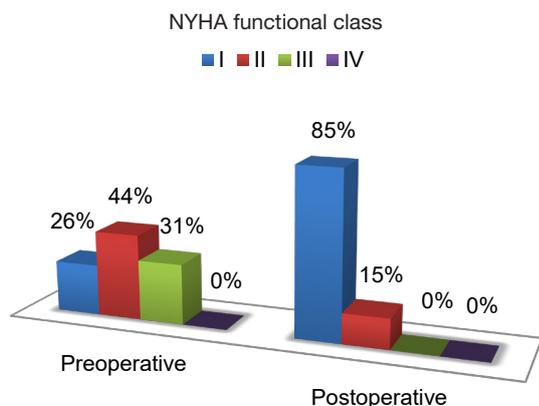


Figure 6 New York Heart Association functional class.

of longer instruments, the restricted operative field, replacement of direct vision by a two dimensional image and the change in tactile feedback represent a challenge to the surgeons. During the last decade, a variety of minimal-access approaches have been developed to access the mitral valve, from limited sternotomies to minithoracotomy assisted by videothoracoscopy. All had the same goals: to decrease surgical trauma and surgical complications leading to better patient recovery (20). The introduction of the DaVinci[®] robotic system significantly improved what can be visualized in the operative field, providing a three-dimensional image with ten times magnification. The use of the DaVinci[®] robotic system allows for better precision whilst performing cardiac surgeries through smaller incisions.

The most commonly performed procedure in this series was robotic mitral valve surgery. Mitral valve repair was possible in 15 patients, with a variety of techniques used: leaflet resections and reconstructions, chordal procedures (including chordal transfers and polytetrafluorethylene neochord constructions). Ring annuloplasty was performed in every case. The DaVinci robotic system represents a truly endoscopic approach to perform mitral valve surgery. The instrument tips are telemanipulated through seven degrees of ergonomic freedom that enable surgeons to perform complex mitral valve repairs through port incisions, providing access to the entire subvalvular chordal apparatus, papillary muscles and all leaflet topography (8). However, mitral valve repair is not always possible, particularly in severely calcified valves of rheumatic origin. In this series, five mitral valves were considered irreparable and replacement was performed. The benefits of robotic mitral valve replacement include smaller thoracotomy, better visualization and exposure of the valve, and easier suture

handling in difficult anatomy (11).

The second most commonly performed procedure in this series was atrioseptoplasty. Patients had a shorter length of hospital stay and no documented complications, which is in accordance with the literature (21). The less common procedures in this series also benefited from robotic assistance. The robotic technique for coronary artery bypass surgery demonstrated reduction in surgical trauma whilst maintaining thoracic integrity. Many studies show reduction of the early postoperative time period with a faster return to full activity (14). Robotic technique for intra-cardiac tumors allowed for complete resection of the mass without sternotomy. Hybrid ablation of atrial fibrillation was facilitated by the robotic equipment, with successful maintenance of sinus rhythm in a period of 26 months of follow-up. Whilst most procedures discussed are suitable for robotic approaches, pericardiectomy is only possible when no significant calcification of the pericardium is present, as robotic instruments are delicate and sectioning thicker tissues may be difficult. In this series, pericardiectomy was easily performed for selected patients as no significant calcification was present in any case.

The major complications documented in this series were related to bleeding and arrhythmias. In three cases, bleeding with need for re-exploration occurred. In these cases, the original surgical access ports were used to perform surgical hemostasis, which was promptly achieved in all cases. In two cases, the source of bleeding was identified as being the intercostal vein. As a result of this finding, careful examination of all port incisions for haemostasis at the final revision is recommended. Atrial fibrillation with rapid ventricular response was documented in two cases; however both were successfully chemically reverted with intravenous amiodarone. Recent studies have documented the rate of post-operative atrial fibrillation to range from 5% to 40% in TECAB, and from 37% to 50% in heart valve surgery (22).

When comparing operative times, robotic cardiac surgery had prolonged times when compared with sternotomy, perhaps reflecting the complexity and learning curve associated with this new technology. Charland and colleagues (23) established that this learning curve can be overcome with adequate planning, gain of surgical volume and experience, with both approaches eventually comparable in duration (24). In this series, the greater CPB and cross-clamp times were not related to increased morbidity or complications. In fact, this series documented shorter length of ICU stay as well as total hospital stay. Furthermore, the majority of patients were extubated in

the operating room. A downward trend in operative times in the last ten cases was observed as the surgeon evolved on the learning curve.

Despite advancements in cardiac surgery over recent decades, very little has been done to improve minimally invasive cardiac surgery (16). Surgical techniques are in constant change and no delay to follow this progress should be tolerated. However, the expansion of robotic cardiac surgery in Brazil and in other developing countries depends on public and private initiatives to create a health system guided by innovation and driven by quality.

There are important limitations to this study, primarily relating to its small, non-randomized, single-center cohort. Further studies involving larger randomized cohorts and long-term follow up should be considered to assess this promising and evolving technology.

In summary, this study supports the role of robotic cardiac surgery as a feasible, safe and effective alternative to sternotomy for the surgical correction of various intra- and extra-cardiac pathologies. Adopting the robotic system has been a challenge in Brazil, where the lack of specific training and high initial costs of technology are the main impediments for a wide adoption. Nevertheless, overcoming these obstacles is essential to ensure that robotic cardiac surgery, as a commonplace in Brazil, may soon become a reality.

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Footnote

Conflicts of Interest: The authors have no conflicts of interest to declare.

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