

# Robotically assisted mitral valve replacement

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**Objective:** In the present study, we determined the safety and efficacy of robotic mitral valve replacement using robotic technology.

**Methods:** From January 2007 through March 2011, more than 400 patients underwent various types of robotic cardiac surgery in our department. Of these, 22 consecutive patients underwent robotically assisted mitral valve replacement. Of the 22 patients with isolated rheumatic mitral valve stenosis (9 men and 13 women), the mean age was  $44.7 \pm 19.8$  years (range, 32–65). Preoperatively, all patients underwent a complete workup, including coronary angiography and transthoracic echocardiography. Of the 22 patients, 15 had concomitant atrial fibrillation. The surgical approach was through 4 right-side chest ports with femoral perfusion. Aortic occlusion was performed with a Chitwood crossclamp, and antegrade cardioplegia was administered directly by way of the anterior chest. Using 3 port incisions in the right side of the chest and a 2.5- to 3.0-cm working port, all the procedures were completed with the da Vinci S robot.

**Results:** All patients underwent successful robotic surgery. Of the 22 patients, 16 received a mechanical valve and 6 a tissue valve. The mean cardiopulmonary bypass time and aortic crossclamp time was  $137.1 \pm 21.9$  minutes (range, 105–168) and  $99.3 \pm 17.9$  minutes (range, 80–133), respectively. No operative deaths, stroke, or other complications occurred, and no incisional conversions were required. After surgery, all the patients were followed up echocardiographically.

**Conclusions:** Robotically assisted mitral valve replacement can be performed safely in patients with isolated mitral valve stenosis, and surgical results are excellent. (*J Thorac Cardiovasc Surg* 2012;143:S64-7)

Minimally invasive cardiac surgery has emerged as an interesting alternative for a wide variety of surgical procedures. Less-invasive valve surgery has proved even more promising than less-invasive coronary bypass procedures because simpler anastomoses are required. The feasibility and safety of robotic mitral valve replacement (MVR) has been proven in several studies,<sup>1-3</sup> with the main goals of improved cosmesis and less postoperative discomfort while maintaining the same level of safety and efficacy as with conventional surgery. However, robotically assisted MVR demands different surgical skills in the implant of the prosthesis. Whenever introducing a new technique, it is necessary to determine the incremental risk to the patient compared with the benefit achieved. Few data have been published concerning the outcomes of robotically assisted MVR.<sup>4</sup>

We report the early results for the first 22 consecutive patients who underwent robotically assisted MVR.

## METHODS

### Patients

From January 2007 through March 2011, more than 400 patients underwent various types of robotic cardiac surgery in our department. Of these patients, 22 consecutive patients underwent robotically assisted MVR. The institutional review board approved the present study, and the 22 patients provided informed consent. Of the 22 patients with isolated rheumatic mitral valve stenosis (9 men and 13 women), the mean age was  $44.7 \pm 19.8$  years (range, 32–65). Preoperatively, all patients underwent a complete workup, including coronary angiography and transthoracic echocardiography. All 22 patients had good left ventricular function. The preoperative characteristics are listed in the Table 1. Two patients were in New York Heart Association class I, 16 in class II, and 4 in class III; 15 patients had concomitant atrial fibrillation (Table 1).

### Surgical Technique

After induction of general anesthesia, all patients were intubated for single-lung ventilation, and a transesophageal echocardiography (TEE) probe and arterial pressure monitoring line were inserted. Both a central catheter and a 15F venous drainage cannula were placed percutaneously into the right internal jugular vein. External defibrillator patches were paced to subtend the maximum cardiac mass. Each patient was positioned with the right side of the chest elevated approximately 30° and with right arm tucked at the side. After systemic heparinization, femoral arterial (18F or 20F) and venous (21F or 23F) cannulation (Metronic, Minneapolis, Minn) were performed through a 2-cm transverse right groin incision cut-down with the Seldinger guidewire method and TEE guidance. Bicaval venous drainage was instituted through the jugular and femoral/inferior vena cava cannulas.

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Disclosures: Changqing Gao, Ming Yang, Cangsong Xiao, Gang Wang, Yang Wu, Jiali Wang, and Jiachun Li have nothing to disclose with regard to commercial support.

Presented at The American Association for Thoracic Surgery Mitral Conclave, New York, New York, May 5-6, 2011.

Received for publication April 27, 2011; revisions received Dec 3, 2011; accepted for publication Jan 16, 2012; available ahead of print Feb 3, 2012.

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0022-5223/\$36.00

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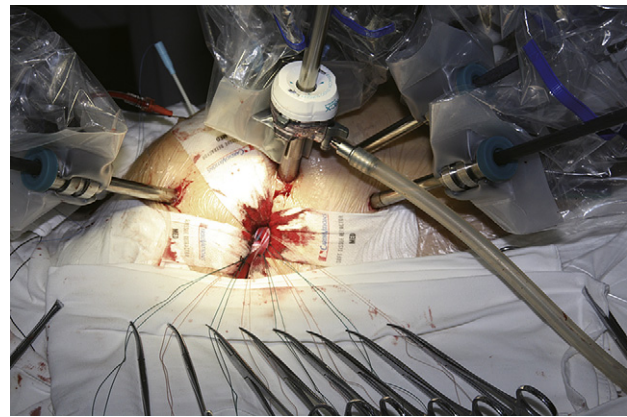
doi:10.1016/j.jtcvs.2012.01.045

### Abbreviations and Acronyms

ICS	= intercostal space
MVR	= mitral valve replacement
TEE	= transesophageal echocardiography

The camera cannula was placed in the right side, 2 to 3 cm lateral to the nipple in the fourth intercostal space (ICS). A 2.5- to 3.0-cm incision was used as a working port in the same ICS for the patient-side surgeon. The soft tissue retractor was inserted into this port, and no other rib retractor was used (Figure 1). The da Vinci instrument arms were inserted through three 0.8-cm trocar incisions in the right side of the chest. The right instrument arm was generally positioned lateral to the working port in the sixth ICS. The left instrument arm was positioned medially and cephalad to the working port in the second ICS. The fourth trocar arm was placed in the midclavicular line in the fifth ICS. All resection and suturing of the mitral valve and atrial closure were completed with the da Vinci S robot.

Cardiopulmonary bypass was initiated with kinetically assisted bicaval venous drainage. The aorta was occluded with a Chitwood crossclamp (Scanlan International, Minneapolis, Minn) using the midaxillary line in the fourth ICS. Antegrade cold blood cardioplegic solution was administered directly through the anterior chest (second ICS) with a 14F angiocatheter, by which de-airing was conducted, and repeat doses could be given when necessary. TEE was used routinely to monitor the position of the angiocatheter and assist with de-airing. Carbon dioxide was insufflated continuously into the operative field for air displacement. The left atrium was then opened parallel to the interatrial septum. All MVRs were performed using standard techniques. The diseased mitral valve was completely excised, and the posterior leaflets were preserved whenever possible. Appropriate sizing was performed, and everting, double-armed, mattress sutures with Teflon pledgets were placed counterclockwise from the 11-o'clock position. Every stitch was fixed sequentially outside with a small hemostat. Usually 10 to 12 sutures were needed (Figure 1). Once the sutures were placed in the prosthesis sewing ring outside the chest, the prosthesis was lowered into the chest and positioned, and the knots were tied using the knot pusher through the incision (Figure 2). The atriotomy was closed with 4-0 polytetrafluoroethylene running suture (W. L. Gore & Associates, Flagstaff, Ariz). Before the Chitwood crossclamp was released, meticulous intracardiac de-airing was conducted through the angiocatheter of cardioplegia, and then the aorta was unclamped. The patient was weaned from cardiopulmonary bypass, and the cardioplegia site was closed with an



**FIGURE 1.** da Vinci setup for mitral valve replacement. Every stitch was fixed sequentially outside with a small hemostat.

extracorporeal knot tied through the working port, and chest tubes were inserted. The prosthesis function was confirmed by TEE.

The data are shown as the mean  $\pm$  standard error of the mean (Figures 1 and 2).

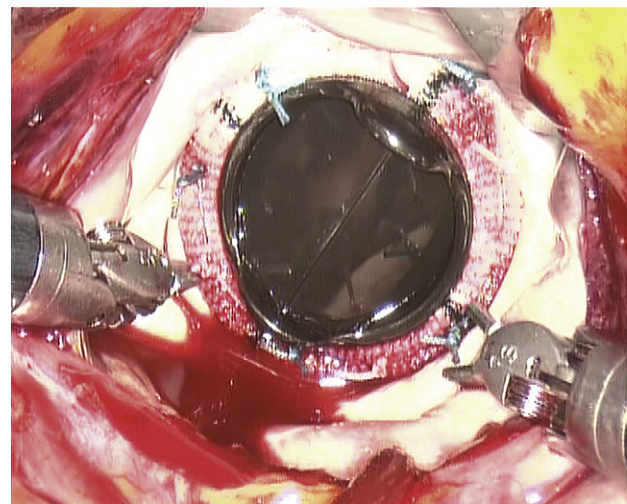
### RESULTS

All patients underwent successful robotically assisted MVR. Of the 22 patients, 16 received a mechanical valve (St Jude Medical, St Paul, Minn) and 6 received a bioprosthetic valve (Cosgrove-Edwards; Edwards Lifesciences, Irvine, Calif), and all patients recovered uneventfully. The mean cardiopulmonary bypass time and aortic crossclamp time was  $137.1 \pm 21.9$  minutes (range, 105–168) and  $99.3 \pm 17.9$  minutes (range, 80–133), respectively. As shown in Figure 2, both the crossclamp and bypass times decreased significantly with experience. Intraoperative TEE revealed appropriate prosthesis function in all patients. One patient had effusion in the right hemithorax. The mean intensive

**TABLE 1. Patient characteristics (n = 22)**

Characteristic	Value
Age (yr)	
Mean	44.7 $\pm$ 9.8
Range	32–65
Gender (n)	
Male	9 (35)
Female	13 (65)
Mean ejection fraction (%)	50.2 $\pm$ 13.8
New York Heart Association class	
I	2 (10)
II	14 (70)
III	4 (20)
Sinus rhythm (n)	5 (25)
Atrial fibrillation (n)	15 (28)

Data in parentheses are percentages.



**FIGURE 2.** The mechanical valve is set into the right position and tied through the working port.

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