# Surgical management of very late paravalvular leaks after mitral valve replacement: a single institutional experience

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**Abstract:** Most mitral paravalvular leaks (PVLs) occur during the first year after mitral valve replacement (MVR). This report describes the surgical management of 6 patients who developed very late mitral PVLs. The median interval between MVR and initial diagnosis of PVL was 16.5 years. All patients presented with congestive cardiac failure and haemolytic anaemia. The median EuroSCORE II was 9.5%. Two patients (33%) had failed attempts at transcatheter closure. Five patients underwent suture repair of the PVL. One patient underwent MVR after removal of the previous prosthesis. No in-hospital mortalities occurred. At latest follow-up (median 3.3 years), 5 patients (83%) were asymptomatic with no residual PVL. Haemolytic anaemia persisted in 1 patient with a mild residual PVL. PVL occurring decades after MVR is a rare but serious complication. Reoperative surgery can be performed in these high-risk patients with satisfactory early and midterm outcomes.

Keywords: Heart valve prosthesis; paravalvular leak (PVL); mitral valve replacement (MVR)

Submitted Jul 03, 2016. Accepted for publication Jul 24, 2016. doi: 10.21037/jtd.2016.08.58 View this article at: http://dx.doi.org/10.21037/jtd.2016.08.58

#### Introduction

The incidence of paravalvular leak (PVL), defined as a regurgitant jet between the prosthetic sewing ring and native tissues on transoesophageal echocardiography (TEE), occurs in 5-32% of patients following mitral valve replacement (MVR) (1,2). Most cases of PVL remain clinically silent, but patients presenting with persistent symptoms of congestive cardiac failure, haemolytic anaemia, or a combination of both may benefit from surgical repair. The majority of PVLs occur during the first postoperative year (2). It is extremely rare for patients to develop PVL decades after MVR (1). Previous reports have cited the longest period between MVR and the diagnosis of a new PVL to be 27 years (3). This retrospective review was conducted at our tertiary center to study the outcomes of 6 patients who underwent reoperative surgery for mitral PVLs occurring more than 10 years after the latest MVR.

### **Case presentation**

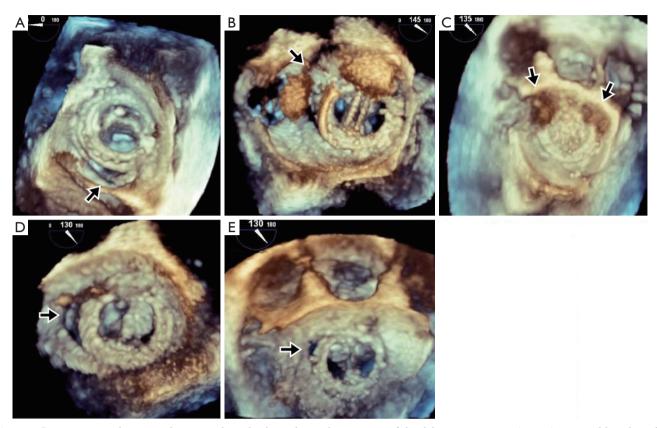
All patients had initially undergone MVR with mechanical prostheses for rheumatic mitral stenosis between 1979 and 1998. Patient characteristics are summarized in *Table 1*. Three patients (50%) had undergone at least 2 previous MVRs. The median time interval between the most recent MVR and initial echocardiographic diagnosis of PVL was 16.5 years (range: 11 to 29 years).

All patients presented with New York Heart Association (NYHA) class III to IV symptoms and haemolytic anaemia (median haemoglobin level 8.3 g/dL, range: 5.7 to 9.0 g/dL, median lactate dehydrogenase level 4,157 unit/L, range 1,484 to 8,268 unit/L). All 6 patients had haptoglobin levels <0.10 g/L. Preoperative transoesophageal echocardiography showed the degree of PVL to be moderate in 1 patient (16.7%), moderate to severe in 2 patients (33.3%) and severe in 3 patients (50.0%). Three-dimensional TEE

Patient	Gender	. Age	Number of previous MVR	Number Interval of since previous last MVR MVR (years)	Number Interval of since Preoperative EuroSCORE Severity of previous last MVR NYHA class II PVL MVR (years)	:uroSCORE II	Severity of PVL	Site of leak	Surgery	Concomitan <sup>.</sup> procedure	Site of leak Surgery Concomitant Complications Follow-up procedure (months)	Follow-up (months)	Latest NYHA class	Residual PVL	Follow-up Latest Residual Persistent (months) Class PVL Haemolysis
-	Male	62	-	13	≥	8.9	Severe	6 to 8 o'clock	Direct closure	AVT	Nil	33	_	None	No
N	Female	49	σ	0	2	42.2	Severe	7 to 11 o'clock Direct closure	Direct closure	Z	(I) Re-exploration for bleeding; (II) Renal failure <sup>§</sup> ; (III) Sepsis; (IV) Respiratory failure <sup>¶</sup>	4	=	Mild	Yes
С	Female	65	2	18	≥	10.0	Severe	10 to 2 o'clock Direct closure	Direct closure	AVR	Nii	45	_	Mild	No
4	Female	58	-	15	≡	4.0	Moderate- severe	8 to 9 o'clock	Direct closure	Nil	Nii	58	_	None	No
Ω	Female	73	-	22	≡	15.4	Moderate	11 o'clock	Direct closure	AVT	Nii	23	_	None	No
9	Female	51	С	11	≡	6.0	Moderate- severe	Moderate- Multiple sites severe	MVR	AVT	Nil	34	_	None	No
<sup>§</sup> , renal York H	failure re eart Asso	equirir sciatio	n; PVL, p	eplacemen aravalvular	$^{\rm s}$ , renal failure requiring renal replacement therapy; $^{\rm t}$ , respiratory failure requiring York Heart Association; PVL, paravalvular leak; TVA, tricuspid valve annuloplasty.	spiratory fail uspid valve	ure requirin; annuloplasty	g tracheostomy y.	. AVR, ao	rtic valve re	, renal failure requiring renal replacement therapy; <sup>¶</sup> , respiratory failure requiring tracheostomy. AVR, aortic valve replacement; MVR, mitral valve replacement; NYHA, New ork Heart Association; PVL, paravalvular leak; TVA, tricuspid valve annuloplasty.	mitral valv	e replac	tement; N	IYHA, New

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**Figure 1** Preoperative 3-dimensional transesophageal echocardiographic imaging of the dehiscent segments (arrows), oriented based on the surgeon's view from the left atrium. (A) Paravalvular leak at the 6 to 8 o'clock position; (B) paravalvular leak at the 7 to 11 o'clock position; (C) paravalvular leak at the 10 to 2 o'clock position; (D) paravalvular leak at the 8 to 9 o'clock position; (E) paravalvular leak at the 11 o'clock position.

images were available for 5 patients (*Figure 1*). The median EuroSCORE II was 9.5% (range: 4.0% to 42.2%). The median left ventricular ejection fraction (LVEF) was 61% (range: 55% to 67%).

Two patients had undergone unsuccessful percutaneous closure of PVL, which failed due to either significant residual regurgitation or the closure device interfering with the existing mitral prosthesis. One patient who received 2 occluder devices required emergency surgery due to cardiogenic shock, resulting from a closure device being caught between the leaflets of the mitral prosthesis. Five patients with discrete PVLs in the presence of a calcified mitral annulus underwent direct suture repair without removal of the existing mitral prosthesis. One patient with multiple PVLs at various sites along the sewing ring underwent MVR after extirpation of the previous prosthetic valve. Concomitant procedures were performed in 4 patients (*Table 1*). The median aortic cross-clamp and cardiopulmonary bypass times were 50 minutes (range,

42 to 119 minutes) and 99 minutes (range, 72 to 196 minutes), respectively.

There were no in-hospital mortalities. Postoperative complications are listed in *Table 1*. No patients suffered from postoperative stroke, myocardial infarction or required insertion of a permanent pacemaker. Upon discharge from hospital, five patients (83.3%) had no residual PVL and 1 patient (16.7%) had mild residual PVL.

All patients were reviewed at our institution during scheduled clinic consultations. The median follow up period was 3.3 years (range: 1.9 to 4.8 years). There were no late mortalities. Five patients (83.3%) were in NYHA class I and had no recurrence of haemolytic anaemia. Four patients (66.7%) had no recurrence of PVL and two patients (33.3%) had mild PVL during follow-up echocardiographic assessment (*Table 1*). In the patient who underwent salvage surgery following failed transcatheter repair, haemolysis persisted despite surgical correction, due to a turbulent jet flowing through a mild residual PVL. The patient opted for conservative management and declined any further high-risk PVL closure.

#### Discussion

Congestive cardiac failure and symptomatic haemolytic anaemia are two major manifestations of PVL. Surgery is indicated for patients experiencing progressive and recurrent heart failure symptoms refractory to optimal medical therapy and/or persistent haemolysis requiring increasingly frequent blood transfusions (3,4). Compared to conservative management, surgical repair has been shown to relieve symptoms, decrease the need for blood transfusion and improve survival (1).

Hwang and associates (2) reported that 74% of PVLs occur during the first postoperative year. It is extremely rare for PVLs to occur more than 20 years after MVR (1,3,5). The longest interval between the latest MVR and diagnosis of a new PVL in our series was 29 years, which exceeds the longest interval (27 years) reported to date (3).

The majority of late PVLs are not associated with infective endocarditis (IE) (2). Accordingly, none of our patients had clinically or microbiologically proven IE, thus suggesting a different cause of their PVLs. Based on the findings of significant mitral annular calcification in most of our patients, we postulate that very late PVL occurred as a result of annular tissue under the sewing cuff undergoing long-term degenerative change and calcification, leading to progressive dehiscence at vulnerable areas along the sewing ring. A less likely mechanism could have been an occult infective process which healed spontaneously, leading to a small area of detachment around the sewing ring.

Reoperative surgery for correction of mitral PVL remains a high-risk procedure. Operative mortality ranges from 6% to 13 % (1-4). The median EuroSCORE II of our cohort (9.5%) falls within the range (7.5% to 9.6%) described by other contemporary series (3,4). Residual mitral PVL has been reported in 2% to 22% of patients following surgical correction (1,3,4). Recurrence of PVL has been cited to occur at a rate of 4% per patient-year (3). The number of previous MVRs has been shown to be associated with the likelihood of recurrent PVL after surgical repair (1-3). This association was evident in our cohort whereby both patients with residual mild PVLs had undergone at least 2 previous MVRs. The choice of direct suture repair versus repeat MVR was not a risk factor for recurrence of PVL (2,3).

Preoperative 3-dimensional TEE has emerged as

a valuable adjunct in delineating the anatomy and morphology of PVL. The intraoperative findings observed in our patients were consistent with preoperative TEE images (*Figure 1*). Intraoperative TEE is mandatory in these cases to evaluate the presence and degree of residual PVL before the conclusion of surgery. Contrary to previous reports describing PVLs occurring predominantly along the posterior annulus or posteromedial commissure, the PVLs in our series were distributed at various sites along the entire circumference of the sewing ring (2,3).

The majority of our patients had a single leak which was amenable to direct suture closure. To reduce the likelihood of residual PVL, we deliberately avoided implanting a new mitral prosthesis in patients with a heavily calcified mitral annulus. Extirpation of the mitral prosthesis and repeat MVR was required in one patient due to the presence of PVLs at multiple sites of dehiscence along the sewing ring. Fortunately, this patient had a minimally calcified mitral annulus.

Percutaneous transcatheter closure of PVLs has emerged as a promising, less-invasive alternative to surgical treatment. Although long-term durability has yet to be established, a recent meta-analysis has shown successful transcatheter PVL correction to be associated with reduced all-cause mortality and improved functional class in patients deemed unsuitable for surgical correction (6). The success of percutaneous closure of PVL is largely determined by anatomical and devicerelated factors. Contraindications to percutaneous closure of PVL include large defects involving more than one third of the prosthetic annulus, an unstable or rocking prosthesis, active endocarditis, and intracardiac thrombi (7). Deployment of occluder devices may result in either interference with the prosthetic leaflets or residual regurgitation. There is an increased risk of interference with mechanical disc mobility if more than once device is deployed. As demonstrated in our series, salvage surgery performed emergently after failed percutaneous PVL closure may be associated with increased risk and morbidity.

#### Conclusions

Very-late PVLs occurring more than 10 years after MVR are rare but serious complications which warrant early attention. Clinicians must be cognizant of this problem in patients who have undergone valve replacement many years ago and have a high index of suspicion in the presence of a new murmur or haemolytic anaemia. Reoperative surgery can be performed in these high risk patients with

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satisfactory early and midterm outcomes.

## Acknowledgements

The authors thank Clara Zhang and Selena Chew for their editorial assistance.

## Footnote

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

*Informed Consent:* The local institutional review board (SingHealth CIRB 2016/2696), along with a waiver of informed consent.

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**Cite this article as:** Pang PY, Zhu J, Sin YK, Chua YL. Surgical management of very late paravalvular leaks after mitral valve replacement: a single institutional experience. J Thorac Dis 2016;8(9):E952-E956. doi: 10.21037/jtd.2016.08.58

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