#### Check for updates

#### **OPEN ACCESS**

EDITED BY Enyi Shi, China Medical University, China

REVIEWED BY Antonino S. Rubino, University of Campania Luigi Vanvitelli, Italy Alberto Albertini, Maria Cecilia Hospital, Italy

\*CORRESPONDENCE Yefan Jiang ☑ jyf5930829@163.com Si Chen ☑ sichen@hust.edu.cn Wenjing Zhang

🖂 wenjing.zhang.0120@163.com

<sup>†</sup>These authors have contributed equally to this work

#### SPECIALTY SECTION

This article was submitted to Heart Surgery, a section of the journal Frontiers in Surgery

RECEIVED 19 September 2022 ACCEPTED 01 December 2022 PUBLISHED 06 January 2023

#### CITATION

Wang S, Zhou T, Bian J, Li G, Zhang W, Chen S and Jiang Y (2023) Clinical outcomes following surgical mitral valve plasty or replacement in patients with infectious endocarditis: A metaanalysis.

Front. Surg. 9:1048036.

doi: 10.3389/fsurg.2022.1048036

#### COPYRIGHT

© 2023 Wang, Zhou, Bian, Li, Zhang and Jiang. This is an open-access article distributed under the terms of the Creative Commons Attribution License (CC BY). The use, distribution or reproduction in other forums is permitted, provided the original author(s) and the copyright owner(s) are credited and that the original publication in this journal is cited, in accordance with accepted academic practice. No use, distribution or reproduction is permitted which does not comply with these terms.

## Clinical outcomes following surgical mitral valve plasty or replacement in patients with infectious endocarditis: A meta-analysis

# Song Wang<sup>1,2†</sup>, Ting Zhou<sup>3†</sup>, Jinhui Bian<sup>2†</sup>, Geng Li<sup>4</sup>, Wenjing Zhang<sup>5,6\*</sup>, Si Chen<sup>4\*</sup> and Yefan Jiang<sup>2,4\*</sup>

<sup>1</sup>Department of Cardiovascular Surgery, The Affiliated Taizhou People's Hospital of Nanjing Medical University, Taizhou School of Clinical Medicine, Nanjing Medical University, Nanjing, China, <sup>2</sup>Department of Cardiovascular Surgery, The First Affiliated Hospital of Nanjing Medical University, Nanjing, China, <sup>3</sup>Health Management Center, The Central Hospital of Wuhan, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, China, <sup>4</sup>Department of Cardiovascular Surgery and Heart Transplantation, Union Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, China, <sup>5</sup>Department of Ultrasound Medicine, The Second Afliated Hospital of Harbin Medical University, Harbin, China, <sup>6</sup>Department of Ultrasound Medicine, Union Hospital, Tongji Medical College, Huazhong University of Science and Technology, Wuhan, China

**Background:** For degenerative mitral disease, more and more evidences support that mitral valve plasty (MVP) has much better clincial outcomes than mitral valve replacement (MVR). However, the advantages of MVP in patients suffering from infectious endocarditis (IE) are unclear. To evaluate the appropriateness of MVP in IE patients, we conducted this meta-analysis. Based on the difference between active and healed phase, we not only compared the result of patients with IE, but also identified the subgroup with active IE.

**Methods:** We systematically searched the clinical trials comparing clinical outcomes of MVP and MVR in patients suffering from IE. Relevant articles were searched from January 1, 2000 to March 18, 2021 in Pubmed and Cochrane Library. Studies were excluded if they were with Newcastle–Ottawa Scale (NOS) score less than 6 or lacking of direct comparisons between MVP and MVR.

**Results:** 23 studies were involved and 25,615 patients were included. Pooled analysis showed fewer adverse events and early or long-term death in the MVP group. However, more reoperations existed in this patient group. And the reinfection rate was close between two groups. Similar results were observed after identifying active IE subgroup, but there is no difference in the freedom from reoperation due to all-events.

**Conclusions:** Although limitimations exited in this study, patients suffering from IE can benefit from both MVP and MVR. For surgeons with consummate skills, MVP can be the preferred choice for suitable IE patients.

#### KEYWORDS

infectious endocarditis, mitral valve plasty, mitral valve replacement, clinical outcomes, meta-analysis

## Introduction

Treatment for infective endocarditis consists of antibiotic therapy and surgery (1). Surgical intervention is needed in about half of patients suffering from IE (2). The indications for surgery in IE are well defined but the choice of surgical procedures is less defined, especially with regard to MVP vs. MVR (3). For non-infected mitral valvular diseases, such as myxomatous, ischemic or degenerative valve disease, MVP is prefered due to fewer reoperation, thromboembolism, and valve infection events (4). While for patients with IE, although several studies have confirmed a better survival after MVP compared with MVR in patients with IE, the small sample size and lacking of randomized controlled trials bring the bias and reduce the confidence level. Besides, so far only short and long-term survival were mentioned in published systematic review and the core issues of reinfection and reoperation were ignored (1, 5, 6). Therefore, in this paper, we performed this meta-analysis to evaluate whether MVP has better clinical outcomes than MVR for patients suffering from IE. Not only the survival but also core complications such as reinfection and reoperation were analyzed. To get more detailed information, the subgroup of patients with active IE was also analyzed.

#### **Methods**

#### Search strategy

Pubmed and Cochrane Library were searched for journals. "MVP", "mitral valve plasty", "mitral valve repair", "mitral valve annuloplasty", "mitral reconstruction", "MVR", "mitral valve replacement", "infective endocarditis", and "IE" were used either alone or in combination. The reference list of relevant articles and reviews were identified manually to find additional studies.

#### Eligibility criteria

The inclusion criteria were as follows: (i) direct comparison of MVP vs. MVR; (ii) clinical outcomes (early survival, long-term survival, event-free survival, freedom from reoperation due to all-events, reinfection events) had to be provided in sufficient details to allow the extraction of hazard ratios(HR) or odd ratios(OR), and their standard errors or Kaplan–Meier curves. Two independent authors (Song Wang and Ting Zhou) extracted data from studies. Disagreements were resolved by a discussion with a superior (Dr. Yefan Jiang). Studies met the inclusion criteria were rated based on the NOS (7). Studies with a NOS score of 5 or lower were excluded. Paper quality was checked by Song Wang and Ting Zhou independently.

#### Statistical analysis

Summary HR for event-free survival, freedom from reoperation due to all-events or reinfection, long-term survival, and OR for reinfection events and early mortality were obtained as weighted averages of measures from the induvidual studies, with inverse variances used as weights. We used methods provided by Parmer, Williamson and Tierney (8–10) to calculate the estimated HR and variance. Besides, we used a *Q*-statistic and  $I^2$  (index of inconsistency) test to quantify the heterogeneity degree and when significant heterogeneity (P < 0.1 or  $I^2 > 50\%$ ) existed, a random effects model was applied. We omitted each study included in sequence to conduct the sensitivity analyses and visual inspection of funnel plots was used to assess the publication bias. RevMan 5.3 was used to analyze the data.

#### Results

#### Study search

The selection strategy in shown in **Figure 1**. 23 retrospective studies met the inclusion criteria. 25,615 patients were included in the final analysis, 10,719 of whom received MVP and 14,896 received MVR from 1980 to 2017. **Table 1** presents the individual studies' characteristics.

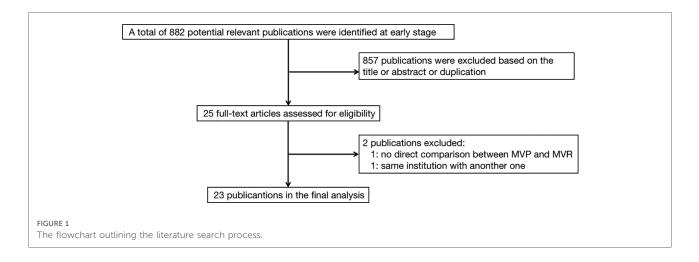
#### Early mortality

Early death refers to in-hosptial death or death occurring within 30 days after operation. All 23 studies contained related details on it and a random effects model was used because of the relatively high heterogeneity ( $I^2 = 57\%$ , P = 0.0004) among studies included. Lower early mortality was apparent in MVP group. Among those 23 studies, 16 studies (12, 14–16, 18–20, 22–24, 26, 27, 29, 31–33) reported patients with active IE. The advantages on early mortality of MVP over MVR was also obvious, when only patients with active IE were included. [MVP vs. MVR: OR: 0.45, 95% CI: 0.33–0.61, Figure 2A; MVP vs. MVR (active IE): OR: 0.53, 95% CI: 0.42–0.65, Figure 2B].

Among those 23 studies, 3 studies contain 90% percentage of patients (21, 23, 32), the exclusion of any one or two or all of those three studies didn't change the overwhelming of MVP. (As shown in **Supplementary Figure S1**).

#### Long-term survival

15 studies (12–14, 16, 19, 22, 24–29, 31, 32) provided related data. Although there was significant heterogeneity( $I^2 = 47\%$ , P = 0.02) among those studies, heterogeneity can be accepted



after omiting the study of Eric Shang et al. (17), and deletion of that study did not change the overall results. 11 studies (12, 14, 16, 19, 22, 24, 26, 27, 29, 31, 32) documented details of patients with active IE. The results presented that patients receiving MVP were with a decreased long-term risk of death irrespective of whether only patients with active IE were included. [MVP vs. MVR: HR: 0.61, 95% CI: 0.49–0.75, **Figure 3A**; MVP vs. MVR (active IE): HR: 0.65, 95% CI: 0.55–0.76, **Figure 3B**].

#### **Event-free survival**

Here event-free means freedom from cerebravascular disease, valvular reoperation, recurrence of endocarditis, and death (33). Of the 23 included studies, 6 (12, 18, 20, 31-33) provided information to allow the determination of event-free survival but significant heterogeneity existed ( $I^2 = 48\%$ , P =0.09), much smaller heterogeneity could be calculated after removing the study of Michele Musci et al. (12), and the deletion of that study did not change the overall results. All of the 6 studies also provided details related to patients with active IE. Same as the analysis mentioned above, significant heterogeneity existed and the deletion of the study by Michele Musci et al. (12), could reduce the heterogeneity and didn't change the overall results. The analyses demonstrated higher event-free survival in the MVP group for both the whole patients and active IE subgroup. [MVP vs. MVR: HR: 0.72, 95% CI: 0.60-0.86, Figure 4A; MVP vs. MVR(active IE): HR: 0.73, 95% CI: 0.61–0.88, Figure 4B].

#### Freedom from reoperation due to allevents

9 studies (12, 16, 19, 22, 24, 26, 28, 29, 32) presented details on the analysis of freedom from reoperation due to all-events, 8 (12, 16, 19, 22, 24, 26, 29, 32) of which documented information related to active IE. While the summary HR suggested that reoperation rate due to all events was lower in patients with IE following MVR, no significant differences existed in patients with active IE after MVR and MVP. [MVP vs. MVR: HR: 1.46, 95% CI: 1.01–2.10, Figure 5A; MVP vs. MVR (active IE): HR: 1.48, 95% CI: 0.97–2.25, Figure 5B].

#### **Reinfection events**

Information on reinfection events was obtained from 14 studies (12, 14, 16, 17, 19, 20, 22, 25–28, 30, 31, 33). 10 (12–14, 16, 19, 20, 26, 27, 31, 33) of which provided details related to patients with active IE. No differences in reinfection events between those two groups were observed no matter whether only active IE patients were included. [MVP vs. MVR: OR: 1.10, 95% CI: 0.67–1.81, Figure 6A; MVP vs. MVR (active IE): OR: 0.96, 95% CI: 0.52–1.78, Figure 6B].

And there is still no difference between in reinfection between MVP group and MVR group when we take time into account and calculate the HR value of reinfection. 4 studies (26, 28, 29, 32) provided related details and 3 (26, 29, 32) of which provided details related to patients with active IE. [MVP vs. MVR: HR: 1.91, 95% CI: 0.53–6.87, **Supplementary Figure S2A**; MVP vs. MVR (active IE): HR: 1.39, 95% CI: 0.18–10.63, **Supplementary Figure S2B**].

#### Discussion

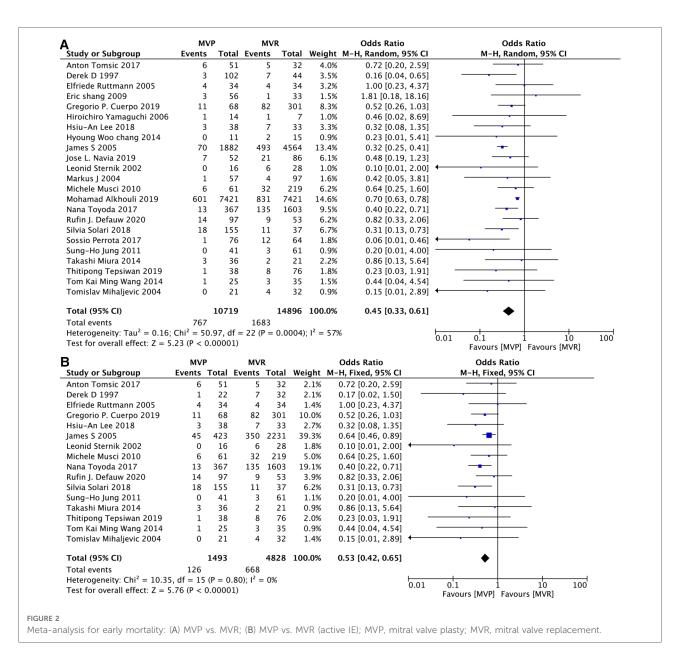
The incidence of IE is gradually increasing (34). IE patients with symptoms of severe valve dysfunction, heart failure, recurrent systemicembolization, et al. should receive surgical treatmen (35). Although the diagnosis, medical treatment and surgical techniques have improved, mortality and morbidity still remain high (2).

Study	Country	Study period		Surgery	/		Active/ Healed		n age ars)	М	ale
			M۷	′P	M١	/R		MVP	MVR	MVP	MVR
Hyoung Woo chang 2014 (11)	South Korea	2004-2011	11		1	5	Both	38.2	48.3	9/2	11/4
Michele Musci 2010 (12)	Germany	1996–2007	61		16	56	Active	47.7	56.2	40/21	134/85
Markus J 2004 (13)	Switerland	1980-1996	57	7	9	7	Both	-	-	-	-
Tomislav Mihaljevic 2004 (14)	United States	1992-2002	21		3	2	Active	66	50	16	24
Leonid Sternik 2002 (15)	United States	1986-1999	16	5	2	8	Active	-	-	-	-
Thitipong Tepsiwan 2019 (16)	Tailand	2006-2017	38	3	7	6	Active	44.1	47.5	21	57
Eric shang 2009 (17)	United States	2002-2007	56	5	3	3	Both	4	·8	5	59
Derek D 1997 (18)	United States	1985–1995	Both	102		44	Both	51	1.9	1	00
			Active	26		32					
			Healed	72		12					
Takashi Miura 2014 (19)	Japan	1999–2012	36	5	2	1	Active	57	55	35	27
Hsiu-An Lee 2018 (20)	Taiwan China	2005-2015	38	3	3	3	Active	42.3	53.7	24	20
Mohamad Alkhouli 2019 (21)	United States	2003-2016	745	51	272	204	Both	53	55	5000	16,513
Anton Tomsic 2017 (22)	Netherlands	2000-2016	51		3	2	Active	55	60	38	17
James S 2005 (23)	United States	1994–2003	Both	1882	2 4	1565	Both	5	6	39	956
			Active	423	2	2231		5	4	15	547
			Treated	1459	9 2	2111		5	8	21	84
Silvia Solari 2018 (24)	Belgium	1991–2015	15	5	3	7	Active	60.1	64.6	109	18
Sossio Perrota 2017 (25)	Sweden	2000-2015	76	5	6	4	Both	60	62	55	45
Rufin J. Defauw 2020 (26)	Netherlands	2000-2017	97	7	5	3	Active	57	61	20	22
Gregorio P. Cuerpo 2019 (27)	Spain	2008-2016	68	3	30	)1	Active	-	-	-	-
Jose L. Navia 2019 (28)	United States	1988-2017	52	2	8	6	Both	55	58	44	52
Tom Kai Ming Wang 2014 (29)	New Zealand	2005-2011	25	5	3	5	Active	43.1	52.1	14	21
Hiroichiro Yamaguchi 2006 (30)	Japan	1999–2005	14	ļ	7	7	Both	58	53		
Sung-Ho Jung 2011 (31)	South korea	1994–2009	41		6	1	Active	34.4	43.1	19	33
Nana Toyoda 2017 (32)	United States	1998–2014	36	7	16	03	Active	56.9	54.9	243	900
Elfriede Ruttmann 2005 (33)	Austria	1992-2004	34	ł	3	4	Active	51.5	53.2	22	17

#### TABLE 1 The characteristics of the individual studies.

MVP, mitral valve plasty; MVR, mitral valve replacement.

Repair and replacement are both candidate therapeutic methods for valve diseases. More and more surgeons prefer MVP as the first choice for patients with myxomatous, ischemic or degenerative mitral valve diseases, due to MVP's advantages of lower early morbidity and operative mortality, higher long-term survival, and fewer reoperation events (4). However, the superiority of MVP in patients with IE still needs identification furtherly. So far related papers are still



limited. In published papers of this topic, several limitations do exist. Firstly, there were only retrospective studies. Secondly, only small number of cases were included in some single-center studies (36). Thirdly, in multi-center studies, patients were included through clinical codes and selection bias was large. Besides, the gap of medical level between different hospitals reduced the reliability of the final results and the rate of successful MVP in active IE varied significantly among previous literatures from 15% up to 100% (16).

MVP is much more complicated in the presence of IE accompanied with the valves' primary pathology like degeneration, calcification, et al. (36). Concomitant cardiac abnormalities such as aortic valve IE, coronary artery diseases,

et al. also contribute (36, 37). For patients with active IE, the reconstructive surgery in inflammatory tissue may be difficult and recurrence of infection is relatively common (5). For patients with healed IE, MVP may not be feasible because of the valvular destruction and heart failure resulting from the infectious process (1). Therefore, two concerns arise on the surgical strategy choice of patients with IE: reinfection for patients with active IE and reoperation for patients with healed IE (26). Previous systematic reviews focused less on those two concerns and had no distinguishing between clinical outcomes of patients with active IE and healed IE (1, 5). Only two published papers provided details related to MVP and MVR in healed IE subgroups. Therefore we can only infer the relevant conclusions from comparison of

Study or Subgroup	log[Hazard Ratio]	SE	Weight	IV, Fixed, 95% CI	IV, Fixed, 95% CI
Anton Tomsic 2017	-2.52			0.08 [0.01, 0.50]	
Gregorio P. Cuerpo 2019	-0.7			0.50 [0.28, 0.89]	
Jose L. Navia 2019	-0.03			0.97 [0.61, 1.55]	
Markus J 2004	-0.36			0.70 [0.37, 1.33]	
Michele Musci 2010	-0.44	0.23	10.7%	0.64 [0.41, 1.01]	
Nana Toyoda 2017	-0.34	0.11		0.71 [0.57, 0.88]	<b>=</b>
Rufin J. Defauw 2020	-0.55	0.29	6.8%	0.58 [0.33, 1.02]	
Silvia Solari 2018	-0.68	0.4	3.6%	0.51 [0.23, 1.11]	
Sossio Perrota 2017	-1.28	0.34	4.9%	0.28 [0.14, 0.54]	
Sung-Ho Jung 2011	-0.67	0.68	1.2%	0.51 [0.13, 1.94]	
Takashi Miura 2014	0.23	0.85	0.8%	1.26 [0.24, 6.66]	
Thitipong Tepsiwan 2019	-0.81	0.62	1.5%	0.44 [0.13, 1.50]	
Tom Kai Ming Wang 2014	-0.48	0.83	0.8%	0.62 [0.12, 3.15]	
Tomislav Mihaljevic 2004	0.38	0.95	0.6%	1.46 [0.23, 9.41]	
Total (95% CI)			100.0%	0.65 [0.56, 0.75]	•
10(a) (95% C)					
	0. $df = 13 (P = 0.16)$ :	$I^2 = 2$	8%		
Heterogeneity: Chi <sup>2</sup> = 18.0		I <sup>2</sup> = 2	8%		
Heterogeneity: Chi <sup>2</sup> = 18.0		I <sup>2</sup> = 2	8%		0.01 0.1 1 10 100 Favours (MVP) Favours (MVR)
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 1		I <sup>2</sup> = 2	8%	Hazard Ratio	
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 5 Study or Subgroup				Hazard Ratio IV, Fixed, 95% CI	Favours [MVP] Favours [MVR]
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 5	5.75 (P < 0.00001)	SE	Weight		Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = Study or Subgroup	5.75 (P < 0.00001) log[Hazard Ratio]	<b>SE</b> 0.93	Weight 0.8%	IV, Fixed, 95% CI	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 5 Study or Subgroup Anton Tomsic 2017	5.75 (P < 0.00001) log[Hazard Ratio] -2.52	<b>SE</b> 0.93 0.3	Weight 0.8% 7.9%	IV, Fixed, 95% CI 0.08 [0.01, 0.50]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 5 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7	<b>SE</b> 0.93 0.3 0.23	Weight 0.8% 7.9% 13.4%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 5 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010 Nana Toyoda 2017	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44	SE 0.93 0.3 0.23 0.11	Weight 0.8% 7.9% 13.4% 58.8%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 1 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010 Nana Toyoda 2017 Rufin J. Defauw 2020	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44 -0.34	SE 0.93 0.23 0.11 0.29	Weight 0.8% 7.9% 13.4% 58.8% 8.5%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01] 0.71 [0.57, 0.88]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 1 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010 Nana Toyoda 2017 Rufin J. Defauw 2020 Silvia Solari 2018	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44 -0.34 -0.55	SE 0.93 0.23 0.11 0.29 0.4	Weight 0.8% 7.9% 13.4% 58.8% 8.5% 4.4%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01] 0.71 [0.57, 0.88] 0.58 [0.33, 1.02]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 1 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010 Michele Musci 2010 Nana Toyoda 2017 Rufin J. Defauw 2020 Silvia Solari 2018 Sung-Ho Jung 2011	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44 -0.34 -0.55 -0.68	SE 0.93 0.23 0.11 0.29 0.4 0.68	Weight 0.8% 7.9% 13.4% 58.8% 8.5% 4.4% 1.5%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01] 0.71 [0.57, 0.88] 0.58 [0.33, 1.02] 0.51 [0.23, 1.11]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 5 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44 -0.34 -0.55 -0.68 -0.67	SE 0.93 0.23 0.11 0.29 0.4 0.68 0.85	Weight 0.8% 7.9% 13.4% 58.8% 8.5% 4.4% 1.5% 1.0%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01] 0.71 [0.57, 0.88] 0.58 [0.33, 1.02] 0.51 [0.23, 1.11] 0.51 [0.13, 1.94]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 1 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010 Nana Toyoda 2017 Rufin J. Defauw 2020 Silvia Solari 2018 Sung-Ho Jung 2011 Takashi Miura 2014	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44 -0.34 -0.55 -0.68 -0.67 0.23	SE 0.93 0.23 0.11 0.29 0.4 0.68 0.85 0.62	Weight 0.8% 7.9% 13.4% 58.8% 8.5% 4.4% 1.5% 1.0% 1.8%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01] 0.71 [0.57, 0.88] 0.58 [0.33, 1.02] 0.51 [0.23, 1.11] 0.51 [0.13, 1.94] 1.26 [0.24, 6.66]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 1 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010 Nana Toyoda 2017 Rufin J. Defauw 2020 Silvia Solari 2018 Sung-Ho Jung 2011 Takashi Miura 2014 Thitipong Tepsiwan 2019 Tom Kai Ming Wang 2014	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44 -0.34 -0.55 -0.68 -0.67 0.23 -0.81	SE 0.93 0.23 0.11 0.29 0.4 0.68 0.85 0.62 0.83	Weight 0.8% 7.9% 13.4% 58.8% 8.5% 4.4% 1.5% 1.0% 1.8% 1.0%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01] 0.71 [0.57, 0.88] 0.58 [0.33, 1.02] 0.51 [0.23, 1.11] 0.51 [0.13, 1.94] 1.26 [0.24, 6.66] 0.44 [0.13, 1.50]	Favours (MVP) Favours (MVR) Hazard Ratio
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 1 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010 Nana Toyoda 2017 Rufin J. Defauw 2020 Silvia Solari 2018 Sung-Ho Jung 2011 Takashi Miura 2014 Thitipong Tepsiwan 2019	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44 -0.34 -0.55 -0.68 -0.67 0.23 -0.81 -0.81 -0.48	SE 0.93 0.23 0.11 0.29 0.4 0.68 0.85 0.62 0.83	Weight 0.8% 7.9% 13.4% 58.8% 8.5% 4.4% 1.5% 1.0% 1.8% 1.0% 0.8%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01] 0.71 [0.57, 0.88] 0.58 [0.33, 1.02] 0.51 [0.23, 1.11] 0.51 [0.13, 1.94] 1.26 [0.24, 6.66] 0.44 [0.13, 1.50] 0.62 [0.12, 3.15] 1.46 [0.23, 9.41]	Favours (MVP) Favours (MVR) Hazard Ratio IV, Fixed, 95% Cl
Heterogeneity: Chi <sup>2</sup> = 18.0 Test for overall effect: Z = 1 Study or Subgroup Anton Tomsic 2017 Gregorio P. Cuerpo 2019 Michele Musci 2010 Nana Toyoda 2017 Rufin J. Defauw 2020 Silvia Solari 2018 Sung-Ho Jung 2011 Takashi Miura 2014 Thitipong Tepsiwan 2019 Tom Kai Ming Wang 2014 Tomislav Mihaljevic 2004	5.75 (P < 0.00001) log[Hazard Ratio] -2.52 -0.7 -0.44 -0.34 -0.55 -0.68 -0.67 0.23 -0.81 -0.48 0.38	SE 0.93 0.23 0.11 0.29 0.4 0.68 0.85 0.62 0.83 0.95	Weight 0.8% 7.9% 13.4% 58.8% 8.5% 4.4% 1.5% 1.0% 0.8% 100.0%	IV, Fixed, 95% CI 0.08 [0.01, 0.50] 0.50 [0.28, 0.89] 0.64 [0.41, 1.01] 0.71 [0.57, 0.88] 0.58 [0.33, 1.02] 0.51 [0.23, 1.11] 0.51 [0.13, 1.94] 1.26 [0.24, 6.66] 0.44 [0.13, 1.50] 0.62 [0.12, 3.15]	Favours (MVP) Favours (MVR) Hazard Ratio IV, Fixed, 95% Cl

FIGURE 3 Meta-analysis for long-term survival: (A) MVP vs. MVR; (B) MVP vs. MVR (active IE); MVP, mitral valve plasty; MVR, mitral valve replacement.

A .				Hazard Ratio	Hazard Ratio
Study or Subgroup	log[Hazard Ratio]	SE	Weight	IV, Fixed, 95% CI	I IV, Fixed, 95% CI
Derek D 1997	-0.82	0.38	6.0%	0.44 [0.21, 0.93]	]
Elfriede Ruttmann 2005	-0.41	0.55	2.9%	0.66 [0.23, 1.95]	]
Hsiu-An Lee 2018	-0.32	0.71	1.7%	0.73 [0.18, 2.92]	]
Nana Toyoda 2017	-0.3	0.1	86.4%	0.74 [0.61, 0.90]	]
Sung-Ho Jung 2011	-0.14	0.53	3.1%	0.87 [0.31, 2.46]	]
Total (95% CI)			100.0%	0.72 [0.60, 0.86]	」 ◆
Heterogeneity: $Chi^2 = 1.9$	90, df = 4 (P = 0.75);	$I^2 = 0$	%		
Test for overall effect: Z =	= 3.55 (P = 0.0004)				0.01 0.1 1 10 10 Favours [mvp] Favours [mvr]
В				Hazard Ratio	Hazard Ratio
Study or Subgroup	log[Hazard Ratio]	SE	Weight	IV, Fixed, 95% CI	I IV, Fixed, 95% CI
Derek D 1997	-0.67	0.56	2.8%	0.51 [0.17, 1.53]	]
Elfriede Ruttmann 2005	-0.41	0.55	3.0%	0.66 [0.23, 1.95]	]
Hsiu-An Lee 2018	-0.32	0.71	1.8%	0.73 [0.18, 2.92]	]
Nana Toyoda 2017	-0.3	0.1	89.3%	0.74 [0.61, 0.90]	]
Sung-Ho Jung 2011	-0.14	0.53	3.2%	0.87 [0.31, 2.46]	]
Total (95% CI)			100.0%	0.73 [0.61, 0.88]	」 ◆
Heterogeneity: $Chi^2 = 0.5$	56, df = 4 (P = $0.97$ );	$I^2 = 0$	%		
Test for overall effect: Z =	, , ,,				0.01 0.1 1 10 10 Favours [mvp] Favours [mvr]

Study or Subgroup	log[Hazard Ratio]		Weight	IV, Fixed, 95% CI		IV, Fixed, 95% CI
Anton Tomsic 2017	-0.78		1.3%	0.46 [0.02, 11.41]		
Jose L. Navia 2019	0.33	0.38	23.9%	1.39 [0.66, 2.93]		
Michele Musci 2010		0.32	33.8%	2.01 [1.08, 3.77]		
Nana Toyoda 2017		0.45	17.1%	1.95 [0.81, 4.72]		
Rufin J. Defauw 2020	-0.44		10.3%	0.64 [0.21, 2.01]		
Silvia Solari 2018	-0.36	0.75	6.1%	0.70 [0.16, 3.03]		
Takashi Miura 2014		1.31	2.0%	1.23 [0.09, 16.08]		
Thitipong Tepsiwan 2019		3.26		20.91 [0.04, 12450.08]		
Tom Kai Ming Wang 2014	0.2	0.82	5.1%	1.22 [0.24, 6.09]		
Total (95% CI)			100.0%	1.46 [1.01, 2.10]		◆
Heterogeneity: $Chi^2 = 5.64$ ,	$df = 8 (P = 0.69); I^2$	= 0%				0,1 1 10 100
Test for overall effect: $Z = 2$	2.03 (P = 0.04)				0.01	0.1 İ 10 100 Favours (MVP) Favours (MVR)
3				Hazard Ratio		Hazard Ratio
Study or Subgroup	log[Hazard Ratio]	SE	Weight	IV, Fixed, 95% CI		IV, Fixed, 95% CI
Anton Tomsic 2017	-0.78	1.64	1.7%	0.46 [0.02, 11.41]	-	
Michele Musci 2010	0.7	0.32	44.4%	2.01 [1.08, 3.77]		
Nana Toyoda 2017	0.67	0.45	22.5%	1.95 [0.81, 4.72]		
Rufin J. Defauw 2020	-0.44	0.58	13.5%	0.64 [0.21, 2.01]		
Silvia Solari 2018	-0.36	0.75	8.1%	0.70 [0.16, 3.03]		
Takashi Miura 2014	0.21	1.31	2.6%	1.23 [0.09, 16.08]		
Thitipong Tepsiwan 2019	3.04	3.26	0.4%	20.91 [0.04, 12450.08]		
Tom Kai Ming Wang 2014	0.2	0.82	6.8%	1.22 [0.24, 6.09]		
Total (95% CI)			100.0%	1.48 [0.97, 2.25]		•
Heterogeneity: $Chi^2 = 5.62$ ,	$df = 7 (P = 0.59); I^2$	= 0%				
Test for overall effect: $Z = 1$					0.01	0.1 İ 10 100 Favours (MVP) Favours (MVR)

clinical outcomes between all patients and subgroup of active IE patients following different mitral strategies.

Analysis showed that early survival, late survival and eventfree survival were higher in the MVP group no matter whether only patients with active IE were included. It does make sense. Firstly, MVP can provide preservation of the subvalvular apparatus and protection of the left ventricular function, which can avoid death resulting from impaired left ventricular function. Secondly, MVP usually accompanies with much more physiologic fitting hemodynamics which can make contributions to the recovery of left ventricular function. Thirdly, the possiblity of valve-related events and perivalvular leakage was reduced by the existence of autogenous valves. Fourthly, it helps patients avoding thromboembolism and hemorrhage with no need for long-term use of anticoagulant therapy (4). Besides, surgeons' preference for MVP in patients with superior body condition and less damaged valves also contribute to fewer complications and death, which can be reflected by the fact that patients receiving MVP are normally younger.

No difference exists in reinfection between MVP group and MVR group no matter whether patients with active IE were regarded as a subgroup. Traditionally, completion of a full course of antibiotics before surgery was recommended (38). While for patients with refractory congestive heart failure, uncontrolled sepsis, et al., mindless delay of surgery should be avoided (39). Moreover, recent literatures have suggested that

early intervention may bring benefits like fewer cardiovascular events and less damaged valvular structure (17, 40). Actually more and more patients are receiving surgeries during active phase. MVR has been the standard procedure for patients with acute IE due to the demand of complete excision of infective tissue (18). However, we can't neglect the fact that the grafts themselves are susceptible to becoming the source of infection (6). There is still debate on the implantation of artificial material in active IE (13, 33). Two adverse events may occur. One is bacterial colonization prior to endothelization of prosthetic materials, and the other is late prosthetic valve IE (22). Some scholars even suggested that, when possible, all artifical materials should be avoided to reduce the recurrence of IE (41, 42). However, there is still no difference between in reinfection between MVP group and MVR group when we take time into account and calculate the HR value of reinfection. Sung-Ho Jun and Moon MR indicated that the complete excision of infective material is much more important than avoiding prosthetic material in preventing recurrence (31, 43). Anyway, we should not ignore either the role of residual infected tissue or artificial implants in reinfection. A successful MVP for patients of IE should comprise of thorough resecting of infected tissue and minimal artificial material implantation at the same time, which requires experience in repair.

Although surgery is important, pre- and post-operative antibiotic treatment is also essential for IE therapy, especially

Študy or Subgroup	Events	Total	Events	Total	Weight	M-H, Fixed, 95% Cl		M-H, Fixed, 95% Cl	
Anton Tomsic 2017	0	51	0	32		Not estimable			
Elfriede Ruttmann 2005	1					0.18 [0.02, 1.59]	·	<b>_</b>	
Eric shang 2009	2								
Gregorio P. Cuerpo 2019	2								
Hiroichiro Yamaguchi 2006	0					Not estimable			
Hsiu-An Lee 2018	0	38	0	33		Not estimable	1		
lose L. Navia 2019	4	52	7	86	16.3%	0.94 [0.26, 3.38]	1		
Michele Musci 2010	3	61							
Rufin J. Defauw 2020	5	97	1	53	4.1%				_
Sossio Perrota 2017	8	76	4	64	13.0%				
Suna-Ho Juna 2011	3	41	0	61	1.2%	11.18 [0.56, 222.46]			<b>→</b>
Takashi Miura 2014	1								
Thitipong Tepsiwan 2019	0	38	5	76	12.2%				
Tomislav Mihaljevic 2004	2	21	1	32	2.4%	3.26 [0.28, 38.48]			
Total (95% CI)		683		999	100.0%	1.10 [0.67, 1.81]		•	
								T	
Total events Heterogeneity: Chi <sup>2</sup> = 9.99, o Test for overall effect: Z = 0.		P = 0.4	43 4); I <sup>2</sup> = 0					0.1 1 10 Favours [MVP] Favours [MVR]	100
Heterogeneity: Chi <sup>2</sup> = 9.99, 6 Test for overall effect: Z = 0.	df = 10 (	P = 0.4		)%		Odds Ratio			100
Heterogeneity: Chi <sup>2</sup> = 9.99, Test for overall effect: Z = 0. <b>}</b>	df = 10 (i .39 (P = 0 MVP	P = 0.4 ).70)	4); $I^2 = 0$ MVR	)%	Weight	Odds Ratio M-H, Fixed, 95% Cl		Favours [MVP] Favours [MVR]	100
Heterogeneity: Chi <sup>2</sup> = 9.99, , Test for overall effect: Z = 0. <b>}</b> Study or Subgroup	df = 10 (i .39 (P = 0 MVP	P = 0.4 ).70)	4); $I^2 = 0$ MVR	)%	Weight			avours [MVP] Favours [MVR] Odds Ratio	100
Heterogeneity: Chi <sup>2</sup> = 9.99, Test for overall effect: Z = 0. <b>}</b>	df = 10 ( .39 (P = ( MVP Events	P = 0.4 ).70) <b>Total</b>	4); I <sup>2</sup> = 0 MVR Events	)% Total	<u>Weight</u> 23.6%	M-H, Fixed, 95% Cl		avours [MVP] Favours [MVR] Odds Ratio	100
Heterogeneity: Chi <sup>2</sup> = 9.99, d Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005	df = 10 ( .39 (P = 0 <u>MVP</u> <u>Events</u> 0	P = 0.4 ).70) Total 51	4); $I^2 = 0$ MVR Events 0	)% Total 32		M-H, Fixed, 95% CI Not estimable		avours [MVP] Favours [MVR] Odds Ratio	100
Heterogeneity: Chi <sup>2</sup> = 9.99, d Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019	df = 10 (( .39 (P = ( <u>MVP</u> <u>Events</u> 0 1	P = 0.4 ().70) Total   51 34	$4); I^{2} = 0$ $MVR$ Events $0$ 5	70% Total 32 34	23.6%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59]		avours [MVP] Favours [MVR] Odds Ratio	100
Heterogeneity: Chi <sup>2</sup> = 9.99, d Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019 Hsiu-An Lee 2018	df = 10 (( .39 (P = ( <u>MVP</u> <u>Events</u> 0 1 2	P = 0.4 ().70) Total 1 51 34 68	4); I <sup>2</sup> = 0 MVR Events 0 5 7	70% Total 32 34 301	23.6%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27]		avours [MVP] Favours [MVR] Odds Ratio	100
Heterogeneity: Chi <sup>2</sup> = 9.99, o Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019 Hsiu-An Lee 2018 Michele Musci 2010	df = 10 (( .39 (P = ( <b>MVP</b> <b>Events</b> 0 1 2 0	P = 0.4 ().70) Total   51 34 68 38	4); I <sup>2</sup> = 0 MVR Events 0 5 7 0	<b>Total</b> 32 34 301 33	23.6% 12.2%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27] Not estimable		avours [MVP] Favours [MVR] Odds Ratio	100
Heterogeneity: Chi <sup>2</sup> = 9.99, o Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019 Hsiu-An Lee 2018 Michele Musci 2010 Rufin J. Defauw 2020	df = 10 (( .39 (P = ( <u>MVP</u> <u>Events</u> 0 1 2 0 3	P = 0.4 0.70) Total 51 34 68 38 61	4); I <sup>2</sup> = 0 MVR Events 0 5 7 0 13	<b>Total</b> 32 34 301 33 166	23.6% 12.2% 32.3% 6.0%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27] Not estimable 0.61 [0.17, 2.21]		avours [MVP] Favours [MVR] Odds Ratio	100
Heterogeneity: Chi <sup>2</sup> = 9.99, d Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019 Hsiu-An Lee 2018 Michele Musci 2010 Rufin J. Defauw 2020 Sung-Ho Jung 2011	df = 10 (( .39 (P = ( <u>MVP</u> <u>Events</u> 0 1 2 0 3 5	P = 0.4 0.70) Total 1 34 68 38 61 97	4); I <sup>2</sup> = 0 <b>MVR</b> <u>Events</u> 0 5 7 0 13 1	Total         32           34         301           33         166           53         53	23.6% 12.2% 32.3% 6.0%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27] Not estimable 0.61 [0.17, 2.21] 2.83 [0.32, 24.85]		avours [MVP] Favours [MVR] Odds Ratio	
Heterogeneity: Chi <sup>2</sup> = 9.99, o Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019 Hsiu–An Lee 2018 Michele Musci 2010 Rufin J. Defauw 2020 Sung–Ho Jung 2011 Takashi Miura 2014	df = 10 (( .39 (P = ( <b>MVP</b> <b>Events</b> 0 1 2 0 3 5 3	P = 0.4 0.70) Total 1 51 34 68 38 61 97 41	4); I <sup>2</sup> = 0 <b>MVR</b> Events 0 5 7 0 13 1 0	Total         32           34         301           33         166           53         61	23.6% 12.2% 32.3% 6.0% 1.8%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27] Not estimable 0.61 [0.17, 2.21] 2.83 [0.32, 24.85] 11.18 [0.56, 222.46]		avours [MVP] Favours [MVR] Odds Ratio	
Heterogeneity: Chi <sup>2</sup> = 9.99, d Test for overall effect: Z = 0. <b>}</b> Study or Subgroup Anton Tomsic 2017	df = 10 (( .39 (P = ( MVP Events 0 1 2 0 3 5 3 1	P = 0.4 D.70) Total 1 51 34 68 38 61 97 41 36	4); I <sup>2</sup> = C MVR Events 0 5 7 0 13 1 0 0	Total         32           34         301           33         166           53         61           21         21	23.6% 12.2% 32.3% 6.0% 1.8% 2.9%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27] Not estimable 0.61 [0.17, 2.21] 2.83 [0.32, 24.85] 11.18 [0.56, 222.46] 1.82 [0.07, 46.63]		avours [MVP] Favours [MVR] Odds Ratio	
Heterogeneity: Chi <sup>2</sup> = 9.99, of Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019 Hsiu-An Lee 2018 Michele Musci 2010 Rufin J. Defauw 2020 Sung-Ho Jung 2011 Takashi Miura 2014 Thitipong Tepsiwan 2019	df = 10 (( .39 (P = ( MVP Events 0 1 2 0 3 5 3 1 0	P = 0.4 D.70) Total 1 51 34 68 38 61 97 41 36 38	4); I <sup>2</sup> = C MVR Events 0 5 7 0 13 1 0 0 5	70% Total 32 34 301 33 166 53 61 21 76 32	23.6% 12.2% 32.3% 6.0% 1.8% 2.9% 17.8%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27] Not estimable 0.61 [0.17, 2.21] 2.83 [0.32, 24.85] 11.18 [0.56, 22.46] 1.82 [0.07, 46.63] 0.17 [0.01, 3.13]		avours [MVP] Favours [MVR] Odds Ratio	
Heterogeneity: Chi <sup>2</sup> = 9.99, o Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019 Hsiu-An Lee 2018 Michele Musci 2010 Rufin J. Defauw 2020 Sung-Ho Jung 2011 Takashi Miura 2014 Thitipong Tepsiwan 2019 Tomislav Mihaljevic 2004	df = 10 (( .39 (P = ( MVP Events 0 1 2 0 3 5 3 1 0	P = 0.4 0.70) Total   1 51 34 68 38 61 97 41 36 38 21	4); I <sup>2</sup> = C MVR Events 0 5 7 0 13 1 0 0 5	70% Total 32 34 301 33 166 53 61 21 76 32	23.6% 12.2% 32.3% 6.0% 1.8% 2.9% 17.8% 3.5%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27] Not estimable 0.61 [0.17, 2.21] 2.83 [0.32, 24.85] 11.18 [0.56, 222.46] 1.82 [0.07, 46.63] 0.17 [0.01, 3.13] 3.26 [0.28, 38.48]		avours [MVP] Favours [MVR] Odds Ratio	
Heterogeneity: Chi <sup>2</sup> = 9.99, o Test for overall effect: Z = 0. Study or Subgroup Anton Tomsic 2017 Elfriede Ruttmann 2005 Gregorio P. Cuerpo 2019 Hsiu-An Lee 2018 Michele Musci 2010 Rufin J. Defauw 2020 Sung-Ho Jung 2011 Takashi Miura 2014 Thitipong Tepsiwan 2019 Tomislav Mihaljevic 2004 Total (95% CI)	df = 10 (( .39 (P = ( <u>MVP</u> <u>Events</u> ) 1 2 0 3 5 3 1 0 2 17	P = 0.4 0.70) Total 1 51 34 68 38 61 97 41 36 38 21 485	4); I <sup>2</sup> = 0 MVR Events 0 5 7 0 13 1 0 0 5 1 32	70% Total 32 34 301 33 166 53 61 21 76 32 809	23.6% 12.2% 32.3% 6.0% 1.8% 2.9% 17.8% 3.5%	M-H, Fixed, 95% Cl Not estimable 0.18 [0.02, 1.59] 1.27 [0.26, 6.27] Not estimable 0.61 [0.17, 2.21] 2.83 [0.32, 24.85] 11.18 [0.56, 222.46] 1.82 [0.07, 46.63] 0.17 [0.01, 3.13] 3.26 [0.28, 38.48] 0.96 [0.52, 1.78]		avours [MVP] Favours [MVR] Odds Ratio	

Meta-analysis for reinfection events between MVP and MVR; (A) MVP vs. MVR; (B) MVP vs. MVR (active IE); MVP, mitral valve plasty; MVR, mitral valve replacement.

for those patients of active phase. It is necessary for patients to receive effective antimicrobial treatment which may decrease the positive cultures from explanted valves after surgery (44). Increasingly invasive health care intervention has changed the distribution of bacteremia in IE patients which makes culture result more important (45). Song Wan reported that 85% of IE cases were culture-positive before operations. And operative specimens are particularly valuable for those culture-negative ones (35, 36). 6-weeks duration of intravenous antimicrobial treatmet is recommended following surgical intervention and longer treatment is necessary when invasive infections, difficult-to-treat microorganisms and prosthetic materials infections exist (35).

While the reoperation rate due to all events was lower in patients with IE following MVR, no significant differences exist in patients with active IE following mitral intervention. It can be concluded that the reoperation rate due to all events was lower in patients with healed IE following valve replacement rather than repair. And mitral insufficiency accounts for most of those reoperations (46). For patients with active IE, similar reoperation rate is due to less damaged valvular structure and better physical and cardiac condition in

early infective stage (39). While for patients with healed IE, more reoperation events in MVP group may be the result of the more extensive destruction of infective valve tissue, annulus and/or the subannular apparatus (47). Wan and associators reported that while vegetation was the most common pathogen in acute IE, valve prolapse resulted from chordae rupture was the most common one in healed IE (36). Repairs in the presence of healed but destructive valves are inevitably more complicated.

Here we have to mention annular abscess as a special pathogen. It represents much more serious situation, but fortunately, it is uncommon. Wan reported that annular or paravalvular abscess was reported in less than 10% of patients (36). Due to the complexity of the anatomy, radical debridement, sterilization and drainage of the infected area are more difficult. Plasty needs more experience and skills, and patches must be generous to minimize tension on the suture lines. Replacement without additional reconstruction is also a good choice. The valve prosthesis could be anchored to the ventricular muscle or to the reconstruction patch in a way that prevents leakage and pseudoaneurysm development beneath the prosthesis. And of course, annular reconstruction

is still needed when it is necessary during replacement procedures (48).

The MVP approach is patient-specific (11). Surgeons performs valvular repairs for IE patients mostly based on their experience of noninfectious valve disease repair (36). No guidelines have been raised and repairs in infectious valves are much more complicated. The ratio and underlying patients of MVP varied greatly among different surgical teams and centres. While some centres can perform difficult repair, including annular abscesses (33), some other centres only perform MVP in patients with vegetations alone (14). This does increase the heterogeneity of our research. The surgeon's experience in MVP is crucial. Only surgeons with rich experience and sufficient skills are capable of repairing severely destructed mitral valves (33). At the same time, strict patient selection is also essential. It's important to distinguish patients that are likely to be repaired from the patients that may have poor repair results.

#### Limitations

Several limitations exist in this analysis. Firstly, only retrospective studies were included in this study. Secondly, the operative years were of a broader range which might have implications in terms of the comparability of studies included in this analysis. Thirdly, there were only a few patients in some analyses which may resulted in higher selection bias. Fourthly, the mechods and technics of MVP were varied in all studies, mainly depanding on surgeons' experience. Fifthly, no direct comparison between MVP and MVR in patients with healed IE.

## Conclusions

To sum up, MVP and MVR are both beneficial for patients suffering from IE. Both two therapeutic methods have their own advantages and disadvantages. It is still hard to determine which one is better. For surgeons with consummate skills, MVP can be the preferred choice for suitable patients suffering from IE. And active phase is not a disadvantageous factor in MVP. More randomized controlled trials should be conducted.

## References

1. Feringa HH, Shaw LJ, Poldermans D, Hoeks S, van der Wall EE, Dion RA, et al. Mitral valve repair and replacement in endocarditis: a systematic review of literature. *Ann Thorac Surg.* (2007) 83(2):564–70. doi: 10.1016/j.athoracsur.2006.09.023

## Data availability statement

The raw data supporting the conclusions of this article will be made available by the authors, without undue reservation.

## Author contributions

YJ, SC designed research; SW, TZ, JB conducted research; GL, WZ analyzed data; SW, SC wrote the paper; YJ, SC, WZ had primary responsibility for final content. All authors contributed to the article and approved the submitted version.

#### Funding

This study was supported by Natural Science Foundation of China (82200409).

## **Conflict of interest**

The authors declare that the research was conducted in the absence of any commercial or financial relationships that could be construed as a potential conflict of interest.

## Publisher's note

All claims expressed in this article are solely those of the authors and do not necessarily represent those of their affiliated organizations, or those of the publisher, the editors and the reviewers. Any product that may be evaluated in this article, or claim that may be made by its manufacturer, is not guaranteed or endorsed by the publisher.

#### Supplementary material

The Supplementary Material for this article can be found online at: https://www.frontiersin.org/articles/10.3389/fsurg. 2022.1048036/full#supplementary-material.

<sup>2.</sup> Duval X, Delahaye F, Alla F, Tattevin P, Obadia JF, Le Moing V, et al. AEPEI Study group. Temporal trends in infective endocarditis in the context of prophylaxis guideline modifications: three successive population-based surveys. *J Am Coll Cardiol.* (2012) 59(22):1968–76. doi: 10.1016/j.jacc.2012.02.029

<sup>3.</sup> Antunes MJ. Mitral valve repair versus replacement for infective endocarditis. What is better in the "real world"? *J Thorac Cardiovasc Surg.* (2018) 156 (4):1471–2. doi: 10.1016/j.jtcvs.2018.05.010

<sup>4.</sup> Jiang Y, Wang C, Li G, Chen S. Clinical outcomes following surgical mitral valve repair or replacement in patients with rheumatic heart disease: a meta-analysis. *Ann Transl Med.* (2021) 9(3):204. doi: 10.21037/atm-20-3542

5. Harky A, Hof A, Garner M, Froghi S, Bashir M. Mitral valve repair or replacement in native valve endocarditis? Systematic review and meta-analysis. *J Card Surg.* (2018) 33(7):364–71. doi: 10.1111/jocs.13728

6. Liu JZ, Li XF, Miao Q, Zhang CJ. Surgical treatment of active native mitral infective endocarditis: a meta-analysis of current evidence. *J Chin Med Assoc.* (2018) 81(2):147–54. doi: 10.1016/j.jcma.2017.08.017

7. Parmar MK, Torri V, Stewart L. Extracting summary statistics to perform meta-analyses of the published literature for survival endpoints. *Stat Med.* (1998) 17(24):2815–34. doi: 10.1002/(SICI)1097-0258(19981230)17:24<2815:: AID-SIM110>3.0.CO:2-8

8. Williamson PR, Smith CT, Hutton JL, Marson AG. Aggregate data metaanalysis with time-to-event outcomes. *Stat Med.* (2002) 21(22):3337–51. doi: 10. 1002/sim.1303

9. Tierney JF, Stewart LA, Ghersi D, Burdett S, Sydes MR. Practical methods for incorporating summary time-to-event data into meta-analysis. *Trials.* (2007) 8:16. doi: 10.1186/1745-6215-8-16

10. Kim JB, Kim HJ, Moon DH, Jung SH, Choo SJ, Chung CH, et al. Long-term outcomes after surgery for rheumatic mitral valve disease: valve repair versus mechanical valve replacement. *Eur J Cardio-Thorac.* (2010) 37(5):1039–46. doi: 10.1016/j.ejcts.2009.11.019

11. Chang HW, Kim KH, Hwang HY, Kim JS. Role of mitral valve repair in infective endocarditis. *J Heart Valve Dis.* (2014) 23(3):350–9.

12. Musci M, Hübler M, Pasic M, Amiri A, Stein J, Siniawski H, et al. Surgery for active infective mitral valve endocarditis: a 20-year, single-center experience. *J Heart Valve Dis.* (2010) 19(2):206–14; discussion 215.

13. Wilhelm MJ, Tavakoli R, Schneeberger K, Hörstrupp S, Reuthebuch O, Seifert B, et al. Surgical treatment of infective mitral valve endocarditis. *J Heart Valve Dis.* (2004) 13(5):754–9.

14. Mihaljevic T, Paul S, Leacche M, Rawn JD, Aranki S, O'Gara PT, et al. Tailored surgical therapy for acute native mitral valve endocarditis. *J Heart Valve Dis.* (2004) 13(2):210–6.

15. Sternik L, Zehr KJ, Orszulak TA, Mullany CJ, Daly RC, Schaff HV. The advantage of repair of mitral valve in acute endocarditis. *J Heart Valve Dis.* (2002) 11(1):91–7; discussion 97–8.

16. Tepsuwan T, Rimsukcharoenchai C, Tantraworasin A, Taksaudom N, Woragidpoonpol S, Chuaratanaphong S, et al. Comparison between mitral valve repair and replacement in active infective endocarditis. *Gen Thorac Cardiovasc Surg.* (2019) 67(12):1030–7. doi: 10.1007/s11748-019-01132-4

17. Shang E, Forrest GN, Chizmar T, Chim J, Brown JM, Zhan M, et al. Mitral valve infective endocarditis: benefit of early operation and aggressive use of repair. *Ann Thorac Surg.* (2009) 87(6):1728–33. doi: 10.1016/j.athoracsur. 2009.02.098

18. Muehrcke DD. Cosgrove DM 3rd, lytle BW. Is there an advantage to repairing infected mitral valves? *Ann Thorac Surg.* (1997) 63(6):1718–24. doi: 10.1016/S0003-4975(97)00271-3

19. Miura T, Hamawaki M, Hazama S, Hashizume K, Ariyoshi T, Sumi M, et al. Outcome of surgical management for active mitral native valve infective endocarditis: a collective review of 57 patients. *Gen Thorac Cardiovasc Surg.* (2014) 62(8):488–98. doi: 10.1007/s11748-014-0377-y

20. Lee HA, Lin CY, Chen YC, Chen SW, Nan YY, Liu KS, et al. Surgical interventions of isolated active mitral valve endocarditis: predisposing factors and impact of neurological insults on final outcome. *Medicine (Baltimore)*. (2018) 97(11):e0054. doi: 10.1097/MD.000000000010054

21. Alkhouli M, Alqahtani F, Berzingi C, Cook CC. Contemporary trends and outcomes of mitral valve surgery for infective endocarditis. *J Card Surg.* (2019) 34(7):583–90. doi: 10.1111/jocs.14116

22. Tomšic A, Versteegh MIM, Ajmone Marsan N, van Brakel TJ, Klautz RJM, Palmen M. Early and late results of surgical treatment for isolated active native mitral valve infective endocarditis. *Interact Cardiovasc Thorac Surg.* (2018) 26 (4):610–6. doi: 10.1093/icvts/ivx395

23. Gammie JS, O'Brien SM, Griffith BP, Peterson ED. Surgical treatment of mitral valve endocarditis in North America. *Ann Thorac Surg.* (2005) 80 (6):2199–204. doi: 10.1016/j.athoracsur.2005.05.036

24. Solari S, De Kerchove L, Tamer S, Aphram G, Baert J, Borsellino S, et al. Active infective mitral valve endocarditis: is a repair-oriented surgery safe and durable? *Eur J Cardiothorac Surg.* (2019) 55(2):256–62. doi: 10.1093/ejcts/ezy242

25. Perrotta S, Fröjd V, Lepore V, Schersten H, Jeppsson A, Svensson G. Surgical treatment for isolated mitral valve endocarditis: a 16-year single-centre experience. *Eur J Cardiothorac Surg.* (2018) 53(3):576–81. doi: 10.1093/ejcts/ezx416

26. Defauw RJ, Tomšič A, van Brakel TJ, Marsan NA, Klautz RJM, Palmen M. A structured approach to native mitral valve infective endocarditis: is repair better than replacement? *Eur J Cardiothorac Surg.* (2020) 58(3):544–50. doi: 10.1093/ejcts/ezaa079

27. Cuerpo GP, Valerio M, Pedraz A, Rodríguez-Abella H, Mestres CA, Obrador EQ, et al. GAMEs study group. Mitral valve repair in infective endocarditis is not inferior to valve replacement: results from a spanish nationwide prospective registry. *Gen Thorac Cardiovasc Surg.* (2019) 67(7):585–93. doi: 10.1007/s11748-019-01063-0

28. Navia JL, Elgharably H, Hakim AH, Witten JC, Haupt MJ, Germano E, et al. Long-term outcomes of surgery for invasive valvular endocarditis involving the aortomitral Fibrosa. *Ann Thorac Surg.* (2019) 108(5):1314–23. doi: 10.1016/j. athoracsur.2019.04.119

29. Wang TK, Oh T, Voss J, Gamble G, Kang N, Pemberton J. Valvular repair or replacement for mitral endocarditis: 7-year cohort study. *Asian Cardiovasc Thorac Ann*. (2014) 22(8):919–26. doi: 10.1177/0218492314521613

30. Yamaguchi H, Eishi K, Yamachika S, Hisata Y, Tanigawa K, Izumi K, et al. Mitral valve repair in patients with infective endocarditis. *Circ J.* (2006) 70 (2):179–83. doi: 10.1253/circj.70.179

31. Jung SH, Je HG, Choo SJ, Song H, Chung CH, Lee JW. Surgical results of active infective native mitral valve endocarditis: repair versus replacement. *Eur J Cardiothorac Surg.* (2011) 40(4):834–9. doi: 10.1016/j.ejcts.2011.01.016

32. Toyoda N, Itagaki S, Egorova NN, Tannous H, Anyanwu AC, El-Eshmawi A, et al. Real-world outcomes of surgery for native mitral valve endocarditis. *J Thorac Cardiovasc Surg.* (2017) 154(6):1906–1912.e9. doi: 10.1016/j.jtcvs.2017. 07.077

33. Ruttmann E, Legit C, Poelzl G, Mueller S, Chevtchik O, Cottogni M, et al. Mitral valve repair provides improved outcome over replacement in active infective endocarditis. *J Thorac Cardiovasc Surg.* (2005) 130(3):765–71. doi: 10. 1016/j.jtcvs.2005.03.016

34. Pant S, Patel NJ, Deshmukh A, Golwala H, Patel N, Badheka A, et al. Trends in infective endocarditis incidence, microbiology, and valve replacement in the United States from 2000 to 2011. *J Am Coll Cardiol.* (2015) 65(19):2070–6. doi: 10.1016/j.jacc.2015.03.518

35. AATS Surgical Treatment of Infective Endocarditis Consensus Guidelines Writing Committee Chairs, Pettersson GB, Coselli JS, Writing Committee, Pettersson GB, Coselli JS, Hussain ST, et al. 2016 The American association for thoracic surgery (AATS) consensus guidelines: surgical treatment of infective endocarditis: executive summary. *J Thorac Cardiovasc Surg.* (2017) 153 (6):1241–1258.e29. doi: 10.1016/j.jtcvs.2016.09.093

36. Hu YN, Wan S. Repair of infected mitral valves: what have we learned? Surg Today. (2018) 48(10):899-908. doi: 10.1007/s00595-018-1637-8

37. García-Cabrera E, Fernández-Hidalgo N, Almirante B, Ivanova-Georgieva R, Noureddine M, Plata A, et al. Neurological complications of infective endocarditis: risk factors, outcome, and impact of cardiac surgery: a multicenter observational study. *Circulation*. (2013) 127(23):2272–84. doi: 10.1161/CIRCULATIONAHA. 112.000813

38. Beynon RP, Bahl VK, Prendergast BD. Infective endocarditis. Br Med J. (2006) 333(7563):334–9. doi: 10.1136/bmj.333.7563.334

39. Nishimura RA, Otto CM, Bonow RO, Carabello BA, Erwin 3rd JP, Guyton RA, et al. 2014 AHA/ACC guideline for the management of patients with valvular heart disease: executive summary: a report of the American college of cardiology/ American heart association task force on practice guidelines. *Circulation*. (2014) 129(23):2440–92. doi: 10.1161/CIR.00000000000029

40. Carrel T. Early valve repair or replacement is not generally contraindicated in patients with infective endocarditis and stroke with or without intracranial haemorrhage. *Eur J Cardiothorac Surg.* (2016) 50(2):383–4. doi: 10.1093/ejcts/ezw090

41. Pektok E, Sierra J, Cikirikcioglu M, Müller H, Myers PO, Kalangos A. Midterm results of valve repair with a biodegradable annuloplasty ring for acute endocarditis. *Ann Thorac Surg.* (2010) 89(4):1180–5. doi: 10.1016/j.athoracsur. 2010.01.041

42. Musci M, Hübler M, Amiri A, Stein J, Kosky S, Weng Y, et al. Repair for active infective atrioventricular valve endocarditis: 23-year single center experience. *Clin Res Cardiol.* (2011) 100(11):993–1002. doi: 10.1007/s00392-011-0331-2

43. Moon MR, Miller DC, Moore KA, Oyer PE, Mitchell RS, Robbins RC, et al. Treatment of endocarditis with valve replacement: the question of tissue versus mechanical prosthesis. *Ann Thorac Surg.* (2001) 71(4):1164–71. doi: 10.1016/S0003-4975(00)02665-5

44. Mekontso Dessap A, Zahar JR, Voiriot G, Ali F, Aissa N, Kirsch M, et al. Influence of preoperative antibiotherapy on valve culture results and outcome

of endocarditis requiring surgery. J Infect. (2009) 59(1):42-8. doi: 10.1016/j.jinf. 2009.04.009

45. Murdoch DR, Corey GR, Hoen B, Miró JM, Fowler Jr VG, Bayer AS, et al. Clinical presentation, etiology, and outcome of infective endocarditis in the 21st century: the international collaboration on endocarditis-prospective cohort study. *Arch Intern Med.* (2009) 169(5):463–73. doi: 10.1001/archinternmed.2008.603

46. Katan O, Michelena HI, Avierinos JF, Mahoney DW, DeSimone DC, Baddour LM, et al. Incidence and predictors of infective endocarditis in mitral

valve prolapse: a population-based study. Mayo Clin Proc. (2016) 91(3):336–42. doi: 10.1016/j.mayocp.2015.12.006

47. Huuskonen A, Vento A, Raivio P. Outcome of early vs. delayed surgery for infective mitral endocarditis. *J Card Surg.* (2019) 34(8):700–7. doi: 10.1111/jocs. 14137

48. Pettersson GB, Hussain ST. Current AATS guidelines on surgical treatment of infective endocarditis. *Ann Cardiothorac Surg.* (2019) 8(6):630–44. doi: 10. 21037/acs.2019.10.05