

Systematic review and meta-analysis of surgical outcomes comparing mechanical valve replacement and bioprosthetic valve replacement in infective endocarditis

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Background: Infective endocarditis (IE) is an infection involving either native or prosthetic heart valves, the endocardial surface of the heart or any implanted intracardiac devices. IE is a rare condition affecting 3–15 patients per 100,000 population. In-hospital mortality rates in patients with IE remain high at around 20% despite treatment advances. There is no consensus recommendation favoring either bioprosthetic valve or mechanical valve implantation in the setting of IE; patient age, co-morbidities and preferences should be considered selecting the replacement prosthesis.

Methods: A systematic review and meta-analysis of studies reporting the outcomes of patients undergoing bioprosthetic or mechanical valve replacement for infective endocarditis with data extracted for overall survival, valve reinfection rates and valve reoperation.

Results: Eleven relevant studies were identified, with 2,336 patients receiving a mechanical valve replacement and 2,057 patients receiving a bioprosthetic valve replacement. There was no significant difference for overall survival between patients treated with mechanical valves and those treated with bioprosthetic valves [hazard ratio (HR) 0.94, 95% confidence interval (CI): 0.73–1.21, P=0.62]. There was no significant difference in reoperation rates between patients treated with a bioprosthetic valve and those treated with a mechanical valve (HR 0.82, 95% CI: 0.34–1.98, P=0.66) and there was no significant difference in the rate of valve reinfection rates (HR 0.95, 95% CI: 0.48–1.89, P=0.89).

Conclusions: The presence of infective endocarditis alone should not influence the decision of which type of valve prosthesis that should be implanted. This decision should be based on patient age, co-morbidities and preferences.

Keywords: Valve replacement; tissue valve; bioprosthesis; bioprosthetic; mechanical; infective endocarditis



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Introduction

Infective endocarditis (IE) is an infection involving either native or prosthetic heart valves, the endocardial surface of the heart or any implanted intracardiac devices (1). The intact endocardium is an effective barrier to infection, with the risk of endocarditis rising for individuals with congenital intracardiac defects, acquired degenerative valvular lesions and implanted intra-cardiac prostheses (2); however, it remains an important condition with significant associated morbidity and mortality. IE is a rare condition affecting 3–15 patients per 100,000 population (3–5). Infective endocarditis is commonly diagnosed according to the modified Duke criteria (6) which determines the likelihood of a patient having IE based on organisms detected on blood culture, echocardiographic images and clinical examination. In patients with prosthetic valve replacement, the risk of developing prosthetic endocarditis is 2.2–3.7% (7–9). Aggressive surgical treatment is often required with surgery indicated in up to half of patients with native valve endocarditis (10,11). In-hospital mortality rates in patients with IE remain high at around 20% despite treatment advances (11).

Indications for surgical intervention for left-sided IE are relatively well defined with urgent intervention suggested for patients without adequate control of sepsis despite appropriate antibiotics, for patients with evidence of intra-cardiac abscess formation or other paravalvular extension, for patients with heart failure secondary to valve destruction or fistula formation, and to prevent catastrophic embolic phenomena in the presence of large mobile vegetations (12,13). Where possible, valve repair is preferred over replacement, particularly for right-sided and mitral valve IE. There is no consensus recommendation for either bioprosthetic valve or mechanical valve in the setting of IE and patient age, co-morbidities and preferences should be considered in making the decision for valve selection (12,14).

In the general population, the European Society of Cardiology suggest treatment with mechanical valves under the age of 60 for a valve in the aortic position or under the age 65 for a valve in the mitral position (15). The American Heart Association guidelines recommend a mechanical valve prosthesis in patients under the age of 50 and a bioprosthetic valve prosthesis over the age of 70. In between these ages, patient factors and preferences will determine prosthesis type (16). This meta-analysis was therefore undertaken to evaluate the surgical outcomes

comparing mechanical valve replacement and bioprosthetic valve replacement in infective endocarditis.

Methods

Literature search strategy

This meta-analysis was performed in accordance with PRISMA recommendations and guidance. The search strategy was employed to search electronic databases EMBASE, Ovid Medline, the entire Cochrane Central Register of Controlled Trials (CCRCT), Cochrane Database of Systematic reviews (CDSR), the Database of Abstracts of Reviews of Effects (DARE) and the ACP journal club from their inception to 03 March 2019. The search strategy included search terms for “(tissue valve OR bioprosthesis OR biological valve OR bioprosthetic valve OR xenograft OR mechanical valve) AND endocarditis”. The references of previous systematic reviews were assessed to ensure no additional publications were missed.

Selection criteria

Eligibility for inclusion in this systematic review and meta-analysis included papers that assessed the outcomes of patients undergoing conventional valve replacement with a bioprosthetic or mechanical valve prosthesis undertaken in adult patients for the treatment of infective endocarditis. In order to ensure sufficient center experience, papers were only included if more than 15 cases were reported in each arm. English language papers with at least 1-year follow-up were considered for assessment. Studies were excluded if there was inadequate data regarding the outcomes and for patients treated with homograft as a bioprosthesis. If centers reported outcomes of overlapping patient series, then the most contemporary series was analyzed. Conference abstracts, case reports, editorials, expert opinion, reviews and expert opinion were excluded.

Data extraction

For the assessed papers, data was extracted from the reviewed text, tables and figures. Data was extracted independently by two of the authors (CD Flynn, NP Curran) and any discrepancies were reviewed and discussed until consensus was reached. The recorded parameters were: number of cases in series, procedure undertaken, urgency of procedure, average age, average follow-up, early

death, late death, valve reinfection, valve reoperation.

Statistical analysis

Meta-analysis of incidence rates of post-operative complications including valve reinfection, reoperation and overall survival. Hazard ratios were calculated from source Kaplan-Meier curves according to Tierney *et al.* methods (17) with mechanical valve being the reference group. Incidence data was assessed using Comprehensive Meta-analysis v3.3 (Biostat, Englewood, USA). Comparative outcomes were assessed using Review Manager v5.3 (Cochrane Collaboration, Copenhagen, Denmark). Due to the varied patient populations, a random effects model was chosen for analysis. Data heterogeneity was assessed with the Cochrane Q statistic with P value <0.05 being significant and the I^2 test statistic an I^2 value greater than 50% denoting significant heterogeneity. Publication bias was assessed by generation of funnel plot and assessment using Egger's test produced with Comprehensive Meta-analysis. Sensitivity analysis was performed using a "leave-one-out" analysis. Individual patient survival data, reinfection and reoperation data were reconstructed using an iterative algorithm that was applied to digitized source Kaplan-Meier curves and subsequently aggregated and graphed (18) using R (R Foundation for Statistical Computing, Vienna, Austria).

Results

The search strategy revealed 4,053 citations for review after duplications were removed with an additional two studies added for full text review on reviewing reference lists. After full review, 11 papers met the pre-determined inclusion criteria (19-29) (*Figure S1*). The included publications had a total analyzed patient population of 4,393 patients with 2,336 patients receiving a mechanical valve replacement and 2,057 patients receiving a bioprosthetic valve replacement. The overall study characteristics are detailed in *Table 1*. An assessment of bias was made for all studies which demonstrated significant risks of bias in all included studies, largely based on study design (*Figure S2*).

The mean age of patients receiving bioprosthetic valves calculated from six studies reporting mean age of patients (19,20,22,23,28,29) was 59.2 years old (95% CI: 58.2–60.1). The mean age of patients receiving mechanical valves was 52.1 years old (95% CI: 51.4–52.9). Overall, patients receiving a mechanical valve were younger by 7.8 years (95%

CI: 6.6–9.0, $P<0.001$).

Ten studies reported overall survival data (19,21-29), drawing outcomes data from 3,884 patients (1,792 patients receiving bioprosthetic valves and 2,092 patients receiving mechanical valves). Meta-analysis of derived hazard ratio (HR) demonstrated no significant difference for overall survival between patients treated with mechanical valves and those treated with bioprosthetic valves (HR 0.94, 95% CI: 0.73–1.21; $P=0.62$) (*Figure 1*). There was significant heterogeneity Cochrane Q-statistic P value <0.001, $I^2=75\%$. Post-hoc subgroup analysis performed to explore sources of heterogeneity demonstrated that analyzing outcomes of left sided infective endocarditis (21-23,25-29) showed no significant difference in overall survival, regardless of valve type, HR 0.97 (95% CI: 0.67–1.41, $P=0.89$), there was still significant heterogeneity in data, Cochrane q-statistic $P=0.005$, $I^2=68\%$ (*Figure 1*). There was a possible benefit for overall survival with bioprosthetic valve replacement in the treatment of prosthetic valve endocarditis (21,25,27) (HR 1.37, 95% CI: 1.01–1.85, $P=0.04$) (*Figure 1*). Composite survival curves demonstrated overall survival for bioprosthetic valves of 71.8%, 57.3%, 49.2%, and 37.5% at 1-, 3-, 5-, and 10-year, respectively, and 73.2%, 56.8%, 49.3%, and 34.0% at 1-, 3-, 5-, and 10-year, respectively, for mechanical valves (*Figure 2*). Leave-one-out sensitivity analysis did not demonstrate any significant impact on overall outcomes with any one study being removed. There was no evidence of publication bias on visual inspection of a funnel plot of the standard error by log hazard ration for overall survival (*Figure S3*), nor by Egger's test ($P=0.358$).

Four studies reported data on reoperation on valve prosthesis (22,23,25,29), drawing outcomes data from 531 patients (302 patients receiving bioprosthetic valves and 229 patients receiving mechanical valves). Meta-analysis of derived HR demonstrated no significant difference between reoperation rates between patients treated with a bioprosthetic valve and those treated with a mechanical valve (HR 0.82, 95% CI: 0.34–1.98, $P=0.66$) There was moderate heterogeneity with the Cochrane q statistic $P=0.12$, $I^2=49\%$ (*Figure 3*). Composite survival curves demonstrated freedom from reoperation for bioprosthetic valves of 90.1%, 80.7%, 79.2%, and 56.9% at 1-, 3-, 5-, and 10-year, respectively, and 94.1%, 91.3%, 86.0%, and 75.6% at 1-, 3-, 5-, and 10-year, respectively, for mechanical valves (*Figure 4*). Leave-one-out sensitivity analysis did not demonstrate any significant difference in overall effect on outcomes. There was no evidence of publication bias on visual inspection of a funnel plot of the standard error by

Table 1 Study characteristics

Study	Recruitment years	Location of institution	Study design	Patient population	Number of patients (n)	Follow up (median) [range]	Patient age (years) [range]
Reul 1989	1979–1984	Houston, Texas (USA)	Single-centre retrospective	NVE and PVE	Total =185 BV =88 MV =97	BV =22 months [6–68 months] MV =18 months [6–70 months]	BV =49 [8–77] MV =46.6 [11–78]
Lytle 1996	1975–1992	Cleveland, Ohio (USA)	Single-centre retrospective (registry data)	PVE	Total =128 BV =52 Mv =76	–	–
Moon 2001	1964–1995	Stanford, California (USA)	Single-centre retrospective	Left-sided valve IE (NVE and PVE)	Total =286 BV =221 MV =65	Overall =15.3 years [0–29 years]	Overall =49
Fedoruk 2009	1975–2000	British Columbia, Vancouver (Canada)	Multi-centre retrospective (registry data)	NVE	Total =358 BV =189 MV =169	Overall =5.5±5.5 years BV =6.1±6.1 years MV =4.9±4.5 years	Overall =48.8±15.9 [18–88] BV =51.6±17.3 MV =45.6±13.5
Musci 2010	1986–2008	Berlin (Germany)	Single-centre retrospective	PVE	Total =210 BV =165 Mv =45	–	–
Nguyen 2010	1998–2000	France	Multicentre prospective	Aortic valve IE (NVE and PVE)	Total =140 BV =31 MV =109	–	Overall =57.9±13.0 BV =63.2±13.6 MV =57.3±11.6
Jassar 2012	2000–2010	Philadelphia, Pennsylvania (USA)	Single-centre retrospective	ARR for IE	Total =98 BV =43 MV =55	BV =31.7±28.5 months MV =27.4±22.1 months	BV =62.7±15.1 MV =51.8±12.7
Leither 2013	2004–2007	USA	Multi-centre retrospective (registry data) ¹	Renal-replacement therapy patients with NVE affecting aortic or mitral valve	Total =1267 BV =561 MV =706	–	–
Greason 2014	1974–2009	Rochester, Minnesota (USA)	Single-centre retrospective (registry data)	PVE affecting the MV	Total =36 BV =18 MV =18	Overall =5.7 years [0.6–24 years]	Overall =68 [15.8–82.3]
Delahaye 2015	2000–2006	Global	Multicentre prospective	Patients with IE undergoing valve surgery in the acute phase	Total =1,467 MV =917 MV =917	–	Overall =56.6±15.7 BV =61.6±15.2 MV =53.6±15.2
Kim 2016	2002–2014	Boston, Massachusetts (USA)	Multicentre prospective	Adult patients with AVE	Total =218 BV =139 MV =79	Overall =29.4 months [4.7–72.6 months]	BV =59.8±14.6 MV =47.2±14.5

NVE, native valve endocarditis; PVE, prosthetic valve endocarditis; IE, infective endocarditis; BV, bioprosthetic valve; MV, mechanical valve.

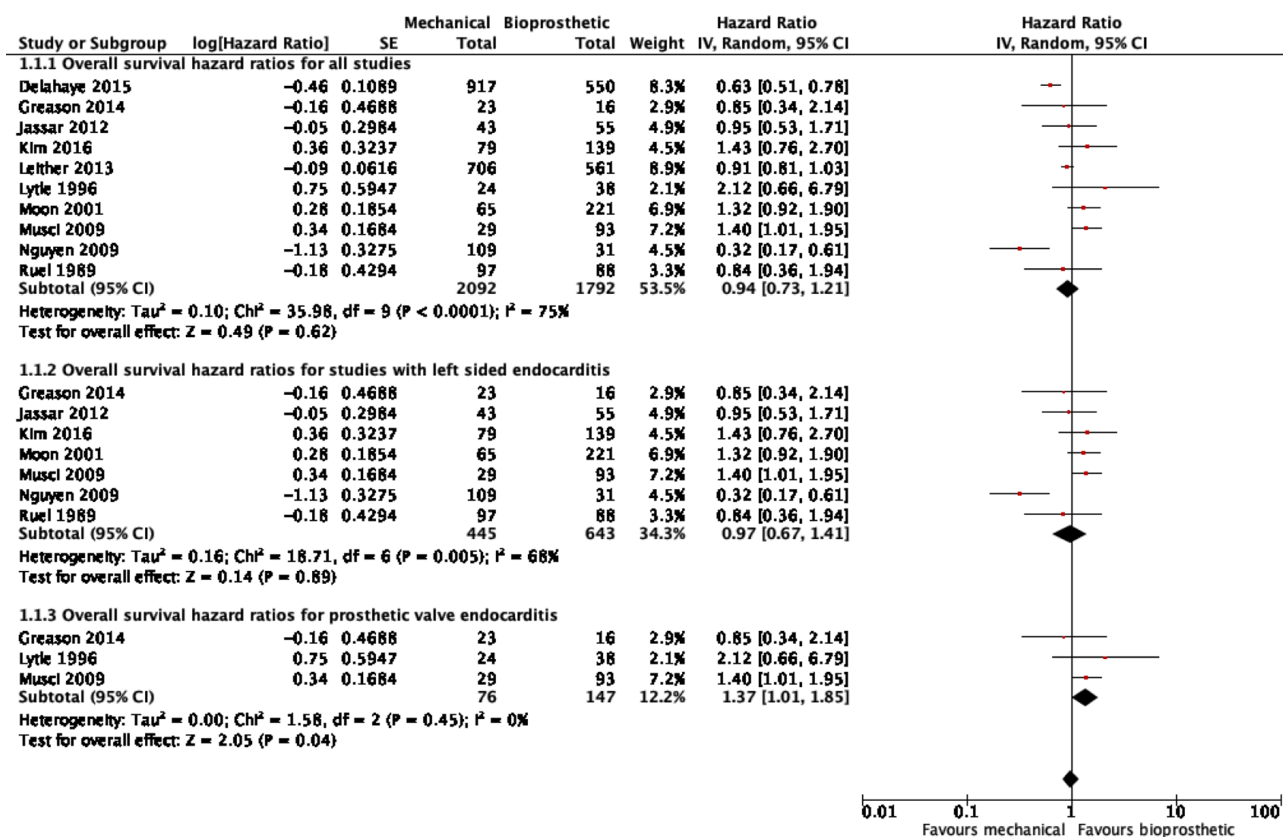


Figure 1 Forest plot detailing survival outcomes.

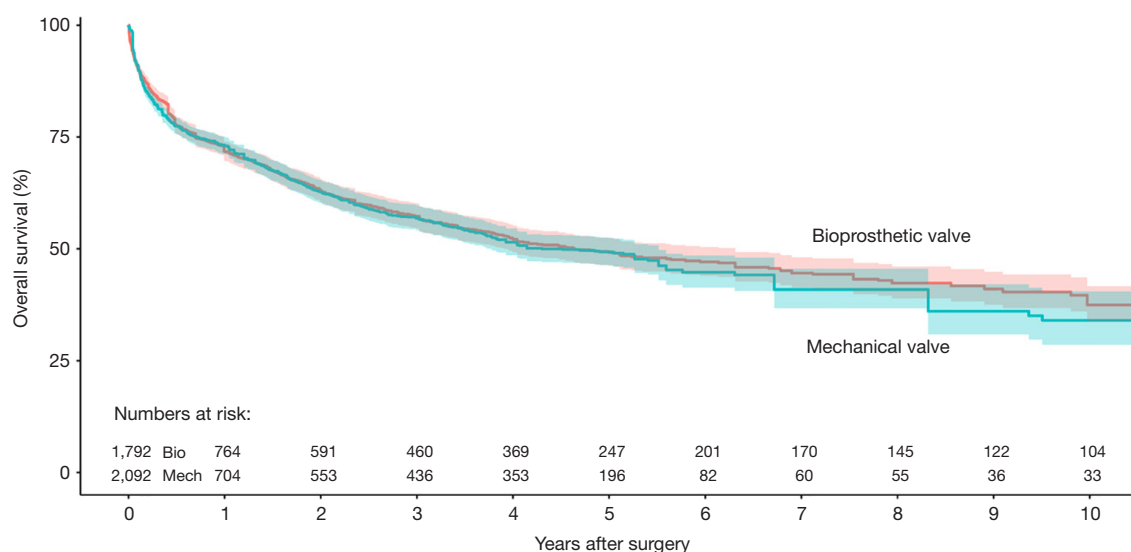


Figure 2 Composite Kaplan-Meier curve detailing overall survival for patients receiving bioprosthetic and mechanical valve for infective endocarditis (shaded region represents 95% confidence intervals).

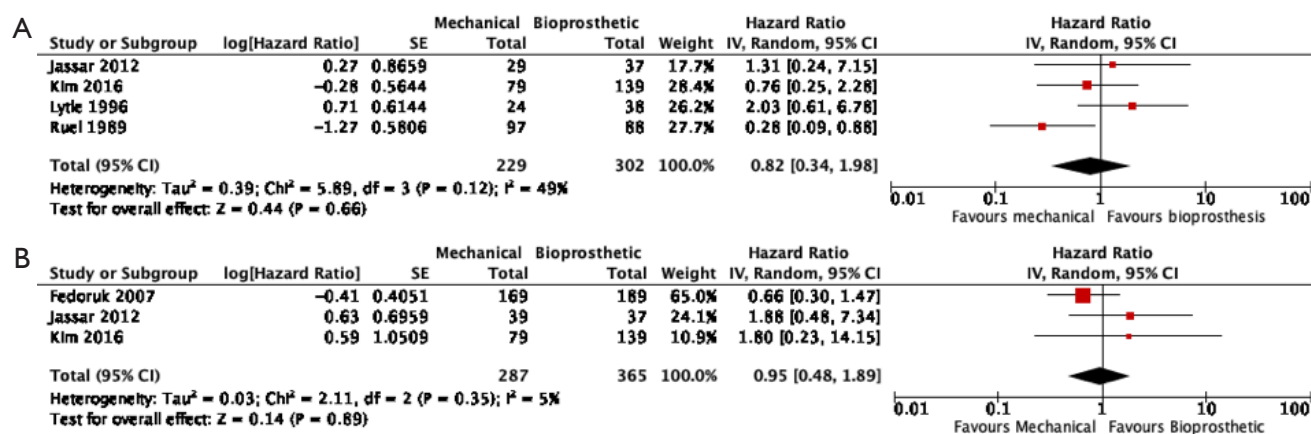


Figure 3 Forest plots detailing outcomes of (A) valve reoperation; (B) valve reinfection.

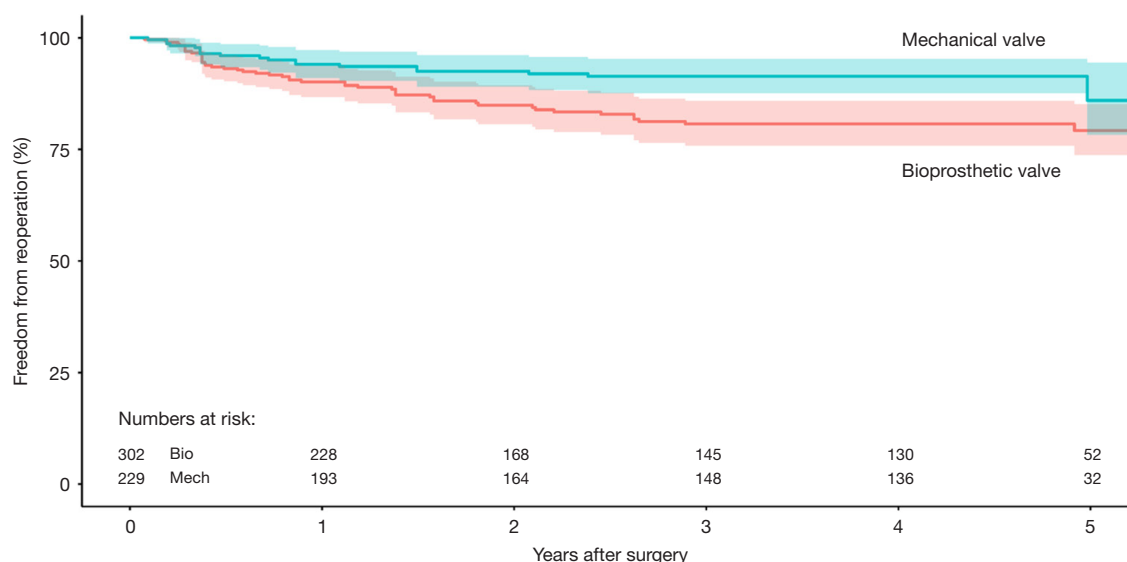


Figure 4 Composite Kaplan-Meier curve detailing freedom from reoperation for patients receiving bioprosthetic and mechanical valve for infective endocarditis (shaded region represents 95% confidence intervals).

log hazard ration for reoperation rates (Figure S4). Egger's test was not applied due to the small number of studies being assessed.

Three studies reported data on valve reinfections (20,22,23) drawing on data from 652 patients (365 patients receiving bioprosthetic valves and 287 patients receiving mechanical valves). Meta-analysis of derived hazard ratios demonstrated no significant difference in the rate of valve reinfection rates (HR 0.95, 95% CI: 0.48–1.89; $P=0.89$). There was minimal heterogeneity in this small group of studies with the Cochrane Q-statistic P value of 0.35,

$I^2=5\%$ (Figure 3B). Composite survival curves demonstrated a freedom from valve reinfection for bioprosthetic valves of 97.0%, 91.4%, 88.3%, and 79.0% at 1-, 3-, 5-, and 10-year, respectively, and 97.1%, 93.8%, 92.3%, and 86.6% at 1-, 3-, 5-, and 10-year, respectively, for mechanical valves (Figure 5). Leave-one-out sensitivity analysis did not demonstrate any significant difference in overall effect on outcomes. There was no evidence of publication bias on visual inspection of a funnel plot of the standard error by log hazard ration for valve reinfection rates (Figure S5). Egger's test was not applied due to the small number of

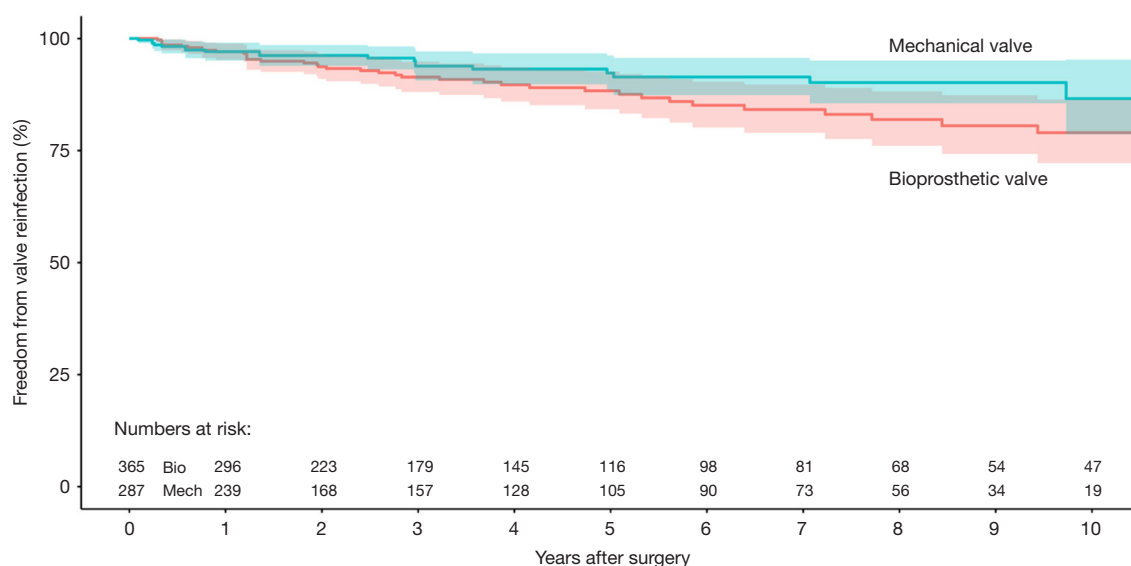


Figure 5 Composite Kaplan-Meier curve detailing freedom from valve reinfection for patients receiving bioprosthetic and mechanical valve for infective endocarditis (shaded region represents 95% confidence intervals).

studies being assessed.

Discussion

The evidence supporting indications for valve selection in infective endocarditis is currently limited. This meta-analysis suggests that at present there is no clear evidence that one prosthetic type should be selected over any other, which is nonetheless an important finding. There was significant heterogeneity when reviewing data from all analyses, which may relate to the varied patient populations being examined with variation in valve choices and operations. It was not possible to correct for severity of infective endocarditis, urgency status, concomitant operations or co-morbidities. An important consideration is that two of the three prospective studies (19,28) demonstrated a significant survival advantage for patients receiving mechanical valves, suggesting there is possibly an area of future research to develop more robust evidence.

It may have been expected that bioprosthetic valves would have worse overall survival as this patient cohort was significantly older compared to those receiving mechanical valves. Several studies reported that increasing age was an independent risk factor for overall mortality in patients with endocarditis (19,21,27,28). Nguyen *et al.* (28) note that advancing age increased the risk for overall 5-year mortality in infective endocarditis (HR 1.03; CI: 1.00–1.05, $P=0.037$),

which was most pronounced in the small number of patients who received a bioprosthetic valve under the age of 65. These patients had a markedly increased 5-year mortality compared to patients receiving a mechanical valve (HR 4.14; 95% CI: 1.27–13.45, $P=0.018$). However, patients receiving bioprosthetic valves in that study were older overall and had more medical co-morbidities including liver disease (OR 21.86) and severe left ventricle (LV) impairment (OR 14.11), therefore confounding the interpretation of the results. Musci *et al.* (27) also note age at operation as independent risk factor for early mortality. In contrast to Nguyen *et al.*, Moon *et al.* (26) found no difference in overall mortality between mechanical and bioprosthetic valves in patients under the age of 60. Unfortunately, there was insufficient data to perform adequate meta-regression to correct for patient age, co-morbidities or pre-operative status in this meta-analysis.

The choice of valve in the general population is determined by balancing the risk of anticoagulation-related and thromboembolic complications that may be associated with mechanical valves versus the risk of structural valve degeneration and the requirement for redo valve surgery for patients receiving bioprosthetic valves. The cut-off for patients receiving a mechanical valve versus a bioprosthetic valve is yet to be firmly established with conflicting evidence regarding long-term outcomes for the two valves. Some evidence refutes an excess of mortality in middle-

aged patients receiving bioprosthetic valves, including a large propensity-matched study of non-elderly patients. McClure *et al.* (30) demonstrated no significant difference in late mortality at 18 years with a cumulative survival estimated at 18 years of 60% for bioprosthetic valve and 51% for mechanical valves, though not unsurprisingly they found an increase in the requirement for reoperation for bioprosthetic valves. Similarly, although Kulick *et al.* (31) demonstrated no difference in survival for patients aged between 50 to 65 years requiring aortic or mitral valve replacement, mechanical valves were associated with a significant increased risk of thromboembolism (HR 4.1; CI: 1.3–12.7, $P=0.01$) and an increased risk of reoperation associated with bioprosthetic valves (HR 7.1; CI: 1.8–27.8 $P=0.005$). However, a recent meta-analysis of five studies by Diaz *et al.* (32), assessed valve replacement in patients between the age of 50 to 70, demonstrated a statistically significant reduction in long-term mortality for patients receiving mechanical valves (HR 0.86; 95% CI: 0.76–0.97, $P=0.01$). They did note an increased risk of major bleeding with mechanical valve recipients.

There remains no simple answer for valve choice, particularly in middle aged patients. It appears that the presence of IE alone should not influence the decision outside other patient factors. Newer-generation bioprosthetic valves with novel preservation techniques could optimistically reduce the rate of structural valve degeneration, with ongoing trials currently at 4 years follow-up and no episodes of structural valve degeneration to date (33), although long-term follow-up is required to determine if this truly is an improvement on current technologies. Another addition to the decision-making algorithm is the ever-increasing experience with valve-in-valve transcatheter aortic valve replacement which has offered equivalent mid-term survival rates with reduced early morbidity compared to conventional reoperative surgery (34–36). However, an option for valve-in-valve replacement is questionable for valves smaller than 21 mm, suggesting that expected valve size and the consequences for future management should play a role in prosthesis choice. On the other hand, the latest generation mechanical valves (On-x valve) have been proven to be safe with reduced doses of anticoagulation and have an approved target INR range of 1.5–2.0. This reduction in anticoagulation has significantly reduced major and minor bleeding events without causing a significant increase in thromboemboli (37). This may address some of the major drawbacks associated with mechanical valve replacement.

There are several limitations that must be considered when interpreting results from the present review. Data has been collected from a relatively small number of studies that have been drawn from markedly different patient populations. Furthermore, all studies are at a high risk of selection bias given the non-randomized and non-blinded nature of the study. In particular, many studies have stark differences between the bioprosthetic and mechanical valve patient populations making accurate analysis challenging. Unfortunately, there is insufficient published data to attempt to correct for some of these differences.

Conclusions

This meta-analysis did not identify any evidence of a significant survival advantage, reduction in valve reinfection or reoperation rate for any particular conventional prosthesis choice in the treatment of patients with infective endocarditis. It is clear that evidence guiding our decision making in this area is currently limited and is an area that would benefit from future research. The choice of prosthesis should continue to be made with the conscientious consideration to the patients age, co-morbidities and patient preferences.

Expert opinion: choice of prosthetic valve in infective endocarditis

Zegri-Reiriz, Tauron

In patients with infective endocarditis and indication for surgery, the general recommendations regarding the choice of prosthesis (biological or mechanical) do not differ with respect to the general population. Both have shown similar rates of post-operative mortality and reinfection.

The European guidelines recommend mechanical prostheses for those patients under 60 years in the aortic position and for those under 65 years in the mitral position (12). According to the American Heart Association, mechanical prostheses are recommended in patients <50 years and biological prostheses in <70 years (13).

However, it should be noted that these patients deserve some important considerations. In many cases, they have a serious clinical condition which is the result of uncontrolled infection, acute heart failure or septic embolism.

In this scenario, the risk of bleeding and the implications of anticoagulation in the short and medium term should be carefully evaluated. Thus, some young patients with a

high risk of bleeding could benefit from a bioprosthesis if carefully selected.

In worse cases, there is a risk of severe coagulopathy, hepatopathy or in an embolic stroke, due to the risk of hemorrhagic transformation. In this population of critically ill patients who may need mechanical circulatory support, a bioprosthesis would be preferable. On the other hand, a sutureless bioprosthesis could be a reasonable alternative for surgical treatment of prosthetic aortic valve endocarditis with significant involvement of the aortic annulus (38).

These decisions should be individualized in each case, balancing the risk of bleeding due to anticoagulation against the possible benefit of implanting a mechanical prosthesis in young patients.

Expert opinion: choice of valve prosthesis in patients undergoing valve surgery for infective endocarditis

Pettersson, Coselli

In this issue of the *Annals of Cardiothoracic Surgery*, Flynn and coworkers present a systematic review and meta-analysis comparing outcomes after surgery for left sided infective endocarditis (IE) using mechanical prosthesis versus bioprosthesis. Both choices produced very similar outcomes with slight trends favoring the mechanical valve with regard to freedom from reoperation and risk of recurrent endocarditis; but a possible benefit for overall survival with bioprosthetic valve replacement.

Current AATS Guidelines (39) for surgical treatment of endocarditis makes recommendations related to the choice of replacement valve: valve repair is preferable when possible. When replacement is required and the disease is confined to cusps, leaflets, or valve prosthesis, choice of replacement valve—mechanical or tissue prosthesis should be based on usual criteria.

With regard to risk of recurrent endocarditis the guidelines say: “For patients requiring valve replacement, there is little evidence that risk of recurrent infection is different between mechanical and tissue prostheses.” Also, in Flynn and coworkers’ paper there is no significant difference in risk of recurrent endocarditis, however, in recent registry studies from Sweden and Denmark when comparing incidence of endocarditis following valve replacement of non-infected pathologies, bioprostheses were associated with higher risk of endocarditis than mechanical prostheses (40,41).

The main limitation with Flynn and coworkers’ meta-analysis is that it doesn’t take into consideration all the nuances, reasons for one choice over the other, information about the pathology and complications (e.g., stroke or intracranial bleed), and other factors guiding the surgeons’ and patients’ choice of replacement valve. It is still true that “Patients with IE are often very sick and have suffered strokes, so using allografts or bioprosthetic valves simplifies management and avoids postoperative anticoagulation, lowering the risk of hemorrhagic conversion of strokes and other bleeding complications.” When the disease is invasive and the annulus destroyed, extensive debridement, reconstruction and usually root replacement are required. In this situation many surgeons, including us believe that an allograft is a better choice than a prosthetic valve conduit, be it mechanical or bioprosthetic valve.

Flynn and coworkers should be thanked and congratulated for confirming that current guidelines and practices for choice of prosthetic type for patients with infective endocarditis are adequate and produce very comparable outcomes with both mechanical and bioprosthetic valves.

Expert opinion: individuality²

Misfeld

In the current issue of the *Annals of Cardiothoracic Surgery*, Flynn *et al.* present a systematic review and meta-analysis of eleven studies, published between 1989 and 2016, about outcomes of mechanical versus bioprosthetic valve substitutes in infective endocarditis (IE). The analysis did not identify a significant survival advantage, reduction in valve reinfection or reoperation with one of the valve types.

There is no clear agreement on the best prosthetic valve choice, neither for non-infective nor for IE. This is represented by different recommendations of the European and American guidelines (39,42). Despite current recommendations, mechanical valves compared to bioprosthetic valve substitutes are probably used in the ratio 1:10, with more tissue valves being implanted even in younger patients (43,44). Bioprostheses are increasingly implanted not only because anticoagulation and valve noise with mechanical valves remaining important aspects for most patients, but also because interventional options, such as valve-in-valve procedures in tissue valves, have become an attractive second step when biological valves fail (43).

Whatever guidelines recommend, neither the patient,

nor the surgeon can foresee, what kind of impact the choice of a valve (biological valve: risk of reoperation, but patient will “forget” about having a valve substitute in between; mechanical valve: valve noise and regular reminder because of regular anticoagulation checks) has for the patient during daily life.

Therefore, it remains an individual decision, which valve type should be used, independent of the underlying disease (valve surgery with/without IE). This individual decision is not only a patient related aspect, but also surgeon related. Patients with IE, especially if complicated by cerebral embolism, will most likely have a tissue valve, despite patient age. On the other hand, surgeon related factors, i.e., not being experienced with the use of allografts, may also influence the type of valve used.

As none of the valve substitutes available for the treatment of IE has been shown to have a clear advantage over the others, surgeons should use the valve substitute they are most familiar and experienced with under the background of patient related factors.

Expert opinion: prosthetic valve endocarditis, still a dilemma

Antunes

In the paper published in this issue of the *Annals of Cardiothoracic Surgery*, Flynn *et al.* analyse the incidence of prosthetic valve endocarditis (PVE) in patients with biological and mechanical prostheses. They performed a meta-analysis of studies reporting the outcomes of patients undergoing valve replacement for infective endocarditis (IE), both native and prosthetic. The authors found that at 10-year follow-up, there were no differences in overall survival, and freedom from valve reinfection and reoperation between the two types of prostheses. However, the series included in the study are mostly historical and include both native and prosthetic endocarditis, and many types of prostheses of both groups.

The results are not surprising, but I cannot completely agree with the conclusion derived by the authors that “the presence of infective endocarditis alone should not influence the decision of which type of valve prosthesis that should be implanted”. Although the data is not provided by the authors, the fact is that there was a significant incidence of PVE after valve replacement in IE. There is no consensus with regard to favouring either bioprosthetic or mechanical valve implantation in the setting of IE. Hence, it is not

a question of which of these two prostheses but if there are alternatives; and, in my view, there is an alternative, especially useful in the aortic position with significant disruption of the annulus—the homograft.

Several works appear to indicate that homografts may be more resistant to infection than either biological or mechanical prostheses (45,46). My own experience also appears to demonstrate that superiority (47). However, others have shown no difference in the rates of re-infection (23). Implantation of homografts may be technically more demanding and, similarly to bioprostheses, they have the problem of biodegradation. In any case, the pliability of the homograft facilitates the treatment of the extensively disrupted aortic annuli often seen in PVE (48). Some authors have also proposed the use of stentless xenografts for the same indications (49). Finally, the Ross operation has also been advocated in these situations (50), but it is a surgically challenging procedure to be used in an already complex anatomical situation.

Expert opinion: do not radically position yourself in the surgery for infective Endocarditis

Mestres, Quintana

In this issue of the *Annals of Cardiothoracic Surgery*, Flynn *et al.* used meta-analysis to understand surgical outcomes comparing mechanical and bioprosthetic valve replacement in infective endocarditis (IE). They found no significant difference in overall survival, reoperation and valve reinfection rates between patients with mechanical or bioprosthetic valves.

The decision of what type of valve to use in the setting of IE has been discussed for over sixty years since the first surgical replacements (51,52), and remains a challenging and controversial topic. In addition to considerations of expected anticoagulation-related events and device reoperations (53), IE is another factor to consider when choosing a prosthesis as this population has higher surgical risk. Current IE guidelines do not support one device over another due to the lack of evidence on superiority (54).

In their analysis, the authors found substantial heterogeneity in the studies and were unable to correct for severity of IE, urgency status, concomitant operations or co-morbidities. There are additional limitations in the Flynn *et al.* meta-analysis (51). It was impossible to separate by valve positions and type of device. It is difficult to understand if under the term “bioprostheses”, stentless

valves and homografts were included. Most homograft implants in IE correspond to the aortic position. Patients implanted with a homograft are at higher risk than those implanted with stented bioprosthesis as they have extravalvular spread of the infection, meaning the infection is more destructive, requiring extensive reconstruction, especially in reoperations for prosthetic valve IE (55,56).

Considering the complexity of IE and the process of choosing a valve, and difficulty in producing high quality clinical data for this surgical population, we should not adopt radical positions, with or without practice recommendations. Guidelines are recommendations, not the law, and all options must be carefully individualized. The treating team makes considered decisions based on preoperative condition, comorbidities, age among other variables, integrating this and acting according to experience, expertise and local resources.

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Footnote

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

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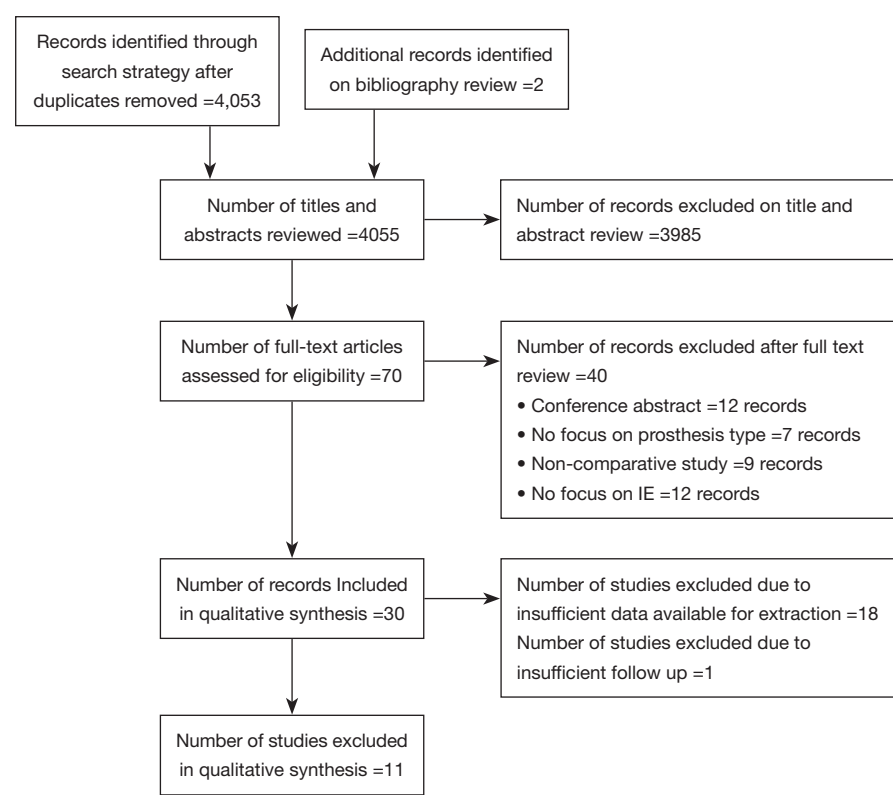


Figure S1 Study selection. PRISMA flow diagram detailing the systematic review to identify studies reporting the outcomes of bioprosthetic valve replacement compared to mechanical valve replacement for infective endocarditis.

	Random sequence generation (selection bias)	Allocation concealment (selection bias)	Blinding of participants and personnel (performance bias)	Blinding of outcome assessment (detection bias)	Incomplete outcome data (attrition bias)	Selective reporting (reporting bias)	Other bias
Delahaye 2015	+	+	+	+	+	+	+
Fedoruk 2009	+	+	+	+	+	?	?
Greason 2014	+	+	+	+	+	+	
Jassar 2012	+	+	+	+	+	+	+
Kim 2016	+	+	+	+	?	?	+
Leither 2013	+	+	+	+	+	?	?
Lytle 1996	+	+	+	+	?	?	+
Moon 2001	+	+	+	+	+	+	?
Musci 2010	+	+	+	+	+	+	?
Nguyen 2010	+	+	+	+	+	?	?
Reul 1989	+	+	+	+	+	?	

Figure S2 Risk of bias table. Thorough assessment of biases for all studies (red = high risk, yellow = uncertain risk, green = low risk).

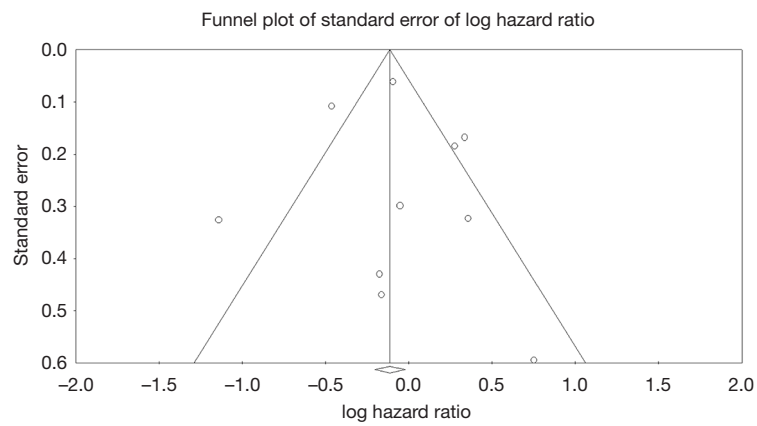


Figure S3 Funnel plot of standard error of log hazard ratio for overall survival.

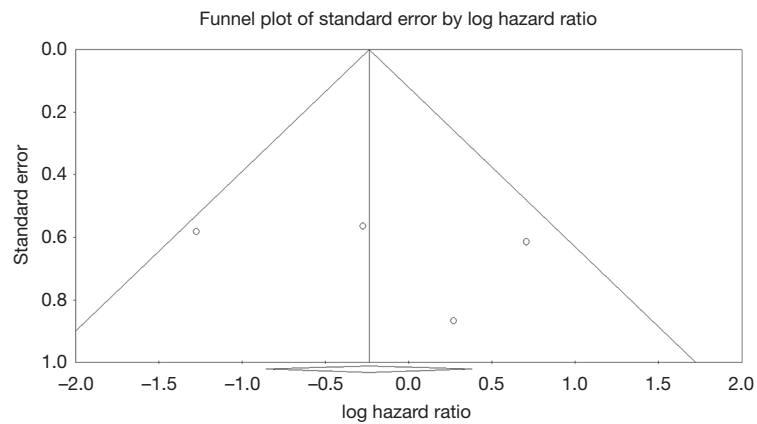


Figure S4 Funnel plot of standard error of log hazard ratio for reoperation rates.

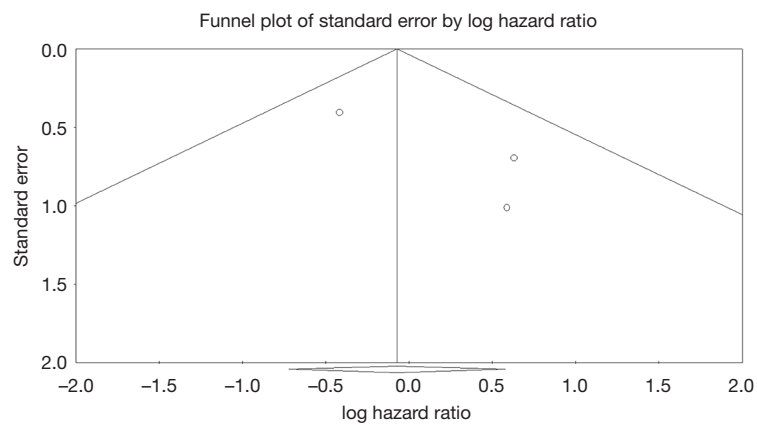


Figure S5 Funnel plot of standard error of log hazard ratio for valve reinfection rates.