REVIEW

Current controversies in infective endocarditis [version 1; peer review: 3 approved]

Thomas J. Cahill¹, Bernard D. Prendergast²

¹Department of Cardiology, Oxford University Hospitals, Oxford, UK

²Department of Cardiology, St Thomas's Hospital, London, UK

V1 First published: 18 Nov 2015, 4(F1000 Faculty Rev):1287
<https://doi.org/10.12688/f1000research.6949.1>

Latest published: 18 Nov 2015, 4(F1000 Faculty Rev):1287
<https://doi.org/10.12688/f1000research.6949.1>

Abstract

Infective endocarditis is a life-threatening disease caused by a focus of infection within the heart. For clinicians and scientists, it has been a moving target that has an evolving microbiology and a changing patient demographic. In the absence of an extensive evidence base to guide clinical practice, controversies abound. Here, we review three main areas of uncertainty: first, in prevention of infective endocarditis, including the role of antibiotic prophylaxis and strategies to reduce health care-associated bacteraemia; second, in diagnosis, specifically the use of multimodality imaging; third, we discuss the optimal timing of surgical intervention and the challenges posed by increasing rates of cardiac device infection.

Keywords

Infective endocarditis , Bacteraemia , Antibiotic prophylaxis , Transcatheter valves , Multimodality imaging , Cardiac device infection

Open Peer Review

Approval Status

	1	2	3
version 1 18 Nov 2015			

Faculty Reviews are review articles written by the prestigious Members of **Faculty Opinions**. The articles are commissioned and peer reviewed before publication to ensure that the final, published version is comprehensive and accessible. The reviewers who approved the final version are listed with their names and affiliations.

1. **Ralph Corey**, Duke University, Durham, USA
2. **Xavier Duval**, Hôpital Bichat-Claude Bernard, Paris, France
3. **Ron Blankstein**, Brigham and Women's Hospital, Boston, USA

Any comments on the article can be found at the end of the article.

Corresponding author: Bernard D. Prendergast (Bernard.Prendergast@gstt.nhs.uk)

Competing interests: The authors declare that they have no competing interests.

Grant information: The author(s) declared that no grants were involved in supporting this work.

Copyright: © 2015 Cahill TJ and Prendergast BD. This is an open access article distributed under the terms of the [Creative Commons Attribution License](#), which permits unrestricted use, distribution, and reproduction in any medium, provided the original work is properly cited.

How to cite this article: Cahill TJ and Prendergast BD. **Current controversies in infective endocarditis [version 1; peer review: 3 approved]** F1000Research 2015, 4(F1000 Faculty Rev):1287 <https://doi.org/10.12688/f1000research.6949.1>

First published: 18 Nov 2015, 4(F1000 Faculty Rev):1287 <https://doi.org/10.12688/f1000research.6949.1>

Introduction

In 1885, William Osler described ‘malignant’ endocarditis as an infection on the scarred heart valves of young adults with rheumatic heart disease¹. At the time, infective endocarditis (IE) was caused mainly by microorganisms originating in the oral cavity (oral streptococci). In 2015 in the developed world, IE looks dramatically different². Rheumatic fever is now rare, and in 25% of cases the causative infection is health care-acquired^{3,4}. With the exception of intravenous drug users and those with congenital heart disease, patients typically are elderly. The focus of infection is frequently on prosthetic material within the heart: for example, cardiac devices (pacemakers, implantable cardioverter defibrillators and cardiac resynchronisation therapy) or prosthetic heart valves. In parallel, the bacteria underlying IE have changed: highly destructive staphylococci have now overtaken oral streptococci as the most common cause^{4,5}. With an incidence of 3 to 10 per 100,000, IE is rare but carries an in-hospital mortality of 20%². Improving outcomes for patients with this complex and heterogeneous disease is challenging. In this review, we provide a focused update on current controversies in disease prevention, diagnosis and management.

Prevention of infective endocarditis

Bacteria enter the bloodstream from the mouth in the course of daily life, and poor dental hygiene has long been recognised as a risk factor for IE⁶. The significance of bacteraemia that occurs with dental extraction is still debated, as is the role of oral antibiotic prophylaxis to prevent it. In hospital, bacteraemia occurs as a complication of invasive procedures and indwelling venous catheters. Bacterial adherence, inflammation of the cardiac endothelium and thrombus deposition lead to formation of an infected ‘vegetation’⁷.

Oral antibiotic prophylaxis

Poor oral hygiene is a risk factor for bacteraemia and subsequent IE⁸. In addition to maintaining good hygiene, oral antibiotic prophylaxis has traditionally been prescribed for those at risk of IE prior to dental and surgical procedures. In 2008, the UK National Institute for Health and Care Excellence recommended against further use of prophylaxis, considering the potential hazards of widespread antibiotic use to outweigh the individual risks⁹. The European Society of Cardiology (ESC) and American College of Cardiology/American Heart Association continued to recommend prophylaxis but only for patients at highest risk: those with previous IE, prosthetic valves and cyanotic congenital heart disease¹⁰.

The effect of restricting antibiotic prophylaxis has now been examined in several studies^{11–13}. Recently, a UK study found that over a 5-year period from 2008 there had been a small but significant increase in the (already-rising) number of IE cases alongside a fall in prescriptions for prophylaxis¹². These findings have been highly controversial. A causal link between IE cases and the withdrawal of systematic prophylaxis cannot be established by observational data, and the findings may be confounded by increased rates of bacteraemia or a growing at-risk population. Importantly, microbiological data were not available, so it is unclear whether the additional cases were caused by oral streptococci (the target of prophylaxis). In the absence of a randomised controlled trial (RCT)—which faces

logistical, funding and ethical barriers—this study may tip the balance back toward prophylaxis for high-risk groups.

Reducing health care-associated bacteraemia

At least a quarter of cases of IE are now acquired in the health-care setting¹⁴. Mortality for health care-acquired IE exceeds 40% and this is due to the susceptible population, often with multiple comorbidities, and higher rates of *Staphylococcus aureus* infection¹⁵. Health care-associated bacteraemia, the upstream cause of IE, is the target of several preventative strategies. Basic hand hygiene, aseptic technique, sterile barrier clothing and avoidance of the femoral route for venous catheters are all effective at reducing bacteraemia, but getting staff to adhere to best practice can be challenging^{16,17}. ‘Practice-changing’ approaches, such as the use of a checklist to improve adherence to sterile technique, are powerful interventions which have been shown to reduce catheter-related infections^{18,19}.

Infection during the implantation of cardiac devices is an important and preventable cause of IE and is reduced by use of perioperative antibiotics²⁰. Vaccination against *S. aureus*, which directly damages the cardiac endothelium to cause IE, is an attractive theoretical strategy²¹. Unfortunately, a number of phase II and III trials have shown negative results, and the vaccine was associated with increased mortality in a 2012 study in patients undergoing cardiotoracic surgery²². Other approaches in preclinical studies include novel materials designed to prevent bacterial adherence and agents that target bacterial biofilm production^{23,24}.

Diagnosis of infective endocarditis

Diagnosis of IE requires evaluation of clinical presentation, microbiology results and cardiac imaging². Reaching a definitive diagnosis can be challenging, particularly in suspected prosthetic valve IE and cardiac device infection (CDI), in which findings can be non-specific. The modified Duke criteria are useful for defining a positive diagnosis but were not intended as a clinical tool and cannot be rigidly applied to the individual patient²⁵. Strategies that facilitate earlier diagnosis, risk stratification and therapy are key to improving survival.

Multimodality imaging for diagnosis of infective endocarditis

Transthoracic and transoesophageal echocardiography (TOE) are the mainstay of cardiac imaging for IE—for diagnosis, detection of complications and follow-up (Figure 1A). Even after TOE, however, diagnosis is inconclusive in 10% to 20% of patients because of limited resolution or image artefacts²⁶, and adjunctive imaging techniques are showing promise in this group.

Cardiac computed tomography (CT) (Figure 1B) is emerging as a valuable modality for detecting and defining IE complications in the aortic root (for example, formation of abscess/pseudoaneurysm)^{27,28}. Identification of a paravalvular lesion (abscess, pseudoaneurysm or fistula) on cardiac CT is now a major criterion for the diagnosis of IE in the newly updated 2015 ESC IE guidelines²⁹. CT is less sensitive than transoesophageal echocardiography at detecting valvular perforations and may miss small vegetations, and interpretation can be challenging in the early postoperative period. Three-dimensional TOE can assist with detection of leaflet perforation and define the anatomy of valvular dehiscence (Figure 1C)^{30,31}.

Figure 1. Multimodality imaging in diagnosis and detection of complications in infective endocarditis (IE). (A) Echocardiography remains the core imaging modality in IE. Here, a vegetation (arrow) is visualised on the aortic valve by transoesophageal echocardiography. (B) Computed tomography (CT) is excellent at defining the anatomical extent of complex endocarditis. A large paravalvular abscess (asterisk) can be seen complicating a case of prosthetic valve IE. (C) Three-dimensional transoesophageal echocardiography provides a reconstructed view of the valve and here demonstrates dehiscence of a prosthetic mitral valve (arrow), an indication for surgical intervention. (D) Positron emission tomography/CT has shown value for diagnosis of prosthetic valve IE and cardiac device infection. A focus of fludeoxyglucose- ^{18}F (^{18}F -FDG) uptake (arrow) can be seen at the site of a prosthetic aortic valve, separate from the myocardium, consistent with prosthetic valve IE. Adapted from Teoh *et al.*³⁴.

^{18}F -FDG-PET/CT (fludeoxyglucose- ^{18}F /positron emission tomography/CT)—which shows uptake of radiolabelled glucose by metabolically active tissues—can visualise the infected vegetations of IE and has been successfully used in patients with suspected prosthetic valve IE or CDI, in which interpretation of TOE can be challenging (Figure 1D)^{32–34}. In a recently published study of 92 patients with suspected prosthetic valve IE or CDI, hybrid PET-CT enabled reclassification of 90% of patients (45 out of 50) with *possible* IE by Duke criteria and provided a conclusive diagnosis (definite/rejected) in 95% of cases³⁵. Addition of PET-CT to the Duke criteria raised the diagnostic sensitivity from 52% to 91%. As such, positive PET-CT (or single-photon emission computed tomography-CT) signal at the site of a prosthetic valve (at least 3 months after implant) has been included as a new major criterion for diagnosis of prosthetic valve IE in the 2015 ESC guidelines²⁹.

Beyond the heart, CT, magnetic resonance imaging (MRI) and PET imaging are facilitating early detection of embolism³⁶. Systematic MRI of the brain detects abnormalities in up to 80%, and imaging evidence of embolism is now a minor diagnostic criterion in the 2015 ESC guidelines³⁷. PET-CT imaging is useful for detection of

peripheral vascular complications such as abscess, mycotic aneurysm and emboli³⁵. Given the range of imaging options now available, the next challenge is to define the optimal imaging strategy for specific patient subsets.

Management of infective endocarditis

The infective endocarditis team

Clinical care of patients with IE is complex and is reviewed in detail elsewhere². Management revolves around prolonged parenteral antibiotic therapy (typically 4 to 6 weeks in duration), surgical intervention for those at high risk or not responding to antibiotics, and surveillance for complications. Input is required from a wide range of specialists, including cardiologists, cardiothoracic surgeons, microbiologists and specialists in cardiovascular imaging and infectious diseases. Coordinating care can be a logistical challenge leading to delayed definitive management. Formation of a dedicated IE multidisciplinary team (MDT) is a simple strategy to improve clinical care. In a 2013 study from Italy, introduction of an IE MDT reduced in-hospital mortality from 28% to 13%³⁸. Recent UK guidelines advocate the formation of an IE team in every major centre³⁹.

Early surgery

Surgery is currently performed in 40% to 50% of patients with IE⁴⁰. There are three main indications: valve dysfunction causing heart failure, uncontrolled infection, and prevention of embolism. Although seemingly clear-cut, the appropriateness and timing of surgery for an individual patient sometimes carry considerable uncertainty. In 2012, a landmark RCT compared early surgery (within 48 hours) with conventional treatment in stable patients with native valve IE, echocardiographically severe valvular regurgitation and large vegetations⁴¹. The cohort was young (mean age of 47 years), was infected mainly by oral streptococci and had little comorbidity. The 'early surgery' group had a significant reduction in a composite endpoint of in-hospital death or embolism, driven by a reduction in the rate of embolism. This trial has led to a trend toward earlier surgery, but enthusiasm has been dampened by concerns that early surgery does not have the same benefits in the older, frailer IE population in much of the industrialized world. In prosthetic valve IE, retrospective studies looking at early surgery in patients have failed to find a benefit^{42,43}.

Surgery to prevent embolism remains the most controversial indication, as the embolic risk for a specific individual is often difficult to predict. In general, surgery should be considered for those in the highest-risk groups: those with very large vegetations (>30 mm) or persistent large vegetations (>10 mm) after an embolic event and despite antibiotics²⁹. Mobile vegetations, staphylococcal infection and mitral valve location are also associated with increased embolic risk⁴⁴. The risk is highest in the first 2 weeks of antibiotic therapy and so intervention for prevention of embolism must be performed early to optimise the risk-to-benefit ratio of major surgery.

Infective endocarditis on cardiac devices and transcatheter valves

The indications and implantation rates for cardiac devices have expanded dramatically in the last decade. Cardiac resynchronisation therapy is used widely for patients with advanced heart failure, and implantable (intracardiac) cardioverter defibrillators are commonly inserted for prevention of ventricular arrhythmias⁴⁵. Similarly, in the last 15 years, transcatheter aortic valve implantation (TAVI) has gone from concept to clinical reality, and over 100,000 valves have now been implanted worldwide⁴⁶. Cardiac device infection (CDI) currently accounts for roughly 10% of IE, and as use of cardiac devices and valve prostheses increases further, cardiologists should expect increasing rates of IE³.

CDI affects 1% to 2% of patients in the first 5 years following device implantation and may involve the pocket housing the generator box, the leads or the endocardial surface²⁰. There are limited studies to direct clinical management of CDI, but UK guidelines outlining current best practice and recommendations for research have recently been published²⁰. Unless infection is superficial and limited to the healing wound, device extraction is required, and this

can usually be achieved percutaneously but carries a small risk of mortality. The optimal duration of antibiotic therapy and risks of further infection after device re-implantation are unknown.

Reports of small cohorts of patients with IE following TAVI are starting to emerge^{47–49}. A recently published multicentre registry identified a cohort of 53 patients with IE post-TAVI, representing a frequency of 0.5% at 1 year⁴⁸. The mean time to IE from TAVI procedure was 6 months. Interestingly, use of the self-expanding CoreValve system (Medtronic, Dublin, Ireland) was independently associated with increased risk of IE (hazard ratio 3.12, 95% confidence interval 1.37 to 7.14, $P = 0.007$). In-hospital mortality for this group was 47%, reflecting the high rates of staphylococcal infection and frailty of the study population. Further studies are required to better define the incidence, risk factors and clinical outcomes of IE following TAVI.

Conclusions

IE is a rare and multifaceted disease whose heterogeneity is a barrier to well-powered research trials. As a consequence, much of the evidence base is derived from observational studies. Despite ongoing uncertainty, some conclusions can be drawn. Given the increasing use of intracardiac prostheses and devices, an ongoing focus on IE prevention strategies is warranted. Reducing time to diagnosis and definitive management requires set-up of IE clinical teams and full use of multimodality imaging alongside echocardiography. The importance of abnormal CT and hybrid PET-CT imaging is now reflected in the 2015 ESC guidelines on IE, in which abnormal imaging findings are novel diagnostic criteria. Finally, research networks should move toward a focus on multicentre trials, for example to address uncertainties in the timing of surgery and management of CDI.

Abbreviations

CDI, cardiac device infection; CT, computed tomography; ESC, European Society of Cardiology; IE, infective endocarditis; MDT, multidisciplinary team; MRI, magnetic resonance imaging; PET, positron emission tomography; RCT, randomised controlled trial; TAVI, transcatheter aortic valve implantation; TOE, transoesophageal echocardiography.

Competing interests

The authors declare that they have no competing interests.

Grant information

The author(s) declared that no grants were involved in supporting this work.

Acknowledgements

The authors thank Jim Newton, Andrew Kelion and Nik Sabharwal for assistance with Figure 1.

References

1. Osler W: **The Gulstonian Lectures, on Malignant Endocarditis.** *Br Med J.* 1885; 1(1264): 577–9.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
2. Cahill TJ, Prendergast BD: **Infective endocarditis.** *Lancet.* 2015; pii: S0140-6736(15)00067-7.
[PubMed Abstract](#) | [Publisher Full Text](#)
3. **F** Murdoch DR, Corey GR, Hoen B, *et al.*: **Clinical presentation, etiology, and outcome of infective endocarditis in the 21st century: the International Collaboration on Endocarditis-Pro prospective Cohort Study.** *Arch Intern Med.* 2009; 169(5): 463–73.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#) | **F1000 Recommendation**
4. **F** Selton-Suty C, Célard M, Le Moing V, *et al.*: **Preeminence of *Staphylococcus aureus* in infective endocarditis: a 1-year population-based survey.** *Clin Infect Dis.* 2012; 54(9): 1230–9.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
5. Fowler VG, Miro JM, Hoen B, *et al.*: ***Staphylococcus aureus* endocarditis: a consequence of medical progress.** *JAMA.* 2005; 293(24): 3012–21.
[PubMed Abstract](#) | [Publisher Full Text](#)
6. **F** Lockhart PB, Brennan MT, Sasser HC, *et al.*: **Bacteremia associated with toothbrushing and dental extraction.** *Circulation.* 2008; 117(24): 3118–25.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#) | **F1000 Recommendation**
7. Werdan K, Dietz S, Löffler B, *et al.*: **Mechanisms of infective endocarditis: pathogen-host interaction and risk states.** *Nat Rev Cardiol.* 2014; 11(1): 35–50.
[PubMed Abstract](#) | [Publisher Full Text](#)
8. Lockhart PB, Brennan MT, Thornhill M, *et al.*: **Poor oral hygiene as a risk factor for infective endocarditis-related bacteremia.** *J Am Dent Assoc.* 2009; 140(10): 1238–44.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
9. Richey R, Wray D, Stokes T: **Prophylaxis against infective endocarditis: summary of NICE guidance.** *BMJ.* 2008; 336(7647): 770–1.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
10. Habib G, Hoen B, Tornos P, *et al.*: **Guidelines on the prevention, diagnosis, and treatment of infective endocarditis (new version 2009): the Task Force on the Prevention, Diagnosis, and Treatment of Infective Endocarditis of the European Society of Cardiology (ESC). Endorsed by the European Society of Clinical Microbiology and Infectious Diseases (ESCMID) and the International Society of Chemotherapy (ISC) for Infection and Cancer.** *Eur Heart J.* 2009; 30(19): 2369–413.
[PubMed Abstract](#) | [Publisher Full Text](#)
11. **F** Pasquali SK, He X, Mohamad Z, *et al.*: **Trends in endocarditis hospitalizations at US children's hospitals: impact of the 2007 American Heart Association Antibiotic Prophylaxis Guidelines.** *Am Heart J.* 2012; 163(5): 894–9.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#) | **F1000 Recommendation**
12. Dayer MJ, Jones S, Prendergast B, *et al.*: **Incidence of infective endocarditis in England, 2000–13: a secular trend, interrupted time-series analysis.** *Lancet.* 2015; 385(9974): 1219–28.
[PubMed Abstract](#) | [Publisher Full Text](#)
13. **F** Pant S, Patel NJ, Deshmukh 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.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
14. **F** Fernández-Hidalgo N, Almirante B, Tornos P, *et al.*: **Contemporary epidemiology and prognosis of health care-associated infective endocarditis.** *Clin Infect Dis.* 2008; 47(10): 1287–97.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
15. **F** Lomas JM, Martínez-Marcos FJ, Plata A, *et al.*: **Healthcare-associated infective endocarditis: an undesirable effect of healthcare universalization.** *Clin Microbiol Infect.* 2010; 16(11): 1683–90.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
16. O'Grady NP, Alexander M, Burns LA, *et al.*: **Guidelines for the prevention of intravascular catheter-related infections.** *Clin Infect Dis.* 2011; 52(9): e162–93.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
17. Pratt RJ, Pellowe CM, Wilson JA, *et al.*: **Guidelines for the prevention of healthcare-associated infections in NHS hospitals in England.** *J Hosp Infect.* 2007; 65(Suppl 1): S1–64.
[PubMed Abstract](#) | [Publisher Full Text](#)
18. **F** Pronovost P, Needham D, Berenholtz S, *et al.*: **An intervention to decrease catheter-related bloodstream infections in the ICU.** *N Engl J Med.* 2006; 355(26): 2725–32.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
19. **F** Blot K, Bergs J, Vogelaers D, *et al.*: **Prevention of central line-associated bloodstream infections through quality improvement interventions: a systematic review and meta-analysis.** *Clin Infect Dis.* 2014; 59(1): 96–105.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#) | **F1000 Recommendation**
20. Sandoe JA, Barlow G, Chambers JB, *et al.*: **Guidelines for the diagnosis, prevention and management of implantable cardiac electronic device infection. Report of a joint Working Party project on behalf of the British Society for Antimicrobial Chemotherapy (BSAC, host organization), British Heart Rhythm Society (BHRS), British Cardiovascular Society (BCS), British Heart Valve Society (BHVS) and British Society for Echocardiography (BSE).** *J Antimicrob Chemother.* 2015; 70(2): 325–59.
[PubMed Abstract](#) | [Publisher Full Text](#)
21. **F** Fowler VG Jr, Proctor RA: **Where does a *Staphylococcus aureus* vaccine stand?** *Clin Microbiol Infect.* 2014; 20(Suppl 5): 66–75.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#) | **F1000 Recommendation**
22. **F** Fowler VG, Allen KB, Moreira ED, *et al.*: **Effect of an investigational vaccine for preventing *Staphylococcus aureus* infections after cardiothoracic surgery: a randomized trial.** *JAMA.* 2013; 309(13): 1368–78.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
23. Beloin C, Renard S, Ghigo JM, *et al.*: **Novel approaches to combat bacterial biofilms.** *Curr Opin Pharmacol.* 2014; 18: 61–8.
[PubMed Abstract](#) | [Publisher Full Text](#)
24. Desrousseaux C, Sautou V, Descamps S, *et al.*: **Modification of the surfaces of medical devices to prevent microbial adhesion and biofilm formation.** *J Hosp Infect.* 2013; 85(2): 87–93.
[PubMed Abstract](#) | [Publisher Full Text](#)
25. Prendergast BD: **Diagnostic criteria and problems in infective endocarditis.** *Heart.* 2004; 90(6): 611–3.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#)
26. **F** Bruun NE, Habib G, Thuny F, *et al.*: **Cardiac imaging in infectious endocarditis.** *Eur Heart J.* 2014; 35(10): 624–32.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
27. **F** Feuchtnner GM, Stolzmann P, Dichtl W, *et al.*: **Multislice computed tomography in infective endocarditis: comparison with transesophageal echocardiography and intraoperative findings.** *J Am Coll Cardiol.* 2009; 53(5): 436–44.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
28. Fagman E, Perrotta S, Bech-Hanssen O, *et al.*: **ECG-gated computed tomography: a new role for patients with suspected aortic prosthetic valve endocarditis.** *Eur Radiol.* 2012; 22(11): 2407–14.
[PubMed Abstract](#) | [Publisher Full Text](#)
29. **F** Authors/Task Force Members, Habib G, Lancellotti P, *et al.*: **2015 ESC Guidelines for the management of infective endocarditis: The Task Force for the Management of Infective Endocarditis of the European Society of Cardiology (ESC) Endorsed by: European Association for Cardio-Thoracic Surgery (EACTS), the European Association of Nuclear Medicine (EANM).** *Eur Heart J.* 2015; pii: ehv319.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
30. Thompson KA, Shiota T, Tolstrup K, *et al.*: **Utility of three-dimensional transesophageal echocardiography in the diagnosis of valvular perforations.** *Am J Cardiol.* 2011; 107(1): 100–2.
[PubMed Abstract](#) | [Publisher Full Text](#)
31. Kronzon I, Sugeng L, Perk G, *et al.*: **Real-time 3-dimensional transesophageal echocardiography in the evaluation of post-operative mitral annuloplasty ring and prosthetic valve dehiscence.** *J Am Coll Cardiol.* 2009; 53(17): 1543–7.
[PubMed Abstract](#) | [Publisher Full Text](#)
32. **F** Sarrazin JF, Philippon F, Tessier M, *et al.*: **Usefulness of fluorine-18 positron emission tomography/computed tomography for identification of cardiovascular implantable electronic device infections.** *J Am Coll Cardiol.* 2012; 59(18): 1616–25.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
33. **F** Saby L, Laas O, Habib G, *et al.*: **Positron emission tomography/computed tomography for diagnosis of prosthetic valve endocarditis: increased valvular ¹⁸F-fluorodeoxyglucose uptake as a novel major criterion.** *J Am Coll Cardiol.* 2013; 61(23): 2374–82.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
34. Teoh EJ, Backhouse L, Chandrasekaran B, *et al.*: **Mycotic aneurysm of the superior mesenteric artery and other sequelae of prosthetic valve endocarditis on ¹⁸F-FDG PET/CT.** *Eur J Nucl Med Mol Imaging.* 2014; 41(10): 1993–4.
[PubMed Abstract](#) | [Publisher Full Text](#)
35. **F** Pizzi MN, Roque A, Fernández-Hidalgo N, *et al.*: **Improving the Diagnosis of Infective Endocarditis in Prosthetic Valves and Intracardiac Devices With ¹⁸F-Fluorodeoxyglucose Positron Emission Tomography/Computed Tomography Angiography: Initial Results at an Infective Endocarditis Referral Center.** *Circulation.* 2015; 132(12): 1113–26.
[PubMed Abstract](#) | [Publisher Full Text](#) | **F1000 Recommendation**
36. Vilacosta I, Graupner C, San Román JA, *et al.*: **Risk of embolization after institution of antibiotic therapy for infective endocarditis.** *J Am Coll Cardiol.* 2002; 39(9): 1489–95.
[PubMed Abstract](#) | [Publisher Full Text](#)

37. **F** Duval X, Jung B, Klein I, *et al.*: **Effect of early cerebral magnetic resonance imaging on clinical decisions in infective endocarditis: a prospective study.** *Ann Intern Med.* 2010; **152**(8): 497–504, W175.
[PubMed Abstract](#) | [Publisher Full Text](#) | [F1000 Recommendation](#)
38. **F** Chirillo F, Scotton P, Rocco F, *et al.*: **Impact of a multidisciplinary management strategy on the outcome of patients with native valve infective endocarditis.** *Am J Cardiol.* 2013; **112**(8): 1171–6.
[PubMed Abstract](#) | [Publisher Full Text](#) | [F1000 Recommendation](#)
39. Chambers J, Sandoe J, Ray S, *et al.*: **The infective endocarditis team: recommendations from an international working group.** *Heart.* 2014; **100**(7): 524–7.
[PubMed Abstract](#) | [Publisher Full Text](#)
40. Prendergast BD, Tornos P: **Surgery for infective endocarditis: who and when?** *Circulation.* 2010; **121**(9): 1141–52.
[PubMed Abstract](#) | [Publisher Full Text](#)
41. **F** Kang DH, Kim YJ, Kim SH, *et al.*: **Early surgery versus conventional treatment for infective endocarditis.** *N Engl J Med.* 2012; **366**(26): 2466–73.
[PubMed Abstract](#) | [Publisher Full Text](#) | [F1000 Recommendation](#)
42. **F** Lalani T, Chu VH, Park LP, *et al.*: **In-hospital and 1-year mortality in patients undergoing early surgery for prosthetic valve endocarditis.** *JAMA Intern Med.* 2013; **173**(16): 1495–504.
[PubMed Abstract](#) | [Publisher Full Text](#) | [F1000 Recommendation](#)
43. **F** Chirouze C, Alla F, Fowler VG Jr, *et al.*: **Impact of early valve surgery on outcome of *Staphylococcus aureus* prosthetic valve infective endocarditis: analysis in the International Collaboration of Endocarditis-Prospective Cohort Study.** *Clin Infect Dis.* 2015; **60**(5): 741–9.
[PubMed Abstract](#) | [Publisher Full Text](#) | [Free Full Text](#) | [F1000 Recommendation](#)
44. **F** Hubert S, Thuny F, Resseguier N, *et al.*: **Prediction of symptomatic embolism in infective endocarditis: construction and validation of a risk calculator in a multicenter cohort.** *J Am Coll Cardiol.* 2013; **62**(15): 1384–92.
[PubMed Abstract](#) | [Publisher Full Text](#) | [F1000 Recommendation](#)
45. The Task Force on cardiac pacing and resynchronization therapy of the European Society of Cardiology (ESC). Developed in collaboration with the European Heart Rhythm Association (EHRA), Brignole M, Esquivias G, *et al.*: **2013 ESC Guidelines on cardiac pacing and cardiac resynchronization therapy.** *Rev Esp Cardiol (Engl Ed).* 2014; **67**(1): 58.
[PubMed Abstract](#) | [Publisher Full Text](#)
46. Ruparel N, Prendergast BD: **TAVI in 2015: who, where and how?** *Heart.* 2015; **101**(17): 1422–31.
[PubMed Abstract](#) | [Publisher Full Text](#)
47. **F** Olsen NT, De Backer O, Thyregod HG, *et al.*: **Prosthetic valve endocarditis after transcatheter aortic valve implantation.** *Circ Cardiovasc Interv.* 2015; **8**(4): pii: e001939.
[PubMed Abstract](#) | [Publisher Full Text](#) | [F1000 Recommendation](#)
48. **F** Amat-Santos IJ, Messika-Zeitoun D, Eltchaninoff H, *et al.*: **Infective endocarditis after transcatheter aortic valve implantation: results from a large multicenter registry.** *Circulation.* 2015; **131**(18): 1566–74.
[PubMed Abstract](#) | [Publisher Full Text](#) | [F1000 Recommendation](#)
49. Puls M, Eiffert H, Hünlich M, *et al.*: **Prosthetic valve endocarditis after transcatheter aortic valve implantation: the incidence in a single-centre cohort and reflections on clinical, echocardiographic and prognostic features.** *EuroIntervention.* 2013; **8**(12): 1407–18.
[PubMed Abstract](#) | [Publisher Full Text](#)

Open Peer Review

Current Peer Review Status:

Editorial Note on the Review Process

Faculty Reviews are review articles written by the prestigious Members of **Faculty Opinions**. The articles are commissioned and peer reviewed before publication to ensure that the final, published version is comprehensive and accessible. The reviewers who approved the final version are listed with their names and affiliations.

The reviewers who approved this article are:

Version 1

- 1. Ron Blankstein**
Cardiovascular Division, Brigham and Women's Hospital, Boston, MA, 02115, USA
Competing Interests: No competing interests were disclosed.
- 2. Xavier Duval**
Service de Maladies Infectieuses et Tropicales, Hôpital Bichat-Claude Bernard, Paris, France
Competing Interests: No competing interests were disclosed.
- 3. Ralph Corey**
Department of Medicine, Duke University, Durham, NC, USA
Competing Interests: No competing interests were disclosed.

The benefits of publishing with F1000Research:

- Your article is published within days, with no editorial bias
- You can publish traditional articles, null/negative results, case reports, data notes and more
- The peer review process is transparent and collaborative
- Your article is indexed in PubMed after passing peer review
- Dedicated customer support at every stage

For pre-submission enquiries, contact research@f1000.com

F1000Research