Infective endocarditis & OPAT: an overview



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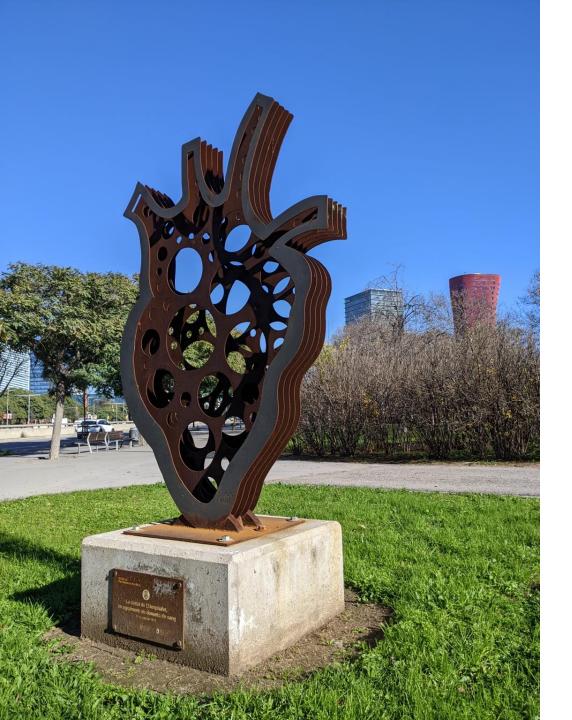
OPAT lead for St Bartholomew's Hospital

4th September 2025 – BSAC Regional OPAT workshops 2025

Disclaimer #1

Do you see IE patients in your service?





Is my OPAT service set up to deal with IE patients?

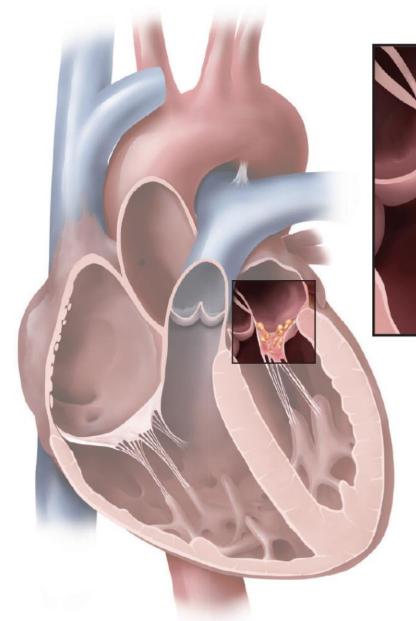
Introduction

Case

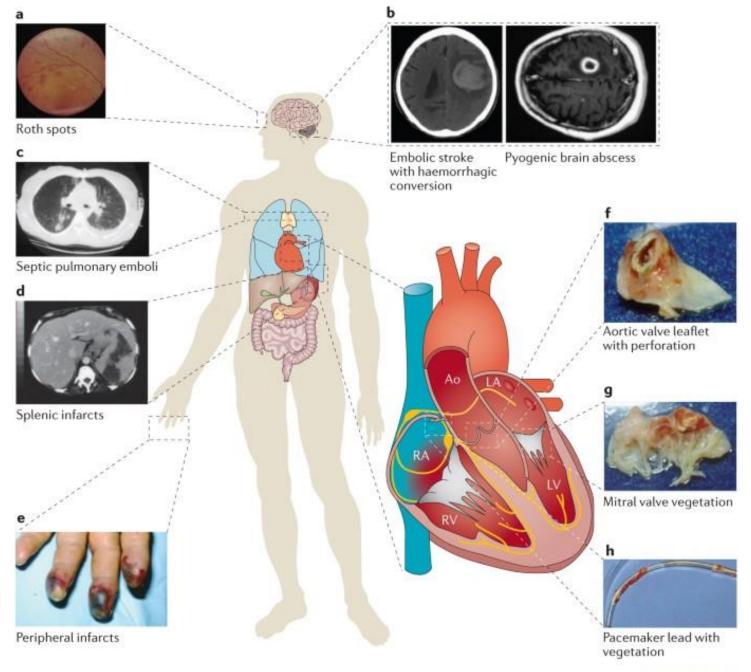
The OPAT - IE cohort at SBH 2022 - 2025

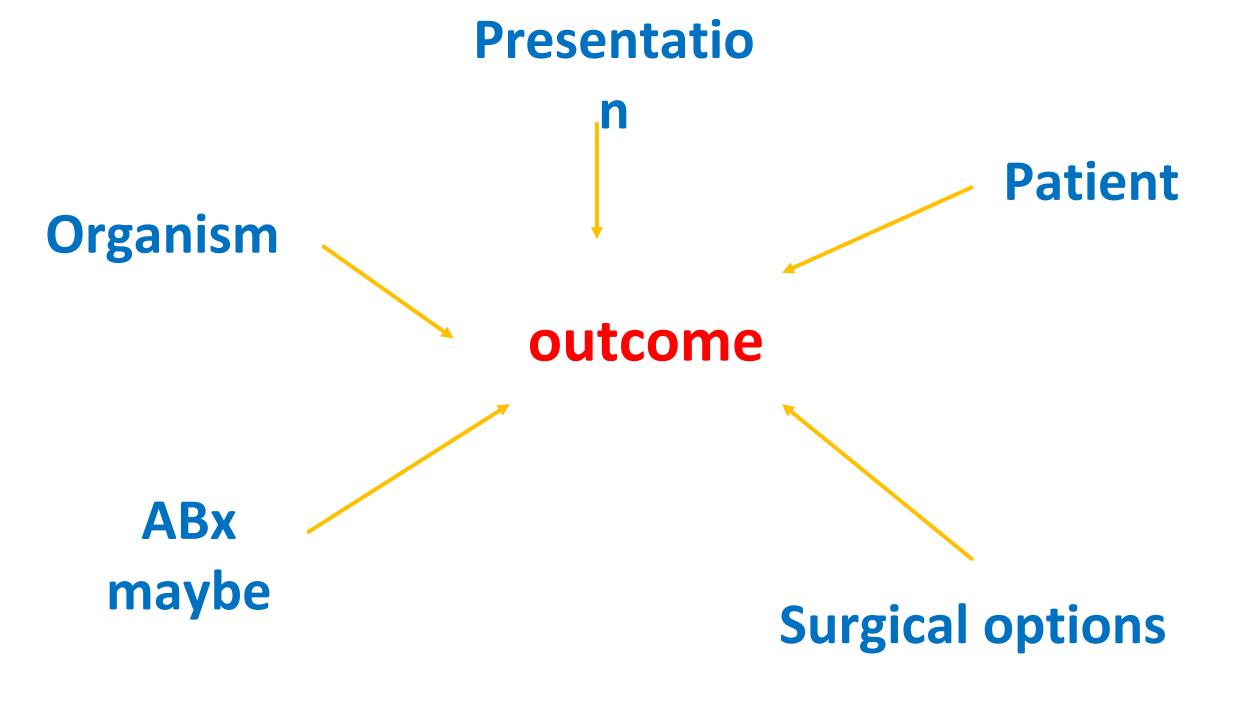
Perspectives





Damaged valve with bacteria growths





Young
Oral streptococci, low pen MIC
No embolic disease
Low surgical risk, has an operation
OR surgery is not needed

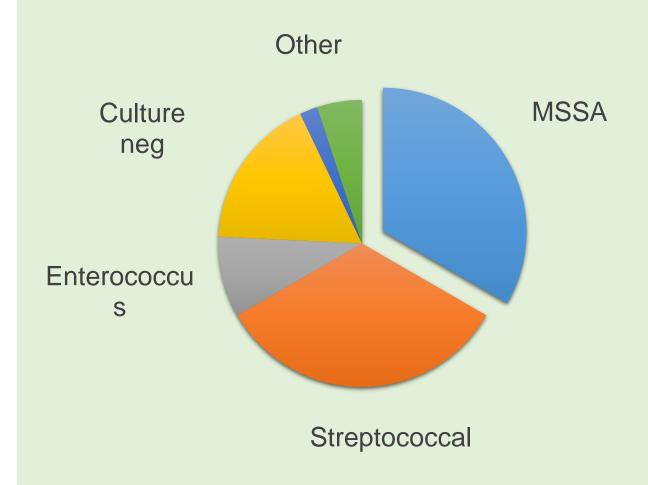
Elderly
Comorbidities
Strep pneumoniae
High surgical risk
Not operable

INFECTIVE ENDOCARDITIS at SBH

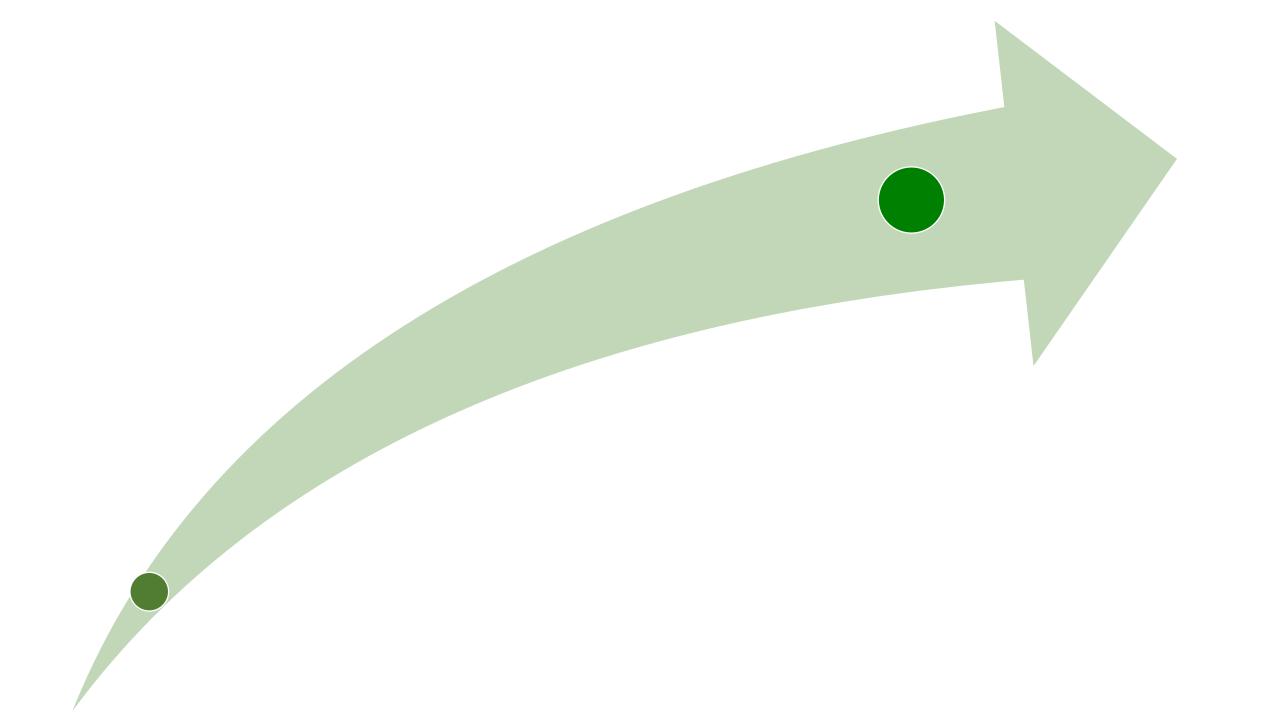
150 episodes / year LOS 27 days (17 – 43) 1 – 2 OPAT / week

OPAT since 2022

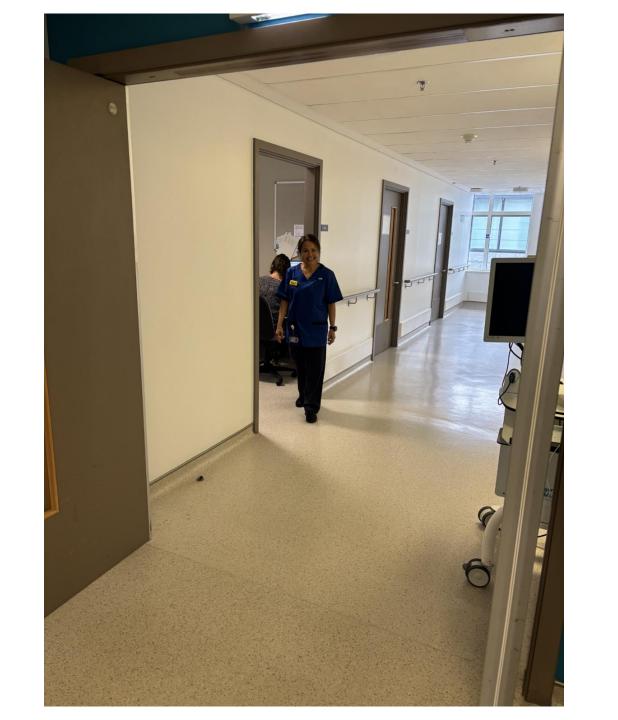
Native Valve IE: organisms



Data: courtesy of Maria Cue and Christopher Primus IE MDT at SBH







71 yo male

Jan 2024 Aortic root replacement

Feb 2024 VRE bacteraemia – short treatment (7 days)

Dapto – eosinophilia

Dec 2024 Prosthetic aortic valve IE by E. faecium (vanc S)

Large vegetation, aortic root involvement

BC = E faecium 20 and 21/12

High risk surgery but potentially operable

Patient not keen in surgery

Referred to OPAT



ABx Vancomycin – 5 days to therapeutic levels

Vanc 30/12 - 13/1 - good levels

gent x 3 weeks in total

OPAT renal adjusted teicoplanin * 3 loading doses, 12 mg/kg (800 mg)

then half dose daily (400 mg)

intermittent face-to-face review *

17/1	day1	OPAT
T//T	uayı	UPAI

17/1 teic level – low

21/1 teic level – low *

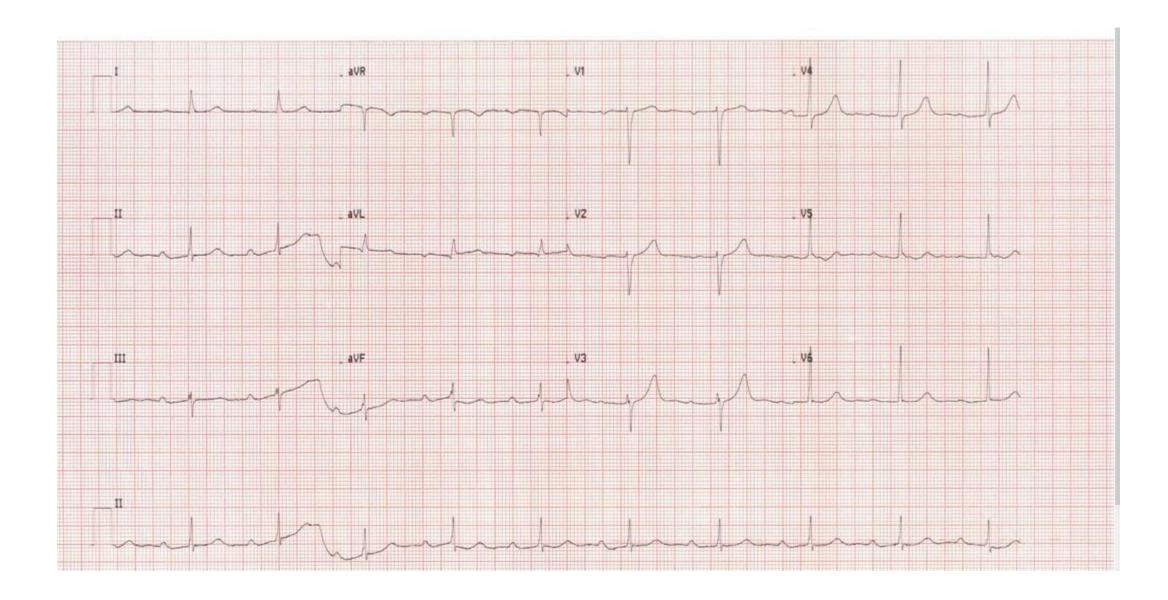
teicoplanin increased to 500 mg

29/1 new level, teic up to 800 mg awaiting levels

29/1 teic level back – still low

30/1 OPAT review





Known possible aortic abscess
Subtherapeutic teic
Not surgery
Prolonging QT



? Evolving aortic abscess



Admission (same day)
Vanc
CT + echo



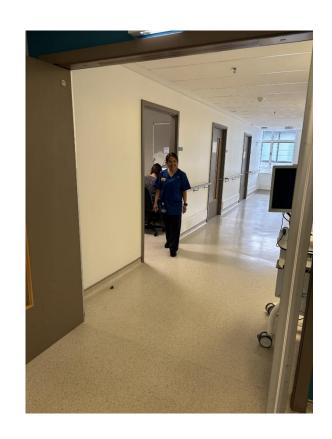
#1

Complications

Potentially severe



May need to act on URGENTLY



IE MDT
Readily available
Responsive
Integrated

#2

Face-to-face review

Frequency?

Clinical review

Post-op/wound/HF...

MDT review if needed



Hospital-based

Introduction

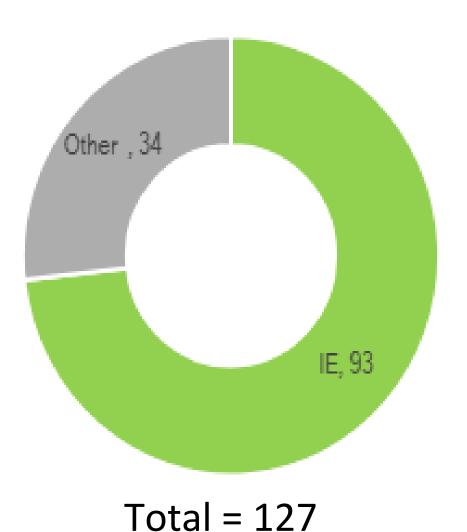
Case

The OPAT - IE cohort at SBH 2022 - 2025

Perspectives

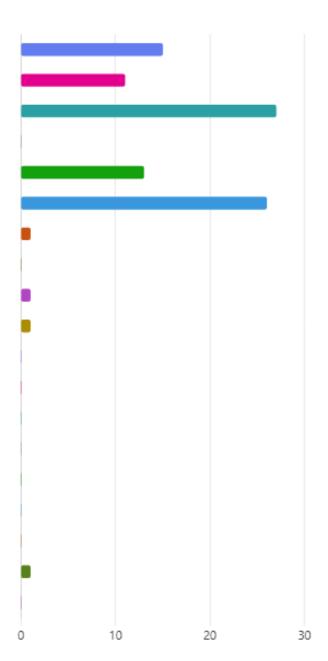


OPAT diagnoses (2022 - 2025)



9. Antibiotic to be used for OPAT

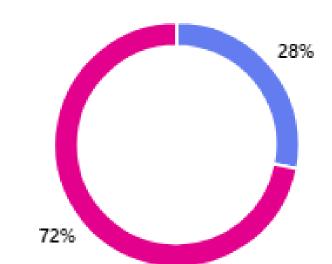
•	Benzylpenicillin	15
•	Flucloxacillin	11
•	Teicoplanin	27
•	Vancomycin	0
•	Daptomycin	13
•	Ceftriaxone	26
•	Ceftazidime	1
•	Piperacillin-tazobactam	0
•	Meropenem	1
•	Ertapenem	1
•	Amikacin	0
•	Gentamicin	0
•	Tigecyclin	0
•	Ambisome	0
•	Micafungin	0
•	Ganciclovir	0
•	Dalbavancin	0
•	Other	1
•	I don't know	0



12. Elastomeric device

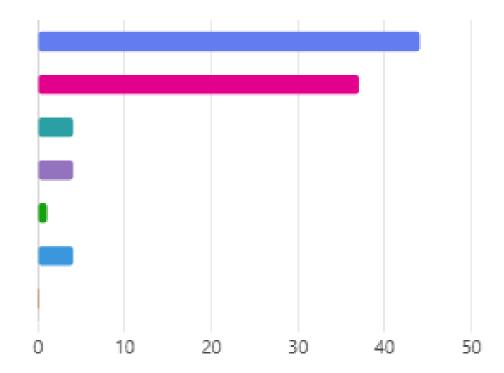
Yes 25

No 64



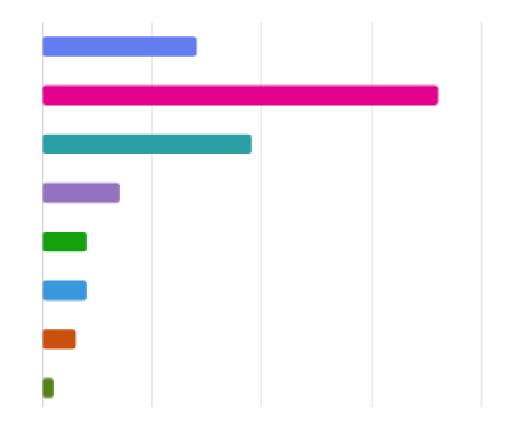
15. Method of administration

•	Self-admin	44
•	DN-admin	37
•	Third party (Baxter subcontract) - admin	4
•	Ambulatory care/ antibiotic centre	4
•	Dialysis	1
•	Virtual ward	4
•	Other	0



16. Indication for OPAT at SBH

•	Endocarditis, native, no surgery	14
•	Endocarditis, native, post-surgery	36
•	Endocarditis, prosthetic valve/conduit/GUCH no surgery	19
•	Endocarditis, prosthetic valve/conduit/GUCH, post-surgery	7
•	Device (pacemaker, ICD) infection, no explantation	4
•	Device (pacemaker, ICD) infection, post- explantation	4
•	Vascular aortic graft (no valve) infection, includes GUCH, no surgery	3
•	Vascular aortic graft (no valve), includes GUCH, post-surgery	1



Median time on IVs (OPAT)

9 days

Median follow up (OPAT)

26 days

Q75

35 days

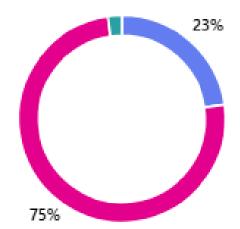
Max

177 days



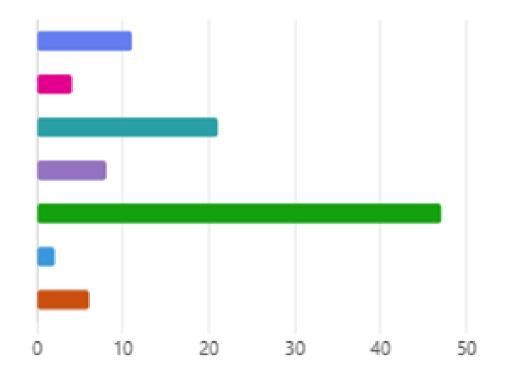
No 69

Lost



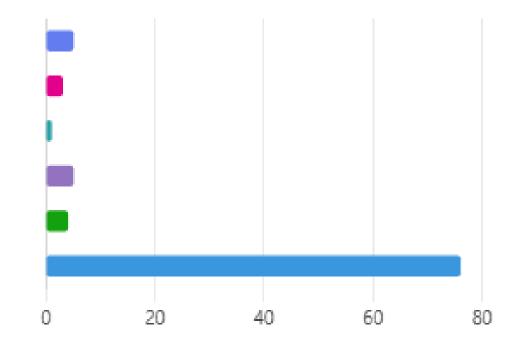
24. Complications during OPAT episode, any

0	Yes, cardiac/ endovascular/ surgery related	11
•	Yes, extra-cardiac/ distant infection (consequence of IE)	4
•	Yes, other typically OPAT related (antibiotic/line infection thrombosis /administration etc)	21
•	Yes, other not typically OPAT - related (UTI, fracture, fall, non line DVT), includes other NEW	8
•	No	47
•	Lost	2
•	Other	6

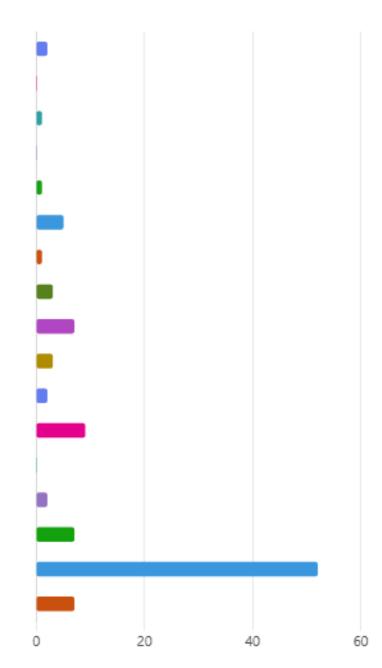


25. If new/unexpected cardiac /endovascular complications, select

•	ECG / rhythm	5
•	Valve - related (HF/ abscess)	3
•	Surgical site - related (abscess /seroma)	1
•	Uncontrolled endovascular infection	5
•	Other	4
•	No complications	76

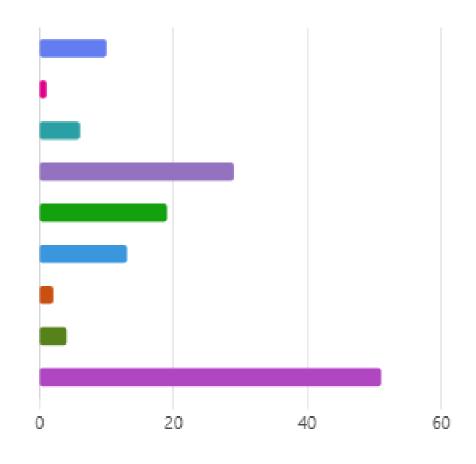


•	Extra cardiac / distant infection (IE - related)	2
•	Line - misplaced	0
•	Line - infection	1
•	Line- thrombosis	0
•	Line - blockage	1
•	Line, other	5
•	Antibiotic - infusion related	1
•	Antibiotic - cytopenia	3
•	Antibiotic - renal	7
•	Antibiotic - liver / cholecystitis	3
•	Antibiotic - rash or other HS	2
•	Antibiotic - other	9
•	Claudication of OPAT system in place	0
•	Other New infection (no IE-related)	2
•	Other	7
•	No complications	52
•	Other	7



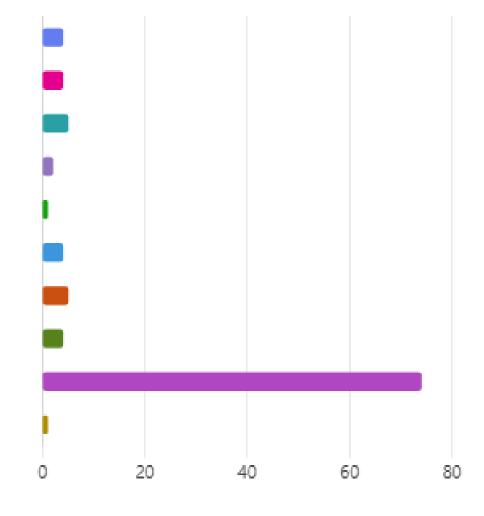
32. Cardiac/Endovascular- related procedures (during OPAT and within 3 months after OPAT completion)

 Surgery (planned/unplanned) 	10
 Washout/ non-surgical debridement 	1
Pacemaker	6
Re-imaging	29
 Re-discussion at MDT (other than opat) 	19
 Change of antibiotic duration / strategy 	13
Death	2
Other	4
No	51



31. Readmission during planned opat episode

•	Yes - sepsis/uncontrolled infection includes septic emboli from IE	4
•	Other new infection (line or non-IE related)	4
•	Yes - heart/valve/graft mechanical failure	5
•	Yes - rhythm issues	2
•	Yes - echo findings	1
•	Yes - failure of treatment	4
•	Yes - other complications from antibiotic treatment (cytopenia, reactions, renal/liver etc)	5
0	Planned readmission eg for surgery	4
•	No	74
•	Other	1

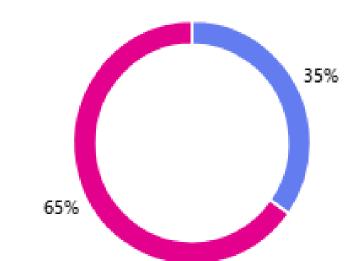


27. Was the parent/ IE team opinion/intervention required during OPAT episode (subjective opinion as per OPAT consultant)



No 58

Not sure / maybe
 0

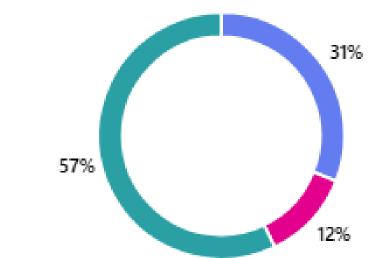


28. Did the patient actually receive any parent / IE team input during OPAT episode



No 10

Not required 48



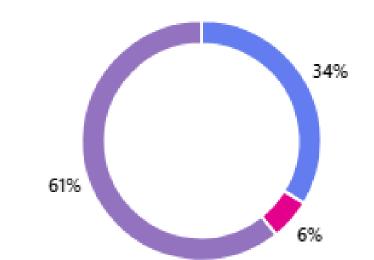
29. If there were complications that required IE/parent team input. Did the OPAT team get involved in managing the complications - to any extent (full management, delegation of tasks, hand over to team etc)



No 5

A bit unclear

There were no complications 54



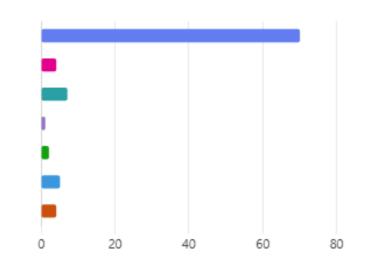


Median "involvement" (1 to 10 scale)

7

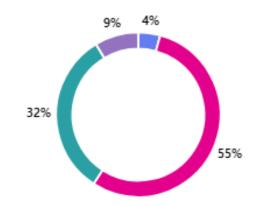
33. Outcome 3 months after end of OPAT follow up

Alive, cured	70
Alive, relapse within 3 months	4
Alive, chronic infection on suppression	7
Alive, chronic infection not on suppression	1
Death	2
 Not 3 months follow up yet 	5
• Lost	4



34. Outcome 1 year after end of OPAT follow up





#3

There will be complications

Who will identify

with them?

them and deal

OPAT team?

Vs

Very close followup by the parent team

Safety and clinical outcomes of outpatient parenteral antibiotic therapy for infective endocarditis in Christchurch, New Zealand: A retrospective cohort study

Patrick O Campbell ¹, Kate Gallagher ², Simon C Dalton ², Sarah C L Metcalf ², Nicholas M Douglas ³, Stephen T Chambers ⁴

Affiliations + expand

PMID: 37331565 DOI: 10.1016/j.ijid.2023.06.008

Free article

Abstract

Objectives: We examined the safety and clinical outcomes of outpatient parenteral antibiotic therapy (OPAT) for patients with infective endocarditis (IE) in Christchurch, New Zealand.

Methods: Demographic and clinical data were collected from all adult patients treated for IE over 5 years. Outcomes were stratified by receipt of at least partial OPAT vs entirely hospital-based parenteral therapy.

Results: There were 172 episodes of IE between 2014 and 2018. OPAT was administered in 115 cases (67%) for a median of 27 days after a median of 12 days of inpatient treatment. In the OPAT cohort, viridans group streptococci were the commonest causative pathogens (35%) followed by Staphylococcus aureus (25%) and Enterococcus faecalis (11%). There were six (5%) antibiotic-related adverse events and 26 (23%) readmissions in the OPAT treatment group. Mortality in OPAT patients was 6% (7/115) at 6 months and 10% (11/114) at 1 year and for patients receiving wholly inpatient parenteral therapy was 56% (31/56) and 58% (33/56), respectively. Three patients (3%) in the OPAT group had a relapse of IE during the 1-year follow-up period.

Conclusion: OPAT can be used safely in patients with IE, even in selected cases with complicated or difficult-to-treat infections.

Keywords: Christchurch; Home intravenous antibiotic treatment; Infective endocarditis; New Zealand; Outpatient parenteral antibiotic treatment.

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Clinical predictors of outcome in patients with infective endocarditis receiving outpatient parenteral antibiotic therapy (OPAT)

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Highlights

- Outpatient parenteral antimicrobial therapy (OPAT) is increasingly used to treat infective endocarditis (IE).
- · Pre-existing renal failure and multimorbidity were associated with OPAT failure.
- Previous IE and cardiac complication were associated with poor long-term outcomes; cardiac surgery was a protective factor.
- OPAT is safe and effective for treating IE, including cases deemed to be at increased risk of complications.
- · We examined risk factors for treatment failure and poor outcomes in patients with IE managed with OPAT.

Higher rate of complications

than general OPAT patient

Introduction

Case

The OPAT - IE cohort at SBH 2022 - 2025

Perspectives



Our cardiologists say....

The OPAT service has revolutionised the way we treat patients

Clearly important to be hospital-based or have very closed links with cardiology

The team should be united, with the patient being assessed by both teams or preferably, a team working together

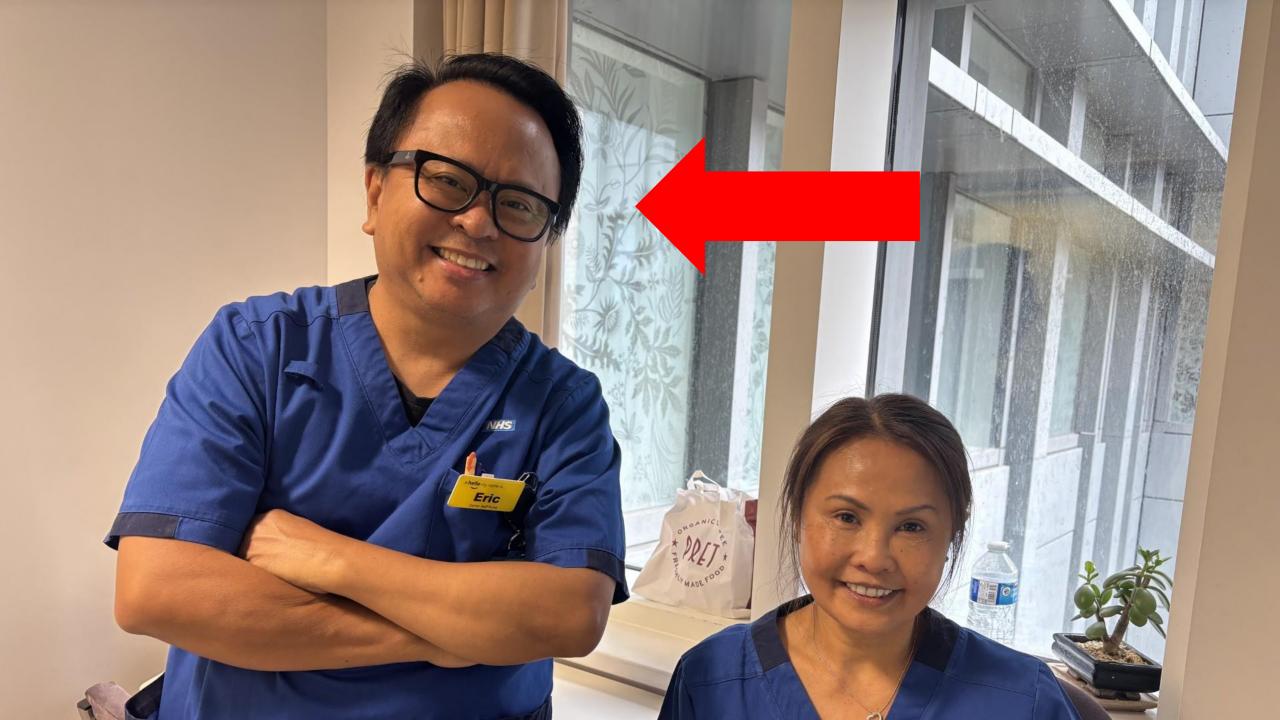
Once the risk of late emboli / complications has fallen (2 weeks)

Our patient Peter valued

Feeling safe and under close supervision and control – despite not being in a hospital bed

Having easy access to the information one needs at different points in the journey





Our CNS says: consider the following...

Complex and long journeys for patients -> closer relationship Physical and psychological impact

- Advocacy
- Navigating the system
- Counselling

Middle person

- Importance of multidisciplinary approach
- Parent team accountability

Pharmacy point of view

Complex patients, polypharmacy, comorbidities

More likely toxicity

Levels, interactions

Logistics - longer treatments than usual

#4

More demanding for ALL the members of the team



Do you have the capacity?

Is your service well resourced for this?

Complex

Demanding

MDT

Training & knowledge

Increased nursing time

Know your cohort

Work in teams

Adapt your OPAT service

Is your service ready?



