

CLINID conference
Hunter Ratliff
06/12/2025

*Ages, dates, and other identifying information may have been changed
I have no conflict of interest in relation to this presentation*

Case #1

Case 1: HPI



A **60 y/o F** with PMH including ESRD (iHD via AVF), s/p TAVR, afib, severe pulm HTN & COPD p/w **chills**.

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 - They ordered blood cultures 3 days ago

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- **Five days prior to admission** was having **chills** at dialysis center
 - They ordered blood cultures 3 days ago
- **Two weeks ago left ear fullness** (no pain, discharge, tinnitus, or hearing loss)
- **Chronic cough** (no dyspnea or change in quantity/quality of sputum)
- No issues with her RUE AVF (aside from **frequent bleeding at HD**, which is chronic)

Case 1: Medical history

A **60 y/o F** with PMH including ESRD (iHD via AVF), s/p TAVR, afib, severe pulm HTN & COPD p/w **chills**.

- Five days of **chills** & **malaise**
- Two weeks ago left ear fullness (resolved)
- Chronic cough
- No issues with her RUE AVF aside from chronic bleeding

Medical / Surgical History

- ESRD via RUE AVF (10+ years)
- Severe pulm HTN (PCWP 45)
 - Group 2 & 3 (from heart/lungs)
- COPD, HFpEF (5L home O2)
- Atrial fibrillation
 - Warfarin (bleeding on DOAC)
 - Also on dronedarone
- Hx TAVR (2021)
- DM (A1c 6.8)
- Mobility: Ambulates with walker

Case 1: Social history, exposures, & risk factors



Geographic & Travel	<ul style="list-style-type: none">• Lives in West Virginia• No travel
Occupational	<ul style="list-style-type: none">• Retired. Used to work at convenience store
Substance & needles	<ul style="list-style-type: none">• No EtOH, tobacco, drugs• No needle exposures
Animals	<ul style="list-style-type: none">• Pet dog at home• Daughter has kittens
Exposures & hobbies	<ul style="list-style-type: none">• None

Case 1: Physical exam



Vitals: 115/65 | 93 bpm | 36.8 °C | 94% | **42.68** kg/m²

Gen: alert and oriented, NAD

ENT: EOMI grossly, sclera color: anicteric sclerae; MMM

Resp: normal respiratory effort **on 5L O2**, symmetric chest rise

CV: irregular but **no murmurs**; extremities perfused

GI: non-distended; no rebound or guarding

Ext: no clubbing or cyanosis; **1+ BLE edema**; RUE AVF w/ thrill but **no TTP, calor, or discharge**

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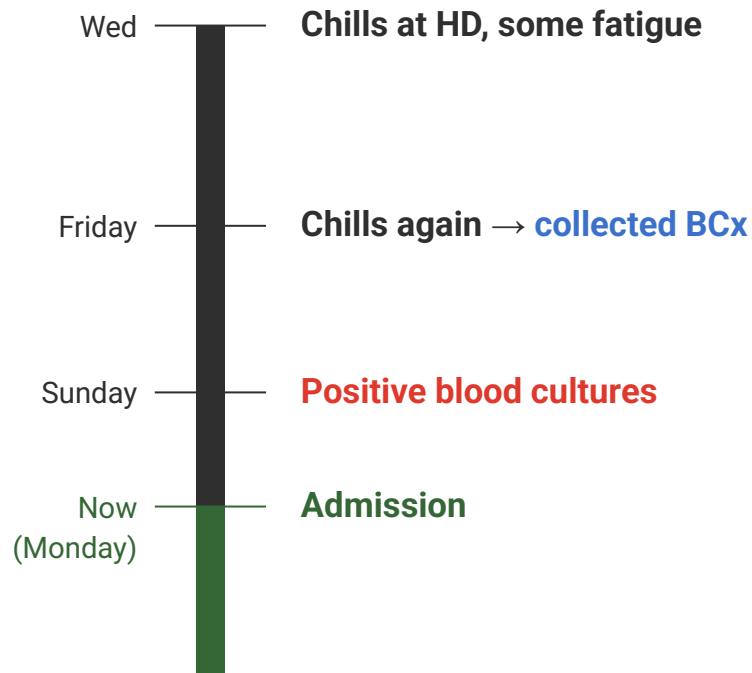
Blood cultures from HD were positive → Direct admission

Case 1: Summary

A **60 y/o F** with PMH including ESRD (iHD via AVF), s/p TAVR, afib, severe pulm HTN & COPD p/w **5 days** of **chills & malaise** and was admitted for **positive blood cultures**.

Info that may (or may not) be helpful:

- Left ear fullness (2 weeks ago, resolved)
- Chronic cough



Case 1: Micro data

GRAM STAIN

!

Yeast

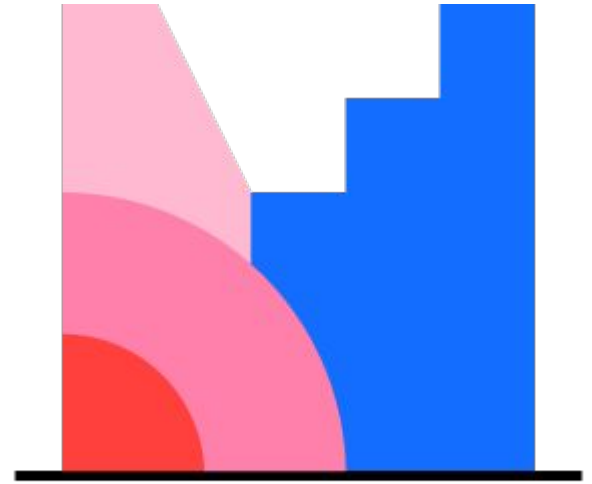
Aerobic Bottle

[Q1.1] Initial treatment



Initial treatment choice

(multiple choice)



Mentimeter

Case 1: Micro data

GRAM STAIN

!

Yeast

Aerobic Bottle

Case 1: Micro data

BLOOD CULTURE,
ROUTINE

Abnormal Stain !!

Candida glabrata !

For susceptibility, see previous report.

GRAM STAIN

!

Yeast

Aerobic Bottle

Case 1: Cardiac workup



Transthoracic echo

Quality: Technically difficult study due to limited acoustic windows

Indications: Fungemia

Mitral Valve: Anterior and posterior mitral valve leaflets appear calcified. Moderate mitral annular calcification. Moderate mitral stenosis. Mild mitral regurgitation

Aortic Valve: There is a 23 mm transcatheter valve (Sapien III) in the aortic position. The aortic prosthesis demonstrates a normal transvalvular gradient for valve type and size. There is no evidence of paravalvular aortic regurgitation.

Conclusions: No change was seen from previous echocardiogram.

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Conclusions: No change was seen from previous echocardiogram.

Medical / Surgical History

- ESRD via RUE AVF
- Severe pulm HTN, COPD, HFpEF
 - PCWP 45
 - On 5L home O2
- Hx TAVR (2021)

Do we need transesophageal echo?

Anesthesia + pulmonary hypertension = bad time

Case 1: Cardiac workup



Transesophageal echo

Quality: The study images were of technically good quality

- S/P #23 Sapien S3 Ultra Valve. Leaflet cusps are thickened and restricted. Mild transvalvular regurgitation. No paravalvular regurgitation. There is linear mobile density **measuring 1.1 x 0.2 cm** attached to the cusp in the right coronary position. In the setting of **bacteremia** fungemia this may represent vegetation.
- Mitral leaflets are calcified and restricted. Mild mitral stenosis. Moderate mitral regurgitation

Case 1: Surgical candidacy

All imaging was reviewed and discussed with the cardiac surgery team and the patient was seen and evaluated with Dr. [REDACTED]. Based on TEE imaging low suspicion for vegetation on bioprosthetic aortic valve, however we are unable to definitely rule out in the setting of fungemia.

Given the patient's multiple chronic medical conditions along with poor functional status she is a prohibitive risk for any cardiac surgical intervention. We recommend continuing antifungals per ID recommendations along with life long suppression. Can also consider follow up with cardiology after completion of IV antifungals with repeat TTE, no need for SCT follow up.

STS Risk Score: <https://acsdriskcalc.research.sts.org/calculation>

Isolated AVR

Mortality **18.5%**

Morbidity and Mortality **37.1%**

Case 1: Susceptibilities

BLOOD CULTURE,
ROUTINE

Abnormal Stain !!
Candida glabrata !

We all agree on the IV therapy, but what about **after IV therapy**?

Amphotericin B	0.5	None
Rezafungin	0.016	Suscept
Anidulafungin	0.03	Suscept
Micafungin	0.016	Suscept
Voriconazole	0.12	None
Isavuconazole	0.12	None
Posaconazole	1	None
Itraconazole	0.5	None
Fluconazole	4	SDD
Caspofungin	0.12	Suscept

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- Atrial fibrillation
 - Warfarin (bleeding on DOAC)
 - Also on dronedarone
- Hx TAVR (2021)

QTC Calculation 450 ms

D

Dronedarone
Fluconazole (QT-prolonging Moderate CYP3A4 Inhibitors (Moderate Risk))

D

Warfarin (Vitamin K Antagonists)
Fluconazole

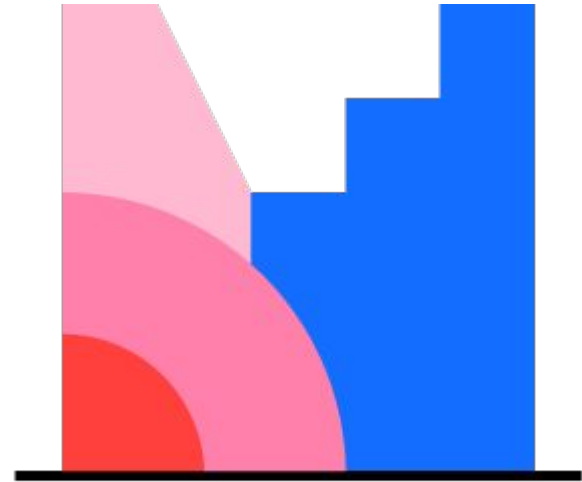
Coverage: Wv Medicaid i.e. paperwork

[Q1.2] What to do at the end?



PO transition

(multiple choice, has wrong answers)



Mentimeter

Case 1: The transition to Cresemba



A **60 y/o F** with PMH including recent ***C glabrata* prosthetic aortic valve endocarditis**, afib (on warfarin & dronedarone), ESRD, severe pulm HTN (on 5L oxygen)

1. Talk with cardiology → **won't change their meds** (patient does not want to go off warfarin)

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1. Talk with cardiology → won't change their meds (patient does not want to go off warfarin)
2. You do some **paperwork**

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2. You do some paperwork
 - And **some more paperwork...**

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1. Talk with cardiology → won't change their meds (patient does not want to go off warfarin)
2. You do some paperwork
 - And some more paperwork...
 - And **some more** (she has **finished 6 weeks** at this point, but still getting OPAT)

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1. Talk with cardiology → won't change their meds (patient does not want to go off warfarin)
2. You do some paperwork
 - And some more paperwork...
 - And some more (she has finished 6 weeks at this point, but still getting OPAT)
3. **Cresemba approved!!!**





But then...

Case 1: The transition to Cresemba

60 y/o F with PMH including recent *C glabrata* prosthetic aortic valve endocarditis, afib (on warfarin & dronedarone), ESRD, severe pulm HTN (on 5L oxygen) s/p 6+ weeks of mica

Two weeks after starting Cresemba, developed profound emesis, followed by persistent diarrhea



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Two weeks after starting **Cresemba**, developed profound **emesis**, followed by **persistent diarrhea**

- **3-4 watery bowel movements** per day
 - Associated **stomach cramping**
 - Thinks symptoms are worse in the **hours after taking Cresemba**



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- **Missed a few HD sessions** due to GI symptoms
 - I can poop while I'm at dialysis (very reasonable)



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- Brief admission to Ruby, **GI biofire negative** (**no C diff testing**)



Case 1: The transition to Cresemba

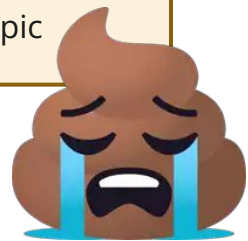
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Warfarin management

- Warfarin can no longer be managed by Ruby pharmacist
- Her family medicine doc is doing the warfarin management now
- They don't use Epic



[Q1.3] Refractory diarrhea



GI intolerance

(multiple choice, has wrong answers)



Case 1: The transition to ~~Cresamba~~ fluconazole

60 y/o F with PMH including recent *C glabrata* prosthetic aortic valve endocarditis, afib (on warfarin & dronedarone), ESRD, severe pulm HTN (on 5L oxygen) s/p 6+ weeks of mica who developed **intractable GI symptoms** 2 weeks after starting **Cresamba**. She does not want to consider DOAC again due to bleeding issues

- Order **C diff testing**
- Sent **fluconazole** to her pharmacy (on a Thursday)

Case 1: The transition to ~~Cresamba~~ fluconazole

60 y/o F with PMH including recent **C glabrata** prosthetic aortic valve endocarditis, afib (on warfarin & dronedarone), ESRD, severe pulm HTN (on 5L oxygen) s/p 6+ weeks of mica who developed **intractable GI symptoms 2 weeks after starting Cresamba**. She does not want to consider DOAC again due to bleeding issues

- Order **C diff testing** → PCR positive , Toxin negative
- Sent **fluconazole** to her pharmacy (on a Thursday)
 - **Medicaid denies fluconazole** (despite it being the *preferred formulary* option)

Case 1: The transition to ~~Cresamba~~ fluconazole

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- Tell her to **keep taking Cresamba** until gets fluconazole
 - Eventually gets **admitted to Ruby** because of symptoms & delay with the switch

Case 1: The transition to ~~Cresamba~~ fluconazole

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 - Got some **PO vanco**, but also got **PO fluconazole**

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- Tell her to **keep taking Cresamba** until gets fluconazole
 - Eventually gets **admitted to Ruby** because of symptoms & delay with the switch
 - Got some **PO vanco**, but more importantly, got **PO fluconazole**
- Things got **better for her GI system!!!**



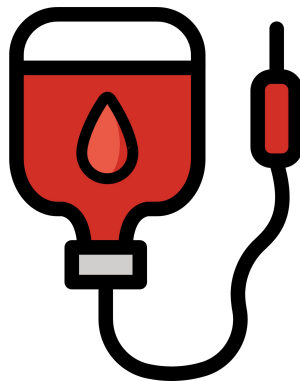


But then...

Case 1: Fluconazole doesn't play nice

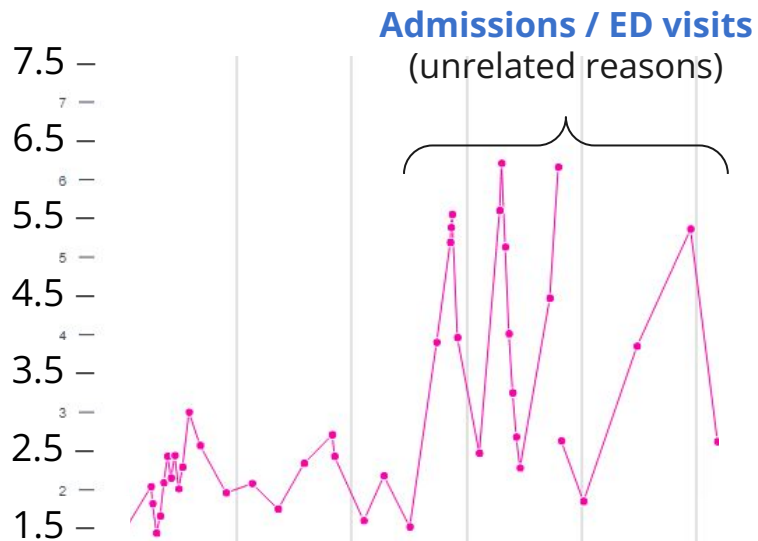
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You can probably guess what happens...



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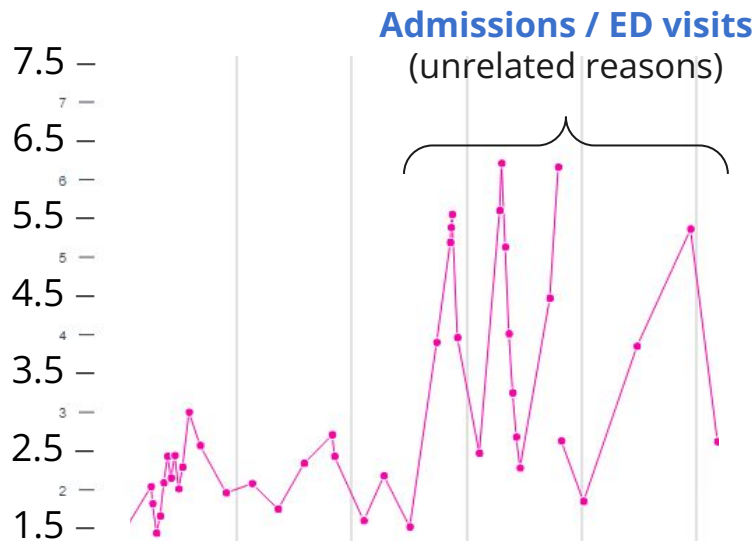


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QTC Calculation 450 ms

QTC Calculation 486 ms



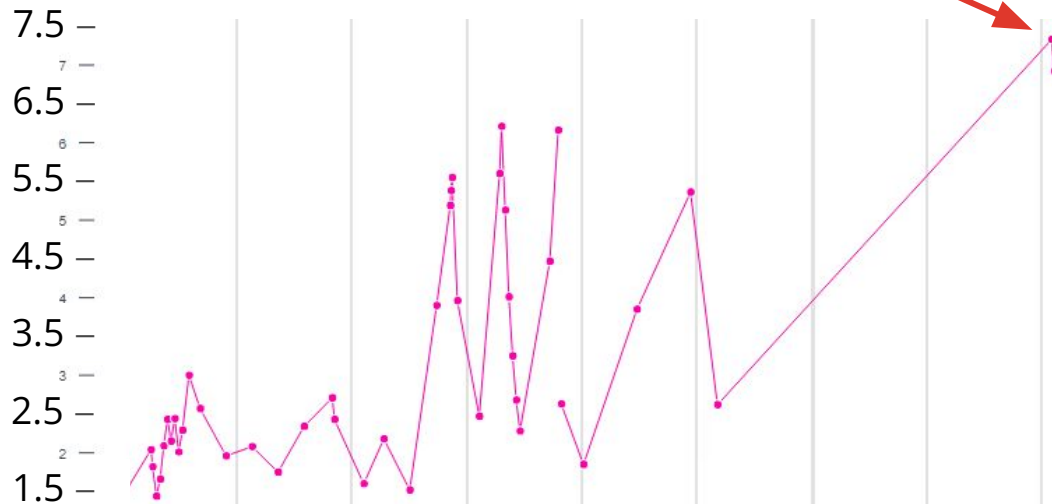
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QTC Calculation 486 ms

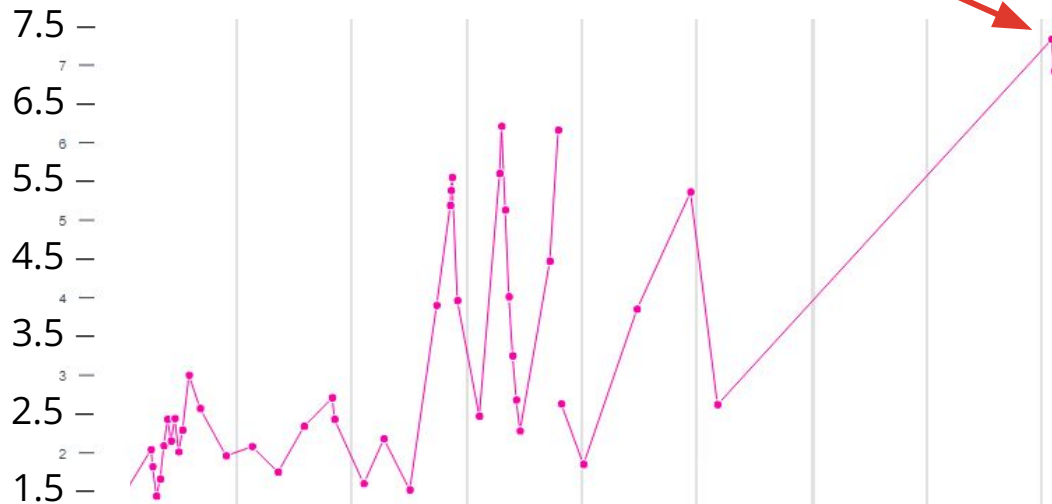
QTC Calculation 505 ms



Case 1: Fluconazole doesn't play nice

60 y/o F with PMH including recent *C glabrata* prosthetic aortic valve endocarditis, afib (on warfarin & dronedarone), ESRD, severe pulm HTN (on 5L oxygen) s/p 6+ weeks of mica who didn't Cresemba (GI Sx) and is now on fluconazole—now admitted for bleeding from fistula

- Inpatient team switched back to Cresemba
- Tolerated it well



Case 1: Hospital course

Later in that admission, had a rapid clinical decline prompting **MICU transfer**



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- Acute on chronic **respiratory failure** → hypotension **requiring pressors**



Case 1: Hospital course

Later in that admission, had a rapid clinical decline prompting **MICU transfer**

- Acute on chronic **respiratory failure** → hypotension **requiring pressors**
 - Resp Biofire: **coronavirus HKU1**
 - All **other micro data was unrevealing** (including **negative blood cultures**)



Case 1: Hospital course

Later in that admission, had a rapid clinical decline prompting **MICU transfer**

- Acute on chronic **respiratory failure** → hypotension **requiring pressors**
 - Resp Biofire: **coronavirus HKU1**
 - All **other micro data was unrevealing** (including **negative blood cultures**)
- Given comorbidities, patient/family did not want escalation of care
- Eventually **CMO**



Case #2

Case 2: HPI



A 30 y/o F with PMH including **substance use**, Hx tricuspid endocarditis **transferred with...**

Case 2: HPI

A 30 y/o F with PMH including **substance use**, Hx tricuspid endocarditis **transferred with...**



Case 2: HPI

A 30 y/o F with PMH including **substance use**, Hx tricuspid endocarditis **transferred with...**



Case 2: Medical Hx

A 30 y/o F with PMH including **substance use**,
Hx tricuspid endocarditis **transferred with**

5 years ago - Ruby

- MRSA bacteremia
- 1.4 cm TV vegetation
- Left AMA (but came back later)
- Got TV repair (but then left AMA again)

-5 years

MRSA native TV IE s/p repair

Treated with vanc → AMA

Returned → TV repair → AMA 4 days later

Case 2: Medical Hx

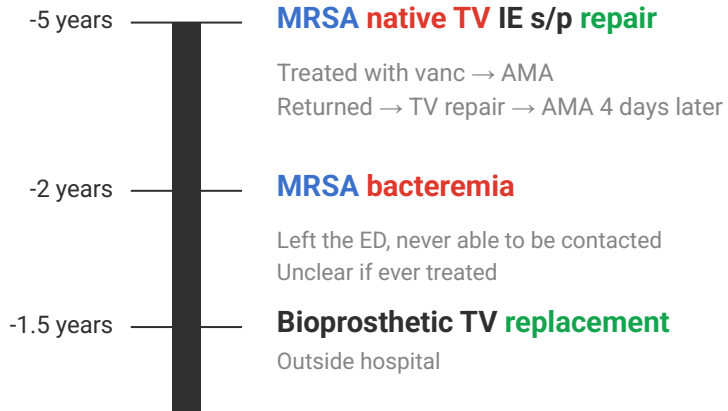
A 30 y/o F with PMH including **substance use**,
Hx tricuspid endocarditis **transferred with**

24 months ago - Satellite ED

- BCx in the ED grew MRSA
- Patient left before being admitted
- Never was able to get in contact with her

18 months ago - OSH

- Got Bioprosthetic TV replacement
- Details unclear

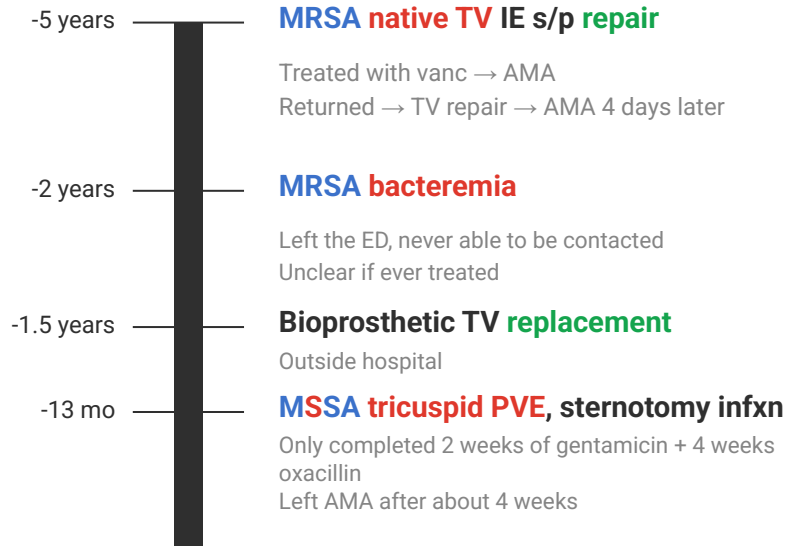


Case 2: Medical Hx

A 30 y/o F with PMH including **substance use**,
Hx tricuspid endocarditis **transferred with**

13 months ago - Ruby

- Sternotomy got infected
- MSSA bacteremia
 - TEE showed TV veggie
 - Also septic arthritis
- Plan for 6 week DOT
 - 2 weeks gentamicin + oxacillin
 - 4 weeks of oxacillin
- Left AMA before finishing last 2 weeks of oxacillin

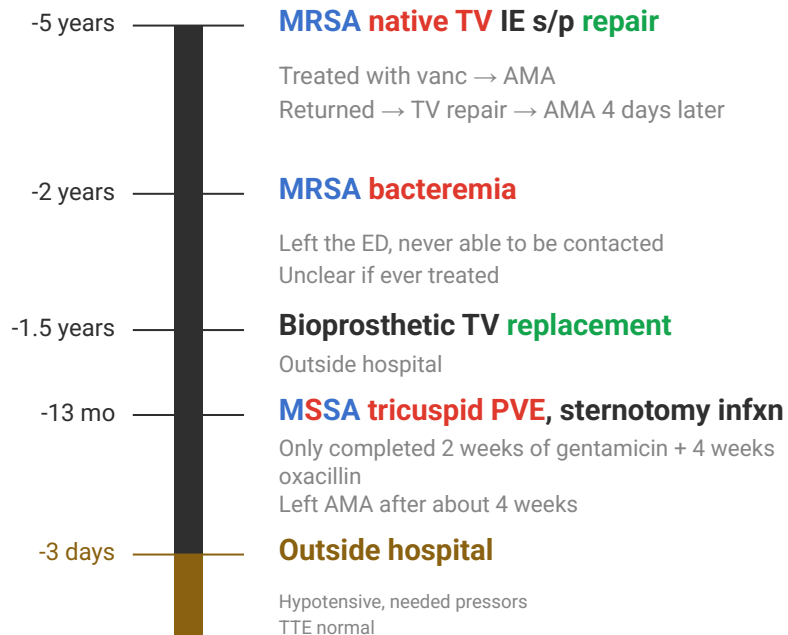


Case 2: Outside hospital

A 30 y/o F with PMH including **substance use**,
Hx tricuspid endocarditis **transferred with**

Review of outside records

- P/W chest pain
 - “Felt like prior episodes” of IE
- Admitted for 3 days
- Hypotensive, only needed levophed
- TTE normal
- Persistent fevers
- Blood cultures positive for...



Case 2: Micro data

30 y/o F with PMH including hx
PWID, Hx native TV S. aureus IE s/p
TV replacement (-1.5 yr), HCV who
p/w “feeling like prior IE” and was
admitted to OSH for 3 days with
MRSA bacteremia & septic shock

	MRSA	INTERP
	MIC	
AMP/SULBACTAM	<8/4	R*
AMOX/K CLAV'ATE	>4/2	R*
AZITHROMYCIN	>4	R
CEFAZOLIN	<8	R*
CEFTAROLINE	<0.5	S
CLINDAMYCIN	0.5	S
ERYTHROMYCIN	>4	R
GENTAMICIN	<4	S
LEVOFLOXACIN	<1	S
OXACILLIN	<0.25	R*
PENICILLIN (a)	>8	R*
RIFAMPIN (b)	<1	S
TETRACYCLINE	<4	S
TRIMETH/SULFA	<0.5/9.5	S
VANCOMYCIN	2	S

Case 2: Micro data

30 y/o F with PMH including hx PWID, Hx native TV S. aureus IE s/p TV replacement (-1.5 yr), HCV who p/w “feeling like prior IE” and was admitted to OSH for 3 days with **MRSA bacteremia** & septic shock

Fevering despite...

- Vancomycin
- Gentamicin
- Rifampin
- Ceftaroline

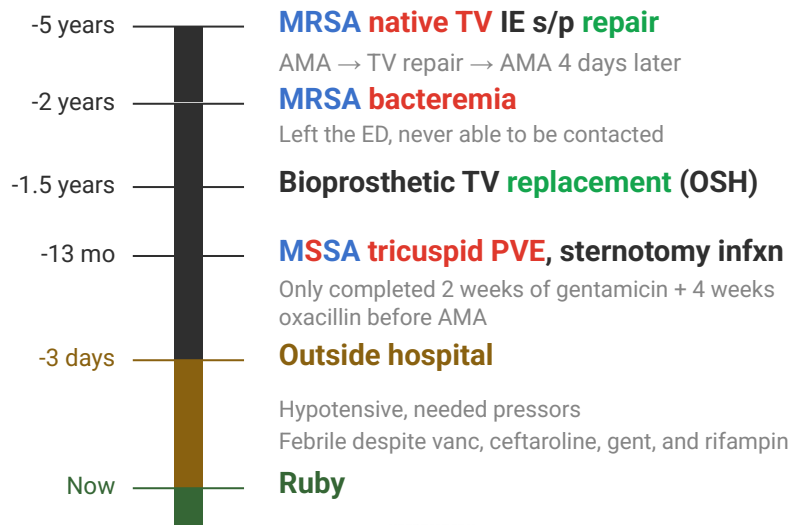
	MRSA	INTERP
	MIC	
AMP/SULBACTAM	<8/4	R*
AMOX/K CLAV'ATE	>4/2	R*
AZITHROMYCIN	>4	R
CEFAZOLIN	<8	R*
CEFTAROLINE	<0.5	S
CLINDAMYCIN	0.5	S
ERYTHROMYCIN	>4	R
GENTAMICIN	<4	S
LEVOFLOXACIN	<1	S
OXACILLIN	<0.25	R*
PENICILLIN (a)	>8	R*
RIFAMPIN (b)	<1	S
TETRACYCLINE	<4	S
TRIMETH/SULFA	<0.5/9.5	S
VANCOMYCIN	2	S

Case 2: Summary

30 y/o F with PMH including hx PWID, Hx native TV *S. aureus* IE **s/p TV replacement** (-1.5 yr), HCV who p/w “**feeling like prior IE**” and was admitted to OSH for 3 days with **MRSA bacteremia** & septic shock

Fevering despite...

- Vancomycin
- Gentamicin
- Rifampin
- Ceftaroline



	MRSA	
	MIC	INTERP
CEFTAROLINE	<0.5	S
CLINDAMYCIN	0.5	S
ERYTHROMYCIN	>4	R
GENTAMICIN	<4	S
LEVOFLOXACIN	<1	S
OXACILLIN	<0.25	R*
PENICILLIN (a)	>8	R*
RIFAMPIN (b)	<1	S
TETRACYCLINE	<4	S
TRIMETH/SULFA	<0.5/9.5	S
VANCOMYCIN	2	S

[Q2.1-3] Initial treatment

Initial treatment choice

(multiple choice)

Part A: (vanc vs dapto) (+/- gent)

Part B: Additional agents (stratified by part A)



Case 2: Hospital course

- Started on **vancomycin** & **gentamicin** on transfer to Ruby
 - Okay, so I may have **lied about the AKI** (but did have septic pulmonary emboli)

Case 2: Hospital course

- Started on **vancomycin** & **gentamicin** on transfer to Ruby
- Blood **cultures did clear** at OSF (on cultures collected prior to transfer)
 - I guess the kitchen sink approach does work!



Case 2: Hospital course



- Started on **vancomycin** & **gentamicin** on transfer to Ruby
- Blood **cultures did clear** at OSF (on cultures collected prior to transfer)
- **Added on rifampin** once blood cultures cleared

Ruby's lab	MIC	Susceptibility
Ceftaroline	KB	Susceptible
Daptomycin	1	Susceptible
Vancomycin	2	Susceptible
Linezolid	KB	Susceptible

Case 2: Hospital course

- Started on **vancomycin** & **gentamicin** on transfer to Ruby
- Blood **cultures did clear** at OSF (on cultures collected prior to transfer)
- **Added on rifampin** once blood cultures cleared
- TTE: Inconclusive

Case 2: Hospital course

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- Blood **cultures did clear** at OSF (on cultures collected prior to transfer)
- **Added on rifampin** once blood cultures cleared
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 - Interestingly **no left sided endocarditis**



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- TEE: **Vegetation** on bioprosthetic *tricuspid* valve
 - Interestingly **no left sided endocarditis**
- Transferred to **Hospitalist 7**





But then...

Case 2: Hospital course

Transferred to **Hospitalist 7** → **AKI** on vanco + gent



Case 2: Hospital course

Transferred to **Hospitalist 7** → **AKI** on vanco + gent

- **Gentamicin stopped 3 days early** due to AKI (see, I didn't totally lie!)



Case 2: Hospital course

Transferred to **Hospitalist 7** → **AKI** on vanco + gent

- **Gentamicin stopped 3 days early** due to AKI (see, I didn't totally lie!)
- Vanco switched to **dapto** (at **week 2.5** of Tx) for AKI as well
- **Rifampin stopped** here as well



Case 2: Hospital course

Transferred to **Hospitalist 7** → **AKI** on vanco + gent

- **Gentamicin stopped 3 days early** due to AKI
- Vanco switched to **dapto** (at week 2.5 of Tx)
- **Rifampin stopped** here as well
- She **stayed for the full treatment** this time
- Discharged to Hope & Healing



Case 2: Hospital course

Transferred to **Hospitalist 7** → **AKI** on vanco + gent

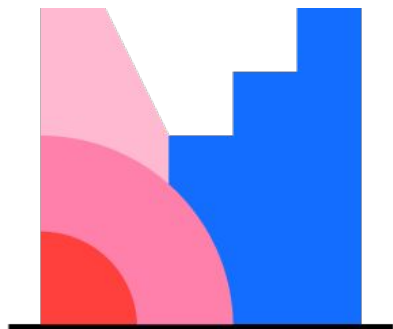
- **Gentamicin stopped 3 days early** due to AKI
- Vanco switched to **dapto** (at week 2.5 of Tx)
- **Rifampin stopped** here as well
- She **stayed for the full treatment** this time
- Discharged to Hope & Healing
- One week Zyvox → **PO suppressive doxycycline**



Case 2: A counterfactual

What would you do if doxycycline were not an option for suppression?

For the sake of argument, assume this applies to doxy & mino too



Mentimeter

	MRSA	
	MIC	INTERP
AMP/SULBACTAM	<8/4	R*
AMOX/K CLAV'ATE	>4/2	R*
AZITHROMYCIN	>4	R
CEFAZOLIN	<8	R*
CEFTAROLINE	<0.5	S
CLINDAMYCIN	0.5	S
ERYTHROMYCIN	>4	R
GENTAMICIN	<4	S
LEVOFLOXACIN	<1	S
OXACILLIN	<0.25	R*
PENICILLIN (a)	>8	R*
RIFAMPIN (b)	<1	S
TETRACYCLINE	>16	R
TRIMETH/SULFA	<0.5/9.5	S
VANCOMYCIN	2	S

[Q2.4] Suppression

MRSA suppression for endovascular infections
(multiple choice, all that apply)



Discussion



Links to articles discussed
here



Suppressive antimicrobial therapy in endocarditis

- Review the **current guidelines** for suppressive antimicrobial therapy (SAT) in endocarditis
 - This won't take long
- Examine **RCT & large prospective cohort studies** of SAT in endocarditis
- Appraise some **studies from France**

AHA / IDSA guidelines [1]



AHA / IDSA (2015)

A 2-phase treatment of fungal IE has evolved. The initial or induction phase consists of control of infection. Treatment includes a combination of a parenteral antifungal agent, usually an amphotericin B-containing product, and valve surgery. Valve surgery should be done in most cases of fungal IE. Results of a meta-analysis that included 879 cases of *Candida* IE demonstrated a marked reduction in death (prevalence odds ratio, 0.56; 95% confidence interval, 0.16–1.99) among those who underwent adjunctive valve surgery.²⁴⁴ In addition, patients who were treated with combination therapy including amphotericin B and flucytosine had reduced mortality compared with those who received antifungal monotherapy.

AHA / IDSA guidelines [1]



AHA / IDSA (2015)

Antifungal therapy usually is given for >6 weeks. After completion of this initial therapy, long-term (lifelong) suppressive therapy with an oral azole is reasonable.^{243,244,246} Suppressive therapy has been used in 2 populations. First, because of the high relapse rate of fungal IE and the prolonged

AHA / IDSA guidelines [1]



AHA / IDSA (2015)

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European Society of Cardiology [2]

ESC (2023)

7.11. Fungi

Fungi are most frequently observed in PVE and in IE affecting PWID or immunocompromised patients.³⁸⁶ *Candida* and *Aspergillus* spp. predominate, the latter resulting in BCNIE.^{387,388} Mortality is very high (>50%), and treatment necessitates combined antifungal administration and with a low threshold for surgery.^{278,387,388} Antifungal therapy for *Candida* IE in-

European Society of Cardiology [2]

ESC (2023)

and treatment necessitates combined antifungal administration and with a low threshold for surgery.^{278,387,388} Antifungal therapy for *Candida* IE includes an echinocandin at high doses or liposomal amphotericin B (or other lipid formulations) with or without flucytosine. for *Aspergillus* IE, voriconazole is the drug of choice. Some experts recommend the addition of an echinocandin or amphotericin B.^{278,387–390} Suppressive long-term treat-

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European Society of Cardiology [2]

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Strong wording!



Suppressive antimicrobial therapy in endocarditis

- Review the **current guidelines** for suppressive antimicrobial therapy (SAT) in endocarditis
- Examine **RCT & large prospective cohort studies** of SAT in endocarditis
 - This won't take long either as **there aren't any**
- I guess we are left talking about **large case series** then...

Horne et al, 2024 [3]

Good review on “The Use of Long-term Antibiotics for Suppression of Bacterial Infections”

- It won't tell you anything you don't already know
- It's a review, not a *systematic* review



Clinical Infectious Diseases



Horne et al, 2024 [3]

Number of studies published

As of 11/2023

	PJI	VGI	CIED	OM
Prospective cohort	3	0	0	0
Case control	1	0	0	0
Retrospective cohort	18	7	6	3
	Type of infection			

Abbreviations:

- **PJI** = Periprosthetic Joint Infections
- **VGI** = Vascular Graft Infections
- **CIED** = Cardiac Implantable Electronic Device Infections
 - *Most are LVADs*
- **OM** = Osteomyelitis and Spinal Hardware Infections

Adapted from Horne (2024)

Horne et al, 2024 [3]

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French studies (Beaumont, Lemmet)

In France, they have **multidisciplinary endocarditis teams** (METs) at many of their tertiary care centers

- Comprised of infectious diseases, cardiology, surgery, radiology, pharmacy, and other specialists (e.g. microbiology, neurology)
- Have regularly scheduled meetings (like **tumor board for IE**)



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functional status she is a prohibitive risk for any cardiac surgical intervention. We recommend continuing antifungals per ID recommendations along with life long suppression. Can also consider follow up with cardiology after completion of IV antifungals with repeat TTE, no need for SCT follow up.

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Relevance for this literature review:

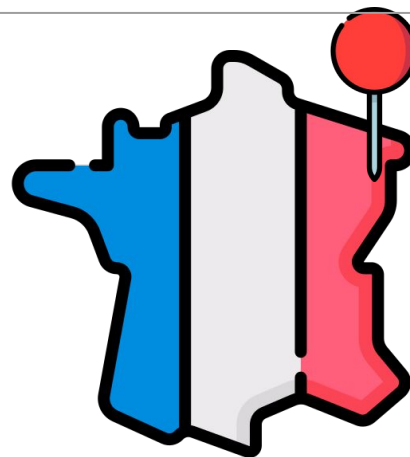
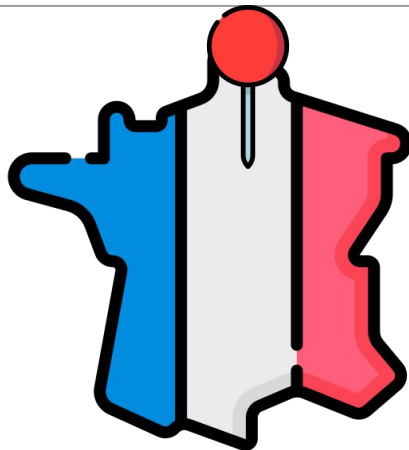
- They keep **large registries** of their patients
- **Less intra-consultant variability** on management decisions



French studies



	Beaumont, 2024	Lemmet, 2024
Timeframe	Paris, France (2016-2022)	Strasbourg, France (2020-2023)
Design	Descriptive case series from single referral centers in France	



French studies



	Beaumont, 2024	Lemmet, 2024
Timeframe	Paris , France (2016-2022)	Strasbourg , France (2020-2023)
Design	Descriptive case series from single referral centers in France	
Inclusion	Finished initial therapy , were recommended to start SAT (& started it); Could be native -or- prosthetic valve IE -or- CIED	
Dx criteria	Definite or possible IE via 2023 Duke-ISCVID criteria	IE based on 2023 ECS criteria
Exclusion	LVADs, vascular graft, Q fever	---

Abbreviations

SAT = Suppressive antimicrobial therapy

IE = Infective endocarditis

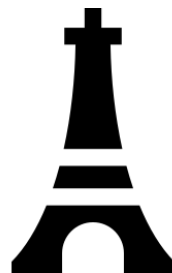
CIED = Cardiac implantable electronic device

French studies



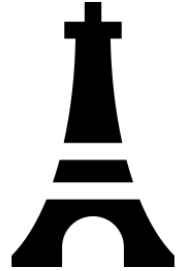
	Beaumont (Paris, 2016-2022)	Lemmet (Strasbourg, 2020-2023)
Patients	IE or CIED (n=42)	IE or CIED (n=22)
Outcomes	Mortality Relapse (infection with same pathogen) Side effects of SAT	
Duration of follow up	3 years	2 years
Other notes	Supplemented mortality analysis with a national database to capture out of hospital deaths (but couldn't ascertain reason for death)	

Beaumont (2024): Demographics [4]



- **42 patients** included
 - Majority **male** (86%)
 - Median **age 73** (IQR 61-82)
 - High charlson comorbidity index **median = 3**
- **95%** had at **least one prosthetic** cardiac device
- Only **one PWID** (person who injects drugs)

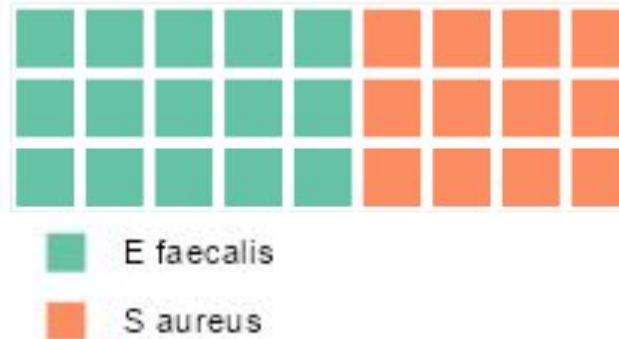
Beaumont (2024): Pathogens [4]



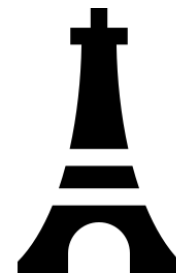
All 42 cases were **community-acquired**

- Enterococcus faecalis (36%)
- Staphylococcus aureus (29%)
 - **11 of these 12 cases** were **MSSA**

Pathogens



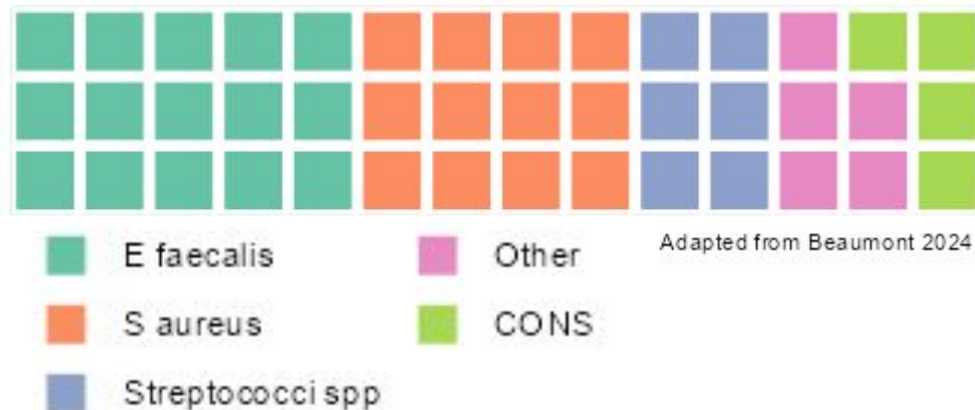
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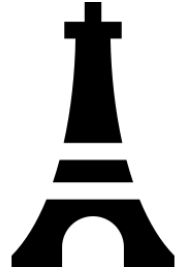
All 42 cases were **community-acquired**

- Enterococcus faecalis (36%)
- Staphylococcus aureus (29%)
 - **11 of these 12 cases** were **MSSA**
- Streptococci spp (14%)
- Coagulase-negative staph (9%)
- Other (12%)
 - Only one (2%) fungal (C albicans)
 - Only one (2%) gram negative (ESBL E cloacae; Tx w/ mino)

Pathogens



Beaumont (2024): Suppressive agents [4]



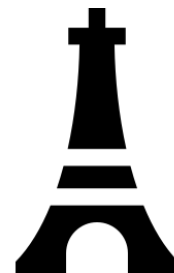
- Doxycycline (**45%**)
- Amoxicillin (**45%**)
- Bactrim (**5%**, n=2)
- Fluconazole & minocycline (**2%**, n=1)

Pathogens



Adapted from Beaumont 2024

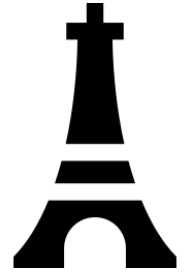
Beaumont (2024): Surgical indications [4]



38/42 cases (90%) had a theoretical indication for surgery

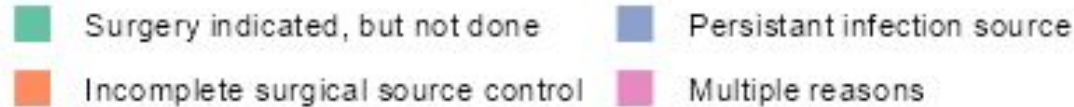
- 50% of the indicated surgeries were valvular
- 42% CIED extraction
 - Extraction considered surgical as $\frac{3}{4}$ had material implanted ≥ 4 years
 - High risk of conversion to sternotomy
- 8% had both indications

Beaumont (2024): SAT indications [4]

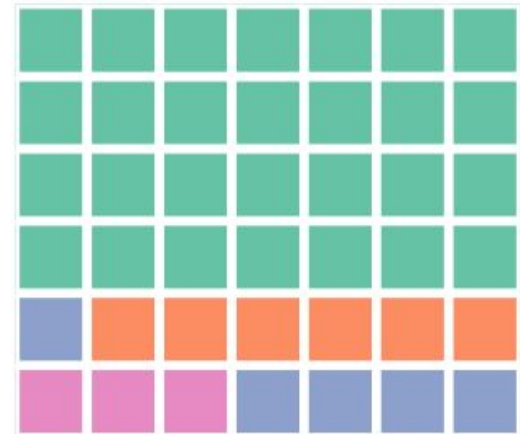


Suppressive antimicrobial therapy (**SAT**) indications:

- Surgery theoretically indicated but not done (**67%**)
- Had surgery, but incomplete source control (**14%**)
- Persistent infection source (**12%**)
- Multiple reasons (**7%**)




SAT indication



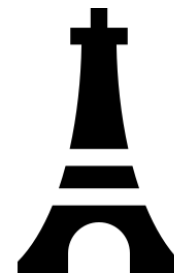
Adapted from Beaumont 2024

Beaumont (2024): Side effects [4]



12% had **adverse events** of SAT **at 1 year** (Kaplan-Meier estimator)

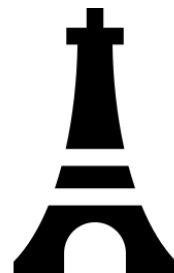
- Most **side effects were mild**, without therapy interruption



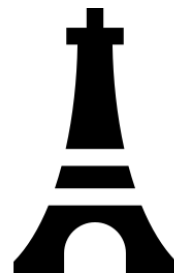
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 - Mild diarrhea
 - Doxy: epigastric burning or moderate thrombocytopenia
 - Mino: blue coloration of scars
- No C diff (across entire cohort)



Beaumont (2024): Side effects [4]



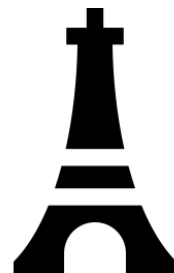
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- No C diff (across entire cohort)

Severe AEs

- The only severe adverse event was AKI
 - From **Bactrim** (big surprise)
- Resolved after stopping (switched to doxy + Augmentin)
- Recall, *only two patients received bactrim*

Beaumont (2024): Interruption & compliance [4]



13% had **interruption** of SAT **at 1 year** (Kaplan-Meier estimator)

- **Two thirds were inadvertent** (SAT was discontinued because of oversight regarding the indication)
- None resulted in recurrence

12% of patients had **poor compliance** (judged by the EMR)

Beaumont (2024): Recurrence [4]

Two had reinfections (4.8%, 1 in 21)

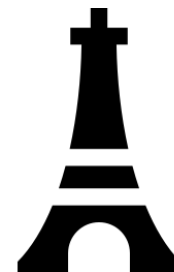
1. One was a transplant patient
 - CRBSI with S epi → switched to mino
2. One was the PWID
 - Reinfected with group B strep

Terms used here

Relapse = Infection with same pathogen

Reinfection = Infection w/ **another** pathogen

Recurrence = Relapse + reinfection



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PWID (only one in this cohort)

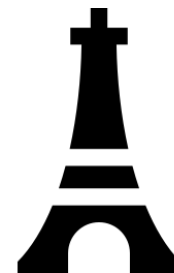
- ❖ Had an index infection with *S epi*
- ❖ Tx with Bactrim
 - Not the AKI patient
- ❖ PVE (aortic & mitral) + CIED
- ❖ They were the only death that could be attributed to infection
 - *More on this later*

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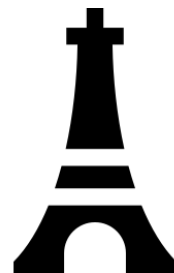


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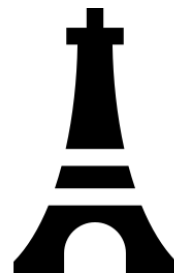
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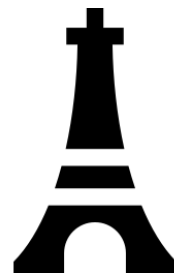
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 - Increased Amox 2g → 3g (no relapses)



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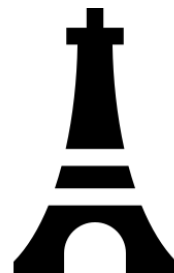
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 - 4 hospitalizations

Beaumont (2024): Recurrence [4]



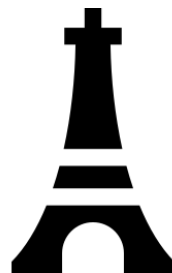
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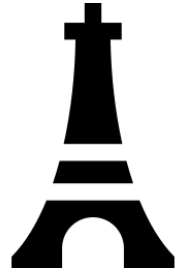
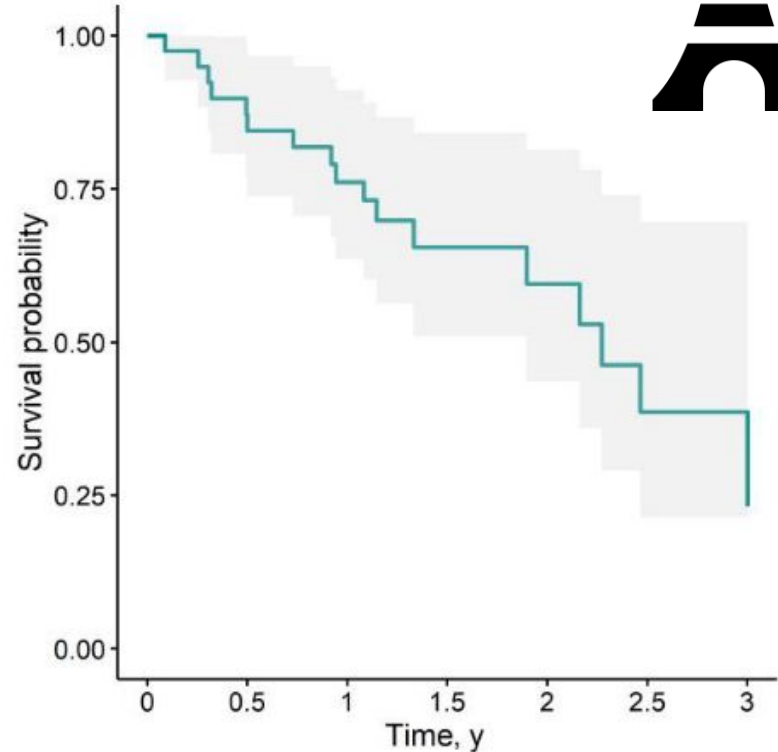
Strange thing is...

All of the isolates from **relapses** were **still susceptible to their SAT**, but had developed **resistance** to the **induction therapy**

Beaumont (2024): Mortality [4]

84.3% **1 year survival** (Kaplan-Meier estimator)

- **One in three died** during follow-up
 - Median time to mortality **13.5 months** (IQR 6.7 - 25.3)



Beaumont (2024): Mortality [4]

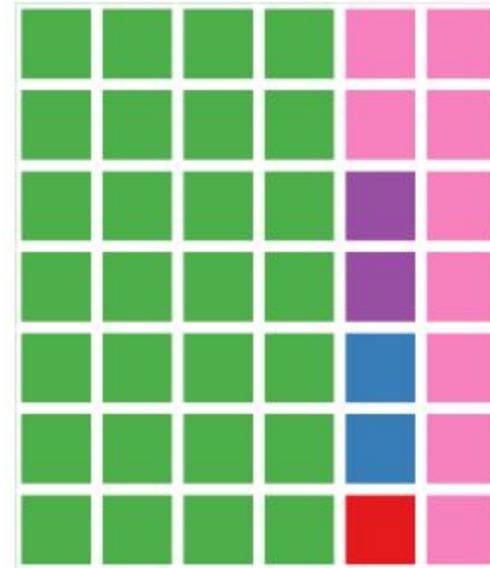
84.3% 1 year survival

- **One in three died** during follow-up
 - Median time to mortality **13.5 months**

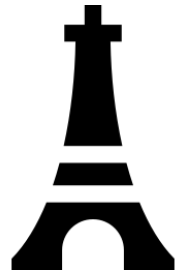
Of the **five deaths documented** in EMR (9 unknown causes of death)

- **One** died from **reinfection** (PWID)
- **Two** died from **unrelated causes**
- **Two** died from **likely unrelated causes**
 - i.e. end-stage heart failure, without any bacterial documentation on blood cultures

Cause of death



Adapted from Beaumont 2024



French studies



	Beaumont (Paris, '16-22) n=42	Lemmet (Strasbourg, '20-23) n=22
Timeframe	Paris, France (2016-2022)	Strasbourg, France (2020-2023)
Design	Descriptive case series from single referral centers in France	
Inclusion	Finished initial therapy , were recommended to start SAT (& started it); Could be native or prosthetic valve	
Dx criteria	Definite or possible IE -or- CIED via <u>2023 Duke-ISCVID</u> criteria	IE based on 2023 ECS criteria
Follow up	3 years	2 years
Outcomes	Mortality Relapse Side effects of SAT	

Lemmet (2024): Similarities & differences [5]



Similarities

- Similar rates & types of prosthetic devices
- Same rate of surgery being indicated but not done (90%)
 - For similar reasons
- Similar rates of inadvertent disruption in SAT

Differences

Lemmet had:

- Higher CCI: Mean of 6.6 (vs median of 3)
- About 5 years older
- **No PWID**

Pathogens	Beaumont	Lemmet
Enterococcus spp	15 (36%)	6 (27%)
Staph aureus	12 (29%)	6 (27%)
Streptococcus spp	6 (14%)	5 (23%)
CONS	4 (9%)	4 (18%)
Fungal	1 (2%)	0
Gram negative	1 (2%)	1 (4.5%)



Lemmet (2024): Suppressive agents [5]



Amoxicillin (41%) - Used for all *E faecalis* (n=4) & *Strep spp* (n=5)

Doxy (13%) - Staph (both *S aureus* & *S epi*)

Clinda (4.5%, n=1) - *Staph aureus*

Parenteral agents (4.5%, n=1) - Teicoplanin (given after HD for *E faecium*)



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Bactrim (32%, n=7) - Mainly for Staph (both *S aureus* & *S epi*)

- The **median age** for Bactrim patients is **88 years old**!
- They've have a 88 years old on 1 DS BID for over 14 months
- Over half of their cohort has CKD (not on HD)

Lemmet (2024): Side effects [5]



14% (n=3) had **side effects** from SAT

1. **Rash** attributed to **Bactrim** (non-severe) at day 8
 - Switched to Doxy
2. **Elevated LFTs** attributed to **Bactrim**
3. **Tooth discoloration** attributed to **amoxicillin**
 - Switched to Bactrim

Lemmet (2024): Recurrence & Mortality [5]



Two patients (9%) had **recurrence**

1. **E. faecium** (native MV) on **Bactrim SS daily** @16 months
 - Thought to be from cholangitis
2. **MSSA** (CIED) on **Bactrim SS daily** @10 months

Lemmet (2024): Recurrence & Mortality [5]



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75% **1 year survival** (Kaplan-Meier estimator)

- **Four deaths** were **unrelated** to IE
- One was unknown

Learning points & take aways



Learning points & take aways

- **Data on SAT** (suppressive antimicrobial therapy) for **prosthetic valve endocarditis is lacking**, to say the least
- **Multidisciplinary Endocarditis Teams** (METs) are used in France, would maybe not be a bad idea here too
- Observational data implies **high mortality risk** in these **comorbid patients**
 - **Often unrelated to their endocarditis** (but perhaps related to why they are not surgical candidates)
- **Side effects & disruptions in SAT** are common
 - Side effects often mild
 - Disruptions often unintentional, but some folks do fine off of SAT
- Some places in **France must really like Bactrim** (Cotrimoxazole)
 - They gave Bactrim 1 DS BID to an **88 year old** (who likely had CKD!)



Slides available on hunteratcliff1.com/talk/; Citations available via QR code or via the “citations” button on the website

Case #3

Case 3: HPI



A **35 y/o M** with PMH including opiate use disorder (PWID, with recent relapse), multiple episodes of mitral valve endocarditis (MRSA **8 years ago** s/p repair of native MV; E faecalis s/p repair & annuloplasty **2.5 years ago**)

- Clinic follow up for ***Serratia marcescens* bacteremia**
- TEE: 1.5 cm vegetation on **prosthetic MV ring**, c/w a prosthetic ring vegetation

Allergies

Sulfa: Urticaria

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Worried about long term side effects from antibiotics (understandably)

Susceptibility

	<i>Serratia marcescens</i> MIC SUSCEPTIBILITY	PHENO MIC SUSCEPTIBILITY
Amikacin		<=4 mcg/mL Sensitive
Amoxicillin/clavulanate	Resistant	
Ampicillin	Resistant	
Aztreonam		2 mcg/mL Sensitive
Cefazolin	Resistant	
Cefepime		<=1 mcg/mL Sensitive
Ceftazidime		<=1 mcg/mL Sensitive
Ceftriaxone		1 mcg/mL Sensitive
Ciprofloxacin		<=0.25 mcg/mL Sensitive
Gentamicin		<=1 mcg/mL Sensitive
Levofloxacin	<=0.12 mcg/mL Sensitive	
Piperacillin/Tazobactam		<=4 mcg/mL Sensitive
Tetracycline	>=16 mcg/mL Resistant	
Tobramycin		8 mcg/mL Intermediate
Trimethoprim/Sulfamethoxazole	<=20 mcg/mL Sensitive	