



Objectives

- Describe dalbavancin use for severe Staphylococcus aureus infections at our institution
- Evaluate clinical outcomes for patients treated for a full or partial course with dalbavancin for severe *S. aureus* infections

Background

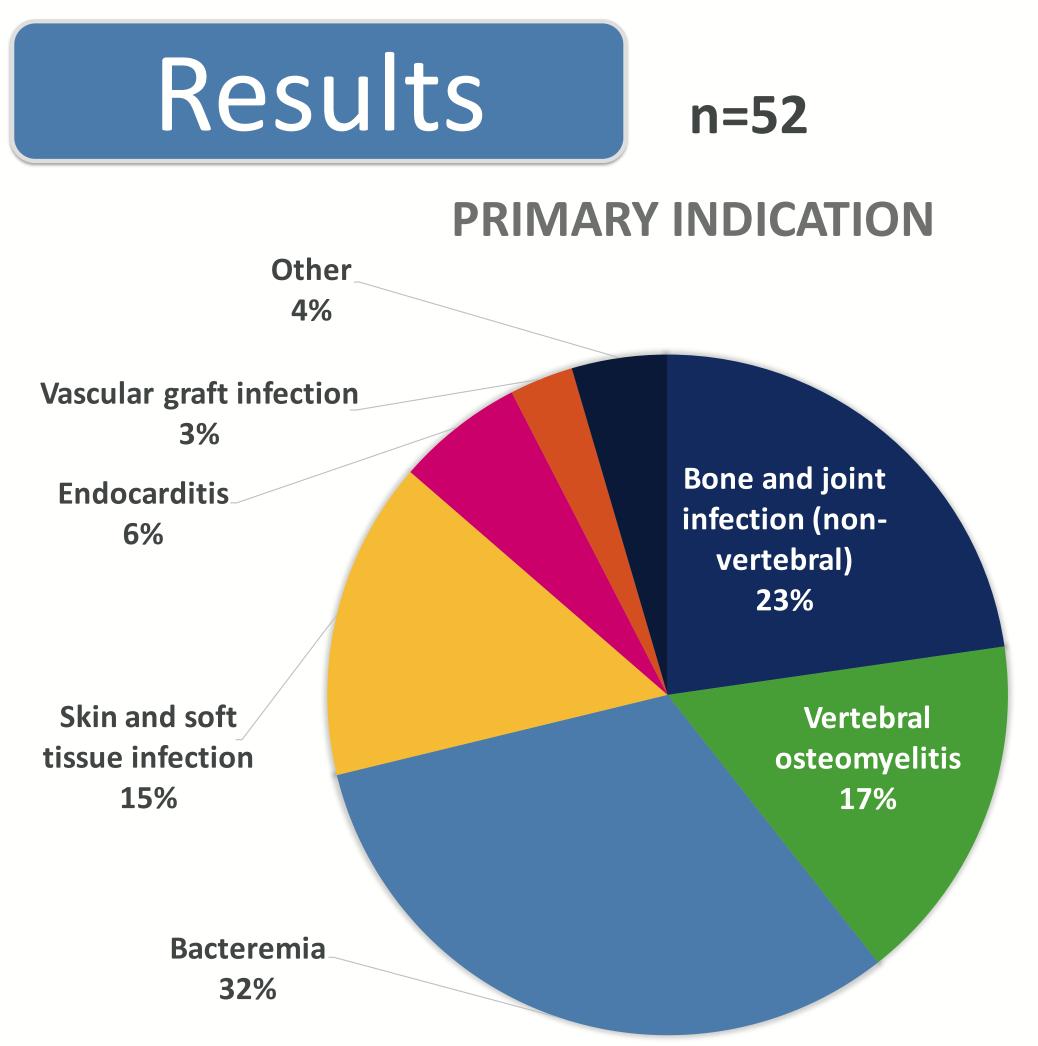
- Dalbavancin is a lipoglycopeptide antibiotic active against gram-positive organisms. It's extended half-life allows for weekly dosing that can last 4 to 6 weeks with 2 doses
- Although only FDA-approved for treating skin and soft tissue infections, adequate bone concentrations and clinical outcomes have been reported for the treatment of osteomyelitis^{1,2}
- Use of for complicated infections is appealing, particularly when daily antibiotics are impractical, but clinical outcomes data for other complicated infections is limited^{3,4}

Methods

- Retrospective study of dalbavancin use for severe infections including bacteremia caused by *S. aureus*
- We identified all patients ≥18 years old who received at least 1 dose of dalbavancin in any setting
- Infectious Disease faculty reviewed charts for clinical characteristics and outcomes of the infections

Dalbavancin for the Treatment of Infections due to Staphylococcus aureus

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*26 (50%) of patients had positive blood cultures

Dosing Regimens Utilized	n (%)
1500 mg x 1	29 (55)
1500 mg x 2	13 (25)
1000 mg x 1	3 (6)
1000 mg x1, followed by 500 mg weekly	3 (6)
1500 mg x1, followed by 1000 mg x1	1 (2)
1000 mg weekly	1 (2)
1000 mg x1, followed by 375 mg weekly	1 (2)
760 mg x 1, followed by 375 mg x1	1 (2)

Diagnostic Imaging Performed in Workup	n (%)
TTE	29 (56)
TTE resulted as no vegetation or unexplained regurgitation	19 (66)
TEE	2 (4)
Cross-sectional Spine Imaging (CT or MRI)	17 (33)
Vascular imaging (venous duplex ultrasound)	13 (25)
Other cross-sectional imaging	18 (35)
Any additional imaging to assess for metastatic infection	5 (10)

Demog

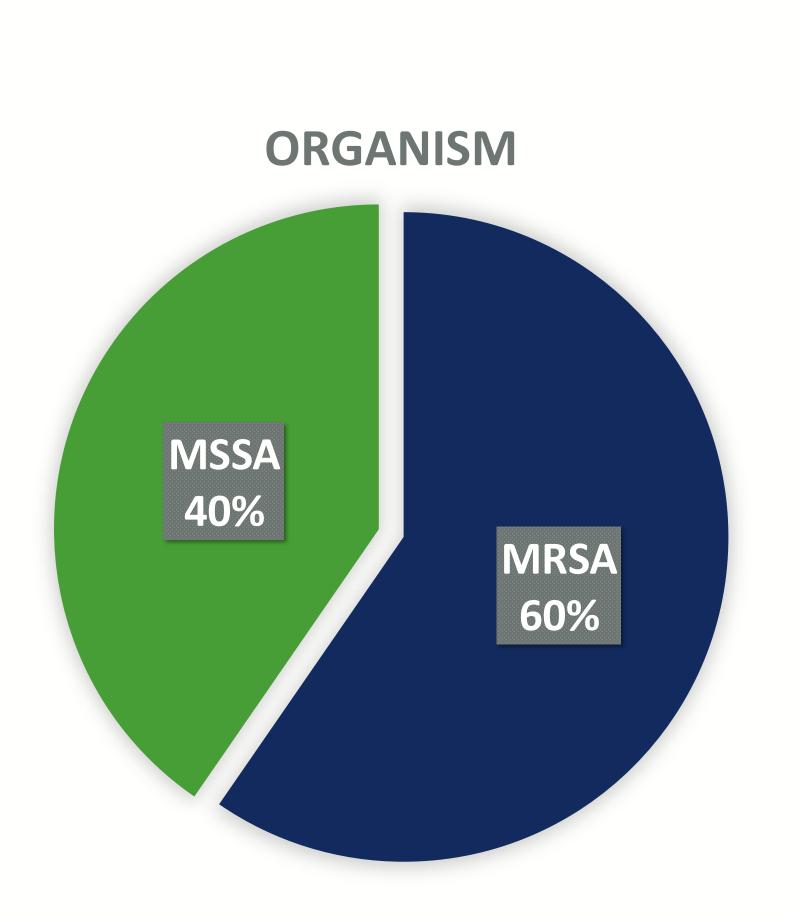
Age (Years Gender (I History of

Reason

History of Lack of sa receive da Prior non-**Clinical** co antibiotic

Adverse r Lack of alt funding o Substance Inability o Patient re antibiotic **Prior histo** Dischargi accommo **Prior trea**

Unclear Dalbavancin was selected for one or more of the below reasons, all reasons given in medical record were noted so the denominator is > 52



rs) Mean 45.5 (STD Female) 15 (29%)) 13.5)	
$Eomalo \qquad 1E(200/)$		
	15 (29%)	
of IVDU 27 (52%)		
for Selection		
of IV drug use	25	
afe home environment in which to laily IV antibiotics	11	
n-adherence to outpatient antibiotics	11	
ontraindications to alternative cs	7	
reaction to initial outpatient antibiotic	5	
Iternative outpatient options due to or insurance issues	5	
ce use, not IV drug use	3	
of patient to physically manage PICC	2	
efused PICC or daily outpatient IV cs	2	
tory of contaminated/manipulated PICC	2	
ing to a setting that could not odate daily IV antibiotics	2	
atment failure	1	
	1	

Clinical End

Loss to follow

Readmission

Readmissior days 30-90

Dalbavancin

Readmission

Recurrence c day 30 Recurrence c between days

30-day morta

90-day morta

Conclusion

While our results suggest dalbavancin is well tolerated, questions about relapse rates in the treatment of severe S. *aureus* infections remain and caution is warranted in clinical situations without optimal control of the infection.

Further research is needed to evaluate clinical outcomes for dalbavancin compared to standard of care antibiotics for *S. aureus* infections.



- efficacy and safety. OFID 6:1-9.

dpoints	n (%)
w-up by day 90	8 (15)
for any reason by day 30	13 (25)
n for any reason between	1 (2)
-related adverse effects	2 (4)
due to adverse effects	0
or relapse of infection by	11 (21)
or relapse of infection ys 30-90	5 (10)
ality	0
ality	0

References

1. Dunne MW, Puttagunta S, Sprenger CR, Rubino C, Van Wart S, Baldassarre J. 2015 Extended-duration dosing and distribution of dalbavancin into bone and articular tissue. AAC 59:1849-1855.

2. Rappo U, Puttagunta S, Shevchenko V, et al. 2018. Dalbavancin for the treatment of osteomyelitis in adult patients: a randomized clinical trial of

3. Raad I, Darouiche R, Vazquez J, et al. 2005. Efficacy and safety of weekly dalbavancin therapy for catheter-related bloodstream infection caused by gram-positive pathogens. CID 40:374-380.

4. Tobudic S, Forstner C, Burgmann H, et al. 2018. Dalbavancin as primary and sequential treatment for gram-positive infective endocarditis: 2-year experience at the General Hospital of Vienna. CID 67:795-798.