

No Vein, No Gain? Think Again. Transitioning to Oral Antibiotics

Sara Azimi, PharmD, BCIDP

Nebraska Infectious Diseases Conference

28 August 2026

Learning Objectives

1. Identify the benefits of transitioning from IV-to-oral antimicrobial therapy
2. Describe eligibility for IV-to-oral antimicrobial conversion
3. Discuss a patient case to determine the most appropriate oral antimicrobial therapy



**PROVIDERS WHEN I ASK TO SWITCH
TO PO FOR COMPLICATED INFECTION**



imgflip.com



The Evidence: PO is just as effective as IV



- PJI, osteomyelitis, VOM
- IV to PO switch after ~7 days of treatment
- 1y treatment failure: 13.2% (PO) vs 14.6% (IV)



The Evidence: PO is just as effective as IV



- PJI, osteomyelitis, VOM
- IV to PO switch after ~7 days of treatment
- Treatment failure: 13.2% (PO) vs 14.6% (IV)

NONINFERIOR



The Evidence: PO is just as effective as IV



- PJI, osteomyelitis, VOM
- IV to PO switch after ~7 days of treatment
- Treatment failure: 13.2% (PO) vs 14.6% (IV)

NONINFERIOR



- Left-sided endocarditis
- IV to PO switch when stable & after ~10d of treatment
- 6mo all-cause mortality & associated adverse events: 9.0% (PO) vs 12.1% (IV)



The Evidence: PO is just as effective as IV



- PJI, osteomyelitis, VOM
- IV to PO switch after ~7 days of treatment
- Treatment failure: 13.2% (PO) vs 14.6% (IV)

NONINFERIOR



- Left-sided endocarditis
- IV to PO switch when stable after ~10d of treatment
- All-cause mortality & associated adverse events: 9.0% (PO) vs 12.1% (IV)

NONINFERIOR



SABATO



The Evidence: PO is just as effective as IV



- PJI, osteomyelitis, VOM
- IV to PO switch after ~7 days of treatment
- Treatment failure: 13.2% (PO) vs 14.6% (IV)

NONINFERIOR



- Left-sided endocarditis
- IV to PO switch when stable after ~10d of treatment
- All-cause mortality & associated adverse events: 9.0% (PO) vs 12.1% (IV)

NONINFERIOR



- Uncomplicated *S. aureus* bloodstream infection
- IV to PO switch 5-7d of treatment
- 90d complication rate: 13% (PO) vs 12% (IV)



The Evidence: PO is just as effective as IV



- PJI, osteomyelitis, VOM
- IV to PO switch after ~7 days of treatment
- Treatment failure: 13.2% (PO) vs 14.6% (IV)

NONINFERIOR



- Left-sided endocarditis
- IV to PO switch when stable after ~10d of treatment
- All-cause mortality & associated adverse events: 9.0% (PO) vs 12.1% (IV)

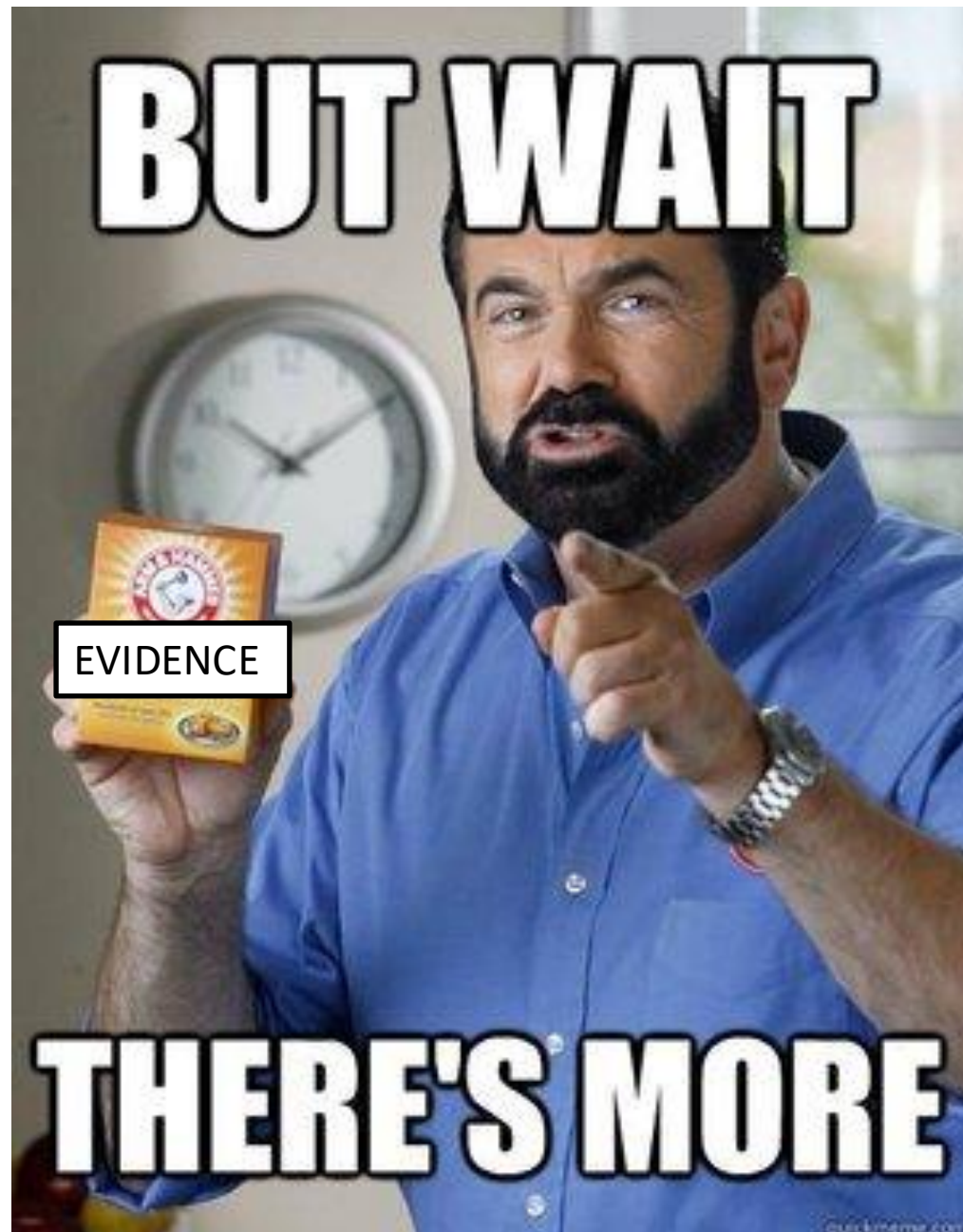
NONINFERIOR



- Uncomplicated *S. aureus* blood stream infection
- IV to PO switch 5-7d of treatment
- SDO complication rate: 13% (PO) vs 12% (IV)

NONINFERIOR





Much, much, more...

CLINICAL RESEARCH STUDY



Oral Is the New IV. Challenging Decades of Blood and Bone Infection Dogma: A Systematic Review



Noah Wald-Dickler, MD,^{a,b,c} Paul D. Holtom, MD,^{a,b} Matthew C. Phillips, MD,^a Robert M. Centor, MD,^{d,e} Rachael A. Lee, MD,^{d,e} Rachel Baden, MD,^a Brad Spellberg, MD^a

^aLos Angeles County + University of Southern California Medical Center, Los Angeles; ^bDepartment of Medicine, Keck School of Medicine, University of Southern California, Los Angeles; ^cDepartment of Preventive Medicine, Keck School of Medicine, University of Southern California, Los Angeles; ^dDepartment of Medicine, University of Alabama at Birmingham School of Medicine, Birmingham; ^eBirmingham Veterans Affairs (VA) Medical Center, Birmingham, Ala.

Visit <https://www.bradspellberg.com/oral-antibiotics> for a comprehensive list of articles supporting early IV to PO switch



Additional Benefits of IV to PO

Improved safety outcomes

- Reduces line-related complications

Cost effective

- Decreased hospital length of stay, supply costs, labor

Increased patient satisfaction

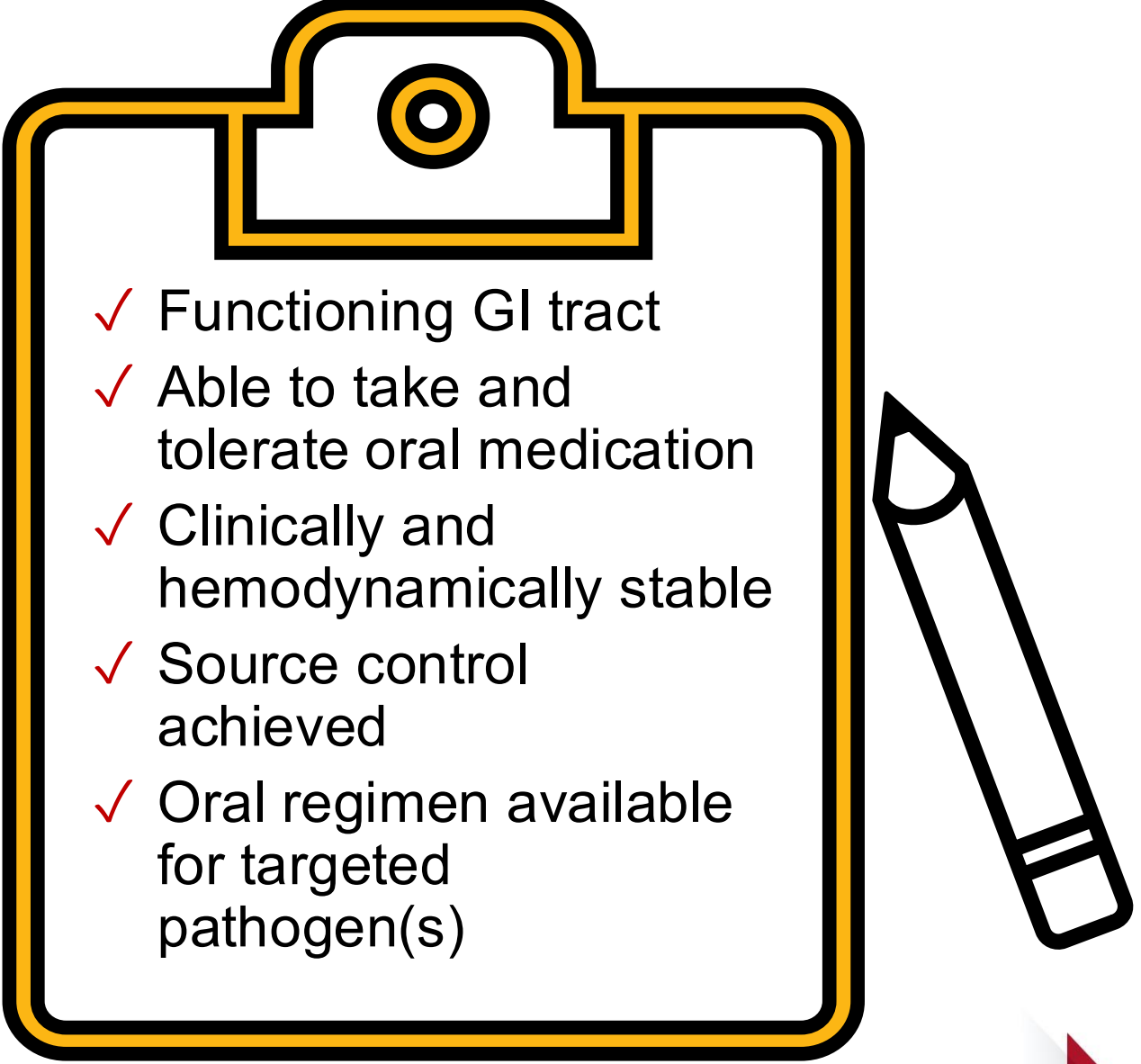
- Greater mobility, comfort, convenience

Environmentally Friendly

- Less waste production
- Antibiotic Waste Impact Calculator:
<https://ecorxchoice.com/?page=home>



Considerations for Oral Step-Down

- 
- ✓ Functioning GI tract
 - ✓ Able to take and tolerate oral medication
 - ✓ Clinically and hemodynamically stable
 - ✓ Source control achieved
 - ✓ Oral regimen available for targeted pathogen(s)



Selecting an Antibiotic

Empiric therapy → use your local antibiogram

Pathogen identified → use reported susceptibilities

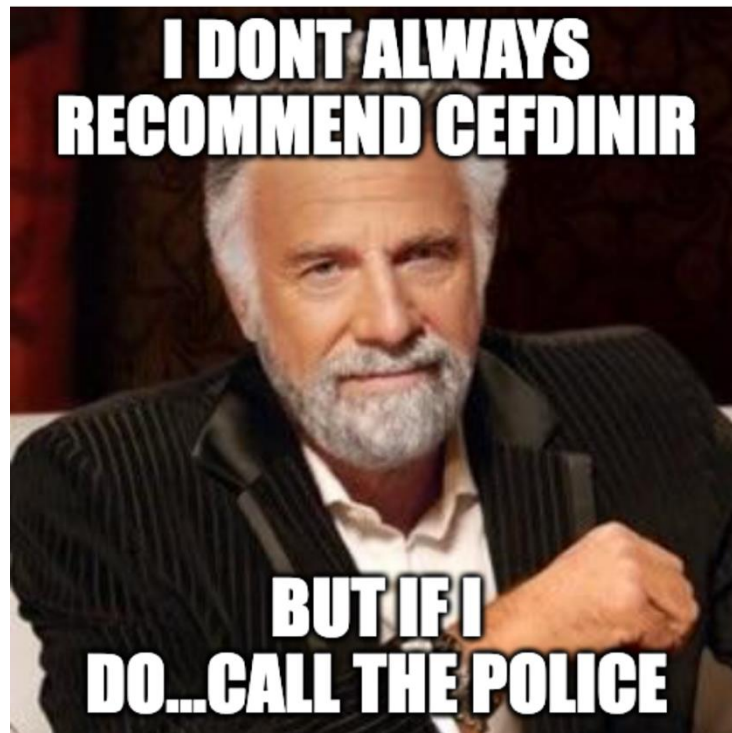
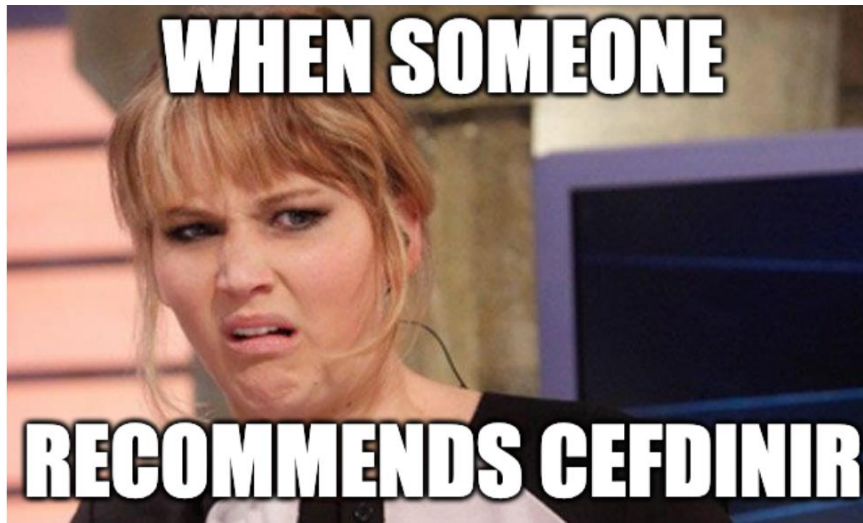
Very Highly
Bioavailable
(~90-100%)

- Linezolid
- Levofloxacin
- Doxycycline
- Cephalexin
- Metronidazole

Highly
Bioavailable
(~80-90%)

- Amoxicillin
- Ciprofloxacin
- Bactrim





Patient Case

70yo M with PMH of penicillin allergy (reaction: anaphylaxis) and obesity (BMI 32), s/p L knee TKA ~2 months ago, now presenting with increased L knee erythema concerning for PJI. A DAIR was performed, cultures have finalized with MSSA, and current plan is for 6wk of cefazolin 2g q8h. It is now post-op day 10 and patient's only barrier to discharge is finding a facility that will accept q8h dosing.

Staphylococcus aureus		
	MIC	Interpretation
Clindamycin	4	Resistant
Daptomycin	0.5	Susceptible
Linezolid	1	Susceptible
Oxacillin	1	Susceptible
Tetracycline	≤ 2	Susceptible
Trimethoprim -Sulfa.	$\leq 0.5/9.5$	Susceptible
Vancomycin	2	Susceptible

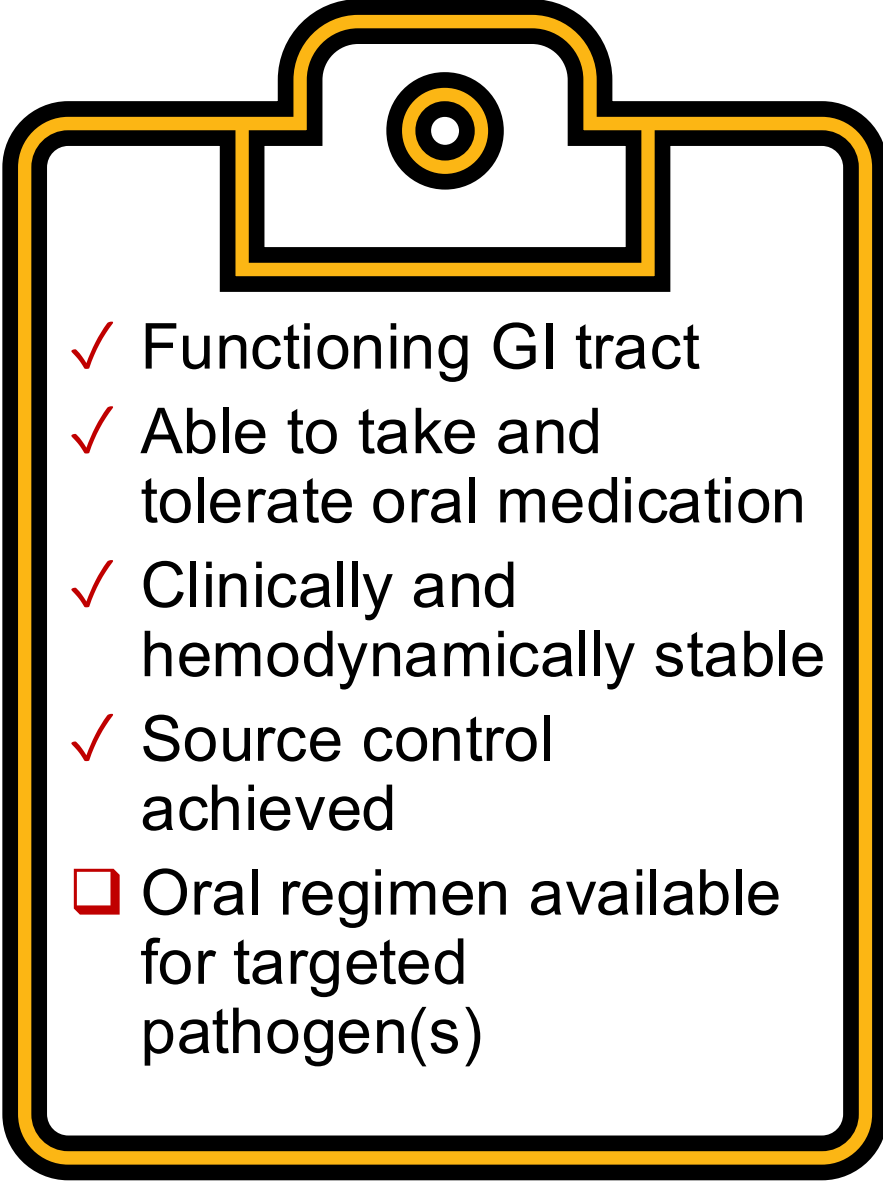
Is this patient eligible for early IV to PO switch?

TKA: total knee arthroplasty

PJI: prosthetic joint infection

DAIR: debridement, antibiotics, and implant retention



- 
- ✓ Functioning GI tract
 - ✓ Able to take and tolerate oral medication
 - ✓ Clinically and hemodynamically stable
 - ✓ Source control achieved
 - ☐ Oral regimen available for targeted pathogen(s)

70yo M with PMH of penicillin allergy (reaction: anaphylaxis) and obesity (BMI 32), s/p L knee TKA ~2 months ago, now presenting with increased L knee erythema concerning for PJI. A DAIR was performed, cultures have finalized with MSSA, and current plan is for 6wk of cefazolin 2g q8h. It is now post-op day 10 and patient's only barrier to discharge is finding a facility that will accept q8h dosing.

TKA: total knee arthroplasty

PJI: prosthetic joint infection

DAIR: debridement, antibiotics, and implant retention



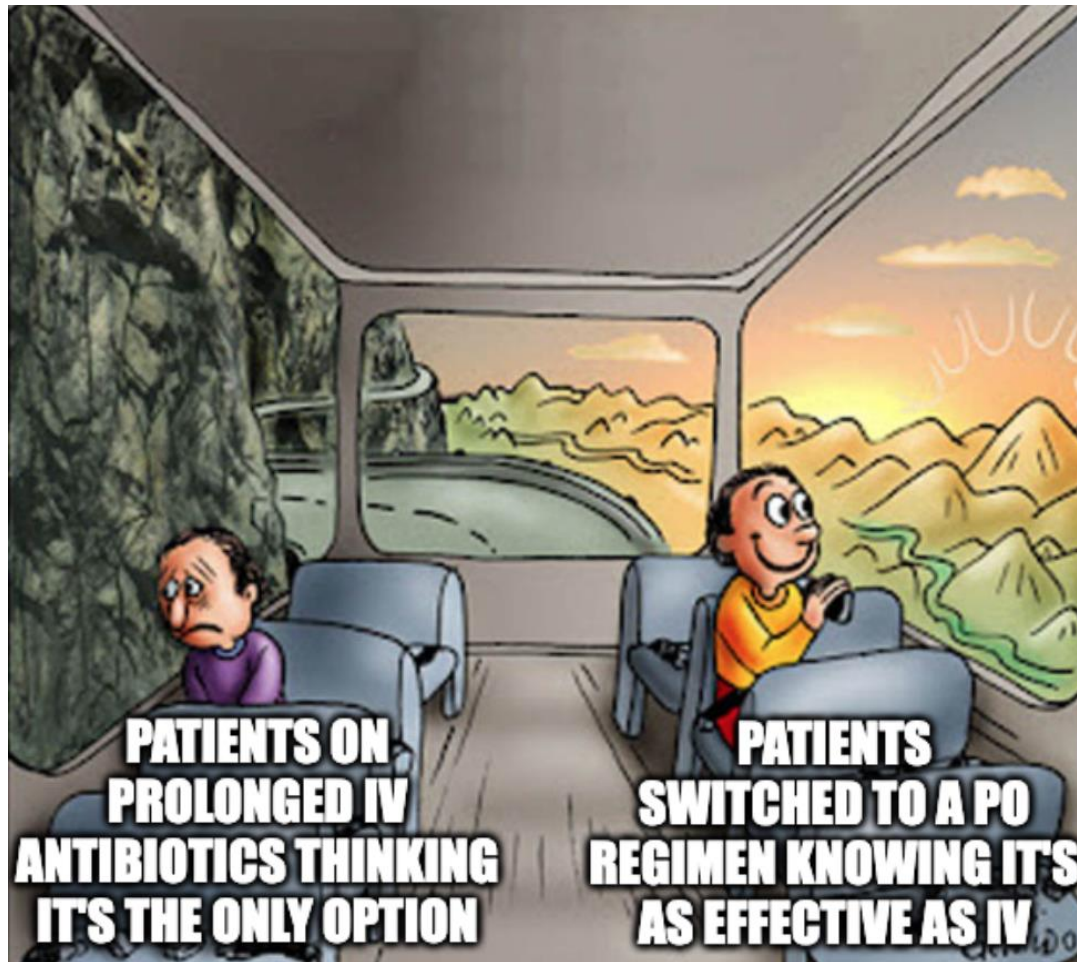
- ✓ Functioning GI tract
- ✓ Able to take and tolerate oral medication
- ✓ Clinically and hemodynamically stable
- ✓ Source control achieved
- ✓ Oral regimen available for targeted pathogen(s)

Staphylococcus aureus		
	MIC	Interpretation
Clindamycin	4	Resistant
Daptomycin	0.5	Susceptible
Linezolid	1	Susceptible
Oxacillin	1	Susceptible
Tetracycline	≤2	Susceptible
Trimethoprim -Sulfa.	≤0.5/9.5	Susceptible
Vancomycin	2	Susceptible

Allergies: Penicillin (anaphylaxis)



Patient-Centered Decision Making



The Pharmacist-Physician Partnership

Pharmacist

- Recommend appropriate PO agent and dosing
- Patient medication counseling



Physician

- Assess and monitor clinical stability
- Document plan
- Patient follow-up appointments (if needed)

► Review IV antibiotics daily to identify candidates for early PO switch ◀

**Successful IV-to-PO programs rely on
multidisciplinary collaboration**



Take Home Messages

- ✓ Review IV antibiotics daily
- ✓ Early oral conversion is safe and evidenced-based
- ✓ IV to PO switch reduces complications, costs, and length of stay
- ✓ Pharmacists and physicians are essential partners in antimicrobial stewardship





**Nebraska[®]
Medicine**

Email: sazimi@nebraskamed.com



References

- 1) Li HK, Rombach I, Zambellas R, Walker AS, McNally MA, Atkins BL, Lipsky BA, Hughes HC, Bose D, Kümin M, Scarborough C, Matthews PC, Brent AJ, Lomas J, Gundle R, Rogers M, Taylor A, Angus B, Byren I, Berendt AR, Warren S, Fitzgerald FE, Mack DJF, Hopkins S, Folb J, Reynolds HE, Moore E, Marshall J, Jenkins N, Moran CE, Woodhouse AF, Stafford S, Seaton RA, Vallance C, Hemsley CJ, Bisnauthsing K, Sandoe JAT, Aggarwal I, Ellis SC, Bunn DJ, Sutherland RK, Barlow G, Cooper C, Geue C, McMeekin N, Briggs AH, Sendi P, Khatamzas E, Wangrangsimakul T, Wong THN, Barrett LK, Alvand A, Old CF, Bostock J, Paul J, Cooke G, Thwaites GE, Bejon P, Scarborough M; OVIVA Trial Collaborators. Oral versus Intravenous Antibiotics for Bone and Joint Infection. *N Engl J Med*. 2019 Jan 31;380(5):425-436. doi: 10.1056/NEJMoa1710926. PMID: 30699315; PMCID: PMC6522347.
- 2) Iversen K, Ihlemann N, Gill SU, Madsen T, Elming H, Jensen KT, Bruun NE, Høfsten DE, Fursted K, Christensen JJ, Schultz M, Klein CF, Fosbøll EL, Rosenvinge F, Schønheyder HC, Køber L, Torp-Pedersen C, Helweg-Larsen J, Tønder N, Moser C, Bundgaard H. Partial Oral versus Intravenous Antibiotic Treatment of Endocarditis. *N Engl J Med*. 2019 Jan 31;380(5):415-424. doi: 10.1056/NEJMoa1808312. Epub 2018 Aug 28. PMID: 30152252.
- 3) Kaasch AJ, López-Cortés LE, Rodríguez-Baño J, Cisneros JM, Dolores Navarro M, Fätkenheuer G, Jung N, Rieg S, Lepeule R, Coutte L, Bernard L, Lemaignan A, Kösters K, MacKenzie CR, Soriano A, Hagel S, Fantin B, Lafaurie M, Talarmin JP, Dinh A, Guimard T, Boutoille D, Welte T, Reuter S, Kluytmans J, Martin ML, Forestier E, Stocker H, Vitrat V, Tattevin P, Rommerskirchen A, Noret M, Adams A, Kern WW, Hellmich M, Seifert H; SABATO study group. Efficacy and safety of an early oral switch in low-risk *Staphylococcus aureus* bloodstream infection (SABATO): an international, open-label, parallel-group, randomised, controlled, non-inferiority trial. *Lancet Infect Dis*. 2024 May;24(5):523-534. doi: 10.1016/S1473-3099(23)00756-9. Epub 2024 Jan 17. PMID: 38244557.
- 4) Wald-Dickler N, Holtom PD, Phillips MC, Centor RM, Lee RA, Baden R, Spellberg B. Oral Is the New IV. Challenging Decades of Blood and Bone Infection Dogma: A Systematic Review. *Am J Med*. 2022 Mar;135(3):369-379.e1. doi: 10.1016/j.amjmed.2021.10.007. Epub 2021 Oct 27. PMID: 34715060; PMCID: PMC8901545.
- 5) Cole A, Aspin J, Laird S, Aciri F, Galley S, Collins M. The environmental impact of intravenous antimicrobial therapies: a comparison of OPAT and inpatient administration care pathways. *JAC Antimicrob Resist*. 2025 Mar 11;7(2):dlaf030. doi: 10.1093/jacamr/dlaf030. PMID: 40070892; PMCID: PMC11894253.
- 6) Landersdorfer CB, Gwee A, Nation RL. Clinical pharmacological considerations in an early intravenous to oral antibiotic switch: are barriers real or simply perceived? *Clin Microbiol Infect*. 2023 Sep;29(9):1120-1125. doi: 10.1016/j.cmi.2023.04.009. Epub 2023 Apr 12. PMID: 37059222.

