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BACKGROUND

- One-third of antibiotic prescriptions in hospitals involve potential prescribing problems such as giving an antibiotic without proper testing or evaluation, prescribing an antibiotic unnecessarily, or giving an antibiotic for too long.¹
- The Centers for Disease Control and Prevention (CDC) established guidelines for appropriate use of vancomycin, which includes serious infections caused by beta-lactam resistant gram-positive bacteria or patients with serious allergies to betalactam antimicrobials.²
- A 2016 CDC study found that rates of vancomycin use increased by 32% within United States hospitals from 2006-2012.³
- Studies have shown vancomycin is commonly prescribed inappropriately, and often is continued without proper clinical indications. 4,5

STUDY PURPOSES

- Evaluate the indication/clinical appropriateness of the continuation of empiric vancomycin therapy.
- Confirm the final microbiology results requiring vancomycin therapy.
- Examine a 72-hour time-out as an effective de-escalation intervention.

METHODS

- Retrospective cohort study from January 2018- October 2018.
- Data collected from the electronic medical record from two community hospitals, Chandler Regional Medical Center and Mercy Gilbert Medical Center.
- Institutional Review Board approved.



Evaluation of Empiric Vancomycin Utilization at 72 Hours Post Admission: is De-escalation of Vancomycin Appropriate?

Positive culture for methicillinresistant Staphylococcus aureus (MRSA)



Presence of infection with or without defined sources with systemic signs of infection*

White Blood Cells <4,000 or >12,000 and/or Temperature \ge 37.5°C

RESULTS

Figure 2. Vancomycin Indications at Day 3 Post-Admission				
	Day 3 Post-Admission			
Vancomycin Indication	Number Patients (N= 160) (%)	Appropriate per criteria (%)	Wound/tissue/ sputum cultures pending (%)	
Positive blood culture	20 (13%)	17 (85%)	N/A	
Skin and Soft Tissue Infection (SSTI)	82 (51%)	54 (65%)	23 (28%)	
Pneumonia (PNA)	37 (23%)	30 (81%)	11 (29%)	
Osteomyelitis	5 (3%)	5 (100%)	2 (40%)	
Joint Infection	6 (4%)	4 (66%)	2 (66%)	
Urinary Tract Infection (UTI)	3 (2%)	2 (66%)	N/A	
Intra-abdominal Infection	4 (2%)	3 (75%)	2 (50%)	
Other	3 (2%)	3 (100%)	1 (33%)	

Figure 3. Final Microbiology Results				
NumberPathogenNumberOfPatients(%)		Pathogen	Number of Patients (%)	
MRSA/MRSE*	36 (23%)	Gram-negative organisms	7 (4%)	
Other Staphylococcus spp.§	28 (18%)	Other (i.e. Micrococcus spp.)	3 (2%)	
Streptococcus spp.	6 (3%)	No pathogen isolated	80 (50%)	

*methicillin-resistant Staphylococcus epidermidis (MRSE), § methicillin-sensitive Staphylococcus spp.

Figure 4. Multivariate Logistic Regression Model Predicting Inappropriate use of Vancomycin at Day 3 Post-Admission				
	Estimate	Odds Ratio (95% Confidence Interval)	P-value	
American Indian/Alaskan Native	1.102	3.01 (1.21, 7.53)	0.0174	
SSTI Diagnosis	1.056	2.87 (1.24, 6.80)	0.0147	



- patients:
- hospital admission (p=0.82).
- assessment.

- hours.
- hours.
- discontinue vancomycin.
- 72-hour antibiotic time-out.



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RESULTS (continued)

• 118 of 160 patients (74%) met appropriate criteria for vancomycin continuation at 3 days post-admission. Of these 118

- 91 (77%) had an infectious diseases consult by day 3 of

-72 patients (61%) had MRSA risk factors (p= 0.99).

- 21 patients (13%) had a culture that was obtained after 72 hours of empiric vancomycin therapy.

• 44 of 160 patients (28%) had pending cultures at 72 hours, and 23 patients (14%) had a known non-MRSA pathogen at time of

Baseline patient characteristics were well-matched between appropriate versus inappropriate per criteria except for Alaskan Native/Native American race (16 patients [14%; p= 0.02]).

CONCLUSIONS

Approximately 25% of patients receiving empiric vancomycin therapy did not meet clinical criteria for continuation beyond 72

Skin and soft tissue infections (SSTI) were the indication most associated with continued vancomycin utilization beyond 72

Although patients were identified as meeting criteria for appropriate continuation of vancomycin at 72 hours post admission, there still may have been an opportunity to

Timely culture obtainment and antibiotic de-escalation

intervention once another pathogen is identified are possible strategies to ensure effective vancomycin de-escalation at a

 Our results identified indications in which empiric vancomycin prescribing can be optimized, and a 72-hour antibiotic time-out may be warranted as a stewardship intervention.

REFERENCES

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