

# Emergency Department Pharmacist Led Methicillin Resistant Staphylococcus Aureus Polymerase Chain Reaction Assay for Vancomycin in Pneumonia

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## Background

- Antimicrobial regimen for methicillin-resistant *Staphylococcus aureus* (MRSA) coverage with antibiotics such as vancomycin are recommended for empiric use in the treatment of suspected MRSA (1-2)
- Nasal MRSA Polymerase Chain Reaction (PCR) assay has shown high negative predictive value for MRSA pneumonia
- Previous retrospective studies in the inpatient setting have demonstrated no difference in hospital mortality or decreased exposure to broad-spectrum antimicrobials (vancomycin) with MRSA PCR utilization for de-escalation of therapy(3-5)
- Early utilization of MRSA PCR by pharmacists in the ED will lead to early de-escalation or avoidance of vancomycin in patients with suspected pneumonia and MRSA risk factors

## Objectives

Utilize MRSA PCR assay by pharmacist in the ED for early de-escalation or avoidance of vancomycin in patients with suspected pneumonia and MRSA risk factors

## Methods

### Design:

Single center, retrospective cohort study

### Groups:

**Control Group:** Historical cohort, no pharmacist MRSA PCR intervention and received IV vancomycin

**Intervention Group:** Pharmacist initiated MRSA PCR assay

### Inclusion Criteria:

- Patients presenting the Emergency Department between 8-1-2019 and 9-30-2020
- ≥ 18 years
- Radiographic diagnosis of pneumonia
- Patients with empiric vancomycin ordered

### Exclusion Criteria:

- Currently on chemotherapy for malignancy or with neutropenic fever
- Patients with lung transplant or cystic fibrosis
- Prior positive MRSA in a blood, sputum culture, or suspected MRSA infection elsewhere
- Patients with concomitant empiric agents with MRSA activity (e.g., linezolid, ceftaroline)

## Methods



### Sample Size:

38 patients (patient enrollment ongoing)

### Primary Endpoints:

- Number of patients who received only one dose of vancomycin prior to MRSA PCR result

### Secondary Endpoints:

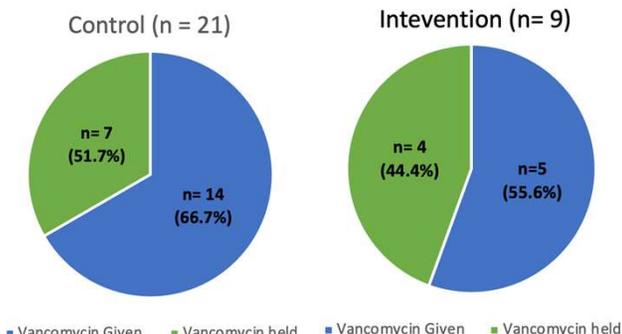
- Number of patients in whom empiric vancomycin was avoided in the Emergency Department
- Number of patients who had positive MRSA sputum or blood culture despite negative MRSA PCR results
- Hospital length of stay and hospital mortality
- Need for vancomycin level monitoring

## Results

	Control (N=29)	Intervention (N=9)	P-Value
Age (yr), m (IQR)	70 (61-82)	83(65-87)	0.19
Sex (male), n (%)	15(51.7)	3(33.3)	0.33
History of tracheostomy, n (%)	4 (13.8)	1(11.1)	1.00
Hemodialysis at admission, n (%)	10 (34.5)	3(33.3)	1.00
Sepsis in the ED, n (%)	9 (31.0)	4 (44.4)	1.00
Septic Shock in the ED, n (%)	4(44.4)	2(50)	1.00
Indication for vancomycin, n (%)			
CAP	12 (8.5)	3 (60)	-
HCAP	1 (7.1)	1 (20)	-
HAP	0 (0)	1 (20)	-
VAP	1 (7.1)	0 (0)	0.27
<b>MRSA PCR ordered in the ER, n (%)</b>	<b>12 (41.4)</b>	<b>8 (88.9)</b>	<b>0.01</b>
MRSA PCR turn around time (min), m (IQR)	129.5 (120-218)	146 (112-159)	0.3
<b>Vancomycin avoided in the ER, n (%)</b>	<b>0 (0)</b>	<b>1 (11.1)</b>	<b>.02</b>
Positive blood or sputum cultures for MRSA, n (%)	0	0	-
Vancomycin level drawn, n (%)	12 (41.4)	2 (22.2)	0.3
Hospital length of stay (days), m (IQR)	7 (3-12)	5 (4-6)	0.97
Death during admission, n (%)	6 (20.7)	1 (11.1)	0.52

## Results Continued

### Number of patients who received one dose of Vancomycin prior to MRSA PCR Result



## Conclusion

- More patients in intervention group had empiric dose of vancomycin held prior to MRSA PCR result
- More patients in the control group had vancomycin levels drawn
- Data collection is currently ongoing, and results presented are preliminary data

## Contact Information

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## Disclosure

The authors have nothing to disclose concerning possible financial or personal relationships with commercial entities

## References

- Metlay JP, et al. Am J Respir Crit Care Med 2019;200(7):e45-67
- Kalil AC, et al. Clin Infect Dis 2016;63(5):61-111
- Cowley MC, et al. Chest 2019;155(1):53-9
- Baby N, et al. Antimicrob Agents Chemother 2017;61(4):02432-16
- Dunaway, et al. International Journal of clinical pharmacy 2018;40:526-32