

Clinical Safety & Effectiveness Cohort # 10

Improving Weight-Based Vancomycin Dosing and Monitoring



SAN ANTONIO

Educating for Quality Improvement & Patient Safety

Financial Disclosure

Elizabeth A. Walter, MD, has no relevant financial relationships with commercial interests to disclose.

Heta Javeri, MD, MPH Fellow, has no relevant financial relationships with commercial interests to disclose.

The Team

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What We Are Trying to Accomplish?

OUR AIM STATEMENT

- 1 a. To improve the use of initial weight-based vancomycin dosing in hospitalized patients by implementing and encouraging the use of a vancomycin dosing order set in Sunrise.
 - Currently, 66% of patients with weight > 100 kg are inappropriately dosed with vancomycin and we aim to decrease this to 50%.
- 1 b. To improve the timing of initial vancomycin trough levels to ensure rapid achievement and maintenance of therapeutic drug levels (TDM).

Project Milestones

 Team Created 	January 2012
 Aim statement created 	January 2012
 Weekly Team Meetings 	February 2012
 Background Data, Brainstorm Sessions, 	March 2012
Workflow and Fishbone Analyses	
 Interventions 	March 2012
 Data Analysis 	May 2012
 CS&E Presentation 	June 2012

Background: Dosing in Obesity

- San Antonio, Texas is one of the most obese cities in the nation.
 - San Antonio obesity rate of 28.2%
 - United States average of 27%¹
- A multicenter evaluation of vancomycin dosing found:
 - 86% of overweight patients (BMI= 25-29.9 kg/m²)
 - 91% of obese patients (BMI ≥ 30 kg/m²) with gfr > 60 mL/min
 - Received a fixed dose of 2 g daily divided into two doses⁴
- A pilot study (n=65) conducted at out institution revealed that only 33% of patients > 100 kg received weight-based vancomycin dosing greater than 30 mg/kg/day.

Background: Guidelines



- Doses of 15-20 mg/kg actual body weight every 8-12 hr are optimal for most patients with normal renal function to achieve the suggested serum concentration.
- A loading dose of 25-30 mg/kg (based on ABW) in seriously ill patients to achieve a more rapid target trough concentration³.
- Trough serum vancomycin concentrations of 15–20 mg/L are recommended for complicated infections (bacteremia, endocarditis, osteomyelitis, meningitis and hospital acquired pneumonia) caused by MRSA.
- Maintain trough > 10 mg/L, based on evidence suggesting that strains with VISA like characteristics (hVISA) may develop³.

Current Status of Therapeutic Drug Monitoring (TDM)

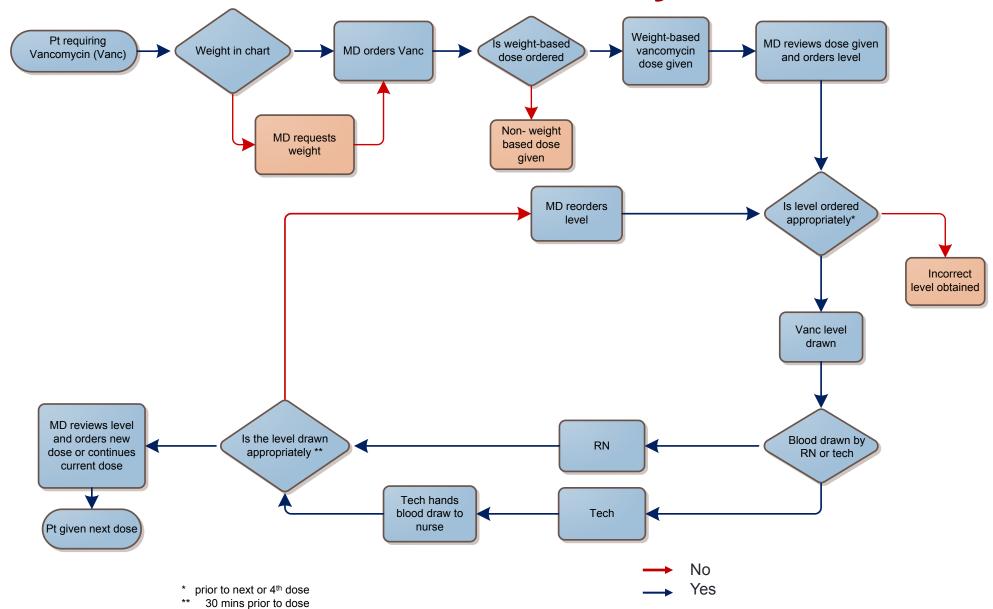
- Large number of vancomycin trough levels incorrectly ordered.
- Difficulty in interpreting results of inappropriately drawn levels.
- Increases unnecessary costs from additional ordering of levels.
- Potential increase length of stay due to inability to ensure target serum concentration prior to patient discharge.

Background: Review of Literature

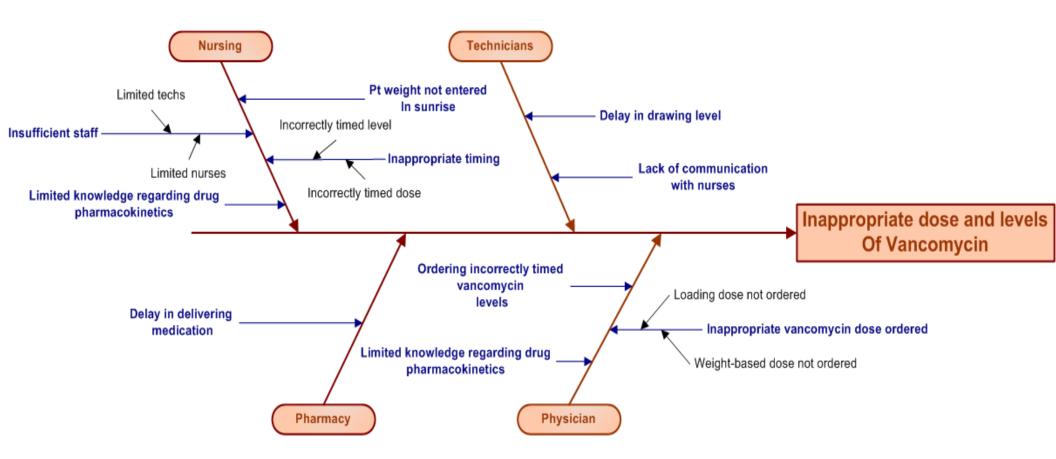
Effect of the implementation of vancomycin dosing and TDM guidelines in computerized prescriber-order-entry (CPOE) system:

- Traugott, et al.
 - Demonstrated a significant increase in the number of appropriately obtained serum vancomycin levels (58% to 68%, p = 0.02)⁵
- McCluggage, et al.
 - Observed a significant increase in the percentage of patients with an initial optimal vancomycin regimen that met nomogram recommendations (36% versus 24%, p = 0.0028)⁶
- · Li, et al.
 - Demonstrated that patients in the post education group on vancomycin dosing protocol had significantly higher, initial median weight-based doses (12.5 mg/kg vs 20.0 mg/kg, p < 0.001), trough concentration (6.8 mg/L to 10.1 mg/L, p= 0.013) and AUC/MICs (262.5 to 365.0, p= 0.001) when compared with the pre-intervention group⁷

Pre-intervention Process Analysis Tool



Decision Making Tool



Plan: Intervention

We implemented the following intervention:

- A vancomycin dosing order set within the computerized prescriber-order-entry (CPOE) system.
- Education of physicians and nursing staff on vancomycin dosing, use of CPOE system and accession of appropriate vancomycin level when indicated.
- Assessment of the effect on vancomycin dosing and therapeutic drug monitoring (TDM).

- 1. System Changes within Sunrise
- Meetings were held with pharmacy staff and sunrise informatics specialists to develop the new order set within Sunrise.
- The newly developed order set was tested on a sample patient list within Sunrise to identify flaws in functioning.
- The order set was then reviewed, accepted, implemented by the P &T committee and incorporated into Sunrise on March 1st, 2012.

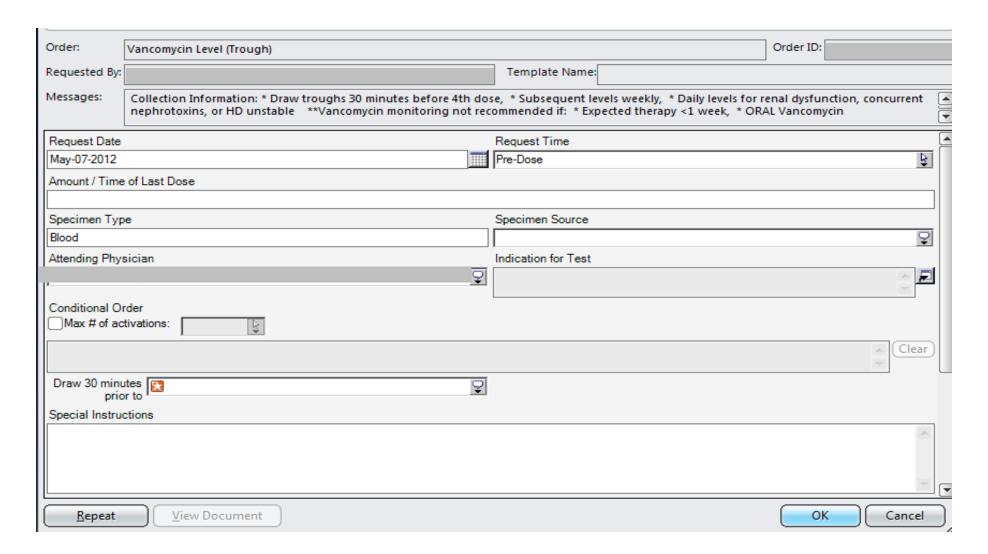
- The order set provided weight-based dosing.
 - Skin/soft tissue infections
 - Serious infections (bacteremia, endocarditis, osteomyelitis, meningitis and hospital acquired pneumonia)
 - Incorporated patient's renal function
- Added a loading dose for serious infections.
- Linked the order for vancomycin trough level to the order set.
 - Default time for vancomycin trough level eliminated
 - Facilitated providers to self-select times for trough levels

Odd dosing and continuous infusion order set were retained.

Order Set

Vancomycin Adu	lt for Serious Inf	ections [2 orders	of 3 are selecte	ed]					
Warnings/Additional	Information								
Pneumonia									
Medications									
Order		Dose	Route	Frequency	PRN PRN Reason	Start Date	Administration Instruction	s	
					?				
Loading Dose - 1						I-			
☐ Vancomycin I		g	IV Piggy Back	Once			If "Red Man" syndrome de	velops, infuse at a lower	
Maintenance Dos		- I-	IN/ Diagram David			M 07 2012	It "D - I M - " I	and and the first of the first	
<u> </u>	n 1 g IV Piggy Back	1 g	IV Piggy Back	Every 8 Hours		May-07-2012	If "Red Man" syndrome de	evelops, infuse at a lower	
	Labs								
	⊞ ✓ Vand	omycin Level (Trou	gh) Pre-Dose [*	Draw 30 minutes p	prior to] [Special Instru	ctions]			
									Relevant Results
Measurements Ht (in)	Ht (cm)	Wt (lb)	Wt (kg)	BS (m)	SA Cr (mg/dL)	ce (Estimated (Cockcroft-(CrCl (es	st) "	Actual • Estimated	Creatinine Serum: 0.66 White Blood Cell Count: 10.7 RBC Count: 3.39
68	172.7	242	109.6	Ž.2	21				Hemoglobin: 10.5
_									
/ancomycin Adult fo	or Skin and Soft Tis	sue Infections [0 or	ders of 2 are select	ted]					
Warnings/Additional Infor	mation								
Medications Order	I D	ose Route	Frequency	PRN PRN Re	eason Start Date	Administration Instruction	ns		
				?					
< 95 kg - 1 item(s)	V Diagraphy	IV/Di-	D1- E 1211		T	K "D - I M - " I	handana fafara at a laura		
Vancomycin 1 g l¹ ⇒ 95 kg - 1 item(s)	V Piggy Back	g IV Piggy	Back Every 12 Ho	ours 🔳		If "Red Man" syndrome d	evelops, infuse at a lower		
Vancomycin 1 g l	V Piggy Back	g IV Piggy	EIV Piggy Back y 8 Hou	ırs 🔲	Т	If "Red Man" syndrome d	evelops, infuse at a lower		
								D D II	
Measurements				Creatinine C	Clearance (Estimated (Cockcro	nft-Gault))		Relevant Results Creatinine Serum: 0.66	
Ht (in)	Ht (cm)	Wt (lb)	Wt (kg)	BSA Cr (ma/dL)	CrCI	(est)	O Actual	White Blood Cell Count: 10.7	
68	172.7	242	109.6	(m²) 2.21	150).5	Estimated	RBC Count: 3.39 Hemoglobin: 10.5	
00	172.7	272	100.0					Hemoglobin: To.5	

Order Set



2. Staff education

≻Nursing

- Nursing educators for floor 8, 9 and MICU were contacted and education method and materials were discussed with them.
- On an average 3 in-service (training sessions) each lasting 10 minutes were scheduled for each of the floors prior to either the morning or evening shift change.
- The training sessions were conducted by members of the team for the initial 3 sessions and then were followed by nurse educators who were present during the initial training.

- Participation of all staff was ensured by an attendance sign in sheet placed prior to all sessions.
- Entering patient height and weight
- Appropriate charting of the time of medication administration and level drawn
- Check out time of vancomycin level lab draw to oncoming nurses at the time of hand off.
- Schedule lab draw prior to X th dose as ordered by physician (e.g next, 4th, etc).
- All levels to be drawn by RN only

>Physicians

- Practitioners were educated on the need for weight-based dosing, especially for patients weighing ≥ 95 kg and the need for loading dose in serious infections.
- A 15 minute presentation was made to the Internal medicine housestaff prior to their daily noon conference.
- A similar presentation was also made to the family medicine housestaff prior to their weekly didatic session.

Study: Determining Change

Types of Measures

- Percent of patients who have weight entered into Sunrise
- Percent of physicians/HCWs using the order set
- Comparing pre and post-intervention values of:
- ➤ Patients not receiving weight-based (<30 MG/KG/D) vancomycin dosing
 - Patients weighing less than 95 kg and greater than 95 kg
- > Patients appropriately receiving loading dose
 - Serious infections
- Patients with appropriate timing of initial vancomycin level
- Time till first appropriate trough level
- >Time till first appropriately timed level

Performance Improvement Design

- Retrospective chart review
- Collection of baseline, pre-intervention data
 - Patients initiated on vancomycin from Dec 1, 2011 to Feb 29, 2012
- Collection of post-intervention data
 - Patients initiated on vancomycin
 from April 1, 2012 to April 30, 2012

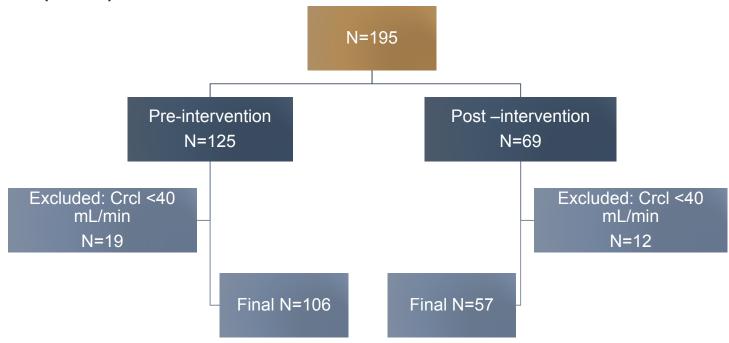


Subjects

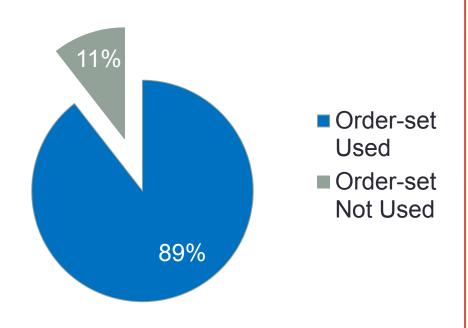
- Adult inpatients at University Hospital (UHS).
 - 18 years or older
 - Admitted to UHS Medicine, Family Medicine or ICU teams
 - Receiving at least 1 dose of vancomycin
- Potential subjects were identified through pharmacy records.
- Information collected:
 - Age, gender, race, height, weight, sCr
 - Diagnosis with culture results
 - Initial dosing regimen, loading dose
 - Data on trough level (timing, level, number)
 - Length of vancomycin therapy
 - Length of stay

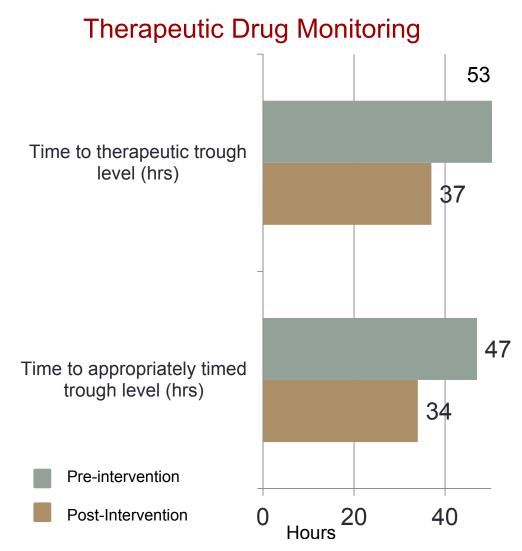
Subjects

- Exclusion criteria:
- Age <18 years</p>
- Hemodialysis and chronic kidney disease with creatinine clearence (Crcl) <40 mL/min

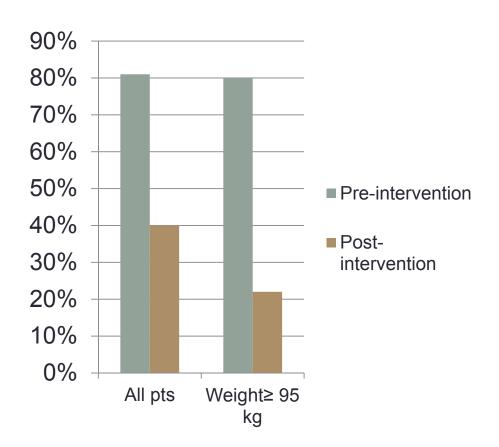


No. of patients prescribed vancomycin using the order set

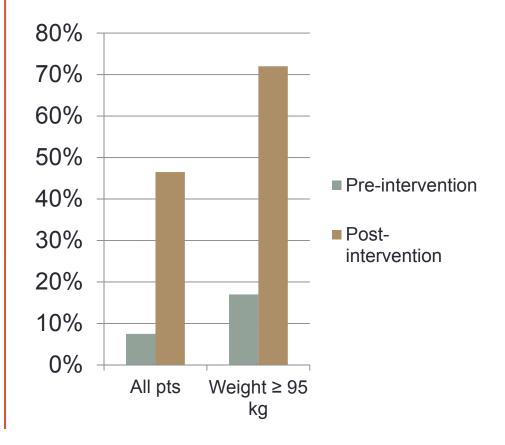




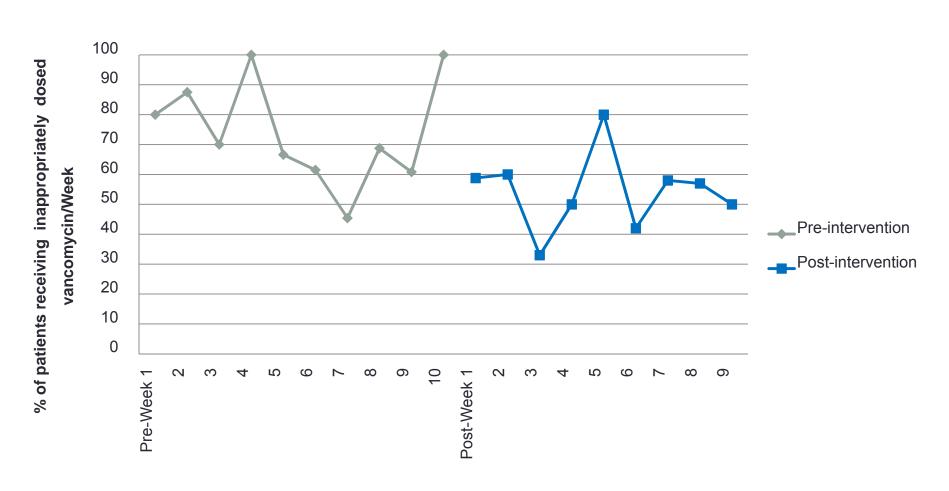
Lower dose vancomycin (1 gm Q12H)



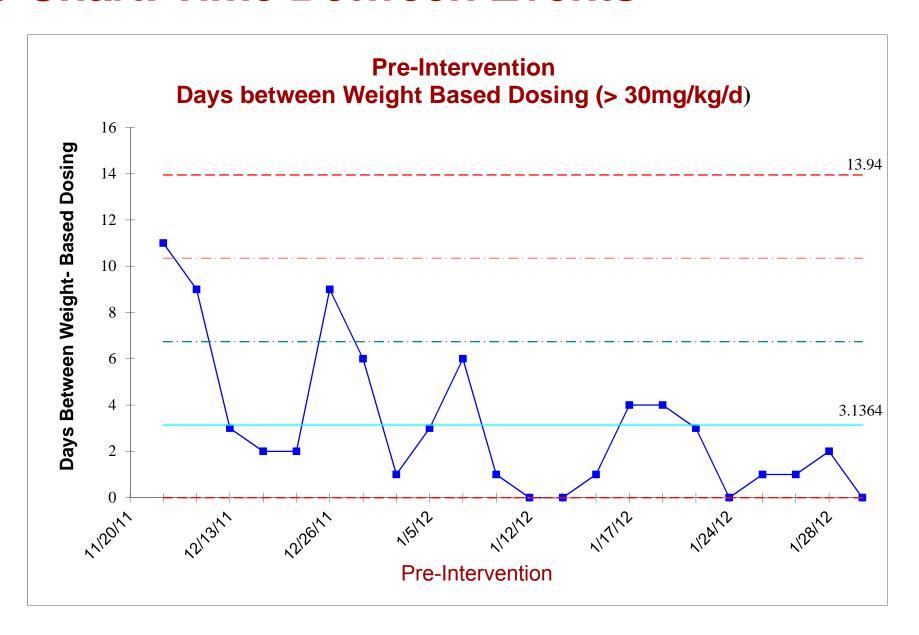
Higher dose vancomycin (1 gm Q8H)

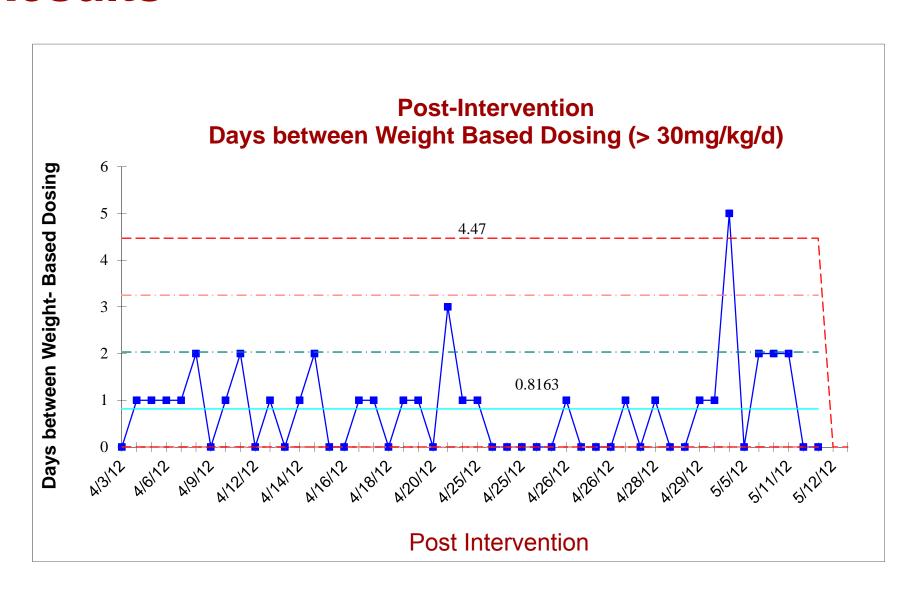


Inappropriate Weight-based (30 mg/kg/d) Vancomycin dosing



G Chart: Time Between Events





• Length of stay was 2 days shorter in the post-intervention group.

Return on Investment

Estimated Project C	Costs	Estimated Project Savings		
Project labor (IT personnel)	\$514	Increased revenue from decreased length of stay	\$31,200	
Implementation costs (RN time)	\$7,000	Saved costs from decreased levels ordered	\$2,543	
		Soft savings from reduced LVN time	\$5,120	

ROI Calculation:

Internal Rate of Return: 109%

Modified Internal Rate of Return: 56%

Act: The Next Step...

- Continuing education.
- Plan to expand staff and physician education to other floors and services of the hospital.
- Post intervention survey.

Limitations

- Retrospective nature of the study may have led to inaccuracies in data collection.
- Small sample size.
- Educational intervention may not have captured all physicians and nursing staff.
- Ensuring continuing education of all staff.
- Caution should be exercised in the population with renal impairment as this were not evaluated in this study.

Conclusion

- The incorporation of a vancomycin dosing order set within the CPOE system in concurrence with provider and nursing staff education led to:
- ➤ Increased the rate of appropriate weight-based dosing.
- ➤ Shortened mean time to achieving appropriate, target serum trough concentrations.
- Decreased overall length of stay.

References

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Thank you!



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