Appropriate Use of Anti-MRSA Antibiotics

Measure Descrip	tion					
Percentage of patients with empiric anti-MRSA antibiotics discontinued when no resistant						
Staphylococcus aureus isolates are present in sterile site cultures						
Measure Compo	nents					
Numerator Statement	Number of denominator eligible patients who have sterile site cultures negative for resistant <i>Staphylococcus aureus</i> isolates <u>AND</u> discontinuation of intravenous anti-MRSA antibiotic at or before 72-of therapy					
	Definitions:Anti-MRSA antibiotic – For the purposes of this measure, anti-MRSA therapy includes Ceftaroline, Dalbavancin, Daptomycin, Linezolid, Oritavancin, Tedizolid, Telavancin, Tigecycline, VancomycinSterile site – For the purposes of this measure, sterile sites include blood, cerebrospinal fluid, pleural fluid, pericardial fluid, peritoneal fluid, joint/synovial fluid, 					
	Numerator Quality-Data Coding Options for Reporting Satisfactorily: IV anti-MRSA antibiotic discontinued at or before 72-hours of therapy when sterile site cultures are negative for resistant <i>Staphylococcus aureus</i> isolates					
	Performance Met: GXXXX:Documentation of discontinuation of IV anti-MRSA antibiotic at or before 72-hours of therapy after sterile site cultures are negative for resistant Staphylococcus aureus isolates					
	<u>OR</u>					
	IV anti-MRSA antibiotic NOT discontinued at or before 72-hours of therapy when					
	sterile site cultures are negative for resistant <i>Staphylococcus aureus</i> isolates					
	Medical Performance Exclusion: GXXXX:Documentation of medical reasons for not discontinuing IV anti-MRSA antibiotic at or before 72-hours of therapy after sterile site cultures are negative for resistant Staphylococcus aureus isolates					
Denominator	Inpatients age 18 years or older with the RxNorm Code for Vancomycin, Linezolid,					
Statement	Daptomycin, Tigecycline, Oritavancin, Dalbavancin, Telavancin, Tedizolid, Ceftaroline Iniectable Solution					
Denominator	- Patients with beta-lactam antibiotic allergies					
Exclusion	 Patients who expire prior to clinical isolate results Patients who transfer to a different hospital prior to obtaining clinical isolate results Pediatric specific units and free standing pediatric hospitals 					

Denominator	N/A				
Exception					
Rationale for	Resistant Staphylococcal aureus infections have increased to an epidemic level,				
the Measure	affecting many countries in the healthcare and community settings. While infections				
	caused by resistant and non-resistant S. aureus are both of grave concerns,				
	methicillin-resistant S. aureus (MRSA) strains have been associated with lethal				
	infections and worse clinical outcomes when compared to methicillin-sensitive S.				
	aureus strains. Empirical therapy for suspected S. aureus infections have been				
	antibiotics are not efficacious against various common S aureus infections, such as				
	MRSA. Although vancomycin is the preferred treatment for serious MRSA infection.				
	prolonged, persistent, or recurrent bacteremia during therapy, high rates of				
	microbiological and clinical failures, nephrotoxicity, and increasing prevalence of non-				
	susceptible strains limit vancomycin's effectiveness. New antimicrobial discovery and				
	development is essential but is unlikely to resolve drug resistance. The overuse and				
	misuse of antibiotics is clearly a contributing factor to antimicrobial resistance and to				
	effectively impede the rise of resistance more rapid diagnostics, a better				
	understanding of pathogenesis of staphylococcal disease, and non-antimicrobial				
	approaches, antimicrobial stewardship, to prevent and treat infections are urgently				
	needed.				
	Chambers HF. Delo FR. Waves of Resistance: Staphylococcus aureus in the Antibiotic Era. Nat				
	Rev Microbiol. 2009; 7: 629-641.				
Supporting	"Vancomycin has been the mainstay of parenteral therapy for MRSA infections. However, its				
Guideline &	efficacy has come into question, with concerns over its slow bactericidal activity, the emergence of resistant strains, and possible "MIC green" among susceptible strains, "				
Other Evidence					
	"Vancomycin kills staphylococci more slowly than do β -lactams in vitro, particularly at higher				
	inocula (107–109 colony-forming units) and is clearly inferior to β -lactams for MSSA				
	bacteremia and infective endocarditis."				
	 Eld, C et al. Clinical Practice Guidelines by the Infectious Diseases Society of America for the Treatment of Methicillin-Resistant Staphylococcus aureus Infections in Adults and Children. Clin 				
	Infect Dis 2011; 52.				
	"Though the current recommendations continue to strengly roly on vancomycin as a standard				
	empiric choice in the setting of severe/invasive infections, alternative therapies exist with				
	studies supporting their non-inferiority. This includes the use of linezolid in pneumonia and				
	severe skin and skin structure infections (SSSI) and daptomycin for MRSA bacteremia,				
	endocarditis, SSSIs and bone/joint infections. Additionally, concerns continue to arise in				
	regards to vancomycin, such as increasing isolate MICs, and relatively high rates of clinical failures with vancomycin. Thus, the growing interest in vancomycin alternatives, such as				
	ceftaroline, ceftobribole, dalbavancin, oritavancin, and tedizolid, and their potential role in				
	treating MRSA infections."				
	VanEperen AS, Segreti J. Empricial therapy in Methicillin-resistant Staphylococcus Aureus				
	infections: An Up-To-Date approach. J Infect Chemother. 2016; 22:351-9.				
	"Daily review of administration of antimicrobials targeting MRSA was highly effective in				
	improving clinical outcomes by optimizing early antimicrobial therapy."				
	Niwa T, et al. Early optimization of antimicrobial therapy improves clinical outcomes of patients administered ecentre to reating as the life excitate to Clearly				
	auministered agents targeting methicillin-resistant staphylococcus aureus. J Clin Pharm Ther. 2016; 41:19-25.				

Measure Importance							
Relationship to	The desired outcome of this measure is the appropriate use of antibiotics for the						
Desired	treatmer	nt of non	-resistant Staph aureus infections. The Clinical Practice Guidelines by				
Outcome	the Infectious Diseases Society of America for the Treatment of Methicillin-Resistant						
	Staphylo	Staphylococcus aureus Infections in Adults and Children cites multiple studies					
	highlight	ing the i	nferior efficacy of vancomycin against non-resistant S. aureus				
	infection	infections (MSSA) compared to beta-lactam antibiotics. If the administration of					
	powerfu	powerful antibiotics indicated for resistant S. aureus, such as vancomycin, is					
	continued without the presence of resistant S. aureus, it can contribute to bacterial						
	for E coli	resistance and render the current pipeline of antibiotics ineffective, which is the case					
	antibioti	for E.coll bacteria carrying the <i>mrc-1</i> gene. The Appropriate Use of Anti-MRSA					
	anumous metric encourages clinicians to assess proad-spectrum antibiotic therapy						
	according to the clinician's medical expertise. The appropriate use of anti-MRSA						
	antibiotic therapy will limit patient exposure to unnecessary drugs as well as improve						
	the efficacy of treatment.						
Opportunity	Antibiotic resistance is a major global public health threat that has emerged with the						
for	inappropriate prescription and use of antibiotics [1, 2]. The Center for Disease Control						
Improvement	and Prevention's Antibiotic Resistance Threats in the United States, 2013 report states						
	that up to 50% of all antibiotic prescriptions are not needed or optimally effective as						
	prescribed. In 2014, the CDC's Antibiotic Resistance Threats in the United States, 2013						
	report as well as the Report to the President on Combating Antibiotic Resistance,						
	Septemb	September 2014 has highlighted the urgency for combating antibiotic resistance with					
		clinical actions that encourage the judicious use of antibiotics.					
	1. Deuster S. Roten I. Muehlebach S: Implementation of treatment guidelines to support judicious use						
	of ant	of antibiotic therapy. Journal of Clinical Pharmacy and Therapeutics 2010, 35(1):71-78.					
	2. Gooss	2. Goossens H, Ferech M, Vander Stichele R, Elseviers M: Outpatient antibiotic use in Europe and					
Exception	association with resistance: a cross-national database study. The Lancet 2005, 365(9459):579-587.						
Justification							
Measure Designa	ation						
Measure Purpose		•	⊠Quality improvement				
		•	⊠Accountability				
		•	Пмос				
Type of Measure	9	•	⊠Process				
		•	□Outcome				
		•					
National Quality		•	⊠Clinical Process-Effectiveness				
Strategy		•	⊠ Patient Safety				
Priority/CMS		•	Patient Experience				
Measure Domain		•	Care Coordination				
(cneck all that apply)		•	⊠Efficiency: Overuse				
		•	⊠Efficiency: Cost				
		•	Population & Community Health				

Level of	• Individual clinicians
Measurement	• 🛛 Clinician groups
(check all that apply)	 Hospitals Outpatient/ED
Care setting	Emergency Departments
(check all that apply)	• Urgent Care
	 Harmonize with other care settings
	 Physician Office Based Measures
	 ○ ⊠Hospital Level Measures
Data source	• Electronic Health Record (EHR) data
(check all that apply)	 Administrative Data/Claims (inpatient, outpatient, or multiple-
	source claims)
	 Paper medical record/chart abstracted
	• 🗆 Registry