# **Antimicrobial Stewardship Program Retreat**November 1, 2024

## Agenda

- Year in Review
- 2024 Priority Matrix
- Current Projects Review
- NHSN updates
  - AU/AR Data Review, Benchmarking
- Prospective Audit and Feedback
- Micro updates
- IDTS
- New Potential Projects
- 2025 Priority Matrix



## Year in Review

## **Accomplishments**

- Blood culture shortage
- Critical Shortage IV fluids
- Perioperative guidance and data
- Nudges
- Next Day Clinic Pathways
- Educational efforts: multiple grand rounds, competency-based modules (nursing)
- Updated HS 1444
- CDPH SDOH Study, submitting data
- Pemibivart implementation
- Caspofungin --> Micafungin

- Submitted CDPH honor roll
- Successful AR validation for NHSN
- Enrollment in AHRQ/JHU project for ASP in Telehealth
- BCID approved!
- Vaccine Oversight Committee
- Intraamniotic infection updated guidance
- Karius paper



## **UC –Wide Beta-Lactam Allergy Project**

Beta-Lactam Utilization Dashboard was created in collaboration with UCDWH since July 2022 to date and compared between FY23 and FY24 YTD.

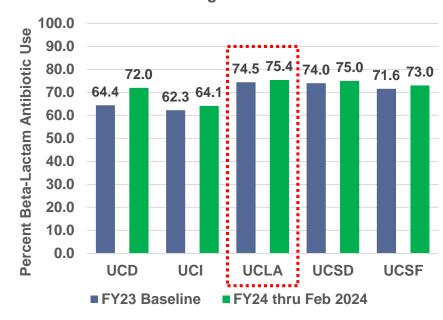
**Numerator**: # of encounters with betalactam allergy history that received any beta-lactam antibiotic dose

**Denominator**: # of encounters with betalactam antibiotic flag where the patient received any antibiotic dose.

Progress: all 5 UC sites improved/ increased beta-lactam antibiotic use in beta-lactam allergic patients in FY24 to date.

	Change since FY23
UCD	<b>↑</b> 7.6%
UCI	<b>1.9%</b>
UCLA	<b>1.0%</b>
UCSD	<b>1.0%</b>
UCSF	<b>1.7%</b>

## Beta-Lactam Antibiotic Use in Beta-Lactam Allergic Patients





## UC Collaborative ID: July 2023-June 2024 Highlights

Accomplishments	New ASP program manager - Bushra Rahman     Pemivibart roll out     Caspo/Mica BPA     Caspofungin to Micafungin conversion
UCLA Health	NHSN AR implementation Next Day Clinic Guideline Development Olorofim/Fosmanogepix Expanded Access IRB protocol Firstline clinical pathway overhaul Ortho-ID case conferencing Competency based modules for staff Allergy Delabeling in Postpartum Increasing WBC threshold for urinalysis to reflex to urine culture (collaboration with laboratory stewardship, microbiology, infection prevention, antimicrobial stewardship) Mycobacterium mucogenicum outbreak investigation, UCLA Santa Monica Medical Center (collaboration with infection prevention, facilities, microbiology) CPO screening for UCLA patients (collaboration with infection prevention, microbiology) Urine retention management standardization (collaboration with infection prevention, nursing, urology) Creation of physician mini-root cause analyses for C difficile, CAUTI and CLABSIs at UCLA (infection prevention, health system CMOs) Foley catheter exchange prior to collecting urine cultures (infection prevention, nursing, Information Services & Solutions, urology)

### <u>Awards</u>

- UCLA Apple for Preceptor Award
  - Ethan Smith, Christine Pham, Meganne Kanatani
- Ethan Smith Nominated for and accepted chair position for CDPH Antimicrobial Stewardship/Antimicrobial Resistance Subcommittee of the Hospital-Acquired Infection Committee
- Ethan Smith UCLA ID Fellow Class of 2024 "Fellows First Award



## UC Collaborative ID: July 2023-June 2024 Highlights



UCLA Health	Tara Vijayan, MD, MPH	<ul> <li>"#9: Arts &amp; Grafts". Febrile: A Cultured Podcast</li> <li>Sabbath Town Hall on Covid Vaccines</li> <li>AHA sponsored Inglewood Active Community Town Hall</li> <li>Covid Vaccine Townhall</li> <li>Carnegie Science Center Covid-19 Vaccine Panel Series</li> <li>Work for various decarceration groups (prison law firms, ACLU, Federal Defenders of San Diego) to reduce COVID in prisons</li> </ul>
	Ishminder Kaur, MD	<ul> <li>Speaker for California Association of Neonatologists and AAP District IX Section on Neonatal-Perinatal Medicine Cool Topics in Neonatology Conference</li> <li>Speaker for Tower Health Maternal Child Health Virtual Conference</li> <li>AAP Speaker for California Chapter</li> <li>Speaker for California Pediatric ASP Meeting</li> </ul>
	Matthew Davis, PharmD	<ul> <li>Remdesivir SIDP Video</li> <li>American College of Medical Toxicology Webinar</li> <li>Contagion Live Interview</li> <li>HHSP Covid Therapeutics Update</li> </ul>



## UC Los Angeles Posters

- Jasdhaul M, Fulgentes GE, Hsueh L, Brewer G, Dogan T, **Prabaker K**. Interdisciplinary Urine Culture Stewardship: Its Impact on Urine Cultures and CAUTI Rate [Abstract #87428]. National Teaching Institute & Critical Care Exposition, Philadelphia 2023
- Parti U, Lewis V, Ison RP, **Uslan DZ**, Bisht A, Walton S, Turay S, Wellbaum D, Mugford Y, de St. Maurice A, **Prabaker K**. Efficacy of Empiric Contact Precautions for Patients from High Risk Facilities [Abstract #97]. Society of Healthcare Epidemiology of America, Houston 2024
- Jasdhaul M, Dogan T, Bennett K, Fulgentes G, **Prabaker K**. Improving Catheter Associated Urinary Tract Infection (CAUTI) Outcomes Through Physician Engagement [Podium presentation]. UCLA Research and Evidence-Based Practice Conference, Los Angeles 2024
- Zamora, R, Dickson, D, **Vijayan, T**. "A Case of Pneumocystis Pneumonia Diagnosed Via Plasma Cell-Free DNA." Unusual Chest Infections Rapid Fire Case Reports. American College of Chest Physicians Annual Meeting, Hawaii, October 2023.
- Shields R, Cifuentes R, Claeys K, DeSear K, Gallagher J, Gregory E, Heil E, Hickey C, Klatt M, Kline E, Kubat R, Kufel W, Lee J, Lim A, Lingg T, MacDougall C, Mathers A, McCreary E, Moore W, Olson S, Oxer J, Pearson J, Pham C, Polk C, Satlin M, Satola S, Shah S, Solanki Y, Tamma P, Vega A, Veena V, Veve M, Wangchinda W, Witt L, Wu J, Pogue J. Efficacy of Ceftazidime-Avibactam versus Ceftolozane-Tazobactam for multidrug-resistant Pseudomonas aeruginosa infections in the United States (CACTUS) among immunocompromised. Poster Abstract. ECCMID, Barcelona, Spain, 2024. -Shields R, Cifuentes R, Claeys K, DeSear K, Gallagher J, Gregory E, Heil E, Hickey C, Klatt M, Kline E, Kubat R, Kufel W, Lee J, Lim A, Lingg T, MacDougall C, Mathers A, McCreary E, Moore W, Olson S, Oxer J, Pearson J, Pham C, Polk C, Satlin M, Satola S, Shah S, Solanki Y, Tamma P, Vega A, Veena V, Veve M, Wangchinda W, Witt L, Wu J, Pogue J.
- De La Reyes G, **Smith E**, Clarito L. Vancomycin Cerebrospinal Fluid (CSF) Concentration in Correlation with Standard Area Under the Curve (AUC) Monitoring Utilizing Bayesian Modeling Software for Patients with Central Nervous System (CNS) Infections. Vizient, Anaheim, California, 2023.
- Angela Castro, **Christine Pham**, Adonia Eskandari, Diep Phan, **Ishminder Kaur**. Retrospective Study on the Impact of Ongoing or Prior IV Vancomycin Therapy on MRSA Nares Test Results: A Time Based Analysis. Poster Abstract. Vizient, Anaheim, California, 2023.
- Beaulac K, Mentler P, Nagel J, Postelnick M, Patel PC, McCoy C, Kinn P, Casaus D, **Smith EA**. Benchmarking Clinical Outcomes by Antimicrobial Spectrum in Patients with Community-Acquired Pneumonia. IDWeek 2024. Los Angeles, California, 2024.
- De Los Reyes G, Blanco MB, Vespa P, **Smith E**, Wherry C, Johnson-Black P, Almahmoud M, Chaudhari A, Ibekwe E, Clarito L. Vancomycin Cerebrospinal Fluid (CSF) Concentrations Achieved with Bayesian Model-Guided Area Under the Curve (AUC) Dosing for Central Nervous System (CNS) Infections. Neurocritical Care Society, San Diego, California, 2024.
- Davis M, Kufel J, Kufel W, Ross J, Oleksiuk L, Ours R, **Smith E, Pham C**, Trisler M, Tverdek F. Infectious Diseases Pharmacist Curbsides: Questions Infectious Diseases Providers Ask Infectious Diseases Pharmacists. IDWeek 2024. Los Angeles, California, 2024.



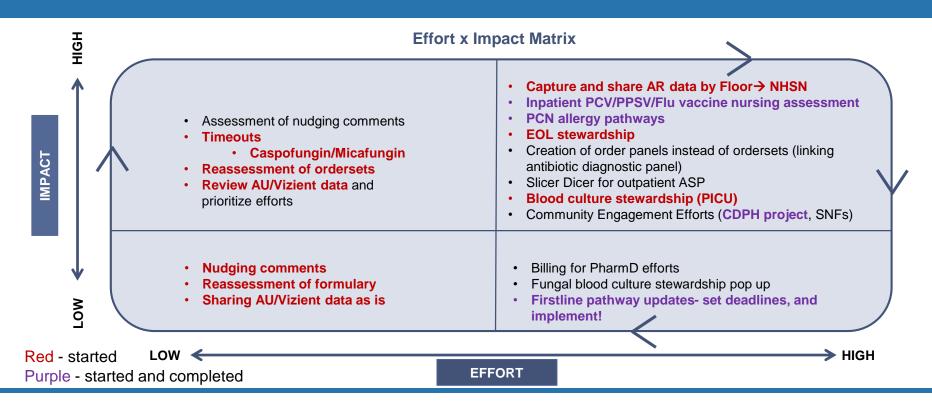
## UC Los Angeles Publications

- Daniel Karlin, Christine Pham, Daisuke Furukawa, Ishminder Kaur, Emily Martin, Olivia Kates, Tara Vijayan. State-of-the-Art Review: Use of Antimicrobials at the End of Life. Clinical Infectious Diseases, Feb 2024: ciad735, <a href="https://doi.org/10.1093/cid/ciad735">https://doi.org/10.1093/cid/ciad735</a>
- Daniel Karlin, Christine Pham, Daisuke Furukawa, Ishminder Kaur, Emily Martin, Olivia Kates, Tara Vijayan. Executive Summary: State-of-the-Art Review: Use of Antimicrobials at the End of Life. Clin Infect Dis. 2024 Feb 1:ciad737. doi: 10.1093/cid/ciad737. Epub ahead of print. PMID: 38301074.
- Melgar M, Abrams JY, Godfred-Cato S, Shah AB, Garg A, Strunk A, Narasimhan M, Koptyev J, Norden A, Musheyev D, Rashid F, Tannenbaum R, Estrada-Y-Martin RM, Patel B, Karanth S, Achenbach CJ, Hall GT, Hockney SM, Caputo M, Abbo LM, Beauchamps L, Morris SB, Cifuentes RO, de St Maurice A, Bell DS, Prabaker KK, Sanz Vidorreta FJ, Bryant E, Cohen DK, Mohan R, Libby CP, SooHoo S, Domingo TJ, Campbell AP, Belay ED. A multicenter retrospective cohort study to characterize patients hospitalized with MIS-A and COVID-19 in the United States, 2020-2021. Clin Infect Dis. 2023 Nov 17;77(10):1395-1405. doi: 10.1093/cid/ciad374. PMID: 37384794: PMCID: PMC10654854
- Gray HK, Beaird OE, **Smith EA**, Schaenman JM, Yang S. Domestically Acquired NDM-1–Producing Pseudomonas aeruginosa, Southern California, USA, 2023. EID Journal. 2023;29(11):2382-5.
- Vijayan, T, Currier, J. "Realising Long-Acting ART as Firstline Therapy." Lancet HIV, invited, published September 10, 2023 PMID: 3756720 Vijayan T. What Did I Miss? Clin Infect Dis. 2023 Oct 4 PMID: 37791971
- Kang D, Huang CX, Yuen AD, Norris KC, Vijayan T. Do hospitals that participate in COVID-19 research differ from non-trial hospitals? A cross-sectional study of US hospitals. Trials. 2023 Aug 7:24(1):504. doi: 10.1186/s13063-023-07450-6. PMID: 37550662: PMCID: PMC10408090.
- Olivo-Freites C, Davar K, Gallardo-Huizar O, Vijayan T, Younes R. Case Report: Cardiovascular Manifestations due to Flea-Borne Typhus. Am J Trop Med Hyg. 2023 Dec 4:110(1):150-154. doi: 10.4269/aitmh.22-0794. PMID: 38052087: PMCID: PMC10793017.
- Sherwood, K and Vijayan, T. "Mycobacterium Avium Complex." ClinicalKey Website. Elsevier, 2023
- **Prabaker, KK.** Vascular Catheter Infections. In: Margaret Hessen. Clinical Overviews. Elsevier. (Accessed on September 8, 2023.) Smith EA, Jariwala R. Precision in Practice: Optimizing Probability of Beta-Lactam Target Attainment with Prolonged Infusions and Therapeutic Drug Monitoring. Pharmacy Times: 21 March 2024.
- Smith EA, Jariwala R. Precision in Practice: Optimizing Probability of Beta-Lactam Target Attainment with Prolonged Infusions and Therapeutic Drug Monitoring. Pharmacy Times. 21 March 2024.
- Smith EA, Jariwala R. Standardized Stewardship" Navigating the new TJC Standards for Antimicrobial Stewardship. Vizient VerifiedRx Podcast –
   https://podcasts.apple.com/dk/podcast/standardized-stewardship-navigating-the-new-tjc/id1544167927?i=1000642695995.

   2024 January.



## 2024 Prioritization Matrix





## **Current Projects**

## **Current Projects: Pharmacy**

- Inpatient Penicillin Delabeling (Ob, expanding)
- Optimizing desensitization ordersets
- Updating penicillin allergy alerts based on side chain similarities vs. Beta-lactam class
- EA: fosmanogepix, olorofim, ibrexafungerp, clofazimine
- Optimizing Intraperitoneal antimicrobials order set
- Inpatient and outpatient vaccine standardization
  - Example GSK contracting: Heplisav, Pediatric DTaP-IPV
- PrecisePK integration with EPIC for 2025 and getting a UC system wide contract discount
  - Updated flowsheet/note templates for pharmacists
- Neutropenic fever guidance
- Standalone antibiotic panels and automation for MRSA nares w/ PNA

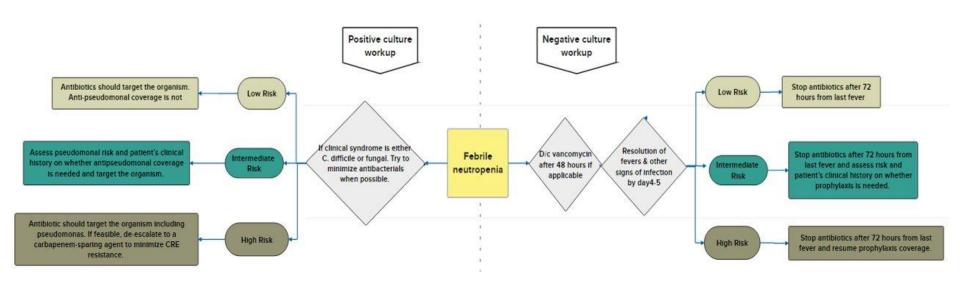


## **Micafungin BPA Data**

- BPA performance
  - In the last 6 months, the BPA has fired 350 times on 86 patients
    - Of the 350 fires, an order was placed 102 times on 58 patients
      - BPA is accepted but ID consult order is not placed
  - BPA allows users to defer multiple times (no maximum- cannot be built in)
  - BPA logic to wait 72 hours was not working in the beginning (time frame confirmation pending)
- Next steps:
  - Add 'discontinue micafungin' button to the BPA
  - Add link to paging interface within BPA
  - Exclude indication "prophylaxis, immunocompromised host" from BPA



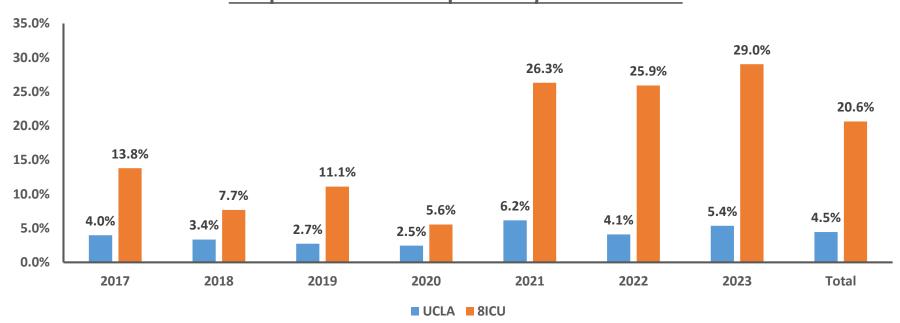
## **Febrile Neutropenia Guidance**





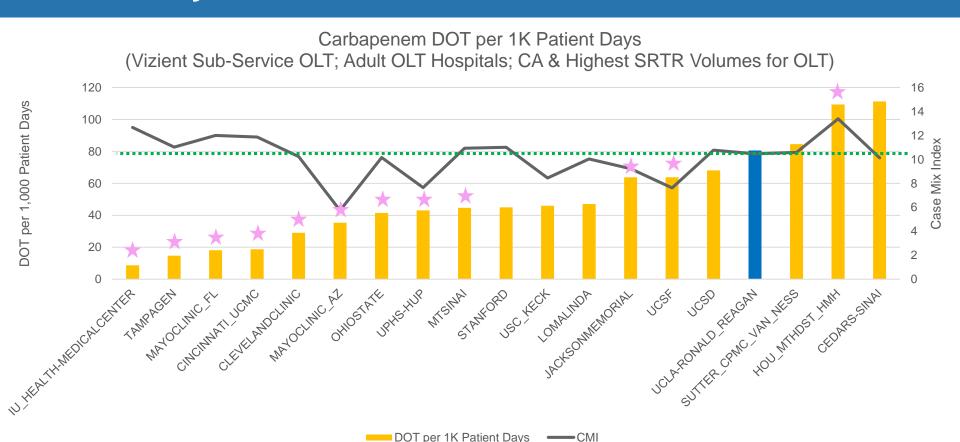
## **8ICU – Planning for Enhanced ASP Activities**





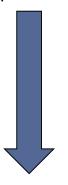


# Vizient: Liver Transplant Carbapenem DOT per 1,000 Patient Days



# OLT Carbapenem Mean Duration Per Case (Vizient) and Hospital Resistance Rates (Personal Communication)

Low Vizient Carbapenem DOTs

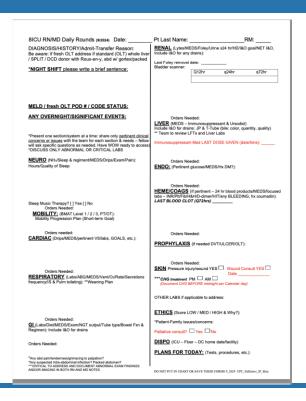


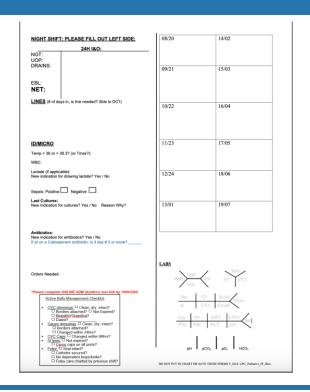
High Vizient Carbapenem DOTs

Hospital	Mean Duration per Case	ESBL%*	CRE%*
IU_HEALTH-MEDICALCENTER	11.5	17	3
TAMPAGEN	9.4	19	
MAYOCLINIC_FL	9.6	21	1
CINCINNATI_UCMC	7.7	18	1
CLEVELANDCLINIC	8.7	8.5	1.5
MAYOCLINIC_AZ	6.4	19	2.5
<b>OHIOSTATE</b>	5.5		
<mark>UPHS-HUP</mark>	8.5	10.1	
<mark>MTSINAI</mark>	16.2	35.5	5.5
STANFORD	13.6	13.5	1
USC_KECK	9.8	28	
LOMALINDA	5.4	19	5
JACKSONMEMORIAL	14.4		
UCSF	6.3	23	2.5
UCSD	11.7	26.5	
UCLA-RONALD_REAGAN	16.2	23.5	3.5
SUTTER_CPMC_VAN_NESS	11.3	12	
HOU_MTHDST_HMH	17.8	21.5	3
CEDARS-SINAI	16.6	(↓)	(↓)



## **Meropenem in Liver Transplant Unit**







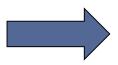
## Perioperative guidance

### Confounders/Mediators



### Exposure:

- Patient factors (DM, BMI, co-morbid, allergies)
- Surgery Factors: length of surgery, vasopressor use
- Hospital factors: infections in hospital
- Equity Factors



## Primary Outcomes/

Exposure for Secondary outcomes

- Duration of periop abx
- Periop abx choice



## **Secondary Outcomes**

Incident 90d C.diff infection

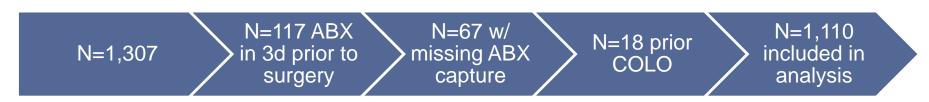


Confounders

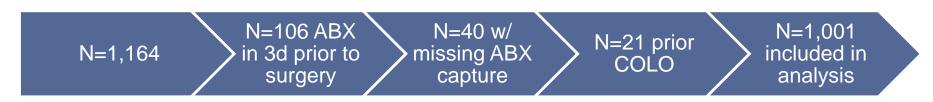


## **Perioperative Antibiotic Guideline Update**

### Pre-Intervention Cohort (2/1/22-2/28/23):



### Post-Intervention Cohort (3/1/23-2/2/24):



## **Perioperative Antibiotic Guideline Update**

Characteristics	Pre: 2/1/22-2/28/23 (n = 1,110)	Post: 3/1/23-2/2/24 (n = 1,001)
Age, mean (years)	64.1	63.7
Procedure, n (%)		
COLO	357 (32.5)	335 (33.5)
HPRO	424 (38.2)	383 (38.3)
KPRO	329 (29.6)	283 (28.3)
Surgery Duration, mean (hours)	2.6	2.8
Wound Classification, n (%)		
Clean	785 (70.7)	644 (64.3)
Clean-Contaminated	293 (26.4)	260 (26.0)
Contaminated	16 (1.4)	22 (2.2)
Dirty/Infected	6 (0.5)	19 (1.9)
No Classification Recorded	10 (0.9)	56 (5.6)
Time from ABX to Surgery Start, mean (minutes)	23.1	22.5
LOS, mean (days)	4.0	4.5
Post-Op Infection, n (%)	14 (1.3)	22 (2.2)
Received Post-Op ABX, n (%)	21 (1.9)	20 (2.0)
Post-Op ABX Duration, mean (hours)	85.2	128.6

## **AU/AR Data Review**

## NHSN AR Reporting Validated for RRMC, NPH, SMH!





### Antimicrobial Resistance - Numerator CDA Results

/ tireiiiiiiii			,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,		
Facility	Date	Date	Validation	Issue(s) found and to be	Production Ready
racility	Received	Tested	Results	corrected	Production Ready
RRMC	4/23/2024	4/23/2024	Pass	None	Yes
NPH	4/24/2024	4/25/2024	Pass	None	Yes
SMH	4/23/2024	4/25/2024	Pass	None	Yes

### Antimicrobial Resistance - Denominator CDA Results

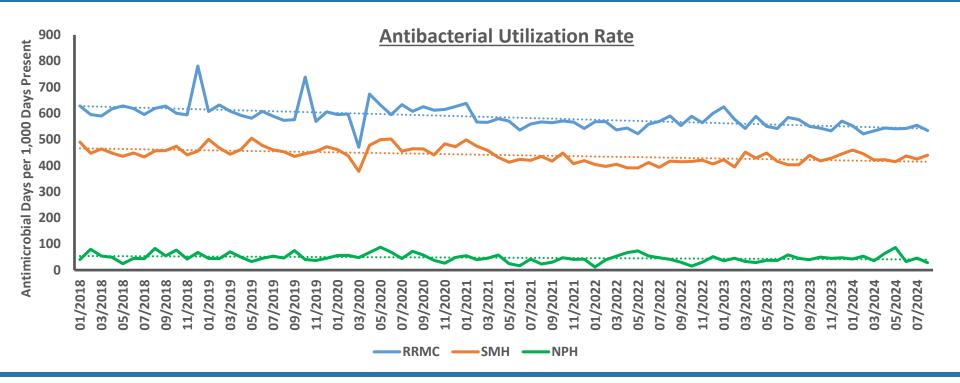
Antininciosia	resistance	Demoninator	CDA NEJUIC	•	
Facility	Date	Date	Validation	Issue(s) found and to be	Production Ready
racility	Received	Tested	Results	corrected	Production Ready
RRMC	4/23/2024	4/23/2024	Pass	None	Yes
NPH	4/24/2024	4/25/2024	Pass	None	Yes
SMH	4/23/2024	4/25/2024	Pass	None	Yes

## **Antimicrobial Use – SAAR by Unit Type**

Facility	LocationGroup	SAARTypeCat	Antimicrobial Days	Predicted Antimicrobial Days	Days Present	Location SAAR	95% Confidence Interval
		BSHO	6816	4545.759		1.499	1.464, 1.535
		GRAMPOS	3475	2552.985		1.361	1.316, 1.407
	ICHE	ANTIFGL	3208	1042.63	12716	3.077	2.972, 3.185
	1003	CDI	2315	2300.954	137 10	1.006	0.966, 1.048
	ICUS	BSCA	1831	1502.938		1.218	1.163, 1.275
		NSBL	575	Imicrobial Days         Antimicrobial Days         Days Present         Location SAAR Integration         95% Col Integration           6816         4545.759         1.499         1.464,           3475         2552.985         1.361         1.316,           3208         1042.63         3.077         2.972,           2315         2300.954         1.006         0.966,           1831         1502.938         1.218         1.163,           575         1832.64         0.314         0.289,           345         606.028         0.569         0.512,           332         450.26         0.737         0.661,           188         337.858         4075         0.422         0.360,           158         122.11         1.294         1.104,           155         651.401         0.238         0.203,           5200         5945.122         0.875         0.851,           5072         4682.772         [1.083]         1.054,           4859         5404.558         41104         0.998         0.970,           2402         3714.371         0.647         0.621,	0.289, 0.340		
	RMC STEPDOWN	CDI	345	606.028		0.569	0.512, 0.632
		BSCA	332	450.26		0.737	0.661, 0.820
DDMC	STEDDOWN	NSBL	188	337.858	4075	0.556	0.481, 0.640
KKIVIC	MC STEPDOWN GAB	GRAMPOS	160	378.957	4075	0.422	0.360, 0.492
		ANTIFGL	158	122.11		1.294	1.104, 1.508
		BSHO	155	651.401		0.238	0.203, 0.278
		CDI	5200	5945.122		0.875	0.851, 0.899
		NSBL	5072	4682.772		[1.083]	1.054, 1.113
	WADDE	BSHO	4859	5404.558	41104	0.899	0.874, 0.925
WARDS B	BSCA	4776	4784.357	41104	0.998	0.970, 1.027	
		GRAMPOS	2402	3714.371		0.647	0.621, 0.673
		ANTIFGL	1839	1045.927		1.758	1.679, 1.840



## RRMC vs. SMH vs. NPH: AU by Class, Antibacterials



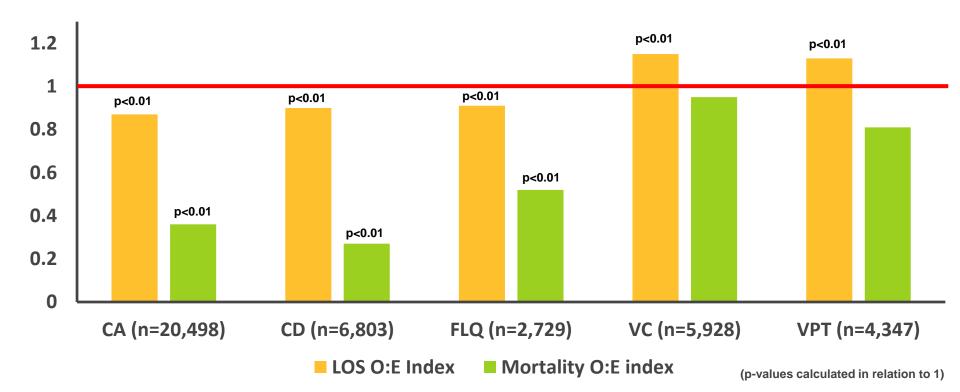


## **Vizient CAP Study**

### **Demographics:**

- 118 Academic Medical Centers
- 46,585 CAP encounters (with eLOS 3-7 days)

O:E index for LOS and mortality (NS ABX vs. BS ABX regimen; >1,000 encounters)



## **ASP Audit and Feedback**

## **RR Prospective Audit and Feedback**

Antimicrobial Audit Ad	cceptance	- RRUMO	C
Interventions from October 30	0, 2023 to Octo	ber 29, 2024	
Intervention Type	Proposed	Approved	Rate
Bug-Drug Mismatch	3	3	100.0%
De-Escalation	9	9	100.0%
Drug Information Consult	21	21	100.0%
Dose Change	64	63	98.4%
Drug-Lab Mismatch	29	27	93.1%
Duplicate Coverage	1	1	100.0%
Duration	3	3	100.0%
Home Health	3	3	100.0%
ID Consult	3	2	66.7%
Interactions	5	5	100.0%
IV to PO Conversion	10	9	90.0%
Regimen Change	30	30	100.0%
Restricted Med	3	0	0.0%
OVERALL	184	176	95.7%



## SMH and NPH Prospective Audit and Feedback

Antimicrobial Aud	lit Acceptanc	e – SMH	
Interventions from Octobe	er 30, 2023 to Oct	ober 29, 2024	ļ
Intervention Type	Proposed	Approved	Rate
Bug-Drug Mismatch	1	1	100.0%
De-Escalation	2	2	100.0%
Drug Information Consult	9	9	100.0%
Dose Change	19	19	100.0%
Drug-Lab Mismatch	9	9	100.0%
Duplicate Coverage	N/A	N/A	N/A
Duration	2	2	100.0%
Home Health	5	5	100.0%
ID Consult	3	3	100.0%
Interactions	2	2	100.0%
IV to PO Conversion	2	2	100.0%
Regimen Change	5	5	100.0%
Restricted Med	N/A	N/A	N/A
OVERALL	59	59	100.0%

Antimicrobial Audit	Acceptanc	e – NPH	
Interventions from October 3	30, 2023 to O	tober 29, 202	24
tervention Type Proposed Approved Rate			
Bug-Drug Mismatch	N/A	N/A	N/A
De-Escalation	N/A	N/A	N/A
Drug Information Consult	N/A	N/A	N/A
Dose Change	N/A	N/A	N/A
Drug-Lab Mismatch	N/A	N/A	N/A
Duplicate Coverage	N/A	N/A	N/A
Duration	N/A	N/A	N/A
Home Health	N/A	N/A	N/A
ID Consult	N/A	N/A	N/A
Interactions	N/A	N/A	N/A
IV to PO Conversion	N/A	N/A	N/A
Regimen Change	N/A	N/A	N/A
Restricted Med	N/A	N/A	N/A
OVERALL	N/A	N/A	N/A



## Microbiology Updates

## **Nudges and panels**

- Amp C
- Shigella
- Ongoing steno updates
- Use of pip-tazo for ESBL
- Fosfomycin disc added to urine cultures with ESBL E.coli and cipro/TMP resistance
- Vitek GN panel approved N814 for urine, N810 for non urine

Drug	N809	N814	N806	N807	N808	N810	N811	N812	GN99 Current Card	T
amikacin			İ			İ	İ			Ì
amoxicillin/clav.acid										1
ampicillin										1
ampicillin/sulbactam										1
aztreonam										1
cefazolin										t
cefepime										1
cefotaxime										1
cefoxitin										1
cefpodoxime										1
ceftazidime										1
ceftazidime/avibactam										1
ceftolozane/tazobactam										1
ceftriaxone										1
cefuroxime										
ciprofloxacin										
delafloxacin										1
doxycycline										1
ertapenem										1
erayacycline										٩
esbl										+
fosfomycin						_				4
gentamicin										4
										-
imipenem							_	_		
imipenem/relabactam										4
levofloxacin										-
meropenem										4
meropenem/vaborbctam	-									4
minocycline							_			4
moxifloaxacin										4
nitrofurantoin										
pip/tazo										4
polymyxin b	-									4
tigecycline										4
tetracycline										
tobramycin										
trimethoprim/sulfa										4
	IMP	IMP	IMP	IMP		IMP	IMP	IMP		4
					ERT					7
	тов	Gent	тов	тов	тов	Gent	Gent			4
Missing Drugs	108	AMK	AMK	AMK	108	AMK	AMK	AMK	<del>                                     </del>	1
	Amox/Clav		Amox/Clav	Amox/Clav		Amox/Clav	Amox/Clav			1
		P/T						P/T		1
						Nitro	Nitro	Nitro		4
	CAZ	CAZ					CAZ	CAZ		4
										+

# Infectious Diseases Transition Service (IDTS)

## What's new in 2023







### What

- Intravenous antibiotic medication used to fight off a bacterial infection
- · Types of infection include: bloodstream, heart, lung, skin and bone infections

### How

- . Given intravenously over 1 to 2 hours through a type of
- Dose and frequency may change based on your weekly



- . The length of time you take this medication will change depending on the type of infection you have and how severe it is.
- · Your doctor will decide how long you need to take this medication.

### Side effects

- Kidney toxicity
  - o Tip: Drink plenty of fluids; watch for decreased urine and swollen legs or ankles
- Infusion-related reaction
  - o Tip: Watch for redness, flushing, and itchiness shortly after intravenous infusion
- Ringing of the ears
- · Low white blood cell and platelet counts
- · Diarrhea, nausea, fatique

### **Tips**

- · If you are having side effects, please inform your nurse, doctor, and pharmacist.
- . Drug monitoring required: It is important to record the time you were given the medication and the time that your blood was drawn (should be drawn before the antibiotic is given).

### **Antibiotics:**

Penicillin Oxacillin Ampicillin Piperacillin-tazobactam Cefazolin Ceftriaxone Ceftazidime Cefepime Ertapenem

Imipenem-cilastatin Meropenem

...And more!



**UCLA Infectious Diseases Outpatient Parenteral Antimicrobial Therapy** Team

> Pharmacist: Lvnn Chan Contact: (310)267-1328

Nurse: Prest Oshodi Contact: (310)694-1230



### What

- Beta-lactams are a class of antibiotics that can be given intravenously to fight off bacterial infection.
- · Types of infection include: bloodstream, heart, lung, skin, and bone infections.

### How

- Given intravenously over 30 minutes or as continuous infusion over 24 hours
- Dose and frequency vary depending on the beta-lactam antibiotic selected for your type of infection

### Duration

- · The length of time you take this medication will change depending on the type of infection you have and how severe it is.
- · Your doctor will decide how long you need to take this

### Side effects

- · Side effects may vary depending on the type of betalactam antibiotic; ask your doctor or pharmacist for more
- Decrease red blood cell, white blood cell, or platelet
- Diarrhea, nausea, fatique

### Tips

- · If you have any side effects, please inform your nurse, doctor, and pharmacist.
- Let your doctor or pharmacist know if you have a betalactam allergy (for example: a penicillin allergy).

**UCLA Infectious Diseases Outpatient Parenteral Antimicrobial Therapy** Team

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## **Brainstorming**

### Executive Summary: (Antimicrobial Stewardship Program)

Date: July 17, 2024

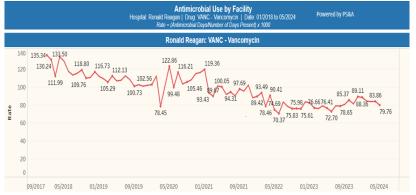
Presenter(s): Tara Vijayan, MD MPH



### Overview:

Current Fiscal Year Goals	Baseline Data	Target	Current Data	Interventions	Met/Not Met
Integrated AR NHSN	Not submitted	submit	submitting	Integrated AR data	Met, ongoing
CDPH SDOH data submission	Not submitted	submit	submitting	Data pull for CDPH	Met, ongoing
Update pathways	Needs updating	update	updated	Periop guidance Quickstart	Met, ongoing
Vancomycin reduce AU	SAAR	Reduce	See below	MRSA nares BCID planned	Met, ongoing
Document OPAT efforts	23% 30d readmission	20%	19%	RN oversight; PharmD	Met, ongoing

### **Data Highlights**



### Accomplishments:

- Multiple educational interventions (competency based modules, AAW, grand rounds)
- · Multidisciplinary collaborations with BMT, liver transplant, orthopedics
- · Updating guidance
- Streamlining CARE Connect processes: ordersets, antimicrobial indications, nudges
- Academic productivity (>30 publications in 1 year)
- Reduction in some antimicrobial use
- · Implementation of Time outs, allergy de-labeling pathways
- Reduction in readmissions documented (OPAT/IDTS program)
- · Integrated AR NHSN

### Challenges / Barriers / Lessons Learned:

We continue to have staffing shortages

Program Manager took another job, new Program Manager joining July 8, 2024 (50%)

We need dedicated time and staff to help us with measurements (much of our efforts are clinically oriented)

### **Sustainability Plans for Improvements:**

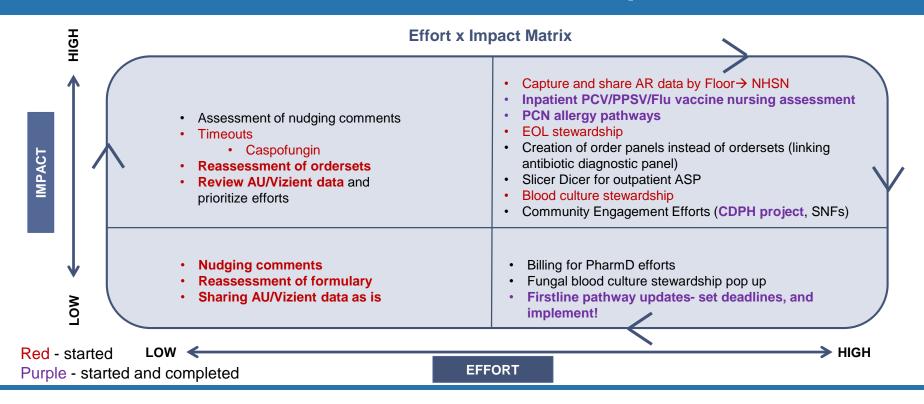
Continue twice monthly meetings to ensure progress Advocate for continued support

Future Goals (for next 12- month period)	Interventions	Metrics	Baselin e Data	Target	Dept. / Org. Goal Alignment
Reduce meropenem use in 8ICU	Time out	Reduction in DOT			Safety (reduce CRE rates)
Reduction of meropenem in BMT	Febrile neutropenia guidance	Reduction in DOT			Safety (reduce CRE rates)
Penicillin allergy pathways	Measure impact	Increase de- labeling			Safety (better antibiotics)
Nudging comments in micro	Measure impact	Adherence to nudges			Improve clinician experience)
PCV nursing updates	Measure adherence to guidance	Adherence to guidance			Safety

## **Pharmacist FTEs**

- Ongoing discussions about need for outpatient pharmD FTE
  - Nursing home collaboratives
  - Outpatient Stewardship
  - Managing complex outpatient antimicrobial therapy (COpAT)
- Grant support for research projects

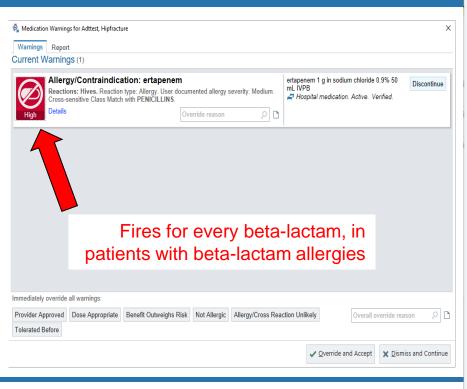
## 2024 Prioritization Matrix: What to Keep?



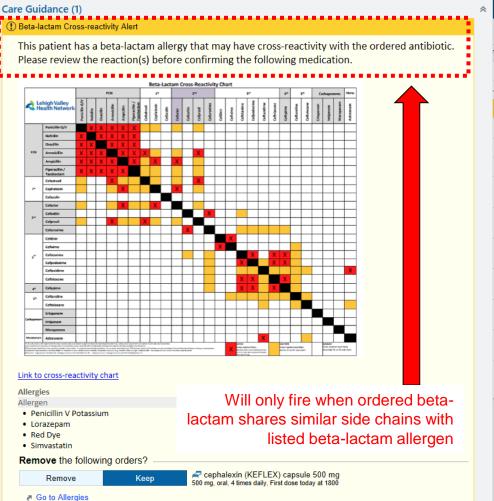


## **New Potential Projects**

## **Beta-Lactam Allergy Project – Next Steps**



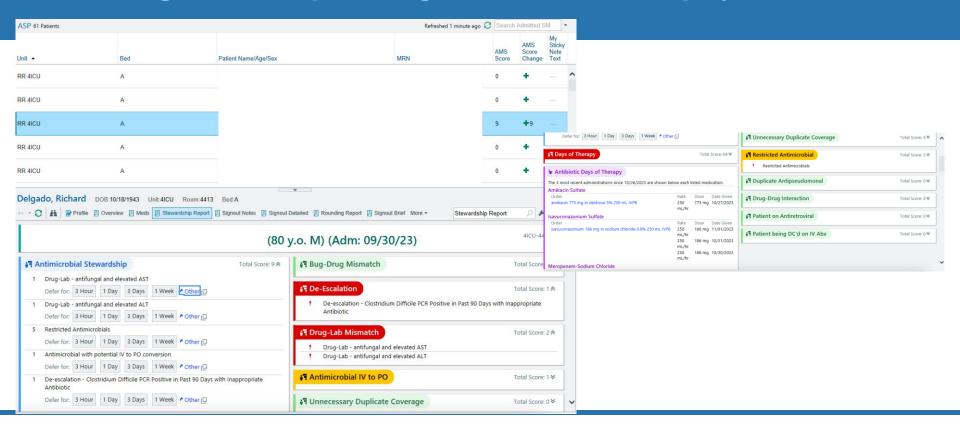




Accept

Dismiss

## AS Navigator – Expanding utilization to ID physicians





## **SEP-1** Bundle and Time to Appropriate Antibiotics

Clinical Infectious Diseases





Improving Sepsis Outcomes in the Era of Pay-for-Performance and Electronic Quality Measures: A Joint IDSA/ACEP/PIDS/SHEA/SHM/SIDP Position Paper

Chanu Rhee, <sup>1,2,0</sup> Jeffrey R. Strich, <sup>3</sup> Kathleen Chiotos, <sup>4</sup> Andre C. Kalil, <sup>9</sup> David N. Gilbert, <sup>10</sup> Henry Masur, <sup>3</sup> Edw and Michael Klompas <sup>1,2,1</sup>

¹Department of Population Medicine, Harvard Medical School/Harvard and Women's Hospital, Boston, Massachusetts, USA. ²Dritical Care Is and Women's Hospital, Boston, Massachusetts, USA. ²Dritical Care Is and Women's Hospital of Philadelphia and Ur of Medicine, Diladelphia, Pennsy Baltimore, Manyland, USA. ²Dociety of Hospital Medicine, Philadelphia, Pennsy Baltimore, Manyland, USA. ²Dociety of Hospital Medicine, Pilose Diseases, Department of Medicine, Oregon Health and Science University, Port Pulmorary and Oritical Care Medicine, Department of Medicine, Univ. Washington University School of Medicine, Washington D.C., USA.² URA ¹Department of Tempenery Medicine, University of Ptitsburgh S.

## PART 1: REASONS TO RETIRE SEP-1 RATHER THAN MAKE IT A PAY-FOR-PERFORMANCE MEASURE

## Real-world Evidence Indicates That SEP-1 Has Not Improved Patient Outcomes

Several time-series analyses using detailed clinical data from hundreds of hospitals elucidate the real-world impact of SEP-1 on patient outcomes (Table 1) [12–15]. Rhee et al analyzed 117 150 patients admitted to 114 academic and community hospitals with suspected sepsis between 2013 and 2017 and found that SEP-1 implementation in October 2015 was associated with an immediate increase in lactate testing but no improvement in the combined outcome of hospital death or discharge to hospice [12]. These findings persisted in several sensitivity anal-

pay more attention to the full breadth of sepsis care and stimulate further innovations in diagnosis and treatment. Hospitals could still choose to emphasize early resuscitation bundles based on internal assessments of gaps in care but they should not be forced to do so.

### PART 2: RECOMMENDATIONS TO IMPROVE THE eCQM SEPSIS MORTALITY MEASURE

We support CMS's plan to implement a risk-adjusted sepsis outcome measure. Although there are multiple patient-centered sepsis outcomes that could be candidates, we believe that a focus on mortality is the right place to start. We also applaud CMS's plan to make the measure fully electronic, as this will improve efficiency, scalability, and objectivity compared to the current manual SEP-1 abstraction process which is highly resource-intensive and often variably applied [39–41].

The draft specification for the eCQM sepsis mortality measure identifies sepsis using three criteria (Table 2): (1) systemic inflammatory response syndrome (SIRS) criteria, defined using vital signs and white blood cell counts, (2) suspected infection, defined as antibiotic administrations  $\sigma$ r the use of

ummenueu consequences

#### Remove SIRS Criteria From the eCQM

SIRS criteria are common and nonspecific. They are present in up to 50% of hospitalized patients at some point during their stay, most of whom do not have sepsis [42]. Another study found that 18% of ED patients met SIRS criteria, but only 26% of that group had an acute infection [43]. SIRS criteria are also insensitive; one in eight critically ill patients with sepsis do not meet SIRS criteria [44]. Limiting the eCQM to patients with SIRS criteria therefore risks both over-detection and under-detection of sepsis.

Anchoring the eCQM to SIRS also risks promoting overreliance on SIRS as a screening tool. Using an insensitive and non-specific trigger cannot drive improvements in care. Indeed, the evidence suggests SIRS-based alerts in the ED increase antibiotic use and Clostridioides difficile infections but do not improve mortality [45, 46]. SIRS-based prompts for sepsis recognition in the intensive care unit (ICU) or inpatient setting have also not improved patient outcomes in randomized trials [47–49]. These limitations of SIRS led to their exclusion from current international consensus criteria for sepsis (Sepsis-3) [50].



## Vizient Benchmarks – Planning for Regular Reporting

### **Upcoming CDB Trainings:**

- Vizient CDB Learning Center <a href="https://learning.vizientinc.com/cdb/Pages/default.aspx">https://learning.vizientinc.com/cdb/Pages/default.aspx</a>
- CDB Orientation Session 1: Welcome to the CDB 2/7/2023 9:30 AM CT
- CDB Orientation Session 2: Introduction to Generating Reports 12/7/2023 12:00 PM CT
- Vizient CDB Manual <a href="https://learning.vizientinc.com/cdb/Clinical/Clinical\_Data\_Base\_User\_Manual\_UI.pdf">https://learning.vizientinc.com/cdb/Clinical/Clinical\_Data\_Base\_User\_Manual\_UI.pdf</a>

### What Benchmarks to Report?:

- Carbapenem utilization (specific populations)?
- Unit specific reports?
- Perioperative antibiotic utilization?
- Specific diagnoses to evaluate?
- What external comparators do we want to use?

### Cadence:

Recommend no more frequently than quarterly given delays with reporting data to CDB



## Other potential projects

- Community Engagement Efforts: Engagement of SNFs
- Antibiotic Time Outs: Expand to other antibiotics?
- Ongoing engagement of nursing and hospital pharmacists
- Outpatient ASP
- Vizient and NHSN data how do we better utilize this information
- Implementation of bacteriophage therapy?
- Improving Streptococcus pneumoniae isolation
- Order panels/Ordersets
  - New ordersets:
    - Splenectomy vaccine order set?
    - IT amphotericin
    - Intraperitoneal antimicrobials



## 2025 Prioritization Matrix

