

Enhancing Antimicrobial Stewardship Programs Through Collaboration With the Microbiology Laboratory

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July 19, 2023

Disclosures

• The presenters have no real or perceived conflicts of interest related to this presentation

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Learning Objectives

At the end of this session, participants should be able to:

- 1. Identify potential obstacles to forming a successful collaboration between stewardship programs and microbiology labs.
- 2. Recall three microbiology reporting approaches that can influence prescribing practices.
- 3. Describe the potential benefits of collaboration from a patient, laboratory and stewardship perspective.







Nichole Neville, PharmD, BCIDP

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- Antimicrobial resistance (AMR) is a pressing worldwide concern
 - Estimated 4.95 million deaths were associated with drug-resistant infections in 2019 global systematic analysis
 - 1.27 million deaths *directly* attributable to bacterial AMR
- 2022 CDC Special Report indicated historic gains made on antibiotic stewardship were reversed during the COVID-19 pandemic

Available data show an alarming increase in resistant infections starting during hospitalization, growing at least 15% from 2019 to 2020.

- Carbapenem-resistant Acinetobacter (†78%)
- Antifungal-resistant Candida auris (+60%)*
- Carbapenem-resistant Enterobacterales (+35%)
- Antifungal-resistant Candida (†26%)

- ESBL-producing Enterobacterales (†32%)
- Vancomycin-resistant Enterococcus (+14%)
- Multidrug-resistant P. aeruginosa (†32%)
 - Methicillin-resistant Staphylococcus aureus (†13%)

Source: Antimicrobial Resistance Collaborators. Global burden of bacterial antimicrobial resistance in 2019: a systematic analysis. Lancet. 2022 Feb 12;399(10325):629-655. doi: 10.1016/S0140-6736(21)02724-0. Epub 2022 Jan 19. Erratum in: Lancet. 2022 Oct 1;400(10358):1102.

Source: CDC. COVID-19: U.S. Impact on Antimicrobial Resistance, Special Report 2022. Atlanta, GA: U.S. Department of Health and Human Services, CDC; 2022. https://www.cdc.gov/drugresistance/covid19.html. Accessed 5/23/2023.











- Varied resistant mechanisms, delays in testing, out-of-date breakpoints, lack of understanding risk factors for resistant infections etc., all impact antibiotic therapy and lead to delays in administration of effective treatment
- Delayed appropriate therapy is associated with worse outcomes
 - Increased total in-hospital costs
 - \circ Increased lengths of stay
 - 20% increased in-hospital mortality
- Time to appropriate antibiotic therapy shown to be an independent predictor of 30-day mortality in patients with certain resistant organisms

Source: Bassetti M, Kanj SS, Kiratisin P, Rodrigues C, Van Duin D, Villegas MV, Yu Y. Early appropriate diagnostics and treatment of MDR Gram-negative infections. JAC Antimicrob Resist. 2022 Sep 13;4(5):dlac089. doi: 10.1093/jacamr/dlac089.

Source: Bonine NG, Berger A, Altincatal A, Wang R, Bhagnani T, Gillard P, Lodise T. Impact of Delayed Appropriate Antibiotic Therapy on Patient Outcomes by Antibiotic Resistance Status From Serious Gram-negative Bacterial Infections. Am J Med Sci. 2019 Feb;357(2):103-110. doi: 10.1016/j.amjms.2018.11.009. Epub 2018 Nov 22.

Source: Falcone M, Bassetti M, Tiseo G, Giordano C, Nencini E, Russo A, Graziano E, Tagliaferri E, Leonildi A, Barnini S, Farcomeni A, Menichetti F. Time to appropriate antibiotic therapy is a predictor of outcome in patients with bloodstream infection caused by KPC-producing Klebsiella pneumoniae. Crit Care. 2020 Jan 30;24(1):29. doi: 10.1186/s13054-020-2742-9.





• Ability to fight and halt further resistance is directly related to the capacity of the microbiology lab's ability to perform:



Source: Wenzler, E, Maximos M, Asempa TE, Biehle L, Schuetz AN, Hirsch EB. Antimicrobial susceptibility testing: an updated primer for clinicians in the era of antimicrobial resistance: insights from the Society of Infectious Diseases Pharmacists. Pharmacotherapy. 2023 Feb 24. Doi: 10.1002/phar.2781. Epub ahead of print.

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- Challenges for microbiology labs in optimizing AST
 - Emergence of new resistance mechanisms
 - $\,\circ\,$ Development of new antibiotics
 - $\,\circ\,$ Revisions to established methods and breakpoints

Labs may interpret AST results using obsolete breakpoints:

- Serious patient safety concerns and ramifications
- Hinders ability to track AMR





Need to update breakpoints not required & often under recognized or not prioritized due to lack of resources

Laboratories faced with numerous challenges in AST capabilities & accurate interpretation

Source: Simner PJ, Rauch CA, Martin IW, et al.. Raising the bar: Improving antimicrobial resistance detection by clinical laboratories by ensuring use of current breakpoints. *Open Forum Infect Dis.* 2022;9 (3). doi.org/10.1093/ofid/ofac007

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13.3% of laboratories

are unaware of need

to update breakpoints

37.9-70.5% of U.S.

laboratories are using

obsolete breakpoints

- Well informed clinicians can help bridge the gap between AST challenges and frontline experience to improve patient outcomes in a world of rapidly changing/increasing AMR
- The microbiology laboratory's capabilities paired with clinician expertise and collaboration are essential in this AMR fight to:
 - Decrease mortality
 - $\circ~$ Improve clinical and economic outcomes
 - $\circ~$ Reduce delays in time to effective and optimal antimicrobial therapy





Forming a Successful Collaboration Between the Microbiology Lab & Stewardship Teams

Kelsey Melander, PharmD, BCIDP

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Formation of Laboratory/Stewardship Committee



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Milestone

Centralization & the Necessity of Collaboration

Case example: Implementation of updated cefazolin breakpoints for Enterobacterales

- 1. Recognizing the need for microbiological and clinical expertise
- 2. Including the right voices
- 3. Appreciating different perspectives





Cefazolin/Enterobacterales susceptibility reporting

• Cascade reporting based on cefazolin susceptibility







Cefazolin/Enterobacterales susceptibility reporting

• Cascade reporting based on cefazolin susceptibility



confounded in setting of use of outdated breakpoints.

In 2019, a survey of 1,490 CAP-accredited clinical laboratories demonstrated a low rate of compliance with current breakpoints (29.5%–62.1%). Breakpoints surveyed had been implemented as of 2010.

Source: Simner PJ, Rauch CA, Martin IW, et al.. Raising the bar: Improving antimicrobial resistance detection by clinical laboratories by ensuring use of current breakpoints. *Open Forum Infect Dis.* 2022;9 (3). doi.org/10.1093/ofid/ofac007

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Cefazolin/Enterobacterales susceptibility reporting

 Cefazolin susceptibilities were not in alignment with CLSI or FDA and outside of the Vitek[®]2 calling range

	Cefazolin Breakpoint Information									
	Concentration	0.5	1	2	4	8	16	32	64	128
VITEK Calling Range										
CLSI (systemic) 2019 M100S				S (<=2)	1	R (>=8)				
FDA Enterobacteriaceae			S	I.	R					
VITEK 2 CLSI Breakpoints (Urine)							S	1	R	

Source: Humphries RJ, Abbott AN, Hindler JA. Understanding and addressing CLSI breakpoint revisions: a primer for clinical laboratories. *J Clin Microbiol*. 2019;57(6):e00203-19. doi: 10.1128/JCM.00203-19

biomerieux. (2018). Vitek[®]2 AST-GN80.



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Clinical Expertise

- Recognition of breakpoints most in alignment with clinical and PK/PD data
- Communicate potential clinical impact of requested change
- Recommending the need for automated vs. by request only work-up

Microbiological Expertise

- Knowledge of validation procedures to update breakpoints
- Working within confines of technology limitations
- Balance efficient laboratory workflow vs. accommodate all clinician requests
- Knowledge of multiple microbiological techniques to form creative solutions

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Including the Right Voices







Appreciating Different Perspectives

- Create a bridge between clinical and laboratory personnel
 - Weighing the clinical benefit against the "lift" required for implementation
 - Timeline for implementation and setting reasonable expectations (updated cefazolin breakpoints: 2018–2020; reporting with limitations)
- Communication with providers
 - Effectiveness of implementation: automated work-up vs. by request only
 - $\circ~$ Supplemental education
 - $\circ~$ Comments included in microbiology results





Implementation of Updated Cefazolin/Enterobacterales Breakpoints



Change and optimization of reporting takes time.

Routine re-evaluation of the effectiveness of new processes.

Implementation without provider education will have a limited impact.

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Assessment Question 1

How did the example of implementing updated cefazolin/enterobacterales breakpoints expose the obstacles to forming a successful partnership between stewardship programs and microbiology labs?

- A. Based on the pharmacokinetics of cefazolin, two separate breakpoints exist depending on site of infection (systemic vs. cystitis). This represents need for clinical expertise.
- B. Based on technology limitations associated with automated susceptibility testing platforms, the microbiology lab was unable to fully align susceptibility reporting with current breakpoints. This represents the need for microbiological expertise.
- C. Implementation of Kirby Bauer discs for Enterobacterales (limited to blood source) highlighted the ability to align breakpoints with current recommendations demonstrated the ability to accommodate clinical and laboratory needs.
- D. All of the above





Assessment Question 1 | Answer...

How did the example of implementing updated cefazolin/enterobacterales breakpoints expose the obstacles to forming a successful partnership between stewardship programs and microbiology labs?

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- D. All of the above







Influential Microbiology Reporting Approaches Nichole Neville, PharmD, BCIDP

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Selective Reporting

Clarity of Reporting





• Cascade Reporting (CR):

- Reporting results for specific antimicrobial agents based on the AST profile/pattern of the isolate
- $\circ~$ Utilized to encourage appropriate antimicrobial use
- $\circ~$ Recommended in CLSI M100 33 $^{\rm rd}$ edition



Source: CLSI. Performance Standards for Antimicrobial Susceptibility Testing. 33rd ed. CLSI supplement M100. Clinical and Laboratory Standards Institute; 2023.

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- Cascade Reporting (CR):
 - Laio, et al.:
 - Decreased cefepime utilization in using CR based on ceftriaxone susceptibility
 - Mean DOT among patients who received cefepime decreased from 1.229 days during baseline period to 0.813 days post-CR (p <.0001)
 - Significantly lowers the lengths of stay (14.139 days to 10.882 days) during the post-CR period compared to baseline (p <.0001)
 - No change in inpatient mortality
 - Langford, et al.:
 - In outpatients aged >65 yo, reporting antibiotic susceptibility on urine cultures was associated with increased odds of prescribing the reported antibiotic (adjusted OR 1.23, 95%CI 1.13–1.33, per 25% increase in reporting)

Source: Liao S, Rhodes J, Jandarov R, DeVore Z, Sopirala MM. Out of Sight-Out of Mind: Impact of Cascade Reporting on Antimicrobial Usage. Open Forum Infect Dis. 2020 Jan 8;7(2):ofaa002. doi: 10.1093/ofid/ofaa002.

Source: Langford BJ, Daneman N, Diong C, Marchand-Austin A, Adomako K, Saedi A, Schwartz KL, Johnstone J, MacFadden DR, Matukas LM, Patel SN, Garber G, Brown KA. Antibiotic susceptibility reporting and association with antibiotic prescribing: a cohort study. Clin Microbiol Infect. 2021 Apr;27(4):568-575. doi: 10.1016/j.cmi.2020.10.001. Epub 2020 Oct 12.





- Cascade Reporting (CR):
 - Weichman, et al.:
 - Implemented CR for Enterobacterales in urine cultures at a network of urgent care clinics
 - Interrupted time series analysis (ITSA) measuring monthly antibiotic prescriptions per 1000 patient encounters showed 38% reduction in fluoroquinolone prescribing rates post CR (p < .0001)

Based on the evidence displaying substantial benefits for CR, the Denver Market of the Continental Division implemented widespread use of CR

Source: Weichman, B., Bushman, A., Rogers, K., & Rosa, R. (2022). Impact of fluoroquinolone cascade reporting of urine samples on antibiotic prescribing rates in a network of urgent care clinics. *Antimicrobial Stewardship & Healthcare Epidemiology, 2*(1), E97. doi:10.1017/ash.2022.227





Cascade Reporting: Denver Market Examples

<i>Escherichia coli:</i> Non-cascaded susceptibility re	eport		Es Ca	<i>cherichia coli:</i> scaded susceptibility report		
1. ESCHERICHIA COLI CEFAZOLIN 1. ESCHERICHIA COLI RX may vary depending on ta AMPICILLIN AMP/SULBACTAM CEFTRIAXONE TRIMETH/SULFA	KIRBY BAU 	er INTERP s and dose INTERP S S S S S		1. ESCHERICHIA COLI ESBL RX may vary depending on ta CEFTRIAXONE GENTAMICIN MEROPENEM PIPERACILLIN/TAZOBACTAM TOBRAMYCIN TRIMETH/SULFA	MIC 	and dose INTERP R R S S R S S

- Organism is pan-sensitive
- Susceptibility report does not cascade

Resistant Ceftriaxone fires the cascade



Cascade Reporting: Denver Market Examples

Carbapenem Resistant Organi Pseudomonas aeruginosa	sm (CRO):		Cascaded (CRO Pseudomonas aer	uginosa	susceptib	ility report
				1. PSEUDOMONAS AERUGINOSA CRO CEFIDEROCOL	N KIRBY BAU	JER INTERP S	
 PSEUDOMONAS AERUGINOSA O RX may vary depending on ta 	RO Arget, route, MIC	and dose INTERP		1. PSEUDOMONAS AERUGINOSA CRO RX may vary depending on targ) get, route, MIC	and dose INTERP	
CEFEPIME GENTAMICIN LEVOFLOXACIN MEROPENEM PIPERACILLIN/TAZOBACTAM TOBRAMYCIN	32 <=1 >=8 >=16 >=128 <=1	R S R R R S		CEFTAZIDIME/AVIBACTAM 1. PSEUDOMONAS AERUGINOSA CRO RX may vary depending on tary CEFTOLOZANE/TAZOBACTAM 1. PSEUDOMONAS AERUGINOSA CRO	8/4 get, route, MIC 1.5/4	S and dose INTERP S	
				RX may vary depending on targ	get, route, MIC 12/4	and dose INTERP R	

- Resistant meropenem fires the automated cascaded
 report
- Microbiology automatically begins ASTs for expanded antimicrobials and reports when complete

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Selective Reporting



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• Selective Reporting (SR):

- Utilizes reporting results for specific antimicrobial agents based on specific criteria instead of AST results
- $\circ~$ May occur for the following reasons:
 - Specific bug/drug combination is inappropriate for a given site of infection
 - Drug is not on the facility formulary
 - A drug may be inappropriate for the source of infection and/or particular patient populations
 - Specific testing method may have bug/drug limitations
 - Literature suggests suboptimal outcomes with a particular drug in the treatment of specific resistance phenotypes and/or types of infections

Source: CLSI. *Performance Standards for Antimicrobial Susceptibility Testing.* 33rd ed. CLSI supplement M100. Clinical and Laboratory Standards Institute; 2023. Source: Selective reporting of antimicrobial susceptibility testing results: a primer for antibiotic stewardship programs. Centers for Disease Control and Prevention website. <u>https://www.cdc.gov/antibiotic-use/pdfs/Selective-Reporting-508.pdf</u> CAa. Published 2020. Accessed May 23, 2023.

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Selective Reporting (SR)

- Aims to guide antimicrobial prescribing to the most appropriate, guideline-recommended antimicrobials
- Recommended in French, European and international guidelines to reduce the number of inappropriately prescribed antibiotics
- Recommended in CLSI M100 33rd edition

• Found to decrease broad spectrum antibiotic prescriptions

Source: Le Dref G, Simon M, Bocquier A, Fougnot S, Kivits J, Duda A, Pulcini C, Thilly N; ANTIBIO-ciblés Scientific Committee[‡]. Selective reporting of antibiotic susceptibility testing results for urine cultures: feasibility and acceptability by general practitioners and laboratory professionals in France. JAC Antimicrob Resist. 2023 Feb 11;5(1):dlad013. doi: 10.1093/jacamr/dlad013. Source: CLSI. *Performance Standards for Antimicrobial Susceptibility Testing*. 33rd ed. CLSI supplement M100. Clinical and Laboratory Standards Institute; 2023.

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Microbiology Reporting Approaches That Can Influence Prescribing Practices Selective Reporting (SR)

- Simon, et al.:
 - Prospective, multi-center, controlled before/after intervention study
 - Assessed impact of SR of AST results for urine cultures positive with *Escherichia coli* on prescription of broad spectrum antibiotics with a higher risk of selection for resistance
 - $\circ~$ Conclusion:
 - Decreased proportion of broad-spectrum antibiotic prescriptions was significantly higher for SR of AST
 - Attributable decrease in prescription proportion of 3rd gen cephalosporins (-8.5% for selective reporting vs. -0.1% for complete reporting, p< 0.001)

Source: Simon M, Fougnot S, De Monchy P, Duda A, Thilly N, Pulcini C; ANTIBIO-ciblés Scientific Committee. Impact of selective reporting of antibiotic susceptibility testing results for urinary tract infections in the outpatient setting: a prospective controlled before-after intervention study. Clin Microbiol Infect. 2023 Mar 13:S1198-743X(23)00126-X. doi: 10.1016/j.cmi.2023.03.012. Epub ahead of print.





Selective Reporting (SR): Denver Market Examples

- Fluoroquinolones
 - 2019 HCA Continental Division Denver Market implemented SR by masking fluoroquinolones from susceptibility reports for Enterobacterales
 - Providers must contact pharmacy or call the microbiology lab for susceptibility information, if needed

URINE AEROBIC CULTURE Fina	1
Organism 1	ESCHERICHIA COLI
COLONY COUNT	GREATER THAN 100,000 CFU'S/ML
1. ESCHERICHIA COLI	
RX may vary depending on	1 target, route, and dose
	MIC INTERP
AMPICILLIN	8 S
CEFAZOLIN	<=4 S
CEFAZOLIN CEFTRIAXONE	<=4 S <=1 S
CEFAZOLIN CEFTRIAXONE GENTAMICIN	<=4 S <=1 S <=1 S
CEFAZOLIN CEFTRIAXONE GENTAMICIN NITROFURANTOIN	<=4 S <=1 S <=1 S <=16 S





Continental Division: Denver Market Data



- Selective Reporting implemented 12/2019
- Most hospitals decreased fluoroquinolone spend by 72–89%

• *Non-transplant facilities* decreased fluoroquinolone Average DOT by 35–60%

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Clarity of Reporting





Reporting additional information to susceptibility reports can benefit prescribing practices by:

Educating clinicians about the most appropriate antimicrobial dose for a susceptible-dose-dependent (SDD) isolate



Notifying prescribers when specific organisms requiring broad-spectrum antibiotics are not present



Informing clinicians when the laboratory is no longer working up an isolate





Clarity of Reporting: Incorporation of SDD designation

Enterococcus

- Daptomycin treatment failures have been demonstrated in Enterococci with elevated MICs
- Standard doses of 4-6mg/kg may not attain pharmacodynamic targets for selected Enterococci
- CLSI approved SDD breakpoint range for Enterococcus spp in 2018
 - SDD breakpoint based on dosage regimen of 8–12mg/kg every 24 hours for serious Enterococci infections

Source: CLSI. Performance Standards for Antimicrobial Susceptibility Testing. 33rd ed. CLSI supplement M100. Clinical and Laboratory Standards Institute; 2023.





Clarity of Reporting: Incorporation of SDD designation

Adema, et al.:

- Mixed-methods study combined a clinician survey with a retrospective pre-post prescribing analysis
- Found SDD reporting for Enterococcus spp was associated with a change in definitive daptomycin dosing
 - Daptomycin dosage following susceptibility testing found to be significantly higher post-SDD compared with pre-SDD (8.5 mg/kg vs 6.4 mg/kg; P < .001)
- Showed ID clinicians had more confidence in SDD category compared to internal medicine clinicians
 - Significant difference between knowledge and interpretation of SDD interpretive category when applied to daptomycin for the treatment of Enterococci

Source: Adema JL, Lake LN, Stevens RW, Hogan BM, Schuetz AN, Tande AJ, Mara KC, Eberly AR, Rivera CG. Understanding and Application of Daptomycin-Susceptible Dose-Dependent Category for *Enterococcus*: A Mixed-Methods Study. Open Forum Infect Dis. 2022 Jan 10;9(1):ofab611. doi: 10.1093/ofid/ofab611.





Clarity of Reporting

- Denver Market incorporation of Susceptible Dose Dependent (SDD) designation
 - Added SDD designation with specified MIC
 - Provided comment regarding the dosing regimen that SDD is based on
 - Included recommendation to consult
 Infectious Diseases

1. ENTEROCOCCUS FAECIUM, VRE		
RX may vary depending on targe	et, route,	and dose
	MIC	INTERP
AMPICILLIN	>=32	R
GENTAMICIN SYNERGY SCREEN	SYN-S	S
LINEZOLID	2	S
STREPTOMYCIN SYNERGY SCREEN	SYN-S	S
VANCOMYCIN	>=32	R
1. ENTEROCOCCUS FAECIUM, VRE		
RX may vary depending on targe	et, route,	and dose
	MIC	INTERP
DAPTOMYCIN	4	SDD

The breakpoint for SDD (susceptible-dose dependent) is based on a dosage regimen of 8-12 mg/kg and is intended for serious infections due to E.faecium. Consult with infectious disease is recommended.

S=SUSCEPTIBLE I=INTERMEDIATE R=RESISTANT N/R=NOT REPORTED BLANK=DATA NOT AVAILABLE, OR DRUG NOT ADVISABLE OR TESTED

MIC=mcg/ml (mg/L)

Source: CLSI. Performance Standards for Antimicrobial Susceptibility Testing. 33rd ed. CLSI supplement M100. Clinical and Laboratory Standards Institute; 2023.





 Simple behavioral strategies in microbiology reporting can improve antimicrobial de-escalation and/or discontinuation of unnecessary broadspectrum antibiotics





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Open Forum Infectious Diseases

MAJOR ARTICLE



Microbiology Comment Nudge Improves Pneumonia Prescribing

Musgrove, et al.:

- Single pre-test, post-test quasi experiment in four-hospital health system
 - Pre-test: reports stated "commensal respiratory flora"
 - Post-test: reports stated "Commensal respiratory flora only: No S. aureus/MRSA or P. aeruginosa"
- Found 5.5-fold increased odds of de-escalation in patients being treated with anti-MRSA and antipseudomonal antibiotics for respiratory infections when the laboratory report indicated no MRSA or PSA for respiratory cultures where neither was cultured and there was no dominant organism growth

Source: Musgrove MA, Kenney RM, Kendall RE, Peters M, Tibbetts R, Samuel L, Davis SL. Microbiology Comment Nudge Improves Pneumonia Prescribing. Open Forum Infect Dis. 2018 Jul 10;5(7):ofy162. doi: 10.1093/ofid/ofy162.

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 Implementation of a Denver Marketwide comment regarding no PSA on respiratory cultures where the organism was not isolated

RESPIRATORY AEROBIC CULTURE	Final	05/24/23-0914	COE
Organism 1 QUANTITATION	NORMAL UPPER RESPIRATORY FLO ABUNDANT	RA	





- Based on favored response from previously mentioned clarifications of reporting, working group implemented additional explanations in reporting:
 - Sent out for susceptibility testing vs. "No further work-up"

BLOOD CULTURE Prelimina GS: 05/23/23 1003 1	ry (PREVIOUS RPT) AMLB.TS>>GNB	05/24/23-	3-1302
Organism 1 BOTTLES(S) STATUS	SALM. ENTERICA SSP AEROBIC AND ANAERON SUSCEPTIBILITY RESU	ENTERICA BIC BOTTLE POSITIVE JLTS TO FOLLOW	
	BL	OOD CULTURE Final Organism 1 BOTTLES(S) STATUS	05/18/23-1103 BACILLUS SPS, NOT ANTHRACIS AEROBIC BOTTLE POSITIVE NO FURTHER WORK-UP WILL BE PERFORMED
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Clarity of Reporting

Clarification of organism

- Specific reporting for Carbapenem Resistance Organisms (CROs)
- Specific reporting for Extended Spectrum Beta Lactamase organisms (ESBLs)
- Provides crucial information to clinicians for treatment considerations
- Provides necessary information to Infection Prevention teams for appropriate patient isolation requirements



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Assessment Question 2

What are three microbiology reporting approaches that can influence prescribing practice?

- A. Selective Reporting (SR) Can assist in influencing prescribers to not prescribe a specific antimicrobial that literature suggests suboptimal outcomes in the treatment of specific resistance phenotypes
- B. Cascade Reporting (CR) Can aim to guide antimicrobial prescribing to the most appropriate guideline-recommended antimicrobials
- C. Clarity of Reporting Can help guide prescribers to potential de-escalation and/or discontinuation of unnecessary broad spectrum antibiotics if specific organisms are not isolated
- D. All of the above

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Assessment Question 2 | Answer...

What are three microbiology reporting approaches that can influence prescribing practice?

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Benefits of Collaboration

Kelsey Melander, PharmD, BCIDP

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Benefits of Collaboration — Patient Perspective

Receive antimicrobial therapy that is based on validated,

local data

- Empiric therapy: Antibiogram data
- Preliminary targeted therapy: Rapid diagnostics

 Verigene[®] Bloodstream Infection Testing Panels
 MALDI-TOF
- Targeted therapy: Implementation of current breakpoints



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Benefits of Collaboration — Laboratory Perspective

- Create a space where conversation can occur
- Understand the "why" and help microbiologists see the clinical impact of the work they do
- Creation of a market-wide decision-making body
- Find a balance between workflow logistics and accommodation of clinician requests







Benefits of Collaboration — Stewardship Perspective

- Automation of processes improves timeliness of results
 - $\circ~$ Automation of susceptibility results
 - Cefazolin Kirby Bauer disks on blood isolates
 - Ertapenem reporting for ESBL susceptible isolates (resistant isolates require confirmatory testing)
 - Azole susceptibility testing for candidemia
 - MDRO cascade upon identification of carbapenem resistance
 - $\,\circ\,$ Labeling of resistant isolates with "ESBL" or "CRO"
 - Aids in identification from an infection prevention perspective
 - Faster time to finalization of blood cultures (60% improvement upon implementation of Virtuo[®] automated system)
- Serving as the intermediary between laboratory practices and front-line clinicians enhances clinical impact

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Benefits of Collaboration — Best Practices

Clinical & Laboratory Standards Institute (CLSI):

 Annual CLSI 2023 AST webinar's emphasis on collaboration with local stewardship teams in decisions affecting breakpoints, ESBL testing, comments, and cascade reporting

CDC's Core Elements (2019)

 4 of 7 core elements have potential for microbiology involvement: hospital leadership commitment, action, tracking, reporting

Source: Humphries RM, Bobenchik AM. 2023, April 5. *What's new in the 2023 CLSI standards antimicrobial susceptibility testing (AST)*. CLSI. https://learn.clsi.org/products/clsi-m100-ed33-2023-ast-update-462023-day-2#tab-product_tab_overview Source: Core elements of hospital antibiotic stewardship programs. Centers for Disease Control and Prevention website. p. 1–40. https://www.cdc.gov/antibiotic-use/healthcare/pdfs/hospital-core-elements-H.pdf. Published 2021. Accessed March 17, 2023.



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Benefits of Collaboration: Regulatory



2021: Colorado HQIP Survey

Source: The Joint Commission. (2022). Available at https://www.jointcommission.org/-/media/tjc/documents/standards/r3-reports/r3_antibioticstewardship_july2022_final.pdf Accessed 3/17/2023. Centers for Medicare and Medicaid Services, Conditions of Participation for Hospitals, Infection Prevention and Control and Antibiotic Stewardship Programs, 42 C.F.R §482.42 (2019). CHASE: 2023 Hospital Quality Incentive Payment (HQIP) Program. (2023). Available at: https://hcpf.colorado.gov/sites/hcpf/files/2023%20CO%20HQIP%20Measure%20Details_June%202022_0.pdf. Accessed 5/31/2023. Simner PJ, Rauch CA, Martin IW, et al.. Raising the bar: Improving antimicrobial resistance detection by clinical laboratories by ensuring use of current breakpoints. *Open Forum Infect Dis.* 2022;9 (3). doi.org/10.1093/ofid/ofac007

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Future of Collaboration



Through the microbiology/stewardship collaboration, we have maintained a focus on providing optimal patient care. Taking these steps has helped up adhere to best practices outlined by national organizations and newly implemented regulatory requirements.

Continuation of this partnership will be necessary as breakpoints evolve, new rapid diagnostics are developed, and new antimicrobials are brought to the market.

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Assessment Question 3

In what way(s) does a patient benefit from the collaboration between stewardship programs and the microbiology lab?

- A. Higher likelihood of receiving optimal antimicrobial therapy
- B. Lower medical bill
- C. Enhanced understanding of diagnostic processes
- D. All of the above





Assessment Question 3 | Answer...

In what way(s) does a patient benefit from the collaboration between stewardship programs and the microbiology lab?

- A. Higher likelihood of receiving optimal antimicrobial therapy
- B. Lower medical bill
- C. Enhanced understanding of diagnostic processes
- D. All of the above





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Thank you...

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