

# Antibiograms and Infection Prevention Efforts to Combat Antimicrobial Resistance Post COVID-19

### Welcome!

- All lines are muted, so please ask your questions in Q&A
- For technical issues, chat to the 'Technical Support' Panelist
- Please actively participate in polling questions that pop up on the lower right-hand side of your screen

# We will get started shortly!

### Antibiograms and Infection Prevention Efforts to Combat Antimicrobial Resistance Post COVID-19



Darrell Childress, PharmD, Antimicrobial Stewardship Coordinator Brooke Bailey, Director of Infection Prevention East Alabama Medical Center



The Quality Improvement Services Group of ALLIANT HEALTH SOLUTIONS



### **Collaborators:**

Alabama Hospital Association
Alliant Quality
Comagine Health
Georgia Hospital Association
KFMC Health Improvement Partners
Konza

### **Hospital Quality Improvement**



# Welcome from all of us!











The Quality Improvement Services Group of ALLIANT HEALTH SOLUTIONS

## **Featured Speakers**



Darrell Childress, PharmD

Antimicrobial Stewardship Coordinator
& Pharmacy Residency Director
East Alabama Medical Center



**Brooke Bailey**Director of Infection Prevention
East Alabama Medical Center

### **Learning Objectives**

### • Learn Today:

- Evaluate the impact of COVID-19 on hospital acquired infections (HAI) and changes in resistance patterns
- Review antibiograms and the importance of stratifying based on specific hospital locations
- Discuss EAMC's antimicrobial stewardship efforts in combating antimicrobial resistance in a post COVID environment

### • Use Tomorrow:

Analyze MDRO and HAI data to enhance and stratify antibiograms

### **East Alabama Medical Center**

- 314-bed acute care regional referral center
- Antimicrobial Stewardship Team
  - Infectious Diseases (ID) Physician, ID Pharmacist , clinical microbiologists, and infection prevention nurses





### **Antimicrobial Stewardship**

- Diversion of resources for COVID-19
  - Testing and Reporting
  - Monitoring inpatients
  - Medication procurement
  - Assisting with ID rounds
- Surveillance cultures for patients receiving immunosuppressive therapy for COVD-19

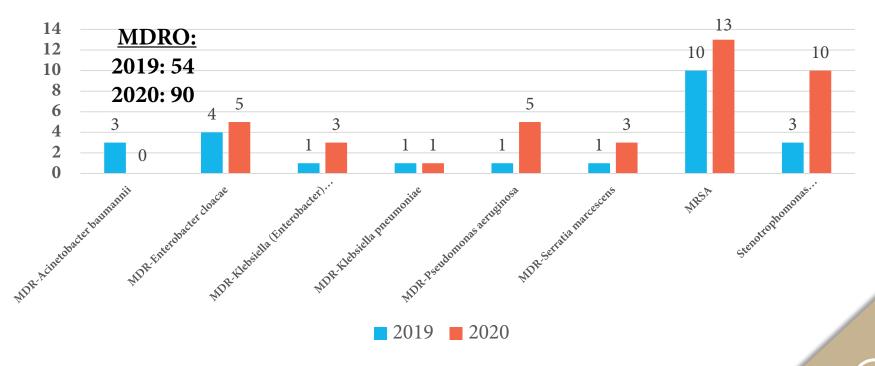
### "We haven't seen that organism before"

- Multi-Drug resistant Organisms (MDRO)
  - Ewingella americana
  - Stenotrophomonas maltophilia
  - Carbapenem Resistant Pseudomonas
  - Carbapenem Resistant Acinetobacter
  - Aspergillus fumigatus

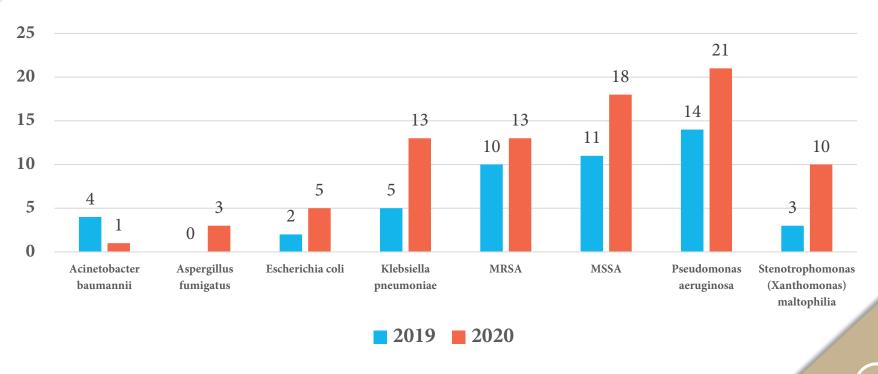
## **Respiratory Organisms 2019-2020**



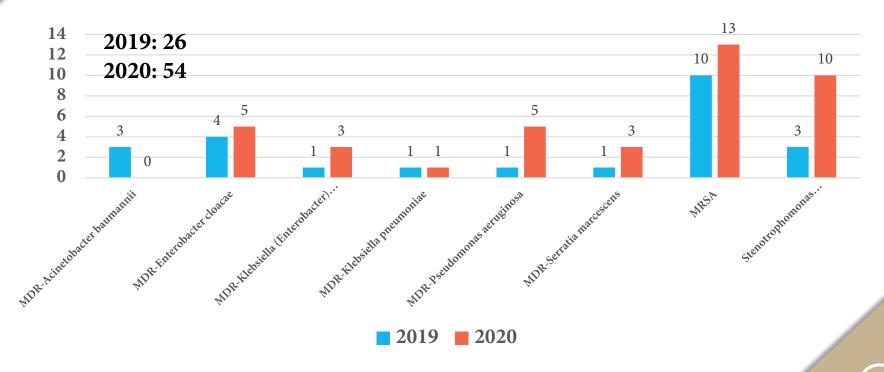
## **MDRO Respiratory 2019-2020**



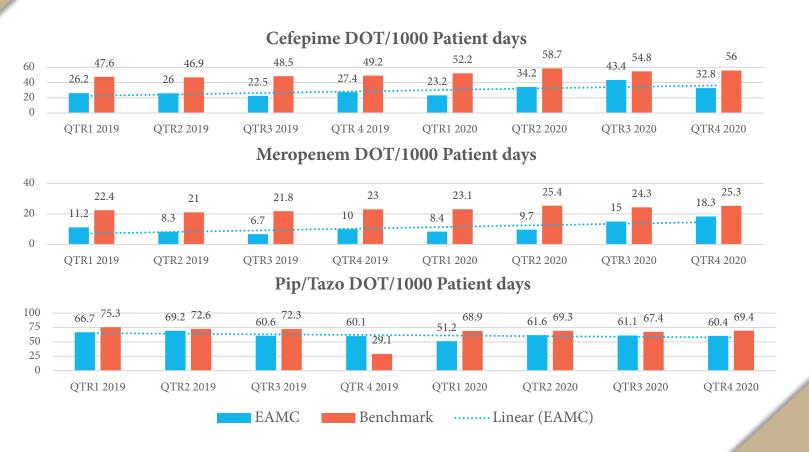
## **ICU Pathogens**



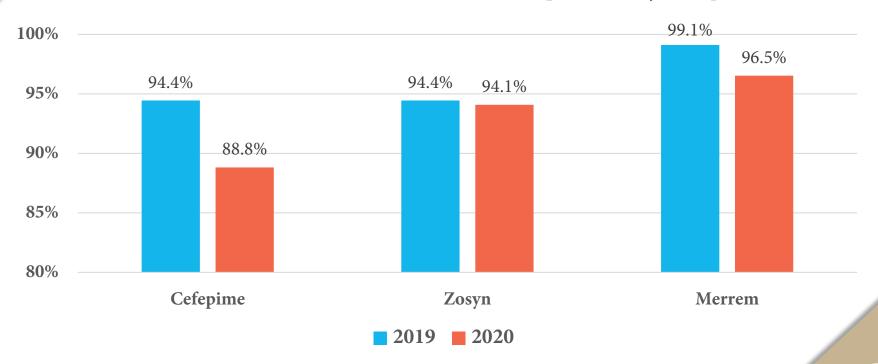
### **MDRO-ICU**



### **Broad Spectrum Antibiotic Usage**



## **Susceptibility of β-Lactams**



# Polling: Is this happening at your facility?

- Is your facility seeing an increase in antimicrobial resistance or MDROs?
  - A. Yes
  - B. No

### **COVID-19's Effect on HAI**

- Grasselli et al., Hospital-acquired infections in critically-ill COVID-19 patients.
  - 46% Patients developed HAI
    - 35% MDRO
    - 50% VAP
    - 34% BSI
    - 10% CLABSI

## **CLABSI and CAUTI during the Pandemic**

Coronavirus disease 2019 (COVID-19) pandemic, central-line–associated bloodstream infection (CLABSI), and catheter-associated urinary tract infection (CAUTI): The urgent need to refocus on hardwiring prevention efforts

- CLABSI rates increased by 51.0% during the pandemic
  - Hospitals with high monthly COVID-19 patients (>10%)
     had SIR 2.38 times higher than hospitals with <5%</li>
- No significant changes were identified for CAUTI

### **Possible Reasons for HAI in COVID-19**

- Unnecessary antimicrobial administration
  - ~80% patients received antibiotics on admission
- Prolonged mechanical ventilation
- Higher device utilization
- Disruption in infection prevention strategies
  - Assessment of central lines, etc.
- Treatment options for COVID
  - Steroids
  - IL-6/IL-1 inhibitors
- Redeployment of staff

### **CLABSI and CAUTI 2019 vs 2020**

	CY2	2019	CY2020						
-	# OF INFECTIONS	SIR	RATE per 1000 catheter days	# OF NFECTIONS	SIR	RATE per 1000 catheter days			
CLABSI									
ALL CMS REPORTED UNITS	0		0	8	1.292	1.06			
ICUs only	0		0	8	2.009	1.74			
CAUTI									
ALL CMS REPORTED UNITS	1	0.17	0.14	8	1.225	1.06			
ICUs only	1	0.395	0.35	5	1.443	1.29			

### **Other Contributing Factors**

- Disruption of Infection Prevention Strategies
- Patient census and acuity levels
  - Higher device utilization
  - Prolonged ventilation
- Existing staff reallocation
- Large amount of contract staff
- Supply deficient
- Alternate care sites

# EAMC's Efforts to Combat HAI Post COVID-19 Peak

## **Infection Prevention Strategies**

- Back to Basics
  - Interdisciplinary collaboration
  - Device utilization/Bundle compliance
  - Appropriate utilization of PPE
  - More frequent rounding
  - Staffing consistency

### **Antibiogram Development**

- Published by the Clinical Laboratory Standards Institute (CLSI)
- Recommendations
  - Report number of isolates tested per timeframe
  - Report species with  $\ge$  30 isolates
  - Report only the first isolate per patient during timeframe
  - Present percentage susceptible



# Polling: Who publishes your antibiogram?

- At your facility, who is in charge of publishing the antibiogram?
  - A. Microbiology
  - **B.** Pharmacy
  - C. A & B
  - D. Other

### **Enhanced Antibiogram**



### Antibiogram and Empiric Treatment Guide 2021

INFECTION TYPE	PATHOGENS	Duration	EMPIRIC TREATMENT	ALTERNATIVE					
RESPIRATORY									
	MRSA/MSSA	7	Vancomycin (PTD)	ICU/CVU:					
	K. pneumonia	7	AND	Recommend ID					
HAP/VAP*	P. peruginosa	7	Cefepime 2 gm fV Q8hrs	consultation due to					
	5. maîtaphilia	7	AND	increased MDRO from					
			Tobramycin (PTD)	COVID-19					
CAP	MRSA/MSSA	7	Ceftriaxone						
	5. pneumaniae	5	2 gm IV Q24hrs	Levaquin 750 mg IV					
	P. aeruginosa	5	AND	Q24hrs					
	H. influenzoe	5	Azithromycin 500 mg IV	4,24ms					
			O24brs	I					



#### Antibiogram and Empiric Treatment Guide 2021

INFECTION TYPE	PATHOGENS	Duration	EMPIRIC TREATMENT	ALTERNATIVE					
RESPIRATORY									
HAP/VAP*	MRSA/MSSA K. pneumonia P. oeruginosa S. maltaphilia	7 7 7	Vancomycin (PTD) AND Cefepime 2 gm IV Q8hrs AND	Recommend ID consultation due to increased MDRO from					
			Tobramycin (PTD)	COVID-19					
CAP	MRSA/MSSA S. pneumaniae P. aeruginosa H. influenzae	7 5 5	Ceftriaxone 2 gm IV Q24hrs AND Azithromycin 500 mg IV Q24hrs	Levaquin 750 mg IV Q24hrs					
INTRA- ABDOMINAL	E. Coli P. mirabilis Anaerobes	4-7 4-7	Piperacillin-Tazobactam 3.375 gm IV QBhrs (4 br infusions)	Cefepime 1 gm IV Q6hr AND Flagyl 500 mg IV Q8hrs					
Clastridioides difficile		10	Vancomycin 125mg PO QID	Vancomycin 125 mg PO QID					
BACTEREMIA*	MRSA/MSSA E. Coli K. pneumoniae	7-14 7-14 7-14	Vancomycin (PTD) AND Piperacillin-Tazobactam 3.375 gm IV Q8hrs (4hr infusions)	Penicillin Allergy: Vancomycin ( <u>PTD)</u> AND Aztreonam 2 gm N Q8hrs					
CELLULITIS	MRSA/MSSA Group A/B Strep	10 10	Vancomycin (PTD)						
DIABETIC FOOT INFECTION	MRSA/MSSA Group B Strep K. pneumoniae E. Coll P. aeruginosa	14-42 14-42 14-42 14-42 14-42	Vancomycin ( <u>PTD)</u> AND Piperacillin-Tazobactam 3.375 gm (V QBhrs 44 br (infusions)	Penicillin Allergy: Vancomycin ( <u>PTD)</u> AND — Aztreonam 2 gm IV Q8hrs					
MENINGITIS†	S. pneumaniae N. meningitides	7	Vancomycin ( <u>PTD)</u> AND Ceftriaxone 2 gm IV Q12hrs	CSF Shunts: Vancomycin ( <u>PTD)</u> AND MEDIOS, 2 gm IV Q8hrs					
URINE E. Coll  K. pneumonipe P. minablis		7 7 7	Ceftriaxone 1 gm IV Q24hrs	Gentamicin (PTD)					

PTD- Pharmacy to Dose

### **Antibiogram Stratifications**

- EAMC stratifies based on:
  - Community versus Hospital acquired
  - Unit\*
    - ICU versus Non-ICU
  - Urine versus non-Urine
    - Respiratory (HAP/VAP/CAP)
    - Intra-abdominal infections (IAI)
    - Skin/Soft Tissues infections (SSTI)
    - Meningitis

### **Rapid Diagnostics**

- Current arsenal
  - MALDI-TOF mass spectrometry
  - Clostridioides difficile PCR
  - BIOFIRE® respiratory panel (2.1)
- Future armament
  - FUNGITELL®
  - BIOFIRE® Pneumonia and Blood culture (BCID2)

### **Summary**

- EAMC is not unique to other healthcare facilities
- Re-education and getting back to basics is key
- Be creative with Antibiograms

### Resource

• CDC's The Core Elements of Hospital Antibiotic Stewardship Programs and Program Assessment Tool (Checklist)

https://www.cdc.gov/antibiotic-use/coreelements/hospital.html

### References

- 1. Grasselli G, Scaravilli V, Mangioni D, et al. Hospital-acquired infections in critically-ill COVID-19 patients. CHEST. 2021 April. https://doi.org/10.1016/j.chest.2021.04.002
- 2. Fakih MG, Bufalino A, Strum L, et. al. Coronavirus disease 2019 (COVID-19) pandemic, central-line–associated bloodstream infection (CLABSI), and catheter-associated urinary tract infection (CAUTI): The urgent need to refocus on hardwiring prevention efforts. Infection Control & Hospital Epidemiology. 2021. Feb. doi:10.1017/ice.2021.70
- 3. Smith L, Karaba SM, Amoah J, et al. Hospital-acquired infections among adult patients admittedfor coronavirus disease 2019 (COVID-19). . Infection Control & Hospital Epidemiology. 2021. doi:10.1017/ice.2021.148



- Learn Today:
  - Evaluate the impact of COVID-19 on hospital acquired infections and changes in resistance patterns
  - Review antibiograms and the importance of stratifying based on specific hospital locations
  - Discuss EAMC's antimicrobial stewardship efforts in combating antimicrobial resistance in a post COVID environment
- Use Tomorrow:
  - Analyze MDRO and HAI data to enhance and stratify antibiograms

How will this change what you do? Please tell us in the poll...

### **Questions?**



Email us at <a href="mailto:HospitalQuality@AlliantQuality.org">HospitalQuality@AlliantQuality.org</a> or call us 678-527-3681

## **Questions?**



	Discharge: Selection & Duration					
Infection	Empiric	Duration of				
Diagnosis	Treatment	Therapy				
	Urinary Tract Infections					
Uncomplicated UTI (cystitis)	Nitrofurantoin (NFT) 100mg BID (crcle30) β-lactams Fosfomycin 3g x 1 dose TMP/SMX (SMT) 1 DS tab BID	NFT: 5 β-lactams: 5-7 Fos: 1 SMT: 3				
Complicated UTI (pyelonephritis)	Levofloxacin 750mg QDay β-lactams TMP/SMX 1-2 DS tab BID	FQs: 5-7 TMP/SMX: 10 β-lactams: 10				
Asymptomatic bacteriuria	Treatment is <b>NOT</b> recommended unless pregnant or GU procedure	0				
	Skin & Soft Tissue Infections					
Non-purulent cellulitis	Cephalexin 500 mg QID Clindamycin 300 mg QID <sup>†</sup>	5				
Purulent cellulitis (after I&D)	TMP/SMX 1 DS tab BID Doxycycline 100mg BID	5				
Animal bite wound	Amox/Clav 875/125mg BID	7-10				
Without comorbidities <sup>††</sup> or risk factors for DRSP	Community Acquired Pneumonia (CAP) Doxycycline 100mg BID Z-Pak Amoxicillin 1000mg TID	5				
Comorbidities or risk factors for DRSP	Amox/Clav 2000/125mg BID + Z-Pak or Doxycycline Cefuroxime 500mg BID + Z-Pak or Doxycycline Levofloxacin 750mg QDay	7 Z-Pak: 5				
	Upper Respiratory Tract Infections					
Bacterial Rhinosinusitis	98% Viral Antibiotics not recommended	Watchful waiting				
Bronchitis	~90% Viral Antibiotics not recommended	Watchful waiting				
Pharyngitis	> 90% Viral Antibiotics not recommended unless GAS pharyngitis	0				
Acute COPD exacerbation	Doxycycline 100mg BID Z-Pak	5-7				
Influenza	Oseltamivir 75mg BID	5				
† Alternative for <i>severe</i> β-lactar †† Comorbidities include chronic heart	m allergy , lung, liver, or renal disease; diabetes mellitus; alcoholism; malignanc	y; or asplenia.				

CO	M	Μ	U	NI	ΤY	' Α	C	Qι	JII	RΕ	D							
All Sources Except Urine																		
GRAM-POSITIVE Community-Aquired	tes	IN G	N.	N	IYCIN	YCIN	OMYCIN	OXACIN	QI	7	rc LINE	J	YCIN					
Percent Susceptible	# of isolates	PENICILLIN	OXACILLIN	AMPICILLIN	CLINDAMYCIN	DAPTOMYCIN	ERYTHROMYCIN	LEVOFLOXACIN	LINEZOLID	RIFAMPIN	TETRACYCLINE	TMP-SMX	VANCOMYCIN					
Enterococcus faecalis	86	100		95		100			100				98					
Enterococcus spp (all)	73	100	_	100		100			100				98					
Staphylococcus aureus	301	6	54		81	100	42		100	100	93	97	100					
Staphylococcus aureus - MRSA	136	0	0		80	100	16		100	100	94	94	100					
Staphylococcus aureus - MSSA	165	12	100		81	100	64		100	100	92	100	100	l.				
Staphylococcus coagulase negative - ALL	69	12	47		69	100	40		100	98	76	81	100					
Streptococcus agalactiae (Group B)	67	100			47			98										
All Sources Except Urine					_	W												
GRAM-NEGATIVE Community-Aquired	# of isolates	NCIN	GENTAMICIN	FOBRAMYCIN	AMOXICILLINICA	AMP/SULBACT AM	AMPICILLIN	20	CEFAZOLIN	NILDX	CEFTAZIDIME	CEFTRIAXONE	JIME	MEROPENEM	AZTREONAM	EVOFLOXACIN	WX	
Percent Susceptible	# of is	AMIKACIN	GENT	TOBR	AMOX	AMP/9	AMPIC	PIP/TAZO	CEFA	CEFOXITIN	CEFT,	CEFTI	CEFEPIME	MERC	AZTR	LEVO	TMP-SMX	
Escherichia coli	155	99	87	89	83	57	54	98	87	96	92	92	94	100	93	85	76	
Klebsiella pneumoniae	73	100	87	87	86	76	0	97	86	93	86	84	86	100	84	97	79	
Proteus mirabilis	58	100	96	96	91	100	86	98	100	100	100	100	100	100	96	84	84	
Pseudomonas aeruginosa	92	94	90	98				97			96		94	100	85	83		
Urine Isolates Only																		
GRAM-POSITIVE Community-Acquired	tes	S NI	z	NI	YCIN	XACIN	Q	MITROFURANTOIN	-	CLINE		YCIN						
Percent Susceptible	# of isolates	PENICILLIN G	OXACILLIN	AMPICILLIN	DAPTOMYCIN	LEVOFLOXACIN	LINEZOLID	NITROFU	RIFAMPIN	TETRACYCLINE	TMP/SMX	VANCOMYCIN						
Enterococcus faecalis	122	100		99	100	71	100	100		27		96						
Enterococcus spp (all)	129	100		96	100	69	100	99		28		95						
Staphylococcus coagulase negative -(all)	68		44		100	77	100	100	100	83	77	100						
Staphylococcus epidermidis	40		50		100	70	100	100	100	82	67	100						
Streptococcus agalactiae (Group B)	31	100	_			100							L					
Urine Isolates Only																		
			×	CIN	-IN/CA	AMP/SULBACTAM	z		2		ME	ONE		ЕМ	W	CACIN	NITROFURANTOIN	
GRAM-NEGATIVE	ş				. ⊒	"	3	0	딩	Ē		SIAX	M	EN	혽	0	ä	×
GRAM-NEGATIVE Community-Acquired Percent Susceptible	of isolates	MIKACIN	ENTAM	OBRAM	MOXIC	MP/SU	MPICI	P/TA2	EF AZ	EF0)	149	EFT	99	IEROF	ZTREC	EVOF	ITROF	MS/dW.
Community-Acquired	4 of isolates	AMIKACIN	S GENTAMICIN	TOBRAMYCIN	MOXICILLIN/CA	_	SAMPICILLIN	₩ PIP/TAZO	CEFAZOLIN	S CEFOXITIN	S CEFTAZIDIME	8 CEFTRIAXONE	S CEFEPIME	MEROPENEM	2 AZTREONAM	EVOFLOXACIN	-	-
Community-Acquired Percent Susceptible  Escherichia coli	#	100	92	92	84	56	S AMPICI	98	83	92	90	89	92	100	91	140A31 81 84	8 8 NITROF	73
Community-Acquired Percent Susceptible	700	_	_		_	_			1		_	_	_	•	1	81	96	73 8'

# **HQIC** Goals



# Behavioral Health Outcomes & Opioid Misuse

- ✓ Promote opioid best practices
- ✓ Decrease high dose opioid prescribing and opioid adverse events in all settings
- ✓ Increase access to behavioral health services



### **Patient Safety**

- ✓ Reduce risky medication combinations
- ✓ Reduce adverse drug events
- ✓ Reduce *C. diff* in all settings



# **Quality of Care Transitions**

- ✓ Convene community coalitions
- ✓ Identify and promote optical care for super utilizers
- ✓ Reduce community-based adverse drug events

## **Upcoming Events**



June 22, 2021, 2:00 p.m. EST

Monoclonal Antibody Therapy for

High Risk COVID Patients

<a href="https://bit.ly/3tXjdUS">https://bit.ly/3tXjdUS</a>

July 27, 2021, 2:00 p.m. EST
Opioid Pain Management
<a href="https://bit.ly/2RZJYLo">https://bit.ly/2RZJYLo</a>



### **Collaborators:**

Alabama Hospital Association
Alliant Quality
Comagine Health
Georgia Hospital Association
KFMC Health Improvement Partners
Konza

### **Hospital Quality Improvement**













The Quality Improvement Services Group of ALLIANT HEALTH SOLUTIONS



### **Collaborators:**

Alabama Hospital Association
Alliant Quality
Comagine Health
Georgia Hospital Association
KFMC Health Improvement Partners
Konza

### **Hospital Quality Improvement**

This material was prepared by Alliant Quality, the quality improvement group of Alliant Health Solutions (AHS), the Hospital Quality Improvement Contractor (HQIC) under contract with the Centers for Medicare & Medicaid Services (CMS), an agency of the U.S. Department of Health and Human Services. The contents presented do not necessarily reflect CMS policy. Publication No. AHSHOIC-TO3H-21-572



The Quality Improvement Services Group of ALLIANT HEALTH SOLUTIONS