



2024 Multi-site Gram-Negative Surveillance Initiative (MuGSI) Healthcare-Associated Infections Community Interface (HAIC) Case Report

NOTE: Enter all dates as mm/dd/yyyy

Form Approved
OMB No. 0920-0978

PATIENT'S NAME: _____				PHONE NO.: _____	
ADDRESS: _____				MRN: _____	
ADDRESS TYPE: _____		HOSPITAL: _____			
----Patient Identifier information is not transmitted to CDC----					
DEMOGRAPHICS					
1. STATE: _____	2a. COUNTY: _____	2b. PLANNING REGION: _____		3. STATE ID: _____	
4a. LABORATORY ID WHERE INCIDENT SPECIMEN IDENTIFIED: _____			4b. FACILITY ID WHERE PATIENT TREATED: _____		
5. DATE OF BIRTH: _____	7. SEX: _____ Male Female Missing value	8a. ETHNIC ORIGIN: _____ Hispanic or Latino Not Hispanic or Latino Unknown	8b. RACE: (Check all that apply) American Indian or Alaska Native Asian Black or African American Native Hawaiian or Other Pacific Islander White Unknown		
6. AGE: _____ Days Mos Yrs					
9a. DATE OF INCIDENT SPECIMEN COLLECTION (DISC): _____	10. ORGANISM: _____ Carbapenem-Resistant <i>Enterobacteriales</i> (CRE) <i>Escherichia coli</i> <i>Klebsiella pneumoniae</i> <i>Klebsiella oxytoca</i> <i>Klebsiella aerogenes</i> <i>Enterobacter cloacae</i> Extended-Spectrum Beta-Lactamase-producing <i>Enterobacteriales</i> (ESBL-E) <i>Escherichia coli</i> <i>Klebsiella pneumoniae</i> <i>Klebsiella oxytoca</i> Carbapenem-Resistant <i>A. baumannii</i> (CRAB) Invasive <i>Escherichia coli</i> (iEC) (not CRE or ESBL-E)				
9b. TIME OF DISC: _____					
11. SPECIMEN COLLECTION SITE(S): Blood Bone Bronchoalveolar lavage (CRAB only, complete Q23c) CSF Internal body site (specify): _____ Muscle Peritoneal fluid Pericardial fluid Pleural fluid Joint/synovial fluid Sputum (CRAB only, complete Q23c) Tracheal aspirate (CRAB only, complete Q23c) Urine (complete 22a–22c) Wound (specify): _____ (CRAB only) Other LRT site (specify): _____ (CRAB only, complete Q23c) Other normally sterile site (specify): _____					
12. LOCATION OF SPECIMEN COLLECTION:			13. WHERE WAS THE PATIENT LOCATED ON THE 3RD CALENDAR DAY BEFORE THE DISC?		
OUTPATIENT Facility ID: _____ Emergency room Clinic/Doctor's office Dialysis center Surgery Observational/Clinical decision unit Other outpatient			INPATIENT Facility ID: _____ ICU OR Radiology Other inpatient		
LTCF Facility ID: _____ LTACH Facility ID: _____ Autopsy Other (Specify): Unknown			Private residence LTCF Facility ID: _____ Hospital inpatient Facility ID: _____ Was the patient transferred from this hospital? Yes No Unknown		
LTACH Facility ID: _____ Homeless Incarcerated Other (specify): Unknown					
14. WAS THE PATIENT HOSPITALIZED ON THE DAY OF OR IN THE 29 CALENDAR DAYS AFTER THE DISC? Yes No Unknown					
IF YES, DATE OF ADMISSION: _____					
15a. WAS THE PATIENT IN AN ICU IN THE 7 DAYS BEFORE THE DISC? Yes No Unknown					
IF YES, DATE OF ICU ADMISSION: _____ OR Date unknown					
15b. WAS THE PATIENT IN AN ICU ON THE DAY OF INCIDENT SPECIMEN COLLECTION OR IN THE 6 DAYS AFTER THE DISC? Yes No Unknown					
IF YES, DATE OF ICU ADMISSION: _____ OR Date unknown					
Public reporting burden of this collection of information is estimated to average 28 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC/ATSDR Information Collection Review Office, 1600 Clifton Road NE, MS D-74, Atlanta, Georgia 30329; ATTN: PRA (0920-XXXX).					

Survived	DATE OF DISCHARGE: _____ OR	IF SURVIVED, DISCHARGED TO:	
Died	Date unknown	Private residence	Other (specify): _____
Unknown	Left against medical advice (AMA)	LTCF, Facility ID: _____	Unknown
		LTACH, Facility ID: _____	
DATE OF DEATH: _____ OR Date unknown			

Abscess, not skin	Decubitus/pressure ulcer	Pneumonia (CRAB cases, complete Q23c)	Skin abscess
AV fistula/graft infection	Empyema	Pyelonephritis (complete 22a–22c)	Surgical incision infection
Bacteremia	Endocarditis	Sepsis	Surgical site infection (internal)
Bursitis	Epidural abscess	Urosepsis	Traumatic wound
Catheter site infection (CVC)	Meningitis	Septic arthritis	Urinary tract infection (complete 22a–22c)
Cellulitis	Osteomyelitis	Septic emboli	Other (specify):
Chronic ulcer/wound (not decubitus)	Peritonitis	Septic shock	

Yes No Unknown

CHRONIC LUNG DISEASE Cystic fibrosis Chronic pulmonary disease	IMMUNOCOMPROMISED CONDITION HIV infection AIDS/CD4 count < 200 Primary immunodeficiency Transplant, hematopoietic stem cell Transplant, solid organ: _____	NEUROLOGIC CONDITION Cerebral palsy Chronic cognitive deficit Dementia Epilepsy/seizure/seizure disorder Multiple sclerosis Neuropathy Parkinson's disease Other (specify): _____	SKIN CONDITION Burn Decubitus/pressure ulcer Surgical wound Other chronic ulcer or chronic wound Other (specify): _____
CHRONIC METABOLIC DISEASE Diabetes mellitus With chronic complications	LIVER DISEASE Chronic liver disease Ascites Cirrhosis Hepatic encephalopathy Variceal bleeding Hepatitis C Treated, in SVR Current, chronic	PLEGIAS/PARALYSIS Hemiplegia Paraplegia Quadriplegia	OTHER Connective tissue disease Obesity or morbid obesity Pregnant
CARDIOVASCULAR DISEASE CVA/Stroke/TIA Congenital heart disease Congestive heart failure Myocardial infarction Peripheral vascular disease (PVD)	GASTROINTESTINAL DISEASE Diverticular disease Inflammatory bowel disease Peptic ulcer disease Short gut syndrome	RENAL DISEASE Chronic kidney disease Lowest serum creatinine: _____ mg/DL Unknown or not done	MUGSI CONDITIONS Urinary tract problems/abnormalities Premature birth Spina bifida
	MALIGNANCY Malignancy, hematologic Malignancy, solid organ (non-metastatic) Malignancy, solid organ (metastatic)		

Yes No Unknown

	DUD/ ABUSE	MODE OF DELIVERY (Check all that apply)			
Marijuana, cannabinoid (other than smoking)	DUD or abuse	IDU	Skin popping	Non-IDU	Unknown
Opioid, DEA schedule I (e.g., heroin)	DUD or abuse	IDU	Skin popping	Non-IDU	Unknown
Opioid, DEA schedule II-IV (e.g., methadone, oxycodone)	DUD or abuse	IDU	Skin popping	Non-IDU	Unknown
Opioid, NOS	DUD or abuse	IDU	Skin popping	Non-IDU	Unknown
Cocaine	DUD or abuse	IDU	Skin popping	Non-IDU	Unknown
Methamphetamine	DUD or abuse	IDU	Skin popping	Non-IDU	Unknown
Other (specify): _____	DUD or abuse	IDU	Skin popping	Non-IDU	Unknown
Unknown substance	DUD or abuse	IDU	Skin popping	Non-IDU	Unknown

20. RISK FACTORS: <i>(Check all that apply)</i>		None	Unknown
WAS INCIDENT SPECIMEN COLLECTED 3 OR MORE CALENDAR DAYS AFTER HOSPITAL ADMISSION?		Yes	No
PREVIOUS HOSPITALIZATION IN THE YEAR BEFORE DISC		Yes	No Unknown
IF YES, DATE OF DISCHARGE CLOSEST TO DISC: _____ OR, DATE UNKNOWN		Facility ID: _____	
OVERNIGHT STAY IN LTCF IN THE YEAR BEFORE DISC:		Yes	No Unknown Facility ID: _____
OVERNIGHT STAY IN LTACH IN THE YEAR BEFORE DISC:		Yes	No Unknown Facility ID: _____
SURGERY IN THE YEAR BEFORE DISC:		Yes	No Unknown
CURRENT CHRONIC DIALYSIS:		Yes	No Unknown
IF YES, TYPE: Hemodialysis Peritoneal Unknown			
IF HEMODIALYSIS, TYPE OF VASCULAR ACCESS: AV fistula/graft Hemodialysis central line Unknown			
CENTRAL LINE IN PLACE ON THE DISC (UP TO THE TIME OF COLLECTION), OR AT ANY TIME IN THE 2 CALENDAR DAYS BEFORE DISC:		Yes	No Unknown
Check here if central line in place for > 2 calendar days			
URINARY CATHETER IN PLACE ON THE DISC (UP TO THE TIME OF COLLECTION), OR AT ANY TIME IN THE 2 CALENDAR DAYS BEFORE DISC:		Yes	No Unknown
IF YES, CHECK ALL THAT APPLY:			
Indwelling Urethral Catheter Condom Catheter			
Suprapubic Catheter Other (specify): _____			
ANY OTHER INDWELLING DEVICE IN PLACE ON THE DISC UP TO THE TIME OF COLLECTION), OR AT ANY TIME IN THE 2 CALENDAR DAYS BEFORE DISC:		Yes	No Unknown
IF YES, CHECK ALL THAT APPLY:			
ET/NT Tube NG Tube Nephrostomy Tube			
Gastrostomy Tube Tracheostomy Other (specify): _____			
PATIENT TRAVELED INTERNATIONALLY IN THE YEAR BEFORE DISC:		Yes	No Unknown
COUNTRY(IES): _____			
PATIENT HOSPITALIZED WHILE VISITING COUNTRY(IES) ABOVE:		Yes	No Unknown
21a. WEIGHT:		21b. HEIGHT:	
_____ lbs. _____ oz. OR _____ kg Unknown		_____ ft. _____ in. OR _____ cm Unknown	
		21c. BMI:	
		_____ Unknown	
Complete questions 22a-22c for all MuGSI cases from urine cultures or where UTI or pyelonephritis is marked in question 17a:			
22a. WAS THE URINE COLLECTED THROUGH AN INDWELLING URETHRAL CATHETER?		Yes	No Unknown
22b. RECORD THE COLONY COUNT: _____			
22c. ASSOCIATED SIGNS AND SYMPTOMS:			
Please indicate if any of the following symptoms were reported during the 5 day time period including the 2 calendar days before through the 2 calendar days after the DISC.			
None	Fever [temperature ≥ 100.4 °F (38 °C)]	Symptoms for patients ≤ 1 year of age only:	
Unknown	Frequency	Apnea	Lethargy
Costovertebral angle pain or tenderness	Suprapubic tenderness	Bradycardia	Vomiting
Dysuria	Urgency		
Complete questions 23a-23b ONLY for A. BAUMANNII cases:			
23a. DID THE PATIENT HAVE A SPUTUM CULTURE POSITIVE FOR CRAB IN THE 30 DAYS BEFORE THE DISC?		Yes	No Unknown N/A
23b. RISK FACTORS PRIOR TO CRAB DISC: <i>(Check all that apply)</i>			
Non-invasive positive pressure ventilation (CPAP or BiPAP) at any time in the 7 calendar days before the DISC			
Nebulizer treatment at any time in the 7 calendar days before the DISC			
Mechanical ventilation at any time in the 7 calendar days before the DISC			
None of the above			

Complete question 23c ONLY for A. BAUMANNII cases from LRT site cultures or for non-LRT cultures where pneumonia is marked in question 17a.

23c. CHEST RADIOLOGY FINDINGS: *(Check all that apply)*

Not done	Ground glass opacities/infiltrates	Consolidation	Nodules
No report available	Bronchopneumonia/pneumonia	Infiltrate	No evidence of pneumonia
Acute respiratory distress syndrome (ARDS)	Cannot rule out pneumonia	Pleural effusion	
Air space density/opacity	Cavitation		

24a. IS ANTIMICROBIAL USE (IV OR ORAL) IN THE 30 DAYS BEFORE THE DISC DOCUMENTED?

Yes No Unknown

24b. IF YES, CHECK ALL ANTIMICROBIALS USED IN THE 30 DAYS BEFORE THE DISC: *(Check all that apply)*

Unknown

Amikacin	Ceftazidime	Fidaxomicin	Rifaximin
Amoxicillin	Ceftazidime/avibactam	Fosfomycin	Tedizolid
Amoxicillin/clavulanic acid	Ceftizoxime	Gentamicin	Telavancin
Ampicillin	Ceftolozane/tazobactam	Imipenem/cilastatin	Tigecycline
Ampicillin/sulbactam	Ceftriaxone	Levofloxacin	Tobramycin
Azithromycin	Cefuroxime	Linezolid	Trimethoprim
Aztreonam	Cephalexin	Meropenem	Trimethoprim/sulfamethoxazole
Cefadroxil	Ciprofloxacin	Meropenem/vaborbactam	Vancomycin
Cefazolin	Clarithromycin	Metronidazole	IV
Cefdinir	Clindamycin	Moxifloxacin	PO
Cefepime	Dalbavancin	Nitrofurantoin	Other (specify):
Cefiderocol	Daptomycin	Omadacycline	
Cefixime	Delafloxacin	Oritavancin	
Cefotaxime	Doripenem	Penicillin	Other (specify):
Cefoxitin	Doxycycline	Piperacillin/tazobactam	
Cefpodoxime	Ertapenem	Polymyxin B	
Ceftaroline	Eravacycline	Polymyxin E (colistin)	

REMINDER: Any prior antimicrobial use that is not noted above should be documented in the other (specify) field.

25a. DID THE PATIENT HAVE A POSITIVE TEST(S) FOR SARS-CoV-2 (MOLECULAR ASSAY, ANTIGEN, OR OTHER VIRAL TEST, EXCLUDING SEROLOGY) IN THE 90 DAYS BEFORE OR DAY OF THE DISC?

Yes No Unknown

25b. SPECIMEN COLLECTION DATES FOR POSITIVE TESTS IN THE 90 DAYS BEFORE OR THE DAY OF THE DISC:

First positive test: _____ or Date unknown **Most recent positive test:** _____ or Date unknown

25c.COVID-NET CASE ID IN THE YEAR BEFORE OR DAY OF DISC: _____ None or N/A

26. WAS THE INCIDENT SPECIMEN POLYMICROBIAL?

Yes No Unknown

27a. WAS THE INCIDENT SPECIMEN TESTED FOR CARBAPENEMASE GENES?

Yes No Laboratory not testing Unknown

27b. IF YES, WHAT TESTING METHOD WAS USED? *(Check all that apply)*

Non-Molecular Test Methods:

Molecular Test Methods:

CarbaNP	Modified Hodge Test (MHT)
Carbapenemase Inactivation Method (CIM)	RAPIDEC
CPO Detect	Other (specify):
Disk Diffusion/ROSCO Disk	
E-test	Unknown
Modified Carbapenemase Inactivation Method (mCIM)	

Automated Molecular Assay	Streack ARM-D
Carba-R	Other (specify):
Check Points	
MALDI-TOF MS	Unknown
Next Generation Nucleic Acid Sequencing	
PCR	

27c. IF TESTED, WHAT WAS THE TESTING RESULT?

Non-Molecular Test Results:

Molecular Test Results:

Positive	NDM	Pos	Neg	Ind	Unk
Indeterminate	KPC	Pos	Neg	Ind	Unk
Negative	OXA (specify):	Pos	Neg	Ind	Unk
Unknown					
	VIM	Pos	Neg	Ind	Unk
	IMP	Pos	Neg	Ind	Unk
	Other carbapenemase gene (specify):	Pos	Neg	Ind	Unk

28a. WAS THE INCIDENT SPECIMEN TESTED FOR ESBL PRODUCTION OR OTHER BETA-LACTAMASE GENES?

Yes
No
Laboratory not testing
Unknown

28b. IF TESTED, WHAT TESTING METHOD WAS USED? (Check all that apply):

Broth Microdilution (ATi detection)

ESBL well

Expert rule (ATi flag)

Unknown

Broth Microdilution (Manual)

Disk Diffusion

E-test

Molecular test (specify): _____

Gene variant (specify): _____

Other non-molecular test (specify): _____

28c. IF TESTED, WHAT WAS THE RESULT?

Pos Neg Ind Unk

Pos Neg Ind Unk

Pos Neg Ind Unk

Pos Neg Ind Unk

Pos Neg Ind Unk

Pos Neg Ind Unk

Pos Neg Ind Unk

Pos Neg Ind Unk

29. SUSCEPTIBILITY RESULTS:

Please complete the table below based on the information found in the indicated data source.

No susceptibility data from the medical record are available

Antibiotic	Data source:	Data source:	Data source:	Data source:	Data source:	Data source:
	MIC or zone diameter	Interpretation	MIC or zone diameter	Interpretation	MIC or zone diameter	Interpretation
Amikacin						
Amoxicillin/Clavulanate						
Ampicillin						
Ampicillin/Sulbactam						
Aztreonam						
Cefazolin						
Cefepime						
Cefiderocol						
Cefotaxime						
Cefoxitin						
Ceftazidime						
Ceftazidime/Avibactam						
Ceftolozane/Tazobactam						
Ceftriaxone						
Cephalothin						
Ciprofloxacin						
Colistin						
Doripenem						
Doxycycline						
Eravacycline						
Ertapenem						
Fosfomycin						
Gentamicin						
Imipenem						
Imipenem-relebactam						
Levofloxacin						
Meropenem						
Meropenem-vaborbactam						
Minocycline						
Moxifloxacin						
Nitrofurantoin						
Omadacycline						
Piperacillin/Tazobactam						
Plazomicin						
Polymyxin B						
Rifampin						
Tetracycline						
Tigecycline						
Tobramycin						
Trimethoprim-sulfamethoxazole						

30a. WAS THE CASE FIRST IDENTIFIED THROUGH AN AUDIT?

Yes
No

30b. CRF STATUS:

Complete
Pending
Chart unavailable after 3 requests
Complete – pending data

30c. SO INITIALS:

30d. DATE OF ABSTRACTION:

30e. COMMENTS: