



MDRO or CDI Infection Event

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*Required for saving Facility ID:		**Required for completion Event #:	
*Patient ID:		Social Security #:	
Secondary ID:		Medicare #:	
Patient Name, Last:		First:	Middle:
*Gender: M F Other		*Date of Birth:	
Ethnicity (Specify):		Race (Specify):	
Event Details			
*Event Type: [For Event Type = BSI, PNEU, SSI, or UTI use the event specific from]		*Date of Event:	
Post Procedure Event: Yes No		Date of Procedure:	
MDRO/CDI Infection Surveillance: Yes	NHSN Procedure Code:	ICD-10-PCS or CPT Procedure Code:	
*Specific Organism Type: (Select up to 3) <input type="checkbox"/> MRSA <input type="checkbox"/> MSSA <input type="checkbox"/> VRE <input type="checkbox"/> CephR- <i>Klebsiella</i> <input type="checkbox"/> CRE- <i>E. coli</i> <input type="checkbox"/> CRE- <i>Enterobacter</i> <input type="checkbox"/> CRE- <i>Klebsiella</i> <input type="checkbox"/> MDR- <i>Acinetobacter</i> <input type="checkbox"/> <i>C. difficile</i>			
*Date Admitted to Facility:		*Location:	
*Specific Event Type (used only for CDC defined events): Specify Criteria Used (check all that apply)			
<u>Signs and Symptoms</u>		<u>Laboratory or Diagnostic Testing</u>	
<input type="checkbox"/> Abscess	<input type="checkbox"/> Heat	<input type="checkbox"/> Dysuria	<input type="checkbox"/> Organism(s) identified
<input type="checkbox"/> Apnea	<input type="checkbox"/> Hypotension	<input type="checkbox"/> Fever	<input type="checkbox"/> Not cultured
<input type="checkbox"/> Bradycardia	<input type="checkbox"/> Hypothermia	<input type="checkbox"/> Bilious aspirate	<input type="checkbox"/> Organism(s) identified from blood specimen ⁺
<input type="checkbox"/> Cough	<input type="checkbox"/> Lethargy	<input type="checkbox"/> Erythema or redness	<input type="checkbox"/> Other positive laboratory tests ⁺
<input type="checkbox"/> Vomiting	<input type="checkbox"/> Nausea	<input type="checkbox"/> Suprapubic tenderness	<input type="checkbox"/> > 15 colonies cultured from IV cannula tip using semiquantitative culture method
<input type="checkbox"/> Abdominal distension			<input type="checkbox"/> Pneumatosis intestinalis by radiograph
<input type="checkbox"/> Pain or tenderness			<input type="checkbox"/> Portal venous gas (Hepatobiliary gas) by radiograph
<input type="checkbox"/> Drainage or material ⁺			<input type="checkbox"/> Pneumoperitoneum by radiograph
<input type="checkbox"/> Wheezing, rales or rhonchi			<input type="checkbox"/> Imaging test evidence of infection ⁺
<input type="checkbox"/> Diarrhea ⁺			
<input type="checkbox"/> Swelling or inflammation			
<input type="checkbox"/> Occult or gross blood in stools (with no rectal fissure)			
<input type="checkbox"/> Surgical evidence of extensive bowel necrosis (>2 cm of bowel affected)			
<input type="checkbox"/> Surgical evidence of pneumatosis intestinalis with or without intestinal perforation			<u>Clinical Diagnosis</u>
<input type="checkbox"/> Other evidence of infection found on invasive procedure, gross anatomic exam, or histopathologic exam ⁺			<input type="checkbox"/> Physician diagnosis of this event type ⁺
<input type="checkbox"/> Other signs and symptoms ⁺			<input type="checkbox"/> Physician institutes appropriate antimicrobial therapy ⁺
⁺ Per specific site criteria			
<i>Clostridioides difficile</i> Infection			
*Admitted to ICU for CDI complications: Yes No		*Surgery for CDI complications: Yes No	
* Secondary Bloodstream Infection: Yes No			
**Died: Yes No		Event contributed to death? Yes No	
Discharge Date: ___ / ___ / ___		*Pathogens Identified: Yes No If yes, specify on Page 2	
<small>Assurance of Confidentiality: The voluntarily provided information obtained in this surveillance system that would permit identification of any individual or institution is collected with a guarantee that it will be held in strict confidence, will be used only for the purposes stated, and will not otherwise be disclosed or released without the consent of the individual, or the institution in accordance with Sections 304, 306 and 308(d) of the Public Health Service Act (42 USC 242b, 242k, and 242m(d)). Public reporting burden of this collection of information is estimated to average 30 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, Reports Clearance Officer, 1600 Clifton Rd., MS D-74, Atlanta, GA 30333, ATTN: PRA (0920-0666). CDC 57.126 (Front) Rev 6 V. 8.6</small>			

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Pathogen #	Gram-positive Organisms								
_____	<i>Staphylococcus</i> coagulase-negative (specify species if available):		VANC SIRN						
_____	_____ <i>Enterococcus faecium</i>	DAPTO SNSN		GENTHL^s SRN		LNZ SIRN		VANC SIRN	
_____	_____ <i>Enterococcus faecalis</i>								
_____	_____ <i>Enterococcus</i> spp. (Only those not identified to the species level)								
_____	<i>Staphylococcus aureus</i>	CIPRO/LEVO/MOXI SIRN		CLIND SIRN		DAPTO SNSN		DOXY/MINO SIRN	
		ERYTH SIRN		GENT SIRN		LNZ SRN			
		OX/CEFOX/METH SIRN		RIF SIRN		TETRA SIRN		TIG SNSN	
		TMZ SIRN		VANC SIRN					
Pathogen #	Gram-negative Organisms								
_____	<i>Acinetobacter</i> (specify species)	AMK SIRN		AMPSUL SIRN		AZT SIRN		CEFEP SIRN	
		CEFTAZ SIRN		CIPRO/LEVO SIRN		COL/PB SIRN			
		GENT SIRN		IMI SIRN		MERO/DORI SIRN		PIP/PIPTAZ SIRN	
		TMZ SIRN		TOBRA SIRN		TETRA/DOXY/MINO SIRN			
_____	<i>Escherichia coli</i>	AMK SIRN		AMP SIRN		AMPSUL/AMXCLV SIRN		AZT SIRN	
		CEFAZ SIRN		CEFEP SIRN		CEFTAZ SIRN		CIPRO/LEVO/MOXI SIRN	
		CEFTAZ SIRN		CEFUR SIRN		CEFOX/CETET SIRN		COL/PB[†] SRN	
		ERTA SIRN		GENT SIRN		IMI SIRN		MERO/DORI SIRN	
		TIG SIRN		TMZ SIRN		TOBRA SIRN		PIPTAZ SIRN	
		TETRA/DOXY/MINO SIRN							
_____	<i>Enterobacter</i> (specify species)	AMK SIRN		AMP SIRN		AMPSUL/AMXCLV SIRN		AZT SIRN	
		CEFAZ SIRN		CEFEP SIRN		CEFTAZ SIRN		CIPRO/LEVO/MOXI SIRN	
		CEFTAZ SIRN		CEFUR SIRN		CEFOX/CETET SIRN		COL/PB[†] SRN	
		ERTA SIRN		GENT SIRN		IMI SIRN		MERO/DORI SIRN	
		TIG SIRN		TMZ SIRN		TOBRA SIRN		PIPTAZ SIRN	
		TETRA/DOXY/MINO SIRN							
_____	_____ <i>Klebsiella pneumoniae</i>	AMK SIRN		AMP SIRN		AMPSUL/AMXCLV SIRN		AZT SIRN	
		CEFAZ SIRN		CEFEP SIRN		CEFTAZ SIRN		CIPRO/LEVO/MOXI SIRN	
		CEFTAZ SIRN		CEFUR SIRN		CEFOX/CETET SIRN		COL/PB[†] SRN	
		ERTA SIRN		GENT SIRN		IMI SIRN		MERO/DORI SIRN	
		TIG SIRN		TMZ SIRN		TOBRA SIRN		PIPTAZ SIRN	
		TETRA/DOXY/MINO SIRN							
_____	_____ <i>Klebsiella oxytoca</i>	AMK SIRN		AMP SIRN		AMPSUL/AMXCLV SIRN		AZT SIRN	
		CEFAZ SIRN		CEFEP SIRN		CEFTAZ SIRN		CIPRO/LEVO/MOXI SIRN	
		CEFTAZ SIRN		CEFUR SIRN		CEFOX/CETET SIRN		COL/PB[†] SRN	
		ERTA SIRN		GENT SIRN		IMI SIRN		MERO/DORI SIRN	
		TIG SIRN		TMZ SIRN		TOBRA SIRN		PIPTAZ SIRN	
		TETRA/DOXY/MINO SIRN							

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Pathogen #	Gram-negative Organisms (<i>continued</i>)									
_____	<i>Pseudomonas aeruginosa</i>	AMK S I R N	AZT S I R N	CEFEP S I R N	CEFTAZ S I R N	CIPRO/LEVO S I R N	COL/PB S I R N	GENT S I R N		
		IMI S I R N	MERO/DORI S I R N	PIP/PIPTAZ S I R N	TOBRA S I R N					
Pathogen #	Fungal Organisms									
_____	<i>Candida</i> (specify species if available)	ANID S I R N	CASPO S N S N	FLUCO S S-DD R N	FLUCY S I R N	ITRA S S-DD R N	MICA S N S N	VORI S S-DD R N		
Pathogen #	Other Organisms									
_____	Organism 1 (specify)	Drug 1 S I R N	Drug 2 S I R N	Drug 3 S I R N	Drug 4 S I R N	Drug 5 S I R N	Drug 6 S I R N	Drug 7 S I R N	Drug 8 S I R N	Drug 9 S I R N
_____	Organism 1 (specify)	Drug 1 S I R N	Drug 2 S I R N	Drug 3 S I R N	Drug 4 S I R N	Drug 5 S I R N	Drug 6 S I R N	Drug 7 S I R N	Drug 8 S I R N	Drug 9 S I R N
_____	Organism 1 (specify)	Drug 1 S I R N	Drug 2 S I R N	Drug 3 S I R N	Drug 4 S I R N	Drug 5 S I R N	Drug 6 S I R N	Drug 7 S I R N	Drug 8 S I R N	Drug 9 S I R N

Result Codes

S = Susceptible I = Intermediate R = Resistant NS = Non-susceptible S-DD = Susceptible-dose dependent N = Not tested

[§] **GENTHL results: S = Susceptible/Synergistic and R = Resistant/Not Synergistic**

[†] **Clinical breakpoints have not been set by FDA or CLSI, Sensitive and Resistant designations should be based upon epidemiological cutoffs of Sensitive MIC ≤ 2 and Resistant MIC ≥ 4**

Drug Codes:

AMK = amikacin	CEFTRX = ceftriaxone	FLUCY = flucytosine	OX = oxacillin
AMP = ampicillin	CEFUR= cefuroxime	GENT = gentamicin	PB = polymyxin B
AMPSUL = ampicillin/sulbactam	CETET= cefotetan	GENTHL = gentamicin –high level test	PIP = piperacillin
AMXCLV = amoxicillin/clavulanic acid	CIPRO = ciprofloxacin	IMI = imipenem	PIPTAZ = piperacillin/tazobactam
ANID = anidulafungin	CLIND = clindamycin	ITRA = itraconazole	RIF = rifampin
AZT = aztreonam	COL = colistin	LEVO = levofloxacin	TETRA = tetracycline
CASPO = caspofungin	DAPTO = daptomycin	LNZ = linezolid	TIG = tigecycline
CEFAZ= ceftazidime	DORI = doripenem	MERO = meropenem	TMZ = trimethoprim/sulfamethoxazole
CEFEP = cefepime	DOXY = doxycycline	METH = methicillin	TOBRA = tobramycin
CEFOT = cefotaxime	ERTA = ertapenem	MICA = micafungin	VANC = vancomycin
CEFOX= ceftaxime	ERYTH = erythromycin	MINO = minocycline	VORI = voriconazole
CEFTAZ = ceftazidime	FLUCO = fluconazole	MOXI = moxifloxacin	

