



Primary Bloodstream Infection (BSI)

*required for saving **required for completion

Facility ID:		Event #:	
*Patient ID:		Social Security #:	
Secondary ID:		Medicare #:	
Patient Name, Last:		First:	Middle:
*Sex: F M		*Date of Birth:	
Ethnicity: Hispanic or Latino Not Hispanic or Latino Unknown Declined to respond		Race (Select all that apply): American Indian or Alaska Native Asian Black or African American Middle Eastern or North African Native Hawaiian or Pacific Islander White Unknown Declined to respond	
Language: (Specify)		Interpreter Needed: Yes No Declined to Respond Unknown	
*Event Type: BSI		*Date of Event:	
Post-procedure BSI: Yes No		Date of Procedure:	
NHSN Procedure Code:		ICD-10-PCS or CPT Procedure Code:	
*MDRO Infection Surveillance: ☐ Yes, this infection's pathogen & location are in-plan for Infection Surveillance in the MDRO/CDI Module ☐ No, this infection's pathogen & location are not in-plan for Infection Surveillance in the MDRO/CDI Module			
*Date Admitted to Facility:		*Location:	
Risk Factors			
*If ICU/Other locations, Central line: Yes No *If Specialty Care Area/Oncology, Permanent central line: Yes No Temporary central line: Yes No *If NICU, Central line, including umbilical catheter Yes No Birth weight (grams)		Check all that apply: Yes ☐ No ☐ *Any hemodialysis catheter present Yes ☐ No ☐ *Extracorporeal life support present (ECLS or ECMO) Yes ☐ No ☐ *Ventricular-assist device (VAD) present Yes ☐ No ☐ *Known or suspected Munchausen Syndrome by Proxy during current admission Yes ☐ No ☐ *Observed or suspected patient injection into vascular line(s) within the BSI infection window period Yes ☐ No ☐ *Epidermolysis bullosa during current admission Yes ☐ No ☐ *Matching organism is identified in blood and from a site-specific specimen, both collected within the infection window period and pus is present at one of the following vascular sites from which the specimen was collected: ☐ Arterial catheter ☐ Arteriovenous fistula ☐ Arteriovenous graft ☐ Atrial lines (Right and Left) ☐ Hemodialysis reliable outflow (HERO) catheter ☐ Intra-aortic balloon pump (IABP) device ☐ Non-accessed central line (not accessed inserted during the admission) ☐ Peripheral IV or Midline catheter Location of Device Insertion: _____ Date of Device Insertion: ____ / ____ / ____	
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 42 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 H21-8, Atlanta, GA 30333, ATTN: PRA (0920-0666).

CDC 57.108 (Front) Rev. 11 v9.4

Page 2 of 4

Event Details		
*Specific Event: Laboratory-confirmed		
*Specify Criteria Used:		
Signs & Symptoms (check all that apply)		
Any Patient	≤ 1 year old	Underlying conditions for MBI-LCBI (check all that apply):
☐ Fever	☐ Fever	☐ Allo-SCT with Grade ≥ 3 GI GVHD
☐ Chills	☐ Hypothermia	☐ Allo-SCT with diarrhea
☐ Hypotension	☐ Apnea	☐ Neutropenia (WBC or ANC < 500 cells mm ³)
	☐ Bradycardia	
		Laboratory (check one)
		☐ Recognized pathogen from one or more blood specimens
		☐ Common commensal from ≥ 2 blood specimens
**Died: Yes No		BSI Contributed to Death: Yes No
Discharge Date:		*Pathogens Identified: Yes No
COVID-19: Yes No		*If Yes, specify on pages 2-3.
If Yes: ☐ Confirmed ☐ Suspected		

Pathogen #	Gram-positive Organisms																								
	<table border="0"> <tr> <td><i>Staphylococcus coagulase-negative</i></td> <td>CEFOX/IOX SRN</td> <td>VANC SIRN</td> </tr> <tr> <td>(specify species if available):</td> <td></td> <td></td> </tr> </table>	<i>Staphylococcus coagulase-negative</i>	CEFOX/IOX SRN	VANC SIRN	(specify species if available):																				
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CDC 57.108 (Back)



HSN National Healthcare Safety Network			TOBR A S I R N								
	<i>Enterobacter</i> (specify species) _____	AMK S I R N CIPRO/LEVO/ MOXI S I R N MERVAB S I R N	AZT S I R N COL/ PB[†] I R N PIPTAZ S I R N	CEFTAZ S I R N DORI/IMI/ MERO S I R N TIG S I R N	CEFOT/CEFTRX S I R N DOXY/MINO/ TETRA S I R N TMZ S I R N	CEFE P S I/S- DDR N ERTA S I R N TOBR A S I R N	CEFTA VI S R N GENT S I R N	CEFTOTA Z S I R N IMIREL S I R N			
Pathogen #	Gram-negative Organisms (continued)										
	_____ <i>Klebsiella pneumoniae</i> _____ <i>Klebsiella oxytoca</i> _____ <i>Klebsiella aerogenes</i>	AMK S I R N CEFTAVI S R N GENT S I R N	AMPSUL/AMXCLV S I R N CEFTOTAZ S I R N IMIREL S I R N	AZT S I R N CIPRO/LEVO/MOXI S I R N MERVAB S I R N	CEFAZ S I R N COL/PB[†] I R N PIPTAZ S I R N	CEFTAZ S I R N DORI/IMI/MERO S I R N TIG S I R N	CEFOT/CEFTRX S I R N DOXY/MINO/TETRA S I R N TMZ S I R N	CEFEP S I/S- DDR N ERTA S I R N TOBRA S I R N			
	<i>Pseudomonas aeruginosa</i>	AMK S I R N COL/PB S I R N	AZT S I R N DORI/IMI/MERO S I R N	CEFTAZ S I R N GENT S I R N	CEFEP S I R N PIPTAZ S I R N	CEFTAVI S R N TOBRA S I R N	CEFTOTAZ S I R N	CIPRO/LEVO S I R N			
Pathogen #	Fungal Organisms										
	<i>Candida</i> (specify species if available) _____	ANID S I R N	CASPO S I R N	FLUCO S S-DDR N	MICA S I R N	VORI S I 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	
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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 are based on CLSI M100-ED30:2020, Intermediate MIC ≤ 2 and Resistant MIC ≥ 4

Drug Codes:			
AMK = amikacin	CEFTAR = ceftaroline	GENT = gentamicin	OX = oxacillin
AMP = ampicillin	CEFTAVI = ceftazidime/avibactam	GENTHL = gentamicin –high level test	PB = polymyxin B
AMPSUL = ampicillin/sulbactam	CEFTOTAZ = ceftolozane/tazobactam	IMI = imipenem	PIPTAZ = piperacillin/tazobactam
AMXCLV = amoxicillin/clavulanic	CEFTRX = ceftriaxone	IMIREL = imipenem/relebactam	RIF = rifampin



ANID = anidulafungin	CIPRO = ciprofloxacin	LEVO = levofloxacin	TETRA = tetracycline
AZT = aztreonam	CLIND = clindamycin	LNZ = linezolid	TIG = tigecycline
CASPO = caspofungin	COL = colistin	MERO = meropenem	TMZ = trimethoprim/sulfamethoxazole
CEFAZ = ceftazidime	DAPTO = daptomycin	MERVAB = meropenem/vaborbactam	TOBRA = tobramycin
CEFEP = cefepime	DORI = doripenem	METH = methicillin	VANC = vancomycin
CEFOT = cefotaxime	DOXY = doxycycline	MICA = micafungin	VORI = voriconazole
CEFOX = ceftazidime	ERTA = ertapenem	MINO = minocycline	
CEFTAZ = ceftazidime	FLUCO = fluconazole	MOXI = moxifloxacin	

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Page 4 of 4

Custom Fields			
Label		Label	
Comments			