

Surgical Site Infection (SSI)

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*required for saving **required for completion

Facility ID:	Event #:	
*Patient ID:	Social Security #:	
Secondary ID:	Medicare #:	
Patient Name, Last:	First:	Middle:
*Gender: F M Other	*Date of Birth:	
Sex at Birth: F M Unknown	Gender Identity (Specify):	
Ethnicity (Specify):	Race (Specify):	
*Event Type: SSI	*Date of Event:	
*NHSN Procedure Code:	ICD-10-PCS or CPT Procedure Code:	
*Date of Procedure:	*Outpatient Procedure: Yes No	

*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:
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Event Details

*Specific Event:

- | | |
|---|--|
| <input type="checkbox"/> Superficial Incisional Primary (SIP) | <input type="checkbox"/> Deep Incisional Primary (DIP) |
| <input type="checkbox"/> Superficial Incisional Secondary (SIS) | <input type="checkbox"/> Deep Incisional Secondary (DIS) |
| <input type="checkbox"/> Organ/Space (specify site): _____ | |

*Infection present at the time of surgery (PATOS): Yes No

*Specify Criteria Used (check all that apply):

<u>Signs & Symptoms</u>		<u>Laboratory</u>
<input type="checkbox"/> Drainage or material [†]	<input type="checkbox"/> Sinus tract	<input type="checkbox"/> Organism(s) identified
<input type="checkbox"/> Pain or tenderness	<input type="checkbox"/> Hypothermia	<input type="checkbox"/> Culture or non-culture based testing not performed
<input type="checkbox"/> Swelling or inflammation	<input type="checkbox"/> Apnea	<input type="checkbox"/> Organism(s) identified from blood specimen
<input type="checkbox"/> Erythema or redness	<input type="checkbox"/> Bradycardia	<input type="checkbox"/> Organism(s) identified from ≥ 2 periprosthetic specimens
<input type="checkbox"/> Heat	<input type="checkbox"/> Lethargy	<input type="checkbox"/> Other positive laboratory tests [†]
<input type="checkbox"/> Fever	<input type="checkbox"/> Cough	<input type="checkbox"/> Imaging test evidence of infection
<input type="checkbox"/> Incision deliberately opened/drained	<input type="checkbox"/> Nausea	
<input type="checkbox"/> Wound spontaneously dehisces	<input type="checkbox"/> Vomiting	
<input type="checkbox"/> Abscess	<input type="checkbox"/> Dysuria	
<input type="checkbox"/> Other evidence of infection found on invasive procedure, gross anatomic exam, or histopathologic exam [†]		
<input type="checkbox"/> Other signs & symptoms [†]		

Clinical Diagnosis

Physician diagnosis of this event type
 Physician institutes appropriate antimicrobial therapy[†]

[†]per specific site criteria

- *Detected: A (During admission) P (Post-discharge surveillance)
 RF (Readmission to facility where procedure performed)
 RO (Readmission to facility other than where procedure was performed)

*Secondary Bloodstream Infection: Yes No	**Died: Yes No	SSI Contributed to Death: Yes No
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Discharge Date:	*Pathogens Identified: Yes No	*If Yes, specify on pages 2-3.
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*COVID-19: Yes No

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 35 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 57.120 (Front) Rev 7, v8.6

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Pathogen #	Gram-positive Organisms							
	<i>Staphylococcus coagulase-negative</i> (specify species if available):	CEFOX/OX S R N	VANC SIR N					
	<u>Enterococcus faecium</u> <u>Enterococcus faecalis</u> <u>Enterococcus</u> spp. (Only those not identified to the species level)	DAPTO S I/S-DD NS R N	GENTHL^s S R N	LNZ SIR N	VANC SIR N			
	<i>Staphylococcus aureus</i>	CEFOX/METH/OX S R N	CEFTAR S S-DD I R N	CIPRO/LEVO/MOXI SIR N	CLIND SIR N	DAPTO S NS N	DOXY/MINO SIR N	GENT SIR N
		LNZ S R N	RIF SIR N	TETRA SIR N	TMZ SIR N	VANC SIR N		
Pathogen #	Gram-negative Organisms							
	<i>Acinetobacter</i> (specify species)	AMK SIR N	AMPSU L SIR N	CEFE P SIR N	CEFTAZ/CEFOT/CEFTRX SIR N	CIPRO/LEVO SIR N	COL/PB S R N	DORI/MERO SIR N
		DOXY/MINO SIR N	GENT SIR N	IMI SIR N	PIPTAZ SIR N	TMZ SIR N	TOBRA SIR N	
	<i>Escherichia coli</i>	AMK SIR N	AMP SIR N	AMPSUL/AMXCLV SIR N	AZT SIR N	CEFAZ SIR N	CEFEP S I/S-DD R N	CEFOT/CEFTRX SIR N
		CEFTAVI I SIR N	CEFTA Z SIR N	CEFTOTAZ SIR N	CIPRO/LEVO/MOXI SIR N	COL/PB[†] I R N	DORI/IMI/MERO SIR N	DOXY/MINO/TETRA SIR N
		ERTA SIR N	GENT SIR N	IMIREL SIR N	MERVAB SIR N	PIPTAZ SIR N	TIG SIR N	TMZ SIR N
		TOBRA SIR N						
	<i>Enterobacter</i> (specify species)	AMK SIR N	AZT SIR N	CEFEP S I/S-DD R N	CEFOT/CEFTRX SIR N	CEFTAVI SIR N	CEFTAZ SIR N	CEFTOTAZ SIR N
		CIPRO/LEVO/MOXI SIR N	COL/PB[†] I R N	DORI/IMI/MERO SIR N	DOXY/MINO/TETRA SIR N	ERTA SIR N	GENT SIR N	IMIREL SIR N
		MERVAB SIR N	PIPTAZ SIR N	TIG SIR N	TMZ SIR N	TOBRA SIR N		

Pathogen #	Gram-negative Organisms (continued)							
	____ <i>Klebsiella pneumoniae</i>	AMK SIR N	AMPSUL/ AMXCLV SIR N	AZT SIR N	CEFAZ SIR N	CEFEP S I/S-DD R N	CEFOT/CEFTRX SIR N	CEFTAV I SIR N
	____ <i>Klebsiella oxytoca</i>	CEFTA Z SIR N	CEFTOTAZ SIR N	CIPRO/LEVO/ MOXI SIR N	COL/PB [†] IR N	DORI/IMI/ MERO SIR N	DOXY/MINO/ TETRA SIR N	ERTA SIR N
	____ <i>Klebsiella aerogenes</i>	GENT SIR N	IMIREL SIR N	MERVAB SIR N	PIPTAZ SIR N	TIG SIR N	TMZ SIR N	TOBRA SIR N
Pathogen #	Fungal Organisms							
	<i>Candida</i> (specify species if available)	ANID SIR N	CASPO SIR N	FLUCO S S-DD R N	MICA SIR N	VORI SIR N		
Pathogen #	Other Organisms							
	Organism 1 (specify)	Drug 1 SIR N	Drug2 SIR N	Drug3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 7 SIR N
	Organism 1 (specify)	Drug 1 SIR N	Drug2 SIR N	Drug3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 7 SIR N
	Organism 1 (specify)	Drug 1 SIR N	Drug2 SIR N	Drug3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 8 SIR 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 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 acid	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 = cefazolin	DAPTO = daptomycin	MERVAB = meropenem/vaborbactam	TOBRA = tobramycin
CEFEP = cefepime	DORI = doripenem	METH = methicillin	VANC = vancomycin
CEFOT = cefotaxime	DOXY = doxycycline	MICA = micafungin	VORI = voriconazole
CEFOX = cefoxitin	ERTA = ertapenem	MINO = minocycline	
CEFTAZ = ceftazidime	FLUCO = fluconazole	MOXI = moxifloxacin	



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Label	/	/	Label
Comments			