

# Urinary Tract infection (UTI)

Page 1 of 4

\*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:	
Ethnicity (Specify):	Race (Specify):	
*Event Type: UTI	*Date of Event:	
Post-procedure UTI: 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**
**\*Urinary Catheter status:**

- In place – Urinary catheter in place > 2 days on the date of event or present for any portion of the calendar day       Removed – Urinary catheter in place > 2 days and removed the day before the date of event       Neither – Not catheter associated – Neither in place nor removed

 Location of Device Insertion: \_\_\_\_\_ Date of Device Insertion: \_\_\_\_ / \_\_\_\_ / \_\_\_\_  
 If NICU, birth weight (gms): \_\_\_\_\_

**Event Details**

 \*Specific Event:  Symptomatic UTI (SUTI)  Asymptomatic Bacteremic UTI (ABUTI)  Urinary System Infection (USI)

**\*Specify Criteria Used: (check all that apply)**
Signs & Symptoms

<u>Any Patient</u>		<u>≤ 1 year old</u>	<u>Laboratory &amp; Diagnostic Testing</u>
<input type="checkbox"/> Fever	<input type="checkbox"/> Urgency	<input type="checkbox"/> Fever	<input type="checkbox"/> Positive culture with no more than 2 species of organisms, at least one of which is a bacterium of $\geq 10^5$ CFU/ml
<input type="checkbox"/> Frequency	<input type="checkbox"/> Dysuria	<input type="checkbox"/> Hypothermia	<input type="checkbox"/> Organism(s) identified from fluid or tissue from affected site (excluding urine)
<input type="checkbox"/> Pain or tenderness	<input type="checkbox"/> Abscess	<input type="checkbox"/> Apnea	<input type="checkbox"/> Organism(s) identified from blood specimen
<input type="checkbox"/> Acute pain, swelling, or tenderness of testes, epididymis, or prostate		<input type="checkbox"/> Bradycardia	<input type="checkbox"/> Imaging test evidence of infection
<input type="checkbox"/> Suprapubic tenderness		<input type="checkbox"/> Lethargy	
<input type="checkbox"/> Costovertebral angle pain or tenderness		<input type="checkbox"/> Vomiting	
<input type="checkbox"/> Purulent drainage from affected site		<input type="checkbox"/> Suprapubic tenderness	
<input type="checkbox"/> Other evidence of infection found on invasive procedure, gross anatomic exam, or histopathologic exam <sup>‡</sup>			

<sup>‡</sup> per specific site criteria

 \*Secondary Bloodstream Infection: Yes No COVID-19: Yes No  
 If Yes: Suspected  Confirmed 

 \*\*Died: Yes No UTI Contributed to Death: Yes No  
 Discharge Date: \*Pathogens Identified: Yes No \*If Yes, specify on pages 2-4.

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 20 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.



National Healthcare  
Safety Network

Page 1 of 4

ATTN: PRA (0920-0666). CDC 57.114 (Front) Rev 12, v8.8

Form Approved  
OMB No. 0920-0666  
Exp. Date: 12/31/2022  
[www.cdc.gov/nhsn](http://www.cdc.gov/nhsn)

\*required for saving \*\*required for completion

## Urinary Tract infection (UTI)

Page 2 of 4

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

## Urinary Tract infection (UTI)

Page 3 of 4

Pathogen #	<b>Gram-negative Organisms (continued)</b>									
_____	<i>Pseudomonas aeruginosa</i>	AMK SIR N	AZT SIR N	CEFEP SIR N	CEFTAZ SIR N	CIPRO/LEVO SIR N	COL/PB SIR N	GENT SIR N		
_____		IMI SIR N	MERO/DORI SIR N		PIP/PIPTAZ SIR N	TOBRA SIR N				
Pathogen #	<b>Fungal Organisms</b>									
_____	<i>Candida</i> (specify species if available)	ANID SIR N	CASPO S NS N	FLUCO S S-DD R N	FLUCY SIR N	ITRA S S-DD R N	MICA S NS N	VORI S S-DD R N		
Pathogen #	<b>Other Organisms</b>									
_____	Organism 1 (specify)	Drug 1 SIR N	Drug 2 SIR N	Drug 3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 7 SIR N	Drug 8 SIR N	Drug 9 SIR N
_____	Organism 1 (specify)	Drug 1 SIR N	Drug 2 SIR N	Drug 3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 7 SIR N	Drug 8 SIR N	Drug 9 SIR N
_____	Organism 1 (specify)	Drug 1 SIR N	Drug 2 SIR N	Drug 3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 7 SIR N	Drug 8 SIR N	Drug 9 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 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	CEPTRX = ceftriaxone	FLUCY = flucytosine	OX = oxacillin
AMP = ampicillin	CEFUR= cefuroxime	GENT = gentamicin	PB = polymyxin B
AMPSUL = ampicillin/sulbactam	CTET= 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= cefazolin	DORI = doripenem	MERO = meropenem	TMZ = trimethoprim/sulfamethoxazole
CEFEP = cefepime	DOXY = doxycycline	METH = methicillin	TOBRA = tobramycin
CEFOT = cefotaxime	ERTA = ertapenem	MICA = micafungin	VANC = vancomycin
CEFOX= cefoxitin	ERYTH = erythromycin	MINO = minocycline	VORI = voriconazole
CEFTAZ = ceftazidime	FLUCO = fluconazole	MOXI = moxifloxacin	



## **Urinary Tract infection (UTI)**

Page 4 of 4

Custom Fields			
Label		Label	
_____ _____ _____ _____ _____	/ ____/ _____ _____ _____ _____	_____ _____ _____ _____ _____	/ ____/ _____ _____ _____ _____
Comments			