

## Instructions for Completion of Urinary Tract Infection (UTI) Form (CDC 57.114)

Data Field	Instructions for Data Collection/Entry
Facility ID	The NHSN-assigned facility ID will be auto-entered by the computer.
Event #	Event ID number will be auto-entered by the computer.
Patient ID	Required. Enter the alphanumeric patient ID number. This is the patient
	identifier assigned by the hospital and may consist of any combination of
	numbers and/or letters.
Social Security #	Optional. Enter the 9-digit numeric patient Social Security Number.
Secondary ID	Optional. Enter the alphanumeric ID number assigned by the facility.
Medicare #	Optional. Enter the patient's Medicare number.
Patient name	Optional. Enter the last, first, and middle name of the patient.
Gender	Required. Check Female, Male, or Other to indicate the gender of the
	patient.
Sex at Birth	Optional. Select the patient's sex assigned at birth.
(Birth Sex)	Male
	Female
	Unknown
Gender Identity	Optional. Specify the gender identity/identities which most closely
	matches how the patient self-identifies. Multiple selections are allowed,
	except when selecting 'Asked but unknown.'
	Male
	Female
	Male-to-female transgender
	Female-to-male transgender
	Identifies as non-conforming
	Other
	Asked but unknown
Date of birth	Required. Record the date of the patient birth using this format:
	MM/DD/YYYY.
Ethnicity	Optional. Specify if the patient is either Hispanic or Latino, or Not Hispanic
	or Not Latino.
Race	Optional. Specify one or more of the choices below to identify the
	patient's race:
	American Indian/Alaska Native
	Asian
	Black or African American
	Native Hawaiian/Other Pacific Islander
	White



Data Field	Instructions for Data Collection/Entry
Event type	Required. UTI.
Date of event	Required. The date when the <i>first</i> element used to meet the UTI infection criterion occurred for the first time, during the Infection Window Period. Enter date of this event using this format: MM/DD/YYYY. NOTE: If a device has been discontinued on the first day of the month in a location where there are no other device days in that month, and a device-associated infection develops after the device is discontinued, use the last day of the previous month as the Date of Event.
Post-procedure UTI	Optional. Check Y if this event occurred after an NHSN-defined procedure but before discharge from the facility, otherwise check N.
Date of procedure	Conditionally required. If Post-procedure UTI = Y, Record the date when the NHSN procedure started.
NHSN procedure code	Conditionally required. If Post-procedure UTI = Y, enter the appropriate NHSN procedure code. <b>NOTE:</b> A UTI cannot be "linked" to an operative procedure unless that procedure has already been added to NHSN. If the procedure was previously added, and the "Link to Procedure" button is clicked, the fields pertaining to the operation will be auto-entered by the computer.
ICD-10-PCS and CPT procedure code	Optional. The <u>ICD-10-PCS</u> or <u>CPT</u> code may be entered here instead of (or in addition to) the NHSN Procedure Code. If the ICD-10-PCS or CPT code is entered, the NHSN procedure code will be auto-entered by the computer. If the NHSN code is entered first, you will have the option to select the appropriate ICD-10-PCS or CPT code. In either case, it is optional to select the ICD-10-PCS or CPT code. The NHSN ICD-10-PCS and CPT codes are found in the "Operative Procedure Code Documents" section of the <u>Surgical Site Infection (SSI) Events</u> page on the NHSN website.
MDRO Infection Surveillance	<ul> <li>Required. Enter "Yes", if the pathogen is being followed for Infection</li> <li>Surveillance in the MDRO/CDI Module in that location as part of your</li> <li>Monthly Reporting Plan: MRSA, MSSA (MRSA/MSSA), VRE, CephR-</li> <li><i>Klebsiella</i>, CRE (<i>E. coli, Klebsiella pneumoniae, Klebsiella oxytoca, Klebsiella aerogenes</i>, or <i>Enterobacter</i>), MDR-<i>Acinetobacter</i>, or <i>C. difficile</i>.</li> <li>If the pathogen for this infection happens to be an MDRO but your facility is not following the Infection Surveillance in the MDRO/CDI Module in your Monthly Reporting Plan, answer "No" to this question.</li> </ul>
Location	Required. Enter the inpatient location to which the patient was assigned on the date of the UTI event. If the date of the UTI occurs on the day of transfer/discharge or the next day, indicate the transferring/ discharging location, not the current location of the patient, in accordance with the Transfer Rule (see <u>Key Terms section</u> ).

Data Field	Instructions for Data Collection/Entry
Date admitted to facility Date admitted to facility Risk factor: Urinary catheter status on the date of event	<ul> <li>Required. Enter date patient admitted to an inpatient location using this format: MM/DD/YYY.</li> <li>NOTES:         <ul> <li>When determining a patient's admission dates to both the facility and specific inpatient location, the NHSN user must consider any days spent in an inpatient location as an "observation" patient before being formally admitted as an inpatient to the facility, as these days contribute to exposure risk. Therefore, all such days are included in the counts of admissions and patient days for the facility and specific location, and facility and admission dates must be moved back to the first day spent in the inpatient location. All inpatient locations are eligible for use with determining date admitted to the facility.</li> <li>When reporting a UTI which occurs on the day of or day after discharge use the previous date of admission as admission date.</li> </ul> </li> <li>Required. Check one of the following:         <ul> <li>"In place" if a urinary catheter that had been in place in the inpatient location for more than 2 consecutive calendar days was present for any portion of the calendar day on the date of event</li> <li>"Removed" if a urinary catheter that had been in place in the inpatient location for more than 2 consecutive calendar days was removed the day before the date of event</li> <li>"Neither" if:                 <ul> <li>Patient has/had an indwelling urinary catheter, but it has/had not been in place more than 2 consecutive days in the inpatient location on the date of event</li> <li>OR</li> <li>Patient did not have a urinary catheter in place in the inpatient location on the date of event</li> <li>OR</li> <li>Patient did not have a urinary catheter in place in the inpatient location on the date of event</li> <li>OR</li> <li>Patient did not have a urinar</li></ul></li></ul></li></ul>
Location of device insertion	Optional. Enter the patient location where the IUC was inserted.
Date of device insertion	Optional. Enter the date the IUC was inserted.
Event details:	Required. Check Symptomatic UTI (SUTI), Asymptomatic Bacteremic UTI
Specific event: UTI	(ABUTI) type you are reporting.
Event details: UTI	Required. Check each of the elements of the criteria that were used to
Specify criteria used	identify the specific type of UTI being reported.
Event Details: Secondary	Required. Check Y if there is a bloodstream infection (BSI) identified
bloodstream infection	related to UTI, otherwise check N. For detailed instructions on identifying whether the blood specimen identification represents a secondary BSI,
	refer to the Secondary BSI Guide (Appendix B of the <u>BSI</u> protocol).

Data Field	Instructions for Data Collection/Entry
COVID-19	Required. Check Y if the patient met the definition of confirmed COVID-19
	on the date of event; otherwise, check N.
	Confirmed: A patient with a positive COVID-19 (SARS-CoV-2) laboratory
	viral test indicating current infection (NOTE: this does not include serology
	testing for antibody).
	<ul> <li>Answer COVID-19 as 'YES' if the patient's lab test confirmed</li> </ul>
	COVID-19 prior to or on the date of event. Keep in mind that
	patients may undergo repeat testing post-treatment and may
	move from a 'confirmed' to 'negative' COVID-19 status.
	<ul> <li>Answer COVID-19 as 'NO' if the most recent lab test prior to or on</li> </ul>
	the date of event is negative.
Event Details:	Required. Check Y if patient died during the hospitalization, otherwise
Died	check N.
Event Details:	Conditionally required. If patient died, check Y if such evidence is available
UTI contributed to death	(for example death/discharge note, autopsy report, etc.).
Event Details:	Optional. Date patient discharged from facility.
Discharge date	
Event Details:	Required. Enter Y if pathogen identified, otherwise check N. If Y, specify
Pathogens identified	organism name on reverse.
Pathogen # for specified	Up to two urine pathogens may be reported for the initial UTI event.
Gram-positive Organisms,	If secondary DCI nother and are entered, they should be entered only often
Gram-negative Organisms, or Other	If secondary BSI pathogens are entered, they should be entered only after site-specific pathogens are entered. A third pathogen field is available in
Organisms	cases of secondary BSI to primary UTI events when additional eligible
organisms	organisms are identified in the same blood specimen as the matching
	pathogen, or when a new urine pathogen is identified in the Repeat
	Infection Timeframe.
	Enter the pathogen judged to be the most important cause of infection as
	#1, the next most as #2, and the least as #3 (usually this order will be
	indicated on the laboratory report). If the species is not given on the lab
	report or is not found on the NHSN organism list, then select the genus for
	example <i>Bacillus natto</i> is not on the list so would be reported as <i>Bacillus</i> .
Antimicrobial agent and	Conditionally required if Pathogen Identified = Y.
susceptibility results	<ul> <li>For those organisms shown on the back of an event form,</li> </ul>
	susceptibility results are required only for the agents listed.
	<ul> <li>For organisms that are not listed on the back of an event form, the optimum of susceptibility regults is optional.</li> </ul>
	entry of susceptibility results is optional. Circle the pathogen's susceptibility result using the codes on the event
	forms.
	For each box listing several drugs of the same class, at least one drug
	susceptibility must be recorded.

Data Field	Instructions for Data Collection/Entry
Custom Fields	Optional. Up to 50 fields may be customized for local or group use in any combination of the following formats: date (MM/DD/YYYY), numeric, or
	alphanumeric.
	NOTE: Each Custom Field must be set up in the Facility/Custom Options
	section of the application before the field can be selected for use.
Comments	Optional. Enter any information on the event.