Urinary Tract Infection (Catheter-Associated Urinary Tract Infection [CAUTI] and Non-Catheter-Associated Urinary Tract Infection [UTI]) and Other Urinary System Infection [USI]) Events

Introduction: Urinary tract infections (UTIs) are the fourth most common type of healthcare-associated infection, with an estimated 93,300 UTIs in acute care hospitals in 2011. UTIs additionally account for more than 12% of infections reported by acute care hospitals\(^1\). Virtually all healthcare-associated UTIs are caused by instrumentation of the urinary tract.

Approximately 12%-16% of adult hospital inpatients will have an indwelling urinary catheter at some time during their hospitalization, and each day the indwelling urinary catheter remains, a patient has a 3%-7% increased risk of acquiring a catheter-associated urinary tract infection (CAUTI).\(^2,3\)

CAUTI can lead to such complications as prostatitis, epididymitis, and orchitis in males, and cystitis, pyelonephritis, gram-negative bacteremia, endocarditis, vertebral osteomyelitis, septic arthritis, endophthalmitis, and meningitis in patients. Complications associated with CAUTI cause discomfort to the patient, prolonged hospital stay, and increased cost and mortality\(^4\). It has been estimated that each year, more than 13,000 deaths are associated with UTIs.\(^5\)

Prevention of CAUTI is discussed in the CDC/HICPAC document, Guideline for Prevention of Catheter-associated Urinary Tract Infection.\(^6\)

Settings: Surveillance may occur in any inpatient location(s) where denominator data can be collected, such as critical intensive care units (ICU), specialty care areas (SCA), step-down units, wards, inpatient rehabilitation locations, and long term acute care locations. Neonatal ICUs may participate, but only off plan (not as a part of their monthly reporting plan). A complete listing of inpatient locations and instructions for mapping are located in the CDC Locations and Descriptions chapter.

Note: Surveillance for CAUTI after the patient is discharged from the facility is not required. However, if discovered, any CAUTI with a date of event (DOE) on the day of discharge or the next day is attributable to the discharging location and should be included in any CAUTIs reported to NHSN for that location (see Transfer Rule Chapter 2). No additional indwelling catheter days are reported.
Refer to the NHSN Patient Safety Manual, Chapter 2 Identifying Healthcare Associated Infections in NHSN and Chapter 16 NHSN Key Terms for definitions of the following universal concepts for conducting HAI surveillance.

I. Date of event (DOE)
II. Healthcare associated infection (HAI)
III. Infection window period (IWP)
IV. Present on admission (POA)
V. Repeat infection timeframe (RIT)
VI. Secondary BSI attribution period (SBAP)
VII. Location of Attribution (LOA)
VIII. Transfer rule

Definitions:

Urinary tract infections (UTI) are defined using Symptomatic Urinary Tract Infection (SUTI) criteria, Asymptomatic Bacteremic UTI (ABUTI), and Urinary System Infection (USI) criteria. (See Table 1 and 2 and Figure2).

Note: UTI is a primary site of infection and cannot be considered secondary to another site of infection.

Indwelling catheter: A drainage tube that is inserted into the urinary bladder through the urethra, is left in place, and is connected to a drainage bag (including leg bags). These devices are also called Foley catheters. Condom or straight in-and-out catheters are not included nor are nephrostomy tubes, ileoconduits, or suprapubic catheters unless a Foley catheter is also present. Indwelling urethral catheters that are used for intermittent or continuous irrigation are included in CAUTI surveillance.

Catheter-associated UTI (CAUTI): A UTI where an indwelling urinary catheter (IUC) was in place for >2 calendar days on the date of event, with day of device placement being Day 1*, AND an indwelling urinary catheter was in place on the date of event or the day before. If an indwelling urinary catheter was in place for > 2 calendar days and then removed, the date of event for the UTI must be the day of discontinuation or the next day for the UTI to be catheter-associated.

*If the IUC was in place prior to inpatient admission, the catheter day count that determines device –association begins with the admission date to the first inpatient location. This allows for consistency with device denominator count (see Table 3 Denominator Data Collection Methods)
Example of Associating Catheter Use to UTI:

A patient in an inpatient unit has a Foley catheter inserted and the following day is the date of event for a UTI. Because the catheter has not been in place >2 calendar days on the date of event, this is not a CAUTI. However, depending on the date of admission, this may be a healthcare-associated UTI. Please refer to SUTI 1b: Non-CAUTI.

Notes:
- SUTI 1b and USI cannot be catheter-associated.
- SUTI 1b cannot be met in a patient > 65 years of age with fever >38°C as the only element within the Infection Window Period

Indwelling urinary catheters that are removed and reinserted: If, after indwelling urinary catheter removal, the patient is without an indwelling urinary catheter for at least 1 full calendar day (NOT to be read as 24 hours), then the urinary catheter day count will start anew. If instead, a new indwelling urinary catheter is inserted before a full calendar day has passed, the urinary catheter device day count, to determine eligibility for a CAUTI, will continue uninterrupted.

Figure 1: Associating Catheter Use to UTI

<table>
<thead>
<tr>
<th></th>
<th>March 31 (Hospital day 3)</th>
<th>April 1</th>
<th>April 2</th>
<th>April 3</th>
<th>April 4</th>
<th>April 5</th>
<th>April 6</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patient A</td>
<td>Foley Day 3</td>
<td>Foley</td>
<td>Foley removed (Foley Day 5)</td>
<td>Foley replaced (Foley Day 6)</td>
<td>Foley Day 7</td>
<td>Foley removed Day 8</td>
<td>No Foley</td>
</tr>
<tr>
<td>Patient B</td>
<td>Foley Day 3</td>
<td>Foley</td>
<td>Foley removed (Foley Day 5)</td>
<td>No Foley</td>
<td>Foley replaced (Foley Day 1)</td>
<td>Foley Day 2</td>
<td>Foley Day 3</td>
</tr>
</tbody>
</table>

Rationale: NHSN surveillance for infection is not aimed at a specific device. Instead surveillance is aimed at identifying risk to the patient that is the result of device use in general.

Notes:
- In the examples above, Patient A is eligible for a CAUTI beginning on March 31, through April 6th, since a Foley was in place for some portion of each calendar day until April 6th. A UTI with date of event on April 6th would be a CAUTI since the catheter had been in place greater than 2 days and was removed the day before the date of event.
• Patient B is eligible for a CAUTI on March 31 (Foley Day 3) through April 3. The catheter had been in place > 2 days and an HAI occurring on the day of device discontinuation or the following calendar day is considered a device-associated infection.
• If the patient did not have a CAUTI by April 3, the patient is not eligible for a CAUTI until April 6, when the second indwelling urinary catheter had been in place for greater than 2 days.
Table 1. Urinary Tract Infection Criteria

<table>
<thead>
<tr>
<th>Criterion</th>
<th>Urinary Tract Infection (UTI)</th>
</tr>
</thead>
<tbody>
<tr>
<td><strong>Symptomatic UTI (SUTI)</strong></td>
<td>Must meet at least one of the following criteria:</td>
</tr>
<tr>
<td>Patient must meet 1, 2, and 3 below:</td>
<td></td>
</tr>
<tr>
<td>1. Patient had an indwelling urinary catheter that had been in place for &gt; 2 days on the <strong>date of event</strong> AND was either:</td>
<td></td>
</tr>
<tr>
<td>• Present for any portion of the calendar day on the date of event†,</td>
<td></td>
</tr>
<tr>
<td>OR</td>
<td></td>
</tr>
<tr>
<td>• Removed the day before the date of event‡</td>
<td></td>
</tr>
<tr>
<td>2. Patient has at least one of the following signs or symptoms:</td>
<td></td>
</tr>
<tr>
<td>• fever (&gt;38.0°C): To use fever in a patient &gt; 65 years of age, the indwelling urinary catheter needs to be in place &gt; 2 calendar days on <strong>date of event</strong>.</td>
<td></td>
</tr>
<tr>
<td>• suprapubic tenderness*</td>
<td></td>
</tr>
<tr>
<td>• costovertebral angle pain or tenderness*</td>
<td></td>
</tr>
<tr>
<td>• urinary urgency ^</td>
<td></td>
</tr>
<tr>
<td>• urinary frequency ^</td>
<td></td>
</tr>
<tr>
<td>• dysuria ^</td>
<td></td>
</tr>
<tr>
<td>3. Patient has a urine culture with no more than two species of organisms identified, at least one of which is a bacterium of ≥10^5 CFU/ml (See <strong>Comments</strong>). All elements of the SUTI criterion must occur during the IWP (See IWP Definition <strong>Chapter 2 Identifying HAIs in NHSN</strong>).</td>
<td></td>
</tr>
</tbody>
</table>

† When entering event into NHSN choose “INPLACE” for Risk Factor for Urinary Catheter
‡ When entering event into NHSN choose “REMOVE” for Risk Factor for Urinary Catheter
*With no other recognized cause (see **Comments**)
^ These symptoms cannot be used when catheter is in place. An indwelling urinary catheter in place could cause patient complaints of “frequency” “urgency” or “dysuria”.

**Note:**
• Fever is a non-specific symptom of infection and cannot be excluded from UTI determination because it is clinically deemed due to another recognized cause.
SUTI 1b  
Non-Catheter-associated Urinary Tract Infection (Non-CAUTI) in any age patient

Patient must meet 1, 2, and 3 below:

1. One of the following is true:
   - Patient has/had an indwelling urinary catheter but it has/had not been in place >2 calendar days on the date of event†
   - OR
   - Patient did not have a urinary catheter in place on the date of event nor the day before the date of event †

2. Patient has at least one of the following signs or symptoms:
   - fever (>38°C) in a patient that is ≤ 65 years of age
   - suprapubic tenderness*
   - costovertebral angle pain or tenderness *
   - urinary frequency ^
   - urinary urgency ^
   - dysuria ^

3. Patient has a urine culture with no more than two species of organisms identified, at least one of which is a bacterium of ≥10⁵ CFU/ml. (See Comments) All elements of the SUTI criterion must occur during the IWP (See IWP Definition Chapter 2 Identifying HAIs in NHSN).

† When entering event into NHSN choose “NEITHER” for Risk Factor for Urinary Catheter
*With no other recognized cause (see Comments)
^These symptoms cannot be used when catheter is in place. An indwelling urinary catheter in place could cause patient complaints of “frequency” “urgency” or “dysuria”.

Note:
- Fever is a non-specific symptom of infection and cannot be excluded from UTI determination because it is clinically deemed due to another recognized cause.
Patient must meet 1, 2, and 3 below:

1. Patient is ≤1 year of age (with‡ or without an indwelling urinary catheter)

2. Patient has at least one of the following signs or symptoms:
   • fever (>38.0°C)
   • hypothermia (<36.0°C)
   • apnea*
   • bradycardia*
   • lethargy*
   • vomiting*
   • suprapubic tenderness*

3. Patient has a urine culture with no more than two species of organisms identified, at least one of which is a bacterium of \( \geq 10^5 \) CFU/ml. (See Comments) All elements of the SUTI criterion must occur during the IWP (See IWP Definition Chapter 2 Identifying HAIs in NHSN).

‡ If patient had an indwelling urinary catheter in place for >2 calendar days, and catheter was in place on the date of event or the previous day the CAUTI criterion is met. If no such indwelling urinary catheter was in place, UTI (non-catheter associated) criterion is met.

*With no other recognized cause (See Comments)

Note: Fever and hypothermia are non-specific symptoms of infection and cannot be excluded from UTI determination because they are clinically deemed due to another recognized cause.
“Mixed flora” is not available in the pathogen list within NSHN. Therefore, it cannot be reported as a pathogen to meet the NHSN UTI criteria. Additionally, “mixed flora” represent at least two species of organisms. Therefore, an additional organism recovered from the same culture would represent >2 species of microorganisms. Such a specimen also cannot be used to meet the UTI criteria.

The following excluded organisms cannot be used to meet the UTI definition:

- *Candida* species or yeast not otherwise specified
- mold
- dimorphic fungi or
- parasites

An acceptable urine specimen may include these organisms as long as one bacterium of greater than or equal to 100,000 CFU/ml is also present. Additionally, these non-bacterial organisms identified from blood cannot be deemed secondary to a UTI since they are excluded as organisms in the UTI definition.

- Suprapubic tenderness whether elicited by palpation (tenderness-sign) or provided as a subjective complaint of suprapubic pain (pain-symptom), documentation of either found in the medical record is acceptable as a part of SUTI criterion if documented in the medical record during the Infection Window Period.

- Lower abdominal pain or bladder or pelvic discomfort are examples of symptoms that can be used as suprapubic tenderness. Generalized “abdominal pain” in the medical record is not to be interpreted as suprapubic tenderness as there are many causes of abdominal pain and this symptom is too general.

- Left or right lower back or flank pain are examples of symptoms that can be used as costovertebral angle pain or tenderness. Generalized "low back pain" is not to be interpreted as costovertebral pain.
Table 2. Urinary System Infection Criteria

<table>
<thead>
<tr>
<th><strong>Catheter – associated ABUTI or Non-catheter associated ABUTI in any age patient</strong></th>
<th><strong>Patient must meet 1, 2, and 3 below:</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>1. Patient with* or without an indwelling urinary catheter has no signs or symptoms of SUTI 1 or 2 according to age (Note: Patients &gt; 65 years of age with a non-catheter-associated ABUTI may have a fever and still meet the ABUTI criterion)</td>
</tr>
<tr>
<td></td>
<td>2. Patient has a urine culture with no more than two species of organisms identified, at least one of which is a bacterium of ( \geq 10^5 ) CFU/ml (see Comment section below)</td>
</tr>
<tr>
<td></td>
<td>3. Patient has organism identified** from blood specimen with at least one matching bacterium to the bacterium identified in the urine specimen, OR meets LCBI criterion 2 (without fever) and matching common commensal(s) in the urine. All elements of the ABUTI criterion must occur during the IWP (See IWP Definition Chapter 2 Identifying HAIs in NHSN).</td>
</tr>
</tbody>
</table>

*Patient had an indwelling urinary catheter in place for >2 calendar days on the date of event, and the indwelling urinary catheter was in place on the date of event or the day before. Catheter - associated ABUTI is reportable if the location is in the facility’s reporting plan. |

**Organisms identified by a culture or non-culture based microbiologic testing method which is performed for purposes of clinical diagnosis or treatment (for example, not Active Surveillance Culture/Testing (ASC/AST). |

**Comments**

A urine specimen with “Mixed flora” cannot be used to meet the urine criterion. Additionally, the following excluded organisms cannot be used to meet the UTI definition:
- *Candida* species or yeast not otherwise specified
- mold
- dimorphic fungi or
- parasites

An acceptable urine specimen may include these excluded organisms as long as one bacterium of greater than or equal to 100,000 CFU/ml is also present. Additionally, these non-bacterial organisms identified from blood cannot be deemed secondary to a UTI since they are excluded as organisms in the UTI definition.
<table>
<thead>
<tr>
<th>Criterion</th>
<th>Urinary System Infection (USI) (kidney, ureter, bladder, urethra, or perinephric space)</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td>Other infections of the urinary system must meet at least one of the following criteria:</td>
</tr>
<tr>
<td>1.</td>
<td>Patient has organisms identified** from fluid (excluding urine) or tissue from affected site</td>
</tr>
<tr>
<td>2.</td>
<td>Patient has an abscess or other evidence of infection on gross anatomical exam, during invasive procedure, or on histopathologic exam</td>
</tr>
<tr>
<td>3.</td>
<td>Patient has at least one of the following signs or symptoms:</td>
</tr>
<tr>
<td></td>
<td>• fever (&gt;38.0°C)</td>
</tr>
<tr>
<td></td>
<td>• localized pain or tenderness*</td>
</tr>
<tr>
<td>And at least one of the following:</td>
<td></td>
</tr>
<tr>
<td>a)</td>
<td>purulent drainage from affected site</td>
</tr>
<tr>
<td>b)</td>
<td>organisms identified** from blood and imaging test evidence of infection (e.g., ultrasound, CT scan, magnetic resonance imaging [MRI], or radiolabel scan [gallium, technetium]) which if equivocal is supported by clinical correlation (i.e., physician documentation of antimicrobial treatment for urinary system infection).</td>
</tr>
<tr>
<td>4.</td>
<td>Patient ≤1 year of age has at least one of the following signs or symptoms:</td>
</tr>
<tr>
<td></td>
<td>• fever (&gt;38.0°C)</td>
</tr>
<tr>
<td></td>
<td>• hypothermia (&lt;36.0°C)</td>
</tr>
<tr>
<td></td>
<td>• apnea*</td>
</tr>
<tr>
<td></td>
<td>• bradycardia*</td>
</tr>
<tr>
<td></td>
<td>• lethargy*</td>
</tr>
<tr>
<td></td>
<td>• vomiting*</td>
</tr>
<tr>
<td>And at least one of the following:</td>
<td></td>
</tr>
<tr>
<td>a)</td>
<td>purulent drainage from affected site</td>
</tr>
<tr>
<td>b)</td>
<td>organisms identified** from blood and imaging test evidence of infection, (for example, ultrasound, CT scans, magnetic resonance imaging [MRI], or radiolabel scan [gallium, technetium])</td>
</tr>
</tbody>
</table>
* With no other recognized cause

** Organisms identified by a culture or non-culture based microbiologic testing method which is performed for purposes of clinical diagnosis or treatment (e.g., not Active Surveillance Culture/Testing (ASC/AST)).

Notes:

- Fever and hypothermia are non-specific symptoms of infection and cannot be excluded from USI determination because they are clinically deemed due to another recognized cause.
- All elements of the USI criterion must occur during the IWP (See IWP Definition Chapter 2 Identifying HAIs in NHSN).

<table>
<thead>
<tr>
<th>Comments</th>
</tr>
</thead>
<tbody>
<tr>
<td>• Report infections following circumcision in newborns as SST-CIRC.</td>
</tr>
<tr>
<td>• If patient meets USI criteria and they also meet UTI criteria, report UTI only, unless the USI is a surgical site organ/space infection, in which case, only USI should be reported.</td>
</tr>
<tr>
<td>• For NHSN reporting purposes, Urinary System Infection (USI) cannot be catheter associated, therefore, USI will only present as specific event type if urinary catheter status is marked “Neither”.</td>
</tr>
</tbody>
</table>
Figure 2: Identifying SUTI and ABUTI Flowchart

Identifying Symptomatic Urinary Tract Infections (SUTI) & Asymptomatic Bacteremic Urinary Tract Infections (ABUTI)

Positive urine culture with no more than 2 species of organisms, at least one of which is a bacterium of ≥10⁵ CFU/ml. All elements of the UTI criterion must occur during the Infection Window Period (Note: if none of the organisms present at ≥10⁵ cfu/ml are bacteria, answer = No)

No

Had an indwelling urinary catheter that had been in place for > 2 days, AND was either:
1. Still present for any portion of the calendar day on date of event
2. Removed day before date of event?

No

Yes

Does not meet UTI criteria

At least one of the following signs or symptoms?
• suprapubic tenderness⁶
• costovertebral angle pain⁷
• urgency⁸
• frequency⁹
• dysuria⁸
• fever (≥ 38.0°C)—In a patient that is ≤ 65 years of age
* With no other recognized cause
^ These symptoms cannot be used when catheter is in place.

No

Yes

At least one of the following signs or symptoms?
• Any age patient: fever (≥ 38.0°C), suprapubic tenderness⁶, costovertebral angle pain⁷, urgency⁸, dysuria⁸, frequency⁹
• Patients ≤ 1 year of age: fever (≥ 38.0°C), hypothermia (≤36.0°C), suprapubic tenderness⁶, costovertebral angle pain⁷, spines⁸, bradycardia⁸, lethargy¹⁰, or vomiting¹⁰
* With no other recognized cause
^ These symptoms cannot be used when catheter is in place.

Yes

No

Organism identified⁶ from blood specimen with at least one matching bacterium to bacterium in the urine at ≥100,000 cfu/ml?

Yes

No

Meets criteria for non-catheter associated SUTI

Organism identified from blood specimen with at least one matching bacterium to bacterium in the urine at ≥100,000 cfu/ml?

Yes

No

Meets criteria for non-catheter associated ABUTI

Does not meet UTI criteria

Meets criteria for catheter-associated SUTI (CAUTI)

Meets criteria for catheter-associated ABUTI (CAUTI)

Does not meet UTI criteria
Monthly Summary Data

**Numerator Data:** The *Urinary Tract Infection (UTI)* form (CDC 57.114) is used to collect and report each CAUTI that is identified during the month selected for surveillance. The *Instructions for Completion of Urinary Tract Infection form* include brief instructions for collection and entry of each data element on the form. USIs are never included in CAUTI data and are reported separately on the *HAI Custom Event Form*. The UTI form includes patient demographic information and information on whether or not an indwelling urinary catheter was present. Additional data include the specific criteria met for identifying the UTI, whether the patient developed a secondary bloodstream infection, whether the patient died, and the organisms isolated from cultures and their antimicrobial susceptibilities.

**Reporting Instructions:**
If no CAUTIs are identified during the month of surveillance, the “Report No Events” box must be checked on the appropriate denominator summary screen, (for example, *Denominators for Intensive Care Unit (ICU)/Other Locations (Not NICU or SCA/ONC)*).

**Denominator Data:** Device days and patient days are used for denominators (See *Key Terms* chapter). The method of collecting device-day denominator data may differ depending on the location of patients being monitored. The following methods may be used:

Table 3: Denominator Data Collection Methods

<table>
<thead>
<tr>
<th>Denominator Data Collection Method</th>
<th>Details</th>
</tr>
</thead>
<tbody>
<tr>
<td>Manual, Daily</td>
<td>Denominator data (patient days and device days) should be collected at the same time, every day, for each location performing surveillance to ensure that differing collection methods don’t inadvertently result in device days being &gt; patient days.</td>
</tr>
<tr>
<td>(specifically, collected at the same time <em>every day</em> of the month)</td>
<td>The <em>Instructions for Completion of Denominators for Intensive Care Unit (ICU)/Other Locations (Not NICU and SCA/ONC)</em> and <em>Instructions for Completion of Denominators for Specialty Care Areas (SCA)/Oncology (ONC)</em> contain brief instructions for collection and entry of each data element on the form.</td>
</tr>
<tr>
<td>Indwelling urinary catheter days, which are the number of patients with an indwelling urinary catheter device, are collected daily, at the same time each day, according to the chosen location using the appropriate form (CDC 57.117 and 57.118). These daily counts are summed and only the total for the month is entered into NHSN. Indwelling urinary catheter days and patient days are collected separately for each of the locations monitored.</td>
<td></td>
</tr>
</tbody>
</table>
### Denominator Data Collection Method

**Manual, sampled once/week** (collected at the same time on the same designated day, *once per week*)

To reduce staff time spent collecting surveillance data, once weekly sampling of denominator data to generate estimated urinary catheter days may be used as an alternative to daily collection in non-oncology ICUs and wards (see Notes below). Sampling may not be used in SCA/ONC locations or NICUs. During the month, the number of patients in the location (patient-days) and the number of patients with an indwelling urinary catheter (urinary catheter-days) is collected on a designated day each week (for example, every Tuesday), at the same time during the month.

Evaluations of this method have repeatedly shown that use of Saturday or Sunday generate the least accurate estimates of denominator data, and, therefore, these days should not be selected as the designated day.\(^7,9\) If the day designated for the collection of sampled data is missed, collect the data on the next available day instead.

The following must be collected and entered into NHSN:

1. The monthly total for patient-days, based on collection daily
2. The sampled total for patient-days
3. The sampled total urinary catheter-days

When these data are entered, the NHSN application will calculate an estimate of urinary catheter-days.

### Notes:

- To ensure the accuracy of estimated denominator data obtained by sampling, only ICU and ward location types with an average of 75 or more urinary catheter-days per month are eligible to use this method. A review of each location’s urinary catheter denominator data for the past 12 months in NHSN will help determine which locations are eligible.
- The accuracy of estimated denominator data generated by sampling can be heavily influenced by incorrect or missing data. Careful implementation of data collection following the guidance in this protocol is essential to avoid erroneous fluctuations in rates or Standardized Infection Ratios (SIRs).
**Device-associated Module**

**UTI**

<table>
<thead>
<tr>
<th>Denominator Data Collection Method</th>
<th>Details</th>
</tr>
</thead>
<tbody>
<tr>
<td>Electronic</td>
<td>For <em>any</em> location, denominator data from electronic sources (for example, urinary catheter days from electronic charting), may be used after validation of a minimum 3 months proves the data to be within 5% (±) of the manually-collected, once a day counts. Perform the validation of electronic counts separately for each location conducting CAUTI surveillance.</td>
</tr>
</tbody>
</table>

**Data Analyses:** The Standardized Infection Ratio (SIR) is calculated by dividing the number of observed infections by the number of predicted infections. The number of predicted infections is calculated using probabilities from negative binomial regression models constructed from 2015 NHSN data, which represents a standard population.

**Notes:**
The SIR will be calculated only if the number of predicted CAUTIs (numPred) is ≥1 to help enforce a minimum precision criterion.

While the CAUTI SIR can be calculated for single locations, the measure also allows you to summarize your data by multiple locations, adjusting for differences in the incidence of infection among the location types. For example, you will be able to obtain one CAUTI SIR adjusting for all locations reported. Similarly, you can obtain one CAUTI SIR for all ICUs in your facility.

The SUR, or Standardized Utilization Ratio, is a risk-adjusted summary measure for device use. Similar to the SIR, the SUR can be calculated for single locations as well as be summarized across multiple locations.

The CAUTI rate per 1000 urinary catheter days is calculated by dividing the number of CAUTIs by the number of catheter days and multiplying the result by 1000. The Urinary Catheter Utilization Ratio is calculated by dividing the number of urinary catheter days by the number of patient days. These calculations will be performed separately for the different types of ICUs, specialty care areas, and other locations in the institution, except for neonatal locations.

Descriptive analysis output options of numerator and denominator data, such as line listings, frequency tables, and bar and pie charts are available in the NHSN application. SIRs, SURs, and CAUTI rates and run charts are also available. Guides on using NHSN analysis features are available at: [www.cdc.gov/nhsn/PS-Analysis-resources/reference-guides.html](http://www.cdc.gov/nhsn/PS-Analysis-resources/reference-guides.html).

Table 3. CAUTI Measures Available in NHSN
<table>
<thead>
<tr>
<th>Measure</th>
<th>Calculation</th>
<th>Application</th>
</tr>
</thead>
<tbody>
<tr>
<td>CAUTI SIR</td>
<td>The number of Observed CAUTIs&lt;br&gt;The number of Predicted CAUTIs</td>
<td>Both location specific and summarized measure</td>
</tr>
<tr>
<td>CAUTI Rates</td>
<td>The number of CAUTIs for a location x 1000&lt;br&gt;The number of Urinary Catheter Days for a location</td>
<td>Location specific measure only</td>
</tr>
<tr>
<td>Urinary Catheter SUR</td>
<td>The number of Observed Urinary Catheter Days&lt;br&gt;The number of Predicted Urinary Catheter Days</td>
<td>Both location specific and summarized measure</td>
</tr>
<tr>
<td>DUR</td>
<td>The Urinary Catheter Days for a location&lt;br&gt;The Patient Days for that location</td>
<td>Location specific measure only</td>
</tr>
</tbody>
</table>
REFERENCES


