

ASID (HICSIG)/AICA Position Statement: Preventing catheter-associated urinary tract infections in patients

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Abstract. Catheter-associated urinary tract infections (CAUTIs) occur frequently in healthcare settings. The insertion and maintenance of indwelling urinary catheters is a routine element of healthcare. In order to prevent CAUTI, it is important that healthcare professionals providing catheter care understand the indications for catheter use and the correct procedure for insertion and maintenance of catheters. This paper reviews and summarises three recent key publications on the prevention of CAUTIs and proposes the use of a care bundle and checklist for catheter indications, insertion and maintenance, and quality improvement.

Process of position statement development

The Tasmanian Infection Prevention and Control Unit (TIPCU) examined and synthesised three catheter-associated urinary tract infections (CAUTI) reviews in the current literature. From this process a checklist/care bundle was developed. Substantial input was then received from the Australasian Society for Infectious Disease (ASID) members, including members of the ASID Healthcare Infection Control Special Interest Group (HICSIG), and the Australian Infection Control Association (AICA). The authors responded to all comments received.

Background

Catheter-associated urinary tract infections (CAUTIs) are common in healthcare settings. Around 20% of all healthcare-

associated infections are urinary tract infections,¹ and most of these are associated with some form of instrumentation of the urinary tract.²

Between 15% and 25% of hospitalised patients receive a short-term indwelling urinary catheter.^{3–6} The duration of catheterisation is the most important risk factor for development of infection,⁷ while additional risk factors include female sex, older age and not maintaining a closed drainage system.⁸

The limitations and heterogeneity of definitions used in various studies present major challenges to defining clinical and surveillance definitions of CAUTI.² The three articles reviewed in this paper do not provide definitions, but rather defer to National Healthcare Safety Network⁹ (NHSN) definitions, or in the case of Tenke *et al.* (2009) definitions are not discussed. NHSN definitions were recently revised and now provide additional clarity and uniformity in the

distinguishing of asymptomatic and symptomatic CAUTI and the time period for follow-up surveillance following catheter removal.²

Most microorganisms causing CAUTI derive from the patient’s own colonic and perineal flora or from the hands of healthcare workers.^{10,11} Bacteria can enter the urinary tract at the time of catheter insertion via the catheter tip or subsequently once the patient is catheterised via the extraluminal or intraluminal route. Common inoculation source-sites include the catheter tip, when the perineum and distal urethra are inadequately cleaned;¹² the taps of the drainage bag which afford bacteria access to the catheter and eventually the bladder;⁷ and residual urine remaining in the collection bag, which acts as a focus for organisms to spread via the intraluminal route.¹² While infections may arise via the intraluminal route when a closed system is not maintained or the collection bag is contaminated, it is believed that the majority of infections arise from extraluminal (via the outside of the catheter) bacterial migration.¹⁰

To reduce the risk of CAUTI, there is a need to develop and implement a range of prevention measures. Over the past 2 years, three major publications have reviewed the literature and developed recommendations on the prevention of CAUTIs, namely:

- European and Asian Guidelines on management and prevention of catheter-associated urinary tract infections;⁷

- Strategies to Prevent Catheter-Associated Urinary Tract Infections in Acute Care Hospitals Practice Recommendation, Society for Healthcare Epidemiology of America (SHEA) and Infectious Diseases Society of America;⁸
- Guidelines for Prevention of Catheter-Associated Urinary Tract Infections 2010, Healthcare Infection Control Practices Advisory Committee (HICPAC).²

The purpose of this paper is to propose a clear approach to the prevention of CAUTIs for adoption across Australia applicable to all healthcare settings including residential and aged care. The approach described is consistent with an Australian context, using recommendations from the publications described above and the NHMRC Australian Guidelines for the Prevention and Control of Infection in Healthcare.¹³ A suggested checklist has also been developed for the insertion of indwelling urinary catheters.

Recommendations

Practices to minimise the risk of CAUTIs can be summarised into three distinct areas: insertion, maintenance and quality improvement. This paper provides recommendations supported by the literature in each of these areas. A summary of key recommendations, including their grading, are detailed in Tables 1 and 2.

Table 1. Summary of key graded recommendations and the level of evidence supporting them as used in the major CAUTI review papers

Recommendation	EPIC (2007) ²⁴		HICPAC (2010) ²		European (2008) ⁷		SHEA/IDSA (2009) ⁸	
	Y/N	Grade	Y/N	Grade	Y/N	Grade	Y/N	Grade
Catheter indications								
Evaluate necessity	Y	Class D	Y	1A	Y	x	Y	A
Alternative methods	ND		Y	1B	Y	x	Y	AI
Insertion								
Documentation of insertion	Y	Class D	Y	II	Y	B	Y	AII
Trained staff	Y	Class D	Y	1C	ND	x	Y	BIII
Train patients/family	Y	Class D	Y	1C	ND	x	Y	AIII
Hand hygiene	ND (implicit)	x	Y	1B	Y	A	Y	A
Select catheter material	ND	x	ND	x	ND (directly)	x	ND	x
Smallest gauge	Y	Class D	Y	II	Y	B	Y	BIII
Aseptic/sterile technique	Y	Class D	Y	1C	Y	B	Y	AIII
Barrier precautions for insertion	ND	x	Y	1C	ND	x	Y	AIII
Antiseptic cleaning of meatus	N	Class D	N	1B	N	A	N	AI
Review ongoing need/time minimal	Y	Class D	Y	1B	Y	A	Y	A
Maintenance								
Use closed system	Y	Class A	Y	1B	Y	A	Y	AI
Urine samples aseptically	Y	Class D	Y	1C	ND	x	Y	AIII
Replace if break in asepsis	ND	x	ND	x	ND	x	ND	x
Do not change catheter routinely	Y	Class D	Y	1B	Y	B	Y	AII
Perform routine meatus care	Y	Class A	Y	1B	ND	x	Y	AI
Avoid irrigation	Y	Class A	Y	1B	ND	x	Y	AI
Cohort patients	ND		Y	II	ND	x	ND	x
Quality improvement								
Compliance with training	ND	x	ND	x	ND	x	ND	x
Compliance with control	ND	x	Y as below	x	Y as below	x	Y as below	x
Compliance with removal	ND	x	Y as below	x	Y as below	x	Y as below	x
Monitor rates	ND	x	Y	II	ND		Y	AII

ND, not discussed.

Table 2. Summary of grading systems used

IDSA/SHEA Practice Recommendations**Strength**

- A – Good evidence to support a recommendation for use.
- B – Moderate evidence to support a recommendation for use.
- C – Poor evidence to support a recommendation.

Quality

- I – Evidence from >1 properly randomised, controlled trial.
- II – Evidence from >1 well designed clinical trial, without randomisation; from cohort or case-control analytic studies (preferably from >1 centre), from multiple time series; or from dramatic results from uncontrolled experiments.
- III – Evidence from opinions of respected authorities, based on clinical experience, descriptive studies, or reports of expert committees.

HICPAC Guidelines

Category IA – Strongly recommended for implementation and supported by well-designed experimental, clinical or epidemiologic studies.

Category IB – Strongly recommended for implementation and supported by certain experimental, clinical or epidemiologic studies and a strong theoretical rationale.

Category IC – Required by state or federal regulation or representative of an established standard for which data are not available.

Category II – Suggested for implementation and supported by suggestive clinical or epidemiologic studies or a theoretical rationale.

No recommendation – Unresolved issue; practices for which insufficient evidence or no consensus regarding efficacy exist.

European & Asian Guideline**Level of evidence**

- Ia – Evidence obtained from meta-analysis of randomised trials.
- Ib – Evidence obtained from at least one randomised trial.
- Ila – Evidence obtained from one well-designed, controlled study without randomisation.
- Ilb – Evidence obtained from at least one other type of well-designed, quasi-experimental study.
- III – Evidence obtained from well-designed, non-experimental studies such as comparative studies, correlation studies and case reports.
- IV – Evidence obtained from expert committee reports or opinions or clinical experience of respected authorities.

Grade of guideline recommendation

- A – Based on clinical studies of good quality and consistency addressing the special recommendations and including at least one randomised trial.
- B – Based on well-conducted clinical studies, but without randomised clinical trials.
- C – Made despite the absence of directly applicable clinical studies of good quality.

EPIC 2 Guidelines**Level of evidence**

- 1++ High-quality meta-analyses, systematic reviews of randomised controlled trials (RCT), or RCT with a very low risk of bias.
- 1+ Well-conducted meta-analyses, systematic reviews of RCT, or RCT with a low risk of bias.
- 1 – Meta-analyses, systematic reviews of RCT, or RCT with a high risk of bias.
- 2++ High-quality systematic reviews of case-control or cohort studies. High-quality case-control or cohort studies with a very low risk of confounding, bias or chance and a high probability that the relationship is causal.
- 2+ Well conducted case-control or cohort studies with a low risk of confounding, bias or chance and a moderate probability that the relationship is causal.
- 2 – Case-control or cohort studies with a high risk of confounding bias, or chance and a significant risk that the relationship is not causal.
- 3 – Non-analytic studies (for example, case reports, case series).
- 4 – Expert opinion, formal consensus.

Class A – Systematic review of RCT or a body of evidence that consists principally of studies rated as 1+, is directly applicable to the target population and demonstrates overall consistency of results. Evidence drawn from a National Institute of Clinical Excellence technology appraisal.

B – A body of evidence that includes studies rated as 2++, is directly applicable to the target population and demonstrates overall consistency of results, or Extrapolated evidence from studies rated as 1++ or 1+.

C – A body of evidence that includes studies rated as 2+, is directly applicable to the target population and demonstrates overall consistency of results, or Extrapolated evidence from studies rated as 2++.

D – Evidence level 3 or 4, or Extrapolated evidence from studies rated as 2+, or Formal consensus. D (GPP). A good practice point (GPP) is a recommendation.

Insertion

The first and foremost goal of CAUTI prevention is a reduction in the overall use of indwelling catheters, especially in those at high risk of complications from CAUTIs. Those at high risk include women, the elderly and patients with impaired immunity, severe underlying illness, diabetes, renal dysfunction or incontinence.² Wherever possible, alternatives, such as suprapubic catheters, condom drainage systems or intermittent catheterisation should always be

considered where appropriate.^{2,7} To assist in this process, there is a need to establish clear indications for the use of indwelling catheters, as summarised in Table 3. Low quality evidence suggested a benefit of alternative catheterisation methods in selected population groups.²

When inserting a urethral catheter, the correct procedure must be followed to minimise the risk of introducing bacteria into the bladder. Recommendations relating to the insertion of indwelling catheters are summarised as follows.

Table 3. Indications for indwelling catheter use

a) Patient has acute urinary retention or obstruction
b) Urinary output monitoring in critically ill patients
c) Peri-operative use for selective surgical procedures [†] <ul style="list-style-type: none"> • Urological surgery or surgery involving other contiguous structures of genitourinary tract • Prolonged duration of surgery (removed in recovery) • Large volumes of infusions or diuretics administered intra-operatively • Operative patients with urinary incontinence • Need for intraoperative monitoring of output
d) Healing of wounds (sacral/perianal) in incontinent patients (as part of a holistic plan)
Patient
e) Patient requiring prolonged immobilisation (e.g. potentially unstable thoracic or lumbar spine, multiple traumatic injuries such as pelvic fractures) [‡]
f) Exceptional circumstances, e.g. comfort at end of life

[†]In these settings the catheter should be removed within 24 h post-operatively where possible.

[‡]Avoid using indwelling urinary catheters for the management of incontinence. This includes residential aged care residents.

- Provide and follow written guidelines for catheter use, insertion and maintenance.⁸
- Consider the use of a portable ultrasound device to assess urine volume in patients undergoing intermittent catheterisation.^{2,8}
- Implement a system for documenting the following information in the patient record: indications for catheter insertion, date and time of catheter insertion, details of the individual who inserted catheter, and date and time of catheter removal.⁸
- Ensure that only trained, competent personnel insert urinary catheters.⁸ This may include hospital personnel, family members, or patients themselves who know the correct technique of aseptic catheter insertion and maintenance.²
- Perform hand hygiene immediately before insertion of the catheter.^{2,8}
- Clean (i.e. non-sterile) equipment is appropriate for chronic intermittent catheterisation in a non-acute care setting.^{7,14,15}
- Insert an indwelling catheter using an aseptic non-touch technique, using sterile equipment. Equipment required includes: sterile gloves, a drape, sterile swabs, a sterile solution for cleaning the urethral meatus, and a single-use packet of sterile lubricant jelly for insertion.^{2,8,13} Antiseptic lubricants need not be used routinely to prevent CAUTI.^{13,14} While there is very low-quality evidence suggesting a reduced risk of CAUTI from using lubricants during catheter insertion, several studies have found no significant difference between antiseptic lubricants and non-antiseptic lubricants in preventing CAUTI.²
- Use the smallest catheter as possible, consistent with proper drainage, to minimise urethral trauma.^{2, 8}
- Do not use antibiotic-impregnated catheters routinely as there is no evidence that they decrease symptomatic infection.⁷
- Do not use systemic antimicrobials routinely as prophylaxis in patients requiring either short or long-term catheterisation unless indications exist.^{2,8} There are studies suggesting the incidence of catheter-associated bacteruria may be reduced by antimicrobial prophylaxis;^{11,16} however, this protective effect is transient (lasts only a few days) and is

associated with the selection of resistant organisms. Prophylaxis is not indicated for patients at low risk for acquired bacteruria and in whom the sequelae of catheter-associated infections are infrequent.¹⁶

- Urinary catheter systems with pre-connected, sealed catheter-tubing junctions may reduce the risk of CAUTI compared with unsealed catheter systems.²

Maintenance

The following suite of strategies relating to the maintenance of a catheter will aid in the reduction of infection risk.

- Properly secure indwelling catheters after insertion to prevent movement and urethral traction.^{2,8}
- Perform hand hygiene before and after any manipulation of the catheter site or apparatus,^{2,8} consistent with the 'five moments' of hand hygiene.¹⁷
- Use standard precautions as appropriate during any manipulation of the catheter or collecting system.²
- The catheter must be maintained as a sterile, continuously closed drainage system, with unobstructed urine flow.^{2,7,8}
- Do not disconnect the catheter and drainage tube unless the catheter has to be irrigated.
- Keep the collecting bag below the level of the bladder at all times, but do not rest the bag on the floor.^{2,8, 13}
- If breaks in aseptic non-touch technique, disconnection, or leakage occur, replace the catheter and collecting system using aseptic non-touch technique and sterile equipment.^{2,8}
- Avoid catheter irrigation if possible, and do not perform continuous irrigation of the bladder with antimicrobials as a routine infection prevention measure. If obstruction is anticipated, closed continuous irrigation may be used to prevent it. To relieve obstruction due to clots, mucus or other causes, an intermittent method of irrigation may be used.⁸
- The use of topical antiseptics or antibiotics applied to the catheter, urethra or meatus is not recommended;⁷ routine hygiene is adequate.¹³
- Chronic antibiotic suppressive therapy is generally not recommended.⁷

- For examination of fresh urine, a small sample should be collected by aspirating urine from the sampling port with a sterile needle and syringe after cleansing the port with disinfectant.^{2,8} Obtain larger volumes of urine for special analyses (not culture) aseptically from the drainage bag.^{2,8}
- Do not screen for asymptomatic bacteruria (ASB) in catheterised patients using a dipstick or a laboratory test (M&C&S) (note that ASB is no longer included in the NHSN surveillance definitions for UTI).^{2,8}
- Do not treat ASB in catheterised patients except before invasive urologic procedures.⁸
- Do not change short-term catheters routinely.^{2,8}
- Long-term indwelling catheters should be changed at intervals adapted to the individual patient.⁷ There is insufficient evidence to support the practice of changing catheters at fixed intervals.⁷ It is suggested therefore, that the decision to change catheters and drainage bags should be based on clinical indications such as infection, obstruction or a break in the closed system.²
- Clamping indwelling catheters before removal is unnecessary.^{2,13}
- Minimise the duration of catheterisation; that is, catheterisation should be only for as long as indications exist.^{2,7,8}

Quality improvement

Positioning CAUTI prevention strategies within an infection prevention and control framework allows for targeted monitoring, assessment and evaluation of interventions. Furthermore, it requires the identification of responsibilities and roles for addressing the prevention of CAUTIs. Under the umbrella of quality improvement, the following recommendations provide guidance to organisations to reduce the risk of CAUTI.

1. A hospital or healthcare organisations' Chief Executive Officer and senior management should be responsible for ensuring that the healthcare system supports an infection prevention and control program and that this program reduces the risks of CAUTI and the transmission of epidemiologically significant pathogens.⁸ An infection control program should include the provision of education about CAUTI, other complications of urinary catheterisation and alternatives to indwelling catheters.^{2,8}
2. Healthcare organisations should consider implementing quality improvement (QI) programs to enhance appropriate use of indwelling catheters and to reduce the risk of CAUTI.² Such QI programs could include the development and implementation of institutional policies requiring continual (daily) review of the need for continued catheterisation.⁸ Electronic or other types of reminders may be useful in assisting in complying with such a recommendation. Examples of strategies to monitor the continued need for urinary catheters include: automatic stop orders requiring renewal of the order for continuation of the indwelling catheter; standardised

reminders placed into the patient record; and daily ward rounds by clinical staff to review all patients with urinary catheters and to ascertain continuing necessity of the catheter.⁸

3. A surveillance program (either process or outcome) would assist in monitoring the rates of urinary tract infections. Such a program could be developed based on a facility risk assessment.² Examples of programs related to CAUTI surveillance and QI could include the establishment of a system for analysing and reporting data on catheter use and adverse events from the catheter, including stratification by relevant risk factors (e.g. sex, age, ward and duration).^{2,8}

Unresolved issues

Either a lack of compelling evidence or a limited number of studies make it difficult to form recommendations addressing several clinical and administrative aspects of catheter use. To elicit appropriate guidance and resolve these issues, further work is required in the following areas:

- Evidence on effect of silver alloy catheters in reducing the risk of symptomatic and asymptomatic bacteruria has been reported in a small number of studies.^{18–21} However, reviews of catheter materials have concluded that the use of silver alloy impregnated catheters has no significant advantages and these are not recommended for routine use. The question of the benefit of using other types of antimicrobial-coated catheters remains unresolved as does the question of identifying which catheter materials best delay the onset of bacteruria, bacterial adherence and bacterial growth.^{7,8}
- The benefit of using an antiseptic solution versus the use of sterile saline for meatal cleaning before catheter insertion remains unresolved.^{2,8}
- The role of periodic (e.g. night time) use of condom catheters in incontinent male patients is unresolved.^{2,8}
- The use of catheters to prevent skin breakdown is unresolved.^{2,8}
- There is limited evidence that post-operative intermittent catheterisation reduces the risk of bacteruria compared with an indwelling catheter.⁷
- Finally, there are no standardised surveillance measures of CAUTI that would aid in inter-regional assessments and external reporting.⁸

Suggested care bundle/checklist

To elicit a consistent and pro-active response to the guidelines summarised here, we recommend the adoption of a care bundle/checklist (Figs 1, 2). Primarily, this would serve as a checklist of considerations taken to ensure catheter use is limited. If catheterisation is indicated, the checklist provides evidence-based guidelines to follow to reduce the risk of acquiring a CAUTI. Secondly, the use of a checklist provides a source of catheter documentation, this being an

INSERTION OF INDWELLING URINARY CATHETER			
Patient Name:	Completed form to be sent to infection control/S&Q unit (where appropriate) out		
Patient ID:			
Date:			
Insertion Date:			
Inserted by (name & position):			
Insertion checklist completed by:			
	Yes	No	Comments
1. Alternatives for catheterisation are explored (such as suprapubic catheters, condom drainage systems, or intermittent catheterisation).			
2. Clinical indication for catheterisation is met (mark which one):			
a) Acute urinary retention or obstruction			
b) Urinary output monitoring in critically ill patients			
c) Peri operative use for selective surgical procedures			
• Urological surgery or surgery to other contiguous – structures of genitourinary tract			
• Prolonged duration of surgery (removed in recovery)			
• Large volumes of infusions or diuretics administered intraoperatively			
• Operative patients with urinary incontinence			
• Need for intraoperative monitoring of output			
d) Healing of wounds (sacral/perianal) in incontinent patients (as part of a holistic plan)			
e) Patient requiring prolonged immobilisation (e.g., potentially unstable thoracic or lumbar spine, multiple traumatic injuries such as pelvic fractures)			
f) Exceptional circumstances e.g. comfort at end of life			
The person inserting the catheter has received training to do so			
The person inserting the catheter has performed hand hygiene immediately before undertaking the procedure			
The small size catheter has been selected			
Gloves, a drape, and sponges; a sterile or antiseptic solution for cleaning the urethral meatus; and a single-use packet of sterile lubricant jelly was used during insertion			
The catheter was inserted using aseptic non touch technique and sterile equipment			
Immediately after insertion:			
Documentation was made according to local policy			
The catheter was placed off of the floor, below the bladder			
Patient was given the following advice:			
a) Keep the catheter below the bladder, but keep the catheter bag off the floor			
b) The expected time the catheter will be inserted for			

Fig. 1. Example of a checklist for catheter insertion. This has been adapted from a CAUTI bundle and checklist developed by Health Protection Scotland²⁵ and Institute for Healthcare Improvement²⁶ and with consideration and input from ASID/AICA.

MAINTENANCE INDWELLING URINARY CATHETER								
Patient Name:						Completed form to be sent to infection control/S&Q unit (where appropriate)		
Patient ID:								
Date:								
Insertion Date:								
Type of catheter:								
Compliance Task	Wk comm	Day 1	Day 2	Day 3	Day 4	Day 5	Day 6	Day 7
There is daily documented assessment of the need for the UC								
A clinical indication for catheterisation remains								
The catheter has been continuously connected								
The patient is aware of his/her role in minimising the risk of developing a urinary tract infection, or daily meatal hygiene has been performed								
Action: Request removal OR Leave <i>in situ</i>								
Name of person completing check								
Initial, date & time of person completing check								

Fig. 2. Example of a checklist for the maintenance of an indwelling catheter.

important step in the monitoring and analysis of CAUTIs. The checklist could exist as a preliminary stand-alone measure before the implementation of a more holistic institutional policy, or as an adjunct to such a policy, such as a clinical care pathway. Regardless of where the checklist is positioned within institutional policy, in order to achieve holistic, systemic improvements to the way catheter care is administered, the checklist should be underpinned by leadership, culture change and measurement.^{22,23} Further, the checklist could easily be broadened to include post-insertion information such as the onset of bacteruria, any other adverse events resulting from catheter use and the eventual duration of catheterisation. The checklist provided can be added to or be modified according to organisational needs.

Summary and future work

After reviewing key publications on the prevention of CAUTIs, it appears there is consensus on the major recommendations and issues relating to the insertion, maintenance and quality improvement practices needed to reduce the incidence of CAUTIs. This AICA/ASID position statement reflects these recommendations. One of the key principles in the prevention of CAUTIs is to minimise both the number of indwelling catheters inserted and also

the duration of catheterisation of individuals. Quality improvement activities form a fundamental part of CAUTI prevention, with one key QI activity being surveillance. The issue of CAUTI surveillance is an area that particularly requires further work and exploration, and the AICA and ASID would encourage this work to occur.

Conflicts of interest

The authors have no conflicts to declare.

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