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Policies for replacing long-term indwelling urinary catheters in adults (Review)

Cooper FPM, Alexander CE, Sinha S, Omar MI

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[Intervention Review]

Policies for replacing long-term indwelling urinary catheters in adults

Fergus PM Cooper[1], Cameron Edwin Alexander[1], Sanjay Sinha[2], Muhammad Imran Omar[1]

[1]Academic Urology Unit, University of Aberdeen, Aberdeen, UK. [2]Apollo Hospital, Hyderabad, India

Contact address: Fergus PM Cooper, Academic Urology Unit, University of Aberdeen, Health Sciences Building, Foresterhill, Aberdeen, AB25 2ZD, UK. fergus_cooper_1992@hotmail.com.

A B S T R A C T

Background

Long-term indwelling catheters are used commonly in people with lower urinary tract problems in home, hospital and specialised health-care settings. There are many potential complications and adverse effects associated with long-term catheter use. The effect of health-care policies related to the replacement of long-term urinary catheters on patient outcomes is unclear.

Objectives

To determine the effectiveness of different policies for replacing long-term indwelling urinary catheters in adults.

Search methods

We searched the Cochrane Incontinence Specialised Trials Register, which contains trials identified from the Cochrane Central Register of Controlled Trials (CENTRAL), MEDLINE, MEDLINE In-Process, MEDLINE Epub Ahead of Print, CINAHL, ClinicalTrials.gov, WHO ICTRP and handsearching of journals and conference proceedings (searched 19 May 2016), and the reference lists of relevant articles.

Selection criteria

All randomised controlled trials investigating policies for replacing long-term indwelling urinary catheters in adults were included.

Data collection and analysis

At least two review authors independently performed data extraction and assessed risk of bias of all the included trials. Quality of evidence was assessed by adopting the GRADE approach. Any discrepancies were resolved by discussion between the review authors or an independent arbitrator. We contacted the authors of included trials to seek clarification where required.

Main results

Three trials met the inclusion criteria, with a total of 107 participants in three different health-care settings: A USA veterans administration nursing home; a geriatric centre in Israel; and a community nursing service in Hong Kong. Data were available for three of the pre-stated comparisons. Priefer and colleagues evaluated different time intervals between catheter replacement (n = 17); Firestein and colleagues evaluated the use of antibiotic prophylaxis at the time of replacement (n = 70); and Cheung and colleagues compared two different types of cleaning solutions (n = 20).

All the included trials were small and under-powered. The reporting of the trials was inadequate and as a result, risk of bias assessment was judged to be unclear for the majority of the domains in two out of the three trials. There was insufficient evidence to indicate that (i) there was a lower incidence of symptomatic UTI in people whose catheter was changed both monthly and when clinically indicated (risk ratio (RR) 0.35, 95% confidence interval (CI) 0.13 to 0.95; very low quality evidence) compared to only when clinically indicated, (ii) there was not enough evidence to assess the effect of antibiotic prophylaxis on reducing: positive urine cultures at 7 days (RR 0.91, 95% CI 0.79 to 1.04); infection (RR 1.41, 95% CI 0.55 to 3.65); or death (RR 2.12, 95% CI 0.20 to 22.30; very low quality evidence), (iii) there was no statistically significant difference in the incidence of asymptomatic bacteruria at 7 days (RR 0.80, 95% CI 0.42 to 1.52) between people receiving water or chlorhexidine solution for periurethral cleansing at the time of catheter replacement. However, none of the 16 participants developed a symptomatic catheter-associated urinary tract infection (CAUTI) at day 14.

The following outcomes were considered critical for decision-making and were also selected for the 'Summary of findings' table: (i) participant satisfaction, (ii) condition-specific quality of life, (iii) urinary tract trauma, and (iv) formal economic analysis. However, none of the trials reported these outcomes.

None of the trials compared the following comparisons: (i) replacing catheter versus other policy e.g. washouts, (ii) replacing in the home environment versus clinical environment, (iii) clean versus aseptic technique for replacing catheter, (iv) lubricant A versus lubricant B or no lubricant, and (v) catheter user versus carer versus health professional performing the catheter replacement procedure.

Authors' conclusions

There is currently insufficient evidence to assess the value of different policies for replacing long-term urinary catheters on patient outcomes. In particular, there are a number of policies for which there are currently no trial data; and a number of important outcomes which have not been assessed, including patient satisfaction, quality of life, urinary tract trauma, and economic outcomes. There is an immediate need for rigorous, adequately powered randomised controlled trials which assess important clinical outcomes and abide by the principles and recommendations of the CONSORT statement.

PLAIN LANGUAGE SUMMARY

Policies for replacing long-term indwelling urinary catheters in adults

Background information

A urinary catheter is a tube that is inserted into the bladder from the end of the urethra to drain urine from the bladder. Usually, urinary catheters are only required for a few days, such as after an operation. However, there are some medical conditions that may require bladder drainage on a long-term basis. There are many different ways to care for and maintain a long-term urinary catheter. In this review we refer to these different care methods as health-care 'policies'. Examples of policies that relate to the replacement of a long-term catheter include: time between catheter replacements; use of antibiotics during replacement; use of cleaning solutions or lubricants during replacement; and personnel, environment and techniques used at replacement. This review aimed to identify which policies at the time of long-term catheter replacement were most effective in improving patient care.

The main findings of the review

This review identified that there is currently insufficient high-quality evidence which evaluates the effectiveness of different policies for replacing long-term urinary catheters. Only three randomised clinical trials, which included a total of 107 participants, were eligible and included in this review.

These trials evaluated: (i) different time intervals for catheter replacement, (ii) the use of antibiotics to prevent infection and (iii) the use of different cleaning solutions. There was insufficient evidence to suggest that replacing the catheter monthly and when there was a clinical reason to do so reduced bacteria in the urine compared to replacing the catheter only when there was a clinical reason to do so. However, there was not enough evidence to say whether using antibiotics at the time of replacing the catheter for prevention of infection was effective or whether using water to cleanse during catheter replacement was as effective as an anti-bacterial washing solution.

Adverse effects

None of the trials reported any adverse effects relating to the policies investigated.

Any limitations of the review

All three trials which were included in this review were very small with methodological flaws. Therefore new trials are needed in order to definitely answer this research question. The evidence in this review is current up to 19 May 2016.

SUMMARY OF FINDINGS FOR THE MAIN COMPARISON [*Explanation*]

Monthly & PRN catheter replacement versus PRN catheter replacement for replacing long-term indwelling urinary catheters in adults

Patient or population: adults with replacing long-term indwelling urinary catheters
Settings: USA veterans administration nursing home
Intervention: Monthly & PRN catheter replacement versus PRN catheter replacement

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE)	Comments
	Assumed risk	Corresponding risk				
	Control	Monthly & PRN catheter replacement versus PRN catheter replacement				
Symptomatic CAUTI as defined by trialists	857 per 1000	300 per 1000 (111 to 814)	**RR 0.35** (0.13 to 0.95)	17 (1 study)	⊕○○○ **very low**[1,2]	
Participant satisfaction - not reported	See comment	See comment	Not estimable	-	See comment	
Condition-specific quality-of-life measures - not reported	See comment	See comment	Not estimable	-	See comment	
Adverse effects (Urinary tract trauma) - not reported	See comment	See comment	Not estimable	-	See comment	
Adverse effects (Death) - not reported	See comment	See comment	Not estimable	-	See comment	

Formal economic analysis - not reported	See comment	See comment	Not estimable	-	See comment

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).

CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence

High quality: Further research is very unlikely to change our confidence in the estimate of effect.

Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.

Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.

Very low quality: We are very uncertain about the estimate.

[1] Downgraded two levels: Sequence generation, allocation concealment and blinding of outcome assessment was judged to be unclear. Blinding of participants and personnel was judged to be at high risk of bias.The outcome was reported by one under-power study.

[2] Downgraded one level: As the sample size and the event rate is small.

PRN: from the Latin "pro re nata", meaning " as needed".

BACKGROUND

Description of the condition

Long-term indwelling urinary catheters can be used in the care of people with urinary incontinence or urinary retention. These conditions can arise from a variety of underlying pathologies that cause dysfunction of bladder storage or emptying, such as multiple sclerosis and benign prostatic hyperplasia. The prevalence of long-term catheter use varies between countries and health-care settings (Stensballe 2005). A study by Sørbye and colleagues found that, in Italy, 35.9% of male and 27.4% of female elderly home care clients used either an indwelling, intermittent or condom catheter. However, rates of use of these catheters were lower in men in Finland (2.9%) and in women in the Czech Republic (0.6%) (Sørbye 2009). There are many well recognised and common complications associated with the use of long-term urinary catheters, including urinary tract infections, encrustation due to mineral deposits, and peri-catheter leakage. The majority of micro-organisms that cause catheter-associated urinary tract infections (CAUTIs) are derived from the patient's own colonic and perineal flora and the hands of health-care professionals involved in catheter insertion or management (Maki 2001). Studies have shown a 5% to 8% increase in the risk of developing bacteriuria for each day following catheterisation, and that almost all users will have bacteriuria four weeks after long-term catheter insertion (Gould 2010; Nicolle 2001; Stamm 1991). Leakage, blockage, pain and dislodgement of long-term catheters are also recognised as commonly occurring problems that often require additional use of health-care resources and services (Wilde 2013). These complications will undoubtedly have an impact on patient quality of life and satisfaction.

Description of the intervention

A number of different policies can be employed during the time of catheter replacement. These policies are described below.

Time between replacements

The recommended time between catheter replacement depends on local policies and varies significantly between centres (Palka 2014; Willson 2009). This discrepancy in clinical practice reflects a lack of evidence to support the early or late replacement of long-term urinary catheters in the reduction of adverse outcomes. The Infectious Diseases Society of America (IDSA) states there is currently insufficient data to recommend a specific time interval between long-term catheter or suprapubic catheter replacement (Hooton 2010). A shorter time interval between catheter replacements may reduce the development of a biofilm that can act to harbour bacteria, and may also reduce the likelihood of mechanical blockage.

However, the tissue disruption caused by more frequent catheter replacement could contribute to the development of CAUTI and other adverse outcomes.

Antibiotics as prophylaxis

For people requiring a long-term indwelling catheter, antibiotics may be given prophylactically in an attempt to prevent CAUTI development or at the time of developing the symptoms associated with CAUTI. However, prophylactic antibiotics have been shown to permit the development of resistant organisms (Hooton 2010). A comparison of prophylactic, clinically indicated and microbiologically indicated antibiotics has been previously explored in another Cochrane Review (Lusardi 2013). This concluded that data was limited and that there was insufficient evidence to determine best practice.

Catheter replacement environments, personnel and techniques

Long-term indwelling urinary catheters can be replaced by personal carers, health-care professionals or by the catheter users themselves. Furthermore, the procedure may be carried out by the catheter user in their own home, by a health-care professional visiting the user at home, or by a health-care professional in a clinical environment or nursing home. Catheter replacement can be performed using either a clean or an aseptic approach. A clean approach involves the use of non-sterile gloves and the cleansing of the external urethral meatus and surrounding area with a non-antiseptic solution. An aseptic approach involves the use of sterile gloves, sterile barriers, antiseptic cleaning solutions and a non-touch technique. Clinical practice worldwide varies according to local policies, individual preferences and the specific clinical setting (Willson 2009). A previous Cochrane Review concluded that existing data does not provide convincing evidence that any specific technique (aseptic or clean), catheter type (coated or uncoated), method (single use or multiple use), or person (self or other) can be considered as the gold standard in the use of intermittent catheters (Moore 2007) .

Cleaning solutions

There are various cleaning solutions which can be used when replacing catheters, including sterile water, chlorhexidine gluconate and povidone-iodine. The cleaning procedure may include cleaning the perineal area as well as the periurethral area. Catheters may also be cleaned instead of replaced. Washout policies for catheters have been evaluated by a previous Cochrane review (Hagen 2010). The review indicated that "the evidence was too scanty to conclude whether or not washouts were beneficial".

Lubricants

Lubricants are commonly used in most types of urethral catheterisation to allow for easier insertion and patient comfort and typically contain local anaesthetic and antiseptic. The administration of lubricants can be incorporated into either a clean or an aseptic catheterisation technique.

Other catheter reviews

This review is one of a series of Cochrane Reviews addressing different issues in the use of catheters for long-term management of catheter users for a variety of conditions. These include:

- Washout policies in long-term indwelling urinary catheterisation in adults (Hagen 2010)
- Types of indwelling urinary catheters for long-term bladder drainage in adults (Jahn 2007)
- Long-term bladder management by intermittent catheterisation in adults and children (Moore 2007)
- Urinary catheter policies for long-term bladder drainage (Niël-Weise 2005).

Why it is important to do this review

Long-term indwelling urinary catheters are used commonly in health care. Guidelines and protocols exist worldwide but are not supported by extensive or high-quality evidence. It is important that policies for replacing long-term indwelling urinary catheters are updated based on all of the existing high-quality evidence. There is an immediate need to improve the rates of adverse effects and optimise patient quality of life, patient and carer satisfaction and resource management associated with long-term catheter use. The review will also identify the specific needs for future research in this area.

O B J E C T I V E S

To determine the effectiveness of different policies for replacing long-term indwelling urinary catheters in adults.

M E T H O D S

Criteria for considering studies for this review

Types of studies

Randomised controlled trials (RCTs) or quasi-RCTs.

Types of participants

Adults (over the age of 18 years) with long-term (> 14 days) indwelling urinary or suprapubic catheters that are anticipated to require replacement.

Types of interventions

The following types of interventions were compared with each other:

1. One interval versus another interval between catheter replacement
2. Antibiotic prophylaxis versus no prophylaxis at the time of catheter replacement
3. Replacing catheter versus other policy e.g. washouts
4. Replacing in home environment versus clinical environment
5. Clean versus aseptic technique for replacing catheter
6. Cleaning solution A versus cleaning solution B
7. Lubricant A versus lubricant B or no lubricant
8. Catheter users versus carer versus health professional performing the catheter replacement procedure

This review did not include: type/material of catheter (Jahn 2007), washout versus no washout in long-term indwelling urinary catheters (Hagen 2010), long-term antibiotics (Niël-Weise 2005), and the use of intermittent catheters (Moore 2004), as these areas have been addressed in other Cochrane Reviews.

Types of outcome measures

Primary outcomes

- Participant satisfaction
- Symptomatic catheter-associated urinary tract infection (CAUTI) as defined by trialists

Secondary outcomes

Participant-reported quality of life

- Generic quality-of-life measures
- Condition-specific quality-of-life measures
- Psychological outcome measures

Clinician-reported outcomes

- Clinician satisfaction
- Number of participants requiring more frequent replacements (than per protocol)
- Duration of use of single catheter

Carer-reported outcomes

- Carer satisfaction

Adverse effects

- Urinary tract trauma
- Pain/discomfort
- Haematuria
- Asymptomatic bacteruria
- Systemic infection (septicaemia)
- Encrustation/breakdown of catheter
- Pericatheter leakage
- Stricture formation
- Failure to achieve catheter replacement
- Hospitalisation
- Bladder calculi
- Bladder cancer

Economic outcomes

- Cost of intervention
- Resource implications
- Formal economic analysis (cost effectiveness)

Other outcomes

Any other outcomes considered to be important if reported in trials.

Search methods for identification of studies

We did not impose any restrictions, for example language or publication status, on the searches described below.

Electronic searches

This review drew on the search strategy developed for Cochrane Incontinence. We identified relevant trials from the Cochrane Incontinence Specialised Trials Register. For more details of the search methods used to build the Specialised Register please see the Group's module in *The Cochrane Library*. The Register contains trials identified from the Cochrane Central Register of Controlled Trials (CENTRAL), MEDLINE, MEDLINE In-Process, MEDLINE Epub Ahead of Print, CINAHL, ClinicalTrials.gov, WHO ICTRP , UK Clinical Research Network Portfolio and hand-searching of journals and conference proceedings. Many of the trials in the Cochrane Incontinence Specialised Register are also contained in CENTRAL. The date of the last search was 19 May 2016.

The terms used to search the Cochrane Incontinence Specialised Register are given in Appendix 1.

Searching other resources

We searched the reference lists of relevant articles.

Data collection and analysis

Selection of studies

We included only randomised and quasi-randomised controlled trials. At least two review authors independently screened the list of titles and abstracts generated by our search. We retrieved full-text articles of potentially relevant studies. At least two review authors independently assessed the full-text articles for eligibility. We contacted study investigators for additional information when required. We resolved any differences of opinion by discussion or involvement of a third party. We listed studies formally considered for the review that were subsequently excluded along with the reasons for their exclusion.

Data extraction and management

Two of the review authors independently extracted data of the included studies by using a standardised form. Any disagreement was resolved by discussion or by consulting a third party. We contacted study authors when there was insufficient information regarding the primary outcome in the published reports or when additional information was required. We used Review Manager software (RevMan 5.3) for data entry. We processed the included trial data according to the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011).

Assessment of risk of bias in included studies

We assessed the risk of bias in the included studies using the Cochrane 'Risk of bias' assessment tool (Higgins 2011). This included assessment of:

- sequence generation
- allocation concealment
- blinding of participants or therapists
- blinding of outcome assessors
- completeness of outcome data
- selective outcome reporting
- other potential sources of bias

Two of the review authors independently assessed these domains and rated each as 'high risk', 'low risk' or 'unclear risk'. We resolved any differences of opinion by consensus or by consulting a third party.

Measures of treatment effect

We based analysis on available data from all included trials relevant to the comparisons and outcomes of interest. For trials with multiple publications, only the most up-to-date or complete data for each outcome were included.

For categorical outcomes we related the numbers reporting an outcome to the numbers at risk in each group to calculate a risk ratio (RR) with 95% confidence interval (CI). For continuous variables we planned to use means and standard deviations to calculate a mean difference (MD) with 95% CI; however no continuous variables were encountered.

If we had found similar outcomes reported on different scales, we would have calculated the standardised mean difference (SMD). We would have reversed the direction of effect, when necessary, to ensure consistency across trials. If data to calculate RRs or MDs were not given, we would have utilised the most detailed numerical data available to calculate the actual numbers or means and standard deviations (for example test statistics, P values).

Unit of analysis issues

In simple parallel group designs, when participants are individually randomised, the primary analysis was per participant randomised. We analysed studies with multiple treatment groups by treating each pair of arms as a separate comparison, as appropriate. We planned to undertake analysis of studies with non-standard designs, such as cross-over trials and cluster-randomised trials, as described in the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011); however no such trials were included.

Dealing with missing data

We analysed data on an intention-to-treat basis, as far as possible, meaning that all participants were analysed in the groups to which they are randomised. We made attempts to obtain missing data from the original trialists. Where this was not possible, we reported the data as given in the studies.

If trials had reported sufficient detail to calculate MDs but gave no information on associated standard deviations (SD), we would have assumed the outcome had an SD equal to the highest SD from other trials within the same analysis. We did not need to do this as meta-analysis was not performed.

Assessment of heterogeneity

We did not perform any assessment of heterogeneity as none of the pre-specified comparisons were addressed by more than one trial. We had intended to combine trials only if they were thought to be clinically similar. We would have assessed heterogeneity between studies by visual inspection of the plots of data, the Chi² test for heterogeneity and the I²statistic (Higgins 2011). We would have defined the thresholds for interpretation of the I² statistic according to the *Cochrane Handbook for Systematic Reviews of Interventions* (Higgins 2011).

Assessment of reporting biases

In view of the difficulty of detecting and correcting for publication bias and other reporting biases, we aimed to minimise their potential impact by ensuring a comprehensive search for eligible studies and by being alert to duplication of data. We encountered no duplication of data as no comparison was investigated by more than one trial.

Data synthesis

We intended to combine trials if interventions were similar, based on clinical criteria; however we did not find more than one study addressing any pre-specified comparison. We would have conducted a meta-analysis with a fixed-effect model to combine trial data unless there had been evidence of heterogeneity across studies.

Subgroup analysis and investigation of heterogeneity

We intended to perform sub-group analysis in order to explore the impact of these sub-groups on the interventions. We selected the following sub-groups:

1. Different catheter materials

Catheter material is a key determinant of encrustation (and presumably blockage) and this is directly linked to the interval between replacements. Antimicrobial impregnation of the catheter or use of special low-friction catheters could impact the need for antibiotic prophylaxis or a lubricant.

2. Participants over 75 versus participants under 75

Elderly people have unique anatomical and functional changes, both local and systemic. Elderly individuals are more likely to have voiding dysfunction, altered lower urinary tract anatomy, co-morbidities, impaired immune function and poor local hygiene. These could have an impact on antimicrobial policy, health-care setting for catheter change or use of an aseptic technique. The authors have chosen 75 years as a marker for the age at which these issues would likely be most relevant.

3. Sex: male versus female

Sex of the individual can influence the interventions under study. The unique anatomy of the female perineum makes women more prone to urinary infections but also renders catheter change technically easy. Men with gross prostatic enlargement could pose technical problems in catheter change demanding greater skill and a hospital setting.

4. Level of care: community/self-care versus residential/assisted living

The level of care such as community care, self care or an assisted facility could potentially have an impact on the policies chosen with regard to catheter change in people requiring long-term urinary catheters.

5. Condition requiring catheterisation

Non-neurological versus neurological reason for indwelling catheter

The underlying condition could have an impact on the chosen policy. For example catheter change in people with neurogenic lower urinary tract dysfunction could be associated with distinct challenges. People with neurogenic bladder on long-term indwelling catheters are usually clinically advised to receive intermittent catheterisation. Many of these people have poor hand function, quadriplegia or impaired cognition and these factors are more likely to be associated with poor local hygiene, impaired nutrition and immunity. Some people with neurogenic dysfunction might have elevated bladder pressures in the form of severe neurogenic detrusor overactivity or poor compliance with consequent risk to the upper urinary tracts. This could conceivably make them more prone to sepsis during urethral manipulations.

People with retention versus those with incontinence

People with incontinence requiring an indwelling catheter are more likely to have cognitive impairment, limited ambulation or neurological disease such as Parkinsonism or stroke; while those with intractable retention are more likely to be suffering from severe cardiovascular or pulmonary disease. All these can influence the interventions under study in unique ways.
However, subgroup analysis could not be performed due to the lack of included trials.

Sensitivity analysis

We intended to conduct sensitivity analyses by including or excluding trials we judged as high risk of bias. We did not conduct sensitivity analyses because meta-analysis was not performed.

Quality of evidence

Primary and secondary outcomes, as defined above, were classified by the review authors as 'critical', 'important' or 'not important' for decision making from the patients' perspective (Gould 2010; Guyatt 2011a; Guyatt 2011b; Guyatt 2013a; Guyatt 2013b). The GRADE Working Group recommend including up to seven outcomes in a systematic review. In this systematic review, GRADE methodology was adopted for assessing the quality of the evidence for the following outcomes classified as critical:

- Symptomatic CAUTI as defined by trialists
- Participant satisfaction
- Condition-specific quality-of-life measures
- Adverse effects (urinary tract trauma)
- Adverse effects (death)
- Formal economic analysis

R E S U L T S

Description of studies

Results of the search

We screened a total of 1125 records, identified by the literature search, for this review. From these records, we considered 61 for full text screening, of which we deemed three were suitable for inclusion in this review (Cheung 2008; Firestein 2001; Priefer 1982). The flow of literature through the assessment process is shown in the PRISMA flowchart (Figure 1).

Figure 1. PRISMA study flow diagram

1125 records identified through database searching

0 additional records identified through other sources

1125 records after duplicates removed

1125 records screened

1064 records excluded

61 full-text articles assessed for eligibility

58 reports of 53 studies excluded, with reasons given in the Characteristics of excluded studies table

3 reports of 3 studies included in qualitative synthesis

0 studies included in quantitative synthesis (meta-analysis)

Included studies

Cheung 2008 randomised 20 subjects (6 males, 14 females) in a Hong Kong community nursing service to either sterile water or conventional 0.05% chlorhexidine gluconate for periurethral cleansing prior to insertion of a replacement of long-term urinary catheter. Subjects were excluded if they had a symptomatic UTI at the time of the replacement, were on antibiotic treatment or if they had only been using a long-term catheter for less than one month. The mean age of the subjects was 78.4 (SD = 11.8) years, and 55% lived in a nursing home. Urine cultures were taken from participants at baseline and then 1, 7 and 14 days after catheter replacement. Reported outcomes were colonisation counts greater than 10^5 cfu/mL and incidence of symptomatic CAUTIs.

Firestein 2001 randomised 70 subjects (21 males, 49 females) in a geriatric centre in Israel to receive either 1 g of intravenous meropenem 30 minutes before long-term catheter replacement or no antibiotic before replacement. Subjects were excluded if their urinary catheter had been in place for less than 4 weeks or if there had been antibiotic use within the 2-week period before enrolment. The mean age of the subjects was 79.3 (SD = 9.6) years. Urine cultures were taken from participants at baseline and then 3, 7, 14 and 28 days after catheter replacement. Reported outcomes were positive urine cultures, incidence of infection, incidence of bacteraemia and death.

Priefer 1982 randomised 17 men in a USA veterans administration nursing home to catheter replacements either: only for obstruction and/or infection or monthly as well as when indicated by obstruction and/or infection. Subjects who required transfer to a hospital for acute problems unrelated to the urinary tract were excluded. The mean age of subjects was 77.1 (SD = 16.3) years in the control group and 83.4 (SD = 7.9) years in the intervention group. Participants were observed over a six-month period and the following outcomes were reported: total number of irrigations required; number of catheter changes per month required; incidence of symptomatic CAUTI over the six months.

Excluded studies

We have listed the excluded studies along with reasons for exclusion in the Characteristics of excluded studies table. The most common reason for excluding studies was that the intervention did not relate to replacing long-term catheters.

Risk of bias in included studies

Detailed results of the 'Risk of bias' assessment are provided in Figure 2; Figure 3 and judgement of individual domains are summarised below.

Figure 2.   Risk of bias graph: review authors' judgements about each risk of bias item presented as percentages across all included studies.
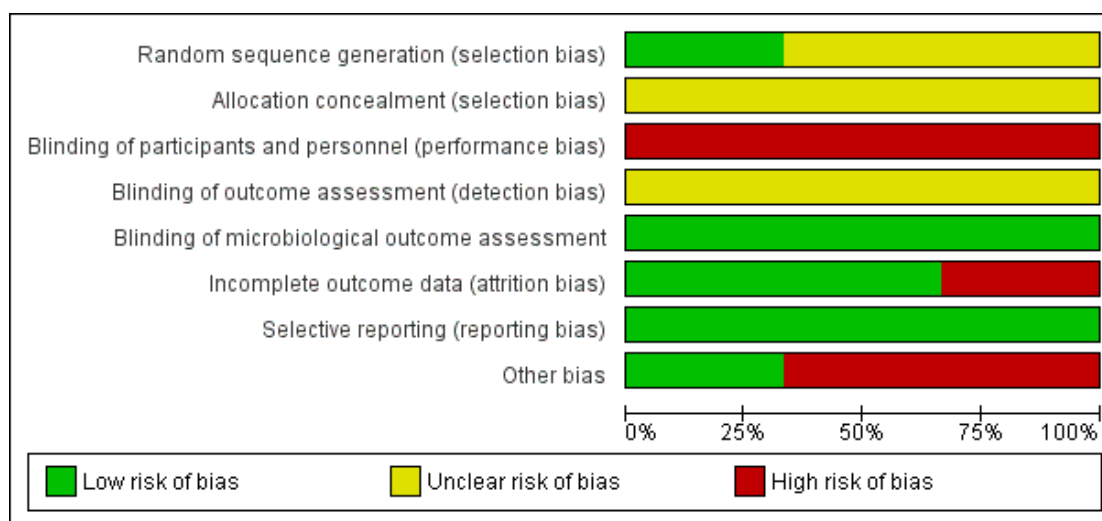
Figure 3.   Risk of bias summary: review authors' judgements about each risk of bias item for each included study.

Allocation

Random sequence generation

Cheung 2008 provided details of sequence generation and was judged to be at low risk of bias. We judged the two remaining trials to be unclear (Firestein 2001; Priefer 1982)

Concealment of allocation

Methods of allocation concealment were inadequately described in all three trials and we judged them to be unclear (Cheung 2008; Firestein 2001; Priefer 1982)

Blinding

Blinding of participants and personnel in two studies are not described and are assumed to be impossible due to the nature of the interventions (Firestein 2001; Priefer 1982).
Blinding participants in one study to their method of periurethral cleansing would be possible, but is not described and therefore assumed not to occur (Cheung 2008). Additionally, this study provides no information on blinding of personnel. We assumed the outcomes in these studies were reported by the same unblinded personnel administering the intervention and they are therefore also classed as high risk.
Microbiological outcomes for the three studies are classed as low risk, as we assumed microbiologists analysing urine cultures to be blind to participants.

Incomplete outcome data

We judged two studies to be at low risk of bias (Cheung 2008; Priefer 1982). We judged Firestein 2001 to be at high risk of bias for this domain.

Selective reporting

All three trials reported all outcomes intended to be investigated by the trial and have been classed as low risk. Priefer 1982 does not report microbiological outcome, but this was not possible as urine cultures were not taken during this trial. Firestein 2001 reports incidence of CAUTI in the two groups, but not number of CAUTIs per subject over the time period.

Other potential sources of bias

We classed two studies as high risk for other areas of bias. In one study, the participants were all male (Priefer 1982). This was unavoidable due to the nature of the study setting and this could have an impact on the outcome and therefore it was judged to be high risk. There is also a wide age range in participants. In the other study, there is a calculation error, inconsistent data and a positive urine culture is not defined (Firestein 2001). We tried to contact the author but did not receive any response.

Effects of interventions

See: **Summary of findings for the main comparison** Monthly & PRN catheter replacement versus PRN catheter replacement for replacing long-term indwelling urinary catheters in adults; **Summary of findings 2** Antibiotics at time of replacement versus no antibiotics at time of replacement for replacing long-term indwelling urinary catheters in adults; **Summary of findings 3** Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement for replacing long-term indwelling urinary catheters in adults

1. One interval versus another interval between catheter replacement

One small trial found that replacing the catheter monthly as well as when clinically indicated resulted in fewer symptomatic CAUTIs than replacing the catheter monthly (RR 0.35, 95% CI 0.13 to 0.95; very low quality evidence; Analysis 1.1) (Priefer 1982). However, when expressed as number of symptomatic CAUTIs per subject over the six months, there was no significant difference between the two interventions. Participants in the 'monthly as well as clinically indicated' group required more catheter replacements but fewer irrigations than those in the clinically indicated group. However, the trial was too small (hence under-powered) to be reliable.

2. Antibiotic prophylaxis versus no prophylaxis at the time of catheter replacement

One small trial found that there was no statistically significant difference in incidence of positive urine culture between giving 1 g of intravenous meropenem 30 minutes before catheter replacement and no antibiotic: (i) 3 days after replacement (RR 0.95, 95% CI 0.80 to 1.13); (ii) 7 days after replacement (RR 0.91, 95% CI 0.79 to 1.04); (iii) 14 days after replacement (RR 0.94, 95% CI 0.85 to 1.04); or (iv) 28 days after replacement (RR 0.94, 95% CI 0.84 to 1.05; Analysis 2.1) (Firestein 2001). Additionally, there was no statistically significant difference in incidence of infection including urosepsis, soft tissue, pneumonia and unknown infection (RR

1.41, 95% CI 0.55 to 3.65; Analysis 2.2); bacteraemia (Analysis 2.3); or death (RR 2.12, 95% CI 0.20 to 22.30; very low quality evidence; Analysis 2.4) between the two groups. However, the trial was too small (hence under-powered) to be reliable.

3. Replacing catheter versus other policy e.g. washouts

No trials were identified that addressed this comparison.

4. Replacing in home environment versus clinical environment

No trials were identified that addressed this comparison.

5. Clean versus aseptic technique for replacing catheter

No trials were identified that addressed this comparison.

6. Cleaning solution A versus cleaning solution B

There was also no statistically significant difference in incidence of symptomatic CAUTI up to 14 days after replacement (Analysis 3.1). However, the trial was too small (hence under-powered) to be reliable. One small trial found that there was no statistically significant difference in rates of asymptomatic bacteriuria between conventional 0.05% chlorhexidine gluconate (CHG) or sterile water periurethral cleansing solutions (i) 0 days after replacement (RR not estimable; 95% CI not estimable), (ii) 7 days after replacement (RR 0.80, 95% CI 0.42 to 1.52), (iii) 14 days after replacement (RR 0.91, 95% CI 0.66 to 1.24; Analysis 3.2) (Cheung 2008).

7. Lubricant A versus lubricant B or no lubricant

No trials were identified that addressed this comparison.

8. Patient versus carer versus health professional performing the catheter replacement procedure

No trials were identified that addressed this comparison.

ADDITIONAL SUMMARY OF FINDINGS [*Explanation*]

Antibiotics at time of replacement versus no antibiotics at time of replacement for replacing long-term indwelling urinary catheters in adults

Patient or population: adults with replacing long-term indwelling urinary catheters
Settings: Geriatric Centre in Israel
Intervention: Antibiotics at time of replacement versus no antibiotics at time of replacement

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE) Comments
	Assumed risk	**Corresponding risk**			
	Control	**Antibiotics at time of replacement versus no antibiotics at time of replacement**			
Symptomatic CAUTI - not reported	See comment	See comment	Not estimable	-	See comment
Participant satisfaction - not reported	See comment	See comment	Not estimable	-	See comment
Condition-specific quality of life measures - not reported	See comment	See comment	Not estimable	-	See comment
Adverse effects (Urinary tract trauma) - not reported	See comment	See comment	Not estimable	-	See comment
Adverse effects (Death)	**28 per 1000**	**59 per 1000** (6 to 619)	**RR 2.12** (0.20 to 22.20)	70 (1 study)	⊕◯◯◯ **very low**[1,3]

Formal economic analysis - not reported	See comment	See comment	Not estimable

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).
CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence
High quality: Further research is very unlikely to change our confidence in the estimate of effect.
Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: We are very uncertain about the estimate.

[1] Downgraded two levels: Sequence generation, allocation concealment and blinding of outcome assessment was judged to be unclear. Blinding of participants and personnel, and incomplete outcome data was judged to be at high risk of bias.
[2] Downgraded one level: 95% CI is wide (0.79 to 1.04)
[3] Downgraded one level: As the sample size and the event rate is small.

Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement for replacing long-term indwelling urinary catheters in adults					

Patient or population: adults with replacing long-term indwelling urinary catheters
Settings: Hong Kong Community Nursing Service
Intervention: Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement

Outcomes	Illustrative comparative risks* (95% CI)		Relative effect (95% CI)	No of Participants (studies)	Quality of the evidence (GRADE) Comments
	Assumed risk	**Corresponding risk**			
	Control	**Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement**			
Symptomatic CAUTI at 14 days after replacement	Study population		Not estimable	16 (1 study)	⊕◯◯◯ **very low**[1,2]
	0 per 1000	0 per 1000 (0 to 0)			
Participant satisfaction - not reported	See comment	See comment	Not estimable	-	See comment
Condition-specific quality of life measures - not reported	See comment	See comment	Not estimable	-	See comment
Adverse effects (Urinary tract trauma) - not reported	See comment	See comment	Not estimable	-	See comment
Adverse effects (Death) - not reported	See comment	See comment	Not estimable	-	See comment

| Formal economic analysis - not reported | See comment | See comment | Not estimable | - | See comment |

*The basis for the **assumed risk** (e.g. the median control group risk across studies) is provided in footnotes. The **corresponding risk** (and its 95% CI) is based on the assumed risk in the comparison group and the **relative effect** of the intervention (and its 95% CI).
CI: Confidence interval; **RR:** Risk ratio;

GRADE Working Group grades of evidence
High quality: Further research is very unlikely to change our confidence in the estimate of effect.
Moderate quality: Further research is likely to have an important impact on our confidence in the estimate of effect and may change the estimate.
Low quality: Further research is very likely to have an important impact on our confidence in the estimate of effect and is likely to change the estimate.
Very low quality: We are very uncertain about the estimate.

[1] Downgraded two levels: Allocation concealment, and blinding of outcome assessment was judged to be unclear. Blinding of participants and personnel was judged to be at high risk of bias. The outcome was reported by one under-power study.
[2] Downgraded one level: As the sample size and the event rate is small.

DISCUSSION

Summary of main results

We identified three trials eligible for inclusion in this review (Cheung 2008; Firestein 2001; Priefer 1982). All the included trials were small and under-powered with inadequate reporting to permit judgement.

There was insufficient evidence to indicate that there was a lower incidence of symptomatic UTI in people whose catheter was replaced both monthly and when clinically indicated (RR 0.35, 95% CI 0.13 to 0.95; very low quality evidence) compared to only when clinically indicated,

There was not enough evidence to assess the effect of antibiotic prophylaxis on reducing: positive urine cultures at 7 days (RR 0.91, 95% CI 0.79 to 1.04); infection (RR 1.41, 95% CI 0.55 to 3.65); or death (RR 2.12, 95% CI 0.20 to 22.30; very low quality evidence).

There was no statistically significant difference in the incidence of asymptomatic bacteriuria at 7 days (RR 0.80, 95% CI 0.42 to 1.52) between participants receiving water or chlorhexidine solution for periurethral cleansing at the time of catheter replacement. However, none of the 16 participants developed a symptomatic catheter-associated urinary tract infection (CAUTI) at day 14.

Main results are summarised in Summary of findings for the main comparison; Summary of findings 2; Summary of findings 3

Overall completeness and applicability of evidence

None of the trials addressed the following comparisons: (i) replacing catheter versus other policy e.g. washouts, (ii) replacing in home environment versus clinical environment, (iii) clean versus aseptic technique for replacing catheter, (iv) lubricant A versus lubricant B or no lubricant, and (v) patient versus carer versus health professional performing the catheter replacement procedure. As for the policies that have been studied in this review, there are several areas of incompleteness. Two out of three studies report rates of CAUTI, an important clinical outcome. However, all the trials were small and the results were imprecise. The trials did not report outcomes such as cost, participant satisfaction, quality of life and other adverse effects such as encrustation. If a policy can be shown to be favourable in relation to these additional outcomes, it would be more likely to be implemented in worldwide practice.

Each trial only explored one alternative intervention in relation to the control. Therefore there is only data on one antibiotic, one cleaning solution and one replacement timing regime. Furthermore, sub-group analysis was not carried out in any of the trials. The settings of these studies represent likely settings where people with long-term catheters would undergo replacements: the community, a geriatric centre and a care home.

Quality of the evidence

We assessed the quality of the evidence by examining the methodology in the report of each included trial. Since the quality of effect estimates are influenced by our judgement of the quality of methodology, the standard of reporting in the included trials directly influences the size of the effect estimates.

Only one trial gave any indication as to randomisation techniques (Cheung 2008). The randomisation method is not described in any detail in the report, but we found details of it through a reference provided and there was sufficient information to award this study low risk for selection bias. None of the three included trials gave any information on blinding of participants to the intervention. Depending on the policy, blinding of participants to their intervention at the time of catheter replacement can be impossible, for example in the trial assessing different times of replacement (Priefer 1982) or giving an antibiotic (Firestein 2001). However, blinding of participants may have been possible for using different cleaning solutions but was not detailed in Cheung 2008. We scored all three trials as high risk for performance bias as, in addition to the lack of participant blinding, we found no descriptions of blinding of personnel in any of the trials.

In regard to detection bias, the three trials do not describe accurately their methods of detecting outcomes. Therefore, we do not know if the personnel assessing these outcomes are the same as the personnel who administered the intervention. We have judged all three trials to be unclear in this domain. We only found one trial to have high risk of attrition bias: this was due to the unexplained dropouts in each arm of the study (Firestein 2001). Priefer 1982 had no dropouts and Cheung 2008 describes dropouts in detail so we judged these studies to have low risk for attrition bias.

One study has some calculation errors in its results as well as discrepancy between different areas of the report (Firestein 2001). We tried to contact the authors of this study for clarification. While we have no clarification, this study has been classed as high risk for other bias.

GRADE quality of evidence was very low for all the assessed outcomes.

Potential biases in the review process

We searched the relevant databases with no language restriction imposed. We acknowledge that these databases may not contain all the trials that could have potentially been included in this review.

Agreements and disagreements with other studies or reviews

Current guidelines on policies regarding long-term urinary catheters show considerable variation regarding this common clinical situation. Most of the available guidance comes from documents that do not specifically address policies with regard to uri-

nary catheters but deal with conditions where long-term catheterisation might be necessary (Gould 2009; Grabe 2015; NICE Clinical Guideline CG97 2010; NICE Clinical Guideline CG139 2012; NICE Clinical Guideline CG148 2012; Saskatchewan Guidelines 2013). Guidelines are often unclear as to whether the comments with regard to use of long-term catheters are more widely applicable.

Timing of replacement of catheters

Existing guidelines take one or more of five different approaches with regard to the replacement of long-term catheters:

1. Most guidelines recommend that catheters should be changed when there is either infection or obstruction (ANZUNS Guideline 2013; Gould 2009; NICE Clinical Guideline CG139 2012; Saskatchewan Guidelines 2013). This is the approach taken by the CDC Guideline on Catheter Associated Urinary Tract Infections that makes a Category II recommendation (implying a weak recommendation) (Gould 2009).

2. In contrast the EAU guideline on Urinary Infections recommends that change should be scheduled in anticipation of obstruction presumably based on previous need for change (Grade B, not based on randomized controlled trials) (Grabe 2015). Such a patient-centric approach is also recommended by the NICE Guideline (NICE Clinical Guideline CG139 2012).

3. A third approach has been to recommend that catheter changes should additionally be guided by the manufacturer's instruction (ANZUNS Guideline 2013; NICE Clinical Guideline CG139 2012). Presumably this would imply usage for the maximum duration that the manufacturer allows.

4. A fourth approach has been taken by two guidelines that refer to specific durations. The UK Royal Cornwall Hospitals NHS Guidance recommends that catheters could be left in place for up to 12 weeks while recognising that this might not always be feasible (Royal Cornwall Hospitals NHS Trust 2015).

5. In contrast the NICE Guideline on Urinary Incontinence in Neurological Disease states that recurrent block occurring before six weeks should be regarded as a red flag and one must reassess for secondary problems (NICE Clinical Guideline CG148 2012). Clearly, catheters are ordinarily expected to remain in situ without change for a longer duration than that. The final approach is a conspicuous lack of any comment on this subject in a guideline where one would ordinarily have expected guidance (NICE Clinical Guideline CG97 2010).

Cleaning and antibiotic policies while changing catheters

1. The UK Royal Cornwall Hospitals Guideline states that cleaning the meatus with saline is suitable since there is no evidence of benefit with antiseptic solutions but makes no mention regarding use of antibiotic prophylaxis during catheter change (Royal Cornwall Hospitals NHS Trust 2015).

2. The Guidelines for the Prevention and Treatment of Urinary Tract Infections (UTIs) in Continuing Care Settings from the Canadian Saskatchewan Government recommends against the use of prophylaxis during catheter change (Saskatchewan Guidelines 2013).

3. Two different NICE Guidelines recommend selective use of antibiotics only in those individuals with a history of recurrent infection during catheter change or on occasions when there has been trauma during catheter change (NICE Clinical Guideline CG139 2012; NICE Clinical Guideline CG148 2012).

4. The EAU Guideline on Urinary Infections recommends against screening for asymptomatic bacteriuria before catheter change (Level of evidence 4, based on expert opinion) (Grabe 2015). The CDC Guideline makes a strong recommendation against the use of prophylactic antibiotics but it is unclear whether the guideline is referring to continued use of prophylaxis in people with on-going long-term catheters or whether this refers to prophylaxis only during catheter replacement (Gould 2009).

The need for additional research on use of single dose antibiotic prophylaxis has also been noted (NICE Clinical Guideline CG139 2012).

AUTHORS' CONCLUSIONS

Implications for practice

There is insufficient evidence to support catheter replacement monthly as well as when clinically indicated compared to only when clinically indicated. Further research with a larger number of participants could turn an insignificant reduction in symptomatic CAUTIs in the monthly replacement group to a significant reduction. It is unclear which policy is more favourable economically, as performing monthly replacements in addition to replacements when clinically indicated incurs more replacements, but less irrigations. As these figures were also insignificant further research is needed, particularly including the need for treatment of any symptomatic CAUTI.

There is not enough evidence to suggest whether antibiotics should be used at catheter replacement in patients with long-term catheters. However, not all variants of this intervention or outcomes have been assessed.

There is not enough evidence to suggest whether sterile water is as effective as 0.05% CHG solution for periurethral cleansing between catheter replacements for reducing rates, and delaying onset, of bacteriuria and symptomatic CAUTIs. However, other factors such as patient comfort remain unexplored. A formal economic analysis should be performed to determine whether this is a more cost-effective approach.

Implications for research

Randomised controlled trials with higher numbers of participants and of higher quality are required in order to make recommendations for changing policies for replacing long-term catheters. Further research into the areas studied by the trials featured in this review would help strengthen the evidence to support their conclusions. Different types of antibiotics, different cleaning solutions and different periods between replacements could also be studied in addition to the interventions described in this protocol where no trials were found. Future research should aim to explore additional outcomes to asymptomatic bacteriuria and symptomatic CAUTI such as participant satisfaction and cost-effectiveness. A validated questionnaire should be developed for assessing participant satisfaction and quality of life measures in order to reduce heterogeneity across trials. It is also important to identify which of these outcomes not yet studied are most crucial to decision making regarding policies for replacing long-term catheters in order to guide future research. Sub-group analysis would give valuable data as to whether certain policies are more effective in sub-groups such as females or younger participants. We did not identify any long-term follow-up data. It is paramount that future trials report long-term follow-up data as this is also valuable evidence.

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* *Indicates the major publication for the study*

CHARACTERISTICS OF STUDIES

Characteristics of included studies *[ordered by study ID]*

Cheung 2008

Methods	**Study design:** Randomised Controlled Trial **Setting:** Community nursing service centre in Hong Kong. **Study dates:** Not specified
Participants	**Population:** People residing in one community nurse service centre in Hong Kong (living at home or in a nursing home) **Inclusion criteria:** aged 18 years or older, receiving community nursing services, requiring long-term indwelling latex urinary catheter and already under community nursing service and requiring long-term catheter care for at least 1 month **Exclusion criteria:** subjects with a symptomatic urinary tract infection on the day of baseline urine collection **Age (mean):** 78.4 (SD = 11.8) **Number of participants:** • **Eligible:** 26 • **Randomised:** 22 • **Reported:** 20 **Dropouts (number of participants & reasons):** 4 subjects refused to participate on the day of catheter change. 2 subjects were excluded from data analysis: 1 due to urinary tract infection and 1 due to antibiotic treatment for fever After 7 days, 2 subjects dropped out from the intervention group because of urinary catheter removal as prescribed by a physician and were admitted to the hospital for a respiratory problem. After 14 days, 1 subject dropped out from the control group because of urinary catheter removal as prescribed by physician. 1 subject dropped out from the intervention group because of admission to the hospital for general deterioration in condition
Interventions	**Control group** (12)**:** Conventional 0.05% chlorhexidine gluconate periurethral cleansing **Intervention group** (8)**:** Sterile water periurethral cleansing.
Outcomes	**Symptomaic bacteruria** 14 days after catheter replacement: Control 0/9; Intervention 0/7 **High (>10⁵ cfu/mL) urine culture colonisation count** 0 days after catheter replacement: Control 0/12 ; Intervention 0/8 7 days after catheter replacement: Control 6/10 ; Intervention 6/8 14 days after catheter replacement: Control 8/9 ; Intervention 7/7
Funding/Sponsorship	No information provided.
Notes	Randomisation method referenced as "Simon" with a link that does not work. This link does work: http://www.bmj.com/rapid-response/2011/10/28/simple-approach-randomisation

Risk of bias

Cheung 2008 (*Continued*)

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Low risk	"The simple randomization method suggested by Simon was used to allocate subjects to either the sterile water group or the conventional 0.05% CHG group"
Allocation concealment (selection bias)	Unclear risk	No information given.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Nurses performed cleansing and were therefore not blind to intervention Participants may be able to sense differences between the two cleansing agents and thus determine their intervention
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	"Nurses also followed the standard protocol for urine collection: all urine specimens were collected through a sterile syringe into a sterile bottle to avoid contamination, and the specimens were kept in a cooler and sent to the laboratory within 2 hours" No information regarding detection of CAUTI in participants.
Blinding of microbiological outcome assessment	Low risk	The microbiologists analysing samples are assumed to be blind to the treatment of the participants
Incomplete outcome data (attrition bias) All outcomes	Low risk	All participants who dropped out were accounted for. See **Participants** above.
Selective reporting (reporting bias)	Low risk	All outcomes intended to be investigated are reported. However, the study protocol was not assessed
Other bias	Low risk	Appears to be free of other sources of bias.

Firestein 2001

Methods	**Study design:** Randomised Controlled Trial **Setting:** The Shoham Geriatric Center, Israel. A 970-bed multilevel geriatric centre **Study dates:** November 1998 to August 1999.
Participants	**Population:** Residents of a geriatric medical centre. **Inclusion criteria:** Residents with a long-term urinary catheter. **Exclusion criteria:** Urinary catheter in place for less than 4 weeks or antibiotic use within the 2-week period before enrolment

Firestein 2001 *(Continued)*

	Age (mean): 79.3 +/− 9.6 years. **Number of participants:** **Eligible:** Not specified. **Randomised:** Not specified. **Reported:** 70 (36 control group, 34 intervention group) with various unexplained dropouts at different points of time in outcome measures. **Dropouts (number of participants & reasons):** Not specified.	
Interventions	**Control group** (36)**:** No antibiotic at time of catheter replacement. **Intervention group** (34)**:** 1 g IV Meropenam 30 minutes before catheter replacement.	
Outcomes	**Positive urine culture** 1 to 3 days after replacement: Control 32/35 ; Intervention 27/31 7 days after replacement: Control 35/36 ; Intervention 30/34 14 days after replacement: Control 34/34 ; Intervention 31/33 28 days after replacement: Control 28/28 ; Intervention 30/32 **Infection** Up to 28 days after replacement: Control 6/36 ; Intervention 8/34 **Bacteraemia** Up to 28 days after replacement: Control 0/36 ; Intervention 0/34 **Death** Up to 28 days after replacement: Control 1/36 ; Intervention 2/34	
Funding/Sponsorship	No information provided.	
Notes	The contact person for this study was contacted regarding inconsistencies in reported results	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	No information provided on randomisation techniques.
Allocation concealment (selection bias)	Unclear risk	No information provided on allocation concealment.
Blinding of participants and personnel (performance bias) All outcomes	High risk	No information provided on blinding of participants or personnel. It is assumed there is no blinding as the intervention is invasive
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No information provided.

Firestein 2001 *(Continued)*

Blinding of microbiological outcome assessment	Low risk	The microbiologists analysing samples are assumed to be blind to the treatment of the participants
Incomplete outcome data (attrition bias) All outcomes	High risk	There are unexplained dropouts in each arm of the study at each time period. The number of infections per subject is not reported
Selective reporting (reporting bias)	Low risk	All outcomes intended to be investigated were reported. However, the study protocol was not assessed
Other bias	High risk	A positive urine culture is not defined and therefore the cut-off for a positive outcome may be different from similar studies. There are calculation errors in the results section

Priefer 1982

Methods	**Study design:** Randomised Controlled Trial **Setting:** Veterans Administration hospital-based nursing home in Wisconsin **Study dates:** "Over a 6 month period"
Participants	**Population:** Males in a Veterans Administration hospital-based nursing home with long-term indwelling urethral catheters **Inclusion criteria:** Not specified. **Exclusion criteria:** Transfer to a hospital for acute problems other than those related to the urinary tract **Age (mean):** Control group: 77.1 +/− 16.3 ; Intervention group: 83.4 +/− 7.9 years **Number of participants** • **Eligible:** Not specified • **Randomised:** Not specified • **Reported:** 17 (7 in control group, 10 in intervention group). **Dropouts (number of participants & reasons):** Not specified.
Interventions	**Control group** (7)**:** Catheter replacement only when indicated by infection (as defined in the study) or obstruction **Intervention group** (10)**:** Catheter replacement monthly as well as when indicated by infection (as defined in the study) or obstruction
Outcomes	**Development of symptomatic CAUTI in 6-month period:** Control 6/7 ; Intervention 3/10 (Also expressed as number per subject: Control 1.0 +/− 0.6 ; Intervention 0.4 +/− 0.7) **Number of catheter replacements:** (Expressed as number per subject) Control 0.64 +/− 0.4 ; Intervention 1.3 +/− 0.6 **Number of catheter irrigations:**

Priefer 1982 (*Continued*)

	(Expressed as number per subject) Control 1.5 +/− 2.3 ; Intervention 0.9 +/− 1.7 **Organism found at time of CAUTI:** No difference. Data not available.	
Funding/Sponsorship	Not specified.	
Notes	Inconsistency between results table and results in text for number of irrigations. The results in the text are correct (clarified with the author 27.06.15)	

Risk of bias

Bias	Authors' judgement	Support for judgement
Random sequence generation (selection bias)	Unclear risk	No information provided on randomisation techniques
Allocation concealment (selection bias)	Unclear risk	No information provided on allocation concealment.
Blinding of participants and personnel (performance bias) All outcomes	High risk	Participants and personnel are assumed not to be blinded to intervention
Blinding of outcome assessment (detection bias) All outcomes	Unclear risk	No information given on who was responsible for recording number of irrigations or replacements
Blinding of microbiological outcome assessment	Low risk	The microbiologists analysing samples are assumed to be blind to the treatment of the participants
Incomplete outcome data (attrition bias) All outcomes	Low risk	There were no dropouts and there is no missing data. Some outcomes are expressed as number per subject, however this is appropriate
Selective reporting (reporting bias)	Low risk	All outcomes intended to be investigated were reported. However, the study protocol was not assessed
Other bias	High risk	This trial only studied male participants. There was a wider range in age of participants compared to similar studies

Characteristics of excluded studies *[ordered by study ID]*

Study	Reason for exclusion
Airaksinen 1979	Deals with bladder irrigation and does not address replacement of long-term catheters
Andersson 1986	Does not compare washout to replacement.
Bach 1990	Washout policy. Not related to replacement of long-term catheters
Bergqvist 1979	Compares types of catheters.
Boccola 2011	Not related to replacement of long-term catheters.
Bruun 1978	Compares types of intermittent irrigation. Does not address replacement
Cai 2014	Intervention not relevant. The study is not about the policies for replacing long-term indwelling urinary catheters
Carapeti 1996	Erratum message to authors of RCT which should not be included as it is not about long-term catheters
Chancellor 1994	Not an RCT and does not study catheters.
Christensen 1983	Deals with short-term catheters.
Chung 2007	Intervention is related to first time catheterisation and temporary catheters
Clark 1973	Participants' indwelling catheters are post-operative and not long-term
Classen 1991a	Policy is not related to replacing indwelling catheters in this study
Cleland 1971	Does not address policies for replacing long term catheters.
Cornia 2003	Does not address replacement of long term-catheters.
Darouiche 2005	Does not address replacement of long-term catheters.
Darouiche 2011	Intervention not relevant. The study compares HU2117 versus sterile saline
Darouiche 2014	Studies policies for replacement when indicated by CAUTI, not routine replacement
Davies 1987	Compares washout regimens and does not address replacement of catheter
Eddeland 1983	Participants have long-term indwelling catheters, but intervention (allopurinol vs placebo) is not administered at time of replacement. This study does explore requirement for catheter change as an adverse outcome, however
Ehrenkranz 1991	Not an RCT.

Eid 1995	Does not address catheter replacement.
Flack 1993	Not an RCT.
Fryklund 1991	Not related to replacement of long-term catheters.
Hayward 2012	Not related to replacement of long-term catheters.
Lee 2015	Intervention not relevant.
Li 2014	Not related to replacement of long-term catheters.
Moore 2004	Intervention not relevant. The study is about washouts.
Nalinthip 1996	Study is on intermittent, not long-term catheters.
NCT01785966, 2013	Does not address long-term catheter replacement.
NCT01797146, 2013	Short term catheters.
NCT02196987, 2014	Participants do not have long-term catheters.
Norton 1987	Does not address catheter replacement.
Oberst 1981	Participants' indwelling catheters are post-operative and not long-term
Obolensky 1975	Participants have short-term catheters.
Pickard 1996	Study only deals with new catheter insertion and not replacement
Platt 1983	Particpants' catheters are not long-term and replacement of catheter is not investigated
Raz 2000	Studies policies for replacing long-term catheter for UTI, not routine replacement
Reid 1982	Not an RCT.
Samimi 2010	Deals with bladder washouts and does not address replacement of catheters
Savage 1982	Does not address catheter replacement.
Schneeberger 1992	Only addresses catheter removal.
Shimpuku 2013	Catheters in these subjects are not long-term.
Siderias 2004	Participants catheters' are not long-term.
Sperling 2014	Participants have suprapubic catheters.

Sweet 1985	Intervention not related to replacement of catheter.
Taweesangsuksalul 2005	Intermittent catheters.
Webster 2000	Participants' catheters are only temporary.
Webster 2006	Patients with long term-catheters were excluded from this study
Wilde 2011	Replacement of long-term catheter not studied.
Wilde 2015	Intervention not relevant.
Williamson 1982	Intervention is for removal and not replacement of catheters
Zhao 1994	Catheters in these subjects are not long-term.

D A T A   A N D   A N A L Y S E S

Comparison 1.   Monthly & PRN (as needed) catheter replacement versus PRN (as needed) catheter replacement

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Symptomatic CAUTI (up to 6 months after replacement)	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Comparison 2.   Antibiotics at time of replacement versus no antibiotics at time of replacement

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Positive urine culture	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
1.1 3 days after replacement	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.2 7 days after replacement	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.3 14 days after replacement	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
1.4 28 days after replacement	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
2 Infection	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only
3 Bacteraemia	1	70	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
4 Death	1		Risk Ratio (M-H, Fixed, 95% CI)	Subtotals only

Comparison 3.   Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement

Outcome or subgroup title	No. of studies	No. of participants	Statistical method	Effect size
1 Symptomatic CAUTI (up to 14 days after repacement)	1	16	Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
2 Asymptomatic bacteruria (as defined by positive urine culture)	1		Risk Ratio (M-H, Fixed, 95% CI)	Totals not selected
2.1 0 day after replacement	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.2 7 days after replacement	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]
2.3 14 days after replacement	1		Risk Ratio (M-H, Fixed, 95% CI)	0.0 [0.0, 0.0]

Analysis 1.1. Comparison 1 Monthly & PRN (as needed) catheter replacement versus PRN (as needed) catheter replacement, Outcome 1 Symptomatic CAUTI (up to 6 months after replacement).

Review: Policies for replacing long-term indwelling urinary catheters in adults

Comparison: 1 Monthly % PRN (as needed) catheter replacement versus PRN (as needed) catheter replacement

Outcome: 1 Symptomatic CAUTI (up to 6 months after replacement)

Study or subgroup	Monthly % PRN replacement n/N	PRN replacement n/N	Risk Ratio M-H,Fixed,95% CI	Weight	Risk Ratio M-H,Fixed,95% CI
Priefer 1982	3/10	6/7			0.35 [0.13, 0.95]

Test for subgroup differences: Not applicable

```
              0.01   0.1    1    10    100
        Favours monthly % PRN      Favours PRN
```
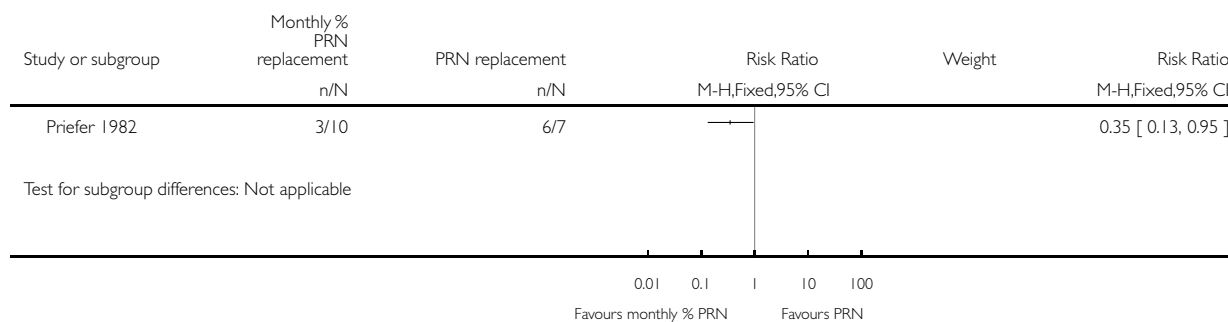
Analysis 2.1. Comparison 2 Antibiotics at time of replacement versus no antibiotics at time of replacement, Outcome 1 Positive urine culture.

Review: Policies for replacing long-term indwelling urinary catheters in adults

Comparison: 2 Antibiotics at time of replacement versus no antibiotics at time of replacement

Outcome: 1 Positive urine culture

Study or subgroup	Meropenem n/N	No antibiotic n/N	Risk Ratio M-H,Fixed,95% CI	Risk Ratio M-H,Fixed,95% CI
1 3 days after replacement				
Firestein 2001	27/31	32/35		0.95 [0.80, 1.13]
2 7 days after replacement				
Firestein 2001	30/34	35/36		0.91 [0.79, 1.04]
3 14 days after replacement				
Firestein 2001	31/33	34/34		0.94 [0.85, 1.04]
4 28 days after replacement				
Firestein 2001	30/32	28/28		0.94 [0.84, 1.05]

```
           0.05   0.2    1    5    20
             meropenem      no antibiotic
```
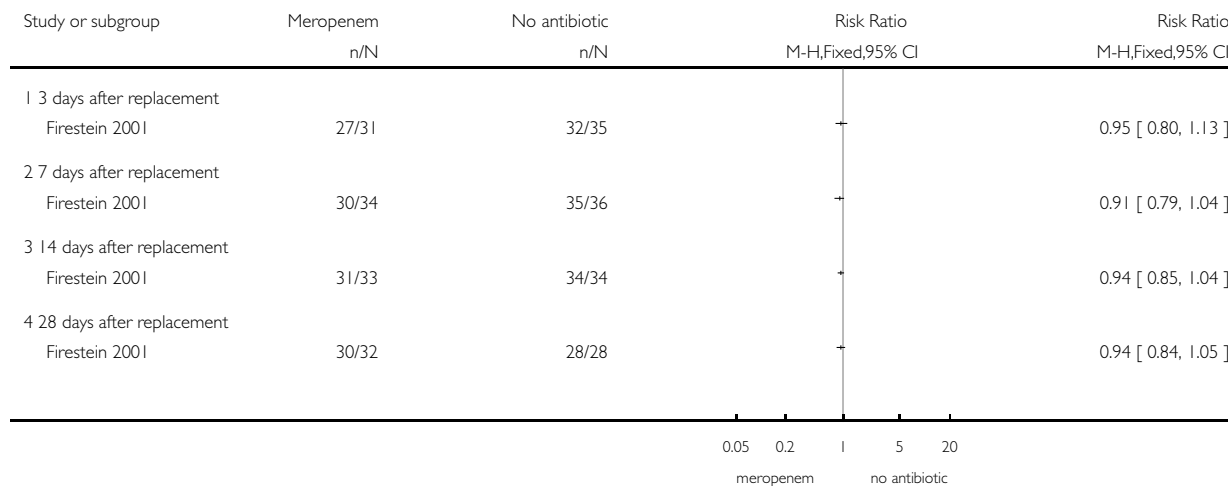
Analysis 2.2. Comparison 2 Antibiotics at time of replacement versus no antibiotics at time of replacement, Outcome 2 Infection.

Review: Policies for replacing long-term indwelling urinary catheters in adults

Comparison: 2 Antibiotics at time of replacement versus no antibiotics at time of replacement

Outcome: 2 Infection

Study or subgroup	Meropenem	No antibiotic	Risk Ratio	Weight	Risk Ratio
	n/N	n/N	M-H,Fixed,95% CI		M-H,Fixed,95% CI
Firestein 2001	8/34	6/36			1.41 [0.55, 3.65]

Test for subgroup differences: Not applicable

```
         0.01   0.1     1     10    100
      Favours meropenem    Favours no antibiotic
```
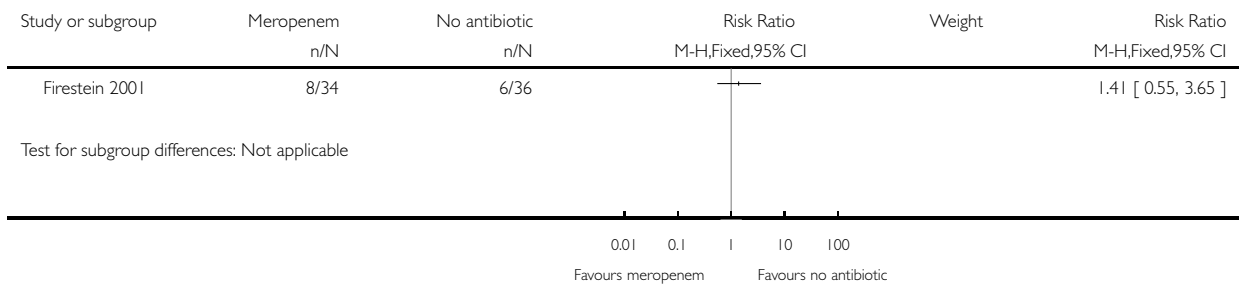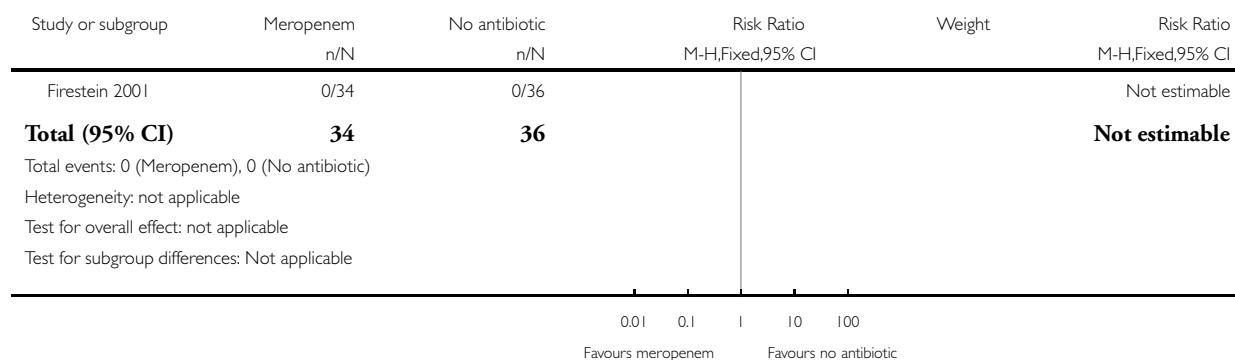
Analysis 2.3. Comparison 2 Antibiotics at time of replacement versus no antibiotics at time of replacement, Outcome 3 Bacteraemia.

Review: Policies for replacing long-term indwelling urinary catheters in adults

Comparison: 2 Antibiotics at time of replacement versus no antibiotics at time of replacement
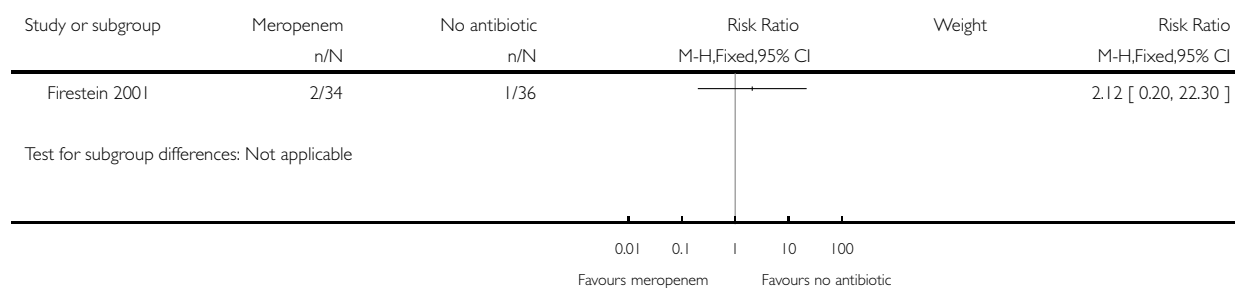
Outcome: 3 Bacteraemia

Study or subgroup	Meropenem n/N	No antibiotic n/N	Risk Ratio M-H,Fixed,95% CI	Weight	Risk Ratio M-H,Fixed,95% CI
Firestein 2001	0/34	0/36			Not estimable
Total (95% CI)	**34**	**36**			**Not estimable**

Total events: 0 (Meropenem), 0 (No antibiotic)

Heterogeneity: not applicable

Test for overall effect: not applicable

Test for subgroup differences: Not applicable

```
        0.01   0.1    1    10   100
      Favours meropenem   Favours no antibiotic
```

Analysis 2.4. Comparison 2 Antibiotics at time of replacement versus no antibiotics at time of replacement, Outcome 4 Death.

Review: Policies for replacing long-term indwelling urinary catheters in adults

Comparison: 2 Antibiotics at time of replacement versus no antibiotics at time of replacement

Outcome: 4 Death

Study or subgroup	Meropenem n/N	No antibiotic n/N	Risk Ratio M-H,Fixed,95% CI	Weight	Risk Ratio M-H,Fixed,95% CI
Firestein 2001	2/34	1/36			2.12 [0.20, 22.30]

Test for subgroup differences: Not applicable

```
        0.01   0.1    1    10   100
      Favours meropenem   Favours no antibiotic
```
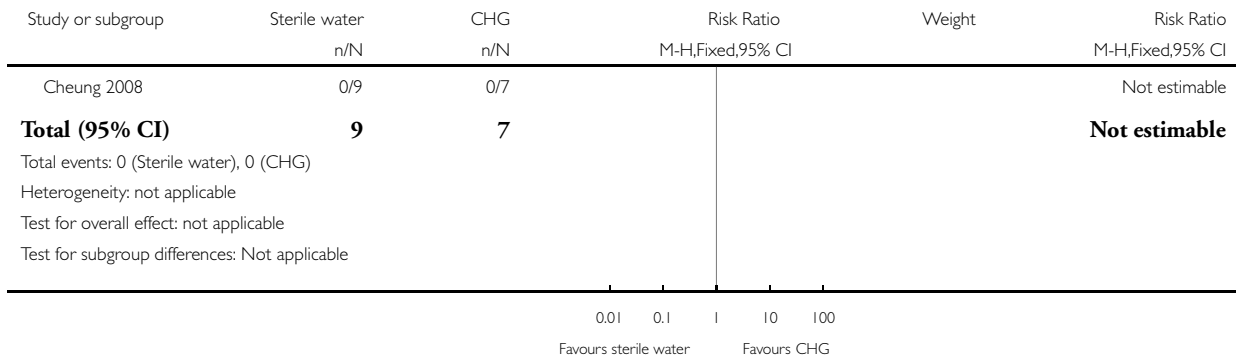
Analysis 3.1. Comparison 3 Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement, Outcome 1 Symptomatic CAUTI (up to 14 days after repacement).

Review: Policies for replacing long-term indwelling urinary catheters in adults

Comparison: 3 Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement

Outcome: 1 Symptomatic CAUTI (up to 14 days after repacement)

Study or subgroup	Sterile water	CHG	Risk Ratio	Weight	Risk Ratio
	n/N	n/N	M-H,Fixed,95% CI		M-H,Fixed,95% CI
Cheung 2008	0/9	0/7			Not estimable
Total (95% CI)	**9**	**7**			**Not estimable**

Total events: 0 (Sterile water), 0 (CHG)
Heterogeneity: not applicable
Test for overall effect: not applicable
Test for subgroup differences: Not applicable

```
        0.01   0.1    1    10   100
      Favours sterile water   Favours CHG
```
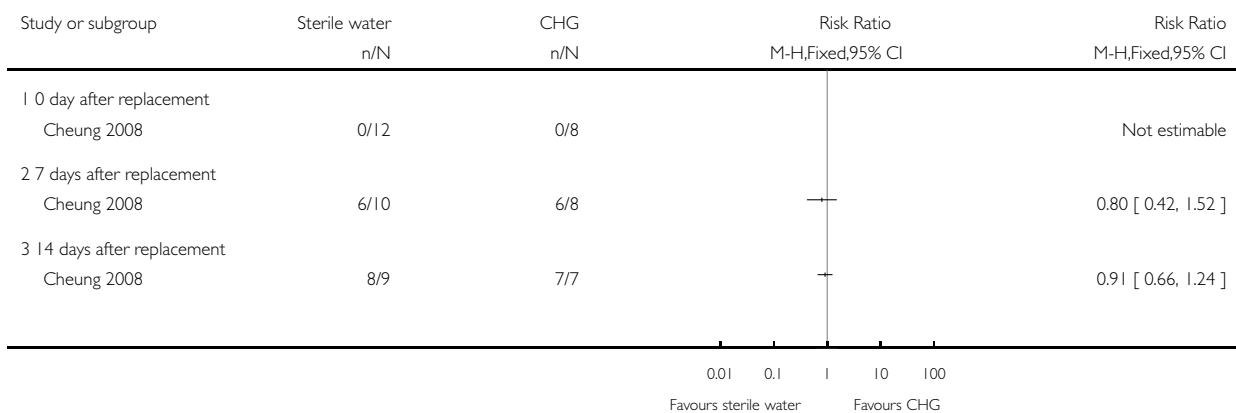
Analysis 3.2. Comparison 3 Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement, Outcome 2 Asymptomatic bacteruria (as defined by positive urine culture).

Review: Policies for replacing long-term indwelling urinary catheters in adults

Comparison: 3 Sterile water versus 0.05% chlorohexidine gluconate for periurethral cleansing during replacement

Outcome: 2 Asymptomatic bacteruria (as defined by positive urine culture)

Study or subgroup	Sterile water	CHG	Risk Ratio	Risk Ratio
	n/N	n/N	M-H,Fixed,95% CI	M-H,Fixed,95% CI
1 0 day after replacement				
Cheung 2008	0/12	0/8		Not estimable
2 7 days after replacement				
Cheung 2008	6/10	6/8		0.80 [0.42, 1.52]
3 14 days after replacement				
Cheung 2008	8/9	7/7		0.91 [0.66, 1.24]

```
        0.01   0.1    1    10   100
      Favours sterile water   Favours CHG
```

APPENDICES

Appendix 1. Cochrane Incontinence Group Specialised Register search strategy

The terms used to search the Incontinence Group Specialised Register are given below:
(({DESIGN.CCT*} OR {DESIGN.RCT*}) AND {INTVENT.MECH.CATH*})
(All searches were of the keyword field of Reference Manager 2012).

CONTRIBUTIONS OF AUTHORS

Fergus PM Cooper (FC), Cameron Edwin Alexander (CEA), Sanjay Sinha (SS), and Muhammad Imran Omar (MIO) were responsible for the conception of the protocol. FC, CEA and SS were responsible for abstract screening. All review authors performed full-text screening, data extraction and analysis. FC and MIO assessed the quality of evidence. MIO also provided methodological expertise. FC took lead in drafting the first version of the review. All review authors contributed in writing the final manuscript.

DECLARATIONS OF INTEREST

Fergus PM Cooper: None known

Cameron Edwin Alexander: None known

Sanjay Sinha was on the Advisory Board for Ranbaxy. The extent of the relationship is very limited both in terms of any monetary compensation as well as in terms of time or responsibility. As such, it does not constitute a conflict of interest for the proposed project.

Muhammad Imran Omar: None known

DIFFERENCES BETWEEN PROTOCOL AND REVIEW

We used "urinary tract infection" as one of the primary outcomes while writing the protocol. However, we realised symptomatic catheter-associated urinary tract infection (CAUTI) is clinically more relevant and important. Therefore, primary outcome was changed to "symptomatic catheter-associated urinary tract infection (CAUTI) as defined by trialists".

The searches of ClinicalTrials.gov and WHO ICTRP are now fully incorporated into the search for the Cochrane Incontinence Specialised Register and were therefore not run separately.