Infection: prevention and control of healthcare-associated infections in primary and community care (partial update of NICE clinical guideline 2)

NICE guideline Draft for consultation, July 2011

If you wish to comment on this version of the guideline, please be aware that all the supporting information and evidence for the 2012 recommendations and the 2003 recommendations that have not been updated is contained in the full version of the 2012 guideline. Evidence considered in 2003 for recommendations that have been updated can be found in the appendix in the full version of the 2012 guideline.

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This guidance is a partial update of NICE clinical guideline 2 (published June 2003) and will replace it.

New recommendations have been added about the prevention and control of healthcare-associated infections in primary and community care.

Where recommendations are shaded in grey and end [2003] the evidence has not been updated since the original guideline. Yellow shading in these recommendations indicates where wording changes have been made for the purposes of clarification only. Recommendations have been labelled [2003, amended 2012] if the evidence has not been reviewed but changes have been made that alter the meaning of the recommendation, such as incorporated guidance being updated or equality issues. These changes are described in appendix C.

You are invited to comment on the new and updated recommendations in this guideline only. These are marked as [2012] if the evidence has been reviewed but no change has been made to the recommendation or [new 2012] if the evidence has been reviewed and the recommendation has been added or updated.

Appendix C contains recommendations from the 2003 guideline that NICE proposes deleting in the 2012 update. This is because the evidence has been reviewed and the recommendation has been updated or because NICE has updated other relevant guidance and has replaced the original recommendations. Where there are replacement recommendations, details are provided. Where there is no replacement recommendation, an explanation for the proposed deletion is given. You are invited to comment on the deleted recommendations as part of the consultation on the 2012 update.

The original NICE guideline and supporting documents are available from www.nice.org.uk/guidance/CG2

Introduction

Clinical context

A wide variety of healthcare is being delivered in primary and community care settings. Healthcare-associated infections arise across a wide range of clinical conditions and can affect patients of all ages. Healthcare workers, family members and carers are also at risk of acquiring an infection as a result of exposure to infections when caring for patients.

Healthcare-associated infections are commonly linked with invasive procedures or devices. For example, indwelling urinary catheters are the most common cause of urinary tract infections, and bloodstream infections are often associated with vascular access devices.

Healthcare-associated infections are caused by a wide range of microorganisms. These are often carried by the patients themselves, and have taken advantage of a route into the body provided by an invasive device or procedure. Healthcare-associated infections can exacerbate existing or underlying conditions, delay recovery and adversely affect quality of life.

Patient safety has become a cornerstone of care, and preventing healthcare-associated infections remains a priority in the patient safety agenda. It is estimated that 300,000 patients a year in England acquire a healthcare-associated infection as a result of care within the NHS. In 2007, meticillin-resistant *Staphylococcus aureus* (MRSA) bloodstream infections and *Clostridium difficile* infections were recorded as the underlying cause of, or a contributory factor in, approximately 9000 deaths in hospital and primary care in England.

Healthcare-associated infections are estimated to cost the NHS approximately £1 billion a year; £56 million of this is estimated to be incurred after the discharge of patients from hospital. In addition to increased costs, each one of these infections means additional use of NHS resources, greater patient discomfort and a decrease in patient safety. A 'no tolerance' attitude is now prevalent in relation to avoidable healthcare-associated infections.

Rationale for the update

Since the publication of the NICE clinical guideline on the prevention of healthcare-associated infection in primary and community care in 2003, many changes have occurred within the NHS that place the patient firmly at the centre of all activities. First, the NHS Constitution for England defines the rights and pledges that every patient can expect regarding their care. To support this, the Care Quality Commission (CQC), the independent regulator of all health and adult social care in England, ensures that health and social care is safe and monitors how providers comply with established standards. In addition, the legal framework that underpins the guidance has changed since 2003.

New guidance is needed to reflect the fact that, as a result of the rapid turnover of patients in acute care settings, complex care is increasingly being delivered in the community. New standards for the care of patients and the management of devices to prevent related healthcare-associated infections are needed that will also reinforce the principles of asepsis.

This clinical guideline is a partial update of 'Infection control: prevention of healthcare-associated infection in primary and community care' (NICE clinical guideline 2; 2003), and addresses areas in which clinical practice for preventing healthcare-associated infections in primary and community care has changed, where the risk of healthcare-associated infections is greatest or where the evidence has changed. Where high-quality evidence is lacking the Guideline Development Group (GDG) has highlighted areas for further research.

Audience

The population covered in this guideline is all adults and children receiving healthcare for which standard infection-control precautions apply in primary and community care. This guideline is commissioned for the NHS, but people providing healthcare in other settings, such as private settings, may find the guidance relevant.

This guideline applies to all healthcare workers employed in primary and community care settings, including ambulance services, and should ensure safe practice if applied consistently. Much care is also delivered by informal carers and family members, and this guideline is equally applicable to them.

Healthcare settings covered by this guideline are:

- Primary care settings, such as general practices, dental clinics, health centres and polyclinics. This also includes care delivered by the ambulance service.
- Community care settings, such as care homes, the patient's own home,
 schools and prisons, where NHS healthcare is provided or commissioned.

Use of 'must' in recommendations

The GDG recognised that there is a legal duty to implement some of the recommendations in this guideline in order to comply with legislation. The word 'must' is used in these recommendations and details of the relevant legislation are given in footnotes to the recommendations.

The GDG was also aware that the consequences of not implementing some other recommendations on patient safety would be very serious – that is, there would be a greatly increased risk of adverse events, including death. The GDG therefore concluded that the use of the word 'must' in these recommendations is justified, in line with the guidance in chapter 9 of 'The guidelines manual (2009)'.

Drugs

The guideline will assume that prescribers will use a drug's summary of product characteristics to inform decisions made with individual patients.

This guideline recommends some drugs for indications for which they do not have a UK marketing authorisation at the date of publication, if there is good evidence to support that use. Where recommendations have been made for the use of drugs outside their licensed indications ('off label use'), these drugs are marked with a footnote in the recommendations.

Patient-centred care

This guideline offers best practice advice on the prevention and control of healthcare-associated infections in primary and community care.

Patients have the right to expect that those who provide their care meet appropriate standards of hygiene and follow the correct procedures to minimise the risk of healthcare-associated infection. Treatment and care should also take into account patients' needs and preferences, and patients should have the opportunity to make informed decisions about their care and treatment, in partnership with their healthcare professionals. If patients do not have the capacity to make decisions, healthcare professionals should follow the Department of Health's advice on consent (available from www.dh.gov.uk/consent) and the code of practice that accompanies the Mental Capacity Act (available from available from www.dh.gov.uk). In Wales, healthcare professionals should follow advice on consent from the Welsh Assembly Government (available from www.wales.nhs.uk/consent).

If the patient is under 16, healthcare professionals should follow the guidelines in 'Seeking consent: working with children' (available from www.dh.gov.uk/consent).

Good communication between healthcare professionals and patients is essential. It should be supported by evidence-based written information tailored to the patient's needs. Treatment and care, and the information patients are given about it, should be culturally appropriate. It should also be accessible to people with additional needs such as physical, sensory or learning disabilities, and to people who do not speak or read English.

If the patient agrees, families and carers should have the opportunity to be involved in decisions about treatment and care.

Families and carers should also be given the information and support they need.

Key priorities for implementation

The following recommendations have been identified as priorities for implementation.

Standard principles: general recommendations

- Everyone involved in providing care should be:
 - educated about the standard principles of infection prevention and control and
 - trained in hand decontamination, the use of personal protective equipment and the safe use and disposal of sharps. [2012] [1.1.1.1]
- Wherever care is delivered, healthcare workers must have available appropriate supplies of:
 - materials for hand decontamination
 - sharps containers
 - personal protective equipment. [new 2012] [1.1.1.2]
- Educate patients and carers about:
 - the benefits of effective hand hygiene
 - the correct techniques and timing of hand decontamination
 - when it is appropriate to use liquid soap and water or handrub
 - the availability of hand decontamination facilities
 - their role in maintaining standards of healthcare workers' hand hygiene. [new 2012] [1.1.1.3]

¹ In accordance with current health and safety legislation (at the time of consultation on the quideline [July 2011]): Health and Safety at Work Act 1974, Management of Health and Safety at Work Regulations 1999, Health and Safety Regulations 2002, Control of Substances Harmful to Health Regulations 2002, Personal Protective Equipment Regulations 2002, and Health and Social Care Act 2008.

Standard principles for hand hygiene

- Hands must be decontaminated in all of the following circumstances:
 - immediately before every episode of direct patient contact or care
 - immediately after every episode of direct patient contact or care
 - immediately after any exposure to body fluids
 - immediately after any other activity or contact with a patient's surroundings that could potentially result in hands becoming contaminated
 - immediately after removal of gloves. [new 2012] [1.1.2.1]

Long-term urinary catheters

- Select the type and gauge of an indwelling urinary catheter based on an assessment of the patient's individual characteristics, including:
 - age
 - any allergy or sensitivity to catheter materials
 - gender
 - history of symptomatic urinary tract infection
 - patient preference and comfort
 - previous catheter history
 - reason for catheterisation. [new 2012] [1.2.3.3]
- Offer non-coated intermittent catheters for multiple use² to patients³ except in the following circumstances, when a choice of single-use hydrophilic or gel reservoir catheters should be offered:
 - if the patient is unable to wash and dry catheters
 - if no suitable facilities to wash, dry and store catheters are readily available
 - if catheterisation is performed by a healthcare worker or anyone else other than the patient or a close family member. [new 2012] [1.2.3.4]

² Refer to the manufacturer's instructions for advice on the use, cleaning and storage of catheters. See also recommendation 1.2.5.13 about the cleaning of reusable intermittent

catheters. ³ Do not offer multiple-use catheters for use in children or young people of 16 years or under (see recommendation 1.2.3.5).

- All catheterisations carried out by healthcare workers should be aseptic procedures. After training, healthcare workers should be assessed for their competence to carry out these types of procedures. [2003] [1.2.4.1]
- When changing catheters in patients with a long-term indwelling urinary catheter:
 - do not offer antibiotic prophylaxis routinely
 - consider antibiotic prophylaxis⁴ for patients who:
 - have a history of symptomatic urinary tract infection after catheter change or
 - ♦ experience trauma⁵ during catheterisation. [new 2012] [1.2.5.12]

Vascular access devices

- Healthcare workers caring for a patient with a vascular access device⁶ should be trained, and assessed as competent, in using and consistently adhering to the infection prevention practices described in this guideline. [2003, amended 2012] [1.4.1.2]
- Decontaminate the skin at the insertion site with chlorhexidine gluconate in 70% alcohol before inserting a peripheral vascular access device or a peripherally inserted central catheter. [new 2012] [1.4.3.1]

attempts of catheterisation.

⁴ At the time of consultation on the guideline (July 2011), no antibiotics have a UK marketing authorisation for this indication. Informed consent should be obtained and documented. The GDG defined trauma as frank haematuria following catheterisation or two or more

⁶ The updated recommendation contains 'vascular access device' rather than 'central venous catheter'. This change has been made because peripherally inserted catheters were included in the scope of the guideline update.

1 Guidance

The following guidance is based on the best available evidence. The full guideline ([hyperlink to be added for final publication]) gives details of the methods and the evidence used to develop the guidance.

Terms used in this guidance

Direct patient care 'Hands on' or face-to-face contact with patients. Any physical aspect of the healthcare of a patient, including treatments, self-care and administration of medication.

Hand decontamination The physical removal of blood, body fluids and transient microorganisms from the hands, for example by handwashing or using a handrub.

Handrub A preparation designed for application to the hands for reducing the number of viable microorganisms. This guideline refers to handrubs compliant with British standards (BS EN1500; standard for efficacy of hygienic handrubs using a reference of 60% isopropyl alcohol).

Healthcare worker Any person employed by the health service, social services, a local authority or an agency to provide care for a sick, disabled or elderly person.

Healthcare waste In this guideline, healthcare waste refers to any waste produced by, and as a consequence of, healthcare activities.

Personal protective equipment Equipment that is intended to be worn or held by a person to protect them from risks to their health and safety while at work. Examples include gloves, aprons and eye and face protection.

1.1 Standard principles

1.1.1 General recommendations

- 1.1.1.1 Everyone involved in providing care should be:
 - educated about the standard principles of infection prevention and control and
 - trained in hand decontamination, the use of personal protective equipment and the safe use and disposal of sharps. [2012]
- 1.1.1.2 Wherever care is delivered, healthcare workers must⁷ have available appropriate supplies of:
 - materials for hand decontamination
 - sharps containers
 - personal protective equipment. [new 2012]
- 1.1.1.3 Educate patients and carers about:

2002, and Health and Social Care Act 2008.

- the benefits of effective hand hygiene
- the correct techniques and timing of hand decontamination
- when it is appropriate to use liquid soap and water or handrub
- the availability of hand decontamination facilities
- their role in maintaining standards of healthcare workers' hand hygiene. [new 2012]

⁷ In accordance with current health and safety legislation (at the time of consultation on the guideline [July 2011]): Health and Safety at Work Act 1974, Management of Health and Safety at Work Regulations 1999, Health and Safety Regulations 2002, Control of Substances Harmful to Health Regulations 2002, Personal Protective Equipment Regulations

1.1.2 Hand hygiene

- 1.1.2.1 Hands must be decontaminated in all of the following circumstances:
 - immediately before every episode of direct patient contact or care
 - immediately after every episode of direct patient contact or care
 - immediately after any exposure to body fluids
 - immediately after any other activity or contact with a patient's surroundings that could potentially result in hands becoming contaminated
 - immediately after removal of gloves. [new 2012]
- 1.1.2.2 Decontaminate hands preferably with a handrub (conforming to current British standards⁸), except in the following circumstances, when liquid soap and water must be used:
 - when hands are visibly soiled or potentially contaminated with body fluids or
 - in clinical situations where there is potential for the spread of alcohol-resistant organisms (such as norovirus, Clostridium difficile or organisms that cause diarrhoeal illness). [new 2012]
- 1.1.2.3 Healthcare workers should ensure that their hands can be decontaminated throughout the duration of clinical work by:
 - being bare below the elbow⁹
 - removing wrist and hand jewellery
 - making sure that fingernails are short, clean and free of nail polish
 - covering cuts and abrasions with waterproof dressings. [new 2012]

⁸ At the time of consultation on the guideline (July 2011): BS EN 1500: 1997.

⁹ For the purposes of this guideline, the GDG considered bare below the elbow to mean: not wearing false nails or nail polish; not wearing a wrist-watch or stoned rings; wearing shortsleeved garments or being able to roll or push up sleeves when delivering direct patient care and performing hand hygiene.

- 1.1.2.4 An effective handwashing technique involves three stages: preparation, washing and rinsing, and drying. Preparation requires wetting hands under tepid running water **before** applying liquid soap or an antimicrobial preparation. The handwash solution must come into contact with **all** of the surfaces of the hand. The hands must be **rubbed** together vigorously for a minimum of 10–15 seconds, paying particular attention to the tips of the fingers, the thumbs and the areas between the fingers. Hands should be rinsed thoroughly before drying with good quality paper towels. [2003]
- 1.1.2.5 When decontaminating hands using an alcohol handrub, hands should be free from dirt and organic material. The handrub solution must come into contact with all surfaces of the hand. The hands must be **rubbed** together vigorously, paying particular attention to the tips of the fingers, the thumbs and the areas between the fingers, until the solution has evaporated and the hands are dry.
 [2003]
- 1.1.2.6 An emollient hand cream should be applied regularly to protect skin from the drying effects of regular hand decontamination. If a particular soap, antimicrobial hand wash or alcohol product causes skin irritation an occupational health team should be consulted.
 [2003]

1.1.3 Use of personal protective equipment

1.1.3.1 Selection of protective equipment must¹⁰ be based on an assessment of the risk of transmission of microorganisms to the patient, and the risk of contamination of the healthcare worker's clothing and skin by patients' blood, body fluids, secretions or excretions. [2003]

¹⁰ In accordance with current health and safety legislation (at the time of consultation on the guideline [July 2011]): Health and Safety at Work Act 1974, Management of Health and Safety at Work Regulations 1999, Health and Safety Regulations 2002, Control of Substances Harmful to Health Regulations 2002, Personal Protective Equipment Regulations 2002 and Health and Social Care Act 2008.

- 1.1.3.2 Gloves used for direct patient care must¹¹ conform to current European Community (CE) standards¹² and should be appropriate for the task. [new 2012]
- 1.1.3.3 Gloves must¹¹ be worn for invasive procedures, contact with sterile sites and non-intact skin or mucous membranes, and all activities that have been assessed as carrying a risk of exposure to blood, body fluids, secretions or excretions, or to sharp or contaminated instruments. [2003]
- 1.1.3.4 Gloves must¹¹ be worn as single-use items. They must be put on immediately before an episode of patient contact or treatment and removed as soon as the activity is completed. Gloves must be changed between caring for different patients, and between different care or treatment activities for the same patient. [2003]
- 1.1.3.5 Gloves that have been used for direct patient care or exposed to body fluids must be disposed of as clinical waste in accordance with current national legislation¹³ or local policies (see section 1.1.5). [new 2012]
- 1.1.3.6 Alternatives to natural rubber latex gloves must¹¹ be available for patients, carers and healthcare workers who have a documented sensitivity to natural rubber latex. **[2012]**
- 1.1.3.7 Do not use polythene gloves for clinical interventions. [new 2012]

¹¹ In accordance with current health and safety legislation (at the time of consultation on the guideline [July 2011]): Health and Safety at Work Act 1974, Management of Health and Safety at Work Regulations 1999, Health and Safety Regulations 2002, Control of Substances Harmful to Health Regulations 2002, Personal Protective Equipment Regulations 2002 and Health and Social Care Act 2008.

At the time of consultation on the guideline (July 2011): BS EN 455-2: 2009.
 For guidance see (at the time of consultation on the guideline [July 2011]): 'Safe management of healthcare waste' (2011); available from https://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH
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- 1.1.3.8 When delivering direct patient care:
 - wear a disposable plastic apron if there is a risk that clothing may be exposed to blood, body fluids, excretions or secretions
 - wear a full-body fluid-repellent gown if there is a risk of extensive splashing of blood, body fluids, excretions or secretions onto skin or clothing. [2012]
- 1.1.3.9 When using disposable plastic aprons or gowns:
 - use them as single-use items, for one procedure or one episode of direct patient care and
 - ensure they are disposed of correctly (see section 1.1.5). [2012]
- Face masks and eye protection must¹⁴ be worn where there is a 1.1.3.10 risk of blood, body fluids, secretions or excretions splashing into the face and eyes. [2003]
- 1.1.3.11 Respiratory protective equipment, for example a particulate filter mask, must¹⁴ be used when clinically indicated. [2003]

1.1.4 Safe use and disposal of sharps

- 1.1.4.1 Sharps should 15 not be passed directly from hand to hand, and handling should be kept to a minimum. [2003, amended 2012]
- 1.1.4.2 Used needles should not be recapped, bent or broken before disposal. If recapping or disassembly is unavoidable:
 - a risk assessment should be undertaken and
 - appropriate safety devices should be used. [new 2012]

¹⁴ In accordance with current health and safety legislation (at the time of consultation on the quideline [July 2011]): Health and Safety at Work Act 1974, Management of Health and Safety at Work Regulations 1999, Health and Safety Regulations 2002, Control of Substances Harmful to Health Regulations 2002, Personal Protective Equipment Regulations 2002 and Health and Social Care Act 2008.

¹⁵ The updated recommendation contains 'should' rather than 'must' (which is in the 2003 guideline) because the GDG considered that this is not covered by legislation (in accordance with the NICE guidelines manual, 2009).

1.1.4.3 Used sharps must be discarded immediately by the person generating the sharps waste into a sharps container conforming to current standards¹⁶. [new 2012]

1.1.4.4 Sharps containers:

- must¹⁷ be located in a safe position that avoids spillage, is at a height that allows the safe disposal of sharps, is away from public access areas and is out of the reach of children
- must not¹⁷ be used for any other purpose than the disposal of sharps
- must not 17 be filled above the fill line
- must¹⁷ be disposed of when the fill line is reached
- should be temporarily closed when not in use
- should be disposed of every 3 months even if not full, by the licensed route in accordance with local policy. [new 2012]
- 1.1.4.5 Use sharps safety devices if a risk assessment has indicated that they will provide safer systems of working for healthcare workers, carers and patients. [new 2012]
- 1.1.4.6 Train and assess all users in the correct use and disposal of sharps. [new 2012]

1.1.5 Waste disposal

1.1.5.1 Healthcare waste must be segregated immediately by the person generating the waste into colour-coded storage bags or containers, as defined by current national legislation¹⁷ and local policies.

[new 2012]

<u>www.dn.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuldance/DH</u>
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At the time of consultation on the guideline (July 2011): UN3291 and BS 7320.
 For guidance see (at the time of consultation on the guideline [July 2011]): 'Safe management of healthcare waste' (2011); available from www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH

- 1.1.5.2 Healthcare waste must be labelled, stored, transported and disposed of in accordance with current national legislation¹⁸ and local policies. [new 2012]
- 1.1.5.3 Educate patients and carers about the correct handling, storage and disposal of healthcare waste. [new 2012]

1.2 Long-term urinary catheters

1.2.1 Education of patients, their carers and healthcare workers

- 1.2.1.1 Patients and carers should be educated about and trained in techniques of hand decontamination, insertion of intermittent catheters where applicable, and catheter management before discharge from hospital. [2003]
- 1.2.1.2 Community and primary healthcare workers must be trained in catheter insertion, including suprapubic catheter replacement and catheter maintenance. [2003]
- 1.2.1.3 Follow-up training and ongoing support of patients and carers should be available for the duration of long-term catheterisation.[2003]

1.2.2 Assessing the need for catherisation

- 1.2.2.1 Indwelling urinary catheters should be used only after alternative methods of management have been considered. [2003]
- 1.2.2.2 The patient's clinical need for catheterisation should be reviewed regularly and the urinary catheter removed as soon as possible.

 [2003]

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¹⁸ For guidance see (at the time of consultation on the guideline [July 2011]): 'Safe management of healthcare waste' (2011); available from https://www.dh.gov.uk/en/Publicationsandstatistics/Publications/PublicationsPolicyAndGuidance/DH_126345

1.2.2.3 Catheter insertion, changes and care should be documented.[2003]

1.2.3 Catheter drainage options

- 1.2.3.1 Following assessment, the best approach to catheterisation that takes account of clinical need, anticipated duration of catheterisation, patient preference and risk of infection should be selected. [2003]
- 1.2.3.2 Intermittent catheterisation should be used in preference to an indwelling catheter if it is clinically appropriate and a practical option for the patient. [2003]
- 1.2.3.3 Select the type and gauge of an indwelling urinary catheter based on an assessment of the patient's individual characteristics, including:
 - age
 - any allergy or sensitivity to catheter materials
 - gender
 - history of symptomatic urinary tract infection
 - patient preference and comfort
 - previous catheter history
 - reason for catheterisation. [new 2012]

- 1.2.3.4 Offer non-coated intermittent catheters for multiple use¹⁹ to patients²⁰ except in the following circumstances, when a choice of single-use hydrophilic or gel reservoir catheters should be offered:
 - if the patient is unable to wash and dry catheters
 - if no suitable facilities to wash, dry and store catheters are readily available
 - if catheterisation is performed by a healthcare worker or anyone else other than the patient or a close family member. [new 2012]
- 1.2.3.5 Do not offer multiple-use catheters for use in children or young people of 16 years or under. [new 2012]
- 1.2.3.6 In general, the catheter balloon should be inflated with 10 ml of sterile water in adults and 3–5 ml in children. [2003]
- 1.2.3.7 In patients for whom it is appropriate, a catheter valve may be used as an alternative to a drainage bag. [2003]

1.2.4 Catheter insertion

- 1.2.4.1 All catheterisations carried out by healthcare workers should be aseptic procedures. After training, healthcare workers should be assessed for their competence to carry out these types of procedures. [2003]
- 1.2.4.2 Intermittent self-catheterisation is a clean procedure. A lubricant for single-patient use is required for non-lubricated catheters. [2003]
- 1.2.4.3 For urethral catheterisation, the meatus should be cleaned before insertion of the catheter, in accordance with local guidelines/policy.[2003]

¹⁹ Refer to the manufacturer's instructions for advice on the use, cleaning and storage of catheters. See also recommendation 1.2.5.13 about the cleaning of reusable intermittent catheters

catheters.

²⁰ Do not offer multiple-use catheters for use in children or young people of 16 years or under (see recommendation 1.2.3.5).

1.2.4.4 An appropriate lubricant from a single-use container should be used during catheter insertion to minimise urethral trauma and infection. [2003]

1.2.5 Catheter maintenance

- 1.2.5.1 Indwelling catheters should be connected to a sterile closed urinary drainage system or catheter valve. [2003]
- 1.2.5.2 Healthcare workers should ensure that the connection between the catheter and the urinary drainage system is not broken except for good clinical reasons, (for example changing the bag in line with the manufacturer's recommendations). [2003]
- 1.2.5.3 Healthcare workers must decontaminate their hands and wear a new pair of clean, non-sterile gloves before manipulating a patient's catheter, and must decontaminate their hands after removing gloves. [2003]
- 1.2.5.4 Patients managing their own catheters, and their carers, must be educated about the need for hand decontamination²¹ before and after manipulation of the catheter, in accordance with the recommendations in the standard principles section (section 1.1). [2003, amended 2012]
- 1.2.5.5 Urine samples must be obtained from a sampling port using an aseptic technique. [2003]
- 1.2.5.6 Urinary drainage bags should be positioned below the level of the bladder, and should not be in contact with the floor. [2003]
- 1.2.5.7 A link system should be used to facilitate overnight drainage, to keep the original system intact. **[2003]**

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²¹ The highlighted text replaces 'Carers and patients managing their own catheters must wash their hands...' in the 2003 guideline. This has been amended to reflect input from the NICE Patient and Public Involvement Programme: recommendations cannot be made directly about what patients and carers must do.

- 1.2.5.8 The urinary drainage bag should be emptied frequently enough to maintain urine flow and prevent reflux, and should be changed when clinically indicated. [2003]
- 1.2.5.9 The meatus should be washed daily with soap and water. [2003]
- 1.2.5.10 To minimise the risk of blockages, encrustations and catheterassociated infections for patients with a long-term indwelling urinary catheter:
 - develop a patient-specific care regimen
 - do not use bladder instillations or washouts
 - consider approaches such as reviewing the frequency of planned catheter changes and increasing fluid intake
 - document catheter blockages. [new 2012]
- 1.2.5.11 Catheters should be changed only when clinically necessary or according to the manufacturer's current recommendations. [2003]
- 1.2.5.12 When changing catheters in patients with a long-term indwelling urinary catheter:
 - do not offer antibiotic prophylaxis routinely
 - consider antibiotic prophylaxis²² for patients who:
 - have a history of symptomatic urinary tract infection after catheter change or
 - experience trauma²³ during catheterisation. [new 2012]
- 1.2.5.13 Reusable intermittent catheters should be cleaned with water and stored dry in accordance with the manufacturer's instructions. [2003]

of catheterisation.

²² At the time of consultation on the guideline (July 2011), no antibiotics have a UK marketing authorisation for this indication. Informed consent should be obtained and documented. The GDG defined trauma as frank haematuria after catheterisation or two or more attempts

1.3 Care during enteral feeding

1.3.1 Education of patients, their carers and healthcare workers

- 1.3.1.1 Patients and carers should be educated about and trained in the techniques of hand decontamination, enteral feeding and the management of the administration system before being discharged from hospital. [2003]
- 1.3.1.2 Healthcare workers should be trained in enteral feeding and management of the administration system. [2003]
- 1.3.1.3 Follow-up training and ongoing support of patients and carers should be available for the duration of home enteral tube feeding.[2003]

1.3.2 Preparation and storage of feeds

- 1.3.2.1 Wherever possible pre-packaged, ready-to-use feeds should be used in preference to feeds requiring decanting, reconstitution or dilution. [2003]
- 1.3.2.2 The system selected should require minimal handling to assemble, and be compatible with the patient's enteral feeding tube. [2003]
- 1.3.2.3 Effective hand decontamination must be carried out before starting feed preparation. [2003]
- 1.3.2.4 When decanting, reconstituting or diluting feeds, a clean working area should be prepared and equipment dedicated for enteral feed use only should be used. [2003]
- 1.3.2.5 Feeds should be mixed using cooled boiled water or freshly opened sterile water and a no-touch technique. [2003]
- 1.3.2.6 Feeds should be stored according to the manufacturer's instructions and, where applicable, food hygiene legislation. [2003]

1.3.2.7 Where ready-to-use feeds are not available, feeds may be prepared in advance, stored in a refrigerator, and used within 24 hours. [2003]

1.3.3 Administration of feeds

- 1.3.3.1 Use minimal handling and a no-touch technique to connect the administration system to the enteral feeding tube. [new 2012]
- 1.3.3.2 Ready-to-use feeds may be given for a whole administration session, up to a maximum of 24 hours. Reconstituted feeds should be administered over a maximum 4-hour period. [2003]
- 1.3.3.3 Administration sets and feed containers are for single use and must be discarded after each feeding session. [2003]

1.3.4 Care of insertion site and enteral feeding tube

- 1.3.4.1 The stoma should be washed daily with water and dried thoroughly.

 [2003]
- 1.3.4.2 To prevent blockages, flush the enteral feeding tube before and after feeding or administering medications using single-use syringes or single-patient-use (reusable) syringes according to the manufacturer's instructions. Use:
 - fresh tap water for patients who are not immunosuppressed
 - either cooled freshly boiled water or sterile water from a freshly opened container for patients who are immunosuppressed.
 [new 2012]

1.4 Vascular access devices

1.4.1 Education of patients, their carers and healthcare workers

- 1.4.1.1 Before discharge from hospital, patients and their carers should be taught any techniques they may need to use to prevent infection and safely manage a vascular access device²⁴. [2003, amended 2012]
- 1.4.1.2 Healthcare workers caring for a patient with a vascular access device²⁴ should be trained, and assessed as competent, in using and consistently adhering to the infection prevention practices described in this guideline. [2003, amended 2012]
- 1.4.1.3 Follow-up training and support should be available to patients with a vascular access device²⁴ and their carers. [2003, amended 2012]

1.4.2 General asepsis

- 1.4.2.1 Hands must be decontaminated (see section □) before accessing or dressing a vascular access device. [new 2012]
- 1.4.2.2 An aseptic technique, such as Aseptic Non Touch Technique (ANTT), must be used for vascular access device catheter site care and when accessing the system. [new 2012]

1.4.3 Vascular access device site care

1.4.3.1 Decontaminate the skin at the insertion site with chlorhexidine gluconate in 70% alcohol before inserting a peripheral vascular access device or a peripherally inserted central catheter. [new 2012]

1.4.3.2 Use a sterile transparent semipermeable membrane dressing to cover the vascular access device insertion site. [new 2012]

²⁴ The updated recommendation contains 'vascular access device' rather than 'central venous catheter'. This change has been made because peripherally inserted catheters were included in the scope of the guideline update.

- 1.4.3.3 Only consider a sterile gauze dressing covered with a sterile transparent semipermeable membrane dressing if the patient has profuse perspiration, or if the vascular access device insertion site is bleeding or oozing. If a gauze dressing is used:
 - change it every 24 hours, or sooner if it is soiled and
 - replace it with a sterile transparent semipermeable membrane dressing as soon as possible. [new 2012]
- 1.4.3.4 Change the transparent semipermeable membrane dressing covering a central venous access device insertion site every
 7 days, or sooner if the dressing is no longer intact or moisture collects under it. [2012]
- 1.4.3.5 Leave the transparent semipermeable membrane dressing applied to a peripheral cannula insertion site in situ for the life of the cannula, provided that the integrity of the dressing is retained. [new 2012]
- 1.4.3.6 Dressings used on tunnelled or implanted central venous catheter sites should be replaced every 7 days until the insertion site has healed, unless there is an indication to change them sooner. [2003]
- 1.4.3.7 Healthcare workers should ensure that catheter-site care is compatible with catheter materials (tubing, hubs, injection ports, luer connectors and extensions) and carefully check compatibility with the manufacturer's recommendations. [2003]
- 1.4.3.8 Decontaminate the central venous catheter insertion site and surrounding skin during dressing changes using chlorhexidine gluconate in 70% alcohol, and allow to air dry. Consider using an aqueous solution of chlorhexidine gluconate if the manufacturer's recommendations prohibit the use of alcohol with their catheter.
 [2012]

1.4.3.9 Individual sachets of antiseptic solution or individual packages of antiseptic-impregnated swabs or wipes should be used to disinfect the dressing site. [2003]

1.4.4 General principles for catheter management

- 1.4.4.1 Decontaminate the injection port or vascular access device catheter hub before and after accessing the system using chlorhexidine gluconate in 70% alcohol. Consider using an aqueous solution of chlorhexidine gluconate if the manufacturer's recommendations prohibit the use of alcohol with their catheter. [new 2012]
- 1.4.4.2 In-line filters should not be used routinely for infection prevention.

 [2003]
- 1.4.4.3 Antibiotic lock solutions should not be used routinely to prevent catheter-related bloodstream infections (CRBSI). [2003]
- 1.4.4.4 Systemic antimicrobial prophylaxis should not be used routinely to prevent catheter colonisation or CRBSI, either before insertion or during the use of a central venous catheter. [2003]
- 1.4.4.5 Preferably, a single lumen catheter should be used to administer parenteral nutrition. If a multilumen catheter is used, one port must be exclusively dedicated for total parenteral nutrition, and all lumens must be handled with the same meticulous attention to aseptic technique. [2003]
- 1.4.4.6 Preferably, a sterile 0.9 percent sodium chloride injection should be used to flush and lock catheter lumens. **[2003]**
- 1.4.4.7 When recommended by the manufacturer, implanted ports or opened-ended catheter lumens should be flushed and locked with heparin sodium flush solutions. [2003]
- 1.4.4.8 Systemic anticoagulants should not be used routinely to prevent CRBSI. [2003]

- 1.4.4.9 If needleless devices are used, the manufacturer's recommendations for changing the needleless components should be followed. [2003]
- 1.4.4.10 When needleless devices are used, healthcare workers should ensure that all components of the system are compatible and secured, to minimise leaks and breaks in the system. [2003]
- 1.4.4.11 When needleless devices are used, the risk of contamination should be minimised by decontaminating the access port with either alcohol or an alcoholic solution of chlorhexidine gluconate before and after using it to access the system. [2003]
- 1.4.4.12 In general, administration sets in continuous use need not be replaced more frequently than at 72-hour intervals unless they become disconnected or a catheter-related infection is suspected or documented. [2003]
- 1.4.4.13 Administration sets for blood and blood components should be changed every 12 hours, or according to the manufacturer's recommendations. [2003]
- 1.4.4.14 Administration sets used for total parenteral nutrition infusions should generally be changed every 24 hours. If the solution contains only glucose and amino acids, administration sets in continuous use do not need to be replaced more frequently than every 72 hours. [2003]
- 1.4.4.15 Avoid the use of multidose vials, in order to prevent the contamination of infusates. [new 2012]

2 Notes on the scope of the guidance

NICE guidelines are developed in accordance with a scope that defines what the guideline will and will not cover. The scope of this guideline is available from http://guidance.nice.org.uk/CG/WaveR/85/Scope

How this guideline was developed

NICE commissioned the National Clinical Guideline Centre to develop this guideline. The Centre established a Guideline Development Group (see appendix A), which reviewed the evidence and developed the recommendations. An independent Guideline Review Panel oversaw the development of the guideline (see appendix B).

There is more information about how NICE clinical guidelines are developed on the NICE website (www.nice.org.uk/HowWeWork). A booklet, 'How NICE clinical guidelines are developed: an overview for stakeholders, the public and the NHS' (fourth edition, published 2009), is available from NICE publications (phone 0845 003 7783 or email publications@nice.org.uk and quote reference N1739).

3 Implementation

NICE has developed tools to help organisations implement this guidance (see www.nice.org.uk/guidance/CG)'.

4 Research recommendations

The Guideline Development Group has made the following recommendations for research, based on its review of evidence, to improve NICE guidance and patient care in the future. The Guideline Development Group's full set of research recommendations is detailed in the full guideline (see section 5).

4.1 Standard principles of infection prevention and control

What are the barriers to compliance with the standard principles of infection prevention and control that patients and carers experience in their own homes?

Why this is important

Recent changes to the delivery of healthcare mean that care is increasingly delivered within a patient's home environment. Infection prevention in this setting is just as important as in hospital. There are currently approximately 6 million unpaid carers in the UK, a number that is likely to increase with an aging population. The association between carer training and infection rates is unknown. Current mandatory surveillance of healthcare-associated infections indicates that the majority are attributable to areas other than acute health provision. Currently, no evidence from the UK exists in this area.

A qualitative study is needed to investigate the themes surrounding the barriers to patient and carer compliance with the standard principles of infection prevention in their own homes. It would be important to assess whether lack of awareness or knowledge is a barrier. If patients and carers have received education, this should be assessed to see if this was applicable to the patient's home setting. Areas of low compliance in the home environment need to be identified. The findings could have far-reaching implications for discharge planning and duty of care.

4.2 Hand hygiene

When clean running water is not available, what is the clinical and cost effectiveness of using wipes, gels, handrubs or other products to remove visible contamination?

Why is this important?

Community healthcare workers often encounter challenges in maintaining hand hygiene when there is no access to running water. This particularly affects ambulance service staff, who often provide emergency care at locations where running water is not available. No evidence from randomised controlled trials is available on the most effective way for community-based healthcare workers to remove physical contamination, such as blood, from their hands in the absence of running water. In recent years, hand hygiene products that can be used without running water, such as gels, handrubs and wipes, have become available. However, their efficacy and suitability in actual clinical practice for use with visibly dirty hands has not been determined. A

randomised controlled trial is required to compare hand wipes (detergent and disinfectant), hand gels and other hand hygiene products that can be used without running water, to determine the most effective way to remove physical dirt in the absence of running water, in order to make a recommendation for their use in real situations. The primary outcome measure should be colony-forming units on the basis of the adenosine triphosphate (ATP) surface test.

4.3 Indwelling urinary catheters: catheter selection

For patients using long-term indwelling urinary catheters, what is the clinical and cost effectiveness of impregnated versus hydrophilic versus silicone catheters in reducing symptomatic urinary tract infections, encrustations and/or blockages?

Why is this important?

Long-term indwelling catheters are commonly used in both hospital and community care settings. Long-term catheterisation carries a significant risk of symptomatic urinary tract infection, which can lead to more serious complications. Several different types of impregnated and hydrophilic long-term indwelling catheters on the market claim to be more effective than non-coated catheters, but are also more expensive.

The clinical evidence review for the guideline revealed an absence of evidence for the effectiveness of indwelling catheters over the long term. A comparison of impregnated (for example, with silver), hydrophilic and silicone catheters is needed. The primary outcome measures should be symptomatic urinary tract infections, encrustations, blockages, cost/resource use and quality of life. Secondary outcome measures should include the mean number of days the catheter remains in situ (mean dwell time) and patient comfort.

4.4 Indwelling urinary catheters: antibiotic prophylaxis

When recatheterising patients who have long-term indwelling urinary catheters, what is the clinical and cost effectiveness of single-dose antibiotic prophylaxis in reducing symptomatic urinary tract infections in patients with a history of urinary tract infections associated with catheter change?

Why is this important?

The immediate clinical and economic impact of urinary tract infection is so great that patients at risk of infection are sometimes offered the option to receive prophylactic antibiotics. However, the widespread use of antibiotics, including their prophylactic use, has been identified as a major factor in the increasing levels of antibiotic resistance observed across England and Wales. There is currently an absence of evidence about the short-term and long-term effects of prophylactic antibiotic use during catheter change. The GDG identified this as an important area for research to establish the benefits and harms of this practice in order to develop future guidance (the recommendation on this topic in the current guideline was based on GDG consensus).

A randomised controlled trial or cohort trial comparing single-dose antibiotic prophylaxis with selected major antibiotic groups is needed. The primary outcome measures should be symptomatic urinary tract infection, cost and quality of life. This is an important area for patients as it could minimise the inappropriate use of antibiotics

4.5 Vascular access devices: skin decontamination

What is the clinical and cost effectiveness of chlorhexidine 2% in alcohol versus chlorhexidine 0.5% in alcohol versus chlorhexidine 2% aqueous solution versus chlorhexidine 0.5% aqueous solution for cleansing skin (before insertion of peripheral vascular access devices [VADs] and during dressing changes of all VADs) in reducing VAD-related bacteraemia and VAD site infections?

Why is this important?

The effective management of VADs is important for reducing phlebitis and bacteraemia. In the community, compliance is improved when a single solution is used for all aspects of VAD-related skin care. There is no direct evidence comparing different percentages of chlorhexidine in aqueous and alcohol solutions, and little evidence on the use of such solutions in the community. A randomised controlled trial is required to compare the clinical and cost effectiveness of the different solutions available. The trial should

enrol patients in the community with a VAD. The protocol would need to use the same skin preparation technique regardless of solution, and could also investigate the effects of decontamination technique and drying time. The primary outcome measures should be rate of VAD-related bacteraemia, rate of VAD site infections, mortality, cost and quality of life. Secondary outcome measures should include visual infusion phlebitis (VIP) score, insertion times and skin irritation.

5 Other versions of this guideline

5.1 Full guideline

The full guideline, 'Infection: prevention and control of healthcare-associated infections in primary and community care' contains details of the methods and evidence used to develop the guideline. It is published by the National Clinical Guideline Centre, and is available from our website (www.nice.org.uk/guidance/CG[XX]/Guidance).

5.2 NICE pathway

The recommendations from this guideline have been incorporated into a NICE pathway, which is available from http://pathways.nice.org.uk/pathways/[xxx]

5.3 'Understanding NICE guidance'

A summary for patients and carers ('Understanding NICE guidance') is available from www.nice.org.uk/guidance/CGIXXI/PublicInfo

For printed copies, phone NICE publications on 0845 003 7783 or email publications@nice.org.uk (quote reference number N[XXXX]).

We encourage NHS and voluntary sector organisations to use text from this booklet in their own information about prevention and control of healthcare-associated infections in primary and community care.

Note: these details will apply when the guideline is published.

6 Related NICE guidance

Published

- Tuberculosis. NICE clinical guideline 117 (2011). Available from www.nice.org.uk/guidance/CG117
- Lower urinary tract symptoms. NICE clinical guideline 97 (2010). Available from www.nice.org.uk/guidance/CG97
- Needle and syringe programmes. NICE public health guidance 18 (2009).
 Available from www.nice.org.uk/guidance/PH18
- Surgical site infection. NICE clinical guideline 74 (2008). Available from www.nice.org.uk/guidance/CG74
- Prophylaxis against infective endocarditis. NICE clinical guideline 64 (2008). Available from www.nice.org.uk/guidance/CG64
- Urinary tract infection in children. NICE clinical guideline 54 (2007).
 Available from www.nice.org.uk/guidance/CG54
- Urinary incontinence. NICE clinical guideline 40 (2006). Available from www.nice.org.uk/guidance/CG40
- Nutrition support in adults. NICE clinical guideline 32 (2006). Available from www.nice.org.uk/guidance/CG32

Under development

NICE is developing the following guidance (details available from www.nice.org.uk):

- Urinary incontinence in neurological disease. NICE clinical guideline.
 Publication expected October 2012.
- Stroke rehabilitation. NICE clinical guideline. Publication expected April 2012.
- Healthcare-associated infections in secondary care settings. NICE advice.
 Publication expected November 2011.
- Intravenous fluid therapy in adults in hospital. NICE clinical guideline.
 Publication date to be confirmed.

7 Updating the guideline

NICE clinical guidelines are updated so that recommendations take into account important new information. New evidence is checked 3 years after publication, and healthcare professionals and patients are asked for their views; we use this information to decide whether all or part of a guideline needs updating. If important new evidence is published at other times, we may decide to do a more rapid update of some recommendations. Please see our website for information about updating the guideline.

Appendix A: The Guideline Development Group, National Collaborating Centre and NICE project team

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Appendix B: The Guideline Review Panel

The Guideline Review Panel is an independent panel that oversees the development of the guideline and takes responsibility for monitoring adherence to NICE guideline development processes. In particular, the panel ensures that stakeholder comments have been adequately considered and responded to. The panel includes members from the following perspectives: primary care, secondary care, lay, public health and industry.

Guideline Review Panel (2012)

NICE to add

[Name; style = Unnumbered bold heading]

[job title and location; style = NICE normal]

Guidelines Advisory Committee (2003)

Professor Martin Eccles (Chairman of the Committee)

Professor of Clinical Effectiveness, Centre for Health Services Research, University of Newcastle upon Tyne

Miss Amanda Wilde

Association of British Healthcare Industries (ABHI) representative

Mrs Joyce Cormie

Lay representative

Mrs Judy Mead

Head of Clinical Effectiveness, Chartered Society of Physiotherapy

Dr Marcia Kelson

Director, Patient Involvement Unit for NICE, College of Health, London

Appendix C: Changes to recommendations

Recommendations from NICE clinical guideline 2 to be deleted

Recommendation in 2003	Comment		
guideline			
Hands that are visibly soiled, or	Replaced by:		
potentially grossly contaminated with dirt or organic material, must be washed with liquid soap and water. (Recommendation 1.1.2.2 in 2003 guideline)	1.1.2.2 Decontaminate hands preferably with a handrub (conforming to current British standards ²⁵), except in the following circumstances, when liquid soap and water must be used:		
	 when hands are visibly soiled or potentially contaminated with body fluids or 		
	 in clinical situations where there is potential for the spread of alcohol-resistant organisms (such as norovirus, Clostridium difficile or organisms that cause diarrhoeal illness). 		
Hands must be decontaminated,	Replaced by:		
preferably with an alcohol-based handrub unless hands are visibly soiled, between caring for different patients and between different care activities for the same patient. (Recommendation 1.1.2.3 in	1.1.2.2 Decontaminate hands preferably with a handrub (conforming to current British standards ²⁵), except in the following circumstances, when liquid soap and water must be used:		
	 when hands are visibly soiled or potentially contaminated with body fluids or 		
2003 guideline)	 in clinical situations where there is potential for the spread of alcohol-resistant organisms (such as norovirus, Clostridium difficile or organisms that cause diarrhoeal illness). 		
Bladder instillations or washouts	Replaced by:		
must not be used to prevent catheter-associated infection. (Recommendation 1.2.5.11 in 2003 guideline)	1.2.5.10 To minimise the risk of blockages, encrustations and catheter-associated infections for patients with a long-term indwelling urinary catheter:		
	develop a patient-specific care regimen		
	do not use bladder instillations or washouts		
	 consider approaches such as reviewing the frequency of planned catheter changes and increasing fluid intake 		
	document catheter blockages.		
Hands that are visibly soiled or	Replaced by:		
contaminated with dirt or organic material must be washed with	1.1.2.2 Decontaminate hands preferably with a		

 $^{^{\}rm 25}$ At the time of consultation on the guideline (July 2011): BS EN 1500: 1997.

soap and water before using an alcohol handrub. (Recommendation 1.4.2.3 in 2003 guideline)	handrub (conforming to current British standards ²⁶), except in the following circumstances, when liquid soap and water must be used:	
	when hands are visibly soiled or potentially contaminated with body fluids or	
	 in clinical situations where there is potential for the spread of alcohol-resistant organisms (such as norovirus, Clostridium difficile or organisms that cause diarrhoeal illness). 	
	The GDG did not consider it necessary to repeat this hand hygiene recommendation in the vascular access device chapter.	
Following hand antisepsis, clean gloves and a no-touch technique or sterile gloves should be used when changing the insertion site dressing.	Replaced by: 1.4.2.1 Hands must be decontaminated (see section □) before accessing or dressing a vascular access device.	
(Recommendation 1.4.2.4 in 2003 guideline)	1.4.2.2 An aseptic technique, such as Aseptic Non Touch Technique (ANTT), must be used for vascular access device catheter site care and when accessing the system.	

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Amended recommendations (change to meaning)

Recommendations have been labelled [2003, amended 2012] if the evidence has not been reviewed but changes have been made (indicated by highlighted text) that change the meaning of the recommendation.

Recommendation in 2003 guideline	Recommendation in current guideline	Comment
1.1.4.1 Sharps must not be passed directly from hand to hand, and handling should be kept to a minimum.	1.1.4.1 Sharps should not be passed directly from hand to hand, and handling should be kept to a minimum.	The updated recommendation contains 'should' rather than 'must' because the GDG considered that the use of 'must' in the 2003 version is not covered by legislation (in accordance with the NICE guidelines manual, 2009).
1.2.5.4 Carers and patients managing their own catheters must wash their hands before and after manipulation of the catheter, in accordance with the recommendations in the standard principles section (Section 1.1).	1.2.5.4 Patients managing their own catheters, and their carers, must be educated about the need for hand decontamination before and after manipulation of the catheter, in accordance with the recommendations in the standard principles section (section 1.1).	This recommendation has been amended to reflect input from the NICE Patient and Public Involvement Programme: recommendations cannot be made directly about what patients and carers must do.
1.4.1.1 Before discharge from hospital, patients and their carers should be taught any techniques they may need to use to prevent infection and safely manage a central venous catheter. 1.4.1.2 Community healthcare personnel caring for a patient with a central venous catheter should be trained, and assessed as competent, in using and consistently adhering to the infection prevention practices described in this guideline. 1.4.1.3 Follow-up training and support should be available to patients with central venous catheters and their carers.	1.4.1.1 Before discharge from hospital, patients and their carers should be taught any techniques they may need to use to prevent infection and safely manage a vascular access device. 1.4.1.2 Healthcare workers caring for a patient with a vascular access device should be trained, and assessed as competent, in using and consistently adhering to the infection prevention practices described in this guideline. 1.4.1.3 Follow-up training and support should be available to patients with a vascular access device and their carers.	The updated recommendations contain 'vascular access device' rather than 'central venous catheter'. This change has been made because peripherally inserted catheters were included in the scope of the guideline update.

Amended recommendations (no change to meaning)

Recommendation in current guideline

- 1.1.3.1 Selection of protective equipment must¹ be based on an assessment of the risk of transmission of microorganisms to the patient, and the risk of contamination of the healthcare worker's clothing and skin by patients' blood, body fluids, secretions or excretions.
- 1.2.1.2 Community and primary healthcare workers must be trained in catheter insertion, including suprapubic catheter replacement and catheter maintenance.
- 1.2.4.1 All catheterisations carried out by healthcare workers should be aseptic procedures. After training, healthcare workers should be assessed for their competence to carry out these types of procedures.
- 1.2.5.2 Healthcare workers should ensure that the connection between the catheter and the urinary drainage system is not broken except for good clinical reasons, (for example changing the bag in line with the manufacturer's recommendations).
- 1.2.5.3 Healthcare workers must decontaminate their hands and wear a new pair of clean, non-sterile gloves before manipulating a patient's catheter, and must decontaminate their hands after removing gloves.
- 1.3.1.2 Healthcare workers should be trained in enteral feeding and management of the administration system.
- 1.4.1.2 Healthcare workers caring for a patient with a vascular access device should be trained, and assessed as competent, in using and consistently adhering to the infection prevention practices described in this guideline.
- 1.4.3.7 Healthcare workers should ensure that catheter-site care is compatible with catheter materials (tubing, hubs, injection ports, luer connectors and extensions) and carefully check compatibility with the manufacturer's recommendations.
- 1.4.4.10 When needleless devices are used, healthcare workers should ensure that all components of the system are compatible and secured, to minimise leaks and breaks in the system.

Comment

All instances of 'healthcare personnel' have been changed to 'healthcare worker'. This is for consistency with other NICE clinical guidelines and is considered a more modern term. The GDG considered the term 'healthcare workers' to include a wider group of people than healthcare professionals, which they considered to be only those staff with professional qualifications.

In recommendation 1.3.1.2, 'community staff' has been changed to 'healthcare workers', for consistency with this terminology.

In recommendation 1.4.1.2, 'community healthcare personnel' has been changed to 'healthcare workers', for consistency with this terminology.