

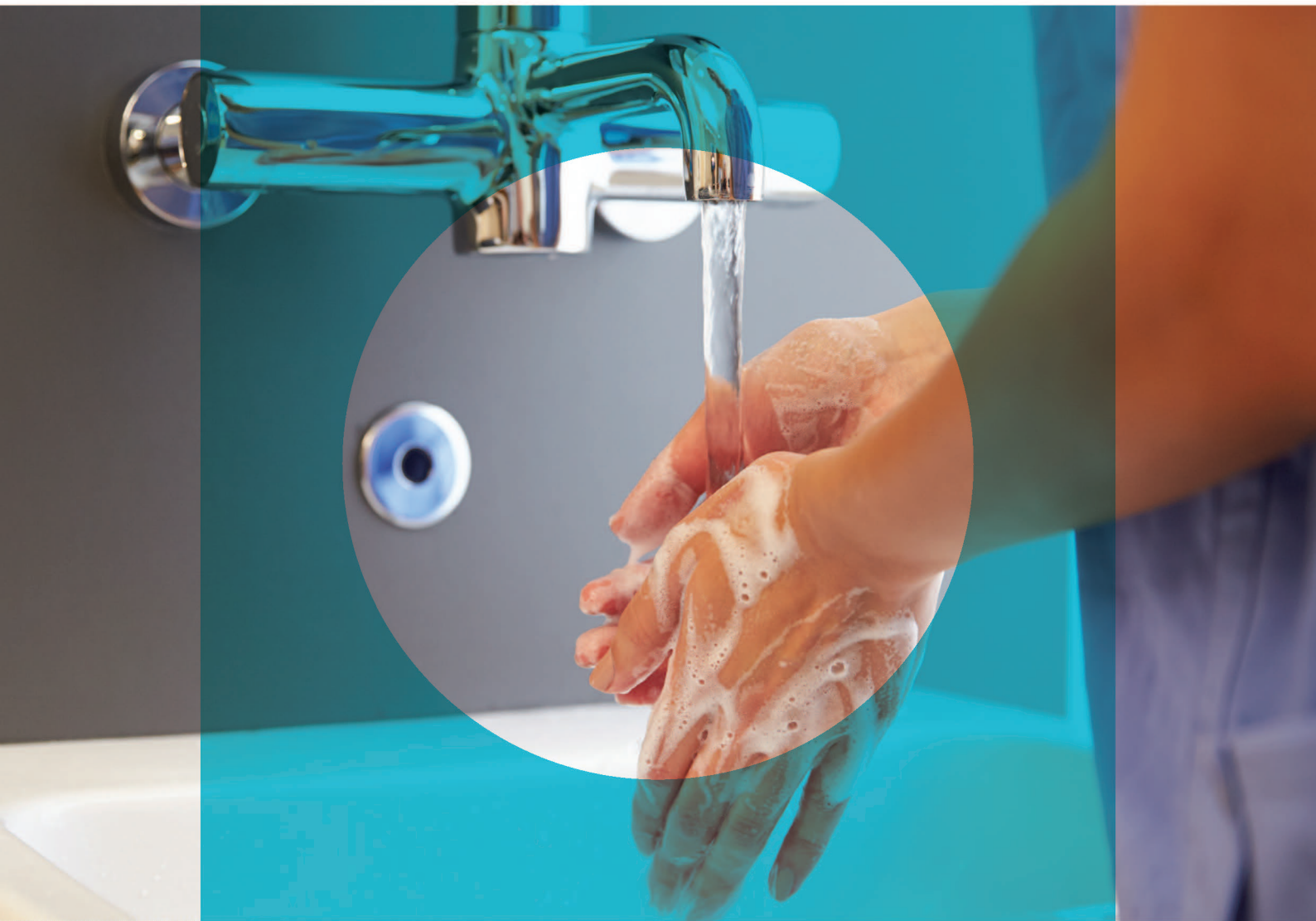


Australian Government

National Health and Medical Research Council

Australian Commission on Safety and Quality in Health Care

Australian Guidelines for the Prevention and Control of Infection in Healthcare



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**AUSTRALIAN COMMISSION
ON SAFETY AND QUALITY IN HEALTH CARE**

The Guidelines provide a nationally accepted approach to infection prevention and control, focusing on core principles and priority areas for action. They provide a basis for healthcare workers and healthcare facilities to develop detailed protocols and processes for infection prevention and control specific to local settings.

This approach is underpinned by a risk-management framework to ensure the basic principles of infection prevention and control can be applied to a wide range of healthcare settings including hospitals, day procedure units, office-based practice, long-term care facilities, remote area health services, home and community nursing and emergency services. It is recognised that the level of risk may differ according to the different types of facility and therefore some recommendations should be justified by risk assessment. When implementing these recommendations all healthcare facilities need to consider the risk of transmission of infection and implement according to their specific setting and circumstances.

The evidence base for the Guidelines addresses the highest level of risk of infection transmission in the healthcare setting, and has predominantly been drawn from the acute-care setting. The recommendations should be read in the context of the evidence base and the advice on the practical application of the recommendations.

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3. Standard and transmission-based precautions

3.1 Standard precautions

3.1.1 Hand hygiene

 **Recommended** 

1. It is recommended that routine hand hygiene is performed:

- before touching a patient
- before a procedure
- after a procedure or body substance exposure risk
- after touching a patient
- after touching a patient's surroundings.

Hand hygiene must also be performed before putting on gloves and after the removal of gloves.

 **Practice Statement** 

2. It is good practice for patients to perform hand hygiene and be educated about the benefits of hand hygiene for infection prevention and control.

Patients should be involved in hand hygiene and offered the opportunity to clean their hands when appropriate, including before meals and after using the toilet, commode or bedpan/urinal. Patient preferences for hand hygiene products may differ, and they should be provided with the option of alcohol-based hand rubs, hand wipes or access to hand wash basins, based on any specific needs.

 **Recommended** 

3. It is recommended that alcohol-based hand rubs that contain between 60% and 80% v/v ethanol or equivalent should be used for all routine hand hygiene practices.

 **Statutory Requirement** 

4. It is good practice that alcohol-based hand rubs that meet the requirements of European Standard EN 1500 are used for all routine hand hygiene practices.

Note: This advice aligns with mandatory requirements as set by Australia's Therapeutic Goods Administration

regarding testing standards for bactericidal effect (*Therapeutic Goods Act 1989*).

Recommended Updated

5. It is recommended that soap and water should be used for hand hygiene when hands are visibly soiled.

Conditional recommendation Updated

6. It is suggested that hand hygiene is performed in the presence of known or suspected *Clostridioides difficile* and non-enveloped viruses such as norovirus as follows:

- If gloves have not been worn, if gloves have been breached or if there is visible contamination of the hands despite glove use, use soap and water to facilitate the mechanical removal of spores. After washing, hands should be dried thoroughly with a single-use towel.
- If gloves have been worn, a lower density of contamination of the hands would be expected and alcohol-based hand rub remains the agent of choice for hand hygiene.

3.1.2 Use and management of sharps, safety engineered devices and medication vials

Statutory Requirement Updated

7. It is good practice to follow safe sharp handling practices including:

- not passing sharps directly from hand to hand
- keep handling to a minimum
- not recapping, bending or breaking needles after use.

Note: This advice reflects best practice as advised by expert consensus and available evidence. Healthcare workers must also consider relevant state or territory legislation that controls the management of clinical and related waste (including sharps) and Commonwealth workplace health and safety legislation (Work Health and Safety Act 2011).

Practice Statement Updated

8. It is good practice to dispose of single-use sharps immediately into an approved sharps container at the point-of-use.

The person who has used the single-use sharp must be responsible for its immediate safe disposal. Sharps containers must not be filled above the mark that indicates the maximum fill level.

3.1.3 Routine management of the physical environment

Practice Statement Updated

9. It is good practice to routinely clean surfaces as follows:

- Clean frequently touched surfaces with detergent solution at least daily, when visibly soiled and after every known contamination.
- Clean general surfaces and fittings when visibly soiled and immediately after spillage.

Practice Statement Updated

10. It is good practice for shared clinical equipment to be cleaned with a detergent solution between patient uses, and disinfected where indicated.

Exceptions to this should be justified by risk assessment.

Practice Statement Updated

11. It is good practice that surface barriers are used to protect clinical surfaces (including

equipment) that are:

- touched frequently with gloved hands during the delivery of patient care
- likely to become contaminated with blood or body substances
- difficult to clean.

Exceptions to this should be justified by risk assessment. Equipment should be appropriately cleaned between patients or uses, regardless of whether a surface barrier has been used.

Conditional recommendation Updated

12. It is suggested that site decontamination should occur after spills of blood or other potentially infectious materials.

Spills of blood or other potentially infectious materials should be promptly cleaned as follows:

- wear gloves and other personal protective equipment appropriate to the task
- confine and contain spill, clean visible matter with disposable absorbent material and discard the used cleaning materials in the appropriate waste container
- clean the spill area with a cloth or paper towels using detergent solution.

Use of Therapeutic Goods Administration-listed hospital-grade disinfectants with specific claims or a chlorine-based product such as sodium hypochlorite should be based on assessment of risk of transmission of infectious agents from that spill (see Section 3.1.3). The decision to use disinfectants should be dependent upon the compatibility of the disinfectant with the materials where the spill occurred.

Practice Statement New

13. It is good practice to use a chlorine-based product such as sodium hypochlorite or a Therapeutic Goods Administration-listed hospital-grade disinfectant with specific claims in addition to standard cleaning practices to effectively manage norovirus specific outbreaks.

3.1.3.1 Emerging disinfection methods

Conditional recommendation New

14. It is suggested that sodium hypochlorite disinfection be used as an adjunct to standard cleaning in healthcare facilities.

The use of sodium hypochlorite disinfection in addition to a detergent solution is suggested for terminal cleans of rooms of patients known or suspected to have *C. difficile* associated disease or multi-drug resistant organisms.

The use of sodium hypochlorite disinfection in addition to detergent solution is suggested to terminate outbreaks of *C. difficile*.

Conditional recommendation against New

15. The effectiveness of hydrogen peroxide vapour disinfection as an adjunct to routine cleaning in healthcare facilities is yet to be established. Therefore routine use is not suggested in healthcare facilities.

Hydrogen peroxide vapour may be considered in high-risk settings and during outbreaks when other disinfection options have been exhausted.

Conditional recommendation against New

16. The effectiveness of ultra-violet light disinfection as an adjunct to routine terminal cleaning in healthcare facilities is yet to be established. Therefore routine use is not

suggested in healthcare facilities.

Ultra-violet light disinfection may be considered in high-risk settings and during outbreaks when other disinfection options have been exhausted.

Conditional recommendation against New

17. The effectiveness of ultra-violet light disinfection in combination with sodium hypochlorite for terminal cleaning in healthcare facilities is yet to be established. Therefore routine use is not suggested in healthcare facilities.

Ultra-violet light disinfection in combination with sodium hypochlorite may be considered in high-risk settings and during outbreaks when other disinfection options have been exhausted.

Conditional recommendation against New

18. The effectiveness of surfaces, fittings or furnishing containing materials with antimicrobial properties in healthcare facilities is yet to be established. Therefore routine use is not suggested in healthcare facilities.

3.1.4 Reprocessing of reusable medical devices

3.1.5 Respiratory hygiene and cough etiquette

3.1.6 Aseptic technique

Conditional recommendation Updated

19. It is suggested that sterile gloves are used for aseptic procedures and contact with sterile sites.

3.1.7 Waste management

3.1.8 Handling of linen

3.2 Transmission-based precautions

3.2.1 Application of transmission-based precautions

3.2.2 Contact precautions

Conditional recommendation Updated

20. It is suggested that contact precautions, in addition to standard precautions, are implemented in the presence of known or suspected infectious agents that are spread by direct or indirect contact with the patient or the patient's environment.

Conditional recommendation Updated

21. It is suggested that appropriate hand hygiene be undertaken and personal protective equipment worn to prevent contact transmission.

It is suggested that when working with patients who require contact precautions, healthcare workers should:

- perform hand hygiene
- put on gloves and gown upon entry to the patient-care area
- if performing multiple tasks whilst in the patient-care area, apply the principles of standard precautions and remove gloves, perform hand hygiene and apply clean gloves between tasks when required to minimise risk of infection transmission
- ensure that clothing and skin do not contact potentially contaminated environmental surfaces
- remove gown and gloves and perform hand hygiene before leaving the patient-care area.

Conditional recommendation Updated

22. It is suggested that patient-dedicated equipment or single-use patient-care equipment be used for patients on contact precautions.

If common use of equipment for multiple patients is unavoidable, clean the equipment and allow it to dry before use on another patient.

3.2.3 Droplet precautions

Conditional recommendation Updated

23. It is suggested that droplet precautions, in addition to standard precautions, are implemented for patients known or suspected to be infected with agents transmitted by respiratory droplets that are generated by a patient when coughing, sneezing or talking.

Conditional recommendation Updated

24. It is suggested that a surgical mask should be worn when entering a patient-care environment to prevent droplet transmission.

Practice Statement Updated

25. It is good practice to place patients who require droplet precautions in a single-patient room.

3.2.4 Airborne precautions

Recommended Updated

26. It is recommended that airborne precautions, in addition to standard precautions, are implemented in the presence of known or suspected infectious agents that are transmitted person-to-person by the airborne route.

Conditional recommendation Updated

27. It is suggested that a correctly fitted P2 respirator is worn when entering the patient-care area when an airborne-transmissible infectious agent is known or suspected to be present.

Practice Statement Updated

28. It is good practice to place patients on airborne precautions in a negative pressure room (Class N/Type 5) with bathroom facilities or in a room from which air does not circulate to other areas.

Exceptions to this should be justified by risk assessment.

3.3 Personal protective equipment

Conditional recommendation Updated

29. It is suggested that clean aprons/gowns should:

- be appropriate to the task being undertaken
- be worn for a single procedure or episode of patient care where contamination with body substances is likely.

The used apron/gown should be removed in the area where the episode of patient care takes place.

Conditional recommendation Updated

30. It is suggested that face and eye protection should be worn during procedures that generate splashes or sprays of blood and body substances into the face and eyes.

Conditional recommendation Updated

31. It is suggested that single-use, fit for purpose gloves are worn for:

- each invasive procedure
- contact with sterile sites and non-intact skin or mucous membranes
- activity that has been assessed as carrying a risk of exposure to blood and body substances.

Hand hygiene should be performed prior to donning gloves and after gloves are removed.

Gloves must be changed between patients and after every episode of individual care.

3.3.1 Other items of clothing

3.4 Management of multi-resistant organisms and outbreak situations

3.4.1 Multi-resistant organisms

Conditional recommendation Updated

32. It is suggested that contact precautions be considered for all patients colonised or infected with a multi-resistant organism (MRO) where there is anticipated patient and/or environmental contact, including:

- performing hand hygiene and putting on gloves and gowns before entering the patient-care area
- using patient-dedicated or single-use non-critical patient-care equipment
- using a single-patient room or, if unavailable, cohorting patients with the same strain of MRO in designated patient-care areas (upon approval from the healthcare facility's Infection Control Team)
- ensuring consistent cleaning and disinfection of surfaces in close proximity to the patient and those likely to be touched by the patient and healthcare workers.

Practice Statement New

33. It is good practice for healthcare facilities to maintain a surveillance system to record the presence of all multi-resistant organisms.

3.4.2 Outbreak investigation and management

Practice Statement New

34. It is good practice for all outbreaks, however minor, to be investigated promptly and thoroughly and the outcomes of the investigations documented.

3.4.2.1 Infection control strategies to contain an outbreak

Practice Statement New

35. It is good practice to consider the use of early bay closures to control known or suspected norovirus outbreaks rather than ward/unit closures.

Rather than closing an entire ward or unit to manage an outbreak of norovirus in a healthcare facility, it may be more efficient to control an outbreak through cohorting symptomatic patients in bays. If taken, this approach needs to be implemented promptly and early (within three days of the first case becoming ill) in combination with adequate infection control strategies.

3.5 Applying standard and transmission-based precautions during procedures

3.5.1 Taking a risk-management approach to procedures

3.5.2 Invasive medical devices

Practice Statement New

36. It is good practice for healthcare facilities to develop, implement and review processes to address the insertion, use and maintenance, and removal of invasive medical devices. These processes should be centred on the principles of only using devices if they are deemed essential, and removing them as soon as no longer needed.

Healthcare facilities should undertake a risk assessment to assist with determining appropriate procedures and timing for the removal of invasive medical devices and for the surveillance and

management of invasive medical devices.

3.5.2.1 Indwelling urinary devices

3.5.2.2 Intravascular access devices

3.5.2.3 Ventilation

3.5.2.4 Enteral feeding tubes

3.5.3 Surgical procedures

3.5.3.1 Preventing surgical site infections (SSIs)

3.5.4 Aerosol-generating procedures (AGPs)

4. Organisational support

4.1 Management and clinical governance

4.1.1 Clinical governance in infection prevention and control

Practice Statement New

37. It is good practice for healthcare facilities to have effective clinical handover processes in place that includes infection risks.

Healthcare facilities should develop and implement a structured system for clinical handover, including documented policies and protocols.

4.1.2 Roles and responsibilities

4.1.3 Infection prevention and control program

4.1.4 Risk management

Practice Statement New

38. It is good practice to use chlorhexidine in appropriate situations and only when clinically indicated.

Healthcare professionals should consider the appropriateness of using chlorhexidine in every clinical situation, as discussed in these Guidelines.

Chlorhexidine-containing products, devices or solutions must never be used on or around patients with known chlorhexidine sensitivity.

Practice Statement New

39. It is good practice to include chlorhexidine in a healthcare facility's chemical register. Any adverse reactions to chlorhexidine should be maintained in an organisational risk

register and reported to the Therapeutic Goods Administration.

4.1.5 Taking an organisational systems approach to infection prevention quality and safety

4.2 Staff health and safety

4.2.1 Health status screening and immunisation

Statutory Requirement New

40. It is recommended that all healthcare workers to be vaccinated in accordance with the recommendations for healthcare workers in the *Australian Immunisation Handbook*.

Note: The advice reflects recommended practice supported by strong evidence. Healthcare facilities must also consider relevant state, territory and/or Commonwealth legislation regarding mandatory vaccination programs for healthcare workers.

4.2.2 Exclusion periods for healthcare workers with acute infections

Practice Statement Updated

41. It is good practice for healthcare workers and visitors to adhere to norovirus exclusion periods.

Healthcare workers should not be at work from symptom onset until 48 hours after symptom resolution. On returning to the healthcare facility, healthcare workers should adhere to appropriate hand hygiene practices.

4.2.3 Managing exposures to occupational hazards

4.2.4 Healthcare workers with specific circumstances

4.2.5 Exposure-prone procedures

4.3 Education and training

Practice Statement Updated

42. It is good practice for infection control professionals to partake in ongoing professional development in order to gain the necessary expertise to fulfil their role. Infection prevention and control staff at all levels should be supported to access formal and informal education and training relevant to their role.

4.3.1 Education strategies

4.4 Healthcare associated infection surveillance

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1. Introduction

Effective infection prevention and control is central to providing high quality healthcare for patients and a safe working environment for those that work in healthcare settings.

Healthcare associated infection is preventable

Healthcare associated infections (HAIs) are infections acquired as a direct or indirect result of healthcare [10]. There is international evidence to suggest a considerable infection burden exists among long-term care residents however in Australia there are few published studies on the rate of infection [1] [2] [3] [4]. In Australian acute healthcare facilities, there are around 165,000 HAIs each year [7]. This makes HAIs the most common complication affecting patients in hospital. As well as causing unnecessary pain and suffering for patients and their families, these adverse events prolong hospital stays and are costly to the health system. Approximately 7% of hospitalised patients will acquire a HAI, with an estimated increase to the cost of a patient's admission of 8.6% [8]. The problem does not just affect patients and workers in hospitals — HAIs can occur in any healthcare setting, including office-based practices (e.g. general practice clinics, dental clinics, community health facilities), the setting in which paramedics work and long-term care facilities. Any person working in or entering a healthcare facility is at risk. However, healthcare associated infection is a potentially preventable adverse event rather than an unpredictable complication and it is possible to significantly reduce the rate of HAIs through effective infection prevention and control.

Infection prevention and control is everybody's business

Understanding the modes of transmission of infectious organisms and knowing how and when to apply the basic principles of infection prevention and control is critical to the success of an infection control program. This responsibility applies to everybody working in and visiting a healthcare facility, including administrators, staff, patients and carers. Infection prevention and control is integral to clinical care and often requires a range of strategies to be successful. It should not be considered as an additional set of practices but as part of standard care.

Successful approaches for preventing and reducing harms arising from HAIs involve applying a risk-management framework to manage 'human' and 'system' factors associated with the transmission of infectious agents. This approach ensures that infectious agents, whether common (e.g. gastrointestinal viruses) or evolving (e.g. influenza or multi-resistant organisms), can be managed effectively.

Involving patients and their carers is essential to successful infection prevention and control in clinical care. Patients need to be sufficiently informed to be able to participate in reducing the risk of transmission of infectious agents.

Aim

By assisting healthcare workers to improve the quality of the care they deliver, these Guidelines aim to promote and facilitate the overall goal of infection prevention and control: the creation of safe healthcare environments through the implementation of evidence-based practices that minimise the risk of transmission of infectious agents.

Scope

The Guidelines provide a nationally accepted approach to infection prevention and control, focusing on core principles and priority areas for action. They provide a basis for healthcare workers and healthcare facilities to develop detailed protocols and processes for infection prevention and control specific to local settings. The Guidelines have been developed to specifically support improved infection prevention and control in acute health settings. While some of the principles and recommendations described in the Guidelines may be applicable to other health settings, all healthcare facilities should consider the risk of transmission of infection in their setting and implement the Guidelines and its recommendations according to their specific setting and circumstances.

This approach is underpinned by a risk-management framework to ensure the basic principles of infection prevention and control can be applied to a wide range of healthcare settings including hospitals, day procedure units, office-based practice, long-term care facilities, remote area health services, home and community nursing and emergency services. It is recognised that the level of risk may differ according to the different types of facility, the type of infection, developments in immunisation, and the availability of emerging technologies. When implementing the recommendations provided in these Guidelines, all healthcare facilities need to consider the risk of transmission of infection in their specific setting and circumstances and implement recommendations accordingly.

The evidence base for the Guidelines addresses the highest level of risk of infection transmission in the healthcare setting and has predominantly been drawn from the acute-care setting. The recommendations should be read in the context of the evidence base and the advice on the practical application of the recommendations. The Guidelines provide guidance on risk management and risk mitigation. Case studies giving examples of risk assessments have been included to help illustrate how these recommendations can be applied to other settings.

The Guidelines make reference to but do *not* include detailed information on:

- infectious diseases
- pandemic planning
- the reprocessing of reusable medical instruments or devices
- work health and safety
- hospital hotel services such as food services, laundry services or waste disposal
- engineering/health facility design.

Target Audience

The Guidelines are for use by all those working in acute health care settings —this includes healthcare workers, management and support staff.

As some of the principles and recommendations described in this guideline may be applicable to other health care settings, individuals working in other health care settings may also find the guidelines useful.

Structure of the Guidelines

The Guidelines are based around the following core principles:

- an understanding of the modes of transmission of infectious agents and of risk management
- effective work practices that minimise the risk of transmission of infectious agents
- governance structures that support the implementation, monitoring and reporting of infection prevention and control work practices
- compliance with legislation, regulations and standards relevant to infection control.

The sections of the Guidelines are based on these core principles and are organised according to the likely readership:

Section 2 presents background information that should be read by everyone working in healthcare (for example as orientation or as part of annual review)—this includes important basics of infection prevention and control, such as the main modes of transmission of infectious agents and the application of risk-management principles. This part of the guidelines does not include recommendations.

Section 3 is specific to the practice of healthcare workers and support staff, and outlines effective work practices that minimise the risk of transmission of infectious agents.

Section 4 describes the responsibilities of management of healthcare facilities, including governance structures that support the implementation, monitoring and reporting of effective work practices. The chapters outline the main components of a systems approach to facility-wide infection prevention and control, giving guidance on management and staff responsibilities, protection of healthcare workers, requirements for education and training of all staff, considerations for facility design, and other important activities such as surveillance and antimicrobial stewardship.

Appendices 1-3 provide additional advice on putting the recommendations into practice, risk-management case studies and resources, and the guideline development process.

Evidence Base

The Guidelines are based on the best available evidence and knowledge of the practicalities of clinical procedures. They draw from other work in this area, including the previous national infection control guidelines, international infection control guidelines, systematic literature reviews, literature reviews and horizon scans conducted to inform the development of these Guidelines, work on HAI prevention from the Australian Commission on Safety and Quality in Health Care (ACSQHC), national discipline-based infection control guidelines, and Australian Standards relevant to infection prevention and control. Australian data are used wherever available. Further information is available in *Appendix 3: Process Report*.

Recommendations, Practice Statements and Statutory Requirements

All recommendations are based on systematic reviews, with the GRADE (Grading of Recommendations Assessment, Development and Evaluation) approach providing the evidence to decision framework which determined the structure and final wording of each recommendation. Each recommendation has an accompanying *strength* which reflects the quality of the evidence underpinning the recommendation and additional factors relating to the harms and benefits of the intervention:

Strong Recommendation: Confident that the desirable effects of adherence to a recommendation outweigh the undesirable effects. Overall the recommendation is based on high quality evidence and is *strongly* recommended for implementation.

Weak/Conditional Recommendation: Concludes that the desirable effects of adherence to a recommendation *probably* outweigh the undesirable effects. Overall the recommendation is based on supportive evidence and a strong theoretical rationale and is recommended for implementation.

The Guidelines also include Practice Statements and Statutory Requirements:

Practice Statement: Set for areas which are not covered by a systematic review of the evidence, but where the provision of clinical guidance is deemed important. The development of practice statements is primarily based on best practice as advised by expert consensus and aligned with the GRADE approach where available evidence and judgements are considered together however a *strength* is not assigned.

Statutory Requirement: This advice reflects a practice statement or recommendation. The terminology '*statutory requirement*' is used to further indicate where there is also a mandated requirement/s by the Commonwealth or the States/Territories, which must be considered when implementing

the advice at the local level. It is important to note that statutory requirements vary across states and territories, and in their applicability to health service delivery sectors and settings.

The formatting of each recommendation and practice statement is as follows:

Research Evidence: Defines the research question underpinning the recommendation in the PICO (population, intervention, comparator, outcomes) format.

Key Information: Captures key information relating to the certainty of the evidence, harms and benefits of the intervention, values and preferences of the target population, resource and other considerations.

Rationale: Provides overarching justification for the stated advice.

Practical Information: Provides practical guidance on how each recommendation or practice statement can be put into practice.

Adaptation: The GRADE approach allows for the adoption, adaptation or creation of de novo recommendations from existing quality guidelines. This section captures the adoption or adaption of guidance from the 2010 edition of NHMRC's *Australian Guidelines for the Prevention and Control of Infection in Healthcare* [9].

References: Provides the full citations for the interactive references cited within each recommendation or practice statement.

Supporting Resources

Supporting resources are available for healthcare workers, patients and health facility managers to assist with implementation of the Guidelines. These materials are available to download under 'Companion Resources' at the end of these guidelines or from the [NHMRC website](#).



Australian Government

National Health and Medical Research Council

Australian Commission on Safety and Quality in Health Care

2. Basics of infection prevention and control

Summary

What are healthcare associated infections?

- Healthcare associated infections (HAIs) are infections acquired as a direct or indirect result of healthcare [1].
- HAIs can occur as a result of the provision of healthcare in any setting. While the specific risks may differ, the basic principles of infection prevention and control apply regardless of the setting.
- In order to prevent HAIs, it is important to understand how infections occur in healthcare settings and then institute ways to prevent them. If effectively implemented, the two-tiered approach of standard and transmission-based precautions recommended in these guidelines provides high-level protection to patients, healthcare workers and other people in healthcare settings.

This section covers:

- **2.1** — The basics of infection prevention and control including an overview of standard and transmission-based precautions
- **2.2** — The risk-management approach to infection prevention and control
- **2.3** — The importance of involving patients and their carers in infection prevention and control

The information presented in this part is relevant to everybody employed by a healthcare facility, including management, healthcare workers and support service staff.

2.1 Companion resources

A number of companion resources and consumer factsheets have been developed to support the implementation of these guidelines.

Companion documents

- [Summary of the guidelines](#)
- [Clinical Educator's Guide](#)
- [Frequently asked questions](#)

Consumer factsheets

- [Clostridioides difficile \(CDI\) consumer factsheet brochure](#)
- [Healthcare association infection \(HAI\) consumer factsheet brochure](#)
- [Methicillin-resistant Staphylococcus aureus \(MRSA\) consumer factsheet brochure](#)
- [Vancomycin resistant enterococci \(VRE\) consumer factsheet brochure](#)

2.2 Infection prevention and control in the healthcare setting

Summary

- Infectious agents (also called pathogens) are biological agents that cause disease or illness to their hosts. Many infectious agents are present in healthcare settings.
- Infection includes six elements - causative agent (pathogen), reservoir, portal of exit, means of transmission, portal of entry, and a susceptible host.
- Patients and healthcare workers are most likely to be sources of infectious agents and are also the most common susceptible hosts. Other people visiting and working in healthcare may also be at risk of both infection and transmission. In some cases, HAIs are serious or even life threatening.
- In healthcare settings, the main modes for transmission of infectious agents are contact (including bloodborne), droplet and airborne.

Contracting a healthcare associated infection

Most infectious agents are microorganisms. These exist naturally everywhere in the environment, and not all cause infection (e.g. 'good' bacteria present in the body's normal flora). Parasites, prions and several classes of microorganism—including bacteria, viruses, fungi and protozoa—can be involved in either colonisation or infection, depending on the susceptibility of the host:

- With **colonisation**, there is a sustained presence of replicating infectious agents on or in the body, without causing infection or disease.
- With **infection**, invasion of infectious agents into the body results in an immune response, with or without symptomatic disease.

Transmission of infectious agents within a healthcare setting requires all of the following elements:

- causative agent (pathogen)
- reservoir
- portal of exit
- means of transmission
- portal of entry
- a susceptible host.

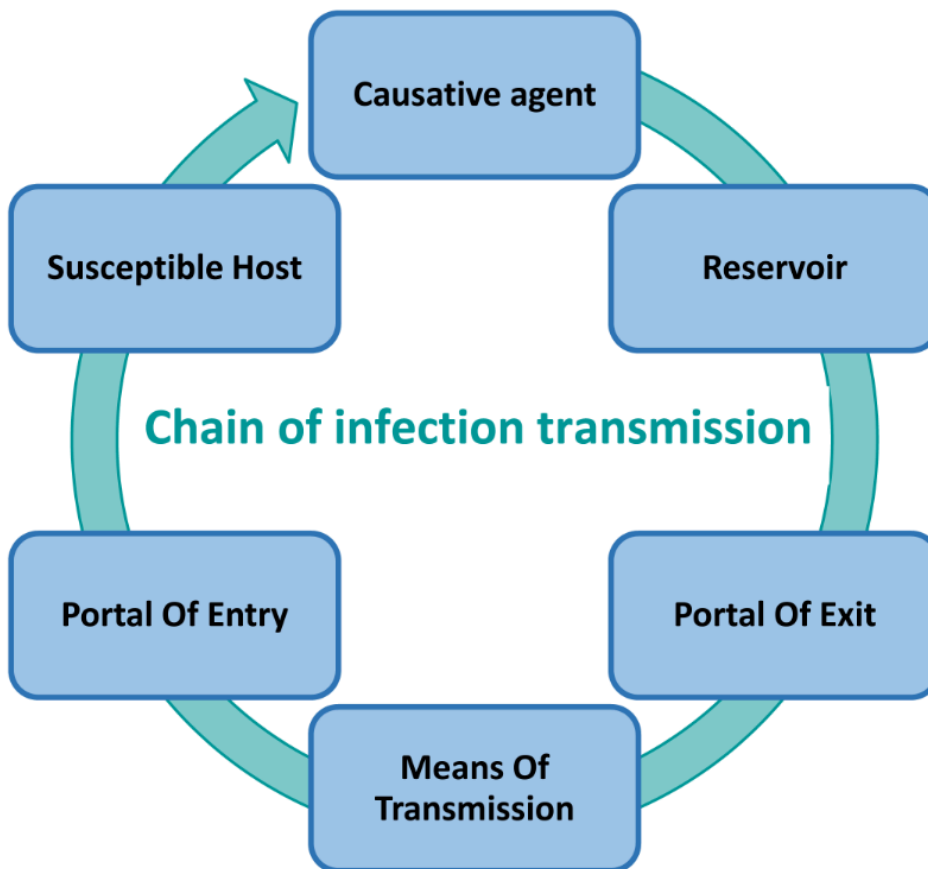


Figure 1. The chain of infection transmission

Infectious agents transmitted during healthcare come primarily from human sources, including patients, healthcare workers and visitors. Source individuals may be actively ill, may have no symptoms but be in the incubation period of a disease, or may be temporary or chronic carriers of an infectious agent with or without symptoms.

Infection is the result of a complex interrelationship between a host and an infectious agent and people vary in their response to exposure to an infectious agent:

- Some people exposed to infectious agents never develop symptomatic disease while others become severely ill and may die.
- Some individuals may become temporarily or permanently colonised but remain asymptomatic.
- Others progress from colonisation to symptomatic disease either soon after exposure, or following a period of asymptomatic colonisation.

Important predictors of an individual's outcome after exposure include:

- his or her immune status at the time of exposure (including whether immune status is compromised by medical treatment such as immunosuppressive agents or irradiation)
- the person's age (e.g. neonates and elderly patients are more susceptible)
- their health status (e.g. when a patient has other underlying disease such as diabetes or is a smoker)

- the virulence of the agent
- other factors that increase the risk of transmission of infection (e.g. undergoing surgery, requiring an indwelling device such as a catheter or remaining in hospital for lengthy periods).

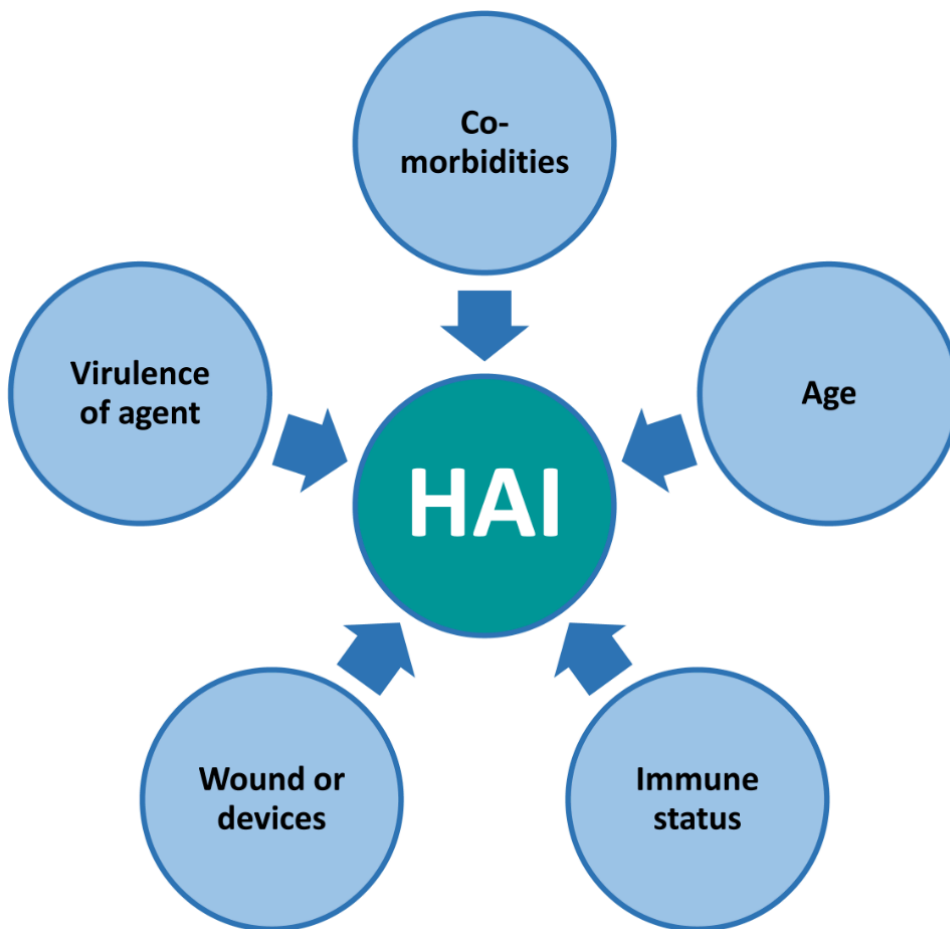


Figure 2. Factors influencing healthcare associated infection

In healthcare settings, the most common susceptible hosts are patients and healthcare workers.

- **Patients** may be exposed to infectious agents from themselves such as bacteria residing within the skin, in the respiratory or gastrointestinal tract (*endogenous infection*) or from other people, instruments and equipment, or the environment (*exogenous infection*). The level of risk relates to the healthcare setting (specifically, the presence or absence of infectious agents), the type of healthcare procedures performed, adherence to hand hygiene, immunisation status and the susceptibility of the patient.
- **Healthcare workers** may be exposed to infectious agents from infected or colonised patients, instruments and equipment, or the environment. The level of risk relates to the type of clinical contact healthcare workers have with potentially infected or colonised patient groups, instruments or environments, adherence with standard and transmission based precautions, and the health status of the healthcare worker (e.g. immunised or immunocompromised).

In healthcare settings, the main modes of transmission of infectious agents are contact (including blood borne), droplet and airborne. The modes of transmission vary by type of organism. In some cases the same organism may be transmitted by more than one route (e.g. norovirus, influenza and respiratory syncytial virus (RSV) can be transmitted by contact and droplet routes).

Routes of transmission

Contact transmission

Contact is the most common mode of transmission, and usually involves transmission by touch or via contact with blood or body substances. Contact may be direct or indirect:

- *Direct transmission* occurs when infectious agents are transferred from one person to another—for example, a patient's blood entering a healthcare worker's body through an unprotected cut in the skin.
- *Indirect transmission* involves the transfer of an infectious agent through a contaminated intermediate object or person—for example, a healthcare worker's hands transmitting infectious agents after touching an infected body site on one patient and not

performing proper hand hygiene before touching another patient, or a healthcare worker coming into contact with fomites (e.g. bedding) or faeces and then with a patient.

Examples of infectious agents transmitted by contact include multi-resistant organisms (MROs), *Clostridioides difficile* (*Clostridium difficile* or *C. difficile*), norovirus and pathogens which cause highly contagious skin infections/infestations (e.g. impetigo, scabies).

Droplet transmission

Droplet transmission can occur when an infected person coughs, sneezes or talks, and during certain procedures. Droplets are infectious particles larger than 5 microns in size. Respiratory droplets transmit infection when they travel directly from the respiratory tract of the infected person to susceptible mucosal surfaces (nasal, conjunctival or oral) of another person, generally over short distances. Droplet distribution is limited by the force of expulsion and gravity and is usually no more than 1 metre.

Examples of infectious agents that are transmitted via droplets include influenza virus and *Neisseria meningitidis* (meningococcal infection).

Airborne transmission

Airborne transmission may occur via particles containing infectious agents that remain infective over time and distance. Small-particle aerosols (often smaller than 5 microns) are created during breathing, talking, coughing or sneezing and secondarily by evaporation of larger droplets in conditions of low humidity. Aerosols containing infectious agents can be dispersed over long distances by air currents (e.g. ventilation or air conditioning systems) and inhaled by susceptible individuals who have not had any contact with the infectious person. These small particles can transmit infection into small airways of the respiratory tract.

An example of infectious agents primarily transmitted via the airborne route are *M. tuberculosis* and rubeola virus (measles).

Standard and transmission-based precautions

Successful infection prevention and control involves implementing work practices that reduce the risk of the transmission of infectious agents through a two-tiered approach, including:

- Routinely applying basic infection prevention and control strategies to minimise risk to both patients and healthcare workers, such as hand hygiene, appropriate use of personal protective equipment, cleaning and safe handling and disposal of sharps (*standard precautions*).
- Effectively managing infectious agents where standard precautions may not be sufficient on their own—these specific interventions control infection by interrupting the mode of transmission (*transmission-based precautions*; formerly referred to as *additional precautions*).

Standard precautions

All people potentially harbour infectious agents. Standard precautions refer to those work practices that are applied to everyone, regardless of their perceived or confirmed infectious status and ensure a basic level of infection prevention and control. Implementing standard precautions as a first-line approach to infection prevention and control in the healthcare environment minimises the risk of transmission of infectious agents from person to person, even in high-risk situations.

Standard precautions are used by healthcare workers to prevent or reduce the likelihood of transmission of infectious agents from one person or place to another, and to render and maintain objects and areas as free as possible from infectious agents. Guidance on implementing standard precautions is given in Sections 3.1, 6.2 & 6.3.

How standard precautions are implemented:

- Personal hygiene practices, particularly **hand hygiene**, aim to reduce the risk of contact transmission of infectious agents (see Section 3.1.1).
- Appropriate use of **personal protective equipment**, which may include gloves, gowns, plastic aprons, masks/face-shields and eye protection, aims to prevent exposure of the healthcare worker and patients to infectious agents (see Section 3.3).
- Safe **handling and disposal of sharps** assists in preventing transmission of blood-borne diseases to healthcare workers (see Section 3.1.2).
- **Environmental controls**, including cleaning and spills management, assist in preventing transmission of infectious agents from the environment to patients (see Sections 3.1.3 and 4.6.1).
- Appropriate **reprocessing of reusable equipment and instruments**, including appropriate use of disinfectants, aims to

prevent patient-to-patient transmission of infectious agents (see Section 3.1.4).

- Practising **respiratory hygiene and cough etiquette** reduces risk of transmission of infection (see Section 3.1.5).
- **Aseptic technique** aims to prevent microorganisms on hands, surfaces or equipment from being introduced into a susceptible site (see Sections 3.1.6).
- Appropriate handling of **waste and linen** assists in reducing transmission of infectious agents (see Sections 3.1.7 and 3.1.8).

Transmission-based precautions

Any infection prevention and control strategy should be based on the use of standard precautions as a minimum level of control. Transmission-based precautions are recommended as additional work practices in situations where standard precautions alone may be insufficient to prevent transmission. Transmission-based precautions are also used in the event of an outbreak (e.g. gastroenteritis), to assist in containing the outbreak and preventing further infection.

Transmission-based precautions should be tailored to the particular infectious agent involved and its mode of transmission. This may involve a combination of practices.

Guidance on when and how to implement transmission-based precautions is given in Sections 3.2, 3.5 and Appendix 2 (Section 6.3).

Types of transmission-based precautions:

- **Contact precautions** are used when there is known or suspected risk of direct or indirect contact transmission of infectious agents that are not effectively contained by standard precautions alone (see Section 3.2.2).
- **Droplet precautions** are used for patients known or suspected to be infected with agents transmitted over short distances by large respiratory droplets (see Section 3.2.3).
- **Airborne precautions** are used for patients known or suspected to be infected with agents transmitted person-to-person by the airborne route (see Section 3.2.4).

Strategies for implementing transmission-based precautions:

- dynamic risk assessment in the pre-hospital (emergency) setting to anticipate and communicate the potential need for transmission-based precautions on patient arrival
- allocating a single room inclusive of bathroom facilities and closing door to patient with a suspected or confirmed infection (isolation)
- placing patients colonised or infected with the same infectious agent and antibiogram in a room together (cohorting)
- wearing specific personal protective equipment
- providing patient-dedicated equipment
- using sodium hypochlorite or an appropriate Therapeutic Goods Administration-listed hospital-grade disinfectant with specific claims
- using specific air handling techniques
- restricting the movement of both patients and healthcare workers.

2.3 Overview of risk management in infection prevention and control

Summary

- Identifying and analysing risks associated with health care is an integral part of successful infection prevention and control.
- Adopting a risk-management approach at all levels of the facility is necessary. This task requires the full support of the facility's management as well as cooperation between management, healthcare workers and support staff.
- Differing types and levels of risk exist in different healthcare settings. In developing local policies and procedures each healthcare facility should conduct its own risk assessment (i.e. how to avoid, identify, analyse, evaluate and treat risks in that setting) and also refer to discipline-specific guidance where relevant.

Risk management basics

In the context of these guidelines, 'risk' is defined as the possibility of microorganism colonisation or infection in patients or healthcare workers arising from activities within a healthcare facility. Risk management is the basis for preventing and reducing harms arising from healthcare

associated infection.

A successful approach to risk management occurs on many levels within a healthcare facility:

- *Facility wide*—for example, providing support for effective risk management through an organisational risk-management policy, staff training, follow-up of outcomes, monitoring and reporting.
- *Ward or department based*—for example, embedding risk management into all policies so that risks are considered in every situation.
- *Individual*—for example, considering the risks involved in carrying out a specific procedure and questioning the necessity of the procedure as part of clinical decision-making, attending education sessions (e.g. hand hygiene or respirator fit testing).

As healthcare settings differ greatly in their day-to-day function, it is not possible to provide a one size fits all approach to risk management. Even within a single setting (e.g. primary care), increasingly complex care is delivered by a range of health professionals with diverse qualifications and training. All healthcare facilities need to be able to determine the risks in their own context and select the appropriate course of action. Therefore it is necessary for facilities to regularly conduct infection prevention risk assessments within their facility and ensure that all staff understand their responsibility in managing these risks.

Standard AS ISO 31000: 2018 Risk management – Guidelines outline a stepwise approach to risk management:

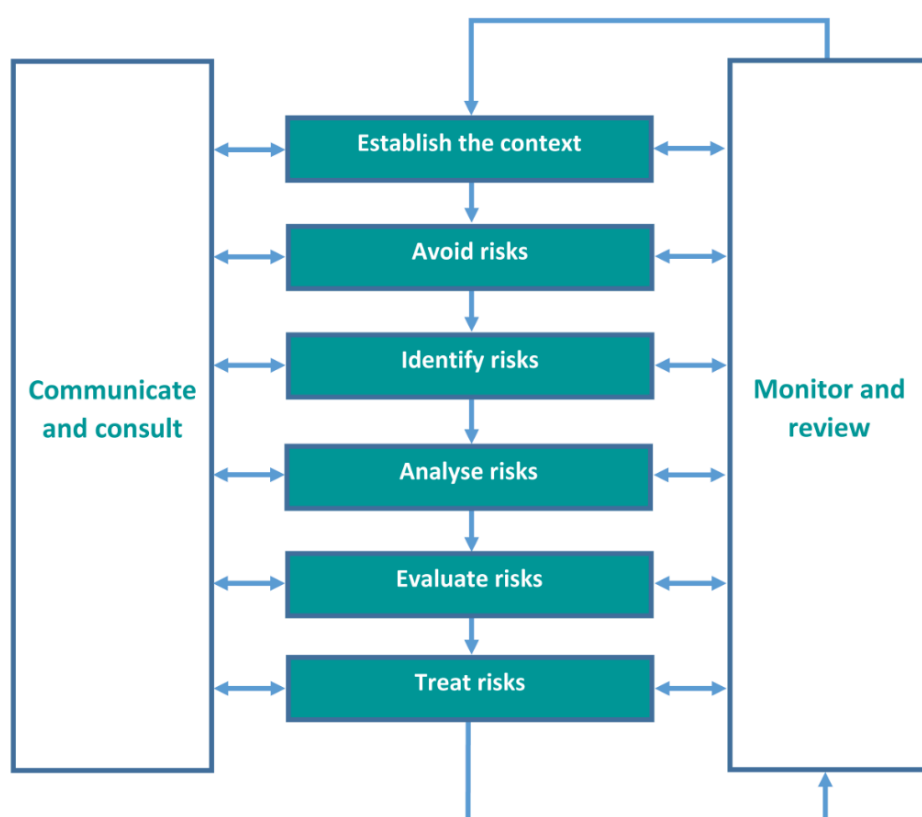


Figure 3. Risk management

1. **Establishing context** — identifying the basic parameters in which risk must be managed e.g. type of health facility, extent of and support for the facility's infection prevention and control program.
2. **Avoiding risk** — establishing whether there is a risk and whether the potential risk can be averted e.g. by questioning whether a procedure is necessary.
3. **Identifying risks** — a systematic and comprehensive process that ensures that no potential risk is excluded from further analysis and treatment e.g. using root cause analysis.
4. **Analysing risks** — considering the sources of risk, their consequences, the likelihood that those consequences may occur, and factors that affect consequences and likelihood e.g. existing controls (see *Figure 4. Risk analysis matrix* below).
5. **Evaluating risks** — comparing the level of risk found during the analysis process with previously established risk criteria, and assessing available options for ease of implementation and impact resulting in a prioritised list of risks for further action.
6. **Treating risks** — implementing appropriate management options for dealing with identified risk e.g. modifying procedures, protocols or work practices; providing education; and monitoring compliance with infection prevention and control procedures.

Monitoring and review is an essential component of the risk-management process. This ensures that:

- new risks are identified
- analysis of risk is verified against real data, if possible, and
- risk treatment is implemented effectively.

Communication and consultation are also key elements of clinical risk management. An interactive exchange of information between management, healthcare workers, patients and other stakeholders provides the basis for increased awareness of the importance of infection prevention and control, identification of risks before they arise and prompt management of risks as they occur.

| Likelihood | Consequences | | | | |
|----------------|---------------|--------|----------|---------|--------------|
| | Insignificant | Minor | Moderate | Major | Catastrophic |
| Almost certain | Medium | High | High | Extreme | Extreme |
| Likely | Medium | Medium | High | High | Extreme |
| Possible | Low | Medium | Medium | High | High |
| Unlikely | Low | Low | Medium | Medium | High |
| Rare | Low | Low | Low | Medium | Medium |

| | |
|--------------|---|
| Low risk | Manage by routine procedures. |
| Medium risk | Manage by specific monitoring or audit procedures. |
| High risk | This is serious and must be addressed immediately. |
| Extreme risk | The magnitude of the consequences of an event, should it occur, and the likelihood of that event occurring, are assessed in the context of the effectiveness of existing strategies and controls. |

Figure 4. Risk analysis matrix

Using a risk analysis matrix may assist with risk analysis and provide input into evaluation and decision making on whether the risks need action, and what the most appropriate risk mitigation strategies and methods may be.

Applying the risk management process

The following case study gives an example of applying the risk-management process in a primary care setting. Case studies giving examples of how to use this process in primary, acute and long-term care settings, including relevant considerations in specific situations, are included in Appendix 1. While the basic process of risk management applies regardless of setting, all healthcare facilities should develop risk-management policies and procedures that are appropriate to the setting.

Case study: measles (rubeola) virus outbreak

State health authorities notify a general practice of an outbreak of measles, and will assist the practice with advice about management of potential exposures. Information about the outbreak is communicated to clinicians and practice staff.

1. Establish the context

- The context is a large general practice in the outer suburbs of Sydney which caters for a diverse group of patients with many young families.

2. Avoid risks

- In this scenario, the risk cannot necessarily be avoided so it must be managed.

3. Identify risks

- The infection risk will be dependent upon the volume of non-immune patients or staff in the general practice, and also the appropriateness of infection prevention practices in place.

4. Analyse risk

- The risk has been identified as a possible airborne transmission of measles, primarily from infectious patients in the waiting room.
- The infection can be transmitted to any susceptible person breathing the same air as the infectious patient for up to 30 minutes after the patient has left the area.
- The infection also has the potential for contact transmission if droplets are transmitted indirectly to surfaces.
- If appropriate infection prevention measures are already in place, then the risk may be manageable.

5. Evaluate risk

- The balance of likelihood and consequences identify this as a 'very high risk' situation, requiring an immediate response.

6. Treat the risk

- Suspected cases should be treated as though they are confirmed cases whilst awaiting laboratory results.

Immediate measures may include:

- placing signs at the entrance to the practice advising patients to phone if they suspect they have measles
- examine suspected cases in their own home where possible or arrange for the to be seen last on the list
- displaying patient information at reception warning about the suspected measles cases
- identify and manage any patient that presents at the practice with suspected measles or similar symptoms e.g. suspected cases should avoid the waiting room
- suspected cases should be fitted with a surgical mask and taken to a separate room where they can be assessed by staff using airborne precautions
- perform the consultation in a room which can remain vacant for at least 30 minutes post consultation with suspected cases
- identify any known at-risk patients (e.g. immunocompromised) who may have an appointment at the general practice and consider potential for exposure
- respiratory etiquette and hand hygiene can be encouraged through posters and staff
- thorough surface and environmental cleaning and disinfection
- confirmed cases should be urgently reported to the local public health unit.

Longer-term measures may include:

- providing additional education to staff on measles identification and management including the process for reporting this nationally notifiable disease to appropriate health authorities and use of airborne precautions
- establish a staff vaccination policy
- at-risk staff who are not vaccinated can be identified and encouraged to be immunised.

Consider which risks need to be actively managed, why, how this will be achieved, and prioritise which actions to take based on the impacts. Use a risk evaluation matrix (Table 1 below) to determine the ease and impact of possible strategies when deciding which to implement. Note that priority must be given to activities that address risks that are high and which could have a potentially catastrophic outcome.

Table 1. Risk evaluation matrix

| Example | Ease | Analysis | Impact |
|---|------|--|--------|
| Clean and disinfect surfaces. | Easy | Potential for measles contact transmission if droplets are transmitted indirectly to surfaces. | High |
| Use of surgical mask and isolation of suspected cases. | Easy | Wearing of correctly-fitted surgical masks by coughing patients prevents dispersal of respiratory secretions into the air. | High |
| Provide ABHR in waiting, clinical rooms and consultation rooms. | Easy | Shown to improve compliance with hand hygiene, which has an impact on the spread of HAI. | High |
| Change linen between each patient in consultation rooms. | Hard | Linen not a high risk cause of measles transmission. | Low |
| Educate infectious patients to report their infectious state prior to attending practice. | Hard | May reduce the incidence of iatrogenic infection. | High |

7. Monitor and review

- The number of cases identified should be reported to the local public health unit as a notifiable disease.
- Monitor and/or follow up with any known at-risk patients e.g. immunocompromised.
- Provide feedback to staff.

Source: Adapted from RACGP (2014) *Infection Prevention and control standards (5th edition)* [379]; CDNA (2015) *National Guidelines for Public Health Units—Measles* [12], and; SA Health (2014) *Measles: Management Guidelines for General Practice* [13].

2.4 A patient-centred approach

Summary

- Healthcare facilities need to take an organisational approach to involving patients in their care.
- A patient-centred health system is known to be associated with safer and higher quality care.
- A two-way approach that encourages patient participation is essential to successful infection prevention and control.

Patient-centred healthcare

Patient-centred healthcare is respectful of, and responsive to, the preferences, needs and values of the patients and consumers [15]. People receiving healthcare increasingly expect to be given information about their condition and treatment options and this extends to their rights and responsibilities as users of healthcare services. Although patient satisfaction with health services in Australia is generally high, patients' experiences are not always valued and their expectations are not always met. While this does not necessarily lead to poor outcomes for the individuals concerned, the best possible outcomes are more likely where patient-centred health care is a priority of the healthcare facility and a strong and consistent effort is made to respect patients' rights and expectations.

The Australian Commission on Safety and Quality in Health Care developed the ***Australian Charter of Healthcare Rights*** [14], which recognises that people receiving care and people providing care all have important parts to play in achieving healthcare rights. The Charter allows patients, families, carers and services providing health care to share an understanding of the rights of people receiving health care. The Charter stipulates that all Australians have the right to:

- access services that address their healthcare needs
- receive safe and high quality health services, provided with professional care, skill and competence.
- receive care that shows respect to them and their culture, beliefs, values and personal characteristics
- receive open, timely and appropriate communication about their health care in a way they can understand
- join in making decisions and choices about their care and about health service planning
- have their personal privacy, personal health and other information properly handled
- comment on or complain about their care and have their concerns dealt with properly and promptly.

Patient-centred care cannot just be 'added on' to usual care. The rights, experiences and views of patients should be at the centre of the care process and drive the way in which care is delivered. In most healthcare facilities, a significant culture change is necessary to embed patient-centred care principles into the philosophy and practices of the organisation. Healthcare workers and organisations need to acknowledge and understand the Charter of Healthcare Rights and work to ensure that patients' rights are integral to the care process.

How does patient-centred care relate to infection prevention and control?

Infection prevention and control is ultimately about people. Effective infection prevention and control is central to providing high quality patient-centred health care.

Putting patients at the centre of infection prevention and control and enabling them to participate in the care process is not just about explaining the risks of treatments, but involves considering patients' needs at every level. However, this has to be balanced with the requirement to maintain an environment where care can be delivered in a safe manner which minimises the spread of infection. This ranges from designing the facility to maximise patient comfort and safety to having a range of processes to engage patients in their care. Healthcare workers need to listen to and act on their patients' feedback as well as provide the patient with education and support so that they can be involved in looking after themselves.

To support a two-way approach to infection prevention and control and encourage the patient participation required to minimise cross-infection or transmission, it is important to:

- take patients' perspectives into account when developing policies and programs
- familiarise patients with the infection prevention and control strategies that are employed in healthcare facilities to protect them, the people caring for them and the healthcare environment
- discuss with patients the specific risks associated with their medical and/or surgical treatment
- encourage patients to disclose their health or risk status if there is a potential risk or source of infection to healthcare workers or others within the healthcare facility

- provide opportunities for patients to identify and communicate risks and encourage them to use feedback procedures for any concerns that they have about infection prevention and control procedures
- provide educational materials about infection prevention and control using a variety of media (e.g. posters in waiting rooms, printed material and educational videos)
- inform patients about the protocols for protecting their privacy and confidentiality.

For more information, see the NSQHS Standard 1: Clinical Governance [16] and Standard 2: Partnering with Consumers [17].

2.4.1 Involving patients in their care

Involving patients in their care

Patients and visitors should be informed on what they can do to prevent the spread of infection and keep themselves infection-free in healthcare situations. Healthcare organisations should provide specific information to patients to assist them in becoming involved in identifying and reducing risks related to standard and transmission-based precautions.

Healthcare workers should, where possible:

- explain the processes of infection prevention and control (e.g. importance of hand hygiene, reasons for wearing personal protective equipment (PPE), importance of appropriate handling and disposing of sharps) to patients and their carers
- engage patients and their carers in the decision-making process regarding their care and how it is delivered
- ensure all patients and their carers are aware that they can ask questions of healthcare professionals.

Written material (such as brochures and posters) can be used to reinforce verbal discussions with patients as part of their care. Patient information supporting this Guideline is available from the [NHMRC Website](#) which aims to inform patients, visitors, families and carers about healthcare associated infection, what activities healthcare facilities may have in place to make sure infections are prevented as much as possible, and what they can do to limit the spread of infections. There is also specific patient information available on Methicillin Resistant *Staphylococcus aureus*, Vancomycin Resistant Enterococci, *Clostridioides difficile* and carbapenemase-producing *Enterobacterales* (CPE).

For further information, see the NSQHS Standard 2 on Partnering with Consumers [17].

Some examples of the types of information that should be provided to patients are below:

| Use of personal protective equipment |
|--|
| <ul style="list-style-type: none"> • Wearing of appropriate PPE such as gowns, gloves and masks is a routine part of infection prevention and control in healthcare—it is used for everybody's safety. • The use of PPE alone is not enough—healthcare workers should perform hand hygiene before putting on and after removing the protective items. • PPE is used in the patient care area only—healthcare workers remove the equipment before they leave the area to reduce the risk of spreading infection. • Gowns or aprons are used so that the healthcare worker's clothing or skin does not become contaminated. • Healthcare workers wear an appropriate mask if there is risk of them inhaling an infectious agent. • Appropriate masks, eye protection or face-shields are worn by a healthcare worker in situations where the patient's body substances may splash onto his or her face. • Healthcare workers wear gloves when they will have direct hand contact with blood or body substances, mucous membranes or wounds or if there is a chance that touching the patient could transmit infection. • Patients who are sensitive or allergic to latex should tell their healthcare workers so that an alternative glove type can be used. • It is important to note that it is not unusual for infection prevention and control practices to change over the course of care based on risk assessments made by individual healthcare workers. • It is okay to question a healthcare worker about whether they should be using protective personal equipment or whether they are using it properly. |

Handling and disposing of sharps

- Healthcare workers are at risk of injury and infection when using sharp equipment such as needles and scalpels.
- Healthcare workers take measures to handle sharp devices in a way that prevents injury to the user and to others who may encounter the device during or after a procedure.
- Special containers are used for the disposal of sharp devices.
- It is okay to question a healthcare worker about the way in which they are handling or disposing of sharp devices.
- Patients will be educated before discharge from hospital about how to safely dispose of sharps used in the home so there is no risk of injury to community members.

Management of outbreak situations

Outbreak situations may require patients to be aware of changes to infection prevention and control activities within the healthcare facility.

- In hospitals, staff must respond quickly to an outbreak of an infection to contain the infection and stop it spreading further. Actions may include testing patients to see who may carry the infection, placing patients in single rooms or with other patients who have the infection, and limiting movements of people around the facility.
- Hand hygiene is the most important part of preventing transmission of an infection.
- If infected patients are transferred, they may be asked to wear a mask.
- Infected patients should avoid unnecessary movement around other parts of the healthcare facility.
- To minimise transmission of infection in hospitals, visitors should perform hand hygiene using an alcohol-based hand rub before entering or exiting the patient care area. They may also be asked to wear gloves and gowns while they are with the patient.

3. Standard and transmission-based precautions

Summary

- The use of standard precautions is the primary strategy for minimising the transmission of healthcare associated infections.
- Standard precautions must be used regardless of known or suspected pathogens being transmitted via the contact, droplet or airborne route.
- Transmission-based precautions are used in addition to standard precautions, where the suspected or confirmed presence of infectious agents represents an increased risk of transmission.
- The application of transmission-based precautions is particularly important in containing multi-resistant organisms and in outbreak management.
- Medical and dental procedures increase the risk of transmission of infectious agents. Effective work practices to minimise risk of transmission of infection related to procedures require consideration of the specific situation, as well as appropriate use of standard and transmission-based precautions.

This section covers:

- **3.1** — Standard precautions
- **3.2** — Transmission-based precautions
- **3.3** — Personal protective equipment (PPE)
- **3.4** — Management of multi-resistant organisms and outbreak situations
- **3.5** — Applying standard and transmission-based precautions during procedures

The information presented in this part is particularly relevant to healthcare workers and support staff. It outlines effective work practices that minimise the risk of transmission of infectious agents.

Patient-care tip

In applying standard and transmission-based infection prevention and control strategies as part of day-to-day practice, healthcare workers should ensure that their patients understand why certain practices are being undertaken, and that these practices are in place to protect everyone from infection. Patients and visitors should also be aware of their role in minimising risks by following basic hand hygiene and respiratory hygiene and cough etiquette and informing staff about aspects of their care or services if necessary.

A risk mitigation approach should be used to reduce the risk of transmission of infection within healthcare facilities as not all clinical environments or activities lend themselves to the optimal conditions to prevent cross infection. It is important to risk assess patients to determine where aspects of standard precautions need to be applied and whether the addition of transmission-based precautions are required.

Where there is a concern that a clinical activity is either an infection or patient safety risk, it may be beneficial to map out the clinical activity to identify points of risk and implement steps to reduce the transmission of organisms without compromising patient safety including ensuring resources are available for cleaning and hand hygiene.

3.1 Standard precautions

Summary

Section 3.1 describes standard precautions used at all times to minimise the risk of transmission of infectious agents.

A checklist of standard precautions for procedures is in **Appendix 2—Section 6.2**.

It is essential that standard precautions are applied at all times. This is because:

- people may be placed at risk of infection from others who carry infectious agents
- people may be infectious before signs or symptoms of disease are recognised or detected, or before laboratory tests are confirmed in time to contribute to care
- people may be at risk from infectious agents present in the surrounding environment including environmental surfaces or from equipment
- there may be an increased risk of transmission associated with specific procedures and practices.

Standard precautions consist of:

- hand hygiene, as consistent with the 5 moments for hand hygiene [57]
- the use of appropriate personal protective equipment
- the safe use and disposal of sharps
- routine environmental cleaning
- reprocessing of reusable medical equipment and instruments
- respiratory hygiene and cough etiquette
- aseptic technique
- waste management
- appropriate handling of linen.

Standard precautions should be used in the handling of: blood (including dried blood); all other body substances, secretions and excretions (excluding sweat), regardless of whether they contain visible blood; non-intact skin; and mucous membranes.

3.1.1 Hand hygiene

Effective hand hygiene is the single most important strategy in preventing healthcare associated infections (HAIs). Ease of access to hand washing facilities (soap and water) and alcohol-based hand rubs can influence the transmission of HAIs. Washing hands with soap and water is required if hands are visibly soiled while either product can be used if hands are visibly clean. Each hand hygiene method is discussed in further detail in Recommendations 1 - 6.

What are the risks?

Any infectious agent transmitted by the contact or droplet route can potentially be transmitted by touch.

Microorganisms are either present on the hands most of the time (resident flora) or acquired during activities such as healthcare (transient flora). Hands can become contaminated through contact with respiratory secretions when coughing or sneezing. Contaminated hands can lead to the cross-transmission of infectious agents in non-outbreak situations [19] [34] and contribute to outbreaks involving organisms such as methicillin-resistant *Staphylococcus aureus* (MRSA), vancomycin-resistant enterococci (VRE) and multi-resistant Gram-negative (MRGN) microorganisms, such as *Acinetobacter* spp [34].

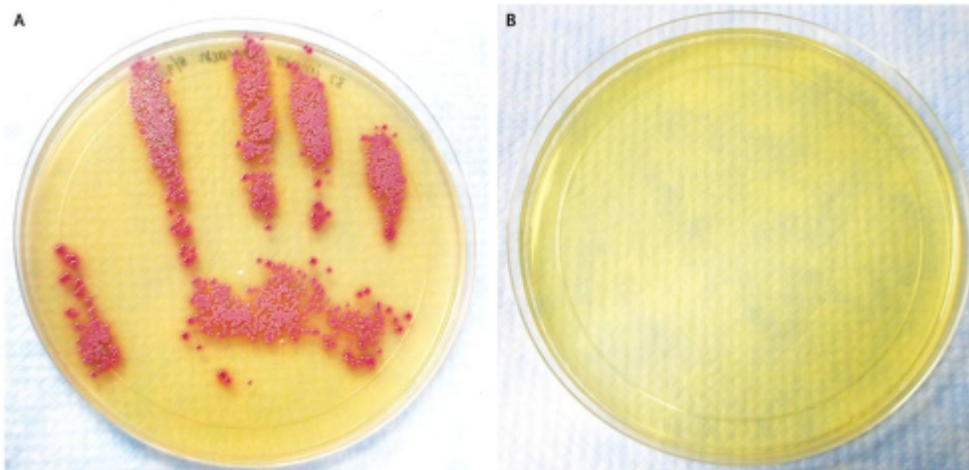


Figure 5. Importance of hand hygiene

Image source: Donskey CJ & Eckstein BC (2009) [66]

Figure 5 illustrates the critical importance of hand hygiene in caring for patients, including those not known to carry antimicrobial-resistant organisms. An imprint of a healthcare worker's ungloved hand was obtained after routine abdominal examination of a patient with no history of MRSA infection but found on routine surveillance to have MRSA colonisation. The resultant culture shows MRSA colonies (image on left). Another hand print obtained after the worker's hand had been cleaned with alcohol-based hand rub was negative for MRSA (image on right).

Improved hand hygiene practices have been associated with:

- sustained decreases in the incidence of infections caused by MRSA and VRE [46] [1] [47]
- reductions in healthcare associated infections of up to 45% in a range of healthcare settings [22] [47] [51]
- greater than 50% reduction in the rates of nosocomial disease associated with MRSA and other multi-resistant organisms,

after 1-2 years [57] [27].

Hand hygiene practices alone are not sufficient to prevent and control infection and need to be used as part of a multifactorial approach to infection control.

Recommended

Updated

1. It is recommended that routine hand hygiene is performed:

- before touching a patient
- before a procedure
- after a procedure or body substance exposure risk
- after touching a patient
- after touching a patient's surroundings.

Hand hygiene must also be performed before putting on gloves and after the removal of gloves.

Practical info

When should hand hygiene be performed?

Hands can become contaminated with infectious agents through contact with a patient, patient surroundings, the environment, or other healthcare workers. Cross-contamination can occur from one site to another in the same patient, between healthcare worker and patient, between patient or healthcare worker and the environment, or between healthcare workers. Practising hand hygiene before every episode of patient contact (including between caring for different patients and between different care activities for the same patient) and after any activity or contact that potentially results in hands becoming contaminated (such as removal of gloves) reduces the risk of cross-contamination.

The '5 moments for hand hygiene'

The '5 moments for hand hygiene' developed by the World Health Organization and recommended by the National Hand Hygiene Initiative (NHHI) Implementation Guide [57]:

- protects patients against acquiring infectious agents from the hands of the healthcare worker
- helps to protect patients from infectious agents (including their own) entering their bodies during procedures
- protects healthcare workers and the healthcare surroundings from acquiring patients' infectious agents.

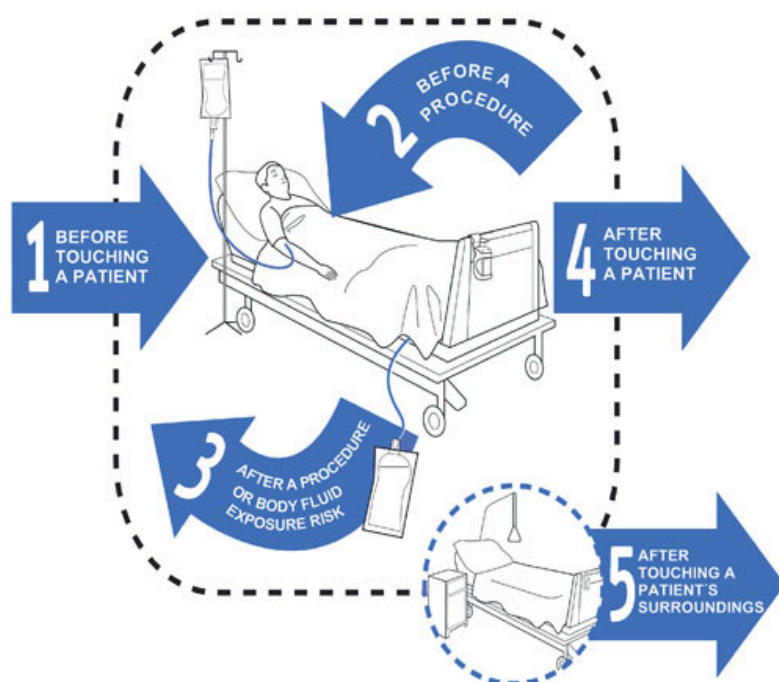


Figure 6. 5 moments for hand hygiene [57]

Note: Hand hygiene is also performed before donning and after the removal of gloves.

While Figure 6 illustrates application of the 5 moments in an acute-care setting, the 5 moments are still generally applicable to other healthcare settings including primary care. The key emphasis in any setting is to perform hand hygiene before and after any procedure, and after each consultation with a patient.

In addition to the 5 moments, hand hygiene should be performed in a range of other situations (see Table 2).

Table 2. Additional situations when hand hygiene should be performed

| Before | After |
|---|--|
| <ul style="list-style-type: none"> Starting/leaving work Eating/handling of food/drinks Using computer keyboard, tablet or mobile device in a clinical area Putting on gloves | <ul style="list-style-type: none"> Hands becoming visibly soiled Eating/handling of food/drinks Visiting the toilet Using a computer keyboard, tablet or mobile device in a clinical area Being in patient-care areas during outbreaks of infection Removing gloves Handling laundry/equipment/waste Blowing/wiping/touching nose and mouth Smoking |
| Before touching a patient | After touching a patient |
| <ul style="list-style-type: none"> Contact with patients, particularly immuno-compromised patients | <ul style="list-style-type: none"> After touching a patient, particularly patients being cared for in isolation or having transmission-based precautions applied due to the potential for spread of infection to others |
| After touching a patient's surroundings | |
| | <ul style="list-style-type: none"> Entering/leaving clinical areas Touching inanimate objects (e.g. equipment, items around the patient) and the patient environment, particularly if within an isolation room or where transmission-based precautions are applied Blood/body substance contamination |

Technique

Effective hand hygiene relies on appropriate technique as much as on selection of the correct product. Inappropriate technique can lead to failure of hand hygiene measures to appropriately remove or kill microorganisms on hands, despite the superficial appearance of having complied with hand hygiene requirements.

Key factors in effective hand hygiene and maintaining skin integrity include [19]:

- the duration of hand hygiene measures
- the exposure of all surfaces of hands and wrists to the preparation used [55]
- the use of rubbing to create friction
- ensuring that hands are completely dry.

Individual actions for reducing risk:

- Follow the 5 moments for hand hygiene, even when it seems that there is not enough time.
- Become familiar with your facility policy on hand hygiene and follow it.
- Use the appropriate product for the situation and use it as directed.

- Follow facility policy on cuts and abrasions, fingernails, nail polish and jewellery.
- Use hand-care products provided by your organisation; your own products may not be compatible with the hand hygiene products provided.
- Minimise physical contact with patient surroundings.
- Lead by example and champion hand hygiene in your setting.
- Attend hand hygiene education sessions regularly to refresh your knowledge and skills.
- Contact the person with designated responsibility for occupational health or infection prevention and control if you have a reaction to hand hygiene and hand-care products used in your setting.
- If alcohol-based hand rub is not readily accessible at key points of care in a patient-care area, consider approaching management.

Other aspects of hand hygiene

Each healthcare facility should develop policies on the wearing of jewellery, artificial fingernails or nail polish by healthcare workers. To enable optimal hand hygiene, the information below discusses several factors that should be considered.

As intact skin is a natural defence against infection, cuts and abrasions reduce the effectiveness of hand hygiene practices. Breaks or lesions of the skin are possible sources of entry for infectious agents [31] and may also be a source of them. Similarly, the presence of fingernail disease may reduce the efficacy of hand hygiene and result in the transmission of pathogens [19]. To reduce the risk of cross-transmission of infectious agents, cuts and abrasions should be covered with waterproof dressings.

The type and length of fingernails can have an impact on the effectiveness of hand hygiene [19] [33]. Artificial or false nails have been associated with higher levels of infectious agents, especially Gram-negative bacilli and yeasts, than natural nails [41] [23] [26] [37] [40] [19] [25] [18]. Fingernails should therefore be kept short (e.g. the length of the finger pad) and clean, and artificial fingernails should not be worn. Studies have also demonstrated that chipped nail polish may support the growth of organisms on the fingernails [57]. It is good practice to not wear nail polish, particularly as chipped nail polish may support the growth of organisms on the fingernail.

It is also encouraged that health care workers should wear short-sleeved clothing when delivering patient care, as this ensures their hands can be decontaminated effectively [34]. This concept is known as 'bare below the elbow.' When not engaged in patient care, some staff members may wish to cover their forearms due to religious, cultural or safety reasons. These staff must ensure they are wearing clothing with sleeves which can be pushed back securely when they are engaged in direct patient care activity [34].

Although there is less evidence concerning the impact of jewellery on the effectiveness of hand hygiene, rings can interfere with the technique used to perform hand hygiene resulting in higher total bacterial counts [19]. Hand contamination with infectious agents is increased with ring wearing [19] [53], although no studies have related this practice to healthcare worker-to-patient transmission. The consensus recommendation is to strongly discourage the wearing of watches, rings or other jewellery during health care; however if jewellery must be worn in clinical areas it should be limited to a plain band (e.g. wedding ring) and this should be moved about on the finger during hand hygiene practices. In high-risk settings such as operating suites/rooms, any jewellery, even a plain band, should not be worn.

For further resources and services to support the education of healthcare workers in the importance of hand hygiene, see the [National Hand Hygiene Initiative \(NHHI\) Implementation Guide](#) [57].

Hand care

The main type of skin irritation associated with hand hygiene, irritant contact dermatitis, includes symptoms such as dryness, irritation, itching and sometimes cracking and bleeding. Allergic contact dermatitis is rare and represents an allergy, which may be to some ingredient in a hand hygiene product.

Generally, alcohol-based hand rubs cause significantly less skin reaction or irritation than hand hygiene with plain or antiseptic soaps [46]. Expert advice concludes that [19] [57]:

- common causes of irritant contact dermatitis include skin cleaners, antiseptic washes, repeated exposure to water, sweating and glove powder
- damaged skin can lead to easier penetration of allergens and an increased likelihood of infection transmission
- the irritant and drying effects of hand preparations are one reason why healthcare workers fail to adhere to hand hygiene guidelines
- appropriate use of hand lotion or moisturisers added to hand hygiene preparations is an important factor in maintaining skin integrity, encouraging adherence to hand hygiene practices and assuring the health and safety of healthcare workers
- healthcare workers should be educated about the risk of irritant contact dermatitis and other skin damage.

Use of hand cream

An emollient hand cream should be applied regularly, such as after performing hand hygiene before a break or going off duty, and when off

duty. Hand hygiene technique should be reviewed if skin irritation occurs. If the irritation persists or if it caused by a particular soap, antiseptic agent or alcohol-based product, the person with designated responsibility for infection control or occupational health should be consulted.

For further information on hand care issues, see the National Hand Hygiene Initiative (NHII) Implementation Guide [57].

For practical information on applying appropriate hand hygiene, see Case study 5.1.

For an example of education in practice for hand hygiene, see Example 5.3.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of hand hygiene clearly outweigh any undesirable effects. A major route for transmission of infection is through contaminated hands. A National Institute for Health and Clinical Excellence (2012) systematic review [217] confirmed the association between hand decontamination and reductions in infection, and this is supported by the consensus of experts. An Australian study [62] also showed that the implementation of the National Hand Hygiene Initiative was associated with sustained hand hygiene compliance and a decrease in the incidence of methicillin-resistant *Staphylococcus aureus* bacteraemia.

The benefits of following the World Health Organization (WHO) 5 moments for hand hygiene includes protections for patients and healthcare workers against cross infection of pathogenic organisms. The potential harms are an increase in cracked skin and skin allergies from continual hand washing. These harms can be mitigated and are clearly outweighed by the benefits.

Certainty of the evidence

High

The evidence supporting this recommendation is from experimental, clinical, or epidemiological studies and these were judged as either being well designed and/or based on a strong theoretical rationale. The WHO 2009 [57] guideline strongly recommends following the five moments for hand hygiene.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections during any episode of care.

There may be challenges around complying with the 'imposition' of increased vigilance regarding hand washing, but no one is expected to challenge the reason for this culture change.

No variations expected, nor impact on health equity.

Resources and other considerations

No important issues with the recommended alternative

Sustained improvement in compliance with hand hygiene initiatives has been reported in parallel with reduced healthcare associated infection (HAI) [47]. Typically, these initiatives have involved the supply of alcohol based hand rubs, increased messaging and signage, and staff education [5]. Available evidence highlights the fact that multimodal intervention strategies lead to improved hand hygiene and a reduction in healthcare associated infection (HAI) [65].

HAIs are the most common complication affecting patients, and some of these add to costs of treatment, requiring expensive medicines, additional care and additional resources (e.g. laboratory testing and technology) to diagnose and manage HAIs [390].

Rationale

Routine hand hygiene is justified to reduce healthcare associated infection.

Routine hand hygiene is a relatively simple, low cost intervention that should be acceptable to all stakeholders. An already lengthy commitment to improving hand hygiene in Australia across public and private health care settings (beginning with the National Hand Hygiene Initiative in 2000) has resulted in improving compliance and falling infection rates. There is also an international strategy facilitated by the World Health Organization's 2005 Clean Care is Safer Care campaign, relaunched as Save Lives: Clean Your Hands in 2009.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Practice Statement

New

2. It is good practice for patients to perform hand hygiene and be educated about the benefits of hand hygiene for infection prevention and control.

Patients should be involved in hand hygiene and offered the opportunity to clean their hands when appropriate, including before meals and after using the toilet, commode or bedpan/urinal. Patient preferences for hand hygiene products may differ, and they should be provided with the option of alcohol-based hand rubs, hand wipes or access to hand wash basins, based on any specific needs.

Practical info

Involving patients in hand hygiene

Healthcare facilities are encouraged to take a patient-centred approach to infection prevention and control, as outlined in Section 2.3. Appropriate hand hygiene is one of the most well-established and supported measures for reducing HAIs, so extending education on hand hygiene practices and involving patients in this aspect of their care is encouraged.

The following information may be provided to patients to assist them in becoming involved in identifying and reducing risks related to poor hand hygiene.

- Hand hygiene is the most important aspect of reducing the risk of infection—this applies to everyone including healthcare workers, patients and visitors.
- The '5 moments for hand hygiene' tell healthcare workers, patients and visitors when hand hygiene should be performed to reduce the risk of infection.
- Healthcare workers generally use alcohol-based hand rub as it is effective and easy to use but, if their hands are visibly dirty, they need to use soap and water first.
- Performing hand hygiene regularly reduces the risk of infection to you and others. If in hospital, remind your visitors to use alcohol-based hand rub when they come into the ward and before they leave.
- No matter what product you use to clean your hands, the solution should come into contact with all surfaces.
- After hand hygiene, the hands should be dry. If alcohol-based hand rub is used, the solution will dry on the hands. After hand hygiene with soap and water, hands should be patted dry.
- Healthcare workers should have short, clean fingernails and not wear artificial fingernails, nail polish or jewellery.
- It's okay to question healthcare workers about their hand hygiene practices.

Evidence to decision

Benefits and harms A multi-modal approach to increasing hand hygiene compliance that involves patient education and engagement is now recommended as the best approach for ensuring sustainable improvement in healthcare associated infection (HAI) rates [57]. The World Health Organization Guide to Implementation recommends that patients are aware and understand the importance of hand hygiene, as engaged patients are more likely to appropriately request that healthcare workers clean their hands [64].

Certainty of the evidence The epic3 guidelines [34] found three studies that described interventions to improve patient hand hygiene, however none met the quality criteria for inclusion in their systematic review.

The National Institute for Health and Clinical Excellence (NICE) [217] guidelines recommend that patients and carers should be educated about hand hygiene, including the benefits, correct techniques and their role in maintaining standards of healthcare workers' hand decontamination.

Values and preferences It is expected that patient preferences and values would align with this intervention to reduce HAIs. Hand hygiene is considered to have a high impact on outcomes that are important to patients [217].

Resources and other considerations There are no significant resource considerations for patient involvement in hand hygiene besides any costs incurred to provide extra hand hygiene supplies such as alcohol-based hand rubs or hand wipes to patients.

Rationale

This advice is based on limited evidence, but on sound theoretical principles and supported by expert advice. Involving patients in their care is identified as a key component of infection prevention and control.

Recommended

Updated

3. It is recommended that alcohol-based hand rubs that contain between 60% and 80% v/v ethanol or equivalent should be used for all routine hand hygiene practices.

Practical info

Alcohol-based hand rubs

One advantage of alcohol-based hand rubs is that they are easily accessible at point of care. They have [57]:

- excellent antimicrobial activity against Gram-positive and Gram-negative vegetative bacteria, *Mycobacterium tuberculosis* and a wide range of fungi
- generally good antimicrobial activity against enveloped viruses
- lesser and/or variable antimicrobial activity against non-enveloped viruses (such as norovirus)
- no activity against protozoan oocysts and bacterial spores (such as *C. difficile*) (see Section 3.2.2).

The range of antimicrobial activity in alcohol-based hand rubs varies with the alcohol compound (ethanol, isopropanol or n-propanol) used. Different alcohol species have different levels of activity (60% v/v n-propanol is approximately equivalent to 70% v/v isopropanol and to 80% v/v ethanol) and many commercial formulations consist of blends of different alcohol species. Most published clinical studies that have demonstrated reductions in healthcare associated infections with the use of alcohol-based hand rubs have been associated with products that contain at least 70% alcohol (isopropanol), 0.5% chlorhexidine and a skin emollient [57]. However the efficacy of alcohol-based hand hygiene products is affected by a number of factors including the type of alcohol used, concentration of alcohol, contact time, volume of product used, and whether the hands are wet when the product is applied. These factors are generally assessed through testing standards for skin disinfectants, for which the Therapeutic Goods Administration (TGA) is the regulatory body responsible for approving products for use in Australia.

Choosing an alcohol-based hand rub

It is necessary to choose products:

- that have excellent antimicrobial efficacy combined with good user acceptability and skin tolerability (dermal tolerance, fragrance, colour, texture and ease of use)
- that are TGA approved as a hand hygiene product
- meet the requirements of EN1500 testing standards for bactericidal effect (which are currently referred to by TGA).

Healthcare worker acceptance of alcohol-based hand rub is a crucial factor in the success of any program to improve hand hygiene practice. Several studies showed that user acceptability and skin tolerability tend to be determined by the overall hand rub composition (e.g. consistency as gel or rub, texture, fragrance) and by emollient additives, but both are largely independent of a formulation's antimicrobial activity [50] [29][24] [57]. Even where emollient agents are present in the product, ready access to a moisturising skin-care product is essential (see Section 3.1.1). The selected alcohol-based hand rubs, soaps and moisturising lotions should be chemically compatible, to minimise skin reactions and ensure that the decontaminating properties of the hand hygiene product are not deactivated. It is advisable to purchase hand hygiene and hand-care products from a range made by a single manufacturer, as this ensures compatibility between the products.

Different healthcare workers and healthcare settings have different preferences, and the choice between a gel or liquid needs to be evaluated on an individual basis [35] [34]. In some healthcare facilities, it may be useful to offer both liquid and gel alongside each other, in order to provide a choice that suits a wide range of healthcare workers [34] [52] [24]. Some studies have noted that gel and foam formulations have generally significantly less antimicrobial activity than liquid alcohol-based hand rub formulations, even if the total alcohol content is similar [44] [29] [43]. However, if gel and foam formulations are more acceptable to healthcare workers and more frequently used than liquid formulations, it is important to ensure effectiveness by choosing an appropriate product (as per standards noted above), using a sufficient amount of product which allows complete coverage of the hands and allowing the hands to remain wet for the recommended amount of time, as per manufacturer instructions [57] [70] [71] [72].

The [National Hand Hygiene Initiative \(NHHI\) Implementation Guide](#) [57] recommends having alcohol-based hand rub readily available in all work areas and near patients to increase accessibility. It also outlines the following alcohol-based hand rub features as important in influencing acceptability:

- fragrance and colour—these may increase the initial appeal but may cause allergic reactions, and are therefore discouraged
- emollient agent(s) in the alcohol-based hand rub—these should prevent skin drying and irritant skin reactions, but not leave a sticky residue on hands
- drying characteristics—in general, solutions have lower viscosity than gels and therefore tend to dry more quickly
- risk of skin irritation and dryness—proactive and sympathetic management of this problem is vital

There is some evidence to suggest that gels are preferred to solutions [57], however it is important for staff to evaluate products themselves before implementation where possible. Even where emollient agents are present in the product, ready access to a moisturising skin-care product is essential. All hand hygiene products should be chemically compatible. It is advisable that hand hygiene and hand-care products are from a range made by a single manufacturer, as this ensures compatibility between the products.

Further information on acceptable dosage forms and strengths are available on the [Therapeutic Goods Administration](#) website.

Other issues associated with alcohol-based hand rubs

Other factors that should be considered when choosing products include cost issues, availability, convenience and functioning of dispenser, and ability to prevent contamination. Consideration should also be given to occupational health and safety issues associated with alcohol-based hand rubs. Alcohols are flammable, and healthcare workers handling alcohol-based preparations should respect safety standards. Accidental and intentional ingestion and dermal absorption of alcohol-based products used for hand hygiene have also been reported [49] [20]. The risk of these issues can be mitigated by appropriate placement of dispensers within the facility (see Section 4; in addition, the [National Hand Hygiene Initiative risk assessment form](#) outlines the safety issues in more detail).

Technique

Effective hand hygiene relies on appropriate technique as much as on selection of the correct product. Inappropriate technique can lead to failure of hand hygiene measures to appropriately remove or kill microorganisms on hands, despite the superficial appearance of having complied with hand hygiene requirements.

Key factors in effective hand hygiene and maintaining skin integrity include [19]:

- the duration of hand hygiene measures
- the exposure of all surfaces of hands and wrists to the preparation used [55]
- the use of rubbing to create friction
- ensuring that hands are completely dry.

Use of alcohol-based hand rub:

- apply the amount of alcohol-based hand rub recommended by the manufacturer onto dry hands
- rub hands together so that the solution comes into contact with all surfaces of the hand, paying particular attention to the tips of the fingers, the thumbs and the areas between the fingers
- continue rubbing until the solution has evaporated and the hands are dry.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of using alcohol-based hand rubs (ABHR) at 60-80% ethanol clearly outweigh any undesirable effects.

There are many level III and/or experimental studies that indicate that within the active range of about 60-90%, ABHRs with higher concentrations fare better than those with lower concentrations. There are multiple entries in the tables and references of the World Health Organization (WHO) and Centers for Disease Control and Prevention (CDC) Guidelines that document this.

The only undesirable effects of this recommendation are using ABHR with inadequate concentration of alcohol that would not pass antimicrobial testing. Specifying the 60-80% v/v ethanol or equivalent concentration accommodates that. This range was reached based on the Committee recommendation of 'at least 70% v/v ethanol' and the Council of the National Health and Medical Research Council considering and advising on the 60-80%v/v ethanol range.

Certainty of the evidence

Moderate

The evidence base and consistency are graded as moderate - one or two level II studies with a low risk of bias or a Systematic Review and/or several level III studies with a low risk of bias.

The Committee advised that the evidence base for this is probably much better than it appears from the small number of studies that were included in the formal analysis. Many studies of level III that are cited and listed in the tables of the guidelines (WHO, CDC), but were not included in the main Evidence Statement show essentially the same results.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections during any episode of care.

The provision of ABHR in health care facilities has facilitated better hand hygiene practices [5].

Resources and other considerations

No important issues with the recommended alternative

Sustained improvement in compliance with hand hygiene initiatives has been reported in parallel with reduced healthcare associated infection (HAI) [47]. Typically, these initiatives have involved the supply of alcohol based hand rubs, increased messaging and signage, and staff education [5]. Available evidence highlights the fact that multimodal intervention strategies lead to improved hand hygiene and a reduction in HAI [65].

At least half of Australia's HAIs are preventable. HAIs are the most common complication affecting patients, and some of these add to costs of treatment, requiring expensive medicines, additional care and additional resources (e.g. laboratory testing and technology) to diagnose and manage HAIs [390].

Rationale

Routine use of alcohol-based hand rub (ABHR) at 60-80% v/v ethanol is justified to reduce healthcare associated infection. ABHR is the gold standard of care for hand hygiene practice in healthcare settings while hand washing is preferred for situations when the hands are visibly soiled, or when gloves have not been worn in the care of a patient with *C. difficile* [57].

This is a relatively simple, low cost intervention that should be acceptable to all stakeholders.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Statutory Requirement

Updated

4. It is good practice that alcohol-based hand rubs that meet the requirements of European Standard EN 1500 are used for all routine hand hygiene practices.

Note: This advice aligns with mandatory requirements as set by Australia's Therapeutic Goods Administration regarding testing standards for bactericidal effect (Therapeutic Goods Act 1989).

Practical info

Choosing an alcohol-based hand rub

Healthcare facilities should choose products that meet the requirements of EN 1500 testing for bactericidal effect.

Mandatory requirements for an effective over-the-counter medicines applications, as set by Australia's Therapeutic Goods Administration (TGA) (*Therapeutic Goods Act 1989*), state that hygienic hand rubs must meet the requirements of European Standard EN 1500 testing standard for bactericidal effect.

The EN 1500 standard has two sub-categories:

- formulations that pass the standard in a contact time of 60 seconds (such as the reference alcohol)
- alcohol-based hand rubs that already achieve the sufficient level of microbial reduction in 30 seconds.

Products should also:

- have excellent antimicrobial efficacy combined with good user acceptability and skin tolerability (dermal tolerance, fragrance, colour, texture and ease of use)
- be TGA approved for skin antiseptis.

Evidence to decision

| | |
|---|--|
| Benefits and harms | The benefits of using alcohol-based hand rubs that meet the EN 1500 standards outweighs any harms. |
| Certainty of the evidence | Mandatory requirements for an effective over-the-counter medicines applications, as set by the Australian Therapeutic Goods Administration (<i>Therapeutic Goods Act 1989</i>), state that hygienic hand rubs must meet the requirements of European Standard EN 1500 testing standard for bactericidal effect. |
| Values and preferences | <p>It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections during any episode of care.</p> <p>The provision of alcohol-based hand rubs in health care facilities has facilitated better hand hygiene practices [6].</p> |
| Resources and other considerations | <p>Sustained improvement in compliance with hand hygiene initiatives has been reported in parallel with reduced healthcare associated infection (HAI) [37]. Typically, these initiatives have involved the supply of alcohol based hand rubs, increased messaging and signage, and staff education [6]. Available evidence highlights the fact that multimodal intervention strategies lead to improved hand hygiene and a reduction in HAI [56].</p> <p>At least half of Australia's HAIs are preventable. HAIs are the most common complication affecting patients, and some of these add to costs of treatment, requiring expensive medicines, additional care and additional resources (e.g. laboratory testing and technology) to diagnose and manage HAIs [285].</p> |

Rationale

Routine use of alcohol-based hand rubs (ABHR) at 60-80% v/v ethanol is justified to reduce healthcare associated infection. ABHR is the gold standard of care for hand hygiene practice in healthcare settings while hand washing is preferred for situations when the hands are visibly soiled, or when gloves have not been worn in the care of a patient with *C. difficile* [57].

Mandatory requirements for an effective over-the-counter medicines applications, as set by Australia's Therapeutic Goods Administration (*Therapeutic Goods Act 1989*), state that hygienic hand rubs must meet the requirements of European Standard EN 1500 testing standard for bactericidal effect.

The passing of stringent antimicrobial testing standards is probably an even better criterion than the total alcohol content. This is derived from the fact that different alcohol species have different levels of activity (60% v/v n-propanol is approximately equivalent to 70% v/v isopropanol and to 80% v/v ethanol) and that many commercial ABHR formulations consist of blends of different alcohol species.

One caveat when judging alcohol content is that percent v/v, w/v and w/w are different, although v/v is most commonly used. A suitable standard to be adopted would be the European EN 1500 standard, which is more stringent (i.e. has a greater safety margin in healthcare) than its US counterpart. The EN 1500 standard has two sub-categories:

- formulations that pass the standard in a contact time of 60 seconds (such as the reference alcohol)
- ABHRs that already achieve the sufficient level of microbial reduction in 30 seconds.

Of the two, the 30 seconds category is more stringent and requires greater antimicrobial activity.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice point to be reassigned as a statutory requirement. All considerations in adopting or adapting this statutory requirement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Recommended

Updated

5. It is recommended that soap and water should be used for hand hygiene when hands are visibly soiled.

Practical info

Plain soap and water

Hand washing refers to the appropriate use of a non-antimicrobial soap and water to the surface of the hands. Plain soaps act by mechanical removal of microorganisms and have no antimicrobial activity. They are sufficient for general social contact and for cleansing of visibly soiled hands. They are also used for mechanical removal of certain organisms such as *C. difficile* and norovirus.

Antimicrobial soaps are used to decontaminate hands however, when alcohol-based hand rub is available in the healthcare facility for hygienic hand antisepsis, the use of antimicrobial soap is not recommended [57]. Antimicrobial soap is associated with skin care issues and it is not necessary for use in everyday clinical practice [19] [58].

Neutral hand-wipe products may be considered in instances where hygienic access to soap and water is not readily available, such as in community care settings. Alcohol-based hand rubs are also suitable for use in resource-limited or remote areas with lack of accessibility to sinks or other facilities for hand hygiene (including clean water, towels etc.).

Technique

Effective hand hygiene relies on appropriate technique as much as on selection of the correct product. Inappropriate technique can lead to failure of hand hygiene measures to appropriately remove or kill microorganisms on hands, despite the superficial appearance of having complied with hand hygiene requirements.

Key factors in effective hand hygiene and maintaining skin integrity include [19]:

- the duration of hand hygiene measures
- the exposure of all surfaces of hands and wrists to the preparation used [55]
- the use of rubbing to create friction
- ensuring that hands are completely dry.

Using soap (including antimicrobial soap) and water:

- Wet hands under tepid running water and apply the recommended amount of liquid soap.
- Rub hands together for a minimum of 20 seconds [60] so that the solution comes into contact with all surfaces of the hand, paying particular attention to the tips of the fingers, the thumbs and the areas between the fingers.
- Rinse hands thoroughly under running water, then pat dry with single-use towels.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of using soap on visibly soiled hands clearly outweighs any undesirable effects.

Plain soap can loosen and remove transient flora. If visible soiling is not removed, the effect of any alcohol-based hand rub is minimised and effective hand hygiene is threatened.

Certainty of the evidence

High

The evidence supporting the recommendation is from experimental, clinical or epidemiological studies and these were judged as either being well designed and/or based on a strong theoretical rationale [57] [215].

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections during any episode of care. This would include maximising the potential effects of all types of hand hygiene.

Resources and other considerations

No important issues with the recommended alternative

Appropriate hand hygiene practices have an extremely high clinical impact across Australia's healthcare system. Practices are easy and feasible to implement. To maximise effectiveness, most healthcare facilities use a wide range of promotional and educational campaigns/signage.

Rationale

Washing visibly soiled hands with soap is justified to reduce healthcare associated infections.

This recommendation is based on high quality evidence, and has a profound clinical impact across healthcare settings. It is easy and feasible to implement.

The assessment is based on the fact that alcohol-based hand rubs are not effective against all microorganisms and do not remove dirt or soiling.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Conditional recommendation

Updated

6. It is suggested that hand hygiene is performed in the presence of known or suspected *Clostridioides difficile* and non-enveloped viruses such as norovirus as follows:

- If gloves have not been worn, if gloves have been breached or if there is visible contamination of the hands despite glove use, use soap and water to facilitate the mechanical removal of spores. After washing, hands should be dried thoroughly with a single-use towel.
- If gloves have been worn, a lower density of contamination of the hands would be expected and alcohol-based hand rub remains the agent of choice for hand hygiene.

Practical info

When *C. difficile* and non-enveloped viruses such as norovirus are suspected or known to be present and gloves have not been worn, a combination of hand hygiene strategies may be required to reduce transmission of these organisms. This should include hand washing with soap and water for at least 20 seconds [60] to facilitate the mechanical removal of spores. Longer hand washing is likely to be required if visible soiling is present.

If gloves are worn during the care of patients in settings where *C. difficile* or non-enveloped viruses are suspected or known to be present, spore contamination of the hands will be minimal and alcohol-based hand rub remains the agent of choice for hand hygiene [52]. The use of alcohol-based hand sanitisers may also be useful in controlling nosocomial transmission of norovirus, in combination with other infection control strategies [63].

Evidence to decision

| | |
|---------------------------|---|
| Benefits and harms | <p>The benefits of washing hands with soap and water when <i>C. difficile</i> or non-enveloped viruses are known or suspected clearly outweigh any undesirable effects.</p> <p>Meticulous hand washing and drying facilitates the mechanical removal of <i>C. difficile</i> spores. Plain soaps act by mechanical removal of <i>C. difficile</i> and non-enveloped virus microorganisms, but have no antimicrobial activity.</p> <p>Use of alcohol-based hand rubs (ABHR) alone may not be sufficient to reduce transmission of <i>C. difficile</i> as they can affect vegetative forms of <i>C. difficile</i>, but not the spores. ABHR may also not affect non-enveloped viruses.</p> |
|---------------------------|---|

| | |
|----------------------------------|---|
| Certainty of the evidence | This recommendation is supported by some experimental, clinical, or epidemiological studies and a strong theoretical rationale. |
|----------------------------------|---|

| | |
|-------------------------------|--|
| Values and preferences | It is expected that all patients and staff of Australian healthcare facilities would highly value minimising <i>C. difficile</i> infections and those caused by non-enveloped viruses such as norovirus. |
|-------------------------------|--|

| | |
|---|---|
| Resources and other considerations | There are no significant resourcing considerations associated with this intervention. |
|---|---|

Rationale

This advice is based on supportive evidence and strong theoretical principles. The use of alcohol based hand rubs alone may not be sufficient to reduce transmission of *Clostridioides difficile* and non-enveloped viruses such as norovirus as they can affect vegetative forms of *C. difficile*, but not the spores.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice point to be reassigned as a GRADE weak recommendation. All considerations in adopting or adapting this weak recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

3.1.2 Use and management of sharps, safety engineered devices and medication vials

What are the risks?

The use of sharp devices exposes healthcare workers to the risk of injury and potential exposure to blood borne infectious agents, including hepatitis B virus, hepatitis C virus and human immunodeficiency virus (HIV) [75] [76].

Sharps injuries can occur in any healthcare setting, including non-hospital settings such as in office-based practices, home health care and long-term care facilities. Injuries most often occur [75]:

- during use of a sharp device on a patient (41%)
- after use and before disposal of a sharp device (40%)
- during or after appropriate or inappropriate disposal of sharp devices (15%).

There are many possible mechanisms of injury during each of these periods.

Hollowbore needles are of particular concern, especially those used for blood collection or intravascular catheter insertion, as they are likely to contain residual blood and are associated with an increased risk for blood borne virus transmission. Non-hollowbore sharps such as glass vials and suture needles have also been involved in sharps incidents [222].

Table 3. Examples of sharps associated with sharps injuries in healthcare settings

| Examples of hollowbore sharps | Non-hollowbore sharps |
|---|----------------------------|
| Disposable needles/syringes | Glass vials |
| Steel-winged (butterfly) needles | Dental probes |
| Intravenous catheter stylets | Scalpel blades |
| Multi-sample blood collection needles | Suture needles |
| Arterial blood collection syringe needles | Retractors |
| Aspiration needles | Skin or bone hooks |
| Injector pen needles | Sharp electrosurgical tips |

Eliminating workplace hazard and risk is a fundamental principle of all work health and safety legislation in Australia. To limit the risk of sharps injuries, the hierarchy of controls method is a well-recognised approach to prevent sharps injuries [75].

The first priority is to eliminate and reduce the use of needles and other sharps where possible. Next is to isolate the hazard, thereby protecting an otherwise exposed sharp, through the use of an engineering control.

When these strategies are not available or will not provide total protection, the focus shifts to work-practice controls and personal protective equipment. An organisational approach to reducing sharps injuries is discussed in Section 4.1.5 and sharps injuries and post-exposure prophylaxis (PEP) in Section 4.2.3.

Safety-engineered devices

A broad range of devices have been designed with built-in safety features that reduce the risk of injury involving a sharp. Examples include devices such as needles with guards, sliding sheaths, shields, blunted tips or retracting needles, blunt suture needles and surgical blades with protective covers.

Safety-engineered devices can be classified into two broad categories - Passive and Active safety devices. Passive safety devices are single handed devices where the safety device activates automatically with no extra action being required by the user, such as a retractable spring loaded safety syringe. Active safety devices, such as needles with guards, require manual activation of the safety feature by the user. Passive safety devices should always be used in preference when available [94]

The use of devices with safety-engineered protective features (e.g. safety or retractable devices) has been mandated in the US, France, Spain,

most Canadian provinces and all European Union member countries including the United Kingdom [77]. Their use has shown a reduction in the rate of incidence of needlestick injuries [78] [92].

As technology advances, safety-engineered devices are going to become more readily available and safer to use.

Needleless devices

Needleless devices (e.g. connectors, vascular access devices, access ports) provide an easy access point for intravascular infusion connections. Needleless devices do not use needles for procedures such as the collection or withdrawal of body substances after initial venous or arterial access is established, or administering medication or fluids.

Since their adoption in healthcare facilities, needleless devices have contributed to a decrease in percutaneous injuries among healthcare workers [78]. While it is difficult to assess the overall effect of needleless devices because of the wide variety of devices and systems that are in use, some studies have shown an increased risk of bloodstream infections (BSI) among patients [84] [85] [93].

Unfamiliarity with the use of these complex devices, together with inadequate disinfection procedures, may contribute to increased BSI rates. The Centers for Disease Control and Prevention (CDC) recommends that [82]:

- the needleless components are changed at least as frequently as the administration set
- caps are changed no more frequently than every 3 days or according to manufacturer's recommendations
- all components of the system are compatible to minimise leaks and breaks
- contamination risk is minimised by wiping the access port with an appropriate antiseptic and accessing the port only with sterile devices.

Disinfection of needleless connectors with chlorhexidine/alcohol or povidone-iodine has been shown to significantly reduce external contamination [74]. See *Practice Statements 38 and 39* for further information on the appropriate use of chlorhexidine and potential for adverse reactions. Section 3.5.2.2 provides further information on caring for the patient's hub or insertion site.

Retractable devices

The use of retractable safety devices on sharps has been associated with a significant reduction in needlestick injury in healthcare settings [83] [88], although their direct impact is difficult to determine because their introduction is often accompanied by other interventions (e.g. training and education, overarching hospital policies and other technologies) that in isolation could also cause a reduction in needlestick injuries [89].

Retractable technology is only one example of the broad range of safety-engineered medical devices that have been designed and produced to assist in reducing the risk of occupational exposure to blood borne pathogens in healthcare.

Implementation of safety-engineered devices must be accompanied by appropriate training and education for healthcare workers in the use of the new technology to achieve successful reduction in percutaneous injury rates [88].

For more information on the handling and disposal of sharps, see **Standard AS 3825: 2020**, **Standard AS 23907:2023** or relevant international standard e.g. **ISO 23907: 2019**.

Medication vials

Single-dose vials

Medications or solutions that come into contact with normally sterile tissue should be sterile. The most effective way to avoid cross-infection via injection of medication is through the use of single-dose vials or ampoules and single-use sterile injecting equipment. Single-dose vials or ampoules, or prefilled syringes, should be used wherever these are available. These include the use of a sterile, single-use needle and syringe for each injection given, and adherence to practices that prevent contamination of injection equipment and medication.

Multi-dose vials

A multi-dose vial is one that contains more than one dose of medication. Multi-dose vials are not commonly used in Australian healthcare facilities as most vaccines are now available as single dose preparations.

However, some injectable products (e.g. Bacillus Calmette-Guérin [BCG] and botulinum toxin) are only available in multi-dose vials. When single-dose vials or ampoules are not available, there is a high risk of cross-contamination if injectable products are used on multiple patients. Steps should be taken to ensure these become available in single dose vials, however the risk of infectious disease transmission may be mitigated by [261]:

- restricting the vial to single patient use wherever possible
- establishing a separate secure area designated for the placement of these medications away from any work area
- compliance with manufacturer's recommendations (adhere to instructions for refrigeration, storage, use within a specified time, expiry date)
- using a sterile needle and syringe to draw up the required dose from the vial or ampoule on every occasion
- using a sterile needle to draw up all the contents of the container into individual syringes before administering to patients

- having only the current patient’s medication in the immediate working environment
- discarding any open ampoule(s) at the end of each procedure
- discarding product if sterility or product integrity is compromised or questionable.

The use of multi-dose vials has been associated with the transmission of infectious diseases including HIV [257], hepatitis B [253] [256] [262], hepatitis C [266] [260] [253] [254] [265] [264] [259], *Staphylococcus aureus* [258] and *Streptococcus pyogenes* [270] [269]. International agencies such as the CDC and World Health Organization recommend that single-dose vials be used for parenteral additives or medications whenever possible, especially when medications will be administered to multiple patients [255] [263].

There may be some exceptional circumstances where for short periods (e.g. a few months) multi-dose vials may be the only way to deliver vaccines or drugs to a large proportion of the population in a timely fashion. An example would be when a health emergency is declared because of an infection that has a high associated mortality and rapid spread (e.g. smallpox outbreak) and when there may be a delay in single-dose vaccines or drugs becoming available for a period of time.

Table 4. Summary of processes for appropriate use of devices

| Device | Process |
|--|---|
| Injection equipment | <ul style="list-style-type: none"> • Avoid contamination of the needle. |
| Single-use items | <ul style="list-style-type: none"> • Do not use the same needle, cannula or syringe for more than one patient nor to access a medication or solution that might be used for a subsequent patient. • Do not administer medications from a single syringe to multiple patients, even if the needle or cannula on the syringe is changed. |
| Single-patient items | <ul style="list-style-type: none"> • Use single-patient items for one patient only and dispose of them appropriately. |
| Single-use medications | <ul style="list-style-type: none"> • Only use single-dose vials when administering drugs, therapeutic agents and vaccines to multiple patients. • Do not administer medications from single-dose vials or ampoules to multiple patients or combine leftover contents for later use. |
| Multi-dose vials | <ul style="list-style-type: none"> • Multi dose vials should not be used except where they are intended solely for the exclusive use of an individual patient (e.g. insulin). |
| Fluid infusion and administration sets (i.e. intravenous bags, tubing and connectors) | <ul style="list-style-type: none"> • Use for one patient only and dispose of appropriately after use. • Do not use bags or bottles of intravenous solution as a common source of supply for multiple patients. • Consider syringes or needles/cannulae as contaminated once they have been used to enter or connect to a patient’s intravenous infusion bag or administration set. • Use closed intravenous delivery devices as standard practice. • Use premixed intravenous bags of medication wherever possible, in order to reduce the risk of contamination or infection during mixing, dilution or preparation. • Avoid disconnection of administration sets if possible to minimise the potential of contamination of IV lines. • Should be changed on a regular basis, depending on their use (see Section 3.5.2). |

Statutory Requirement

Updated

7. It is good practice to follow safe sharp handling practices including:

- not passing sharps directly from hand to hand
- keep handling to a minimum
- not recapping, bending or breaking needles after use.

Note: This advice reflects best practice as advised by expert consensus and available evidence. Healthcare workers must also consider relevant state or territory legislation that controls the management of clinical and related waste (including sharps) and Commonwealth workplace health and safety legislation (Work Health and Safety Act 2011).

Practical info

Handling of sharps

All healthcare workers should take precautions to prevent injuries caused by needles, scalpels and other sharp instruments or devices: during procedures; when cleaning used instruments; during disposal of used needles; and when handling sharp instruments after procedures.

Safety devices should be considered where appropriate to minimise risk of injury to healthcare workers. Standard measures to avoid sharps injuries include handling sharp devices in a way that prevents injury to the user and to others who may encounter the device during or after a procedure (for example the use of specifically designed scalpel blade removers to remove a used scalpel).

Examples include [75]:

- using instruments, rather than fingers, to grasp needles, retract tissue, and load/unload needles and scalpels
- using appropriately designed single-handed devices to unload needles and scalpels
- giving verbal announcements when passing sharps
- avoiding hand-to-hand passage of sharp instruments by using a Hands Free Passing Technique (HFPT) or neutral zone
- using round-tipped scalpel blades instead of pointed sharp-tipped blades.

The extent to which gloves protect healthcare workers from transmission of blood borne infectious agents following a needlestick or other puncture that penetrates the glove has not been determined [87]. Although gloves may reduce the volume of blood on the external surface of a sharp [80], the residual blood in the lumen of a hollowbore needle would not be affected; therefore, the effect on reduction of transmission risk is not quantifiable [87].

In dentistry, recapping or disassembling sharps may be unavoidable. If so, a risk assessment must be undertaken and safety devices should be used where appropriate [90].

Healthcare facilities should have sharps safety programs which include consideration of notifiable incidents as defined in Section 38 of the *Work Health and Safety Act 2011*.

Individual actions for reducing the risk:

- Explain to patients the risks to healthcare workers and others involved in the use and disposal of sharps and the measures taken to reduce these.
- Become familiar with facility protocols on handling and disposal of sharps, and legislated notifiable incidents.
- Use the appropriate product for the situation and use it as directed - as available, passive safety devices should be considered where appropriate to minimise risk of injury [94] [11].
- Avoid using needles where safe and effective alternatives are available.
- Before using any sharp medical device such as needles or scalpels, always plan for their safe handling and immediate disposal at the point-of-use.
- Make sure every used sharp medical device such as needles, scalpels etc. are disposed of properly in puncture-resistant sharps containers located at the point-of-use.
- Report any needlestick or sharps-related injuries promptly as relevant (e.g. to infection control or occupational health and safety professional, management, insurer) and ensure that you receive appropriate follow-up care.
- Ensure that you are vaccinated against blood-borne viruses such as hepatitis B.
- Participate in education sessions and professional development sessions on handling sharps, as well as those on new safety devices and how to use them.

For practical information on reducing sharp injuries, see Case Study 5.4

Evidence to decision

Benefits and harms The benefits of safe handling of sharps clearly outweigh any undesirable effects.

Eliminating workplace hazard and risk is a fundamental principle of all workplace health and safety legislation in Australia.

Certainty of the evidence This advice is based on limited evidence, but on sound theoretical principles and supported by expert advice. National and international guidelines are consistent in the advice regarding the importance of the safe use and disposal of sharps. The Epic Guidelines [79] recommend that sharps should not be passed

directly from hand to hand, and that needles must not be bent, broken or recapped.

Healthcare workers must also consider relevant state or territory legislation that controls the management of clinical and related waste (including sharps) and Commonwealth workplace health and safety legislation (*Work Health and Safety Act 2011*).

Values and preferences It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections through the safe handling of sharps during any episode of care.

Resources and other considerations Following appropriate sharps handling processes has minimal impact on resources – this is more about safe work practices and handling.

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Healthcare workers must also consider relevant state or territory legislation that controls the management of clinical and related waste (including sharps) and Commonwealth workplace health and safety legislation (*Work Health and Safety Act 2011*).

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Practice Statement

Updated

8. It is good practice to dispose of single-use sharps immediately into an approved sharps container at the point-of-use.

The person who has used the single-use sharp must be responsible for its immediate safe disposal. Sharps containers must not be filled above the mark that indicates the maximum fill level.

Practical info

Disposal of single-use sharps

Any person who has used a disposable sharp instrument or equipment must be responsible for its safe management and immediate disposal after use. Patients who use sharps, such as diabetics, should have access to suitable sharps container at the point of use.

After they are used, single-use syringes and needles, scalpel blades and other sharp items such as capillary tubes, glass and dental wires, should be placed in an appropriate container. These containers should be clearly labelled, puncture and leak proof, and conform to the **Australian Standard AS 23907:2023** or relevant international standard e.g. **ISO 23907: 2019**. The containers should be located at the point of use or, if this is not possible, as close as practical to the use area. Reusable sharps requiring transport to a reprocessing area must be placed in a puncture-resistant lidded container.

Sharps containers must be appropriately placed so that they are at an accessible height for the healthcare worker but out of reach of children and others to prevent hands and fingers entering the disposal unit. They should also be placed in a secure position or mounted on the wall to prevent tipping (approx. 1300 mm minimum off the ground) [91]. Placement of wall-mounted units should be away from general waste bins to minimise the risk of incorrect disposal.

There are numerous safety devices available that assist with safe removal and disposal of sharps (e.g. scalpel blade removers). Local protocol and procedures need to be developed to outline their appropriate use.

Reducing risks if a sharps injury is sustained:

- Seek care immediately if you sustain a sharps injury.
- If skin is penetrated, wash the affected area immediately with soap and water. Alcohol-based hand rub can be used to clean the area if soap and water are not available.
- Do not squeeze the affected area.
- Report the incident immediately to your supervisor.
- Ask about follow-up care, including post-exposure prophylaxis, which is most effective if implemented soon after the incident.
- Complete an accident / incident report form, including the date and time of the exposure, how it happened, and name of the source individual (if known).
- If a sharps injury happens to you, you can be reassured that only a small proportion of accidental exposures result in infection. Taking immediate action will lower the risk even further.

Evidence to decision

Benefits and harms The benefits of safely disposing of sharps clearly outweigh any undesirable effects. Eliminating workplace hazard and risk is a fundamental principle of all work, health and safety legislation in Australia.

The immediate disposal of sharps will greatly reduce the likelihood of needlestick injury [79].

Certainty of the evidence This advice is based on limited evidence, but on sound theoretical principles and supported by expert advice.

Values and preferences It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections through the safe handling and disposal of sharps during any episode of care.

Resources and other considerations This practice would entail the provision of approved sharps containers that conform to Australian Standards or relevant international standard e.g. ISO or EN.

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Safe disposal of single-use sharps is justified to reduce healthcare associated infection.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

3.1.3 Routine management of the physical environment

What are the risks?

Infectious agents can be widely found in healthcare settings and there is a body of clinical evidence, derived from case reports and outbreak investigations, suggesting an association between poor environmental hygiene and the transmission of infectious agents in healthcare settings [101] [186]. Transmission of infectious agents from the environment to patients may occur through direct contact with contaminated equipment, or indirectly, for example, in the acute-care setting, via hands that are in contact with contaminated equipment or the environment and then touch a patient [99].

Environmental surfaces can be safely decontaminated using less rigorous methods than those used on medical instruments and devices. The level of cleaning required depends on the objects involved and the risk of contamination—for example, surfaces that are likely to be contaminated with infectious agents (e.g. shared clinical equipment) require cleaning between patient uses, which is more often than general surfaces and fittings. However, all surfaces require regular cleaning. Thorough cleaning of all surfaces is necessary after spills and between patient uses of a room or patient-care area, especially in acute-care settings.

Managing the physical environment across healthcare settings

The cleaning practices discussed in these Guidelines are applicable to all healthcare settings. In certain circumstances, such as the setting in which paramedics work, care may be provided outside of a controlled environment and this should be considered when providing patient care e.g. surface barriers such as blue sheets may be used in uncontrolled environments where environmental cleaning is difficult to perform prior to patient contact.

Intensive care units and isolation areas require additional levels of cleaning, especially where there is a risk of multi-resistant organism transmission (see Section 3.4). The Australian Commission on Safety and Quality in Health Care has developed a number of [resources on environmental cleaning and infection prevention and control](#) that support health service organisations to implement the recommendations relating to environmental cleaning in this guideline.

[101][186][99]

Practice Statement

Updated

9. It is good practice to routinely clean surfaces as follows:

- Clean frequently touched surfaces with detergent solution at least daily, when visibly soiled and after every known contamination.
- Clean general surfaces and fittings when visibly soiled and immediately after spillage.

Practical info

Routine environmental cleaning

General surfaces and the cleaning requirements for each can be divided into two groups:

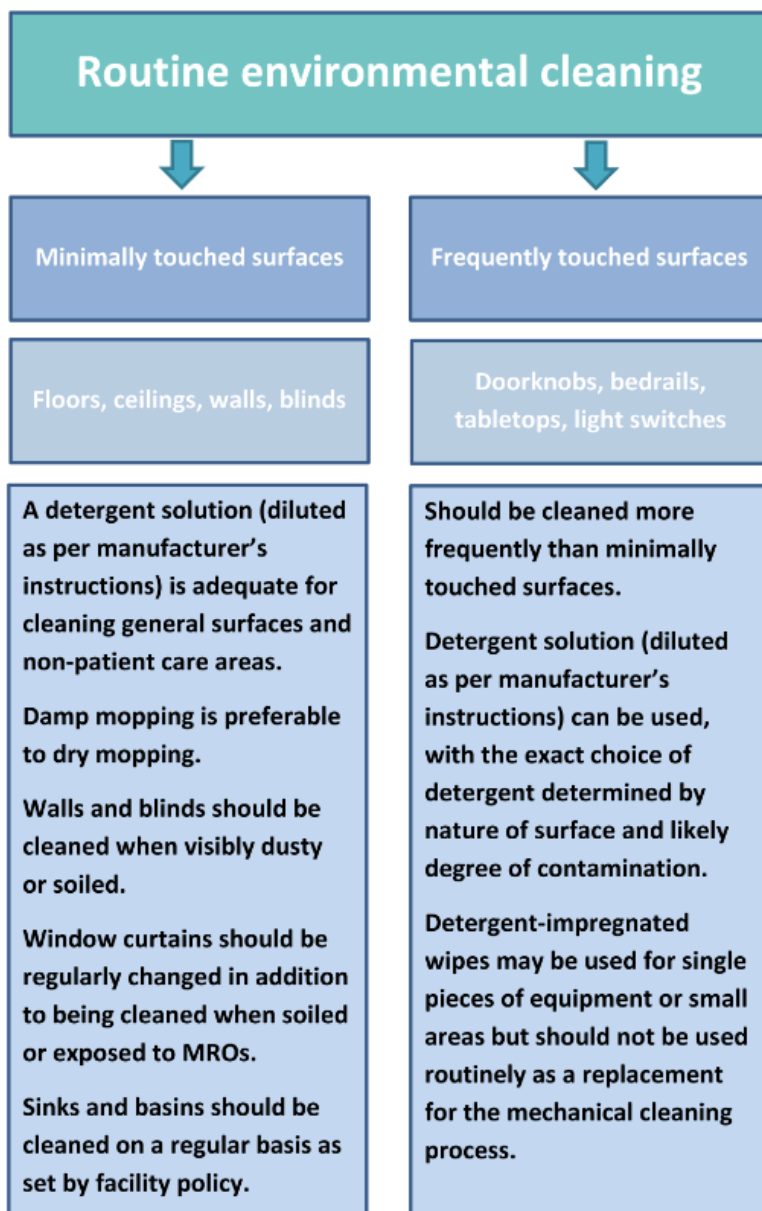


Figure 7. Cleaning requirements for routine environmental cleaning

Risk assessment

The methods, thoroughness and frequency of cleaning and the products used for different surfaces are determined by risk analysis and reflected in healthcare facility policy. Infection control professionals typically use a risk-assessment approach to identify frequently touched surfaces and then coordinate an appropriately thorough cleaning strategy and schedule with the housekeeping staff.

A detergent solution is recommended for routine cleaning. When multi-resistant organisms (MROs) are suspected or known to be present, routine cleaning is intensified and the use of a detergent solution is followed by the use of a disinfectant so that surfaces are cleaned and disinfected (see Section 3.4.1, *Recommendation 32*).

Cleaning method and product choice

Routine cleaning with detergent and water, followed by rinsing and drying, is the most useful method for removing germs from surfaces. Detergents help to loosen the germs so that they can be rinsed away with clean water. Mechanical cleaning (scrubbing the surface) physically reduces the number of germs on the surface. Rinsing with clean water removes the loosened germs and any detergent residues from the surface, and drying the surface makes it harder for germs to survive or grow.

Disinfectants are usually only necessary if a surface that has already been cleaned with detergent and water is suspected or known

to have been contaminated by MROs and/or other potentially infectious material including blood and other bodily fluids. Most germs do not survive for long on clean surfaces when exposed to air and light, and routine cleaning with detergent and water should be enough to reduce germ numbers. Disinfectants might be used after routine cleaning during an outbreak of, for example, a gastrointestinal disease.

When choosing an appropriate product/s the following factors should be considered:

- cleaning products used on different surfaces should be determined by risk assessment
- initial mechanical cleaning with a suitable detergent followed by disinfection with Therapeutic Goods Administration (TGA)-listed hospital-grade disinfectant with specific claims or a chlorine-based product such as sodium hypochlorite, where indicated for use
- the intended purpose of the product as per manufacturer's instructions
- that manufacturer's instructions are able to be complied with in the facility
- the suitability of the product to the surface or setting
- the practical application of using the product or technology with available resources including trained staff
- the effectiveness of the product against particular organisms including microbiological activity and contact time to kill microorganisms.

Cleaning schedules

The recommendations outlined for cleaning should be justified by the risk of transmission of infection within a particular healthcare facility. All organisations should have a documented cleaning schedule that outlines clear responsibilities of staff, a roster of duties and the frequency of cleaning required and the products that should be used to clean specific areas. Organisations should also facilitate job or task-specific education and training by accredited bodies for general and special cleaning of the physical environment. More detailed information about recommended cleaning schedules for different healthcare settings is in Appendix 2—Section 6.1.

If cleaning is outsourced to cleaning service providers, all cleaning service delivery procedures should be documented, including details of how the cleaning service will be undertaken. The procedures must include the following [116].

- *Minimum cleaning frequencies and methods:* cleaning service providers are required to provide cleaning services at whatever frequencies are deemed necessary in order to meet required standards. Appendix 2—Section 6.1 provides a guide for minimum frequencies for cleaning within a healthcare facility providing acute care. It can be used as a guide for other settings.
- *Staffing:* including rosters for full-time, part-time and relief staffing members, as well as for management and supervisory positions.
- *Equipment:* including provision of consumable items (such as cleaning fluids and toilet paper) and facilities to be used to deliver each cleaning service.
- *Management of the cleaning service:* how the cleaning services will be managed and controlled at the service level, including specific details of the on-site management functions.

The risk of transmission of particular infections should be assessed and the cleaning schedule should be adjusted if a known infectious agent is present (e.g. an outbreak of *C. difficile* requires surfaces to be disinfected with sodium hypochlorite after cleaning with detergent).

Usual environmental cleaning of frequently touched surfaces, such as handles, toilets, curtains and bedsheets, should be used to control and reduce the spread of non-enveloped viruses such as norovirus [96].

Housekeeping Rooms/Closets

It is important that staff who perform housekeeping duties in healthcare facilities have access to dedicated housekeeping rooms or closets. All housekeeping rooms/closets should be maintained in accordance with good hygiene practices, and should not be used for the storage of personal clothing or grooming supplies [106]. All housekeeping rooms/closets also should:

- have appropriate personal protective equipment available
- have an appropriate water supply and sink/floor drain
- be appropriately sized and well ventilated, with suitable lighting and locks fitted to all doors
- have chemical storage facilities.

All cleaning equipment must be well maintained, clean and in good repair. Cleaning equipment should be cleaned and dried between uses, and mop heads should be laundered daily.

Cleaning carts should [106]:

- have a separation between clean and soiled items
- never contain personal clothing or grooming supplies, food or beverages
- be thoroughly cleaned at the end of the day.

In long-term care homes, cleaning carts should be equipped with a locked compartment for storage of hazardous substances and each cart should be locked at all times when not attended.

Cleaning implements and solutions

Part of the cleaning strategy is to minimise contamination of cleaning solutions and cleaning tools. Proper procedures for effective use of mops, cloths, and solutions should be followed:

- prepare cleaning solutions daily or as needed, and replace with fresh solution frequently according to facility policy
- clean mops and cloths after use and allow to dry before reuse, or use single-use mop heads and cloths.

Carpet

The use of carpet in patient care areas is not suggested [396]. However, if used, carpets in public areas and in general patient-care areas should be vacuumed daily with well-maintained equipment fitted with high efficiency particulate air (HEPA) filters to minimise dust dispersion (see also Section 4.6.1). After a spill has been removed as much as possible (see practical info for *Recommendation 12*), the carpet should be cleaned using the hot water extraction method, which is recognised by **Standard AS/NZS 3733: 2018** to minimise chemical and soil residue.

Carpets should undergo thorough cleaning on a regular basis as set by facility policy, using a method that minimises the production of aerosols, leaves little or no residue and is recommended by Australian Standards and manufacturer's recommendations.

For more information about carpets, see Section 4.6.1—Mechanisms for influencing healthcare associated infection through environmental design.

Use of disinfectants

In acute-care settings where there is uncertainty about the nature of soiling on the surface (e.g. blood or body substance contamination versus routine dust or dirt) or the presence of MROs (including *C. difficile*) or other infectious agents requiring transmission-based precautions (e.g. pulmonary tuberculosis) is known or suspected, surfaces should be physically cleaned with a detergent solution, followed or combined with a hospital-grade disinfectant with specific claims listed on the Australian Register of Therapeutic Goods (ARTG) or a chlorine-based product such as sodium hypochlorite, where indicated for use (as per *Recommendation 14*). This process must involve either:

- **2-step clean**—a physical clean using a detergent followed by disinfection with a TGA-listed hospital-grade disinfectant with specific claims or a chlorine-based product such as sodium hypochlorite, where indicated for use i.e. physically clean with detergent, then physically disinfect.
- **2-in-1 clean**—a physical clean using a combined detergent and TGA-listed hospital-grade disinfectant with specific claims or a chlorine-based product such as sodium hypochlorite, where indicated for use i.e. a combined detergent/disinfectant wipe or solution could be used if this process also involves physical cleaning.

Physical (mechanical or manual) cleaning is the most important step in cleaning. Sole reliance on a disinfectant without physical cleaning is therefore not recommended. Given this, the routine use of a combined detergent and TGA-listed hospital-grade disinfectant with specific claims or a chlorine-based product such as sodium hypochlorite (2-in-1 clean) should include a risk analysis.

To kill germs, any disinfectant must:

- have enough time in contact with the surface to kill the germs (as per the manufacturer's instructions)
- be used at the right concentration
- be applied to a clean, dry surface
- be effective against those particular germs.

Hard surface disinfectants

Hard surface disinfectants include hospital, household and commercial grade disinfectants. Hard surface disinfectants are regulated by the TGA and form part of the group of products referred to as "other therapeutic goods" (OTGs). OTGs are subject to the requirements under **TGA Order 104 (Standard for Disinfectants and Sanitary Products)** of the *Therapeutic Goods Act*

1989.

Hospital-grade disinfectants with specific claims must comply with TGA Order 104 and must be 'listed' on the ARTG before they can be supplied in Australia. The term 'specific claims' covers virucidal, sporicidal, tuberculocidal, fungicidal or other biocidal activity. Except where claims of activity against fungi (yeast and mould) for excluded products are concerned, these claims mean a product is regulated as a listed OTG.

Healthcare facilities should refer to **TGA Order 104 (Standard for Disinfectants and Sanitary Products)** for more information about disinfectants and sterilants. It is best practice to refer to product safety data sheets prior to purchase for work health and safety information.

Evidence also demonstrates that a chlorine-based disinfectant products such as sodium hypochlorite can be used in addition to a detergent solution for terminal cleans of rooms of patients known to have *C. difficile* associated disease, or MROs, or to terminate outbreaks of *C. difficile*. More information is available at Section 3.1.3.1 Emerging disinfection methods, and *Recommendations 14 to 18*.

In office-based practice and less acute patient-care areas (e.g. long-term care facilities), the risk of contamination, mode of transmission and risk to others should be used to determine whether disinfectants are required.

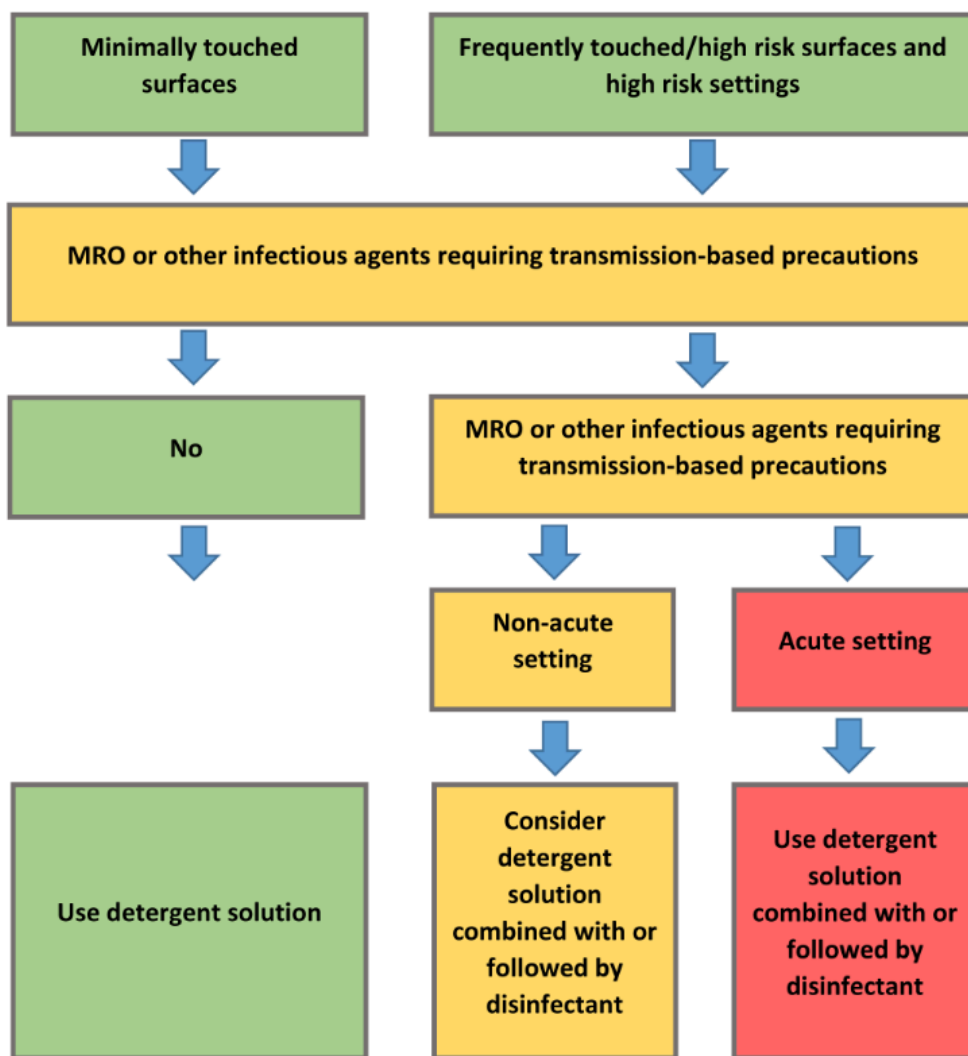


Figure 8. Processes for routine cleaning and product choice

High-level disinfectants or liquid chemical sterilants are not appropriate for general cleaning; such use is counter to manufacturers' instructions for these hazardous chemicals. Alcohol should not be used to disinfect large environmental surfaces, given the risk of additional hazards such as flammability.

Technologies in this area are evolving and new technologies for cleaning and decontaminating the healthcare environment have become available over the past ten years. More information is available in Section 3.1.3.1 Emerging disinfection methods.

Checking, auditing and environmental sampling

Healthcare facilities use a variety of systems to ensure that cleaning standards are met. These include checklists, colour coding to reduce the chance of cross infection, cleaning manuals, model cleaning contracts, infection control guidance, and monitoring strategies. Some states and territories have cleaning standards that are applied to healthcare facilities regardless of whether cleaning services are contracted or performed in-house. Users should adopt the cleaning policies appropriate to their state or territory.

Auditing of cleaning can be performed through a variety of different methods, including Process Testing and Outcome Testing. Audits of environmental cleanliness can also facilitate education programs and motivate staff to strive for improvements in routine cleaning practices [124].

Table 5. Methods for evaluating environmental cleanliness in healthcare facilities [103]

| Type | Method | Definition | Advantages | Disadvantages |
|-----------------|-------------------------|--|---|---|
| Process Testing | Visual inspection | An individual trained in the auditing of cleaning inspects an area to assess the level of cleanliness. Primary method used in healthcare facilities. | Can detect obvious soiling of the environment. Most cost-effective method and most rapid for detecting major cleaning issues. | Cannot detect microorganisms that are invisible to the naked eye. |
| | Fluorescent gel marker | An invisible gel that can only be detected with UV light is applied to surfaces. The effectiveness of cleaning processes can be determined by shining UV light to determine if the gel has been adequately removed through a cleaning process. | Can allow for an efficient and timely cleaning evaluation on a large scale. | Does not assess environmental contamination or bioburden. |
| Outcome testing | ATP bioluminescence | A swab of a surface is taken which is placed into a detection device that will catalyse a reaction with ATP. Testing the surface for ATP measures the amount of organic residue on a surface. | ATP testing provides rapid results and requires no specific laboratory training to be undertaken. | The test can produce false positives, and cannot identify the source of the ATP. The residue of some cleaning products may alter the results. |
| | Microbiological testing | Involves swabbing, dipslides, air sampling or settle plates to detect the presence of a specific microbiological organism on a surface or object. | Can provide an accurate indication of infection risk from the environment. | Expensive, labour intensive, requires specific expertise and access to a microbiology laboratory. Only recommended in the management of specific situations such as an outbreak or an unidentified cause of infections. |

Audit tools

Some organisations have developed tools and templates to assist with an environmental cleaning audit. Some examples include:

- NSW Clinical Excellence Commission Environmental Cleaning Audit Tool - <http://www.cec.health.nsw.gov.au/patient-safety-programs/assurance-governance/healthcare-associated-infections/environment-cleaning>

- CDC Environmental Cleaning Evaluation Worksheet - <https://www.cdc.gov/hai/toolkits/evaluating-environmental-cleaning.html>

- Department of Health and Human Services, Tasmania Environmental Assessment Cleaning Protocol - http://www.dhhs.tas.gov.au/publichealth/tasmanian_infection_prevention_and_control_unit/evaluating_environmental_cleanliness

Evidence to decision

| | |
|---|---|
| Benefits and harms | The benefits of routine cleaning of surfaces clearly outweigh any undesirable effects. The cleaning of environmental surfaces is fundamental in reducing the potential contribution to the incidence of healthcare associated infections [111]. |
| Certainty of the evidence | The research evidence for routine cleaning is largely limited to ecological studies and weak quasi-experimental and observational study designs. |
| Values and preferences | It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections through safe and effective cleaning of general surfaces as well as those more frequently touched or handled. |
| Resources and other considerations | This practice would entail the provision of approved cleaning agents, which has been part of standard precautions for many years. Staff training and/or tools such as checklists in wards, may be necessary to ensure uptake of advice. |

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Routine cleaning of surfaces is justified to reduce healthcare associated infection.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Practice Statement

Updated

10. It is good practice for shared clinical equipment to be cleaned with a detergent solution between patient uses, and disinfected where indicated.

Exceptions to this should be justified by risk assessment.

Practical info

Shared clinical equipment

While shared clinical equipment comes into contact with intact skin only and is therefore unlikely to introduce infection, it can act as a vehicle by which infectious agents are transferred between patients [207]. Examples of possible contaminated surfaces on shared medical equipment include knobs or handles on haemodialysis machines, x-ray machines, instrument trolleys, stethoscopes, axillary temperature monitoring probes, blood pressure cuffs, commodes and dental units [111]. Cleaning frequencies for specific shared clinical equipment is outlined in Appendix 2—Section 6.1.

Shared equipment should be cleaned with a detergent solution after each use with cleaning agents compatible with the piece of equipment being cleaned, as per manufacturer instructions. Where indicated, disinfection may also be required following routine cleaning. It is best practice to refer to the manufacturer instructions and product safety data sheet prior to using disinfectants. Choosing a disinfectant that is compatible with the surface material is integral in order to avoid damage to the equipment. All exceptions to this should be justified by risk assessment [58].

Adequate cleaning supplies should be available at or close to the point of care to enable routine management of the physical environment. The same standard procedures for the cleaning of shared equipment should apply across all healthcare settings including home health care, community settings and outpatient settings.

To reduce the risk of environmental contamination and the need to clean items between patients, disposable equipment, including

thermometers and blood pressure cuffs, should be used when caring for patients requiring contact precautions (e.g. those with *C. difficile*).

Evidence to decision

| | |
|---|---|
| Benefits and harms | <p>The benefits of cleaning shared clinical equipment between patient uses clearly outweigh any undesirable effects.</p> <p>Healthcare associated infections are a major threat to patient safety and the appropriate cleaning of shared clinical equipment can help to reduce the risk of infection.</p> |
| Certainty of the evidence | <p>There is little high quality evidence to support the cleaning of shared clinical equipment, however expert advice and work health and safety principles support this intervention.</p> |
| Values and preferences | <p>It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections through safe and effective cleaning of general surfaces as well as those more frequently touched or handled.</p> |
| Resources and other considerations | <p>This practice would entail the provision of approved cleaning agents, which has been part of standard precautions for many years.</p> <p>Staff training and/or tools such as checklists in wards may be necessary to ensure uptake of advice.</p> |

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Cleaning shared clinical equipment between uses is justified to reduce healthcare associated infections.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Practice Statement

Updated

11. It is good practice that surface barriers are used to protect clinical surfaces (including equipment) that are:

- touched frequently with gloved hands during the delivery of patient care
- likely to become contaminated with blood or body substances
- difficult to clean.

Exceptions to this should be justified by risk assessment. Equipment should be appropriately cleaned between patients or uses, regardless of whether a surface barrier has been used.

Practical info

Surface barriers (e.g. clear plastic wrap, bags, sheets, tubing or other materials impervious to moisture) help prevent contamination of surfaces and equipment. Surface barriers on equipment (e.g. air water syringes, bedboards, computer keyboards) need to be placed carefully to ensure that they protect the surfaces underneath and should be changed and cleaned between patients. Cleaning clinical surfaces including equipment should always occur between patients or uses, regardless of whether a surface barrier has been used or not.

For specialised equipment which is difficult to clean and the application of detergent directly onto the device is not recommended by the manufacturer, a custom surface barrier should be used e.g. intraoral camera. Any custom surface barrier used on such equipment should be disposed of after each patient treatment and replaced with a new custom surface barrier.

Evidence to decision

| | |
|---|---|
| Benefits and harms | <p>The benefits of cleaning shared clinical equipment clearly outweigh any undesirable effects. The use of surface barriers can help to reduce the spread of infection.</p> <p>Healthcare associated infections are a major threat to patient safety and the appropriate use of surface barriers to protect clinical surfaces can help to reduce the transmission of infection.</p> |
| Certainty of the evidence | <p>The evidence for this practice statement is limited or inconsistent. This practice statement is based on the advice of experts and trends in clinical practice.</p> |
| Values and preferences | <p>It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections through the use of surface barriers to protect clinical contact surfaces.</p> |
| Resources and other considerations | <p>The net benefits of this intervention are worth the cost.</p> <p>This practice would entail the provision of appropriate barriers, at a cost to the healthcare facility.</p> |

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. The use of surface barriers is justified to reduce healthcare associated infections.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Conditional recommendation

Updated

12. It is suggested that site decontamination should occur after spills of blood or other potentially infectious materials.

Spills of blood or other potentially infectious materials should be promptly cleaned as follows:

- wear gloves and other personal protective equipment appropriate to the task
- confine and contain spill, clean visible matter with disposable absorbent material and discard the used cleaning materials in the appropriate waste container
- clean the spill area with a cloth or paper towels using detergent solution.

Use of Therapeutic Goods Administration-listed hospital-grade disinfectants with specific claims or a chlorine-based product such as sodium hypochlorite should be based on assessment of risk of transmission of infectious agents from that spill (see Section 3.1.3). The decision to use disinfectants should be dependent upon the compatibility of the disinfectant with the materials where the spill occurred.

Practical info

Management of blood and body substance spills

Prompt removal of spots and spills of blood and body substance followed by cleaning and disinfection of the area contaminated is a sound infection control practice and meets occupational health and safety requirements [111].

In circumstances where emergency procedures or urgent transport are under way, spills should be attended to as soon as it is safe to do so. Further advice should be sought from discipline specific guidelines.

Process of spills management

Strategies for decontaminating spills of blood and other body substances (e.g. vomit, urine) differ based on the setting in which they occur and the volume of the spill:

- healthcare workers can manage small spills by cleaning with detergent solution
- for spills containing large amounts of blood or other body substances, workers should contain and confine the spill by:
 - removing visible organic matter with absorbent material (e.g. disposable paper towels)
 - removing any broken glass or sharp material with forceps
 - soaking up excess liquid using an absorbent clumping agent (e.g. absorbent granules).

If spillage has occurred on soft furnishings, a detergent solution can be used to clean the area thoroughly. Do not clean soft furnishings with a disinfectant such as sodium hypochlorite.

Soft furnishings can also be wet vacuumed. Following cleaning of soft furnishings, they must be allowed to dry before reuse.

Alcohol solutions should not be used to clean spillages.

Table 6. Appropriate processes for managing spills

| Volume of spill | Process |
|--|--|
| Spot cleaning | <ul style="list-style-type: none"> • Select appropriate personal protective equipment (PPE) • Wipe up spot immediately with a damp cloth, tissue or paper towel • Discard contaminated materials • Perform hand hygiene |
| Small spills (up to 10cm diameter) | <ul style="list-style-type: none"> • Select appropriate PPE • Wipe up spill immediately with absorbent material • Place contaminated absorbent material into impervious container or plastic bag for disposal • Clean the area with warm detergent solution, using disposable cloth or sponge • Wipe the area with sodium hypochlorite and allow to dry • Perform hand hygiene |
| Large spills (greater than 10cm diameter) | <ul style="list-style-type: none"> • Select appropriate PPE • Cover area of the spill with an absorbent clumping agent and allow to absorb • Use disposable scraper and pan to scoop up absorbent material and any unabsorbed blood or body substances • Place all contaminated items into impervious container or plastic bag for disposal • Discard contaminated materials • Mop the area with detergent solution • Wipe the area with sodium hypochlorite and allow to dry • Perform hand hygiene |

Choosing a disinfectant (when required)

The use of sodium hypochlorite is not necessary for routinely managing spills but it may be used in specific circumstances. There is evidence supporting the use of sodium hypochlorite to inactivate various blood borne and gastrointestinal viruses [110], and to clean rooms of patients known or suspected to be infected with bacteria such as *C. difficile* or multi-resistant organisms. The consideration to use sodium hypochlorite should be based on risk assessment of the environment, the spill, risk of transmission of disease, and the surface area and potential hazards with using the product.

If a disinfectant is required, particularly during the implementation of transmission-based precautions, a Therapeutic Goods Administration (TGA)-listed hospital grade disinfectant with specific claims or sodium hypochlorite (where indicated for use as per *Recommendations 13 and 14*) must be used. The disinfectant chosen must have label claims against the organism of concern.

Choosing a disinfectant that is compatible with the surface material where the spill has occurred is integral in order to avoid damage to the surface.

Spill kit

A spill kit should be readily available in each clinical area and should include a scoop and scraper, single-use gloves, protective apron, surgical mask and eye protection, absorbent agent, clinical waste bags and ties, and detergent. All parts should be disposable to ensure that cross-contamination does not occur.

For practical information on spills management, see Case Study 5.5.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of decontaminating after a spill clearly outweigh any undesirable effects.

Healthcare associated infections are a major threat to patient safety and appropriate site decontamination after spills of blood or other potentially infectious materials can help to reduce the spread of infection.

Certainty of the evidence

Low

There is supportive evidence and a strong theoretical rationale for this intervention.

The Centers for Disease Control and Prevention guidelines [115] give Category 1B and 1C evidence in support of this recommendation.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections by decontaminating after a spill.

Resources and other considerations

No important issues with the recommended alternative

This practice would entail the provision of approved cleaning agents, which has been part of standard care for many years.

Staff training and/or tools such as checklists in wards, may be necessary to ensure uptake of advice.

Rationale

The prompt decontamination of an area contaminated by either blood or body substances is a sound infection control practice and a work health and safety requirement.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Practice Statement

New

13. It is good practice to use a chlorine-based product such as sodium hypochlorite or a Therapeutic Goods Administration-listed hospital-grade disinfectant with specific claims in addition to standard cleaning practices to effectively manage norovirus specific outbreaks.

Practical info

Norovirus epidemiology

Norovirus is the most frequently occurring cause of community-acquired acute gastroenteritis in people of all ages. It is one of the most common causes of outbreaks in healthcare settings, affecting both long-term care facilities and acute care hospitals [96]

Norovirus belongs to the family Caliciviridae and are a single-stranded RNA, non-enveloped virus that can cause gastroenteritis in humans [118]. Noroviruses are divided into at least six genogroups (GI-GVI) and further subdivided into more than 38 genotypes based on phylogenetic analysis of the major capsid protein [118] [122]. Currently, human noroviruses belong to one of three norovirus genogroups which are further divided into more than 25 genetic clusters [119].

Human noroviruses cannot be grown in cell culture [123], therefore, diagnostic methods focus on detecting viral RNA or antigen. 17 studies (81%) identified that NoV genotype GII.4 have caused the majority of clinical outbreaks in healthcare settings during the past decade.

Norovirus infections generally have a shorter incubation period and are characterised by acute onset of nausea, vomiting, abdominal pain and diarrhoea [63]. The mean duration of symptoms is 2-3 days.

Transmission pathways

Transmission for norovirus infections in healthcare settings mainly occur by the faecal–oral route, either through person to person contact or through exposure to contaminated food.

Whilst some observational studies have suggested there is a possibility of viral transmission via aerosols, there is no data or determination criteria to support this assumption.

It appears that Genotype GII.4 is more often associated with transmission mediated by person-to-person contact than with other types of transmission.

Individuals may shed norovirus more than 21 days after the resolution of symptoms, possibly acting as a possible source for nosocomial transmission. However no data has been reported on ongoing transmission or secondary cases.

Environmental cleaning

During a norovirus outbreak, routine cleaning should be intensified with the use of a detergent solution followed by the use of a Therapeutic Goods Administration-listed hospital-grade disinfectant with specific claims so that surfaces are cleaned and disinfected. When using sodium hypochlorite (i.e. bleach) as a disinfectant in addition to standard cleaning practice to manage norovirus outbreaks, a minimum dilution factor of 1:10 should be used, equivalent to 0.1% active chlorine.

When using sodium hypochlorite to disinfect hard surfaces, the following should be considered:

- environmental surfaces should be clean and free of matter
- allow sufficient time to kill the virus – at least 10 minutes surface contact time
- dilution of sodium hypochlorite (10% bleach solution) should be made up fresh, just before use (1:10 solution = 1 part bleach for every 9 parts water).

This concentration and advice is aligned with the advice provided in the *Infection control guidelines for the public health management of gastroenteritis outbreaks due to norovirus or suspected viral agents in Australia* [361].

Evidence to decision

| | |
|----------------------------------|--|
| Benefits and harms | Whilst the benefits of sodium hypochlorite use outweighs the harms, there are potential adverse effects involved in using such products. However, these can be minimised by safe handling in accordance with the Safety Data Sheets. |
| Certainty of the evidence | The evidence for this practice statement is limited or inconsistent. This practice statement is based on the advice of experts and trends in clinical practice. |
| Values and preferences | There are no acceptability considerations which directly impact the patient for the use of sodium hypochlorite. |

The routine use of sodium hypochlorite should not impact on patient health equity.

Resources and other considerations Sodium hypochlorite is inexpensive and its feasibility has been demonstrated by widespread existing use.

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. The use of sodium hypochlorite to manage norovirus-specific outbreaks is justified to reduce healthcare associated infection.

3.1.3.1 Emerging disinfection methods

Emerging modes of disinfection

Some modes of disinfection have emerged and undergone further development for use in healthcare facilities in recent years.

These include:

- sodium hypochlorite (considered emerging as it is not part of routine cleaning procedures)
- ultra-violet light
- hydrogen peroxide vapour
- electrolysed water.

With the exception of using sodium hypochlorite during terminal cleans of rooms contaminated with known or suspected *C. difficile* and multi-resistant organisms, the evidence of the effects of these emerging disinfection methods on clinical outcomes remains sparse. If emerging disinfectants are used in healthcare facilities, this should always be used in addition to standard cleaning practices.

There is also an emerging trend towards using steam and microfiber clothes for environmental cleaning as an alternative to other disinfection methods. The evidence on the use of steam and microfiber cloths on patient-centred outcomes including patient colonisation and healthcare associated infection were not systematically reviewed and therefore no recommendation on use is made in this Guideline. However, concerns that steam technology may only be practical to use on specific surfaces, and may spread infectious organisms to the nearby environment leading to further contamination, are documented in the literature [126]. Healthcare facilities using microfiber cloths and steam should ensure they have appropriate monitoring systems in place.

Healthcare facilities that choose to use steam and microfibre cloths for environmental cleaning need to ensure staff are well educated and trained in its use, and that infection prevention staff and other healthcare professionals understand the methodology and how to implement it.

Further information on emerging environmental cleaning technologies can be found [here](#).

Antimicrobial surfaces

Other disinfectant modalities have emerged or undergone further development for use in healthcare facilities subsequent to the review of the evidence for the 2010 Guidelines. This includes the effects of self-disinfecting materials used to coat or impregnate surfaces in patient care areas. These materials include heavy metal alloys (copper and silver), light activated antimicrobial coatings, and surfaces with altered topography designed to inhibit bacterial growth.

Currently, there is only sufficient evidence to examine the effectiveness of copper-based surfaces compared to standard surfaces on hospital acquired infection. However, the quality of the evidence is very low; as a result, copper-coated surfaces are currently not recommended for use, nor are other antimicrobial surfaces or fittings.

Future research

The inclusion of infection and colonisation outcomes in future studies is key to determining whether these interventions have a clinically important impact; this is an important gap in existing research which has largely focused on whether emerging disinfectants reduce bacterial contamination. Studies among high-risk populations are also needed (e.g. oncology, burns), as investment in emerging disinfectant modalities is expensive and their use is therefore likely to be prioritised in areas of highest risk.

The optimal design to assess the causal effects of an intervention is one which involves random allocation of individuals or clusters of individuals to treatment groups. Individually randomised trials are unlikely to be possible in this setting because of the risk of

contamination between treatment groups. For example, patients allocated to control rooms may be moved to intervention rooms during the course of their admission. Cluster randomisation by ward may reduce contamination between treatment groups, but will not resolve the issue completely because of movement of patients between wards. Therefore, the optimal design would be a cluster randomised trial where clusters are hospitals.

For further information on choosing an appropriate disinfectant, see *Recommendations 9 and 12*.

Conditional recommendation

New

14. It is suggested that sodium hypochlorite disinfection be used as an adjunct to standard cleaning in healthcare facilities.

The use of sodium hypochlorite disinfection in addition to a detergent solution is suggested for terminal cleans of rooms of patients known or suspected to have *C. difficile* associated disease or multi-drug resistant organisms.

The use of sodium hypochlorite disinfection in addition to detergent solution is suggested to terminate outbreaks of *C. difficile*.

Practical info

Sodium hypochlorite

Despite the emergence of new disinfection products and technologies, sodium hypochlorite remains a commonly used and accessible chlorine-based disinfectant with broad spectrum antimicrobial properties. The evidence suggests that when the dilution factor is sufficient for sporicidal activity ($\geq 5000\text{ppm}$ free chlorine), sodium hypochlorite is effective against *C. difficile*. There is also evidence to suggest that sodium hypochlorite disinfection is effective for managing norovirus outbreaks [128] —see Section 3.1.3 *Practice Statement 13*.

From a work health and safety perspective, sodium hypochlorite should be used as per manufacturer instructions as it may cause irritation to the skin, eyes and other mucous membranes. It can also corrode metals and discolour or stain fabrics.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

Whilst the benefits of sodium hypochlorite use outweighs the harms, there are potential adverse effects involved in using such products. However, these can be minimised by safe handling in accordance with the Safety Data Sheets.

Certainty of the evidence

Very low

The dilution of sodium hypochlorite in the studies suitable for inclusion in the systematic review may not be what is being used in practice and the variation of concentrations used in the studies may affect the outcome. The overall quality of evidence was rated as very low [128]. Three very low quality studies reviewed infection acquisition rates. One randomised trial examined the effects of sodium hypochlorite for terminal room disinfection on infection and colonisation incidence rates together, however there was serious imprecision and consistency cannot be assessed [128].

Values and preferences

No substantial variability expected

There are no acceptability considerations which directly impact the patient for the use of sodium hypochlorite.

The routine use of sodium hypochlorite should not impact on patient health equity.

Resources and other considerations

No important issues with the recommended alternative

Sodium hypochlorite is inexpensive and its feasibility has been demonstrated by widespread existing use.

Rationale

The evidence on the effects of sodium hypochlorite disinfection (daily and terminal disinfection) on clinical outcomes is sparse. Three very low quality studies reviewing infection acquisition rates and one moderate quality study reviewing infection and colonisation incidence rates suggest the effect on *C. difficile* and multi-resistant organisms (MROs) of more frequent cleaning with sodium hypochlorite compared to terminal cleaning is not known, however sodium hypochlorite may be used in terminal cleans due to its effectiveness on *C. difficile* and MROs [128].

Low quality evidence supports the use of routine sodium hypochlorite cleaning, compared to standard cleaning with detergents, in terminating outbreaks of *C. difficile* [128].

The body of evidence informing this recommendation used a minimum of 1:10 dilution, equivalent to 0.1% active chlorine, preceded by a detergent solution. This concentration and advice is aligned with the advice provided in the Infection control guidelines for the public health management of gastroenteritis outbreaks due to norovirus or suspected viral agents in Australia [308].

Conditional recommendation against

New

15. The effectiveness of hydrogen peroxide vapour disinfection as an adjunct to routine cleaning in healthcare facilities is yet to be established. Therefore routine use is not suggested in healthcare facilities.

Hydrogen peroxide vapour may be considered in high-risk settings and during outbreaks when other disinfection options have been exhausted.

Practical info

Current use of emerging disinfectants

Overall, the evidence of the effects of these emerging disinfection methods on clinical outcomes remains sparse. If emerging disinfectants are used in healthcare facilities, this should always be used in addition to standard cleaning practices [128][130].

Hydrogen peroxide vapour

Hydrogen peroxide has microbicidal properties against multiple pathogens, including *C. difficile*. Automated (no touch) systems for producing hydrogen peroxide vapour and hydrogen peroxide mist are designed to disinfect by dispersing vapour or mist evenly across a room. As with ultra-violet light, the systems can only be used when rooms are vacated [125]. Rooms and ventilation systems must be sealed to prevent exposure, and hydrogen peroxide must be monitored to ensure safe levels outside the room during disinfection, and within the room before re-entering.

There is not yet enough high quality evidence to determine whether the benefits of using hydrogen peroxide vapour for infection prevention and control outweigh the harms.

Evidence to decision

Benefits and harms

Small net benefit, or little difference between alternatives

As there is limited evidence on the effects of hydrogen peroxide vapour, the benefits of its use for infection prevention and control in addition to standard cleaning procedures cannot be evaluated. There are potential risks of using hydrogen peroxide vapour, however these can be minimised by safe handling in accordance with Safety Data Sheets and manufacturer's instructions for use. If hydrogen

peroxide vapour disinfection is used, risk mitigation strategies should be implemented to ensure risks to staff and patients are reduced.

Certainty of the evidence

Very low

The overall quality of evidence for the use of hydrogen peroxide vapour in healthcare facilities is very low [128]. None of the three included studies in the review were randomised trials, and one study was at high risk of bias due to industry and financial ties and reported no safeguards to protect against bias [128].

Values and preferences

No substantial variability expected

There are no acceptability considerations for hydrogen peroxide vapour which directly impact patients.

Routine use of hydrogen peroxide vapour should not impact on patient health equity.

Resources and other considerations

Important issues, or potential issues not investigated

There are significant costs associated with hydrogen peroxide vapour, and the cost-effectiveness of this practice is yet to be conclusively established. These factors in addition to implementation issues affect its feasibility.

Rationale

The evidence on the effects of hydrogen peroxide vapour disinfection on clinical outcomes is sparse. There is very low quality evidence (n=1) reviewing infection acquisition rates and high cost associated with the use of hydrogen peroxide vapour, therefore routine use is not recommended [128].

Overall, the evidence on infection and colonisation incidence rates contributing to this review was of very low quality, due to the small number of studies (n=3) and serious risk of bias. However, the limited evidence does suggest that hydrogen peroxide vapour may be considered in high risk settings in outbreaks [128] [130].

Conditional recommendation against

New

16. The effectiveness of ultra-violet light disinfection as an adjunct to routine terminal cleaning in healthcare facilities is yet to be established. Therefore routine use is not suggested in healthcare facilities.

Ultra-violet light disinfection may be considered in high-risk settings and during outbreaks when other disinfection options have been exhausted.

Practical info

Current use of emerging disinfectants

Overall, the evidence of the effects of these emerging disinfection methods on clinical outcomes remains sparse. If emerging disinfectants are used in healthcare facilities, this should always be used in addition to standard cleaning practices [128][130].

Ultra-violet light

Ultra-violet light in the UV-C wavelength range (200 to 270 nanometers) has microbicidal properties against multiple

pathogens, including *C. difficile* and other healthcare associated pathogens.

Technologies have been developed for automated (no-touch) disinfection of hospital rooms using ultra-violet light. The technologies only disinfect areas directly in the ultra-violet light and can only be used when rooms are vacated, partly because of the potentially harmful effects of ultra-violet exposure [125].

Whilst there is some evidence to demonstrate the ultra-violet light disinfection can be effective for infection prevention and control, the magnitude of the benefits is yet to be established and it is unknown whether these outweigh the harms [128][130].

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

There is evidence of the benefit of ultra-violet light, however there are known risks involved when using ultra-violet light as a disinfection method. Ultra-violet light can be harmful to healthcare workers if they enter the room during operation as toxic levels can be reached within three seconds of exposure.

Certainty of the evidence

Low

The evidence for the effectiveness of ultra-violet light for terminal cleaning in healthcare facilities is low [128].

One study [127] was high quality and suggested that ultraviolet light can be effective in reducing infection rates in hospitalised populations. However, the study was graded down due to imprecision from large confidence intervals. These findings require replication in other contexts, and data for high risk groups are needed.

Values and preferences

No substantial variability expected

There are no acceptability considerations related to ultra-violet light which would directly impact a patient.

The routine use of ultra-violet light should not impact on patient health equity.

Resources and other considerations

Important issues, or potential issues not investigated

The magnitude of benefit of ultra-violet light is yet to be established, so the cost effectiveness of these interventions is likely to vary according to the baseline risks of healthcare associated infections due to multi-resistant organisms. These factors in addition to the upfront costs and implementation issues affect the feasibility of using ultra-violet light. The feasibility of its use also depends on the design of the healthcare facility.

Rationale

The evidence on the effects of ultra-violet light disinfection on clinical outcomes is sparse despite the recent publication of a low quality, large cluster randomised trial [127] reviewing infection acquisition rates. This single study suggests that ultra-violet light may be effective for multi-resistant organisms (MRO) disinfection but its implementation has significant workplace health and safety considerations and feasibility issues relating to facility design and upfront costs.

Overall the evidence on infection and colonisation incidence rates contributing to this review was of low quality, due to the small number of studies (n=2) and serious imprecision [128], however the limited evidence does suggest that ultra-violet light may be considered in high risk settings in MRO outbreaks for terminal cleaning when other disinfection options have not been effective [128][130]. Ultra-violet light is not shown to be effective for *C. difficile* by comparison to the more inexpensive disinfectant, sodium hypochlorite.

Conditional recommendation against

New

17. The effectiveness of ultra-violet light disinfection in combination with sodium hypochlorite for terminal cleaning in healthcare facilities is yet to be established. Therefore routine use is not suggested in healthcare facilities.

Ultra-violet light disinfection in combination with sodium hypochlorite may be considered in high-risk settings and during outbreaks when other disinfection options have been exhausted.

Practical info

Current use of emerging disinfectants

Overall, the evidence of the effects of these emerging disinfection methods on clinical outcomes remains sparse. If emerging disinfectants are used in healthcare facilities, this should always be used in addition to standard cleaning practices [128].

Ultra-violet light with sodium hypochlorite

Ultra-violet light in the UV-C wavelength range (200 to 270 nanometers) has microbicidal properties against multiple pathogens, including *C. difficile* and other healthcare associated pathogens.

Technologies have been developed for automated (no-touch) disinfection of hospital rooms using ultra-violet light. The technologies only disinfect areas directly in the ultra-violet light and can only be used when rooms are vacated, partly because of the potentially harmful effects of ultra-violet exposure [152].

Sodium hypochlorite is a commonly used chlorine-based disinfectant with broad spectrum antimicrobial properties. Sodium hypochlorite may cause irritation to the skin, eyes and other mucous membranes. It can also corrode metals and discolour or stain fabrics.

Evidence to decision

Benefits and harms

Small net benefit, or little difference between alternatives

There is evidence of the benefit for the combined use of ultra-violet disinfection and sodium hypochlorite. However, there are known risks involved in the use of both ultra-violet disinfection and sodium hypochlorite individually, so risk management strategies addressing their concurrent use need to be put in place.

Certainty of the evidence

Moderate

One study in the systematic review [128] investigated the effectiveness of using ultra-violet light disinfection and sodium hypochlorite. However, this study reviewed the composite outcome of infection and colonisation (rated as an important outcome), unlike the other interventions in the review which provided evidence for the critical outcome of infection only.

Values and preferences

No substantial variability expected

There are no acceptability considerations which directly impact the patient.

The combined use of ultra-violet light disinfection and sodium hypochlorite should not impact on patient health equity.

Resources and other considerations

Important issues, or potential issues not investigated

The effectiveness of the combined use of ultra-violet light disinfection and sodium hypochlorite is yet to be conclusively established. Sodium hypochlorite is inexpensive, so if it is effective it is likely to be a cost saving intervention. However, ultra-violet light is expensive and its effectiveness is likely to vary according to the baseline risks of healthcare associated infections due to multi-resistant organisms. Upfront costs, implementation issues and facilities design also affect the feasibility of using ultra-violet light disinfection in combination with sodium hypochlorite.

Rationale

The evidence on the effects of ultra-violet light disinfection in combination with sodium hypochlorite disinfection on clinical outcomes is sparse despite the recent publication of a moderate quality large cluster randomised trial [127] reviewing infection and colonisation incidence rates. This single study suggests the combination of ultra violet light disinfection and sodium hypochlorite may be effective for multi-resistant organism (MRO) disinfection but its implementation has significant work health and safety considerations and feasibility issues relating to facility design and upfront costs.

Overall the evidence on infection and colonisation incidence rates contributing to this review [128] was of moderate quality, with the limited evidence (n=1) suggesting that ultra-violet light disinfection in combination with sodium hypochlorite disinfection may be considered in high risk settings in MRO outbreaks for terminal cleaning when other disinfection options have been exhausted. It is important to note that in isolation, ultra-violet light disinfection is not shown to be effective for *C. difficile* by comparison to the less expensive disinfectant, sodium hypochlorite.

Conditional recommendation against

New

18. The effectiveness of surfaces, fittings or furnishing containing materials with antimicrobial properties in healthcare facilities is yet to be established. Therefore routine use is not suggested in healthcare facilities.

Practical info

Current use of antimicrobial surfaces

Overall, the evidence of the effects of antimicrobial surfaces on clinical outcomes remains sparse. If antimicrobial surfaces are used in healthcare facilities, this should always be used in addition to standard cleaning practices [129].

Antimicrobial surfaces

The use of surfaces, fittings or furnishing containing materials with antimicrobial properties has been suggested to reduce the concentration of bacteria on surfaces, in turn reducing environmental exposure to pathogens. Self-disinfecting materials that are considered for use in healthcare facilities include the use of heavy metal alloy coatings on fittings (e.g. copper or silver coatings for bed rails, tray tables or IV stands). There is currently limited evidence to support the use of environmental fittings with antimicrobial properties to prevent infection [129].

Evidence to decision

Benefits and harms

Small net benefit, or little difference between alternatives

The limited evidence on the use of copper-surfaced objects in health care facilities means that the potential harms and benefits cannot be assessed.

Certainty of the evidence

Very low

The evidence for the effectiveness of copper-surfaced fittings is of very low quality [129]. Only two studies were identified for inclusion, both of which were non-randomised and had high risks of bias

and serious imprecision [129]. One study reviewed infection acquisition rates while the other considered infection and colonisation incidence rates together.

There were no eligible studies identified of any other antimicrobial surfaces.

Values and preferences

No substantial variability expected

No important uncertainty or variability was identified, and no acceptability considerations that directly impact the patient were identified.

Resources and other considerations

Important issues, or potential issues not investigated

The effectiveness of antimicrobial surfaces is yet to be established, so the cost effectiveness of these interventions is unknown. The actual cost and feasibility may vary depending on the context - i.e. new buildings versus retrofitting hospitals and wards.

Rationale

The evidence on the effects of copper surfaces on healthcare associated infection is sparse. With only one non-randomised trial reviewing infection acquisition rates with uncertain results, it is not possible to draw conclusions from this evidence [129].

Overall the evidence on infection and colonisation incidence rates contributing to this review was of very low quality [129], due to the small number of studies (n=2) and imprecise estimates of effect observed in both included studies. Both were non-randomised studies and were at risk of biases that further reduced certainty about the effects observed.

There were no eligible studies identified of any other antimicrobial surfaces [129].

3.1.4 Reprocessing of reusable medical devices

This section gives core principles for reprocessing of reusable medical devices (RMDs - instruments and equipment) in any healthcare setting. Unless exempt, reusable medical devices must be 'included' onto the Australian Register of Therapeutic Goods (ARTG) before they may be supplied in or exported from Australia. Healthcare facilities should develop local policies and procedures relevant to their setting and may also need to consult relevant Australian Standards and discipline-specific guidelines for further advice on reprocessing requirements. Further information is contained in **Standard AS 5369:2023** *Reprocessing of reusable medical devices and other devices in health and non-health related facilities*, or equivalent international standards e.g. International Organisation for Standardisation (ISO) or European Standard (EN).

What are the risks?

Any infectious agents introduced into the body can establish infection. In all healthcare settings, reusable medical devices should be handled in a manner that will prevent patient, healthcare worker and environmental contact with potentially infectious material.

Principles of reprocessing reusable medical devices include [133]:

- Only Therapeutic Goods Administration (TGA)-included reusable medical devices should be used; before purchase, healthcare facilities should ensure that manufacturer's reprocessing instructions are provided and are able to be followed by the healthcare facility.
- All reusable medical devices and patient-care equipment used in the clinical environment must be reprocessed according to their intended use and manufacturer's advice.
- Single-use medical devices must not be reprocessed.
- If a healthcare facility makes a decision to reprocess single-use devices, the facility must be licensed by the TGA and:
 - will be considered a manufacturer under section 41BG(2) of the *Therapeutic Goods Act 1989*
 - will be subject to audit for conformance.

Assessing the degree of risk

Any medical device (instruments and equipment) that is to be reused and requires reprocessing—cleaning, disinfection and/or sterilisation. The minimum level of reprocessing required for reusable instruments and equipment depends on the individual situation and manufacturer

instructions (i.e. the body site and the nature by which the instrument will be used).

The approach to disinfection and sterilisation of patient-care items and equipment devised by Spaulding in 1957 has been retained and refined and is still successfully used by infection control professionals and others when planning methods for disinfection or sterilisation [132]. The system is based on instruments and items for patient care being categorised into critical, semi-critical and non-critical, according to the degree of risk for infection involved in use of the items.

Table 7. Categories of items for patient care

| Category | Description |
|----------------------|--|
| Critical | These items confer a high risk for infection if they are contaminated with any microorganism and must be sterile at the time of use. This includes any objects that enter sterile tissue or the vascular system, because any microbial contamination could transmit disease. |
| Semi-critical | These items come into contact with mucous membranes or non-intact skin, and should be single use or sterilised after each use. If this is not possible, high-level disinfection is the minimum level of reprocessing that is acceptable. |
| Non-critical | These items come into contact with intact skin but not mucous membranes. Thorough cleaning is sufficient for most non-critical items after each individual use, although either intermediate or low-level disinfection may be appropriate in specific circumstances. |

The Australian Commission on Safety and Quality in Health Care has issued [Advisory DI21/04](#) to provide guidance on the disinfection requirements for reusable semi-critical medical devices for the Diagnostic Imaging Accreditation Scheme Standards. The Advisory and these Guidelines are consistent with guidance issued by the [Australian Society for Ultrasound in Medicine \(ASUM\)](#) about categorisation of ultrasound probes used in diagnostic imaging as semi-critical reusable medical devices that require high-level disinfection after each use.

High-level disinfection can be achieved with a disinfectant included on the [Australian Register of Therapeutic Goods \(ARTG\)](#) as a Class IIb medical device. The disinfectant may be in the form of a liquid, spray, wipe, or aerosol intended to be used on medical devices and consistent with the manufacturer’s instructions for use.

As per the ASUM guidance, due to the reported breakage rate of transducer covers, for maximum safety, cleaning and high-level disinfection of the transducer is recommended between each use.

Computers, portable mobile devices and personal digital assistants used in patient care are classified as non-critical patient care items [132]. It is important that these items are included in policies for cleaning non-critical items.

Surfaces barriers such as keyboard covers and washable keyboards that can be easily cleaned may help prevent contamination of surfaces and equipment. These should be correctly used, as per the manufacturer instructions and changed or cleaned between patients. Further information on surface barriers is available in Section 3.1.3 *Practice Statement 11*.

Cleaning

Cleaning is the removal of foreign material (e.g. soil and organic material) from objects and is normally carried out using detergent, water and physical action.

Cleaning to remove organic material must always precede high-level disinfection and sterilisation of critical and semi-critical instruments and devices as residual proteinaceous material reduces the effectiveness of the disinfection and sterilisation processes. If an item cannot be cleaned, it cannot be disinfected or sterilised.

Instruments should be cleaned as soon as practical after use (e.g. preferably at the point of use) before soiled materials become dried onto the instruments. Dried or baked materials on the instrument make the removal process more difficult, the disinfection or sterilisation process less effective or ineffective and can damage the RMD.

Instruments that can be disassembled must be disassembled before the cleaning and the disinfection/sterilisation process.

Methods of cleaning

Automated

Automated cleaners (ultrasonic cleaners and washer-disinfectors) reduce the handling of instruments and are recommended for cleaning basic instruments that can withstand the process.

- Ultrasonic cleaners work by subjecting instruments to high frequency, high-energy sound waves, thereby loosening and dislodging dirt.
- Washer-disinfectors use cleaning agents at predetermined high temperatures and time periods to clean reusable medical devices. When a washer-disinfectant is used, care should be taken in loading instruments: hinged instruments should be opened fully to allow adequate contact with the cleaning agent; over loading of reusable medical devices in washer disinfectors should be avoided; items should be placed in a way that prevents tipping and collection of water; and instruments should be disassembled.

Manual

Cleaning is done manually for fragile or difficult-to-clean reusable medical devices and in settings without automatic units. Where manual cleaning methods are used, these should comply with **AS 5369:2023**, or equivalent international standards e.g. International Organisation for Standardisation (ISO) or European Standard (EN), whichever is relevant to the type of healthcare facility.

The two essential components of manual cleaning are:

- friction—rubbing/scrubbing the soiled area with an appropriately sized soft brush
- fluidics—use of fluids to remove soil and debris from internal channels after brushing with an appropriately sized brush and when the design does not allow passage of a brush through a channel.

Healthcare workers should wear appropriate PPE for the task—plastic apron, utility gloves and face protection (protective eyewear and mask or face shield). Care should be taken to prevent aerosols, splashes to mucous membranes or penetration of the skin by sharp instruments.

Cleaning agents

The cleaning agent and style must be appropriate for each instrument and piece of equipment. The manufacturer's instructions will guide the type of cleaning agent required. Different cleaning agents may be used for manual cleaning, as opposed to cleaning in a washer-disinfector or ultrasonic cleaner. Standard AS 5369:2023 provides further guidance for all cleaning agents used for reprocessing of reusable medical devices. A cleaning agent should only be used if it is listed on the ARTG and intended for use on medical devices. Cleaning agents should be neutral pH or mildly alkaline as such solutions generally provide the best material compatibility profile and good soil removal; mildly acidic solutions may damage instruments. Where multiple chemicals are used, they should be compatible with each other. Where possible, the cleaning agent should be biodegradable.

Enzymes, usually proteases (enzymes active on proteins), are sometimes added to neutral pH solutions to assist in removing organic material such as blood and pus. Cleaning agents can also contain lipases (enzymes active on fats) and amylases (enzymes active on starches). Enzymatic cleaners are not disinfectants, and enzymes present in cleaning agents may be inactivated when used in conjunction with some germicidal compounds.

As with all chemicals, enzymes must be rinsed from the equipment or adverse reactions could result.

Checking effectiveness of cleaning

The most common means of monitoring the efficacy of the cleaning process for reusable medical devices is by thorough visual inspection following cleaning. However, for complex reusable medical devices, visual inspection may be insufficient to monitor cleaning efficacy. Commercially available surrogate devices carrying test soil (e.g. protein, blood) may be used to challenge the cleaning process. Alternatively, medical devices may be sampled for residual contaminants (e.g. protein, ATP) immediately after reprocessing. AS 5369:2023 requires that cleaning processes are validated by both visual and objective means.

International Standard **ISO 15883-5: 2005 (Washer-Disinfectors, Part 5)**, outlines specific test methods to check the effectiveness of cleaning to verify manual and automated processes. At a minimum, all instruments should be individually inspected (with magnification where possible) and be visibly clean. More information on checking the effectiveness of cleaning is available in 3.1.3 - Routine management of the physical environment.

Disinfection

Disinfection is a process that inactivates non-sporing infectious agents. Disinfection processes may include automated, high-level disinfection systems using either, chemical, heat or light-based methods. Items need to be cleaned before being disinfected.

(**source:** *Therapeutic Goods Regulations 1990*, Statutory rules No.394, 1990. (accessed 16 August 2023) legislation.gov.au)

Instruments should be removed from the disinfectant after reprocessing and stored dry. To preserve the surfaces of the instruments, dissimilar metals should be separated before cleaning.

- *Thermal disinfection* uses heat and water, at temperatures that destroy infectious agents and is appropriate for items that are heat and moisture resistant and do not require sterilisation. Thermal disinfection, is the simplest, most efficient and cost-effective method of disinfection. It can be achieved in an automated thermal washer-disinfector by choosing the appropriate cycle.
- *Chemical disinfection* can be achieved with a compatible TGA-included sterilant or medical device disinfectant, used alone or together with an automated washer-disinfector. A chemical disinfectant used to reprocess a reusable medical device or other device shall be labelled as an "instrument grade disinfectant". Disinfectants need to be listed on the ARTG and for high-level disinfection, only disinfectants listed as Class IIb devices should be used. Other classes of chemical disinfectants, e.g. hospital-grade disinfectant, must not be used to reprocess a reusable medical or other devices (source: AS 5369:2023). Chemical disinfectants include alcohols, chlorine and chlorine compounds, formaldehyde, hydrogen peroxide, phenolics and quaternary ammonium compounds. Commercial formulations based on these chemicals are considered unique products and must be 'included' onto the Australian Register of Therapeutic Goods (ARTG). In most instances, each product is designed for a specific purpose; therefore, users should read labels carefully to ensure the correct product is selected for the intended use and applied efficiently.

There are three levels of disinfection, depending on the intended use of the instruments:

- **High-level disinfection**—a disinfectant that kills all microbial pathogens, except large numbers of bacterial endospores when used as recommended by manufacturer. High-level disinfection is the minimum grade disinfectant used for disinfection of a semi-critical reusable medical devices (instruments and equipment).
- **Intermediate-level disinfection**—a disinfectant that kills all microbial pathogens except bacterial endospores, when used as recommended by the manufacturer. This process is effective against most bacterial, viral and fungal spores (source: AS 5369:2023).
- **Low-level disinfection**—a disinfectant that rapidly kills most vegetative bacteria as well as medium sized lipid containing viruses, when used as recommended by the manufacturer. It cannot be relied upon to destroy, within a practical period, bacterial endospores, mycobacteria, fungi or all small nonlipid viruses. Intermediate and low-level disinfection is the minimum grade disinfectant used for disinfection of a non-critical reusable medical or other devices where required.

Disinfection is not a sterilising process. Wherever possible, sterilise items to be used in semi-critical sites, or employ single-use items.

Sterilisation

Sterilisation destroys all microorganisms on the surface of an instrument or device, to prevent disease transmission associated with the use of that item. While the use of inadequately sterilised critical items represents a high risk of transmitting infectious agents, documented transmission associated with an inadequately sterilised critical item is rare. This is probably due to the wide safety margin associated with the sterilisation processes used in healthcare facilities.

- Reprocessing of heat resistant items is recommended by steam sterilisation due to the safety margin, reliability, validity and lethality.
- Reprocessing heat and moisture-sensitive items requires use of a low-temperature sterilisation technology (e.g. ethylene oxide, hydrogen peroxide plasma, peracetic acid and aldehyde).

Sterilisation methods are designed to give a sterility assurance level (SAL) of at least 10^{-6} , provided the sterilisation process is validated by the user. Records of sterilisation must also be kept to verify that an appropriate reprocessing system is in place according to state and federal legislation. Details of the documentation required can be found in **Standard AS 5369:2023**.

In this rapidly changing area, reprocessing standards evolve to accommodate changes in equipment design and emerging technologies in sterilisation.

Medical device sterilants and disinfectants

Only sterilant or medical device disinfectants listed on the ARTG should be used on medical devices. To be considered as ARTG listed, sterilants and disinfectants must comply with the essential principles for quality, safety and performance through using appropriate testing regimes for performance. Refer to [TGA website](#).

Traceability

Traceability is the ability to trace the reprocessing history, use and location of a reusable medical device. At a minimum, traceability systems are required to enable reusable medical devices and patients to be linked at the point of use, and to allow recall of non-conforming products (source: AS 5369:2023).

At a minimum, the traceability process should clearly identify and record the:

- Patient for whom the device/s was used on
- Type of procedure, including date and time
- Equipment, instruments and devices that were used during the procedure
- Reprocessing history.

Processes for traceability should be included in the organisation's policies and procedures for reprocessing of critical and semi-critical reusable medical devices and other devices.

Traceability processes apply to all reusable medical devices and other devices classified as:

- Critical reusable medical devices and other devices undergoing sterilisation, including loan and trial sets, implantable devices/equipment
- Semi-critical reusable medical devices and other devices undergoing high-level disinfection, such as trans-rectal ultrasound probe, anaesthetic equipment/airway equipment

For more information on Traceability, refer to AS 5369:2023 Section 2.5.3 Identification and traceability of product.

Storage and maintenance

All items must be stored in a way that maintains their level of reprocessing (e.g. sterile, high level disinfected). Dry, sterile, packaged instruments and equipment should be stored to prevent environmental contamination in a designated storage area. The storage areas should be clean and dry, to protect instruments from direct sunlight, heat, moisture dust, vermin and sharp objects that may damage the packaging. Further information on handling, transport and storage of reprocessed medical devices is available in **Standard AS 5369:2023**, or equivalent international standards.

Equipment and instrument surfaces should be regularly examined for breaks in integrity that would impair either cleaning or disinfection/sterilisation (this should be a documented process). Equipment that no longer functions as intended or cannot be adequately cleaned and disinfected or sterilised should be repaired or discarded.

Table 8. General criteria for reprocessing and storage of equipment and instruments in healthcare settings

| Level of risk | Process | Examples | Storage |
|--|--|--|--|
| <p>*Critical Entry or penetration into sterile tissue, cavity or blood stream</p> | <ul style="list-style-type: none"> Clean thoroughly as soon as possible after using. Sterilise after cleaning by steam under pressure. If heat or moisture sensitive, sterilise through an automated low temperature chemical sterilant system, other liquid chemical sterilants or ethylene oxide sterilisation. Ensure critical items are sterilised between each patient use. | <ul style="list-style-type: none"> Invasive surgical and dental equipment e.g. surgical oral instruments, arthroscopes, laparoscopes, rigid and flexible bronchoscopes, heat stable scopes. Implants and ultrasound probes used in sterile body cavities. | <p>Sterility must be maintained:</p> <ul style="list-style-type: none"> Packaged items must go through a drying cycle and then be checked to ensure drying has taken place before use or storage. The integrity of the wrap must be maintained. Wraps act as an effective biobarrier during storage. Unpackaged sterile items must be used immediately (without contamination in transfer from steriliser to site of use) or resterilised. All endoscopic instruments (except those in sterile packaging) should be stored in a TGA-approved forced-air drying cabinet. |
| <p>Semi-critical Contact with intact mucous membranes or non- intact skin</p> | <ul style="list-style-type: none"> Clean thoroughly as soon as possible after using. Steam sterilisation is preferable. If the equipment will not tolerate steam use a high level TGA-included chemical or thermal sterilant or medical device disinfectant. | <ul style="list-style-type: none"> Respiratory therapy and anaesthesia equipment, some endoscopes, vaginal speculae, laryngoscope blades, cystoscopes, anorectal manometry catheters, diaphragm fitting rings. Probes including transoesophageal echocardiogram, transrectal ultrasound and transvaginal probes. | <ul style="list-style-type: none"> Store to prevent environmental contamination. All endoscopic instruments (except those in sterile packaging) should be stored in a TGA-approved forced-air drying cabinet or reprocessed within set timeframes prior to use. |
| <p>Non-critical Contact with intact skin</p> | <ul style="list-style-type: none"> Clean as necessary with detergent solution. If decontamination is necessary, disinfect with compatible low or intermediate level TGA- included sterilant or medical device disinfectant after cleaning. | <ul style="list-style-type: none"> Stethoscopes, sphygmomanometers, blood pressure cuffs, mercury thermometers, non-invasive ultrasound probes. Intravenous pumps and ventilators. Noninvasive ultrasound probes (<i>not used in contact with non-intact skin or mucous membranes</i>). Commodes, bedpans, blood pressure cuffs, and crutches. | <ul style="list-style-type: none"> Store in a clean, dry place to prevent environmental contamination. |

Source: Rutala & Weber (2008) [108]

Reprocessing of flexible endoscopes

Outbreaks associated with flexible endoscopy most commonly occur due to errors in reprocessing. Reprocessing of endoscopes should be

according to the manufacturer's instructions. Staff should be aware of the number of channels and valves within the endoscope.

Table 9. Class of device and associated reprocessing method

| Class | Use | Device | Reprocessing method (minimum requirement) |
|----------------------|---|--|--|
| Semi-critical | Touches mucous membranes and non-intact skin. | Gastrointestinal endoscope (e.g. colonoscope and duodenoscope) Cystoscope | Cleaning, followed by high level disinfection. |
| Critical | Enters sterile tissue, vascular system or body space. | Snares/loops Arthroscopes Laparoscopes Bronchoscopes | Cleaning followed by sterilisation. |

Source: Public Health Agency of Canada (2010) [237]

All flexible endoscopes with channels must undergo microbiological testing. For more information on the specific reprocessing and microbiological testing requirements for flexible endoscopes, see the Gastroenterological Society of Australia's (GESA) *Infection Control in Endoscopy (2021) Guidelines* and the accompanying 2017 Consensus statements—<https://www.gesa.org.au/resources/infection-control-in-endoscopy/> [134].

Reprocessing of loan sets and privately owned sets

Loan sets and private sets need to be processed prior to use by the healthcare facility as there are no ways to verify claims that the device has been previously reprocessed. Healthcare facilities may not always have the facilities or capacity to reprocess loan sets immediately, so the reprocessing unit should be contacted prior to confirming the ability to reprocess and organising scheduling requirements [345]. This process should be informed by risk assessment.

Loan sets should undergo routine cleaning and sterilisation before being returned to the loaner, as per manufacturer instructions. Loan sets should be transported in fit-for-purpose containers to minimise the risk of damage, contamination and injury to handlers.

For more information, see **Standard AS 5369:2023**.

Routine testing of disinfectant

Concentration of a disinfectant or sterilising agent is critical to its effectiveness in infection prevention. The concentration and temperature of the disinfectant and the contact time with the instrument must adhere with the disinfectant's stated inclusion on the Australian Register of Therapeutic Goods (ARTG). This information should be reflected in the manufacturer's instructions on the disinfectant's label.

Reusable disinfectants will gradually reduce in their effectiveness over time, and the appropriate number of reuses must be determined by testing that the solution is at or above its minimum effective concentration. This should be checked daily or more frequently according to the number of instruments being reprocessed [102].

Further considerations

Steam sterilisation and the other methods listed above are not sufficient for reprocessing items potentially contaminated with infectious agents such as prions. Prions cause diseases such as Creutzfeldt-Jakob disease (CJD), and single-use items should be used wherever possible and subsequently destroyed by incineration if there is a risk of prion contamination.

For further information on infection control issues relating to CJD and the reprocessing of reusable medical devices, refer to the Department of Health *Infection Control Guidelines* at <http://www.health.gov.au/internet/main/publishing.nsf/Content/icg-guidelines-index.htm> [135].

For further information on reprocessing ultrasound probes, refer to the Australasian Society for Ultrasound in Medicine *Guidelines for Reprocessing Ultrasound Transducers* at <http://onlinelibrary.wiley.com/doi/10.1002/ajum.12042/full> [136].

Individual actions for reducing risk:

- Become familiar with AS, ISO and EN standards and facility protocols on cleaning, disinfecting and sterilising.
- Use the appropriate product for the situation and use it as directed.
- Participate in education sessions and professional development sessions on reprocessing instruments and equipment, particularly when new sterilising or disinfecting equipment is introduced.

For practical information on the reprocessing of equipment, see Case Study 5.6.

3.1.5 Respiratory hygiene and cough etiquette

Respiratory hygiene and cough etiquette must be applied as a standard infection control precaution at all times. Covering sneezes and coughs prevents infected persons from dispersing respiratory secretions into the air. Hands must be washed with soap and water after coughing, sneezing, using tissues, or after contact with respiratory secretions or objects contaminated by these secretions.

Table 10. Steps in respiratory hygiene and cough etiquette

Anyone with signs and symptoms of a respiratory infection, regardless of the cause, should follow or be instructed to follow respiratory hygiene and cough etiquette as follows:

- Cover the nose/mouth with disposable single-use tissues when coughing, sneezing, wiping and blowing noses.
- Use tissues to contain respiratory secretions.
- Dispose of tissues in the nearest waste receptacle or bin after use.
- If no tissues are available, cough or sneeze into the inner elbow rather than the hand.
- Practice hand hygiene after contact with respiratory secretions and contaminated objects/materials.
- Keep contaminated hands away from the mucous membranes of the mouth, eyes and nose.
- In healthcare facilities, patients with symptoms of respiratory infections should sit as far away from others as possible. If available, healthcare facilities may place these patients in a separate area while waiting for care.

Healthcare workers should also assist patients (e.g. elderly, children) who need assistance with containment of respiratory secretions. Those who are immobile will need a receptacle (e.g. plastic bag) readily at hand for the immediate disposal of used tissues and will need to be offered hand hygiene facilities.

A risk assessment should be performed if patients present with aerosol-generating behaviours, such as persistent or severe coughing, screaming, or shouting, heavy breathing and panting, and if there are environmental factors that increase risk, such as poor ventilation. If tolerated, encourage the patient to wear an appropriate mask as a means of source control.

Healthcare workers with viral respiratory tract infections should remain at home until their symptoms have resolved.

Respiratory hygiene and cough etiquette are particularly important for patients on droplet precautions (see Section 3.2.3).

3.1.6 Aseptic technique

Aseptic technique

Aseptic technique is a set of practices aimed at minimising contamination and is particularly used to protect the patient from infection during procedures [12]. Many of the other work practices that form standard precautions are required for aseptic technique, however, adherence to these practices alone does not constitute aseptic technique. Sterile single-use equipment or instruments must be used according to manufacturer's instructions and in such a way that the sterility of the item is maintained.

Commercial frameworks to assist with the implementation of aseptic technique are available and may be practiced in some healthcare facilities.

The five essential principles of aseptic technique are:

1. Sequencing:

- Performing a risk assessment
- Pre-procedure preparation
- Performing the procedure

- Post procedure practices, handover and documentation
2. Environmental control:
- Prior to aseptic procedures, healthcare workers must ensure there are no avoidable nearby environmental risk factors, such as bed making or patients using commodes
3. Hand hygiene:
- Perform hand hygiene before a procedure and after a procedure or body fluid exposure
4. Maintenance of aseptic fields:
- Cleaning and/or disinfection of equipment and patient prior to procedure(s)
 - Establishing an aseptic field
 - Use of sterile equipment
 - Maintenance of the aseptic field, including protecting the key sites and key parts
 - Use of a non-touch technique
5. PPE:
- Correct selection and use of sterile and non-sterile PPE

Example of an Aseptic Technique: Aseptic Non-Touch Technique (ANTT®)

The NHMRC and the Australian Commission on Safety and Quality in Health Care does not specifically endorse ANTT, but recognises ANTT as a best practice example of a clinical practice and education framework for aseptic technique.

ANTT is defined as, 'A specific type of aseptic technique with a unique theory and practice framework.'^[90] ANTT principles are intended for use in all clinical specialities and care settings - from the operating theatre to the community^[426]. Since the late 1990's, The Association for Safe Aseptic Practice (ASAP) originated and has subsequently overseen the development and dissemination of ANTT internationally, and supported hospitals and community health organisations to implement ANTT and help maintain safe standards of aseptic technique.^[428] There are two types of ANTT: Standard-ANTT and Surgical-ANTT and the ANTT Clinical Practice Framework is summarised below ^[427]:

Terminology

Historically, the practice of protecting patients from contamination and infection during clinical procedures has generated an inaccurate and confusing paradigm based on the terminology of undefined sterile, aseptic and clean techniques. The use of accurate terminology is important in order to promote clarity in practice.

Sterile 'Free from microorganisms' [140]

Due to the natural multitude of organisms in the atmosphere it is not possible to achieve a sterile technique in a typical healthcare setting. Near sterile techniques can only be achieved in controlled environments such as a laminar air flow cabinet or a specially equipped theatre. The commonly used term 'sterile technique', i.e. the instruction to maintain sterility of equipment exposed to air, is obviously not possible and is often applied inaccurately.

Asepsis 'Freedom from infection or infectious (pathogenic) material' [140]

An aseptic technique aims to prevent pathogenic organisms, in sufficient quantity to cause infection, from being introduced to susceptible sites by hands, surfaces and equipment. Therefore, unlike sterile techniques, aseptic techniques are possible and can be achieved in typical hospital and community settings.

Clean 'Free from dirt, marks or stains' [142]

Although cleaning followed by drying of equipment and surfaces can be very effective, it does not necessarily meet the quality standard of asepsis ^[138]. However, the action of cleaning is an important component in helping render equipment and skin aseptic, especially when there are high levels of contaminants that require removal or reduction. As such, to be confident of achieving asepsis, an application of a skin or hard surface disinfectant is required either during cleaning or afterwards.

Consequently, the aim of ANTT is asepsis.

ANTT in practice

ANTT is based upon an original concept of Key-Part and Key-Site Protection to protect the procedure Key-Parts and patient Key-Sites from contamination by microorganisms that could cause infection. In ANTT, asepsis is ensured by identifying and then protecting all Key-Parts and Key-Sites by hand hygiene, non-touch technique, using new sterilised equipment and/or cleaning existing Key-Parts to a standard that renders them aseptic prior to use.

Core infection control components of ANTT

Both types of ANTT first demand standard precautions such as hand hygiene and personal protective equipment (PPE). Thereafter, Key-Parts and Key-Sites are protected using a combination of non touch technique and aseptic fields – depending on the type of ANTT utilised.

Risk assessment

While the principles of ANTT remain constant for all clinical procedures, the level of aseptic practice will differ depending upon ANTT risk assessment. The health care worker will consider the technical difficulty of achieving asepsis by appraising a range of procedure variables including user competence. The healthcare worker assesses whether the procedure can be performed easily without touching Key-Parts and Key-Sites directly. If no, Surgical-ANTT is utilised. If yes, Standard -ANTT is utilised. See below.

Aseptic technique or ANTT cannot always be applied due to emergency and/or uncontrolled environmental conditions. Where this occurs, healthcare workers should aim to utilise the principles of ANTT where practical and safe to do so. Where there has been a breach, this should be documented and included in handover and the infection risks mitigated as soon as possible.

Key-Part and Key Site Protection

The health care worker will always apply the Key-Part and Key-Site rule: 'Aseptic Key Parts must only come into contact with other aseptic Key-Parts and/or Key-Sites'.

Hand hygiene

Effective hand hygiene is an essential component of aseptic technique. In Standard-ANTT, hand hygiene should be performed as outlined in Section 3.1.1. In Surgical-ANTT, a surgical hand scrub is required [143] (see Section 3.5.3).

Glove use

It is known that hand hygiene is not always correctly performed and that even correctly performed hand hygiene cannot always remove all pathogenic organisms. Therefore, identifying Key-Parts and Key-Sites and not touching them directly or indirectly using a non touch method - is a vital component of achieving asepsis. In Standard-ANTT, non-sterile gloves are typically worn. In some instances the additional rationale is to protect the user from body fluids or exposure to chemicals. For Surgical-ANTT, sterile gloves are always worn.

Aseptic fields

Even well cleaned hospitals can be said to be 'dirty'—as they are busy and dynamic environments that typically harbour unusual antibiotic-resistant organisms. Consequently, aseptic fields are important in providing a controlled aseptic working space to help promote or ensure the integrity of asepsis during clinical procedures. It is also important that aseptic fields are fit for purpose. In aseptic technique, aseptic fields are increased in size and sterilised drapes are added on the basis of procedure complexity; for example, in IV therapy, 'mobile' aseptic fields such as plastic trays should be large enough and with high sides to provide an adequate working space to contain equipment, sharps and spillages.

ANTT employs three types of aseptic field that require different management depending on whether the primary purpose is to promote or ensure asepsis.

Critical Aseptic Fields; ensuring asepsis

Critical Aseptic Fields are used when Key-Parts and/or Key-Sites, usually due to their size or number, cannot easily be protected individually at all times with sterilized covers and caps, or handled at all times using non-touch technique (such as in peripherally inserted central venous catheters (PICC lines, urinary catheter insertion, complex wound care etc.), or when particularly open and invasive procedures demand large aseptic working areas for long durations, as in the operating room. In such cases, the Critical Aseptic Field demands to be managed as a Key-Part (i.e. only equipment that has been sterilised can come into contact with it). Such a Critical Aseptic Field demands the use of sterilised gloves and, often, full barrier precautions [34]. Large main Critical Aseptic Fields are used in Surgical-ANTT, generally reflecting more complex and longer duration aseptic procedures.

A sub-type of a Critical Aseptic Field is the Micro Critical Aseptic field. Traditional aseptic/ clean techniques have protected Key-Parts by syringe caps, sheathed clean needles, covers or packaging etc. This often-understated approach is given particular emphasis in ANTT, because the inside of such caps and covers have been sterilised and thus can provide an optimum all-encompassing aseptic field for Key-Parts.

General Aseptic Fields; promoting asepsis

General Aseptic Fields are used in Standard-ANTT when Key-Parts can easily and optimally be protected by Micro Critical

Aseptic Fields and a non touch method. The main General Aseptic Field isn't managed as a Key-Part and is essentially promoting, rather than ensuring, asepsis. Subsequently, the procedure is considerably simplified and typically involves non-sterile gloves.

Environmental control

Prior to aseptic procedures, healthcare workers must ensure that there are no avoidable nearby environmental risk factors, such as bed making or patients using commodes.

Sequencing

ANTT practice is sequenced to ensure an efficient, logical and safe order of procedure events. Section 5.11 provides examples of how to perform ANTT for peripheral and central access intravenous therapy, and for wound care.

Surgical-ANTT or Standard-ANTT?

Differentiation between Standard-ANTT and Surgical-ANTT is intended to provide clarity and structure to aid understanding, but not polarise practice. Sequenced procedure guidelines help standardise practice, technique and equipment levels.

- **Standard-ANTT**—Clinical procedures managed with Standard-ANTT will characteristically be technically simple to achieve asepsis, short in duration (approximately less than 20 minutes), and involve a relatively small Key-Site and few, small Key-Parts. Standard-ANTT requires a main General Aseptic Field and typically non-sterile gloves. The use of Micro Critical Aseptic Fields and a non touch method is essential to protect Key-Parts and Key-Sites individually.
- **Surgical-ANTT**— is demanded when procedures are technically complex to achieve asepsis. Procedures involve extended periods of time, a large open Key-Site and large or numerous Key-Parts. To counter these risks, a main Critical Aseptic Field and sterile gloves are required and often full barrier precautions [34]. Surgical-ANTT should still utilise Micro Critical Aseptic Fields where practical to do so.

Table 11. Selection of Standard-ANTT or Surgical-ANTT for specific procedures

| Procedure | Type of ANTT | Rationale/Typical Procedure |
|----------------------------------|-------------------------------|---|
| IV therapy | Standard-ANTT | Asepsis is typically straight forward to achieve: Key-Parts can typically be protected by optimal Micro Critical Fields and non-touch technique. Key-Sites are small. Procedures are technically simple and <20 mins duration. |
| Simple wound dressings | Standard-ANTT | Key-Parts and Key-Sites can be protected by Micro Critical Fields and non-touch technique. Procedures are technically simple and <20 mins duration. |
| Complex or large wound dressings | Surgical-ANTT | The complexity, duration or number of Key-Parts generally demands a Critical Aseptic Field. |
| Urinary catheterisation | Surgical-ANTT / Standard-ANTT | An experienced healthcare worker can perform catheterisation achieving asepsis, by managing Key-Parts individually, with the use of a main General Aseptic Field, Micro Critical Aseptic Fields and non touch technique. However, less experienced healthcare workers may require a Critical Aseptic Field. |
| Cannulation | Surgical-ANTT / Standard-ANTT | Although technically quite simple, the close proximity of healthcare worker hands to the puncture site and Key-Parts may demand sterile gloves – dependent upon the healthcare worker's competency and the difficulty of vein access. |
| PICC/CVC insertion | Surgical-ANTT | The size of the CVC or PICC line, invasiveness, numerous Key-Parts and equipment and duration will demand a Critical Aseptic Field and full barrier precautions. |
| Surgery | Surgical-ANTT | Surgical access involves deep or large exposed wounds, numerous Key-Parts and equipment and long duration procedures. Standard operating room precautions required. |

Conditional recommendation

Updated

19. It is suggested that sterile gloves are used for aseptic procedures and contact with sterile sites.

Practical info

Sterile gloves are indicated for any surgical procedure, vaginal delivery, invasive radiological procedures, performing vascular access and procedures (central lines), preparing total parental nutrition and chemotherapeutic agents. For further information on selection of glove type, Section 3.3 *Recommendation 31*.

Evidence to decision

| | |
|---|---|
| Benefits and harms | The benefits of wearing sterile gloves clearly outweigh any undesirable effects. |
| Certainty of the evidence | This advice is based on limited empirical evidence, but on sound theoretical principles and is supported by expert advice. |
| Values and preferences | It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections during any episode of care through the use of sterile gloves. |
| Resources and other considerations | The provision of sterile gloves presents costs to healthcare facilities, additional to that of normal gloves. In some areas of primary care, it may be difficult to implement the use of sterile gloves. |

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. The use of sterile gloves for aseptic procedures and contact with sterile sites is justified to reduce healthcare associated infection.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

3.1.7 Waste management

As there is currently no national definition of clinical waste in Australia, healthcare facilities, including community healthcare settings, need to conform to relevant state or territory legislation and regulations on the management of clinical and related wastes. Healthcare facilities should also refer to **Standard AS/NZS 3816: 2018** and the Waste Management Association of Australia's industry code of practice. Waste management resources for each state and territory are provided below in Table 12.

When handling waste:

- apply standard precautions to protect against exposure to blood and body substances during handling of waste; wash hands following procedure
- segregation should occur at the point of generation
- waste should be contained in the appropriate receptacle (identified by colour and label) and disposed of according to the facility waste management plan
- healthcare workers should be trained in the correct procedures for waste handling.

Regardless of where waste is generated (e.g. from isolation rooms/patients versus routine patient-care areas), the principles of determining whether it is to be treated as clinical or general waste remain the same.

Table 12. State and territory resources for waste management

| | |
|------------------------------|---|
| Australian Capital Territory | http://www.legislation.act.gov.au/a/1990-5/ |
| New South Wales | http://www.health.nsw.gov.au/environment/clinicalwaste/Pages/default.aspx |
| Northern Territory | https://ntepa.nt.gov.au/waste-pollution/guidelines/guidelines |
| Queensland | https://www.ehp.qld.gov.au/assets/documents/regulation/pr-gl-clinical-and-related-waste.pdf |
| South Australia | http://www.epa.sa.gov.au/community/waste_and_recycling/medical_waste |
| Tasmania | https://epa.tas.gov.au/regulation/waste-management/controlled-waste/handling-controlled-waste-in-tasmania/handling-transport-requirements-for-particular-controlled-wastes/clinical-and-related-waste |
| Victoria | http://www.epa.vic.gov.au/business-and-industry/guidelines/waste-guidance/clinical-waste-guidance |
| Western Australia | https://ww2.health.wa.gov.au/About-us/Policy-frameworks/Public-Health/Mandatory-requirements/Environmental-Health-Management/Clinical-and-Related-Waste-Management-Policy |

3.1.8 Handling of linen

Healthcare facilities must have documented policies on the collection, transport and storage of linen. Healthcare facilities that process or launder linen must have documented operating policies consistent with **Standard AS/NZS 4146: 2000**.

All used linen should be handled with care to avoid dispersal of microorganisms into the environment and to avoid contact with staff clothing. The following principles apply for linen used for all patients (i.e. whether or not transmission-based precautions are required):

- Appropriate personal protective equipment is worn during handling of soiled linen to prevent exposure of skin and mucous membrane to blood and body substances.
- Used linen is 'bagged' at the location of use into an appropriate laundry receptacle.
- Used linen must not be rinsed or sorted in patient-care areas or washed in domestic washing machines.
- Linen soiled with body substances should be placed into leak-proof laundry bags for safe transport.
- Hand hygiene is performed following the handling of used linen.

Clean linen must be stored in a clean and dry place that prevents contamination by aerosols, dust, moisture and vermin, and is separate from used linen.

Patient items

Domestic-type washing machines must only be used for a patient's personal items (not other linen). Washing must involve the use of an appropriate detergent and hot water. If hot water is not available, only individual patient loads can be washed at one time. Clothes dryers should be used for drying.

3.2 Transmission-based precautions

Summary

Section 3.2 outlines transmission-based precautions to guide staff in the presence of suspected or known infectious agents that represent an increased risk of transmission.

A summary of recommended precautions for specific infectious agents can be found in **Appendix 2 — Section 6.4**.

- Transmission-based precautions are applied in addition to standard precautions.
- The aim of instituting transmission-based precautions early, is to reduce further transmission opportunities that may arise due to the specific route of transmission of a particular pathogen.
- While it is not possible to prospectively identify all patients needing transmission-based precautions, in certain settings recognising an increased risk warrants their use while confirmatory tests are pending.

Patient-care tip

When transmission-based precautions are applied during the care of an individual patient, there is potential for adverse effects such as anxiety, mood disturbances, perceptions of stigma and reduced contact with clinical staff. Clearly explaining to patients why these precautions are necessary may help to alleviate these effects.

3.2.1 Application of transmission-based precautions

What are the risks?

- Indirect or direct **contact transmission**: when a healthcare workers' hands or clothing become contaminated, patient-care devices are shared between patients, infectious patients have contact with other patients, or environmental surfaces are not regularly decontaminated.
- **Droplet transmission**: when healthcare workers' hands become contaminated with respiratory droplets and are transferred to susceptible mucosal surfaces such as the eyes; when infectious respiratory droplets are expelled by coughing, sneezing or talking, and come into contact with another's mucosa (eyes, nose or mouth), either directly or via contaminated hands.
- **Airborne transmission**: when attending healthcare workers or patients inhale small particles that contain infectious agents.

When are transmission-based precautions applied?

Transmission-based precautions are applied to patients suspected or confirmed to be infected with agents transmitted by the contact, droplet or airborne routes.

The combination of measures used in transmission-based precautions depends on the route(s) of transmission of the infectious agent involved, as outlined in Sections 3.2.2, 3.2.3 and 3.2.4 below. In the acute-care setting, this will involve a combination of the following measures:

- continued implementation of standard precautions
- appropriate use of personal protective equipment (PPE) (including gloves, apron or gowns, surgical masks or P2 respirators, and protective eyewear)
- patient-dedicated equipment
- allocation of single rooms or cohorting of patients
- appropriate air handling requirements
- enhanced cleaning and disinfecting of the patient environment
- restricted transfer of patients within and between facilities.

For diseases that have multiple routes of transmission, more than one transmission-based precaution category is applied. Whether used singly or in combination, transmission-based precautions are always applied in addition to standard precautions. Transmission-based precautions remain in effect for limited periods of time until signs and symptoms of the infection have resolved, or according to recommendations from infection control professionals specific to the infectious agent (see Appendix 2—Section 6.4).

The mode of transmission of infectious agents is the same in primary care or office-based practice as it is in the acute-care setting. However, the risk of transmission may differ due to the population groups and the nature of care provided.

Considering the following will help to establish the risk of infection in primary care and office-based practice:

- **Patient population**—this will influence the nature of care required and the type of potential infectious agents (i.e. some populations have a higher incidence of tuberculosis).
- **The profile of care**—this includes the level of training of staff, what forms of invasive procedures are performed and whether equipment is reprocessed or single use.
- **Local infrastructure**—this influences water quality, food availability and access to other health services (i.e. rural vs urban).

In developing policies and procedures for a healthcare facility it is useful to refer to discipline-specific guidelines to inform practice on specialised areas.

An overview of risk-management principles and processes is given in Section 2.2.

Individual actions for reducing risk:

- Consult with infection control professionals to ensure that appropriate transmission-based precautions are applied and that they

remain in place until the risk of transmission of the infectious agent has passed.

- Remember that transmission-based precautions are applied AS WELL as standard precautions.
- Advise patients why particular measures are needed to control infection.
- Become familiar with local policy on appropriate PPE, and when it should be put on and taken off, when attending patients on transmission-based precautions.
- Make sure you know which type of mask is needed in different situations and how to check that they are properly fitted.
- Always contain or cover the infected or colonised areas of a patient subject to contact precautions before moving them from one patient-care area to another.
- Explain the purpose and process of respiratory hygiene and cough etiquette to patients on droplet precautions.
- Ask patients on droplet or airborne precautions to wear a surgical mask if they are being moved from one patient-care area to another.
- If patients are moved to a single-patient room (contact or droplet precautions) or negative pressure room (airborne precautions) explain why this is necessary to prevent transmission of infection.
- Make sure you are fully immunised against vaccine-preventable diseases as recommended in the *Australian Immunisation Handbook* [353].

Environmental cleaning

In acute-care areas where the presence of infectious agents requiring transmission-based precautions is suspected or known, surfaces should be physically cleaned with a detergent solution. A Therapeutic Goods Administration (TGA)-listed hospital-grade disinfectant with specific claims (or sodium hypochlorite if indicated for use as per *Recommendation 14*) should then be used (e.g. physical 2-step clean or 2-in-1 clean) as outlined in Section 3.1.4. In office-based practices and non-acute-care areas (e.g. long-term care facilities), the risk of contamination, mode of transmission and risk to others should be used to determine whether disinfectants are required. Further information on environmental cleaning product selection can be found [here](#).

Crockery and utensils used by patients on transmission-based precautions do not require containment and should be treated in the same manner as those used for non-infectious patients (i.e. washed in a dishwasher). Disposable crockery and utensils are not necessary.

This section does not provide specific guidance on cleaning. Appendix 2 (Section 6.1) provides guidance on frequency of cleaning of specific items in low, medium and high-risk settings. Further information on the considerations required when developing cleaning schedules is provided in Section 3.1.3 Practice Statement 9 practical information.

3.2.2 Contact precautions

What are the risks?

There is clear evidence that certain infectious agents are transmitted by direct or indirect contact during patient care.

Direct transmission occurs when infectious agents are transferred from one person to another person without a contaminated intermediate object or person. For example, blood or other body substances from an infectious person may come into contact with a mucous membrane or breaks in the skin of another person [162] [151].

Indirect transmission involves the transfer of an infectious agent through a contaminated intermediate object (fomite) or person. Contaminated hands of healthcare workers have been shown to be important contributors to indirect contact transmission [153] [152] [158]. Other opportunities for indirect contact transmission include:

- when clothing becomes contaminated while caring for a patient colonised or infected with an infectious agent, which can then be transmitted to subsequent patients [161] [166]
- when contaminated patient-care devices are shared between patients without cleaning and disinfection between patients [154] [156] [164]
- when environmental surfaces become contaminated (see Section 3.1.3 *Practice Statement 9* practical information on routine environmental cleaning and Appendix 2 (Section 6.1) on frequency of cleaning of specific items).

Direct or indirect contact transmission of microorganisms during patient care is responsible for the majority of healthcare associated infections in patients and healthcare staff.

Conditional recommendation

Updated

20. It is suggested that contact precautions, in addition to standard precautions, are implemented in the presence of known or suspected infectious agents that are spread by direct or indirect contact with the patient or the patient's environment.

Practical info

When should contact precautions be implemented?

Contact precautions are used when there is a risk of direct or indirect contact transmission of infectious agents (e.g. *C. difficile*, or highly contagious skin infections/infestations such as methicillin-resistant *Staphylococcus aureus* (MRSA) that are not effectively contained by standard precautions alone (see Section 3.1).

Information about which precautions to apply for specific conditions is given in Appendix 2—Section 6.4.

Patient placement

Single-patient rooms

A single-patient room is recommended for patients who require contact precautions. Note that single-patient rooms are also an effective precaution to prevent droplet and airborne transmission of infection, and can also protect immunocompromised patients. Rooms with ensembles and anterooms are preferred. Anterooms increase the effectiveness of single-patient rooms by reducing the potential escape of airborne infectious particles in the corridor.

The rationale for single-patient rooms include:

- it can facilitate greater frequency of cleaning and decontamination, as there is limited impact on neighbouring patients
- access to an ensuite bathroom can reduce the spread of *C. difficile* and other infectious agents
- it can reduce the spread of multi-resistant organisms
- the greater prominence of sinks or hand hygiene dispenser is likely to improve hand hygiene compliance.

Other points relevant to patient placement include the following:

- keep patient notes outside the room
- keep patient bedside charts outside the room
- disinfect hands upon leaving room and after writing in the chart
- keep doors closed where safe to do so (this may not be possible for patients requiring high visualisation)
- make sure rooms are clearly signed.

When determining the number and type of single rooms in a healthcare facility, project planning teams should consider [396]

- trends in disease in the general population and the particular population serviced
- demographic trends in the population served
- specialities of the healthcare facility
- project changes in future clinical activities.

When a single-patient room is not available, consultation with infection control professionals is recommended to assess the various risks associated with other patient placement options (e.g. cohorting).

Cohorting

If cohorting takes place, it is recommended that patient beds are separated by approximately one metre or more to reduce opportunities for the inadvertent sharing of items between patients [164].

If it is necessary to place a patient who requires contact precautions in a room with a patient who is not infected or colonised:

- avoid placing these patients with patients who are at increased risk of an adverse outcome from infection (e.g. patients who are immunocompromised, have open wounds or have anticipated prolonged lengths of stay)
- change protective attire and perform hand hygiene between contact with patients in the same room, regardless of whether one or both patients are on contact precautions.

Transfer of patients

Limiting transfer of a patient on contact precautions reduces the risk of environmental contamination. If transfer within or between facilities is necessary, it is important to ensure that infected or colonised areas of the patient's body are contained and covered. Contaminated personal protective equipment (PPE) should be removed and disposed of and hand hygiene performed before the patient is moved. Clean personal protective equipment should be put on before assisting the patient at the destination.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of using contact precautions clearly outweigh any undesirable effects.

Certainty of the evidence

Low

There is supportive evidence and a strong theoretical rationale to support this intervention [164]. This intervention is also supported by work, health and safety principles.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value use of contact precautions to minimise infections when there is a known infectious agent spread by direct or indirect contact.

Resources and other considerations

No important issues with the recommended alternative

The net benefits of implementing contact precautions are worth the cost.

This practice would entail standard hand hygiene practices, as well as additional steps such as personal protective equipment, and consideration of patient placement and patient transport.

Staff training and/or tools such as checklists in wards, may be necessary to ensure uptake of advice.

Rationale

There is supportive evidence and a strong theoretical rationale to support the use of contact precautions for patients known or suspected to be infected with an infectious agent spread via the contact route. This intervention is also supported by work, health and safety principles.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Conditional recommendation

Updated

21. It is suggested that appropriate hand hygiene be undertaken and personal protective equipment worn to prevent contact transmission.

It is suggested that when working with patients who require contact precautions, healthcare workers should:

- perform hand hygiene
- put on gloves and gown upon entry to the patient-care area
- if performing multiple tasks whilst in the patient-care area, apply the principles of standard precautions and remove gloves, perform hand hygiene and apply clean gloves between tasks when required to minimise risk of infection transmission
- ensure that clothing and skin do not contact potentially contaminated environmental surfaces
- remove gown and gloves and perform hand hygiene before leaving the patient-care area.

Practical info

How should contact precautions be applied?

Contact precautions are applied in addition to standard precautions.

The key aspects of applying contact precautions relate to:

- use of appropriate personal protective equipment (PPE)
- special handling of equipment
- patient placement
- minimising patient transfer or transport.

Hand hygiene and PPE

Effective hand hygiene is particularly important in preventing contact transmission and the 5 moments for hand hygiene outlined in Section 3.1.1 *Recommendation 1* should be followed at all times. When the presence of *C. difficile* or non-enveloped viruses is known or suspected, use of alcohol-based hand rubs alone may not be sufficient to reduce transmission of these organisms (see Section 3.1.1)

Putting on both gloves and gown upon entering the patient-care area helps to contain infectious agents, especially those that have been implicated in transmission through environmental contamination (e.g. *C. difficile*, norovirus and other intestinal tract pathogens, respiratory syncytial virus) [155][157][158] [159][160] [165]. If you are going to perform multiple tasks whilst in the patient zone, remember to remove contaminated gloves after a task, perform hand hygiene and apply clean gloves before starting the next task with that patient to reduce the risk of infection transmission. Considerations in selecting a gown appropriate to the situation are outlined in Section 3.3, *Recommendation 29*.

A surgical mask and protective eyewear or face shield must be worn if there is the potential for generation of splashes or sprays of blood and body substances into the face and eyes.

Hand hygiene compliance is likely to be improved through greater prominence of sinks or alcohol-based hand rub dispensers. More information is at Section 4.6.1.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of using hand hygiene and personal protective equipment (PPE) clearly outweigh any undesirable effects.

Certainty of the evidence

Low

There is supportive evidence and a strong theoretical rationale to support this intervention. The Centers for Disease Control and Prevention [164] provides 1B evidence to support the practice of wearing gloves and gowns for transmission-based precautions.

This intervention is also supported by work, health and safety principles.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value use of hand hygiene and PPE to prevent contact transmission of infections.

Resources and other considerations

No important issues with the recommended alternative

Any marginal increases in costs of compliance to hand hygiene and PPE is offset by the corresponding reduction in infection rates [217].

Rationale

The practice of wearing personal protective equipment is suggested for transmission-based precautions, including contact precautions. Personal protective equipment can protect healthcare workers against contaminated environmental surfaces and medical equipment, so gloves and gowns should be put on before entry into a patient's room and removed after leaving the patient-care area.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Conditional recommendation

Updated

22. It is suggested that patient-dedicated equipment or single-use patient-care equipment be used for patients on contact precautions.

If common use of equipment for multiple patients is unavoidable, clean the equipment and allow it to dry before use on another patient.

Practical info

Single-use or patient-dedicated equipment

Standard precautions concerning patient-care equipment (see Section 3.1.3 *Practice Statements 9-11 & Recommendation 12*) are very important in the care of patients on contact precautions. If patient-care devices (e.g. blood pressure cuffs, nebulisers, mobility aids) are shared between patients without being reprocessed between uses, they may transmit infectious agents [154][156][163]. Where common use of equipment for multiple patients is unavoidable, a risk assessment should be performed and cleaning carried out according to the manufacturer's instructions.

Any medical device (instruments and equipment) that is to be reused requires reprocessing—cleaning, disinfection and/or sterilisation. The minimum level of reprocessing required for reusable instruments and equipment depends on the individual situation (i.e. the body site, presence of multi-resistant organisms and the nature by which the instrument will be used). For further information on the reprocessing of reusable medical devices, see Section 3.1.4.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of using patient dedicated equipment clearly outweigh any undesirable effects.

Certainty of the evidence

Low

There is supportive evidence and a strong theoretical rationale for this intervention. The Centers for Disease Control and Prevention suggest (Category 1B evidence) [164] that patient-dedicated or single-use equipment be used as part of contact precautions.

This intervention is also supported by work health and safety principles.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value use of dedicated patient equipment to prevent contact transmission of infections.

Resources and other considerations

No important issues with the recommended alternative

The cost effectiveness and feasibility of this practice would depend on the availability of such equipment in a healthcare facility.

Staff training and/or tools such as checklists in wards, may be necessary to ensure uptake of advice.

Rationale

Evidence shows that using patient-dedicated equipment has been beneficial for preventing transmission of infectious agents [164].

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

3.2.3 Droplet precautions

What are the risks?

A number of infectious agents are transmitted through respiratory droplets (i.e. large-particle droplets >5 microns in size) that are generated by a patient who is coughing, sneezing or talking. Transmission via large droplets requires close contact as the droplets do not remain suspended in the air and generally only travel short distances. They can however, contaminate horizontal surfaces close to the source patient, and the hands of healthcare workers can become contaminated through contact with those surfaces. For this reason consideration should be given to the need for additional personal protective equipment (PPE) (see Section 3.3.).

Droplet precautions are based on evidence that shows that:

- Hand hygiene is effective in preventing transmission of viruses and reducing the incidence of respiratory infections both within and outside healthcare settings [167] [173] [168].
- Physical interventions are highly effective against the spread of a broad range of respiratory viruses [170] [172].
- Surgical masks protect the wearer from droplet contamination of the nasal or oral mucosa [174]. See *Recommendation 30* for further information on surgical masks.
- Physical proximity of less than one metre has been associated with an increased risk for transmission of some infections via

the droplet route.

- Placing surgical masks on coughing patients can also prevent infected patients from dispersing respiratory secretions into the air [174].

How should droplet precautions be applied?

The key aspects of applying droplet precautions relate to:

- standard precautions
- use of appropriate PPE
- special handling of equipment
- patient placement
- minimising patient transfer or transport.

Conditional recommendation

Updated

23. It is suggested that droplet precautions, in addition to standard precautions, are implemented for patients known or suspected to be infected with agents transmitted by respiratory droplets that are generated by a patient when coughing, sneezing or talking.

Practical info

When should droplet precautions be implemented?

Droplet precautions are intended to prevent transmission of infectious agents spread through close respiratory or mucous membrane contact with respiratory secretions. As these microorganisms do not travel over long distances, special air handling and ventilation are not required. Infectious agents for which droplet precautions are indicated include influenza, norovirus, pertussis, meningococcus.

Hand hygiene and droplet precautions

Some infectious agents transmitted by the droplet route may also be transmitted by contact [198]. Hand hygiene is therefore an important aspect of droplet precautions and the 5 moments for hand hygiene outlined in Section 3.1.1 should be followed.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of using droplet precautions for patients known or suspected to be infected with agents transmitted by respiratory droplets clearly outweigh any undesirable effects.

Certainty of the evidence

Low

There is supportive evidence and a strong theoretical rationale (CDC Category 1B evidence) [174] for the use of droplet precautions for patients known or suspected to be infected with agents transmitted by respiratory droplets.

This intervention is also supported by work health and safety principles.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value use of droplet precautions (such as hand hygiene combined with personal protective equipment (PPE), special handling of equipment, patient placement and minimising patient transport) to prevent transmission of infections.

Resources and other considerations

No important issues with the recommended alternative

This practice would entail standard hand hygiene practices, as well as additional steps such as PPE, and consideration of patient placement and patient transport.

Staff training and/or tools such as checklists in wards, may be necessary to ensure uptake of advice.

Rationale

Droplets precautions are intended to prevent the transmission of pathogens that are spread through respiratory or mucous membrane contact with respiratory secretions.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Conditional recommendation

Updated

24. It is suggested that a surgical mask should be worn when entering a patient-care environment to prevent droplet transmission.

Practical info

Use of surgical masks to prevent droplet transmission

There is insufficient evidence to support the use of P2 respirators for reducing the risk of infections transmitted by the droplet route. Although surgical masks do not protect the wearer from infectious agents that are transmitted via the airborne route, surgical masks that meet Australian Standards are fluid impervious and protect the wearer from droplet contamination of the nasal or oral mucosa. The surgical mask is generally put on upon room entry, with hand hygiene practised before putting on the mask and after taking off the mask. For further information on characteristics and levels of surgical masks, see *Recommendation 30* and *Table 17*.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of wearing a surgical mask for patients known or suspected to be infected with agents transmitted by respiratory droplets clearly outweigh any undesirable effects.

Certainty of the evidence

Moderate

There is evidence (CDC Category 1B) [164] to support the recommendation that masks should be worn upon entry into a patient-care environment.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value use of a surgical mask as a personal protective equipment to prevent droplet transmission of infections.

Resources and other considerations

No important issues with the recommended alternative

This practice would require the provision of surgical masks.

Staff training and/or tools such as checklists in wards may be necessary to ensure uptake of advice.

Rationale

More studies are needed to improve understanding of droplet transmission under various circumstances.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Practice Statement

Updated

25. It is good practice to place patients who require droplet precautions in a single-patient room.

Practical info

Placement of patients on droplet precautions

Placing patients on droplet precautions in a single-patient room reduces the risk of patient-to-patient transmission. When single-patient rooms are in short supply, the following principles apply in decision-making on patient placement:

- prioritise patients who have excessive cough and sputum production for single-patient room placement
- consider patients ability to perform hand hygiene and follow appropriate cough etiquette
- place together in the same room (cohort) patients who are infected with the same pathogen and are suitable roommates.

If it becomes necessary to place patients who require droplet precautions in a room with a patient who does not have the same infection:

- avoid placing patients on droplet precautions in the same room with patients who have conditions that may increase the risk of adverse outcomes from infection or that may facilitate transmission (e.g. those who are immunocompromised, have anticipated prolonged lengths of stay, have cystic fibrosis, cardiac conditions or muscular dystrophy)
- ensure that patients are physically separated (> one metre apart) from each other and draw the privacy curtain between beds to minimise opportunities for close contact.

If a patient requires care under droplet precautions but an aerosol generating procedure is undertaken, then droplet precautions should be increased to airborne precautions for at least the duration of the procedure. The procedure should be undertaken in a treatment room, away from other patients (if the patient is cohorted with others).

In all cases, the importance of respiratory hygiene and cough etiquette should be explained to patients on droplet precautions (see Section 3.1.5).

Where clinical need permits and is not compromised, it may be possible in some settings (e.g. interventional radiology) to schedule patients who require droplet precautions to the end of the list.

In primary care and other office-based practice, examples of appropriate implementation of droplet precautions include segregation in waiting rooms for patients with violent or frequent coughing, and the availability of tissues, alcohol-based hand rub and a waste bin so that patients can practice respiratory hygiene and cough etiquette.

For further information on isolation and cohorting of patients, see Section 3.4.2.1.

Transfer of patients on droplet precautions

When transfer of a patient on droplet precautions within or between facilities is necessary, there is the potential for other patients and healthcare workers to come in contact with infectious agents when the patient coughs or sneezes. This can be addressed by asking the patient to wear a fluid resistant surgical mask while they are being transferred and to follow respiratory hygiene and cough etiquette. Children should wear a correctly fitting mask when they are outside an isolation room. The child's oxygen saturation should be monitored.

Evidence to decision

| | |
|---|---|
| Benefits and harms | The benefits of placing patients who require droplet precautions in a single-patient room outweighs any harms. However, it is important to consider the potential adverse outcomes associated with isolation rooms. |
| Certainty of the evidence | The evidence in this area is limited or inconsistent. The practice statement is based on current expert advice and trends in clinical practice. |
| Values and preferences | It is expected that all patients and staff of Australian healthcare facilities would highly value placing patients who require droplet precautions in a single room to prevent the spread of infection. |
| Resources and other considerations | Placing patients on droplet precautions in a single-patient room requires the availability of such rooms. In smaller facilities, or those in more remote or regional areas, making such a room available may not be possible. |

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Placing patients on droplet precautions in single-patient rooms is justified to reduce healthcare associated infection.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

3.2.4 Airborne precautions

Why are airborne precautions important?

Certain infectious agents are disseminated through airborne droplet nuclei or small particles in the respirable size range that remain infective over time and distance.

Airborne precautions are based on evidence that shows that:

- the use of P2 respirators prevents the inhalation by healthcare workers of small particles that may contain infectious agents transmitted via the airborne route
- the use of negative pressure rooms reduces the transmission of infection
- wearing of correctly-fitted surgical masks by coughing patients prevents dispersal of respiratory secretions into the air (close monitoring is required to minimise respiratory risk) [87].

Implementing airborne precautions in different healthcare settings

Healthcare workers should prioritise the use of P2 / N95 respirators and other respiratory protection, where available.

In situations when personal protective equipment may be limited and a patient is exhibiting the signs of a respiratory disease,

consider placing a mask or tissue over the patient's mouth as tolerated.

Recommended

Updated

26. It is recommended that airborne precautions, in addition to standard precautions, are implemented in the presence of known or suspected infectious agents that are transmitted person-to-person by the airborne route.

Practical info

When should airborne precautions be implemented?

Airborne precautions prevent transmission of microorganisms that remain infectious over time and distance when suspended in the air. These agents may be inhaled by susceptible individuals who have not had face-to-face contact with (or been in the same room as) the infectious individual.

Infectious agents for which airborne precautions are indicated include measles (rubeola), chickenpox (varicella) and *M. tuberculosis*.

Information about which precautions to apply for specific conditions is given in Appendix 2—Section 6.4.

Implementing airborne precautions in different healthcare settings

In situations when personal protective equipment may be limited (such as the setting in which paramedics work) and a patient is exhibiting the signs of a respiratory disease, consider placing a mask or tissue over their mouth as tolerated.

Healthcare workers should also prioritise the use of N95 respirators and other respiratory protection, where available.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of implementing airborne precautions for patients known or suspected to be infected with infectious agents transmitted person-to-person by the airborne route clearly outweigh any undesirable effects.

Certainty of the evidence

Moderate

The Centers for Disease Control and Prevention [164] strongly recommends (Category 1A) applying airborne precautions.

The Public Health Agency of Canada [237] strongly recommends the implementation of airborne precautions for patients known or suspected to be infected with infectious agents transmitted person-to-person by airborne route, such as infectious pulmonary tuberculosis.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value the implementation of airborne precautions to stop the spread of infectious agents.

Resources and other considerations

No important issues with the recommended alternative

This practice would require implementing standard precautions, as well as minimising exposure of other patients and staff to the infectious agent. This requires the provision of appropriate personal protective equipment and training for staff on their correct use.

Rationale

The implementation of airborne precautions in the presence of known or suspected infectious agents that are transmitted by the airborne route is an effective measure to reduce the spread of infection.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Conditional recommendation

Updated

27. It is suggested that a correctly fitted P2 respirator is worn when entering the patient-care area when an airborne-transmissible infectious agent is known or suspected to be present.

Practical info

How should airborne precautions be applied?

The key aspects of applying airborne precautions relate to:

- standard precautions, including respiratory hygiene and cough etiquette (see Section 3.1.5)
- use of appropriate personal protective equipment (PPE) (particularly correctly-fitted P2 respirators)
- minimising exposure of other patients and staff members to the infectious agent.

Specialist procedural areas should refer to their discipline-specific guidelines for detailed advice on applying airborne precautions relevant to the field of practice.

Personal protective equipment

The need for PPE should be based on the precautions required to protect against infectious agents based on the mode of transmission. In the majority of situations where standard respiratory protection is needed, a single use surgical mask (minimum level 2 barrier) is appropriate [177]. See *Recommendation 30, Practical Info, Table 17* for further information on surgical mask levels.

When there is a high probability of airborne transmission due to the infectious agent or procedure (e.g. bronchoscopy), sound scientific principles support the use of P2 respirators to prevent transmission. P2 respirators are designed to help reduce the wearer's respiratory exposure to airborne contaminants such as particles, gases or vapours.

For example, influenza does not usually require the routine use of a P2 respirator except when there is a pandemic strain of influenza.

Standard AS/NZS 1715: 2009 outlines a range of respiratory protective equipment, which provide different levels of protection dependent upon the nature of the microorganism, the mode of transmission and procedure being undertaken.

While the terms 'P2 respirator' and 'N95 respirator' are often used interchangeably in the healthcare setting, they are required to meet different standards. In Australia, the requirements for P2 respirators are stated in **Standard AS/NZS 1716: 2012**. The United States (US) National Institute of Occupational Safety and Health (NIOSH) specifies N95 respirator requirements. See *Table 13* for further information.

Table 13. Properties of different types of mask

| Properties | P2 respirators | N95 respirators |
|------------------------------|---|---|
| Other names | N95 respirator, respiratory protection device, particulate respirator | P2 respirator, respiratory protection device, particulate respirator |
| Characteristics [170] | <ul style="list-style-type: none"> • Raised dome or duckbill | <ul style="list-style-type: none"> • Raised dome or duckbill |

| | | |
|-----------------------------|---|--|
| | <ul style="list-style-type: none"> • 4–5 layers (outer polypropylene, central layers electret [charged polypropylene]) • Filtration through mechanical impaction and electrostatic capture • Designed to provide a good facial fit to minimise aerosol contamination of the mucous membranes of the nose and mouth <p>P2 particulate filtering respirators/masks must have a filter efficiency of at least 94% when tested with Sodium Chloride aerosol at a flow rate of 95 litres/minute.</p> <p>Under the EN system, aerosol testing is similar to Standard AS/NZS 1716: 2012, but have additional filter efficiency testing with paraffin oil aerosol that must also meet the minimum 94% filter efficiency to be classified as P2.</p> <p>The particle size of this aerosol has a mass median diameter of 0.3 to 0.6 microns with a range of particles in the 0.02 to 2 micron size range.</p> | <ul style="list-style-type: none"> • 4–5 layers (outer polypropylene, central layers electret [charged polypropylene]) • Filtration through mechanical impaction and electrostatic capture • Designed to provide a good facial fit to minimise aerosol contamination of the mucous membranes of the nose and mouth <p>NIOSH classified N95 particulate filtering respirators/masks must have a filter efficiency of at least 95% when tested with Sodium Chloride aerosol at a flow rate of 85 litres/minute.</p> <p>N95 respirator masks can only be used for oil free aerosols.</p> <p>The particle size of this aerosol ~0.3 micron.</p> |
| Sealing [170] | <ul style="list-style-type: none"> • Ties at crown and bottom of head, pliable metal nose bridge • Fit testing and fit checking required | <ul style="list-style-type: none"> • Ties at crown and bottom of head, pliable metal nose bridge • Fit testing and fit checking required |
| Australian Standards | Standard AS/NZS 1715: 2009 Standard AS/NZS 1716: 2012 | Set by the US NIOSH classification (NIOSH Guidelines – Procedure No. TEB-APR-STP-0059) |
| Intended use | <ul style="list-style-type: none"> • Routine care of patients on airborne precautions • High-risk procedures such as bronchoscopy when the patient’s infectious status is unknown • Procedures that involve aerosolisation of particles that may contain specific known pathogens | <ul style="list-style-type: none"> • Routine care of patients on airborne precautions • High-risk procedures such as bronchoscopy when the patient’s infectious status is unknown • Procedures that involve aerosolisation of particles that may contain specific known pathogens |
| Notes | <ul style="list-style-type: none"> • Care must be taken if placing respirators on patients and must suit clinical need (i.e. if the patient has chronic obstructive airways disease [COAD] or is in respiratory distress, the respirator will exacerbate symptoms). | <ul style="list-style-type: none"> • Care must be taken if placing respirators on patients and must suit clinical need (i.e. if the patient has chronic obstructive airways disease [COAD] or is in respiratory distress, the respirator will exacerbate symptoms). |

Particulate filtration respirators (PFRs) - fit testing and fit checking

In order for a PFR, such as a P2 respirator to offer the maximum desired protection it is essential that the wearer is properly fitted and trained in its safe use. Healthcare workers are encouraged to actively observe each other’s mask fitting and immediately advise of any fitting issues to maximise healthcare worker and patient safety. A risk-management approach should be applied to ensure that staff working in high-risk areas are trained in appropriate fit of the P2 respirator and how to perform a fit check at the point of use. This may also include fit testing of the mask as outlined below.

Fit testing

The purpose of fit testing is to identify which size and style of P2 respirator is suitable for an individual, and to ensure that it is worn correctly. It also provides an opportunity to ensure healthcare workers are properly trained in the correct use of the mask.

When fit testing is to be undertaken, it should be done so based on relevant state/territory jurisdictional requirements in conjunction with a risk assessment with relevance to the healthcare setting.

While fit testing may be complex and resource intensive, it is a valuable practice which provides an opportunity to educate healthcare professionals. Fit testing programs may be considered:

- at the commencement of employment for employees who will be working in clinical areas where there is a significant risk of exposure to

infectious agents transmitted via the airborne route— assessment of the significance of risk will involve consideration of the location (e.g. risk is higher in an intensive care unit) and activities to be undertaken (e.g. a physiotherapist performing induced sputum is at risk of exposure to infectious aerosols);

- when there is a significant change in the wearer’s facial characteristics that could alter the facial seal of the respirator (e.g. significant change in body weight, facial surgery); and
- at regular intervals — **Standard AS/NZS 1715: 2009** recommends annual fit testing. Healthcare facilities should ensure that they have a respiratory protection program that regularly evaluates the risk to which healthcare workers are exposed and determines which employees are required to undertake fit testing.

Employers must ensure that their employees have the medical ability to wear a respirator. Medical evaluations are required for both positive pressure and negative pressure respirators.

There are two types of facial fit test—qualitative and quantitative. Qualitative fit tests are fast and simple but can be influenced by the wearer. Quantitative fit tests require the use of specialised equipment used by a trained operator. **Standard AS/NZS 1715: 2009** outlines the method by which fit testing is conducted.

Fit checking

Healthcare workers must perform fit checks every time they put on a P2 respirator to ensure it is properly applied. No clinical activity should be undertaken until a satisfactory fit has been achieved. Fit checks ensure the respirator is sealed over the bridge of the nose and mouth and that there are no gaps between the respirator and face. Healthcare workers must be informed about how to perform a fit check.

The procedure for fit checking includes (see Figure 9):

- placement of the respirator on the face
- placement of the headband or ties over the head and at the base of the neck
- compressing the respirator to ensure a seal across the face, cheeks and the bridge of the nose
- checking the positive pressure seal of the respirator by gently exhaling. If air escapes, the respirator needs to be adjusted
- checking the negative pressure seal of the respirator by gently inhaling. If the respirator is not drawn in towards the face, or air leaks around the face seal, readjust the respirator and repeat process, or check for defects in the respirator.

The manufacturer’s instructions for fit checking of individual brands and types of P2 respirator should be referred to at all times.

Health service organisations should have a respiratory protection program in place to support healthcare workers to access and use properly fitting face masks including particulate filtration respirators (PFRs).

A range of different respirators should be considered to ensure reasonable respiratory protection for healthcare workers.

Healthcare workers who have facial hair (including a 1–2 day beard growth) must be aware that an adequate seal cannot be guaranteed between the PFR and the wearer’s face.

- It is recommended that the face be smooth and/or clean-shaven to achieve a good airtight seal
- If facial hair cannot be removed for cultural, religious or medial reasons, an alternative type of PFR should be considered
- PAPRs may be suitable for healthcare workers with facial hair and those who fail fit testing for PFRs.

Wearing a P2 respirator

Considerations when using a P2 respirator include [176]:

- if a good facial seal cannot be achieved (e.g. the intended wearer has a beard or long moustache), an alternative respirator such as a powered air-purifying respirator (PAPR) should be used
- respirators should not be touched while being worn
- respirators should be changed when they become moist
- respirators should never be reapplied after they have been removed
- respirators should not be left dangling around the neck
- hand hygiene should be performed upon touching or disposing of a used respirator.

Removal of a P2 respirator

Correct removal of a P2 respirator is important as there is a risk of contamination to the user if not removed correctly. Considerations when removing a P2 respirator include (see Figure 9):

- removal of respirators should be by the straps from the back of the head
- respirators should be removed outside the patient-care area and disposed of in a closed receptacle [263].

Fitting a P2 respirator

P2 respirators are available in several different designs, and only one is shown here.



- Position respirator over mouth and nose



- Position tapes above and below ears at back of head



- Fit snugly at bridge of nose and under chin by using the adjusters

Removing and disposing of respirator



- With clean hands, grasp tapes at back of head and remove by only handling the tapes, then discard in appropriate waste



- Wash hands

Note: P2 respirators are available in several different designs, and only one is illustrated above.

Figure 9. Process for fitting and removing a P2 respirator

Source: Australian Government Department of Health [178].

Powered air-purifying respirators (PAPRs)

Powered air-purifying respirator (PAPR) devices are not generally recommended for healthcare workers for respiratory protection [439]. The use of a PAPR is a local decision, based on a risk assessment that considers the transmission route, the type of care being provided to the patient, equipment availability, and staff training [439]. PAPRs are associated with greater fatigue among healthcare workers, and may limit physical interaction and care delivery [440].

If PAPRs are required, the device:

- Must be included in the ARTG as a medical device in accordance with the *Therapeutic Goods Act 1989*.
- Should conform to **Standard AS/NZS 1715:2009** and **Standard AS/NZS 1716:2012** and must only be used by healthcare workers who are trained in their use.
- Should be cleaned, decontaminated and maintained in accordance with the manufacturer's instructions.

Evidence to decision

Benefits and harms The benefits of wearing a correctly fitted P2 respirator when an airborne transmissible agent is known or suspected clearly outweighs any undesirable effects.

Certainty of the evidence The overall quality of evidence regarding the use of face masks is low due to poorly controlled studies.

The epic3 Guidelines [79] present Class D evidence (general practice statements) stating that it is best practice to use correctly fitted respirators in situations where a risk-assessment deems it necessary.

The Public Health Agency of Canada [237] present CII evidence (general practice statements) stating that it is best practice for healthcare workers to wear respirators when caring for patients with airborne infections.

| | |
|-------------------------------|--|
| Values and preferences | It is expected that all patients and staff of Australian healthcare facilities would highly value the use of a P2 respirator to prevent airborne transmission of infections. |
|-------------------------------|--|

| | |
|---|---|
| Resources and other considerations | This practice would require the provision of P2 respirators and fit testing of respirators to the healthcare staff. |
|---|---|

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Wearing a correctly fitted P2 respirator when entering the patient-care area of a patient under airborne precautions is justified to reduce healthcare associated infection.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Practice Statement

Updated

28. It is good practice to place patients on airborne precautions in a negative pressure room (Class N/ Type 5) with bathroom facilities or in a room from which air does not circulate to other areas.

Exceptions to this should be justified by risk assessment.

Practical info

Patient placement

When patients with suspected or confirmed airborne infection require treatment using nebulisers, or procedures such as bronchoscopy or nasoendoscopy, it is suggested that a negative pressure room, or a room from which air does not circulate to other areas, should be used. Healthcare workers should be aware it is important to place patients in an area that can be contained. Patients should also be asked to wear a surgical mask while not in a single room, until advised to remove it by attending staff. The door to the room must remain closed for any patient who requires airborne precautions. Where possible only staff and visitors who have confirmed immunity (evidenced by serological immunity or vaccination history) to the specific infectious agent should enter the room, see Section 4.2.1 for further information. While appropriate personal protective equipment should be worn by all staff and visitors, those with unknown immunity or non-immune healthcare workers should be extra vigilant. While there is a paucity of evidence to confirm their effectiveness, negative pressure rooms may reduce the transmission of airborne infection within healthcare settings [174].

Standardised transmission-based precautions signage should identify the isolation room and include the necessary precautions to be adopted.

Prior to placing a patient in a negative pressure room, the pressure differential should be checked. When negative pressure rooms are in use, the pressure differential should be checked regularly, preferably daily, even if a continuous differential pressure sensing device is in use. For further information on negative pressure rooms (Class N/Type 5) see the *Australasian Health Facility Guidelines* [396], **Standard AS 1324.1: 2001, Standard AS 1324.2: 2003 and Standard AS 1668.2: 2024**.

Visitors should be restricted and screened by nursing staff, with visitors' names recorded either in a log book or in the case notes.

Transfer of patients

If transfer of the patient outside the negative pressure room is necessary, asking the patient to wear a correctly fitted surgical mask while they are being transferred and to follow respiratory hygiene and cough etiquette, as well as covering any skin lesions associated with the condition (e.g. chickenpox [varicella]) will reduce the risk of cross-transmission. Children should wear a correctly fitting mask when they are outside an isolation room. The child's oxygen saturation should be monitored.

Evidence to decision

| | |
|---|---|
| Benefits and harms | The benefits of placing patients on airborne precautions in a negative pressure room or similar clearly outweighs any undesirable effects. |
| Certainty of the evidence | This practice is supported by suggestive clinical or epidemiologic studies, and based on sound theoretical principles. |
| Values and preferences | It is expected that all patients and staff of Australian healthcare facilities would value placing patients in a room matched to their known or suspected airborne precautions. |
| Resources and other considerations | This practice would require having either a negative pressure room or a room that does not circulate air to other areas available. This may be difficult in smaller or more regional facilities, as well as in primary care facilities. |

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Placing patients on airborne precautions in a negative pressure room is justified to reduce healthcare associated infection.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

3.3 Personal protective equipment

What are the risks?

Any infectious agent transmitted by the contact or droplet route can potentially be transmitted by contamination of healthcare workers' hands, skin or clothing. Cross-contamination can then occur between the healthcare worker and other patients or healthcare workers, or between the healthcare worker and the environment. Infectious agents transmitted through droplets can also come into contact with the mucous membranes of the healthcare worker.

Personal protective equipment (PPE) refers to a variety of barriers, used alone or in combination, to protect mucous membranes, airways, skin and clothing from contact with infectious agents. PPE may include aprons, gowns, gloves, surgical masks, protective eyewear and face shields or a combination of several of these items.

In Australia, the Therapeutic Goods Administration regulates PPE if a manufacturer makes any of the following claims:

- The PPE is to be used for preventing the transmission of disease between people
- The PPE is suitable for therapeutic use, such as surgical, clinical or medical use, or use in other health services.

See the *Therapeutic Goods Administration* for further information on regulations for PPE in Australia.

The selection of PPE should be based on a risk assessment that considers the type of patient interaction, known or possible infectious agents, and/or the likely mode(s) of transmission.

When used as part of standard precautions, PPE protects against anticipated blood and body fluid exposure.

When used as part of transmission-based precautions, PPE serves as a physical barrier against the specific route(s) of transmission.

There have been few controlled clinical studies evaluating the relationship between the use of PPE and risk of healthcare associated infections. However, the use of barriers reduces opportunities for transmission of infectious agents [58]. PPE also protects patients from exposure to infectious agents in the surrounding environment carried by healthcare workers.

Decision-making about personal protective equipment

Selection of protective equipment must be based on assessment of the risk of transmission of infectious agents to the patient or carer, and the risk of contamination of the clothing or skin of healthcare workers or other staff by patients' blood, body substances, secretions or excretions.

Local policies and current health and safety legislation should also be taken into account [221].

Factors to be considered are:

- probability of exposure to blood and body substances
- type of body substance involved
- probable type and probable route of transmission of infectious agents.

Appropriate sequences and procedures for putting on and removing PPE are shown below.

All PPE must meet relevant Therapeutic Goods Administration criteria for listing on the Australian Register of Therapeutic Goods or equivalent and relevant Australian Standards. PPE should also be used in accordance with manufacturer's recommendations.

Where to wear PPE

PPE is designed and issued for a particular purpose in a protected environment and should not be worn outside that area. Protective clothing provided for staff in areas where there is high risk of contamination (e.g. operating suite/room) must be removed before leaving the area. Even where there is a lower risk of contamination, protective clothing that has been in contact with patients should not be worn outside the patient-care area. Similarly, to reduce the risk of disease transmission between patients, PPE should be removed and if necessary, replaced, before attending to another patient. When providing patient care in a room where patients are cohorted together, PPE should be changed when moving between patients in these rooms. The extended use of PPE items can increase the risk of cross infection for patients and healthcare workers and the risk of environmental contamination. Any decision to allow extended use of PPE must be made in consultation with the infection prevention and control service. Inappropriate wearing of PPE (e.g. wearing operating suite/room attire in the public areas of a hospital or wearing such attire outside the facility) may also lead to a public perception of poor practice within the facility.

Managing breaches in PPE

Health service organisations should have local policies and procedures, which at a minimum, include guidance on the immediate management of occupational exposures including breaches in PPE.

A breach in PPE may include:

- Damaged, torn or broken PPE which can result in the healthcare worker's skin, eyes, other mucous membranes or clothing being exposed or potentially contaminated
- The healthcare worker touching their face mask/ eye protection with gloved hands and potentially contaminating themselves
- The healthcare worker not wearing appropriate PPE for the level of exposure.

If there is a concern about a potential breach in PPE or potential self-contamination, notify the direct supervisor for advice on next steps. The health service organisation must have systems in place that provide healthcare workers with access to:

- Immediate first aid, if required
- Expert advice and confidential rapid assessment of the infection risk
- Ongoing support and follow-up, as required
- Access to reporting systems for workplace health and safety incidents
- Quality improvement processes to reduce the risk of occupational exposures, such as ensuring access to appropriate PPE, and staff training in the use of PPE.

Review of processes as part of an incident management review should include members of the workforce, and timely feedback on the outcomes from this review should be provided to the relevant individuals. Refer to *Section 4.2.3 Managing exposures to occupational hazards* for further information.

Patient-care tip

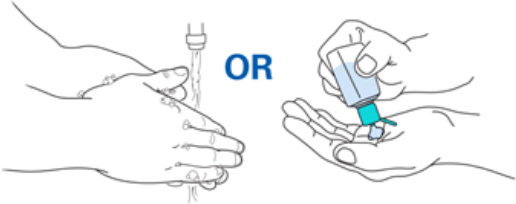
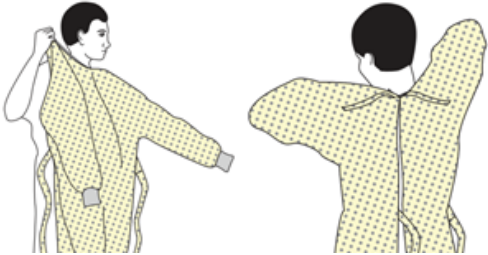


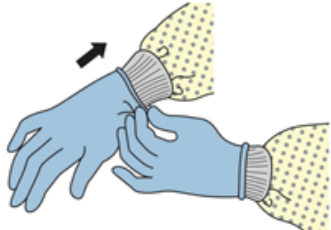
In certain settings, patients may also be required to wear PPE. However, there may be issues around compliance when dealing with specific

patient groups such as paediatric patients, or patients with dementia or claustrophobia. In these cases, other infection control measures should be applied.

Sequence for putting on and removing PPE

To reduce the risk of transmission of infectious agents, PPE must be used appropriately. The following table outlines sequences and procedures for putting on and removing PPE.

Table 14. Putting on and removing PPE

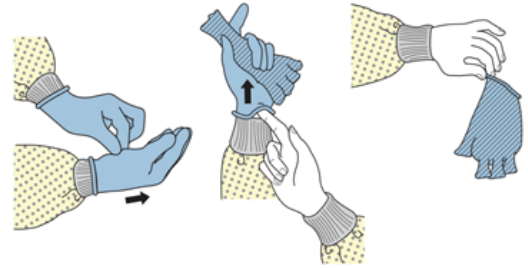
| SEQUENCE FOR PUTTING ON PPE | |
|--|---|
| Put on PPE before patient contact and generally before entering the patient room | |
| <p>1. PERFORM HAND HYGIENE</p> <ul style="list-style-type: none"> Wash hands or use an alcohol based hand rub. |  |
| <p>2. PUT ON GOWN</p> <ul style="list-style-type: none"> Fully cover torso from neck to knees, arms to end of wrists, and wrap around the back. Fasten at the back of neck and waist. |  |
| <p>3. PUT ON MASK</p> <ul style="list-style-type: none"> Secure ties or elastic bands at middle of head and neck. |  |
| <p>4. PUT ON PROTECTIVE EYEWEAR OR FACE SHIELD</p> <ul style="list-style-type: none"> Place over face and eyes and adjust to fit. |  |
| <p>5. PUT ON GLOVES</p> <ul style="list-style-type: none"> Extend to cover wrist of isolation gown. |  |

SEQUENCE FOR REMOVING PPE

Begin PPE removal at the doorway or in the anteroom

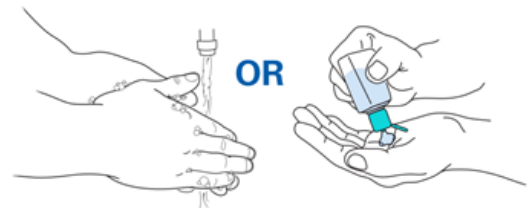
1. REMOVE GLOVES

- Outside of gloves is contaminated!
- Grasp outside of glove with opposite gloved hand; peel off.
- Hold removed glove in gloved hand.
- Slide fingers of ungloved hand under remaining glove at wrist.
- Peel glove off over first glove.
- Discard gloves in waste container.



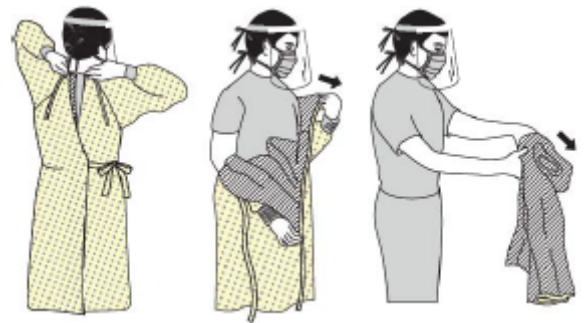
2. PERFORM HAND HYGIENE

- Wash hands or use an alcohol based hand rub.



3. REMOVE GOWN

- Gown front and sleeves are contaminated!
- Unfasten ties.
- Pull away from neck and shoulders, touching inside of gown only.
- Turn gown inside out.
- Fold or roll into a bundle and discard.



Alternatively gloves and gown can be removed as one step. Then perform hand hygiene.

4. REMOVE PROTECTIVE EYEWEAR OR FACE SHIELD

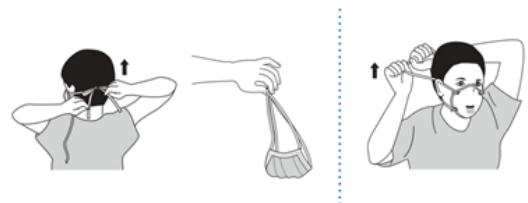
- Outside of eye protection or face shield is contaminated!
- To remove, handle by head band or ear pieces.
- Place in designated receptacle for reprocessing or in waste container.



At a minimum, perform hand hygiene if the removed PPE is contaminated.

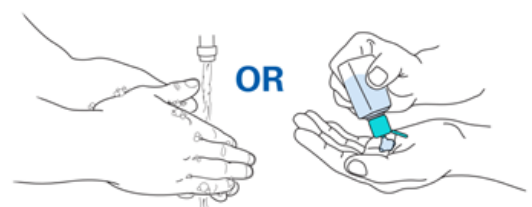
5. REMOVE MASK

- Front of mask is contaminated—DO NOT TOUCH!
- Grasp bottom, then top ties or elastics and remove.
- Discard in waste container.



6. PERFORM HAND HYGIENE

- Wash hands or use an alcohol based hand rub immediately after removing all PPE.



Adapted from CDC Guideline for Isolation Precautions [215].

Removal of PPE should be done at the doorway (just prior to leaving patient’s room) or immediately outside patient room.

Surgical or particulate masks should be removed outside of the patient room or >1 metre from symptomatic patients [87].

To remove a P2 respirator, remove gloves and gown, perform hand hygiene and step outside the room or into an anteroom. Remove protective eyewear or faceshield before removing and disposing of the respirator in a closed container and performing hand hygiene again.

At a minimum, hand hygiene should be performed after the removal of gloves as well as after the removal of any other individual item of contaminated PPE.

Sequencing may differ across healthcare settings and any variations to sequencing should be based on a risk assessment relevant to the setting and task being undertaken. Note that for surgical procedures and dentistry, the sequence for putting on PPE differs. In these situations, masks and protective eyewear are applied first prior to hand preparation. Gown and gloves are then put on (see Section 3.5.3).

Sequencing may also be different in unique circumstances such as in an Ebola outbreak. When an outbreak is detected, advice from the healthcare facilities Infection Control Team should be followed.

Conditional recommendation

Updated

29. It is suggested that clean aprons/gowns should:

- be appropriate to the task being undertaken
- be worn for a single procedure or episode of patient care where contamination with body substances is likely.

The used apron/gown should be removed in the area where the episode of patient care takes place.

Practical info

Aprons and gowns

International guidelines recommend that protective clothing (apron or gown) be worn by all healthcare workers when [58] [215]:

- close contact with the patient, materials or equipment may lead to contamination of skin, uniforms or other clothing with infectious agents
- there is a risk of contamination with blood, body substances, secretions or excretions (except sweat).

The type of apron or gown required depends on the degree of risk, including the anticipated degree of contact with infectious material and the potential for blood and body substances to penetrate through to clothes or skin. Gowns and aprons used in clinical areas should be fluid impervious.

Gowns and aprons should be changed between patients.

Aprons/gowns are routinely used upon entering the room of a patient requiring contact precautions. Clean aprons or gowns should be appropriate to the task being undertaken. This is discussed further in Section 3.2.2 Recommendation 21 practical information.

Table 15. Recommended use and characteristics of aprons/gowns

| Type | Recommended use | Characteristics |
|----------------------|--|--|
| Plastic apron | Worn for general use when there is the possibility of sprays or spills or exposure to blood or body substances during low-risk procedures [58] [215]. Worn during contact precautions when patient contact is likely. | <ul style="list-style-type: none"> • Fluid impervious • Single-use, for one procedure or episode of patient care • Disposable |
| Gown | Worn to protect the healthcare worker’s exposed body areas and | <ul style="list-style-type: none"> • Fluid impervious • Single-use |

| | | |
|-----------------------|--|---|
| | prevent contamination of clothing with blood, body substances, and other potentially infectious material [183] [193] [198] [218]. | <ul style="list-style-type: none"> • Disposable • Choice of sleeve length depends on the procedure being undertaken, the extent of risk of exposure of the healthcare worker's arms, the volume of body substances likely to be encountered, and the probable time and route of transmission of infectious agents |
| Full body gown | <p>Worn when there is a risk of contact of the healthcare worker's skin with a patient's broken skin, extensive skin to skin contact (e.g. lifting a patient with scabies), or a risk of contact with bloody and body substances which are not contained (e.g. vomiting).</p> <p>Worn when there is the possibility of extensive splashing of bloody and body substances or there is a risk of exposure to large amounts of body substances (e.g. in some operative procedures).</p> | <ul style="list-style-type: none"> • Fluid impervious • Single-use • Long sleeved so clothing and exposed upper body areas are protected • Always worn in combination with gloves and other PPE where indicated |
| Sterile gown | Worn for procedures that require an aseptic field | <ul style="list-style-type: none"> • Pre-packaged |

*Some types can be re-used. These need to be laundered or reprocessed according to **Standard AS/NZS 4146: 2000** — Laundry Practice

Note: Clinical and laboratory coats or jackets worn over personal clothing for comfort and/or purposes of identity are not considered to be PPE. These items of clothing need to be changed dependent on activity and the extent of exposure to potential pathogens.

Removing aprons and gowns

Removal of aprons and gowns before leaving the patient-care area (e.g. in the room or anteroom) prevents possible contamination of the environment outside the patient's room. Aprons and gowns should be removed in a manner that prevents contamination of clothing or skin. The outer, 'contaminated', side of the gown is turned inward and rolled into a bundle, and then discarded into a designated container for waste or linen to contain contamination (see Sections 3.1.7 & 3.1.8).

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of wearing aprons or gowns clearly outweigh any undesirable effects.

The wearing of aprons and gowns helps prevent the spread of infection to other patients through protecting healthcare workers from contamination while providing care.

A possible harm is that healthcare workers may rely on the use of an apron/gown and not undertake other appropriate infection control practices, however this can be mitigated through education and training.

Certainty of the evidence

Low

This recommendation is supported by some experimental, clinical, or epidemiological studies and a strong theoretical rationale.

A systematic review (n=4) found that acquisition rates of vancomycin-resistant enterococci (VRE) were lower when gloves and gowns were worn. All studies were observational and deemed very low quality [217].

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections during any episode of care and that a recommendation to wear aprons/gowns should be appropriate for most healthcare workers.

Resources and other considerations

No important issues with the recommended alternative

The provision and laundering of cloth gowns presents a cost to healthcare facilities, as does the provision and disposal of single use gowns/aprons. These costs would likely be outweighed by the costs and consequences of not wearing aprons and gowns.

Rationale

Wearing of gowns and aprons is justified to reduce healthcare associated infection.

Wearing of personal protective clothing (gowns and aprons) is an internationally accepted practice when:

- healthcare workers are in close contact with the patient, materials or equipment that may lead to contamination of skin, uniforms or other clothing with infectious agents
- there is a risk of contamination with blood, body substances, secretions or excretions (except sweat)

Gowns are used to protect the healthcare worker’s exposed body areas and prevent contamination of clothing with blood, body substances, and other potentially infectious material.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the ‘key info’ tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Conditional recommendation

Updated

30. It is suggested that face and eye protection should be worn during procedures that generate splashes or sprays of blood and body substances into the face and eyes.

Practical info

Face and eye protection

The mucous membranes of the mouth, nose and eyes are portals of entry for infectious agents, as are other skin surfaces if skin integrity is compromised (e.g. by acne, dermatitis) [213] [212] [197] [195].

Face and eye protection reduces the risk of exposure of healthcare workers to splashes or sprays of blood and body substances [186] [58] and is an important part of standard precautions. Procedures that generate splashes or sprays of blood and body substances, require either a face shield or a mask worn with protective eyewear [187] [191] [214] [219] [190] [206] [179].

Face and eye protection is worn as part of transmission-based precautions as discussed in Sections 3.2 - 3.2.4.

Table 16: Use of face and eye protection as part of standard precautions

| Type of care | Examples | Face and eye protection required |
|---------------------|--|--|
| Routine care | General examination (e.g. medical, physiotherapy, nursing) Routine observations | Not required unless caring for a patient on droplet precautions (surgical mask) (see Section 3.2.3) or airborne precautions (P2 respirator) (see Section 3.2.4). |

| | | |
|---|--|---|
| Procedures that generate splashes or sprays | Dental procedures Nasopharyngeal aspiration Emptying wound or catheter bag | Protective eyewear/full-length face shield Surgical mask |
| Procedures involving the respiratory tract (including the mouth) | Intubation Nasopharyngeal suction | Protective eyewear Surgical mask |

Surgical masks

Surgical masks are loose fitting, single-use items that cover the nose and mouth. They are used as part of standard precautions to keep splashes or sprays from reaching the mouth and nose of the person wearing them. They also provide some protection from respiratory secretions and are worn when caring for patients on droplet precautions. Surgical masks can be placed on coughing patients to limit potential dissemination of infectious respiratory secretions from the patient to others (see Section 3.2.3, *Recommendation 24*).

Considerations when using a surgical mask include:

- masks should be changed between patients and when they become soiled or wet
- masks should never be reapplied after they have been removed
- masks should not be left dangling around the neck
- touching the front of the mask while wearing it should be avoided
- hand hygiene should be performed upon touching or discarding a used mask.

Children should wear a specifically designed child mask and their oxygen saturation should be monitored.

Surgical masks can be categorized into three types based on the level of protection provided.

Table 17. Types of surgical masks

| Characteristics* | Level 1 barrier | Level 2 barrier | Level 3 barrier |
|---|--|--|--|
| Application | For general purpose medical procedures, where the wearer is not a risk of blood or body fluid splash, or to protect staff and/or the patient from droplet exposure to microorganisms | For use in emergency departments, dentistry, changing dressings on small wounds or healing wounds where minimal blood droplet exposure may occur | For all surgical procedures, major trauma first aid or in any area where the healthcare worker is at risk of bloody or body fluid splash |
| Bacterial filtration efficiency (BFE), % | ≥95 | ≥98 | ≥98 |
| Differential pressure, mm, H₂O/cm² | <4.0 | <5.0 | <5.0 |
| Resistance to penetration by synthetic blood, minimum pressure in mmHg for pass result | 80 mmHg | 120 mmHg | 160 mmHg |

*Note that these characteristics are based on unworn masks, and may differ or not meet performance expectations due to individual fit characteristics.

Source: Standard AS 4381: 2015

More information about different types of masks is at Section 3.2.4 - Airborne precautions.

Eye protection

Goggles with a manufacturer's anti-fog coating provide reliable, practical eye protection from splashes and sprays from multiple angles. Newer styles of goggles fit adequately over prescription glasses with minimal gaps (to be efficacious, goggles must fit snugly, particularly from the corners of the eye across the brow).

Other types of protective eyewear include safety glasses with side-shield protection, which are widely used in dentistry and other specialties that use operating microscopes [399]. While effective as eye protection, goggles and safety glasses do not provide splash or spray protection to other parts of the face.

Personal eyeglasses and contact lenses are not considered adequate eye protection. Newer styles of goggles fit adequately over prescription glasses with minimal gaps (to be efficacious, goggles must fit snugly, particularly from the corners of the eye across the brow).

Face shields

Single-use or reusable face shields may be used in addition to surgical masks, as an alternative to protective eyewear. Compared with other forms of protective eyewear, a face shield can provide protection to other parts of the face as well as the eyes. Face shields extending from chin to crown provide better face and eye protection from splashes and sprays; face shields that wrap around the sides may reduce splashes around the edge of the shield.

If healthcare workers wish to use prescription protective eyewear, the eyewear needs to meet the appropriate standard for impact as described in AS/NZS 1337.6.2012 – *Prescription eye protectors against low and medium impact*. Prescription eyewear is considered to provide appropriate eye protection for blood or body fluid splash or droplet exposure if:

- the eyewear is close fitting, particularly at the corners of the eye and across the brow, and there is minimal gaps between the frame and the face
- provides full coverage of the ocular region
- includes side protection that is indirectly vented
- can be cleaned and disinfected between use, in accordance with the facility’s equipment cleaning protocols.

Additional protective eyewear does not need to be worn with prescription protective eyewear that have these features. If prescription eyewear does not have these features, additional protective eyewear needs to be worn.

Additional facial protection may need to be considered, such as a mask or faceshield, if a large volume of blood or body fluid exposure is likely.

Removing face and eye protection

Removal of a face shield, protective eyewear and surgical mask can be performed safely after gloves have been removed and hand hygiene performed. The ties, earpieces and/or headband used to secure the equipment to the head are considered ‘clean’ and therefore safe to touch with bare hands. The front of a mask, protective eyewear or face shield is considered contaminated.

Cleaning reusable face and eye protection

Reusable face shields and protective eyewear should be cleaned according to the manufacturer’s instructions, generally with detergent solution, and be completely dry before being stored. If they are to be disinfected, they should be disinfected using either a TGA-included sterilant or medical device disinfectant - low level, or by heat as per **Standard AS 5369:2023**.

Individual actions for reducing the risk:

- Before putting on personal protective equipment (PPE) explain to the patient that it is a routine part of infection prevention and control.
- Assess the risk of spraying or splashing in the specific situation and choose PPE accordingly.
- If you have a sensitivity or allergy to latex, inform your manager and ensure you always use an alternative glove type.
- Follow appropriate sequence and procedure for putting on and removing PPE outlined in Table 14.
- Remove PPE before leaving the patient-care area and follow the sequence and procedure outlined in Table 14.
- Lead by example and champion the appropriate use of PPE in your setting.

Evidence to decision

Benefits and harms

Substantial net benefits of the recommended alternative

The benefits of wearing masks and protective eyewear when indicated clearly outweigh any undesirable effects

The benefits of reducing healthcare associated infection (HAI) transmission are clearly established. Face and protective eyewear must be worn appropriately and fitted correctly for the full benefit to be realised.

Certainty of the evidence

Low

There is little high quality evidence to support the use of masks and protective eyewear in healthcare facilities. The use of masks is supported by a strong theoretical rationale and occupational health and safety principles.

Values and preferences

No substantial variability expected

It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections during any episode of care.

Resources and other considerations

No important issues with the recommended alternative

The main resource implications of masks and protective eyewear are the purchasing costs. Indirect costs include the increased time involved in fit-testing and checking. The benefits of using masks and protective eyewear on reduced HAIs, however, outweighs these costs.

Consideration to how an adequate supply of masks will be maintained during a potential outbreak or pandemics must be given.

Rationale

Whilst there is little high quality evidence to support the use of face and protective eyewear, their use is supported by work health and safety principles and expert advice.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Conditional recommendation

Updated

31. It is suggested that single-use, fit for purpose gloves are worn for:

- each invasive procedure
- contact with sterile sites and non-intact skin or mucous membranes
- activity that has been assessed as carrying a risk of exposure to blood and body substances.

Hand hygiene should be performed prior to donning gloves and after gloves are removed.

Gloves must be changed between patients and after every episode of individual care.

Practical info

Gloves

Gloves can protect both patients and healthcare workers from exposure to infectious agents that may be carried on hands [189]. As part of standard precautions, they are used to prevent contamination of healthcare workers' hands when [215]:

- anticipating direct contact with blood or body substances, mucous membranes, non-intact skin and other potentially infectious material
- handling or touching visibly or potentially contaminated patient-care equipment and environmental surfaces [19] [181] [189].

The capacity of gloves to protect healthcare workers from transmission of blood borne infectious agents following a needlestick or other puncture that penetrates the glove barrier has not been determined [215].

Appropriate use of gloves is an essential component of standard precautions (see Section 2.1 and 3.1).

When and how should gloves be worn?

As with all personal protective equipment (PPE), the need for gloves is based on careful assessment of the task to be carried out, the related risk of transmission of microorganisms to the patient; and the risk of contamination of the healthcare worker's clothing and skin by the patient's blood and body substances [215] [58]. Risk assessment includes consideration of:

- who is at risk (whether it is the patient or the healthcare worker)
- whether sterile or non-sterile gloves are required, based on contact with susceptible sites or clinical devices and the aspect of care or treatment to be undertaken
- the potential for exposure to blood or body substances
- whether there will be contact with non-intact skin or mucous membranes during general care and invasive procedures
- whether contaminated instruments will be handled.

When gloves are worn in combination with other PPE, they are put on last (see Table 14).

What type of gloves should be worn?

Non-sterile single-use medical gloves are available in a variety of materials, the most common being natural rubber latex (NRL) and synthetic materials (e.g. nitrile). NRL remains the material of choice due to its efficacy in protecting against blood borne viruses and properties that enable the wearer to maintain dexterity [207] [221]. However, sensitivity to NRL in patients, carers and healthcare workers may occur (see below) and must be documented. A local policy is required on using alternative glove types when patients have latex allergies.

The selection of glove type for non-surgical use is based on a number of factors [202] [182] [200] [210]:

- the task to be performed (i.e. glove type should be fit for purpose and aim to avoid interference with dexterity, friction, excessive sweating or finger and hand muscle fatigue)
- anticipated contact with chemicals and chemotherapeutic agents
- personal factors, such as latex sensitivity and size.

Facility policies for creating a latex-free environment should also be taken into account.

Table 18. Selection of glove type

| Glove | Indications for use | Examples |
|--|---|--|
| Non-sterile gloves | <ul style="list-style-type: none"> • Potential for exposure to blood, body substances, secretions or excretions • Contact with non-intact skin or mucous membranes • Anticipated contact with chemicals and chemotherapeutic agents* | <ul style="list-style-type: none"> • Venepuncture • Vaginal examination • Dental examination • Emptying a urinary catheter bag • Naso-gastric aspiration • Management of minor cuts and abrasions • Handling and administration of chemotherapeutic agents* • Handling chemicals, such as cleaning agents/wipes* |
| *Gloves used for the handling or in the administration of chemotherapeutic agents must be compliant with EN ISO 374-5:2016 and ASTM D6978-05(2019) | | |
| Sterile gloves (see 3.1.8 Aseptic technique for further detail) | <ul style="list-style-type: none"> • Potential for exposure to blood, body substances, secretions or excretions • Contact with susceptible sites or clinical devices where sterile conditions should be maintained | Surgical aseptic technique procedures e.g. <ul style="list-style-type: none"> • Urinary catheter insertion • Complex dressings • Central venous line insertion site dressing • Lumbar puncture • Clinical care of surgical wounds or drainage sites • Dental procedures requiring a sterile field |
| Reusable utility gloves† | <ul style="list-style-type: none"> • Indicated for <u>non-patient-care</u> activities • Utility gloves may be decontaminated for re-use (according to the glove manufacturer's directions) | <ul style="list-style-type: none"> • Worn for cleaning the environment or cleaning and disinfecting patient care equipment • Instrument cleaning in sterilising |

| | | |
|--|--|---------------|
| | provided the integrity of the glove is not compromised | services unit |
| <p>† The use of reusable utility gloves is preferable for the handling of cleaning products and supports the sustainable use of equipment in health service organisations. Reusable utility gloves should be reserved for use during environmental cleaning or in reprocessing units. Non-sterile, single use gloves may also be suitable for the handling of cleaning products (e.g. neutral detergent and cleaning wipes etc) in patient care areas, if not contraindicated by the cleaning product manufacturer's Instructions for Use, or work health and safety requirements for the safe handling of the cleaning product.</p> | | |
| Gloves suitable for clinical use | | |
| NRL (latex) gloves | <ul style="list-style-type: none"> • Preferable for clinical procedures that require manual dexterity and/or will involve more than brief patient contact • Select powder-free latex gloves to minimise the risk of latex sensitivity or allergies | |
| Synthetic gloves (e.g. nitrile) | <ul style="list-style-type: none"> • Procedures involving high risk of exposure to blood-borne virus and where high barrier protection is needed • Provides suitable alternative to latex if there are no issues with glove fit or sensitivity | |

Sources: Derived from Kotilainen et al 1989 [204]; Korniewicz et al 1989 [203]; Korniewicz et al 1993 [201]; Rego & Roley 1999 [211]; Loveday et al 2014 [79]; Korniewicz et al 2002 [199]; Sehulster & Chinn 2007 [111]; Siegel et al 2007 [215].

When should gloves be changed?

Gloves (other than utility gloves) should be treated as single-use items, and discarded, not washed. This is because infectious agents cannot be reliably removed from glove surfaces and continued glove integrity cannot be ensured.

International guidance suggests that changing of gloves is necessary:

- between episodes of care for different patients, to prevent transmission of infectious material [58] [215]
- during the care of a single patient, to prevent cross-contamination of body sites [19] [215]
- if the patient interaction involves touching portable computer keyboards, other portable devices or any other mobile equipment that is transported from room to room [215].

Prolonged and indiscriminate use of gloves should be avoided for a number of important reasons. A failure to change gloves between procedures has been found to increase the risk of cross-transmission [225] [226] and has been associated with transmission of methicillin-resistant *Staphylococcus aureus* (MRSA) and Gram- negative bacilli [188] [208] [209]. Prolonged glove use may also cause adverse reactions and skin sensitivity [58] [215].

Further, the use of gloves must be in addition to standard hand hygiene practices. Hand hygiene should always be performed before putting on gloves and after removing them [209] [216] [19].

Latex allergy

Latex allergy is a reaction to certain proteins in latex rubber. The amount of latex exposure needed to produce sensitisation or an allergic reaction is unknown. However, current understanding of latex allergy is as follows: [220]

- increasing the exposure to latex proteins increases the risk of developing allergic symptoms— most people who are allergic to latex have had frequent exposure to latex over many years; the majority are nurses, doctors, dentists or patients who have had a number of operations
- in sensitised people, symptoms usually begin within minutes of exposure; but they can occur hours later and can be quite varied—mild reactions involve skin redness, rash, hives, or itching; more severe reactions may involve respiratory symptoms such as runny nose, sneezing, itchy eyes, scratchy throat, and asthma (difficult breathing, coughing spells, and wheezing); and rarely, shock may occur although a life-threatening reaction is seldom the first sign of latex allergy
- the risk of latex allergy is influenced by the amount of protein/allergen and powder in the latex glove; not by powder alone [196].

Healthcare workers with latex allergies should inform their managers to ensure that their work areas can be latex free.

If latex gloves are used, they should be non-powdered due to the risks associated with aerosolisation and an increased risk of latex allergies.

Removing and disposing of gloves

When removing gloves, care should be taken not to contaminate the hands. After gloves have been removed, hand hygiene should be performed in case infectious agents have penetrated through unrecognised tears or have contaminated the hands during glove removal.

Gloves should be disposed of as soon as they are removed, with disposal complying with local policies and standards.

Individual actions for reducing the risk:

- Before putting on PPE explain to the patient that it is a routine part of infection prevention and control.
- Assess the risk of spraying or splashing in the specific situation and choose PPE accordingly.
- If you have a sensitivity or allergy to latex, inform your manager and ensure you always use an alternative glove type.
- Follow appropriate sequence and procedure for putting on and removing PPE outlined in Table 14.
- Remove PPE before leaving the patient-care area and follow the sequence and procedure outlined in Table 14.
- Lead by example and champion the appropriate use of PPE in your setting.

For practical information on the use of gloves, see Case Study 5.2.

Evidence to decision

Benefits and harms The benefits of wearing gloves clearly outweigh any undesirable effects. Gloves can protect both patients and healthcare workers from exposure to infectious agents that may be carried on hands. Gloves are an essential component of standard precautions.

Whilst the benefits of appropriate glove usage in reducing the transmission of infection are well-documented, evidence from numerous studies demonstrates that the rate of compliance with hand hygiene is often worse when gloves are worn [223] [224]. The reason for this association, however, has not been investigated thoroughly.

Certainty of the evidence This advice is based on limited evidence, but on sound theoretical principles and supported by expert advice.

Values and preferences It is expected that all patients and staff of Australian healthcare facilities would highly value minimising infections during any episode of care.

Resources and other considerations The provision of gloves presents cost to healthcare facilities.

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. The use of gloves is justified to reduce healthcare associated infection.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 good practice points to be reassigned as a GRADE practice statement. All considerations in adopting or adapting this practice statement are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

3.3.1 Other items of clothing

Ties and lanyards

Where possible, the wearing of lanyards and neckties should be avoided as evidence indicates these pieces of clothes may facilitate transmission of infection [205] [227] [228] [229] [230].

Footwear

Footwear suitable for the duties being undertaken must be worn and preferably be designed to minimise the risk of injury from dropped sharps, as well as minimise risk of exposure to blood and body substances.

Uniforms

In areas of clinical practice where there is a high risk of repeated exposure to blood and other body substances, it is recommended that uniforms be worn as well as the appropriate personal protective equipment.

Whilst no clinical studies that have demonstrated cross-transmission of healthcare associated pathogens via standard apparel, a number of small prospective trials have demonstrated that the uniforms of healthcare professionals can become contaminated with a variety of pathogens [180].

Healthcare workers should wear a clean uniform for each shift. Uniforms should be washed at home, separately from other items. If the uniform has been contaminated with blood or body substances, the hospital facility laundry must be used [207].

3.4 Management of multi-resistant organisms and outbreak situations

Summary

Section 3.4 outlines approaches to the management of multi-resistant organisms (MROs) or outbreak situations.

For the purpose of these Guidelines, MROs are taken to include: methicillin-resistant *Staphylococcus aureus* (MRSA), vancomycin-resistant enterococci (VRE), and multi-resistant Gram-negative bacteria (MRGN).

- The decision to screen for MROs when a patient is admitted to a healthcare facility should be dependent upon the specific MRO, any identified patient-risk factors and the local epidemiology of the MRO.
- MRO clearance processes are well-defined for MRSA however there is less evidence regarding clearance for MRGNs (including carbapenemase-producing *Enterobacteriales* (CPE)) and VRE, and decisions should be made based on state and territory policy and after consultation with an infection prevention and control expert.
- Applying standard precautions, including hand hygiene, is the most effective measure to prevent and control the spread of MROs.
- It is suggested that transmission-based precautions be considered for all patients colonised or infected with an MRO.
- Maintaining a surveillance system to record the presence of all MROs can assist in the timely reporting and notification of cases or outbreaks.
- Outbreaks of MROs should be managed by a dedicated outbreak control team, and appropriate infection control strategies should be implemented.

Patient-care tip

When a patient is infected or colonised with an MRO or involved in an outbreak, there is potential for adverse effects such as anxiety, mood disturbances, perceptions of stigma and reduced contact with clinical staff. Clearly explaining to patients the measures being undertaken and why they are necessary may help to alleviate these effects. At no time should a patient's MRO status interfere with the provision of appropriate, high quality care.

3.4.1 Multi-resistant organisms

What are the risks?

Multi-resistant organism (MROs), which are predominantly bacteria, are resistant to multiple classes of antimicrobial agents. Antimicrobial resistance increases the morbidity and mortality associated with infections, and contributes to increased costs of care due to prolonged hospital stays and other factors, including the need for more expensive drugs [239]. A major cause of antimicrobial resistance is the exposure of a high-density, high-acuity patient population in frequent contact with healthcare workers to extensive antimicrobial use, along with the attendant risk of cross-infection [232] [233].

For the purpose of these guidelines, MROs are taken to include:

- **all methicillin-resistant *Staphylococcus aureus* (MRSAs)**, which are responsible for up to a third of healthcare associated bloodstream infections (BSI) [232]. Mortality from MRSA related BSI ranged from 10% to 50%, according to the setting [234]
- **all vancomycin-resistant enterococci (VREs)** with mobile resistance determinants (e.g. VanA, VanB)—the ratio of invasive VRE infection to colonisation appears to be proportionately lower than that of MRSAs [232]
- **a range of Gram-negative bacteria (MRGNs)** with multiple classes of drug resistance or resistant mechanisms to critically important antimicrobials—highly transmissible resistance is a particular feature of antimicrobial resistance among the Gram-negative bacteria, especially the *Enterobacteriales*. In particular, an increasing number of carbapenemase-producing *Enterobacteriales* (CPE) cases and outbreaks have been identified in Australia and this is recognised as an emerging public health issue [231]. Multi-drug resistance is also common and increasing among non-fermenting Gram-negative bacteria (e.g. *Pseudomonas aeruginosa* and *Acinetobacter baumannii*) and a number of strains have now been identified that exhibit resistance to essentially all commonly used antimicrobials. These organisms are associated with treatment failure and increased morbidity [232].

A two-level approach is necessary for the prevention and control of MROs. This involves implementation of:

- **core strategies** for MRO prevention and control in any situation where MRO infection or colonisation is suspected or identified (see *Recommendation 33*)
- **organism-based or resistance mechanism-based approaches** if incidence or prevalence of MROs are not decreasing despite implementation of the core strategies (see below for further information).

In the event of an MRO outbreak, investigation and control/containment should be conducted as outlined in Section 3.4.2.

The best practices in these guidelines are based on the assumption that healthcare settings already have basic infection prevention and control systems in place. If this is not the case, healthcare settings will find it challenging to implement the practices recommended for the management of MROs. These healthcare settings must work with organisations that have infection prevention and control expertise, such as: academic health science centres, regional infection control networks, public health units that have professional staff certified in infection prevention and control, and local infection prevention and control associations to develop evidence-based programs [261].

Organism-specific approach

When the incidence or prevalence of MROs is not decreasing despite implementation of the core strategies outlined in *Recommendation 33*, further measures to control transmission need to be considered. A risk management approach focuses on:

- the type of MRO (e.g. prioritisation of available isolation facilities according to MRO)
- the healthcare area (e.g. intensive care or haematology/oncology units have higher risks of transmission)
- patient factors (e.g. whether the consequences of infection are severe)
- available resources (e.g. whether screening a certain patient population is feasible)
- whether interventions to interrupt transmission are available (e.g. decolonisation for MRSA).

Further measures may include:

- *Targeted screening*—timely active screening to identify colonised patients combined with the use of contact precautions for the care of colonised patients has been followed by a significant reduction in the rates of both colonisation and infection of patients with MRSA [245] [246]. Screening involves collecting specimens from the patient and subsequent laboratory analysis of these samples. In a risk assessment approach to screening, considerations include the endemicity of the MRO, the prevalence of MRO infection and the likelihood of MRO carriage. Clinicians and the infection control professional should be informed of both negative and positive screening results promptly. If screening returns a positive sample, contact precautions should be applied and appropriate use of isolation and cohorting facilities should be implemented.
- *Decolonisation*—interventions may be:
 - topical—whole body washes (using chlorhexidine) and topically applied antimicrobial agents
 - systemic—orally administered antimicrobials (tetracyclines, fusidic acid, ciprofloxacin, rifampin and trimethoprim-sulfamethoxazole)
 - combinations of systemic and topical therapy.
- *Surveillance and timely feedback*—increased surveillance may be appropriate to monitor the effect of interventions designed to control particular MROs. Surveillance information should be fed back to health care workers and facility management promptly.

Screening

In acute-care settings, routine screening for MROs for all admitted patients is not encouraged. The decision to screen for specific MROs should be based on the level of risk and the local epidemiology of the specific MRO. Control measures specific to local factors should be determined and endorsed by the healthcare facility management structure, and the screening protocols for MROs should be influenced by the:

- local prevalence of the MRO
- the reason for admission of the patient
- the potential risk of transmission to others
- the risk status of the unit to which the patient is admitted
- the likelihood that the patient is carrying an MRO.

As a minimum standard to reduce the risk of transmission of MROs, it is recommended that the following approaches to screening be implemented. This guidance for screening is based on patient risk factors for these organisms. Other risk groups may be defined by local experience, based on screening initiatives or outbreak epidemiology. Expert direction and resources allocation is required for effective MRO screening.

For example some facilities have found that screening patients who are recent hospital admissions from international facilities into Australian facilities have increasingly been shown to be positive for MRGN. While this is an area for future research, healthcare facilities could currently consider screening these patients on admission, particularly in areas where MROs are found to be prevalent in transferred patients.

Table 19. Suggested approach to routinely screening for MRSA

| Organism | Screen who | Screen when | Sample collection |
|----------|---|--|---|
| MRSA | <p>1. Patients with any of the following risk factors:</p> <ul style="list-style-type: none"> • previous infection or colonisation of MRSA • frequent re-admissions to any healthcare facility • transfers from other acute care facility, particularly one known to have a high prevalence of MRSA • residence in long term care facilities • a wound that has not healed (e.g. burn, surgical incision, chronic wounds) • a number of co-morbidities • locales or populations with a high prevalence of community strains of MRSA • a long-term indwelling medical device. | <ul style="list-style-type: none"> • Screened routinely at the time of admission unless they are being admitted directly to isolation facilities and it is not planned to attempt to clear them of MRSA carriage. | <ul style="list-style-type: none"> • Multiple sites including one from the nose and another mucosal surface. • Reasonable sites to swab include anterior nares, skin lesions and wounds, sites of catheters, catheter urine, ostomy sites, groin/perineum, tracheostomy and other skin break in all patients, and sputum from patients with a productive cough. • Where maximum sensitivity is required, consideration should be given to adding a throat swab. The umbilicus should be sampled in all neonates. |
| | <p>2. Healthcare workers epidemiologically linked to single-strain outbreak in health care facility.</p> | <ul style="list-style-type: none"> • After confirmation of epidemiological evidence. • Two weeks after decolonisation therapy. | |
| | <p>3. High risk units and/or procedures:</p> <ul style="list-style-type: none"> • intensive care unit (ICU)/high dependency unit • spinal unit • burns unit • pre-operative clinics • Patients with planned prosthetic surgery (joint replacement, cardiothoracic surgery) | <ul style="list-style-type: none"> • All patients on admission, discharge and once weekly. | |

Table 20. Suggested approach to routinely screening for VRE and MRGN

| Organism | Screen who | Screen when | Sample collection |
|----------|---|---|--|
| VRE | <p>1. High risk units:</p> <ul style="list-style-type: none"> • ICU | <ul style="list-style-type: none"> • For endemic VRE screen on admission to intensive care unit, | <ul style="list-style-type: none"> • Multiple sites including rectal or perianal swabs. |

| Organism | Screen who | Screen when | Sample collection |
|---|--|---|--|
| | <ul style="list-style-type: none"> nephrology haematology solid organ transplant unit, and patients epidemiologically linked to single-strain outbreak in health care facility. <p>1. Patients at high risk of carriage:</p> <ul style="list-style-type: none"> recent hospitalisation in any health care facility critical illness in intensive care units long duration of stay and severity of illness chronic disease and impaired functional status patients with urinary catheters, and/or prolonged or broad-spectrum antimicrobial use, particularly vancomycin. | <p>at discharge and once weekly.</p> <ul style="list-style-type: none"> For VRE in a haematology/ oncology facility screen periodically every 3-6 months. | <ul style="list-style-type: none"> Reasonable sites include groin, wounds, ostomy sites and respiratory secretions or tracheal aspirates. |
| <p>MRGN, ESBLs, plasmid-mediated AmpC, MR-<i>Pseudomonas aeruginosa</i>, MR-<i>Acinetobacter baumannii</i>, carbapenemase-producing <i>Enterobacteriales</i> (CPE, see below for additional CPE-specific risk factors)</p> | <p>1. High risk units:</p> <ul style="list-style-type: none"> ICU solid-organ transplant unit speciality centres (e.g. burns, neurosurgery), and patients epidemiologically linked to single-strain outbreak in healthcare facility. <p>2. Patients at high risk of carriage:</p> <ul style="list-style-type: none"> those with recent broad spectrum antimicrobial therapy (carbapenem, quinolones, and 3rd and 4th generation cephalosporins) history of recent travel to area of high endemicity long duration of stay and severity of illness chronic disease and impaired functional status, and/or presence of invasive medical devices Admission screening of MRGNs should be based on local epidemiology and consideration of high-risk groups. | <ul style="list-style-type: none"> If screening and isolation are not routine, they may be considered to address a temporary outbreak in a specific unit and discontinued once the outbreak is controlled. | <ul style="list-style-type: none"> Multiple sites including rectal or perianal swabs. Reasonable sites to include are nares, groin, wounds, ostomy sites and respiratory secretions or tracheal aspirates depending on the infectious agent. |
| <p>Carbapenemase-producing <i>Enterobacteriales</i> (CPE)</p> | <p><i>In addition to the above considerations for screening of MRGNs, the following risk</i></p> | <ul style="list-style-type: none"> Patients at high-risk of colonisation or infection with CPE should be actively screened | <ul style="list-style-type: none"> Aspirates from any tubes or drains. Rectal swabs, faeces or urine |

| Organism | Screen who | Screen when | Sample collection |
|----------|---|--|---|
| | <p><i>factors should also be considered for CPE [231].</i> <i>Patients who:</i></p> <ul style="list-style-type: none"> • are hospitalised for a long time • have been hospitalised or had surgery overseas • have had multiple or recent exposures to different antibiotic agents, especially cephalosporins, fluoroquinolones and carbapenems • have diabetes mellitus • are on mechanical ventilation • have an indwelling medical device, and/or • are recipients of an organ or stem cell transplant. | for CPE colonisation or infection upon hospital admission. | from catheterised patients. Specimens from open wounds should also be considered [231]. |

Screening for patients in home care and other community-based settings

The current evidence does not support routine screening for MROs in home care and community-based settings such as residential aged care. In these settings, the use of standard precautions as part of routine practice should assist in minimising the cross-transmission risks of infection, regardless of multi-resistant organism status.

Any patients with risk factors for transmission, such as a discharging wound, should have a risk assessment performed to determine whether any transmission-based precautions should be implemented.

MRO clearance

The evidence is still emerging on the most effective method of demonstrating clearance of a particular MRO. This is an area that warrants further research.

MRSA

The following criteria should be satisfied prior to certifying that a patient has cleared MRSA:

- more than three months elapsed time from the last positive specimen
- all wounds healed, and no indwelling medical devices present (this may differ across the states and territories—refer to local policy for clarification)
- no exposure to any antimicrobial or antiseptic body wash for at least two weeks prior to screening
- no exposure to specific anti-MRSA antimicrobial therapy in the past three months
- consecutive negative screens from above screening sites on two separate occasions.

It is important to note that colonisation of MRSA may continue for a prolonged period of time, and that MRSA can resurface if a patient is hospitalised or prescribed antimicrobials. Patients may need to undertake clearance again if re-admitted to an acute care facility, particularly a high-risk unit, as the previous clearance is only reflective of a patient's MRSA status at the time the swabs were taken.

VRE

There is no agreed protocol for VRE clearance and caution should be applied. Some patients with VRE may appear to 'clear' with time but relapse with antimicrobial therapy.

Patients should be 'cleared' of VRE for the purposes of discontinuing contact precautions in the acute setting on an individual basis. In this instance, it is reasonable to follow the above criteria for MRSA and MRGN. Expert direction from senior infection control experts should be sought on all VRE clearances.

MRGNs (excluding CPE)

There is not yet an agreed protocol for the clearance of MRGN carriage due to a lack of scientific evidence. However, patients can have their MRO alert retired and can be managed with standard precautions if the following criteria are met:

- more than three months elapsed since time from last positive specimen
- all discharging wounds healed
- no indwelling medical devices present
- no enterostomy or tracheostomy present
- no antimicrobial therapy in the past three months
- consecutive negative screens from specified screening sites (above) obtained on separate occasions at least three weeks apart.

Recommendations regarding MRGN clearance are continuing to evolve with emerging evidence, and infection control professionals should be involved in determining appropriate clearance protocols.

CPE

The following recommendations are in place regarding the clearance of CPE [231]:

- a patient colonised with CPE cannot be considered cleared within 12 months of a positive result
- clearance must be assessed based on relevant state and territory policies, and in consultation with an infection prevention and control professional, and a clinical microbiologist or infectious diseases physician
- any person deemed cleared should be rescreened at every subsequent overnight admission to a healthcare facility to identify any relapse in detectable CPE colonisation
- a healthcare facility can consider ceasing contact precautions for a patient readmitted to hospital more than 12 months since a positive result of CPE colonisation if they have no risk factors. This requires three negative screening swabs at least 24 hours apart.

In absence of high-quality evidence to show that clearance of colonisation will occur, a cautious approach to determining clearance for CPE is required.

Emerging MROs

It is acknowledged that Australia has had very few identified cases of *Candida auris* to date. All of the cases found in Australia so far have been in people who were in hospital overseas where *C. auris* is more common.

Antimicrobial stewardship

Safe and appropriate use of antimicrobials is a strategic goal of the clinical governance system [299]. Over the last 50 years, the prevalence of MROs such as MRSA has risen alarmingly, initially mainly in hospitals but now increasingly in the community. There is good evidence that overall rates of antimicrobial resistance correlate with the total quantity of antimicrobials used, as determined by the number of individuals treated, prior exposure and the average duration of each treatment course. In individuals, the risk of colonisation and infection with MROs correlates strongly with previous antimicrobial therapy.

For more information about antimicrobial resistance and antimicrobial stewardship programs, see Section 4.5.

Conditional recommendation

Updated

32. It is suggested that contact precautions be considered for all patients colonised or infected with a multi-resistant organism (MRO) where there is anticipated patient and/or environmental contact, including:

- performing hand hygiene and putting on gloves and gowns before entering the patient-care area
- using patient-dedicated or single-use non-critical patient-care equipment
- using a single-patient room or, if unavailable, cohorting patients with the same strain of MRO in designated patient-care areas (upon approval from the healthcare facility's Infection Control Team)
- ensuring consistent cleaning and disinfection of surfaces in close proximity to the patient and those likely to be touched by the patient and healthcare workers.

Practical info**Core strategies for multi-resistant organisms (MRO) prevention and control**

Successful control of MROs is based on a combination of interventions with a shift over the last decade towards the engagement and participation of patients in infection control strategies. These involve continued rigorous adherence to hand hygiene, appropriate use of personal protective equipment (PPE) and implementation of specific contact-based precautions (isolation of infected or colonised patients, increased environmental cleaning and patient-dedicated equipment) until patients are culture-negative for a target MRO or have been discharged from the facility.

In non-acute healthcare settings, general measures of infection control (particularly hand hygiene by both patients and healthcare workers) may be enough to prevent transmission. However, contact precautions, such as gowns and gloves, may be necessary if the patient is heavily colonised or there is known continuing transmission. Local guidelines and circumstances should determine practice in settings where the patient population is vulnerable [236].

There is emerging evidence which suggests that there are a range of possible negative patient outcomes associated with the use of contact precautions for patients infected or colonised with an MRO which need to be considered. These are discussed below.

Organisational measures—such as staff education on prevention and management of MRO transmission, antimicrobial stewardship programs, and appropriate response to active surveillance cultures—are discussed in Section 4.

Hand hygiene

MROs can be carried from one person to another via the hands of a healthcare worker. Contamination can occur during patient care or from contact with environmental surfaces in close proximity to the patient, particularly when patients have diarrhoea and the reservoir of the MRO is the gastrointestinal tract. Effective hand hygiene is therefore the most important measure to prevent and control the spread of MROs. Alcohol-based hand rub of at least 70% v/v ethanol or equivalent has been shown to be effective against methicillin-resistant *Staphylococcus aureus* (MRSA) and vancomycin-resistant enterococci (VRE) [238].

Personal protective equipment

Both direct patient contact (e.g. routine patient care) and indirect contact (e.g. involving environmental contamination) can lead to contamination of the healthcare worker's hands and clothing. Appropriate use of gloves has been found to be as effective a strategy as patient isolation in containing MROs, particularly when isolation may not be feasible [240] [241]. Evidence suggests the use of gowns is shown to be effective against MRO transmission where contact precautions are being applied [235]. Glove use is more effective when combined with wearing of gowns [242] [243] [244]. However, it must also be acknowledged that implementation issues such as cost and availability of resources may prohibit universal gown use [235].

Recommendation 29 provides further guidance on the selection of appropriate gowns and Recommendation 31 on selection of gloves.

Isolation

Placing colonised or infected patients in single rooms, cohort rooms or cohort areas as a component of a multifaceted infection control policy can reduce acquisition rate and infection with MROs in acute-care settings. Cohorting patients with the same strain of MRO has been used extensively for managing outbreaks of specific MROs, including MRSA, VRE, extended spectrum beta-lactamase (ESBL)-producing bacteria, and *Pseudomonas aeruginosa*.

However, it is not always appropriate to cohort patients with the same MRO species if they have a different resistance mechanism or phenotype (e.g. if one has a community-acquired strain of likely panton-valentine leukocidin (PVL)-positive MRSA and the other has a hospital-acquired strain of MRSA). CPE positive patients should not be grouped together without expert direction from the Infection Control Team [231].

Decisions regarding priority of isolation when demand for single rooms exceeds availability should be made by the Infection Control Team based on the patients, their acuity and the types and strains of MRO present. Priority should always be given to patients requiring airborne precautions. There may also be other competing priorities for single rooms not related to infectious diseases and these may need to be considered when allocating resources e.g. patient security and palliation.

In long-term care facilities, isolation and cohorting may not be possible, so hand hygiene with appropriate routine use of gloves for individual resident and environmental contact is preferred [240].

Due to the varying nature of healthcare facilities, it is not feasible to provide a generic policy on the movement of patients with MROs. This needs to occur at a local level and be relevant to the patient's treatment plan. These policies should not limit access to treatment and should consider the social implications of managing a patient with an MRO.

Information about the reported harms of patient isolation as part of contact precautions are discussed below.

Environmental cleaning

In acute-care areas where the risk of patient vulnerability and risk of cross infection due to the presence of an MRO is high, contact precautions should be followed. This will require all patient surrounds and frequently touched objects (e.g. bedrails, trolleys, bedside commodes, doorknobs, light switches or tap handles, ensuite facilities) to be cleaned with a suitable detergent and disinfected with a Therapeutic Goods Administration (TGA)-listed hospital-grade disinfectant (or sodium hypochlorite if indicated for use as per *Recommendation 14*). As outlined in Section 3.1.3 *Practice Statement 9* practical information, this process must involve either:

- a 2-step clean, which involves a physical clean using detergent solution followed by use of a chemical disinfectant
- a 2-in-1 clean in which a combined detergent/disinfectant wipe or solution is used and mechanical/manual cleaning action is involved.

Sole reliance on a disinfectant without mechanical/manual cleaning is not recommended.

Patient equipment

Standard precautions concerning patient-care equipment are very important in the care of patients with MROs. Patient-care devices (e.g. electronic thermometers) may transmit infectious agents if devices are shared between patients. To reduce the risk of transmission, disposable or patient dedicated equipment is preferred. Section 3.1.4 provides more detailed information on reusable instruments and equipment.

Monitoring

Monitoring of the incidence of target MRO infection and colonisation should continue after these interventions are implemented. If rates do not decrease, more interventions may be needed to reduce MRO transmission.

Management of specific MROs

MRSA

In addition to standard precautions and contact precautions, droplet precautions should be used for patients known to be infected or colonised with MRSA in the lower respiratory tract when patient care activities are likely to expose healthcare workers.

Patients positive for MRSA should have an electronic alert placed on their case record for easy identification on readmission. The Infection Control Team may prescribe a decolonisation program for patients depending on their level of risk from ongoing colonisation and availability for follow-up. Consider topical plus/minus systemic decolonisation for:

- healthcare workers epidemiologically linked to transmission
- patients having prolonged hospitalisation
- patients with chronic conditions likely to be readmitted (e.g. haemodialysis)
- patients prior to undergoing high-risk elective surgery such as cardiac and implant surgery.

VRE

Patients positive for VRE should have an electronic alert placed on their case record for easy identification on readmission.

Management of patients with VRE depends on the potential risks involved.

- It is suggested that patients with suspected or confirmed VRE are managed using contact precautions, including placement in a single room with dedicated toilet facilities. Infection control precautions should concentrate on managing the risk of VRE transmission, including the risk of environmental transmission, and take into account the presence of other patients who may be at high risk of infection with VRE.
- Emerging evidence suggests that in appropriate circumstances, patients positive for VRE may be managed with standard precautions (i.e. hand hygiene, environmental cleaning, cleaning of shared patient equipment between use and frequently touched surfaces, and appropriate use of disinfectants) provided the patient does not have risk factors that facilitate transmission such as diarrhoea or poor compliance with personal hygiene.
- This decision should be based on a risk assessment performed by the Infection Control Team, and in the context of a locally agreed policy relating to management of patients with VRE. This decision should also be clearly outlined in a policy approved through the organisations governance committee. If these processes cannot be adequately completed by the healthcare facility, contact precautions should be implemented for all patients with suspected or confirmed VRE.

Further information on the use of horizontal measures to prevent transmission of MROs is discussed below.

MRGNs

Patients positive for MRGN should have an electronic alert placed on their case record for easy identification on readmission. It is

suggested that patients with suspected or confirmed MRGNs are managed through contact precautions, including placement in a single room with dedicated toilet facilities.

CPE

Patients positive for CPE should have an electronic alert placed on their case record for easy identification on readmission. It is suggested that patients with suspected or confirmed CPE are managed through contact precautions, including placement in a single room with dedicated toilet facilities. Priority for isolation should be dependent upon the risk of transmission for each patient (e.g. patients with diarrhoea, patients with medical devices in situ). These precautions should remain in place until the patient is discharged from the facility. CPE positive patients should not be cohorted without guidance from the Infection Control Team.

Emerging evidence on contact precautions

It is suggested that contact precautions be considered for all patients colonised or infected with an MRO where there is anticipated patient and/or environmental contact. Variations to this should be determined via a risk assessment with consideration for the healthcare setting. If there is no contact with the patient or the patient's environment (e.g. briefly speaking with the patient) then there may not be a need for the healthcare worker to apply contact precautions.

Increasingly, some healthcare facilities are focusing on the use of horizontal measures to prevent transmission of MROs. These include hand hygiene, bathing patients with chlorhexidine (see *Recommendation 38*) and environmental cleaning and disinfection [250]. It is important that healthcare facilities using horizontal measures implement processes and policies surrounding the use for patients infected or colonised with an MRO, and monitor the impact of these measures on the transmission of MROs through process and outcome reporting.

There is emerging evidence that the use of contact precautions for patients colonised or infected with an MRO may be associated with potential negative outcomes such as decreased patient satisfaction with care [249], increase in adverse events such as falls and ulcers [251], reduced patient contact with healthcare workers [250], higher rates of anxiety and depression [249][250], and increased days spent in hospitals [249]. However, not all studies regarding the specific impact of contact precautions on these negative outcomes are conclusive [248], in part due to low quality evidence and poor study design. Potential negative outcomes and mitigation measures should be considered in developing and implementing healthcare facility policies.

It is important that both patients and healthcare workers are provided with education about the purpose and use of contact precautions. Educating healthcare workers on the importance of maintaining similar patient visitation rates for patients in isolation can assist in reducing some of the harms associated with contact precautions.

Patient-care tip

When patients are placed on transmission-based precautions due to infection or colonisation with an MRO, efforts should be made to ensure patients continue to receive adequate medical care, and to counteract potential psychological adverse effects of isolation such as anxiety and depression, and feeling of stigmatisation.

Evidence to decision

Benefits and harms

Small net benefit, or little difference between alternatives

The benefits of implementing contact precautions for patients colonised or infected with a multi-resistant organism (MRO) include a decrease in transmission of methicillin-resistant *Staphylococcus aureus* (MRSA) and vancomycin-resistant enterococci (VRE). However, the evidence on the effectiveness of contact precautions is not conclusive and few studies have directly compared the benefits of contact precautions above and beyond standard precautions.

Some of the reported harms of contact precautions for patients include [248]: reduced patient contact with healthcare workers, increased number of preventable adverse effects, decreased patient satisfaction with their quality of care and delays in access to radiological examinations.

Certainty of the evidence

Low

This recommendation has been adapted from United States Centers for Disease Control and Prevention *Management of Multidrug-Resistant Organisms in Healthcare Settings* (2006).

The quality of evidence is low, and there is significant variability amongst results regarding both the benefits and harms of contact precautions for MROs.

Values and preferences

Substantial variability is expected or uncertain

As contact precautions have been found to be associated with a range of negative patient outcomes including prolonged hospital stays and less patient visits, not all patients and staff may value the implementation of contact precautions for patients colonised or infected with MROs.

Resources and other considerations

No important issues with the recommended alternative

Whilst contact precautions may be cost effective at the individual level, across an entire healthcare facility, there may be significant resource considerations associated with implementing contact precautions. The costs associated with implementing contact precautions may include the cost of equipment and the cost of screening processes.

Rationale

Implementing transmission based precautions when a patient has been colonised or infected with a multi-resistant organism is justified to reduce healthcare associated infection. When deciding whether to implement transmission-based precautions, the benefits and harms should be considered, alongside any associated costs.

Adaptation

The GRADE process provided a consistent and transparent approach which allowed for this 2010 recommendation (developed using the FORM approach) to be reassigned a GRADE recommendation and accompanying strength. All considerations in adopting or adapting this recommendation are captured in the 'key info' tab.

Further information on the application of GRADE can be found in **Appendix 3: Process Report**.

Practice Statement

New

33. It is good practice for healthcare facilities to maintain a surveillance system to record the presence of all multi-resistant organisms.

Practical info

Surveillance of multi-resistant organism (MROs)

Surveillance of MROs is imperative in order to understand the impact, magnitude and distribution of antimicrobial resistance (AMR) in Australia. This allows for the development of policies and programs and can inform a coordinated response to critical antimicrobial resistances (CARs). Surveillance data on MROs in Australia comes mostly from public hospital laboratories. Each health facility should select an appropriate active surveillance strategy, based on their current epidemiology of MRO colonisation and local state or territory policies and legislation.

A National Alert System for Critical Antimicrobial Resistances (CARAlert) has been established by the Australian Commission on Safety and Quality in Health Care as part of the Antimicrobial Use and Resistance in Australia (AURA) Surveillance System. CARAlert collects surveillance data on nationally agreed priority organisms with critical resistance to last-line antimicrobial agents. CARAlert aims to complement existing surveillance programs in providing timely advice to state and territory health authorities on the occurrence of CARs nationally. For further information and to access reports, see <https://www.safetyandquality.gov.au/antimicrobial-use-and-resistance-in-australia/what-is-aura/national-alert-system-for-critical-antimicrobial-resistances-caralert/?section=5>.

All healthcare facilities should also have systems in place to ensure timely reporting of MROs which are classified as notifiable diseases to the relevant national and/or state/territory health authorities.

Further information on surveillance in healthcare facilities is available in Sections 4.4, its subsections, and 4.5.3.

Evidence to decision

| | |
|---|--|
| Benefits and harms | The benefits of establishing a surveillance system for multi-resistant organisms in healthcare facilities clearly outweighs the harms. |
| Certainty of the evidence | The National Safety and Quality Health Service Standards on Governance (Standard 1) and Infection Prevention and Control (Standard 3) require organisations to demonstrate governance mechanisms and risk management for infection prevention and control. |
| Values and preferences | There are no significant issues that impact on patient preferences and values, or on health equity. |
| Resources and other considerations | It is suggested that an existing register within a healthcare facility is used, so this practice should not impact on resourcing besides possible extra training for staff. |

Rationale

Effective surveillance systems should provide meaningful and accessible information so that actions can be taken to prevent and contain multi-resistant organisms.

As coordinated and integrated efforts in antimicrobial resistance are established across Australia, this information will be important for benchmarking activities and in monitoring the emergence of resistance and changes in patterns of resistance.

3.4.2 Outbreak investigation and management

What constitutes an outbreak?

When there are more cases of infection with the same organism than would normally be expected in one area or period of time, this constitutes an outbreak.

An outbreak may be defined as:

- occurrence of more cases of disease than expected in a given area among a specific group of people over a particular period of time
- two or more linked cases of the same illness.

Outbreaks commonly identified in healthcare facilities include:

- methicillin-resistant *Staphylococcus aureus* (MRSA)
- vancomycin-resistant enterococci (VRE)
- multi-resistant Gram negative (MRGN)
- respiratory pathogens (e.g. influenza, respiratory syncytial virus [RSV])
- diarrhoeal pathogens (e.g. norovirus)
- measles (rubeola), chickenpox (varicella)
- scabies
- *C. difficile*.

This section gives principles and overall guidance for managing an outbreak. For specific guidance on particular infections, please refer to national guidelines related to the management of that infection.

Outbreak investigation and management

A suspected outbreak may be identified by a healthcare worker; by laboratory personnel; or by state/territory health authorities conducting routine surveillance, investigating reports of illness and from reportable disease notifications. When an outbreak is detected, the healthcare facility's infection control management system should be notified and an outbreak control team should be formed relevant to the size and seriousness of the outbreak and the healthcare facility involved. There may also be a requirement to notify the state/territory public health unit.

The responsibility for investigation and the extent of investigations will vary according to the outbreak type and circumstances. It is important to investigate an outbreak immediately, as the availability and quality of microbiological evidence and epidemiological data diminishes rapidly with time between illness and investigation.

An outbreak management plan should be developed based on local policy and consultation between the infection control professional, healthcare workers, patients, facility management and state/territory health authorities, as appropriate. Such a plan is multifactorial and its implementation is typically overseen by a person with designated responsibility for infection control, such as an infection control professional, clinical microbiologist or infectious diseases physician.

The outbreak response may differ according to the nature of disease, the virulence of the organism and the vulnerability of the patients concerned. However, the principles that underlie an outbreak investigation are similar: identification of the aetiological agent, the route(s) of transmission, exposure factors and the population at risk.

All healthcare facilities should have systems in place to ensure timely reporting of notifiable diseases to the relevant national and/or state/territory health departments. As patients may present to a healthcare facility and be later confirmed to have a transmissible disease state/territory health authorities need to be notified to enable tracing of contacts of the infected patient in order to initiate appropriate counselling, quarantine and post-exposure prophylaxis. Healthcare facilities may need to identify staff on duty and other patients present who may have been exposed to the infectious patient and be at risk.

One of the important aspects of the outbreak management process is the written and oral communication of findings to the appropriate authorities, the appropriate health professionals and the public. This communication is based on the type and severity of the outbreak. During an outbreak it is important to provide education to the key stakeholders and clinicians about the organism and its mode of transmission.

Within a healthcare facility, effective communication could consist of:

- appropriate signage to limit access to a room or a clinical unit
- electronic alerts on the medical record to manage cases and contacts
- emails and multimedia to target all stakeholders within the healthcare facility
- provision of education and written materials to visitors to inform them of the situation and the infection control measures with which they should comply.

Table 21 outlines the process of outbreak investigation and corresponding management. In practice many steps are taken more or less simultaneously, as the results of investigations and implementation of strategies to contain and control will vary with the availability and timeliness of information and seriousness of the outbreak. In primary care there may be a limited ability to investigate an outbreak, which will be generally conducted by public health authorities once they have been notified. All outbreaks, however minor, should be investigated promptly and thoroughly and the outcomes of the investigations documented.

Note: An infection prevention and control professional should be involved and consulted throughout all stages of an outbreak investigation.

Table 21. Steps in an outbreak investigation

The steps below should be considered during healthcare facility-level investigations of outbreaks.

| Steps | Suggested approach | Responsibilities (dependent on facility and type of outbreak) |
|--|--|--|
| Step 1. Recognise outbreak and prepare to investigate | | |
| Determine existence of the outbreak | <ul style="list-style-type: none"> • Establish background rate of disease. • Consider if observed number of cases is in excess of the usual number, and cases are typical. • Examine surveillance data. | <ul style="list-style-type: none"> • Healthcare workers. • Laboratory personnel. |
| Determine if immediate control measures are needed (refer to Section 3.4.2.1) | <ul style="list-style-type: none"> • Reinforce standard precautions. • Apply appropriate transmission-based precautions. | <ul style="list-style-type: none"> • Healthcare workers—as soon as outbreak is suspected. |

| Steps | Suggested approach | Responsibilities (dependent on facility and type of outbreak) |
|---|---|--|
| <p>Notify and communicate with:</p> <p>Formation of an outbreak investigation/ management team (OMT) – this will vary according to location/ resources, made up of one or more people with designated responsibility</p> | <ul style="list-style-type: none"> Healthcare workers and ancillary staff in immediate area. Infection control professional. Executive. Laboratory. Public health unit (if notifiable disease or required pursuant to public health legislation). <p>Membership may include but is not limited to:</p> <ul style="list-style-type: none"> administrators (medical and nursing) managers of implicated areas infection control professional or designated person with infection control experience clinical Microbiologist infectious diseases physician/epidemiologist/ statistician lead investigator or 'chair' nominated others as defined by circumstances. | <ul style="list-style-type: none"> Healthcare workers—as soon as outbreak is suspected. Laboratory personnel (e.g. routine screening can identify outbreak)—as soon as outbreak is suspected. Management—as soon as notified. |
| <p>Step 2. Verify the diagnosis and confirm that an outbreak exists</p> | | |
| <p>Confirm that there are more than expected number of cases meeting the surveillance case definition of the disease of interest in the period under review</p> <p>Consider likely outbreak definition and whether criteria are met</p> | <ul style="list-style-type: none"> Confirm clinical diagnoses (symptoms and features of illness). Review laboratory data and request additional laboratory tests if necessary, e.g. molecular typing of organisms to confirm clonality. <p>Are there more cases than expected compared to previous weeks/months?</p> <ul style="list-style-type: none"> Review scientific literature. Consider epidemiology of cases—are there two or more linked cases of the same illness? | <ul style="list-style-type: none"> Laboratory personnel to report results. Clinicians to verify clinical diagnosis. OMT representatives (clinical microbiologist, senior clinicians). |
| <p>Step 3. Establish case definition and find cases</p> | | |
| <p>Establish a set of standard criteria to decide whether or not a person has the disease of concern</p> <p>Find cases</p> | <p>Case definition should be based on:</p> <ul style="list-style-type: none"> clinical information about the disease characteristics of the people who are affected information about the location specification of the time period for the outbreak. <p>Case definition can be refined later after collection of primary data.</p> <p>Cases should be classified using the case definition as:</p> <ul style="list-style-type: none"> 'Confirmed' (usually laboratory verification) 'Probable' (usually has typical clinical features) 'Suspect' (usually has fewer typical clinical features). <p>Gather critical information by:</p> <ul style="list-style-type: none"> interview follow-up of disease notification health alerts. | <ul style="list-style-type: none"> OMT representatives (clinical microbiologist, senior clinicians). Healthcare workers. OMT representatives. Healthcare facility management. |

| Steps | Suggested approach | Responsibilities (dependent on facility and type of outbreak) |
|---|---|--|
| Identify and count cases Tabulate information collected on cases investigated and update as new cases appear | Collect the following types of information: <ul style="list-style-type: none"> identifying information demographic information clinical information risk factor information (including environmental tests). • Time—date of onset of illness. • Person—age, sex. • Place—where did the exposure occur? • Other relevant information. | <ul style="list-style-type: none"> OMT representative. OMT representative. |
| Step 4. Characterise outbreak by person, place and time | | |
| Review descriptive epidemiology of all cases | Descriptive epidemiological data includes: <ul style="list-style-type: none"> person: sex, age, occupation, residence place: information that provides detail about possible source of agent and nature of exposure time: date and time of onset, record relevant events in a timeline. | <ul style="list-style-type: none"> OMT representative. |
| Step 5. Determine who is at risk | | |
| Identify groups at risk Initiate precautionary measures | <ul style="list-style-type: none"> Number of people who have developed the disease/condition of interest (numerator). At risk population: number of people likely to be exposed e.g. total number of patients on ward, staff and visitor contact (denominator). Time and place of onset. Personal characteristics. <ul style="list-style-type: none"> Use of standard precautions and appropriate transmission-based precautions. Increase frequency and efficiency of environmental cleaning. Prophylactic treatment/immunisation. Antibiotic restrictions. Exclusion of cases from high risk activities. Isolation and/or cohorting of patients. Restricting movement of patients, staff and visitors. Screening of patients with isolation of patients and cohorting of contacts. Provision of health information and advice. | <ul style="list-style-type: none"> OMT representative. Healthcare workers. Infection control professional. |
| Step 6. Implement ongoing control/prevention measures (this can be done at any time during the outbreak as deemed necessary) | | |
| Review measures initiated for immediate control (Step 1 and Step 5) Implement appropriate ongoing control measures and strategies to prevent further illness (see Section 3.4.2.1) | <ul style="list-style-type: none"> Are infection control measures adequate to reduce risk of transmission? Restrict spread from the case. Interrupt chain of infection. Interrupt transmission or reduce exposure. Reduce susceptibility to infection. Assessment of policy, regulations, standards. | <ul style="list-style-type: none"> Healthcare workers. OMT. Healthcare facility management. Healthcare workers. OMT. Healthcare facility management. |

| Steps | Suggested approach | Responsibilities (dependent on facility and type of outbreak) |
|---|---|--|
| Communicate and coordinate with all stakeholders | <ul style="list-style-type: none"> Electronic flagging of medical records of contacts. Reinforcement of infection control precautions to staff, patients and visitors. | <ul style="list-style-type: none"> Healthcare workers. OMT. Infection control professional. |
| Make plans to evaluate their effectiveness | <ul style="list-style-type: none"> Document type and time of implementation of infection control measures. Monitor factors contributing or affected by outbreak and any associated changes. | <ul style="list-style-type: none"> Healthcare workers. OMT. Infection control professional. |
| Step 7. Communicate findings | | |
| Prepare written report that evaluates methods used for the control of the outbreak | <ul style="list-style-type: none"> Include discussion of factors leading to outbreak, comprehensive timelines, summary of investigation and documented actions. Short and long term recommendations for prevention of similar outbreak. Disseminate to appropriate stakeholders including publication. | <ul style="list-style-type: none"> OMT. Healthcare facility management. |

Table 22. Developing and testing an hypothesis for an outbreak

In addition to Table 21 above, the steps outlined below should be considered during public health level investigations of outbreaks.

| Step 8. Develop hypothesis—the 'how' and 'why' | | |
|---|--|---|
| Develop hypotheses from the factual information gathered to date on potential source, vector, pathogen and route of transmission | <ul style="list-style-type: none"> Data collected by interview. Common links. Plausible exposure. Environmental test results where appropriate. Review literature. | <ul style="list-style-type: none"> OMT representative. |
| Step 9. Test hypothesis with established facts | | |
| Perform epidemiologic study | <ul style="list-style-type: none"> Cohort. Case-control. | <ul style="list-style-type: none"> OMT representative. |
| Analyse the data | <ul style="list-style-type: none"> Compare the risk factors among ill (cases) vs. not ill (controls). Attack rates. Relative risk. | <ul style="list-style-type: none"> OMT representative or outsourced to consultant with knowledge of statistical methods. |
| Carry out further studies if necessary- to support the hypothesis or if analytic studies do not confirm the hypothesis | <ul style="list-style-type: none"> Further study to refine case definition. May involve testing of environment samples, food samples or environmental screening in some situations (e.g. Legionella, Pseudomonas). | <ul style="list-style-type: none"> OMT. |

Individual actions for reducing the risk:

- Become familiar with local policy on the implementation of transmission-based precautions in the event of an outbreak.
- If an outbreak is suspected or identified, implement core strategies for prevention and control and seek advice from an infection control professional or person with designated responsibility for this task regarding intensified strategies appropriate to the specific organism.

34. It is good practice for all outbreaks, however minor, to be investigated promptly and thoroughly and the outcomes of the investigations documented.

Practical info

All healthcare facilities should have an outbreak management plan based on local policy and consultation between the infection control professional, healthcare workers, patients, facility management and state/territory health authorities, as appropriate.

The responsibility and extend of outbreak investigations will vary according to the outbreak type and circumstances. It is important to investigate an outbreak immediately, as the availability and quality of microbiological evidence and epidemiological data diminishes rapidly with time between illness and investigation.

Evidence to decision

| | |
|---|--|
| Benefits and harms | The benefits of promptly and thoroughly managing outbreaks of mulit-resistant organism (MROs) in healthcare facilities clearly outweighs the harms. |
| Certainty of the evidence | The National Safety and Quality Health Service Standards on Governance (Standard 1) and Infection Prevention and Control (Standard 3) require organisations to demonstrate governance mechanisms and risk management for infection prevention and control. |
| Values and preferences | This measure should not impact patient preferences and values nor have any health equity implications. |
| Resources and other considerations | Promptly and thoroughly managing outbreaks of MROs in healthcare facilities would result in economic benefits due to a shorter duration of outbreak and reduction in bed-days lost. |

Rationale

Successful outbreak management is based on the prompt exchange of information to enable the successful management of the outbreak. Management will be based on a combination of standard and transmission-based precautions. Specific interventions will be determined by the infection control professional, based on the mode of transmission of the infectious agent.

3.4.2.1 Infection control strategies to contain an outbreak

Infection control strategies to control/contain an outbreak

These Guidelines provide core principles of infection prevention and control. However, it is the responsibility of healthcare facilities to develop local policies and procedures relevant to their setting, in addition to the Australian Commission on Safety and Quality in Health Care recommendations for certain multi-resistant organism (MROs).

Good governance and administrative or managerial support are crucial to support outbreak management (see Section 4.1). The healthcare worker’s role in outbreak management will include:

- reinforcement of standard precautions, including adherence to the 5 moments for hand hygiene, environmental cleaning protocols and appropriate use of personal protective equipment (PPE)
- implementation of relevant transmission-based precautions, including isolation and cohorting.

The specific precautions required for each infectious agent are listed in Appendix 2—Section 6.4.

Hand hygiene

During an outbreak, adherence to the five moments for hand hygiene can assist in preventing further cases and reducing environmental contamination.

Environmental Cleaning

Frequency and efficiency of environmental cleaning should be increased above the standard for the area to ensure any

contaminants are removed (see Appendix 2—Section 6.1 for guidance on cleaning in high-risk situations). A targeted cleaning regime may be introduced and continued for the duration of the outbreak dependent on the mode of transmission of the infectious agent.

Consideration should be given to whether the surrounding environment will need to be disinfected in addition to being cleaned. Further information is available in Section 3.1.3.

Patient Isolation

The isolation of colonised or infected patients is important when managing an outbreak. Standardised transmission-based precautions signage should identify the isolation room and include the necessary precautions to be adopted. Patients should be isolated to a negative pressure room with bathroom facilities or in a room from which air does not circulate to other areas if available. The door should be kept closed for patients on airborne precautions.

If isolation is not possible, cohorting of patients should occur as advised by an infection control professional.

Single room

Single-patient rooms are always indicated for patients placed on airborne precautions and are preferred for patients who require contact or droplet precautions. In the event of an outbreak, single-patient rooms are preferred for all modes of transmission.

During an outbreak, single patient rooms should be prioritised for patients who have conditions that facilitate transmission of infectious material to other patients (e.g. draining wounds, stool incontinence, uncontained secretions) and for those who are at increased risk of acquisition and adverse outcomes resulting from infection (e.g. immunosuppression, open wounds, indwelling catheters, anticipated prolonged length of stay, total dependence on healthcare workers for activities of daily living). The prioritisation of single rooms may also need to take into account other factors that might warrant the need for single rooms, including patients requiring end of life care or special security.

Cohorting

Cohorting patients who are colonised or infected with the same strain confines their care to one area and prevents contact with other patients. Cohorts are created based on clinical diagnosis, microbiologic confirmation when available, epidemiology, and mode of transmission of the infectious agent. It is generally preferred not to place severely immunosuppressed patients in patient-care areas with other patients. Carbapenemase-producing *Enterobacterales* (CPE) positive patients should also not be grouped together without expert direction from senior infection control experts.

Cohorting allows more efficient use of staff. Cohorting has been used for managing outbreaks of MROs and pandemic influenza, and modelling studies provide additional support for cohorting patients to control outbreaks.

Placement of large numbers of patients

In the event of an outbreak or exposure involving large numbers of patients who require transmission precautions, an infection control professional should be consulted before patient placement.

Appropriate measures may include:

- cohorting of patients in areas of the facility that are away from other patients
- if airborne, using temporary portable solutions (e.g. exhaust fan) to create a negative pressure environment in the converted area of the facility.

Restricting movement within the facility

Restricting movement of patients during an outbreak reduces the risk of further transmission.

If transfer within the facility or transport to another facility is necessary, advice should be sought from an infection control professional. If an infected or colonised patient must be moved the transport service and receiving area or facility should be notified of the nature of the patient's infection or colonisation.

It is important to:

- ensure that infected or colonised areas of the patient's body are covered if relevant
- if the target infection is transmitted by the droplet or airborne route, ask the patient to wear a mask while being moved.

Contaminated PPE should be removed and disposed of and hand hygiene performed before the patient is moved. Clean PPE should be put on before the patient is handled at the destination.

For specific control measures for CPE positive patient movement, see [Australian Commission on Safety and Quality in Health Care, Recommendations for the control of CPE: A guide for acute care health facilities \(2017\) \[231\]](#).

Exclusion Policies

Exclusion policies may also be implemented to restrict the spread of disease throughout a healthcare facility. This could include:

- excluding patients from participating in specific activities
- restricting or cancelling visiting hours for patients in outbreak areas
- excluding staff from work until well if they are implicated in the transmission of infection (e.g. food handlers or clinical staff)
- managing vaccine refusal, contraindication to vaccination and vaccine non-response by ensuring appropriate work placements, work adjustments, work restrictions and exclusions.

In an outbreak of viral gastroenteritis, healthcare workers should not return to work until diarrhoea and vomiting have ceased for 2 days. It is extremely important that healthcare workers comply with appropriate hand hygiene methods and infection control practices upon return to work, given that some studies have shown prolonged viral shedding for up to 21 days.

Information about exclusion periods for healthcare workers with acute infections is in Section 4.2.2.

Patient-care tip

Patients, their families and visitors may experience concern or fear or feel that they are not being given enough information in an outbreak situation.

Clearly explaining the process of outbreak management and the importance of infection control measures may assist them in understanding the situation and improve compliance with infection control directives.

Consider referring the healthcare facilities Infection Control Practitioner to the patient and/or family for more in-depth discussions and information.

Applying transmission-based precautions during an outbreak

Successful outbreak management is based on a combination of standard and transmission-based precautions. Specific interventions will be determined by the infection control professional, based on the mode of transmission of the infectious agent. These include:

- appropriate hand hygiene practices (see *Recommendation 1*)
- use of appropriate PPE (including gloves, apron or gowns, and surgical mask or P2 respirator) (see *Recommendations 29-31*)
- implementing patient-dedicated or single-use non-critical equipment (e.g. blood pressure cuff, stethoscope) and instruments and devices (see *Recommendation 22*)
- following standard procedures for containment, cleaning and decontamination of spills (see *Recommendations 9-12*)
- increasing the frequency of environmental cleaning over the standard for that area, using appropriate products (see Appendix 2, Sections 6.1 & 6.7).

Practice Statement

New

35. It is good practice to consider the use of early bay closures to control known or suspected norovirus outbreaks rather than ward/unit closures.

Rather than closing an entire ward or unit to manage an outbreak of norovirus in a healthcare facility, it may be more efficient to control an outbreak through cohorting symptomatic patients in bays. If taken, this approach needs to be implemented promptly and early (within three days of the first case becoming ill) in combination with adequate infection control strategies.

Practical info

In the past, ward or unit closure was considered as the central control measure for managing outbreaks of norovirus in healthcare facilities. However, a recent literature review [150] undertaken for the development of the 2019 Guidelines found that efficient control may be achieved by the closure of bays. If taken, this approach needs to be implemented promptly and

early (within three days of the first case becoming ill) in an outbreak before extensive transmission has occurred within a clinical area.

Norovirus epidemiology

Norovirus is the most frequently occurring cause of community-acquired acute gastroenteritis in people of all ages. It is one of the most common causes of outbreaks in healthcare settings, affecting both long-term care facilities and acute care hospitals.

Norovirus belongs to the family *Caliciviridae* and is a single-stranded RNA, non-enveloped virus that can cause gastroenteritis in humans [118]. Noroviruses are divided into at least six genogroups (GI-GVI) and further subdivided into more than 38 genotypes based on phylogenetic analysis of the major capsid protein [118] [122]. Currently, human noroviruses belong to one of three norovirus genogroups which are further divided into more than 25 genetic clusters [119].

Human noroviruses cannot be grown in cell culture [123], therefore, diagnostic methods focus on detecting viral RNA or antigen. There were 17 studies (81%) which identified that NoV genotype GII.4 have caused the majority of clinical outbreaks in healthcare settings during the past decade.

Norovirus infections generally have a shorter incubation period and are characterised by acute onset of nausea, vomiting, abdominal pain and diarrhoea [63]. The mean duration of symptoms is 2-3 days.

Transmission pathways

Transmission for norovirus infections in healthcare settings mainly occur by the faecal–oral route, either through person to person contact or through exposure to contaminated food [96].

Whilst some observational studies have suggested there is a possibility of viral transmission via aerosols, there is no data or determination criteria to support this assumption.

It appears that Genotype GII.4 is more often associated with transmission mediated by person-to-person contact than with other types of transmission.

Individuals may shed norovirus for more than 21 days after the resolution of symptoms, possibly acting as a possible source for nosocomial transmission. However no data has been reported on ongoing transmission or secondary cases.

Evidence to decision

| | |
|---|---|
| Benefits and harms | Bay closures have been found to be effective for controlling norovirus outbreaks [120] [121], particularly when this is combined with other infection control measures. |
| Certainty of the evidence | Three studies were found which examined the effectiveness of ward or bay closures [150]. None of the studies were randomised, but their overall methodological quality was rated as high. |
| Values and preferences | This measure should not impact patient preferences and values nor have any health equity implications. |
| Resources and other considerations | Containing patients in bays can reduce the amount of environmental cleaning required. Bay closures can result in a more rapid turnaround of closed areas and removal of restrictions on admissions and discharges. This would result in economic benefits due to a shorter duration of outbreak and reduction in bed-days lost. |

Rationale

Evidence from three studies has shown that bay closures may be a more efficient way to control a norovirus outbreak than ward closures. This measure has resource implications, as it can result in a reduction in the number of bed-days lost during an

outbreak, and a reduction in outbreak frequency, which can in turn result in cost savings.

3.5 Applying standard and transmission-based precautions during procedures

Summary

Section 3.5 outlines processes for risk identification and the application of standard and transmission-based precautions for certain procedures. It is not intended to provide guidance on performing procedures, but outlines the principles involved in the delivery of care that reduces the risk of transmission of infection during the insertion and maintenance of invasive medical devices and for surgery.

Medical and dental procedures increase the risk of transmission of infectious agents between patients and healthcare workers.

- 'Procedure' includes any situation in which there is a potential for contact between the skin of the healthcare worker and the patient's tissues, body cavities or organs, either directly or via surgical instruments or invasive medical devices.
- The more invasive the procedure, the greater the risk of transmission of infection. Before a procedure is undertaken, consideration should be given to whether there is a safer, less invasive alternative.
- The level of perceived infection risk depends on a range of factors including the site and complexity of the procedure and patient characteristics (e.g. age, underlying illness).
- Healthcare workers should be trained and competent in safe procedural techniques and participate in regular education sessions about minimising the infection risk of procedures. If there is any uncertainty, healthcare workers should contact the person with designated responsibility for infection control.

Patient-care tip

Patients and their carers should be offered clear, consistent information and advice throughout all stages of their care. This should include the risks of procedure-related infections, what is being done to reduce them and how they are managed.

3.5.1 Taking a risk-management approach to procedures

All procedures involve some risk of infection. Minimising the infection risk associated with a procedure should be an integral part of considering the overall risks and benefits of that procedure to the patient. The aim should be to perform the procedure with the lowest level of perceived infection risk that will meet the treatment goals for that patient. When performing the procedure, associated infection risks should be identified and minimised.

In developing local policies for a healthcare facility, it is useful to refer to guidelines developed to inform practice in performing specialised procedures.

Risk assessment for pulmonary function tests

Ideally pulmonary function tests including spirometry should be performed in a room that is closed off to other internal spaces and involve only one patient at a time. In situations where respiratory infection is suspected or confirmed, the use of spirometry should be delayed if clinically appropriate. If the test cannot be delayed, healthcare workers should implement transmission-based precautions and conduct a risk assessment using the hierarchy of controls prior to performing the procedure.

Classifying procedures

Procedures can be classified according to the level of perceived risk, by applying the principles of Spaulding's criteria for assessing the risk of medical instruments and equipment according to their intended use (see Section 3.1.4).

Table 23. Level of risk to patients from different types of procedures

| Level of risk | Criteria | Example |
|----------------------------------|---|-------------------------------------|
| High risk (critical site) | Any surgical entry into tissue, body cavities or organs, or repair of traumatic injury. | Abdominal surgery Dental surgery |

| | | |
|--|--|---|
| <p>Medium risk (semi-critical site)</p> | <p>Contact with mucous membranes or non-intact skin.</p> | <p>Respiratory procedure</p> <p>Internal/instrument examination (e.g. ultrasound, endoscopy)</p> <p>Minor skin surgery</p> <p>Minor dental procedures</p> |
| <p>Low risk (non-critical site)</p> | <p>Contact with intact skin.</p> | <p>Non-invasive examinations or procedures (e.g. abdominal ultrasound)</p> <p>Blood pressure measurement, electrocardiogram (ECG), injection through intact skin</p> <p>Extra-oral dental examination</p> |

Appropriate use of devices

Appropriate use of devices is integral to reducing the risk of procedures. Single-use or single-patient items should be used wherever practicable, and items designed for single use must not be used for multiple patients. Healthcare workers should be aware of situations where cross-contamination may occur during routine procedures. See Section 3.1.4 for further information.

Aseptic technique

Aseptic technique protects patients during invasive clinical procedures by employing infection control measures that minimise, as far as practicably possible, the presence of pathogenic microorganisms. While the principles of aseptic technique remain constant for all clinical procedures, the level of practice will change depending upon a standard aseptic technique risk assessment. See Section 3.1.6 for further information.

The care bundle approach

The Institute for Healthcare Improvement (IHI) in the US developed a structured ‘care bundle’ approach to help healthcare workers consistently deliver the safest possible care for patients undergoing treatments with inherent risks. A bundle is a set of evidence-based practices (generally three to five) that, when performed collectively and reliably, improve patient outcomes.

Many bundle elements are well-established practices, combined in a structured protocol that is agreed upon and is the responsibility of the whole clinical team. Bundle characteristics include the following:

- The elements are all necessary and sufficient and make up a cohesive unit of steps that must all be completed to succeed.
- The elements are all based on randomised controlled trial evidence.
- The elements involve all-or-nothing measurement, making implementation clear-cut.
- Bundle elements occur at a specific time and in a specific place (e.g. during morning rounds every day).

Examples of care bundles are given in Section 3.5.2.2 (intravascular device [IVD] care bundle) and Section 3.5.2.3 (ventilator-associated pneumonia [VAP] care bundle). These can be used to monitor, assess and improve performance as well as to increase consistency of care.

Existing care bundles can be used as a tool and be developed by each facility to meet its needs. For more information, refer to the IHI website at www.ihl.org.

3.5.2 Invasive medical devices

| |
|---|
| <p>Summary</p> |
| <ul style="list-style-type: none"> • Invasive medical devices include: <ul style="list-style-type: none"> ◦ catheters inserted for drainage (e.g. urinary catheters) ◦ catheters for intravascular access (e.g. peripheral intravenous catheter, peripherally inserted central venous catheter, central venous catheter) ◦ devices for mechanical ventilation (e.g. intubation) |

- **devices for feeding** (e.g. enteral feeding tube).
- The following sections provide best-practice guidance on strategies for the selection, insertion, maintenance and removal of invasive medical devices.

Practice Statement

New

36. It is good practice for healthcare facilities to develop, implement and review processes to address the insertion, use and maintenance, and removal of invasive medical devices. These processes should be centred on the principles of only using devices if they are deemed essential, and removing them as soon as no longer needed.

Healthcare facilities should undertake a risk assessment to assist with determining appropriate procedures and timing for the removal of invasive medical devices and for the surveillance and management of invasive medical devices.

Practical info

Invasive medical devices are a common source of healthcare associated infections (HAIs) and provide a route for infectious agents to enter the body. Pneumonia, urinary tract infections and bloodstream infection account for around 70% of intensive care unit HAIs, and most of these are associated with invasive devices [272]. The need for appropriate processes and policies in all healthcare facilities that addresses the proper insertion, use, management and removal of invasive medical devices is, therefore, paramount [274].

Aseptic insertion and careful maintenance of devices is critical to reducing infection risk. Information on use of aseptic technique for specific procedures (including invasive medical devices) can be found in Section 3.1.6 and Appendix 1—Section 5.11.

Key concepts in minimising the risk of infection related to the use of invasive medical devices:

- Only use an invasive medical device when clinically indicated and consider the infection-risk during decision making.
- Ensure all staff are adequately trained and competent in the skills required for safe insertion, maintenance and removal of a device.
- Choose the most appropriate device and system for the patient.
- Check the device at every shift and remove as soon as no longer necessary.
- Regularly monitor patients, the insertion site and the device for any signs and symptoms of infection.
- Minimise the period of time a device remains in a patient.
- Provide patient education on the infection risk associated with the insertion of devices and the importance of proper maintenance.
- Clearly document the insertion, maintenance and removal of the device, as well as daily review of device necessity.
- Implement appropriate surveillance systems to monitor infection rates.

Evidence to decision

| | |
|---|---|
| Benefits and harms | The benefits of developing and implementing process for the use of invasive medical devices outweighs any undesirable effects. |
| Certainty of the evidence | Systematic reviews (n=3) have been carried out which investigate the role of quality improvement programs on the risk of infection, and have found that implementing processes and protocols around the insertion and removal of invasive medical devices can be effective [274]. However, many studies in this area are uncontrolled before-after designs and are at risk of bias. |
| Values and preferences | It is expected that all patients and staff of Australian healthcare facilities would highly value the implementation of processes surrounding the use of invasive medical devices in order to reduce infection. |
| Resources and other considerations | There may be costs associated with the implementation of processes and protocols around the use of invasive medical devices, including education and clinical leadership activities. |

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. The development and implementation of processes for using invasive medical devices is justified to reduce healthcare associated infection.

3.5.2.1 Indwelling urinary devices

Indwelling urinary devices

An indwelling urinary catheter is a flexible, tubular instrument passed into the bladder either through the urethra or through the abdominal wall above the symphysis pubis and is used to empty the contents of the bladder. Indwelling urinary catheters are used for a number of reasons, including [287] [284]:

- management of urinary retention or obstruction
- clot retention associated with gross haematuria
- monitoring for sepsis, trauma, renal function, electrolyte or fluid balance
- injury or surgery affecting urinary function and/or involving immobility (including injury, surgery or disease affecting the spinal cord)
- urinary incontinence management associated with wound care, end-of-life care or chemotherapy, if other options available adversely impact patient comfort
- urogenital or bladder management (e.g. management of fistula or haematuria)
- labour and birth management.

What are the risks?

Bacterial infections associated with urinary catheterisation gain access to the urinary tract either through:

- *Extraluminal contamination*—this can occur if there is a break in aseptic technique during insertion of the catheter or servicing the drainage system, from the healthcare worker's hands or from the patient's own colonic or perineal flora.
- *Intraluminal contamination*—this can occur through reflux of bacteria from a contaminated urine drainage bag.

Around 20% of healthcare associated infections are urinary tract infections, with approximately 1.7% of all hospital patients acquiring a urinary tract infection during their stay [280]. A large proportion of urinary tract infections are associated with catheterisation, with up to 97% of urinary tract infections in intensive care units associated with an indwelling catheter. As approximately 25% of patients in hospital receive short-term indwelling urinary catheters [82], it is paramount that best-practice infection prevention processes are followed.

Healthcare associated urinary tract infections are associated with a range of negative outcomes including an increased length of stay in hospitals [280]. The risk of infection is related with the method and duration of catheterisation, the quality of catheter care and host susceptibility. The longer a urinary catheter is in place, the greater the risk of infection.

Minimising the risk from indwelling urinary devices

- **Assessing the need for catheterisation:** Limiting catheter use and minimising duration are primary strategies in reducing the risk of catheter-associated urinary tract infections (CAUTI). Facilities should clearly outline the indications for catheter insertion and the need for insertion of an indwelling urinary device should be reviewed before the procedure is performed.
- **Education of healthcare workers:** Healthcare workers performing catheterisation should be trained and competent in the technique and familiar with policies and procedures for insertion, maintenance and changing regimes of indwelling urinary devices.
- **Educating patients:** It is important to provide patients with information in relation to the need for catheterisation and details about the insertion, maintenance and removal of their catheter.
- **Implementing appropriate surveillance:** Surveillance relating to indwelling catheters is suggested, and can include monitoring for compliance with indications for insertion, and documentation of processes.

Table 24. Process for urethral catheter insertion and maintenance

| Stage | Process |
|-----------|---|
| Insertion | <ul style="list-style-type: none"> • Insert only if clinically indicated. • Ensure documented facility policy on urethral catheter insertion is being followed and that staff members performing the procedure are trained in the specific technique. • Select appropriate catheter and catheter size [286]. |

| Stage | Process |
|--------------------|--|
| | <ul style="list-style-type: none"> • Use sterile equipment (including a sterile drape) and aseptic technique when inserting urinary catheters and connecting to the sterile system. • Clean the urethral meatus before before insertion of the catheter. Consider the emerging evidence available for suitable cleaning solutions [289] [282] [290]. • Use an appropriate sterile, single-use lubricant when inserting the catheter. Male patients may require the application of anaesthetic gel prior to the insertion of the catheter [285]. • After insertion, properly secure the catheter to the drainage device and secure the catheter and the drainage device to the patient [284]. • Document insertion of the device in the patient medical record (detailing device, date, time, product and clinical indication). |
| Maintenance | <ul style="list-style-type: none"> • The need for catheterisation should be assessed at least once daily. • Use an aseptic closed system and avoid breaches to this system (e.g. unnecessary emptying of the urinary drainage bag). • Before manipulation, perform hand hygiene and put on non-sterile gloves. • Position drainage bag to prevent back-flow of urine or contact of bag with the floor. • Regularly check for kinks in tubing and ensure that there is continuous drainage. • Ensure there is a secure connection between the catheter and the drainage device. • Do not add antiseptic or antimicrobial solutions into drainage bags, as studies have shown no reduction in the incidence of bacteraemia when adding hydrogen peroxide or chlorhexidine into drainage bags. • Empty the drainage bag frequently enough to maintain urine flow and prevent reflux. • Use a separate urine collection container for each patient, avoiding contact between the drainage bag and container. Following use, the container should be discarded if single use, or cleaned and sterilised if reusable. • Change drainage bags only when necessary (i.e. according to either manufacturers' recommendations or the patient's clinical needs). • Clamping is unnecessary. • Daily meatal and periurethral hygiene can be maintained through routine bathing or showering. No reduction in bacteraemia has been demonstrated when aseptic/antimicrobial agents are used for meatal care compared with routine bathing or showering [282]. • Document all procedures involving the catheter or drainage system. • Evidence indicates that bladder irrigation, instillation and washout may have local toxic effects and contribute to the development of resistant microorganisms. However, continuous or intermittent bladder irrigation may be indicated during urological surgery or to manage catheter obstruction. |
| Removal | <ul style="list-style-type: none"> • Remove as soon as the need for catheterisation is no longer required. • Systems should be used to prompt early removal of the urinary catheter, as evidence suggests that reminders and stop orders can reduce CAUTI [281]. • Document all information regarding the catheter removal. |

Sources: Tenke et al (2008) [279]; Loveday et al (2014) [274]; NICE (2012) [276]

Patient-care tip

Patients should be provided with information regarding the reason for the catheter and the plan for review and removal.

Given the risk of urinary tract infection associated with urinary catheterisation, it is important that patients and relatives understand about infection prevention, are aware of the signs and symptoms of urinary tract infection and know how to access expert help if difficulties arise.

CAUTI maintenance bundle

An example of a bundle procedure for maintenance of urinary catheters is to:

- perform a daily review of the need for the urinary catheter
- check the catheter has been continuously connected to the drainage system
- ensure patients are aware of their role in preventing urinary tract infection, or if the patient is unable to be made aware, perform routine daily meatal hygiene
- empty urinary drainage bags frequently enough to maintain urine flow and prevent reflux. Use a separate urine collection container for each patient, avoiding contact between the drainage bag and container

- perform hand hygiene and put on gloves and apron before each catheter care procedure; on procedure completion, remove gloves and apron and perform hand hygiene again. These practices can be measured and used to monitor performance by the clinical team.

3.5.2.2 Intravascular access devices

Intravascular access devices

Indwelling intravascular access devices (catheters) provide a route for:

- administering fluids, blood products, nutrients and intravenous medications
- monitoring haemodynamic function
- maintaining emergency vascular access
- obtaining blood specimens.

The main types of intravascular access devices are:

- **Peripheral intravenous catheters (PIVCs)**—which are inserted into peripheral veins (e.g. small veins in the arms) and are the most commonly used intravascular access device in hospitalised patients. They are short term devices.
- **Peripheral inserted central venous catheters (PICCs)**—which are also inserted through a peripheral vein site and can be used for a prolonged period of time (e.g. for long chemotherapy regimes, extended antibiotic therapy or total parenteral nutrition).
- **Central venous catheters (CVCs)**—which are inserted into larger veins within the chest and abdomen and generally remain in place for long periods of time. They are also sometimes called a central venous access device (CVAD) or central venous line.
- **Other vascular access devices**—examples include arterial lines, mid lines, and totally implantable central venous access ports.

What are the risks?

Intravascular access devices provide potential routes for infectious agents to cause local infection or to enter the bloodstream. As a result, despite their important role in diagnostic and therapeutic care, intravascular access devices are a potential source of healthcare associated infections, the most severe form being bloodstream infections (BSIs) associated with the insertion and maintenance of these devices. Intravascular access device related BSIs are associated with significant mortality, worsening of the severity of the patient's underlying ill health, prolonging the period of hospitalisation and increasing the cost of care.

There is risk of infection when the device is inserted and while it remains in situ. The risks inherent in insertion of intravascular access devices include bypassing the skin, which is an important barrier against microorganisms gaining entry to sterile sites such as the bloodstream, and leaving a foreign body in the patient for several days or longer which is likely to become colonised by microorganisms.

Risk factors for intravascular access device related BSI [297]:

- Prolonged hospitalisation before the intravascular access device is inserted.
- Prolonged placement of the intravascular access device.
- Heavy microbial colonisation of the insertion site that contaminate the catheter during insertion and migrate along the cutaneous catheter track and risk of contamination from healthcare workers' hands or equipment during insertion.
- Heavy microbial colonisation of the cannula/catheter hub, usually secondary to contamination from healthcare workers' hands during care interventions such as preparing and administering injections.
- Contamination of fluids, medicines and/or ultrasound gel.

The microorganisms that colonise catheter hubs and the skin adjacent to the insertion site are the source of most intravascular access device related BSIs. Coagulase-negative staphylococci, particularly *Staphylococcus epidermidis*, are the most frequently implicated microorganisms. Other microorganisms commonly involved include *Staphylococcus aureus*, *Candida* species and enterococci.

Minimising the risk from intravascular access devices

Table 25. Minimising the infection risk to patients from intravascular access devices by device type

| | Peripheral Intravenous Catheter (PIVC) | Peripheral Inserted Central Catheter (PICC) | Central Venous Catheter (CVC) |
|---------------------------------|--|---|--|
| Need for catheterisation | All types of intravascular access devices should be used only when clinically indicated and deemed necessary, and when all other alternatives have been considered (such as oral medication). | | |
| | Select the most appropriate device and site for the patient after assessing the need for the device and duration of therapy [314]. | | |
| | | The risk factors associated with inserting central lines should be considered prior to insertion, and all risks should be minimised, where possible. | |
| Skin preparation | Healthcare workers should allow sufficient contact time for site preparation, ensuring the following: <ul style="list-style-type: none"> • remove hair, if necessary, using clippers (not shavers) [316] [317] • clean a site large enough for insertion before applying antiseptics and allowing to dry completely • decontaminate the site using a single-use application of alcohol-based chlorhexidine gluconate solution (2% chlorhexidine gluconate in 70% isopropyl alcohol) [314] [326] • if insertion through or close to mucous membranes is necessary, use aqueous solution supplemented with 2% chlorhexidine • for patients with a history of chlorhexidine sensitivity, use 5% alcohol-based povidone-iodine solution or 10% aqueous povidone-iodine if insertion is close to or through mucous membranes (see <i>Recommendation 39</i> for further information). | | |
| Device selection | Choose the shortest and smallest gauge suitable for the prescribed therapy as this can reduce the risk of phlebitis. This must be well secured to prevent dislodgement. | Use a central catheter with the least number of lumens, connectors and ports possible [311]. Consider the length of time that the catheter is likely to be in situ. If total parenteral nutrition is being administered, a single lumen should be reserved for that use. There is evidence to suggest that antimicrobial coated or impregnated catheters can reduce the risk of bloodstream infection. However, the magnitude of the benefits differs according to the healthcare setting, and significant benefits have primarily been reported in intensive care units [34] [291] [313]. | |
| Site selection | In selecting the best insertion site, consider: <ul style="list-style-type: none"> • using the patient's non-dominant forearm, where possible • using the basilic or brachial veins on the posterior (dorsal) forearm, where possible • that the metacarpal veins on the dorsum of the hand are easiest to visualise but more liable to clot and are prone to vessel damage. Avoid, where possible: <ul style="list-style-type: none"> • using areas of flexion (i.e. wrist and antecubital fossa), as this may | In selecting the best insertion site, consider using the non-dominant arm where possible. | Use a subclavian site, rather than a jugular or a femoral site, in adult patients for CVC placement. |

| | | | |
|---------------------------------------|---|--|---|
| | <p>predispose to phlebitis</p> <ul style="list-style-type: none"> • using areas below previous cannulation, bruised or phlebitic areas • using an infected limb, or a limb with a PICC or implanted venous access device • using the arm on the side of the body where lymph node clearance/fistulas may be located • using lower limbs due to the risk of deep vein thrombosis • using the anterior (ventral) forearm veins, especially the cephalic vein, in patients with chronic renal failure. | | |
| <p>Insertion</p> | <p>All healthcare workers who insert intravascular access devices should be appropriately trained or under the supervision of a trained clinician. Required competencies and accreditation are determined according to healthcare facility policy.</p> <p>Perform hand hygiene immediately prior to the insertion of all catheters, either by washing with antimicrobial hand wash solution and water or using an alcohol-based hand rub.</p> <p>Multiple insertion attempts increase risk of infection, to avoid this each healthcare facility should have a document inserter escalation pathway, and a process for identifying difficult vascular access patients early so that they are referred to the appropriately skilled inserter.</p> | | |
| | <p>Use aseptic technique for the insertion of PIVCs (see Appendix 1, Section 5.11).</p> | <p>Use maximum sterile barrier precautions for insertion of central venous catheters; there is evidence that they can reduce immediate post-insertion skin colonisation.</p> | |
| | | <p>When PICC insertion is done at the bedside (i.e. the patients' room), establish a suitable aseptic field and maintain this throughout the procedure.</p> | <p>Using a two-dimensional ultrasound can offer benefits in safety and quality when compared with an anatomical landmark technique [304]. If using ultrasound guidance, healthcare workers should be appropriately trained in this technique and the principles of asepsis applied throughout the procedure including the use of sterile ultrasound gel. See ultrasound care guidelines for further information [136].</p> <p>Consideration may be given to the use of CVC insertion bundles.</p> |
| <p>Dressing and securement</p> | <p>Use either sterile gauze or sterile, transparent, semi-permeable dressing to cover the catheter site. There is insufficient evidence to suggest the use of one dressing type over the other. Patient preference and</p> | | |

| | | |
|---|--|---------------------------------------|
| | <p>clinician preference are currently acceptable factors to consider when choosing a dressing type.</p> <p>If a patient is diaphoretic or if the site is bleeding or oozing, use gauze dressing until this is resolved.</p> <p>Inspect dressing (device and site) at each shift. If there is any moisture or leaking, or the dressing becomes damp, loose, soiled or lifting then it should be replaced.</p> <p>Replacement of dressings:</p> <ul style="list-style-type: none"> • Gauze dressings should be replaced at least every 24 hours. • CVAD/PICC transparent dressings should be replaced every 7 days. • PIVCs are a short term device and change may not be required until the device is removed. | |
| Maintenance | <p>Use hand antisepsis and aseptic technique for catheter site care and for accessing the system.</p> <p>The safe maintenance of an intravascular access device includes: good practice in caring for the patient's catheter hub and connection port to avoid contamination by staff hands, the use of an appropriate site dressing regime, and using flush solutions to maintain the patency of the line.</p> <p>Examine the dressing, device and site at each shift and promptly remove a catheter that is no longer required.</p> <p>Replace catheter site dressing if it becomes damp, loosened or soiled—do not reinforce a suboptimal dressing with tape.</p> <p>Using chlorhexidine-impregnated dressings at the catheter insertion site has been shown to reduce intravascular access device related bloodstream infection and device colonisation rates. The safety of these dressings has not been established in low birth- weight neonates who may be at risk of skin or systemic toxicity [312] [313].</p> <p>Patients should be educated to alert healthcare staff if they experience any discomfort at the insertion site including pain, burning, swelling or bleeding.</p> <p>When using a needleless connector, the hub should be scrubbed (70% alcohol wipe is the application of choice) before each access to minimise the risk of microbial contamination [293] [297] [318] [319] [320].</p> | |
| Device replacement | <p>All catheters should be checked at each shift and removed when no longer required or if infection is suspected.</p> <p>All catheters inserted in an emergency situation (e.g. by emergency ambulance services or during cardiac arrest) should be removed and replaced when the patient is stable and within 24 hours of insertion.</p> | |
| Replacement of administration sets | <p>Do not routinely replace PIVCs in neonates and children.</p> <p>There are two options for the replacement of PIVCs in adults. See below for more detailed information.</p> | <p>Do not routinely replace CVCs.</p> |

Sources: Queensland Health Guidelines [292] [293] [294] [295]; HICPAC Guidelines [277]; NSW Ministry of Health

Guidelines [301]; Western Australia Guidelines [302]; epic Guidelines [274]; CDC Guidelines [314]

Patient Engagement

Healthcare workers should inform patients of the reason they require an intravascular access device and the plan of care including planned removal. Where appropriate, patients should also be involved in the choice and placement of the intravascular access device in the first place, and educated about keeping the dressing dry.

Replacement of PIVCs

Policies on the replacement of PIVCs should be based on a formal risk assessment that takes into account:

- the availability of staff appropriately trained in the insertion, monitoring, assessment and maintenance of PIVCs on each shift
- the quality of PIVC surveillance in the healthcare facility, including surveillance of regular inspection of the site and device, and of PIVC-related staphylococcus-aureus-bacteraemia (SAB)
- the need for robust documentation and reporting processes on device insertion, maintenance and removal that is supported by the results of audits.

In considering the above factors, healthcare facilities may routinely follow one of the following two options:

Option 1: Replace a PIVC every 72 hours

This practice is based on observational studies that show an increased risk of blood stream infection with PIVCs left in place for more than 72 hours [297].

Option 2: Replace a PIVC based on clinical indication

A strategy of replacing a PIVC when a clinical indication for replacement is identified, rather than routinely at 72 hours, may be considered only when there is:

- Surveillance of PIVC related BSI performed at the facility.
- Comprehensive documentation of insertion, maintenance and removal of PIVCs (audit results demonstrate a sustained compliance with daily PIVC assessment documentation).
- Compliance with competency requirements for insertion and management.

This option is informed by a systematic review, first published in 2011 and updated most recently in 2015, which concluded that rates of bloodstream infection and thrombophlebitis were not significantly different when PIVCs were changed based on clinical indication rather than routinely replaced [310]. The rate of PIVC-related bloodstream infection, however, is approximately 1/1000-3000 patients, so studies with larger patient numbers are required to determine the true impact of this approach on bloodstream infection.

Replacing a PIVC based on clinical indication can be cost saving and may reduce the discomfort for patients associated with regular replacement.

Quality improvement interventions

The implementation of quality improvement interventions can support the appropriate use and management of intravascular access devices and ensure their timely removal [274]. Aspects of quality improvement can include:

- implementing protocols for device insertion and maintenance
- using reminders and prompts to review the use and suitable removal of intravascular access devices
- auditing compliance with intravascular access device protocols and providing feedback to staff
- providing continual professional education to staff regularly engaged in intravascular access device insertion and maintenance.

More information for healthcare workers about intravascular access device insertion, including online training, is available at: <http://www.cec.health.nsw.gov.au/topics/concluded-projects/clab-icu/cli-online-training> [321]; <https://www.health.qld.gov.au/clinical-practice/guidelines-procedures/diseases-infection/infection-prevention/intravascular-device-management> [322].



Figure 10. Personal protective equipment and maximal barrier precautions for insertion of CVCs, PICCs or guidewire exchange (cap, surgical mask (covering mouth and nose), sterile gown, sterile gloves and sterile full body drape) [314]

Intravascular access device care bundles

There are numerous care bundles in use on the management of central and peripheral vascular devices.

Before implementing a care bundle, it is important to identify current practice in the particular area. Gaps in service provision need to be identified, analysed and systematically addressed through the implementation of the bundle. Examples of available bundles include:

- i-CARE <https://www.health.qld.gov.au/clinical-practice/guidelines-procedures/diseases-infection/infection-prevention/intravascular-device-management>
- Health Protection Scotland <http://www.hps.scot.nhs.uk/haic/ic/bundles.aspx>.

3.5.2.3 Ventilation

Certain patients require mechanical ventilatory support by endotracheal tube or tracheostomy. Common medical indications include acute lung injury, chronic obstructive lung disease and acute respiratory acidosis.

When performing intubation and ventilation in the presence of known or suspected infectious agents that are transmitted by the airborne route, airborne precautions are required. For further information refer to Section 3.2.4.

What are the risks?

Ventilator-associated pneumonia (VAP) is a type of healthcare associated pneumonia that can occur in up to 25% of all people who require mechanical ventilation. VAP is a common cause of morbidity and mortality with crude death rates of 5% to 65% as well as increased healthcare costs. VAP can develop at any time during ventilation, but occurs more often in the first few days after intubation, because the intubation process itself contributes to the development of VAP.

VAP primarily occurs because microorganisms colonise the endotracheal or tracheostomy tube and are embolised into the lungs, often in patients who may have underlying lung or immune problems. Bacteria may enter the lungs with procedures such as bronchoscopy.

Many practices have been demonstrated to reduce the incidence of VAP and its associated burden of illness. The first consideration should always be whether intubation is necessary.

Table 26. Summary of strategies for preventing VAP

| Strategy | Summary |
|----------------------------|--|
| Physical strategies | <ul style="list-style-type: none"> • When intubation is necessary, use the orotracheal route as this is associated with a reduction in VAP and a decreased incidence of sinusitis compared to nasotracheal intubation. • Use new circuits for each patient and change these if they become visibly soiled or are |

| | |
|-----------------------------------|---|
| | <p>malfunctioning, as per manufacturer instructions.</p> <ul style="list-style-type: none"> • There is no difference in the incidence of VAP between patients whose airways are humidified using a heat and moisture exchanger, and those whose airways are humidified using a heated humidifier. The decision should be made for each patient, with the aim to ensure adequate moisture output to minimise the risk of airway obstruction. • Change heat and moisture exchangers for each patient every 5–7 days and as clinically indicated. Less frequent changes of heat and moisture exchangers may be associated with a slightly decreased incidence of VAP. • Use a closed endotracheal suctioning system, as safety considerations favour the use of closed systems. The number of disconnections should be minimised to reduce the risk of staff exposure to potentially infected secretions. • Change the endotracheal system for each patient as clinically indicated. Scheduled daily changes of closed systems have no effect on VAP. • Provide endotracheal tubes with subglottic secretion drainage ports for patients likely to require intubation for more than 48 or 72 hours. • Assess patients for sedation, weaning and extubation each day. • Use a microbiological filter to prevent contamination of the ventilator. |
| Positional strategies | <ul style="list-style-type: none"> • Elevate the head of the bed to 30°–45°. Where this is not possible, raise the head of the bed as much as possible. • Semi-recumbent positioning may be associated with a decreased incidence of VAP, but may not be safe for all patients. |
| Pharmacological strategies | <ul style="list-style-type: none"> • Consider the use of the oral antiseptic chlorhexidine. • For patients with severe head injury, consider the use of the oral antiseptic povidone-iodine (in the form of an oropharyngeal rinse). There are insufficient data to make a recommendation in critically ill patients other than those with severe head injuries. |

Sources: Muscedere et al (2008) [275]; Kalil et al (2016) [273]; SHEA (2014) [283]

VAP care bundles

There are numerous care bundles in use for the management and prevention of VAP. Before implementing a care bundle, it is important to identify current practice in the particular area. Gaps in service provision need to be identified, analysed and systematically addressed through the implementation of the bundle.

Examples of available bundles include:

- Scottish Infection Care Society Audit Group VAP Prevention Bundle: <http://www.sicsag.scot.nhs.uk/hai/VAP-Prevention-Bundle-web.pdf>
- IHI Ventilator Bundle: <http://www.ihl.org/resources/Pages/Tools/VentilatorBundleChecklist.aspx>.

3.5.2.4 Enteral feeding tubes

Enteral feeding is usually prescribed for patients in hospital requiring artificial nutrition support for 7–10 days and long-term feeding. Home enteral tube feeding may be considered for patients needing artificial nutrition support for more than 30 days.

What are the risks?

Contamination of feeds is a key concern in both the hospital and community setting, with contamination largely occurring during the preparation or administration of feeds and being linked to serious clinical infection.

Most evidence concerning enteral feeding relates to gastrostomy or percutaneous endoscopic gastrostomies (PEG feeds). However, the principles outlined here are also applicable to nasogastric and jejunostomy feeding.

Table 27. Summary of processes for using enteral feeding tubes

| Activity | Process |
|--------------------|---|
| Preparation | <ul style="list-style-type: none"> • Perform hand hygiene before starting feed preparation. Even closed systems can become decontaminated if hand hygiene is not adequate. |

| | |
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| | <ul style="list-style-type: none"> • Wherever possible, use pre-packaged, ready-to-use feeds. Closed systems (i.e. pre-sterilised, pre-filled, ready-to-use feeds that do not expose feed to the air during assembly) have lower contamination than open systems. The design of the system is also important in order to minimise handling. • If decanting, reconstitution or dilution is required, use a clean working area and equipment dedicated for enteral feed use. • Mix feeds with cooled boiled water or freshly opened sterilised water using an aseptic non-touch technique. |
| Administration | <ul style="list-style-type: none"> • Perform hand hygiene immediately before administration. • Use minimal handling and aseptic non-touch technique to connect the administration system to the enteral feeding tube. • Use aseptic technique for administration of medications. • Discard administration sets and feed containers after each feeding session. |
| Care of insertion site and enteral feeding tube | <ul style="list-style-type: none"> • Perform hand hygiene immediately before commencing. • Wash the stoma daily with water and dry thoroughly. • Flush the enteral feeding tube with fresh tap water before and after feeding or administering medications to help minimise the potential risk of microbial colonisation of the internal and external surfaces. Use cooled boiled water or sterilised water for patients who are immunosuppressed. |

Source: NICE (2012, updated 2017) [276]

Patient-care tip

Patients and carers should be educated in techniques of hand hygiene, enteral feeding and the management of the administration system before being discharged from hospital.

3.5.3 Surgical procedures

The discussion in this section applies to all surgical procedures regardless of setting. While there is less evidence for surgical procedures in office-based practice than in hospitals, the same principles apply.

What are the risks?

The microorganisms that cause surgical-site infections are usually derived from patients (endogenous infection), being present on their skin or from a surgical opening in the body. Exogenous infection occurs when microorganisms from instruments or the operating environment contaminate the site at operation, when microorganisms from the environment contaminate a traumatic wound, or when microorganisms gain access to the wound after surgery, before the skin has sealed.

The risk of surgery-related infection is increased by factors that:

- increase the risk of endogenous contamination (e.g. procedures that involve parts of the body with a high concentration of normal flora such as the bowel)
- increase the risk of exogenous contamination (e.g. prolonged operations that increase the length of time that tissues are exposed)
- diminish the efficacy of the general immune response (e.g. diabetes, malnutrition, or immunosuppressive therapy with radiotherapy, chemotherapy or steroids) or local immune response (e.g. foreign bodies, damaged tissue or formation of a haematoma).

Minimising the risk of surgical procedures

Practices to prevent surgical-site infections are aimed at minimising the number of microorganisms introduced into the operative site—for example by:

- removing microorganisms that normally colonise the skin

- preventing the multiplication of microorganisms at the operative site—for example by using prophylactic antimicrobial therapy
- enhancing the patient’s defences against infection—for example by minimising tissue damage and maintaining normothermia
- preventing access of microorganisms into the incision postoperatively by use of a wound dressing.

This section gives general guidance on preventing surgical infection. More detailed information can be found in the National Institute for Health and Clinical Excellence (NICE) surgical-site infection guidelines (NICE 2008) [326] and the World Health Organization (WHO) Global Guidelines for the Prevention of Surgical Site Infection (2016) [327].

Patient-care tip

Patients and carers require clear, consistent information and advice throughout all stages of their care, including:

- the risks of surgical-site infections, what is being done to reduce them and how they are managed
- how to care for their wound after discharge
- how to recognise a surgical-site infection and who to contact if they are concerned.

An integrated care pathway helps to communicate this information to both patients and all those involved in their care after discharge.

Patients should always be informed if they have been given antibiotics.

Hand hygiene for surgery

Surgical hand preparation should reduce the release of skin bacteria from the hands of the surgical team for the duration of the procedure; in case of an unnoticed puncture of the surgical glove that releases bacteria to the open wound. Surgical hand preparation must eliminate the transient and reduce the resident flora. There are special surgical scrub formulations available for use, although any product used within Australia should preferably be listed on the Australian Register of Therapeutic Goods (ARTG). Current WHO guidelines recommend the use of an alcohol-based formulation for preoperative surgical hand preparation, given its superior antimicrobial efficacy compared to other methods [327] [331]. Specific policies and procedures on products and methods of surgical hand preparation should be developed locally.

PPE for surgical and dental procedures

Personal protective equipment (PPE) is designed and issued for a particular purpose in a protected environment and should not be worn outside that area. For surgical procedures and dentistry, the sequence for putting on PPE differs from that outlined in Section 3.3. In these situations, masks and protective eyewear are applied first prior to hand preparation. Gown and gloves are then put on.

Double-gloving is the process of wearing two sets of gloves to protect against injury from sharps or the transmission of blood-borne infections that can occur in the event of glove perforation. A second pair of gloves protects the inner pair, and should not impact surgical performance [330]. There is little high quality evidence to determine the effectiveness of double-gloving above and beyond single-glove use or no change of gloves; however, many surgeons do prefer to double-glove for their own protection. In the case of double-gloving, it is suggested that the outer gloves are changed routinely during longer surgeries [327].

Information on the use of surgical aseptic technique and standard aseptic technique for wound care can be found in Section 3.1.6 and in Appendix 1—Section 5.11.

3.5.3.1 Preventing surgical site infections (SSIs)

Considerations pre-procedure

Table 28. Summary of perioperative processes

| Stage | Process |
|------------------|--|
| Hand preparation | <ul style="list-style-type: none"> • Operating team members should remove hand jewellery, nail polish and artificial nails. • If hands are visibly soiled, perform hand hygiene with liquid soap prior to scrubbing. • Remove debris from underneath fingernails using a nail cleaner, preferably under running water. • Use a suitable antimicrobial soap or surgical alcohol-based hand rub, preferably with a |

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|---|--|
| | product ensuring sustained activity, as directed and for the length of time recommended by the manufacturer. |
| Operating suite/room or procedure attire | <ul style="list-style-type: none"> Operating team members must wear sterile operation or procedure attire. All operating suite/room staff who are not operating within the critical aseptic field must wear dedicated non-sterile attire in all areas where operations are undertaken. This may contribute to minimising operating environment contamination and reduce the risk of SSIs. Movements in and out of the operating area should be kept to a minimum. |
| Patient preparation | <ul style="list-style-type: none"> Advise patients to shower or have a bath (or help patients to shower, bath or bed bath) using soap, either the day before or on the day of, surgery. Avoid routine removal of hair—if clinical circumstances require hair removal, it should be clipped on the day of surgery or as close as possible to the time of operation. Hair removal should be performed outside of the operating theatre [334] [335]. Shaving hair is strongly discouraged. Provide antibiotic prophylaxis where appropriate, in accordance with the Australian Therapeutic Guidelines. The indication for antibiotics and the choice of antibiotic prophylaxis should be based on these Guidelines. If indicated, it is suggested that antibiotics are administered within 120 minutes before the incision, whilst considering the half-life of the antibiotics [323]. Consider screening for <i>S. aureus</i> and decolonise those with nasal carriage identified before high-risk surgery such as cardiothoracic or orthopaedic. Avoid delaying surgery to provide parenteral nutrition, as there is no high quality evidence to demonstrate the effectiveness of parenteral nutrition in reducing SSIs. The combined use of oral antibiotics and mechanical cleaning in adult patients undergoing colorectal procedures can lead to a reduced risk of SSIs in high-risk patients. Neither oral antibiotics or mechanical cleaning alone are suggested. |

Source: WHO (2016) [323]; SHEA (2013) [332]; SHEA/IDSA (2014) [324]; TGL AEG (2014) [366]; [if not otherwise stated]

Considerations during a surgical procedure

Table 29. Summary of intraoperative processes

| Stage | Process |
|------------------------------------|--|
| Hand hygiene | <ul style="list-style-type: none"> Perform hand hygiene before the first operation on the list using an antiseptic surgical solution, according to the manufacturer's instructions for the product that is being used. Use a single-use brush or pick for the nails, and ensure that hands and nails are visibly clean. Before subsequent operations, perform hand hygiene using an antiseptic surgical solution. If hands are soiled during a procedure, hand hygiene should be performed again with an antiseptic surgical solution. |
| Operating suite/room attire | <ul style="list-style-type: none"> In hospital settings, wear sterile gowns during the procedure. There is no available evidence that double-gloving reduces the risk of SSI or that glove perforation increases the risk of SSI. However, consider wearing two pairs of sterile gloves when there is a high risk of glove perforation and the consequences of contamination of the operative field are high. |
| Patient preparation | <ul style="list-style-type: none"> Prepare the skin at the surgical site immediately before incision using an antiseptic preparation, preferably chlorhexidine. If diathermy is to be used, ensure that antiseptic skin preparations are dried by evaporation and there is no pooling of alcohol-based preparations. Evidence suggests there is no difference in rates of SSI when diathermy is used to make an incision compared with conventional techniques. If an incise drape is required, use an iodophor-impregnated drape unless the patient has an iodine allergy. Do not use non-iodophor-impregnated incise drapes routinely for surgery as they may increase the risk of surgical-site infection. Ensure skin preparation is dry before |

| | |
|-------------------------|---|
| | <p>draping the patient.</p> <ul style="list-style-type: none"> • Preoperative and intraoperative warming can be used to reduce SSI rates. • Administering supplemental oxygen intraoperatively and postoperatively can reduce the risk of SSI in patients undergoing mechanical ventilation. • There is not significant evidence to determine an appropriate re-dosing protocol for prophylactic antimicrobial agents for the prevention of SSIs. |
| Wound management | <ul style="list-style-type: none"> • Avoid routine use of wound irrigation or intracavity antibiotic lavage as measures to reduce surgical site infection as there is no evidence that it reduces the incidence of SSI. There is some evidence that postoperative lavage of the perineal space with povidone-iodine reduces SSI. • There is no robust evidence to support the use of a dressing in the immediate postoperative period for the prevention of SSI. However, it is generally accepted good clinical practice to cover the wound with an appropriate interactive dressing for a period of 2 days unless otherwise clinically indicated—for example, if there is excess wound leakage or haemorrhage. • There is no robust evidence to support the instillation of antibiotics into wounds prior to closure. • Using antimicrobial-coated sutures (included on the ARTG e.g. triclosan-coated sutures) can help to reduce SSI rates**. • There is no robust evidence to support the use of one dressing over another. However, in the majority of clinical situations a semi-permeable film membrane with or without an absorbent island is preferable. • Avoid routine use of intraoperative skin re-disinfection or topical antibiotics as measures to reduce the risk of surgical-site infection in abdominal surgery. • Single or double-ring surgical wound protector devices can reduce the rate of SSIs compared to regular wound protection. The use of wound protector devices should be determined by local need, and the availability and cost of the devices. |

**Information contained in Table 29 should not be interpreted as formal recommendations. Table 29 summarises evidence which has been reported in international guidelines.

Refer to the Summary of Recommendations for all formal NHMRC approved recommendations contained in this Guideline, based on the GRADE methodology.

Source: WHO (2016) [323]; SHEA (2013) [332]; SHEA/IDSA (2014) [324]; TGL AEG (2014) [366]

Considerations post-procedure

Table 30. Summary of postoperative processes

| Stage | Process |
|--|---|
| Dressings | <ul style="list-style-type: none"> • Use aseptic technique for changing or removing surgical wound dressings (see Section 3.1.6 and Appendix 1 Section 5.11 for further information). • Avoid the routine use of topical antimicrobial agents for surgical wounds that are healing by primary intention as measures to reduce the risk of SSI. • Use an appropriate dressing (such as semi-permeable film membrane with or without an absorbent island) to manage surgical wounds that are healing by secondary intention. |
| Cleansings | <ul style="list-style-type: none"> • Use sterile saline for wound cleansing up to 2 days after surgery. |
| Patient care | <ul style="list-style-type: none"> • There is evidence to suggest that blood glucose levels should be controlled during the immediate postoperative period at 180 mg/dL or lower. Intensive postoperative glucose control is not suggested as an effective measure to reduce the risk of SSI, and may in fact lead to a range of serious adverse outcomes. |
| Management of surgical site infection | <ul style="list-style-type: none"> • When surgical-site infection is suspected, take a specimen for culture and then, if clinically indicated, give the patient an antibiotic that covers the likely causative organisms. Consider local resistance patterns in choosing an antibiotic and review the selection in light of results of |

| | |
|--|---|
| | <p>microbiological tests.</p> <ul style="list-style-type: none"> • Antibiotic treatment may not be required for all SSIs: minor infections may respond to drainage of pus (for example, by removal of sutures) and topical antiseptics. Antibiotic therapy carries with it the risk of adverse drug reactions and the development of antimicrobial-resistant bacteria as well as the associated risk of <i>C. difficile</i> diarrhoea. • Avoid the use of Eusol and gauze, or dextranomer or enzymatic treatments for debridement in the management of SSI. |
|--|---|

Source: WHO (2016) [323]; SHEA (2013) [332]; SHEA/IDSA (2014) [324]; TGL AEG (2014) [366]

3.5.4 Aerosol-generating procedures (AGPs)

Aerosol-generating procedures are those procedures that have the potential to create aerosols through inducing coughing or sneezing. They include suctioning, sputum induction, non-invasive ventilation (NIV; for example, continuous positive airway pressure [CPAP]) bronchoscopy and the use of nebulisers (this list is not exhaustive).

If an AGP needs to be performed, the following strategies should be considered regardless of the known infection status:

- Use a single room for the procedure
- Reduce the number of other healthcare workers that enter the room during and after for a set period of time after the procedure (usually approximately 30 minutes)
- Healthcare workers experienced in AGPs should oversee and manage the AGP
- Use appropriate PPE, such as particulate filtration mask (PFR), protective eyewear/ face shields and other PPE, as per standard precautions. Refer to Table 13 in Section 3.2.4

4. Organisational support

Summary

For infection prevention and control to be effective at the clinical level, much organisational support is required. This includes embedding infection control into governance and management structures, initiating procedures (e.g. immunisation programs) to ensure that healthcare workers are protected, instituting processes for surveillance that feed into the overall quality control program, implementing systems for ongoing staff education and training, and incorporating infection control into planning for facility design and maintenance.

Infection control is a health and safety issue, which means that all those working in the healthcare facility—managers, healthcare workers and support staff—are responsible for providing a safe environment for patients and other staff.

Organisational support should aim to ensure that clinical work practices provide patient-centred care—this is not only essential from a safety and quality perspective, but out of consideration for patient preferences. This may require consultation with patients and relevant consumer groups in the development of healthcare services.

The information presented in this part is particularly relevant to managers of healthcare facilities. It outlines responsibilities of management of healthcare facilities, including governance structures that support the implementation, monitoring and reporting of effective work practices. While the focus of the information is acute-care facilities, much of the information is relevant in other healthcare settings.

This section covers:

- **4.1** — Management and clinical governance
- **4.2** — Staff health and safety
- **4.3** — Education and training
- **4.4** — Healthcare associated infection surveillance
- **4.5** — Antimicrobial Stewardship
- **4.6** — Influence of facility design on healthcare associated infection

4.1 Management and clinical governance

Summary

To be effective, infection prevention and control must be a priority in every healthcare facility—this requires total commitment at every level of the organisation.

- Organisational capacity is achieved by having appropriate governance and management structures. This means that managers are aware of the healthcare facility's performance in terms of infection transmission and there are systems in place to prevent the transmission of infection, reduce risk and address problems when they arise.
- The management structure and processes associated with infection control will differ depending on the size of the organisation and the types of healthcare services it delivers. However, the principles of clinical governance apply regardless of the setting and all essential roles and responsibilities should be fulfilled.
- The person in charge of the organisation (e.g. chief executive officer of a hospital, principal of an office-based practice) must have overall responsibility for and direct involvement in the organisation's infection prevention and control program.
- There must be adequate resourcing for dedicated infection control staff, and resources to run the infection prevention and control program including professional development.
- Each organisation should define the outcome measures for monitoring infection prevention and control policies (see Section 4.4.2).
- All employees should understand their roles and responsibilities and have appropriate training to maintain a safe work environment (see Section 4.2).
- Patient-centred healthcare is safer healthcare—patients' healthcare rights must be considered during the development of programs, policies and procedures.

4.1.1 Clinical governance in infection prevention and control

Addressing infection prevention and control requires a facility wide program and is everybody's responsibility. Healthcare facilities have a legal responsibility to provide a safe work environment, safe systems of work and a safe environment for patients and visitors. Clinical governance refers to the system by which managers and clinicians in each healthcare facility share responsibility and are held accountable for patient care. This involves minimising risks to patients and staff, and continuously monitoring and improving the quality of clinical care.

Preventing transmission of infectious agents should be a priority in every healthcare facility. This will involve action to:

- develop a facility-wide strategic plan for infection prevention and control
- establish a system to manage infection prevention and control (such as a committee) with input from across the spectrum of clinical services and management, and a mechanism for considering patients' feedback
- appoint infection control professionals and support their continuing professional development (e.g. attendance at relevant state or national professional organisation meetings)
- incorporate infection prevention and control into the objectives of the facility's patient and occupational safety programs
- provide administrative support, including fiscal and human resources, for maintaining infection prevention and control programs
- provide adequate staff training and protective clothing and equipment, and arrange workplace conditions and structures to minimise potential hazards.

All healthcare workers need to be aware of their individual responsibility for maintaining a safe care environment for patients and other staff.

Practice Statement

New

37. It is good practice for healthcare facilities to have effective clinical handover processes in place that includes infection risks.

Healthcare facilities should develop and implement a structured system for clinical handover, including documented policies and protocols.

Practical info

Clinical communication problems are a major contributing factor in 70% of all hospital sentinel events.

Patient safety and communication between healthcare workers can be maximised when effective clinical handover processes are in place. This should include consideration for transfers between wards and departments, transfers between different healthcare facilities and communication/alerts on re-admission for long-term infection risks such as multi-resistant organism colonisation.

All healthcare facilities should develop and implement an organisational system for structured clinical handover, including documented policies/procedures and protocols, and agreed tools and guidelines. These processes should be evaluated regularly to ensure the effectiveness of clinical handover is maximised.

Refer to National Safety and Quality Health Service *Standard 6 [367]* for more information on clinical handover.

Evidence to decision

Benefits and harms The benefits of effective clinical handover processes, which include improved communication between clinicians and staff, can result in improved patient safety and care, and outweigh any possible harms.

Certainty of the evidence A 2008 literature review of clinical handover processes [338] identified that implementing protocols and education on training around clinical handover processes results in a range of positive outcomes including less technical errors and information omission, and greater staff confidence in undertaking effective handover processes.

Values and preferences There should be no significant impact on patient preferences and values, nor on health equity.

Resources and other considerations Education and training for staff on effective clinical handover processes requires resourcing.

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Having clinical handover processes in place in healthcare facilities is justified to reduce healthcare associated infection.

4.1.2 Roles and responsibilities

Management and clinical governance can have a positive impact on the effectiveness of infection prevention and control by driving continuous quality improvement and promoting a non-punitive culture of trust and honesty [340]. It is important that healthcare managers and clinicians effectively collaborate, and involve patients as partners in their healthcare in order to effect change and achieve the best possible outcomes [339].

The roles and responsibilities described below are most relevant to acute health care settings. However, all the roles described in this section are important for effective infection prevention and control and can be readily adapted to other healthcare settings—for example, with the practice principal fulfilling relevant roles and responsibilities of a Chief Executive Officer (CEO), and the office manager or other staff representative with an interest in infection prevention and control fulfilling the role of infection control professional.

A. Chief Executive Officer/Administrator

The healthcare facility's CEO or designated equivalent administrator should support and promote infection prevention and control as an integral part of the organisation's culture through the following strategies:

- having a performance agreement that includes infection prevention and control outcomes as a key performance indicator
- endorsing the inclusion of specific articulated infection prevention and control roles, responsibilities and accountabilities for relevant staff within the facility's management plan
- attending and participating in each Infection Prevention and Control (IPC) Committee meeting
- ensuring that infection control professionals are resourced:
 - in terms of co-workers, information technology, access to up-to-date information, designated office/work space and tools to meet relevant infection prevention-related legislative, regulatory and accreditation requirements
 - to achieve negotiated healthcare associated infection reduction targets and to perform the essential tasks outlined below
- ensuring that the healthcare facility's IPC program includes involvement of one or more medical practitioners to support and play a shared leadership role
- ensuring that the rights of patients, as articulated in the Australian Charter of Healthcare Rights [14], are integral to the IPC program
- committing to the IPC program vision, mission, priorities, targets and annual infection prevention and control plan with specific, measurable goals for healthcare associated infection risk mitigation and reduction—these should be outlined in an annual business plan which the CEO (or his or her designate) and the infection control professional jointly develop
- supporting an organisational culture that promotes individual responsibility for infection prevention and control among all staff and values the IPC program contribution to the safety of patients, healthcare workers and others. This support includes ensuring IPC program staffing levels are sufficient and incorporating responsibility for infection prevention and control into every staff member's job description
- authorising infection control professionals to:
 - implement IPC program recommendations
 - intervene when clinical or other practices pose infection risks (e.g. halt building and construction activities, close units during outbreaks and guide patient placement for isolation or cohorting)
- recommending remedial action when infection prevention and control measures are compromised or breached.

In some Australian states and territories and internationally, performance against infection control indicators is monitored.

B. Infection control professionals

Infection control professionals should have the skills, experience and qualifications relevant to their specific clinical setting and be able to:

- develop, manage and evaluate governance of infection prevention and control systems, related programs and services
- provide expert infection prevention consultancy and strategic direction to the healthcare facility and external agencies.

Infection control professionals are primarily responsible for designing, coordinating, implementing and undertaking ongoing evaluation of the facility's infection prevention and control program and policies. This includes compliance with the respective state/territory and/or national accreditation, licensing, policy or regulatory requirements. They are also responsible for equipment and product evaluation.

Infection control professionals need to be supported by the facility with resources, authority and time to maintain clinical and professional currency (including support for credentialing and have, preferably, a postgraduate qualification relevant to infection control [see Section 4.3.1]).

Infection control professionals must be involved in decisions on facility construction and design, patient placement ratios (e.g. single rooms, negative pressure rooms) and environmental assessments (see Sections 4.6 & 4.6.1).

The infection control professional's performance should be appraised at least annually, along with negotiation of individual professional development goals, support, opportunities and plan of work.

C. Infection prevention and control committee

A multidisciplinary IPC committee should review and guide the healthcare facility's IPC program, strategies and plans. Membership must include, but not be limited to:

- the CEO or his/her designate
- an executive member with the authority to allocate the necessary resources and take remedial action as needed from time to time
- an infection control professional
- one or more medical practitioners (preferably a clinical microbiologist and/or an infectious diseases physician).

The meeting frequency and content will depend on the facility's size, case-mix complexity and the infection risk of populations serviced. IPC Committee activity should be measured against an operational plan with set priorities to target within key focus areas.

The IPC committee should have a formal mechanism for regularly considering patients' experiences and feedback and modifying the IPC program accordingly.

The IPC committee should have an organisational communication strategy to facilitate day-to-day activities and reporting activities, which should be able to be escalated in response to an incident or outbreak. Regular and ad-hoc communication processes should exist between the IPC team and relevant public health authorities. Healthcare facilities that do not have access to an IPC committee (or infection control professional) should consult with an infection control professional in a larger health service for program advice and support.

D. Infection prevention and control processes in office-based practice

In office-based practice, the processes associated with infection prevention and control will differ although the responsibilities are the same. The principal of the practice is equivalent to the CEO. He or she has overall responsibility for infection prevention and control in the practice and should demonstrate a strong commitment to an agreed infection prevention and control plan based on the identified risks for that practice. Local policies and procedures need to be developed and implemented as part of standard operating procedures. A nominated staff member must take on the role of infection control professional, developing infection prevention and control procedures and overseeing their implementation. This staff member is likely to need additional training and perhaps ongoing external support in managing infection prevention and control issues. Infection prevention and control should be considered at every staff meeting, with discussion of procedures and processes of the practice and any problem areas.

For more information, see The Royal Australian College of General Practitioners' *Infection Prevention and Control Standards for general practice and other office-based and community-based practices* (5th edition, 2022) [379] and Australian Dental Association's *Guidelines for Infection Control* (5th edition, 2024) [341].

4.1.3 Infection prevention and control program

Infection prevention and control (IPC) program

The IPC program is the means by which infection prevention and control practice is implemented in every part of the healthcare facility. IPC programs should be considered across all healthcare settings including acute care hospitals, office-based practices (e.g. general practice clinics, dental clinics, community health facilities), the setting in which paramedics work and long-term care facilities. This should involve the development of a risk-management policy for each healthcare facility (see Section 4.1.4). The World Health Organization [344] has developed recommendations for the core components of an IPC program. These recommendations include:

- the need for a dedicated, trained team within each healthcare facility to run an IPC program
- the development of infection prevention and control policies and procedures that are multimodal and based on national and state/territory Guidelines
- education and training of healthcare workers to improve their understanding of healthcare associated infection (HAI) and antimicrobial resistance, and so they can implement relevant policies and procedures
- the need for facility-based HAI surveillance which includes timely mechanisms for feedback and reporting to relevant healthcare professionals and senior management
- the use of multimodal strategies to address the prevention of HAIs

- regular monitoring and review of healthcare practices to ensure that all policies and procedures are being correctly implemented against key performance indicators
- developing policies and procedures related to staff health and safety, including immunisation policies, and strategies to prevent occupational exposure to infection hazards
- evaluation of chemical disinfectants, products and equipment purchase
- ensuring at the facility level that healthcare environments are clean and appropriate materials and equipment are available to enable appropriate infection prevention and control procedures, see [environmental cleaning and infection prevention and control resources](#).
- provide input during the planning, design and construction of healthcare facilities.

An IPC program may also include antimicrobial stewardship initiatives run in conjunction with the pharmacy department/services.

A useful resource is the National Safety and Quality Health Service *Measurement for Improvement Toolkit*, which provides a set of practical methods to measure the safety and quality of clinical health care services:

<https://www.safetyandquality.gov.au/wp-content/uploads/2012/01/measurement-for-Improvement-toolkit-a.pdf>

Resource allocation

The number of infection prevention and control professionals should ensure that healthcare facilities have the appropriate level of skills and resources required to develop IPC programs and the capacity to respond to of HAIs.

Healthcare facility managers should ensure that there are sufficient resources available to support all aspects of the IPC program, including [342] [343]:

- providing specific infection prevention and control full-time equivalents, determined according to the scope of the IPC program, the complexity of the healthcare facility, the characteristics of the patient population and the needs of the facility and community (office-based practices may choose to attribute responsibilities and functions relating to infection prevention and control to a particular staff member)
- meeting occupational health needs related to infection prevention and control (e.g. provision of appropriate technologies and protective personal equipment, healthcare worker immunisation, post-exposure evaluation and care, counselling services for healthcare workers involved in outbreak management, evaluation and management of healthcare workers with communicable infections)
- in a hospital setting, providing clinical microbiology laboratory support, including a sufficient number of medical technologists trained in microbiology as well as support appropriate: to the healthcare setting; for detecting endemic and emerging pathogens; monitoring transmission of microorganisms; planning and conducting epidemiologic investigations
- funding surveillance cultures, rapid diagnostic testing for viral and other selected pathogens, preparation of antibiotic susceptibility summary reports and trend analysis.

4.1.4 Risk management

Risk management is the basis for preventing and reducing harm arising from healthcare associated infections and underpins the approach to infection prevention and control throughout these guidelines. Within a healthcare facility, a successful approach to risk management includes action at the organisational level (for example providing support for effective risk management through an organisational risk-management policy, staff training and monitoring and reporting) as well as in clinical practice.

Organisational support for risk management

For risk management within an organisation to be effective, there needs to be appropriate infrastructure and culture; a logical and systematic approach to implementing the required steps; and embedding of risk-management principles into the philosophy, practices and business processes of an organisation, rather than it being a separate activity or focus. Factors that support risk management across the organisation include: development of a risk-management policy; staff training in risk management; implementation of a risk register, risk treatment schedule and integrated action plans; monitoring and audit; and risk-management reporting.

An infrastructure and environment that encourages two-way communication between management and healthcare workers and among healthcare workers is an important factor in increasing the level of support for and compliance with infection prevention and control programs. Management should:

- provide direction (e.g. nominate issues for attention that are relevant to the core business of the organisation, such as respiratory hygiene and cough etiquette in general practice, prevention of diarrhoeal disease in paediatrics, appropriate management of urinary catheters in spinal injury care)
- establish and evaluate periodic goals (i.e. nominate reduced rates for performance improvement)
- seek feedback on policy directives particularly in regards to changes in clinical care protocols or new technologies and how patients can be involved in policy formation

- provide information to individuals, self-directed work groups, patients and other stakeholders, with an emphasis on continually improving performance.

Healthcare workers can contribute to the development of risk-management structures, and are integral to the strategies within these. Strategies and examples to assist individual healthcare workers to reduce risk are included in Appendix 1.

New technologies and testing

Before purchasing any new technologies, consultation should occur with the reprocessing unit and infection prevention and control unit.

Advice should be sought on [345]:

- the impact on risk of infection to the patient or other individuals as a result of the product
- whether the product may be implicated in the transmission of infection over time
- whether the product will have infection prevention and control implications for other consumables, equipment or plans
- whether any difficulties in cleaning and reprocessing the product may impact on the product's functionality and safety
- whether any alternative products that are available may present a lower risk of infection
- whether the product is 'listed' (hard surface disinfectants with specific claims), 'included' (disinfectants intended for use on medical devices) or exempt from the Australian Register of Therapeutic Goods.

A risk assessment should be undertaken before purchasing new technologies which should consider:

- the design of the instrument—how this may impact the ease of cleaning
- local capacity and expertise—whether staff will be able to adequately reprocess this instrument.

Practice Statement

New

38. It is good practice to use chlorhexidine in appropriate situations and only when clinically indicated.

Healthcare professionals should consider the appropriateness of using chlorhexidine in every clinical situation, as discussed in these Guidelines.

Chlorhexidine-containing products, devices or solutions must never be used on or around patients with known chlorhexidine sensitivity.

Practical info

Chlorhexidine is an antiseptic antibacterial agent which is widely used in healthcare facilities, general practice and aged care settings. This product is available in numerous different forms: Dressing; Gel/Jelly; Lotion; Solution; Liquid; Pad; Sponge; Cream. Skin cleansing with chlorhexidine plays an important role in reducing the incidence of healthcare associated infections.

It is good practice to use chlorhexidine in situations where there is a clear patient benefit and efforts should be made to ensure it is used correctly.

There is a need for greater understanding of how resistance mechanisms are changing the susceptibility of disease-causing bacteria to chlorhexidine and/or antibiotics. There is also currently no standardised method or definition of chlorhexidine 'resistance'.

For further information on appropriate chlorhexidine use, see the Australian and New Zealand College of Anaesthetists and the Australian Commission on Safety and Quality in Health Care *Joint Safety Statement: Topical application of chlorhexidine and the risks of accidental injection* [351].

Evidence to decision

Benefits and harms It is appropriate to use chlorhexidine in hand hygiene formulations, as an oral care solution, for skin prep solution, in impregnated dressings where benefit is shown including for intravascular devices and in high risk settings and at-risk populations such as intensive care units and oncology, in impregnated or coated central venous catheters. Each of these is detailed in the corresponding sections of this Guideline.

However, it is suggested that healthcare workers limit the use of chlorhexidine and consider the

appropriateness of using chlorhexidine in every clinical situation as this can assist in preventing chlorhexidine resistance.

Certainty of the evidence The majority of studies related to chlorhexidine resistance are controlled laboratory/susceptibility studies (n=24) so the results cannot be generalised and applied to clinical settings.

Values and preferences There are no acceptability considerations which directly impact the patient for the use of chlorhexidine.
Applying caution in the use of chlorhexidine should not impact on patient health equity.

Resources and other considerations There are no significant resource implications of limiting the use of chlorhexidine to clinically appropriate situations.

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. The use of chlorhexidine when clinically indicated and appropriate is justified to reduce healthcare associated infection.

Practice Statement

New

39. It is good practice to include chlorhexidine in a healthcare facility's chemical register. Any adverse reactions to chlorhexidine should be maintained in an organisational risk register and reported to the Therapeutic Goods Administration.

Practical info

Anaphylactic reactions to chlorhexidine are rare but are potentially life-threatening complications. Increasing chlorhexidine usage by consumers and healthcare workers has resulted in a number of different adverse reactions including allergic contact dermatitis, photosensitivity, anaphylaxis and septic shock [346] [347] [348] [349]. Greater recognition of the potential for chlorhexidine-related anaphylaxis is needed, and patients should be prompted for any chlorhexidine reactions/anaphylaxis during the 'allergy history' subjective assessment. Chlorhexidine containing solutions, dressings or impregnated devices should never be used with patients known or suspected to be sensitive to chlorhexidine.

Whilst evidence is limited, it seems the severity of allergic reactions is greatest with repeated exposure and related to chlorhexidine impregnated central venous catheters. The effects of chlorhexidine use are relative to the population group where chlorhexidine is mostly used - which is in perioperative settings. Invasive use is a significant risk factor for anaphylaxis, and reactions are primarily reported in this population group, particularly with repeat and frequent exposure.

Exposure to chlorhexidine in the clinical environment is routine therefore avoidance of recurrent anaphylactic reactions must begin with the identification and substitution of all products containing chlorhexidine. Careful planning and implementation is required to prevent inadvertent exposure to chlorhexidine in the sensitised patient as they move throughout the hospital system.

Systems should be in place in healthcare facilities to ensure that [274]:

- healthcare workers are aware that chlorhexidine can cause anaphylactic reactions
- any known patient allergies are recorded in patient notes
- healthcare workers check for the use of chlorhexidine on any patient who experiences an unexplained reaction.

For further information on appropriate chlorhexidine use, see the Australian and New Zealand College of Anaesthetists and the Australian Commission on Safety and Quality in Health Care *Joint Safety Statement: Topical application of chlorhexidine and the risks of accidental injection* [351].

Evidence to decision

| | |
|---|---|
| Benefits and harms | As chlorhexidine usage can result in a number of adverse reactions including anaphylaxis, there is significant benefit in including chlorhexidine in a healthcare facility's register and recording any adverse reactions. |
| Certainty of the evidence | A recent literature review found that chlorhexidine-related anaphylaxis appears to be a relatively rare event in healthcare. However, the evidence in this area is limited, and the studies available tend to be retrospective and focused specifically in perioperative settings. The limited nature of the evidence makes it difficult to determine the clinical significance of these findings, and it is possible that larger acute care healthcare facilities might encounter one or more anaphylactic events each year. |
| Values and preferences | Recording adverse reactions to chlorhexidine in an organisational risk register should not impact on patient values and preferences, nor health equity. |
| Resources and other considerations | It is suggested that an existing register within a healthcare facility is used, so this practice should not impact on resourcing besides possible extra training for staff. |

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Including chlorhexidine in a chemical register is justified to reduce healthcare associated infection.

4.1.5 Taking an organisational systems approach to infection prevention quality and safety

Addressing infection prevention and control issues requires a multi-component, facility-wide program and is everybody's responsibility. This section gives an outline of a systematic approach that has been shown to be effective, care bundles, together with examples of the organisational support required at a facility level to address two crucial areas of infection prevention and control—reducing sharps injuries to healthcare workers and, in patients, lowering the incidence of bloodstream infections associated with intravascular devices. Sections 4.2 to 4.6 discuss the separate aspects of a systems approach to infection prevention and control.

Care bundles

'Care bundling' is an approach developed by the United States Institute of Healthcare Improvement to improve consistency of practice in healthcare facilities, particularly for conditions and procedures known to increase patients' risk of healthcare associated infections.

Care bundles can be used to monitor care, and care bundle results can provide feedback to clinical staff in order to decrease the rate of healthcare associated infections related to that condition or that procedure. It is important that bundles are designed, implemented and evaluated with measurement designed for quality improvement rather than research or judgment.

Examples of some procedural care bundles are given in Sections 4.4, 3.5.2.2, & 3.5.2.3.

Reducing sharps injuries

Safe handling of sharps is discussed in more detail in Section 3.1.2. A systems approach can support reducing sharps injuries through [352]:

| |
|---|
| <p>Clinical governance</p> <ul style="list-style-type: none"> • Champion a culture of safety underpinned by concepts of patient-centred care. <p>Staff health and safety</p> <ul style="list-style-type: none"> • Adopt and evaluate the use of passive or active safety engineered devices as an alternative to sharps without safety engineered features. • Standardise changes to work practices that will reduce risks (e.g. using instruments, rather than fingers, to grasp needles, retract tissue, and load/unload needles and using appropriately designed single-handed devices to unload needles and scalpels). |
|---|

Education and training

- Provide education on the use of new devices and work practices.

Surveillance

- Ensure comprehensive reporting of injuries and preventive strategies.

Facility design

- Apply engineering controls (e.g. sharps disposal containers and passive or active sharps devices engineered to prevent sharps injury).

Lowering the incidence of intravascular device (IVD)-related bloodstream infections

Section 3.5.2 outlines infection prevention and control guidance for healthcare workers to follow when inserting an invasive medical device such as a central venous catheter—with the first consideration being whether the device is necessary.

The care bundle (see also Section 3.5.1) for central venous catheter insertion stipulates the use of hand hygiene, maximal barrier protection, optimal intravascular catheter site selection, topical chlorhexidine for skin disinfection, and daily review to ensure that catheters are removed as soon as they are no longer necessary. Support and infrastructure requirements to facilitate implementation of these measures include:

Clinical governance

- Champion a culture of safety underpinned by the concepts of patient-centred care.
- Utilise peer networks to promote and increase compliance with best practice techniques.

Education and training

- Develop orientation programs for staff including rigorous grounding in facility policies for infection prevention standard precautions.
- Develop and promote an education program that addresses facility procedures for the insertion, maintenance and removal of IVDs.
- Provide staff with ongoing education to maintain high levels of compliance with facility policy.
- Engage patients so they have the knowledge and skills to be actively involved in their own care.

Surveillance

- Implement a quality adherence tool for best practice IVD insertion, maintenance and removal.
- Measure and evaluate performance through formal and informal audits of clinical practice.
- Provide feedback to staff.
- Measure bloodstream infection rates to monitor performance.

Facility design and equipment

- Ensure appropriate equipment is provided, such as IVD-insertion kits with standardised content to enable a competent healthcare professional to perform the procedures.
- Ensure ready access to handwashing basins and alcohol-based hand rub.

4.2 Staff health and safety

Summary

- Infection protection for all members of the healthcare workforce should be an integral part of the infection prevention and control and work health and safety programs of every health service organisation.
- These programs should include implementing a staff health screening policy, promoting immunisation, management of vulnerable members of the workforce in specific circumstances (e.g. pregnant healthcare workers) and having processes for minimising and managing risk exposure.
- The organisation has a duty of care to protect all members of the workforce and members of the workforce have a responsibility to protect themselves and to not put others at risk.

Roles and responsibilities

Healthcare facilities

States and territories work health and safety legislation places a duty of care on employers to ensure workplace health and safety, including where occupational infectious disease hazards exist.

All members of the workforce, including contractors, volunteers and students, should be informed of their healthcare facilities or training institution's policy on health screening.

Counselling should be provided to any individuals whose ability to undertake work or complete study may be impacted due to transmissible infections.

The privacy and civil rights of all members of the workforce should always be respected and not breached.

The five measures of protection for infection prevention are:

- health status screening (see Section 4.2.1)
- education on safe work practices that minimise the transmission of infection (see Section 4.3)
- safe systems of work, with workplaces that are designed to minimise the transmission of infection (see Section 4.6)
- physical protection, including the use of personal protective equipment (see Section 3.3) and immunisation (see Section 4.2.1)
- reporting systems for compliance and identifying breaches of infection prevention and control protocols.

Members of the health workforce

Any member of the health workforce can become exposed to infectious agents in a number of ways, including:

- through direct contact with an infectious patient
- as a result of a sharps injury
- through eating or drinking in a patient care area.

Members of the health workforce may also put patients at risk of infection if they have an infectious condition.

Members of the health workforce are obligated to follow specific established infection prevention and control policies as part of their employment. Failure to follow policies and procedures may be grounds for disciplinary action.

Members of the health workforce with infections need to manage their condition through seeking medical care, and receiving counselling, where appropriate, about their work options, and staying home if they are unwell.

Information about exclusion periods for members of the workforce with acute infections is in Section 4.2.2.

Information for members of the workforce in specific circumstances (e.g. pregnant healthcare workers) is in Section 4.2.4.

Information about members of the workforce who carry a blood borne virus and how this impacts on their ability to perform exposure-prone procedures is in Section 4.2.5.

4.2.1 Health status screening and immunisation

Staff health screening policies

Healthcare facilities should specify a framework for the assessment, screening and vaccination of members of the workforce to minimise the risk of transmission of vaccine preventable diseases. This must align with relevant state and territory policies and/or legislation.

Before beginning employment, all members of the health workforce, including contractors, volunteers and students undertaking clinical placements, should be assessed and offered testing and/or vaccination for specific infectious diseases before being allowed to work in a healthcare facility. Particular attention should be paid to immune status, skin conditions, pregnancy in staff, as well as risk factors for specific groups of patients. These conditions may vary according to state/territory specific requirements and recommendations.

Routine screening and assessment

Routine screening at the start of employment occurs in three forms:

- personal assessment of disease and immune status—a questionnaire (with recording of information gained) should check for details of medical history, particularly for rubella, measles (rubeola), mumps, chickenpox (varicella), hepatitis B, immune disorders, skin conditions, and for prior exposure to tuberculosis (including working in high-risk settings and high-risk demographic background)
- immunisation

- laboratory and other testing—this may include a tuberculin skin test or blood test (interferon gamma release assay-IGRA).

Immunisation

Pre-vaccination screening

Pre-vaccination screening steps and procedures are outlined in the *Australian Immunisation Handbook*, including a pre-vaccination screening checklist. Healthcare facilities should have education programs to support their immunisation policy and reinforce the need for compliance.

Occupational vaccination program

Employers should take all reasonable steps to ensure that staff members are protected against vaccine-preventable diseases. Where members of the workforce may be at significant occupational risk of acquiring or transmitting a vaccine-preventable disease, a comprehensive occupational vaccination program should be implemented. Such a program should include:

- a vaccination policy
- maintenance of current staff vaccination records
- provision of information about the relevant vaccine-preventable diseases
- the management of vaccine refusal (which should, for example, include reducing the risk of a member of the workforce transmitting disease to a vulnerable patient).

Healthcare facilities should advise members of the workforce of the potential consequences if they refuse reasonable requests for immunisation. Such advice and refusal to comply should be documented. Duties may be modified if members of the workforce have a confirmed infection that may directly affect the risk of transmission of infection during exposure-prone procedures. This is determined at the local facility level.

Vaccine refusal, contraindication to vaccination and vaccine non-response may be managed by ensuring appropriate work placements, work adjustments and work restrictions.

Recommended vaccinations

The most recent edition of the *Australian Immunisation Handbook* [353] provides detailed information on immunisation schedules and vaccines. Staff vaccination programs should comply as much as possible with these schedules, which acknowledge that some circumstances may require special consideration before vaccination.

Members of the workforce should be up to date with routinely recommended vaccines for adults, such as dTpa (diphtheria-tetanus-acellular pertussis)—containing, MMR (measles-mumps-rubella) vaccines and catch-up vaccinations.

Members of the workforce should check which vaccines their state or territory guidelines require them to have and what documentation they need to support this. Contact your state or territory health authority for more details regarding mandatory vaccination programs for all members of the workforce.

Table 31. Recommended vaccinations for all healthcare workers

| Healthcare workers | Disease/Vaccine |
|--|---|
| All members of the workforce: including all workers and students directly involved in patient care or the handling of human tissues, blood or body fluids. | Hepatitis B Influenza Pertussis (dTpa) MMR (if non-immune) Varicella (if non-immune) COVID-19 (as per jurisdictional policy directives) |
| Members of the workforce who work with remote Indigenous communities in Northern Territory, Queensland, South Australia and Western Australia; and other specified healthcare workers in some jurisdictions. | Vaccines listed for 'All healthcare workers', plus hepatitis A |
| Members of the workforce who may be at high risk of exposure to drug-resistant cases of tuberculosis (dependent on state or territory guidelines). | Vaccines listed for 'All healthcare workers', plus consider Bacillus Calmette-Guérin (BCG) vaccine |

Source: Adapted from the *Australian Immunisation Handbook* [353] and jurisdictional health department policies.

Workforce immunisation risk matrix

The National Safety and Quality Health Services (NSQHS) Standards recommend that healthcare facilities implement a risk-based immunisation program to identify, evaluate and address risks that are relevant to the scope of the role and function of the members of the workforce. The risks for vaccine-preventable diseases may differ between work groups or departments within the same organisation.

Workforce immunisation risk factors

Each of the following risk factors need to be considered when assessing risk for workforce

Immunisation requirements and status:

- Can all members of the workforce provide history/evidence of vaccination or immunity to specific vaccine-preventable diseases/infections?
- Does the healthcare facility have processes/services available to assess and screen members of the workforce for their vaccination/immunisation status, such as a serology service?
- Are there processes in place that protect members of the workforce from exposure to vaccine preventable-diseases and infections?

A risk matrix is a qualitative tool used to assess the risk of an event and the consequences associated with the event. The [Workforce Immunisation Risk Matrix](#) (Figure 13) is an example of a tool which healthcare facilities can use to assess the level of risk to members of the workforce and the healthcare facility from vaccine-preventable diseases and infections. This risk matrix will generate one risk rating, either low, medium, high or very high, to describe the level of risk.

A risk assessment should be completed at an organisational level first, but can also be applied at a department/ward level if required.

Risk assessment should be an ongoing process. At the time of completing a risk assessment for VPDs, the incidence of VPDs in the community, including outbreaks and seasonal epidemics, should be considered, along with any outbreaks of VPDs within the healthcare facility. This means that the risk rating may increase if a healthcare facility is responding to changing circumstances either within or external to the organisation for a period of time.

Figure 13. Workforce Immunisation Risk Matrix

| | | Workforce immunity/vaccination status & access to workforce screening | | |
|--------------------------|--|---|---|--|
| | | All members of the workforce have up-to date records of their immune status/vaccination history available AND A workforce screening/vaccination program is in place | Up-to-date immune status/vaccination records are NOT available for all members of the workforce AND A workforce screening/vaccination program is in place | Up-to-date immune status/vaccination records are NOT available for all members of the workforce AND A workforce screening/vaccination program is NOT in place |
| Risk of disease exposure | No increased risk of exposure to disease or infection associated with working in the health service organisation | Low | Medium | High |
| | Members of the workforce have a risk of exposure to vaccine-preventable disease due to the nature of their role | Medium | High | Very high |
| | Hospital-based outbreak of vaccine-preventable disease (no evidence of community transmission) | Medium | High | Very high |
| | Community-wide outbreak of a vaccine-preventable disease | High | Very high | Very high |
| | Disease with no available vaccine | Very high | Very high | Very high |

| | |
|--------------------|---|
| Low risk | Risk is managed by routine procedures. There is minimal risk of harm or injury from the risk |
| Medium risk | Risk is managed by specific monitoring or audit procedures. There is potential for harm or injury from the risk. |
| High risk | There is a serious risk that must be addressed immediately. Consequences to individuals and the organisation are high due to a high potential for harm or injury |

Very high risk

There is a serious risk that must be addressed immediately. The magnitude of the consequences to the individual and organisation of an event, should it occur, are considered very high with potentially significant harm or injury.

Source: Workforce immunisation risk matrix adapted from the *Australian Commission on Safety and Quality in Health Care*
<https://www.safetyandquality.gov.au/sites/default/files/migrated/Workforce-immunisation-risk-matrix1.pdf>

In this example, when performing a risk assessment, you should select the statements that best describe the current situation in the healthcare facility or department, from the options under the heading Workforce immunity/vaccination status and access to workforce screening and the heading Risk of disease exposure to identify the risk level. Once the risk rating has been identified (Low, Medium, High, Very high), the healthcare facility should develop plan of action to prioritise strategies and identify resources to address vaccine-preventable disease risks to members of the workforce.

Staff records

Employers and healthcare facilities need to retain details of screening results and immunisations provided, including vaccine preventable disease history, date and results of serology, record of immunisations consented/refused, date given, batch number, type and brand name of vaccine.

Records need to be secure and accessible by authorised personnel when needed, updated when relevant events occur and maintained in accordance with confidentiality and privacy laws.

Statutory Requirement

New

40. It is recommended that all healthcare workers to be vaccinated in accordance with the recommendations for healthcare workers in the *Australian Immunisation Handbook*.

Note: The advice reflects recommended practice supported by strong evidence. Healthcare facilities must also consider relevant state, territory and/or Commonwealth legislation regarding mandatory vaccination programs for healthcare workers.

Practical info

Healthcare facilities must consider relevant state, territory and/or Commonwealth legislation regarding mandatory vaccination programs for healthcare workers.

Healthcare facilities should maintain a record of healthcare workers who choose not to vaccinate in accordance with the *Australian Immunisation Handbook* [353].

For further information on health status screening and immunisation see Section 4.2.1.

Evidence to decision

Benefits and harms The benefits of vaccination programs clearly outweigh the harms.

The benefits of adhering to recommendations for healthcare workers in the *Australian Immunisation Handbook* [353] include reduced risk of infection and transmission of vaccine-preventable disease to other healthcare workers and to patients.

Certainty of the evidence

This practice is supported by strong clinical and epidemiologic studies. Full details of the review of all relevant information from the latest medical literature can be found in the appendices of the most recent edition of the *Australian Immunisation Handbook* [353]. This also provides detailed information on immunisation schedules and vaccines. Staff vaccination programs should comply as much as possible with these schedules.

Healthcare facilities must also consider relevant state, territory and/or Commonwealth legislation regarding mandatory vaccination programs for healthcare workers.

Values and preferences Regardless of cultural, social and economic background, all healthcare workers should be offered protection against vaccine-preventable diseases.

If vaccinations are refused, contraindications to vaccination are present and/or if vaccine non-response occurs, this should be managed by ensuring appropriate work placements, work adjustments and work restrictions in healthcare facilities.

Resources and other considerations Vaccination has been demonstrated to be one of the most effective and cost-effective public health interventions.

Rationale

Vaccination not only protects individuals, but also protects others in the community by increasing the overall level of immunity in the population and thereby minimising the spread of infection [353].

Australia has one of the most comprehensive publicly funded immunisation programs in the world. All vaccines marketed in Australia are manufactured according to strict safety guidelines and are evaluated by the Therapeutic Goods Administration, to ensure they are efficacious and are of adequate quality and safety, before marketing approval is granted [353].

Healthcare facilities must also consider relevant state, territory and/or Commonwealth legislation regarding mandatory vaccination programs for healthcare workers.

4.2.2 Exclusion periods for healthcare workers with acute infections

Every healthcare facility should have comprehensive written policies regarding disease-specific work restriction and exclusion, which include a statement of authority defining who can implement such policies.

Any member of the health workforce who has an infectious disease has a responsibility to:

- consult with an appropriate medical practitioner to determine that they are capable of performing their tasks without putting patients or other workers at risk
- undergo regular medical follow-up and comply with all aspects of informed clinical management regarding their condition.

These policies should encourage members of the workforce to seek appropriate preventive and curative care and report their illnesses, medical conditions, or treatments that can render them more susceptible to opportunistic infection or exposures.

Employers should not penalise members of the health workforce with loss of wages, benefits, or job status if they are unwell with an infectious disease.

The overarching principle for exclusion periods is that staff members should not come to work if they have signs or symptoms of a potentially infectious disease.

The Communicable Diseases Network Australia (CDNA) provides specific guidance on the management of staff infected with a range of diseases. For more information, see CDNA's *Series of National Guidelines* [359].

Table 32. Staff exclusion periods for infectious illnesses

| Acute infection | Exclusion period |
|--|---|
| Conjunctivitis | Must not provide patient care for the duration of symptoms (i.e. while eye discharge is present). |
| Gastroenteritis* (except norovirus) | Must not come to work while symptomatic (e.g. diarrhoea and/or vomiting) and until 24 hours after symptoms have resolved. If the cause is unknown, possible exclusion for 48 hours until the cause is identified. Healthcare staff who have a food handling role should always be excluded for until 48 hours after symptoms have resolved. |
| Glandular fever | No need for exclusion, even if having direct patient contact, provided staff members are well enough to return to work |

| | |
|--|---|
| | and employ standard precautions. |
| Hand, foot and mouth disease | Healthcare workers should be excluded until all blisters have dried. Those who may have been in contact with someone who has hand, foot and mouth disease do not need to be excluded from work however consideration should be given to those who care for patients who are more susceptible to infection. |
| Herpes Simplex (cold sores) | Must not provide direct care to neonates, newborns, patients in delivery suites, severely immunocompromised patients, burns patients, patients with extensive eczema, or patients in operating room if there is an exposed herpetic lesion. May provide direct patient care to other patients and do not need to wear a mask. However, sores should be covered with a dressing where possible, and hygiene practices to minimise the risk of transmission need to be maintained. |
| Herpes Zoster (Shingles) | Must not provide ANY direct patient care if lesions cannot be covered (e.g. ophthalmic zoster). If active lesions can be covered, can provide care to all patients except for pregnant women, neonates, severely immunocompromised patients, burns patients and patients with extensive eczema. |
| Norovirus | Must not come to work for at least 48 hours after symptoms have stopped (e.g. diarrhoea and/or vomiting) (see practice statement 41). |
| Pertussis (Whooping Cough) | Remain away from work until at least 5 days after commencement of appropriate antibiotic therapy; or for 21 days after the onset of symptoms if not receiving antibiotic treatment; or 14 days after the onset of paroxysmal cough (if the onset is known). |
| Scabies and Lice | Healthcare workers should remain off work until 24 hours after first treatment started. |
| Staphylococcal infection | Any staphylococcal lesions (e.g. boils, wound infections) must be covered with an occlusive dressing while at work. If lesions cannot be covered, must not perform patient care or prepare hospital food until they have received appropriate antibiotic therapy and the infection has resolved. |
| Streptococcal infection | Any healthcare worker with streptococcal lesions (e.g. impetigo, streptococcal tonsillitis) must ensure that lesions are covered with an occlusive dressing while at work. If lesions cannot be covered, healthcare workers must not provide direct patient care nor prepare hospital food until 24 hours after commencement of appropriate antibiotic therapy. Healthcare workers with pharyngitis/tonsillitis should avoid patient contact for at least 24 hours after starting appropriate antibiotic therapy. |
| Tuberculosis (TB) | If TB disease is suspected or is present, TB Services are to be notified of the staff, and the staff treated. Any personnel with pulmonary TB is to be excluded from the workplace until cleared by TB Services. Any active TB must be monitored by TB Services. |
| Viral rashes | Before starting employment, personnel should be screened by completing a pre-employment health assessment for measles, mumps, rubella and varicella. Non immune healthcare workers should be offered vaccination unless contraindicated. Measles (rubeola) —If suspected, must remain off of work until appropriate test results are known. May return to work if they have serological evidence of immunity (i.e. are IgG sero-positive and IgM sero-negative); but must be excluded until 4 days after the appearance of the rash if they develop measles. Mumps —If suspected, must remain off work until appropriate test results are known. May return to work if they have serological evidence of immunity (i.e. are IgG sero-positive and IgM sero-negative). If mumps develop, they must be excluded from work for 9 days after the onset of parotid gland swelling or until the swelling goes down. Rubella (German Measles) —If suspected, must remain off of work until appropriate test results are known. May return to work if they have serological evidence of immunity (i.e. are IgG sero-positive and IgM sero-negative). If they develop Rubella, they must be excluded for at least 4 days after the appearance of the rash. Chickenpox (Varicella) — if healthcare worker develops Varicella, they must be excluded until all blisters have dried (this usually takes at least 5 days). Human Parvovirus B19 (Slapped Face) —does not require exclusion from work, non-infectious once rash develops. |
| Viral respiratory tract infections including influenza, SARS- | Must not come to work while symptomatic. Refer to relevant national, state or territory guidelines for further detail for specific infectious agents. |

CoV-2 and other viruses (e.g. common colds)

*Includes Giardiasis, Shigella infection, Salmonella infection, Campylobacter infection.

Source: Adapted from *NHMRC Staying Healthy: Preventing infectious diseases in early childhood education and care services (Fifth Edition)* [360] and *CDNA National Guidelines For Public Health Units: Influenza Infection* [362] & *Pertussis* [363] and *CDC Guidelines and Recommendations: Prevention Strategies for Seasonal Influenza in Healthcare Settings* [364] and <https://www.health.gov.au/news/ahppc-statement-on-testing-tracing-isolating-and-quarantining-in-high-levels-of-covid-19-community-transmission> May 23 2022

Practice Statement

Updated

41. It is good practice for healthcare workers and visitors to adhere to norovirus exclusion periods.

Healthcare workers should not be at work from symptom onset until 48 hours after symptom resolution. On returning to the healthcare facility, healthcare workers should adhere to appropriate hand hygiene practices.

Practical info

Norovirus infections generally have a shorter incubation period and are characterised by acute onset of nausea, vomiting, abdominal pain and diarrhoea [96]. The mean duration of symptoms is two-three days.

Maximum viral shedding occurs between 24 and 48 hours after exposure, so it is good practice for healthcare workers to be excluded for 48 hours after the resolution of symptoms. Clearance of norovirus is not practical or beneficial as viral excretion can persist for multiple days.

There is evidence to suggest that shedding of norovirus may continue to occur for more than 21 days after the resolution of symptoms. However, no data has been reported on ongoing transmission or secondary cases.

Norovirus epidemiology

Norovirus is the most frequently occurring cause of community-acquired acute gastroenteritis in people of all ages. It is one of the most common causes of outbreaks in healthcare settings, affecting both long-term care facilities and acute care hospitals [115].

Norovirus belongs to the family *Caliciviridae* and are a single-stranded RNA, non-enveloped virus that can cause gastroenteritis in humans [118]. Noroviruses are divided into at least six genogroups (GI-GVI) and further subdivided into more than 38 genotypes based on phylogenetic analysis of the major capsid protein [118] [122]. Currently, human noroviruses belong to one of three norovirus genogroups which are further divided into more than 25 genetic clusters [119].

Human noroviruses cannot be grown in cell culture [123], therefore, diagnostic methods focus on detecting viral RNA or antigen.

Transmission pathways

Transmission for norovirus infections in healthcare settings mainly occur by the faecal–oral route, either through person to person contact or through exposure to contaminated food. Whilst some observational studies have suggested there is a possibility of viral transmission via aerosols, there is no data or determination criteria to support this assumption. It appears that Genotype GII.4 is more often associated with transmission mediated by person-to-person contact than with other types of transmission.

Individuals may shed norovirus more than 21 days after the resolution of symptoms, possibly acting as a possible source for nosocomial transmission. However no data has been reported on ongoing transmission or secondary cases.

Evidence to decision

| | |
|---|--|
| Benefits and harms | The benefits of adhering to norovirus exclusion periods includes reduced risk of transmission of the virus to other healthcare workers and to patients. |
| Certainty of the evidence | <p>The Communicable Diseases Network Australia Guidelines recommend that workers be excluded for 48 hours after symptoms have stopped, and the Centres for Disease Control and Prevention Guidelines provide evidence that ill personnel should be excluded for a minimum of 48 hours after symptom resolution.</p> <p>Evidence from observational studies (n=4) also shows that individuals may shed norovirus for more than 21 days after symptoms resolve, which may act as a source of nosocomial transmission [96].</p> |
| Values and preferences | It is expected that all patients and staff of healthcare facilities would highly value the implementation of exclusion periods for healthcare workers infected with norovirus. |
| Resources and other considerations | The exclusion of healthcare workers may impact upon resourcing and staffing. |

Rationale

Maximum viral shedding of norovirus is thought to occur 24-48 hours after exposure.

There is evidence suggesting shedding of Norovirus for more than 21 days, possibly being a source for nosocomial transmission. However there is no evidence to suggest that ongoing shedding equates to transmission or secondary cases. Despite the virus being present, provided the infected person can maintain appropriate hygiene practices, there is no reason for exclusion once the physical symptoms have resolved. However, it would be good practice to ensure that health professionals who may be shedding are not caring for high risk populations e.g. immunosuppressed.

4.2.3 Managing exposures to occupational hazards

Exposures that might place a healthcare worker at risk of hepatitis B virus, hepatitis C virus, HIV or human T-cell lymphotropic virus type I (HTLV-I) are percutaneous injury (e.g. needlestick or cut with sharp object) or contact of a mucous membrane or non-intact skin (e.g. exposed skin that is chapped, abraded, or affected by dermatitis) with blood, tissue or other potentially infectious body substances.

Each healthcare facility requires a policy on the management of needlestick injuries, and on providing immediate post-exposure advice for sharps injuries and other blood or body substance incidents involving healthcare workers. This is because generic policies may not be relevant to individual settings (e.g. access to care, especially after hours).

Managing exposures

Some general components relevant to all occupational exposures to blood-borne viruses include [366]:

- The healthcare worker should receive immediate care and treatment at the site of exposure.
- A risk assessment of the exposure should be taken—including the type of exposure, type and amount of fluid involved, infectious status of the source, and susceptibility of the exposed healthcare worker.
- If the source of exposure can be identified, they should be tested for HBV surface antigen, HCV antibody and HIV antibody.
- The healthcare worker should have baseline testing, as required.
- Counselling and follow-up should be provided to the healthcare worker.

Treatment protocols include removal of contaminated clothing, thorough washing of the injured area with soap and water; and flushing of affected mucous membranes with large amounts of water.

Post-exposure prophylaxis

Post-exposure prophylaxis (PEP) is the medical response given to prevent the transmission of blood borne pathogens following a potential exposure to HIV. The decision to prescribe PEP should be made on a case-by-case basis and include consideration of

the need for first aid, counselling including the assessment of risk of exposure to the infection, testing, and depending on the outcome of the exposure assessment, the prescription of antiretroviral drugs, with appropriate support and follow-up [365].

When PEP is recommended, it should be prescribed and started as close to the time of exposure as possible, and within 72 hours. Eligibility for PEP and the type of regime prescribed is individualised and determined by a number of factors, including the transmission risk associated with the exposure [365].

A 28-day course of PEP is recommended.

Specific guidance on PEP can be found in the Australian Society of HIV Medicine Guidelines (second edition) [365].

Management of possible exposure to other conditions [365]

- **Hepatitis B**—Healthcare workers with evidence of previous immunity to hepatitis B require no follow up. Non-immune individuals require immunisation and follow up.
- **Hepatitis C**—Healthcare workers potentially at risk require baseline and follow-up testing for hepatitis C. Patients should be informed about the symptoms of hepatitis C and advised to seek medical advice if any are displayed.
- **Tetanus**—Tetanus status should be assessed for any healthcare workers who sustain abrasions or wounds.

Consult the current edition of the *Australian Immunisation Handbook* [353] for further advice and guidance.

4.2.4 Healthcare workers with specific circumstances

Healthcare workers with specific circumstances

Healthcare facilities need to assist healthcare workers who experience circumstances that place them at greater risk of infection to develop management plans that ensure their well-being.

Where a healthcare worker is known to be particularly susceptible to healthcare associated infections, work duties are assessed to ensure that the welfare of that person, patients and other healthcare workers is safeguarded. This may involve appropriate work placements, adjustments or restrictions, or deployment to a role involving less risk. Healthcare workers in this situation may require counselling on what tasks they can perform, what they should avoid and the possible impact of their work on their health.

Pregnant healthcare workers

Employers should provide information on the risks associated with pregnancy and should assist pregnant healthcare workers to avoid infectious circumstances that may present a risk to her or the baby. It is the responsibility of pregnant healthcare workers to advise their doctor and employer of their pregnancy; this information must remain confidential.

All pregnant healthcare workers should adhere to standard and transmission-based precautions and ensure that they are appropriately vaccinated. However, pregnant healthcare workers should be given the opportunity to avoid patients with specific infections.

For more information, refer to the *Australian Immunisation Handbook*.

Immunocompromised healthcare workers

Healthcare workers with immune deficiencies are more at risk of acquiring infections. The type of employment they can undertake should include only duties that will minimise their exposure to infections. Predisposing conditions include neutropenia, disseminated malignancy and infections that produce immunodeficiency (e.g. HIV).

Refer to the *Australian Immunisation Handbook* for guidance on the immunisation of immunocompromised healthcare workers.

Healthcare workers with skin conditions

Skin integrity is the ultimate barrier to transmission of infectious agents. When staff members have damaged skin or weeping skin conditions (e.g. allergic eczema, psoriasis, exfoliating dermatitis), they may be readily colonised by healthcare associated microorganisms and may become a vehicle for disseminating these organisms. Healthcare workers in this situation should be identified by personal history screening when they start employment, and need to be informed of the risks they may pose to patients. Any damaged skin must be appropriately covered before healthcare workers carry out procedures. Consideration must be given to providing these staff members with appropriate, individual PPE such as specific types of gloves, hand hygiene products and moisturising lotion.

Healthcare workers with cystic fibrosis

Healthcare workers with cystic fibrosis may be at greater risk of cross-infection than healthcare workers without cystic fibrosis. They may also pose an infection risk to patients; however, this will vary according to the severity of their disease, the frequency of coughing and the type of cystic fibrosis pathogens evident [369].

It is recommended that healthcare workers with cystic fibrosis do not work with patients or other healthcare workers with cystic fibrosis. Healthcare workers are encouraged to disclose their diagnosis during pre-employment screening to determine the safest workplace arrangements.

Healthcare workers living with a blood-borne virus (BBV)

Healthcare workers living with a BBV, including hepatitis B, hepatitis C and HIV, must be under the care of a treating doctor and must be tested for the respective BBV viral load levels, as well as for other BBVs, in accordance with the *Australian National Guidelines for the Management of Health Care Workers known to be infected with blood-borne viruses 2018* [355].

Healthcare workers living with a BBV deserve a supportive work environment, including retraining if required, counselling and appropriate infection control measures [355].

4.2.5 Exposure-prone procedures

Non-exposure prone procedures (non-EPPs) are procedures where the hands and fingers of the healthcare worker are visible and outside of the body at all times and procedures or internal examinations that do not involve possible injury to the healthcare worker's hands by sharp instruments and/or tissues, provided routine infection prevention and control procedures are adhered to at all times e.g. routine oral examination with appropriate personal protective equipment, insertion and maintenance of intravenous or central lines [355].

Exposure prone procedures (EPPs) are invasive procedures where there is potential for direct contact between the skin, usually finger or thumb of the healthcare worker, and sharp objects or surgical instruments—such as needles, sharp body parts (e.g. fractured bones), spicules of bone or teeth—in body cavities or in poorly visualised or confined body sites, including the mouth of the patient. During EPPs there is an increased risk of transmitting blood borne viruses (BBVs) between healthcare workers and patients.

There are two major risks related to healthcare workers that arise out of EPPs:

1. Healthcare workers can become infected with a BBV.
2. Healthcare workers who already have a BBV may transmit the virus to the patient.

Some procedures that are generally considered not to be EPP may have the potential to escalate to EPPs. These procedures include:

- *Minimally invasive procedures* including laparoscopy, endovascular procedures, thoracoscopic procedures, Natural Orifice Transluminal Endoscopic Surgery, cystoscopic procedures, arthroscopic procedures and robotic surgery.
- *Trauma/emergency situations* where a previously non-EPP may escalate into an EPP.

Table 33 provides advice on EPPs in specific areas of clinical care as well as general procedures that are not considered to be EPPs.

Table 33. EPPs and non-EPPs in specific areas of clinical care

| Area of Clinical Care | Exposure Prone Procedure | Non-Exposure Prone Procedure |
|-----------------------|--------------------------|---|
| General | | <ul style="list-style-type: none"> • Routine non trauma related vaginal or rectal examination in the absence of a sharp. • Insertion and maintenance of arterial or intravenous cannulae whether inserted centrally or peripherally. • Open incision and drainage of superficial abscesses or haematomas. • Percutaneous drainage of abscesses, fluid collections or hematomas under radiation or ultrasound guidance. • Suturing of uncomplicated skin lacerations. |

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| Cardiothoracic | <ul style="list-style-type: none"> • Generally all cardiothoracic procedures. | |
| Dentistry | <ul style="list-style-type: none"> • All maxillofacial surgery. • All oral surgical procedures. • The extraction of teeth (with some exceptions). • Periodontal surgical procedures. • Endodontic surgical procedures. • Implant surgical procedures. | <ul style="list-style-type: none"> • Extraction of highly mobile or exfoliating teeth. • Assessment and management of removable dentures and mouthguards. • Taking impressions of teeth. • Apply decay preventive agents. • Removing dental plaque, calculus and stains. |
| Emergency/trauma | <ul style="list-style-type: none"> • Open head injuries resulting from trauma. • Insertion of intercostal catheter, where the procedure requires insertion of the finger into the pleural cavity in a trauma situation. • Reduction of facial or jaw fractures from within the oral cavity. • Rectal or vaginal examination in the presence of suspected pelvic trauma. • Placement of Thoracic Aortic clamp, packing a deep wound in a body cavity, or deep suturing to arrest haemorrhage. • Internal cardiac massage. | <ul style="list-style-type: none"> • Percutaneous insertion of intercostal catheter (e.g. via Seldinger technique), where the procedure does not require insertion of the finger into the pleural cavity). • Insertion of intercostal catheter, where the procedure requires insertion of the finger into the pleural cavity in a non-trauma situation. • Endotracheal intubation. • Bag-valve-mask ventilation. • Simple suturing under direct vision. |
| General surgery | <ul style="list-style-type: none"> • Open abdominal or thoracic procedures. | <ul style="list-style-type: none"> • Excision of skin lesions. • Breast surgery, where hands remain in view. |
| Gynaecology | <ul style="list-style-type: none"> • Perineal surgery. • Trans-vaginal surgery. • Open abdominal gynaecological surgery. • Local anaesthetic administered to the cervix other than under direct vision (i.e. with fingers concealed in the vagina). | <ul style="list-style-type: none"> • Vaginal examination in absence of a sharp. • Laparoscopy. • Colposcopy. • Surgical insertion of depot contraceptive implant/ devices. • Fitting intrauterine contraceptive devices (coils). • Cone biopsy. • Dilatation and curettage (D&C). |
| Neurosurgery | <ul style="list-style-type: none"> • Any surgical procedures that involve exposure to sharp bone fragments (e.g. trauma and some spinal surgery). | |
| Obstetric or midwifery | <ul style="list-style-type: none"> • Caesarean birth. • Instrumental birth. • Infiltration of the perineum with local anaesthetic. • Episiotomy. • Repair of an episiotomy or perineal/ vaginal tear. • Application of fetal scalp electrodes. • Fetal blood sampling. | <ul style="list-style-type: none"> • Vaginal examination, in absence of a sharp. • Vaginal egg collection provided fingers remain visible at all times when sharp instruments are in use. • Suction termination of pregnancy. |
| Ophthalmology | <ul style="list-style-type: none"> • Orbital surgery. • Oculoplastic and lacrimal surgery where bony reconstruction and bone fragments are involved. | <ul style="list-style-type: none"> • Routine ocular surgery. |
| Orthopaedic | <ul style="list-style-type: none"> • Cutting or fixation of bones or the distant transfer of tissues from a second site (such as in a thumb reconstruction). | <ul style="list-style-type: none"> • Closed fracture reduction. • Diagnostic arthroscopy. • Endoscopic carpal tunnel decompression. |

| | | |
|--------------------------------------|---|---|
| | <ul style="list-style-type: none"> • Open procedures where there is the possibility of: <ul style="list-style-type: none"> ◦ bone fragments and/or bone spicules ◦ mechanical drilling involved ◦ deep tunnelling using sharp instruments. | |
| Otolaryngology, head and neck | <ul style="list-style-type: none"> • Bony facial reconstructive surgery (elective or after trauma). | <ul style="list-style-type: none"> • Otological procedures e.g. stapedectomy/ stapedotomy, insertion of ventilation tubes, insertion of a titanium screw for a bone anchored hearing aid. • Most head and neck cancer operations except where fingers are not visible at all times. • Most rhinological procedures. • Functional endoscopic sinus surgery (FESS). |
| Paediatric surgery | <ul style="list-style-type: none"> • Extensive cosmetic procedures that involve bony reconstruction. • Free tissue transfer involving bone or in the thorax. | <ul style="list-style-type: none"> • Herniorrhaphy. • Orchidopexy. • Superficial procedures. |
| Plastic surgery | <ul style="list-style-type: none"> • Extensive cosmetic procedures that involve bony reconstruction. • Free tissue transfer involving bone or in the thorax. | <ul style="list-style-type: none"> • Excision of superficial lesions. • Superficial skin excision and reconstruction. |
| Podiatry | <ul style="list-style-type: none"> • Procedures undertaken by podiatric surgeons including open surgical procedures on bones and soft tissue of the foot and lower leg. • See also Orthopaedics. | <ul style="list-style-type: none"> • Routine procedures undertaken by podiatrists (including nail avulsion performed in the clinic setting). |
| Urology | <ul style="list-style-type: none"> • Open urological procedures. | <ul style="list-style-type: none"> • Image guided biopsies. • Scrotal procedures. |
| Vascular surgery | <ul style="list-style-type: none"> • Open abdominal or thoracic vascular surgery. | <ul style="list-style-type: none"> • Carotid endarterectomy. • Percutaneous dilatation, stenting or recanalisation of arteries. • Percutaneous treatment of varicose veins. • Diagnostic angiography. • Peripheral embolectomy/ thrombectomy. |

Adapted from the *Australian National Guidelines for the Management of Healthcare Workers Living with Blood Borne Viruses and Healthcare Workers who Perform Exposure Prone Procedures at Risk of Exposure to Blood Borne Viruses (2018)* [355].

National Guidelines

The *Australian National Guidelines for the Management of Healthcare Workers Living with Blood Borne Viruses and Healthcare Workers who Perform Exposure Prone Procedures at Risk of Exposure to Blood Borne Viruses (2018)* (National Guidelines) [355] provide detailed information about the required BBV testing for healthcare workers, and the criteria to be met that allow a healthcare worker with a BBV to return to work. The advice in the National Guidelines reflects the effectiveness of antiviral treatment for hepatitis B virus (HBV), hepatitis C virus (HCV) and HIV. The National Guidelines allow healthcare workers living with a BBV to perform EPPs, provided they comply with the National Guidelines [355].

Responsibilities of healthcare workers

As stated in the National Guidelines, all healthcare workers (including students) who are performing EPPs are required to take reasonable steps to know their BBV status and should be tested for BBVs at least once every three years and are required to have appropriate and timely testing after potential BBV exposures, both occupational and non-occupational [355].

Healthcare workers living with a BBV who are performing EPPs must be under the care of a medical practitioner with relevant expertise, and are expected to comply with the requirements stated in the National Guidelines and relevant legislation in their jurisdiction [355].

Responsibilities of employers

The National Guidelines do not prescribe additional responsibilities on employers of healthcare workers or students living with a BBV. However, employers should support employees who perform EPPs with access to appropriate information, testing, training, counselling and vaccination programs [355].

Healthcare facilities should aim to achieve voluntary compliance and self-disclosure by providing an environment in which healthcare workers know their confidentiality will be maintained. Employers of healthcare workers must consider the relevant public health, antidiscrimination, privacy, industrial relations and equal employment opportunity legislation in their jurisdiction [355].

Depending upon the jurisdiction and/or employment arrangements, employers may need to request evidence of viral clearance from the treating medical practitioner [355]. Similarly, the protection of public health may require that health monitoring information (including viral load and relevant clinical information) may need to be released to a designated person in the workplace in the event of a potential exposure incident to assess the requirement for further public health action [355].

Responsibilities of treating doctors

The treating medical practitioner may be required to inform the appropriate registration board and the relevant area of the jurisdictional health department if the healthcare worker is not in compliance with the National Guidelines [355].

A summary of key requirements is provided below:

| |
|---|
| All healthcare workers and healthcare students [355]: |
| All healthcare workers should be encouraged to undertake regular testing for BBVs. |
| All healthcare workers have the right to access confidential testing, counselling, support and treatment. |
| All healthcare workers should be vaccinated against hepatitis B virus. |

| |
|---|
| Healthcare workers who perform EPPs [355]: |
| Healthcare workers who undertake EPPs must take reasonable steps to know their BBV status and should be tested for BBVs at least once every three years. |
| All registered healthcare workers who undertake EPPs must declare when applying for renewal of registration that they are complying with, and have been tested in accordance with the National Guidelines [355]. |
| All Healthcare workers who undertake EPPs should understand their obligation to report their BBVs status, if required, under jurisdictional legislation and/or policies. |
| Healthcare workers should understand their obligation to report all sharps injuries, whether or not there was a risk of patient exposure. |
| Healthcare workers must also be tested for BBVs after the occurrence of any potential occupational exposure incident. In addition, healthcare workers who are exposed to risks for BBV transmission in non-occupational settings should be aware of national recommendations for testing frequencies that sit outside of the National Guidelines. |
| Healthcare workers must cease performing all EPPs if diagnosed with a BBV until the criteria in the National Guidelines are met. |

| |
|---|
| Healthcare workers living with a BBV [355]: |
| All healthcare workers with a BBV must be under the care of a treating doctor with relevant expertise. |
| All healthcare workers living with one or more BBVs must be tested for the respective BBV viral load levels, in accordance with the National Guidelines [355]. |
| Healthcare workers living with one or more BBVs are permitted to perform EPPs providing the criteria set out in the National Guidelines and specific viral load criteria are met. |

For further information on EPPs and BBVs, including compliance requirements and occupational exposure, see *the National Guidelines [355]*.

4.3 Education and training

Summary

- Education and training underpin efforts to integrate infection prevention and control practices into practice at all levels of every healthcare facility.
- Essential education for all healthcare workers should cover infection prevention and control work practices and their role in preventing the spread of infection. This should be a part of undergraduate education, staff orientation and any continuing

professional development.

- Specific postgraduate education of infection control professionals is strongly recommended.
- Engaging patients, their carers and families in their own healthcare is integral to effective infection prevention and control. All healthcare workers should be informed about the rights and responsibilities of patients and learn how to apply this understanding in the way that they deliver care.

Practice Statement

Updated

42. It is good practice for infection control professionals to partake in ongoing professional development in order to gain the necessary expertise to fulfil their role. Infection prevention and control staff at all levels should be supported to access formal and informal education and training relevant to their role.

Practical info

Postgraduate education

While some states in Australia have requirements for practising as an infection control professional, there is currently no minimum or standardised educational requirement to practice as an infection control professional, or to coordinate an organisational infection prevention and control program. A range of postgraduate education programs are currently available for nurses seeking or establishing a career in infection control in Australia, although the content of these courses is variable. These courses include Graduate Certificates and Masters degrees in Infection Prevention and Control.

Compliance and credentialing

The Australasian College for Infection Prevention and Control is the peak body for infection prevention and control professionals in the Australasian region. The College was formed in 2012 and brought together the state and territory infection control associations formerly in the Australian Infection Control Association. The College recommends that healthcare workers apply for credentialing in infection prevention and control. This is a self-regulatory process to determine and acknowledge that an individual has demonstrated the prescribed competence of the relevant specialist role.

Mentoring, support and networking

While there are no formal mentoring programs in place, many infection control professionals provide mentoring to less experienced staff. Mentoring requires the support of health facility administrators, so that it is recognised as being part of healthcare worker core time, but additional to their workload.

Some healthcare facilities have successfully established linked-nurse programs to support their infection prevention and control programs and to facilitate mentoring of healthcare professionals at varying levels of expertise in infection prevention and control.

Auditing

Auditing of healthcare worker behaviour is important for surveillance and accreditation, and to reinforce positive signs of culture change within the facility. Auditing to measure compliance with infection prevention and control policies and procedures can occur through:

- direct observation
- examining logs and registers of specific activities (e.g. sterilisers)
- monitoring use of personal protective equipment or hand hygiene products.

Timely feedback is a critical aspect of auditing. In acute-care settings, measurement and feedback generally occurs at ward level.

Evidence to decision

Benefits and harms The benefits of having educated infection control professionals clearly outweighs the harms.

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|----------------------------------|--|
| Certainty of the evidence | This practice is supported by suggestive clinical or epidemiologic studies, and based on sound theoretical principles. |
|----------------------------------|--|

| | |
|-------------------------------|---|
| Values and preferences | It is expected that all patients and staff in healthcare facilities would value effective training and education for healthcare workers as part of a multimodal strategy to reduce healthcare associated infection. |
|-------------------------------|---|

| | |
|---|---|
| Resources and other considerations | There are resources and costs associated with professional development for healthcare professionals, however the benefits outweigh the costs. |
|---|---|

Rationale

This advice is based on limited empirical evidence, but on sound theoretical principles and supported by expert advice. Providing infection control professionals with appropriate formal and informal learning is justified to reduce healthcare associated infection.

4.3.1 Education strategies

On-the-job training

All healthcare workers need to understand the basis and importance of infection prevention and control. This information should be provided to healthcare workers and students by the healthcare facility where they work. The basis for this teaching should be the information contained in these Guidelines (see Section 1: Introduction for a summary), ensuring that healthcare workers understand standard and transmission based precautions. This information should be tailored specifically to the healthcare facility, and where necessary, to a healthcare workers specific role in the workplace.

Job-specific training should be provided as: part of orientation, when new procedures affect the employee's occupational exposure, and before rostering to a hazardous area. Healthcare workers competency should be assessed and records should be maintained of their participation in education programs.

University education

Up-to-date information on infection prevention and control basics, policy, procedures, quality assurance and incident monitoring should be included in the curriculum of all undergraduate and postgraduate courses in health-related areas.

Universities and training colleges also have an obligation to inform prospective students about the impact that particular infections may have on their ability to complete the course and engage in the full spectrum of clinical practice after graduation (see Section 4.2, in particular Section 4.2.2). This information should include advice about specific measures, including immunisation, that reduce the risk of acquiring infection.

Effective educational strategies

There are a variety of educational strategies that can be used in healthcare settings to improve knowledge and understanding of infection prevention and control.

These include:

- Multifaceted strategies, which consider the needs of the target group, potential barriers and facilitators and the context in which educational strategies are applied. These are likely to be more effective than single strategies as learning is reinforced through repetition and variation.
- Active educational interventions that are repeated with some frequency have a greater chance of changing behaviour than a single, didactic session. Repetition and interactivity have both been shown to be important factors in achieving behaviour change that is sustained.
- The use of multiple forms of media including printed materials and videos.
- Educational outreach visits have been found to be an effective method, especially when combined with other strategies such as interactive education and printed materials, but are costly to implement. They seem to be most effective when related to prescribing practices of moderate complexity.

Education activities can be integrated into staff orientation programs, credentialing packages, annual training and competency

testing, implementation of policy and procedure manuals, and in decision support tools available on the facility intranet. The infection control professionals' contact details should be readily available to all staff and included in all resources.

E-learning (e.g. interactive web-based training) is being used in some states and territories, and may be a useful addition to other education strategies.

Local programs for education, audit and feedback should be refined regularly and promoted widely in healthcare facilities by senior staff members [274].

4.4 Healthcare associated infection surveillance

Summary

- Appropriate surveillance can substantially reduce healthcare associated infections, morbidity and mortality.
- Both outcome and process measures are used for surveillance in large health facilities; process measures alone can provide a useful alternative, particularly in smaller facilities.
- Timely targeted feedback is critical for effective surveillance.
- All staff involved in surveillance should be appropriately trained in data collection techniques.

4.4.1 Role of surveillance in reducing healthcare associated infection

Many infections can be prevented using approaches based on quality and safety theories such as:

- quality improvement methodologies
- creating a safety culture (individuals taking responsibility for ensuring safety and quality of themselves and others)
- application of systems thinking (i.e. understanding the factors in the system that allow errors to occur).

To be successful, all these approaches need to be based on comprehensive information obtained through surveillance—the ongoing, systematic collection, analysis, interpretation, and dissemination of data regarding a health-related event, for use in public health action to reduce morbidity, mortality and to improve health.

Surveillance is important for wider systems of quality management, but the main purpose of collecting reliable data are to improve quality within a service or facility. Collecting such data can provide the impetus for change and make it possible to evaluate the effectiveness of an intervention [377] [376]. For example, monitoring both hand hygiene compliance and the rate of bloodstream infections, and disseminating this information within the facility, can improve hand hygiene practices.

Surveillance of healthcare associated infections (HAI) draws information about the agent, host, environment and risk factors from a number of data sources and:

- provides baseline information on the frequency and type of HAI
- enables breakdowns in infection prevention and control to be identified
- allows for timely investigation and appropriate infection prevention and control measures to be instituted.

All healthcare facilities, including small acute-care facilities and office practices, should collect data on HAIs, infection prevention and control breaches, outbreaks of infectious disease and antibiotic resistance. Post-discharge surveillance by community-based healthcare practices should also be considered. The surveillance system used by a healthcare facility depends on the type and size of the facility, its case mix, and the resources available.

It is important that the collection of surveillance data are guided by a clear purpose that is understood by all relevant stakeholders [375].

A recent study identified that only half of those involved in collecting and analysing HAI surveillance data had been trained, which subsequently led to poor agreement when identifying HAIs [374]. To drive the success of surveillance, relevant staff members

need appropriate training to ensure consistency in the application of definitions and data collection [375].

4.4.2 Types of surveillance programs

It is not feasible to conduct facility-wide surveillance for all events; therefore, surveillance is often targeted with a focus on specific events, processes, organisms, medical devices or high-risk patient populations. Healthcare associated infections surveillance programs may focus on:

- specific sites of infection (e.g. bloodstream, surgical sites)
- specific populations (e.g. neonates, healthcare worker occupational exposure to blood and body substances)
- specific organisms or types of organisms (e.g. multi-resistant organism [MRO], *C. difficile*, respiratory syncytial virus [RSV], rotavirus)
- specific locations in the healthcare facility or community (e.g. intensive care unit, neonatal intensive care unit, long-term care facility).

For further information and nationally-consistent definitions, see the Australian Commission on Safety and Quality in Health Care's National Surveillance Initiative website —<https://www.safetyandquality.gov.au/our-work/healthcare-associated-infection/national-hai-surveillance-initiative/>.

There are two main methods of surveillance —process and outcome. Process measurements are usually easier to measure, less ambiguous and more widely applicable than outcome indicators. Process surveillance may be an adjunct to outcome surveillance; alternatively, it can entirely replace outcome surveillance for practices or locations that have too few adverse outcomes for statistical analysis (e.g. small facilities where the number of patients at risk of infection may be too small to calculate valid infection rates).

Process surveillance

Process surveillance involves auditing practice against a certain standard, guideline or policy [380].

As no single intervention will prevent every healthcare associated infection, packages of evidence-based interventions have been developed and are increasingly being used in process surveillance (e.g. care bundles, see also Sections 3.5.1 and 4.1.5).

Process measures that are linked by evidence to important outcomes:

- do not require risk adjustment
- can predict outcomes
- can easily be acted on because potential improvements are usually the responsibility of the clinical service
- can be captured quickly
- are sensitive because many episodes of inappropriate care do not cause harm.

Examples of published process indicators of high value include:

- aseptic insertion and management of peripheral or central intravascular devices
- healthcare workers' compliance with hand hygiene and the techniques they used
- perioperative and intraoperative practice such as antibiotic prophylaxis, normothermia, normoglycaemia and appropriate hair removal
- healthcare workers' uptake of immunisation.

Outcome surveillance

Outcome surveillance involves measuring adverse events, a proportion of which are preventable. The sensitivity and specificity of event definitions and the reliability of data collection need to be considered when developing methods to detect adverse events. It is important to create a balance between avoiding false positives (specificity) and picking up true positives (sensitivity), given that true positives are rare events in the overall patient population.

Certain outcome measures — for example, the incidence of healthcare associated methicillin-resistant *Staphylococcus aureus* (MRSA) bacteraemia — appear to be reliable and have driven practice change, leading to significant improvements in patient safety.

Australia currently has no system-wide approach to measurement of patient mortality caused by or associated with healthcare associated infection (HAI). These deaths are unlikely to be reported using existing mechanisms such as adverse event reporting

systems. Mortality from infection may be seen as ‘anticipated’ even though the occurrence of the infection that led to the death was unanticipated.

A further challenge in measuring patient deaths is differentiating between patients who die with a HAI and those who die from a HAI or suffer serious injury due to a HAI (i.e. attributable injury or death). One new approach is to evaluate such patient deaths to determine whether mortality was unexpected, and then analyse the contributing factors to determine preventable root causes that might be modified in future. In this approach, infection events (usually deaths or BSI) are considered and investigated individually. Although mandated by the UK’s National Health Service, evidence of the value of this approach is lacking.

Critical incidents

If there has been a breakdown in an infection prevention and control procedure or protocol, a ‘lookback’ investigation may be necessary to identify, trace, recall, counsel and test patients or healthcare workers who may have been exposed to an infection, usually a blood borne virus.

Lookback investigations must be managed with due regard to ethical and legal considerations. In the event of such an incident (e.g. failure of sterilisation or disinfection), the local public health unit should be advised immediately.

Monitoring of critical incidents and other sentinel events is an important part of surveillance. Root cause analysis of sentinel events is a structured process for identifying the process and contributing factors, exploring and identifying risk reduction strategies and implementing solutions.

4.4.3 Data collection and management

Surveillance involves:

- defining surveyed events precisely
- systematic collection and validation of data
- analysis and interpretation
- communication of findings to relevant people.

The following epidemiologic principles should be applied during healthcare associated infection (HAI) surveillance:

- Use standardised definitions of infection.
- Use laboratory-based data (when available).
- Collect epidemiologically important variables (e.g. clinical service in hospitals and other large facilities, population-specific risk factors, underlying conditions that predispose to serious adverse outcomes).
- Analyse data to identify trends that may indicate increased rates of transmission.
- Feedback information on trends in the incidence and prevalence of HAI, probable risk factors and prevention strategies and their impact, to the appropriate healthcare workers, administrators, and as required by local and state/territory health authorities.

Surveillance data for quality improvement must be of high quality. The characteristics that qualify data as evidence for action include [378]:

- Representativeness—the data fairly represent the thing measured.
- Accuracy—the data reflect what is intended to be measured.
- Precision—the data and the target of measurement correspond closely.
- Authoritativeness—the data are appropriate for drawing a meaningful conclusion.
- Clarity—the data are presented in a form that the target audience can understand.

Data of this nature are more likely to arise from surveillance processes:

- that involve all stakeholders in design and implementation
- for which there are agreed organisational objectives, and processes that are relevant to the population served
- that use trained staff to collect and manage data, and that provide them with appropriate information technology support
- that use definitions of surveillance events that are unambiguous, practical, specific and can be validated
- that have reliable and practical methods for detecting events
- for which the processes that determine an outcome are thoroughly understood
- for which appropriate denominators are collected for risk adjustment
- for which reporting links measurement to prevention efforts, and meets the needs of both clinicians and managers.

For further information on surveillance, refer to the Australian Commission on Safety and Quality in Health Care's National Healthcare Associated Infection Surveillance initiative [382].

4.4.4 Outbreak surveillance

An outbreak may be defined as the occurrence of infections at a rate greater than that expected within a specific geographical area and over a defined period of time.

Ideally, surveillance systems should facilitate the early detection of outbreaks. Increasingly, microbiological data are being relied on for this purpose, although outbreaks may be detected using other sources such as pharmacy records.

In some instances, the occurrence of an outbreak is obvious, such as in an episode of food poisoning that affects both healthcare workers and patients. It is more usual, however, for the outbreak to have an insidious onset that is not immediately apparent. When an outbreak is detected, the infection prevention and control committee should be informed and an outbreak team formed. Depending on the size and severity of the outbreak, it may be necessary to involve occupational health and safety staff, facility administrators, engineers and public health officials. Details on the steps involved in the management of an outbreak are provided in Section 3.4.2.

Legislation requires that the relevant public health authority be informed of outbreaks related to notifiable infections. It may also be prudent to involve public health officers at an early stage if an outbreak is likely to come to the attention of the media.

4.4.5 Disease surveillance in office-based practice

All staff members in office-based practices need to be aware of the possibility that patients will present with suspected or confirmed infectious diseases.

For certain diseases, timely notification to the relevant authority will be required, sometimes by telephone. Systems need to be in place so that authorities are able to trace those with whom infectious patients have been in contact. A staff member should be responsible for checking national and state websites for relevant guidelines [379].

In most office-based practices, there will not be enough procedures performed to undertake outcome surveillance. Process surveillance can be used to evaluate processes and procedures and to monitor sentinel events. Systems should be in place for monitoring for threats of outbreaks (e.g. chickenpox [varicella], measles [rubeola]) and emerging diseases (e.g. *Candida auris* and Hendra virus).

Office-based practices can also play an important role in supporting the surveillance undertaken in acute care facilities by enabling post discharge surveillance [380]. Many surgical site infections, for example, will be identified up to 30 days after surgery, however patients are often discharged during this period. Office-based practices can participate in post-discharge surveillance case findings and report on these findings in order to maintain accurate data on surgical site infections.

4.4.6 Notifiable diseases

Communicable disease surveillance in Australia operates at the national, state/territory and local levels, with the states and territories having primary responsibility for the public health response to events identified by that surveillance.

National notifiable diseases

The Communicable Diseases Network Australia (CDNA) has agreed to a list of communicable diseases which are to be notified nationally and provided to the Commonwealth's National Notifiable Diseases Surveillance System (NNDSS).

See the *CDNA Australian national notifiable diseases and case definitions* [383] for further information.

Human biosecurity

Human biosecurity identifies the pests and communicable diseases that pose the most serious risk to public health in Australia. The *Biosecurity Act 2015* provides for the management of biosecurity threats posed by people, goods, vessels and aircraft at Australia's international borders. The departments of Health and Agriculture and Water Resources have joint responsibility for

administration of the Act, which commenced on 16 June 2016 and fully replaced the *Quarantine Act 1908*.

Under the *Biosecurity Act 2015*, provides for the concept of 'listed human disease'. This identified communicable diseases which require a range of powers and measures to become available to effectively manage significant human health risk.

Listed Human Diseases under the *Biosecurity Act 2015* include:

- human influenza with pandemic potential
- plague
- severe acute respiratory syndrome (SARS)
- Middle East respiratory syndrome
- smallpox
- viral haemorrhagic fevers
- yellow fever.

State and territory health department

Public health legislation in each state and territory mandates the reporting of certain diseases by medical practitioners, hospitals, and/or laboratories to the relevant state or territory Communicable Diseases Unit. Notifications are collected at the state/territory level, and computerised, de-identified records are sent to the Australian Government Department of Health for collation into the NNDSS for analysis at a national level. NNDSS was established in consultation with the CDNA.

- Links to state and territory public health legislation can be found at: <http://www.health.gov.au/internet/main/publishing.nsf/Content/cda-state-legislation-links.htm>
- In addition to those diseases which are notifiable nationally, a list of diseases which are notifiable in Australian states and territories can be found at: <http://www.health.gov.au/internet/main/publishing.nsf/Content/cda-surveil-nndss-casedefs-statedis.htm>

4.5 Antimicrobial Stewardship

Summary

- Resistance to antimicrobials is commonly found in Australian hospitals and increasingly so in the community. This resistance can have a significant impact on morbidity, mortality and treatment costs.
- A significant driver of antimicrobial resistance is the unnecessary or inappropriate use of antimicrobials. Around one third of all antimicrobial use in healthcare is unnecessary or inappropriately prescribed [384].
- Antimicrobial stewardship is a suite of coordinated strategies which together aim to promote the appropriate use of antimicrobials to maximise their benefit, while causing the least harm.
- Appropriate antimicrobial use occurs when antimicrobials are prescribed according to evidence-based guidelines, with drug choice, indication, dose and duration selected to optimise clinical outcomes and minimise adverse consequences (including antimicrobial resistance, toxicity and unnecessary costs).
- Surveillance of antimicrobial usage and resistance can be used to identify areas for improvement and measure the impact of antimicrobial stewardship programs.
- Infection prevention and control is recognised as an essential part of an effective response to antimicrobial resistance.

4.5.1 Antimicrobial resistance in Australia

Antimicrobial resistance

Antimicrobial resistance is recognised as a significant global health priority. The relationship between the unrestrained use of antimicrobials in all human health settings, agriculture and animal husbandry and the emergence of bacterial resistance is well documented.

The drivers of antimicrobial resistance are interlinked, with the use of particular antimicrobial classes associated with the emergence and amplification of specific multi-resistant pathogens, particularly *C. difficile*, methicillin-resistant *Staphylococcus aureus* (MRSA), vancomycin-resistant enterococci (VRE) and multi-resistant Gram negative (MRGN) organisms. If unchecked, high levels of antimicrobial usage increases the number of patients who are colonised or infected with resistant organisms, both in

healthcare facilities and in the community [385] [388].

Whilst the impact of antimicrobial resistance in Australia is yet to be quantified, in the United States it is expected to be responsible for at least 23,000 deaths and more than two million illnesses per year [387]. The World Health Organization (WHO) has called for governments to commit to a comprehensive national plan against antimicrobial resistance and has developed a Global Action Plan (2015) on antimicrobial resistance—<http://www.who.int/antimicrobial-resistance/global-action-plan/en/>.

The additional costs of infections caused by resistant organisms include:

- the need for more expensive and broader spectrum antimicrobials to treat the infections
- the need to isolate patients colonised with resistant organisms in order to minimise cross-infection
- the need for additional requirements such as personal protective equipment etc.

The role of infection prevention and control

Infection prevention and control practices are recognised in Australia and internationally as a key part of an effective response to antimicrobial resistance. Preventing infection reduces the need for antimicrobials and the opportunity for organisms to develop resistance. Vaccination (as discussed in Section 4.2, in particular Section 4.2.1) can also reduce antimicrobial resistance through preventing infectious diseases and reducing the prevalence of primary viral infections, which are often inappropriately treated with antimicrobials.

Antimicrobial use in Australia

Although Australia has seen a decline in antimicrobial use in the community since 2015, Australia remains in the top quarter of countries for antimicrobial use, as measured by defined daily doses (DDD) per 1,000 inhabitants, compared with European countries and Canada [389]. Australian antimicrobial use continues to be above the average of these countries, particularly compared with Scandinavian countries and the Netherlands, which are often seen as benchmark countries for antimicrobial use and resistance [389].

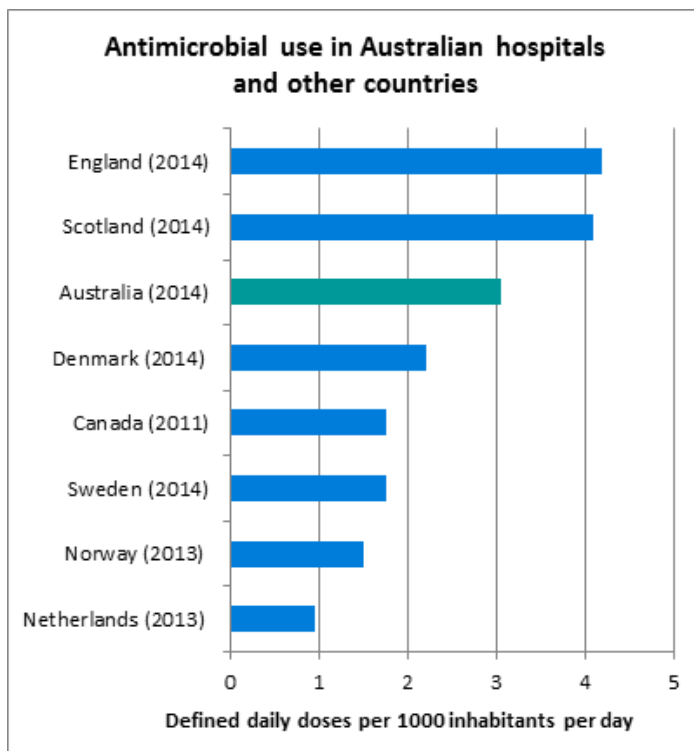


Figure 14. Antimicrobial use in Australian hospitals and other countries [384]

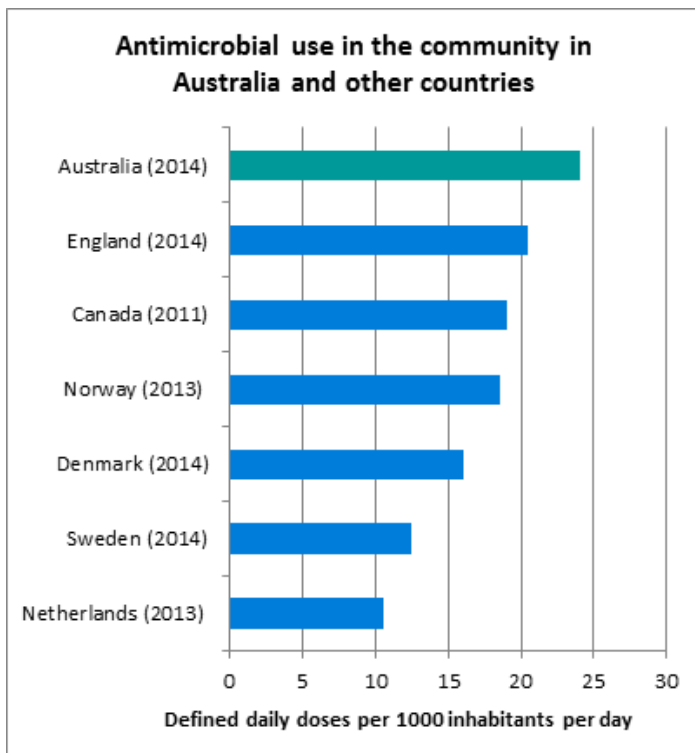


Figure 15. Antimicrobial use in the community in Australia and in other countries [384]

In 2017, 41.5% (n = 10,215,109) of the Australian population had at least one antibiotic dispensed under the Pharmaceutical Benefits Scheme/Repatriation Pharmaceutical Benefits Scheme [389].

The overall rate of inappropriate prescribing in hospitals that participated in the National Antimicrobial Prescribing Survey (a voluntary survey that assesses the appropriateness of antimicrobial use in hospitals) has been static since 2013. In 2017, 23.5% of prescriptions assessed were found to be inappropriate [389]. The conditions for which the highest proportions of prescriptions were assessed as inappropriate were surgical prophylaxis, infective exacerbation of chronic obstructive pulmonary disease and trauma [389].

Antibiotic use in hospitals, measured by voluntary participation in the National Antimicrobial Utilisation Surveillance Program, varied among states and territories, and among peer groups [389]. Antimicrobial use in aged care homes is a persistent concern with almost 1 in 10 residents of aged care homes that participated in the Aged Care National Antimicrobial Prescribing Survey being prescribed at least one antimicrobial; more than half of those were for residents with no signs or symptoms of infection [389].

4.5.2 Antimicrobial stewardship programs

What is antimicrobial stewardship?

Antimicrobial stewardship is a suite of coordinated strategies which together aim to promote the appropriate use of antimicrobials to maximise their benefit, whilst causing the least harm. Antimicrobial stewardship programs should be implemented across the entire spectrum of healthcare facilities - including in community and residential care.

Antimicrobial stewardship programs in hospital/acute settings

Successful antimicrobial stewardship programs have been associated with reduced facility resistance rates as well as reduced morbidity, mortality and associated costs. Some Australian hospitals have also demonstrated significant cost savings through a reduction in drug costs. Antimicrobial stewardship programs require a multidisciplinary approach which utilises the expertise and resources of infectious diseases physicians, clinical microbiologists and pharmacists. Antimicrobial stewardship programs should reside within the hospital's quality improvement and patient safety governance structure. They should be tailored depending on the organisational context and factors such as size of the facility, staffing and resourcing [390]. The literature demonstrates that the success of antimicrobial stewardship programs depends on the support of the hospital administration, the allocation of adequate resources, and the cooperation and engagement of prescribers [386].

The density of antimicrobial use within specialised units such as intensive care units, haematology and oncology units, and solid-organ transplant units is several-fold higher than in other hospital settings. This increased use has been shown to generate high

rates of antimicrobial resistance; therefore, these areas should be a particular focus for surveillance and intervention.

Key requirements of a healthcare facility antimicrobial stewardship program [386][390]

Essential strategies for all hospitals:

- Implement clinical guidelines that comply with the latest version of the *Therapeutic Guidelines: Antibiotic* and that take into account local microbiology and resistance patterns.
- Develop, review and maintain antimicrobial prescribing policies. This should include formulary restriction and approval systems that cover restriction of broad-spectrum antibiotics to those patients where use is clinically justified.
- Review individual antimicrobial prescribing with intervention and direct feedback to the prescriber.
- Monitor antimicrobial prescribing and report on antimicrobial use and outcomes.
- Ensure selective reporting of susceptibility testing results that is consistent with hospital antimicrobial treatment guidelines by the clinical microbiology laboratory.
- Ensure ongoing education and training for health professionals and consumers about antimicrobial stewardship, antimicrobial resistance and safe and appropriate antimicrobial use.
- Use information technology such as electronic prescribing with clinical decision-support or online approval systems.
- Use point-of-care interventions, including streamlining or de-escalation of therapy, dose optimisation, parenteral to oral conversation.
- Annually publish antimicrobial usage and antimicrobial susceptibility data.

The Australian Commission on Safety and Quality in Health Care website has a range of resources and materials to support appropriate antimicrobial prescribing and use, and implementation of antimicrobial stewardship programs in Australian healthcare - see: <https://www.safetyandquality.gov.au/our-work/healthcare-associated-infection/antimicrobial-stewardship/>.

Antimicrobial stewardship in other healthcare settings

There is a strong need to implement Antimicrobial Stewardship Programs in other healthcare settings, particularly in residential aged care facilities where the infection risk for elderly patients is high. There is increasing concern about the widespread use of antimicrobials in aged care settings, with many prescriptions deemed unnecessary or inappropriate [392].

Whilst there may be barriers to implementing antimicrobial stewardship programs in aged care settings including resourcing issues and lack of access to laboratories, Antimicrobial Stewardship activities that are tailored to this setting can be very effective.

Some strategies which may be feasible to implement in aged care settings include [391]:

- educating healthcare staff on appropriate antimicrobial prescribing and antimicrobial resistance and providing feedback on their prescribing practices
- implementing facility wide tracking and reporting on antimicrobial use and resistance
- designating staff members, such as a medical director or director of nursing, to be accountable for promoting and overseeing antimicrobial stewardship activities
- implementing policies that support optimal antimicrobial use.

Further information on antimicrobial stewardship strategies and interventions can be found in *Antimicrobial Stewardship In Australian Health Care 2018* [393].

4.5.3 Antimicrobial stewardship surveillance methods

Healthcare facilities

Continuous surveillance of the appropriateness of antimicrobial prescribing should be the ultimate aim of any stewardship program. There are two main methods of antimicrobial data collection in healthcare facilities, patient level surveillance and population surveillance:

- **Patient level surveillance** involves collecting data about the dose, dosage interval, indication, antimicrobial choice and duration of therapy for individual patients, usually collected as a point prevalence survey. This approach can be used to measure the appropriateness of the use of antimicrobials. Such information is usually only available through reviews of patient records, although electronic prescribing and recording of drug administration will make patient level surveillance more practical in the future.
- **Population surveillance** involves aggregating antimicrobial use data, mostly supplied through pharmacy reports, and summarised at the level of a hospital or unit. Currently, this type of surveillance is the only realistic alternative for ongoing and systematic monitoring of antimicrobial use. In most Australian hospitals, data from issues confronted in wards combined with

individual patient dispensing records are used. Another data collection method is to use pharmacy purchase data; however, this is less accurate than ward issues and individual inpatient supplies.

Participation in jurisdictional or national data collection programs allows for healthcare facilities to monitor antimicrobial use against similar hospitals and access auditing tools. The *National Antimicrobial Utilisation Surveillance Program* (NAUSP) provides hospitals with a method to report on the volume of hospital inpatient antimicrobial usage and reports to the Australian Commission on Safety and Quality in Health Care on an aggregated and Australian Institute for Health and Welfare (AIHW) peer group basis. In excess of 200 hospitals from all states and territories currently contribute to NAUSP; including all principal referral hospitals.

Similarly, the *National Antimicrobial Prescribing Survey* (NAPS) can be used to measure the appropriateness of antimicrobial prescribing practices within Australian healthcare settings. NAPS is a standardised data collection tool that collects data from healthcare facilities on their prescribing behaviours and allows for the audit and review of antimicrobial use. NAPS is a component of the Antimicrobial Use and Resistance in Australia (AURA) Surveillance System.

Community

Measurement of community antimicrobial use is generally based on prescription data. In Australia, this is collected from two sources: Australian Government Department of Human Services pharmacy claim records of prescriptions submitted for payment under the Pharmaceutical Benefits Scheme (PBS) and Repatriation Pharmaceutical Benefits Scheme (RPBS); and Drug Utilisation Subcommittee database. These data also include antimicrobials dispensed to outpatients and discharged patients in most states. NPS MedicineWise MedicineInsight program collects and reports on the appropriateness of antimicrobial use in general practice.

Further information on antimicrobial stewardship strategies and interventions can be found in *Antimicrobial Stewardship In Australian Health Care 2018* [393].

4.6 Influence of facility design on healthcare associated infection

Summary

The design of a healthcare facility can influence the transmission of healthcare associated infections by air, water and contact with the physical environment. Whilst there is little high quality evidence relevant to the effects of specific design features on health outcomes, data from case reports and published literature relating to outbreaks shows that the design of buildings can have an impact on rates of healthcare associated infections. Infection prevention and control requirements need to be taken into account during the planning, design and construction of all healthcare facilities.

Key design features that minimise the transmission of infection include:

- surface finishes that are easy to maintain and clean (floors, walls, benches, fixtures and fittings)
- ventilation, air conditioning, cooling towers and water systems that meet Australian standards for the facility they are to service
- the ability to isolate patients:
 - in a single room (infectious patients)
 - in a negative pressure room (to prevent transmission of airborne pathogens)
 - triaging of patients in waiting rooms with separation of infectious patients
- positive pressure rooms for immunocompromised patients
- appropriate workplace design:
 - separation of procedural and cleaning areas
 - movement of work flow systems
 - ready access to hand hygiene facilities
 - adequate storage for all patient-care items
 - easily accessible storage for personal protective equipment
 - adequate waste management procedures and linen handling
- involvement in demolition, construction and renovation projects of a multidisciplinary team that includes infection prevention and control staff to coordinate preventive measures.

Information about infection prevention and control during construction or renovation of a healthcare facility is at Case Study 5.15 and further information is available in the *Australasian Health Facility Guidelines (2016)* [394].

4.6.1 Mechanisms for influencing healthcare associated infection through environmental design

Many studies indicate that infection rates are lower when there is very good air and water quality, greater physical separation of patients and greater space per patient (with isolation where appropriate).

Reducing airborne transmission

Reservoirs for airborne pathogens include [418]:

- dust (e.g. spores of *Aspergillus*)
- aerosols (e.g. tuberculosis [TB], severe acute respiratory syndrome [SARS], influenza, chickenpox)
- skin scales shed by patients infected or colonised with methicillin-resistant *Staphylococcus aureus* (MRSA).

Approaches to airborne transmission

Approaches to reducing airborne transmission include:

- installation of effective air filtration
- specifying appropriate ventilation systems and air change rates (e.g. negative airflow pressure)
- employing monitoring and control measures during construction or renovation
- using single-bed instead of multi-bed rooms
- the use of anterooms.

Specific examples

In dental practices, engineering rules state there must be separation between inlet air for compressors and air conditioning outlets [399].

Gastrointestinal endoscopy suites should be organised and planned carefully to ensure endoscopes are not re-contaminated or damaged. Endoscopy suites also need good ventilation to minimise staff inhalation of biological aerosols [406].

Filtration

An effective way to prevent infections is to control the source of pathogens. Heating, ventilation and air-conditioning systems control the concentration of airborne particulates in high risk areas, to minimise the risk of infection by means of air pressure, flow control and air filtration (the physical removal of particulates from air). The level of control should be proportional to the risk.

In acute healthcare settings, a commonly used approach to filtration is the high efficiency particulate air (HEPA) filter [417]. There is evidence that there is a lower incidence of infection when immunocompromised and other high-acuity patients are housed in HEPA-filtered isolation rooms. HEPA filters must comply with **Standard AS 1324.1: 2001, Standard AS 1324.2: 2003 and Standard AS 4260**. For further information on HEPA filters, see the current *Australasian Health Facility Guidelines* [394].

Ventilation systems and airflow control

Optimal ventilation rates, airflow patterns and humidity can help to minimise the spread of infection.

The ventilation rate is a measure used to control indoor air quality, and in healthcare facilities is usually expressed as room air changes per hour (ACH). The peak efficiency for particle removal in the air space often occurs between 12 ACH and 15 ACH. **Standard AS 1668.2: 2024** states that isolation rooms should have a minimum of 15 ACH; however, there is a lack of consistency in the minimum ventilation requirements needed for effective infection prevention. A study of 17 Canadian hospitals found that the risk of healthcare workers acquiring TB was strongly linked with exposure to infected patients in rooms with low ACH rates, such as waiting areas [410].

Airflow direction is also important:

- **Negative airflow pressure** is preferred for rooms housing infectious patients to prevent the dispersion of pathogen-laden aerosols (e.g. measles [rubeola], TB, chickenpox [varicella]), dust and skin scales from the locus of the infected patient to other spaces. A review of 40 studies concluded that there is strong evidence to support and recommend the use of negatively pressurised isolation rooms [409].
- **Positive airflow pressure** is desirable in the care of immunocompromised patients (e.g. surgical patients, patients with underlying chronic lung disease, or dialysis patients) and immunosuppressed patients (e.g. transplant patients or cancer patients) to safeguard them from aerial pathogens entering from adjacent spaces.

Maintenance systems

Ventilation and airflow control systems need to be maintained regularly by suitably qualified staff according to an agreed maintenance plan, and accurately documented in a maintenance record.

Maintaining air quality during construction or renovation

Effective control and prevention measures are necessary during construction and renovation within a healthcare facility, because such activities have been frequently implicated in outbreaks of airborne infection. The key to eliminating infections is to minimise the dust generated during the construction activity and to prevent dust infiltration into patient-care areas near the construction. Examples of such measures include installing barriers between patient-care areas and construction/renovation areas, generating negative air pressure for construction/renovation areas relative to patient-care areas, using portable HEPA filters and sealing patient windows.

For more information, refer to the Centers for Disease Control and Prevention Guidelines for Environmental Infection Control in Health-Care Facilities (2004) [414] and Australasian Health Facility Guidelines (2016) [396].

Reducing infections spread through the physical environment

The prevention of contact-spread infections is of paramount importance in healthcare settings. Contact contamination is generally recognised as the principal transmission route of healthcare acquired infections, including pathogens such as MRSA, *C. difficile* and vancomycin-resistant enterococci (VRE), which survive well on environmental surfaces and other reservoirs.

Environmental routes of contact-spread infections include direct person-to-person contact and indirect transmission via environmental surfaces.

Reducing surface contamination through hand-hygiene compliance

Healthcare workers' hands play a key role in both direct and indirect transmission (see Section 3.1.1). Given the importance of maximising hand-hygiene compliance, it is absolutely essential that all areas of the facility are designed to facilitate compliance with hand-hygiene requirements.

Accessibility

Conveniently located alcohol-based product dispensers, sinks and basins can facilitate healthcare worker compliance with hand-hygiene requirements [57].

Hand-hygiene compliance can be increased by providing a greater number of alcohol-based product dispensers, particularly if they are placed in appropriate locations (where clinical care is provided [e.g. bedside] or where indirect care tasks are performed). Other aspects of design that may increase compliance include automated dispensers of hand-hygiene products, electronic monitoring and computerised voice prompts.

Alcohol-based hand rub dispensers need to be suitably located out of the reach of children, or in supervised locations. Placement of dispensers must be carefully considered in mental health facilities and alcohol withdrawal units. Further guidance is available from the [National Hand Hygiene Initiative](#) [57].

Consideration needs to be given to ensuring availability of basins for healthcare workers that are separate from patient bathrooms.

As well as being installed in all patient-care areas, hand-hygiene facilities should be placed in all areas where careful attention to hygiene is essential, such as kitchens, laundries, pharmacies, laboratories and staff amenities areas (e.g. bathrooms, toilets and change rooms). This also includes in specific settings such as treatment/procedure rooms [399].

Personal protective equipment

It is also essential that all areas of the facility are designed to facilitate appropriate use of PPE. All rooms should have dedicated and accessible areas for storage of gowns, aprons, gloves, masks and protective eyewear.

Separation of procedural and cleaning areas

The instrument reprocessing area also requires sufficient physical space to allow separation of soiled instruments from those that have been cleaned or sterilised. Ideally, the area where instruments are cleaned will be physically separated from the areas where cleaned instruments are packaged and sterilised.

Control of surface contamination through material selection

Ease of cleaning should be a key consideration in selecting appropriate floor and furniture coverings. Several design-related factors should be considered to minimise the risk of infection stemming from contaminated surfaces:

- the nature and type of contamination that is likely to occur
- if a suitable cleaning method for that surface can be performed.

Areas that may be in direct contact with blood and body substances (e.g. surfaces such as floors and bench tops) need to be made of

impervious material that is smooth and easy to clean.

Healthcare flooring

A wide range of floor covering materials is used in healthcare settings. These include but are not limited to: ceramic tiling, linoleum, rubber, textile floor covering, vinyl, sheet terrazzo, cork, timber laminates, mats and matting, cementitious toppings, seamless coatings and outdoor flooring.

When selecting floor covering for a health care setting consideration needs to be given to the following:

- Who is at risk of acquiring infection?
- What is the risk of exposure to the infectious agents?
- What is the nature of the possible infectious agents?
- How can the agent be transmitted?—for example, airborne; through cleaning techniques; through contact, especially in environments in which there are young children.

Floor coverings should be clean and easy to repair. In areas subject to frequent wet cleaning, floor materials must be able to tolerate the use of disinfectants. In areas where sterilizing services take place, flooring should be non-slip and have smooth surfaces for cleaning [414].

Carpeting should be avoided in clinical areas where patient care and treatments are undertaken, including areas where [414]:

- spills are likely to occur (e.g. around sinks or in isolation or soiled utility/holding areas)
- patients may have direct contact with contaminated carpets (e.g. children/babies crawling on the floor)
- patients are at greater risk of airborne infections.

However, carpeting may offer advantages unrelated to infection prevention and control, including noise reduction [413]. Carpet may be used in areas within clinical zones such as interview rooms and office areas. Textile floor finishes should not be considered unless there is a comprehensive maintenance and replacement program in place complying with **Standard AS/NZS 3733:**

2018. Manufacturer's recommendations need to be considered in the care and maintenance of floor covering.

Furnishings

It has been identified that fabric-covered furniture can be a source of VRE infection in hospitals and suggested the use of easily cleanable, non-porous material is preferable [412].

A study comparing the performance of a variety of furniture upholstery types with respect to VRE and *Pseudomonas aeruginosa* (PSAE) contamination [419] found that performance was similar across different furniture coverings in terms of reductions in VRE and PSAE after cleaning and the transfer of VRE and PSAE to hands through contact. However, while there were no differences in the ability of different upholstery types to harbour PSAE, the VRE pathogen survived less well or for shorter periods on vinyl [419].

Requirements for finishes of items may vary depending upon the location used e.g. fabric upholstery on chairs may not be permitted in clinical areas.

Blinds and curtains should be easy to clean and discourage the accumulation of dust.

For further information on furniture and fittings, see the current *Australasian Health Facility Guidelines* [394].

Reducing water-borne transmission

Compared with airborne and contact transmission of infection, fewer studies were identified on waterborne transmission in relation to healthcare facility design factors. The literature nonetheless is clear that waterborne infections can be a serious threat to patient safety. Many bacterial and some protozoal microorganisms can proliferate or remain viable in moist environments or aqueous solutions in healthcare settings [414].

Contaminated water systems in healthcare settings (such as inadequately treated wastewater) may lead to the pollution of municipal water systems, enter surface or ground water, and affect people in the community [408].

Sources of water contamination

The CDC/Healthcare Infection Control Practices Advisory Committee (HICPAC) guidelines [414] identify the following categories of environmental routes or sources of waterborne transmission:

- direct contact, such as hydrotherapy [397]
- ingestion of water, such as drinking water [403] [416]
- inhalation of aerosols dispersed from contaminated water sources, such as improperly cleaned or maintained cooling towers, showers [411], respiratory therapy equipment and room air humidifiers

- aspiration of contaminated water.

Approaches to reducing waterborne transmission

Water supply system

The water supply system should be designed and maintained with proper temperature and adequate pressure; stagnation and back flow should be minimised and dead-end pipes should be avoided.

To prevent the growth of *Legionella* and other bacteria, the CDC/HICPAC guidelines [414] recommend that healthcare facilities maintain cold water at a temperature below 20°C, store hot water above 60°C, and circulate hot water with a minimum return temperature of 51°C.

When the recommended standards cannot be achieved because of inadequate facilities that are unable to be renovated, other measures such as chlorine treatment, copper-silver ionisation, or ultraviolet lights are recommended to ensure water quality and prevent infection [414].

Point-of-use fixtures

Water fixtures such as sinks, faucets, aerators, showers, and toilets have been identified as potential reservoirs for pathogenic microorganisms [401] [403] [411] [416]. Such fixtures produce aerosols that can disperse microbes and they have wet surfaces on which moulds and other microorganisms can proliferate. However, empirical evidence linking these fixtures to HAIs is still limited; no consensus has been reached regarding the disinfection or removal of these devices for general use [414].

Regular cleaning, disinfection and preventative maintenance programs should be provided, especially in areas housing immunocompromised patients.

Ice machines

Ice storage receptacles and ice-making machines should be properly maintained and regularly cleaned. Ice and ice-making machines may be contaminated through improper handling of ice by patients and/or staff. Ice for human consumption should be differentiated from ice for first aid or storage of clinical specimens. Pharmaceuticals or medical solutions should not be stored on ice intended for consumption.

Machines that dispense ice are preferable to those that require ice to be removed from bins or chests with a scoop. Ice machines and their dispensers should be flushed and cleaned if they have not been disconnected before anticipated lengthy water disruptions.

All ice-storage chests should be cleaned, disinfected, and maintained on a regular basis as per manufacturers' instructions.

Suggested steps to avoid improper handling of ice include [414]:

- avoiding handling ice directly by hand
- washing hands before obtaining ice
- using a smooth-surface ice scoop to dispense ice
- keeping the ice scoop on a chain short enough that the scoop cannot touch the floor, or keeping the scoop on a clean, hard surface when not in use
- avoiding storing the ice scoop in the ice bin.

Water features

Despite the absence of empirical documentation linking properly maintained fountains to healthcare-acquired infections, it is suggested that fountains not be installed in enclosed spaces in healthcare facilities.

5. APPENDIX 1: Case studies and examples

Summary

The following risk-management case studies and examples provide practical guidance on how to implement infection control recommendations and practice statements in practice.

This section covers:

- **5.1** — Risk-management: Case study for hand hygiene in a neonatal intensive care unit
- **5.2** — Risk-management: Case study for glove use, hand hygiene and seasonal influenza vaccination in an office-based practice
- **5.3** — Example: Education in practice—hand hygiene
- **5.4** — Risk-management: Case study for the prevention of needle-stick injury during surgery at a tertiary referral hospital
- **5.5** — Risk-management: Case study for spills management in a busy paediatric ward
- **5.6** — Risk-management: Case study for reprocessing of instruments in a dental practice
- **5.7** — Risk-management: Case study for Klebsiella pneumoniae sepsis in a neonatal unit
- **5.8** — Risk-management: Case Study for influenza in a long-term care facility
- **5.9** — Risk-management: Case study for M. tuberculosis
- **5.10** — Risk-management: Case study for M. tuberculosis among immunocompromised patients attending outpatient services
- **5.11** — Examples of how to perform aseptic technique
- **5.12** — Risk-management: Case study for norovirus outbreak in a long-term care facility
- **5.13** — Risk-management: Case study for management of confirmed case of CPE
- **5.14** — Risk Management: Case study for VRE outbreak in a large tertiary-care referral hospital
- **5.15** — Risk-management: Case study for infection prevention during renovation of emergency department

5.1 Risk-management: Case study for hand hygiene in a neonatal intensive care unit

The neonatal intensive care unit in a large regional hospital identifies colonisation or infection with Pseudomonas aeruginosa in a number of infants. Surveillance cultures from other infants in the unit, from the hands of staff on the unit and from possible environmental reservoirs are assessed. The cultures show that an additional three infants are colonised. Cultures of environmental specimens are negative but cultures of three of twenty-four healthcare workers are positive. Of these, two have recently joined the unit and received no education on hand hygiene in orientation and the third has artificial fingernails.

Table A1.1. Case study for hand hygiene in a neonatal intensive care unit

| | |
|--------------------------|---|
| Eliminating risks | In this situation, it is not possible to eliminate risk, so it must be managed. |
| Identifying risks | In this case, the risk has been identified as cross-transmission of <i>Pseudomonas aeruginosa</i> . Ongoing surveillance would assist in identifying other infectious agents that may be present in the neonatal intensive care unit. |
| Analysing risks | <p>One source of the risk is the lack of appropriate hand hygiene practices by some staff members. Each time these staff members are involved in the care of an infected or colonised infant, there is potential for spread of the infectious agent (to other infants and to staff members), with the risk continuing until appropriate hand hygiene practices are performed.</p> <p>Further, neonates are at a greater risk of infection than an average patient because their immune system has not properly developed yet.</p> <p>There is no mention in the case study of existing controls to counter the risk (e.g. use of gloves) but these would need to be included in the analysis, as would other possible causes of the risk (e.g. line setup, type of hand hygiene products available, reprocessing of equipment).</p> |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response including daily observations by clinical managers. |
| Treating risks | <p><u>Immediate measures</u> may include the provision of alcohol-based hand rub next to each incubator, introducing clustering of patient-care activities to reduce contact and providing staff education sessions.</p> <p><u>In the longer term</u>, improvements could be made to facility orientation processes. Banning the use of artificial fingernails in neonatal intensive care units may also be considered.</p> |

| | |
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| Monitoring | Hand hygiene compliance could be audited through direct observation by trained observers. |
|-------------------|---|

5.2 Risk-management: Case study for glove use, hand hygiene and seasonal influenza vaccination in an office-based practice

As part of an audit of hand hygiene practices during influenza season in a general medical practice, a general practitioner (GP) identifies a poor rate of compliance with the 5 Moments for Hand Hygiene in her practice when the results are compared with Primary Health Networks benchmarks. The practice comprises five GPs and two part-time practice nurses. Practice policy is that staff members use gloves for patient contact, changing gloves between patients. There is no recommendation in the policy for hand hygiene between different care activities for the same patient or after removing gloves. There is no policy on vaccination of the workforce for annual seasonal influenza vaccination to be offered or assessed.

Table A1.2. Case study for glove use, hand hygiene and seasonal influenza vaccination in an office-based practice

| | |
|--------------------------|---|
| Eliminating risks | <p>Patients with influenza may be infectious both before and after they present with symptoms. In this situation it is not possible to eliminate risk, so it must be managed.</p> <p>Staff in the general practice who have not been vaccinated for seasonal influenza may be at increased risk of contracting influenza from patients or transmitting it to patients if they are at work whilst infectious. This situation is possible and needs to be managed in the general practice. The risk can be minimised by offering or assessing annual seasonal influenza to staff and having a policy where staff who are unvaccinated limit contact with possible cases and use appropriate personal protective equipment (PPE).</p> |
| Identifying risks | <p>The risk has been identified as cross-transmission of influenza, with higher than usual rates occurring. An audit of the healthcare workers to determine who has received annual seasonal influenza vaccination.</p> <p>An audit of cases of other infections that may be transmitted in the healthcare environment would assist in identifying other infectious agents that may also be occurring at high rates in the practice.</p> |
| Analysing risks | <p>Wearing gloves does not replace the need for hand hygiene. One source of the risk has been identified as the lack of hand hygiene before and after use of gloves.</p> <p>Each time a patient carrying influenza is examined, there is potential for the spread of the infectious agent directly to the unvaccinated healthcare worker or from the glove to the healthcare worker's hand and to the gloves worn for subsequent patients. The same applies for other infectious agents spread by contact or droplet routes.</p> <p>Existing controls such as availability of alcohol-based hand rub or hand washing facilities, and other sources of risk such as environmental surfaces would also need investigation.</p> |
| Evaluating risks | <p>The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response.</p> |
| Treating risks | <p><u>Immediate measures</u> may include:</p> <ul style="list-style-type: none"> • additional cleaning of the practice to reduce the levels of transient micro-organisms • increased provision of alcohol-based hand rub or hand washing facilities, as well as signage at all points of care • staff education in hand hygiene, appropriate use of PPE and environmental cleaning • identify if an area in the waiting room can be allocated for segregation of patients with signs and symptoms of influenza-like-illness whilst waiting to see a GP or practice nurses • identify which healthcare staff have been vaccinated for influenza and offer it to any of the workforce who haven't been vaccinated • promote influenza vaccination with patients especially those in identified risk groups. <p><i>For example:</i> With a typical patient, the practitioner should complete appropriate hand hygiene before seeing the patient. If the patient shakes the practitioner's hand, risk should be mitigated by again performing hand hygiene. When attending to the patient, hand hygiene is required between all procedures with the patient, each time gloves are removed and before the practitioner attends to patient notes.</p> |

| | |
|--------------------------|--|
| | <p>The practitioner should also consider that infectious agents which are transferred by droplet routes could also require contact precautions. The practitioner should ensure that both opened and unopened boxes of gloves and other clinical equipment is stored appropriately to prevent exposure to droplets from infected patients.</p> <p><u>Long-term measures</u> may include:</p> <ul style="list-style-type: none"> • Development or revision; and implementation of hand hygiene, PPE, healthcare worker vaccination and environmental cleaning policies. This could be carried out by the GP as practice leader, in consultation with other staff. <p><i>For example:</i> A range of environmental controls should be in place to reduce the risk of transmission of influenza for patients visiting the practice; including, the design of the facility, availability of PPE for staff and patients, routine cleaning of waiting and treatment rooms, and environmental cleaning schedules.</p> |
| <p>Monitoring</p> | <p>GPs and practice nurses should continue to monitor patients for influenza-like illness (ILI) and ongoing ILI surveillance should be heightened during the influenza season.</p> <p>Systems should be in place which allow for reporting, monitoring and rectifying breaches of infection control policies.</p> <p><i>For example:</i> Changes in practice could be monitored through audit of amounts of gloves and alcohol-based hand rub used. Repeating the audit of patient infections at regular intervals would assist in monitoring improvements.</p> |

5.3 Example: Education in practice - hand hygiene

Although the concept of hand hygiene is straightforward, improving hand hygiene practices involves changing attitudes and behaviour among healthcare workers. Numerous barriers to appropriate hand hygiene have been reported, several of which reflect a lack of understanding and knowledge, including the following [62]:

- all hand hygiene agents being thought to cause skin irritation and dryness
- patient needs being perceived to take priority over hand hygiene
- perception that glove use dispenses with the need for additional hand hygiene
- belief that there is insufficient time for hand hygiene, due to high workload
- inadequate knowledge of guidelines or protocols for hand hygiene
- lack of role models
- lack of recognition of the risk of cross-transmission of microbial pathogens.

National Hand Hygiene Initiative

The National Hand Hygiene Initiative aims to implement a national approach to improving hand hygiene and monitoring its effectiveness. In the initiative, healthcare worker education is a key component of a multi-modal intervention strategy, involving basic educational sessions for all healthcare workers, including:

- definition, impact and burden of healthcare associated infection
- common pathways for disease transmission - specifically the role of hands
- prevention of HAIs and the role of hand hygiene
- 5 Moments for Hand Hygiene - with key messages
- when to perform hand hygiene
- use of alcohol-based hand rubs
- use at point of care.

As well as introductory educational sessions, a program of formal regular sessions and updates is recommended, taking the form of specific orientation programs, in-service lectures and/or special workshops. All education sessions are supported by an online training package, DVD, video demonstrations of each of the five moments, and slide presentations.

Other opportunities for education include:

- informal education opportunities in day-to-day activities such as nursing ward rounds, clinical unit meetings, increased presence on the ward by infection prevention and control staff, and prompt feedback of compliance results

- promotional activities to raise awareness, with promotional products (e.g. stickers) or incentives for staff who attend education sessions
- regular scientific presentations at surgical and medical meetings
- a multi-modal approach has also found to be effective, led by hand hygiene champions.

All healthcare workers should be regularly assessed for their hand hygiene compliance and be provided with rapid feedback of results (see 3.5.1).

Since data was first collected in 2009, the national compliance rate for hand hygiene in Australian hospitals has risen from 63.5% to 84.6% in 2017.

Other measures to increase compliance with hand hygiene are discussed in Section 4.6.

5.4 Risk-management: Case study for the prevention of needlestick injury during surgery at a tertiary referral hospital

As part of the revision of infection control policies at a tertiary referral hospital, an analysis of the risk of percutaneous blood and body substance exposure during surgical procedures was undertaken. Separate analyses were conducted for different device types and for different members of the surgical team. Surgeons and first assistants were at highest risk for injury, suffering more than half of injuries in the operating room, followed by scrub nurses and technicians, anaesthetists and circulating nurses. Rates of needle-stick injury increased with estimated blood loss and surgery duration. Suture needle injuries were the most common and mostly occurred during wound closure.

A considerable number of injuries also occurred while passing sharp instruments hand-to-hand. As many as one-third of devices that caused injuries came in contact with the patient after the injury to the healthcare worker. However, only a small proportion of injuries to surgeons (0.5%) involved hollowbore vascular access needles, which are defined as 'high risk'.

Table A1.3. Case study for the prevention of needlestick injury during surgery at a tertiary referral hospital

| | |
|--------------------------|---|
| Eliminating risks | Although the risk of injury varies for different healthcare team members, it is never zero and must be managed. |
| Identifying risks | In this case, the risk has been identified as exposure of healthcare workers to blood and body substances (and potential infection) through suture needle injury. Risks associated with sharps injuries can be increased when the sharp is hollowbore needle (e.g. hypodermic needle) compared to a solid needle or sharp (e.g. suture needle) due to the increased risk of exposure to blood. As a high proportion of devices causing injury came into contact with the patient after injury to the healthcare worker, there could also be a risk of transmission of blood-borne infection to the patient. |
| Analysing risks | The fundamental source of risk is the need to use sharps coupled with the potential for a patient to be a source of infection. The level of risk increases with duration of procedure and amount of blood lost. Other factors that may contribute to the risk are levels of staff training and experience, staffing levels, the existence of a hospital policy for safe use of sharps and compliance with the policy. Other factors that would need to be included in the analysis are existing controls to mitigate risk (e.g. double gloving) and other possible causes (e.g. poor surgical technique increasing blood loss and procedure duration). |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response. |
| Treating risks | All healthcare workers should understand their obligation to report all sharps injuries, whether or not there was a risk of patient exposure. Comprehensive reporting is required to enhance surveillance of possible blood-borne virus (BBV) transmission. Healthcare workers who are BBV positive and who undertake exposure-prone procedures must be familiar with the current standards of infection prevention and control and have an action plan in place in the event of a potential transmission event that includes reporting the event according to local policies. |

| | |
|-------------------|--|
| | <p><u>Immediate measures</u> to treat the risk may include providing staff education, use of blunt suture needles and a neutral zone for passing surgical equipment, and double gloving during long surgery.</p> <p><u>In the longer term</u>, reviewing local policy on the prevention of needlestick injury and raising awareness of measures to reduce injury among staff members might also be considered.</p> |
| Monitoring | Changes in adverse events could be evaluated by repeating the analysis after implementation of changes. |

5.5 Risk-management: Case study for spills management in a busy paediatric ward

A visitor to the paediatric ward in a small regional hospital notices that the child in the next bed is vomiting and has diarrhoea. The ward is extremely busy and the two nurses on duty are fully occupied. The child's mother has cleaned up any spills, but there are still traces of vomit on the bedside table. Later the visitor notices that equipment is being placed on this table. When there is a lull in activity in the ward, the visitor approaches one of the nurses and mentions what she has noticed. The nurse is grateful for the advice and the quiet period is used for more thorough cleaning of surfaces around the vomiting child. The nurse thanks the mother for her assistance and explains to her the importance of thorough cleaning and hand hygiene in the prevention of transmission of infection.

Table A1.4. Case study for spills management in a busy paediatric ward

| | |
|--------------------------|---|
| Eliminating risks | Ideally, this risk can be eliminated through immediate removal and cleaning of spills. However, in many situations it is more likely that the risk will be managed. |
| Identifying risks | The risk has been identified as potential cross-transmission of Norovirus through environmental contamination. |
| Analysing risks | <p>One source of the risk has been identified as inadequate environmental cleaning by a visitor resulting in potential contamination of equipment placed on environmental surfaces (bedside table) or hands touching this surface.</p> <p>There is then potential for direct or indirect spread of infection to other patients, visitors and healthcare workers.</p> <p>There are likely to be other infectious agents that could be transmitted in the same way (e.g. rotavirus).</p> |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'high risk' situation requiring immediate response. |
| Treating risks | <p><u>Immediate measures</u> may include raising patient and visitor awareness of hygiene measures (including hand hygiene as well as environmental cleaning). This could be done through posters and/or discussion with patients and carers on admission.</p> <p><u>Longer-term measures</u> could include revision and implementation of environmental cleaning policies and involvement of patients/visitors in this review.</p> <p>Retrospectively, a review would be undertaken to determine why the patient was or wasn't placed in a single room as per contact precautions.</p> |
| Monitoring | Changes in practice could be monitored through observation of patient/visitor behaviour. |

5.6 Risk-management: Case study for reprocessing of instruments in a dental practice

A patient attends a dental practice for a scaling and cleaning of his teeth. He has moderate periodontal disease with inflamed gingiva (gums). The dentist uses both an ultrasonic scaler (which creates aerosol) and very sharp hand scalers and curettes. Neither the dentist nor the assistant wears a mask. To protect the tongue and cheeks of the patient from being injured by the sharp instruments, a dental mirror is used to retract them. The mirror consists of a handle into which a mirror head is screwed. The handle of the mirror has a corrugated surface so that it doesn't slip. During this procedure the mirror gets covered in blood from the bleeding of the inflamed

diseased gums.

Table A1.5. Case study for reprocessing of instruments in a dental practice

| | |
|--------------------------|---|
| Eliminating risks | Proper reprocessing of the instrument, operator and assistant care in the use of sharp instruments and protecting against possible aerosol exposure has the potential to eliminate the risk. |
| Identifying risks | There is a risk of exposure of other patients to blood borne viruses if the mirror is not reprocessed properly. There is also a risk of exposure of staff to aerosol infectious agents (influenza in particular) and a risk of staff exposure to blood borne viruses through sharps injury (either during the treatment or during reprocessing). |
| Analysing risks | Sources of the risk are difficulties in reprocessing the mirror, the use of multiple sharp instruments in a bloody field and aerosolisation caused by the treatment. |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'medium risk' situation requiring management by specific monitoring or audit procedures. |
| Treating risks | <u>Immediate measures</u> include: making sure that mirror handles are clean before sterilisation, operator care in the use of sharp instruments, use of high volume evacuation to reduce aerosolisation caused by this treatment and, wearing of protective eyewear and face masks by operator and assistant. <u>Longer-term measures</u> could include revising practice, personal protective equipment (PPE) and instrument cleaning and reprocessing policies. |
| Monitoring | Repeated checking of reprocessed instruments, audits of staff sharps injuries and monitoring of PPE use would assist in assessing the level of risk on an ongoing basis. |

5.7 Risk-management: Case study for *Klebsiella pneumoniae* sepsis in a neonatal unit

During a 7-month period, seven infants in a neonatal unit developed septicaemia from multi-resistant extended spectrum β lactamase producing *Klebsiella pneumoniae*, and two babies died. Molecular typing revealed that four of the strains were identical; not all isolates were available for typing. Screening of all babies was not carried out, as it was expected that many would already be colonised, and that babies whose gut was colonised by the bacteria would be the source of infection through the hands of healthcare workers. The outbreak was brought under control by in-service education and improvement of hand hygiene compliance, and wearing of single-use gloves when babies' nappies were being changed. Nurses were declared to be the advocates for the babies, and the nurse caring for each baby was responsible for ensuring that all attending personnel perform hand hygiene before and after handling the baby, with non-compliance being reported to the infection control team.

Table A1.6. Case study for *Klebsiella pneumoniae* sepsis in a neonatal unit

| | |
|--------------------------|---|
| Eliminating risks | In this situation it is not possible to eliminate the risk entirely, so it must be managed. |
| Identifying risks | In this case, the risk has been identified as cross-transmission of <i>Klebsiella pneumoniae</i> . |
| Analysing risks | The major source of the risk is transmission between neonates by healthcare workers' hands, with failure to wear gloves when changing nappies and a lack of appropriate hand hygiene practices by some staff members. In addition, review of the cleaning of frequently touched surfaces in the neonatal intensive care unit to minimise the risk of environmental contamination. |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response. |
| Treating risks | <u>Immediate measures</u> include review of the application of standard precautions and the implementation of contact precautions, with strict enforcement of hand hygiene, appropriate wearing of personal protective equipment (e.g. gloves), changing of gloves between tasks with the same patient and performing hand hygiene after gloves are removed, and provision of in-service education on hand hygiene. |

| | |
|-------------------|--|
| | <u>Longer-term measures</u> might include review of increased frequency of environmental cleaning especially of frequently touched surfaces, performance of surveillance cultures, and cohorting of colonised babies if the outbreak could not be brought under control by immediate measures. |
| Monitoring | Changes in rates of infection could be monitored through ongoing surveillance. |

5.8 Risk-management: Case study for influenza in a long-term care facility

A cluster of cases of confirmed influenza occurred in a long-term care facility, which were observed after a group activity involving dancing was held in the dining room prior to the midday meal. It was observed that a number of residents who had been unwell had attended the group activity and had sat at the dining tables. Due to the lack of waste receptacles in the dining room, used tissues were placed on the dining room tables. It was also noticed that a number of residents remained in the vicinity of the dining room post activity as their rooms were a short distance from the dining room. The shared bathrooms were at the other end of the corridor so it was not known whether hand hygiene was performed prior to meals or the event. Residents reported signs and symptoms consistent with influenza at least two days following the event, which was later confirmed by rapid diagnostic test from four patients and two staff members. The vaccination coverage of the staff was 41.7%.

None of the staff members with influenza symptoms who had assisted in the group activities had been immunised.

Table A1.7. Case study for influenza in a long-term care facility

| | |
|--------------------------|---|
| Eliminating risks | In this situation it is not possible to eliminate the risk entirely, so it must be managed. |
| Identifying risks | In this case, the risk has been identified as cross-transmission of influenza. |
| Analysing risks | <p>One opportunity for transmission is the assembling of large numbers of residents in a confined area in which close contact and droplet transmission occurred such as sneezing, coughing or talking.</p> <p>In addition, the lack of waste receptacles available would have hindered immediate disposal of infectious waste material. Healthcare workers or other residents may have had indirect contact with influenza droplets from the dirty tissues lying on the table, particularly if any of the people other than the ones who used the tissues disposed of them later and surfaces were not cleaned.</p> <p>The lack of hand hygiene facilities in the immediate vicinity could have resulted in poor hand hygiene compliance, with staff or residents not decontaminating their hands prior to eating, after sneezing or coughing. Low levels of staff immunisation also contributed to the spread of the infection.</p> |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response. |
| Treating risks | <p><u>Immediate measures</u> may include:</p> <ul style="list-style-type: none"> • waste receptacles being made available in a common area, so people can dispose of tissues immediately after use • the provision of alcohol-based hand run so residents (and staff) can decontaminate their hands prior to eating, handling food or coughing and sneezing • restricting access to the common areas or group activities for anyone who has signs and symptoms of an influenza-like illness. <p><u>Other measures</u> may include:</p> <ul style="list-style-type: none"> • education of staff and residents on the importance of hand hygiene, respiratory hygiene and cough etiquette • immunisation of residents and staff, and asking sick staff members to stay at home • education of residents in that if they feel unwell, to avoid participating in group activities until they feel better • displaying posters and signage on hand hygiene and respiratory hygiene around the facility on an ongoing basis. |

| | |
|-------------------|---|
| Monitoring | <p>Staff should continue to monitor residents for influenza-like illness (ILI) and ongoing ILI surveillance should be heightened during the influenza season.</p> <p>Immunisation rates among staff and residents could be monitored, as well as monitoring the difference in case numbers from previous influenza outbreaks and outbreaks after the measures have been put in place.</p> |
|-------------------|---|

For further guidance see [Guidelines for the Prevention, Control and Public Health Management of Influenza Outbreaks in Residential Care Facilities in Australia \[420\]](#)

5.9 Risk-management: Case study for M. tuberculosis

A young male patient (patient A) is admitted to a medical ward in a small metropolitan hospital with signs and symptoms of pneumonia. He is placed in a 2 bed room with another patient (patient B) who also has pneumonia. Patient A has just returned from an extended working holiday in Central America, spending time in Mexico, Panama and Bolivia. When he was admitted to hospital via the emergency department, he had a recent history of lower respiratory infection and had a non-productive cough that had been treated overseas with antibiotics and had improved but not resolved. When the antibiotics had been completed, he continued to be tired and still had the cough. He was reported to be a thin young man, with an insignificant medical history who was normally fit but his respiratory infection had made him tired and breathless.

When Patient A is seen by the medical officer in the ward, they ordered a chest x-ray and morning sputum culture for atypical bacterial and viral screening as the patient had a cough. Patient A is taken to x-ray for his chest x-ray and the nursing staff collect the sputum culture the following morning and send it to laboratory. Standard precautions are used for all care provided and Patient A mobilises freely around the clinical area. The laboratory contacts the medical officer that afternoon, to advise the patient's sputum has shown high count of Acid Fast Bacilli (AFBs) when the sputum was examined by microscopy. The provisional diagnosis was pulmonary tuberculosis and the medical officer notified the ward to place the patient in a single room with negative pressure and apply airborne

precautions.

The ward notify Infection Control Team who review the patient, their history, their placement since admission and implement the plan for management of an exposure to a case of tuberculosis (TB) by staff and patients.

Table A1.8. Case study for M. tuberculosis

| | |
|--------------------------|--|
| Eliminating risks | In this situation it is not possible to eliminate the risk entirely, so it must be managed. |
| Identifying risks | In this case, the risk has been identified as possible exposure of healthcare workers and patients to a case of <i>M. tuberculosis</i> from a single patient attending the emergency department and a number of areas within the hospital. |
| Analysing risks | <p>The sources of risk are a failure to consider the possibility of pulmonary tuberculosis and drug resistant pulmonary tuberculosis that occurs in any parts of the world today. Delays in reviewing the placement of the patient when history is taken and tests are requested. These resulted in a lack of transmission-based precautions applied to Patient A, and a source of risk to subsequent patients and healthcare workers who had contact with Patient A. Placing like diagnoses in the same room occurs but it is important to consider that not all pneumonia like illnesses are the same.</p> <p>With cases such as this, it is possible that if this patient had a drug resistant pulmonary tuberculosis this could take an extended time to identify and they could be hospitalized for many months and the social and psychological risks also need to be addressed for the patient being kept in airborne precautions for extended periods.</p> <p>The laboratory reported the microscopy result as smear positive. Culture will take an extended period and should not be waited for to evaluate and respond to the risk for identification of contacts.</p> |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring an immediate response. |
| Treating risks | <p><u>Immediate measures</u> may include advising the patient of their probable diagnosis and what measures are required to care for them and protect other patients and healthcare workers. This includes placing Patient A in a negative pressure single room with airborne precautions applied in addition to standard precautions. Implementing a care plan to minimise risk of exposure to other patients and healthcare workers when tests or other department (x-ray) visits are required. State/territory TB services would also need to be notified, as they would assist in the development of a management plan including to identify contacts outside the hospital.</p> <p><u>Longer-term measures</u> could include implementation of baseline risk factors that may need to have TB screening for and protocols to assist with early diagnosis of active disease. Further measures would include increasing awareness of tuberculosis generally, educating staff about identifying the high-risk patients for a particular facility, and development of specific protocols, such as 'cough protocols'.</p> |
| Monitoring | Ongoing surveillance would assist in reducing the risk of subsequent outbreaks. Retrospective review and screening of other contacts and laboratory typing of <i>M. tuberculosis</i> isolates to identify unrecognised, linked transmission could also inform future actions. |

5.10 Risk-management: Case study for M. tuberculosis among immunocompromised patients attending outpatient services

An investigation into the healthcare-associated transmission of *M. tuberculosis* followed reports of two epidemiologically linked patients (Patient 1 and Patient 2) with haematologic malignancies and active pulmonary tuberculosis (TB). Subsequently it was found that four oncology patients had spent more than an hour in the same room as Patient 1. Patient 1's pulmonary TB was not diagnosed for 3 months as clinical findings were attributed to lower respiratory tract infection from other infectious agents or adverse effects of oncology treatments. Patient 1 was not placed on airborne precautions during this period. The investigation found that delayed TB diagnosis in Patients 1 and 2 ultimately resulted in the transmission of *M. tuberculosis* to 19 patients and staff at three hospitals and a

residential facility.

Source: Based on Malone et al (2004)

Table A1.9. Case study for *M. tuberculosis* among immunocompromised patients attending outpatient services

| | |
|--------------------------|---|
| Eliminating risks | In this situation it is not possible to eliminate the risk entirely, so it must be managed. |
| Identifying risks | In this case, the risk has been identified as cross-transmission of <i>M. tuberculosis</i> from a single patient attending a number of outpatient facilities. |
| Analysing risks | The sources of risk are a failure to consider the possibility of tuberculosis and delays in screening and diagnostic tests. These resulted in a lack of transmission-based precautions applied to Patient 1, and a source of risk to subsequent patients. |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response. |
| Treating risks | <u>Immediate measures</u> may include avoidance of potential exposures in outpatient settings, implementation of airborne precautions and treatment of febrile, coughing patients with pulmonary TB. State/territory TB services would also need to be notified, as they would assist in the development of a management plan. <u>Longer-term measures</u> could include implementation of baseline TB screening for immunocompromised patients and protocols to assist with earlier diagnosis of active disease. Further measures would include increasing awareness of tuberculosis generally, educating staff about identifying the high-risk patients for a particular facility, and development of specific protocols, such as 'cough protocols'. |
| Monitoring | Ongoing surveillance would assist in reducing the risk of subsequent outbreaks. Retrospective review and screening of other contacts and laboratory typing of <i>M. tuberculosis</i> isolates to identify unrecognised, linked transmission could also inform future actions. |

5.11 Examples of how to perform aseptic technique

Aseptic technique for peripheral and central access intravenous therapy (IV)

Typically, IV maintenance procedures will be assessed as requiring aseptic technique with the employment of a main general aseptic field and critical micro aseptic fields.

Table A1.10. Aseptic technique for peripheral and central access IV

| | |
|----------|---|
| 1 | Perform hand hygiene This will interrupt any potential transmission of infection from the clinical ward environment to the clean preparation area/room. Effective hand hygiene is vital to reduce the risk of contaminating key parts/sites. |
| 2 | Use a clean tray or trolley Such a tray or trolley provides a sufficiently large, robust and controlled working area. Reprocess re-usable trays according to local policy. |
| 3 | While the tray is drying, gather equipment Hands are contaminated when gathering equipment from storage cupboards etc. It's important therefore to gather all equipment before performing hand hygiene at Step 4. Gathering equipment at this point also allows the tray to dry properly and saves a little time. |
| 4 | Perform hand hygiene This occurs immediately before assembly of equipment and the preparation of drugs. This way, hands are optimally clean prior to glove application and aseptic technique key part manipulation. |
| 5 | Apply gloves (use sterile gloves if you must touch key parts) |

| | |
|-----------|--|
| | <p>Gloves should be worn as per standard precautions (to protect the HCW from blood or body fluid exposure) or to protect the HCW from exposure to some medicines, such as cytotoxic chemicals [437]. All peripheral and central access IV procedures should be performed without touching key parts or key sites [438]. In most cases, non-sterile gloves are appropriate, however, if there is a risk of contaminating 'key parts' or 'key sites', the HCW should select sterile gloves [438].</p> |
| 6 | <p>Assemble equipment and prepare medications—protect key parts using aseptic technique</p> <p>Aseptic technique is the most important component of aseptic practice because a key part cannot be contaminated directly if it is not touched. Key parts should be protected throughout the procedure when they are not in use. This can be achieved by using sterilised IV bungs or the inside of syringe packets. Both systems provide critical micro aseptic fields around the key part.</p> |
| 7 | <p>User assessment:</p> <p>If gloves become contaminated – remove gloves – decontaminate hands and re-glove</p> <p>This is necessary when it is not possible to proceed from preparation to administration without contaminating gloved hands (e.g. due to prepping a patient).</p> <p>If gloves remain uncontaminated between steps 6 & 7 proceed directly to step 7</p> <p>Where it is possible to retain the asepsis of gloved hands between preparation and administration, the user does not need to decontaminate hands between administration and preparation.</p> <p>This will promote compliance and save time.</p> |
| 8 | <p>Clean key parts</p> <p>70% alcohol wipes are the application of choice [274]. In addition, the benefit of using friction and allowing key parts to dry has been demonstrated [414].</p> <p>Method:</p> <p>A large 70% alcohol wipe should be fully unfolded to provide a suitable working surface area.</p> <p>One side of the wipe should be exposed to the user's gloved hand, the other side should be introduced to the hub.</p> <p>The port tip should be thoroughly wiped hard for 5 seconds—to create friction.</p> <p>This should be repeated 4 times using different parts of the tissue (to remove dirt from the tip). After cleaning the hub clean the sides of the port and line, working away from the port tip.</p> <p>Allowing the hub to air dry promotes asepsis.</p> <p>This technique provides the required level of friction. Using different parts of the wipe ensures any dirt is transferred from the hub to the wipe. The hub must dry before use otherwise it won't be aseptic (if organisms have remained, a wet tip will facilitate their transportation into the patient on injection).</p> |
| 9 | <p>Administer medication using aseptic technique</p> <p>Key parts cannot be contaminated by contact if they are not touched. Aseptic technique should therefore be used even if the user is wearing sterile gloves (because once sterile gloves are open to air they are no longer sterile, and can also be inadvertently contaminated by touch). If necessary, a small sterilised towel can be placed under a patient's line to promote safe handling.</p> |
| 10 | <p>Dispose of sharps and equipment then dispose of gloves</p> <p>Sharps are best disposed of at the bedside if possible (on the basis that the quicker they are disposed of the less chance there is of an accident).</p> |
| 11 | <p>Clean tray</p> <p>Re-usable trays are reprocessed at the end of the procedure to prevent cross infection between patients and staff. Trays are reprocessed according to local policy.</p> |
| 12 | <p>Perform hand hygiene</p> <p>It is essential that the post-procedure hand hygiene is performed immediately after glove removal, i.e. before contact with the environment (because gloves encourage the hands to sweat-out organisms from the skin).</p> |

Aseptic technique for wound care

Wound care procedures are highly variable. Typically, a critical main aseptic field is employed and practice is dictated accordingly.

Table A1.11. Aseptic technique for wound care

| | |
|-----------|--|
| 1 | With clean hands clean trolley surfaces Clean surface according to local policy to reduce the risk of aseptic field contamination. |
| 2 | Gather dressing pack and equipment, place on bottom shelf Hands are contaminated when gathering equipment from storage cupboards etc. It's important therefore to gather all equipment before the next performance of hand hygiene. Gathering equipment at this point also allows the trolley to dry properly and saves a little time. |
| 3 | Perform hand hygiene This occurs immediately before assembly of the aseptic field drape and equipment etc. in order to promote asepsis. |
| 4 | Open pack, place drape on top shelf and position waste bag. |
| 5 | Assemble equipment and position onto top shelf, protecting key parts. |
| 6 | Apply non-sterile gloves Non-sterile gloves are indicated because Steps 7 and 8 do not involve the touching of key sites or key parts. |
| 7 | Position a paper towel or drape under the wound This will promote asepsis and help protect the surrounding environment from contamination. |
| 8 | Remove dressing, expose wound and dispose of dressing into waste bag Disposing of the dressing here limits the movement of contaminated waste, helping to protect the wider clinical or community environment. |
| 9 | Perform hand hygiene Steps 7 and 8 are 'dirty' procedures and hand hygiene will promote asepsis. |
| 10 | Apply sterile gloves Although not essential for some small, minor dressings, sterile gloves at this stage will help promote asepsis of the wound. NB: Sterile gloves are essential at this stage if the wound requires direct touching with gloved hands. |
| 11 | Clean wound To help protect the wound from colonisation or infection. |
| 12 | Dress wound To help protect the wound from colonisation or infection. |
| 13 | Dispose of equipment, waste and gloves Folding the used equipment and waste into the aseptic field drape and disposing of it in the attached waste bag will minimise the movement of waste and protect the wider working environment. |
| 14 | Clean trolley surfaces Cleaning according to local policy will prevent cross infection. |
| 15 | Perform hand hygiene This will help break any chain of potential cross infection. |

5.12 Risk-management: Case study for norovirus outbreak in a long-term care facility

A resident in an aged care home where high and low care residents live reports they have been unwell for over 24 hours with diarrhoea and during this time they have attended the communal dining room as well as group activities. The infectious agent involved is identified as norovirus. The aged care home has a policy in place which requires the immediate implementation of contact precautions in addition to standard precautions. When residents are symptomatic and may have an increased risk of generating airborne droplets, carers should use a mask to minimise risk of contact with droplets generated. The issue for the aged care home is that they only have a communal dining area for meals and do not have the staff to monitor residents' meals in their rooms. The aged care home assesses all the residents and identifies a further two residents who have diarrhoea and/or vomiting. All residents who are symptomatic are confined to their rooms with ensuite facilities.

Healthcare workers from the aged care home pay particular attention to hand hygiene and appropriate use of personal protective equipment. No further cases are identified. Investigation reveals low levels of hand hygiene among residents in aged care facilities. An education program is developed and provided to assist in preventing further infections. Cleaning staff, food services and general care staff are notified to report any signs and symptoms of gastroenteritis. Increased cleaning of frequently touched surfaces is implemented to reduce environmental contamination and risk of transmission.

Table A1.12. Case study for norovirus outbreak in a long-term care facility

| | |
|--------------------------|---|
| Eliminating risks | In this situation, it is not possible to eliminate risk, so it must be managed. |
| Identifying risks | In this case, the risk has been identified as cross-transmission of norovirus by contact (faecal-oral) transmission or by the droplet route if droplets are generated during the symptomatic stage. |
| Analysing risks | One source of the risk is the lack of appropriate hand hygiene practices by some residents. Each time there is social contact between these and other residents, there is potential for cross-transmission. Depending upon hand hygiene practices among residents more broadly, there is potential for the infection to spread through the aged care home. Healthcare workers and visitors are also at risk of cross-contamination. |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response. |
| Treating risks | <p><u>Immediate measures</u> may include:</p> <ul style="list-style-type: none"> • ensure all hand washing facilities have soap, water and disposable hand towel available and encourage hand washing • raising residents' awareness of the highly transmissible nature of norovirus infection, its modes of transmission and the particular need for hand hygiene practices • for healthcare workers, the following hand hygiene practices are recommended: <ul style="list-style-type: none"> ◦ If gloves have not be worn, use soap and water to facilitate the mechanical removal of spores. After washing, hands should be dried thoroughly with a single-use towel ◦ If gloves have been worn, a lower density of contamination of the hands would be expected and alcohol-based hand rub remains the agent of choice for hand hygiene • exclude any symptomatic residents from the common dining area and group activities until they are recovered • implement contact precautions for the residents who are symptomatic. Closely monitor the other residents for signs and symptoms of norovirus. <p><u>Longer-term measures</u> may include:</p> <ul style="list-style-type: none"> • providing education to residents and visitors on hand hygiene and other infection control measures • education for healthcare workers to raise awareness of the high transmissibility of norovirus, and its capacity to spread very rapidly within aged care homes where there are poor or inadequate hygiene practices among residents and staff • visitors should be requested not to enter the facility if they are unwell. |
| Monitoring | Changes in practice could be evaluated by surveying residents on hand hygiene practice and |

| | |
|--|--|
| | reviewing the availability and use of soap and/or alcohol-based hand rubs. |
|--|--|

5.13 Risk-management: Case study for management of confirmed case of carbapenemase-producing Enterobacterales (CPE)

A liver transplant patient residing in a two-bed room at a large tertiary hospital is screened for CPE as part of the hospital's CPE surveillance program. The local infection control team was notified by the laboratory that the patient had CPE, and the patient was transferred immediately to a single room. The date of likely acquisition of CPE for the patient could not be determined and as such, the period of transmission risk was considered to be from one month prior to the date of isolation of CPE, until the patient was identified and placed into the single room.

Adapted from the Victorian guideline on carbapenemase-producing Enterobacteriaceae [421] and the Australian Commission on Safety and Quality in Health Care Recommendations for the control of carbapenemase-producing Enterobacteriaceae [231].

Table A1.13. Case study for management of confirmed case of CPE

| | |
|--------------------------|---|
| Eliminating risks | In this situation, it is not possible to eliminate risk, so it must be managed. |
| Identifying risks | In this case, the risk has been identified as a known case of CPE that may have been transmitted to other patients in the healthcare facility. |
| Analysing risks | One source of the risk is that the patient was sharing a room with multiple other patients. The infected patient is identified as high-risk due to faecal incontinence and wandering behaviours. |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response. |
| Treating risks | <p><u>Immediate measures</u> include: placing the patient in a single patient room with their own bathroom; implementing contact precautions including the appropriate wearing of PPE; where possible, allocate clinical equipment to this patient for the duration of the admission or if not possible, ensuring shared clinical equipment is appropriately cleaned and disinfected before re-use; and limiting patient movement and intensifying routine environmental cleaning, including daily disinfection of the patient room and bathroom and twice daily disinfection of frequently-touched surfaces. All healthcare workers should be provided with education about infection prevention and control strategies for CPE.</p> <p><u>Other measures</u> include conducting a lookback investigation to identify all patients who had shared a room, bathroom or toilet facilities with the infected patients for more than 24 hours. Any patient who is still in the healthcare facility should be placed into single rooms with contact precautions, screened for CPE and provided with information about CPE. Any patient who has since been discharged or transferred to another facility should be sent a letter advising them to seek screening for CPE, and the facility or general practitioner should be notified. An alert should be placed in their medical history so they can be placed into contact precautions and screened if they are readmitted within 12 months.</p> |
| Monitoring | <p>The healthcare facility should implement a surveillance program to monitor the development of transmission of CPE. Healthcare workers adherence to infection prevention strategies should also be monitored.</p> <p>The healthcare facility can also review and monitor their antimicrobial prescription/use trends and use audit systems to identify inappropriate antimicrobial use.</p> |

5.14 Risk Management: Case study for vancomycin-resistant enterococci (VRE) outbreak in a large tertiary-care referral hospital

Two months after the first index case of VRE was detected in the intensive care unit of a large teaching hospital, 68 patients had become either infected or colonised with an epidemic strain of vanB vancomycin-resistant Enterococcus faecium, despite standard infection control procedures. Subsequently, 169 patients in 23 wards were found to be colonised with a single strain of vanB vancomycin-resistant E. faecium. Introducing additional control measures rapidly brought the outbreak under control. Hospital-wide

screening found 39 previously unidentified colonised patients, with only 7 more non-segregated patients being detected in the next 2 months. The outbreak was terminated within 3 months due to a well-resourced, multifaceted approach.

Source: Based on Christiansen et al [232]

Table A1.14. Case study for VRE outbreak in a large tertiary-care referral hospital

| | |
|--------------------------|---|
| Eliminating Risks | In this situation, it is not possible to eliminate risk immediately, so it must be managed. |
| Identifying Risks | In this case, the risk has been identified as cross-transmission of VRE. |
| Analysing Risks | The source of the risk is multidrug resistance coupled with a vulnerable patient population (intensive care unit). Each time there is contact with an infected patient there is potential for cross-transmission to the healthcare workers and/or other patients. |
| Evaluating Risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring immediate response. |
| Treating Risks | <p><u>Immediate measures</u> to control the outbreak may include:</p> <ul style="list-style-type: none"> • formation of a VRE executive group • rapid laboratory identification (30 to 48 hours) using culture and polymerase chain reaction detection of vanA and vanB resistance genes • screening of hospitalised patients with isolation of patients and cohorting of contacts • increased cleaning • electronic flagging of medical records of contacts • antibiotic restrictions (third-generation cephalosporins and vancomycin). <p><u>In the longer-term</u>, hospital policies may be changed to restrict antibiotic use, institute targeted screening and increase environmental cleaning efficiency and frequency.</p> <p>These measures are relevant to a recent outbreak in an area of low endemicity. Some of these approaches may also be relevant in an area of high endemicity.</p> |
| Monitoring | Repeated screening would identify whether the outbreak recurred. |

5.15 Risk-management: Case study for infection prevention during renovation of emergency department

An emergency department in a state healthcare facility requires renovation. The work involves construction of new walls and major cabling activities, so will result in a moderate to high level of dust generation.

Source: based on the Australasian Health Facility Guidelines [394]

Table A1.15. Case study for infection prevention during renovation of emergency department

| | |
|--------------------------|---|
| Eliminating risks | In this situation, it is not possible to completely eliminate risks, so they must be managed. |
| Identifying risks | In this case, the risk will be varied and the activities where there is interruption of ceiling space will have higher risks than others. Use a risk matrix for determining the risk for each activity in renovation and construction. The risk has been identified as invasive Aspergillosis which may be increased by the renovation activities and infect immunosuppressed patients. |
| Analysing risks | The extent of the risk may depend upon the nature of the renovation, the location of the patient population in relation to the renovation site, the type of ventilation systems in place and the identification of possible contaminants (ceiling dust, service shafts etc.) and their locations. |
| Evaluating risks | The balance of likelihood and consequences identify this as a 'very high risk' situation requiring action to mitigate risk being implemented during the planning phase of this project and a system for immediate response to any risks identified during the work. |
| Treating risks | <p>Measures to prevent infections during the renovation period include:</p> <ul style="list-style-type: none"> • establishing a multidisciplinary team including infection control experts and project staff who should be consulted early in the planning phase and then during all stages of the project |

| | |
|------------------|--|
| | <ul style="list-style-type: none"> • induction program to address infection risks for all contractors • ensure all healthcare and construction staff are aware of the necessary contact and airborne precautions as well as relevant workplace health and safety requirements for training and personal protective equipment • completing all renovation barriers before the renovation begins • wet mop or vacuum frequently during periods of renovation activity to minimise tracking • place a dust-mat at the entrances and exits to all work areas • monitoring dust levels in work area and in adjacent areas and where high risk patient services are located • monitor movement of materials and workers into and out of the construction/renovation area. |
| Monitoring risks | Construction workers compliance with procedures can be monitored. Air and water sampling can occur. |

6. APPENDIX 2: Supplementary information

Summary

The following supplementary information provides further background information on infection prevention and control in healthcare.

This section covers:

- **6.1** — Recommended routine cleaning frequencies
- **6.2** — Checklist of standard precautions for procedures
- **6.3** — Use of standard and transmission-based precautions
- **6.4** — Type and duration of precautions for specific infections and conditions
- **6.5** — Collection of pathology specimens
- **6.6** — Allowing animals into healthcare facilities
- **6.7** — Mechanisms of antibacterial action of antiseptics and disinfectants
- **6.8** — Summary table of relevant infection prevention and control resources
- **6.9** — Research gaps

6.1 Recommended routine cleaning frequencies

The following tables (A2.1 **Risk categories for environmental cleaning** and A 2.2 **Minimum cleaning frequency**) include examples of risk categories for environmental cleaning and suggested cleaning frequencies for equipment and environmental surfaces in health service organisations. Health service organisations should use their local risk management framework to determine individual and collective infection risks and to identify priorities and develop management options to reduce the risk of infection.

The methods, thoroughness and frequency of cleaning and the products used for different surfaces should be determined following a risk assessment and specified in local policies and procedures. For guidance on cleaning of spills, see Section 3.1.3 *Recommendation 12 practical information*.

Table A2.1. Risk categories for environmental cleaning

| Risk category | Examples |
|----------------------------|---|
| | <i>These examples are inclusive of any adjoining area, for example, bathrooms, corridors, storerooms, meeting rooms, offices, pan rooms and staff lounges.</i> |
| Very high / Extreme | <ul style="list-style-type: none"> • Operating theatres • Day procedure areas • Medical imaging procedural areas • Intensive care units • Clinical areas with patients in isolation or with patients who are immunosuppressed • Emergency departments • Level 2 and 3 nurseries • Labour and delivery wards |
| High | <ul style="list-style-type: none"> • General wards • Level 1 nurseries • Pharmacy clean areas • Mortuaries, mortuary fridges and body holding rooms • Outpatient clinics and treatment rooms • Sterile stock storage (outside of central sterilising services departments and sterile supply areas) • Emergency ambulances and other rescue vehicles |
| Medium | <ul style="list-style-type: none"> • Day activity areas • Rehabilitation areas, including hydrotherapy • Outpatient clinics, including consulting rooms, ambulatory care • Residential accommodation • Offices in patient/clinical areas • Medical imaging • Pathology and other laboratories |

| | |
|------------|---|
| | <ul style="list-style-type: none"> • General pharmacy • Kitchenette/pantry/other food preparation or storage areas • Main foyer • Cleaners room • Non-emergency patient transport vehicles • All ambulance operational support vehicles |
| Low | <ul style="list-style-type: none"> • Office/administration areas |

Source: Table adapted from:

SA Health. Cleaning Standard for South Australian Healthcare Facilities. [Online] Adelaide: SA Health; 2021 [cited 12 September, 2022]; Available from: [Cleaning Standards \(sahealth.sa.gov.au\)](https://www.sahealth.sa.gov.au)

Clinical Excellence Commission. Cleaning of the Healthcare Environment. [Online] Sydney Clinical Excellence Commission; 2020 [cited 12 September, 2022]; Available from: [Cleaning of the Healthcare Environment \(nsw.gov.au\)](https://www.nsw.gov.au)

Table A2.2. Recommended cleaning frequency

Note: The choice of disinfectant is dependent upon the local epidemiology and a local risk assessment.

| Element | Very high risk/ Extreme risk | | | | Method |
|--|---|---|---|---|---|
| | (e.g. outbreaks of <i>Candida auris</i> or carbapenemse-producing <i>Enterobacterales</i>) | High risk | Medium Risk | Low risk | |
| Alcohol-based hand rub dispenser, bedside | Clean daily | Clean daily | Clean daily | Clean weekly | Detergent |
| Alcohol-based hand rub dispenser, not in patient/ treatment rooms | Clean daily | Clean daily | Clean daily | N/A | Detergent |
| Bath | Clean daily & spot/check clean once daily | Clean daily & spot/check clean once daily | Clean daily & spot/check clean once daily | Clean daily & spot/check clean once daily | Detergent |
| Bed | Clean frame daily | Clean frame daily | Clean frame daily | When visibly soiled & whole on discharge | Detergent |
| | Clean underneath weekly | Clean underneath weekly | Clean underneath weekly | | Detergent + disinfectant for multi-resistant organism (MRO) |
| | Clean whole on discharge | Clean whole on discharge | Clean whole on discharge | | |
| Bed rails | Clean twice daily & after discharge | Clean daily & after discharge | Clean daily & after discharge | Clean weekly & after discharge | Detergent |
| | | | | | Detergent + disinfectant for MRO |
| Bedside table | Clean twice daily & after use | Clean daily & after use | Clean daily | Clean weekly | Detergent |
| | | | | | Detergent + disinfectant for MRO |
| Bidet | Clean three times daily | Clean three times daily | Clean daily | Clean daily | Detergent and disinfectant |
| Blood pressure cuff | Clean after use | Clean after use | Clean after use | Clean after use | Detergent |
| Call bell | Clean Daily | Clean Daily | Clean Daily | Clean Weekly | Detergent |
| | Clean after discharge | Clean after | Clean after | Clean after | |

| Element | Very high risk/ Extreme risk | | | | Method |
|--|---|------------------------------------|---|---|---|
| | (e.g. outbreaks of <i>Candida auris</i> or carbapenamse-producing <i>Enterobacterales</i>) | High risk | Medium Risk | Low risk | |
| Carpet (soft floor) | Clean twice daily | discharge Clean daily | discharge Clean daily | discharge Clean weekly | Vacuum with high efficiency particulate air filter |
| | Clean 6-monthly | Clean 6-monthly | Clean annually | Clean annually | Steam clean (or shampoo) |
| Catheter stand/bracket | Clean daily & after use | Clean daily & after use | Clean before initial use, after use & monthly | Clean before initial use, after use & monthly | Detergent |
| Ceiling | Spot clean daily & wash yearly | Spot clean daily & wash yearly | Spot clean weekly & wash yearly | Spot clean monthly & wash every 3 years | Detergent/Damp dust |
| Chair | Clean twice daily | Clean twice daily | Clean daily | Clean weekly | Detergent Detergent + disinfectant for MRO |
| Chair, dental and surrounds | N/A | N/A | N/A | Clean daily & when visibly soiled | Detergent |
| Cleaning equipment | Clean after use | Clean after use | Clean after use | Clean after use | Detergent Detergent + disinfectant for MRO |
| Clipboard | Clean daily & between patient use | Clean daily & between patient use | Clean daily & between patient use | Clean weekly | Detergent |
| Commode | Clean contact points after use | Clean contact points after use | Clean contact points after use | Clean contact points after use | Detergent Detergent + disinfectant for MRO |
| | Clean whole daily | Clean whole daily | Clean whole daily | Clean whole weekly | |
| Computer & keyboard (general ward use, non-mobile, located outside patient area) | Clean twice daily or when visibly soiled | Clean daily or when visibly soiled | Clean daily or when visibly soiled | Clean weekly or when visibly soiled | Manufacturer's recommendations Install keyboard covers or washable keyboards where feasible Detergent |
| Computer & keyboard (used and/or located in close proximity to patient e.g. patient bay or room) | Clean twice daily or when visibly soiled | Clean daily or when visibly soiled | Clean daily or when visibly soiled | Clean weekly or when visibly soiled | Manufacturer's recommendations |
| | Clean between patients | Clean between patients | Clean between patients | Clean between patients | Install keyboard covers or washable keyboards where feasible |
| | Clean after discharge | Clean after discharge | Clean after discharge | Clean after discharge | |

| Element | Very high risk/ Extreme risk | | | | | Method |
|---|---|---|--|--|--|--|
| | (e.g. outbreaks of <i>Candida auris</i> or carbapenamse-producing <i>Enterobacterales</i>) | High risk | Medium Risk | Low risk | | |
| Curtains and blinds Disposable curtains and blinds | Change as per the manufacturer's instructions for use. | Change as per the manufacturer's instructions for use. | Change as per the manufacturer's instructions for use. | Change as per the manufacturer's instructions for use. | Change as per the manufacturer's instructions for use. | Detergent Follow the manufacturer's recommendations. |
| Curtains and blinds Non-disposable curtains and blinds | Change upon discharge, when visibly soiled or damaged. Change or clean weekly and upon discharge of patient. Change when: <ul style="list-style-type: none"> visibly soiled damaged terminal clean is required | Change when visibly soiled or damaged. Change or clean monthly . Change when: <ul style="list-style-type: none"> visibly soiled damaged terminal clean is required | Change when visibly soiled or damaged. Change or clean biannually . Change when: <ul style="list-style-type: none"> visibly soiled damaged terminal clean is required | Change when visibly soiled or damaged. Change or clean annually . Change when: <ul style="list-style-type: none"> visibly soiled damaged terminal clean is required | Change when visibly soiled or damaged. Change or clean annually . Change when: <ul style="list-style-type: none"> visibly soiled damaged terminal clean is required | Replace with laundered curtains or steam clean while in place. Follow the manufacturer's recommendations. |
| Door knob/handle, general | Clean twice daily | Clean daily | Clean daily | Clean weekly | Clean weekly | Detergent |
| Door knob/handle, patient room | Clean twice daily | Clean daily | Clean daily | Clean daily | Clean daily | Detergent Detergent + disinfectant for MRO |
| Drip/intravenous stands | Clean contact points after use | Clean contact points after use | Clean contact points after use | Clean contact points after use | Clean contact points after use | Detergent Detergent + disinfectant for MRO |
| Fan, patients | Clean daily & between patient use | Clean daily & between patient use | Clean daily & between patient use | Clean weekly & between patient use | Clean weekly & between patient use | Detergent |
| Floor, non-slip | Damp mop twice daily | Damp mop twice daily | Damp mop daily | Damp mop daily | Damp mop daily | Detergent Detergent + disinfectant for MRO |
| Floor, polished | Dust removal & clean twice daily | Dust removal & clean daily | Dust removal & clean daily | Dust removal & clean weekly | Dust removal & clean weekly | Detergent for routine Consider electrostatic mops Detergent + disinfectant for MRO |
| Fridges | Weekly & defrost as required | Weekly & defrost as required | Monthly & defrost as required | Monthly & defrost as required | Monthly & defrost as required | Detergent |

| Element | Very high risk/ Extreme risk | | | | Method |
|---|---|--|--|--|--|
| | (e.g. outbreaks of <i>Candida auris</i> or carbapenamse-producing <i>Enterobacterales</i>) | High risk | Medium Risk | Low risk | |
| Fridge (drug) | Three times daily spot check—clean when necessary Clean weekly | Daily spot check—clean when necessary Clean weekly | Daily spot check—clean when necessary Clean weekly | Daily spot check—clean when necessary Clean weekly | Detergent |
| Glazing, internal (incl. partitions) | Spot clean daily & full clean weekly | Spot clean daily & full clean weekly | Spot clean daily & full clean weekly | Clean weekly | Detergent |
| Hoist, bathroom | Clean contact points after use | Clean contact points after use | Clean contact points after use | Clean contact points after use | Detergent |
| Drip/IV stand & poles | Clean daily & clean contact points after use | Clean daily & clean contact points after use | Clean contact points after use | Clean contact points after use | Detergent + disinfectant for MRO |
| Light switch | Clean daily | Clean daily | Clean weekly | Clean weekly | Detergent |
| Bedside locker | Clean contact points twice daily | Clean contact points twice daily | Clean contact points daily | Clean contact points weekly | Detergent + disinfectant for MRO |
| Manual handling (I.e. hoists) | Clean contact points after use | Clean contact points after use | Clean contact points after use | Clean contact points after use | Detergent + disinfectant for MRO |
| Mattress | Clean when visibly soiled/ bodily fluids & after discharge | Clean when visibly soiled/ bodily fluids & after discharge | Clean when visibly soiled/ bodily fluids & after discharge | Clean when visibly soiled/ bodily fluids & after discharge | Detergent + disinfectant for MRO Preferable that entire mattress has waterproof cover |
| Medical equipment (e.g. IV infusion pumps, pulse oximeters) NOT connected to a patient | Clean daily (when in use) between patient use | Clean daily (when in use) & between patient use | Clean daily (when in use) & between patient use | Clean weekly (when in use) & between patient use | Detergent Detergent + disinfectant for MRO |
| Medical gas equipment | Clean daily | Clean daily | Clean daily | Clean weekly | Detergent Detergent + disinfectant for MRO |
| Microwave | Clean daily | Clean daily | Clean daily | Clean daily | Detergent |
| Nebuliser, portable (when in use) | Clean daily & after use | Clean daily & after use | Clean monthly & after use & before initial use | Clean bi-monthly & after use & before initial use | Detergent |
| Notes folder | Clean daily | Clean daily | Clean weekly | Clean weekly | Detergent |
| Over bed tray table (overway table) | Twice daily | Daily | Daily | Weekly | Detergent Detergent + disinfectant for MRO |
| Oxygen equipment | Clean daily & after use | Clean daily & after | Clean monthly & | Clean monthly & | Detergent |

| Element | Very high risk/ Extreme risk | | | | | Method |
|--|---|--|--|---|--|---|
| | (e.g. outbreaks of <i>Candida auris</i> or carbapenamse-producing <i>Enterobacterales</i>) | High risk | Medium Risk | Low risk | | |
| Patient slide/board | Clean daily & after use | use Clean daily & after use | after discharge & before initial use Clean monthly & after use | after discharge & before initial use Clean monthly & after use | | Detergent + disinfectant for MRO |
| Pillow (waterproof cover) | Clean when visibly soiled/ bodily substances & after discharge | Clean when visibly soiled/ bodily substances & after discharge | Clean when visibly soiled/ bodily substances & after discharge | Clean when visibly soiled/ bodily substances & after discharge | | Detergent + disinfectant for MRO |
| Sharps bin trolley | Clean daily | Clean twice weekly | Clean weekly | Clean monthly | | Detergent |
| Shower | Clean daily & one spot check clean daily | Clean daily & one spot check clean daily | Clean daily | Clean daily | | Detergent + disinfectant for MRO |
| Sink (hand washing) | Clean twice daily & after use | Clean twice daily & after use | Clean daily | Clean daily | | Detergent |
| Surfaces (general horizontal) in patient room e.g. ledges | Clean twice daily & spot clean after use | Clean twice daily & spot clean after use | Clean daily & after discharge | Clean weekly & after discharge | | Detergent + disinfectant for MRO |
| Telephone | Clean daily & spot clean after use | Clean daily & spot clean after use | Clean daily | Clean weekly | | Detergent |
| Toilet | Clean twice daily & spot clean after use | Clean twice daily & spot clean after use | Clean daily | Clean weekly | | Detergent + disinfectant |
| Toilet seat, raised | Clean twice daily & spot clean after use | Clean twice daily & spot clean after use | Clean monthly & before initial use & spot clean after use | Clean monthly & before initial use & spot clean after use | | Detergent for routine Detergent + disinfectant for MRO |
| Trolley, dressing | Clean utilised surfaces before & after use Clean whole trolley weekly | Clean utilised surfaces before & after use Clean whole trolley weekly | Clean utilised surfaces before & after use Clean whole trolley weekly | Clean utilised surfaces before & after use Clean whole trolley monthly | | Detergent + disinfectant for MRO |
| Trolley, linen | Clean contact points daily Clean whole trolley weekly | Clean contact points daily Clean whole trolley weekly | Clean contact points daily Clean whole trolley weekly | Clean contact points weekly Clean whole trolley monthly | | Detergent |
| Trolley, resuscitation | Clean daily | Clean twice weekly | Clean weekly | Clean monthly | | Detergent |
| TV, fixed (out of patient reach) | Clean weekly | Clean weekly | Clean weekly | Clean weekly | | Detergent |
| TV, patient bedside (mobile and within patient reach) | Clean daily & between patients | Clean daily & between patients | Clean weekly and between patients | Clean monthly & between patients | | Detergent/damp dust |
| Walls | Spot clean daily & dust weekly & full clean yearly | Spot clean daily & dust weekly & full clean yearly | Spot clean weekly & full clean yearly | Spot clean weekly & full clean yearly | | Detergent/damp dust |

| Element | Very high risk/ Extreme risk | | | | Method |
|--------------------------|---|---|---|---|----------------------------------|
| | (e.g. outbreaks of <i>Candida auris</i> or carbapenamse-producing <i>Enterobacterales</i>) | High risk | Medium Risk | Low risk | |
| Washbowl, patient | One full clean daily and between patient use | One full clean daily and between patient use | One full clean daily and between patient use | One full clean daily and between patient use | Detergent + disinfectant for MRO |
| Waste receptacle | Clean weekly & spot clean when visibly soiled/bodily substances | Clean weekly & spot clean when visibly soiled/bodily substances | Clean weekly & spot clean when visibly soiled/bodily substances | Clean weekly & spot clean when visibly soiled/bodily substances | Detergent |
| Wheelchair | Clean daily & after use | Clean daily & after use | Clean monthly & after use | Clean monthly & after use | Detergent |

Source: Adapted from National Health Service National Specifications for Cleanliness [423]

6.2 Checklist of standard precautions for procedures

This table outlines the use of standard precautions for a range of example procedures. It is assumed that there is no known or suspected infection. Decision-making about the level of protection required involves a risk assessment of the procedure to be performed; for example, usual wound irrigation is unlikely to require surgical mask and eye protection in primary care, but may be required more often in the hospital setting.

Table A2.3. Standard precautions for procedures

| Procedure | Hand hygiene | Gloves | Sterile gloves | Surgical mask | Eye protection | Gown |
|---|--------------|--|------------------------------------|--|--|--|
| Activities of daily living (washing, toilet etc.) | ✓ | — | — | — | — | — |
| Routine observations (e.g. blood pressure measurement) | ✓ | — | — | — | — | — |
| General medical examination | ✓ | ✓ For contact with broken skin/rash/mucous membrane | — | ✓ If splash risk likely | ✓ If splash risk likely | ✓ If splash risk likely |
| Wound examination/dressing | ✓ | ✓ For contact with body substances | ✓ For direct contact with wound | ✓ For wound irrigation if splash likely | ✓ For wound irrigation if splash likely | ✓ For grossly infected wounds |
| Blood glucose and haemoglobin monitoring | ✓ | ✓ | — | — | — | — |
| Vaginal delivery | ✓ | — | ✓ | — | ✓ | ✓ |
| Intravenous cannula insertion | ✓ | ✓ | — | — | ✓ If splash risk likely | — |
| Intravascular access device insertion | ✓ | — | ✓ | ✓ | ✓ | ✓ Where max. barrier precautions are used |
| Intravascular access device | ✓ | — | ✓ | — | — | — |

| Procedure | Hand hygiene | Gloves | Sterile gloves | Surgical mask | Eye protection | Gown |
|---|--------------|--------|--|------------------------------|---------------------------------|------------------------------|
| care | | | | | | |
| Surgical aseptic technique procedure (e.g. lumbar puncture) | ✓ | — | ✓ | ✓ | ✓ | ✓ |
| Insertion of urinary catheter | ✓ | — | ✓ | ✓ If exposure risk likely | ✓ If exposure risk likely | ✓ If exposure risk likely |
| Urinary catheter care | ✓ | ✓ | — | — | ✓ When emptying drainage bag | ✓ If exposure risk likely |
| Suctioning: endotracheal tube, tracheostomy | ✓ | — | ✓ Dominant hand (open suction system) | ✓ | ✓ | ✓ If exposure risk likely |
| Major dental procedure | ✓ | — | ✓ | ✓ | ✓ | ✓ |
| Routine intra-oral dental procedures | ✓ | ✓ | — | ✓ | ✓ | ✓ If exposure risk likely |
| Pulmonary function tests (for example, spirometry) | ✓ | — | — | ✓ | ✓ | — |

6.3 Use of standard and transmission-based precautions

In certain situations, the use of standard precautions alone may not be enough to limit the spread of infection. When this occurs, transmission-based precautions are required. There are three types of transmission-based precautions: contact, droplet and airborne precautions. One or more types of transmission-based precautions may be required, depending on how an infection is spread between people. See Section 6.4 for further information.

There are different personal protective equipment and patient placement requirements for standard precautions and transmission-based precautions. The table below provides a general overview of these requirements by precaution type. A local risk assessment should always be undertaken to determine the most appropriate personal protective equipment and patient placement for the delivery of care.

Table A2.4. Use of standard and transmission-based precautions

| Overview of the use of standard and transmission-based precautions | | | | | | | | | |
|--|---|---------------------------------------|---|-------|------|--------------------|--------------------------------|--|------------------------------------|
| Type of precautions | Examples of infectious agents | Patient placement | Gloves | Gowns | Mask | Protective eyewear | Handling of shared equipment | Environmental cleaning | Visitors* |
| Standard precautions | Hepatitis B, Hepatitis C, Cytomegalovirus (CMV) | No restrictions on patient placement. | PPE use: * Gloves and gowns to be worn when there is potential exposure to blood or body substances | | | | Single-use, single-patient use | Clean with neutral detergent. If surfaces are contaminated with blood or body fluids, | Hand hygiene, respiratory hygiene, |

| | | | | | | | | | |
|----------------------------|--|---|--|-----------------------------|---|---------------|--|--|--|
| | | | * Mask and protective eyewear to be worn when there is potential for blood or body substances exposure to mucosa (for example, attending to a patient who is coughing and sneezing). | | | | | cleaning should be followed with disinfection. | cough etiquette |
| Contact | Multidrug-resistant organisms, <i>C.difficile</i> , norovirus | Single room, or cohort with same strain of infectious agent. | Yes | Yes | As per standard precautions | or re-process | Neutral detergent and disinfectant are required. | As per standard precautions | |
| Droplet[^] | Norovirus, pertussis [#] , meningococcus | Single room with door open, or cohort with same strain of infectious agent. | As per standard precautions | As per standard precautions | Yes Use surgical mask | | As per standard precautions | Neutral detergent. Use disinfectant if infectious agent is a multidrug-resistant organism or in the event of an outbreak. | Restrict visitor numbers and use same precautions as staff |
| Airborne | Pulmonary TB, rubella [#] , measles [#] and chicken pox [#] | Single room with door closed. Use negative pressure room if available. | | | Use particulate respirator (P2 or N95 mask) | | | Neutral detergent. Use disinfectant if infectious agent is a multidrug-resistant organism or in the event of an outbreak. | Restrict visitor numbers and use same precautions as staff |

Notes:

Standard precautions should be used during all patient care to prevent the likelihood of transmission of infection between the patient and the healthcare worker, as well as prevent contamination of the environment.

Patient placement: Patient placement should be informed by a risk assessment followed by prioritisation of the seriousness of the infection and any competing patient needs. For further information refer to the [Patient placement guide](#) and local infection prevention and control service.

Personal protective equipment (PPE) use:

When used as part of **standard precautions**, PPE protects against potential exposure to blood and body substances.

When used as part of **transmission-based precautions**, PPE is used as a barrier against specific means of transmission of infectious agents.

Protective eyewear is part of standard precautions, and serves to protect the healthcare worker from body fluid exposure to the eyes from droplets generated by patient when coughing and sneezing.

[^] Droplets can contaminate horizontal surfaces close to the source patient, and the hands of healthcare workers can become contaminated through contact with those surfaces.

***Visitors** should be given instruction about correct procedures when transmission-based precautions are applied and given appropriate resources to support them in meeting these requirements.

[#] For vaccine-preventable diseases, where possible, only staff and visitors who have confirmed immunity (evidenced by serological immunity or vaccination history) to the specific infectious agent should enter the room. See Section 4.2.1 of the Australian Guidelines for the Prevention and Control of Infection in Healthcare for further information.

Table adapted from: Section 6.3 Use of standard and transmission-based precautions **Table A2.4. Use of standard and transmission-based precautions**, Australian Guidelines for the Prevention and Control of Infection in Healthcare

6.4 Type and duration of precautions for specific infections and conditions

The information in the table below provides a summary of diseases and the precautions which may be required by healthcare workers. Decisions regarding precautions should be based on a risk assessment performed by the Infection Control Team, and in the context of locally agreed policy relating to management of patients with specific diseases.

REMINDER: Transmission-based precautions are applied in addition to standard precautions.

S = Standard C = Contact D = Droplet A = Airborne

Table A2.5. Precautions for specific infections and conditions

| Syndrome or condition | | | | | |
|---|--------------------------------------|-----------------------------------|--|---|--|
| Syndrome or condition | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| Gastroenteritis Unknown cause | Unknown and / or under investigation | Unknown until cause is determined | S + C | Until cause identified and / or duration of illness | Continue precautions if cause is communicable. Cease precautions if non-communicable cause identified. |
| Meningitis | Unknown or under investigation | Unknown until case is determined | S + D | Until cause identified and / or duration of illness | If cause of meningitis is bacterial, S + D precautions can be ceased after 24 hours of antibiotic treatment is complete. If cause of meningitis is viral, use standard precautions. |
| Respiratory illness Unknown cause | Unknown and / or under investigation | Unknown until cause is determined | S + C + D (+ A as per additional comments) | Until cause identified and/or for duration of illness | Continue precautions if cause is communicable. Cease precautions if non-communicable cause identified. Airborne precautions if pulmonary tuberculosis suspected (see Tuberculosis). |
| Wound infections Major | Bacterial (usually) | Contact | S + C | Duration of illness | Until drainage stops or can be contained by dressing. |

| Disease or Organism | | | | | |
|--|-------------------|---|------------------------------------|--|---|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| Anthrax <i>(Bacillus anthracis)</i> Cutaneous | Bacterial | Inoculation; person-to-person transmission rare | S (+ C as per additional comments) | Duration of illness | Transmission through non-intact skin contact with draining lesions possible, therefore use Contact Precautions if large amount of uncontained drainage. Hand washing with soap and water preferable to use of waterless alcohol-based antiseptics since alcohol does not have sporicidal activity. |
| Bronchiolitis | Viral, bacterial | Contact; droplet | S + C + D | Duration of illness | Use mask according to Standard Precautions. Avoid contact between the person who is ill and infants, young children, frail and elderly people, or immunocompromised patients until the person is feeling well. |
| Burkholderia cepacia | Bacterial | Contact; droplet | S + C + D | Duration of illness | Avoid exposure to other persons with cystic fibrosis. |
| Chickenpox <i>(Varicella Virus)</i> | Viral | Airborne droplets; direct contact with fluid in blisters or nasopharyngeal secretions | S + C + A | Until all lesions dry and crusted over | Screen by history and serology; pre-employment Varicella vaccine. Post-exposure prophylaxis (vaccination, or zoster immunoglobulin (ZIG) in high risk cases and late pregnancy) may be indicated. Susceptible healthcare workers should not enter |

| Disease or Organism | | | | | |
|---|------------------------------------|----------------------|------------------------------------|---|--|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| | | | | | room if immune caregivers are available. |
| Clostridium perfringens Gas gangrene | Bacterial | Contact | S (+ C as per additional comments) | Duration of illness | Adenovirus, enterovirus, Coxsackie A24 most common. Highly contagious; outbreaks in eye clinics, paediatric and neonatal settings, institutions. Eye clinics should follow Standard Precautions when handling patients with conjunctivitis. |
| Conjunctivitis caused by the organisms: Adenovirus Enterovirus Coxsackie A24 | Viral | Contact | S + C | Duration of illness | Highly contagious; outbreaks in eye clinics, paediatric and neonatal settings, institutions. |
| Diphtheria (<i>Corynebacterium diphtheriae</i>) | Bacterial Cutaneous infection | Contact | S + C | All wounds are clinically improving and can be covered by an occlusive waterproof dressing or have healed; and 72 hours of appropriate antimicrobial therapy has been completed. | Diphtheria is notifiable to public health. |
| | Bacterial Respiratory infection | Respiratory droplets | S + C + D | Droplet and contact precautions should remain until two negative nasopharyngeal and throat cultures taken at least 24 hours apart and more than 24 hours after the cessation of appropriate antimicrobial therapy are obtained. | (Source: Diphtheria control guideline) |
| Gastroenteritis caused by the following organisms: | | | | | |
| <i>Clostridioides difficile</i> | Bacterial | Ingestion; contact | S + C | Duration of illness | Discontinue antibiotics if appropriate. Do not share electronic thermometers; ensure consistent environmental cleaning and disinfection. Hypochlorite solutions may be required for cleaning if transmission continues. Use ABHR following glove removal. Wash hands with soap and water if gloves not worn and hands become soiled. |
| Adenovirus | Viral | Ingestion; contact | S (+ C as per additional comments) | Duration of illness | Use Contact Precautions for nappied or incontinent persons for the duration of illness or to control institutional outbreaks. |
| Cryptosporidium | Protozoal | Ingestion; contact | S (+ C as per additional comments) | Duration of illness | |

| Disease or Organism | | | | | |
|--|-------------------|---|--|---|---|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| Escherichia coli (STEC, ETEC) | Bacterial | Ingestion; contact | S (+ C as per additional comments) | Duration of illness | |
| Vibrio cholerae (<i>Cholera</i>) | Bacterial | Ingestion; contact | S (+ C as per additional comments) | Duration of illness | |
| <i>Vibrio parahaemolyticus</i> | Bacterial | Ingestion; contact | S (+ C as per additional comments) | Duration of illness | |
| <i>Yersinia enterocolitica</i> | Bacterial | Ingestion; contact | S (+ C as per additional comments) | Duration of illness | |
| <i>Salmonella spp</i> | Bacterial | Ingestion; contact | S (+ C as per additional comments) | 24 hours after symptoms cease | Use Contact Precautions for nappied or incontinent persons for the duration of illness or to control institutional outbreaks. |
| <i>Shigella spp</i> | Bacterial | Ingestion; contact | S (+ C as per additional comments) | 24 hours after symptoms cease | |
| <i>Campylobacter spp</i> | Bacterial | Ingestion; contact | S (+ C as per additional comments) | 24 hours after symptoms cease | |
| Haemophilus influenzae type b (HiB) Invasive disease - meningitis, epiglottitis, pneumonia | Bacterial | Droplet; contact (rare) | S + D | Until 24 hours after initiation of effective therapy | <i>H. Influenzae</i> presents more commonly in children than in adults. |
| Hendra virus (<i>paramyxovirus</i>) | Viral | Direct contact with respiratory secretions and bodily substances; Flying foxes (fruit bats) to horses then horse to human | S + C + D (+ A as per additional comments) | Duration of illness | Current evidence does not support airborne exposure as a recognised mode of transmission; however, employ airborne precautions for aerosol-generating procedures. |
| Hepatitis A | Viral | Ingestion; Contact | S (+ C as per additional comments) | For 7 days after onset of jaundice; for duration of hospitalisation for children <3 years | Immunise if at high risk; provide hepatitis A vaccine or normal human immunoglobulin (NHIG) post-exposure as recommended. Use Contact Precautions for nappied or incontinent persons for the duration of illness. |
| Hepatitis E | Viral | Ingestion; Contact | S (+ C as per additional comments) | Period of communicability unknown, probably at least 14 days after onset of jaundice | Use Contact Precautions for nappied or incontinent individuals for the duration of illness. |
| Herpes simplex virus infection mucocutaneous, disseminated or primary, severe | Viral | Contact (droplet, fomites, lesions) | S + C | Until lesions dry and crusted | - |

| Disease or Organism | | | | | |
|---|-------------------|---|------------------------------------|---|--|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| Herpes simplex virus infection neonatal | Viral | Contact (droplet, fomites, lesions) | S + C | Until lesions dry and crusted | Also, for asymptomatic, exposed infants delivered vaginally or by C-section and if mother has active infection and membranes have been ruptured for more than 4 to 6 hours until infant surface cultures obtained at 24-36 hours of age negative after 48 hours incubation. |
| Herpes zoster (shingles – disseminated disease) | Viral | Airborne; direct contact with fluid in blisters | S + C + A | Duration of illness | Susceptible healthcare workers must not provide direct patient care. |
| Herpes zoster (shingles – localised disease) | Viral | Direct contact with fluid in blisters | S (+ C as per additional comments) | Duration of illness (if wound lesions, until wounds cease draining, are dry and crusted) | Lesions not covered or dressing does not contain exudate. Susceptible healthcare workers must not provide direct patient care. |
| Human Metapneumovirus | Viral | Contact; droplet | S + C + D | Duration of illness | - |
| Influenza | Viral | Droplet; contact (both direct & indirect) | S + C + D | Until after 72 hours of the patient receiving anti-influenza medication; or five days have elapsed since onset of respiratory symptoms. May be longer for young children, immunosuppressed or ICU patients. | Annual immunisation recommended. |
| Influenza Avian | Viral | Transmission rare (see additional comments) | S + C + D + A | Duration of illness | Transmission of infection from birds to humans is rare. When it has occurred, it is believed to have resulted from close contact with infected poultry or breathing in dust contaminated with their excretions. Transmission has been thought to occur by ingesting uncooked poultry products (including raw blood) from H5N1 infected poultry. The spread of these viruses from one ill person to another through prolonged, unprotected, close contact has been reported very rarely, and has been limited, inefficient and not sustained. |
| Lice (pediculosis) Head | Arthropod | Contact | S + C | Contact until 24 hours after initiation of effective therapy, then Standard | - |
| Measles (rubeola) | Viral | Contact; Airborne | S + C + A | Until 4 days after rash appears: duration of illness in immune compromised patients | Screen by serology; pre-employment measles, mumps, rubella vaccine (MMR) if not |

| Disease or Organism | | | | | |
|---|-------------------|---|---|---|---|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| | | | | | pregnant. Non-immune staff should not care for patient. |
| Meningococcal infection (<i>Neisseria meningitidis</i>) | Bacterial | Droplet | S + D | For 24 hours after beginning effective treatment. | Immunisation possible in outbreaks. Post-exposure prophylaxis if indicated. Follow up of contacts may be required. |
| Middle East Respiratory Syndrome <i>Coronavirus (MERS-CoV)</i> | Viral | Not fully known. Presumed contact; droplet; airborne | S + C + D + A | Transmission-based precautions should be applied throughout any admission; additional isolation precautions should be continued until at least 24 hours after the resolution of symptoms. | - |
| Mpox (Previously known as monkeypox and also abbreviated as MPXV) | Viral - Clade I | Direct contact* (Unknown if also can be transmitted via respiratory route) | S + C + A** (see additional comments) | People are considered infectious for four days before the onset of symptoms and/or the appearance of rash/skin lesions. Precautions should be implemented until all symptoms have resolved, including rash, mucosal lesions and other lesions (this includes the formation and shedding of a scab, leaving fresh intact skin underneath). | To reduce the risk of transmission of MPXV in healthcare settings, patients with suspected or confirmed MPXV infection should be managed with S+C+A precautions until a risk assessment considering the clade type is completed. For more information, refer to the CDNA Series of National Guidelines . *The principal mode of MPXV transmission is thought to be direct contact with MPXV lesions or objects contaminated with lesions, such as clothing and bed linen (fomites). All HCWs should wear disposable gloves when providing care to patients with suspected or confirmed MPXV (including handling of used clothes, bed linen and towels), and practice strict hand hygiene before and after the use of gloves. **Clade I: Clade I appears more easily transmissible than clade II and there remains some uncertainty regarding the mode of transmission. Although likely that clade I is spread by the same route as clade II, until more information is available HCWs should use airborne precautions when caring for patients suspected or confirmed with MPXV clade I infection. ***Clade II: The use of droplet precautions is required during potentially particulate-dispersing activities such as handling used linen or if a patient has respiratory symptoms and potentially lesions in the nasopharynx. The use of either |
| | Viral - Clade II | | S + C + D*** (see additional comments) | | |

| Disease or Organism | | | | | |
|---|-------------------|--|--|--|---|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| | | | | | intermittent or continuous droplet precautions for clade II infections should be based on patient risk assessment. |
| Multi drug resistant organisms - infection or colonisation by the following organisms: | | | | | |
| Carbapenemase - producing Enterobacterales (CPE) | Bacterial | Contact | S + C | Duration of illness | Single room with ensuite for faecally incontinent patients is preferred. |
| Extended-spectrum beta-lactamase producing Enterobacteriaceae | Bacterial | Contact | S (+ C as per additional comments) | Duration of symptomatic illness | The use of contact precautions should be determined locally based on the bacterial species, local epidemiology and in the event of an outbreak. |
| Vancomycin-resistant Enterococcus (VRE) | Bacterial | Contact | S + C | Duration of illness | Single room with ensuite for faecally incontinent patients is preferred. |
| Mumps | Viral | Contact; droplet (respiratory secretions) | S + D | Until 5 days after onset of swelling. Exposed non-immune people should be considered infectious from 12th-25th day after exposure, with or without symptoms. | Screen by serology; pre-employment measles mumps rubella vaccine (MMR) if not pregnant. |
| Necrotising enterocolitis | Bacterial | Usually endogenous | S (+ C as per additional comments) | Duration of illness | Contact precautions when cases clustered temporally. |
| Nipah virus (NiV) (Paramyxoviridae family) | Viral | Contact and Droplet | S + C + D | Until symptoms resolve. | Zoonosis infection. Can transfer from animals to people, and less commonly person to person. In humans, NiV causes encephalitis (inflammation of the brain) or respiratory signs and symptoms. Symptom onset 4 to 18 days from exposure to infected animal. (Source: WHO) |
| Norovirus | Viral | Contact (droplet in certain circumstances) | S + C (+ D if determined to be necessary by risk assessment) | For a minimum of 48 hours after the resolution of symptoms or to control institutional outbreaks | The use of contact and/or droplet precautions may be required for incontinent patients or during outbreaks. This should be based on a risk assessment. Alcohol-based hand hygiene products are less effective than hand washing with soap and water for this infectious agent. |

| Disease or Organism | | | | | |
|--|-------------------|------------------------------------|---|---|--|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| | | | | | <p>Healthcare workers should use a surgical mask while patient is vomiting.</p> <p>Persons who clean areas heavily contaminated with faeces or vomitus may benefit from wearing masks since virus can be aerosolised from these body substances.</p> <p>Cohorting of affected patients to separate airspaces and toilet facilities may help interrupt transmission during outbreaks.</p> |
| Parainfluenza | Viral | Droplet | S + D | Duration of illness | Viral shedding may be prolonged in immunosuppressed patients. |
| Parvovirus B19 Infection (<i>Erythema infectiosum</i>) | Viral | Droplet | S + D | Duration of illness | - |
| Pertussis (Whooping cough) | Bacterial | Droplet | S + D | Until at least 5 days after commencement of appropriate antibiotic therapy, or; for 21 days after the onset of symptoms if not receiving antibiotic treatment, or; for 14 days after the onset of paroxysmal cough (if the onset is known). | Pre-employment booster/ vaccination recommended; post-exposure prophylaxis for healthcare worker in late pregnancy and high risk areas. |
| Plague (<i>Yersinia pestis</i>) pneumonic | Bacterial | Droplet | S + D | Until 48 hours after initiation of effective antibiotic therapy. Patient must be in respiratory isolation room. | Antimicrobial prophylaxis for exposed healthcare workers. |
| Pneumococcal pneumonia <i>Streptococcus pneumoniae</i> (<i>Pneumococcal pneumonia</i>) | Bacterial | Droplet | S (+ D if evidence of transmission within a facility) | Duration of illness | - |
| Pneumonia Adenovirus | Viral | Droplet | S + C + D | Duration of illness | In immunocompromised hosts, extend duration of Droplet and Contact Precautions due to prolonged shedding of virus. |
| Pneumonia <i>Burkholderia cepacia</i> in patients with cystic fibrosis, including respiratory tract colonization | Bacterial | Contact; contaminated medicines | S + C + D | Duration of illness | Avoid exposure to other persons with cystic fibrosis; single room preferred. Criteria for D/C precautions not established. See Cystic Fibrosis Foundation guideline. |

| Disease or Organism | | | | | |
|---|-------------------------|--|-------------------------|--|--|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| Pneumonia <i>Mycoplasma</i> | Bacterial | Droplet | S + D | Duration of illness | - |
| Pneumonia Viral | Viral | Droplet; contact | S + D | Duration of illness | - |
| Poliomyelitis | Viral | Ingestion | S + C | Duration of illness; may be shed in faeces for up to 6 weeks | Healthcare workers should be vaccinated (if have not had childhood vaccinations); non-immune healthcare workers should not care for patient. |
| Pressure ulcer (decubitus ulcer, pressure sore) Infected Major | Bacterial (commonly) | Not transmitted person-to-person | S + C | Duration of illness | - |
| Rabies / Australian Bat Lyssavirus | Viral | Transmitted by animal bites, scratches, or by contamination of mucous membranes or broken skin | S + C | Duration of illness | Person to person transmission rare; if patient bites another person wash exposed area thoroughly and administer post-exposure prophylaxis. |
| Respiratory Syncytial Virus (RSV) | Viral | Contact; droplet | S + C + D | Duration of illness | Use mask according to Standard Precautions. Avoid contact between the person who is ill and infants, young children, frail and elderly people, or immunocompromised patients until the person is feeling well. May be prolonged shedding in immunocompromised patients—as such, extend duration of Contact Precautions. |
| Rhinovirus | Viral | Contact; droplet | S + C + D | Duration of illness | Avoid contact between the person who is ill and infants, young children, frail and elderly people, or immunocompromised patients until the person is feeling well. |
| Rotavirus gastroenteritis | Viral | Ingestion; contact; droplet | S + C | Duration of illness | Alcohol-based hand hygiene products are less effective than hand washing with soap and water for this infectious agent. Ensure consistent environmental cleaning and disinfection and frequent removal of soiled diapers. Prolonged shedding may occur in both immunocompetent and immunocompromised children and the elderly. |
| Rubella | Viral | Contact; droplet | S + D (+ C if touching) | Until 7 days after onset of rash | Screen by serology; pre-employment MMR if not pregnant; |

| Disease or Organism | | | | | |
|---|-----------------------|--|---|---|---|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| | | | respiratory secretions and bodily substances) | | non-immune and pregnant staff should not attend patient. |
| Rubella Congenital | Viral | Vertical | S + C | Until 1 year of age | Standard precautions may be used if nasopharyngeal and urine cultures are repeatedly negative after 3 months of age; non-immune pregnant staff should not attend patient. |
| Scabies (<i>Sarcoptes scabiei</i>) | Arthropod infestation | Contact (skin to skin) or from infested fomites | S + C | Until 24 hours after treatment commenced | Healthcare workers should be excluded from work until effective treatment has been commenced. |
| Severe Acute Respiratory Syndrome (SARS) | Viral | Contact; droplet; airborne | S + C + D + A | Duration of illness + 10 days after resolution of fever, provided respiratory symptoms are absent or improving | P2 / N95 respirator or higher respiratory protection; surgical mask if P2 / N95 unavailable; eye protection (goggles, face shield); aerosol-generating procedures and "super-shedders" highest risk for transmission via small droplet nuclei and large droplets. |
| SARS-CoV-2 Coronavirus Disease 2019 (COVID-19) | Viral | Respiratory droplets or airborne particles | S + D + A* | Duration of illness + 10 days after resolution of fever, provided respiratory symptoms are absent or improving. | Transmission of SARS-CoV-2 occurs via respiratory droplets when an infected person coughs, sneezes, talks or sings, or during aerosol generating-procedures (AGPs). PPE recommended for respiratory viruses generally includes a surgical mask, protective eyewear and face shields. *In some unusual and higher risk situations, a PFR should be used instead of a surgical mask; however, this should be decided by the IPC lead or the person(s) responsible for IPC. |
| Smallpox (variola) | Viral | Contact; droplet; inhalation; contaminated fomites | S + C + A | Duration of illness | Until all scabs have crusted and separated (3-4 weeks). Non-vaccinated healthcare workers should not provide care when immune healthcare workers are available; P2/N95 or higher respiratory protection for both susceptible and successfully vaccinated individuals; post-exposure vaccine within 4 days of exposure is protective. |
| Staphylococcus aureus infections due to Methicillin sensitive Staphylococcus aureus (MSSA): | | | | | |
| Enterocolitis | Bacterial | Contact; Ingestion | S (+ C as per additional comments) | Duration of illness | Use Contact Precautions for nappied or incontinent children for duration of illness. |

| Disease or Organism | | | | | |
|---|---|--------------------|---|---|--|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| Furunculosis | Bacterial | Contact | S (+ C as per additional comments) | Duration of illness | Contact precautions if drainage not controlled. |
| Scalded skin syndrome | Bacterial | Contact | S + C | Duration of illness | - |
| Skin, wound or burns - major | Bacterial | Contact | S + C | Duration of illness for draining wound | No dressing or dressing does not contain drainage adequately. |
| Skin, wound or burn – minor or limited | Bacterial | Contact | S (+ C as per additional comments) | Duration of illness for draining wound | No dressing or dressing does not contain drainage adequately. |
| Staphylococcus aureus infections due to Methicillin resistant <i>Staphylococcus aureus</i> (MRSA) | Bacterial | Contact | S + C (+ D as per additional comments) | Duration of illness | Isolation of patients with MRSA may not be feasible. In each case, the implementation of contact precautions and isolation should be based on an appropriate risk assessment. Use Droplet precautions for patients known to have respiratory infection or colonisation with MRSA. |
| Streptococcal infections caused by Group A Streptococci (e.g. Impetigo, Pharyngitis in infants and young children, Scarlet Fever in infants and young children, Pneumonia) | Bacterial | Contact | S + D | Until 24 hours after treatment commenced | - |
| Streptococcal major skin, wound or burn infections caused by Group A Streptococci | Bacterial | Contact | S + C + D | Until 24 hours after treatment commenced | Until drainage stops or can be adequately contained by dressing. |
| Tuberculosis | <p>Airborne precautions are indicated for all patients where pulmonary TB is suspected or proven. Due to the difficulty associated with induced sputum collection and non-pulmonary TB presentation in the paediatric population, airborne precautions should be used until active pulmonary/laryngeal TB is ruled out in visiting family, caregivers or household contacts or a diagnosis of pulmonary/laryngeal TB has been ruled out. Seek specialist infectious diseases advice and refer to current TB SoNG for further information if caring for paediatric patients.</p> <p>Airborne precautions should also be used if any procedure involving aerosolisation is to be performed and tuberculosis is a diagnostic possibility. Airborne precautions are not necessary for persons with extrapulmonary (where there is no evidence of pulmonary TB as well) and latent TB infection.</p> | | | | |
| Extrapulmonary, draining lesion | Bacterial | Airborne | S + C | Until patient improving clinically and drainage has ceased or 3 consecutive negative cultures | - |
| Extrapulmonary but no draining lesion; meningitis | Bacterial | Airborne | S (+ A as per additional comments for Tuberculosis) | Duration of illness | Examine for pulmonary TB. |

| Disease or Organism | | | | | |
|--|-------------------|--|-------------------------------------|--|---|
| Disease / Organism | Type of infection | Transmission Route | Required precautions | Duration of precautions | Additional comments |
| Pulmonary or laryngeal disease, confirmed | Bacterial | Airborne | S + A | Usually until after 1 week of treatment and 3 sputum smears negative; consult with respiratory physician | - |
| Pulmonary or laryngeal disease, suspected | Bacterial | Airborne | S + A | Until TB excluded; alternate diagnosis or 3 sputum smears AFB negative, each specimen collected 8-24 hours apart and at least one early morning specimen | - |
| Typhoid (<i>Salmonella Typhi</i>) | Bacterial | Faecal-oral route; Ingestion of contaminated water and food | S (+ C as per additional comments) | Duration of illness | Transmission by direct person-to-person contact can occur but is rare. Individual toilet is preferred in addition to Standard Precautions. Use Contact Precautions for nappied or incontinent persons for the duration of illness or to control institutional outbreaks. |
| Viral haemorrhagic fevers (VHF) Crimean-Congo, Ebola, Lassa, Marburg | Viral | Contact with blood or body substances (mucosal, parenteral) either directly or indirectly Lassa fever: aerosols | S + C + D (+ A, organism dependent) | Duration of illness; isolation room | Emphasise: use of sharps safety devices and safe work practices; hand hygiene; barrier protection against blood and body substances upon entry into room—gloves, fluid-resistant or impermeable gowns, face/eye protection with masks, goggles or face shields; appropriate waste handling. |

Source: *Guideline for Isolation Precautions: Preventing Transmission of Infectious Agents in Healthcare Settings (Appendix A updated September 2018) [422]; Series of National Guidelines (SoNGs) [357]*

6.5 Collection of pathology specimens

Best practice in the identification, collection and handling of pathology specimens involves:

- adherence to standard infection prevention precautions is required for all procedures involving the collection and handling of pathology specimens
- specimens that are to go to histopathology should be placed in a pathology biohazard bag, and then into the specimen bucket
- the label should be stuck on the container immediately before the specimen is placed in the specimen bucket
- larger specimens require a clear plastic bag, and should then be placed in a specimen bucket
- following the Australian Commission on Safety and Quality in Health Care's '*Ensuring correct patient, correct site, correct procedure protocol*' to confirm the right specimen has been collected from the right patient and labelled correctly. Double verification of the process should occur.

For more information, see the *National Pathology Accreditation Advisory Council Standards [425]*.

6.6 Allowing animals into healthcare facilities

Assistance, therapeutic and companion animals are increasingly being used in Australian healthcare facilities. Healthcare facilities are encouraged to develop policies on infection control and hand hygiene for animal-assisted interventions, with input from an infection control expert. This may include [430]:

- ensuring regular hand hygiene is performed before and after entering a patient care area, before and after handling an animal, and after toileting an animal
- restricting entrance into clinical areas such as operating theatres, intensive care units, isolation rooms, neonatal and newborn nurseries, and other areas identified specifically by the healthcare facility
- ensuring routine environmental cleaning or disinfection is performed after animal visits
- ensuring that all animals visiting or permanently residing in the healthcare facility have been assessed by a veterinarian, screened for parasites and skin problems, and are fully vaccinated (a veterinary immunisation certificate should be provided).

For further information see the *Australian College for Infection Prevention and Control position statement on animal visits in healthcare facilities* [431].

6.7 Mechanisms of antibacterial action of antiseptics and disinfectants

Antiseptics and disinfectants are used widely in healthcare facilities for a variety of topical and hard-surface applications. Both antiseptics and disinfectants reduce the number of micro-organisms, with the former used on living tissues and the latter on nonliving objects.

The table below provides information on the mode of action of active agents in different antiseptics and disinfectants.

Table A2.6. Active agents in antiseptics and disinfectants

| Target | Antiseptic or disinfectant | Mode of action |
|---|---|---|
| Cytoplasmic (inner) membrane | Disinfectant: Quaternary ammonium compounds (QACs) | Generalised membrane damage involving phospholipid bilayers. Bactericidal action of the quaternaries has been attributed to the inactivation of energy-producing enzymes, denaturation of essential cell proteins, and disruption of the cell membrane. |
| | Disinfectant and antiseptic: Chlorhexidine | Low concentrations affect membrane integrity, high concentrations cause congealing of cytoplasm. |
| | Disinfectant: Phenols | Leakage; some cause uncoupling. In high concentrations, phenol acts as a gross protoplasmic poison, penetrating and disrupting the cell wall and precipitating the cell proteins. Low concentrations of phenol and higher molecular-weight phenol derivatives cause bacterial death by inactivation of essential enzyme systems and leakage of essential metabolites from the cell wall. |
| Cross-linking of macromolecules | Disinfection: Formaldehyde | Cross-linking of proteins, RNA, and DNA. Formaldehyde inactivates microorganisms by alkylating the amine and sulfhydryl groups of proteins and ring nitrogen atoms of purine bases. |
| Denaturing protein, dissolving lipids, membrane damage | Disinfectant and antiseptic: alcohols (isopropyl and ethyl alcohol) | Primary mode of action of alcohols is through membrane damage, rapid denaturing of proteins and dissolving of lipids. Effective against many types of bacterial and viral cells but is not sporicidal. |
| DNA intercalation | Disinfectant and antiseptic: Acridines | Intercalation of an acridine molecule between two layers of base pairs in DNA. |
| Interaction with thiol groups | Disinfectant and antiseptic: Silver compounds | Membrane-bound enzymes (interaction with thiol |

| Target | Antiseptic or disinfectant | Mode of action |
|--|--|---|
| Effects on DNA | Disinfectant and antiseptic: Halogen-releasing agents (sodium hypochlorite), Hydrogen peroxide, silver ions | groups). DNA strand breakage. |
| Oxidizing agents | Disinfectant and antiseptic: Halogen-releasing agents (sodium hypochlorite), Peroxygens | Hydrogen Peroxide: produces destructive hydroxyl free radicals that can attack membrane lipids, DNA and other essential cell components. Hydrogen peroxide: Activity due to formation of free hydroxyl radicals (OH), which oxidize thiol groups in enzymes and proteins. Peracetic acid (PAA): Disruption of thiol groups in proteins and enzymes. |
| Cell envelope (cell wall, outer membrane) | Disinfectant: Ethylenediaminetetraacetic (EDTA), other permeabilizers Disinfectant and high level sterilant: Glutaraldehyde | Gram-negative bacteria: Removal of Mg ²⁺ , release of some of some lipopolysaccharides (LPS). Cross-linking of proteins. As an aqueous solution Glutaraldehyde is acidic and generally in this state not sporicidal. However, if the solution is activated through alkalinating agents (pH 7.5-8.5) the solution becomes sporicidal. |
| Cytoplasmic (inner) membrane | Disinfectant: Diamines Disinfectant and antiseptic: Polyhexanide (polyhexamethylene biguanide; PHMB), Alxidine | The biocidal activity is a result from its alkylation of sulfhydryl, hydroxyl, carboxyl, and amino groups of microorganisms, which alters RNA, DNA, and protein synthesis. Induction of leakage of amino acids. Phase separation and domain formation of membrane lipids. |

Adapted from: Antiseptics and Disinfectants: Activity, Action and Resistance [422]; Disinfection and Sterilisation [429]

6.8 Summary table of relevant infection prevention and control resources

It is acknowledged that the example resources provided below may be updated on a regular basis. To ensure best practice is being followed, ensure the current electronic version is being accessed.

Table A2.7. Infection prevention and control resources

| Topic | Organisation | Resource |
|--|--|--|
| | | National |
| <i>Clostridium difficile</i> (<i>Clostridioides difficile</i>) | Australasian Society for Infectious Diseases (ASID)/Australasian College for Infection Prevention and Control (ACIPC) (2019) | ASID/ACIPC position statement – Infection control for patients with <i>Clostridium difficile</i> infection in healthcare facilities |
| Carbapenemase-producing <i>Enterobacteriales</i> (CPE) | Australian Commission on Safety and Quality in Health Care (2017) | Recommendations for the control of carbapenemase-producing <i>Enterobacteriaceae</i> (CPE): A guide for acute care health facilities |
| Antimicrobial stewardship | Australian Commission on Safety and Quality in Health Care (2011) | Antimicrobial Stewardship in Australian Hospitals |
| Surgical Site Infection surveillance | Australian Commission on Safety and Quality in Health Care (2017) | Approaches to Surgical Site Infection Surveillance: For acute care settings in Australia |
| Dental | Australian Dental Association (2015) | Guidelines for Infection Control |

| Topic | Organisation | Resource |
|--|--|--|
| Influenza Kit for Aged Care | Australian Government Department of Health | Influ-Info Influenza Kit for Aged Care |
| Facility design | Australasian Health Infrastructure Alliance (2016) | Australasian Health Facility Guidelines: Part D - Infection Prevention and Control |
| Reprocessing | Australasian Society for Ultrasound in Medicine (2017) | Guidelines for Reprocessing Ultrasound Transducers |
| Post-exposure prophylaxis | Australasian Society of HIV Medicine (2016) | National guidelines for post-exposure prophylaxis after non-occupational and occupational exposure to HIV |
| Immunisation | Australian Technical Advisory Group on Immunisation (2017) | The Australian Immunisation Handbook, 10th edition |
| PIVC | Clinical Excellence Commission (2013) | Peripheral Intravenous Cannula (PIVC) Insertion and Post Insertion Care in Adult Patients |
| Gastroenteritis outbreaks | Communicable Diseases Network Australia (2010) | Guidelines for the public health management of gastroenteritis outbreaks due to norovirus or suspected viral agents in Australia |
| Healthcare workers positive for blood-borne viruses | Communicable Diseases Network Australia (2018) | Australian National Guidelines for the Management of Health Care Workers known to be Infected with Blood-Borne Viruses |
| Management of Influenza Outbreaks in Residential Care Facilities | Communicable Diseases Network Australia (2017) | Guidelines for the Prevention, Control and Public Health Management of Influenza Outbreaks in Residential Care Facilities in Australia |
| Ebola | Department of Health (2015) | Infection prevention and control principles and recommendations for Ebola virus disease |
| Creutzfeld-Jakob disease | Department of Health (2013) | Creutzfeld-Jakob Disease Infection Control Guidelines |
| Endoscopy | Gastroenterological Nurses College of Australia (2010) | Infection Control in Endoscopy |
| Hand hygiene | Australian Commission on Safety and Quality in Healthcare (2019) | National Hand Hygiene Initiative (NHHI) Implementation Guide |
| Australian Standards | Standards Australia | Australian Standards relevant to healthcare facilities |
| Office-based and community-based practices | The Royal Australian College of General Practitioners (2014) | Infection Prevention and Control Standards: For general practices and other office-based and community-based practices |
| International | | |
| MRSA | Calfee et al (2014) | Strategies to Prevent Methicillin-Resistant Staphylococcus aureas Transmission and Infection in Acute Care Hospitals |
| Intravascular access devices | Centers for Disease Control and Prevention (2011) | Guidelines for the Prevention of Intravascular Catheter-Related Infections |
| Environmental infection control | Centers for Disease Control and Prevention (2003) | Guidelines for Environmental Infection Control in Health-Care Facilities |
| Outpatient settings | Centers for Disease Control and Prevention (2016) | Guide to Infection Prevention for Outpatient Settings: Minimum Expectations for Safe Care |
| Clostridioides difficile (formerly Clostridium difficile) | Cohen et al (2010) | Clinical Practice Guidelines for Clostridium difficile Infection in Adults |
| Norovirus | Hall et al (2011) | Updated Norovirus Outbreak Management and Disease Prevention Guidelines |
| Hospital-acquired and ventilator associated pneumonia | Kalil et al (2016) | The 2016 Clinical Practice Guidelines for the Management of Adults with Hospital-acquired and Ventilator-associated Pneumonia |
| General infection prevention | Loveday et al (2014) | epic3: National Evidence-Based Guidelines for Preventing Healthcare-Associated Infections in NHS Hospitals in England |
| General infection prevention | National Institute for Health | Healthcare-associated infections: prevention and control |

| Topic | Organisation | Resource |
|---|--|--|
| General infection prevention | Care Excellence (2011) Public Health Agency of Canada (2012) | Routine Practices and Additional Precautions for Preventing the Transmission of Infection in Healthcare Settings |
| Infection Prevention and Control Programs | Public Health Ontario (2012) | Best Practices for Infection Prevention and Control Programs |
| Primary and community care | National Institute for Health Care Excellence (2012, updated 2017) | Healthcare-associated infections: prevention and control in primary and community care |
| Antibiotic Resistant Organisms | Provincial Infection Control Network for British Columbia (2013) | Antibiotic Resistant Organisms (ARO) Guidelines |
| Environmental cleaning | Provincial Infection Control Network for British Columbia (2012) | Best Practices for Environmental Cleaning for Prevention and Control of Infections |
| Endoscopy | Public Health Agency of Canada and Canadian Association of Gastroenterology (2013) | Infection Prevention and Control Guidelines for Flexible Gastrointestinal Endoscopy and Flexible Bronchoscopy |
| Catheters | Tenke et al (2008) | European and Asian Guidelines on the Management and Prevention of Catheter-Associated Urinary Tract Infections |
| Surgical Site Infections | The Society for Healthcare Epidemiology of America (2014) | Strategies to Prevent Surgical Site Infections in Acute Care Hospitals: 2014 Update |
| Surgical Site Infections | The World Health Organization (2016) | Global Guidelines on the Prevention of Surgical Site Infections |
| Surgical Site Infections | National Institute for Health and Care Excellence (2008, updated 2017) | Surgical site infections: prevention and treatment |
| Infection Control Programs | The World Health Organization (2016) | Guidelines on core components of infection prevention and control programmes at the national and acute healthcare facility level |

6.9 Research gaps

Norovirus

- More high quality comparative studies that evaluate the effectiveness of infection control strategies are needed in order to extrapolate the findings to clinical practice.

Chlorhexidine resistance

- Studies that investigate the implications of chlorhexidine based decolonisation/universal chlorhexidine use on methicillin-resistant *Staphylococcus aureus* (MRSA) *qacA/B* genes among hospitalised patients, especially intensive care unit patients/whether the carriage of *qacA/B* can account for some of the decolonization failures/role of *qacA/B* and decreased susceptibility to chlorhexidine.
- Studies that examine how to ensure that chlorhexidine is correctly used and applied in clinical practice as an antiseptic and as a disinfectant to control and/or prevent health-care associated infections.
- Studies that examine the role of chlorhexidine in biofilm formation and how best to suppress the biofilm induction.
- Studies that investigate whether a resistance of certain strains of bacteria to chlorhexidine contributes to these strains persisting in the hospital environment.

Chlorhexidine and anaphylaxis

- More research is required on chlorhexidine-related anaphylaxis to better understand the nature, risks and magnitude of the problem.
- Clearer definitions, better reporting and clearer documentation of the exposures at the time of the anaphylactic event, and the exposure prior to the event, are needed in future research.

Antimicrobial surfaces

- Future studies need to include both infection and colonisation as outcomes in order to determine whether antimicrobial surfaces have a clinically important impact on infection prevention and control.
- Future research needs to utilise optimal study designs, such as cluster randomised trials or an interrupted time series designs when randomisation is not possible.

Novel disinfection methods

- Future studies need to include both infection and colonisation as outcomes in order to determine whether antimicrobial surfaces have a clinically important impact on infection prevention and control.
- Studies are needed amongst high risk population groups (e.g. burns or oncology patients) as the resourcing and cost considerations of novel disinfectants may be outweighed by the benefits in areas of high risk.

Contact precautions for patients known to be infected or colonised with a multi-resistant organism (MRO)

- More large scale, high quality trials are needed to assess the efficacy of contact precautions compared to horizontal interventions.
- Further research is also required to determine whether there are specific patient populations who would benefit more from the implementation of contact precautions than others.

Clearance criteria for specific MROs

- There is not yet an agreed protocol for the clearance of multi-resistant Gram negative (MRGN) carriage and vancomycin-resistant enterococci (VRE) carriage due to a lack of empirical evidence.

7. APPENDIX 3: Process report

Introduction

In November 2014, the Australian Health Ministers' Advisory Council (AHMAC) endorsed four criteria for prioritising clinical practice guidelines to be considered for funding. As part of this process, the Australian Commission on Safety and Quality in Health Care (ACSQHC) decided that the *2010 Australian Guidelines for the Prevention and Control of Infection in Healthcare* (the 2010 Guidelines) required review.

Although no formal evaluation of the 2010 Guidelines has been undertaken, ACSQHC advised that the 2010 Guidelines were a key resource for the National Safety and Quality Health Service Standards (Standard 3: Preventing and controlling healthcare associated infections), and are used in healthcare facilities for the development of policies and protocols across a range of acute and non-acute settings. This demonstrated the need for maintaining the currency and accuracy of the Guidelines.

ACSQHC and the National Health and Medical Research Council (NHMRC) signed a memorandum of understanding in early 2016 to update the 2010 Guidelines.

Contributors

The Guidelines were developed utilising a collaborative approach, combining the content expertise of the Infection Control Guidelines Advisory Committee (ICGAC) with the assistance of systematic reviewers and expert methodologists. NHMRC managed the review process and technical writing and representatives from ACSQHC participated as contributing observers throughout the process. ACSQHC provided a significant proportion of the funding for developing these Guidelines.

a. ICGAC

Information on appointment, membership and declaration of interest processes is available in subsection 7.1 to this appendix.

b. NHMRC Project Team

- Cathy Connor
- Emma Breen
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c. ACSQHC Project Team

- Adjunct Professor Kathy Meleady PSM
- Professor Marilyn Cruickshank
- Sue Greig
- Cate Quoye

d. Contractors

- KP Health, Horizon Scan
- Monash University, Systematic Reviewers
- University of South Australia, Literature Reviewers
- University of Sydney, Methodologists
- Health Technology Analysts, Methodologists

Assessment of the evidence

Evidence base

These Guidelines are an evolution of the 2010 Guidelines and build upon their evidence and science base. New evidence was assessed where the ICGAC advised that the prevention or management of infection had strengthened, weakened, or remained unchanged. Where the evidence base was unlikely to have changed substantially, additional systematic reviews were not conducted but references were updated for currency.

The methods used to analyse the evidence were in accordance with international best practice. The main methods are summarised below

with further process information available within each document. The Guidelines were further informed by substantial advances in the methodology for guideline development and usability since publication of the previous edition of the Guidelines. This included the use of the Grading of Recommendations Assessment, Development and Evaluation (GRADE) methodology and the MAGICapp platform for publication.

Randomised controlled trials provide the highest level of evidence however, as with many clinical and public health interventions, these often raise ethical, logistical and economic challenges. This is particularly the case in conducting randomised controlled trials in infection prevention and control, where the evidence may be limited by the inability to conduct certain study designs that are difficult to implement in real practice. Under GRADE, this has implications for rating the quality of evidence and the strength of recommendations as evidence based grading systems generally favour study designs that may not be feasible or ethical to conduct in infection control settings, such as randomised controlled trials. For example, it is unethical to compare the incidence of infection related to surgical instruments by allocating one patient group to have sterilised instruments used on them and one patient group to have non-sterile instruments used. This may result in a weak recommendation due to the available evidence but sterilisation of surgical instruments is universally deemed critical to infection control.

Given that there is limited evidence available to support many routine practices intended to reduce infection risk, practice is based on decisions made upon scientific principles. The relatively high proportion of recommendations graded as weak should not be interpreted as suggesting lack of evidence to help guide practice. Some activities, such as performing hand hygiene between administering care to successive patients, have a credible history to support the routine application of this activity in preventing cross-infection. Others, such as some uniform and clothing requirements, have more to do with the ethos of quality care and workplace culture than with a proven reduction in cross-infection.

Health professionals and the public can be assured that the process of assessing the scientific evidence provides for the best possible advice. It is not acceptable to discontinue practices for which there is a solid scientific basis, even if the level of evidence and subsequent recommendation is not high. Rather, routine practices should continue unless there is sufficient evidence to support alternative procedures. Continuing research is needed to keep evaluating practice, to identify evidence gaps and promote research in these areas, and to ensure that poor practices are replaced with better ones.

Capturing new evidence

Infection prevention and control research is continuously evolving and studies are published regularly. A commissioned horizon scan allowed for the ICGAC to determine topics that required a more extensive literature research and assisted in identifying relevant results from high quality studies (primarily systematic reviews or international guidelines) that were published since the previous Guidelines. While results from these studies were not graded, they were included in the Guideline update to ensure the content was current and applicable to the Australian population.

1. *Horizon scan*

A horizon scan was undertaken by expert consultants from KP Health to identify relevant national and international infection control guidelines, relevant standards and policies produced by Australian jurisdictions since 2009, and to determine gaps between recommendations in the 2010 Guidelines and more recently published guidelines. These gaps or changes primarily related to routine hand hygiene, routine cleaning of surfaces (novel methods), cleaning of shared clinical equipment and core strategies in the control of multi-resistant organisms.

Based on the horizon scan and their expertise in the field, the ICGAC considered priority areas for the revision of the Guidelines and determined topics that required a more extensive literature research, as follows:

- Chlorhexidine
- Norovirus
- Antimicrobial Surfaces, and
- Novel Disinfectants.

2. *Systematic reviews*

Monash University were commissioned by NHMRC in June 2016 to develop research protocols and conduct systematic reviews to evaluate the evidence on:

- i. the effect of environmental surfaces, fittings or fixtures with antimicrobial properties on infection rates in hospital patients compared with standard surfaces
- ii. the effect of ultraviolet light, hydrogen peroxide vapour and/or electrolysed water on infection rates in high risk population groups compared with standard care (cleaning with detergent, disinfection with bleach, or both)
- iii. the effect of disinfection with bleach on infection rates in high risk population groups compared with cleaning with detergent.

Monash University were engaged from NHMRC's Health Evidence Panel for its expertise in systematic review methodology, following a competitive tender process. The evidence review team completed a declaration of interest process before being appointed by NHMRC and no conflicts of interest were identified.

The independent evaluation of the evidence was undertaken in accordance with research protocols that were approved by the NHMRC Project Team, based on advice from the ICGAC. The research protocols outlined the scope, clinical questions, and methodology of the evidence reviews.

The methods and results of the reviews are detailed in the Evidence Evaluation Reports and Technical Reports. These include the research questions using the PICO approach (population, intervention, comparator, outcomes and setting), the search strategy, methods used to select, appraise and summarise the evidence, results, and evidence profiles.

The Evidence Evaluation Reports and Technical Reports were published on the NHMRC website on 16 November 2017, to coincide with the start of targeted consultation on the draft Guidelines.

3. **Literature Reviews**

The University of South Australia were commissioned by NHMRC in June 2016 to develop research protocols and conduct literature reviews on:

- i. the impact of chlorhexidine use on the incidence of anaphylaxis
- ii. the use of chlorhexidine and its contribution to increased resistance to chlorhexidine and/or antibiotics
- iii. current epidemiology and evidence on transmission pathways and infection prevention and control measures for Norovirus Gastroenteritis.

The University of South Australia were engaged from NHMRC's Health Evidence Panel for its expertise in literature review methodology, following a competitive tender process. The evidence review team completed a declaration of interest process before being appointed by NHMRC and no conflicts of interest were identified.

The independent evaluation of the evidence was undertaken in accordance with research protocols that were approved by the NHMRC Project Team, based on advice from the ICGAC. The research protocols outlined the scope, clinical questions and methodology of the evidence reviews.

The methods and results of the literature reviews are detailed in the Evidence Evaluation Reports and Technical Reports. These include the research questions using the PICO approach (population, intervention, comparator, outcomes and setting), the search strategy, methods used to select, appraise and summarise the evidence and results.

The Evidence Evaluation Reports and Technical Reports were published on the NHMRC website on 16 November 2017, to coincide with the start of targeted consultation on the draft Guidelines.

Independent methodological review

Independent methodological review of the Evidence Evaluations were completed to ensure they followed the systematic and rigorous approach specified in the corresponding research protocol.

a. Systematic reviews

Health Technology Analysts were contracted in July 2016 to assess the methodological quality of the commissioned systematic reviews. The methodological reviewers were appropriately qualified in systematic review processes and methodology. Health Technology Analysts successfully tendered to complete this activity through a competitive tender process using the NHMRC Health Evidence Panel. The methodological review team completed a declaration of interest process before being appointed by NHMRC and no conflicts of interest were identified.

Health Technology Analysts completed an independent methodological appraisal of the draft research protocols to provide NHMRC with assurance of the proposed methodology for the Evidence Evaluations. The methodologists comments were provided to the commissioned evidence reviewers who revised the protocols accordingly, based on advice of the ICGAC, prior to the evaluations of the evidence commencing.

Once the Evidence Evaluations were complete, Health Technology Analysts appraised the methodological quality of the Evidence Evaluation Reports and Technical Reports, including whether the rigorous approach prescribed in the review protocols had been adhered to, and whether the interpretation and reporting of the evidence was transparent and thorough.

The ICGAC considered the feedback received from the methodological reviews and were satisfied with the critical appraisal performed.

b. Literature reviews

University of Sydney were contracted in July 2016 to assess the methodological quality of the commissioned literature reviews. The methodological reviewers were appropriately qualified in systematic review processes and methodology. University of Sydney successfully tendered to complete this activity through a competitive tender process using the NHMRC Health Evidence Panel. The methodological review team completed a declaration of interest process before being appointed by NHMRC and no conflicts of interest were identified.

University of Sydney completed an independent methodological appraisal of the draft research protocols to provide NHMRC with assurance of the proposed methodology for the Evidence Evaluations. The methodologists comments were provided to the commissioned evidence reviewers who revised the protocols accordingly, based on advice of the ICGAC, prior to the evaluations of the evidence commencing.

Once the Evidence Evaluations were complete, University of Sydney appraised the methodological quality of the Evidence Evaluation Reports and Technical Reports, including whether the rigorous approach prescribed in the review protocols had been adhered to, and whether the interpretation and reporting of the evidence was transparent and thorough.

The ICGAC considered the feedback received from the methodological reviews and were satisfied with the critical appraisal performed.

Governance and stakeholder involvement

The reports underpinning the systematic and literature reviews were critically appraised by the ICGAC and refined using a consensus approach over a series of meetings/teleconferences during early 2017.

The reports were provided to the Council of NHMRC on 12 October 2017. The Chief Executive Officer (CEO) of NHMRC subsequently agreed to publicly release the final systematic and literature reviews on the NHMRC website, to coincide with the start of targeted consultation on the draft Guidelines on 16 November 2017.

Guideline development

The recommendations in these Guidelines were formulated by the ICGAC through a process of consensus. Recommendations are given when an action is deemed critical to preventing or managing infection. Recommendations are assessed using the GRADE approach, incorporating NHMRC's Levels of Evidence [434]. Practice Statements, previously known as good practice points, have also been included as these are important in outlining actions that are essential to infection prevention and control but where evidence grades cannot be applied.

Table A3.1. Levels of evidence in literature reviews [434]

| Level | Description |
|-------------|---|
| Level I | A systematic review of level II studies |
| Level II | A randomised controlled trial |
| Level III-1 | A pseudorandomised controlled trial |
| Level III-2 | A comparative study with concurrent controls: <ul style="list-style-type: none"> • Non-randomised experimental trial • Cohort study • Case-control study • Interrupted time series with a control group |
| Level III-3 | A comparative study without concurrent controls: <ul style="list-style-type: none"> • Historical control study • Two or more single arm study • Interrupted time series without a parallel control group |
| Level IV | Case series with either post-test or pre-test/post-test outcomes |

Developing new recommendations, practice statements and statutory requirements

The GRADE approach is a system for rating the quality of a body of evidence in systematic reviews and grading recommendations in healthcare [433]. GRADE offers a transparent and structured process for developing and presenting evidence summaries and for carrying out the steps involved in developing recommendations. The GRADE system provided an evidence to decision framework which determined the structure for the ICGAC in developing new recommendations. The final wording of each recommendation was developed by the ICGAC through a consensus approach, within the constraints of GRADE strength of recommendations terminology.

Systematic reviews provide a comprehensive summary of the evidence but do not include recommendations. The commissioned evidence reviewer's role in the GRADE approach terminated after rating the quality of evidence and estimate of effect for outcomes, as per the results in the Evidence Profiles.

The ICGAC was responsible for critically appraising the systematic review Evidence Evaluations, developing recommendations and

completing the subsequent steps of GRADE and the guideline development process. This included explicit consideration of:

- the quality of the evidence (including consideration for the level of evidence)
- benefits and harms
- values and preferences
- resource use
- health inequalities.

These considerations are captured in the 'Key Info' tab accompanying each recommendation. This ensured that the ICGAC's application of GRADE was transparent and a consistent approach was applied in the development of all recommendations.

In drafting practice statements, discussions of the ICGAC were aligned with the GRADE process where available evidence and judgements were considered together. Practice Statements were informed by the systematic reviews, literature reviews and advice of the ICGAC.

The terminology 'statutory requirement' has been used for selected practice statements/recommendations to further indicate where there is also a mandated requirement/s by the Commonwealth or the states/territories, which must be considered when implementing the advice at the local level. It is important to note that statutory requirements vary across states and territories, and in their applicability to health service delivery sectors and settings.

Further information on GRADE can be found in the GRADE Handbook (<http://grade.pro.org>).

Adapting or adopting the 2010 recommendations and practice statements

Following the outcomes of the horizon scan activity and advice of the ICGAC, it was determined that many of the 2010 recommendations were underpinned by established evidence and/or it was unlikely that the body of evidence had significantly changed since 2010 (i.e. no known recent studies investigating a significant relationship).

GRADE allows for guideline developers to [435]:

- adopt existing recommendations from others
- adapt existing recommendations to their own context
- create recommendations de novo.

The GRADE process provided a consistent and transparent approach which allowed the ICGAC to consider the 2010 recommendations (developed using the FORM approach which included recommendations graded from A to D) and good practice points on a case by case basis to reassign a 'GRADE' recommendation or practice statements and its accompanying strength (see Table A3.2). The ICGAC reviewed each recommendation to ensure it followed best practice standards for guideline development, and were required to reach a consensus in agreement when determining the direction and strength of the adopted or adapted recommendation.

These considerations are captured in the 'key info' tab accompanying each adopted or adapted recommendation or practice statement. This ensured that the ICGAC's application of GRADE was transparent and a consistent approach was applied in the development of all recommendations.

Table A3.2. Comparison of FORM and GRADE recommendation descriptors

| FORM Recommendation Descriptor (2010 Guidelines) | GRADE Recommendation Descriptor (2019 Guidelines) |
|---|--|
| Grade A | Strong |
| Grade B | Strong |
| Grade C | Conditional/Weak |
| Grade D | Practice Statement |

Technical writing

NHMRC Project Team assumed the role of technical writers using MAGICapp cloud-based research and innovation platform to author the Guidelines. This aims to facilitate dynamic dissemination and updating of the Guideline, as well as encourage international collaboration through open data sharing.

Targeted consultation

On 12 October 2017 the Council of NHMRC agreed to recommend the CEO of NHMRC release the Guidelines for targeted consultation. Targeted consultation on the draft Guidelines was undertaken from 16 November 2017 to 21 December 2017. Invitations were sent to a number of key public and private healthcare providers in the acute setting, with eight submissions received.

The ICGAC considered the submissions out of session during February 2018 with the majority of comments proposing minor clarifications and/or improvements to content on the following topics:

- practical elements of the retained 2010 Guideline content such as case studies
- aseptic Technique (removal of Aseptic Non-Touch Technique (ANTT) registered trademark terminology)
- multi-Resistant Organisms (screening, clearance, antimicrobial stewardship)
- personal protective equipment (PPE) and when different precautions are required
- management of the physical environment and routine cleaning
- minimising the risk from intravascular access devices (length of time *in situ*)
- clinical governance and organisational support
- transition to the new IT platform - MAGICapp.

Public consultation

On 20 March 2018 the Council of NHMRC agreed to recommend the CEO of NHMRC release the Guidelines for public consultation. Public consultation on the draft Guidelines was undertaken from 13 April 2018 to 15 May 2018. This process was conducted in accordance with Section 13 of the *National Health and Medical Research Council Act 1992*.

Public consultation was advertised on the NHMRC website, NHMRC Tracker, and social media platforms. Invitations were also sent to a large number of key stakeholders and those with a known interest in infection prevention and control. In total, 52 submissions were received, with over 600 individual comments considered. The submissions were received from a variety of stakeholders including individuals, professionals, state and territory governments and industry.

The ICGAC met on 2 of July 2018, 22 October 2018 and 2 November 2018 to consider the submissions. Further information and/or consideration were requested for:

- involving patients in their care
- hand hygiene and glove use
- PPE and transmission based precautions
- routine environmental cleaning: use of detergents and disinfectants and cleaning frequencies
- Therapeutic Goods Administration (TGA)-registered and TGA-listed products (*note: since public consultation, reduced regulation for hard surface disinfectants have been implemented by TGA*)
- use of surface barriers
- management and disposal of sharps
- emerging disinfection methods: ultra-violet light (UVL), hydrogen peroxide vapour (HPV)
- reprocessing of reusable instruments and equipment
- aseptic Technique (removal of Aseptic Non-Touch Technique (ANTT) registered trademark terminology)
- management and surveillance of MROs
- standard precautions versus contact precautions for Vancomycin Resistant Enterococci (VRE)
- minimising the risk from intravascular access devices (length of time *in situ*)
- the appropriate use of chlorhexidine
- exclusion periods and precautions for specific infections and conditions
- transition to MAGICapp and GRADE.

Independent expert review

The Guidelines underwent independent expert review from 23 November 2018 to 20 December 2018 to gain advice, primarily on the evidence base and its translation. The expert reviewers were required to declare any interest as per the process outlined in Section 7.1. Independent expert reviewers are listed in Table A3.3.

Table A3.3. Independent expert review participants

| Name | Affiliation |
|------|-------------|
|------|-------------|

| | |
|------------------------|---|
| Dr Celia Cooper | Head, Infectious Diseases Department, Adelaide Women's and Children's Hospital |
| Dr Ann Koehler | Senior Medical Consultant, Communicable Disease Control Branch, System Performance & Service Delivery, South Australia Health |

Consultation with ACSQHC

The Guidelines were considered by ACSQHC's Inter-Jurisdictional Committee (IJC) from 11 December 2018 to 17 January 2019. All states and territories supported the progression of the Guidelines to the Council of NHMRC.

The Guidelines were noted for release in March 2019 by the IJC and ACSQHC Board.

Council of NHMRC endorsement

The Guidelines were considered by the Council of NHMRC on 14 March 2019 for recommendation to the CEO for issuing. The CEO was pleased to accept the Council's advice and agreed to issue the Guidelines under Section 7(1a) of the *National Health and Medical Research Council Act 1992*.



Australian Government

National Health and Medical Research Council

Australian Commission on Safety and Quality in Health Care

7.1 Membership and Terms of Reference of the Advisory Committee

Appointment process

The Infection Control Guidelines Advisory Committee (ICGAC) was a working committee established by a Chief Executive Officer delegate under Section 39 of the *National Health and Medical Research Council Act 1992 (NHMRC Act)*. The ICGAC was selected to ensure appropriate expertise in the key areas of infectious diseases, antimicrobial resistance and/or microbiology, healthcare associated infection, paediatrics, and evidence-based methodologies, in order to oversee and provide expertise in updating the *2010 Australian Guidelines for the Prevention and Control of Infection in Healthcare*. A person with expertise in the health needs of Aboriginal persons and Torres Strait Islanders and a person with expertise in consumer issues were also sought.

Table A3.4. Members of the Infection Control Guidelines Advisory Committee

| Name | Affiliation | Area of expertise |
|--|--|---|
| Professor Brett Mitchell (Chair) | Professor of Nursing, Faculty of Nursing and Health and Director Lifestyle Research Centre – Avondale College | Infection prevention and control |
| Professor Allen Cheng | Professor of Infectious Diseases Epidemiology and the Director of Infection Prevention and Healthcare Epidemiology Unit at Alfred Health | Influenza, hospital acquired infections, antimicrobial resistance |
| Professor Peter Collignon AM | Executive Director, ACT Pathology Infectious Diseases Physician and Microbiologist, Canberra Hospital Professor, Medical School, Australian National University | Infectious disease physician, clinical microbiology and antibiotic resistance |
| Ms Sylvia Gandossi | Infection Prevention and Control Consultant, Western Diagnostics Pathology; Member, Australasian College for Infection Prevention and Control | Infection prevention and control |
| Associate Professor Thomas Gottlieb | Senior Staff Specialist Microbiology and Infectious Diseases – Concord Hospital; Clinical Associate Professor, Department of Microbiology and Infectious Diseases – University of Sydney | Infectious diseases, microbiology and antimicrobial resistance (AMR) |
| Dr Jan Galton | Senior Project Officer, Australian Commission on Safety | Implementation |

| | | |
|--|--|---|
| | and Quality in Health Care | |
| Ms Linda Henderson (until November 2017) | Infection Control Consultant; Infection Control Nurse Advisor – South Australian Health; Member of the Commission's National Hand Hygiene Advisory Committee | Healthcare associated infections |
| Dr Sharon Liberali | Senior Clinical Lecturer – School of Dentistry at the University of Adelaide; Clinical Examiner – Australian Dental Council; Clinical Supervisor; Chair, Infection Control Committee – Australian Dental Association | Dentistry |
| Dr Deborough Macbeth | Assistant Director of Nursing, Infection Control, Gold Coast Health | Healthcare associated infections, nursing |
| Ms Rebecca McCann | Program Manager, Healthcare Associated Infection Unit, Communicable Disease Control Directorate – Department of Health (Western Australia) | Implementation |
| Professor Peter Morris | Deputy Head, Child Health Division, Menzies School of Health Research; Editor, Acute Respiratory Infections Group of The Cochrane Collaboration | Indigenous health, paediatrics, evidence-based medicine |
| Ms Mary Potter | Active in state and national health issues from a consumer/community stand point | Consumer representative |
| Associate Professor Philip Russo | Research Fellow, Deakin/Alfred Health Centre for Quality and Patient Safety Research (until 2018) Director, Cabrini Monash University Department of Nursing Research (from 2019) | Infection prevention and epidemiology |

Terms of Reference

The ICGAC will oversee and provide expertise in updating the *Australian Guidelines for the Prevention and Control of Infection in Healthcare* (2010 Guidelines) to ensure it reflects the best available evidence and is current and relevant for the Australian context. As per the 2010 Guidelines, the updated guidelines will aim to promote and facilitate the overall goal of infection prevention and control: *"The creation of safe healthcare environments through the implementation of practices that minimise the risk of transmission of infectious agents"*.

In undertaking the update, the ICGAC will:

- Determine the scope of the guideline based on consideration of:
 - The currency, accuracy and relevance of the existing 2010 Guidelines.
 - Feedback provided by the broader community and healthcare sector on the 2010 guidelines, including new areas for potential inclusion.
 - The results of a horizon scan funded by the Australian Commission on Safety and Quality in Health Care (ACSQHC), that identifies and assesses the quality of guidelines published since the 2010 Guidelines.
 - Committee member's knowledge/expertise of current evidence or practice in infection prevention and control in Australia.
 - Any other information deemed relevant by the Office of NHMRC or ACSQHC.
- Advise on the evidence evaluation required to update the 2010 Guidelines by:
 - Identifying and prioritising topics requiring the identification of new evidence published since the search period for the 2010 Guidelines.
 - Advising on the scope and clinical questions for the evaluation/s, and methods to identify and evaluate relevant evidence.
- Oversee the update of the 2010 Guidelines by considering (1) and (2) above. This may include, but is not limited to, providing advice on:
 - developing recommendations using NHMRC's preferred method
 - key stakeholders to be included in consultation activities, and
 - comments received during consultation and expert review on the draft revised Guidelines.

The ICGAC will be effective for the period 25 January 2016 to 31 March 2019 and will report to the Council of NHMRC.

Declarations of Interest

Members of NHMRC committees provide high quality, expert and independent advice that allows NHMRC to fulfil its functions under the NHMRC Act. Members are appointed for their expertise and experience across a diverse range of professions and fields. Appointments are also made with consideration of balancing the benefit of having persons with expertise against the risks of their interests biasing a process.

Appointees to committees of NHMRC are required to disclose their interests consistent with section 42A of the NHMRC Act and section 29 of the *Public Governance, Performance and Accountability Act 2013* (PGPA Act). Prior to appointment decisions being made, NHMRC asked each prospective member of the ICGAC to disclose their interests to NHMRC in writing. Prospective members were specifically asked to identify, to the best of their ability, interests including:

- employment by an entity having commercial or other interests in infection prevention and/or control
- financial interests or relationships including ownership, board membership, honoraria, consultancies, or research funding related to infection prevention and/or control
- publications, speeches or expert testimony regarding infection prevention and/or control
- involvement in the development of guidelines, standards, education material or fact sheets related to infection prevention and/or control
- affiliations or associations with any organisation having an interest with infection prevention and/or control.

Under the PGPA Act, members have a responsibility to declare any interests to the whole committee and members have a joint responsibility to decide on the management of any perceived or real conflict. No unmanageable conflicts were identified by the ICGAC or NHMRC and there were no instances in the development process where members of the ICGAC determined that a disclosed interest warranted a member being absent from a discussion or decision.

Throughout the project, members were reminded of their obligation to consider any interest that may have arisen since the last meeting or with any particular agenda items. All disclosures and determinations about interests were recorded in the minutes of the ICGAC meetings. Member's relevant expertise and a summary of their disclosed interests were accessible on the NHMRC website throughout the duration of the project.

Governance

The ICGAC were responsible for advising on the scope of the review and determining priority areas for review and accompanying research protocols. The ICGAC met to critically appraise the Evidence Evaluations and development recommendations regarding infection prevention and control practices, as per the Grading of Recommendations Assessment, Development and Evaluation (GRADE) approach.

The ICGAC advised on the draft Guidelines, including consideration of feedback received during the targeted consultation period held with key stakeholders from the public and private healthcare sectors from 16 November 2017 to 21 December 2017). Public consultation on the draft Guidelines occurred from 13 April 2018 to 15 May 2018. Following this, the ICGAC met to consider public consultation submissions and advise on any revisions required.

All discussions of the ICGAC were robust and open, and decision-making was consensus-based. Minutes from all meetings were recorded by the NHMRC Project Team.

8. Glossary

This section outlines the way in which certain terms are used in these guidelines.

| Term | Meaning |
|---------------------------|--|
| Acinetobacter | An aerobic Gram-negative bacillus commonly isolated from the hospital environment (especially intensive care units) and hospitalised patients; can cause healthcare associated infections, especially wound infections and pneumonia. |
| Active safety devices | Active safety devices, such as needles with guards, are safety devices that require manual activation of the safety feature by the user. |
| Adaptation | This term is used within this Guidelines recommendations and practice statements. The GRADE approach allows for the adoption, adaptation or creation of de novo recommendations from existing quality guidelines. This section captures the adoption or adaption of guidance from the previous 2010 edition of NHMRC's Australian Guidelines for the Prevention and Control of Infection in Healthcare. |
| Aerosols | Microscopic particles < 5 µm in size that are the residue of evaporated droplets and are produced when a person coughs, sneezes, shouts, or sings. These particles can remain suspended in the air for prolonged periods of time and can be carried on normal air currents in a room or beyond, to adjacent spaces or areas receiving exhaust air. |
| Airborne precautions | A set of practices used for patients known or suspected to be infected with agents transmitted person-to-person by the airborne route. |
| Alcohol-based hand rub | A TGA-listed alcohol-containing preparation designed for reducing the number of viable micro-organisms on the hands without the use or aid of running water, and which is listed on the ARTG as a medicinal product. |
| Allergy | Occurs when a person's immune system reacts to allergens in the environment that are harmless for most people. Typical allergens include some medicines, foods and latex. An allergen may be encountered through inhalation, ingestion, injection or skin contact. A medicine allergy is one type of adverse drug reaction. |
| Anteroom | A small room leading from a corridor into a room. |
| Antibiogram | The result of a laboratory testing for the sensitivity of an isolated bacterial strain to different antibiotics. |
| Antibiotic | A substance that kills or inhibits the growth of bacteria, fungi or parasites. |
| Antimicrobial | A chemical substance that inhibits or destroys bacteria, viruses or fungi, and can be safely administered to humans and animals. |
| Antimicrobial resistance | Failure of an antimicrobial to inhibit a microorganism at the antimicrobial concentrations usually achieved over time with standard dosing regimens. |
| Antimicrobial stewardship | An ongoing effort by a health service organisation to reduce the risks associated with increasing antimicrobial resistance and to extend the effectiveness of antimicrobial treatments. It may incorporate several strategies, including monitoring and review of antimicrobial use. |
| Antisepsis | The use of chemical or physical methods to prevent infection by destroying or inhibiting the growth of harmful microorganisms. |
| Antiseptic | Biocides or products that destroy or inhibit the growth of micro-organisms in or on living tissue. |
| Asepsis | 'Freedom from infection or infectious (pathogenic) material'. |
| Aseptic Technique | Aseptic technique is a set of practices aimed at minimising contamination and is particularly used to protect the patient from infection during procedures. Many of the other work practices that form standard precautions are required for aseptic technique, however, adherence to these practices alone does not constitute aseptic technique. Sterile single-use equipment or instruments must be used according to manufacturer's instructions and in such a way that the sterility of the item is maintained. |
| Bay | Refers to a room or area within a general ward in a healthcare facility that generally accommodates up to six patients. |
| Best practice | When the diagnosis, treatment or care provided is based on the best available evidence, which is used to achieve the best possible outcomes for patients. |
| Bloodstream Infection | The presence of live pathogens in the blood, causing an infection. |
| Bundle | A set of evidence-based practices that have been shown to improve outcomes when performed collectively and consistently. The concept was developed by the Institute for Healthcare Improvement in the United States to |

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| | improve the care process and patient outcomes. |
| Carbapenemase-producing <i>Enterobacterales</i> (CPE) | Members of the Enterobacterales that are resistant to carbapenems, a class of 'last resort' antimicrobials for treating serious infections. <i>Note: Taxonomic studies have narrowed the definition of the family Enterobacteriaceae. Some previous members of this family are now included in other families within the order Enterobacterales, and this term is now used in the Guidelines.</i> |
| Carer | A person who provides personal care, support and assistance to another individual who needs it because they have a disability, medical condition (including a terminal or chronic illness) or mental illness, or they are frail or aged. An individual is not a carer merely because they are a spouse, de facto partner, parent, child, other relative or guardian of an individual, or live with an individual who requires care. A person is not considered a carer if they are paid, a volunteer for an organisation, or caring as part of a training or education program. |
| Catheter | A thin, flexible, hollow tube used to add or remove fluids from the body. |
| Chlorhexidine | A biguanide compound used as an antiseptic agent with topical antibacterial activity. |
| Chronic wound | A wound that makes slow progression through the healing phases or displays delayed, interrupted or stalled healing. Inhibited healing may be due to intrinsic and extrinsic factors that impact on the person, their wound and their healing environment. |
| Clean technique | Clean technique refers to practices that reduce the number of infectious agents, and should be considered the minimum level of infection control for non-invasive patient-care activities. Practices include: personal hygiene, particularly hand hygiene, to reduce the number of infectious agents on the skin; use of barriers to reduce transmission of infectious agents (including proper handling and disposal of sharps); environmental cleaning; and reprocessing of equipment between patient uses. |
| Cleaning agent | A product used to remove dirt, stains and microorganisms from surfaces, objects or environments. Cleaning agents can be in the form of powders, liquids, sprays, gels and often contain a surfactant or other chemicals. |
| Cleaning process | Removing dirt and microorganisms from equipment and environmental surfaces. The most effective way to do this is by rubbing or scrubbing the surface with warm water and detergent, followed by rinsing and drying. When MROs are suspected or known to be present, the cleaning process should include the use of a detergent solution followed by the use of a disinfectant so that surfaces are cleaned and disinfected. |
| Clinical governance | An integrated component of corporate governance of health service organisations. It ensures that everyone—from frontline clinicians to managers and members of governing bodies, such as boards—is accountable to patients and the community for assuring the delivery of safe, effective and high-quality services. Clinical governance systems provide confidence to the community and the healthcare organisation that systems are in place to deliver safe and high quality health care. |
| Clinical handover | The transfer of professional responsibility and accountability for some or all aspects of care for a patient, or group of patients, to another person or professional group on a temporary or permanent basis. |
| Clinical waste | Waste material that consists wholly or partly of human or animal tissue, blood or body substances, excretions, drugs or other pharmaceutical products, swabs/dressings, syringes, needles or other sharp instruments. |
| Clinician | A healthcare provider, trained as a health professional, including registered and nonregistered practitioners. Clinicians may provide care within a health service organisation as an employee, a contractor or a credentialed healthcare provider, or under other working arrangements. They include nurses, midwives, medical practitioners, allied health practitioners, technicians, scientists and other clinicians who provide health care, and students who provide health care under supervision. |
| <i>Clostridioides difficile</i> (formerly <i>Clostridium difficile</i>) | A disease of the large intestine caused by toxins produced by the spore forming bacterium <i>Clostridioides difficile</i> . <i>Note: Reclassified in 2016 as Clostridioides difficile (see Lawson PA, Citron DM, Tyrrell KL, Finegold SM. Reclassification of Clostridium difficile as Clostridioides difficile (Hall and O'Toole 1935) Prévot 1938. Anaerobe. 2016; 40:95–9) [436].</i> |
| Cohorting | a group of individuals (patients) that have certain factors (such as age, gender, infectious disease status) in common. |
| Colonisation | The sustained presence of replicating infectious agents on or in the body without causing infection or disease. |
| Communicable | An infection that can be transferred from one person or host to another. |
| Conditional recommendation | See <i>weak recommendation</i> . |
| Contact | The touching of any patient or their immediate surroundings or performing any procedure. |
| Contact point | The area of direct contact of skin to equipment. |

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| Contact precautions | A set of practices used to prevent transmission of infectious agents that are spread by direct or indirect contact with the patient or the patient's environment. |
| Coronavirus Disease 2019 (COVID-19) | The disease caused by SARS-CoV-2. A viral respiratory infection with symptoms ranging from mild to severe. Symptoms may include fever, coughing, sore throat and shortness of breath (this list is not exhaustive). Refer to Coronavirus Disease 2019 (COVID-19) CDNA National Guidelines for Public Health Units for full case definition. |
| COVID-19 | Coronavirus Disease 2019. |
| Decolonisation | Use of topical and/or systemic antibiotics and/or other measures to eradicate MRO's from colonised persons. |
| Decontamination | Use of physical or chemical means to remove, inactivate, or destroy pathogens on a surface or item so that they are no longer capable of transmitting infectious particles and the surface or item is rendered safe for handling, use, or disposal. |
| Detergent Solution | A detergent product which is intended to be used in the cleaning of surfaces or other medical devices diluted with water as per manufacturer's instructions. |
| Discharge clean | See <i>terminal clean</i> . |
| Disinfectant | Chemical or combination of chemicals used for disinfection (source: AS 5369:2023). |
| Disinfection | Process (by physical or chemical means) used to inactivate viable microorganisms on environmental surfaces or equipment to a level previously specified as appropriate for a defined purpose (source: AS 5369:2023). |
| Droplet precautions | A set of practices used for patients known or suspected to be infected with agents transmitted by respiratory droplets. |
| Droplets | Small particles of moisture generated when a person coughs or sneezes, or when water is converted to a fine mist by an aerator or shower head. These particles, intermediate in size between drops and droplet nuclei, can contain infectious microorganisms and tend to quickly settle from the air such that risk of disease transmission is usually limited to persons in close proximity (e.g. at least 1 metre) to the droplet source. |
| Electrolysed water | One class of emerging disinfectants that functions as a broad-spectrum, aqueous chemical oxidant generated by passing a dilute sodium chlorine solution through an electrolytic cell in a process known as electrolysis. |
| Engineering controls | Removal or isolation of a workplace hazard through technology. |
| Environment | The physical surroundings in which health care is delivered, including the building, fixtures, fittings, and services such as air and water supply. Environment can also include other patients, consumers, visitors and the workforce. |
| Environmental cleaning | See <i>cleaning process</i> . |
| Epidemic | A widespread outbreak of an infectious disease. Many people are infected at the same time. |
| Fit checking | A fit check is the minimum requirement at the point of use for staff using particulate filter respirators (PFRs). A fit check must be performed each time a PFR is used to ensure it is properly applied, that a good seal is achieved over the bridge of the nose and mouth, and there are no gaps between the face and respirator. |
| Fit testing | A quantitative or qualitative fit test is a validated method to determine whether the type of respirator being worn provides an adequate seal with a person's face. The test is done while a person is performing several physical movements and talking exercises while wearing PFR that is attached to a testing unit. Qualitative fit tests are fast and simple but can be influenced by the wearer's ability to smell or taste a test substance. Quantitative fit tests require the use of specialised equipment to detect air leaks from the respirator. Fit testing should comply with the recommendations in Standard AS/NZS 1715: 2009. |
| FORM | National Health and Medical Research Council's evidence approach used for the 2010 <i>Australian Guidelines for the Prevention and Control of Infection in Healthcare</i> . |
| GRADE | <i>Grading of Recommendations Assessment, Development and Evaluation</i>) approach is a system for rating the quality of a body of evidence in systematic reviews and grading recommendations in healthcare. |
| Hand hygiene | A general term applying to processes aiming to reduce the number of microorganisms on hands. This includes: application of a waterless antimicrobial agent (e.g. alcohol-based hand rub) to the surface of the hands; and use of soap/solution (plain or antimicrobial) and water (if hands are visibly soiled), followed by patting dry with single-use towels. |
| Health care | The prevention, treatment and management of illness and injury, and the preservation of mental and physical wellbeing through the services offered by clinicians, such as medical, nursing and allied health professionals. |
| Healthcare facility | Any facility that delivers healthcare services. Healthcare facilities could be: hospitals, general practice clinics, dentistry practices, other community-based office practices, day surgery centres, emergency services, domiciliary nursing services, long-term care facilities, aged care facilities, indigenous medical services, alternative health provider facilities and other community service facilities, such as needle exchanges. |

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| Healthcare setting | The environment which health care is provided. A healthcare setting can include but is not limited to hospitals, residential aged care facilities, community/ primary health settings and private office-based settings. |
| Healthcare workers (HCW) | For the purpose of this guideline a HCW means, a person who works in a healthcare setting, whether paid or unpaid, clinical or non-clinical, permanent or casual (includes visiting, sessional and agency), full-time or part-time, in the facilities or services in scope for this guidance. The terms HCW, health worker and staff are interchangeable. |
| Healthcare associated infections | Infections acquired in healthcare facilities ('nosocomial' infections) and infections that occur as a result of healthcare interventions ('iatrogenic' infections), and which may manifest after people leave the healthcare facility. |
| Health service organisation | A health service organisation is a separately constituted health service that is responsible for implementing clinical governance, administration and financial management of a service unit or service units providing health care at the direction of the governing body. Health service organisations include acute, primary and community healthcare services, as well as other services involved in the delivery of health care. |
| High-level disinfection | Disinfection process that kills all microbial pathogens, except large numbers of bacterial endospores. Minimum treatment recommended for reprocessing instruments and devices that cannot be sterilised for use in semi-critical sites. |
| High-efficiency particulate air (HEPA) filter | An air filter that removes >99.97% of particles > 0.3 microns (the most penetrating particle size) at a specified flow rate of air. |
| High-risk patients | Patients with an increased probability of infection due to their underlying medical condition. Often refers to patients in intensive care units, those receiving total parenteral nutrition, and immunocompromised patients. |
| Horizontal measures | A horizontal approach to infection prevention and control measures refers to broad population level approaches attempting reduction of all infections due to all pathogens. Horizontal approaches are long term and favour hand hygiene, chlorhexidine bathing and care bundles. |
| Hospital-grade disinfectant with specific claims | A TGA-listed disinfectant with specific claims that is suitable for general purpose disinfection of hard surfaces, and purposes not involving instruments or surfaces likely to come into contact with broken skin. Refer to TGA Order 104 (Standard for Disinfectants) for further information. These are therapeutic goods that are included in the part of the ARTG for goods known as Listed goods. |
| Hypochlorite | A chlorine-based disinfectant. |
| Immunocompromised | Having an immune system that has been impaired by disease or treatment. |
| Incidence | The number of new events (e.g. cases of disease) occurring in a population over defined period of time. |
| Infectious agent | An infectious agent (also called a pathogen or germ) is a biological agent that causes disease or illness to its host. Most infectious agents are microorganisms (such as bacteria, viruses, fungi and protozoa), parasites and prions. |
| Intermediate-level disinfection | Disinfection process that kills all microbial pathogens except bacterial endospores. This process is effective against most bacterial, viral and fungal spores (source: AS5369:2023). Minimum treatment recommended for reprocessing instruments and devices for use in non-critical sites, or where there are specific concerns regarding contamination of surfaces with species of mycobacteria (e.g. <i>Mycobacterium tuberculosis</i>). |
| Invasive medical device | Devices which in whole or part enter the body through an orifice or through any surface of the body. This includes penetrating skin, mucous membranes, organs or internal cavities of the body. Examples include surgical instruments, implantable devices, dental equipment, intravascular devices, medical and therapeutic devices. |
| Invasive procedure | Entry into tissues, cavities or organs or repair of traumatic injuries. |
| Key information | This term is used within this Guidelines recommendations and practice statements. It captures key information relating to the quality of the evidence, harms and benefits of the intervention, values and preferences of the target population, resource and other considerations. |
| Key parts | Parts of the procedure equipment or solutions that must remain aseptic throughout clinical procedures, in order to protect the patient from contamination or infection. For example a wound dressing, catheter lubrication, syringe tip, needle etc. In IV therapy, key parts are usually those that come into direct contact with the liquid infusion—for example needles, syringe tips and exposed central line lumens. |
| Key sites | Susceptible open or broken wounds, surgical or intravenous access sites. |
| <i>Klebsiella pneumoniae</i> | Gram-negative bacteria frequently responsible for healthcare associated infections of wounds and urinary tract, particularly in immunocompromised patients; may also cause pneumonia. |
| Long-term care facilities | A range of residential and outpatient facilities designed to meet the bio-psychosocial needs of persons with sustained self-care deficits. |
| Low-level disinfection | Disinfection process that kills all vegetative bacteria (except tubercle bacilli), lipid viruses, some nonlipid viruses, and some fungi, but not bacterial spores (source: AS 5369:2023). An adjunct to cleaning alone when devices for use |

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| | in non-critical sites are reprocessed and when only vegetative bactericidal activity is needed. |
| Manual cleaning | Physical (mechanical or manual) cleaning is the most important step in cleaning. It is a cleaning process which applies friction (rubbing/scrubbing) and fluids (detergent). It intends to remove foreign material (e.g. blood, body substances, micro-organisms and dust) from a surface or an object. |
| Mechanical cleaning | See <i>Manual cleaning</i> |
| Medical device disinfectant | A TGA-included disinfectant which is used to reprocess reusable medical devices. These are therapeutic goods that are 'Included' on the ARTG. |
| Medicine | A chemical substance given with the intention of preventing, diagnosing, curing, controlling or alleviating disease, or otherwise improving the physical or mental wellbeing of people. These include prescription, nonprescription, investigational, clinical trial and complementary medicines, irrespective of how they are administered. |
| Methicillin-Resistant <i>Staphylococcus Aureus</i> (MRSA) | Strains of <i>Staphylococcus aureus</i> that are resistant to many of the antibiotics commonly used to treat infections. Epidemic strains also have a capacity to spread easily from person-to-person. |
| Microorganism | Most infectious agents are microorganisms. These exist naturally everywhere in the environment and not all cause infection e.g. 'good' bacteria present in the body's normal flora. Parasites, prions and several classes of microorganism—including bacteria, viruses, fungi and protozoa—can be involved in either colonisation or infection, depending on the susceptibility of the host. |
| Multidisciplinary team | A team including clinicians from multiple disciplines who work together to deliver comprehensive care that deals with as many of the patient's health and other needs as possible. The team may operate under one organisational umbrella or may be from several organisations brought together as a unique team. As a patient's condition changes, the composition of the team may change to reflect the changing clinical and psychosocial needs of the patient. Multidisciplinary care includes interdisciplinary care. (A discipline is a branch of knowledge within the health system.) |
| Multi-Resistant Organisms (MROs) | In general, bacteria that are resistant to one or more classes of antimicrobial agents and are usually resistant to all but one or two commercially available antimicrobial agents. |
| Needleless devices | Needleless devices (e.g. connectors, vascular access devices, access ports) provide an easy access point for intravascular infusion connections. Needleless devices do not use needles for procedures such as the collection or withdrawal of body substances after initial venous or arterial access is established, or administering medication or fluids. |
| Negative pressure room (also referred to as Class N - negative pressure and/or Type 5 - respiratory isolation) | A single-occupancy patient-care room used to isolate persons with a suspected or confirmed airborne infectious disease. Environmental factors are controlled in negative pressure rooms to minimise the transmission of infectious agents that are usually transmitted from person-to-person by droplet nuclei associated with coughing or aerosolisation of contaminated fluids. The air handling system operates at a lower pressure with respect to adjacent areas such as the anteroom and corridor and is exhausted to the outside. |
| Not Set | This term generally accompanies best practice statements which have followed GRADE but the strength of the evidence has not been determined. |
| Outbreak | An outbreak of an infection is a sudden increase in the number of cases of an infection above what is normally expected in the population. Outbreaks are usually limited to specific geographic areas, and are linked by time, person, and/or place. |
| Outcome | The status of an individual, group of people or population that is wholly or partially attributable to an action, agent or circumstance. |
| Pandemic | An epidemic that is geographically widespread, occurring throughout a region or even throughout the world. |
| Passive safety device | Passive safety devices are single handed devices where the safety device activates automatically with no extra action being required by the user, such as a retractable spring loaded safety syringe. |
| Pathogen | See <i>infectious agent</i> . |
| Patient | A person who is receiving care in a health service organisation. |
| Patient contact | Involves touching the patient and their immediate surroundings, or performing any procedure on the patient. |
| Patient surroundings | All inanimate surfaces that are touched by or in physical contact with the patient (such as bed rails, bedside table, bed linen, invasive devices, dressings, personal belongings and food) and surfaces frequently touched by healthcare workers while caring for the patient (such as monitors, knobs and buttons). |
| Patient-care area | The room or area in which patient care takes place. |
| Percutaneous injury | An injury that results in a sharp instrument/object—for example a needle or scalpel—cutting or puncturing the skin. |

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| Personal protective equipment (PPE) | A variety of barriers used alone or in combination to protect mucous membranes, skin, and clothing from contact with infectious agents. PPE includes gloves, masks, respirators, protective eyewear, face shields, and gowns/aprons. |
| Physical cleaning | See <i>Manual cleaning</i> |
| Practical Information | This term is used within this Guidelines recommendations and practice statements. It provides practical guidance on how each recommendation or practice statement can be put into practice. |
| Practice Statement | Set for areas which are not covered by a systematic review of the evidence, but where the provision of clinical guidance is deemed important. The development of practice statements is primarily based on best practice as advised by expert consensus and aligned with the GRADE approach where available evidence and judgements are considered together however a strength is not assigned. |
| Prevalence | The number of events (e.g. cases of disease) present in a defined population at one point in time. |
| Procedure | An act of care for a patient where there is a risk of direct introduction of a pathogen to the patient. |
| Randomised Controlled Trial (RCT) and non-randomised control trial (NRCT) | A clinical trial where at least two treatment groups are compared, one of them serving as the control group, and treatment allocation is carried out using a random, unbiased method. A non-randomised controlled trial compares a control and treatment group but allocation to each group is not random. Bias is more likely to occur in NRCT. |
| Rationale | This term is used within this Guidelines recommendations and practice statements. It provides overarching justification for the stated advice. |
| Reprocessing | This is a multistep process that includes cleaning, inspection/examination and assembly, functional testing (if applicable), disinfection (if applicable), packaging and labelling, and sterilisation (if applicable) (source: AS 5369). |
| Research evidence | This term is used within this Guidelines recommendations and practice statements. It defines the research question underpinning the recommendation in the PICO (population, intervention, comparator, outcomes) format. |
| Respiratory hygiene and cough etiquette | A combination of measures designed to minimise the transmission of respiratory pathogens via droplet or airborne routes in healthcare settings. |
| Respiratory protection (Masks) | An item of personal protective equipment that is worn over the mouth and nose to protect the wearer from inhaling infectious agents. Masks also protect the wearer from splashes or sprays of blood or body fluid to the mouth and nose. <ul style="list-style-type: none"> • Surgical/procedure mask: a single use fluid repellent mask utilised to cover the nose and mouth to contain respiratory secretions of the wearer, to prevent splashes or sprays of blood or body fluids and to prevent droplet inhalation by the wearer • Reusable mask: Similar in design to surgical masks, includes commercially available reusable cloth or other material mask • Particulate Filter Respirator (PFR): A mask which forms a tight seal around the face. These masks have a higher filtration and are recommended for protection against airborne particles such as fine dust, vapours, smoke and aerosolising or airborne infectious diseases. The most common PFRs are P2 or N95 respirators: P2 respirators are those that comply with the Australian Standard AS/NZS 1716:2012 Selection, use and maintenance of respiratory protective devices. N95 respirators are those that comply with the United States National Institute for Occupational Safety and Health (NIOSH) 42 CFR part 84, which is a less stringent standard • Powered air-purifying respirator (PAPR): A reusable system which includes a hood or helmet that uses a fan to circulate filtered air to the wearer. |
| Reusable medical device (RMD) | Medical device designated or intended by the manufacturer as suitable for reprocessing and reuse (source: AS 5369). |
| Risk | The chance of something happening that will have a negative impact. Risk is measured by the consequences of an event and its likelihood. |
| Risk assessment | Assessment, analysis and management of risks. It involves recognising which events may lead to harm in the future, and minimising their likelihood and consequences. |
| Risk management | The design and implementation of a program to identify and avoid or minimise risks to patients, employees, volunteers, visitors and the organisation. |
| Routine | Performed as part of usual practice (as opposed to the use of additional measures in specific circumstances e.g. where invasive procedures are conducted or in the event of an outbreak). |
| Safety-engineered device | Safety-engineered devices are sharp devices with an integrated safety feature designed to shield the sharp object after use and reduce the risk of injury. See Passive and Active safety devices. |
| SARS-CoV-2 | Severe Acute Respiratory Syndrome Coronavirus 2. |
| Screening | A process of identifying patients who are at risk, or already have a disease or injury. Screening requires enough knowledge to make a clinical judgement. |
| Sharps | Instruments used in delivering healthcare that can inflict a penetrating injury, e.g. needles, lancets and scalpels. |

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| Single patient use equipment | Equipment to be used for one patient only. Single patient use equipment should not be used on more than one patient. ³³⁶ |
| Single use equipment | Equipment to be used on a single occasion for one patient only and then discarded after use. Single use equipment should not be reused on the same patient or any other patient. Single use equipment is labelled with this symbol:  ³³⁶ |
| Specific claims | The term 'specific claims' covers virucidal, sporicidal, tuberculocidal, fungicidal or other biocidal activity. Except where claims of activity against fungi (yeast and mould) for excluded products are concerned, these claims mean a product is regulated as a listed Other Therapeutic Good on the ARTG. |
| Standard precautions | Work practices that constitute the first-line approach to infection prevention and control in the healthcare environment. These are recommended for the treatment and care of all patients. |
| Statutory Requirement | This advice reflects a practice statement or recommendation. The terminology 'statutory requirement' is used to further indicate where there is also a mandated requirement/s by the Commonwealth or the States/Territories, which must be considered when implementing the advice at the local level. It is important to note that statutory requirements vary across states and territories, and in their applicability to health service delivery sectors and settings. |
| Sterile | Free from viable microorganisms; usually described as a probability (e.g. the probability of a surviving microorganism being 1 in 1 million). |
| Sterile technique | Sterile technique aims to eliminate microorganisms from areas and objects, and should be undertaken by all healthcare workers undertaking invasive medical procedures. This includes: ensuring that everything within a defined radius is clean and sterile, or as a minimum is subject to high level chemical or thermal disinfection; use of skin antiseptics and sterile personal protective equipment; and reprocessing of instruments between patient uses. Due to the natural multitude of organisms in the atmosphere it is not possible to achieve a true sterile technique for most invasive procedures in a typical hospital environment (even when wearing sterile gloves). Sterile techniques can only be achieved in controlled environments such as a laminar air flow cabinet or a specially equipped theatre. The commonly used term, 'sterile technique' is therefore inaccurate, as practitioners are not actually achieving their stated objective. |
| Sterilisation | Use of a physical or chemical procedure to destroy all microorganisms including substantial numbers of resistant bacterial spores. |
| Strain | A strain is a genetic variant or subtype of a micro-organism (e.g. a virus, bacterium or fungus). Some strains may be more dangerous or difficult to treat than others. |
| Strong recommendation | Confident that the desirable effects of adherence to a recommendation outweigh the undesirable effects. Overall the recommendation is based on high quality evidence and is strongly recommended for implementation. |
| Surface barrier | Barriers (e.g. clear plastic wrap, bags, sheets, tubing or other materials impervious to moisture) designed to help prevent contamination of surfaces and equipment. |
| Surgical ABHR | Antiseptic hand rub performed preoperatively by the surgical team following the surgical pre-wash to eliminate transient flora and reduce resident skin flora. Such antiseptics often have persistent antimicrobial activity. Use should be in accordance with manufacturers' instructions. Surgical ABHRs are often a waterless and high percent alcohol solution. |
| Surgical hand preparation | The process of eliminating transient and reducing resident flora prior to surgery. This comprises removal of hand jewellery, performing hand hygiene with liquid soap if hands are visibly soiled, removing debris from underneath fingernails and scrubbing hands and forearms using a suitable antimicrobial formulation. |
| Surgical masks | Loose-fitting, single-use items that cover the nose and mouth. These include products labelled as dental, medical procedure, isolation and laser masks. |
| Surgical-site infection | An infection at the site of a surgical operation that is caused by the operation. |
| Surveillance | Disease surveillance is an epidemiological practice by which the spread of disease is monitored in order to establish patterns of progression. The main role of disease surveillance is to predict, observe and minimise the harm caused by outbreak, epidemic and pandemic situations, as well as increase knowledge as to what factors might contribute to such circumstances. |
| Targeted surveillance | A process in which data are collated on the susceptibilities and resistances of disease-causing microbes to various antimicrobial treatments. Targeted surveillance gathers data that is not generated by routine testing: specific species or groups of species are examined in detail to answer important questions that cannot be addressed by passive surveillance. |

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| Terminal clean | Cleaning process required after patient(s) has vacated the room, either through room transfer or discharge. |
| TGA-Listed | Must comply with Therapeutic Goods Order No. 104 (TGO 104 – Standard for Disinfectants). Pre-market evaluation will be conducted on new ingredients and/or new specific claims. |
| Traceability | Ability to trace the reprocessing history, use and location of a reusable medical device. At a minimum, traceability systems are required to enable reusable medical devices and patients to be linked at the point of use, and to allow recall of non-conforming products (source: AS 5369:2023). |
| Transmission-based precautions (formerly additional precautions) | Extra work practices in situations where standard precautions alone may be insufficient to prevent infection (e.g. for patients known or suspected to be infected or colonised with infectious agents that may not be contained with standard precautions alone). |
| Ultra-violet light | Light in the UV-C wavelength range (200 to 270 nanometers) that has microbiocidal properties against multiple pathogens, including <i>C. difficile</i> and other healthcare associated pathogens. |
| Vancomycin Resistant Enterococci (VRE) | Enterococci are Gram-positive bacteria that are naturally present in the intestinal tract of all people. Vancomycin is an antibiotic to which some strains of enterococci have become resistant. These resistant strains are referred to as VRE and are frequently resistant to other antibiotics generally used to treat enterococcal infections. |
| Vertical measures | A vertical approach refers to a narrow program focusing on a single pathogen and aims to reduce infection or colonisation. Vertical approaches are generally indicated when standard measures have been exhausted. Vertical approaches are short term and generally favour active surveillance. |
| Ward | A healthcare unit which consists of a group of inpatient beds (generally 10 – 40 beds) designed for a particular type of service or care. |
| Weak recommendation | Concludes that the desirable effects of adherence to a recommendation probably outweigh the undesirable effects. Overall the recommendation is based on supportive evidence and a strong theoretical rationale and is recommended for implementation. |
| Workforce | All people working in a health service organisation, including clinicians and any other employed or contracted, locum, agency, student, volunteer or peer workers. The workforce can be members of the health service organisation or medical company representatives providing technical support who have assigned roles and responsibilities for care of, administration of, support of, or involvement with patients in the health service organisation. |

9. Abbreviations and acronyms

| Abbreviations and acronyms | Meaning |
|----------------------------|---|
| ACH | air changes per hour |
| ABHR | alcohol-based hand rub |
| ACIPC | Australasian College for Infection Prevention and Control |
| ACSQHC | Australian Commission on Safety and Quality in Health Care |
| AGREE | Appraisal of guidelines research and evaluation |
| AMR | antimicrobial resistance |
| ARTG | Australian Register of Therapeutic Goods |
| AS | Australian Standard |
| AS/NZS | Joint Australian/New Zealand Standard |
| AusHFG | Australasian Health Facility Guidelines |
| BCG | Bacillus Calmette-Guérin |
| BSI | bloodstream infection |
| CA-MRSA | community-acquired methicillin-resistant <i>Staphylococcus aureus</i> |
| CAUTI | catheter-associated urinary tract infection |
| CBIC | Certification Board of Infection Control |
| CCJD | classical Creutzfeldt-Jakob disease |
| CJD | Creutzfeldt-Jakob disease |
| CDC | Centers for Disease Control and Prevention (US) |
| CDNA | Communicable Diseases Network Australia |
| CEO | chief executive officer |
| CHG | chlorhexidine |
| CHG-impregnated | chlorhexidine-impregnated |
| CPE | carbapenemase-producing <i>Enterobacterales</i> |
| CVC | central venous catheters |
| DNA | deoxyribonucleic acid |
| EN | European Standards |
| EPP | exposure-prone procedures |
| ESBL | extended spectrum beta-lactamase |
| GP | general practitioner |
| GRADE | Grading of Recommendations Assessment, Development and Evaluation |
| HAI | healthcare associated infection |
| HBeAg | hepatitis B e antigen |
| HBsAg | HBV surface antigen |
| HBV | hepatitis B virus |
| HCV | hepatitis C virus |
| HEPA | high efficiency particulate air |
| HICPAC | Healthcare Infection Control Practices Advisory Committee |
| HIV | human immunodeficiency virus |
| HTLV-I | human T-cell lymphotropic virus type I |

| Abbreviations and acronyms | Meaning |
|----------------------------|--|
| IHI | Institute for Healthcare Improvement (US) |
| ISO | International Organization for Standardisation |
| IPC | infection prevention and control |
| IVD | intravascular device |
| MMR | measles mumps rubella vaccine |
| MRGN | multi-resistant Gram negative |
| MRO | multi-resistant organism |
| MRSA | methicillin-resistant <i>Staphylococcus aureus</i> |
| NaOH | sodium hydroxide |
| NATA | National Association of Testing Authorities |
| NHHI | National Hand Hygiene Initiative |
| NHIG | normal human immunoglobulin |
| NHMRC | National Health and Medical Research Council |
| NICE | National Institute for Health and Clinical Excellence |
| NNDSS | National Notifiable Diseases Surveillance System |
| NPS | National Prescribing Service |
| NRL | natural rubber latex |
| NSQHS | National Safety and Quality Health Service |
| NZS | New Zealand Standard |
| OMT | outbreak management team |
| PAPR | powered air-purifying respirator |
| PBS | Pharmaceutical Benefits Scheme |
| PEG | percutaneous endoscopic gastrostomies |
| PEP | post-exposure prophylaxis |
| PICC | peripherally inserted central venous catheter |
| PICO | population, intervention, comparator, outcomes (research question) |
| PIVC | peripheral intravenous catheter |
| PPE | personal protective equipment |
| PSAE | <i>Pseudomonas aeruginosa</i> |
| PVL | panton-valentine leukocidin |
| RNA | ribonucleic acid |
| RPBS | Repatriation Pharmaceutical Benefits Scheme |
| RSV | respiratory syncytial virus |
| SAB | <i>staphylococcus-aureus-bacteraemia</i> |
| SAL | sterility assurance level |
| SARS | severe acute respiratory syndrome |
| STEC | shiga toxin-producing <i>Escherichia coli</i> |
| SSI | surgical-site infection |
| TB | tuberculosis |
| TGA | Therapeutic Goods Administration |

| Abbreviations and acronyms | Meaning |
|----------------------------|-----------------------------------|
| VAP | ventilator-associated pneumonia |
| vCJD | variant Creutzfeldt-Jakob disease |
| VRE | vancomycin-resistant enterococci |
| WHO | World Health Organization |
| WHS | workplace health and safety |
| ZIG | Zoster immune globulin |

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11. Amendments to the Guidelines

"Process to maintain the guidelines

NHMRC, in partnership with the Australian Commission on Safety and Quality in Health Care (ACSQHC), updated the Australian Guidelines for the Prevention and Control of Infection in Healthcare (the guidelines) in May 2019. NHMRC established an advisory group from January 2016 to March 2019 to oversee the updating process. This advisory group was dissolved after the guidelines release in May 2019.

NHMRC agreed to maintain the guidelines following their release with technical content support provided by the ACSQHC. To facilitate the maintenance process the guidelines are published and maintained by NHMRC on the MAGICapp platform.

How queries and requests for change are processed

NHMRC's Infection Prevention and Control team maintains a log of all guideline queries received and forwards it to the ACSQHC each month. Upon receiving a query NHMRC will either respond directly to the sender or refer the query to the ACSQHC for a response.

If the ACSQHC and NHMRC identify that clinical content in the guidelines may need to be changed then the following will be considered:

1. If the change is a minor edit or clarification which does not require a review of the evidence, the ACSQHC will provide NHMRC with the correct wording, and the guidelines will be amended in an upcoming edit cycle (currently February, June and October). Edits will be listed in MAGICapp.
2. If the change is more substantial, ACSQHC and NHMRC will discuss an approach and may seek further advice from experts such as ACSQHC's Healthcare Associated Infection Advisory Committee on:
 - a. whether a change needs to be made immediately, or b. should be deferred until a full review of the guidelines takes place.
3. New or emerging evidence, a significant change to the body of evidence underpinning the guidelines, and/or implications for the guidelines' recommendation(s), will be considered as part of a full revision of the guidelines.

When both agencies agree that a full review is warranted, a formal revision process will be activated. This process will include the establishment of an expert committee to advise on the revision and a public consultation process being conducted in accordance with the National Health and Medical Research Council Act (1992).

All edits and changes made to the guidelines will be recorded in the version history log under the About Tab.

Amendments since the guidelines were released in May 2019

As amendments occur, the Table of Amendments below will be revised to ensure users are aware of all changes to the Guidelines.

| Amendment detail | Date updated |
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| <ul style="list-style-type: none"> • Updated Section 6.4 'Type and duration of precautions for specific infections and conditions', Table A2.5 'Precautions for specific infections and conditions' with revised recommendations for diphtheria and new recommendations for Nipah virus. | April 2026 |
| <ul style="list-style-type: none"> • Updated Scope (of AICG) with additional information about level of risk dependent on the environment and situation. • Updated Section 3 "Standard Transmission-based precautions", with additional statement regarding risk assessment of patients. • Updated Section 3.1.4 "Reprocessing", including change in terminology from 'cleaning solution' to 'cleaning agent' in all instances, plus addition of updated information for cleaning agents per Standard AS 5369:2023. • Updated Section 3.1.4 "Reprocessing" to include new section for Traceability per Standard AS 5369:2023. • Updated Section 3.1.4 "Reprocessing" to include updated information for: "Medical device sterilants and disinfectants", and "Storage and maintenance". • Updated Section 3.1.4 "Reprocessing" to include updated information for "Reprocessing of flexible endoscopes", plus update to GESA guidelines date. • Updated Section 3.1.4 "Reprocessing" to include updated information for "Further considerations", regarding Prions. • Updated Section 3.1.5 "Respiratory hygiene and cough etiquette" with additional information about risk assessment for patients presenting aerosol-generating behaviours. • Updated Section 3.2.4 "Airborne precautions", Recommendation 27 to replace 'P2 Respirators' heading with 'Particulate filtration respirators (PFR)' heading and corresponding information • Updated Section 3.2.4 "Airborne precautions", Recommendation 27 to include updated information about use for PFRs, as well as updated recommendations for healthcare workers with facial hair. • Updated Section 3.2.4 "Airborne precautions", to include new section and information about PAPRs. 2x new references added alongside in-text citations (Licina & Silvers 2021, Jun et al 2023) • Updated Section 3.2.4 "Airborne precautions" - Practical Information: Patient Placement, with spelling amendment and replacement of Standard AS 1668:2:2012 and Amendment 2:2016 with Standard AS 1668:2:2024. • Updated Section 3.3 "Personal protective equipment (PPE)" with updated TGA information regarding PPE regulations, and information about risk assessments for PPEs. • Updated Section 3.3 "Personal protective equipment (PPE)" to include new section for "Managing breaches in PPE". • Updated Section 3.5 "Applying standard and transmission-based precautions during procedures", to include new | October 2024 |

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| <ul style="list-style-type: none"> subsection 3.5.4 "Aerosol-generating procedures (AGPs)". Updated Section 4.1.2 "Roles and responsibilities", with minor change to date for Australian Dental Association's Guidelines for Infection Control citation (3rd Edition, 2015 to 5th Edition, 2024) Updated Section 4.6.1 "Mechanisms for influencing healthcare associated infection through environmental design", with updated information regarding Ventilation Rate with reference to Standard AS 1668:2:2024. Updated Section 5.11 "Examples of how to perform aseptic technique", to include updated Step 5 about recommended use of gloves for peripheral and central access IV procedures. 2x new references added (Rowley 2001, Rowley et al 2010) as in-text citations and to reference list. Updated Table A2.5 in Section 6.4 to amend detail for the SARS-CoV-2 information; included new entry for Mpox and corresponding infection type, transmission route, required precautions, duration of precautions, additional comments. Updated reference items 179, 341, 399 (Australian Dental Association Guidelines for Infection Control) to replace 3rd Edition 2015 with 5th Edition 2024. Updated reference item 379 (RACGP Infection Prevention and Control Standards) with new URL and inclusion of 2022 and 2024 dates. Glossary updates for the following terms: Cleaning process, Cleaning agent, Disinfectant, Disinfection, High-level disinfection, Intermediate-level disinfection, Low-level disinfection, Reprocessing, Reusable medical device (RMD), Sterile, Traceability, Coronavirus Disease 2019 (COVID-19), Fit checking, Fit testing, Healthcare setting, Health service organisation, Healthcare worker (HCW), Outbreak, Personal protective equipment (PPE), Respiratory protection (Masks), SARS-CoV-2, COVID-19 | |
| <ul style="list-style-type: none"> Re-formatted Table A.2.4. Use of standard and transmission-based precautions. | April 2024 |
| <ul style="list-style-type: none"> Updated all references to Australian Standards e.g. AS ISO 31000:2018 and AS 5369:2023. Re-formatted Table A.2.4. Use of standard and transmission-based precautions | February 2024 |
| <ul style="list-style-type: none"> Updated Section 3.1.2 on Retractable devices to refer to new Standards: Standard AS 3825: 2020, Standard AS 23907:2023 or relevant international standard e.g. ISO 23907: 2019. Updated Section 3.1.2 on the handling and disposal of sharps to refer to new Standards Australian Standard AS 23907:2023 or relevant international standard e.g. ISO 23907: 2019. | October 2023 |
| <ul style="list-style-type: none"> Updated references to the "National Hand Hygiene Initiative Manual" to the "National Hand Hygiene Initiative (NHHI) implementation guide" Updated link to reflect change | September 2023 |
| <ul style="list-style-type: none"> Changed all mentions of "outbreak potential" to "in the event of an outbreak" in Table A2.4 and Table A2.5. Removed droplet precautions from Table A2.5 Changed link to CDNA SoNG to ICEG Guidance in Table A2.5 Updated wording in Table A2.1 including risk category Extreme renamed to Very high risk/ Extreme. Category also changed in Table A2.2 Added wording on disinfection of reuseable semi-critical medical devices to Section 3.1.4 Updated definition of Disinfection in Section 3.1.4 | August 2023 |
| <ul style="list-style-type: none"> Amended Table 18 Selection of glove type (Section 3.3 Personal Protective Equipment) to include handling of chemotherapeutic agents and add note on sustainable use of gloves under reuseable utility gloves. | March 2023 |
| <ul style="list-style-type: none"> Updated Table 2.2 Minimum cleaning frequency to include instructions to change when visibly soiled or torn and separate advice for disposable and non-disposable curtains and blinds | February 2023 |
| <ul style="list-style-type: none"> Amended table A2.1 and A2.2 in Appendix 2: 6.1 Recommended routine cleaning frequencies to reflect revised risk ratings. | September 2022 |
| <ul style="list-style-type: none"> Amended text in Section 6.3 and Table A2.4 to include an Environmental Cleaning column, and changed the Single room/cohort column to Patient Placement. | August 2022 |
| <ul style="list-style-type: none"> Amended text to include the updated Workforce Immunisation Risk Matrix and corresponding text in Section 4.2 Staff health and safety. Amended text to include COVID-19 in Table 32. Staff exclusion periods for infectious illnesses Amended text on goggles contained in Practical info under recommendation 30 in Personal protective equipment. | June 2022 |
| <ul style="list-style-type: none"> Included meningitis into the syndromic section of Table A2.5 in Section 6.4 when the type of infection is unknown or under investigation. | August 2021 |

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| <ul style="list-style-type: none"> Amended three entries in Table A2.5 in Section 6.4. Corrected the required precautions for parainfluenza to S + D, deleted the pneumonia entry under <i>Staphylococcus aureus</i> as only standard precautions apply and amended the additional comments for SARS-CoV-2 to align with the latest Infection Control Expert Group (ICEG) advice. Added references to the ACSQHC's environmental cleaning resources in sections 3.1.3, 3.1.3.1, 3.2.1 and 4.1.3 that support the AICGs recommendations on this topic. Amended Section 3.3 Personal protective equipment to highlight what it means to use PPE as part of standard or transmission-based precautions. Amended Sections 3.1.2 Use and management of sharps, safety engineered devices and medication vials and 4.1.5 Taking an organisational systems approach to infection prevention quality and safety to further highlight Passive and Active safety device use. Amended Section 1 Introduction to highlight the use of the guidelines in acute health settings. | July 2021 |
| <ul style="list-style-type: none"> Amended Section 3.1.2 Use and management of sharps, safety engineered devices and medication vials, to highlight Passive and Active safety devices, the two broad categories of safety-engineered devices. Also added Standard AS 3825: 2020 to the reference for more information on the handling and disposal of sharps. Added definitions for Safety-engineered device, Passive safety device and Active safety devices to Section 8 Glossary. | April 2021 |
| <ul style="list-style-type: none"> Amended the definition on Cohorting to clarify that the act of cohorting can be applied in different scenarios. Amended Section 3.3 to clarify that the extended use of PPE between patients in a cohort situation is not supported. | February 2021 |
| <ul style="list-style-type: none"> Merged the <i>Haemophilus influenzae</i> type b (HiB) entries relating to pneumonia in Table A2.5 in section 6.4 and added comments on its greater presentation in children than in adults. | January 2021 |
| <ul style="list-style-type: none"> Revised the entry on Respiratory illness in Table A2.5 in section 6.4 to accurately reflect the required precautions. | December 2020 |
| <ul style="list-style-type: none"> Amended the 'Process to maintain guidelines' text in Section 11 to reflect amendments on NHMRC's website. | November 2020 |
| <ul style="list-style-type: none"> Updated Section 3 Summary page to include risk mitigation information where clinical environments or activities don't lend themselves to optimal conditions to prevent cross infection. Amended steps 3 and 4 removing PPE images in Section 3.3 to accurately reflect the presence of a gown or face shield in each step. Updated Table 19 in Section 3.4.1 to clarify the term chronic wound for MRSA's. Updated Table 24 in Section 3.5.2.1 to clarify the evidence available for cleaning the urethral meatus. Revised Table A2.5 in Section 6.4 to improve clarity. Actions included: removal of infections/conditions where only Standard Precautions apply, inclusion of precautions for syndromic presentations and provided further detail in the Additional Comments column. | October 2020 |
| <ul style="list-style-type: none"> Updated Table 14 in Section 3.3 to revise the sequence for doffing PPE Updated terminology for <i>Clostridioides difficile</i> in Recommendation 6 and associated references Amended typographical errors in section 3.4.2 for MRSA, VRE and staphylococcus in outbreak investigation | June 2020 |
| <ul style="list-style-type: none"> Updated terminology in Table A2.5 for Abscesses, ESBL and SARS (clarified P2 masks) Updated Table A2.4 with further clarification on staffing requirements Updated curtains and blinds text in Table A2.2 to distinguish between cloth and disposable curtains Updated glossary for single use items Updated links to the Australian Immunisation Handbook Updated further references to the National Hand Hygiene Initiative (NHHI) and added links | May 2020 |
| <ul style="list-style-type: none"> Added links to companion resources | February 2020 |
| <ul style="list-style-type: none"> Updated all Hand Hygiene Australia references to reflect the change in coordination of the National Hand Hygiene Initiative (NHHI). Updated Table A2.5 – amended wording in the additional comments for Amoebiasis (<i>Entamoeba histolytica</i>) Updated Table A2.5 – transmission route, required precautions, duration of precautions and additional comments for Extended-spectrum beta-lactamase producing Enterobacteriaceae Revised wording on replacing a PIVC based on clinical indication | January 2020 |
| <ul style="list-style-type: none"> Updated Table A2.5 - required precautions for Herpes Zoster (localised shingles). Revised - Section 3.1.6 Aseptic Technique & Section 3.1.6.1 Example – Aseptic Non-Touch Technique (ANTT®). Glossary Update – Aseptic Technique Definition | December 2019 |

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| <ul style="list-style-type: none"> Updated Table A2.5 - required precautions for Rhinovirus. | November 2019 |
| <ul style="list-style-type: none"> Glossary update – single-use and single-use devices. Antimicrobial-coated sutures. Minor grammatical update. | October 2019 |
| <ul style="list-style-type: none"> Updated Table A2.5 – required precautions for Herpes Zoster (localised shingles). | August 2019 |
| <ul style="list-style-type: none"> Updated the position of the members of the Infection Control Guidelines Advisory Committee. Updated Table A2.5 - the required precautions and added additional comments on Bronchiolitis, Respiratory Syncytial Virus (RSV) and Rhinovirus. Minor grammatical update. | June 2019 |

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Annex: All evidence profiles, sorted by sections

1. Introduction

2. Basics of infection prevention and control

2.1. Companion resources

2.2. Infection prevention and control in the healthcare setting

2.3. Overview of risk management in infection prevention and control

2.4. A patient-centred approach

2.4.1. Involving patients in their care

3. Standard and transmission-based precautions

3.1. Standard precautions

3.1.1. Hand hygiene

Clinical question/ PICO

Population: Healthcare Workers

Intervention: Hand hygiene

Comparator: No hand hygiene

Summary

Research question

- **Population:** Healthcare Workers
- **Intervention:** Hand hygiene
- **Comparator:** No hand hygiene
- **Outcome:** Reduced transmission and infection

Clinical question/ PICO

Population: Healthcare Workers

Intervention: ABHRs containing between 60% and 80% v/v ethanol or equivalent

Comparator: ABHRs with lower concentration of ethanol or equivalent

Summary

Research question

- **Population:** Healthcare Workers

- **Intervention:** ABHRs containing between 60% and 80% v/v ethanol or equivalent
- **Comparator:** ABHRs with lower concentration of ethanol or equivalent
- **Outcome:** Decontamination of hands

Clinical question/ PICO

Population: Healthcare Workers
Intervention: Hand hygiene
Comparator: Non-alcohol based products

Summary

Research question

- **Population:** Healthcare Workers
- **Intervention:** Hand hygiene
- **Comparator:** Non-alcohol based products
- **Outcome:** Decontamination of hands

Clinical question/ PICO

Population: Healthcare Workers
Intervention: Hand hygiene comparing different concentrations of alcohol, and of different alcohols e.g. ethyl, methyl, isopropyl, in reducing the risk of transmission of Clostridium difficile and non-enveloped viruses
Comparator: Washing with water and soap/detergent/chlorhexidine, other concentrations of same alcohol, other alcohols on Clostridium difficile and non-enveloped viruses

Summary

Research question

- **Population:** Healthcare Workers
- **Intervention:** Hand hygiene comparing different concentrations of alcohol, and of different alcohols e.g. ethyl, methyl, isopropyl, in reducing the risk of transmission of Clostridium difficile and non-enveloped viruses
- **Comparator:** Washing with water and soap/detergent/chlorhexidine, other concentrations of same alcohol, other alcohols on Clostridium difficile and non-enveloped viruses
- **Outcome:** Decontamination of hands

3.1.2. Use and management of sharps, safety engineered devices and medication vials

Clinical question/ PICO

Population: Healthcare Workers
Intervention: Safety Devices
Comparator: Non-retractable devices

Summary

Research question

- **Population:** Healthcare Workers
- **Intervention:** Safety Devices
- **Comparator:** Non-retractable devices
- **Outcome:** Sharps injuries

Clinical question/ PICO

Population: Healthcare workers

Intervention: Immediate disposal of sharps

Comparator: Non-immediate disposal of sharps

Summary

Research question

- **Population:** Healthcare Workers
- **Intervention:** Immediate disposal of sharps
- **Comparator:** Non-immediate disposal of sharps
- **Outcome:** Sharps injuries

3.1.3. Routine management of the physical environment

Clinical question/ PICO

Population: Bacteria, non-enveloped and enveloped viruses

Intervention: Cleaning agent

Comparator: Frequency of agent use considering survival rates of the organisms

Summary

Research question

- **Population:** Bacteria, non-enveloped and enveloped viruses
- **Intervention:** Cleaning agent
- **Comparator:** Frequency of agent use considering survival rates of the organisms
- **Outcome:** Reduced levels of surface agents

Clinical question/ PICO

Population: Bacteria, non-enveloped and enveloped viruses

Intervention: Environmental cleaning agent

Comparator: Alternative environmental cleaning agents and mode of transmission of organisms

Summary

Research question

- **Population:** Bacteria, non-enveloped and enveloped viruses
 - **Intervention:** Environmental leaning agent
 - **Comparator:** Alternative environmental cleaning agents and mode of transmission of organisms
 - **Outcome:** Reduced levels of surface agents
-

Clinical question/ PICO

Population: Healthcare workers

Intervention: Surface barriers

Comparator: No surface barriers

Summary

Research question

- **Population:** Healthcare workers
 - **Intervention:** Surface barriers
 - **Comparator:** No surface barriers
 - **Outcome:** Reduced levels of surface agents
-

Clinical question/ PICO

Population: Bacteria, non-enveloped and enveloped viruses

Intervention: Site decontamination after spills of blood or other potentially infectious materials

Comparator: No site decontamination

Summary

Research question

- **Population:** Bacteria, non-enveloped and enveloped viruses
 - **Intervention:** Site decontamination after spills of blood or other potentially infectious materials
 - **Comparator:** No site decontamination
 - **Outcome:** Reduced levels of surface agents
-

Clinical question/ PICO

Population: Patients

Intervention: Disinfection/bleach, hand washing/soap/water, personal protective equipment etc

Comparator: Other, alcohol based

Summary

Research question

- **Population:** Patients
- **Intervention:** Disinfection/bleach, hand washing/soap/water, PPE etc.
- **Comparator:** Other, alcohol based
- **Outcome:** Severity of infection, number of people infected, duration of outbreak

3.1.3.1. Emerging disinfection methods

Clinical question/ PICO

Population: Patients

Intervention: Sodium hypochlorite

Comparator: Standard cleaning practice

| Outcome Timeframe | Study results and measurements | Comparator Standard cleaning practice | Intervention Sodium hypochlorite | Certainty of the evidence (Quality of evidence) | Summary |
|--|---|---|-------------------------------------|--|---|
| <p>Incidence rate of healthcare associated infections</p> <p>9 Critical</p> | <p>Based on data from participants in 3 studies. (Observational (non-randomized))</p> | | | <p>Very low Due to serious risk of bias, due to serious indirectness, due to serious imprecision ¹</p> | <p>Three studies of very low methodological quality investigated the effects of sodium hypochlorite disinfection on the incidence rate of hospital-acquired infection, compared to standard cleaning/ disinfection. Whilst one study showed that sodium hypochlorite disinfection led to a clinically important reduction in the rate of hospital-acquired CDAD, it had a serious risk of bias due to concerns that the intervention was not independent of other changes. Another study also demonstrated a clinically important reduction in the rate of hospital-acquired CDAD, however it had a very serious risk of bias due to concerns that the intervention was not independent of other changes. In addition, the pre-intervention period and the first intervention period had few data points, so the observed effects may be biased</p> |

| Outcome Timeframe | Study results and measurements | Comparator Standard cleaning practice | Intervention Sodium hypochlorite | Certainty of the evidence (Quality of evidence) | Summary |
|----------------------|-----------------------------------|---|-------------------------------------|---|---|
| | | | | | <p>due to regression to the mean or overfitting. The third study demonstrated an immediate reduction in the rate of hospital-acquired CDAD, however this effect was not statistically significant.</p> <p>There was serious imprecision due to wide confidence intervals, and a very serious risk of bias due to concerns that the intervention was not independent of other changes, and due to industry ties.</p> |

1. **Risk of Bias: serious.** Inadequate sequence generation/ generation of comparable groups, resulting in potential for selection bias, Incomplete data . **Indirectness: serious.** Due to outcome being hospital acquired infection, arising from any pathogen, so the outcome is not specific to MROs. **Imprecision: serious.** Wide confidence intervals, Only data from one study. **Publication bias: no serious.**

Clinical question/ PICO

Population: Patients

Intervention: Hydrogen peroxide vapour disinfection and standard cleaning/disinfection

Comparator: Standard cleaning practice

| Outcome Timeframe | Study results and measurements | Comparator Standard cleaning practice | Intervention Hydrogen peroxide vapour disinfection and standard cleaning | Certainty of the evidence (Quality of evidence) | Summary |
|--|--|--|--|---|---|
| <p>Incidence rate of healthcare associated infections (CDAD)</p> <p>9 Critical</p> | <p>(CI 95% -0.75 — 0.41) Based on data from participants in 1 studies. (Observational (non- randomized))</p> | <p>0.92 per 1000</p> <p>Difference:</p> | <p>0.74 per 1000</p> <p>0.52 fewer per 1000 CI 95%</p> | <p>Very low Due to very serious risk of bias and serious imprecision. Single study, so consistency cannot be assessed.¹</p> | <p>The effect of HPV disinfection on the rate of hospital acquired CDAD are uncertain due to very low quality evidence. HPV disinfection led to a small immediate reduction in the rate of hospital-acquired CDAD of -0.17/ 1000 patient days (equivalently, a decrease of 19%) but the confidence interval included no reduction, and an increase in CDAD rate as plausible estimates (95%CI -0.75, 0.41). During the intervention period, the</p> |

| Outcome Timeframe | Study results and measurements | Comparator Standard cleaning practice | Intervention Hydrogen peroxide vapour disinfection and standard cleaning | Certainty of the evidence (Quality of evidence) | Summary |
|----------------------|-----------------------------------|---|--|---|---------------------------------------|
| | | | | | trend in the rate of CDAD was stable. |

1. **Risk of Bias: very serious.** Due to concerns that the intervention was not independent of other changes, possible changes to screening during the outbreak period (concurrent with the intervention), and industry ties. In addition, the pre-intervention period had few data points, so the observed effects may be biased due to regression to the mean or overfitting. . **Indirectness: no serious. Imprecision: serious.** Wide confidence intervals.

Clinical question/ PICO

Population: Patients

Intervention: Ultra-violet light disinfection and standard cleaning/disinfection

Comparator: Standard terminal room disinfection (sodium hypochlorite)

| Outcome Timeframe | Study results and measurements | Comparator Standard terminal room disinfection (sodium hypochlorite) | Intervention Ultra-violet light disinfection and standard cleaning | Certainty of the evidence (Quality of evidence) | Summary |
|--|--|--|---|---|---|
| Incidence rate of healthcare associated infections (CDAD) 9 Critical | Relative risk 1 (CI 95% 0.57 — 1.75) Based on data from 5,177 participants in 1 studies. (Randomized controlled) | 36 per 1000 | 38 per 1000 | Low Due to very serious imprecision. Single study, so consistency cannot be assessed. ¹ | Terminal room disinfection with UV light may have little or no effect on the incidence of hospital-acquired CDAD compared to using sodium hypochlorite alone. However, further research is likely to have an important impact on our confidence in the effect estimate and may change the estimate. Addition of UV light for terminal room disinfection did not reduce the rate of hospital-acquired CDAD compared to sodium hypochlorite disinfection alone (RR 1.0, equivalently a decrease of 0% (95%CI: 0.57, 1.75; p=0.997)). |

1. **Indirectness: no serious. Imprecision: very serious.** Wide confidence intervals includes the possibility of a large reduction or a small increase, leading to conflicting interpretation of effects.. **Publication bias: no serious.**

Clinical question/ PICO

Population: Patients
Intervention: Ultra-violet light disinfection and sodium hypochlorite
Comparator: Standard cleaning practices

| Outcome Timeframe | Study results and measurements | Comparator Standard cleaning practices | Intervention Ultra-violet light disinfection and sodium hypochlorite | Certainty of the evidence (Quality of evidence) | Summary |
|---|---|--|---|--|---|
| <p>Incidence rate of infection or colonisation (Multi-Resistant Organisms - MROs)</p> <p>6 Important</p> | <p>Relative risk 0.82 (CI 95% 0.67 — 1) Based on data from 8,403 participants in 1 studies. (Randomized controlled)</p> | <p>5.64 per 1000</p> | <p>4.62 per 1000</p> | <p>Moderate Due to serious imprecision. Single study, so consistency cannot be assessed.¹</p> | <p>The effect of terminal room disinfection with UV light plus sodium hypochlorite on the incidence of hospital-acquired MROs is uncertain. Addition of UV light plus sodium hypochlorite for terminal room disinfection reduced the rate of hospital-acquired MROs by 18% compared to quaternary ammonium disinfection alone (RR 0.82, 95% CI 0.67, 1.00, p=0.048). However, the confidence interval includes the possibility of a clinically important reduction of 33% reduction, or no reduction.</p> |

1. **Indirectness: no serious. Imprecision: serious.** The 95% confidence interval includes the possibility of a clinically important reduction, or no reduction. . **Publication bias: no serious.**

Clinical question/ PICO

Population: Patients
Intervention: Environmental surfaces coated or impregnated with antimicrobial (self-disinfecting) materials
Comparator: Standard-surfaced objects

| Outcome Timeframe | Study results and measurements | Comparator Standard-surfaced objects | Intervention | Certainty of the evidence (Quality of evidence) | Summary |
|---|---|--|---------------------------------|---|---|
| <p>Incidence rate of healthcare associated infections (any pathogen)</p> | <p>Relative risk 0.82 (CI 95% 0.49 — 1.37) Based on data from 515 participants in 1 studies. (Observational (non-randomized))</p> | <p>13 per 1000</p> | <p>10.6 per 1000</p> | <p>Very low Due to serious risk of bias, serious indirectness and serious imprecision¹</p> | <p>The effect of copper-surfaced objects on the rate of hospital acquired infection is uncertain due to very low quality evidence. A small (18%) reduction in the rate of</p> |

| Outcome Timeframe | Study results and measurements | Comparator Standard-surfaced objects | Intervention | Certainty of the evidence (Quality of evidence) | Summary |
|----------------------|-----------------------------------|--|--------------|---|---|
| 9 Critical | | | | | hospital-acquired infection was found in rooms with copper-surfaced objects compared to rooms with standard objects (IRR 0.82 (95% CI: 0.49, 1.37), p=0.41). However, this does not exclude the possibility that the true intervention effect could be a 37% increase in colonisation with copper surfaces. |

1. **Risk of Bias: serious.** Inadequate sequence generation/ generation of comparable groups, resulting in potential for selection bias, Incomplete data . **Indirectness: serious.** Differences between the outcomes of interest and those reported (infection from any pathogen, not specific to MROs). **Imprecision: serious.** Wide confidence intervals. **Publication bias: no serious.**

3.1.4. Reprocessing of reusable medical devices

3.1.5. Respiratory hygiene and cough etiquette

3.1.6. Aseptic technique

Clinical question/ PICO

Population: Healthcare workers

Intervention: Sterile gloves

Comparator: Non-sterile gloves

Summary

Research question

- **Population:** Healthcare workers
- **Intervention:** Sterile gloves
- **Comparator:** Non-sterile gloves

- **Outcome:** Incidence rate of healthcare associated infections

3.1.7. Waste management

3.1.8. Handling of linen

3.2. Transmission-based precautions

3.2.1. Application of transmission-based precautions

3.2.2. Contact precautions

Clinical question/ PICO

Population: Patients

Intervention: Contact Precautions

Comparator: Non-contact Precautions

Summary

Research question

- **Population:** Patients
- **Intervention:** Contact Precautions
- **Comparator:** Non-Contact Precautions
- **Outcome:** Incidence rate of healthcare associated infections

Clinical question/ PICO

Population: Patients

Intervention: Hand hygiene and personal protective equipment

Comparator: No hand hygiene and no personal protective equipment

Summary

Research question

- **Population:** Patients
- **Intervention:** Hand hygiene and personal protective equipment
- **Comparator:** No hand hygiene and no personal protective equipment
- **Outcome:** Incidence rate of healthcare associated infections

Clinical question/ PICO

Population: Patients

Intervention: Patient-dedicated/single-use patient-care equipment

Comparator: Reusable equipment

Summary

Research question

- **Population:** Patients
- **Intervention:** Patient-dedicated/single-use patient-care equipment
- **Comparator:** Reusable equipment
- **Outcome:** Incidence rate of healthcare associated infections

3.2.3. Droplet precautions

Clinical question/ PICO

Population: Patients

Intervention: Droplet precautions

Comparator: Non-droplet precautions

Summary

Research question

- **Population:** Patients
- **Intervention:** Droplet precautions
- **Comparator:** Non-droplet precautions
- **Outcome:** Incidence rate of healthcare associated infections

Clinical question/ PICO

Population: Healthcare workers

Intervention: Masks and protective eyewear

Comparator: No masks and protective eyewear

Summary

Research question

- **Population:** Healthcare workers
- **Intervention:** Masks and protective eyewear
- **Comparator:** No masks and protective eyewear

- **Outcome:** Incidence rate of healthcare associated infections

3.2.4. Airborne precautions

Clinical question/ PICO

Population: Patients

Intervention: Airborne precautions

Comparator: Non-airborne precautions

Summary

Research question

- **Population:** Patients
- **Intervention:** Airborne precautions
- **Comparator:** Non-airborne precautions
- **Outcome:** Incidence rate of healthcare associated infections

Clinical question/ PICO

Population: Healthcare workers

Intervention: Masks and protective eyewear

Comparator: No masks and protective eyewear

Summary

Research question

- **Population:** Healthcare workers
- **Intervention:** Masks and protective eyewear
- **Comparator:** No masks and protective eyewear
- **Outcome:** Incidence rate of healthcare associated infections

3.3. Personal protective equipment

Clinical question/ PICO

Population: Healthcare workers

Intervention: Aprons and Gowns

Comparator: No aprons and gowns

Summary

Research question

- **Population:** Healthcare workers
 - **Intervention:** Aprons and Gowns
 - **Comparator:** No aprons and gowns
 - **Outcome:** Incidence rate of healthcare associated infections
-

Clinical question/ PICO

Population: Healthcare workers

Intervention: Masks and protective eyewear

Comparator: No masks and protective eyewear

Summary

Research question

- **Population:** Healthcare workers
 - **Intervention:** Masks and protective eyewear
 - **Comparator:** No masks and protective eyewear
 - **Outcome:** Incidence rate of healthcare associated infections
-

Clinical question/ PICO

Population: Patients

Intervention: Personal protective equipment

Comparator: No personal protective equipment

Summary

Research question

- **Population:** Patients
- **Intervention:** Personal protective equipment
- **Comparator:** No personal protective equipment
- **Outcome:** Incidence rate of healthcare associated infections

3.3.1. Other items of clothing

3.4. Management of multi-resistant organisms and outbreak situations

3.4.1. Multi-resistant organisms

Clinical question/ PICO

Population: Patients colonised or infected with a Multi-Resistant Organism (MRO)

Intervention: Transmission-based precautions

Comparator: Non--transmission-based precautions

Summary

Research question

- **Population:** Patients colonised or infected with an MRO
- **Intervention:** Transmission-based precautions
- **Comparator:** Non-transmission-based precautions
- **Outcome:** Incidence rate of healthcare associated infections (MROs)

3.4.2. Outbreak investigation and management

Clinical question/ PICO

Population: Patients colonised or infected with a Multi-Resistant Organism (MRO)

Intervention: Transmission-based precautions

Comparator: Non--transmission-based precautions

Summary

Research question

- **Population:** Patients colonised or infected with an MRO
- **Intervention:** Transmission-based precautions
- **Comparator:** Non-transmission-based precautions
- **Outcome:** Incidence rate of healthcare associated infections (MROs)

3.4.2.1. Infection control strategies to contain an outbreak

Clinical question/ PICO

Population: Patients

Intervention: Bay closures

Comparator: Ward closures

Summary

Research question

- **Population:** Patients
- **Intervention:** Bay closures
- **Comparator:** Ward closures

- **Outcome:** Incidence rate of healthcare associated infections

3.5. Applying standard and transmission-based precautions during procedures

3.5.1. Taking a risk-management approach to procedures

3.5.2. Invasive medical devices

Clinical question/ PICO

Population: Patients

Intervention: Systems and processes for device management

Comparator: No systems and processes for device management

Summary

Research question

- **Population:** Patients
- **Intervention:** Systems and processes for device management
- **Comparator:** No systems and processes for device management

- **Outcome:** Post-procedural infection

3.5.2.1. Indwelling urinary devices

3.5.2.2. Intravascular access devices

3.5.2.3. Ventilation

3.5.2.4. Enteral feeding tubes

3.5.3. Surgical procedures

3.5.3.1. Preventing surgical site infections (SSIs)

3.5.4. Aerosol-generating procedures (AGPs)

4. Organisational support

4.1. Management and clinical governance

4.1.1. Clinical governance in infection prevention and control

Clinical question/ PICO

Population: Healthcare workers

Intervention: Policies and procedures for clinical handover

Comparator: No formalised process for clinical handover

Summary

Research question

- **Population:** Healthcare workers
- **Intervention:** Policies and procedures for clinical handover
- **Comparator:** No formalised process for clinical handover

- **Outcome:** Efficient communication of high quality clinical information and improved patient outcomes

4.1.2. Roles and responsibilities

4.1.3. Infection prevention and control program

4.1.4. Risk management

Clinical question/ PICO

Population: Patients

Intervention: Chlorhexidine use

Comparator: No Chlorhexidine use

Summary

Research question

- **Population:** Patients
- **Intervention:** Chlorhexidine use
- **Comparator:** No Chlorhexidine use
- **Outcome:** Resistance to chlorhexidine and/or resistance against antibiotics

Clinical question/ PICO

Population: Patients

Intervention: Risk register for chlorhexidine

Comparator: No risk register for chlorhexidine

Summary

Research question

- **Population:** Patients
- **Intervention:** Risk register for chlorhexidine
- **Comparator:** No risk register for chlorhexidine

- **Outcome:** Documented use of chlorhexidine and incidence of anaphylaxis

4.1.5. Taking an organisational systems approach to infection prevention quality and safety

4.2. Staff health and safety

4.2.1. Health status screening and immunisation

Clinical question/ PICO

- Population:** Healthcare workers
Intervention: Exclusion period
Comparator: Different periods of time

Summary

Research question

- **Population:** Healthcare workers
- **Intervention:** Exclusion period
- **Comparator:** Different periods of time
- **Outcome:** Rates of transmission of infection to healthcare worker or patient

Clinical question/ PICO

- Population:** Healthcare workers
Intervention: Education programs
Comparator: Other education programs

Summary

Research question

- **Population:** Healthcare workers
- **Intervention:** Education programs
- **Comparator:** Other education programs
- **Outcome:** Changes in healthcare worker behaviour

4.2.2. Exclusion periods for healthcare workers with acute infections

Clinical question/ PICO

Population: Healthcare workers

Intervention: Exclusion period

Comparator: Different periods of time

Summary

Research question

- **Population:** Healthcare workers
- **Intervention:** Exclusion period
- **Comparator:** Different periods of time
- **Outcome:** Rates of transmission of infection to healthcare worker or patient

4.2.3. Managing exposures to occupational hazards

4.2.4. Healthcare workers with specific circumstances

4.2.5. Exposure-prone procedures

4.3. Education and training

Clinical question/ PICO

Population: Healthcare workers

Intervention: Education programs

Comparator: Other education programs

Summary

Research question

- **Population:** Healthcare workers
- **Intervention:** Education programs
- **Comparator:** Other education programs

- **Outcome:** Changes in healthcare worker behaviour

4.3.1. Education strategies

4.4. Healthcare associated infection surveillance

4.4.1. Role of surveillance in reducing healthcare associated infection

4.4.2. Types of surveillance programs

4.4.3. Data collection and management

4.4.4. Outbreak surveillance

4.4.5. Disease surveillance in office-based practice

4.4.6. Notifiable diseases

4.5. Antimicrobial Stewardship

4.5.1. Antimicrobial resistance in Australia

4.5.2. Antimicrobial stewardship programs

4.5.3. Antimicrobial stewardship surveillance methods

4.6. Influence of facility design on healthcare associated infection

4.6.1. Mechanisms for influencing healthcare associated infection through environmental design

5. APPENDIX 1: Case studies and examples

5.1. Risk-management: Case study for hand hygiene in a neonatal intensive care unit

5.2. Risk-management: Case study for glove use, hand hygiene and seasonal influenza vaccination in an office-based practice

5.3. Example: Education in practice - hand hygiene

5.4. Risk-management: Case study for the prevention of needlestick injury during surgery at a tertiary referral hospital

5.5. Risk-management: Case study for spills management in a busy paediatric ward

5.6. Risk-management: Case study for reprocessing of instruments in a dental practice

5.7. Risk-management: Case study for Klebsiella pneumoniae sepsis in a neonatal unit

5.8. Risk-management: Case study for influenza in a long-term care facility

5.9. Risk-management: Case study for M. tuberculosis

5.10. Risk-management: Case study for M. tuberculosis among immunocompromised patients attending outpatient services

5.11. Examples of how to perform aseptic technique

5.12. Risk-management: Case study for norovirus outbreak in a long-term care facility

5.13. Risk-management: Case study for management of confirmed case of carbapenemase-producing Enterobacterales (CPE)

5.14. Risk Management: Case study for vancomycin-resistant enterococci (VRE) outbreak in a large tertiary-care referral hospital

5.15. Risk-management: Case study for infection prevention during renovation of emergency department

6. APPENDIX 2: Supplementary information

6.1. Recommended routine cleaning frequencies

6.2. Checklist of standard precautions for procedures

6.3. Use of standard and transmission-based precautions

6.4. Type and duration of precautions for specific infections and conditions

6.5. Collection of pathology specimens

6.6. Allowing animals into healthcare facilities

6.7. Mechanisms of antibacterial action of antiseptics and disinfectants

6.8. Summary table of relevant infection prevention and control resources

6.9. Research gaps

7. APPENDIX 3: Process report

7.1. Membership and Terms of Reference of the Advisory Committee

8. Glossary

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11. Amendments to the Guidelines