A systematic review of the literature addressing asepsis in wound management

Haesler E, Thomas L, Morey P & Barker J

ABSTRACT

There has been extensive ongoing debate on the application of aseptic technique in wound management over the previous decades and changes to the way in which theory is applied to clinical practice have occurred regularly. Clinicians often express confusion over the way various techniques should be applied, particularly when practising in clinical settings in which maintenance of strict asepsis is inherently difficult (for example, community-based wound management). Wound cleansing, use of open but unused

Emily Haesler*

PhD, BN, PostGradDipAdvNurs(Gerontics) Adjunct Associate Professor Curtin University, School of Nursing, Midwifery and Paramedicine, WA, Australia Honorary Associate La Trobe University, Australian Centre for Evidence Based Aged Care, Vic, Australia Email: Emily.Haesler@curtin.edu.au Tel +61 2 6244 2946

Lyn Thomas

RN, NP, BHlthSci(Nurs), MNP Community and Aged Care Services, Greater Newcastle Sector, Hunter New England Local Health District, NSW, Australia

Pam Morey

MN, NP, STN, PhD(C) Nurse Practitioner, Advanced Wound Assessment Service, Silver Chain Nurse Practitioner Course Coordinator Curtin University, WA, Australia

Judith Barker

RN, NP, BHIthSc(Nurs), MNP Nurse Practitioner — Wound Management Adjunct Associate Professor, University of Canberra, Synergy: Research Centre for Nursing and Midwifery Practice, Canberra Canberra Hospital, ACT Health, ACT, Australia

* Corresponding author

wound dressings and storage of wound management equipment are frequent issues on which clinicians request guidance. A systematic review using Joanna Briggs Institute methods was undertaken in order to establish the current state of the scientific literature on this topic and inform the development of recommendations for practice in this field. All levels of evidence were included in the review, including opinion papers. Findings from the 20 quantitative studies were reported in narrative summary and findings from 37 qualitative research papers were aggregated in a thematic synthesis. Although high-level evidence on wound cleansing solutions was identified, the review concluded that there is a paucity of scientific literature on most topics related to asepsis in wound care.

Keywords: Asepsis, wound cleansing, aseptic non-touch technique, handwashing, infection control.

INTRODUCTION

Wound infection has a large impact on individuals and the health care system. Precise incidence rates are difficult to determine due to the many types of wounds and various methods of diagnosing and tracking wound infection. As many as 60% of chronic wounds have infection in the form of demonstrated presence of surface bacteria or invasive biofilm^{1,2}. Rates of surgical site infection vary substantially based on surgical site³; however, recent estimates suggest 10–12% of all surgical wounds become clinically infected⁴. Infection rate in lacerations is cited at 5%⁵, and rate of biofilm in all acute wounds is approximately 6%².

Facility-acquired wound infection is of particular concern given the increasing significance of antibiotic-resistant bacteria. Infection control procedures are first-line strategy to prevent infection spread⁶. Given the impact of wound infection and significance of infection control practices in reducing its incidence, it is important that clinicians understand the implementation of infection control procedures when managing wounds. Historically, there have been major changes to aseptic theory in wound management⁷⁻⁹. Surveys indicate clinicians experience confusion about how to implement aseptic technique and other infection control principles^{10,11}. Within Australia, the introduction of a standard on health care-associated infection¹² and publication of a national infection control policy¹³ led to a demand for updated wound management procedures. Wounds Australia established a working party to develop clinical guidance on procedures associated with prevention and control of wound infection. To inform the development of this document, a systematic review (SR) was undertaken.

AIMS

The objective of this review was to identify the contemporary evidence addressing topics associated with aseptic technique and infection control in wound management. Specific aims were to identify evidence related to cleaning considerations when performing a wound procedure, techniques for wound cleansing, environmental considerations in performing wound management and ways in which wound dressings can be handled and stored aseptically.

REVIEW METHODS

The review was undertaken using methods published by the Joanna Briggs Institute (JBI)14,15. An initial search was conducted in MEDLINE, CINAHL, EMBASE, Current Contents and the Cochrane library. All papers published in English up to October 2015 that related to topics outlined in the aims were eligible for inclusion. All research designs, gualitative research and opinion papers were eligible for inclusion; however, news items, letters and conference abstracts were excluded. Papers related to aseptic technique in the operating room, intravenous therapy or catheterisation were excluded. Search terms and MESH headings included: asepsis, non-touch technique, aseptic technique, steriliz/sation, disinfection, microbial and bacterial contamination, hospital, healthcare and community-acquired infection. These terms were used in combination with terms associated with wound care, wound dressings, equipment storage, cleansing, and equipment recycling. The working party reviewed the search strategy to ensure it captured the intended literature. On review of the evidence it was noted by the working party that significant changes in theory and practice have occurred in the field of aseptic technique. It was determined that inclusion would be limited to papers published between January 2000 to October 2015 in order that the review findings reflect contemporary knowledge. References cited in included manuscripts were also considered for inclusion.

All papers meeting inclusion criteria were critically appraised by two independent reviewers using the JBI suite of appraisal tools. For randomised controlled trials (RCTs) and pseudo-RCTs, critical appraisal evaluated randomisation, blinding, allocation concealment, withdrawals, comparability and equivalent treatment of participants, outcome measurement and statistical analysis¹⁴. Consistent with JBI appraisal, RCTs and pseudo-RCTs were ranked as high quality or lower¹⁵. For descriptive studies and case series, process for randomisation, sample inclusion, outcome measurement, management of confounders, participant withdrawal and data analysis were evaluated¹⁴. These studies received a ranking of low or very low quality¹⁵. For interpretive and critical research, congruity of philosophies, methodology, research methods and analysis was evaluated, as well as reflexivity¹⁴. Qualitative research was ranked as high quality or lower¹⁵. Textual and opinion papers were evaluated based on source and logic of opinion and arguments, focus, referencing and support from peers¹⁴ and ranked as low or very low quality¹⁵.

Data extraction used standardised JBI tools. Quantitative results were not appropriate for meta-analysis as they generally addressed different topics, had heterogeneous methods, or were meta-analyses. These results are reported in a narrative format. Qualitative studies and opinion papers were analysed to identify themes, concepts and meanings within the research¹⁴, with identification of primary findings that were grouped in categories based on similarity in meaning. The categories were meta-aggregated in syntheses.

IDENTIFIED RESEARCH

The Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) flow diagram¹⁶ is presented in Figure 1. The searches initially identified over 2,000 potential studies that was reduced to 57 papers that met inclusion criteria and were critically appraised. As indicated in Figure 1, most studies were excluded in the first review of the flagged references due to having insufficient focus on the topic of this review.

Of the 57 included papers, 14 were quantitative research papers¹⁷⁻³⁰, six were SRs³¹⁻³⁶, three were qualitative research papers^{10,11,37} and 34 papers were non-research articles^{7-9,38-68}. The quantitative research consisted of six RCTs^{18,21,23,25,26,29} (level 1.c evidence) that were of low or moderate quality¹⁵. There was one very low quality before/after study²² (level 2.d evidence), five observational studies^{17,19,20,28,30} (level 2 and 3 evidence), one very low quality cohort study²⁴ (level 3.c evidence) and a very low quality cross-sectional study²⁷ (level 4.b evidence). The SRs³¹⁻³⁶ (level 1 evidence) ranged in quality from low to very high and the qualitative research^{10,11,37} (level 3 evidence) was of moderate to high quality. The majority of findings in this review arose from textual papers providing low and very low quality evidence. Table 1 presents summaries of the research papers.

Quantitative results from the literature *Cleansing solutions and technique*

SRs and studies exploring irrigation fluids received the most attention in quantitative research. Details of the included studies, including their quality and level of evidence, are provided in Table 1 in the supplementary material accessible at www.woundsaustralia.com.au/journal/2404.php. As the individual studies^{18,19,21,23,26,29} in Table 1 were included in identified SRs, only SR results are reported below; however, none of the individual RCTs established significant

Figure 1: PRISMA review flow



differences in infection rates between wounds cleansed with sterile solutions versus tap water^{18,19,21,23,26,29}.

A high quality SR³⁶ compared sterile saline (n=326) to tap water (n=257) for cleansing lacerations, acute and chronic wounds. Pooled results from two RCTs showed no significant difference in wound infection rates, with tap water slightly less likely to result in infection (odds ratio [OR] 0.79, 95% confidence interval [CI] 0.36 to 1.72, p=0.55)³⁶.

A moderate quality Cochrane SR³³ compared cleansing methods for lacerations, open fractures, chronic and surgical wounds. Infection rate in all wound types (3 RCTs) was not significantly different between tap water cleansing and no cleansing (relative risk [RR] 1.06, 95% CI 0.07 to 16.50, p=not significant [ns]). There was no difference in infection rates in all acute sutured wounds (3 RCTs) between tap water versus sterile saline irrigation (RR 0.66, 95% CI 0.42 to 1.04, p=ns). Cost-effective analyses favoured tap water³³. This Cochrane review reached the same conclusions as earlier systematic reviews by the same research team^{34,35}.

A low quality SR compared tap water to sterile saline. Significantly more wounds cleansed with sterile saline became clinically infected (saline 7.1% versus tap water 4.3%, RR 0.62, 95% Cl 0.39 to 1.01, p=0.05). There was no significant difference in wounds with positive cultures (saline 3.1% versus tap water 4.4%, RR 1.53, 95% Cl 0.79 to 2.99, $p=0.21)^{31}$.

A low quality SR compared bathing to no bathing for postsurgical foot wounds. Normal hygiene groups showered at 1–5 days postoperative (n=1,639). Patients abstaining from foot hygiene waited until sutures/staples removal (n=511). There were no significant differences in surgical site infection rates in any study³².

Although confounding factors are noted (for example, administration of saline at cooler temperatures than water) when controlled for these factors the outcomes did not change. Findings from the high-level evidence^{31,33-36} indicated no increase in wound infection rates associated with cleansing wounds in tap water.

Reuse of wound dressing products

A moderate quality observational study investigated rate of contamination of opened hydrogel products. The products were opened after handwashing, using clean gloves and away from direct patient care. After 28 days, one package from 60 random samples returned a positive bacterial culture. The sample collection technique may not reflect clinical practice¹⁷. A very low quality observational study reported contamination rates for opened dressings and reusable equipment stored in different containers in patients' homes. After 14 days, 75% of samples (n=21) were contaminated³⁰. Another low quality observational study investigated contamination rates for randomly selected multi-use saline flasks stored in hospital settings. Approximately half of the samples were found to be contaminated²⁰.

Wound dressing practice

In a low quality RCT, leaving surgical wounds uncovered after surgery (n=235) was compared to wound dressings applied in the operating theatre (n=216). Patients were reviewed after seven days for clinical signs of infection and no significant difference in infection rates was found (exposed wounds 1.7% versus covered wounds 1.4%, p=ns)²⁵.

A very low quality cohort study compared sterile (n=1,070 admissions) and clean (n=963 admissions) dressing procedures for surgical wounds. The outcome measure was positive wound culture established by wound swab. There was no significant difference in surgical site infection rates (0.84% versus 0.83%, p=ns) and the clean procedure was faster (10 minutes versus 13 minutes)²⁴.

Aseptic technique education and behaviours

A very low quality before/after study investigated an education program delivered to medical students. The course was based on principles associated with handwashing and dressing procedures. After 10 weeks, there was a significant decline (p<0.001) in the ratio of students who were able to achieve a pass mark in the assessment, indicating the education had no prolonged influence on practice. Poor role modelling and lack of resources were identified as contributing to poor outcomes²². A very low quality observational study reported clinical practice amongst nurses in community settings. Practice was established through direct observation and validated in interviews with participants. As many as 40% of nurses did not engage in handwashing before a procedure²⁸.

Qualitative results from the literature

Three hundred and eighty-six findings were extracted from qualitative studies and non-research articles. Using the JBI ratings¹⁵, 60 of the findings were rated as unequivocal, 218 were rated as credible and 20 findings were rated as unsupported, generally where an assertion was made without any supporting reference. These findings were grouped in 65 categories and aggregated into 23 syntheses that are reported in full in Figure 2 in the supplementary material accessible at www.woundsaustralia.com.au/journal/2404.php.

Current evidence base

Synthesis 1: Research on wound cleansing and aseptic technique is insufficient and that which is available is poorly translated into practice.

There is a lack of research on aseptic techniques^{9,65}. Inconsistencies in terminology and practice guidance, and ongoing change to theory interpretation has a negative impact on compliance⁷⁻⁹. The need for more research on aseptic technique, including translation to different clinical settings was highlighted^{38,44,48}.

Handwashing practices

Six categories aggregated into two syntheses represented textual findings on handwashing.

Synthesis 2: Liquid alcohol rub, antimicrobial hand wash or soap and water can be used for washing hands. When hands are visibly soiled, use soap and water.

Articles referred to three handwashing solutions: alcoholbased rubs, antiseptic/antimicrobial hand washes and soaps/ detergents. Alcohol rub has broad spectrum activity⁵¹ and is quick to apply without the need for water^{8,40,45,50,51,55,57,59}. A small risk of fire from alcohol exposed to a heat source before complete evaporation⁴⁵ and potential for dry hands are reported. Some texts suggest alcohol-based hand rubs are not appropriate when hands are visibly soiled^{40,55}. Antiseptic or antimicrobial hand wash with water is suggested for cleaning visibly dirty hands^{40,45,51,55} although it may be more expensive or cause irritation⁵¹. Antimicrobial-impregnated towels are an alternative for visibly clean hands, but not a replacement for soap and water^{40,45,55}. Opinion articles agreed that soap and water is appropriate for visibly soiled hands^{40,49,51,55,64}.

Synthesis 3: Handwashing should occur before and after patient contact, regardless of the use of gloves, and consist of vigorous rubbing for at least 15 to 30 seconds.

Hands should be washed before/after patient contact, or after contact with body fluids, to prevent crosscontamination^{7,40,55,56,64}. Use of gloves does not preclude the need to wash hands^{40,55-57,59} because hands may become contaminated when removing gloves⁵⁶. Handwashing should be a vigorous process covering all hand surfaces using soap and water or an alcohol rub^{40,55,59}. Most papers suggested that handwashing should take at least 15 seconds^{40,55}; however, one suggested at least 30 seconds⁵⁹.

Gloves and personal protective equipment

Three syntheses related to the use and selection of gloves, and one related to other personal protective equipment.

Synthesis 4: Gloves are required to prevent contamination and cross-infection; however, they do not replace routine handwashing.

Textual findings highlighted that the primary purpose of gloves is to prevent contamination, between the patient and the nurse, or cross-infection between different anatomical sites on the same patient^{40,44,51,55,59}. It was suggested that gloves are worn when there is a risk of coming into contact with bodily fluids or non-intact skin^{40,55}; when removing old wound dressings⁴⁵; and for invasive activities⁵¹. Use of gloves does not preclude handwashing^{45,57}, regardless of the implementation of double-gloving⁴⁵. Findings suggested gloves be removed immediately following care^{40,55}.

Synthesis 5: Selection of gloves is guided by the procedure to be performed, risk of contamination, latex allergies and cost.

Synthesis 6: Sterile gloves are required for surgical aseptic non-touch technique, surgery and invasive aseptic procedures

and clean gloves are for non-sterile procedures/standard aseptic non-touch technique.

Level of expected direct contact with susceptible sites^{44,53,57} should guide glove selection. Latex allergy influences glove choice^{44,50,59}, and some texts identified the increased cost of sterile gloves as a factor in selection^{59,62}. There was agreement that sterile gloves are required for sterile procedures^{44,48,51,57,61}. Within the literature 'sterile procedures' referred to invasive activities^{44,51} surgical procedures⁴⁴, aseptic technique⁴⁴, aseptic non-touch technique (ANTT) requiring direct contact with key parts⁶¹, and delivering sterile pharmaceuticals⁴⁴. Clean, non-sterile gloves were suggested for removing old wound dressings⁴⁸, performing clean procedures⁴⁸ and performing procedures that do not require direct contact with the key parts⁶¹.

Synthesis 7: Wearing appropriately selected personal protective equipment helps to reduce the risk of cross-infection from exposure to body fluids or airborne contamination.

Personal protective equipment is designed to reduce the risk of contamination for both the patient and clinician^{57,64}. When protective equipment is used, the clinician is protected from body fluid exposure (for example, blood splashes)^{51,57} and the patient is protected from the clinician as a source of infection risk⁵⁹. Textual findings focused on using plastic aprons^{51,59} with selection of equipment based on the level of risk of body fluid exposure⁶⁴.

Wound management environment

Two syntheses addressed the general and specific environment in which wound management is conducted.

Synthesis 8: Actions should be taken to reduce airborne and other infection risks in the home and hospital to ensure wound care is conducted in a clean environment.

The requirement for a clean environment, free from airborne and other infection risks was described^{51,61,64}. The risk posed by carpets, soft furnishings and pets was reported⁶⁴. Strategies to reduce environmental risk included reporting infection risk in the home to authorities⁶⁴, ensuring there are cleaning routines that incorporate the ventilation and water supplies⁶⁴; reducing airborne infection by closing windows, reducing foot traffic and turning off fans⁶⁴; leaving the wound exposed for the shortest time⁶⁴; and disposing of waste promptly and appropriately⁶⁴.

Synthesis 9: A wound management field can be established on a clean surface in a space at low risk of environmental contaminants. Once established, introduction of contaminated external objects should be avoided.

The importance of establishing a sterile local field on a clean surface^{8,57,64} was discussed. Strategies for establishing a sterile field in a clean environment included using a visually clean dressing trolley⁵⁷ or cleaning a hard surface with a broad spectrum disinfectant⁶⁴. In the community, a plastic apron or

lid could be used^{8,64}. Considering the wound, an extension of the wound management field was suggested⁷, as was ensuring the wound management field remains sterile⁵⁷. Findings were consistent that objects external to the wound and field should not contaminate the wound management field^{7,57,61}. Clinicians could use either a critical aseptic field into which only sterile equipment is introduced (for example, for an invasive or extensive procedure) or a general aseptic field in which key parts are individually protected within the field (for example, for a simple wound procedure)⁶¹.

Cleansing solutions and technique

Eighteen categories were aggregated into six syntheses related to wound cleansing.

Synthesis 10: An ideal wound cleanser should adequately clean the wound, not cause cell damage or sensitivity and have a long shelf life.

The principle of doing no harm and preventing infection were highlighted as guiding the choice of wound cleanser⁹. Consideration to the toxicity of a cleanser and its potential to cause sensitivity was highlighted^{42,63}. Using an expired product should be avoided⁵⁷ by selecting a wound cleanser with a long shelf life^{42,63}. The ability to effectively remove organic material and reduce bioburden are other considerations^{42,63}.

Synthesis 11: An assessment should be conducted by the interdisciplinary team to determine if a wound bed should be cleansed, and if so, the cleansing process to use.

Not all wound beds require cleansing as a wound may heal without disruption if there are no visual contaminants or signs of infection^{39,58}. The wound management team could work together to determine the best approach for individual patients^{9,39,58}.

Synthesis 12: Normal saline, potable tap water, sterile water and low concentration antimicrobial solutions are safe and effective wound cleansers. Antiseptics are not a good choice for wound cleansing.

Sterile saline is an isotonic solution that has no impact on tissue repair processes^{42,46}; therefore it is a safe and traditional option⁶², particularly in hospital environments⁴⁶ or for vulnerable wounds⁹. Tap water, sterile water and normal saline were all reported as safe; however, none of these solutions reduces bioburden in the wound^{42,46,56,57,67}. Antimicrobial solutions reduce bioburden^{63,67}, although concentrations should be selected carefully in light of potential cell toxicity^{63,67}. Use of skin cleaners and antiseptics in wound cleansing is warned against^{39,42,58,63}. Cell toxicity^{42,58,63}, potential carcinogenicity⁴¹, insufficient contact time with the wound to effectively reduce bacteria levels³⁹ and association with antibiotic-resistant bacteria³⁹ were concerns.

Synthesis 13: Apply a wound cleanser at a lukewarm temperature with consideration to the potential for cross-infection and using low pressure to irrigate the wound bed.

Irrigation at a low pressure (4 to 15 pounds per square inch) using a syringe or faucet tubing is suggested for promoting debris removal without disrupting granulating tissue^{42,56,62,65}. Applying fluid at lukewarm temperature avoids vasoconstriction that lowers tissue healing capacity^{9,45,46,54,62,66}. The potential for cross-infection between patients, or contaminated water from dirty body areas flowing over a wound are considerations when washing in a shower^{9,54,58}. Directing fluid flow appropriately when irrigating⁵⁴ was noted as another strategy to prevent cross-infection.

Synthesis 14: Good quality tap water is a cost-effective option for cleansing dirty wounds, chronic wounds and wounds with closed or sutured edges, although it may cause pain.

Benefits and disadvantages of tap water were discussed^{39,42,46,54,56,68,62,66}. Water was noted as acceptable for sutured³⁹, sacral/perineal⁵⁶, open traumatic⁵⁶, and chronic⁵⁶ wounds, and wounds with sealed edges³⁹. Ensuring high quality water is important^{54,56,62,66}, although commentators noted that in cities with monitored and drinkable tap water it is sufficiently safe for wounds^{54,56,62,66}. Texts suggested using running tap water for at least 30 seconds⁶² or soaking wounds in a bucket⁵⁸. Higher, constant pressure⁶², large fluid volumes⁶², patient satisfaction⁶⁶ and reduced time⁶⁶ are advantages of water. Lack of additional equipment (for example, syringes) contributes to the cost-effectiveness of water^{46,54,62,66}. However, there is potential that water may cause pain due to increasing osmotic pressure⁴⁶.

Synthesis 15: Precautions can be taken to reduce the risk of potential contamination of water sources.

Another disadvantage is the potential for contaminated tap water^{9,41}. One commentator suggested a risk of acquiring virulent pathogens or biofilm from hospital water⁴¹. This risk may be higher for immunocompromised patients⁴¹. However, precautions can be taken^{41,42,56}. Water filters⁴¹, running taps for a few minutes before using the water⁴² and evaluating the water storage and delivery before use⁹ were suggested.

Selecting wound care technique and equipment

Four syntheses addressed selection of wound care techniques and equipment.

Synthesis 16: Selection of sterile/surgical ANTT or clean/ standard ANTT is determined by the level of risk posed to the patient by his or her health status, the environment, factors associated with the wound and the type of wound management procedure being performed.

The infection risk from the surrounding environment is one consideration in selecting a wound management technique^{7,37,44,57,60,61,68}. Findings illustrated that both the health care setting^{7,37,44,68} and the storage of equipment⁶⁸ influences the ability to maintain a sterile or aseptic environment. The complexity of the procedure is a contributing factor, for example extensive debridement, wound packing and necessity to touch key parts were considered more invasive and requiring greater precautions^{44,51,57,60,61,68}. Patient-related factors (for example, immune status) may also contribute to the risk of infection from a dressing procedure^{7,44,57,60,68}. The chronicity, depth and location of the wound also contribute to selection of a technique^{7,44,50,56,57,68}. Rigorous asepsis was considered to be inappropriate for chronic wounds^{50,56,57,68}.

Synthesis 17: Simple wound management procedures on low-risk patients can be performed with non-sterile but clean equipment, solutions and gloves. More complex procedures or procedures in higher risk patients require surgical aseptic non-touch technique, using sterile gloves, solutions and equipment.

Textual findings referred to clean technique/standard ANTT and aseptic technique/surgical technique/surgical ANTT. The first technique is appropriate for routine dressing changes without surgical conservative debridement and simple procedures lasting less than 20 minutes^{37,61,68}. This technique was reported to involve a clean surface, non-sterile gloves and clean equipment and irrigation fluids (for example, tap water)^{37,61,68}. The surgical ANTT requires sterile gloves and equipment and a sterile irrigation fluid, with a strict aseptic field^{37,56,57,61,68}. The findings suggested this procedure was appropriate for patients at high infection risk, wounds requiring surgical conservative debridement, complex/ invasive procedures with many key parts or procedures lasting longer than 20 minutes^{56,57,61,68}. One commentator suggested that this should be standard practice⁵⁶.

Synthesis 18: Wound management equipment should be single use or cleaned with alcohol preparations. Using cleansers and wound dressings in smaller packages reduces waste and contamination risk.

Ensuring products are cleaned appropriately via sterilisation, disinfection or decontamination is important^{57,61,64}. Using alcohol preparations or wipes and vigorously rubbing equipment to remove visual soiling cleans reusable products^{61,64}, although single-use products may be easier, especially in community settings⁶⁴. Wastage of excess products was noted as a concern, especially from dressing packs with pre-selected materials that are not always appropriate for the procedure^{8,11}. Selecting smaller packages to reduce waste or risk of contamination from reusing products was suggested^{45,65}.

Synthesis 19: When performing surgical ANTT the wound management field must remain free of non-sterile items, including equipment, cleansing fluids and gloved hands that have touched a non-sterile object.

Commentary highlighted the importance of all sterile equipment being free from potentially contaminated objects, including water that had touched surrounding skin during washing or forceps that had touched the wound bed^{7,8,43,51,68}. One text referred to a dirty hand or forceps/a clean hand

or forceps⁷. The difficulty clinicians have in manoeuvring forceps was raised^{7,56}, and using a gloved hand for parts of a procedure was proposed as an optional wound management method^{7,8,56}, if the potentially contaminated hand could be maintained away from the wound management field^{7,68}.

Managing patients with known infection

Synthesis 20: Extra infection control precautions should be taken for people with known infection.

One opinion article addressed infection control for patients with known methicillin-resistant *Staphylococcus aureus* (MRSA)⁴⁹. Findings indicated that clinicians should take additional precautions by thoroughly disinfecting surfaces, putting down plastic sheeting and using disposable equipment when possible⁴⁹. Reusable equipment could be cleaned immediately⁴⁹. Double-bagging waste products before disposal may reduce cross-infection⁴⁹. Diligence is required in handwashing and the use of personal protective equipment⁴⁹.

Product storage

Synthesis 21: Wound management products should be stored in dry, clean environments to reduce risk of contamination and cleansers should be dated on opening and discarded if visually contaminated or according to organisation policy.

Storing dressing products in a clean, dry space was suggested^{56,64}. Wound cleansers should be dated and refrigerated on opening, although there is no set time frame after which they should be disposed⁴⁵. One commentator mentioned discarding fluids if there is visual contamination, or to follow the organisation policy⁴⁵. Risks posed by storing gloves in a manner that attracts mould or contamination were noted^{50,62}.

Structural support

Two syntheses summarised eight categories related to structural support for aseptic technique and infection control.

Synthesis 22: Staff education that incorporates skills practice, simulation learning, theoretical knowledge update and procedures for different clinical settings is essential in promoting best practice in aseptic technique and infection control.

Importance of education was highlighted^{8,11,38,39,44,49-51,57,59,64}. Some references suggested that clinicians develop ritualistic practice and may not fully understand theoretical concepts^{8,11,39}. Qualitative studies indicated that community nurses experience frustration and have fatalistic attitudes¹¹ that may influence the way in which they perform wound care¹⁰. Ongoing reinforcement of knowledge and skills through regular education using simulation learning^{53,57}, visual feedback (for example, dye in handwashing exercises)⁵⁹, hands-on practice with feedback⁶⁴ and didactic lectures⁵³ is suggested.

Synthesis 23: Best practice in aseptic technique and infection control procedures is promoted through development of facility policies, regular risk surveillance, annual auditing of staff practice, engaging with staff and patients and provision of acceptable hand hygiene products.

Regular risk surveillance^{7,50,59,64} promoting a culture of clinicians identifying risks⁵⁰, and root cause analysis⁶⁴ were highlighted as promoting quality improvement. Engaging with staff and patients by working with a wound champion to promote best practice^{38,64}, empowering patients to ask clinicians about hand hygiene⁵⁹ and ensuring adequate staffing levels may promote best practice⁵¹. Commentators proposed incorporating annual handwashing audits into quality improvement programs^{8,38,40,55,59}. The importance of local policies and procedures was raised^{8,48,61}, especially for topics for which there is insufficient evidence to make recommendations⁴⁸. Finally, provision of products that are acceptable to clinicians (for example, low allergen) may promote handwashing^{40,55}.

DISCUSSION

There was general agreement between the quantitative and qualitative/textual findings in this review. Use of potable tap water for irrigation received the most attention, and findings from SRs, RCTs and non-research articles were in agreement that for many wound types, cleansing with good-quality, lukewarm tap water does not increase risk of wound infection^{18,20,21,23-26,29,32-36,39,42,46,54,56,58,62,66}. This evidence should be considered when selecting appropriate and cost-effective wound management techniques.

Limited evidence was available on other topics of interest. No significant evidence was identified regarding strategies for managing opened wound dressing packages and minimal commentary on the advantages and risks of this practice was identified. One moderate quality study¹⁷ suggested reusing hydrogel products may be safe if the product was dispensed in controlled conditions. A very low quality study suggested contamination of opened wound dressings is an issue in community settings³⁰ and commentators suggested using smaller packages to reduce waste^{45,65}. There was also limited evidence on methods of performing aseptic technique, with technique details often derived from unreferenced opinion sources⁶¹. There is a strong need for well-designed studies exploring these issues.

There was a paucity of evidence on environmental factors in conducting wound care identified in the available literature. There was agreement that the environment in which wound care is conducted should be clean^{51,61,64}, with guidance generally focused on strategies to address the potential risk from airborne contamination^{51,61,64}. Practical solutions for maintaining asepsis in home care settings were provided^{8,51,61,64}; however, the evidence supporting these practices was at best minimal. This review is not without limitations. Foremost, the literature search was limited to journal articles. Reports¹² and guidelines^{13,69} also inform this topic; however, these resources are developed from the existing body of evidence and are not specific to wound management. The search terms used for this review focused specifically on asepsis in wound management. It is probable that evidence on some included topics is available in the broader literature. Except for evidence related to irrigation, the evidence was primarily from non-research texts of low and very low quality and this should be considered when evaluating the adoption of the suggestions into practice. It should be noted that theory and translation in this field has changed substantially over time. The reviewers attempted to identify a cut-off date in order to exclude outdated concepts; however, given that much of the findings were opinion, some ideas may be anachronous.

CONCLUSIONS

The findings of this systematic review highlighted the lack of high-level evidence in many clinical areas associated with aseptic wound management practice. There is a need for further research in this field to establish with certainty the procedures that are necessary to prevent and control wound infection. Until such research exists, guidance based on the current evidence base, evidence derived from other clinical procedures (for example, intravenous therapy), broader guidelines^{12,13,69} and expert opinion is required to assist facilities in developing local policies and procedures.

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The voluntary contribution that each reviewer makes contributes greatly to the high standards that Wound Practice and Research strives to achieve.