Antiseptic Use in Orthopaedic Wounds

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Authors’ contributions

This work was carried out in collaboration among all authors. All authors contributed equally to the production of this review. All authors read and approved the final manuscript.

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ABSTRACT

Objectives: The aim of this study is to review the available literature addressing the safety and efficacy of antiseptics in surgical wounds. The different antiseptic solutions, irrigation volumes, time scales and delivery methods have been compared so that evidence-based recommendations on antiseptic use in orthopaedic, foot and ankle surgical procedures can be proposed.

Methods: A literature search was performed using the online databases Medline and EMBase to identify in-vitro and in-vivo studies pertaining to antiseptic use in an orthopaedic context. Terms including antiseptic, irrigation fluid, bacitracin, hydrogen peroxide, povidone-iodine and chlorhexidine were searched. Literature published in English from inception to July 2020 in which the full text was accessible was considered for inclusion. Cellular and animal studies were included on the basis that authors analysed antiseptic efficacy and/or toxic effect of antiseptic on cells present in orthopaedic wounds. Clinical studies that met the criteria for inclusion in this review assessed antiseptic use in a surgical context, with a focus on foot and ankle procedures. These included case reports, case series, case control, prospective and retrospective studies as well as randomised controlled trials. Studies were categorised as in-vitro, animal and human studies. Twenty-three, eleven and forty-four studies were identified as in-vitro, animal and human studies respectively. These have been summarised and presented herein in a narrative format.

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Results: There is strong evidence that skin preparation with antiseptics before orthopaedic procedures reduces the risk of post-operative infection. **Conclusion:** Routine prophylactic intra-operative antiseptic use should be performed with caution as they increase the risk of local and systemic complications. However, there is strong evidence supporting the use of antiseptics pre-operatively when preparing the skin. Determining the best antiseptic preparation remains a matter of debate since a single agent or solution is not effective against all organisms. Further research is therefore needed to assess the efficacy of antiseptics in prevention and treatment of infections.

Keywords: Antiseptic; orthopaedic wounds; prevention; treatment.

1. INTRODUCTION

Deep wound infections are a devastating complication of orthopaedic surgery, particularly in the presence of implants. The bacterial load required to cause clinical infection is markedly reduced in complex orthopaedic wounds [1]. Two-thirds of such infections are mono-microbial, the most commonly isolated bacteria being *Staphylococcus aureus* and *Staphylococcus epidermidis* [2]. Such bacteria are able to colonise the surface of implanted materials by molecular mechanisms of adherence. Microbes then form biofilms which constitute a multi-layered defense mechanism innately resistant to antimicrobial penetration. Prosthetic joint infections are thus notoriously difficult to treat, requiring prolonged courses of systemic antibiotics and implant exchange.

The focus of management should therefore be primary prevention. Peri-operative antibiotics and topical antimicrobial agents are standard practice. Wound debridement and irrigation also have major roles in prophylaxis and management of infections. However, the incidence of complex wound infections remains acceptably high. The problem is compounded further by continuing emergence of multiply resistant bacteria. The challenge therefore remains to find an antiseptic able to eliminate targeted pathogens while being safe for the patient.

This study aims to review available literature addressing the safety and efficacy of antiseptics in surgical wounds. The different antiseptic solutions, time scales and delivery methods have been compared so that evidence-based recommendations on antiseptic use in orthopaedic, foot and ankle surgical procedures can be proposed.

2. In-vitro STUDIES

2.1 Efficacy and Use of Antiseptics

The efficacy of an irrigation solution relies on solution composition and delivery. To be useful in orthopaedic practice, antiseptics must eradicate commonly encountered pathogens and act on both tissues and orthopaedic metalwork. Gainor et al. [3] showed that benzalkonium chloride is more effective than saline to disinfect bovine muscle contaminated with *Staphylococcus aureus* (*S. aureus*), *Staphylococcus epidermidis* (*S. epidermidis*) or *Pseudomonas aeruginosa* (*P. aeruginosa*). Moussa et al. [4] colonised stainless-steel screws with a preformed biofilm of these bacteria then immersed them in benzalkonium chloride solution, confirming its efficacy when used to disinfect orthopaedic hardware. Presterl et al. [5] used biofilms of *S. epidermidis* isolated from patients with catheter-related bacteraemia and cardiac implant infections, showing that hydrogen peroxide 3% and 5% and N-propanol significantly reduced biofilm thickness, while povidone-iodine proved less effective.

Anglen et al. [6] compared various solutions and delivery methods using *S. aureus*-coated stainless-steel screws. Irrigation with saline, liquid soap or antibiotics (bacitracin, neomycin and polymyxin/neomycin) had the greatest effect on *S. aureus* when using liquid soap. Washout by bulb syringe or jet lavage highlighted the importance of administration technique used; bacterial removal increased by over 100-fold with jet lavage, regardless of the solution used. A follow-up study [7] revealed bacterial elimination depends on the interplay between the irrigation solution, species present, and surface involved. Three surface types (cortical bone, stainless-steel and titanium) coated with *S. aureus*, *S. epidermidis* or *P. aeruginosa* were irrigated with normal saline, soap solution, bacitracin or neomycin. Jet lavage using all solutions resulted in considerably fewer bacteria compared to an
unwashed control group. Soap solution proved superior at removing all bacteria from all three surfaces. Authors therefore concluded that when removing some species from metallic surfaces, soap solution may serve as a better irrigation additive, especially with jet lavage delivery.

The efficacy of soap solutions is concentration and time-dependent; Krueger et al. [8] contaminated porcine bone segments with S. aureus. Soaking in normal saline, 2% chlorhexidine, or 4% chlorhexidine then re-imaging after 5, 10, 20, 30, and 60-minute intervals showed significantly fewer bacteria at all time intervals in the chlorhexidine groups compared with the saline group, but no significant difference between chlorhexidine groups. Given the potential cytotoxicity of chlorhexidine, the authors recommended soaking contaminated bone segments with 2% chlorhexidine for 20 minutes.

### 2.2 Cytotoxic Effects of Antiseptics

The clinical suitability of antiseptics in irrigation fluids depends on their ability to remove pathogens from orthopaedic wounds and prostheses without damaging the osteoblasts, chondrocytes, fibroblasts, keratinocytes and lymphocytes. *In-vitro* studies have enabled careful evaluation of the varying degrees of cytotoxicity associated with these cell-types.

Table 1 summarises these adverse effects associated with different antiseptics

### 2.3 Antiseptics and MRSA

Multi-drug-resistant infections pose an even bigger challenge for eliminating pathogens while avoiding adverse effects. Methicillin-resistant *Staphylococcus aureus* (MRSA) is increasingly responsible for intra-operative infection and subsequent morbidity [22]. While vancomycin is the antibiotic of choice for MRSA infections, multidrug-resistant strains with moderate vancomycin resistance have been isolated [23]. This resistance increases post-operative complication rates and is further compounded by MRSA’s tendency to form biofilms. Various studies have looked specifically at reducing the operative risk of MRSA by evaluating the efficacy of irrigation solutions and debridement techniques. Haley et al. [24] found Povidone-iodine to be the most rapidly effective antiseptic against both MRSA and MSSA, maximal effectiveness being evident at 1:100 dilutions, which killed all strains within 15 seconds. The other three antiseptic solutions (chlorhexidine gluconate-alcohol (4%), p-chloro-m-xyleneol (1%), and hexachlorophene (3%)) tested produced 2-log reductions in the MRSA CFU count after 15 seconds of exposure but failed to kill all MRSA, even after 240 seconds of exposure.

More recent studies however, support chlorhexidine as the most effective irrigation solution for eradicating MRSA infection. Schwechter et al. [25] evaluated the efficacy of a number of *in vitro* irrigation and debridement techniques for treating MRSA peri-prosthetic joint infection, using MRSA biofilm-coated titanium alloy discs as model. Irrigation and scrubbing were not shown to be more effective than irrigation alone. Both chlorhexidine and povidone-iodine scrubs reduced MRSA CFU counts pre-incubation, however, chlorhexidine was found to be the most bactericidal while the results for povidone-iodine were not statistically significant. The study therefore concluded that chlorhexidine solutions have the greatest potential to decrease biofilm load on orthopaedic implants. Determining the minimum concentration of chlorhexidine required to eradicate MRSA is important to reduce the risk of antiseptic-induced cytotoxicity. This was evaluated in an *in vitro* study by Smith et al. [26] where a series of MRSA biofilm-coated titanium discs were irrigated with varying concentrations of chlorhexidine. MRSA CFUs were counted before and after a 24-hour re-incubation period. The study showed a significant reduction in CFUs at all irrigation concentrations before re-incubation. However, post re-incubation counts only demonstrated a significant decrease with 4% and 2% chlorhexidine solutions, indicating that 2% was the minimum effective chlorhexidine concentration.

*In-vitro* studies have generated variable and at times conflicting, data for the different irrigation solutions. Many of the *in vitro* studies discussed in this review address either the efficacy of the antiseptic or its potential cytotoxicity which would limit its overall clinical applicability. For example, the efficacy of chlorhexidine against MRSA has been demonstrated in a number of studies [25,26] but other studies have demonstrated potential cytotoxicity to osteoblasts, fibroblasts, stromal cells and keratinocytes which would discourage its use in clinical practice. *In-vitro* studies are limited in that they do not assess the antiseptic in the presence of all cells, chemicals, mediators and other factors that would be present in a living animal or human and which may have a role in the post-operative healing
process, immune response to an invading organism, and reaction to the antiseptic itself. *In-vivo* studies are therefore essential to clarify the optimal choice and use of irrigation solutions.

### 3. ANIMAL STUDIES

In the living organism, factors in addition to those monitored through *in-vitro* studies determine the effect of the irrigating solution. The components of the irrigating solution or the physical forces from pressure lavage may delay wound healing or facilitate wound colonisation by bacteria that may be present. Tissues in contact with the solution may mount an inflammatory response that could lead to complications, including joint fibrosis and prosthesis failure. Absorption of lavage fluid components may also lead to systemic side-effects. *In vivo* studies are therefore important to give a clearer demonstration of how an irrigating solution will perform in clinical practice.

Conroy et al. [27] designed an animal study using rat models, to test the efficacy of a number of wound irrigation solutions which included normal saline, castile soap, benzalkonium chloride and bacitracin. The effectiveness of sequential irrigation with 1L each of benzalkonium, castile soap and normal saline was also tested. Orthopaedic wounds contaminated with *S. aureus* or *P. aeruginosa* were washed with 3L of the specified solution and inspected daily for 14 days. Benzalkonium irrigation alone significantly lowered the rate of *S. aureus* positive wound cultures compared with normal saline but increased the rate of *P. aeruginosa* positive wound cultures. In contrast, irrigation with either castile soap alone or sequential irrigation with benzalkonium/castile soap/saline reduced the rate of *P. aeruginosa* cultures compared to irrigation with normal saline alone. The authors proposed that benzalkonium may cause lysis of gram-negative cells leading to release of endotoxin and producing an inflammatory response. They therefore suggested that in wounds containing *P. aeruginosa*, benzalkonium can only be used safely if combined in a sequential irrigation with castile soap and normal saline.

The effectiveness of a sequential surfactant irrigation protocol in overcoming bacterial adhesion in contaminated orthopaedic implants was explored further in a randomised animal study carried out by Marberry et al. [2]. This study suggested that the sequential surfactant irrigation protocol is an effective method of wound irrigation in orthopaedic implants contaminated with *S. aureus* alone, while in mono-microbial *S. epidermidis* infections normal saline irrigation alone may suffice.

Irrigation devices are also applied in the treatment of open wounds, however there is currently no clear consensus for its use. Owens et al. [28] compared the results of normal saline versus multiple other irrigation solutions using either a bulb syringe or pulsatile lavage. They found that both irrigation devices reduced bacterial counts but these had rebounded to almost pre-treatment levels in the pulsed lavage group while the bulb syringe group showed a significantly lower rebound. They also found that all irrigation solutions initially lowered bacterial counts, however, 48 hours after irrigation, all groups showed a rebound in bacterial counts. This rebound was the lowest with normal saline while the other solutions showed rebounds approaching or even surpassing pre-treatment levels. These findings suggested that although solutions other than normal saline may be better at initial bacterial removal, these solutions may have deleterious effects to the host tissue and these effects may be compounded by use of high-pressure devices used for their delivery. The authors therefore recommended saline solution delivered by a low-pressure device to irrigate wounds in order to reduce the risk of complications.

Povidone-iodine (Betadine) lavage is used prophylactically in various surgical sub-specialities and its efficacy in preventing surgical site infection is well established. There have, however, been concerns about its effect on fibroblasts and its potential to damage intra-capsular tissues. Baird et al. [29] found 0.1% povidone-iodine lavage to be safe to use on equine tendon, as it caused only a mild synovitis similar to that caused by balanced electrolyte solution. However, 0.5% povidone iodine and 0.5% chlorhexidine caused severe synovitis and therefore should not be used for tendon sheath lavage. Moreover, chlorhexidine causes greater detriment to joints, as concentrations of just 0.05% have been shown to cause intense synovitis. Keudell et al. [30] evaluated the potential chondrotoxicity of various povidone-iodine concentrations and exposure times in a cadaveric animal model. The higher concentrations all showed greater chondrotoxicity with increased exposure times, therefore this should be considered in articular cartilage-retaining procedures, such as...
uncompartmental knee arthroplasty, and opting to use saline may be prudent, as reduced chondrocyte viability can contribute to early development of osteoarthritis and osteoarthritis in the residual cartilage, leading to early failure of partial knee arthroplasty. This would not, however, be a problem in total joint arthroplasties where articular cartilage is not retained.

Acute post-operative arthroplasty infection was evaluated in a rabbit prosthetic knee infection model [31]. Metalwork inoculated with S. aureus was inserted into the knee joints for a total of seven days, following which a polyethylene washer exchange was performed and irrigation with either dilute 3.5% povidone-iodine solution or normal saline. Culture of the harvested screw, polyethylene washer, bone and joint tissue with bacterial quantification showed a significant reduction in bacterial burden on both the screw and polyethylene washer that had received povidone-iodine lavage, although there was no difference in soft-tissue growth. The study thus showed that povidone-iodine lavage may be a useful treatment adjunct in acute post-operative arthroplasty infection, and it may improve the rate of component retention.

Chlorhexidine gluconate 0.05% peritoneal lavage has been shown to be effective in reducing the rate of abscess formation following peritoneal contamination [32]. However, its suitability and ideal concentration for use in joint lavage remains in dispute. Sanchez et al showed that bactericidal concentrations of chlorhexidine diacetate are lethal to canine embryonic fibroblasts in-vitro [18]. However, a subsequent study by the same team showed that irrigation with chlorhexidine diacetate 0.05% or 0.005% provided bactericidal activity and improved wound healing compared to irrigation with saline alone, indicating that chlorhexidine diacetate concentrations that are cytotoxic to cultured fibroblasts in-vitro do not interfere with wound healing in-vivo.

Contrastingly, in a study on open fractures, Barwell et al found that both chlorhexidine and saline reduced bacterial loads equally [33]. However, they concluded that although chlorhexidine has an initial bactericidal benefit, the necrotic tissue caused by chlorhexidine exposure may promote a "rebound" of bacterial growth in the wound bed. They therefore suggested that saline remains the best choice for orthopaedic wound irrigation, however if chlorhexidine is preferred, a concentration of 0.05% should be used and the wound should undergo a final saline rinse to remove any residue before closure. The authors also urged particular caution with chlorhexidine lavage in wounds containing large amounts of tissue of borderline viability as such wounds may be more susceptible to the toxic effects of chlorhexidine.

The potential role of the antibacterial allicin in preventing biofilm formation in joint infection was investigated in a rabbit prosthetic joint infection model by Zhai et al. [34]. Biofilm counts of S. epidermidis were found to be lowest in rabbits treated with allicin plus vancomycin. The authors therefore concluded that allicin/vancomycin combination may have a role in treatment of prosthetic joint infections as allicin appears to inhibit biofilm formation and it can have a synergistic bactericidal effect when combined with vancomycin.

The use of different irrigation solutions in preparing bony surfaces before cementing of total joint replacement components was evaluated by Howells et al. [35]. Cement fixation following hydrogen peroxide irrigation was found to be statistically better than that achieved with either povidone iodine or saline. Clinically, dilute povidone-iodine lavage is generally performed following implant insertion and it therefore should not affect the bone-cement interface. However, if performing single-stage revision, the potential benefit of hydrogen peroxide should be considered.

4. HUMAN STUDIES

A number of human studies have evaluated the optimal irrigation fluid composition and volume to prevent or control of the musculoskeletal infection. Though the use of lavage fluid in arthroplasty has been studied extensively, a gold standard of care has yet to be determined. Studies have shown that pre-operative skin preparation with chlorhexidine reduces the incidence of periprosthetic joint infections but its superiority over other agents such as povidone-iodine in periprosthetic joint infection is inconclusive [36]. The National Institute for Health and Care Excellence (NICE, 2019) authorise the pre-operative preparation of skin using chlorhexidine (0.5% chlorhexidine in 70% alcohol solution prior to minor surgical procedures and 2.0% chlorhexidine in 70% alcohol applicators prior to invasive medical procedures), or povidone-iodine if chlorhexidine is contraindicated [37].
## Table 1. Cytotoxic effects established by In-vitro studies

<table>
<thead>
<tr>
<th>Affected cell-type</th>
<th>Bacitracin</th>
<th>Hydrogen Peroxide</th>
<th>Povidone-iodine (PI/Betadine)</th>
<th>Chlorhexidine gluconate (CHX)</th>
<th>Other solutions</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Bone</td>
<td>Osteoblasts unaffected [9, 10].</td>
<td>Cytotoxic to osteoblasts at high concentrations [9, 10].</td>
<td>Osteoblasts: Affected in dose-[11,12] and time-[14] dependent manner. Cytotoxicity was shown at high concentrations. [9,10] At concentrations of up to 5% for up to 1-minute osteoblast activity was not impaired. [11] Osteoclasts: Affected in time-dependent manner [13].</td>
<td>Osteoblasts: Affected in dose-[11,12] and time-[14] dependent manner. Osteoclasts: Affected in time-dependent manner [14]. Safe at concentrations of 0.2% for up to 1 minute and 1% for up to 30 seconds.[11]</td>
<td>Sodium hypochlorite: Limits osteoblast growth in dose dependent manner [11]. Soap solution: Affects osteoblasts and osteoclasts in time dependent manner but preserved the greatest number of osteoclasts compared with PI and CHX [13].</td>
<td>• Kellam et al. [9] • Kaysinger et al. [10] • Verdugo et al. [11] • Cabral et al. [12] • Bhandari et al. [13]</td>
</tr>
<tr>
<td>Cartilage</td>
<td>Not assessed by these reviewers.</td>
<td>Cytotoxic to chondrocytes.[14-16]</td>
<td>Not assessed by these reviewers.</td>
<td>Not assessed by these reviewers.</td>
<td>Not assessed by these reviewers.</td>
<td>• Bates et al. [14] • Asada et al. [15] • Asada et al. [16]</td>
</tr>
<tr>
<td>Connective tissue</td>
<td>Not assessed by these reviewers.</td>
<td>Reduced fibroblast activity with cytotoxicity shown at bactericidal concentrations [18].</td>
<td>Reduced fibroblast activity [18]. Fibroblast cytotoxicity resulted at bactericidal concentrations [19]. Fibroblast viability retained at 1.3 g/L with moderate bactericidal effect [21].</td>
<td>Increased fibroblast proliferative potential at low doses [18]. Fibroblast cytotoxicity resulted at bactericidal concentrations [19-21] Cytotoxic to stromal cells [21].</td>
<td>Silver sulfasalazine: Increased fibroblast proliferative potential at low doses.[18] Therapeutic doses of local antibiotics did not damage fibroblasts.[21]</td>
<td>• Sanchez et al. [18] • Mariotti et al. [19] • Meurs et al. [20] • Damour et al. [21]</td>
</tr>
<tr>
<td>Epidermal</td>
<td>Not assessed by these reviewers.</td>
<td>Not assessed by these reviewers.</td>
<td>Cytotoxicity to keratinocytes was shown at bactericidal concentrations [21].</td>
<td>Cytotoxicity to keratinocytes was shown at bactericidal concentrations [21].</td>
<td>Therapeutic doses of local antibiotics did not damage keratinocytes.[21]</td>
<td>• Damour et al. [21]</td>
</tr>
</tbody>
</table>
Table 2. Efficacy of irrigation solutions, as established by animal studies

<table>
<thead>
<tr>
<th>Animal Model</th>
<th>Irrigation solution</th>
<th>Staphylococcus aureus</th>
<th>Staphylococcus epidermis</th>
<th>Pseudomonas aeruginosa</th>
<th>References</th>
</tr>
</thead>
<tbody>
<tr>
<td>Rat</td>
<td>Normal saline (NS)</td>
<td>✓ [27]</td>
<td>✓ [2]</td>
<td>✓ [27]</td>
<td>• Conroy et al. [27]</td>
</tr>
<tr>
<td></td>
<td>Castile soap (CS)</td>
<td>✓ [27]</td>
<td>✓ [27]</td>
<td>✓ [27]</td>
<td>• Marberry et al. [2]</td>
</tr>
<tr>
<td></td>
<td>Benzalkonium chloride (BzC)</td>
<td>✓ [27]</td>
<td>✓ [27]</td>
<td>✓ [27]</td>
<td>• Penn-Barwell et al. [33]</td>
</tr>
<tr>
<td></td>
<td>Bacitracin</td>
<td>✓ [27]</td>
<td>✓ [27]</td>
<td>✓ [27]</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Chlorhexidine</td>
<td>✓ [27]</td>
<td>✓ [27]</td>
<td>✓ [27]</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Sequential irrigation: BzC + CS + NS</td>
<td>✓ [2]</td>
<td>✓ [2]</td>
<td>✓ [27]</td>
<td></td>
</tr>
<tr>
<td>Rabbit</td>
<td>Normal saline</td>
<td>✓ [31]</td>
<td>✓ [34]</td>
<td>✓ [34]</td>
<td>• Gilotra et al. [31]</td>
</tr>
<tr>
<td></td>
<td>Povidone-iodine</td>
<td>✓ [31]</td>
<td>✓ [34]</td>
<td>✓ [34]</td>
<td>• Zhai et al. [34]</td>
</tr>
<tr>
<td></td>
<td>Allicin</td>
<td>✓ [31]</td>
<td>✓ [34]</td>
<td>✓ [34]</td>
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<tr>
<td></td>
<td>Vancomycin</td>
<td>✓ [31]</td>
<td>✓ [34]</td>
<td>✓ [34]</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Allicin + Vancomycin</td>
<td>✓ [31]</td>
<td>✓ [34]</td>
<td>✓ [34]</td>
<td></td>
</tr>
<tr>
<td>Goat</td>
<td>Normal saline</td>
<td>✓ [28]</td>
<td>✓ [28]</td>
<td>✓ [28]</td>
<td>• Owens et al. [28]</td>
</tr>
<tr>
<td></td>
<td>Castile soap</td>
<td>✓ [28]</td>
<td>✓ [28]</td>
<td>✓ [28]</td>
<td></td>
</tr>
<tr>
<td></td>
<td>Benzalkonium chloride</td>
<td>✓ [28]</td>
<td>✓ [28]</td>
<td>✓ [28]</td>
<td></td>
</tr>
</tbody>
</table>

✓: Solution demonstrated statistically significant decrease in bacterial count
✗: Solution did not demonstrate statistically significant decrease in bacterial count
- : Bacteria/solution combination not tested

Table 3. Efficacy of antiseptic solutions in humans

<table>
<thead>
<tr>
<th>Study</th>
<th>Procedure</th>
<th>Variant</th>
<th>Indication</th>
<th>Antiseptic(s)</th>
<th>Type of Study</th>
<th>Outcomes Assessed</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Patrick et al.</td>
<td>Elective spinal surgery</td>
<td>Pre-op preparation</td>
<td>Primary prophylaxis</td>
<td>Skin disinfection with sequential povidone-iodine + chlorhexidine vs povidone-iodine alone twice</td>
<td>Randomised controlled trial</td>
<td>Surgical site contamination + internal bacterial contamination</td>
<td>Reduced risk of 30% and 37% in surgical site contamination and internal bacterial contamination with sequential disinfection</td>
</tr>
<tr>
<td>Study</td>
<td>Procedure</td>
<td>Variant</td>
<td>Indication</td>
<td>Antiseptic(s)</td>
<td>Type of Study</td>
<td>Outcomes Assessed</td>
<td>Results</td>
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</tr>
<tr>
<td>Woo et al. [41]</td>
<td>Knee joint replacement</td>
<td>Intra-operative irrigation</td>
<td>Primary prophylaxis</td>
<td>Taurolidine (synthetic broad-spectrum antibiotic) irrigation</td>
<td>Case-control</td>
<td>Lower CRP + ESR values 3 days after surgery but less significant 6 days post-op with taurolidine vs control</td>
<td>Irrigation with taurolidine did not decrease the risk of infection</td>
</tr>
<tr>
<td>Chundamala et al. [42]</td>
<td>General, cardiovascular, orthopaedic spine, and urologic surgery</td>
<td>Intra-operative irrigation</td>
<td>Primary prophylaxis</td>
<td>Povidone-iodine vs normal saline irrigation</td>
<td>Meta-analysis</td>
<td>Post-operative infection</td>
<td>Reduced risk of infection with povidone-iodine (p = 0.015 and p = 0.007, p &lt; 0.05)</td>
</tr>
<tr>
<td>Brown et al. [43]</td>
<td>Hip and knee joint replacement</td>
<td>Intra-operative irrigation</td>
<td>Primary prophylaxis</td>
<td>Povidone-iodine vs normal saline irrigation</td>
<td>Case-control</td>
<td>Post-operative infection</td>
<td>Significant reduction with povidone-iodine (0.15 vs 0.97%)</td>
</tr>
<tr>
<td>Frisch et al. [44]</td>
<td>Hip and knee joint replacement</td>
<td>Intra-operative irrigation</td>
<td>Primary prophylaxis</td>
<td>Chlorhexidine 0.05% vs povidone-iodine (&lt;2%) for hip replacement, and chlorhexidine 0.05% vs 0.9% saline for knee replacement</td>
<td>Case-control</td>
<td>Deep and superficial site infections</td>
<td>No significant difference</td>
</tr>
<tr>
<td>Ulivieri et al [45].</td>
<td>Elective spinal surgery</td>
<td>Intra-operative irrigation</td>
<td>Primary prophylaxis</td>
<td>Combination irrigation protocol (soak with 10 cc of 10% povidone-iodine, 5cc of water and 1cc of hydrogen peroxide and after am minute irrigation with copious sterile saline)</td>
<td>Case-control</td>
<td>Post-operative infection</td>
<td>Significant reduction with combination irrigation protocol (0% vs 1.5%)</td>
</tr>
<tr>
<td>Kosashvili et al. [46]</td>
<td>Multiple and first time revision spinal</td>
<td>Intra-operative irrigation</td>
<td>Primary prophylaxis</td>
<td>Combination irrigation protocol (povidone-iodine, hydrogen)</td>
<td>Case-control</td>
<td>Post-operative infection</td>
<td>Reduced risk of infection with combination irrigation protocol (2.14% in multiple)</td>
</tr>
<tr>
<td>Study</td>
<td>Procedure</td>
<td>Variant</td>
<td>Indication</td>
<td>Antiseptic(s)</td>
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<tr>
<td>Patterson et al. [47]</td>
<td>Surgery</td>
<td>Pin dressing</td>
<td>Primary prophylaxis</td>
<td>Peroxide and bacitracin</td>
<td>Half-strength study</td>
<td>Pin-site infection</td>
<td>Lower infection rate with half-strength peroxide and Xeroform dressing</td>
</tr>
<tr>
<td>Woolridge et al. [48]</td>
<td>High-grade soft tissue sarcoma resection</td>
<td>Adjuvant local irrigation</td>
<td>Primary prophylaxis</td>
<td>Hydrogen peroxide solution</td>
<td>Retrospective case-control</td>
<td>Local recurrence and surgical site infection</td>
<td>Reduced local control and infection rates</td>
</tr>
<tr>
<td>Anglen et al. [49]</td>
<td>Lower limb open fracture</td>
<td>Open wound irrigation</td>
<td>Primary prophylaxis</td>
<td>Bacitracin solution vs a non-sterile castile soap solution</td>
<td>Randomised controlled trial</td>
<td>Post-operative infection</td>
<td>No advantage and may increase the risk of wound-healing problems with bacitracin solution</td>
</tr>
<tr>
<td>Conroy et al. [27]</td>
<td>Complex contaminated orthopaedic wound</td>
<td>Open wound irrigation</td>
<td>Primary prophylaxis</td>
<td>Castile soap vs benzalkonium chloride vs anracin</td>
<td>Case-control</td>
<td>Post-operative infection</td>
<td>No benefit of antibiotic solutions over normal saline</td>
</tr>
<tr>
<td>Bhandari et al. [50]</td>
<td>Open fracture</td>
<td>Wound irrigation</td>
<td>Primary prophylaxis</td>
<td>Castile soap versus normal saline irrigation and its pressure</td>
<td>Randomised controlled trial</td>
<td>Re-operation rate</td>
<td>Reduced with saline regardless of irrigation pressure</td>
</tr>
<tr>
<td>Yung Han et al. [51]</td>
<td>Contaminated anterior cruciate ligament grafts</td>
<td>Wound irrigation</td>
<td>Treatment of contaminated ligament grafts</td>
<td>3L 2% chlorhexidine power irrigation</td>
<td>Controlled laboratory study</td>
<td>Post irrigation disinfection and weakness of the anterior cruciate ligament.</td>
<td>Disinfection using 3L 2% chlorhexidine power irrigation does not significantly weaken the tendon.</td>
</tr>
<tr>
<td>Burd et al. [52]</td>
<td>Contaminated Achilles tendon-calcaneous allografts</td>
<td>Wound irrigation</td>
<td>Treatment of contaminated tendon grafts</td>
<td>Benzalkonium chloride vs castile soap vs castile soap followed by benzalkonium chloride vs triple</td>
<td>Case-control</td>
<td>Post-irrigation infection</td>
<td>2% chlorhexidine irrigation solution is effective in decontaminating bone-tendon allografts infected with <em>Staphylococcus aureus, Staphylococcus</em></td>
</tr>
<tr>
<td>Study</td>
<td>Procedure</td>
<td>Variant</td>
<td>Indication</td>
<td>Antiseptic(s)</td>
<td>Type of Study</td>
<td>Outcomes Assessed</td>
<td>Results</td>
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<tr>
<td>Ostrander et al. [53]</td>
<td>Procedures involving the hallux, toes and tibia.</td>
<td>Pre-op preparation</td>
<td>Primary prophylaxis</td>
<td>antibiotic vs chlorhexidine gluconate vs chlorhexidine gluconate/triple antibiotic combination</td>
<td>Prospective study</td>
<td>Epidermis, pseudomonas aeruginosa or Klebsiella pneumoniae.</td>
<td>All three solutions were effective. Of the three, ChloraPrep was most effective at bacterial elimination (p &lt; 0.0001).</td>
</tr>
<tr>
<td>Saltzman et al. [54]</td>
<td>Shoulder surgery</td>
<td>Pre-op preparation</td>
<td>Primary prophylaxis</td>
<td>DuraPrep (0.7% iodine and 74% isopropyl alcohol) vs Techni-Care (3.0% chloroxylenol) vs ChloraPrep (2% chlorhexidine gluconate and 70% isopropyl alcohol)</td>
<td>Prospective study</td>
<td>Post-preparation infection of the hallux nailfold, web spaces between the 2nd/3rd and 4th/5th digits, and the anterior tibia (control)</td>
<td>ChloraPrep was most effective at bacterial elimination (p &lt; 0.0001).</td>
</tr>
<tr>
<td>Hunter et al. [55]</td>
<td>Foot and ankle surgery</td>
<td>Pre-op preparation</td>
<td>Primary prophylaxis</td>
<td>70% isopropyl alcohol then 4% chlorhexidine gluconate vs 4% chlorhexidine gluconate then 70% isopropyl alcohol</td>
<td>Randomised prospective study</td>
<td>Post-preparation colonisation of the operative site.</td>
<td>The combination of 70% isopropyl alcohol followed by 4% chlorhexidine gluconate was more effective.</td>
</tr>
<tr>
<td>Mankovecky et al. [56]</td>
<td>Ankle septic arthritis</td>
<td>Arthroscopic joint irrigation</td>
<td>Septic arthritis</td>
<td>Irrigation with lactated Ringer's solution impregnated with Bacitracin sterile powder</td>
<td>Case series</td>
<td>Infection control</td>
<td>Successful elimination of infection in all 15 cases</td>
</tr>
<tr>
<td>Study</td>
<td>Procedure</td>
<td>Variant</td>
<td>Indication</td>
<td>Antiseptic(s)</td>
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<tr>
<td>Acello et al.</td>
<td>Open foot and ankle fractures</td>
<td>Open wound irrigation</td>
<td>Primary prophylaxis</td>
<td>50,000 units bacitracin vs 25 mg polymyxin per L vs sterile saline</td>
<td>Retrospective study</td>
<td>Post-operative infection</td>
<td>Irrigation with bacitracin or polymyxin is beneficial in reducing infection in open fracture repairs.</td>
</tr>
<tr>
<td>Wu et al.</td>
<td>Foot and ankle external fixation</td>
<td>Pre-op preparation</td>
<td>Primary prophylaxis</td>
<td>Chlorhexidine gluconate-impregnated polyurethane patches vs standard pin care.</td>
<td>Pilot trial</td>
<td>Post-operative infection</td>
<td>There were significantly lower infection rates in the group receiving chlorhexidine patches on the pin sites (0% versus 25%, P = 0.047)</td>
</tr>
<tr>
<td>Ruder et al.</td>
<td>Total joint arthroplasty</td>
<td>Open wound irrigation</td>
<td>Primary prophylaxis</td>
<td>17.5ml 10% povidone-iodine diluted in 500-1000cc normal saline</td>
<td>Expert opinion report</td>
<td>Post-operative infection</td>
<td>In primary total joint arthroplasty, 10% povidone-iodine reduces the risk of infection.</td>
</tr>
<tr>
<td>George et al.</td>
<td>Hip and knee arthroplasty</td>
<td>Open wound irrigation</td>
<td>Prosthetic joint infection</td>
<td>1% povidone iodine and a 50:50 dilution of 3% hydrogen peroxide</td>
<td>Case series</td>
<td>Post-operative infection</td>
<td>There were no recurrences of infection in 11 hips at a mean of 5 years and 28 knees at a mean of 6.5 years</td>
</tr>
<tr>
<td>Riesgo et al.</td>
<td>Total joint arthroplasty</td>
<td>Open wound irrigation</td>
<td>Prosthetic joint infection</td>
<td>Vancomycin povidone-iodine + irrigation and debridement with modular component and linear exchange (IDLE) vs IDLE alone.</td>
<td>Case-control</td>
<td>Post-operative infection</td>
<td>Vancomycin povidone-iodine is effective in treating prosthetic joint infection with irrigation and debridement. Its use resulted in failure (infection) in 16.7% (6/36), lower than the control group (37%, i.e. 14/38).</td>
</tr>
</tbody>
</table>
Table 4. Adverse effects associated with antiseptic fluids

<table>
<thead>
<tr>
<th>Antiseptic fluid</th>
<th>Adverse Effect</th>
</tr>
</thead>
<tbody>
<tr>
<td>Chlorhexidine</td>
<td>Chondrolysis [62,63]</td>
</tr>
<tr>
<td>Hydrogen peroxide</td>
<td>Chondrolysis [15-17], pneumocephalus and air embolism [64-68]</td>
</tr>
<tr>
<td>Povidone-iodine</td>
<td>Acute kidney injury [69-72], induce hyperthyroidism [69,73-75]</td>
</tr>
<tr>
<td>Alcohol-based solutions: Chlorhexidine, povidone-iodine</td>
<td>Electrocautery-associated surgical fire [76-79]</td>
</tr>
</tbody>
</table>

Although no “gold standard” of irrigation fluid currently exists for the management of open fracture wounds, a survey completed by 1764 surgeons found that the majority of surgeons favor both normal saline and low-pressure lavage as part of the initial treatment [38]. In a recent document from the 2nd International Consensus Meeting on Musculoskeletal Infection no particular recommendation was made with regards the use of dilute povidone-iodine (betadine) irrigation or other antiseptic irrigation solutions during total ankle arthroplasty (TAA) or other foot and ankle procedures due to lack of good quality evidence [39].

Hansen and Parvizi [40] described a specific irrigation and debridement regimen involving five steps to eradicate periprosthetic infection: (1) soak the surgical site in Dakin’s solution or hydrogen peroxide for 3 minutes, (2) irrigate wound thoroughly with 3 litres of sterile saline, (3) add 0.3% dilute betadine for another three minutes while continuing to mechanically scrub and debride the surgical site, (4) irrigate surgical site again with 3 litres of sterile saline, and (5) irrigate site with 3 litres of saline containing 500,000 units of polymyxin B and 50,000 units of bacitracin. As yet, there is no secondary peer-reviewed research to affirm this protocol, although studies do describe debridement and copious irrigation of the surgical site for operative management of infection. Table 3 summarises efficacy of antiseptic solutions in humans.

4.1 Adverse Effects of Irrigation

Table 4 summarises adverse effects associated with different antiseptic solutions in human.

5. CONCLUSION

Routine intraoperative irrigation with antiseptic solutions for the prevention or treatment of established infection should be undertaken with caution. It is advisable to consider a number of circumstances where it may be detrimental, in particular with the use of Chlorhexidine and Hydrogen Peroxide within native joint cavities in view of their chondrotoxicity. Systemic complications such as thyroid [68,72-74] and kidney [68-71] dysfunction, air embolism and even death have been reported [64-67]. There is strong evidence that skin preparation with antiseptics before orthopaedic procedures reduces the risk of post-operative infection. Although the best antiseptic preparation remains a matter of debate. A single agent or solution is also not effective against all organisms (Table 2) leading to the development of some complex regimens that are difficult to validate.

The social and economic burden of bone and joint infections associated with implants are high and life altering for the afflicted individual. The treatment invariably requires multiple procedures, prolonged inpatient stay and leads to poor functional outcomes. This combined with the emergence of resistant strains of microorganisms and ever more compromised hosts make prevention an essential tool in the armamentarium of the surgeon. Therefore, further research is required to assess efficacy of different antiseptic solutions for prevention and treatment of infection. Current knowledge based on available literature is insufficient to strongly recommend a particular antiseptic and its volume in clinical practice. However, we have addressed the National Institute for Health and Care Excellence (NICE) recommendation and the “Second International Consensus Meeting on Musculoskeletal Infection” as the latest recommendations regarding antiseptic use in orthopaedic practice.

ACCESS TO DATA

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CONSENT

It is not applicable.

ETHICAL APPROVAL

It is not applicable.

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COMPETING INTERESTS

Authors have declared that no competing interests exist.

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