Wisconsin Division of Public Health Supplemental Guidance for the Prevention of Surgical Site Infections: An Evidence-Based Perspective

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Charles E. Edmiston, Jr., PhD, CIC, FIDSA, FSHEA, FAPIC  
Emeritus Professor of Surgery  
Medical College of Wisconsin  
Milwaukee  

Gwen Borlaug, MPH, CIC  
Director, Healthcare-Associated Infections Prevention Program  
Wisconsin Division of Public Health  
Madison  

Jeffrey P. Davis, MD, FIDSA  
Chief Medical Officer and State Epidemiologist for Communicable Diseases  
Wisconsin Division of Public Health  
Madison  

Jon C. Gould, MD, FACS (National Surgical Quality Improvement Program Champion)  
Alonzo P. Walker Professor in General Surgery  
Chief, Division of General Surgery  
Medical College of Wisconsin  
Milwaukee  

Michael Roskos, MD, FACS (National Surgical Quality Improvement Program Champion)  
General Surgery  
Franciscan Healthcare in La Crosse - Mayo Clinic Health System  
La Crosse  

Gary R. Seabrook, MD, FACS (Perioperative Surgical Champion)  
Professor and Chief, Division of Vascular Surgery  
Medical College of Wisconsin  
Senior Medical Director, Surgical Services  
Froedtert Hospital  
Milwaukee  

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Introduction

Surgical site infections (SSIs) are the most frequently reported healthcare-associated infection (HAI) in Wisconsin, and approximately 900 SSIs were reported annually to the Wisconsin Division of Public Health (DPH) during 2013-2015. About 1.5 percent of surgical procedures performed in Wisconsin are complicated by an SSI, and the Centers for Disease Control and Prevention (CDC) reports that mortality associated with SSIs is as high as 25 percent nationally. Furthermore, the fiscal burden of these adverse events can approach $10 billion annually in the United States.¹⁻⁵

During 2017, the CDC will finalize the draft Healthcare Infection Control Practices Advisory Committee (HICPAC) Guidelines for the Prevention of SSIs (HICPAC SSI Prevention Guidelines), which will be the first update since publication of the 1999 SSI prevention guidelines.⁶ Because the evidence on which the HICPAC SSI Prevention Guidelines are based is limited to randomized controlled trials published prior to 2015, DPH determined that supplemental guidance incorporating current evidence-based data from well-designed laboratory studies, prospective cohort clinical studies, case-control studies, randomized controlled trials, systematic reviews, and meta-analyses was necessary to provide surgical teams with the most recent and relevant SSI prevention strategies available.

The 2017 Wisconsin Division of Public Health Supplemental Guidance for the Prevention of Surgical Site Infections: An Evidence-Based Perspective (WDPH SSI Prevention Guidance) was written by a statewide panel of content experts and was reviewed by three distinguished national and international surgical care experts. This guidance is intended to enhance, not replace, the HICPAC SSI Prevention Guidelines. DPH recommends that surgical teams follow the HICPAC SSI Prevention Guidelines, but the WDPH SSI Prevention Guidance supersedes the HICPAC SSI Prevention Guidelines in areas where the WDPH SSI Prevention Guidance provides stronger, more current evidence for certain SSI prevention interventions.

The HICPAC SSI Prevention Guidelines contain two sections. The Core Section describes recommendations that should be applied to all surgical procedures, and addresses five specific content areas: antimicrobial prophylaxis (AMP), glycemic control, normothermia, oxygenation, and antiseptic prophylaxis.
The Prosthetic Joint Arthroplasty Section contains additional recommendations for these frequently performed procedures that can result in SSIs causing significant human and economic burden. This section addresses blood transfusion, systemic immunosuppressive therapy, intra-articular corticosteroid injection, anticoagulation, orthopedic space suits, and biofilms. Each topic in the two sections of the HICPAC SSI Prevention Guidelines was graded according to the strength of evidence described in the table below.

**Table I. CDC SSI Guidelines Evidence-Based Criteria Grade**

<table>
<thead>
<tr>
<th>Category</th>
<th>Description</th>
</tr>
</thead>
<tbody>
<tr>
<td>Category IA</td>
<td>A strong recommendation supported by high- to moderate-quality evidence suggesting net clinical benefits or harms.</td>
</tr>
<tr>
<td>Category IB</td>
<td>A strong recommendation supported by low-quality evidence suggesting net clinical benefits or harms, or an accepted practice, supported by low- to very low-quality evidence.</td>
</tr>
<tr>
<td>Category IC</td>
<td>A strong recommendation required by state or federal regulation.</td>
</tr>
<tr>
<td>Category II</td>
<td>A weak recommendation supported by any quality evidence suggesting a tradeoff between clinical benefits and harms.</td>
</tr>
<tr>
<td>No recommendation/unresolved issue</td>
<td>An unresolved issue for which there is either low- to very low-quality evidence with uncertain tradeoffs between benefits and harms or no published evidence on outcomes deemed critical to weighing the risks and benefits of a given intervention.</td>
</tr>
</tbody>
</table>

The HICPAC SSI Prevention Guidelines and strength of evidence for each recommendation are included in this document, and are followed by the WDPH SSI Prevention Guidance with the corresponding evidence-based references validating the recommendations. The WDPH SSI Prevention Guidance also addresses the evidence supporting staphylococcal surveillance and decolonization, and implementation of a surgical care bundle. Neither of these topics is included in the HICPAC SSI Prevention Guidelines.

**Introduction Citations**


6. CDC-HICPAC Guidelines – No Citation Yet


**Wisconsin Division of Public Health Supplemental Guidance for the Prevention of Surgical Site Infections: An Evidence-Based Perspective**  
**January 2017**

### Core Considerations

## Interventions for all surgical procedures

### Antimicrobial Prophylaxis

<table>
<thead>
<tr>
<th>HICPAC SSI Prevention Guidelines</th>
<th>WDPH SSI Prevention Guidance</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Administer preoperative antimicrobial agents only when indicated, based on published clinical practice guidelines (<strong>Category 1B</strong>).</td>
<td>1. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>2. Administer the appropriate parenteral prophylactic antimicrobial agent prior to skin incision in all cesarean sections (<strong>Category 1A</strong>).</td>
<td>2. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>3. Adjust the prophylactic antimicrobial agent dose based on the patient’s weight in obese and morbidly obese patients (<strong>Category 1B</strong>).</td>
<td>3. Follow the 2013 American Society of Health-System Pharmacists (ASHP) guidelines for antimicrobial prophylaxis in surgery. Administer prophylactic antibiotic agents based on the patient’s Body Mass Index (BMI) or the patient’s weight in kilograms. For example, patients with a BMI &lt;30 (or &lt;120 kg) should receive 2 grams of a beta-lactam agent; patients with a BMI ≥ 30 (or &gt;120 kg) should receive 3 grams.</td>
</tr>
<tr>
<td>4. No recommendation can be made regarding the safety and effectiveness of intraoperative re-dosing of parenteral prophylactic antimicrobial agents for the prevention of SSI (<strong>No recommendation/unresolved issue</strong>).</td>
<td>4. Base re-dosing of antibiotic agents on the drug half-life and duration of surgery.</td>
</tr>
<tr>
<td>5. In clean and clean-contaminated procedures, do not administer additional prophylactic antimicrobial agent doses after the surgical incision is closed in the operating room, even in the presence of a drain (<strong>Category 1A</strong>).</td>
<td>5. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>HICPAC SSI Prevention Guidelines</td>
<td>WDPH SSI Prevention Guidance</td>
</tr>
<tr>
<td>----------------------------------</td>
<td>-------------------------------</td>
</tr>
<tr>
<td>6. This issue not addressed.</td>
<td>6. Include preoperative oral antibiotics in combination with mechanical bowel preparations (OA-MBP) as a safe and effective adjunctive strategy for reducing the risk of infection following colorectal surgery. Current peer-reviewed evidence indicates that OA-MBP should be part of a comprehensive colorectal surgical care bundle. 10-14</td>
</tr>
</tbody>
</table>

Antimicrobial Prophylaxis Citations


**Glycemic Control**

<table>
<thead>
<tr>
<th>HICPAC SSI Prevention Guidelines</th>
<th>WDPH SSI Prevention Guidance</th>
</tr>
</thead>
</table>
| 1. Implement perioperative glycemic control and blood glucose target levels of <200 mg/dl in diabetic and non-diabetic surgical patients (Category 1A). | 1. Maintain a mean perioperative blood glucose level <200 mg/dl in diabetic and non-diabetic surgical patients.  
15,16 |
| 2. No recommendation can be made regarding the safety and effectiveness of lower or narrower blood glucose target levels and SSI (No recommendation/unresolved issue). | 2. Avoid increased risk of hypoglycemic events and increased mortality associated with tight glycemic control (81 to 108 mg/dl).  
17,18 |
| 3. No recommendation can be made regarding hemoglobin A1C target levels and risk of SSI in diabetic and non-diabetic patients (No recommendation/unresolved issue). | 3. Maintain hemoglobin A1C level <6.7. This has been shown to minimize postoperative infectious complications in surgical patients.  
19,20 |

**Glycemic Control Citations**


Normothermia

<table>
<thead>
<tr>
<th><strong>HICPAC SSI Prevention Guidelines</strong></th>
<th><strong>WDPH SSI Prevention Guidance</strong></th>
</tr>
</thead>
<tbody>
<tr>
<td>1. Maintain perioperative normothermia <em>(Category 1A).</em></td>
<td>1. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>2. No recommendation can be made regarding the safety or effectiveness of strategies to achieve and maintain normothermia, the lower limit of normothermia, or the optimal timing and duration of normothermia <em>(No recommendation/unresolved issue).</em></td>
<td>2. Consider use of forced-air warming (FAW) to reduce incidence of SSIs.</td>
</tr>
<tr>
<td></td>
<td>Based on 67 trials (45 of which were randomized controlled trials) with 5,438 participants, a Cochrane Collaboration found that FAW reduced incidence of SSIs and complications among patients undergoing abdominal surgery.(^{21}) It was also beneficial in preventing major cardiovascular complications in patients with substantial cardiovascular disease.(^{21}) It has been suggested that use of FAW in laminar air flow operating rooms during orthopedic procedures may pose a risk for intraoperative wound contamination, however, there are no definitive clinical studies suggesting that FAW increases the risk of postoperative surgical site infections.(^{22,23}) Normothermia should be maintained in the preoperative, intraoperative and in the postoperative environment.(^{24})</td>
</tr>
</tbody>
</table>

**Normothermia Citations**


## Oxygenation Guidelines

<table>
<thead>
<tr>
<th>HICPAC SSI Prevention Guidelines</th>
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</thead>
<tbody>
<tr>
<td>1. For patients with normal pulmonary function undergoing general anesthesia with endotracheal intubation, administer an increased fraction of inspired oxygen (FiO2) both intraoperatively and post-extubation in the immediate postoperative period. To optimize tissue oxygen delivery, maintain perioperative normothermia and adequate volume replacement (Category IA).</td>
<td>1. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>2. No recommendation can be made regarding the safety and effectiveness of administering a perioperative increased fraction of inspired oxygen (FiO2) for the prevention of SSI in patients with normal pulmonary function undergoing either general anesthesia without endotracheal intubation or neuraxial anesthesia (i.e., spinal, epidural, or local nerve blocks) (No recommendation/unresolved issue).</td>
<td>2. Consider use of high oxygen supplementation as an SSI risk reduction strategy during colorectal procedures. The use of high oxygen supplementation as an SSI risk reduction strategy is controversial. However, oxygen supplementation (80% FiO₂) during the perioperative period has been documented to reduce the risk of SSI in patients undergoing colorectal surgeries.²⁵,²⁶ In heterogeneous patient populations comprised of abdominal, gynecological, breast-related or bariatric patient populations, supplemental oxygen administration demonstrated no SSI reduction benefit.²⁷-²⁹</td>
</tr>
<tr>
<td>3. No recommendation can be made regarding the safety and effectiveness of administering increased fraction of inspired oxygen (FiO2) via facemask or nasal cannula only during the postoperative period for the prevention of SSI in patients with normal pulmonary function (No recommendation/unresolved issue).</td>
<td></td>
</tr>
<tr>
<td>4. No recommendation can be made regarding the optimal target level, duration, and delivery method of the fraction of inspired oxygen (FiO2) for the prevention of surgical site infection (No recommendation/unresolved issue).</td>
<td></td>
</tr>
</tbody>
</table>

## Oxygenation Citations


# Antisepsis Prophylaxis

<table>
<thead>
<tr>
<th>HICPAC SSI Prevention Guidelines</th>
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<tbody>
<tr>
<td>1. Perform intraoperative skin preparation with an alcohol-based antiseptic agent, unless contraindicated (Category IA).</td>
<td>1. Use 2% chlorhexidine gluconate (CHG) with 70% alcohol as the preferred intraoperative skin preparation agent. CHG is also a safe and effective antiseptic agent for obstetrical and gynecologic procedures.</td>
</tr>
<tr>
<td>2. Advise patients to shower or bathe (full body) with either soap (antimicrobial or non-antimicrobial) or an antiseptic agent on at least the night before the operative day (Category IB).</td>
<td>2. Ensure that all patients undergoing elective surgical procedures involving skin incisions undergo a standardized preadmission shower/cleansing with 4% aqueous or 2% (cloth coated) CHG.</td>
</tr>
<tr>
<td>3. No recommendation can be made regarding the optimal timing of the preoperative shower or bath, the total number of soap or antiseptic agent applications, or the use of chlorhexidine gluconate washcloths for the prevention of SSI (No recommendation/ unresolved issue).</td>
<td>3. Standardize the preadmission shower or cleansing process according to the protocols below. Recent randomized controlled trials have documented that high skin surface concentrations of CHG can be obtained by standardization of the preadmission shower or cleansing process using 4% aqueous chlorhexidine gluconate (CHG) or 2% CHG coated on a disposable polyester cloth.</td>
</tr>
</tbody>
</table>

### 4% Aqueous CHG Shower Protocol
- Remind patients to perform the CHG shower regimen with a text message, email, or voicemail.
- Provide patients with both oral and written instructions regarding the standardized CHG shower regimen.
- Instruct patients to take two showers, one the evening before surgery, and one the morning of surgery.
- Instruct patients to pause for one minute after applying the CHG and before rinsing.
- Ensure patients use a total volume of 4 oz. of CHG for each shower.

### 2% CHG Polyester Cloth Cleansing
- Remind patients to perform the CHG shower regimen with a text message, email, or voicemail.
- Provide patients with both oral and written instructions regarding the standardized CHG cloth cleansing, emphasizing gentle application of the cloths to the skin.
<table>
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<tr>
<td>- Instruct patients to use a total of 12 cloths per cleansing—6 cloths the night before surgery, and another 6 cloths the morning of surgery. Ensure patients understand they should use both sides of the cloth to maximize release of the CHG onto the skin.</td>
<td>4. Consider use of intraoperative irrigation with aqueous 0.05% CHG. Current laboratory and animal studies suggest that aqueous 0.05% CHG is an effective intraoperative wound irrigation solution for reducing the risk of SSI. A prospective, randomized controlled trial is currently underway to assess the clinical efficacy of 0.05% CHG intraoperative irrigation for open laparotomies.</td>
</tr>
<tr>
<td>4. Consider intraoperative irrigation of deep or subcutaneous tissues with aqueous iodophor solution for the prevention of SSI. Intra-peritoneal lavage with aqueous iodophor solution in contaminated or dirty abdominal procedures is not necessary (Category II).</td>
<td>4. Consider use of intraoperative irrigation with aqueous 0.05% CHG.</td>
</tr>
<tr>
<td>5. No recommendation can be made regarding the safety and effectiveness of soaking prosthetic devices in antiseptic solutions prior to implantation for the prevention of SSI (No recommendation/unresolved issue).</td>
<td>5. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>6. Use of plastic adhesive drapes with or without antimicrobial properties, is not necessary for the prevention of SSI (Category II).</td>
<td>6. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>7. No recommendation can be made regarding the safety and effectiveness of repeat application of antiseptic agents to the patient’s skin immediately prior to closing the surgical incision to prevent SSIs (No recommendation/unresolved issue).</td>
<td>7. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>8. Consider use of triclosan-coated sutures to prevent SSIs (Category II).</td>
<td>8. Use triclosan-coated antimicrobial sutures to close surgical wounds. All surgical wounds are contaminated at the time of closure. The risk of infection is related to several comorbid factors, including presence of a foreign body (e.g., necrotic tissue, hematin and sutures) in the wound at closure. Triclosan-coated sutures have been clinically shown to be safe for wound closure in adult and pediatric populations. Triclosan-coated sutures are effective against both Gram-positive and Gram-negative surgical wound pathogens. Several recent clinical trials, systematic reviews, and</td>
</tr>
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</table>


<table>
<thead>
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</tr>
</thead>
<tbody>
<tr>
<td>meta-analysis have determined that the use of triclosan antimicrobial sutures for closure of surgical wounds represents Category 1 clinical evidence in prevention of SSI.</td>
<td></td>
</tr>
</tbody>
</table>

Recommendations for the use of triclosan-coated sutures for wound closure are also included in the 2016 World Health Organization Global guidelines on the prevention of surgical site infection and the American College of Surgeons and Surgical Infection Society: Surgical Site Infection, 2016 Update.

Two recent meta-analyses and one clinical study have suggested that use of staples for wound closure is associated with an increased risk of wound complication, including infection in selective surgical disciplines (orthopedic and obstetrical). Although further studies are warranted to validate this risk, clinicians should be aware of the current clinical findings when considering wound closure.

### Antisepsis Prophylaxis Citations


## Interventions for Prosthetic Joint Arthroplasty

<table>
<thead>
<tr>
<th>HICPAC SSI Prevention Guidelines</th>
<th>WDPH SSI Prevention Guidance</th>
</tr>
</thead>
<tbody>
<tr>
<td>1. No recommendation can be made regarding the perioperative management of blood transfusions for the prevention of SSI after prosthetic joint arthroplasty <em>(No recommendation/unresolved issue).</em></td>
<td>1. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>2. Do not withhold transfusion of necessary blood products from surgical patients as a means to prevent SSI <em>(Category IB).</em></td>
<td>2. Balance the risk of complications from post-operative anemia with the potential increased risk of SSI following administration of blood products. Although some studies suggest that perioperative blood transfusion is associated with increased risk of SSI after selective pediatric and adult surgical procedures, this risk should be balanced with the undesirable complication of postoperative anemia.</td>
</tr>
<tr>
<td>3. No recommendation can be made regarding the perioperative management of systemic corticosteroid or other immunosuppressive therapy for the prevention of SSI after prosthetic joint arthroplasty <em>(No recommendation/unresolved issue).</em></td>
<td>3. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>4. No recommendation can be made regarding the management of preoperative intra-articular corticosteroid injection for the prevention of SSI after prosthetic joint arthroplasty <em>(No recommendation/unresolved issue).</em></td>
<td>4. No difference in guidance recommendation. The concern that intra-articular steroid injection for postoperative pain management is a risk factor for SSI is at present controversial. However, the risk may be influenced by the presence of co-morbid risk factors; further studies are warranted.</td>
</tr>
<tr>
<td>5. No recommendation can be made regarding the perioperative management of venous thromboembolism prophylaxis for the prevention of SSI in prosthetic joint arthroplasty <em>(No recommendation/unresolved issue).</em></td>
<td>5. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>6. No recommendation can be made regarding the safety and effectiveness of orthopedic space suits or the health care personnel who should wear them for the prevention of SSI after prosthetic joint arthroplasty <em>(No recommendation/unresolved issue).</em></td>
<td>6. No difference in guidance recommendation.</td>
</tr>
<tr>
<td><strong>HICPAC SSI Prevention Guidelines</strong></td>
<td><strong>WDPH SSI Prevention Guidance</strong></td>
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<tr>
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</tr>
<tr>
<td>7. In prosthetic joint arthroplasty, clean and clean-contaminated procedures, do not administer additional prophylactic antimicrobial agent doses after the surgical incision is closed in the operating room, even in the presence of a drain <em>(Category IA)</em>.</td>
<td>7. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>8. No recommendation can be made regarding the safety and effectiveness of cement modifications and the prevention of biofilm formation or SSI in prosthetic joint arthroplasty <em>(No recommendation/ unresolved issue)</em>.</td>
<td>8. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>9. No recommendation can be made regarding the safety and effectiveness of prosthesis modifications for the prevention of biofilm formation or SSI after prosthetic joint arthroplasty <em>(No recommendation/unresolved issue)</em>.</td>
<td>9. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>10. No recommendation can be made regarding the safety and effectiveness of vaccines for the prevention of biofilm formation or SSI after prosthetic joint arthroplasty <em>(No recommendation/unresolved issue)</em>.</td>
<td>10. No difference in guidance recommendation.</td>
</tr>
<tr>
<td>11. No recommendation can be made regarding the safety and effectiveness of biofilm control agents such as biofilm dispersants, quorum-sensing inhibitors, or novel antimicrobial agents for the prevention of biofilm formation or SSI after prosthetic joint arthroplasty <em>(No recommendation/unresolved issue)</em>.</td>
<td>11. No difference in guidance recommendation.</td>
</tr>
</tbody>
</table>

**Blood Transfusion Citations**


Intraarticular Corticosteroid Injection Citations


General Comments Regarding Biofilms and SSIs

The global impact of SSIs on healthcare systems is considerable and it has been estimated that as many as 80% of SSIs may be related to the formation of a microbial biofilm. Biofilm-mediated infections exhibit resistance to host defenses and often contribute to an excessive or inappropriate local inflammatory response. This leads to complement activation and formation of immune complexes, which in turn lead to tissue injury. Unfortunately, the incidence of biofilm-associated SSIs is likely to increase because of the expanding use of implanted medical devices. Although investigators are currently focusing on biofilm-resistant polymers and other surface coatings that discourage microbial attachment, these efforts are in the initial stages and are unlikely to significantly alter SSI risk during the immediate future. Prevention of intraoperative contamination offers the greatest benefit for patients receiving an implantable medical device. Therefore, meticulous surgical technique, use of perioperative care bundles and awareness of the various possible avenues of intraoperative contamination that can occur at the time of implantation are rational strategies for improving surgical patient outcomes.

Finally, every institution should have specific policies and procedures in place for the management, sterilization, storage, and handling of biomedical devices prior to surgical implantation.

Biofilm Citations


Interventions Omitted from Consideration in the HICPAC SSI Prevention Guidelines

Although staphylococcal surveillance and use of surgical care bundles are not included in the HICPAC SSI Prevention Guidelines, members of the WDPH SSI Prevention Expert Panel recommend these strategies in addition to the interventions described above, as part of a comprehensive surgical care improvement program.

**Staphylococcal Surveillance**

Results of several published studies suggest that suppression of the methicillin-sensitive *Staphylococcus aureus* (MSSA) and methicillin-resistant *S. aureus* (MRSA) carrier state is effective in reducing the occurrence of SSIs caused by these surgical wound pathogens. Nasal mupirocin (twice daily for 5 to 7 days) with a minimum of two 4% aqueous CHG showers has been widely used for the suppression of nasal carriage of MSSA and MRSA. Although mupirocin has been viewed as the “gold standard” for suppressing staphylococci in the nares, the suppression of organisms in the nares on the morning of surgery using a swab coated with 5% or 10% povidone iodine (0.5% available iodine) has been shown to be an effective alternative. Considering the current evidence-based literature, the following are justified:

a. Selection of an efficacious (risk-reducing, cost effective) active screening strategy should be based on the relative risk of MSSA or MRSA healthcare-associated infections among “at risk” surgical patients.

b. In the absence of targeted or universal screening, routine topical mupirocin or systemic antimicrobial agents is not currently recommended for the suppression of MSSA or MRSA carriage among surgical patients.

c. In the case of targeted screening, preoperative suppression may be considered for MSSA and MRSA colonized patients undergoing “at risk” surgical procedures, such as cardiovascular and vascular procedures with implantation of prosthetic grafts and orthopedic total joint procedures. The benefit of targeted screening and preoperative suppression in other device-related surgical procedures (i.e., implantation of neurosurgical hardware, hernia repair with mesh, etc.) is unknown and currently not supported by data.
d. Although the optimal suppression regimen is unclear, the following is recommended: a standardized regimen of topical nasal mupirocin (twice a day for 5-7 days) or an alternative approach involving the use of a nasal swab containing 5% or 10% povidone iodine applied to the nares 1 to 2 hours prior to surgery, along with a 2% or 4% chlorhexidine gluconate body cleansing/shower (once a day for 2 days) prior to surgical admission.

**Staphylococcal Surveillance Citations**


Surgical Care Bundles (SCB)

Recent peer-reviewed literature has documented the benefit of combining selective evidence-based interventional practices to form a comprehensive surgical care bundle for reducing the risk of postoperative infections. Surgical care bundles have been developed for colorectal, cardiothoracic, OB/GYN, vascular, and orthopedic procedures. SCBs should be developed in collaboration with the surgical team (surgeons and OR nursing), infection preventionists and pharmacy personnel. Implementation of a SCB requires close monitoring to ensure 100 percent compliance, because poor compliance diminishes the preventive benefits of the SCB.

Table II. Selective elements of the surgical care bundle from the evidence-based literature

<table>
<thead>
<tr>
<th>Appropriate antimicrobial prophylaxis</th>
<th>Antimicrobial (triclosan) sutures</th>
</tr>
</thead>
<tbody>
<tr>
<td>Weight-based dosing</td>
<td>Smoking cessation</td>
</tr>
<tr>
<td>Glycemic control</td>
<td>Staphylococcal surveillance (cardiothoracic and orthopedic procedures)</td>
</tr>
<tr>
<td>Normothermia</td>
<td>Oral antibiotics plus mechanical bowel preparation (colorectal)</td>
</tr>
<tr>
<td>Appropriate hair removal</td>
<td>Minimally invasive surgery</td>
</tr>
<tr>
<td>Supplemental O₂ (colorectal procedures)</td>
<td>Short duration of surgery</td>
</tr>
<tr>
<td>Use of wound edge protectors</td>
<td>Glove change prior to fascia and skin closure</td>
</tr>
<tr>
<td>Dedicated wound closure tray for fascia and skin</td>
<td>Limit traffic in the operating room</td>
</tr>
<tr>
<td>Pre-operative 4% CHG shower or 2% CHG cleansing</td>
<td>CHG cleansing of surgical wound</td>
</tr>
<tr>
<td>70% alcohol with 2% CHG perioperative skin preparation</td>
<td>Keep sterile dressing intact for first 48 hours</td>
</tr>
</tbody>
</table>
Surgical Care Bundles Citations


