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Impact of universal disinfectant cap implementation on central line–associated bloodstream infections

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Central line
Bloodstream
Compliance
Cost
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Infection prevention

Background: Central line–associated bloodstream infections (CLABSIs) result in increased length of stay, cost, and patient morbidity and mortality. One CLABSI prevention method is disinfection of intravenous access points. The literature suggests that placing disinfectant caps over needleless connectors decreases CLABSI risk.

Methods: A quasi-experimental intervention study was conducted in a >430-bed trauma I center. In addition to an existing standard central line bundle, a new intervention consisting of a luer-lock disinfectant cap with 70% alcohol was implemented in all intravenous (IV) needleless connectors on patients with peripheral and central lines. Compliance to the disinfectant cap was monitored weekly. A generalized linear model using a Poisson distribution was fit to determine if there were significant relationships between CLABSIs and disinfectant cap use. Impacts on costs were also examined.

Results: The rate of CLABSI decreased following implementation of the disinfectant cap. The incidence rate ratios (.577, \(P = .004\)) for implementing the disinfectant caps was statistically significant, indicating that the rate of patient infections decreased by >40%. Increased compliance rates were associated with lower infection rates. Disinfectant cap use was associated with an estimated savings of almost $300,000 per year in the hospital studied.

Conclusions: Use of a disinfectant cap on IV needleless connectors in addition to an existing standard central line bundle was associated with decreased CLABSI and costs.

The magnitude of central line–associated bloodstream infections (CLABSIs) in the United States and abroad is staggering. It is estimated that 200,000–400,000 episodes of bloodstream infections occur annually in U.S. hospitals, resulting in increased length of stay, cost, and patient morbidity and mortality.1 Nearly 1 in every 10 hemodialysis catheters fail each month as a result of CLABSI, and the numbers are even greater in nontunneled catheters used in the acute care setting. Researchers report that mortality rates are between 12% and 25% from CLABSIs and estimate that annual costs associated with treatment exceed $2 billion.2,3

The goal of hospital infection prevention programs is to eliminate CLABSI or decrease it as much as feasible given the patient population.2 Methods to reduce CLABSI consist of implementing several techniques referred to as a bundle of interventions that specify recommendations for proper insertion and appropriate handling of central lines.4 These multimodal approaches have been successful in decreasing CLABSI.5,6

One specific aspect of the central line bundle is to minimize infection from intravenous (IV) access points. Most IV tubing contains several needleless connectors for the purposes of medication delivery and blood draws. Nurses access these needleless connectors several times a day, potentially increasing the possibility of contamination and subsequent infection.

To prevent infection in patients with IV access devices, the Centers for Disease Control and Prevention strongly recommend appropriate disinfectant of needleless connectors prior to access.2
Although time and friction are recommended for disinfection, there are no specific recommendations on the amount of time needed to reach optimal disinfection (e.g., 10, 15, or 30 seconds). Additionally, some argue that proper disinfection practices are difficult to maintain given the increased work load of nurses.7

In a 2006 in vitro study, researchers reported that the use of 70% alcohol prior to needleless connector entry is not sufficient protection against microbial contamination; therefore, the use of a disinfectant cap was recommended.8 The literature suggests that a disinfectant cap placed over IV needleless connectors decreases colonization on the connector, therefore lowering the risk of CLABSI.9 Another quality improvement study found that the use of a closed luer-lock disinfectant cap significantly decreased hemodialysis catheter infections in pediatric patients.10 In a more recent study, the implementation of a disinfectant cap decreased line contamination, bacterial density, and CLABSI rates.11 Based on these promising results, additional research is needed to test the generalizable effectiveness of widespread implementation of a disinfectant cap to decrease or eliminate CLABSI.

The purpose of this study is to analyze the effect of universal IV needleless connector disinfectant cap implementation on the rate and type of CLABSI and estimated costs in a large tertiary care center using a standard central line bundle. Additionally, the relationship between disinfectant cap compliance and CLABSI rates is explored.

METHODS

Setting and design

This quasi-experimental short interrupted time series12 intervention study was conducted in a 430-bed tertiary care trauma I center in the U.S. Mountain West. A luer-lock disinfectant cap with 70% alcohol was implemented in all patients (newborn to adults) with peripheral and central lines residing on 13 inpatient units at 1 hospital beginning in January 2012. Excluded from this study were patients in the emergency department, ambulatory care, surgical services, labor and delivery, and well-baby nursery and patients who were postpartum.

Intervention

The disinfectant cap (Curos Disinfecting Port Protector, Curos, San Diego, CA) is a plastic-threaded device that contains 70% isopropyl alcohol. This device received a 510(k) clearance from the U.S. Food and Drug Administration for 1-time use.13 It is effective within 3 minutes of application and may be used up to 7 days.

Prior to implementation, the hospital product review committee and the institutional review board approved the use of the product for this research study. Nursing staff were introduced to the proper use of the cap through onsite training by the vendor, an education fact sheet, or required online training. Staff also participated in individual 1-on-1 follow up.

In this study, the disinfectant cap was placed universally on all IV needleless connectors (central, peripheral, and IV tubing) when the connectors were not in use. Compliance to the disinfectant cap was monitored weekly and reported to each unit to encourage use of the disinfectant cap.

Data collection and analysis

Compliance

Following implementation of the disinfectant cap, compliance was determined by audits conducted 1-2 times per week beginning in February 2012 and lasting throughout the study period. The number of disinfectant caps present was divided by the number of total available needleless connectors to result in an overall compliance rate per central line patient. The data were then aggregated to the nursing department level and reported to each manager as a weekly nursing department compliance rate.
CLABSI

The presence of CLABSI was defined as “a primary laboratory confirmed bloodstream infection in a patient with a central line at the time of (or within 48 hours prior to) the onset of symptoms and the infection is not related to an infection from another site.” All positive blood cultures in the hospital were reviewed to determine if they met the definition for CLABSI. The rate of CLABSI was calculated per 1,000 central line catheter days.

Impact of CLABSI

Cost, estimated case fatality, and additional ICU length of stay were estimated to represent the impact of CLABSI. Cost of CLABSI was estimated at $25,000 per case, a low estimate based on the literature, which estimates CLABSI results at $25,000–$55,000 per incident. Estimated case fatality was based as 6%, with 4 additional days in the hospital; both are low estimates from the CDC literature. The number of CLABSI cases was multiplied by the estimated cost, mortality, and length of stay before and after implementation of the disinfectant cap. The cost of implementation supplies (disinfectant cap and no-port tubing) were also taken into consideration.

Analysis

The rate of CLABSI and costs were compared for 12 months before (2011) and after the intervention (2012). Data were analyzed using IBM SPSS version 22.0 (SPSS Inc, Chicago, IL). A generalized linear model using a Poisson distribution was fit to determine if there was a significant difference in CLABSI rates following implementation of the disinfectant cap. The model was adjusted for the number of line days per patient.

RESULTS

The rate of CLABSI per 1,000 central line days decreased following implementation of the disinfectant cap (before implementation: mean ± SD, 1.5 ± .37) (after implementation: mean ± SD, .88 ± .62) (Fig 1). The robust estimator option was used as the deviance statistic (.679) and was somewhat <1 as assumed when using the Poisson distribution. The incidence rate ratio (IRR = .93; 95% confidence interval, .88–.97), as seen in Table 1. In addition, during implementation, nurses expressed frustration that uncovered needleless connectors high in the IV tubing that were never used were counted against compliance rates. As a result of this feedback, the hospital purchased no-port tubing, further protecting the patient from potential entries into the system. Creating a feedback loop proved to be an important strategy for increasing compliance, thereby preventing infection as is supported by the quality improvement plan-do-check-act cycle.

Another important finding in this study is the relationship with implementation of a disinfectant cap and a significant decrease in CLABSI rates. These findings are timely because they support the growing body of knowledge that disinfectant caps are one method to prevent CLABSI. This study also showed that disinfectant caps are one aspect of aseptic techniques for preventing CLABSI that is easier to monitor for compliance than other infrequent procedures (eg, dressing change, line insertion) that are not readily apparent with visual surveillance.

The treatment of CLABSI varies by patient; however, most clinicians agree that CLABSI results in an increased hospital length of stay and, in some cases, death. This study used low estimations for length of stay and mortality; even so, implementation of the disinfectant cap might have decreased hospital stays by 1 day per week and eliminated 1 death during the study period. Using the CLABSI opportunity estimator published by Johns Hopkins Quality Safety Research Group would have doubled these numbers.

Although improved patient outcomes alone should be a sufficient motivator to prevent health care–acquired infections, it is often the financial impact that receives the most attention. In a comparative study, the costs of hospitalized patients who acquired CLABSI were matched with similar patients who did not experience CLABSI. They concluded that the occurrence of CLABSI resulted in a significant impact on hospital operating costs.

When infections are prevented, the impact is calculated in terms of cost avoidance. After considering the costs of purchasing disinfectant caps and no-port tubing, this current study also identified a significant cost avoidance by decreasing CLABSI. Cost avoidance is considered soft money and can be difficult to track and justify. It is important to emphasize cost avoidance when considering the investment in the purchase of disinfectant caps and no-port tubing. This study was successful in part because of the support of hospital administration and the cooperation of the hospital products committee who understood the overall patient safety ramifications and cost savings.

<table>
<thead>
<tr>
<th>Table 1</th>
<th>Disinfectant cap intervention, CLABSI rates, and compliance parameter estimates</th>
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<tbody>
<tr>
<td>Model</td>
<td>Parameter</td>
</tr>
<tr>
<td>CLABSI</td>
<td>Intercept</td>
</tr>
<tr>
<td></td>
<td>Intervention</td>
</tr>
<tr>
<td>CLABSI</td>
<td>Intercept</td>
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<tr>
<td>Compliance</td>
<td>Compliance</td>
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</tbody>
</table>

*Statistically significant result.

Table 2 | Effect of disinfectant cap on central line—associated bloodstream infection costs |
<table>
<thead>
<tr>
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</thead>
<tbody>
<tr>
<td>Item</td>
<td>Estimated costs in 2011 (before implementation)</td>
</tr>
<tr>
<td>Supplies</td>
<td>NA</td>
</tr>
<tr>
<td>Annual BSI cost (estimated at $25,000 per episode)</td>
<td>$1,050,000</td>
</tr>
<tr>
<td>Totals</td>
<td>$1,050,000</td>
</tr>
</tbody>
</table>

BSI, bloodstream infection; NA, not applicable.
Disinfection of the IV needleless connectors is only one aspect of CLABSI prevention. Other aspects of prevention (eg, insertion, dressing changes) are important to consider for those striving for zero infection rates. Unfortunately, compliance to these important aseptic techniques might be more difficult to measure than compliance to a disinfectant cap.

LIMITATIONS

There are some limitations to this study. Ongoing education was implemented simultaneously by the hospital, which might have affected the CLABSI rates. Further, use of the disinfectant cap may have resulted in an increased vigilance to compliance to the central line bundle, which was not measured as part of the study. Cost estimates were based on projections reported in the literature. Although costs were about 50% lower than other reports, they might not reflect true costs.

CONCLUSION

CLABSI is a serious, preventable, health care–acquired infection. This study found a relationship between implementation of a disinfectant cap and reduced rates of CLABSI, cost, length of stay, and mortality. As this increasing body of knowledge becomes available, infection preventionists might need to consider that some time honored traditions (eg, scrub the hub) should now be replaced with new product technology. Further, this study found success in implementation of a quality improvement feedback loop and found that compliance rates resulted in prevention of CLABSI. This improvement model might prove successful in other infection prevention campaigns.

References