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Hospital-wide Impact of Mandatory Infectious Disease Consultation for Staphylococcus aureus Septicemia

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Introduction

Staphylococcus aureus bacteremia (SAB) is a major human pathogen that causes a wide-range of clinical infections.⁶ It is a leading cause of bacteremia and infective endocarditis, as well osteoarticular, skin and soft tissue, and device-related infections.⁶ It is also one of the main contributors to both hospital and community-onset bloodstream infections worldwide.⁵ SAB continues to grow in number and complexity, which is why it is presently associated with a 10%-30% mortality.^{1,7}

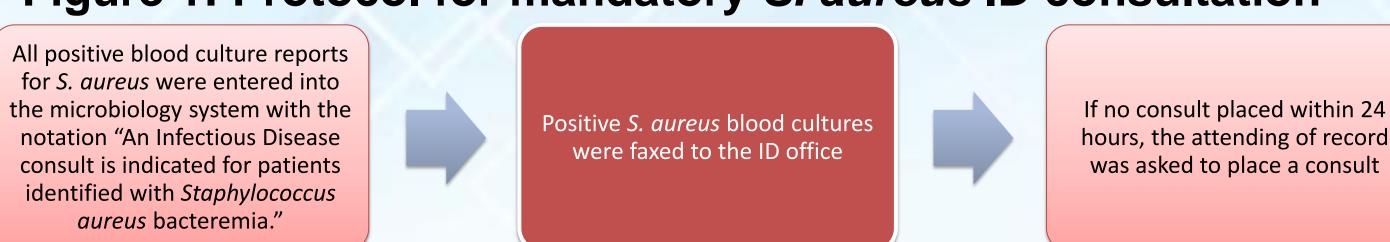
This study focused on determining if mandatory Infectious Disease (ID) consultations:

- 1) improved adherence to published guidelines on quality-of-care indicators, and
 - 2) improved outcomes of patients with Staphylococcus aureus bacteremia (SAB)

Methods

- A total of 276 patients from Lehigh Valley Cedar Crest (CC) and Muhlenberg (MHC) campuses had positive S. aureus blood cultures and met the predefined study inclusion criteria during this study.
 - 179 patients from January 1st 2013 to December 31st 2013, prior to the institution of mandatory ID SAB consultation
 - 197 patients from May 12, 2014 to May 11, 2015, the intervention period.
- Patients with positive blood cultures for methicillin-resistant or methicillin-susceptible Staphylococcus aureus (MRSA/MSSA) were identified using ICD-9 codes as well as from lists of culture results provided by the microbiology laboratory.
- The patients' medical records were reviewed for demographic data, features of *S. aureus* bacteremia, details regarding treatment, and outcomes.
- All variables were compared and analyzed using the Mann-Whitney test or t test on Minitab 17 to determine significance.

Figure 1: Protocol for mandatory S. aureus ID consultation



Results

Table 1: Characteristics of Patients with Staphylococcus aureus bacteremia

| | Pre-Intervention Period (n=179) | Intervention Period (n=197) | P-Value |
|-----------------------------|---------------------------------|--------------------------------|---------|
| Age, median years (IQR) | 64 (54-78) | 68 (52-76) | 0.937 |
| Female | 83 (46) | 65 (33) | 0.007 |
| White | 152 (85) | 180 (91) | 0.404 |
| APACHE II Score, mean | 13.4 | 13.5 | 0.995 |
| Pitt bacteremia score, mean | 1.53 | 1.26 | 0.134 |
| How-acquired | | | |
| Community-acquired | 80 (45) | 122 (62) | 0.003 |
| Hospital-acquired | 25 (14) | 12 (6) | 0.190 |
| Healthcare-associated | 74 (41) | 62 (31) | 0.104 |
| Penicillin allergy | 28 (16) | 28 (14) | 0.811 |

Figure 2: Source of *Staphylococcus aureus* Infection during the Pre-Intervention and Intervention Periods



Table 2: Infectious Diseases (ID) Consultation and Adherence to Quality-of-Care (QCI) Indicators

| | Pre-Intervention Period | Intervention Period | P-Value |
|---------------------------------------|-------------------------|---------------------|---------|
| | (n=179) | (n=197) | |
| ID Consult | 176 (98) | 193 (98) | 0.953 |
| Days to ID consult, mean | 0 (0-1) | 0 (0-1) | 0.072 |
| Follow-up blood cultures obtained | 159 (89) | 151 (77) | 0.042 |
| Early source control ⁹ | 46 (82) | 29 (85) | 0.718 |
| Echocardiography performed | 149 (83) | 182 (92) | 0.126 |
| TTE | 143 (80) | 150 (76) | 0.531 |
| TEE | 66 (37) | 108 (56) | 0.0026 |
| Treatment duration, median days | 28 (14-42) | 29 (19-42) | 0.363 |
| (IQR) | | | |
| Appropriate duration§ | 109 (72) | 152 (91) | 0.009 |
| Appropriate deescalation [±] | 66 (37) | 105 (53) | 0.012 |
| Hours to deescalation, median | 64 (42-83) | 41 (20-60) | 0.0001 |
| $(IQR)^{\circ}$ | | | |

Table 3: Outcomes of Patients with Staphylococcus aureus Bacteremia

| | Pre-Intervention Period (n=179) | Intervention Period (n=197) | P-Value |
|---|---------------------------------|--------------------------------|---------|
| ength of hospitalization, median days (IQR) | 12 (7-9) | 13 (8-21) | 0.329 |
| Attributable mortality | 28 (48) | 25 (66) | 0.149 |
| All-cause morality | 58 (32) | 38 (19) | 0.0281 |
| 14 days | 18 (31) | 16 (42) | 0.295 |
| 30 days | 8 (13) | 8 (21) | 0.510 |

Discussion

- Most of the baseline characteristics were similar between the two groups. There was a female-predominance in the pre-intervention group (46% vs 33%, p=0.007), and an increased the number of community-acquired infections in the intervention period (62% vs 45%, 0.003) (Table 1). This is most likely a result of the study's small data size.
- There was no difference in the incidence of ID consultation between the pre-intervention and intervention periods (98% vs. 98%, p=0.953), nor in days to ID consult (median, 0 (0-1) vs 0 (0-1), p=0.072, Table 2).
- There was a significant increase in the amount of SAB patients with appropriate antibiotic duration (72% vs. 91%, p=0.009) and the amount of methicillin-sensitive SAB with appropriate deescalation (p=0.012) (Table 2).
- Interestingly, there was better adherence to follow up blood cultures being obtained 48-96 hours later during the pre-intervention period versus the intervention period (89% vs. 77%, p=0.042, Table 2). This is likely secondary to the specific time frame noted and the data will be reevaluated to look for repeat blood cultures without the time constraints.
- Four patients in the intervention group never received an ID consult (Table 2). This is likely secondary to human error.
- There was an increase in the number transesophageal echocardiograms (TEE) performed (Table 2).
- There were significantly more endocarditis patients in the intervention period than there were during the pre-intervention period (28% vs 21%, p=0.009, Figure 2).
- There was no significant difference in outcome measures between the two groups for hospital length of stay, attributable mortality (48% vs 66%, p=0.149) and 14-day and 30-day mortality. However, there was a significant difference in all-cause mortality between the two cohorts (32% vs 19%, p=0.0281) (Table 3). This is likely a reflection of the short duration of follow-up with patients from the intervention period.

Future Analyses

- The mandatory ID consultation protocol is still in effect, allowing LVHN to continue improving on its adherence to quality-of-care indicators.
 - An auto-fax system to alert the ID consultants earlier to patients identified with SAB will go into effect shortly.
 - There is still a 24-hour hold in place on ID consultations which needs to be evaluated further.
- This protocol warrants future investigations into continued compliance with QCI's in SAB patients.

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