

CLABSIs & BMT: Challenges and Strategies for Prevention

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Session Objectives

- Describe current CLABSI definitions with attention to implications for patients undergoing blood and marrow transplant
- Evaluate clinical practice strategies to prevent CLABSIs
- Discuss quality improvement strategies to reduce CLABSIs

- “The very first requirement in a hospital is that it should do the sick no harm.”

- Florence Nightingale



The Cost of CLABSIs

- Significant source of morbidity and mortality
 - Incidence of CLABSIs among BMT patients as high as 7.4 /1000 catheter days
- Approximate cost: \$7,000 - \$29,000/infection
 - Annual cost up to \$2.68 billion
- Pressure for public reporting of and pay for performance incentives
 - Centers for Medicare and Medicaid Services no longer reimbursing hospitals for care required to treat CLABSIs

CLABSI Definition

- A laboratory-confirmed bloodstream infection (LCBI) where a central line was in place for >2 calendar days on the date of event, with day of device placement being Day 1, **and**
- a central line was in place on the date of the event or the day before.
 - Source: http://www.cdc.gov/nhsn/pdfs/pscmanual/4psc_clabscurrent.pdf

Defining Laboratory-Confirmed Bloodstream Infections (LCBIs)

- Criterion 1
 - Patient has a recognized pathogen cultured from one or more blood cultures **and**
 - organism cultured from blood is not related to an infection at another site
- Criterion 2
 - Patient has at least one of the following signs or symptoms: fever (>38°C), chills, or hypotension **and**
 - organism cultured from blood is not related to an infection at another site **and**
 - the same common commensal is cultured from two or more blood cultures drawn on separate occasions

Defining Laboratory-Confirmed Bloodstream Infections (LCBIs)

- Criterion 3
 - Patient \leq 1 year of age has at least one of the following signs or symptoms: fever ($>38^{\circ}\text{C}$ core), hypothermia ($<36^{\circ}\text{C}$ core), apnea, or bradycardia **and**
 - positive laboratory results are not related to an infection at another site **and**
 - the same common commensal is cultured from two or more blood cultures drawn on separate occasions

Audience Response Question

- A patient who is Day + 5 following an allogeneic hematopoietic stem cell transplant develops a fever to 38.1°C . Blood cultures are obtained from both lumens of the double lumen tunneled catheter. One of the blood cultures becomes positive for *E. coli*.
- Does this represent a CLABSI?
 - Yes
 - No

Audience Response Question

- A patient who is Day + 5 following an allogeneic hematopoietic stem cell transplant develops a fever to 38.1°C . Blood cultures are obtained from both lumens of the double lumen tunneled catheter. One of the blood cultures becomes positive for *E. coli*.
- Does this represent a CLABSI?
- Yes – Because *E. coli* is a recognized pathogen, this event represents a CLABSI.

Audience Response Question

- A patient who is Day + 5 following an allogeneic hematopoietic stem cell transplant develops a fever to 38.1°C . Blood cultures are obtained from both lumens of the double lumen tunneled catheter. One of the blood cultures becomes positive for *Staphylococcus epidermidis*.
- Does this represent a CLABSI?
 - Yes
 - No

Audience Response Question

- A patient who is Day + 5 following an allogeneic hematopoietic stem cell transplant develops a fever to 38.1°C . Blood cultures are obtained from both lumens of the double lumen tunneled catheter. One of the blood cultures becomes positive for *Staphylococcus epidermidis*.
- Does this represent a CLABSI?
- No – because *Staphylococcus epidermidis* is a common contaminant, it must be present in both cultures for the event to represent a CLABSI

Mucosal Barrier Injury Laboratory-Confirmed Bloodstream Infections (MBI-LCBI)

- First introduced in 2013
- New to 2014: required to indicate which of the underlying conditions of the MBI-LCBI criterion was met, if any
 - All CLABSI, whether LCBI or MBI-LCBI, must be reported if CLABSI is part of the institution's Monthly Reporting Plan.

Defining MBI-LCBI

- Criterion 1
 - Patient of any age meets criterion 1 for LCBI with at least one blood culture growing an MBI organism with no other organisms isolated
 - See: NHSN Organisms Lists (All Organisms, Top Organisms, Common Commensals, MBI Organisms, & Uropathogens) – accessible at <http://www.cdc.gov/nhsn/acute-care-hospital/clabsi/index.html>
- **and**

Defining MBI-LCBI

- One of the following is met:
 - Is an allogeneic hematopoietic stem cell transplant recipient within the past year with one of the following documented during same hospitalization as positive blood culture
 - Grade III or IV gastrointestinal graft-versus-host disease
 - ≥ 1 liter diarrhea in a 24-hour period (or ≥ 20 mL/kg in a 24-hour period for patients < 18 years of age) with onset on or within 7 calendar days before the date the positive blood culture was collected
- **OR**

Defining MBI-LCBI

- Is neutropenic, defined as at least 2 separate days with values of absolute neutrophil count (ANC) or total white blood cell count (WBC) < 500 cells/mm³ within a seven-day time period which includes the date the positive blood culture was collected (Day 1), the 3 calendar days before *and* the 3 calendar days after

Example MBI Organisms

- *Bacteroides* spp., *Candida* spp., *Clostridium* spp., *Enterococcus* spp., *Fusobacterium* spp., *Peptostreptococcus* spp., *Prevotella* spp., *Veillonella* spp., or Enterobacteriaceae
- Note: Refer to the current NHSN Organisms Lists (All Organisms, Top Organisms, Common Commensals, MBI Organisms, & Uropathogens)

Defining MBI-LCBI

- Criterion 2
 - Patient of any age meets criterion 2 for LCBI when the blood cultures are growing only viridans group streptococci with no other organisms isolated **and one of the following**
 - Allogeneic hematopoietic stem cell transplant criteria for MBI-LCBI criterion 1
- **OR**
- Neutropenia criteria for MBI-LCBI criterion 1

Defining MBI-LCBI

- Criterion 3
 - Patient of any age meets criterion 3 for LCBI when the blood cultures are growing only viridans group streptococci with no other organisms isolated **and one of the following**
 - Allogeneic hematopoietic stem cell transplant criteria for MBI-LCBI criterion 1
- **OR**
- Neutropenia criteria for MBI-LCBI criterion 1

Audience Response Question

- A patient who has an ANC of 0 and is Day + 5 following an allogeneic hematopoietic stem cell transplant develops a fever to 38.1°C. Blood cultures are obtained from both lumens of the double lumen tunneled catheter. One of the blood cultures becomes positive for *Enterococcus faecalis*. The other blood culture is positive for *Pseudomonas aeruginosa*.
- Does this event meet MBI criteria?
- Yes
- No

Audience Response Question

- A patient who has an ANC of 0 and is Day + 5 following an allogeneic hematopoietic stem cell transplant develops a fever to 38.1°C. Blood cultures are obtained from both lumens of the double lumen tunneled catheter. One of the blood cultures becomes positive for *Enterococcus faecalis*. The other blood culture is positive for *Pseudomonas aeruginosa*.
- Does this event meet MBI criteria?
- No – although the patient meets the neutropenia criterion and has one MBI organism present, *Pseudomonas aeruginosa* is not an MBI organism

Audience Response Question

- A patient who has an ANC of 0 and is Day + 5 following an allogeneic hematopoietic stem cell transplant develops a fever to 38.1°C. Blood cultures are obtained from both lumens of the double lumen tunneled catheter. Both cultures are positive for viridans group *Streptococcus*.
- Does this event meet MBI criteria?
- Yes
- No

Audience Response Question

- A patient who has an ANC of 0 and is Day + 5 following an allogeneic hematopoietic stem cell transplant develops a fever to 38.1°C. Blood cultures are obtained from both lumens of the double lumen tunneled catheter. Both cultures are positive for viridans group *Streptococcus*.
- Does this event meet MBI criteria?
- Yes – This event meets the neutropenia and organism criteria to be classified as an MBI event

Defining Single Positive Blood Culture (SPBC) Events

- A common commensal organism (e.g. coagulase-negative staphylococci [including *S. epidermidis*], viridans group streptococci) is cultured from a single blood culture with no other organisms isolated
 - Note: Refer to the current NHSN Organisms Lists (All Organisms, Top Organisms, Common Commensals, MBI Organisms, & Uropathogens)

Defining Secondary Bloodstream Infections

- LCBI related to an infection at another site
 - Blood and site-specific specimen cultures match for at least one organism in a patient suspected of having an infection
 - Blood and site-specific specimen cultures do not have to match if the site-specific culture is an element used to meet the infection site criterion and the blood isolate is also an element used to meet another criterion at the same infection site
 - Source: Appendix 1. Secondary Bloodstream Infection (BSI) Guide (not applicable to Ventilator-associated Events [VAE])
http://www.cdc.gov/nhsn/pdfs/pscmanual/4psc_clabscurrent.pdf

Preventing CLABSIs

- “Let whoever is in charge keep this simple question in her head (not, how can I always do this right thing myself, but) how can I provide for this right thing to be always done?”
 - Florence Nightingale

Organizational Obligations

- Empower staff to speak up if proper procedures are not being followed
- “Bundle” supplies (e.g., in a kit) to ensure items are readily available for use
- Ensure efficient access to hand hygiene
- Monitor and provide prompt feedback for adherence to hand hygiene
<http://www.cdc.gov/handhygiene/Measurement.html>
- Provide recurring education sessions on central line insertion, handling and maintenance

CLABSI Prevention Strategies

- Bundles
 - Groups of evidence-based interventions that, when implemented together, result in better outcomes than when implemented individually
- Maintenance care bundles
 - Accessing the line
 - Hand hygiene
 - Scrub access port or hub prior to each access
 - Dressings
 - Change frequency based on dressing type
 - Change promptly if wet, soiled, or dislodged
 - Perform dressing changes under aseptic technique
 - Administration sets and connectors
 - Routine tubing replacement
 - Routine needleless connector replacement

O’Grady, et al. (2011). Guidelines for the prevention of intravascular catheter-related infections. Centers for Disease Control and Prevention. http://www.cdc.gov/hicpac/pdf/guidelines/bsi-guidelines_2011.pdf

Checklist for Prevention of Central Line Associated Blood Stream Infections

© 2011 CDC. All rights reserved. This document is available at http://www.cdc.gov/hicpac/pdf/guidelines/bsi-guidelines_2011.pdf

- For Clinicians:**
 Promptly remove unnecessary central lines
- Perform daily audits to assess whether each central line is still needed
- Follow proper insertion practices**
- Perform hand hygiene before insertion
 - Adhere to aseptic technique
 - Use maximal sterile barrier precautions (i.e., mask, cap, gown, sterile gloves, and sterile full body sheet)
 - Perform skin antisepsis with 70% alcohol/methane with alcohol
 - Change the lead wire to minimize friction and mechanical complications
 - Avoid removal sites in adult patients
 - Avoid removal sites in adult patients
 - Cover the site with sterile gauze or sterile transparent, semipermeable dressings
- Handle and maintain central lines appropriately**
- Comply with hand hygiene requirements
 - Scrub the connector or hub immediately prior to each use with an appropriate antiseptic (e.g., chlorhexidine, povidone iodine, or iodophor) or 70% alcohol
 - Access catheters only with sterile device
 - Replace dressings that are wet, soiled, or dislodged
 - Perform dressing changes under aseptic technique using clean or sterile gloves
- For Facilities:**
- Empower staff to stop non-emergent insertion if proper procedures are not followed
 - Bundle supplies (e.g., kits) to ensure items are readily available for use
 - Provide the checklist above to clinicians to ensure all insertion practices are followed
 - Provide efficient access to hand hygiene
 - Monitor and prompt feedback for adherence to hand hygiene
 - Provide recurring education sessions on central line insertion, handling and maintenance
- Supplemental strategies for consideration:**
- 2% Chlorhexidine bathing
 - Antiseptic-impregnated catheters
 - Chlorhexidine impregnated dressings

National Center for Emerging and Zoonotic Infectious Diseases
http://www.cdc.gov/hicpac/pdf/guidelines/bsi-guidelines_2011.pdf

CLABSI Prevention Strategies

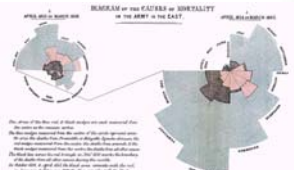
- Chlorhexidine bathing
 - Daily washing with disposable 2% CHG-impregnated washcloths
- Outcomes
 - Reduction in bacteremias involving coagulase-negative staph in critically ill adults and children (Climo et al., 2013; Milstone et al., 2013)
 - No reduction in bacteremias involving MBI organisms (Climo et al., 2013; Milstone et al., 2013)
 - Mixed findings regarding VRE colonization (Bass et al., 2013; Climo et al., 2009)
- Limitations
 - No well designed trials involving adult or pediatric BMT or oncology patients

CLABSI Prevention Strategies

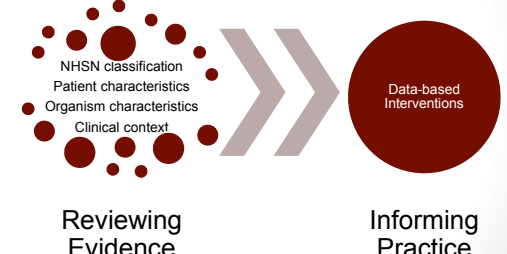
- Chlorhexidine dressings
 - CHG-impregnated sponge placed around the central line insertion site prior to placing the dressing
- Outcomes
 - No reduction in CLABSIs in trials involving critically ill children (Levy, 2005; Miller et al., 2011)
 - Mixed findings among critically ill adults (Arvaniti et al., 2012; Timsit et al., 2012)
- Limitations
 - No well-designed trials involving adult or pediatric BMT or oncology patients
 - No well-designed trials involving adult or pediatric patients with tunneled central venous catheters

Reviewing Data

- “To understand God's thoughts we must study statistics, for these are the measure of His purpose.”
 - Florence Nightingale



Know (and Own) Your Data



Risks for Bacteremia in BMT Patients

- Allogeneic vs. autologous transplant
- Donor type
- Myeloablative conditioning regimen
- Total body irradiation
- Prolonged neutropenia
- Underlying disease process
- Graft-versus-host disease
- Mucositis
- Immunosuppressive therapy
- Compromised skin integrity

Leading Sources of Bacteremia

Autologous Transplant <ul style="list-style-type: none"> • Coagulase negative <i>Staphylococcus</i> • <i>Enterococcus</i> species • <i>Corynebacterium nos</i> 	Allogeneic Transplant <ul style="list-style-type: none"> • Coagulase negative <i>Staphylococcus</i> • <i>Enterococcus</i> species • <i>Pseudomonas</i> species
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Source: Bock et al. (2013). Bacteremia in blood or marrow transplantation patients: Clinical risk factors for infection and emerging antibiotic resistance. *Biology of Blood and Marrow Transplantation*, 19, 102-108.

Leading Sources of Bacteremia

First 30 Days of Transplant <ul style="list-style-type: none"> • Coagulase-negative <i>Staphylococcus</i> • <i>Enterococcus</i> species • α-hemolytic <i>Streptococci</i> • <i>Escherichia coli</i> • <i>Pseudomonas</i> species 	31 to 100 Days Post-Transplant <ul style="list-style-type: none"> • Coagulase-negative <i>Staphylococcus</i> • <i>Enterococcus</i> species • <i>Pseudomonas</i> species • <i>Escherichia coli</i>
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Source: Bock et al. (2013). Bacteremia in blood or marrow transplantation patients: Clinical risk factors for infection and emerging antibiotic resistance. *Biology of Blood and Marrow Transplantation*, 19, 102-108.

Use Data to Guide Practice

- Review individual cases on a regular basis
- Implement interventions that match identified problems
- Evaluate outcomes of interventions
- Revise and refine interventions

Patient Characteristics

Length of hospitalization

Immune function

Skin integrity

Presence of GVHD

Clinical Context

Positive blood culture classification

Type of organism

Line type

Adherence to standard of care

Reviewing Data

- Review all positive blood culture events
 - All positive blood cultures can be clinically significant regardless of classification
 - Focusing solely on CLABSIs can minimize the significance of non-CLABSI events
 - Eliminating non-CLABSI positive blood cultures can result in missed opportunities to:
 - Understand the root cause of these events
 - Identify interventions for improvement
- Review and report CLABSI rates with and without MBI cases included
- Review inpatient and ambulatory events

Keys to Improvement

- “To be “in charge” is certainly not only to carry out the proper measures yourself but to see that every one else does so too; to see that no one either willfully or ignorantly thwarts or prevents such measures. It is neither to do everything yourself nor to appoint a number of people to each duty, but to ensure that each does that duty to which he is appointed.”
 - Florence Nightingale

Keys to Improvement

- “Structured, organization-wide approach to understanding and improving work processes”
 - Specific, measurable mission or goal statement
 - Multi-departmental/disciplinary involvement
 - Resource and educational materials
 - Reward/incentive programs
 - Internal and external stakeholders
 - Project champion
 - Feedback-based process and outcome measurement

Source: Compas et al. (2008). Best practices in implementing and sustaining quality of care: A review of the quality improvement literature. *Research in Gerontological Nursing*, 1, 209-216.

Keys to Improvement

- Leadership support
 - Ensure financial resources
 - Support project leaders with the training and time to commit to the project
 - Recognize and support the time needed to conduct a successful project
 - Reinforce expectations as needed as component of organization’s commitment to safety

Source: Hughes, R. (2008). *Patient safety and quality: An evidence-based handbook for nurses*. Rockville, MD: Agency for Healthcare Research and Quality. Accessible online at <http://www.ahrq.gov/QUAL/nursesdbk/>.

Keys to Improvement

- Effective teams
 - Identified leader and core team members
 - Assigned roles and responsibilities
 - Ensure input/feedback representing various perspectives
 - Team meetings used for specific objectives
 - Work accomplished outside of meetings
 - Clear communication
 - Accountability of team members

Proposing Interventions

- “It is often thought that medicine is the curative process. It is no such thing; medicine is the surgery of functions, as surgery proper is that of limbs and organs. Neither can do anything but remove obstructions; neither can cure; nature alone cures. ... And ***what nursing has to do in either case, is to put the patient in the best condition for nature to act upon him.***”
 - Florence Nightingale

Proposing Interventions

- Is the proposed intervention consistent with sound nursing practice or current best practice guidelines for BMT patients?
- Does the proposed intervention address the identified problem?
- Does the proposed intervention represent a departure from current institutional practice?
- What additional resources will be required to implement the proposed intervention?

- “The amount of relief and comfort experienced by the sick after the skin has been carefully washed and dried, is one of the commonest observations made at a sick bed.”
 - Florence Nightingale

Data-Based Intervention

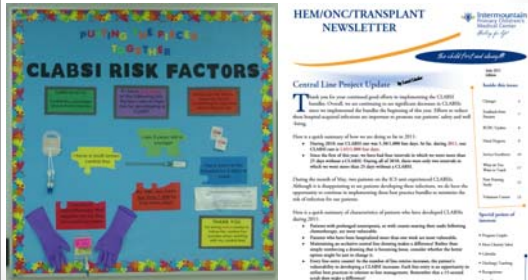
- Emphasis on supportive cares
 - 1-2-3 initiative
 - Skin care
 - Preventing constipation
- Outcomes
 - Inpatient CLABSI rates
 - 2012: 3.78/1,000 line days
 - 2013: 2.47/1,000 line days
 - 38% reduction in all inpatient positive blood culture events



Promoting Staff Engagement

- Communicate on a regular basis
 - Facilitating questions and answers related to the project
- Interact with staff on an ongoing basis
 - Ensuring staff expectations and understanding related to the project
- Promote personal commitment to the project
 - Linking project outcomes to annual goals
 - Sharing personal success stories

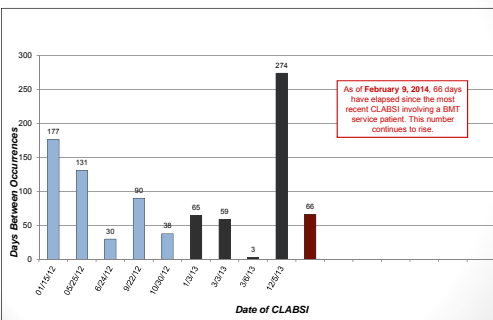
Promoting Staff Engagement



Promoting Staff Engagement

- Presenting data related to the project
 - Present data in a consistent, meaningful manner
 - Provide resources for interpreting data
- Relating data to practice
 - Express in outcomes relevant to practice
 - Present in a manner that illustrates benefits to patients
 - Share lessons learned that can be applied to clinical practice

Calendar days between CLABSIs involving BMT Service Patients (MBI included) 2012 - 2014 YTD



Areas for Ongoing Development

- CLABSI reporting
 - Consistency in application of NHSN CLABSI definitions (Gaur et al., 2013)
 - Documenting MBI-LCBI criteria
- Refining practice
 - Research evaluating interventions for CLABSI reduction in BMT settings
 - Institution-based quality improvement efforts
- Institutional commitment
 - Availability of resources
 - Facilitating implementation of best practice

Summary

- Applying MBI-LCBI criteria can help delineate the nature of bloodstream infections occurring in BMT and other immune compromised patients
- Understanding characteristics of CLABSIs occurring in BMT patients can inform strategies for reducing these infections
- Developing data-based quality improvement projects can engage staff and improve overall care, including reducing CLABSIs

In closing ...

- “So never lose an opportunity of urging a practical beginning, however small, for it is wonderful how often in such matters the mustard-seed germinates and roots itself.”
 - Florence Nightingale

