

THE PREVENTION OF BLOODSTREAM INFECTIONS IN ALL VASCULAR ACCESS DEVICES

A Lecture for APIC San Diego, CA
July 10, 2024

Robert Garcia, MT(ASCP), CIC, FAPIC
President & Senior Infection Prevention Consultant,
Enhanced Epidemiology LLC



Disclosures

Clinical Advisor/Consultant/Speaker affiliations:

Bravida

Entrotech

Ansell

Kurin

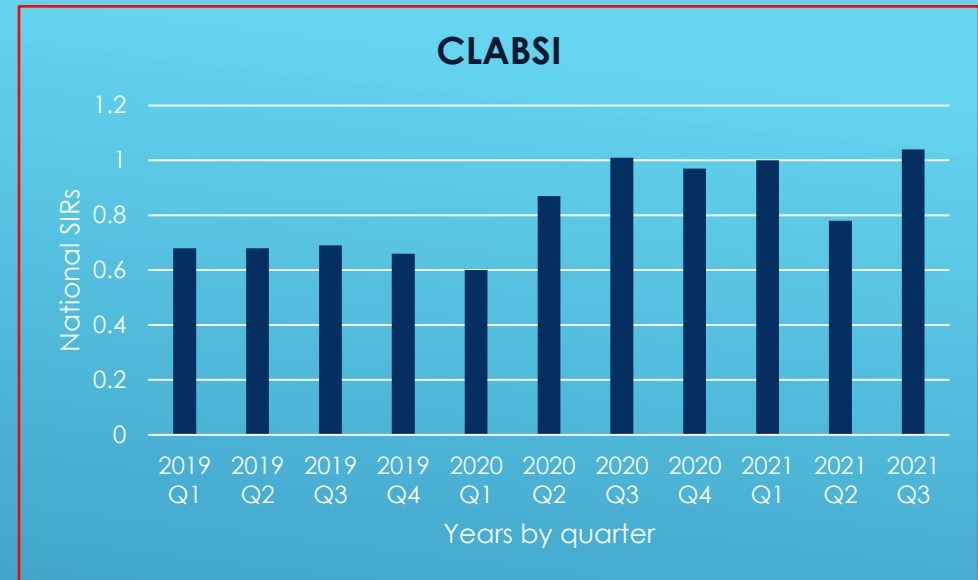
A decorative graphic consisting of several parallel white lines of varying lengths and thicknesses, arranged diagonally from the bottom right towards the top right of the slide.

Learning Objectives

At the conclusion of this session, participants will be able to:

- Discuss the main mechanism of development of catheter-related BSIs
 - Describe the impact of the Great Resignation and the COVID-19 pandemic on rates of central line-associated bloodstream infection (CLABSI)
 - Discuss the pending national requirements on infection surveillance of all-cause Hospital Onset Bloodstream Infections
 - Evaluate the findings of recent studies on new strategies for the prevention of Hospital Onset BSIs
- 

Background



- ▶ CDC NHSN data indicated a **47% increase** in CLABSI during the COVID 19 pandemic
- ▶ Rates have decreased by 9% between 2021 and 2022
- ▶ *What does this data not tell us?*
 - ▶ *There was considerable prevention efforts that were needed prior to the pandemic*
 - ▶ *Does not include BSI rates on other VADs*

The Great Resignation

Daily Briefing

The 'Great Resignation' is over. So why doesn't it feel like it?

Posted on January 26, 2024

Updated on January 25, 2024

According to a new analysis published in *NEJM Catalyst*, employment in healthcare settings is now higher than its pre-pandemic levels, and real wage growth is higher than inflation. However, hospitals are still struggling with employee burnout, difficulty hiring, and more — signs of deeper troubles that have not been offset by employment and wage growth.

TOPICS

Workforce

- ▶ During the pandemic, **100,000** registered nurses left the workforce
- ▶ By 2027, projections are that **900,000** more nurses intend to leave the profession
- ▶ *What does this data not tell us?*
 - ▶ *Although healthcare worker hiring has increased in the last year, what the data excludes is the impact on the application of proper IP practices via the loss of experienced front-line workers*

Need for Expansion of Surveillance and Prevention Initiatives (I)



ONGOING DISCUSSION WITHIN
IP PROGRAMS TO **EXPAND BSI
SURVEILLANCE** BEYOND CLABSI



NEED TO INSTITUTE
STANDARDIZED INTERVENTIONS
TO PREVENT BSIS IN **ALL VADS**



NATIONAL BSI RATES
ASSOCIATED WITH **NON-CLABSI
VADS** IS UNKNOWN

Need for Expansion of Surveillance and Prevention Initiatives (II)



Centers for Medicare and Medicaid Services (CMS) has proposed a new metric, **Hospital Onset Bacteremia and Fungemia (HOB)**

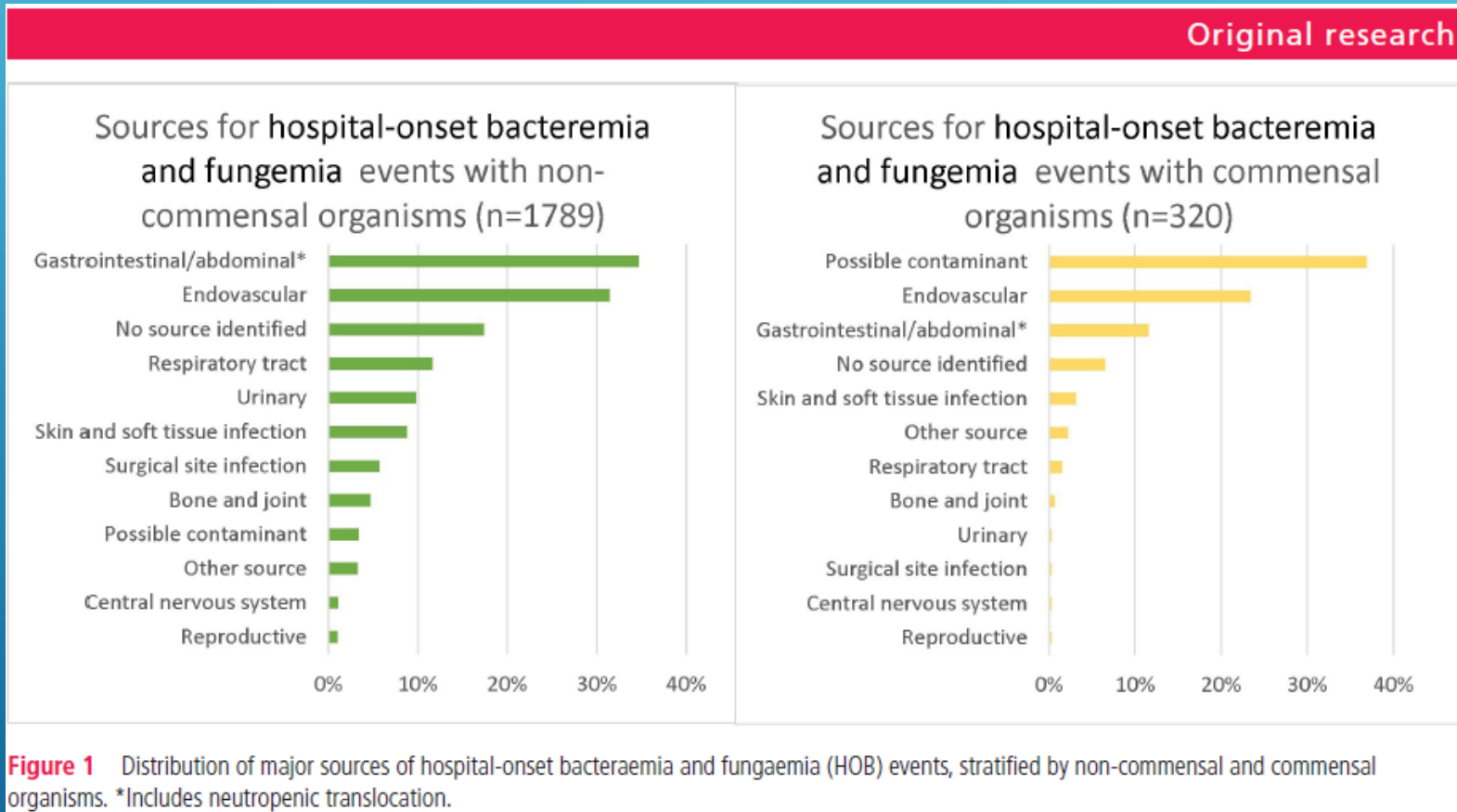
The **HOB** proposal would require hospitals to expand surveillance with the intent on broader reduction of BSIs regardless of causative organism or association with a medical device

WHY THE EVOLUTION TO HOB?

- ▶ *“The transition from a focus on a parochial quality measure and “on paper” improvement with CLABSI to true and broader-scale harm reduction with HOB has precedent in the transition from ventilator-associated pneumonia to ventilator-associated events: **a simpler and more objective definition, a broadening of the scope of harm, and new science demonstrating the preventability of these adverse events.**”*

SOURCES OF HOB

- ▶ Study of HOB events at 10 academic and 3 community hospitals in USA



Leekha S, et al. Evaluation of hospital-onset bacteraemia and fungaemia in the USA as a potential healthcare quality measure: a cross-sectional study. *BMJ Qual Saf* 2024. Epub ahead of print.

HOB Metric and Complementary Metrics

Category/Use	Event	Numerator	Denominator
	Measure Event		
<i>Primary Metric</i>	HOB Event	Pathogenic bacteria or fungi from blood culture on hospital day ≥ 4 , (excluding patients with prior matching cultures and HOB events)	Inpatient Admissions
	Complementary Metrics		
<i>Risk Adjustment, Diagnostic Stewardship, QI</i>	Blood Culture Utilization	<u>Testing Prevalence</u> : Admissions with at least 1 blood culture <u>Testing Intensity</u> : Total blood cultures among patients with at least 1 blood culture	
<i>QI</i>	Blood Culture Contamination	Skin commensal organism in 1 of 2 blood culture sets	Total blood culture sets
<i>Risk Adjustment</i>	Community-Onset Bacteremia & Fungemia Event	Pathogenic bacteria or fungi from blood culture prior to hospital day 4, (excluding patients with prior matching cultures and COB events)	Inpatient Admissions
<i>Risk Adjustment, QI</i>	Matching Commensal Bacteremia Event	Skin commensal from ≥ 2 blood cultures, AND ≥ 4 days of antibiotic treatment	Inpatient Admissions
<i>Risk Adjustment, QI</i>	Non-Measure HOB Event	HOB events among patients with conditions that highly predict non-preventability	Inpatient Admissions

#APIC2023

***“Prevention of Vascular Access Device-Associated Hospital Onset Bacteremia and Fungemia:
A Review of Emerging Perspectives and Synthesis of Technical Aspects”***

E-pub publication in



- ▶ Authored by IPs, ID, and VA specialists
 - ▶ Intent is to provide a comprehensive review of strategies to optimize VAD lifecycle practices, while providing new perspectives to enhance current prevention programs and assist hospitals in preparation for expansion of surveillance and prevention of VAD-HOB
-
- Robert Garcia, MT(ASCP), CIC, FAPIC
 - Edward J. Septimus, MD
 - Jack LeDonne, MD, VA-BC
 - Lisa K. Sturm, MPH, CIC, FAPIC
 - Nancy Moureau, PhD, RN, CRNI, CPUI, VA-BC
 - Michelle DeVries, MPH, CIC, VA-BC, CPHQ, FAPIC
 - Barbara DeBaun, MSN, RN, CIC

***“Prevention of Vascular Access Device-Associated Hospital Onset Bacteremia and Fungemia:
A Review of Emerging Perspectives and Synthesis of Technical Aspects”***

Identifies and provides insights on emerging issues in the mitigation of HOB

➤ Technical Aspects (Article 1):

- Lifecycle of a VAD
- Guideline Recommendations
- Patient Decolonization
- Create/Expand VATs
- Ultrasound Transducers
- Peripheral Intravenous Catheters
- Advanced Antimicrobial Dressings

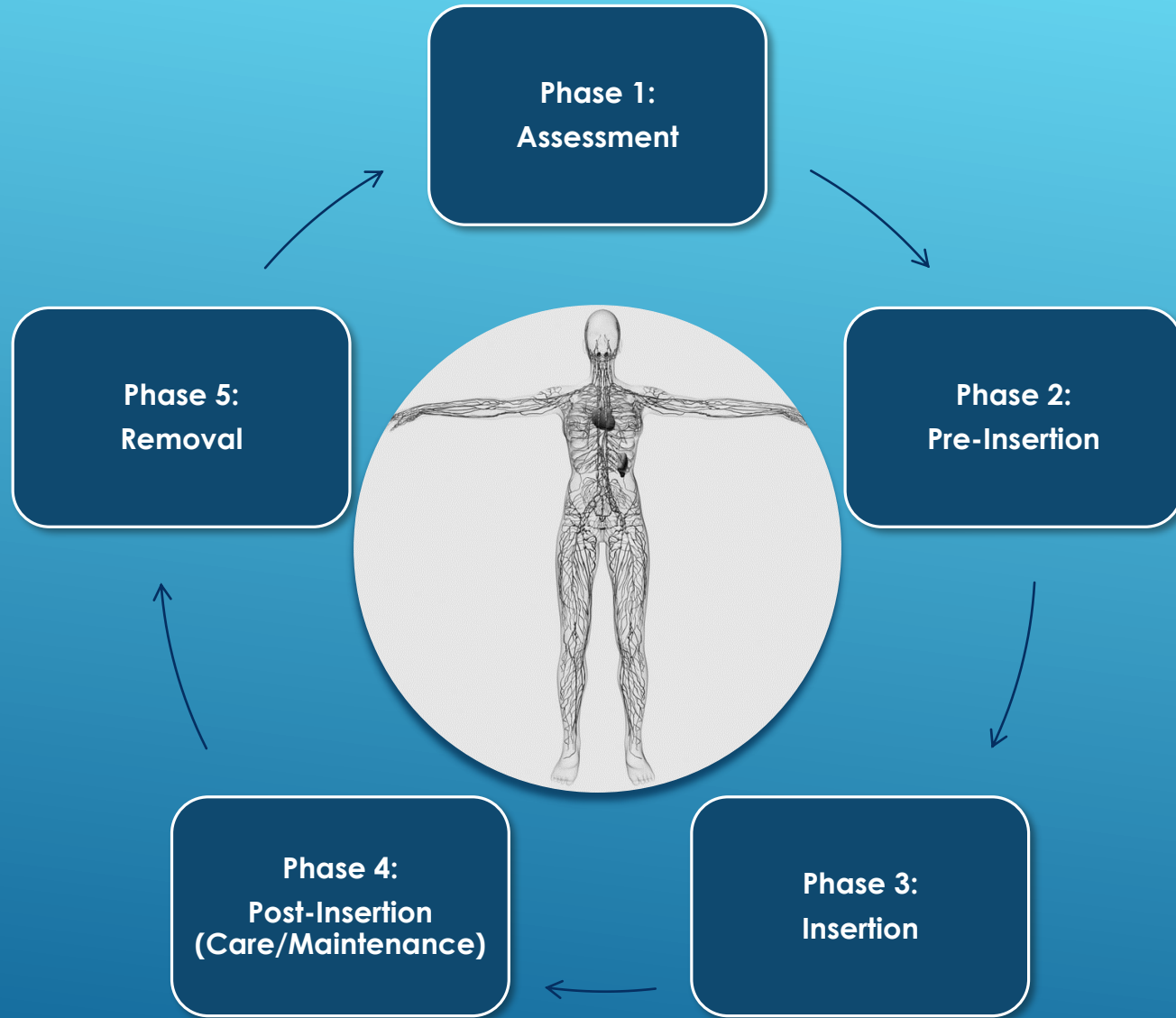
➤ Implementation Aspects (Article 2):

- Leadership
- Staffing
- Diagnostic Stewardship
- Systems and Human Factors Engineering
- Approaches for Prevention
- Bundle Compliance
- Data Comprehension
- Education of VA/IP Specialists



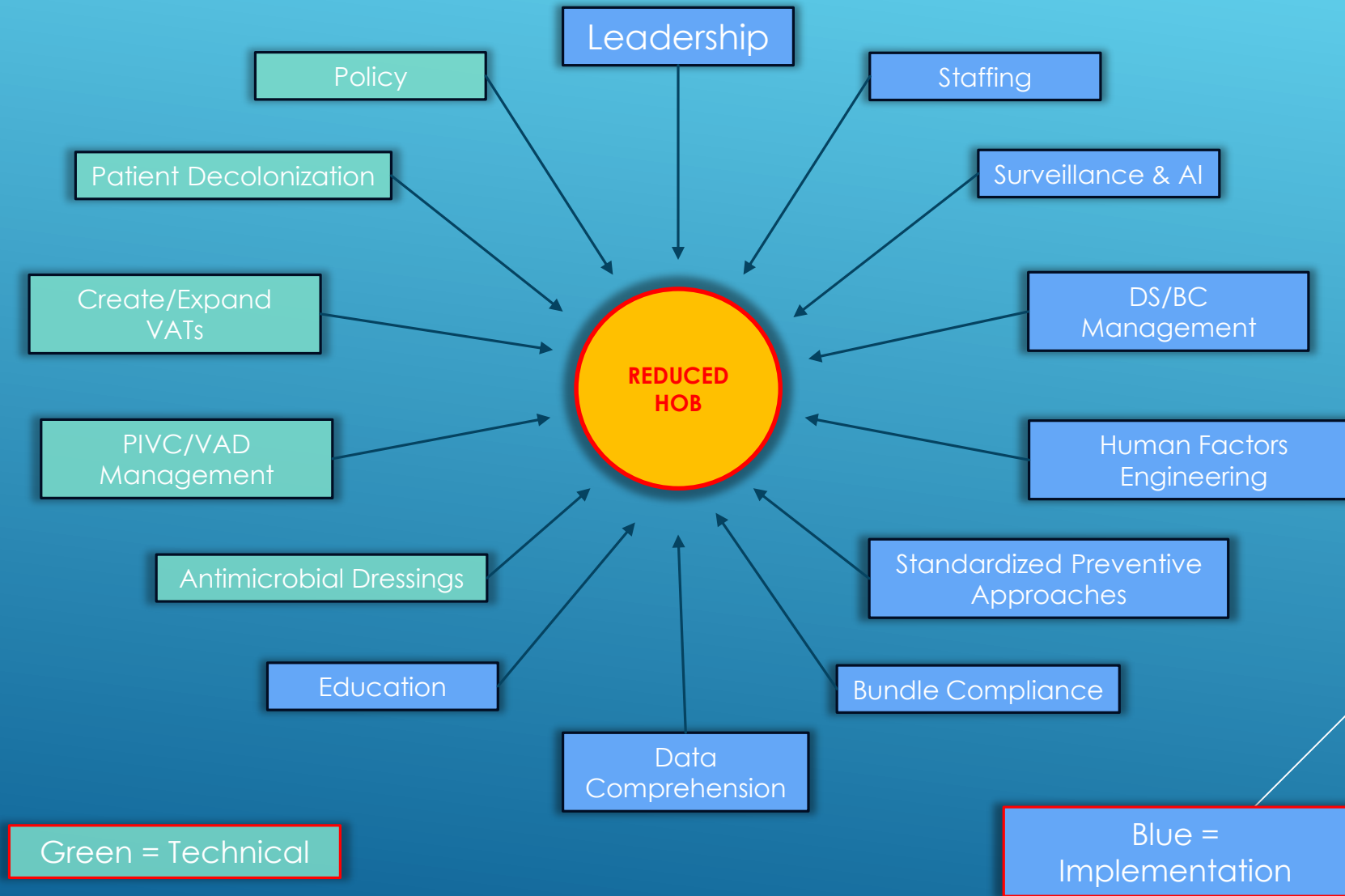
EMERGING PERSPECTIVES: TECHNICAL ASPECTS

Lifecycle of a Vascular Access Device



While specific practices in each phase may differ among various types of VAD, the primary message is that each phase represents an opportunity for infection prevention

CORE COMPONENTS OF HOB PREVENTION PROGRAMS



Lifecycle of a VAD: Time Frames

Phase	Time frame
1. Assessment	From start of patient assessment to determination of need and type of VAD for insertion
2. Pre-Insertion	From hand hygiene through gathering of supplies, donning of barriers, field setup, and patient skin prepping
3. Insertion	From start of patient draping to application of insertion site dressing
4. Post-Insertion	Starts after application of initial dressing through VAD removal
5. Removal	Starts when daily assessment process indicates device is no longer needed through actual removal of the device Note: the clinician may decide to initiate an alternate device

Vascular Health & Preservation (VHP)

J Vasc Access 2012;13(3):351-356
DOI: 10.5301/jva.5000042

ORIGINAL ARTICLE

Vessel health and preservation (Part 1): A new evidence-based approach to vascular access selection and management

Nancy L. Moureau¹, Nancy Trick², Thomas Nifong³, Cathy Perry⁴, Cheryl Kelley⁵, Ruth Carrico⁶, Michael Leavitt⁷, Steven M. Gordon⁸, Jessica Wallace⁹, Monte Harvill¹⁰, Connie Biggar¹¹, Michael Doll¹², Loreli Papke¹³, Lori Benton¹⁴, Deborah A. Phelan¹⁵

¹Excellence, Inc., Hartwell, GA - USA
²Teleflex, Reading, PA - USA
³Pennsylvania State University
⁴Vanderbilt Medical Center, Nashville, TN
⁵University of Louisville, KY
⁶University of Michigan Medical Center, Ann Arbor, MI
⁷Cleveland Clinic, Cleveland, OH
⁸University of California, Los Angeles, Los Angeles, CA
⁹Detroit Medical Center, Detroit, MI
¹⁰Munson Medical Center, Traverse City, MI
¹¹Geisinger Medical Center, Danville, PA
¹²Texas A&M Health Science Center, Houston, TX
¹³Mercy Hospital, Portland, ME

ABSTRACT

Vascular access for the infusion of medications is a common procedure. Traditional vascular access selection is based on individual clinical pathways. A Vessel Health and Preservation (VHP) program implemented within 24 hours of device selection that provides a systematic approach to device selection and management is the ultimate goal of vessel health.

Key words: Central Venous Catheter, Peripherally Inserted Central Catheter

Accepted: November 28, 2011

INTRODUCTION

Venous access for infusion of medications is a common procedure. Without venous access, few medications can be administered. Given the high incidence of intravenous device placement, it is necessary to ensure the delivery of medications. In spite of emphasis on safety from various regulatory agencies,

Vessel Health and Preservation: The Right Approach for Vascular Access

Nancy L. Moureau
Editor



Springer Open

Emerging perspectives in the mitigation of HOB BSI stress the necessity of ensuring for each patient the most appropriate device selection, optimal insertion technique, followed by standardization of post-insertion care. The concept of **Vascular Health and Preservation**, a vascular access framework, includes the primary goal to “...drive vascular access care, regardless of the point of entry into a healthcare facility, based on a system of evidence-based practices, standards, and guidelines by means of collaborative agreement by all disciplines and care providers.”

Focus on CVAD Practices

Phase 1 – Assessment:

- the clinician determines whether peripheral IV access appropriate or if a CVAD is necessary
- determines which device is appropriate
- evaluates the condition of the various veins of the neck, chest, extremities

Phase 2 – Pre-Insertion:

- starts with hand hygiene until maximum sterile draping is applied
- To achieve an optimal level of practice in Phase 2, each element that comprises the pre-insertion bundle must be standardized by the institution among all designated inserters

Focus on CVAD Practices

Phase 3 – Insertion: encompasses the period between the application of the sterile drapes and the application of the dressing at the end of the insertion

****What is done in Phase 3 Insertion has a profound impact on optimal Phase 4 Care and Maintenance practice****

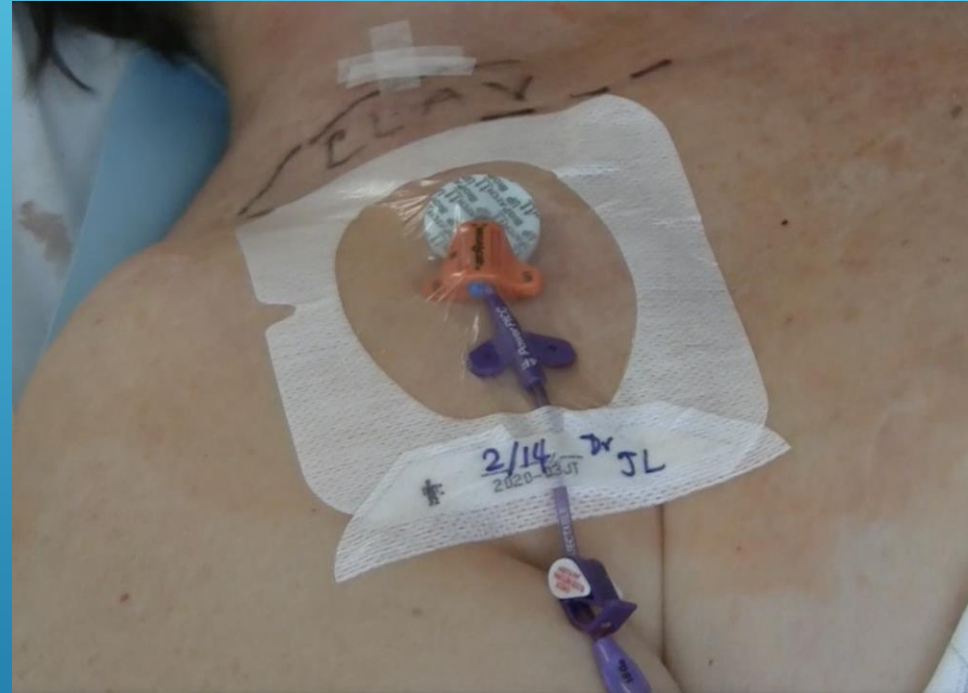
A proper CVAD insertion consists of the following:

- i. minimal attempts
- ii. the tip of the catheter must be positioned in the vicinity of the Cavoatrial Junction (this location minimizes the risk of thrombosis)
- iii. proper exit site
- iv. sutureless securement
- v. antimicrobial protection (coated catheters and dressings)
- vi. the application of a transparent dressing that remains intact for at least 7 days

Focus on CVAD Practices



Upward and on the Neck



Downward and on the Chest

Focus on CVAD Practices

Phase 4 – Post-Insertion (Dressing and Care and Maintenance):

- Starts with the first dressing application and ends when the catheter is removed
- An emphasis on **maintenance of asepsis** during all VAD manipulations should be considered one of the strongest recommendations. This concept applies to all catheter access via hubs, connectors, or injection ports.
- Unscheduled (less than 7 days) dressing changes increase the risk of infection and add nursing time and supply cost to an unreimbursed procedure



Focus on CVAD Practices

Phase 5 – Removal:

- **Removal** of unnecessary medical devices is a fundamental intervention in preventing HAIs
- **Strategies** advocated to assist in removal of VADs include daily assessment of clinical need, establishing criteria to assist in clinical determination (e.g., unresolved complication, discontinuation of infusion therapy, when no longer necessary for the plan of care), and avoidance of scheduled replacement based on dwell time
- The **2024 INS Standards of Practice** provides a comprehensive list of recommendations addressing removal of various VADs including PIVCs, midline catheters, non-tunneled CVADs including PICCs, surgically placed CVADs, and arterial catheters

Infusion Therapy Standards of Practice

Barbara Nickel, APRN-CNS, CCRN, CRNI®
Lisa Gorski, MS, RN, HHCNS-BC, CRNI®, FAAN
Tricia Kleidon, PhD(c), MNSc, RN
Amy Kyes, MSN, APRN, AG-CNS, CV-BC™, CRNI®
Michelle DeVries, MPH, CIC, VA-BC, CPHQ, FAPIC
Samantha Keogh, PhD, RN, FACN
Britt Meyer, PhD, RN, CRNI®, VA-BC, NE-BC
Mary Jo Sarver, MSN, ARNP, AOCN, CRNI®, LNC, VA-BC
Rachael Crickman, DNP, ARNP-CNS, AOCNS, OCN, RN
Jenny Ong, PharmD
Simon Clare, MRes, BA, RGN
Mary E. Hagle, PhD, RN-RB, FAAN

9TH EDITION
REVISED 2024

INS

INFUSION NURSES SOCIETY
ADVANCING THE STANDARDS FOR OUR CARE™
One Edgewater Drive, Norwood, MA 02062
www.ins1.org

Supplementary Table 1: Emerging Perspectives for Optimizing Vascular Access Device Practices

Lifecycle Phase (Time frame)	Vessel Health & Preservation (VHP) Framework	Focus on Central Venous Access Catheters (CVAD)
<p>1. Assessment (from start of patient assessment to determination of need and type of VAD for insertion)</p>	<ul style="list-style-type: none"> Evaluate patient risk, history, comorbidities, vein choices and options to determine if there is a genuine need for vascular access Determine vein quality using a grading system that assists with insertion management Determine medication suitability on the safety of the drugs and assess appropriate VAD Use an algorithm for VAD choice, based on vein quality, drug choice and treatment duration When conducting a risk assessment before insertion of a PIVC, question whether a more suitable alternative is available Select device size based on vein size, risk, and insertion location Verify minimum number of lumens required 	<ul style="list-style-type: none"> During the <i>Assessment Phase</i>, there are three critical decision points to be achieved: <ul style="list-style-type: none"> (i) the clinician should initially determine whether peripheral intravenous access is appropriate or if the patient will require an alternate VAD (ii) the clinician determines, based on the patient's medical condition and comorbidities, which device(s) are <u>appropriate</u> and which are unsuitable (iii) evaluate the condition of the various target veins to establish the likelihood of successful cannulation In a patient with underlying chronic kidney disease that requires central venous access, a PICC would generally be inappropriate to avoid compromising permanent dialysis fistula options. In this example, the clinician would evaluate the condition of the various veins of the neck, chest, and extremities using ultrasound technology [See section "Use and Disinfection of Ultrasound Transducers"]
<p>2. Pre-Insertion (from hand hygiene through gathering of supplies, donning of barriers, field setup, and patient skin prepping)</p>	<ul style="list-style-type: none"> Perform hand hygiene using an alcohol-based hand rub or liquid soap and water Equipment and supplies for the VAD insertion procedure should be standardized in all areas where the procedure is performed Skin decontamination should be standardized and reflect the manufacturer's recommendation for use 	<ul style="list-style-type: none"> To achieve an optimal level of practice in the <i>Pre-Insertion Phase</i>, each element that comprises a pre-insertion bundle must be standardized by the institution for each type of VAD and among all designated inserters (e.g., set-up of aseptic field prior to insertion, ensuring the use of the skin antiseptic as per manufacturer's recommendations, selection and provision of a standardized equipment and supply set in every area where such procedures are conducted) Periodic monitoring for compliance with each element is warranted The original 2001 Institute for Healthcare Improvement (IHI) Central Line Bundle advocated five elements: hand hygiene, chlorhexidine 2% skin preparation, maximal barrier precautions, optimal site selection (i.e., avoid the femoral vein), and daily review of line necessity. In actual practice, a <i>Pre-Insertion Phase</i> needs to be emphasized, a designated period that starts with hand hygiene until the maximum sterile draping is applied

► Guideline Recommendations on the Technical Aspects of HOB BSI Prevention

Supplementary Table 2: Synthesis of Selected Technical Aspects of Vascular Access Devices

Applicable to all Vascular Access Devices				
Lifecycle Phase	Recommendation Topic	CDC Guidelines for the Prevention of Intravascular Catheter-Related Infections, 2011/2017	SHEA/IDSA/APIC Strategies to Prevent Central Line-Associated Bloodstream Infection in Acute-care Hospitals, 2022	INS Standards of Practice, 2024*
1-Assessment	Catheter selection	Select catheters <u>on the basis of</u> the intended purpose and duration of use, known infectious and non-infectious complications (e.g., phlebitis and infiltration), and experience of individual catheter operators (Cat IB)		<p>Infusion therapy and vascular access decisions are collaborative (health care team, the patient/caregiver), with consideration to the patient's diagnosis and clinical presentation, vasculature, device selection, and risk versus benefit of alternative routes of therapy. (Standard)</p> <p>The appropriate vascular access device (VAD), peripheral or central, is selected based on the prescribed therapy or treatment regimen, anticipated duration of therapy, vascular pathway, patient's age, comorbidities, history of infusion therapy and vascular access, patient preference for VAD type and location, overall vascular health (history of difficult intravenous access, vessel, and skin health at insertion site), and ability and resources available to care for the VAD. (Standard)</p>

► Guideline Recommendations on the Technical Aspects of HOB BSI Prevention

Table 3. Summary of Vascular Access Device Recommendations

Device	PIVC	UGPIVC	MIDLINE	PICC	CVC non-tunneled	Antimicrobial CVC	Tunneled CVC	PORT
Definition	Known by several terms such as short peripheral intravenous catheter, Designed for placement in the extremities. Over the needle cannula, sizes 22 to 12 gauge, length 2-7.5cm.	Ultrasound guided peripheral intravenous cannula. Over the needle cannula, sizes 22 to 12 gauge, length usually 4-7.5cm using insertion guided by ultrasound.	Midline catheter, 8-25cm, inserted into veins of the periphery, usually the mid upper arm. Various types of insertions including traditional with modified Seldinger technique 20cm catheter or, accelerated Seldinger technique, 8-15cm over the needle catheter and wire all in one. Terminal tip does not extend into the chest.	Peripherally inserted central catheter usually inserted in the mid upper arm into the basilic or brachial veins. Terminal tip in the distal superior vena cava requires tip confirmation prior to usage either by x-ray, fluoroscopy, or ECG guidance.	Percutaneously catheter inserted most commonly into the internal jugular or subclavian vein and advanced to the distal portion of the superior vena cava. Terminal tip requires confirmation prior to usage.	Antimicrobial impregnated or coated catheter inserted into the distal portion of the superior vena cava either through the arm (PICC) or in the same method as non-tunneled CVC. Reduces risk of infection.	Percutaneously inserted catheter where the external lumen of the catheter is tunneled back to a favorable area reducing risk of bacterial migration along the insertion track. Added stabilization is conferred to this catheter when the skin grows into the Dacron cuff adherent to catheter and coated just under	Subcutaneously implanted port includes both a port reservoir and a catheter attached to the port. The catheter is inserted in the same manner as a tunneled catheter and attached to the port positioned in a pocket with skin closure on top. The port is usually positioned in an area on the

Patient Decolonization

ORIGINAL ARTICLE

Targeted versus Universal Decolonization to Prevent ICU Infection

Susan S. Huang, M.D., M.P.H., Edward Septimus, M.D., Ken Kleinman, Sc.D., Julia Moody, M.S., Jason Hickok, M.B.A., R.N., Taliser R. Avery, M.S., Julie Lankiewicz, M.P.H., Adrijana Gombosov, B.S., Leah Terpstra, B.A., Fallon Hartford, M.S., Mary K. Hayden, M.D., John A. Jernigan, M.D., *et al.*, for the CDC Prevention Epicenters Program and the AHRQ DECIDE Network and Healthcare-Associated Infections Program*

Article	Figures/Media	Metrics	June 13, 2013
40 References	580 Citing Articles	Letters	2 Comments

N Engl J Med 2013; 368:2255-2265
DOI: 10.1056/NEJMoa1207290

Decolonization is an evidence-based intervention that can be used to prevent HAIs including BSIs, with the goal to reduce or eliminate the bioburden on the patient, thereby reducing the risk of a subsequent infection

Since colonization can lead to infection, two overarching approaches to HAI prevention have emerged: (1) **vertical** approaches to reduce colonization or infection due to specific pathogens and (2) **horizontal** strategies to broadly reduce the burden of all pathogens

There have been at least seven trials that utilized chlorhexidine gluconate (CHG) as a skin decolonizer (with and w/o nasal decolonization), all of which demonstrated reductions in either all-cause BSIs or CLABSIs

REDUCE MRSA trial: mupirocin/CHG, universal decolonization, 74 ICUs; Result: **44% decrease in all-cause BSI**

Can we apply this concept to all VADs?

Create/Expand Vascular Access Teams

Since 2002, the CDC has recommended specialized “IV teams” due to their “...unequivocal effectiveness in reducing the incidence of catheter-related infections and associated complications and costs”

A dedicated vascular access team (VAT) provides expert guidance in insertion, maintenance, and removal practices, as well as demonstrating “...extensive knowledge in difficult blood draws, use of ultrasound guidance, dressing protocols, daily evaluation of catheter necessity, and removal of unnecessary catheters, as well as providing recommendations on alternative devices”

Establishment of VATs has been correlated with an overall **47% reduction** in insertion related CLABSI



Use and Disinfection of Ultrasound Transducers

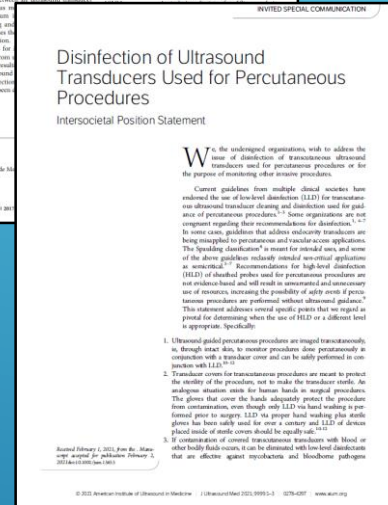
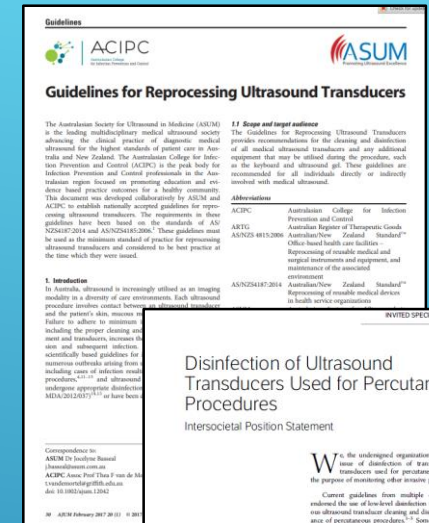
As challenges with difficult access patients increase, use of visualization technologies is necessary to gain successful VAD access

The use of **ultrasound** technology provides extensive benefits including reducing the number of cannulation attempts which may contribute to mitigation of microorganism introduction and subsequent BSI

Disinfection approaches:

Australian Society for Ultrasound in Medicine (ASUM): Based on the **Spaulding Classification**, a system that categorizes medical devices or equipment into three groups: *critical* (e.g., surgical instruments that enter sterile tissues/vascular system; requires sterilization), *semicritical* (e.g., items that contact mucous membranes or non-intact skin; need to be free of all microorganisms except small numbers of spores; use high-level disinfection [HLD] or sterilization), and *non-critical* (e.g., items that touch only intact skin and not mucous membranes; use low-level disinfection [LLD]).

American Institute of Ultrasound in Medicine (AIUM): Based on type of procedure performed, **Percutaneous vs. Endocavitary**. AIUM states "preparation of *external* transducers between patients requires a LLD process", while "preparation of *internal* transducers between patients requires routine mandatory HLD and the use of a high-quality single-use transducer cover during each examination". Endorsed by APIC, AVA, SHEA.



Peripheral Intravenous Catheters (PIVC)

While there has been an intensive focus on infections and complications from CVADs for more than five decades, there has been a growing awareness that PIVCs can also cause significant morbidity and mortality

Researchers have identified that more than a third of the hospital onset *S. aureus* BSIs were associated with PIVCs rather than central lines

Bundled approaches to PIVC insertion and care have been developed by some hospitals, however, additional work is needed to identify an optimal bundle

Pending publication "Peripheral Intravenous Catheter Consensus Document" (AVA, INS, AACN, ECRI)

Table 1. Sources of *Staphylococcus aureus* Bacteremia

Source	Total <i>S. aureus</i> Bacteremia (n = 205), No. (%)	Hospital-Onset <i>S. aureus</i> Bacteremia (n = 45), No. (%)	Community-Onset <i>S. aureus</i> Bacteremia (n = 160), No. (%)
Soft tissue/bone	67 (32.7)	4 (8.9)	63 (39.4)
PVC	18 (8.8)	16 (35.6)	2 (1.3)
CVC or PICC	14 (6.8)	7 (15.6)	7 (4.4)
Hemodialysis	13 (6.3)	2 (4.4)	11 (6.9)
Pulmonary	8 (3.9)	0 (0.0)	8 (5.0)
Endovascular	7 (3.4)	1 (2.2)	6 (3.8)
Biliary	1 (0.5)	0 (0.0)	1 (0.6)
Urinary	3 (1.5)	0 (0.0)	3 (1.9)
Unknown	74 (36.1)	15 (33.3)	59 (36.9)

Abbreviations: CVC, central intravenous catheter; PICC, peripherally inserted central catheter; PVC, peripheral intravenous catheter.

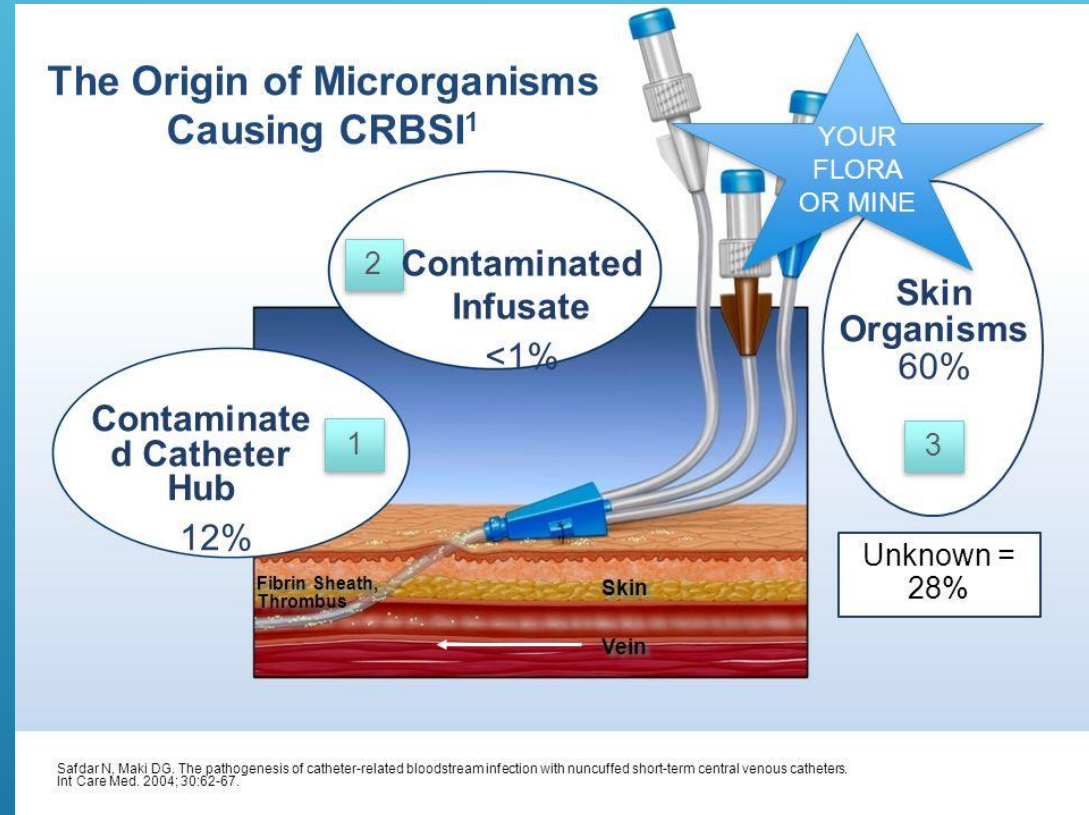


ADVANCED ANTIMICROBIAL INTRAVENOUS DRESSINGS – MECHANISM OF TRANSMISSION

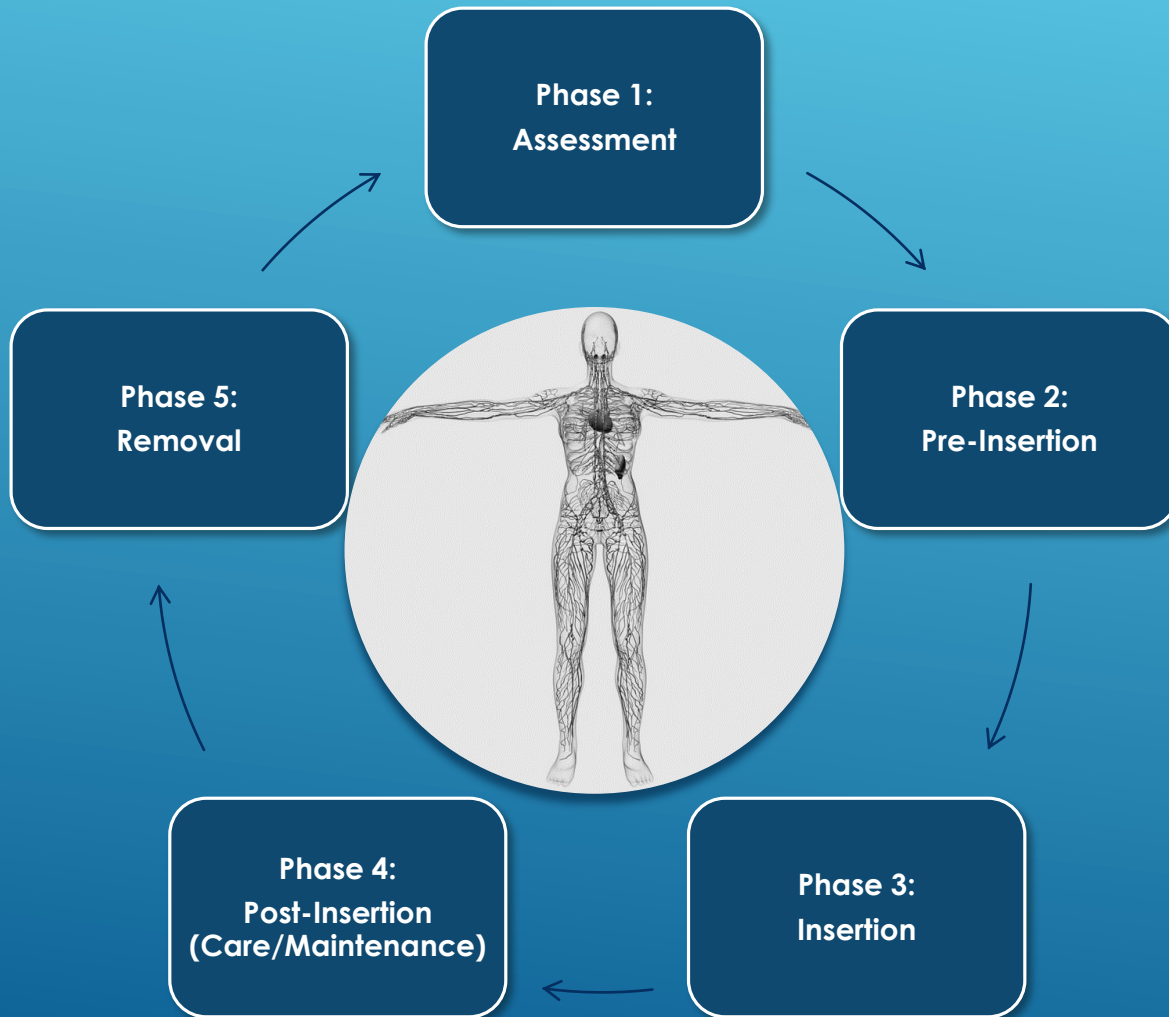
➤ When an IV catheter bypasses the skin barrier, a process starts whereby microorganisms colonize the catheter surface, and may enter the blood vessel, leading to systemic infection

➤ The majority of CRBSI are associated with organisms acquired from skin sources

➤ This process occurs at *all* IV catheter sites soon after insertion, regardless of catheter type and body site




RELATIONSHIP OF VAD LIFECYCLE & IV DRESSING



Total Lifecycle Time (mins):
Assessment, Pre-Insertion,
Insertion = 1%;
Care & Maintenance = 99%

Advanced Antimicrobial Intravenous Dressings - CHG

 NIH Public Access
Author Manuscript
Manuscript available in PMC 2014 December 08
Published in final edited form as:
Crit Care Med. 2014 July; 42(7): 1703-1713. doi:10.1097/CCM.0000000000000319

CHLORHEXIDINE-IMPREGNATED DRESSING FOR PREVENTION OF CATHETER-RELATED BLOODSTREAM INFECTION: A META-ANALYSIS

Nasia Safdar, MD, PhD¹, John C. O'Horo, MD, Aiman Ghufran, MD, Allison Bearden, MD, MPH, Maria Eugenia Didier, MD, Dan Chateau, PhD, and Dennis G. Maki, MD
Section of Infectious Diseases, Department of Medicine, University of Wisconsin School of Medicine and Public Health, Madison, WI (NS, AB, MD, DGM, AG), Department of Pulmonary and Critical Care Medicine, Mayo Clinic, Rochester MN (JCO), and the Biostatistical Consulting Unit (DC), Department of Community Health Sciences, University of Manitoba, Winnipeg, MB (DC)

Abstract

 Review Article
Proc J Public Health, Vol. 44, No. 5, May 2019, pp 196-207

The Effects of Chlorhexidine Dressing on Health Care-Associated Infection in Hospitalized Patients: A Meta-Analysis

Hou-Xing WANG, Shu-Yuan XIE, Hao Wang, Hao-Kai CHU
Department of Emergency Intensive Care Unit, Zhejiang Provincial Hospital, Ningbo, China
*Corresponding Author: Email: whx0241@sina.com
(Received 21 Oct 2018; accepted 19 Jan 2019)

Abstract
Background: To assess the effects of chlorhexidine dressing on health care-associated infection in hospitalized patients.
Methods: We searched for English-language published randomized controlled trials (RCTs) in Cochrane Library, EMBASE and PubMed between January 1998 and January 2018. We used meta-analysis to calculate the risk ratios (RR) and 95% confidence intervals (CI) of the data, and using the P assessment to summarize the heterogeneity of RCTs and the forest plot and Egger regression test to evaluate publication bias.
Results: A total of 13 RCTs were included in our meta-analysis, including 7555 patients and 11,931 catheters. The effects of chlorhexidine dressing on the incidence of catheter-related bloodstream infections (CRBSI) were reported in 10 RCTs, and the incidence of CRBSI was 1.5% (80/4160) in the chlorhexidine group and 2.5% (245/9771) in the control group. We used a forest plot to determine the risk ratio (RR) of chlorhexidine dressing on the incidence of CRBSI, and our results showed that chlorhexidine dressing significantly reduced the incidence of CRBSI (RR 0.55, 95% CI 0.39-0.77, P<0.001). Moreover, we also analyzed the effects of chlorhexidine dressing on the incidence of catheter colonization and catheter-related infections (CRI), and our forest plot results showed that chlorhexidine dressing significantly reduced the incidence of catheter colonization (RR 0.52, 95% CI 0.40-0.67, P<0.001) and the incidence of CRI (RR 0.43, 95% CI 0.28-0.66, P<0.001) in hospitalized patients.
Conclusion: The use of chlorhexidine dressings for hospitalized patients significantly reduce the incidence of CRBSI, catheter colonization and CRI.
Keywords: Chlorhexidine dressing; Catheter-related bloodstream infections; Randomized controlled trials

Introduction
Central venous catheters (CVCs) are an important source of bloodstream infections (BSIs) in hospitalized critically ill patients and are closely related to patients' mortality (1). During the hospitalization, patients complicated with catheter-related bloodstream infections (CRBSIs) and/or catheter-related infection (CRI) caused their illness to worsen, the length of hospital stay was extended, and hospitalization expenses increased (2-4). According to data reported by the Centers for Disease Control and Prevention in US in 2009, the number of CRBSI in the Intensive Care Unit (ICU) was 12,000-18,000, and the medical expenses generated per case were about \$16,550, and the overall mortality rate was increased by 15%-25% (5). At present, due to the limited number of antimicrobial drugs and the emergence of multi-drug

Available at: <http://jghp.tsmu.ac.ir>

Wu et al. BMC Infectious Diseases (2019) 19:429
<https://doi.org/10.1186/s12879-019-4029-9> BMC Infectious Diseases

RESEARCH ARTICLE Open Access

Chlorhexidine-impregnated dressing for the prophylaxis of central venous catheter-related complications: a systematic review and meta-analysis

Li Wei¹, Yan Li¹, Xiaoyan Li¹, Lusheng Bian, Zunjia Wen¹ and Mei Li¹

Abstract
Background: Several randomized controlled trials (RCTs) evaluated the role of Chlorhexidine impregnated dressing for prophylaxis of central venous catheter (CVC) related complications, but the results remained inconsistent.

Infection Control & Hospital Epidemiology (2020), 44, 1288–1295
doi:10.1017/S0950268819001316

Original Article

Effectiveness of chlorhexidine dressings to prevent catheter-related bloodstream infections. Does one size fit all? A systematic literature review and meta-analysis

Mireia Puig-Asensio MD, PhD^{1,2}, Alexandre R. Marrs MD, PhD^{1,2,3,4}, Christopher A. Childs MS⁵, Mary E. Kulkas BSN, RN⁶, Eli N. Pieronecovich MD, MS^{1,4} and Martin L. Schwelzer PhD^{1,4}
University of Iowa Carver College of Medicine, Iowa City, Iowa, United States; ¹Division of Medical Practice, Hospital Israelita Albert Einstein, São Paulo, Brazil; ²Wardens Library for the Health Sciences, University of Iowa Libraries, Iowa City, Iowa, United States and ³Center for Access & Delivery Research and Evaluation (CADRE), Iowa City VA Health Care System, Iowa City, Iowa, United States

Abstract
Objective: To evaluate the effectiveness of chlorhexidine (CHG) dressings to prevent catheter-related bloodstream infections (CRBSI).
Design: Systematic review and meta-analysis.
Methods: We searched PubMed, CINAHL, EMBASE, and ClinicalTrials.gov for studies (randomized controlled and quasi-experimental trials) with the following criteria: patients with short- or long-term catheters; CHG dressings were used in the intervention group and nonantimicrobial dressings in the control group; CRBSI was an outcome. Random-effects models were used to obtain pooled risk ratios (pRR). Heterogeneity was evaluated using the I² test and the Cochran Q statistic.
Results: In total, 20 studies (18 randomized controlled trials; 15,590 catheters) without evidence of publication bias and mainly performed in intensive care units (ICUs) were included. CHG dressings significantly reduced CRBSI (pRR, 0.71; 95% CI, 0.58–0.87), independent of the CHG dressing type used. Benefits were limited to adults with short-term central venous catheters (CVCs), including once-hemodialysis patients. For long-term CVCs, CHG dressings decreased exit-site/tunnel infections (pRR, 0.37; 95% CI, 0.22–0.64). Contact dermatitis was associated with CHG dressing use (pRR, 5.16; 95% CI, 2.09–12.70); especially in neonates and pediatric populations in whom severe reactions occurred. Also, 2 studies evaluated and did not find CHG-acquired mastitis.
Conclusions: CHG dressings prevent CRBSI in adults with short-term CVCs, including patients with an oncology-related disease. CHG dressings might reduce exit-site and tunnel infections in long-term CVCs. In neonates and pediatric populations, proof of CHG dressing effectiveness is lacking and there is an increased risk of serious adverse events. Future studies should investigate CHG effectiveness in non-ICU settings and monitor for CHG resistance.
(Received 16 March 2020; accepted 18 July 2020; electronically published 16 September 2020)

During the past decade, hospitals have made significant progress in preventing catheter-related bloodstream infections (CRBSI). Basic strategies during the insertion and maintenance of catheters have successfully reduced CRBSI and have been incorporated into clinical practice.^{1–3} Despite these advances, however, CRBSI remains a problem. Recent data from the United States showed that CRBSI rates have been steady between 0.56 and 0.67 cases per 1,000 catheter days⁴ and, in European intensive care units (ICUs), rates have oscillated between 1.7 and 4.7 cases per 1,000 catheter days.⁵

A substantial number of patients experience CRBSI, a preventable healthcare-associated infection linked to increased mortality, length of hospital stay, and healthcare costs.⁶ Thus, attention has moved to other preventive measures, including the use of chlorhexidine-impregnated dressing at the catheter insertion site. Chlorhexidine (CHG) is an antiseptic with broad-spectrum antimicrobial activity.⁷ When incorporated into catheter dressings, the burden of microorganisms on the skin decreases and CRBSI caused by an extraluminal route might be reduced.⁸ Previous meta-analyses have suggested that CHG dressings can prevent CRBSI.^{9–12} However, numerous gaps in knowledge remain and tailored recommendations are lacking. It is unclear which patients benefit the most from CHG dressings, to what extent the catheter type impacts CHG effectiveness, or whether the main types of CHG dressing available (CHG-impregnated discs or transparent dressings with an integrated CHG gel pad) are equally effective. Numerous clinical trials have been published in recent years regarding CHG dressings, and it is time to re-evaluate the evidence

Author for correspondence: Mireia Puig-Asensio, E-mail: mireia-puig@uiowa.edu

ADDITIONAL INFORMATION: Results from this study were accepted to be presented at the following scientific meetings: SHEA/IDC, December 6th International Conference on Healthcare Associated Infections and the 30th European Congress of Clinical Microbiology and Infectious Diseases.

Cite this article: Puig-Asensio M, et al. (2020), Effectiveness of chlorhexidine dressings to prevent catheter-related bloodstream infections. Does one size fit all? A systematic literature review and meta-analysis. *Infection Control & Hospital Epidemiology*, 44, 1288–1295. <https://doi.org/10.1017/S0950268819001316>

© 2020 by The Society for Healthcare Epidemiology of America. All rights reserved.

Downloaded from <https://www.cambridge.org/core>. 23 Jul 2021 at 16:02:28, subject to the Cambridge Core terms of use.

- ▶ Four published meta-analyses examined the effectiveness of **chlorhexidine gluconate (CHG)** dressings on reducing colonization of catheters and CRBSI
- ▶ 20 RCTs and 2 quasi-experimental studies
- ▶ Colonization: 6.5% of catheters in groups using CHG dressings and 13.2% in the control groups
- ▶ CRBSI: 1.2% rate in CHG groups and 2.3% in control groups



ADVANCED ANTIMICROBIAL INTRAVENOUS DRESSINGS

- ▶ Study comparing the *in vitro* antimicrobial performance of a chlorhexidine free base (CHX) dressing to a chlorhexidine salt (CHA) dressing
- ▶ Aim was to identify if there are differences in magnitude, rate of action, and overall antimicrobial performance
- ▶ CHX, CHA, and control dressings were evaluated for antimicrobial performance against 12 challenge microorganisms
- ▶ Various dressings were inoculated and incubated, followed by sampling at 1-, 3-, and 7-day time periods
- ▶ Time frames were chosen to reflect the CDC's clinical practice guidelines for maintenance of IV catheters
- ▶ Benchmark: 4.0 log₁₀ reduction

Challenge microorganism species	Inoculum media	Plate media	Incubation temperature (°C)
Gram-positive bacteria			
Methicillin-resistant <i>Staphylococcus aureus</i> (MRSA), ATCC 33591	SCDB	SCDA	30–39
Methicillin-resistant <i>Staphylococcus epidermis</i> (MRSE), ATCC 51625	SCDB	SCDA	30–39
Multiple drug-resistant <i>Enterococcus faecium</i> (MDR), ATCC 51559	SCDB	SCDA	30–39
Vancomycin-resistant <i>Enterococcus faecalis</i> (VRE), ATCC 51299	SCDB	SCDA	30–39
<i>Enterococcus faecium</i> , ATCC 19434	SCDB	SCDA	35–39
Gram-negative bacteria			
<i>Pseudomonas aeruginosa</i> , ATCC 9027	SCDB	SCDA	35–39
<i>Escherichia coli</i> , ATCC 8739	SCDB	SCDA	35–39
<i>Serratia marcescens</i> , ATCC 8100	SCDB	SCDA	30–35
Yeasts			
<i>Candida albicans</i> , ATCC 10231	SDEX	SDEX	20–25
<i>Candida parapsilosis</i> , ATCC 14054	SDEX	SDEX	20–25
<i>Candida tropicalis</i> , ATCC 750	SDEX	SDEX	20–25
Fungus			
<i>Aspergillus brasiliensis</i> , ATCC 16404	SDEX	SDEX	20–25
ATCC—American type culture collection; SCDA—soybean casein digest agar; SCDB—soybean casein digest broth; SDEX—sabouraud dextrose agar			

ADVANCED ANTIMICROBIAL INTRAVENOUS DRESSINGS

Challenge organism		Starting titer (CFU/sample)	Cover film negative control mean log ₁₀ reduction				PrevahexCHX mean log ₁₀ reduction			SurgiClear mean log ₁₀ reduction			Mean log ₁₀ reduction colour temperature scale
			0-Hour	1-Day	3-Day	7-Day	1-Day	3-Day	7-Day	1-Day	3-Day	7-Day	
Gram (+) Bacteria	MRSA	2.4 x 10 ⁶	0	0.66	1.51	2.65	6.05	6.18	6.38	3.56	4.43	2.76	
	MRSE	1.3 x 10 ⁶	-0.18	2.96	1.71	1.97	5.93	6.02	6.1	3.52	5.12	6.1	-2
	<i>E. faecium</i> (MDR)	1.1 x 10 ⁶	-0.04	-0.11	2.41	2.55	6.09	5.63	5.67	2.66	4.69	3.21	-1
	<i>E. faecalis</i> (VRE)	1.8 x 10 ⁶	0	-0.09	4.01	2.4	6.24	6.21	6.44	2.77	4.72	6.44	0
	<i>E. faecium</i>	4.3 x 10 ⁶	-0.01	1.16	2.94	3.71	6.42	6.07	6.55	3.11	6	6.51	1
Gram (-) Bacteria	<i>P. aeruginosa</i>	2.0 x 10 ⁶	-0.05	-1.22	-0.92	1.48	6.3	6.26	6.3	5.21	6.3	6.17	2
	<i>E. coli</i>	4.3 x 10 ⁶	0.01	-0.79	-0.13	3.86	6.22	6.64	6.64	6.64	6.64	6.64	3
	<i>S. marcescens</i>	8.6 x 10 ⁶	-0.01	2.89	4.63	4.63	6.28	6.93	6.46	5.1	6.93	6.93	4
Yeasts	<i>C. albicans</i>	1.2 x 10 ⁶	-0.04	0.14	1.85	3.77	6.07	6.07	6.07	2.23	2.74	6.07	5
	<i>C. parapsilosis</i>	3.3 x 10 ⁶	-0.04	-0.04	-0.21	-0.12	5.11	5.91	5.79	0*	1.64	2.73	6
	<i>C. tropicalis</i>	4.7 x 10 ⁶	0.03	0.17	1.26	1.31	6.49	6.58	6.5	2.69	3.32	3.95	7
Fungus	<i>A. brasiliensis</i>	3.7 x 10 ⁶	0.04	0.41	0.46	0.5	2.99	3.89	3.75	1.72	1.67	1.74	

Fig 1. Summary of the mean *in vitro* log₁₀ reduction values of the two antimicrobial dressing types and the polymer cover film control at 1-, 3-, and 7-day time points. Colour-temperature indicates magnitude of microbial log₁₀ reduction observed. Mean *in vitro* log₁₀ reduction values ≥4.0 log₁₀ are presented in shades of green which increase in darkness with magnitude, while log₁₀ reduction values <4.0 log₁₀ transition from shades of light yellow to red with decreasing magnitude. *Actual experimental value was 'too numerous to count' (TNTC) and is thus presented here as '0.00' for data analysis purposes since the TNTC outcome indicates no organism reduction and the approximation of this value as '0.00' is likely to be a conservative estimate for the actual value

- ▶ **Magnitude:** The CHX dressing demonstrated a superior *in vitro* antimicrobial effect at 67% of the experimental time points than the CHA dressing, with at least equivalent efficacy at all other time points
- ▶ **Rate of Action:** the CHX dressing had a more rapid action than the CHA dressing particularly at the 1-day time point; the CHX dressing achieved a >5 log₁₀ reduction at the 7-day time point, whereas the CHA dressing demonstrated such reduction in only seven test organisms
- ▶ **Chlorhexidine content:** CHA adhesive film has **36% greater chlorhexidine mole content**. So why was it not as effective as CHX?

Advanced Antimicrobial Intravenous Dressings

CDC: For patients aged 18 years and older: **Chlorhexidine-impregnated dressings** with an FDA-cleared label that specifies a clinical indication for reducing catheter-related bloodstream infection (CRBSI) or catheter-associated blood stream infection (CABSI) are recommended to protect the insertion site of short-term, nontunneled central venous catheters. (Category IA)

INS: Change transparent semipermeable membrane (TSM) dressings at **least every 7 days** (except neonatal patients) or immediately if dressing integrity is disrupted (eg, lifted/detached on any border edge or within transparent portion of dressing; visibly soiled; presence of moisture, drainage, or blood) or evidence of compromised skin integrity under the dressing, and following manufacturer's instruction for use. (III)

Infusion Therapy Standards of Practice

Barbara Nicolai, APRN-CNS, CCRN, CRNI®
Lisa Gorski, MS, RN, HMCNS-BC, CRNI®, FAAN
Tritia Kleidon, PhD(c), MNSc, RN
Amy Kyles, MSN, APRN, AG-CNS, CV-BC™, CRNI®
Michelle DeVries, MPH, CIC, VA-BC, CPHQ, FAIC
Samantha Keogh, PhD, RN, FAAN
Britt Meyer, PhD, RN, CRNI®, VA-BC, NE-BC
Mary Jo Sarver, MSN, ARNP, AOCN, CRNI®, LNC, VA-BC
Rachael Crickman, DNP, ARNP-CNS, AOCNS, OCN, RN
Jenny Ong, PharmD
Simon Clare, MRes, BA, RGN
Mary E. Hagle, PhD, RN-BC, FAAN

9TH EDITION
REVISED 2024

INS

INFUSION NURSES SOCIETY
11700 THE UNIVERSITY DRIVE
One Edgewater Drive, Norwood, MA 02062
www.ins1.org

Accessible version: <https://www.cdc.gov/infectioncontrol/guidelines/hai/c-dressings/index.html>



2017 Updated Recommendations on the Use of Chlorhexidine-Impregnated Dressings for Prevention of Intravascular Catheter-Related Infections

Centers for Disease Control and Prevention
National Center for Zoonotic and Emerging Infectious Diseases
Division of Healthcare Quality Promotion

Thomas R. Talbot III, MD, MPH; Erin C. Stone, MA; Kathleen Irwin, MD, MPH; Amanda D. Overholz, MPH; Mahasa Dast, MPH; Alexander Kalkan, MD, MPH, for the Healthcare Infection Control Practices Advisory Committee*

*Vanderbilt University School of Medicine, Vanderbilt University Medical Center, †Division of Healthcare Quality Promotion, National Center for Emerging and Zoonotic Infectious Diseases, Centers for Disease Control and Prevention, ‡Northrop Grumman Corporation, †Time Solutions LLC, and †Contributing IHC PAC Members

Suggested citation: Centers for Disease Control and Prevention. 2017 Recommendations on use of chlorhexidine-impregnated dressings for prevention of intravascular catheter-related infections: An update to the 2011 guidelines for the prevention of intravascular catheter-related infections from the Centers for Disease Control and Prevention, Centers for Disease Control and Prevention, National Center for Emerging and Zoonotic Infectious Diseases, Division of Healthcare Quality and Promotion. DATE. (DHQP website URL to be added)

Disclosures and disclaimers: The authors report no actual or potential conflicts of interest related to the commercial products discussed herein. This document is not protected by the Copyright Act, and copyright ownership cannot be transferred. It may be used and reprinted without special permission.

Last update: July 17, 2017

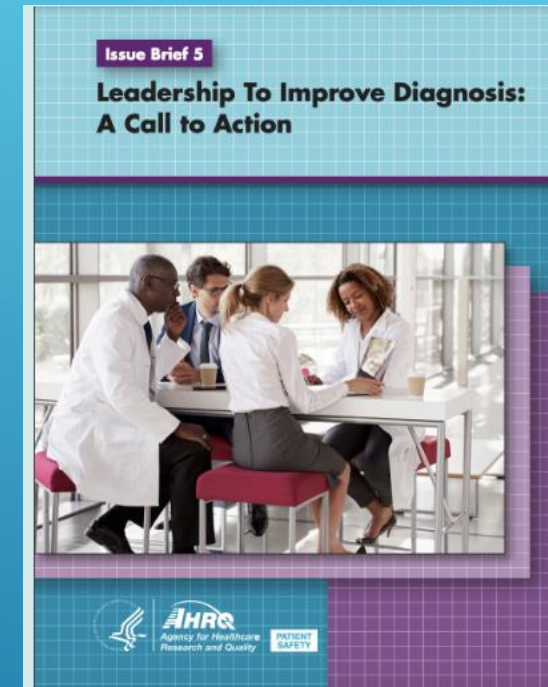
Page 1 of 15



EMERGING PERSPECTIVES: IMPLEMENTATION ASPECTS

Leadership

- ▶ Successful outcomes in IP programs will largely depend on dedicated support provided by hospital executives, managers, and clinical leaders
- ▶ Due to the complexity in controlling HOB in all VADs, several key management practices will need to be included:
 1. setting of **organizational IP priorities** whereby goals are established and shared among managers and front-line staff
 2. establishing an **information-sharing system** within the organization that allows relevant infection data to be relayed, displayed, and discussed in a timely manner with clinicians and those performing direct patient care, and
 3. provides **management coaching** activities that include staff feedback and re-education sessions emphasizing best practices for IP
- ▶ Within IP programs, directors will need to ensure they are adding additional VAD risks to their annual infection control risk assessment. Based on this information, there will be a need to analyze staffing structures, including the skill level of IP staff



Staffing



A potential expansion of reportable LabID events will inevitably trigger the opportunity for prevention of other device-related BSIs



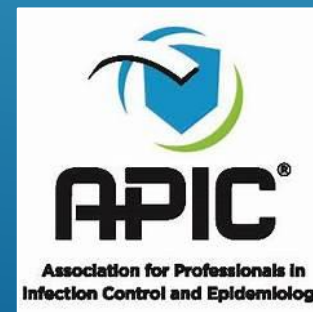
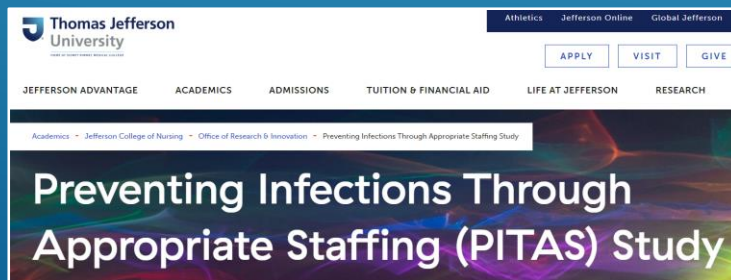
The evolving role of the infection preventionist coupled with national events affecting the healthcare industry provides substantial affirmation for enhancing IP and control resources



The number of IP personnel for required and effective prevention functions in healthcare institutions needs re-evaluation based on increasing roles and responsibilities.



Studies contributing to a better understanding of resource development: MegaSurvey conducted by APIC, the Preventing Infections Through Appropriate Staffing (PITAS) survey



WHAT IS DIAGNOSTIC STEWARDSHIP?

- ▶ **Diagnostic Stewardship** is a coordinated system or user-based interventions designed to promote evidence-based utilization of diagnostic tests, with the primary goals of improving value and care quality and safely reducing costs
- ▶ DS involves modifying the process of ordering, performing, and reporting diagnostic tests in order to direct appropriate antimicrobial therapy
- ▶ The Microbiology laboratory provides information that identifies if a patient is infected, what the pathogen is, and which antibiotics may be effective in treatment of true infection

KEY CONSIDERATIONS

DS

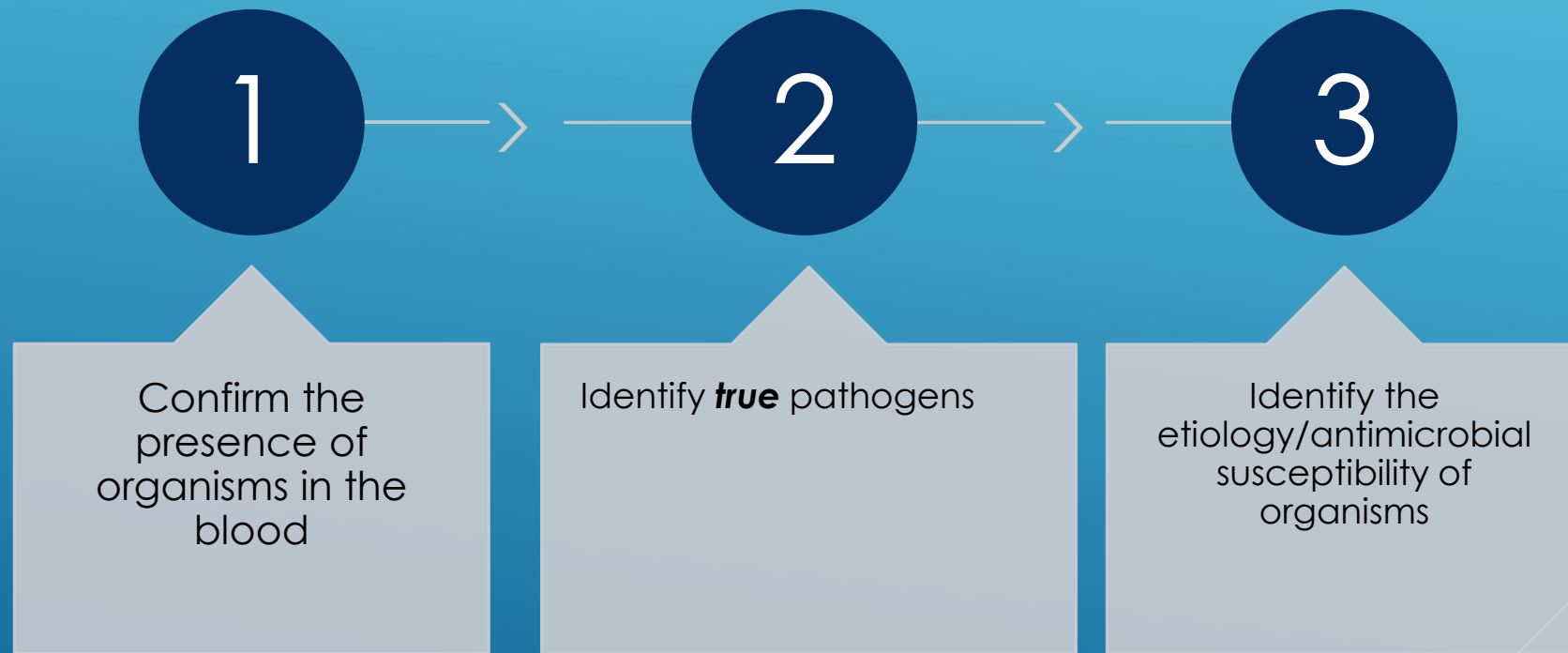
Goal	Key question	Key considerations and potential strategies
Right test	Is the test appropriate for the clinical setting?	Sensitivity and specificity Predictive values Testing volumes Diagnostic yield Laboratory feasibility Cost Clinical impact
Right patient	Will the clinical care of the patient be affected by the test result?	Laboratory test utilization committee Automatic laboratory reflex CPOE decision support Appropriate use criteria Indication selection Prior authorization Benchmarking Specimen rejection
Right time	Will the result be available in time to optimally affect care?	Time to specimen receipt Centralized vs point-of-care testing On-demand vs batched testing Specimen preparation time Run time Result reporting time

AS

Goal	Key question	Key considerations and potential strategies ^a
Right interpretation	Will the clinician understand the test result?	Result report language Selective reporting of relevant results AS prospective audit and feedback AS real-time decision support
Right antimicrobial	Will the clinician appropriately modify antimicrobials based on the test result?	Clinical practice guidelines EMR-based decision support with result reporting AS prospective audit and feedback AS real-time decision support
Right time	Will the clinician act upon the test result promptly?	EMR reporting Results called with readback reporting AS prospective audit and feedback AS real-time decision support

^aAS, antimicrobial stewardship.

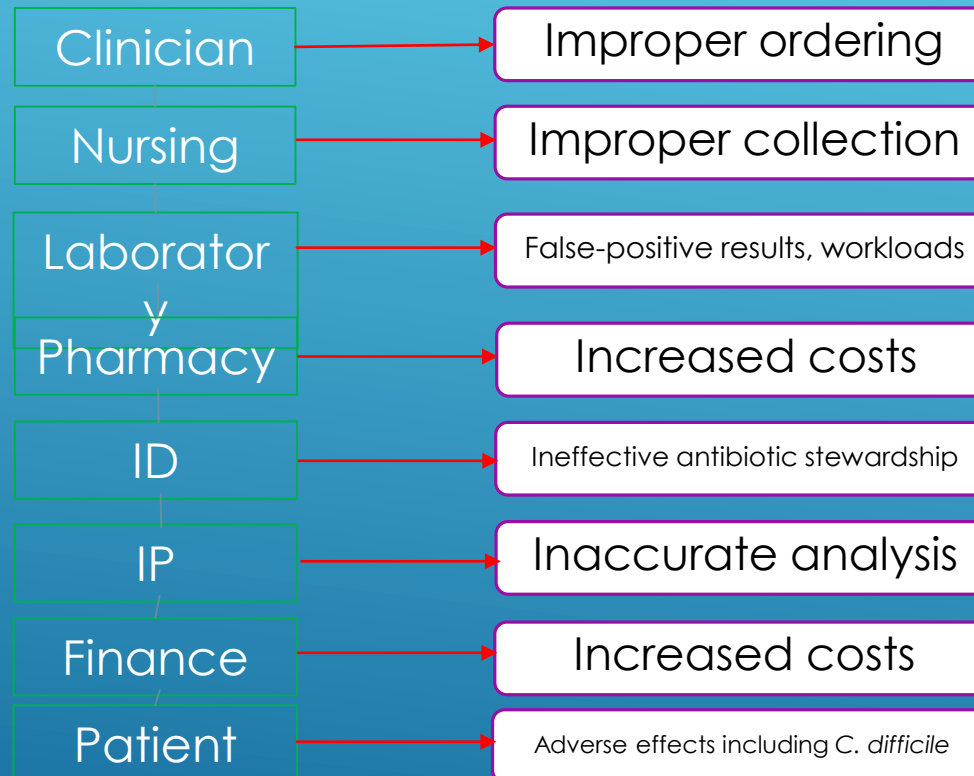
THE PURPOSE OF BLOOD CULTURES



THE EFFECTS ON HEALTHCARE WHEN PROPER BLOOD CULTURE MANAGEMENT IS NOT IMPLEMENTED

- Systematic review on costs (BC Contamination):
- Pharmacy:
 - \$210-\$12,611
 - Labs:
 - \$2397-\$11,151
 - Hospital costs:
 - \$16,200-\$111,627
 - LOS: 1-22 days

Dempsey C, et al.
Economic health care costs of BC contamination, AJIC 2018



PRINCIPAL INTERVENTION # 1

▶ Implementing Evidence-Based Decision Aids

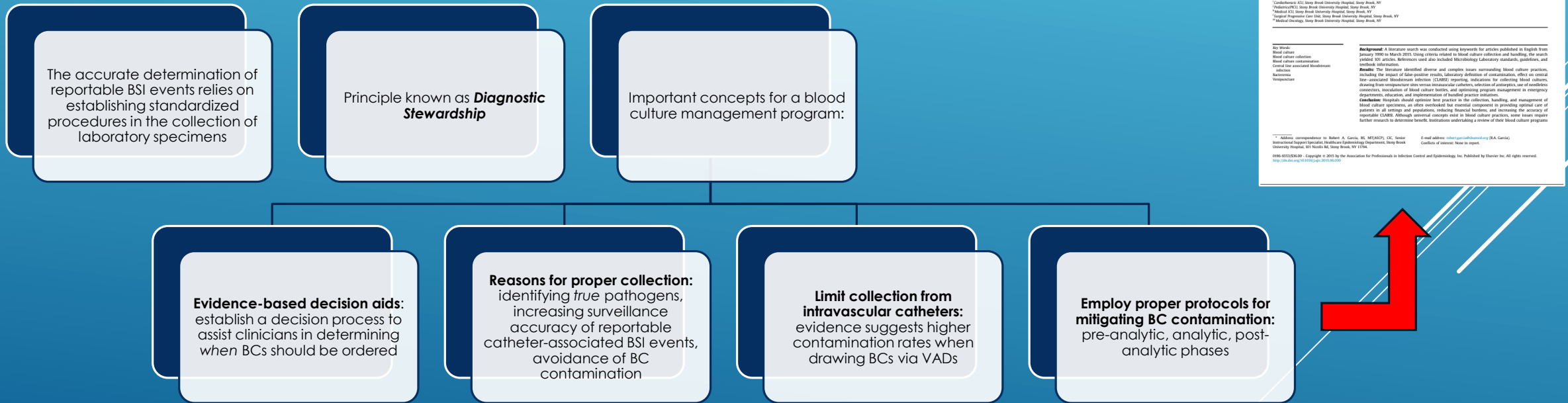
- ▶ An initial, but critical, consideration in optimizing the accuracy of BC results related to VAD HOB surveillance is the establishment of a decision process to assist clinicians in determining *when* BCs should be ordered
- ▶ A 2021 survey of US acute care hospitals to determine prevalence of technical interventions aimed at preventing central line-associated bloodstream infection (CLABSI), indicated that only 23% of respondents reported strategies to reduce routine BCs.
- ▶ There is compelling evidence however, that such interventions improve clinical decision-making when considering BC orders. In the DISTRIBUTE (DIagnostic STewardship Improves Blood cULTures) quality improvement study, an algorithm for ordering BCs was provided to clinicians in conjunction with education concerning the avoidance of orders for solitary BCs. Participants received feedback on BC rates and the appropriateness of their decisions.
- ▶ The algorithm specified that if the BC is being ordered for a new clinical event, BCs are recommended based on the probability of bacteremia (high, moderate, or low). BCs were not recommended for scenarios with a low probability of bacteremia. In this study, BC best practices in a medical intensive care unit (ICU) and medicine wards at a large academic center **reduced BC utilization by 18% and 30%, respectively.**

Pisney L, Campese L, Greene MT, et al. Practices to prevent central line-associated bloodstream infection: A 2012 survey of infection preventionists in US hospitals. *Infect Control Hosp Epidemiol* 2024; Published online 2024:1-5. doi:10.1017/ice.2024.53 Fabre V, Klein E, Salinas AB, et al. A diagnostic stewardship intervention to improve blood culture use among adult nonneutropenic inpatients: The DISTRIBUTE Study. *J Clin Micro* 2020;58:1-8.

PRINCIPAL INTERVENTION #2

- ▶ **Standardizing Evidence-Based Methods for Proper Collection of BCs**
- ▶ Hospitals should review and standardize how BCs are collected in consideration of the primary benefits that can be achieved and its relevance to conducting accurate surveillance for VAD HOB: the recovery of true pathogens (i.e., avoidance of false-negative BCs), increasing the surveillance accuracy of catheter-associated BSI events, and a voidance of blood culture contamination (BCC).
- ▶ BCC triggers a cascade of detrimental consequences that creates further issues of global importance to hospitals: improper antimicrobial treatment of the patient potentially leading to adverse drug reactions, emergence of antibiotic-resistant organisms (AROs), prolonged venous access increasing risk for associated infection, disruption of the natural microbiome leading to *Clostridioides difficile* infection, additional unnecessary testing, increased laboratory and pharmaceutical costs, and lost reimbursement and financial penalties as may occur with misinterpretation of reportable HAIs.

Diagnostic Stewardship & Blood Culture Management



PRINCIPAL INTERVENTION #3

- ▶ **Diversion of the First Portion of Blood Theoretically Removes the Contaminating Organism from the Remaining Aliquot of Blood**
 - ▶ Efforts have been made to implement BCC reduction strategies that unify proper drawing techniques and novel engineering controls. Studies cited in a systematic review and meta-analysis on blood diversion interventions indicate the two most frequently used devices used in addressing BCC are commercially available products or a lithium heparin waste tube. Use of these items across nine studies resulted in reduced BCC rates ranging from 0.0% to 2.6%.
 - ▶ Regardless of the type of device used, the meta-analysis indicated that using a separate item to divert the initial aliquot of blood during venipuncture was associated with a significant reduction in BCC rate compared with a standard procedure of collection.
 - ▶ Current guidelines advocate a new BCC baseline of $\leq 1\%$

GUIDELINES ADDRESSING DIVERSION

CDC Division of Laboratory Systems
EXCELLENT LABORATORIES, OUTSTANDING HEALTH

Preventing Adult Blood Culture Contamination: A Quality Tool for Clinical Laboratory Professionals

Protect Patients during the Diagnostic Process by Monitoring Adult Blood Culture Contamination (BCC) Rates

Laboratory analysis of blood cultures is vital to the accurate diagnosis of infectious diseases. The reliability of your testing depends on clinical compliance with best practices. Inconclusive or incorrect results, false negative blood culture results, misdiagnosis, delay therapy, and put patients at heightened risk of false positives, compromising care by leading to unnecessary treatment. In December 2022, a Centers for Medicare & Medicaid Services proposal for a new patient safety measure to address these topics. CDC developed this quality measure to promote best practices for blood culture collection.

The Clinical Laboratory Improvement Amendments of 1988 when indicated, correct problems identified in their preana tool to calculate the BCC and single-set rates will help meet in addition, this quality measure incorporates best practice Standards Institute (CLSI) and the Infectious Diseases Society of America (IDSA) and the American Society for Microbiology (ASM) place at many laboratories across the nation and have shown significantly reduce incidence of BCC, and limit unnecessary; adopt these practices into your laboratory's standard operating quality management system, and to work with infection control staff on their use.

Follow CLIA Regulations
"Laboratory Requirements," Code of Federal Regulations, Title 42 Part 431 - Quality System for Non-Waived Testing - § 493.124
The laboratory must establish and follow written policies and procedures to correct problems identified in the preanalytic systems specified in 493.124.

Collecting Adult Blood Culture Sets
A blood culture set from an adult patient should consist of one or two bottles, depending on the volume of blood collected.

Collect Multiple Sets to Achieve the Optimal Volume
The volume of blood collected is critically important to the generally requires two or more sets to achieve. In addition, of a commensal organism can be classified as a possible co-infection.

To achieve an optimal volume, the blood culture collection sets from adult patients with a suspected blood stream infection (BSI). Your hospital or clinical setting should instruct health volume of 40-60 mL within a 24-hour period by peripheral venipuncture.

Find this information at: <https://www.cdc.gov/lab-standards/2023-01-10-preventing-adult-blood-culture-contamination>



CLINICAL AND LABORATORY STANDARDS INSTITUTE®

M47

Principles and Procedures for Blood Cultures

2nd Edition

This guideline includes recommendations for collecting, transporting, and processing specimens for blood culture, as well as procedures for recovering pathogens from the blood of patients with suspected bacteremia or fungemia.

A guideline for global application developed through the Clinical and Laboratory Standards Institute consensus process.



Clinical Infectious Diseases

IDSA GUIDELINES

IDSA Infectious Diseases Society of America | **hivma** HIV Medicine Association | **OXFORD**

Guide to Utilization of the Microbiology Laboratory for Diagnosis of Infectious Diseases: 2024 Update by the Infectious Diseases Society of America (IDSA) and the American Society for Microbiology (ASM)*

J. Michael Miller, Matthew J. Brzezinski, Shelton Campbell, Karen C. Carroll, Katherine C. Chappell, Mark D. Gonzalez, Amanda Harrington, Robert C. Jurek, Scott A. Koenig, Klaus H. Lee, J. Robin Penell, Robert S. Plim, Sandy S. Rickman, Barbara Robinson-Dunn, James W. Snyder, Sanjiv S. Talwar, Erica S. Tarr, Richard S. Tammen, Jr., Melissa P. Williams, and Joseph B. Wolf

*Approved 18 February 2024, editorial decision 17 February 2024, published online 5 March 2024. Correspondence: J. M. Miller, Microbiology Technical Services, LLC, PO Box 880232, Columbus, GA 31922 (jmmiller@idsa.org).

This critical nature of the microbiology laboratory in infectious disease diagnosis calls for a close, positive physician and the microbiologist who provide enormous value to the healthcare team. This document, and pediatric laboratory and clinical medicine, provides information on which tests are valuable and to add little or no value for diagnostic decisions. Sections are divided into anatomic systems, including Blood of the Cardiovascular System, Central Nervous System Infections, Ocular Infections, Soft Tissue Infection, Respiratory Infections, Lower Respiratory Tract Infections, Infections of the Gastrointestinal Tract, Joint Infections, Urinary Tract Infections, Central Infections, and Skin and Soft Tissue Infections, and non-anatomic Infections, Viral Syndromes, and Blood and Tissue Parasite Infections. Each section contains key points, and detailed tables that list suspected agents, the most reliable tests to order, the samples preferred, specimen transport devices, procedures, times, and temperatures, and detailed notes on methods, such as when tests are likely to require a specialized laboratory or have prolonged to pediatric needs of specimen management are also addressed. There is redundancy among the tables many choices overlap. The document is intended to serve as a reference to guide physicians in diagnose infectious diseases in their patients.

Keywords: specimen quality; diagnostic accuracy; physician lab interface; optimizing results

© 2024 Infectious Diseases Society of America. All rights reserved. No part of this guide may be reproduced, distributed, or transmitted in any form, or by any means, electronic, mechanical, photocopying, recording, or by any information storage or retrieval system, without the prior written permission of the Infectious Diseases Society of America. All rights reserved. For more information, please contact the Infectious Diseases Society of America at <https://www.idsa.org>.

Clinicians and Microbiologists (Patients)

Infusion Therapy Standards of Practice

Barbara Nickel, APRN-CNS, CCRN, CRNI®
Lisa Gorski, MS, RN, HHCHS-BC, CRNI®, FAAN
Tricia Kleidon, PhD(c), MNISC, RN
Amy Kyles, MSN, APRN, AG-CNS, CV-BC™, CRNI®
Michelle DeVries, MPH, CIC, VA-BC, CPHQ, FAPIC
Samantha Keogh, PhD, RN, FACH
Britt Meyer, PhD, RN, CRNI®, VA-BC, NE-BC
Mary Jo Sarver, MSN, ARNP, AOCN, CRNI®, LNC, VA-BC
Rachael Crickman, DNP, ARNP-CNS, AOCNS, OCN, RN
Jenny Ong, PharmD
Simon Clare, MRes, BA, RGN
Mary E. Hagle, PhD, RN-IB, FAAN

9TH EDITION
REVISED 2024

INS
INFUSION NURSES SOCIETY
SETTING THE STANDARD FOR INFUSION CARE®
One Edgewater Drive, Norwood, MA 02062
www.ins1.org



Systems and Human Factors Engineering

As IP science “...moves from evidence generation to translation into practice, an effective framework to promote, facilitate, and evaluate the implementation of evidence-based strategies in IP is needed”

Human factors engineering (HFE) considers the *complexity* of modern healthcare (e.g., the number of VAD inserters, types of catheters, catheter access) and the *ambiguity* or uncertainty of the system (e.g., the skill levels of VAD inserters, compliance with elements of a prevention bundle)

HFE “...is the scientific discipline concerned with understanding the interactions among humans and other elements of a system in order to improve system performance and well-being”

Approaches for Prevention



How the infection preventionist **translates** knowledge as contained in published guidelines into **actual** practice is fundamental in achieving successful outcomes



Tiered BSI prevention approaches, such as those that consider the “**lifecycle**” of a VAD, start by emphasizing high-quality, low-intensity, and lower cost elements; if these fail to lower infections, interventions requiring more resources and human capital are then introduced



Success in application of practices and interventions using IP implementation concepts and frameworks varies widely depending on **organization factors** such as operational support, informatics resources, experience, willingness to change, and safety culture



Bartles study, CLABSI reduction, 11 hospitals: included a pre-assessment, w/onsite interviews and observations, communicating potential root causes of infection management and staff based on assessment findings, institution of Lean and Six Sigma models to develop and implement a targeted intervention...result: a network-wide **70% reduction** in infection rate

The Art and Science of Infusion Nursing

Using a Comprehensive On-Site Assessment Process to Reduce Central Line-Associated Bloodstream Infection Rates

Rebecca Bartles, DPH, MPH, CIC, IDMC • Andrea Moore, ML, RH, CPHQ, CCRN-K • Rosemary Martin, ASCP (MICA, CLSBB), CIC • Rebecca Clarkson, RH, MHI, CIC • Laura Ehlinger, CIC

ABSTRACT
Central line-associated bloodstream infection (CLABSI) rates increased substantially in the United States following the emergence of COVID-19 and subsequent surges. The pandemic resulted in hospital capacities being exceeded and crisis standards of care being implemented for sustained periods. As COVID-19 rates in the United States began to stabilize, some facilities did not return to previous CLABSI rates, indicating a change in practices that had a longer-term impact on CLABSI prevention. The authors' large health care system observed similar increases in CLABSI following the emergence of COVID-19, prompting investigation and intervention in the form of a quality improvement project. To identify changes related to ongoing increases in CLABSI, an assessment team conducted standardized on-site assessments at 11 facilities. Site assessments were considered an intervention, as they involved

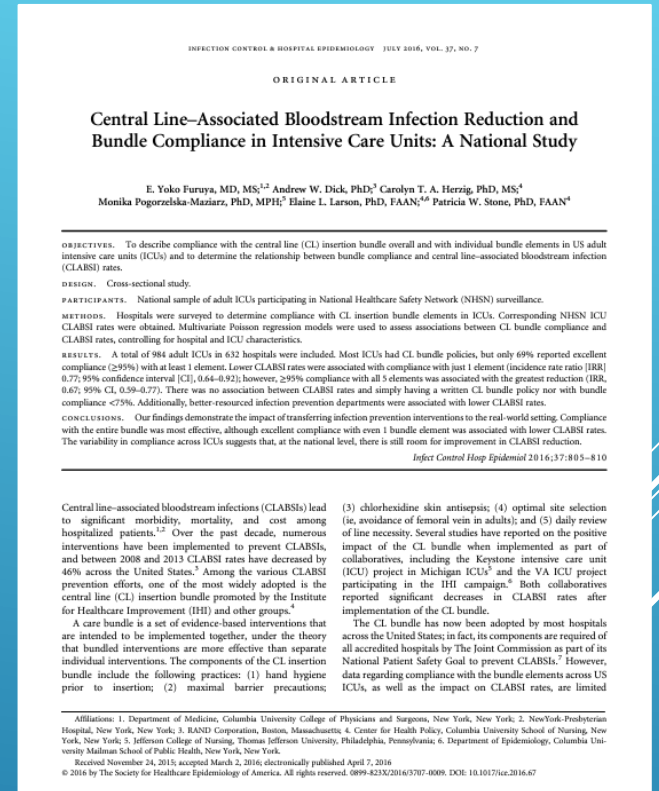
Author Affiliation: Providence, Maple Valley, Washington.
Rebecca Bartles, DPH, MPH, CIC, IDMC, is the executive director of infectious disease management and prevention for Providence. Dr. Bartles has practiced infection prevention for the last 17 years in a variety of health care settings and has numerous publications focused on infection prevention staffing and endoscope safety. She received both her BS in Public Health, Health Education, and her MPH in Epidemiology from East Tennessee State University. She completed her Doctorate in Public Health in 2021 with a dissertation topic of "Measuring Efficacy of an Evidence-Based Chlorhexidine Glucate Screening Tool Using Electronic Medical Record Data." Dr. Bartles also teaches courses at the University of Providence for a Master's in Infection Prevention degree program that she founded in 2016. She has been CIC certified since 2008 and is an Association for Professionals in Infection Control and Epidemiology fellow. Most notably, though, Dr. Bartles is the mother of 4 beautiful daughters, ages 8 to 23. Andrea Moore, ML, RH, CPHQ, CCRN-K, is a senior program manager for nursing practice and quality at Providence Health System. She completed her nursing degree at Georgetown University and her Master's in Nursing from the University of Washington. She holds credentials in both critical care nursing and health care quality. She has over a decade of experience in various health care settings. Currently she works in close partnership with interdisciplinary teams across Providence to drive system-wide quality and practice strategic initiatives. Her work focuses on reduction of nurse sensitive quality indicators, optimizing nursing workflows, and leading practice change to ensure alignment with the latest evidence-based care standards. Rosemary Martin, ASCP (MICA, CLSBB), CIC, is the system infection prevention program manager for Providence Health System, a 54-hospital system across 10 states. She has 30 years of experience as a certified clinical micro-biologist and process improvement consultant. Her projects currently focus on developing a process improvement program for system infection prevention and creating dashboards for the program that aggregates various data at the system, regional, and local level. She provides support to the system team as a subject matter expert in leveraging technological platforms for governance and team management. She holds a degree in microbiology from the University of Washington, is certified as an Infection Preventionist (CIC), Clinical Microbiologist (ACM/CM), and Lean Six Sigma Black Belt (CLSBB). Rebecca Clarkson, RH, MHI, CIC, is a senior infection preventionist with Providence Health System. She is a registered nurse by background and has practiced since 2009 in multiple settings across the hospital, including emergency medicine, cardiac, oncology, acute observation, and charge nurse. Her notable experience includes work in home infusions, business consultation, advancement, and annual educational presentations, curriculum development, and writing set compliance officers. Ms. Clarkson received her MHI certification in 2019, an emphasis in nursing education with an intent to teach nursing school when she first joined Providence in 2013 and realized that this is where she belongs. Her publications and national presentations revolve around topics related to central line-associated bloodstream infection and catheter-associated urinary tract infection prevention. Operating her passion and dedication to patient safety, Laura Ehlinger, CIC, is a system infection preventionist for Providence Health System. She has practiced infection prevention and epidemiology for the last 17 years in several health care settings. She received her BS in psychology through the University of Washington. Ms. Ehlinger worked as an instructional designer for 4 years for a private university creating online modules for master-level programs. She also has her own infection prevention consulting business, which provides consultative expertise to health care facilities and which provides practical and efficient work processes.

The authors have no conflicts of interest to disclose.
Corresponding Author: Rebecca Bartles, DPH, MPH, CIC, IDMC, Infectious Disease Management and Prevention, Providence, 2228 28th Street, Maple Valley, WA 98149 (rebecca.bartles@providence.org). DOI: 10.1097/NAI.0000000000000512

246 Copyright © 2023 Infusion Nurses Society. Journal of Infection Nursing
Copyright © 2023 Infusion Nurses Society. Unauthorized reproduction of this article is prohibited.

Bundle Compliance

- ▶ Drafting, dissemination, and education of IP protocols alone is often insufficient in achieving sustained improvements
- ▶ Long-term success requires measuring the level of **compliance with individual components** that comprise intervention bundles, followed by identifying barriers related to specific elements that are deemed below acceptable standards
- ▶ This contention is well supported in a study conducted in 984 ICUs whereby reductions in CLABSI events were related when a $\geq 95\%$ compliance level with bundle components was reached rather than related to bundle implementation
- ▶ **Data in, Data out.** Documentation in the EMR is crucial for all procedures of a bundle



Data Comprehension and Practice Change

Despite effective audit and feedback of quality data, inconsistencies exist in responses related to clinician behavior.

One goal in program modification should be to increase the **Comprehension** of relevant concepts, e.g., infection rates such as SIRs

Achieving such a goal has potential policy relevance by aiding efforts to make quality metrics more effective in influencing medical decision-making and promoting necessary practice changes

In one study, researchers used an 11-item comprehension instrument that contained questions related to metric assessment (e.g., which is better: a higher or lower SIR?), which helped identify specific factors that when modified into a comprehension scale may prove to be most relevant in driving practice change

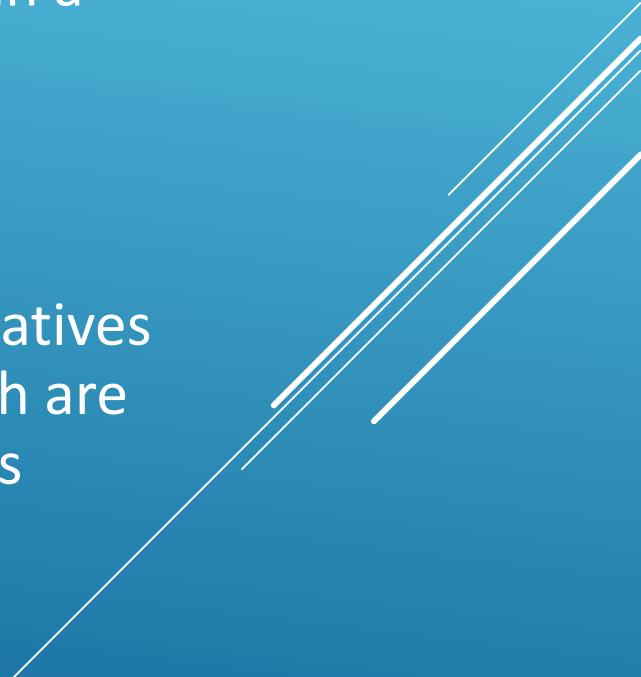


Education of VA Specialists and Infection Preventionists

- ▶ Important education points:
 - ▶ Research shows an inverse relationship between experience and complication rates
 - ▶ To address patient safety and liability concerns, standardized training practices and tools must be developed to ensure practitioner competency with safe practices during invasive procedures
 - ▶ VAS need to participate in HAI Prevention Committees
- ▶ VAS and IPs may obtain certification via Vascular Access Certification Corp
- ▶ The Infusion Nurses Society Certified Registered Nurse Infusionist (CRNI) certification



Conclusion

- The emerging perspectives presented today outline IP and VA topics supported by recent research relevant to the prevention of BSIs in a surveillance setting that includes expanding efforts to all VADs
 - The areas of focus provide insights on potential new avenues of intervention that when integrated into quality improvement initiatives should prove to be beneficial in mitigating VAD HOB events which are associated with serious, and often life-threatening, complications
- 

References

- ▶ CDC NHSN HAI Progress Reports. [Current HAI Progress Report | HAI | CDC](https://www.cdc.gov/hai/data/portal/progress-report.html)
- ▶ The Great Resignation is over, so why doesn't it feel like it? <https://www.advisory.com/daily-briefing/2024/01/26/great-resignation>he 'Great Resignation' is over. So why doesn't it feel like it? (advisory.com)
- ▶ Centers for Medicare and Medicaid Services. FY 2023 Hospital Inpatient Prospective Payment System (IPPS) and Long Term Care Hospitals (LTCH PPS) Proposed Rule - CMS-1771-P. <https://www.cms.gov/newsroom/fact-sheets/fy-2023-hospital-inpatient-prospective-payment-system-ipps-and-long-term-care-hospitals-ltch-pps>.
- ▶ McAlearney AS, Gaughan AA, DePuccio MJ, MacEwan SR, Hebert C, Walker DM. Management practices for leaders to promote infection prevention: lessons from a qualitative study. *Am J Infect Control.* 2021;49:536-541.
- ▶ Agency for Healthcare Research and Quality. Leadership to improve diagnosis: A call to action. AHRQ Publication No. 20(21)-0040-5-EF, June 2021. <https://www.improvediagnosis.org/publications/improvedx-july-2021/from-the-field-coalition-member-ahrq-calls-for-focus-on-healthcare-leadership-to-improve-diagnostic-safety/#:~:text=The%20Agency%20for%20Healthcare%20Research%20and%20Quality%20%28AHRQ%29,healthcare%20leaders%20can%20drive%20improvements%20in%20diagnostic%20safety>
- ▶ Madden GR, Weinstein RA, Sifri CD. Diagnostic Stewardship for healthcare-associated infections: opportunities and challenges to safely reduce test use. *Infect Control Hosp Epidemiol.* 2018;39:214-218.
- ▶ Garcia R, Spitzer E, Beaudry J, Beck C, DiBlasi R, Gilleeny-Blabac M, et al. Multidisciplinary team review of best practices for collection and handling of blood cultures to determine effective interventions for increasing the yield of true-positive bacteremias, reducing contamination, and eliminating false-positive central line-associated bloodstream infections. *Am J Infect Control.* 2015;43:1222-1237.

References

- ▶ Drews FA, Visnovsky LC, Mayer J. Human Factors Engineering contributions to infection prevention and control. *Human Factors*. 2019;61:693-701.
- ▶ Bartles R, Moore A, Martin R, Clarkson R, Ebinger L. Using a comprehensive on-site assessment process to reduce central line-associated bloodstream infection rates. *Infusion Nurses Soc*. 2023;46:266-271.
- ▶ Futuya EY, Dick AW, Herzig CT, Pogorzelska-Maziarz M, Larsen E, Stone P. Central line-associated bloodstream infection reduction and bundle compliance in intensive care units: a national study. *Infect Control Hosp Epidemiol*. 2016;37:805-810.
- ▶ Govindan S, Iwashyna TJ. Do clinicians understand quality metric data? An evaluation in a Twitter-derived sample. *J Hosp Med*. 2017;12:18-22.
- ▶ Huang SS, Septimus E, Platt R. Targeted decolonization to prevent ICU infections. *N Engl J Med*. 2014;369:1470-1471.
- ▶ Summerlin-Long S, DiBlase LM, Padgett M, Mack J, Clark C, Teal L, et al. A helping hand: the impact of a central line insertion support team. *Am J Infect Control*. 2023. Online ahead of print. <https://www.ajicjournal.org/action/showPdf?pii=S0196-6553%2823%2900609-06090.pdf>.
- ▶ Vascular Access Certification Corporation. *Vascular Access Certification and Recertification Candidate Handbook 2023*. <https://www.vacert.org>.
- ▶ Infusion Nurses Certification Corporation. *CRNI Exam*. <https://www.ins1.org/crni/exam/overview>.
- ▶ Mastrandra G, Giuliani R, Graps EA. International good practices on central venous catheters' placement and daily management in adults and on educational interventions addressed to healthcare professionals or awake/outpatients. Results of scoping review compared with the existence Italian good practices. *Front Med*. 2022;9:943164. doi: 10.3389/frmed.2022.943164.
- ▶ Timsit JF, Bouadma L, Ruckly S, Schwebel S, Garrouste-Orgeas M, Bronchard R, et al. Dressing disruption is a major risk factor to catheter related infections. *Crit Care Med*. 2012;40:1707- 1714.

References

- ▶ Brescia F, Pitirutti M, Ostroff M, Spencer TR, Dawson RB. The SIC protocol: A seven-step strategy to minimize complications potentially related to the insertion of centrally inserted central catheters. *J Vasc Access*. 2021;24:185-190.
- ▶ The Australasian Society for Ultrasound in Medicine. Guidelines for Reprocessing Ultrasound Transducers. *AJUM*. 2017;20:30-40.
- ▶ American Institute of Ultrasound in Medicine. Guidelines for Cleaning and Preparing External- and Internal-Use Ultrasound Transducers and Equipment Between Patients as Well as Safe Handling and Use of Ultrasound Coupling Gel. <https://www.aium.org/resources/official-statements/view/guidelines-for-cleaning-and-preparing-external--and-internal-use-ultrasound-transducers-and-equipment-between-patients-as-well-as-safe-handling-and-use-of-ultrasound-coupling-gel>.
- ▶ Centers for Disease Control and Prevention. 2017 Updated Recommendations on the Use of Chlorhexidine-Impregnated Dressings for Prevention of Intravascular Catheter-Related Infections. <https://www.cdc.gov/infectioncontrol/pdf/guidelines/bsi-guidelines-H.pdf>. Accessed December 23, 2023.
- ▶ Holinga G, McGuire JE. *In vitro* antimicrobial effects of chlorhexidine diacetate versus chlorhexidine free base dressings. *J Wound Care*. 2020;29(Sup5A):522-528.

Thank you!

Questions & Answers

Robert Garcia, MT(ASCP), CIC, FAPIC
President & Senior Infection Preventionist
Enhanced Epidemiology, LLC
rgarciaipc1980@gmail.com

