
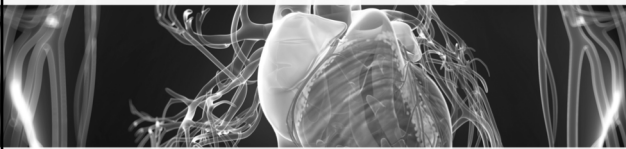


MAYO CLINIC



CLABSI Prevention – It's Not Just for Intensivists



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Disclosures

- Nothing to disclose

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Objectives

- Recognize aspects and terminology of Central Line Associated Blood Stream Infections (CLABSI) reporting
- Understand implications of CLABSIs to an institution
- Identify perioperative strategies that could impact an institution's CLABSI rate

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Overview

- 1 of 14 Hospital Acquired Conditions
- 30,000 to 34,000 (2016)
- Publically Reported Data
 - CDC – National Health Safety Network (NHSN)
 - CMS
- The Leapfrog Group



Definitions

- Primary blood stream infection (BSI)
 - Laboratory Confirmed Bloodstream Infection (LCBI) that is **NOT** secondary to an infection at another site
 - Mucosal Barrier Injury – LCBI
 - Stem Cell transplantation
- Secondary BSI
 - A BSI seeded from another site-specific infection



Definitions

- Secondary BSI Attribution Period
 - Time period where the blood **MUST** be collected
 - 14 to 17 days
- Infection Window Period (IWP)
- Repeat Infection Timeframe (RIT)



Definitions

- Location of Attribution (LOA)
 - Inpatient location assigned to the patient on the date of the event
- Central Lines
 - Tunneled catheters
 - Implanted catheters (port)
 - Temporary catheters (PICC)
 - Umbilical catheters



Devices Not Considered Central Lines

- Arterial catheters
- Arteriovenous fistula
- Arteriovenous graft
- Atrial catheters
- Extracorporeal membrane oxygenation (ECMO)
- Hemodialysis reliable outflow (HERO) dialysis catheter
- Intra-aortic balloon pump (IABP) devices
- Non-accessed central line (not accessed nor inserted during the hospitalization)
- Peripheral IV or Midlines
- Ventricular Assist Device (VAD)



Definitions

- CDC NHSN CLABSI Definition
 - Laboratory confirmed blood stream infection (LCBI)
 - Eligible BSI organism
 - Eligible central line
 - Day of event (DOE) or day before (24hrs)



Definitions

- Catheter-related bloodstream infection (CRBSI)
 - BSI is attributed to the catheter by culture of the tip
 - Time to Differential Positivity - >120 minutes
 - **Reporting agencies do not care!**

Ann Intern Med. 2004;140(1):18-25

Reported Data

- Standardized Infection Ratio (SIR)
 - Observed HAIs / Predicted HAIs
- Predicted HAI
 - National aggregate data for HAI
 - Logistic Regression Model – SSI
 - Negative Binomial Regression Model – CLABSI, VAE, etc.

Ann Intern Med. 2004;140(1):18-25

McStuffinsville Hospital

Year/Quarter	Central Line Days	Number of Infections	Number of Expected Infections	SIR
2017Q1	1561	3	2.185	1.373
2017Q2	1671	5	2.339	2.137
2017Q3	1590	4	2.370	1.687
2017Q4	1715	3	2.230	1.345
2017 YTD	6537	15	9.124	1.64

Ann Intern Med. 2004;140(1):18-25

Case

- 71 y/o male s/p CABG x 2, AVR, and MVR
 - Cardiogenic shock after separation from bypass
 - Extubated POD #2
 - IABP removed on POD #3
 - POD# 5
 - Spikes fever to 38.6°
 - Leukocytosis 12.2 (8.6)
 - Low MAP
 - Norepinephrine
 - Cultures & antibiotics

Bacteria / Candida Culture, Blood #1
 Status: Final result Visible to patient: Yes (Patient Online Services)
 Specimen Information: Blood, Arterial Line; Blood

Component	4mo ago
Bacteria/Candida Culture, Blood	<p>↓</p> <p>STAPHYLOCOCCUS COAGULANS-NEGATIVE Growth after 22 hours 3 of 3 bottles. Critical Results: mrsa detected. Methicillin (mexillin)-resistant coagulase-negative staphylococcus. Possible blood culture contamination (unless isolated from more than one blood culture draw or clinical case suggests pathogenicity). No antibiotic treatment is indicated for blood culture contamination. Susceptibilities not performed on this isolate.</p>

Bacteria / Candida Culture, Blood #2
 Status: Final result Visible to patient: Yes (Patient Online Services)
 Specimen Information: Blood, Short Term CVC; Blood

Component	4mo ago
Bacteria/Candida Culture, Blood	No growth after 5 days of incubation.

Bacterial Culture, Aerobic + Susc
 Status: Final result Visible to patient: Yes (Patient Online Services)
 Specimen Information: Catheter Tip; Device

Component	4mo ago
Bacterial Culture, Aerobic + Susc	No growth after 5 days of incubation.

Bacterial Culture, Aerobic + Susc
 Status: Final result Visible to patient: Yes (Patient Online Services)
 Specimen Information: Catheter Tip, Central Venous; Device

Component	4mo ago
Bacterial Culture, Aerobic + Susc	No growth after 5 days of incubation.

\$45,254

Wait Just a Minute!

- Arterial lines \neq central lines
- A primary BSI **cannot** be secondary to a BSI
- VASC – venous or arterial infection
 - Pus at the site **plus** matching organisms

Infectious risk associated with arterial catheters compared with central venous catheters*

Jean-Christophe Lucet, MD, PhD; Lila Bouadma, MD; Jean-Ralph Zahar, MD; Carole Schwebel, MD; Arnaud Geffroy, MD; Sebastien Pesse, MD; Marie-Christine Heraud, MD; Hakim Hachouache, MD; Christophe Adris, MD; Marie Thuong, MD; Adrien François, RT; Maïté Garrouste-Orgeas, MD; Jean-François Timsit, MD, PhD

	Incidence per 1000 days colonization	Incidence per 1000 days CRI	
Arterial	11.4 (n=127)	0.99 (n=11)	p=0.8
STCVC	11.1 (n=183)	1.09 (n=18)	

Crit Care Med 2010 Vol. 38, No. 4

Arterial catheter-related bloodstream infection: incidence, pathogenesis, risk factors and prevention^{1,2}

N. Safdar^{a,b,*}, J.C. O'Horo^c, D.G. Maki^b

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^cDepartment of Pulmonary and Critical Care Medicine, Mayo Clinic, Rochester, MN, USA

Arterial Catheters	Patients	BSI Control	BSI Treatment	Colonization Control & Treatment
834	542	9	2	109

Wait Just a Minute!

- Transfer Rule
- Exception to the LOA
- Date of event
 - Transfer
 - Discharge
 - Next day
- Infection attributed to the transferring/discharging location
- CLABSI **cannot** be attributed to a non-bedded location
 - Inpatient Dialysis
 - Operating Room
 - Vascular/Interventional Radiology
 - Cardiac Cath/EP Lab
 - GI/Endoscopy Suite



HAI: When in Doubt, Blame Anesthesia. Could They Be Right?
by Ian Yum, MD, and Jeffrey M. Feldman, MD, MSE
JUNE 2016

ASHP
 (P) Anesthesia Patient Safety Foundation 2016

Checklist for Prevention of Central Line Associated Blood Stream Infections
Journal of Clinical Pharmacy and Therapeutics

RESEARCH **Open Access**

Poor adherence to guidelines for preventing central line-associated bloodstream infections (CLABSI): results of a worldwide survey

Dev CrossMark

Giordina Valencia^{1,2}, Naïma Hammami¹, Antonella Agosti³, Alain Lepape⁴, Eduardo Palencia Hercejon⁵, Sijm Blot⁶, Jean-Louis Vincent⁷ and Marie-Laurence Lambert¹

The Children's Hospital, Boston, Massachusetts; ¹Department of Internal Medicine, University of Nebraska Medical Center Omaha, Nebraska; and ²Department of Internal Medicine, Ann Arbor VA Medical Center and University of Michigan, Ann Arbor, Michigan


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Anesthesiology 2008; 109:399-407
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Transmission of Pathogenic Bacterial Organisms in the Anesthesia Work Area

Randy W. Loftus, M.D.,* Matthew D. Koff, M.D.,† Corey C. Burchman, M.D.,‡ Joseph D. Schwartzman, M.D.,§
 Valerie Thorum, M.T. (A.S.C.P.),|| Megan E. Read, M.T. (A.S.C.P.),|| Tamara A. Wood, M.T., (A.M.T.),||
 Michael L. Beach, M.D., Ph.D.¶



Anesthes. 2008;109(3):399-407. doi:10.1097/ALN.0b013e318182c855

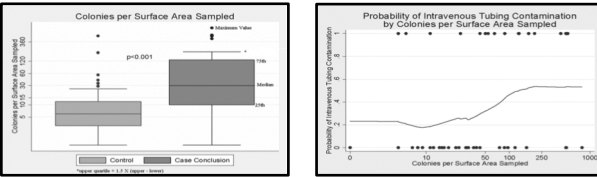


Fig. 2. Box plot illustrating increase in intraoperative contamination on the anesthesia machine as measured by colonies per surface area sampled. $P < 0.001$ for comparison of medians.

Fig. 3. Probability of obtaining bacterial growth in the stopcock as a function of the number of colonies per surface area sampled from the anesthesia machine.

Anesthes. 2008;109(3):399-407. doi:10.1097/ALN.0b013e318182c855

Site	Organism	Number of Colonies	Number of Sites
AA	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1
	Staphylococcus saprophyticus	100	1
	Staphylococcus carnosus	100	1
	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1
	Staphylococcus saprophyticus	100	1
	Staphylococcus carnosus	100	1
	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1
E	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1
	Staphylococcus saprophyticus	100	1
	Staphylococcus carnosus	100	1
	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1
	Staphylococcus saprophyticus	100	1
	Staphylococcus carnosus	100	1
	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1
E	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1
	Staphylococcus saprophyticus	100	1
	Staphylococcus carnosus	100	1
	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1
	Staphylococcus saprophyticus	100	1
	Staphylococcus carnosus	100	1
	Staphylococcus epidermidis	100	1
	Staphylococcus aureus	100	1

- *Staph Epidermidis*
- *VRE*
- *Enterobacter chloacae*

Anesthes. 2008;109(3):399-407. doi:10.1097/ALN.0b013e318182c855

Transmission Dynamics of Gram-Negative Bacterial Pathogens in the Anesthesia Work Area

Randy W. Loftus, MD,* Jeremiah R. Brown, MS, PhD,† Hetal M. Patel, BS,* Matthew D. Koff, MD, MS,* Jens T. Jensen, MS,* Sundara Reddy, MD,‡ Kathryni L. Rusoff, PhD,* Stephen O. Heard, MD,§ Thomas M. Dodds, MD,* Michael L. Beach, MD,* and Mark P. Yeager, MD*

- Identify and examine transmission of gram (-) bacteria
- Examine the link between transmission and 30 day HAIs

Anesthesia & Analgesia. 120(4):819-826, April 2015.

Organism	Hospital site	Patient culture source	Possible transmission link	PFGE confirmed link to culture	Temporal association of link	HCAI
Enterobacter aerogenes	1	Sputum	Attending physician hand	Yes	Provider to patient	Yes
Proteus mirabilis	0	Urine patient case 1	Patient nasopharynx 2nd case	Yes	Patient to patient	Yes
Serratia liquefaciens	0	Sputum	Patient nasopharynx 2nd case	Yes	Patient to self	Yes

HCAI = health care-associated infection; PFGE = pulsed-field gel electrophoresis.

Anesthesia & Analgesia. 120(4):819-826, April 2015.

GUIDELINES

Guidelines for the Prevention of Intravascular Catheter-related Infections

Nasimi P, O'Grady M, Alexander L, Lillan A, Burns T, E. Paschon DeBonger, Jeffrey Garland, Stephen O. Heard, Pamela A. Lipman, Nancy Maslow, Leonard A. Mermel, Michelle L. Pearson, Isaac I. Blank, Adrienne G. Bandelack, Mark E. Ryge, Sanjay Saxe, and the Healthcare Infection Control Practices Advisory Committee (HICPAC) (Appendix 1)

*Critical Care Medicine Department, National Institutes of Health, Bethesda, Maryland; †Infectious Nurses Society, Norwood, Massachusetts; ‡State Island University Hospital, State Island, New York; §Department of Surgery, University of Washington, Seattle, Washington; ¶Department of Pediatrics, Whittier Franciscan Healthcare St. Joseph, Milwaukee, Wisconsin; ¶Department of Anesthesiology, University of Massachusetts Medical Center, Worcester, Massachusetts; ¶Department of Surgery, Johns Hopkins University School of Medicine, Baltimore, Maryland; ¶Division of Infectious Diseases, Warren Alpert Medical School of Brown University and Rhode Island Hospital, Providence, Rhode Island; ¶Office of Infectious Diseases, CDC, Atlanta, Georgia; ¶Department of Infectious Diseases, MD Anderson Cancer Center, Houston, Texas; ¶Department of Anesthesiology, The Children's Hospital, Boston, Massachusetts; ¶Department of Internal Medicine, University of Nebraska Medical Center, Omaha, Nebraska; and ¶¶Department of Internal Medicine, Ann Arbor VA Medical Center and University of Michigan, Ann Arbor, Michigan

CID 2011:52 (1 May)

IV Access Points

- Closed systems
 - Pressure transducers
 - Medication administration
 - IV infusions
- Needless connections
 - Spilt septum
 - Mechanical Valve

Figure 1a and 1b: Split septum used for blood access. (Continued on reverse side of slide)

Figure 1c and 1d: Use of split septum with split septum. (Continued on reverse side of slide)

Journal of Intensive Care Medicine, 33: 2010
 CID 2011;52 (1 May)
 doi:10.1093/cid/cir053

Journal of Intensive Care Medicine, 33: 2010
 CID 2011;52 (1 May)
 doi:10.1093/cid/cir053

Health Care-Associated Bloodstream Infections Associated with Negative- or Positive-Pressure or Displacement Mechanical Valve Needleless Connectors

William S. Jarvis,¹ Colleen Murphy,² Kent K. Holt,³ Pamela J. Fugle,⁴ Tali B. Kachrim,⁵ Clancy Hamilton,⁶ Catherine Delgado,⁷ Yoo Y. Cho,⁸ Carol Gonzalez,⁹ and Robert J. Stobo¹⁰

Table 3. Participating Hospital Bloodstream Infection (BSI) Rates during Split Septum (SS) and Mechanical Valve (MV) Needleless Device Use Period.

Hospital, unit/ward	SS BSI rate ^a	MV BSI rate	Relative risk (95% CI)	P	Post-MV BSI rate	Relative risk (95% CI)	P
A: Adult ICU (n = 4)	6.47	9.84	1.16 (0.94-1.44)	.16	6.10	1.01 (1.18-2.22)	.003
B: Adult ICU (n = 6)	3.09	6.62	2.65 (2.15-3.65)	<.001	5.29	1.67 (1.12-2.40)	.008
C:							
Adult ward	2.48	3.41	1.38 (0.98-1.93)	.05	2.29	1.03 (1.04-2.11)	.03
Adult ICU (n = 4)	3.15	3.47	1.10 (0.61-1.98)	.87	2.89	1.20 (1.14-1.98)	.43
D:							
Adult ICU	0	4.30	NC (0.03-999)	.80			
Adult oncology ward	2.20	6.20	2.20 (2.05-2.71)	.04			
Adult ICU	6.65	14.8	2.24 (1.43-3.49)	<.001			
A-C: Adult ICU (n = 16)	6.15	9.59	1.56 (1.37-1.76)	<.001	6.17 ^b	1.05 (1.28-1.30)	<.001

^a Rates are based on health care-associated BSIs per 1000 central venous catheter-days for all except hospital C, which used 1000 patient-days for the adult ward health care-associated BSI rate that includes the entire hospital.
^b Includes 3 hospitals with 14 adult ICUs. Post-MV rate includes the health care-associated BSI rate only in facilities changing from MV needleless connectors to SS needleless connectors.

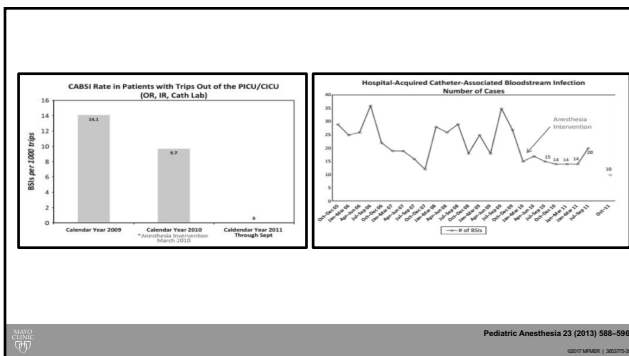
CID 2009-49 (15 December)
 doi:10.1093/cid/cir053

Pediatric Anesthesia
Pediatric Anesthesia 23 (2013) 588-596

ORIGINAL ARTICLE
Modification of anesthesia practice reduces catheter-associated bloodstream infections: a quality improvement initiative

- Seattle Children's Hospital
- 7 days post procedure
- AWE contamination of access points
- Airway management
- Peripheral IV insertion
- Medication administration
 - Scrub hub
 - Manifold
- "Clean" and "Dirty" zones


Pediatric Anesthesia 23 (2013) 588-596



Take Home Points

- CLABSI ≠ CRBSI
- Beyond the bundle
 - Arterial catheters
 - IV access points
 - Closed systems

Pediatric Anesthesia 23 (2013) 588-596


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Chlorhexidine-Impregnated Sponges and Less Frequent Dressing Changes for Prevention of Catheter-Related Infections in Critically Ill Adults
 A Randomized Controlled Trial
JAMA, March 25, 2009—Vol 301, No. 12

60%

Randomized Controlled Trial of Chlorhexidine Dressing and Highly Adhesive Dressing for Prevention of Catheter-related Infections in Critically Ill Adults
JAMA, March 25, 2009—Vol 301, No. 12

Jean-François Timon^{1,2}, Olivier Meunier^{1,2}, Monique Bertrand^{1,2}, Marie-Cécile Orgerat³, Serge Allardard⁴, Gaëtan Planeteaux⁵, Régis Bronckard⁶, Gilles Trochu⁷, Rémy Gauzit⁸, Aurélien Antonia⁹, Emmanuel Canel¹⁰, Julien Bohen¹¹, Alain Lapaque¹², Aurélien Verdin¹³, Xavier Arrault¹⁴, Carole Schwechtl¹⁵, Christophe Adrie¹⁶, Jean-Ralph Zahar¹⁷, Stéphane Ruckly¹⁸, Caroline Tournegros¹⁹, and Jean-Christophe Lucet²⁰

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Incidence of central venous catheter hub contamination
Journal of Critical Care 39 (2017) 162–168

Julie L. Holroyd^{a,*}, Terrie Vasilopoulos^{b,c}, Mark J. Rice^d, Kenneth H. Rand^e, Brenda G. Fahy^{f,g}




Fig. 1. Sanitizing the outside surface of multilumen connector hub with cotton tipped swab.




Fig. 2. Multilumen connector device with multiple ports that allow several catheters to be administered through the same lumen of the catheter.

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