Rebooting Your Unit Culture: Preventing Sepsis by Reducing Health care Acquired Infection





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Disclosures

Consultant-Michigan Hospital Association Keystone Center

Subject matter expert on CAUTI, CLABSI, HAPI, Safety culture for AHA

Consultant and speaker bureau

- \triangle Stryker's Sage business
- \triangle Baxter healthcare
- \triangle Potrero Medical
- \triangle Atlas Lift Tech

Objectives

- Discuss trends in CLABSI & VAP rates during the pandemic
- Discuss strategies for getting back to the basics in resetting the culture of safety
- Identify and detail the evidence-based practices that help prevent CLABSI's & VAP.



Impact of COVID on Healthcare-Associated Infections (HAIs) in 2020 Compared to 2019: Data from NHSN

	2020 Q1	2020 Q2	2020 Q3	2020 Q4
CLABSI	-11.8%	27.9%	46.4%	47.0%
CAUTI	-21.3%	No Change ¹	12.7%	18.8%
VAE	11.3%	133.7%	29.0%	44.8%
SSI: Colon surgery	-9.1%	No Change ¹	-6.9%	-8.3%
SSI: Abdominal hysterectomy	-16.0%	No Change ¹	No Change ¹	-13.1%
Laboratory-identified MRSA bacteremia	-7.2%	12.2%	22.5%	133.8%
Laboratory-identified CDI	-17.5%	-10.3%	-8.8%	-5.5%





Qualitative Feedback on Rationale for Increase

Shortage of personal protective equipment (PPE)	Staffing changesTravelersNon-ICU clinicians	 Reduced frequency of contact Less chlorhexidine bathing Alterations in line care due to IV pumps in the hallway Scrub the hub compliance
Line and dressing integrity gaps related to prone positioning of patients	Increase in line draws for blood cultures	Less line rounding/competing priorities



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journal homepage: www.elsevier.com/locate/ijid

The impact of COVID-19 on health care-associated infections in intensive care units in low- and middle-income countries: International Nosocomial Infection Control Consortium (INICC) findings



Victor D. Rosenthal^{a,b,*}, Sheila Nainan Myatra^c, Jigeeshu Vasishtha Divatia^c,

Table 1

Comparison of pooled HAI rates per 1,000 device days and relative risk in ICUs of 7 INICC hospitals from LMICs in 2019 and 2020.

HAI type	Device days, n	2019 HAIs, n	HAI rate	Device days, n	2020 HAIs, n	HAI rate	RRª	95% Cl ^a	P-value
CLAB ^a	36,652 CLª days	93 CLABs	2.54 CLAB per 1,000 CL days	9,515 CL days	45 CLABs	4.73 CLAB per 1,000 CL days	1.85	1.30-2.65	.0006
VAE ^a	13,801 MVª days	134 VAEs	9.71 VAE per 1,000 MV days	4,611 MV days	58 VAEs	12.58 VAE per 1,000 MV days	1.29	0.95-1.75	.10
CAUTI ^a	24,919 UC ^a days	41 CAUTIs	1.64 CAUTI per 1,000 UC days	7,653 UC days	11 CAUTIs	1.43 CAUTI per 1,000 UC days	1.14	0.58-2.22	.69

Acute care hospitals were located in India, Mongolia, Jordan, Turkey, Egypt, Palestine, and Lebanon.

CAUTI = catheter associated urinary tract infection; CI = confidence interval; CL = central line; CLAB = central line-associated bloodstream infection; HAI = health care-associated infection; ICU = intensive care unit; INICC = International Nosocomial Infection Control Consortium; LMIC = low- and medium-income country; MV = mechanical ventilator; RR = relative risk; UC = urinary catheter; VAE = ventilator-associated event.

Mortality Comparison 2019 to 2020: 15.2% & 23.2% with an increase in mean LOS by almost 2 days

Life after a Crisis

Life is about how much you can take and keep fighting, how much you can suffer and keep moving forward.~ Anderson Silva

Don't dwell on what went wrong. Instead, focus on what to do next. Spend your energies on moving forward toward finding the answer. ~Denis Waitley

If everyone is moving forward together, then success takes care of itself. ~Henry Ford

One day? Or day one. You decide.

From the Field

"We're on our knees here, and it's really difficult and we're all trying the best we can and we don't feel... we feel like we could be doing more, and I know we can't ... We're staying away from our families and we're putting ourselves in danger to try and save other people's loved ones. It feels like a losing battle but it's not, we've all got hope and we're all trying to do what we can."

The Problem is Large

- 3 million central venous catheters (CVCs) inserted each year-15 million catheter days
- ▲ 30,100 CLABSIs in ICU yearly
- More of the CLABSI now occur outside the ICU
- \land 12 to 25%
- 2.27-fold increased risk for mortality

Centers for Disease Control and Prevention. (2015). Bloodstream infection event (central line-associated bloodstream infection and non-central line-associated bloodstream infection). Buetti N, et al. Shea/IDSA/APIC Practice Recommendation. Infection Control & Hosp Epidemiology, 2022;1-17*

Excess Mortality Estimates for Hospital Acquired Conditions (HACs)

	Ν	Range (RR)	Estimates of RR (95% Cl)	Underlying Mortality	Estimates of Excess Mortality (95% CI)
Adverse Drug Events (ADE)	6	0.68–3.09	1.61 (1.14–2.27)	0.020	0.012 (0.003–0.025)
Catheter- Associated Urinary Tract Infections (CAUTI)	4	1.28–1.97	1.50 (1.06–2.11)	0.071	0.036 (0.004–0.079)
Central Line- Associated	5	1.86-4.88	2.72 (1.81–4.10)	0.086	0.150 (0.070–0.270)
Bloodstream Infections (CLABSI)	For Every 1000) in-hospital CLA	BSI cases, there	are 150 excess d	eaths
Falls	1	3.50	3.50 (2.73–4.48)	0.020	0.050 (0.035–0.070)
Obstetric Adverse Events (OBAE)	-	-	-	-	0.005 (0.003–0.013)
Pressure Ulcers	³ For Every 10	00 in-hospital VA	P cases, there ar	re 140 excess dea	oths ^{0.013-0.093)}
Surgical Site Infections (SSI)	3	1.75–5.70	3.32 (1.79–6.18)	0.0114	0.026 (0.009–0.059)
Ventilator- Associated Pneumonia (VAP)	10	0.52–4.90	1.48 (0.64–3.42)	0.300	0.140 (-0.110–0.730)
Venous Thromboembolis m (VTE)	9	1.01–13.63	3.15 (2.02–4.91)	0.020	0.043 (0.040–0.078)
C. difficile Infections (CDI)	13	1.17–9.60	1.60 (1.38–1.87)	0.073 http://www.ahr	0.044 (0.028–0.064) g.gov/hai/pfp/haccost



Additional Inpatient Costs & Mortality for HAC's: Building the Business Case

	Studies (n)	Range of Estimates	Estimate (95% CI)
Adverse Drug Events (ADE)	2	\$1,277-\$9,062	\$5,746 (-\$3,950–\$15,441)
Catheter-Associated Urinary Tract Infections (CAUTI)	6	\$4,694–\$29,743	\$13,793 (\$5,019–\$22,568)
Central Line-Associated Bloodstream Infections (CLABSI)	7	\$17,896–\$94,879	\$48,108 (\$27,232–\$68,983)
Falls	3	\$2,680–\$15,491	\$6,694 (-\$1,277–\$14,665)
Obstetric Adverse Events (OBAE)	2	\$13–\$1,190	\$602 (-\$578–\$1,782)
Pressure Ulcers	4	\$8,573-\$21,075	\$14,506 (-\$14,506–\$41,326)
Surgical Site Infections (SSI)	5	\$11,778-\$42,177	\$28,219 (\$18,237–\$38,202)
Ventilator-Associated Pneumonia (VAP)	5	\$19,325–\$80,013	\$47,238 (\$21,890–\$72,587)
Venous Thromboembolism (VTE)	4	\$11,011-\$31,687	\$17,367 (\$11,837–\$22,898)
C. difficile Infections (CDI)	9	\$4,157–\$32,394	\$17,260 (\$9,341–\$25,180)

Resetting the Culture Notes on Hospitals: 1859

"It may seem a strange principle to enunciate as the very first requirement in a hospital that it should do the sick no harm."

- Florence Nightingale







Protect The Patient From Bad Things Happening on Your Watch





Implement Interventional Patient Hygiene





INTERVENTIONAL PATIENT HYGIENE

- Hygiene...the science and practice of the establishment and maintenance of health
- ▲Interventional Patient Hygiene....nursing action plan directly focused on fortifying the patients host defense through proactive use of evidence-based hygiene care strategies



Incontinence Associated Dermatitis Prevention Program

Pressure Prevention

Bathing & Assessment

INTERVENTIONAL PATIENT HYGIENE(IPH)



Vollman KM. Intensive Crit Care Nurs, 2013;22(4): 152-154

Achieving the Use of the Evidence

Value

Skills thomedoe **Factors Impacting the Ability to Achieve Quality Nursing Outcomes at the Point of Care**

Pesources

¢

- System

Attitude & Accountability

Polling Question

On quality measures, were you able to keep track on CLABSI data during the pandemic? AYes

 $\triangle No$

Chat in: Did it get worse or better?



Teamwork and Fundamental Nursing Interventions











Teamwork

What teamwork strategies learned through the pandemic will help you and your team to getting back on track? Select all that apply!

 \triangle More streamlined communication

 \bigtriangleup Creative communication strategies

 \bigtriangleup Faster development & approval of practice changes

- \triangle Resiliency practices (individual or organizational)
- \triangle Managing change on the quick (Rapid PDCA's)

 \triangle Just do it

 $\bigtriangleup\ensuremath{\operatorname{Nimbleness}}$ that we didn't believe we had



Key Team/Culture Work to Re-Engage

- A Place CAUTI & CLABSI as permanent agenda items for the Multidisciplinary Team/CUSP team
- ▲ Consider revising goals for the work based on the current state
- Creating a plan for re-engagement of staff post-COVID
- Share unit-specific data again with frontline staff, physicians, safety committee
- A Refresh on expectations/clinical practice & documentation
- Reestablish unit-based improvement practices & meetings (improvement huddle, unit-based council mtgs, learn from a defect huddles)

Do you and the staff you work with see HAPI, CAUTI/CLABSI as harm?



Strategies to Link Harm with Nurse Patient Advocacy Role

▲ Do No Harm Rounding

Immediate learn from a deficit

Incorporate action plans and data into daily huddle

Learn from Defects Tool Worksheet CLABSI	Why did it happen
Date:	1) Was patien 2) Were there
What happened? (brief description) Patient with documented CLABSI Infection Control:	patient? [3) Was line ne
Nursing:	4) Was the dr
Significant co markidition	5) Was this lin
Significant co-morbidities.	radiology, e
Location of CLABSI: Unit Date of CLABSI	6) Was the tu
Date of Line Insertion:	7) Was the ca
Type of Line No. of Lumens	If yes, was
Non-tunneled (other than dialysis) Single Double Triple	8) Where the
Tunneled (other than dialysis) Dialysis (tunneled)	9) Anything e
Dialysis (non-tunneled)	If yes, desc
PICC Port	10) Do you fee
Insertion Site	11) Other Com
Chest U SC Femoral Upper Extremity	What prev
What is the indication for the line?	_
Hemodynamic monitoring Poor venous access Long-term Antibiotics	
Vessicants or irritant drugs Chemo Hemodialysis	
Wurupie incompatible iluidsOther	
Lab draws Medication Administration IV Fluid Administration	
TPN Hemodialysis Other	What can we do to Action Plan
	Action Plan

Why did it happen? (what factors contributed) - summarize what happened to cause the defect from below
1) Was patient receiving TPN?
2) Were there any observed breaches of proper hand hygiene by anyone involved in line care for this
patient? Y N Unknown
3) Was line necessity assessed daily?
4) Was the dressing integrity difficult to maintain?
5) Was this line manipulated/used by any other staff besides the unit's physicians/nurses (e.g., anesthesia,
radiology, etc)?
6) Was the tubing changed appropriately for the duration of the line?
7) Was the catheter occluded at any time while the line was in place?
If yes, was TPA used? Y N Unknown
8) Where there any problems drawing off the line prior to the infection date?
9) Anything else, patient factors or otherwise, that may have contributed to the infection?
If yes, describe briefly
10) Do you feel this infection was potentially preventable?
11) Other Comments:
+
Duration of central line catheter # days: (Time of insert
Is the patient being treated for any other infections?
Comments:

/hat can we do to reduce the risk	of it happening with a different per	son?	
ction Plan	Responsible Person	Targeted Date	Evaluation Plan – How will we know risk is reduced?

Data Drive Performance

- of
- Audits of Insertion and Maintenance practices-specific to areas of challenge
- Early & Late Infections
- \Lambda Display of Data

Infection	Days Since Last Infection
CAUTI	150 days
CLABSI	200 days
MRSA transmission	50 days

Measure unit specific instances of CLABSI and report data on a regular basis (high level of evidence)

Buetti N, et al. Shea/IDSA/APIC Practice Recommendation. Infection Control & Hosp Epidemiology, 2022;1-17

Central Line -Associated Blood Stream Infections: Prevention is Key



www.catheterout.org, (Adapted Meddings. Clin Infect Dis 2011)

BSI Prevention Bundle

- A Remove/Avoid unnecessary lines
- \Lambda Hand hygiene
- \Lambda Maximal barrier
- Alcoholic/Chlorhexidine for skin prep
- Subclavian preferred* (high level evidence)



984 Adult intensive care units (ICUs) in 632 hospitals: Bundle compliance on all 5 elements > 95% greatest reduction (33% ↓) Bundle compliance of 1 element > 95 % second best reduction Bundle compliance < 75% no change in rates seen</p>

Furuya EY, et al. Infect Control Hosp Epidemiol, 2016;37:805-810

Grady NP, et al. CDC Guidelines for the Prevention of Intravascular Catheter-Related Infections, 2011. <u>www.cdc.gov</u> Buetti N, et al. Shea/IDSA/APIC Practice Recommendation. Infection Control & Hosp Epidemiology, 2022;1-17*

The Right Catheter for the Right Length of Time for the Right Infusate

Magic Guidelines: Peripherally Compatible Infusate

	Proposed Duration of Infusion				
Device Type	≤5 d	6–14 d	15–30 d	≥31 d	
Peripheral IV catheter	No preference between peripheral IV and US-guided peripheral IV catheters for use ≤5 d				
US-guided peripheral IV catheter	US-guided peripheral IV cathe catheter if proposed	eter preferred to peripheral IV duration is 6–14 d			
Nontunneled/acute central venous catheter	Central venous catheter pro or if hemodynamic moni	eferred in critically ill patients toring is needed for 6–14 d			
Midline catheter	Midline catheter preferred to P	ICC if proposed duration is ≤14 d			
PICC		PICC preferred to mid	line catheter if proposed duration o	of infusion is ≥15 d	
Tunneled catheter				PICC preferred to tunneled catheter and ports for	
Port				infusion 15–30 d	



Chopra V, et al. Annals of Internal Medicine. 2015;suppl

Appropriate

Neutral

Inappropriate

Disagreement

Magic Guidelines: Peripherally Incompatible Infusate

Appropriate

	Proposed Duration of Infusion				
Device Type	≤5 d	6–14 d	15–30 d	≥31 d	
Peripheral IV catheter					
US-guided peripheral IV catheter					
Nontunneled/acute central venous catheter	Central venous catheter pro	eferred in critically ill patients toring is needed for 6–14 d			
Midline catheter					
PICC		PICCs rated as appropriate at all	proposed durations of infusion		
Tunneled catheter		Tunneled catheter neutral for for use ≥15 d	No preference between tun proposed du	neled catheter and PICC for rations ≥15 d	
Port				No preference among port, tunneled catheter, or PICC for ≥31 d	

Neutral

Inappropriate

DIsagreement



Chopra V, et al. Annals of Internal Medicine. 2015;suppl



Line Location

- 2-fold risk of CLABSI's using Jugular vs. Subclavian
- A Higher risk of CLABSI and thrombosis with Femoral vs. Subclavian
- Ultrasound utilization/Simulation training/Subclavian safe
- Different approach to Internal Jugular (Dr Jack LeDonne)
- Ultrasound guided peripheral placements-extended dwell
- Use of Midline catheters (infections less-more mechanical issues)
- No difference in DVT between PICC and Midline

Bing S, et al. JACS, 2020;231(4): suppl 2:p135 DeVries M, et al. Am J of IC. 2019;47:1118-1121 Au AK, et al. Am J Emerg Med. 2012;30(9):1950-1954. Parienti J-J, et al. New England Journal of Medicine. 2015;373(13):1220-1229

After Insertion Bundle

CHG Dressings/Dressing Integrity/Site Securement

▲Bathing

- ▲Accessing the site
- Antimicrobial impregnated CVC & PICCs

Appropriate nursing staff levels in ICUs (high level evidence)

Health Research & Educational Trust (2017). *Central Line-Associated Bloodstream Infections (CLABSI) Change Package: 2017 Update.* Chicago, IL: Health Research & Educational Trust. Accessed at www.hret-hiin.org Buetti N, et al. Shea/IDSA/APIC Practice Recommendation. Infection Control & Hosp Epidemiology, 2022;1-17

Dressing Disruption: A Major Risk Factor for Catheter-Related Infections

- Secondary analysis of an randomized controlled trial (RCT)
- ▲ 1,419 patients (3,275 arterial or central venous catheters)
- ▲ 296-Colonize catheters, 29 major catheter related infections and 23 CLABSI
- ▲ 11,036, dressing changes and
 7,347 (67%) were performed
 before the planned date



Timsit JF, et al Crit Care Med; 2012:1707-1714

Impact of Dressing Disruption

- Dressing cost inversely related to rate of disruption
- ▲ Number of dressing disruptions r/t ↑ risk for colonization of the skin around the catheter at removal (p< .0001)</p>
- A Risk of infection increased threefold after 2nd dressing disruption
- A Risk of infection increased 10-fold when the final dressing was disrupted independently of other risk factors of infection





Meta-analysis on Impact of CHG Dressings on CLABSI's

- △ 20 studies, 18 RCT's and 15,590 catheters
- ▲ No evidence of publication bias
- ▲ Mostly performed in ICU's
- ▲ Independent of CHG dressing type

A Catheter-related bloodstream infections

				Risk Ratio		Risk Ratio		
Study or Subgroup	log[Risk Ratio]	SE	Weight	IV, Random, 95% CI		IV, Random, 95% CI		_
Arpa 2013	-1.147	1.623	0.4%	0.32 [0.01, 7.64]				
Arvaniti 2012	0.445	0.906	1.4%	1.56 [0.26, 9.21]				
Biehl 2016	-0.616	0.335	9.9%	0.54 [0.28, 1.04]				
Chambers 2005	-2.267	1.62	0.4%	0.10 [0.00, 2.48]	•			
Chan 2017	0.216	1.621	0.4%	1.24 [0.05, 29.76]				
Düzkaya 2016	-1.609	1.077	1.0%	0.20 [0.02, 1.65]	-		(C Adve
Ergul 2018	-0.249	0.319	10.9%	0.78 [0.42, 1.46]				
Garland 2001	0.17	0.41	6.6%	1.19 [0.53, 2.65]		-		Study o
Gerçeker 2017	1.54	1.504	0.5%	4.66 [0.24, 88.92]				Arpa 20
Levy 2005	0.246	0.745	2.0%	1.28 [0.30, 5.51]				Biehl 20
Maki 2000	-0.997	0.587	3.2%	0.37 [0.12, 1.17]				Chan 2
Margatho 2018	-0.511	1.203	0.8%	0.60 [0.06, 6.34]				Lew 20
O'Horo 2013	-0.216	0.18	34.3%	0.81 [0.57, 1.15]				Margath
Pedrolo 2014	0.159	0.565	3.5%	1.17 [0.39, 3.55]				Pedrolo
Pivkina 2018	-0.916	0.796	1.8%	0.40 [0.08, 1.90]				Pivkina
Roberts 1998	1.041	1.786	0.3%	2.83 [0.09, 93.83]				Timsit 2
Ruschulte 2009	-0.579	0.275	14.7%	0.56 [0.33, 0.96]				Timsit 2
Timsit 2009	-1.109	0.718	2.2%	0.33 [0.08, 1.35]				Total (9
Timsit 2012	-0.919	0.585	3.2%	0.40 [0.13, 1.26]				Heterog
Yu 2019	0.101	0.667	2.5%	1.11 [0.30, 4.09]				Test for
Total (95% CI)			100.0%	0.71 [0.58, 0.87]				
Heterogeneity: Tau ² =	0.00; Chi ² = 15.3	8, df = 1	9 (P = 0.7	'0); I² = 0%			100	
Test for overall effect:	Z = 3.23 (P = 0.00)1)			0.01	Eavours CHG Eavours control	100	

Use of CHG dressing in all patients > 2 months of age (High level of evidence

C Adverse events

				Risk Ratio	Risk Ratio
Study or Subgroup	log[Risk Ratio]	SE	Weight	IV, Random, 95% CI	IV, Random, 95% Cl
Arpa 2013	2.149	1.48	3.1%	8.58 [0.47, 155.98]	
Biehl 2016	0.051	0.218	25.1%	1.05 [0.69, 1.61]	+
Chan 2017	-0.373	0.287	22.5%	0.69 [0.39, 1.21]	
Garland 2001	3.746	1.43	3.3%	42.35 [2.57, 698.37]	
Levy 2005	1.345	1.106	5.1%	3.84 [0.44, 33.54]	
Margatho 2018	0.182	1.396	3.4%	1.20 [0.08, 18.51]	•
Pedrolo 2014	-0.135	0.254	23.7%	0.87 [0.53, 1.44]	
Pivkina 2018	1.099	1.613	2.7%	3.00 [0.13, 70.84]	
Timsit 2009	2.977	2.195	1.5%	19.63 [0.27, 1449.71]	
Timsit 2012	1.456	0.733	9.6%	4.29 [1.02, 18.04]	
Total (95% CI)			100.0%	1.46 [0.85, 2.50]	
Heterogeneity: Tau ² =	0.26; Chi ² = 18.6	5, df = 9	(P = 0.03)	3); I² = 52%	
Test for overall effect:	Z = 1.36 (P = 0.17	Favours CHS Favours control			

Puig-Asensio, et al. Infection Control & Hospital Epid, 2020'41:1388-1395

Durability and Costs of Different CVC Dressings

Dressing duration was captured prospectively on four different dressings on five critical care units over a 12-month period

▲ 590 CVCs with 1,229 dressing changes

Staff received training on evidence-based CVC dressing practices and a 'how to guide' was implemented

Phase	Months	CVC dressing evaluated	Other securement techniques	
One	I <u>-</u> 4	Standard dressings: sterile, transparent, semi-permeable polyurethane dressings (Opsite IV 3000 and 3M Tegaderm®)	None	
Two	5–8	3M Tegaderm® IV Advanced: sterile, transparent, semi-permeable polyurethane dressings	Dressing with an integrated border around the dressing. Separate Hyperfix® border applied to create a further secure 'window' around the edge of the dressing	
Three	9–12	Sorbaview®: sterile, transparent, semi- permeable polyurethane dressings	Integrated two piece dressing, one part for the site with a wide border and second part with a wide supporting bridge	Richardson A, et al. J of Infection Control. 2015;16(6):256-261

Durability and Costs of Different CVC Dressings: Results Figure 1. Reasons for dressing removal, all CVC (n=590).

- 3-4 dressings lasted < 48hr, 1 dressing a mean of 68hrs
- Mean time to change the dressing:13.5 min
- Cost range: \$2.85 to \$7.20
- Only 3% lasted 7 days

	Dressings removed f	or any reason, n=1229		Dressings removed for non-adherence, clammy skin, or bleeding under dressing n=630			
Dressing Type	Number of dressings observed	Dressing duration (hrs) median [IQR]	z value*	Number of dressings observed	Dressing duration (hrs) median [IQR]	z value**	
Opsite IV 3000	310	43.5 [21–78]	-1.79	160	36.0 [15-67.5]	-1.21	
Tegaderm	237	46.0 [22–85]	-0.33	122	45.5 [22-73.8]	1.17	
IV Advanced	262	40.5 [20-85]	-1.12	143	32.0 [14-69.5]	-1.98	
Sorbaview	116	68.5 [32–105]	4.5 I	42	53.0 [30-95]	3.39	
Unrecorded	304			163			



IQR, inter quartile range; *P < 0.001 and **P = 0.002 for at least one difference between dressings.
Polling Question

- Does your organization have a dressing kit that includes all the necessary components to complete an evidence-based dressing change?
 - \triangle Yes
 - \triangle No



Human Factor Engineering of Central Line Maintenance

10 fold decrease

- 29 month prospective
- \land 95 nurses, 151 patients
- 126 observation pre compared with 90 post intervention procedures (kit use)

AResults

- \triangle Pre CLABSI:2.21/1,000 cath days \triangle Post CLABSI:0/1,000 cath days
- △ Practice Adherence: Better aseptic technique, better CHG scrub, hand sanitization & disinfecting hub
- \triangle Procedure omission \downarrow by 44%





Drews FA, et al. American Journal of Infect Control. 2017;45:1224-30

After Insertion

▲CHG Dressings/Dressing Integrity/Site Securement
▲Bathing

Accessing the site/Disinfecting the hub

Antimicrobial impregnated CVC & PICCs

▲ Appropriate nursing staff levels in ICUs

Health Research & Educational Trust (2017). *Central Line-Associated Bloodstream Infections (CLABSI) Change Package: 2017 Update.* Chicago, IL: Health Research & Educational Trust. Accessed at www.hrethiin.org

Traditional Bathing

Why are there so many bugs in here?



Soap and water basin bath was an independent predictor for the development of a CLABSI

Musuuza JS, et al.. BMC Infect Dis. 2019;19(1):416

CHG Impact on CLABSIs: A Meta-analysis

\land 26 studies

- \triangle 18 non-randomized
- \triangle 8 randomized controlled trials (RCTs) (19 in the ICU)
- \bigtriangleup 18/27 studies used 2% CHG cloth
- \land 861,546 patient-days
- 💪 5259 CLABSI
- Evaluated 5 measure of fidelity
 - $\triangle Adherence$
 - \triangle Exposure/dose
 - \triangle Quality of delivery
 - \triangle Participant responsiveness
 - \triangle Program differentiation
- ▲ 12% all 5, 12% 4, 15% 3, 27% 2, 35% 1
- Adverse events: 16 studies-skin rashes, dryness and pruritus



CHG Group: CLABSI 4.4 per 1000 pt days Comparison Group: 7.5 per 1000 pt days

CHG Bathing: Works Upstream



Differential Effects of Antisepsis Skin Cleansing Methods

Rhee Y, et al. Infect Control Hosp Epidemiol 2018;39:405-411

- Prospective, randomized 2center study with blinded assessment.
- ▲ To determine whether 3 different CHG skin cleansing methods yield similar residual CHG concentrations and bacterial densities on skin.



Method A- 2% CHG cloth Method B- 4% CHG liquid poured onto nonmedicated cloth Method C-4% CHG liquid on cotton wash cloth



Passive Disinfection: Meta-Analysis

▲ To compare the effects of antiseptic barrier cap use and manual disinfection on the incidence of CLABSIs

\Lambda Outcome

- △ Reduction in CLABSIs per 1,000 catheter-days
- △ Studies were included if 1) conducted in a hospital setting, 2) used antiseptic barrier caps on hubs of central lines with access to the bloodstream, and 3) reported the number of CLABSIs per 1,000 catheter-days when using the barrier cap and when using manual disinfection
- $\bigtriangleup\,$ 7 were included in the random effects meta-analysis



Unresolved issue: Whether mechanical disinfection needs to occur is using passive disinfection Buetti N, et al. Shea/IDSA/APIC Practice Recommendation. Infection Control & Hosp Epidemiology, 2022;1-17



Replacement of Administration Sets

A Routine administration sets can be changed at intervals up to seven days

- △This does not include blood, blood products or lipid formulations
- △ Optimal replacement for intermittent used sets is unresolved

Buetti N, et al. Shea/IDSA/APIC Practice Recommendation. Infection Control & Hosp Epidemiology, 2022;1-17

Rickard CM, et al. Lancet, 2021;397:1447-58

01-CHSS Blue Plus-Antimicrobial Impregnated

02-Silver Impregnated-Antimicrobial Impregnated

03-Minocycline-Rifampin-Antibiotic

04-Miconazole & Rifampicin -Antibiotic

05-Benzalkonium Chloride Impregnated-Antimicrobial Impregnated

06-CHSS-Antimicrobial coated

Study	Treatment	Control	Peto OR	Weight	Peto OR
or sub-category	n/N	n/N	95% CI	%	95% CI

01 CHSS Blue Plus - Antimicrobial Impregnate Brun-Buisson 2004 - 27199 Jaeger, 201

2011 CDC Recommendations:

Subtotal (9) Use a chlorhexidine/silver sulfadiazine or Total events Test for het Test for over minocycline/rifampin -impregnated if, after successful 02 Silver Im Boswald, Stoiser, 20 implementation of a comprehensive strategy to reduce Bong. 200 Corral, 200 Moretti, 200 Kalfon, 200 rates of CLABSI, the CLABSI rate is not decreasing. ^{2,3} Subtotal (9) Total events Test for heb Test for our



	1				
3 Minocycline Rifampin - Anfl	biolo				
Raad, 1997	0/130	5/136		2.24	0.14 [0.02, 0.80]
darik B. 1999	0/38	2/39		0.90	0.14 [0.01, 2.20]
Chatanikolaou, 2003	0/66	7/64		3.04	0.12 [0.03, 0.54]
lanna. 2004	3/182	14/174		7.39	0.25 [0.09, 0.65]
eon 2004	6/187	11/180		7.40	0.52 [0.20, 1.37]
ubtotal (95% Ci)	603	593	•	20.97	0.26 [0.15, 0.47]
tal events:9 (Treatment), 39	(Control)		•		
est for heterogeneity: Chill = 3	3.69. df = 4 (P = 0.45). I* =	=0%			
ist for overall effect: Z = 4.52	(P < 0.00001)				
4 Miconazole and Rifampicin	-Antibiotic				
doel. 2004	0/118	1/105		0.45	0.12 [0.00, 6.07]
ubtotal (95% Ci)	118	105		0.45	0.12 [0.00, 6.07]
tal events:0 (Treatment),1 ((Control)	_ = =			
est for heterogeneity; not app	licable				
est for overall effect: Z = 1.06	(P = 0.29)				
Benzalkonium Chloride Imp	pregnated - Antimicrobia	i impregnated			
teger, 2001	1/25	1/25		0.89	1.00 [0.06, 16.45]
ubiotal (95% Ci)	25	25		0.89	1.00 [0.06, 16.45]
zal events: 1 (Treatment), 1 ((Control)				
est for heterogeneity: not app	dicable				
est for overall effect: Z = 0.00	(P = 1.00)				
7 CHSS - Antimicrobial Coate	ed				
ach. 1996b	0/116	3/117		1.35	0.13 [0.01, 1.30]
emberton, 1996	2/32	3/40		2.12	0.83 [0.13, 5.08]
ogghe, 1997	17/338	15/342	+	13.91	1.15 [0.57, 2.35]
aki, 1997	2/208	9/195		4.88	0.25 [0.08, 0.84]
ennenberg, 1997	5/137	9/145		6.07	0.58 [0.20, 1.70]
olin, 1999	1/98	4/139		2.17	0.41 [0.07, 2.46]
annan, 1999	1/174	3/177		1.81	0.37 [0.05, 2.66]
arik A, 1999	1/36	2/39		1.33	0.55 [0.06, 5.43]
heng. 2000	1/113	2/122		1.35	0.55 [0.06, 5.36]
ubtotal (95% CI)	1252	1316	•	34.99	0.62 [0.40, 0.98]
tal events: 30 (Treatment), 5	(Control)		-		
ast for heterogeneity: ChP = 7	7.43, df = 8 (P = 0.49), I* -	-0%			
est for overall effect: Z = 2.06	(P = 0.04)				
		3728	•	100.00	0.49 [0.37, 0.64]
tal /95% Cit	3696				
tal (95% Cl) tal events: 75 (Treatment), 1	3696 (Control)	3724	•		
tal (95% Cl) tal events: 75 (Treatment), 1 ist for heterogeneits: ChP = 2	3696 155 (Control) 28.78.df = 26 (P = 0.32).	P=9.7%	•		
tal (95% Cl) tal events: 75 (Treatment), 1 st for heterogeneity: ChP = 2 st for overall effect: 7 = 5 32	3696 155 (Control) 28.78, df = 26 (P = 0.32), 1 (P < 0.00001)	I*=9.7%	, i		

Favours treatment Favours control

1. Matheos T, et al. Infect Control Hosp Epidemiol 2010;31:295-297

2. Grady NP, et al. CDC Guidelines for the Prevention of Intravascular Catheter-Related Infections, 2011. <u>www.cdc.gov</u>

3. Buetti N, et al. Shea/IDSA/APIC Practice Recommendation. Infection Control & Hosp Epidemiology, 2022;1-17



1

VAP Prevention is Key



Building Blocks to Best Practice in Caring for Mechanically Ventilated Patients & \checkmark VAP

Ventilator Bundle: HOB 30, Deep Vein Thrombosis (DVT) prophylaxis, Peptic Ulcer Disease (PUD) prophylaxis, Sedation interruption, Spontaneous breathing trial, daily care with chlorhexidine

VAP Bundle: HOB 30, Sedation interruption, Spontaneous breathing trial, oral care 6x per day, CHG rinse 2x per day, subglottic secretions drainage if expected to be ventilated > 72hrs

ABCDEF Bundle: Assess & manage pain, Both Spontaneous awakening trial (SAT) & spontaneous Breathing trial(SBT), Choice of Sedation, Delirium Assessment and management, Early Mobility, Family and Patient Engagement

http://www.ihi.org/resources/Pages/Tools/HowtoGuidePreventVAP.aspx www.ICUliberation.org Rawat N, et al. Crit Care Med, 2017;45:1208-1215

ASSESS, PREVENT & MANAGE PAIN



Recommendations/Guidelines

Society of Critical Care Medicine

August 2018

- Severe pain negatively effects ICU patients
- ▲ Vital Signs and behaviors are flags to investigate.
- Recommend use of a protocol based pain assessment and management program
- 🛆 Treat pain first

Use a valid and reliable assessment tool

The American Society of

Pain Management Nursing

July 2011



💪 Poor pain control

▲ Vital signs are not "sensitive"

CPOT is acceptable for the critically ill/unconscious



Treating Acute Pain in the ICU

Situation	Preferred Intervention
Acute pain	Fentanyl IVP until pain resolves
Acute pain that persists/recurs	Fentanyl infusion plus fentanyl IVP for breakthrough
Acute pain in chronic opioid user?	Account for previous opioid use when using IV opioid (may consider ketamine)
Planned transition out of ICU and patient on IV opioid infusion	Start scheduled oral/enteral opioid therapy (e.g., oxycodone) plus intermittent IV opioid (e.g., IVP or PCA)

New Guidelines Recommend: Adjuncts to opioidsacetaminophen, nefopam, ketamine

> www.ICU liberation.org Devlin J. Crit Care Med. 2018 Sep;46(9):e825-e873

Agitation



Light sedation suggested:

- \bigtriangleup Sedative medications should be titrated to maintain **lighter** levels of sedation, unless clinically contraindicated.
- \triangle Use daily awakening or a titrated sedation strategy to maintain patient wakefulness.

Choice of sedative:

△ Suggest using propofol or dexmedetomidine over benzodiazepines to improve clinical outcomes in mechanically ventilated ICU patients.

A Cardiac Surgery Patients

 \bigtriangleup Suggest using propofol over a benzodiapine

Devlin J. Crit Care Med. 2018 Sep;46(9):e825-e873

Barr J. Crit Care Med. 2013;41:263-306

Agitation

- Assess q 4hrs or prn with change in dose or patients condition
- Use validated tool (RASS or SAS)
- RASS target -1 to +1
- SAS target 3 to 4

TABLE 1. RICHMOND AGITATION-SEDATION SCALE

Score	Term	Description
+4	Combative	Overtly combative or violent; immediate danger to staff
+3	Very agitation	Pulls on or removes tube(s) or catheter(s) or has aggressive behavior toward staff
+2	Agitated	Frequent nonpurposeful movement or patient-ventilator dyssynchrony
+1	Restless	Anxious or apprehensive but movements not aggressive or vigorous
0	Alert and calm	
-1	Drowsy	Not fully alert, but has sustained (more than 10 seconds) awakening, with eye contact, to voice
-2	Light sedation	Briefly (less than 10 seconds) awakens with eye contact to voice
-3	Moderate sedation	Any movement (but no eye contact) to voice
-4	Deep sedation	No response to voice, but any movement to physical stimulation
-5	Unarousable	No response to voice or physical stimulation
Procedure		
1. Observe p	atient. Is patient alert and calm (sco	ore 0)?
Does pation	ent have behavior that is consistent cription)?	t with restlessness or agitation (score $+1$ to $+4$ using the criteria listed above,
2. If patient i once if ne	s not alert, in a loud speaking voice cessary. Can prompt patient to con	state patient's name and direct patient to open eyes and look at speaker. Repeat tinue looking at speaker.
Patient ha	s eye opening and eye contact, wh	ich is sustained for more than 10 seconds (score -1).
Patient ha	s eye opening and eye contact, but	this is not sustained for 10 seconds (score -2).
Dettent he	a since an account of the second second	

Patient has any movement in response to voice, excluding eye contact (score -3).
If patient does not respond to voice, physically stimulate patient by shaking shoulder and then rubbing sternum if there is no response to shaking shoulder.
Patient has any movement to physical stimulation (score -4).

Patient has no response to voice or physical stimulation (score -5).

www.iculiberation.org



NKE UP





Daily Sedation Interruption Decreases Duration of Mechanical Ventilation

▲ Hold sedation infusion until patient awake, then restart at 50% of prior dose

▲ "Awake" defined as any 3 of the following:
 △ Open eyes in response to voice
 △ Use eyes to follow investigator on request

- $\bigtriangleup \mathsf{Squeeze}$ hand on request
- $\bigtriangleup Stick$ out tongue on request





- Length of MV 4.9 vs. 7.3 days (P=0.004)
- ICU LOS 6.4 vs. 9.9 days (P=0.02)
- Fewer diagnostic tests to assess changes in mental status
- No increase in rate of agitated-related complications or episodes of patient-initiated device removal
- No increase in PTSD or cardiac ischemia

ABC Trial (RCT Paired Sedation & Vent Weaning Protocols)

Outcome*	SBT	SAT+SBT	<i>P</i> value
Outcome	301	SALISET	/ Value
Ventilator-free days	12	15	0.02
Time-to-event, days			
Successful extubation, days	7.0	5	0.05
ICU discharge, days	13	9	0.02
Hospital discharge, days	19	15	0.04
Death at 1 year, n (%)	97 (58%)	74 (44%)	0.01
Days of brain dysfunction			
Coma	3.0	2.0	0.002
Delirium	2.0	2.0	0.50
*Median, except as noted			

Girard, et al, Lancet. 2008;371:126-34

ABC Trail: Mortality at 1 Year





CDC Prevention Epicenters' Wake Up and Breathe Collaborative





▲16% ¹ in SBTs

81% in SBTs done with sedatives off

▲37% in VACs

▲65% in IVACs

Klompas M. (CDC ABCDE Collaborative) Am J Respir Crit Care Med. 2015;191:292-301.



Making it Happen: Wake Up & Breathe

Process Measure: Daily audit of SAT/SBT compliance or documentation of contraindication

- \bigtriangleup Determine if they meet SAT criteria
- \bigtriangleup Decrease or stop sedation per protocol
- △ Determine if patient meets Readiness to Wean/Resp
- \triangle Determine if meet SBT protocol criteria/Resp
- \bigtriangleup Consider one time of day-coordinate between resp and nursing (white board-EMR-communication tool)
- \bigtriangleup Discuss results in multidisciplinary rounds
- \triangle Include in nurse to nurse handoff/other handoffs
- \bigtriangleup Dedicated RRT in rounds speaking up
- \triangle Ventilator LOS posted/Extubation rates posted

Oral Cavity & VAP

- ▲89 critically ill patients
- Examined microbial colonization of the oropharynx through out ICU stay
- Used pulse field gel electrophoresis to compare chromosomal DNA

AResults:

- Diagnosed 31 VAPs
- 28 of 31 VAP's the causative organism was identical via DNA analysis
 - Garrouste-Orgeas et. al. Am J Respir Crit Care Med. 1997;156:1647-1655

49 elderly nursing home residents admitted to the hospital

- Examined baseline dental plaque scores & microorganism within dental plaque
- Used pulse field gel electrophoresis to compare chromosomal DNA

AResults

- 14/49 adults developed pneumonia
- 10 of 14 pneumonias, the causative organism was identical via DNA analysis



Formation of Biofilm Over 13 Hours



Loesche, W. 2012

http://helios.bto.ed.ac.uk/bto/microbes/biofilm.htm

Does Compliance Make A Difference?

Oral care compliance & use of the ventilator bundle resulted in a 89.7% reduction in VAP



Compliance rates for the years of the study



VAP rates for the years of the study





Delirium: First Focus on Prevention

Pain and sedation scores

Analgesia and Sedative Algorithm

 $\bigtriangleup\ensuremath{\operatorname{Control}}$ pain first, then anxiety

 \triangle Use intermittent meds first before continuous

▲Target RASS + 1 to -1

Daily SAT (spontaneous awakening trial)

Daily SBT (spontaneous breathing trial)

Implement non-pharmacological strategies



Delirium Assessment & Management

- Delirium Assessment:
 - •ICU-CAM
 - •ICU Delirium Screening Checklist
- Frequency:
 - Q shift & prn

Confusion Assessment Method in the ICU



Non-Pharmacological Strategies

Sleep Promotion

- ▲ Appropriate Medications
- \Lambda Bath during day
- ▲ Chair position
- \Lambda Lighting
- ▲ Television
- Hearing/Vision Aids/Dentures
- \Lambda Control Noise

Other

- ▲ Cognitive Stimulation/Music
- ▲ Familiar objects in room/pictures

Mobility Promotion

- ▲ Evaluate for Physical Therapy
- \land Range of Motion
- 💪 Sleep
- \Lambda Work with PT
- \Lambda Spontaneous Awakening Trial

Sedation Holidays

- Sleep Promotion
- Mobility

Pandharipande P et al. (Lorazepam) *Anesthesiology* 2006;104:21–26; Oimet ICM 2007; 33:1007-1013; Pandharipande P et al. (Midazolam) *J Trauma* 2008 Dubois MJ et al., (Morphine) *Intensive Care Med* 2001; 27:1297–

Outcomes of Early Progressive Mobility Program

- ${\scriptstyle \bigtriangleup} \psi$ incidence of skin injury
- $\bigtriangleup \psi$ time on the ventilator
- ${\scriptstyle \bigtriangleup} \psi$ incidence of VAP
- $\Delta \psi$ days of sedation
- $arsigma \psi$ delirium
- ▲↑ ambulatory distance
- Improved function

Staudinger t, et al. Crit Care Med, 2010;38. Abroung F, et al. Critical Care, 2011;15:R6 Morris PE, et al. Crit Care Med, 2008;36:2238-2243 Pohlman MC, et al. Crit Care Med, 2010;38:2089-2094 Schweickert WD, et al. Lancet, 373(9678):1874-82. Thomsen GE, et al. CCM 2008;36;1119-1124 Winkelman C et al, CCN,2010;30:36-60





ABCDEF Bundle: Improving Survival & Reducing Brain Dysfunction

- Ventilated and non-ventilated medical and surgical ICU patients enrolled between January 1, 2014, and December 31, 2014
- ▲ Determine association between ABCDEF bundle compliance/total & partial & outcomes of hospital survival and delirium-free and coma-free days/
- A Patients experienced more days alive and free of delirium and coma with both total bundle compliance and partial bundle compliance



ICU Liberation Collaborative



Prospective, multicenter, cohort study of national collaborative

- ▲ 68 academic, community and federal ICU's collected data-20 months
- Measure complete and proportional bundle performance against 3 groups of outcomes, patient related, symptom related and system related





ICU Liberation Collaborative





Integrating ABCDEF into ICU culture

▲ Talk about all the **ABCDEF** bundle as ONE.

Utilize Change Champions in all aspects of integration

 \triangle Demonstrate/Mentor staff

 \triangle Ground Up

▲ Daily Rounds with Multidisciplinary Team

• Expectation is for RN to speak the language

△ Don't start each intervention separate from the others

△ Group interventions together, demonstrate how they connect and evaluate together



WHEN WOULD NOW BE A GOOD TIME TO DO THIS?

It is not enough to do your best; you must know what to do, and THEN do your best. ~ W. Edwards Deming
Forbid yourself to be deterred by poor odds just because your mind has calculated that the opposition is too great. If it were easy, everyone would do it. HAI prevention courses by Kathleen Vollman

https://www.medbridgeeducation.co m/advancing-nursing







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