

CONVEGNO E VIDEOCONFERENZA

## IL NURSING DEI DISPOSITIVI PER ACCESSO VASCOLARE PER EMODIALISI



Polo Didattico Universitario  
Ospedale Sacco -  
Milano e Videoconferenza



5 Ottobre 2024



**RESPONSABILE SCIENTIFICO:** *Maurizio Gallieni*, Direttore Nefrologia e Dialisi, ASST Fatebenefratelli Sacco, Dipartimento di Scienze Biomediche e Cliniche, UNIMI, Milano

Gemelli



Fondazione Policlinico Universitario A. Gemelli  
Università Cattolica del Sacro Cuore



## BUNDLES PROCEDURALI PER LA SICUREZZA DEL PAZIENTE

GIANCARLO SCOPPETTUOLO

FONDAZIONE POLICLINICO UNIVERSITARIO  
"A. GEMELLI" - IRCCS - ROMA

## The Top Patient Safety Strategies That Can Be Encouraged for Adoption Now

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## *Table 2. Patient Safety Strategies Ready for Adoption Now*

### **Strongly encouraged**

Preoperative checklists and anesthesia checklists to prevent operative and postoperative events

Bundles that include checklists to prevent central line–associated bloodstream infections

Interventions to reduce urinary catheter use, including catheter reminders, stop orders, or nurse-initiated removal protocols

Bundles that include head-of-bed elevation, sedation vacations, oral care with chlorhexidine, and subglottic suctioning endotracheal tubes to prevent ventilator-associated pneumonia

Hand hygiene

The do-not-use list for hazardous abbreviations

Multicomponent interventions to reduce pressure ulcers

Barrier precautions to prevent health care–associated infections

Use of real-time ultrasonography for central line placement

Interventions to improve prophylaxis for venous thromboembolisms

1 October 2005  
Volume 41  
Number 7

Published by  
The University of  
Chicago Press

 IDSA  
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# Clinical Infectious Diseases



 IDSA  
Infectious Diseases Society  
of America

GUIDELINES

## Guidelines for the Prevention of Intravascular Catheter-Related Infections

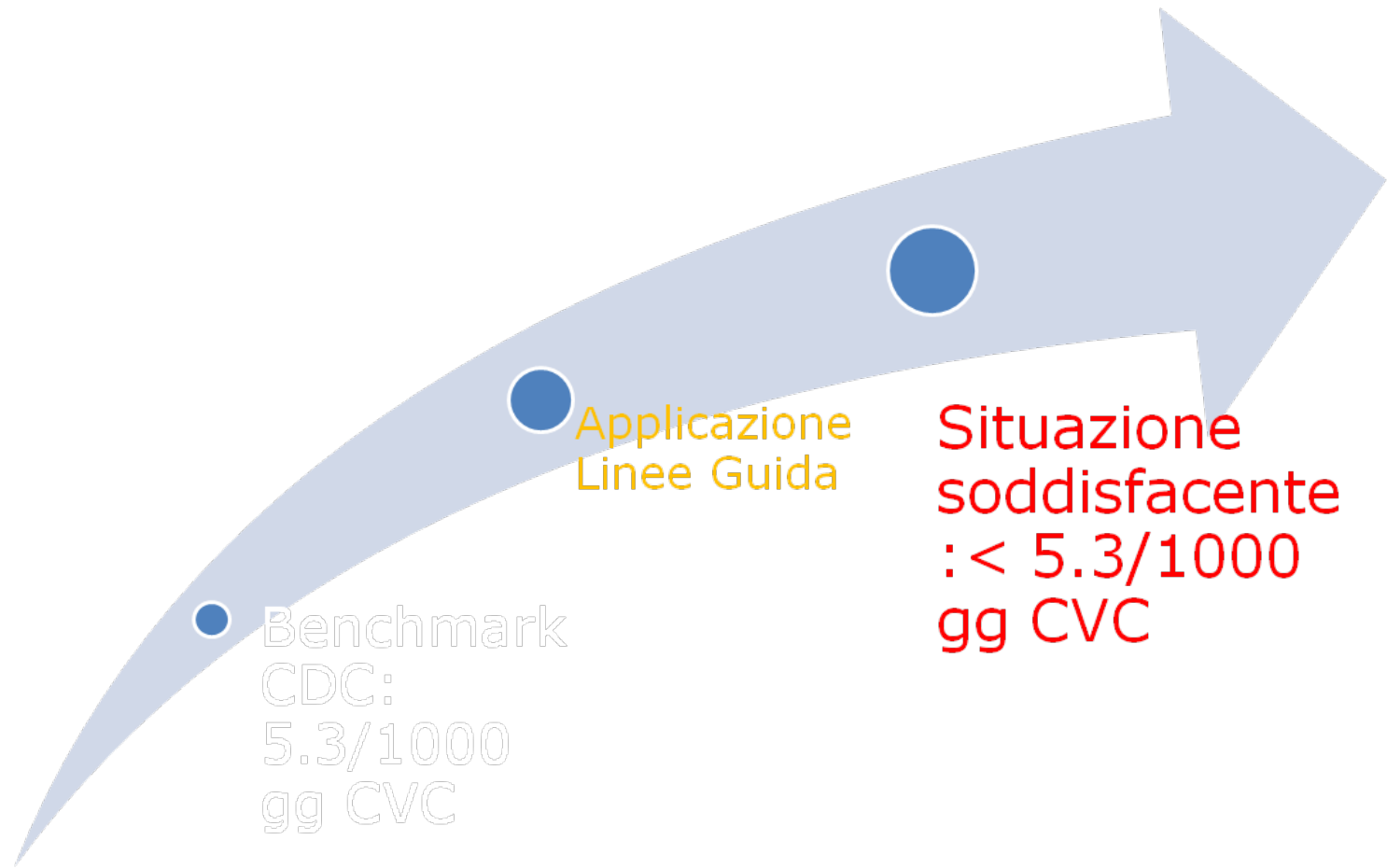
Naomi P. O'Grady,<sup>1</sup> Mary Alexander,<sup>1</sup> E. Patchen Dellinger,<sup>2</sup> Julie L. Gerberding,<sup>3</sup> Stephen O. Heard,<sup>4</sup> Dennis G. Maki,<sup>5</sup> Henry Masur,<sup>6</sup> Rita D. McCormick,<sup>7</sup> Leonard A. Mermel,<sup>8</sup> Michele L. Pearson,<sup>9</sup> Issam I. Raad,<sup>10</sup> Adrienne Randolph,<sup>11</sup> and Robert A. Weinstein<sup>12</sup>

<sup>1</sup>National Institutes of Health, Bethesda, Maryland; <sup>2</sup>Infection Nurses Society, Cambridge; and <sup>3</sup>University of Massachusetts Medical School, Worcester; and <sup>4</sup>The Children's Hospital, Boston, Massachusetts; <sup>5</sup>University of Washington, Seattle; <sup>6</sup>Office of the Director, Centers for Disease Control and Prevention (CDC), and <sup>7</sup>Division of Healthcare Quality Promotion, National Center for Infectious Diseases, CDC, Atlanta, Georgia; <sup>8</sup>University of Wisconsin Medical School and <sup>9</sup>Hospital and Clinics, Madison; <sup>10</sup>Rhode Island Hospital and Brown University School of Medicine, Providence, Rhode Island; <sup>11</sup>M.D. Anderson Cancer Center, Houston, Texas; and <sup>12</sup>Cook County Hospital and Rush Medical College, Chicago, Illinois

**Clinical Infectious Diseases** 2002; 35:1281–307

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1058-4838/2002/3511-0001



Benchmark  
CDC:  
5.3/1000  
gg CVC

Applicazione  
Linee Guida

Situazione  
soddisfacente  
:< 5.3/1000  
gg CVC



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## Guidelines for the Prevention of Intravascular Catheter-Related Infections, 2011

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The goal of an effective prevention program should be the elimination of CRBSI from all patient-care areas. Although this is challenging, programs have demonstrated success, but sustained elimination requires continued effort. The goal of the measures discussed in this document is to reduce the rate to as low as feasible given the specific patient population being served, the universal presence of microorganisms in the human environment, and the limitations of current strategies and technologies.



## *The Risk of Bloodstream Infection in Adults With Different Intravascular Devices: A Systematic Review of 200 Published Prospective Studies*

Dennis G. Maki, Daniel M. Kluger, Christopher J. Crnich

*Mayo Clin Proc.* September 2006; 81 (9): 1159-1171

RISK OF IVD-RELATED BLOODSTREAM INFECTION

TABLE 3. Rates of Intravascular Device-Related Bloodstream Infection Caused by Various Types of Devices Used for Vascular Access\*

Device	No. of studies	No. of catheters	No. of IVD (d)	No. of BSIs	Rates of IVD-related bloodstream infection			
					Per 100 devices		Per 1000 IVD-days	
					Pooled mean	95% CI	Pooled mean	95% CI
Peripheral IV catheters								
Plastic catheters	110	10,910	28,720	13	0.1	0.1-0.2	0.5	0.2-0.7
Steel needles	1	148	350	3	2.0	0.0-4.3	8.6	0.0-18.2
Venous cutdown	1	27	111	1	3.7	0.0-10.8	9.0	0.0-26.6
Midline catheters	3	514	9251	2	0.4	0.0-0.9	0.2	0.0-0.5
Arterial catheters for hemodynamic monitoring	14	4366	21,397	37	0.8	0.6-1.1	1.7	1.2-2.3
Peripherally inserted central catheters								
Inpatient and outpatient	15	3566	105,839	112	3.1	2.6-3.7	1.1	0.9-1.3
Inpatient	6	625	7137	15	2.4	1.2-3.6	2.1	1.0-3.2
Outpatient	9	2813	98,702	97	3.5	2.8-4.1	1.0	0.8-1.2
Short-term noncuffed central venous catheters								
Nonmedicated								
Nontunneled	79	20,226	322,283	883	4.4	4.1-4.6	2.7	2.6-2.9
Tunneled	9	741	20,065	35	4.7	3.2-6.2	1.7	1.2-2.3
Medicated								
Chlorhexidine-silver-sulfadiazine	18	3367	54,054	89	2.6	2.1-3.2	1.6	1.3-2.0
Minocycline-rifampin	3	690	5797	7	1.0	0.3-1.8	1.2	0.3-2.1
Silver impregnated	2	154	1689	8	5.2	1.7-8.7	4.7	1.5-8.0
Silver iontophoretic	2	396	4796	16	4.0	2.1-6.0	3.3	1.7-5.0
Benzalkonium chloride	1	277	2493	12	4.3	1.9-6.7	4.8	2.1-7.5
Pulmonary artery catheters	13	2057	8143	30	1.5	0.9-2.0	3.7	2.4-5.0
Hemodialysis catheters								
Temporary, noncuffed	16	3066	51,840	246	8.0	7.0-9.0	4.8	4.2-5.3
Long-term, cuffed and tunneled	16	2806	373,563	596	21.2	19.7-22.8	1.6	1.5-1.7
Cuffed and tunneled central venous catheters	29	4512	622,535	1013	22.5	21.2-23.7	1.6	1.5-1.7
Subcutaneous venous ports								
Central	14	3007	983,480	81	3.6	2.9-4.3	0.1	0.0-0.1
Peripheral	3	579	162,203	23	4.0	2.4-5.6	0.1	0.1-0.2
Intra-aortic balloon pumps	1	101	414	3	3.0	0.0-6.3	7.3	0.0-15.4
Left ventricular assist devices	3	157	19,653	41	26.1	19.2-33.0	2.1	1.5-2.7

\*BSI = bloodstream infection; CI = confidence interval; IV = intravenous; IVD = intravascular device.





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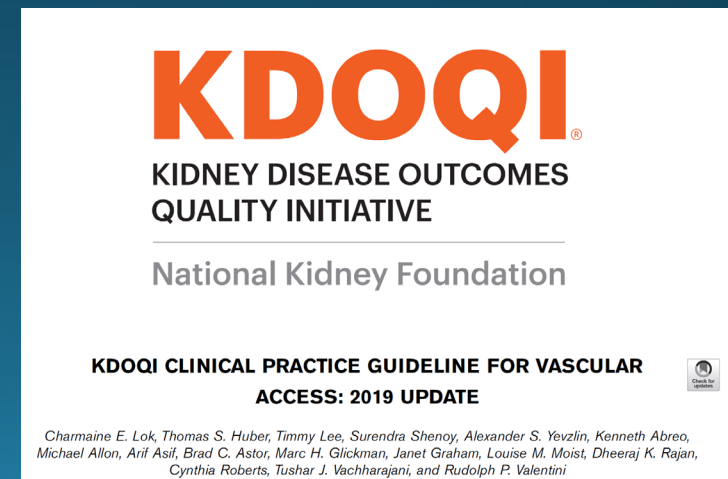
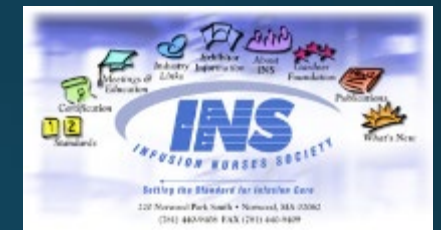
MAUROBIANI 2008

For HAIs, it is widely demonstrated that all are preventable, but some are partly preventable and some others (CLABSI), on the contrary, are completely preventable and avoidable.

According to IHI's  
experiences and  
Campaigns, the best tool  
to Target Zero Infections  
is the "Bundle"

# Linee Guida di Riferimento per la prevenzione delle CRBSI

- CDC Atlanta 2002 (Centers for Disease Control, USA)
- RCN 2005 (Royal College of Nurses, UK)
- INS 2006 (Infusion Nursing Society, USA)
- BCSH 2006 (British Committee for Standards in Hematology, UK)
- **EPIC 2007 (Evidence -Based Practice in Infection Control, UK)**
- **SHEA/IDSA 2008**
- **ESPEN 2009**
- **RCN 2010**
- **INS 2011**
- **CDC 2011**
- **EPIC 3 2014**
- **SHEA 2014**
- **INS 2016**
- **RCN 2016**
- **KDOQI 2019**
- **INS 2021**
- **SHEA/IDSA 2022**
- **INS 2024**



VAD SELECTION  
AND HEALTHCARE WORKERS  
EDUCATION AND TRAINING

INSERTION

CRBSI Prevention

CARE OF EXITE SITE

DISINFECTION OF CATHETER HUBS,  
CONNECTORS AND INJECTION PORTS

# Central Line Bundle



Hand Hygiene



Maximal Barrier Precautions Upon Insertion



Chlorhexidine Skin Antisepsis



Optimal Catheter Site Selection, with Subclavian Vein as the Preferred Site for Non-Tunneled Catheters



Daily Review of Line Necessity with Prompt Removal of Unnecessary Lines

*The* NEW ENGLAND  
JOURNAL *of* MEDICINE

ESTABLISHED IN 1812

DECEMBER 28, 2006

VOL. 355 NO. 26

An Intervention to Decrease Catheter-Related Bloodstream  
Infections in the ICU

Peter Pronovost, M.D., Ph.D., Dale Needham, M.D., Ph.D., Sean Berenholtz, M.D., David Sinopoli, M.P.H., M.B.A.,  
Haitao Chu, M.D., Ph.D., Sara Cosgrove, M.D., Bryan Sexton, Ph.D., Robert Hyzy, M.D., Robert Welsh, M.D.,  
Gary Roth, M.D., Joseph Bander, M.D., John Kepros, M.D., and Christine Goeschel, R.N., M.P.A.

# Checklist for Prevention of Central Line Associated Blood Stream Infections

*Based on 2011 CDC guideline for prevention of intravascular catheter-associated bloodstream infections:  
<http://www.cdc.gov/hicpac/pdf/guidelines/bsi-guidelines-2011.pdf>*

## **For Clinicians:**

### Promptly remove unnecessary central lines

- Perform daily audits to assess whether each central line is still needed

### Follow proper insertion practices

- Perform hand hygiene before insertion
- Adhere to aseptic technique
- Use maximal sterile barrier precautions (i.e., mask, cap, gown, sterile gloves, and sterile full-body drape)
- Perform skin antisepsis with >0.5% chlorhexidine with alcohol
- Choose the best site to minimize infections and mechanical complications
  - Avoid femoral site in adult patients
- Cover the site with sterile gauze or sterile, transparent, semipermeable dressings

### Handle and maintain central lines appropriately

- Comply with hand hygiene requirements
- Scrub the access port or hub immediately prior to each use with an appropriate antiseptic (e.g., chlorhexidine, povidone iodine, an iodophor, or 70% alcohol)
- Access catheters only with sterile devices
- Replace dressings that are wet, soiled, or dislodged
- Perform dressing changes under aseptic technique using clean or sterile gloves



## For Facilities:

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

## Supplemental strategies for consideration:

- 2% Chlorhexidine bathing
- Antimicrobial/Antiseptic-impregnated catheters
- Chlorhexidine-impregnated dressings

National Center for Emerging and Zoonotic Infectious Diseases

Division of Healthcare Quality Promotion





JOHNS HOPKINS  
M E D I C I N E

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QUALITY AND SAFETY  
RESEARCH GROUP

JOHNS HOPKINS QUALITY AND SAFETY RESEARCH GROUP (QSRG)

**ON THE CUSP: STOP BSI**  
CENTRAL LINE-ASSOCIATED BLOOD STREAM INFECTION  
TOOLKIT

## How to Use This Toolkit

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The purpose of this toolkit is to support your efforts to implement evidence-based practices and eliminate Central Line Associated Blood Stream Infections (CLABSIs) in your clinical area. The strategies in this toolkit have nearly eliminated CLABSIs in participating Michigan ICUs (Appendix A). These strategies have been adopted by over 100 ICUs in large and small, academic and community hospitals that we have worked with to date. Most of these ICUs have demonstrated a significant reduction in their CLABSI rates and many have not had a CLABSI in >6 months.

Nevertheless, your leadership is needed to achieve these results in your clinical area. Most of your efforts will be working with staff that insert and assist with the insertion of central lines. We developed a model to help disseminate this, and other, interventions. This model includes 4 stages that answer the following questions:

1. **Engage:** How will this make the world a better place?
2. **Educate:** How will we do this?
3. **Execute:** What do I need to do?
4. **Evaluate:** How will we know we made a difference?

This toolkit provides details of what you should do in each of these stages. In the appendices, we provide all the tools you will need to eliminate CLABSIs in your clinical area; the rest is up to you.

## 'BUNDLE' GAVECELT PER LA PREVENZIONE DELLE INFEZIONI ASSOCIATE A CATETERI VENOSI CENTRALI NON TUNNELLIZZATI A BREVE E MEDIO TERMINE



- Igiene delle mani e Massime precauzioni di barriera durante l'impianto del catetere venoso
- Scelta appropriata del sito di inserzione (in ordine di preferenza: metà braccio, zona sottoclaveare, zona sopraclaveare, collo, inguine)
- Impianto ecoguidato, ovunque possibile, sia per i cateteri a inserzione centrale che per i cateteri a inserzione periferica
- Utilizzo di clorexidina al 2% per la disinfezione cutanea prima dell'inserzione nonché per la disinfezione continua o discontinua dell'exit site
- Impiego di "sutureless devices" per il fissaggio del catetere
- Impiego di medicazioni semipermeabili trasparenti, ovunque possibile
- Rimozione immediata del catetere venoso centrale non più indispensabile



# Targeting zero CLABSI in patients with PICC lines: a case-control study

**G. Scoppettuolo§, L. Dolcetti§, C. Taraschi§, C. Chiarini§, C. Donato§, S. Lardo§, A. La Greca\*, M. Pittiruti\***  
**§ Clinic of Infectious Diseases, \* Dpt. of Surgery, Catholic University, Rome**

RESEARCH

Open Access

## Clinical experience with power-injectable PICCs in intensive care patients

Mauro Pittiruti<sup>1\*</sup>, Alberto Brutti<sup>2</sup>, Davide Celentano<sup>2</sup>, Massimiliano Pomponi<sup>2</sup>, Daniele G Biasucci<sup>2</sup>, Maria Giuseppina Annetta<sup>2</sup> and Giancarlo Scoppettuolo<sup>3</sup>

See related Letter by Zampieri,

<http://ccforum.com/content/16/2/425>

### Abstract

**Introduction:** In the ICU, peripherally inserted central catheters (PICCs) may be an alternative option to standard central venous catheters, particularly in patients with coagulation disorders or at high risk for infection. Some limits of PICCs (such as low flow rates) may be overcome with the use of power-injectable catheters.

**Methods:** We retrospectively reviewed all of the power-injectable PICCs inserted in adult and pediatric patients in the ICU during a 12-month period, focusing on the rate of complications at insertion and during maintenance.

**Results:** We collected 89 power-injectable PICCs (in adults and in children), both multiple and single lumen. All insertions were successful. There were no major complications at insertion and no episodes of catheter-related bloodstream infection. Non-infective complications during management were not clinically significant. There was one episode of symptomatic thrombosis during the stay in the ICU and one episode after transfer of a patient to a non-intensive ward.

**Conclusion:** Power-injectable PICCs have many advantages in the ICU: they can be used as multipurpose central lines for any type of infusion including high-flow infusion, for hemodynamic monitoring, and for high-pressure injection of contrast media during radiological procedures. Their insertion is successful in 100% of cases and is not associated with significant risks, even in patients with coagulation disorders. Their maintenance is associated with an extremely low rate of infective and non-infective complications.

## Targeting zero catheter-related bloodstream infections in pediatric intensive care unit: a retrospective matched case-control study.

[Biasucci DG](#)<sup>1</sup>, [Pittiruti M](#)<sup>2</sup>, [Taddei A](#)<sup>3</sup>, [Picconi E](#)<sup>1</sup>, [Pizza A](#)<sup>1</sup>, [Celentano D](#)<sup>1</sup>, [Piastra M](#)<sup>1</sup>, [Scoppettuolo G](#)<sup>4</sup>, [Conti G](#)<sup>1</sup>.

### Author information

#### Abstract

**INTRODUCTION:** The aim of this study was to evaluate the effectiveness and safety of a new three-component 'bundle' for insertion and management of centrally inserted central catheters (CICCs), designed to minimize catheter-related bloodstream infections (CRBSIs) in critically ill children.

**METHODS:** Our 'bundle' has three components: insertion, management, and education. Insertion and management recommendations include: skin antisepsis with 2% chlorhexidine; maximal barrier precautions; ultrasound-guided venipuncture; tunneling of the catheter when a long indwelling time is expected; glue on the exit site; sutureless securement; use of transparent dressing; chlorhexidine sponge dressing on the 7th day; neutral displacement needle-free connectors. All CICCs were inserted by appropriately trained physicians proficient in a standardized simulation training program.

**RESULTS:** We compared CRBSI rate per 1000 catheters-days of CICCs inserted before adoption of our new bundle with that of CICCs inserted after implementation of the bundle. CICCs inserted after adoption of the bundle remained in place for a mean of 2.2 days longer than those inserted before. We found a drop in CRBSI rate to 10%, from 15 per 1000 catheters-days to 1.5.

**CONCLUSIONS:** Our data suggest that a bundle aimed at minimizing CR-BSI in critically ill children should incorporate four practices: (1) ultrasound guidance, which minimizes contamination by reducing the number of attempts and possible break-down of aseptic technique; (2) tunneling the catheter to obtain exit site in the infra-clavicular area with reduced bacterial colonization; (3) glue, which seals and protects the exit site; (4) simulation-based education of the staff.

# Vascular Access Team



[Antimicrobial Stewardship & Healthcare Epidemiology](#)

## Article contents

[Abstract](#)

## The effectiveness of a dedicated central venous access care team to prevent catheter-related bloodstream infections

Published online by Cambridge University Press: 16 May 2022

[Fortune Charles Fil de Lara](#), [Maria Jesusa Mano](#), [Karl Evans Henson](#), [Jia An Bello](#) and [Cybele Lara Abad](#)

Article

Metrics



### Abstract



Core share and HTML view are not possible as this article does not have html content. However, as you have access to this content, a full PDF is available via the 'Save PDF' action button.

**Background:** Catheter-related bloodstream infection (CRBSI) rates remain high despite the use of an insertion bundle. We hypothesized that line care and maintenance by a dedicated team would help decrease CRBSI rates. This study was conducted in The Medical City (TMC), is a 526-bed, private, tertiary-care center in Pasig City, Philippines. **Methods:** All adult hospitalized patients from October 1, 2020, to October 31, 2021, with a newly inserted temporary central venous catheter (CVC) were eligible for inclusion. CRBSI rates before the intervention (October 2019 to March 2020) and after the intervention (April to October 2021) were compared. The intervention arm consisted of a dedicated central venous access team (CVAT) who provided



# Vascular Access Team Policlinico Gemelli

- Multiprofessionale
  - Vascular Access Team Infermieristico (1 Coordinatrice, 5 Infermieri bedside, 6 Infermieri DH)
  - Centro Interdipartimentale Accessi Venosi Centrali (5 Medici)
- Multidisciplinare
  - Infermieri di varie aree (Oncologia, Pediatria, Ginecologia Oncologica, Radioterapia, Ematologia, Malattie Infettive, Urologia, Neurochirurgia)
  - Medici di varie discipline (2 Chirurghi, 1 Anestesista Rianimatore, 1 Infettivologo, 1 Angiologo)
- Setting
  - Tutti i Reparti del Policlinico, ad eccezione delle Terapie Intensive e della Dialisi (ma collaborazione continuativa con il Nefrologo responsabile della Emodialisi per inpatients)
  - Consulenze per scelta accesso venoso, impianto, gestione, gestione delle complicanze

# Skills Infermieri Vascular Access Team

COMPETENZE SPECIFICHE	MINIMIDLINE	MIDLINE	PICC	CICC/FICC/PORT/PICC PORT
	GRUPPO A	GRUPPO B	GRUPPO C	GRUPPO D
	21 OPERATORI	16 OPERATORI	15 OPERATORI	OPERATORI 6
Indicazione appropriata al dispositivo	X	X	X	x
Tecnica asettica appropriata	X	X	X	x
Tecnica di anestesia locale	X	X	X	x
puntura ecoguidata vene braccio	X	X	X	x
puntura ecoguidata vene centrali				x
puntura ecoguidata all'inguine				x
tecnica di tunnellizzazione			X	x
tip location con ECG intracavitario			X	x
tip location con bubble test ed ecocardiogramma trans toracico				x
confezione della tasca				x
sutura intradermica				x
uso del cianoacrilato	X	X	X	x
tecniche di fissaggio e medicazione	X	X	X	x

# CATHETER STEWARDSHIP

- RIGHT CATHETER
- RIGHT PATIENT
- RIGHT TIME

**INFECTION PREVENTION**



# PROACTIVE VASCULAR ACCESS PLANNING

Reprinted from JOURNAL OF INFUSION NURSING  
Vol. 28 No. 3S May/June 2005  
Copyright © 2005 by Lippincott Williams & Wilkins  
Printed in U.S.A.

Kathy Kokotis, RN, BS, MBA

## Cost Containment and Infusion Services

### Abstract

*The implementation of the Medicare Prospective Payment System (MPPS) has placed pressure on healthcare organizations to decrease patient length of stay without adversely affecting outcomes. This article discusses the impact of the MPPS on clinicians who provide infusion therapy, and examines methods for containing costs related to infusion care such as advanced planning and accurate vascular access device selection.*

The overall makeup of the healthcare system transitioned after the implementation of the Medicare Prospective Payment System in the 1990s. "Revenue production on a hospital inpatient" became a passé label, and "revenue saving on the DRG [diagnosis-related group]" took its place. The new aim is to decrease hospital length of stay (LOS) without altering patient outcome. What impact does this change have on the skilled infusion professional?

A hospital administrator may envision an infusion team as disposable because it is assumed that any nurse can insert a conventional peripheral catheter. But is it true that all nurses will insert a catheter with the same level of skill? A Press Ganey survey of almost 1.8 million patients in more than 1,000 hospitals shows that this is not true.<sup>1</sup> In fact, 58% of patients are dissatisfied with the venipuncture skill level of their nurse, and 52% are not satisfied with the courtesy of the nurse inserting the catheter.<sup>1</sup> Barton et al<sup>2</sup> and Danek and Kilroy<sup>3</sup> from the University of Florida indicate that a clinician requires 2.18 attempts to achieve a successful catheter insertion. Therefore, it appears that

- Cost reduction with Vascular Access Planning
- Cost reduction with increased proficiency in venipuncture
- Using high-tech tools to improve Picc insertion performance
- Revenue saved by reducing Catheter Related Bloodstream Infections
- Salvage of an occluded catheter

MACOV@ 2023



Connect

Kathy Kokotis

Reserve your place



LINEE GUIDA	Rev.: 1
PARTE PRIMA: RACCOMANDAZIONI PER LA SCELTA CORRETTA DELL'ACCESSO VENOSO	LG.029

LINEE GUIDA

Parte prima: Raccomandazioni per la scelta corretta dell'accesso venoso  
 LG.029

	NOME	FUNZIONE	DATA	FIRMA
Redatto da:	Gruppo di Lavoro			
Verificato da:	Giuseppe Vetrugno	Responsabile Risk Management		
	Maurizio Zega	Responsabile SITRA		
	Alberto Fiore	Responsabile Qualità e Accreditamento		
	Patrizia Laurenti	Responsabile Igiene Ospedaliera	10/3/2021	
	Maria Elena D'Alfonso	Direttore Presidio Gemelli	8/3/21	
	CIO			
Approvato da:	Andrea Cambieri	Direttore Sanitario	8/3/21	
	Rocco Bellantone	Direttore Governo Clinico		

Livello organizzativo di applicazione:

- Aziendale
- Dipartimento
- Unità Operativa



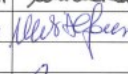




STATO DELLE REVISIONI

Rev. N.	PARAGRAFI REVISIONATI	DESCRIZIONE REVISIONE	DATA
0		Prima Stesura	14/09/2014
1	Intera procedura e allegati	Revisione in coerenza con evidenze scientifiche aggiornate e nuovi processi organizzativi definiti	15/02/2021

LINEE GUIDA	Rev.: 1
PARTE SECONDA: RACCOMANDAZIONI PER IL CORRETTO IMPIANTO DELL'ACCESSO VENOSO	LG.030

LINEE GUIDA

Parte seconda: Raccomandazioni per il corretto impianto dell'accesso venoso  
 LG.030

	NOME	FUNZIONE	DATA	FIRMA
Redatto da:	Gruppo di Lavoro			
Verificato da:	Giuseppe Vetrugno	Responsabile Risk Management		
	Maurizio Zega	Responsabile SITRA		
	Alberto Fiore	Responsabile Qualità e Accreditamento		
	Patrizia Laurenti	Direttore Igiene Ospedaliera	10/3/2021	
	Maria Elena D'Alfonso	Direttore Presidio Gemelli	8/3/21	
	CIO			
Approvato da:	Andrea Cambieri	Direttore Sanitario	8/3/21	
	Rocco Bellantone	Direttore Governo Clinico		

Livello organizzativo di applicazione:

- Aziendale
- Dipartimento
- Unità Operativa


STATO DELLE REVISIONI

Rev. N.	PARAGRAFI REVISIONATI	DESCRIZIONE REVISIONE	DATA
0		Prima Stesura	14/09/2014
1	Intera procedura e allegati	Revisione in coerenza con evidenze scientifiche aggiornate e nuovi processi organizzativi definiti	15/02/2021

LINEE GUIDA	Rev.: 1
PARTE TERZA: RACCOMANDAZIONI PER LA GESTIONE CORRETTA DELL'ACCESSO VENOSO	LG.031

LINEE GUIDA

Parte terza: Raccomandazioni per la gestione corretta dell'accesso venoso  
 LG.031

	NOME	FUNZIONE	DATA	FIRMA
Redatto da:	Gruppo di Lavoro			
Verificato da:	Giuseppe Vetrugno	Responsabile Risk Management		
	Maurizio Zega	Responsabile SITRA		
	Alberto Fiore	Responsabile Qualità e Accreditamento		
	Patrizia Laurenti	Direttore Igiene Ospedaliera	10/3/21	
	Maria Elena D'Alfonso	Direttore Presidio Gemelli	8/3/21	
	CIO			
Approvato da:	Andrea Cambieri	Direttore Sanitario	8/3/21	
	Rocco Bellantone	Direttore Governo Clinico		

Livello organizzativo di applicazione:

- Aziendale
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STATO DELLE REVISIONI

Rev. N.	PARAGRAFI REVISIONATI	DESCRIZIONE REVISIONE	DATA
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1	Intera procedura	Revisione in coerenza con evidenze scientifiche aggiornate e nuovi processi organizzativi definiti	15/02/2021

# Un algoritmo GAVeCeLT 2018



**DAV  
Exp  
GAVeCeLT**

**Sistema esperto  
per la scelta  
del dispositivo  
per accesso venoso**

**Scelta del dispositivo  
per accesso venoso**

Scegli la tipologia di paziente

opzioni:

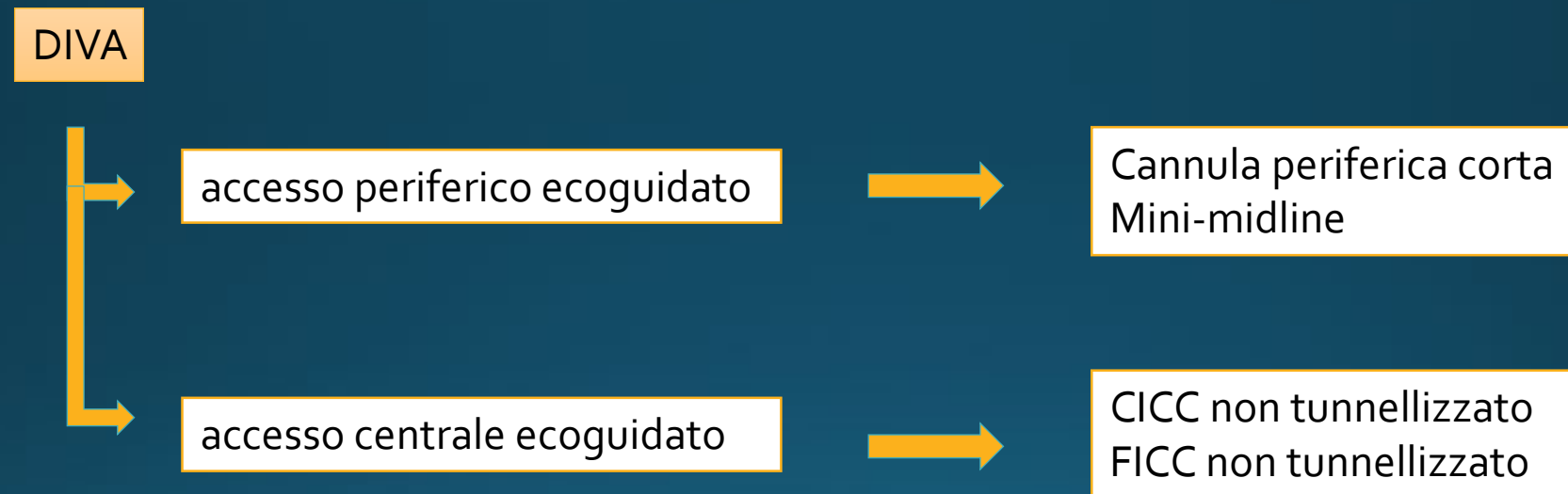
**Bambino**

**Neonato**

**Adulto**

Adulto - Emergenza

Non DIVA → cannula periferica corta 'standard'

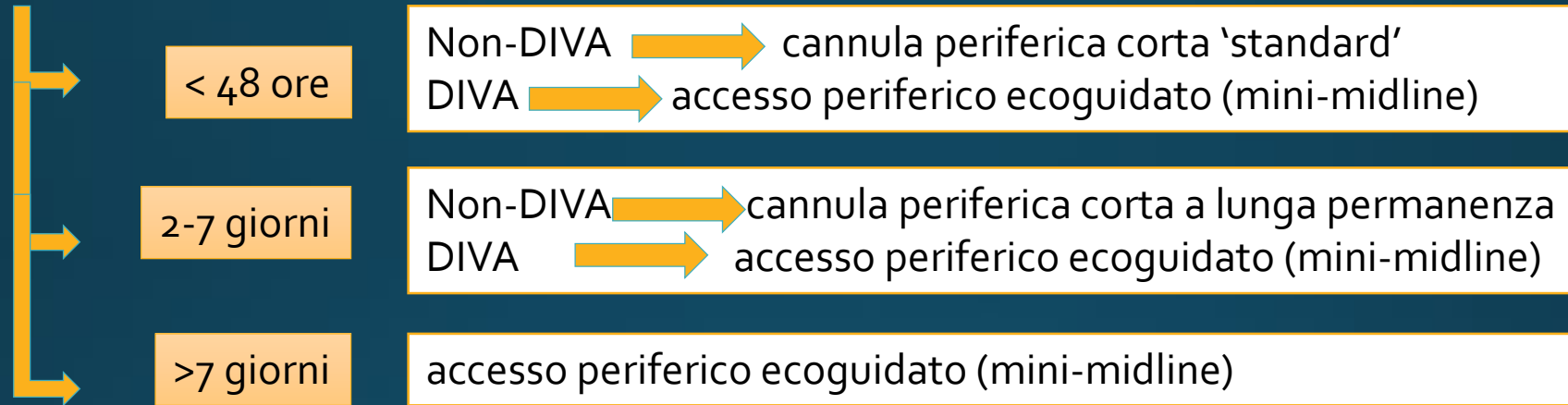


Adulto- Elezione

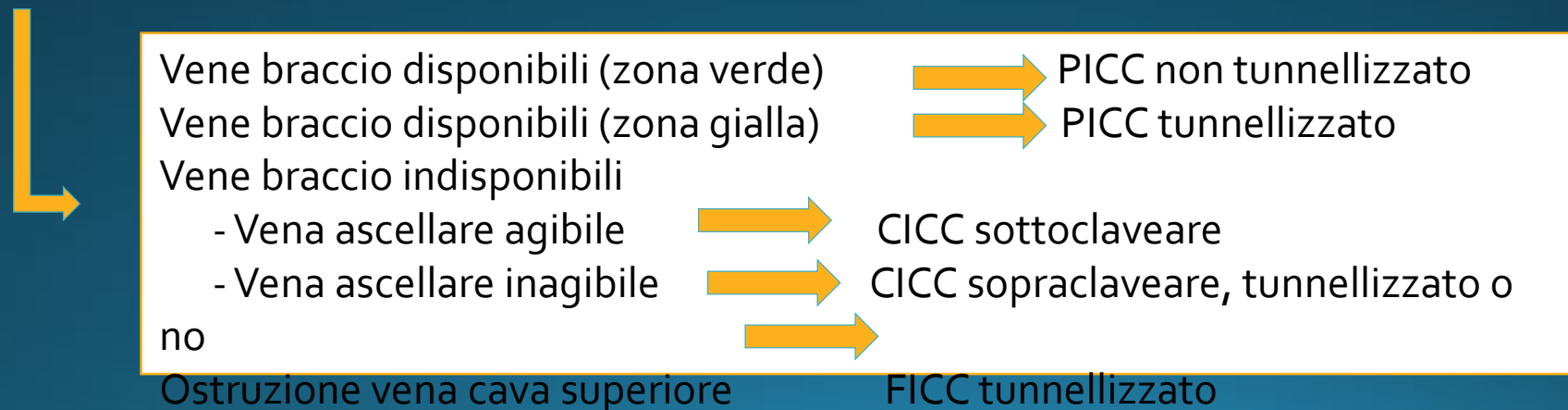
Uso intraospedaliero



### Sufficiente accesso periferico



### Necessità accesso centrale





Adulto- Elezione

Uso extraospedaliero

<4-6 mesi



Sufficiente accesso periferico



< 3-4 sett.: mini-midline  
> 3-4 sett.: midline

Necessità accesso centrale



Vene braccio disponibili : PICC non cuffiato, tunnellizzato o no  
Vene braccio indisponibili: CICC non cuffiato, tunnellizzato  
Ostruzione vena cava sup.: FICC non cuffiato, tunnellizzato

>4-6 mesi



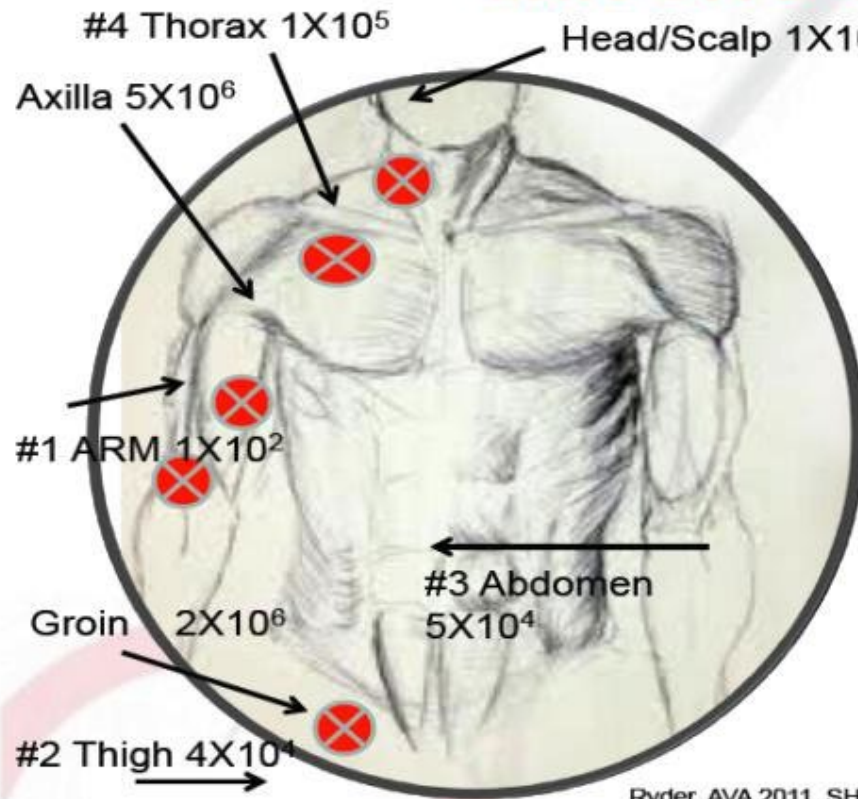
Uso frequente

Vene braccio disponibili : PICC cuffiato opp. tunnellizzato con SAS  
Vene braccio indisponibili: CICC cuffiato opp. tunnellizzato con SAS

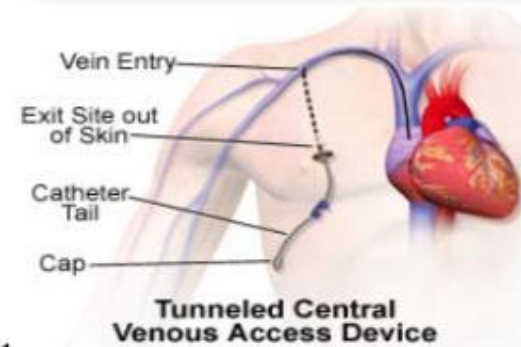
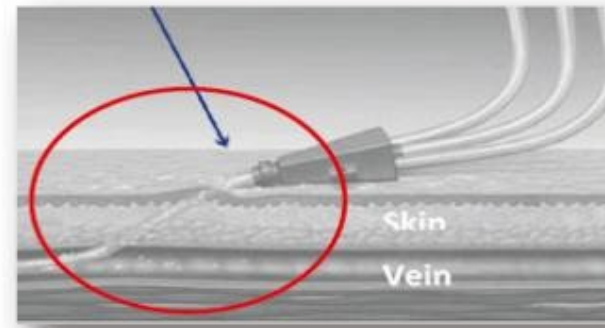
Uso infrequente

Vene braccio disponibili : PICC- port  
Vene braccio indisponibili: port toracico  
Ostruzione vena cava sup.: port femorale oppure  
FICC cuffiato  
opp. tunnellizzato con SAS

# Exit Site Options



Ryder, AVA 2011, SHEA 2011



# Protocollo PIDAV



**1. CORRETTA INDICAZIONE** - Verifica dell'indicazione all'accesso venoso, scelta del dispositivo più appropriato (periferico vs. centrale) e sua rimozione appena non è più indispensabile.



**2. CORRETTA ASEPSI** - Igiene delle mani con gel idroalcolico, prima dell'impianto e prima e dopo ogni manovra di gestione; massime precauzioni di barriera durante l'inserzione di dispositivi per accesso centrale o accesso periferico di lunga durata; antisepsi cutanea con clorexidina 2% in alcool - in applicatori monodose sterili - prima dell'impianto e al momento del cambio della medicazione.



**3. SCELTA CORRETTA DEL SITO DI EMERGENZA** – Per gli accessi periferici, evitare le zone di flessione; per gli accessi centrali, preferire (nell'ordine) il terzo medio del braccio, la zona sottoclaveare e la zona sopraclaveare; evitare il sito di emergenza al collo o all'inguine (tranne che in urgenza); tunnellizzare un dispositivo per accesso venoso centrale se ciò è necessario per ottimizzare il sito di emergenza.



**4. TECNICA CORRETTA DI IMPIANTO** - Utilizzare sempre l'impianto ecoguidato per il posizionamento dei dispositivi centrali e dei dispositivi periferici di lunga durata.

# Protocollo PIDAV



**5. FISSAGGIO APPROPRIATO:** evitare sempre punti di sutura e cerotti; stabilizzare invece il dispositivo con un sistema *sutureless* appropriato (integrato nella medicazione o ad adesività cutanea o ad ancoraggio sottocutaneo).



**6. PROTEZIONE DEL SITO DI EMERGENZA** - Utilizzare membrane trasparenti semipermeabili ad alta traspirabilità, associate a feltrini a rilascio di clorexidina o a sigillo del sito di emergenza con colla al cianoacrilato.



**7. PROTEGGERE LA LINEA INFUSIONALE:** - Disinfettare le porte di accesso strofinando con soluzioni alcoliche alla clorexidina 2% oppure applicando *port protectors* (cappucci disinfettanti) sopra ai connettori senz'ago; lavare e chiudere il sistema soltanto con soluzione fisiologica, usando siringhe preriempite.



**8. FACILITARE L'ADOZIONE DEL BUNDLE** - Utilizzare carrelli dedicati, kit omnicomprensivi e *checklist*, sia per l'impianto che per la gestione.

# CHECKLIST IMPIANTO

Bundle per l'inserzione del catetere venoso centrale

Data 20/11/2023

15:41

## Percentuale di compliance al bundle

### Prima della procedura

Il paziente è stato identificato?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stata verificata la corretta indicazione all'accesso venoso centrale e l'assenza di eventuali controindicazioni (allergie, rischio infettivo, stato coagulativo)?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Il paziente è stato informato sull'indicazione e le complicanze (sottoscrizione consenso informato)?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stata verificata la presenza di tutto il materiale necessario per la procedura?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stato effettuato lo studio ecografico pre-procedurale delle vene del paziente e contrassegnato il sito di inserzione?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Il paziente è stato posizionato correttamente?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stata eseguita l'igiene delle mani secondo protocollo?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stata eseguita antisepsi cutanea con clorexidina 2% in alcool isopropilico al 70% o, in caso di nota intolleranza alla clorexidina, con iodopovidone 10%?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Sono state messe in atto le massime precauzioni di barriera (mascherina non sterile, cuffia non sterile, guanti sterili, camice sterile, telo sterile "full body", coprisonda sterile)?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A

### Durante la procedura

L'utilizzo di anestesia locale e/o sedazione è appropriato secondo indicazione?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
La venipuntura è ecoguidata?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stata confermata ecograficamente la corretta posizione intravenosa della guida e della sua direzione?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stato verificato il posizionamento intravascolare del catetere mediante aspirazione di sangue e lavaggio con soluzione fisiologica?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stato eseguito il controllo intra-procedurale della posizione della punta del catetere mediante ECG intracavitario e/o ecocardiogramma?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stato eseguito il flush e lock del catetere?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stato chiuso il catetere con needlefree connector e applicato il port protector?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stato eseguito il fissaggio con sistema sutureless?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stata utilizzata la colla istocrilica per sigillare il sito di emergenza e per la eventuale chiusura di brecche cutanee?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
È stata coperta con medicazione adesiva semipermeabile trasparente e indicata la data?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A
Confermato il mantenimento del campo sterile per tutta la durata della procedura?	<input type="checkbox"/> Sì	<input type="checkbox"/> No	<input type="checkbox"/> N/A

Note

# CHECKLIST GESTIONE (MEDICAZIONE E LINEA INFUSIONALE)

## Procedura di medicazione

Sono stati utilizzati guanti puliti non sterili?

Sì  No  N/A

È stata eseguita palpazione del sito di emergenza per verificare eventuale dolenzia?

Sì  No  N/A

È stata rimossa la membrana trasparente e il feltrino a rilascio di clorexidina (se presente)?

Sì  No  N/A

È stato ispezionato visivamente il sito di emergenza?

Sì  No  N/A

È stato rimosso il sistema sutureless ad adesività cutanea (se non è presente un sistema ad ancoraggio sottocutaneo)?

Sì  No  N/A

È stata eseguita antisepsi cutanea con clorexidina 2% in alcool isopropilico al 70% o, in caso di nota intolleranza alla clorexidina, con iodopovidone 10%?

Sì  No  N/A

È stato aperto il materiale necessario per la nuova medicazione?

Sì  No  N/A

Sono stati utilizzati guanti sterili, dopo nuova igiene delle mani secondo protocollo?

Sì  No  N/A

È stata applicata nuova medicazione: feltrino a rilascio di clorexidina (se indicato), sistema sutureless ad adesività cutanea (se non è presente un sistema ad ancoraggio sottocutaneo), e membrana adesiva trasparente semipermeabile (con apposizione della data)

Sì  No  N/A

Confermato il mantenimento del campo sterile per tutta la durata della procedura?

Sì  No  N/A

## Procedura di lavaggio

Sono stati utilizzati guanti puliti non sterili?

Sì  No  N/A

È stato effettuato il clampaggio della linea infusionale e la rimozione del needlefree connector

Sì  No  N/A

È stata eseguita la disinfezione del cono di connessione?

Sì  No  N/A

È stata effettuata applicazione di nuovo needlefree connector e declampaggio della linea infusionale

Sì  No  N/A

È stato effettuato il lavaggio pulsante con 10ml di soluzione fisiologica

Sì  No  N/A

È stato applicato il port protector?

Sì  No  N/A

# VASCULAR ACCESS TEAM ED EMODIALISI

## CDC Approach to BSI Prevention in Dialysis Facilities

(i.e., the Core Interventions for Dialysis Bloodstream Infection (BSI) Prevention)

### 1. Surveillance and feedback using NHSN

Conduct monthly surveillance for BSIs and other dialysis events using CDC's National Healthcare Safety Network (NHSN). Calculate facility rates and compare to rates in other NHSN facilities. Actively share results with front-line clinical staff.

### 2. Hand hygiene observations

Perform observations of hand hygiene opportunities monthly and share results with clinical staff.

### 3. Catheter/vascular access care observations

Perform observations of vascular access care and catheter accessing quarterly. Assess staff adherence to aseptic technique when connecting and disconnecting catheters and during dressing changes. Share results with clinical staff.

### 4. Staff education and competency

Train staff on infection control topics, including access care and aseptic technique. Perform competency evaluation for skills such as catheter care and accessing every 6-12 months and upon hire.

### 5. Patient education/engagement

Provide standardized education to all patients on infection prevention topics including vascular access care, hand hygiene, risks related to catheter use, recognizing signs of infection, and instructions for access management when away from the dialysis unit.

### 6. Catheter reduction

Incorporate efforts (e.g., through patient education, vascular access coordinator) to reduce catheters by identifying and addressing barriers to permanent vascular access placement and catheter removal.

### 7. Chlorhexidine for skin antisepsis

Use an alcohol-based chlorhexidine (>0.5%) solution as the first line skin antiseptic agent for central line insertion and during dressing changes.\*

### 8. Catheter hub disinfection

Scrub catheter hubs with an appropriate antiseptic after cap is removed and before accessing. Perform every time catheter is accessed or disconnected.\*\*

### 9. Antimicrobial ointment

Apply antibiotic ointment or povidone-iodine ointment to catheter exit sites during dressing change.\*\*\*

\* Povidone-iodine (preferably with alcohol) or 70% alcohol are alternatives for patients with chlorhexidine intolerance.

\*\* If closed needleless connector device is used, disinfect device per manufacturer's instructions.

\*\*\* See information on selecting an antimicrobial ointment for hemodialysis catheter exit sites on CDC's Dialysis Safety website (<http://www.cdc.gov/dialysis/prevention-tools/core-interventions.html#sites>). Use of chlorhexidine-impregnated sponge dressing might be an alternative.



For more information about the Core Interventions for Dialysis Bloodstream Infection (BSI) Prevention, please visit <http://www.cdc.gov/dialysis>

# Catheter connection and disconnection checklists

## Checklist: Hemodialysis catheter connection

- Wear mask (if required)
- Perform hand hygiene
- Put on new, clean gloves
- Clamp the catheter and remove caps
- Scrub catheter hub with antiseptic
- Allow hub antiseptic to dry
- Connect catheter to blood lines aseptically
- Remove gloves
- Perform hand hygiene



## Checklist: Hemodialysis catheter disconnection

- Wear mask (if required)
- Perform hand hygiene
- Put on new, clean gloves
- Clamp the catheter
- Disconnect catheter from blood lines aseptically
- Scrub catheter hub with antiseptic
- Allow hub antiseptic to dry
- Attach new caps aseptically
- Remove gloves
- Perform hand hygiene





# Hemodialysis Central Venous Catheter Scrub-the-Hub Protocol

This protocol outlines a suggested approach to preparing catheter hubs prior to accessing the catheter for hemodialysis. It is based on evidence where available and incorporates theoretical rationale when published evidence is unavailable.

## Definitions:

**Catheter** refers to a central venous catheter (CVC) or a central line

**Hub** refers to the end of the CVC that connects to the blood lines or cap

**Cap** refers to a device that screws on to and occludes the hub

**Limb** refers to the catheter portion that extends from the patient's body to the hub

**Blood lines** refer to the arterial and venous ends of the extracorporeal circuit that connect the patient's catheter to the dialyzer

## Catheter Connection and Disconnection Steps:

### Connection Steps

1. Perform hand hygiene and don new clean gloves.
2. Clamp the catheter (*Note: Always* clamp the catheter before removing the cap. Never leave an uncapped catheter unattended).
3. Disinfect the hub with caps removed using an appropriate antiseptic (*see notes*).
  - a. (*Optional*) Prior to cap removal, disinfect the caps and the part of the hub that is accessible and discard the antiseptic pad (i.e., use a separate antiseptic pad for the next step).
  - b. Remove the caps and disinfect the hub with a new antiseptic pad for each hub. Scrub the sides (threads) and end of the hub thoroughly with friction, making sure to remove any residue (e.g., blood).
  - c. Using the same antiseptic pad, apply antiseptic with friction to the catheter, moving from the hub at least several centimeters towards the body. Hold the limb while allowing the antiseptic to dry.
  - d. Use a separate antiseptic pad for each hub/catheter limb. Leave hubs "open" (i.e., uncapped and disconnected) for the shortest time possible.

4. Always handle the catheter hubs aseptically. Once disinfected, do not allow the catheter hubs to touch nonsterile surfaces.
5. Attach sterile syringe, unclamp the catheter, withdraw blood, and flush per facility protocol.
6. Repeat for other limb (this might occur in parallel).
7. Connect the ends of the blood lines to the catheter aseptically.
8. Remove gloves and perform hand hygiene.

### Disconnection Steps:

1. Perform hand hygiene and don new clean gloves.
2. Clamp the catheter (*Note: Always* clamp the catheter before disconnecting. Never leave an uncapped catheter unattended).
3. Disinfect the catheter hub before applying the new cap using an appropriate antiseptic (*see notes*).
  - a. (*Optional*) Disinfect the connection prior to disconnection. If this is done, use a separate antiseptic pad for the subsequent disinfection of the hub.
  - b. Disconnect the blood line from the catheter and disinfect the hub with a new antiseptic pad. Scrub the sides (threads) and end of the hub thoroughly with friction, making sure to remove any residue (e.g., blood).
  - c. Use a separate antiseptic pad for each hub. Leave hubs "open" (i.e., uncapped and disconnected) for the shortest time possible.
4. Always handle the catheter hubs aseptically. Once disinfected, do not allow the catheter hubs to touch nonsterile surfaces. Hold the catheter until the antiseptic has dried.
5. Attach the new sterile caps to the catheter aseptically. Use caution if tape is used to secure caps to the catheter (*see notes*).
6. Ensure that catheter is still clamped.
7. Remove gloves and perform hand hygiene.

## Notes/Discussion:

### Antiseptic Use and Selection

As described in the 2011 CDC/Healthcare Infection Control Practices Advisory Committee (HICPAC) Guidelines for the Prevention of Intravascular Catheter-Related Infections, prior to accessing the catheter hub it should be disinfected with an appropriate antiseptic (greater than 0.5% chlorhexidine with alcohol, 70% alcohol, or 10% povidone-iodine). There is not enough evidence to recommend one antiseptic over the others. Generally, antiseptics should be allowed to dry for maximal effect.

If using 70% alcohol, sterile antiseptic pads should be used (sterile pads are labeled sterile and packaging for nonsterile pads often does not state whether the pads are sterile or nonsterile). For practical reasons, pads or similar products might be preferred over other forms of antiseptics (e.g., swabsticks) for disinfecting the catheter as they are malleable and allow for vigorous cleaning of small spaces.

If using an antiseptic that leaves a residue (e.g., chlorhexidine), avoid allowing large amounts of antiseptic to enter the lumen of the catheter to avoid potential toxicities to the patient.

If using chlorhexidine, removing all blood residue is particularly important to maximize the effect of the antiseptic.

### Soaking Caps

The role of soaking caps in an antiseptic prior to removing them is not clear. It is not a CDC/HICPAC recommendation. This procedure is described in the 2000 National Kidney Foundation's Kidney Disease Outcomes Quality Initiative (KDOQI) Vascular Access Guidelines but was not included in the 2006 update.

### Handling Catheter Hubs

Catheter hubs should always be handled aseptically. Once disinfected, the catheter hubs should not be allowed to touch nonsterile surfaces. This might be best performed by holding them until the antiseptic dries. During this time, the staff member performing the procedure should also ensure that the catheter remains clamped.

When disinfecting catheter hubs, clean, nonsterile gloves can be used if aseptic technique is maintained.

### Bloodline Disinfection

When accessing the line, disinfecting the ends of the sterile blood lines is not required if care has been taken not to contaminate the ends of the blood lines (i.e., through careful aseptic technique). Blood lines can become contaminated during connections and disconnections, as well as during the priming process. Contact with contaminated prime waste in prime buckets that have not been properly cleaned and disinfected or through backflow from waste handling ports must be avoided. Disinfecting the bloodlines does not address this issue.

### Disconnection and Line Reversals

Catheter hubs should be disinfected again after disconnecting from bloodlines and before replacing a new cap at the end of a treatment. This should be done in a manner similar to that used when disinfecting the hub prior to accessing. Disinfecting the catheter hub and the end of the extracorporeal blood line should also be performed if, during a treatment, a patient must be disconnected and their blood is re-circulated. Anytime a patient's circuit is disconnected this should be done aseptically and the number of times a patient's catheter is disconnected from the blood lines should be minimized to the extent possible.

### Securing Caps with Tape

Caution should be used if taping caps on to hubs between treatments. Tape can leave residue on the hubs that might make disinfecting them more difficult.

### Use of Masks

Although data supporting the use of masks during catheter accessing/deaccessing to prevent vascular access infections is lacking, this practice is recommended for patients and staff in the 2000 KDOQI guidelines and is included in the Centers for Medicare and Medicaid Services (CMS) End Stage Renal Disease Program Conditions for Coverage Interpretive Guidance.

### Personal Protective Equipment (PPE)

Proper PPE should always be worn by staff to avoid exposure to potentially infectious blood and body fluids when connecting/disconnecting catheters.

### Aseptic Technique

This includes practices that prevent the contamination of clean/sterile items and surfaces. Once tasks requiring aseptic technique have been started, care must be taken to avoid contamination of gloves and other clean/sterile items that can occur when touching dirty surfaces (e.g., positioning patient, using computer keyboard).

### Selected References:

1. National Kidney Foundation. KDOQI Clinical Practice Guidelines and Clinical Practice Recommendations for 2006 Updates: Hemodialysis Adequacy, Peritoneal Dialysis Adequacy and Vascular Access. *Am J Kidney Dis* 2006; 48 (suppl 1):S1-S322.
2. National Kidney Foundation. KDOQI Clinical Practice Guidelines for Hemodialysis Adequacy, 2000. *Am J Kidney Dis* 2001; 37 (suppl 1):S7-S64.
3. O'Grady NP, Alexander M, Burns LM, et al. Guideline for the prevention of intravascular catheter-related infections. *Clin Infect Dis* 2011; 52:e162-e193.



# KEYPOINTS

- IMPIANTO ECOGUIDATO
- ANTISETTICI (CLOREXIDINA GLUCONATO 2% IN IPA 70%)
- PACK PROCEDURALI
- SISTEMI DI FISSAGGIO
- PORT PROTECTORS
- SIRINGHE PRERIEMPITE STERILI PER FLUSH E LOCK DEL CATETERE
- LOCK PROPHYLAXIS



2003Gen15 00:29

Ven  
L25  
IM  
0,7  
ITT  
0,0  
Gen  
MB  
99%



## Real-time ultrasound-guided catheterisation of the internal jugular vein: a prospective comparison with the landmark technique in critical care patients

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Received: 23 May 2006 Revisions requested: 15 Jun 2006 Revisions received: 8 Sep 2006 Accepted: 10 Nov 2006 Published: 17 Nov 2006

Critical Care 2006, 10:R162 (doi:10.1186/cc5101)

This article is online at: <http://ccforum.com/content/10/6/R162>

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Critical Care, 2006

firmed by the present data. We found that the incidence of CVC-BSI in the ultrasound group of patients was significantly lower compared with that documented in the landmark group. The number of CVC-BSIs was significantly correlated to the number of needle passes in the total study population. We could speculate that repeated attempts might lead to a breakdown of aseptic technique and more colonisation of skin-related pathogens [17]. The above findings may be of clinical

**Table 2**

### Outcome measures in the ultrasound group versus the landmark group of patients

Outcome measures	Ultrasound group (n = 450)	Landmark group (n = 450)
Access time (seconds)	17.1 ± 16.5 (11.5 to 41.4)*	44 ± 95.4 (33.2 to 77.5)
Success rate	450 (100%)*	425 (94.4%)
Carotid puncture	5 (1.1%)*	48 (10.6%)
Haematoma	2 (0.4%)*	38 (8.4%)
Haemothorax	0 (0%)*	8 (1.7%)
Pneumothorax	0 (0%)*	11 (2.4%)
Average number of attempts	1.1 ± 0.6 (1.1 to 1.9)*	2.6 ± 2.9 (1.5 to 6.3)
CVC-BSI	47 (10.4%)*	72 (16%)

\*Comparison of the outcome measures between the ultrasound group and the landmark group of patients ( $p < 0.001$ ). Access time and average number of attempts are expressed as mean ± standard deviation (95% confidence interval). Success rate, carotid puncture, haematoma, haemothorax, pneumothorax, and CVC-BSI are expressed as the absolute number of patients and percentage of their group. CVC-BSI, central venous catheter-associated blood stream infection.



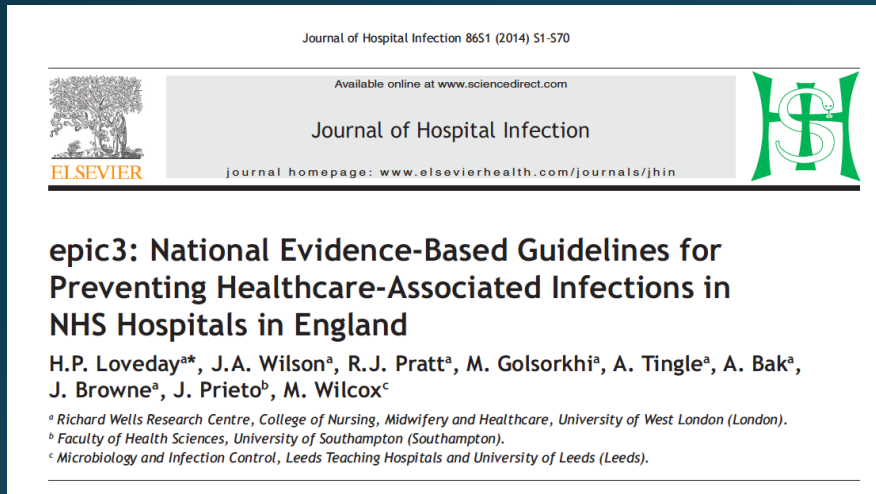
# CDC 2011

7. Usare la guida ecografica per posizionare i cateteri venosi centrali (ovunque questa tecnologia sia disponibile) così da ridurre il numero di tentativi di incannulamento e le complicanze meccaniche da venipuntura. La guida ecografica dovrebbe essere utilizzata da personale pienamente addestrato nell'utilizzo di questa tecnica. [60–64]. Categoria 1B



# Linee Guida EPIC 2014

- "... L'uso dell'ecografia può indirettamente ridurre il rischio di infezione facilitando il posizionamento del catetere venoso centrale..."



## Strategies to Prevent Central Line–Associated Bloodstream Infections in Acute Care Hospitals: 2014 Update

Jonas Marschall, MD;<sup>1,2,a</sup> Leonard A. Mermel, DO, ScM;<sup>3,a</sup> Mohamad Fakih, MD, MPH;<sup>4</sup>  
Lynn Hadaway, MEd, RN, BC, CRNI;<sup>5</sup> Alexander Kallen, MD, MPH;<sup>6</sup> Naomi P. O'Grady, MD;<sup>7</sup>  
Ann Marie Pettis, RN, BSN, CIC;<sup>8</sup> Mark E. Rupp, MD;<sup>9</sup> Thomas Sandora, MD, MPH;<sup>10</sup>  
Lisa L. Maragakis, MD, MPH;<sup>11</sup> Deborah S. Yokoe, MD, MPH<sup>12</sup>

5. Use ultrasound guidance for internal jugular catheter insertion (quality of evidence: II).<sup>99</sup>
  - a. Ultrasound-guided internal jugular vein catheterization reduces the risk of CLABSI and of non-infectious complications of CVC placement.<sup>100</sup>

# Consensus 2012

Intensive Care Med  
DOI 10.1007/s00134-012-2597-x

CONFERENCE REPORTS AND EXPERT PANEL

**Massimo Lamperti**  
**Andrew R. Bodenham**  
**Mauro Pittiruti**  
**Michael Blaivas**  
**John G. Augoustides**  
**Mahmoud Elbarbary**  
**Thierry Pirotte**  
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**David Feller-Kopman**  
**Wolfram Schummer**  
**Roberto Biffi**  
**Eric Desruennes**  
**Lawrence A. Melniker**  
**Susan T. Verghese**

**International evidence-based  
recommendations on ultrasound-guided  
vascular access**



**Table 6** Recommendations regarding sterility using ultrasound guidance and prevention of infectious and mechanical complications using ultrasound-guided cannulation

Sterility during ultrasound vascular procedures				
Domain code	Suggested definition	Level of evidence	Degree of consensus	Strength of recommendation
D8.S1	Sterile techniques should always be used during the placement of a vascular access device, including hand washing; sterile full body drapes; wearing of sterile gowns, gloves, caps and masks covering both the mouth and nose. Probe and cable sterility have to be maintained using sterile gel and appropriate probe and cable shields	A	Very good	Strong
Prevention of infectious and mechanical complications with ultrasound-guided cannulation				
D8.S2	Ultrasound guidance should be used in order to decrease the rate of CRBSI in adults and children	C	Very good	Strong
D8.S3–4	A multi-faceted strategy, including the use of ultrasound guidance with specific preventive and educational measures and the promotion of good practices applied by both medical and nursing staff, is suggested in order to reduce the incidence of CRBSI	B	Good	Strong
D8.S5	Ultrasound guidance should be used to avoid cannulation of thrombotic sites	A	Very good	Strong
D8.S6	Ultrasound guidance, by reducing puncture attempts, technical failure rates and mechanical complications, has to be preferred because of a reduced incidence of catheter-related thrombosis	A	Very good	Strong

## 22. VASCULAR VISUALIZATION

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### Standard

22.1 To ensure patient safety, the clinician is competent in the use of vascular visualization technology for vascular access device (VAD) insertion. This knowledge includes, but is not limited to, appropriate vessels, size, depth, location, and potential complications.

22.2 Vascular visualization technology is used in patients with difficult venous access and/or after failed venipuncture attempts.

22.3 Vascular visualization technology is employed to increase the success with peripheral cannulation and decrease the need for central vascular access device (CVAD) insertion, when other factors do not require a CVAD.

# KEYPOINTS

- IMPIANTO ECOGUIDATO
- ANTISETTICI (CLOREXIDINA GLUCONATO 2% IN IPA 70%)
- PACK PROCEDURALI
- SISTEMI DI FISSAGGIO
- PORT PROTECTORS
- SIRINGHE PRERIEMPITE STERILI PER FLUSH E LOCK DEL CATETERE
- LOCK PROPHYLAXIS

Robert A. Weinstein, Section Editor

# Chlorhexidine: Expanding the Armamentarium for Infection Control and Prevention

**Aaron M. Milstone,<sup>1,3</sup> Catherine L. Passaretti,<sup>2,3</sup> and Trish M. Perl<sup>2,3</sup>**

<sup>1</sup>Division of Pediatric Infectious Diseases, Department of Pediatrics, and <sup>2</sup>Division of Infectious Diseases, Department of Medicine, Johns Hopkins University School of Medicine, and <sup>3</sup>Department of Hospital Epidemiology and Infection Control, The Johns Hopkins Hospital, Baltimore, Maryland

---

**Health care–associated infections (HAIs) result in increased patient morbidity and utilization of health care resources. Rates of HAI are increasing despite advances in health care technology. Limited antimicrobial agents and a dry drug pipeline make novel prevention efforts critical. Chlorhexidine, an antiseptic solution that has been used worldwide since the 1950s, is a safe and effective product with broad antiseptic activity. Novel uses of chlorhexidine-containing products are being implemented to promote antisepsis and prevent bacterial colonization and infection. We review some of the many infection control applications of chlorhexidine in the battle against HAI, such as general skin cleansing, skin decolonization, preoperative showering and bathing, vascular catheter site preparation, impregnated catheter site dressings, impregnated catheters, and oral decontamination. As mandatory public reporting and pay for performance force infection control issues to the forefront, chlorhexidine-containing products may provide a vast armamentarium for the control and prevention of HAI.**

**IVAD14** Decontaminate the skin at the insertion site with a **single-use application** of 2% chlorhexidine gluconate in 70% isopropyl alcohol (or povidone iodine in alcohol for patients with sensitivity to chlorhexidine) and allow to dry prior to the insertion of a central venous access device.

*Class A*

**IVAD15** Decontaminate the skin at the insertion site with a single-use application of 2% chlorhexidine gluconate in 70% isopropyl alcohol (or povidone iodine in alcohol for patients with sensitivity to chlorhexidine) and allow to dry before inserting a peripheral vascular access device.

*New recommendation Class D/GPP*



### **epic3: National Evidence-Based Guidelines for Preventing Healthcare-Associated Infections in NHS Hospitals in England**

H.P. Loveday<sup>a\*</sup>, J.A. Wilson<sup>a</sup>, R.J. Pratt<sup>a</sup>, M. Golsorkhi<sup>a</sup>, A. Tingle<sup>a</sup>, A. Bak<sup>a</sup>, J. Browne<sup>a</sup>, J. Prieto<sup>b</sup>, M. Wilcox<sup>c</sup>

<sup>a</sup> Richard Wells Research Centre, College of Nursing, Midwifery and Healthcare, University of West London (London).

<sup>b</sup> Faculty of Health Sciences, University of Southampton (Southampton).

<sup>c</sup> Microbiology and Infection Control, Leeds Teaching Hospitals and University of Leeds (Leeds).



## Preventing Central Line–Associated Bloodstream Infections

### A Global Challenge, A Global Perspective

 The Joint Commission

 Joint Commission  
Resources

 Joint Commission  
International

The following summarizes current recommendations for skin antisepsis prior to CVC insertion and during dressing changes<sup>13,14,18,19,36</sup>:

- Apply antiseptics to clean skin.
- Apply chlorhexidine/alcohol in a concentration greater than 0.5% in alcohol.
- If there is a contraindication to chlorhexidine, apply tincture of iodine, an iodophor, or alcohol as an alternative.
- Allow the antiseptic solution to dry before placing the catheter.

# ***Infusion Therapy Standards of Practice***

Lisa A. Gorski, MS, RN, HHCNS-BC, CRNI<sup>®</sup>, FAAN

Lynn Hadaway, MEd, RN, NPD-BC, CRNI<sup>®</sup>

Mary E. Hagle, PhD, RN-BC, FAAN

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Barb Nickel, APRN-CNS, CCRN, CRNI<sup>®</sup>

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Mary Alexander, MA, RN, CRNI<sup>®</sup>, CAE, FAAN

8TH EDITION

REVISED 2021

## 33. VASCULAR ACCESS SITE PREPARATION AND SKIN ANTISEPSIS

### Standard

33.1 Skin antiseptics are performed prior to VAD placement.

33.2 The intended VAD insertion site is visibly clean prior to application of an antiseptic solution; if visibly soiled, cleanse the intended site with soap and water prior to application of antiseptic solution(s).

### Practice Recommendations





- A. Remove excess hair at the insertion site if needed to facilitate application of VAD dressings; use single-patient-use scissors or disposable-head surgical clippers; do not shave as this may increase the risk for infection.<sup>1,2</sup> (I)
- B. Evaluate patient history of any allergy or sensitivity to skin antiseptics (see Standard 55, *Catheter-Associated Skin Injury*).<sup>3,4</sup> (V)
- C. Perform skin antiseptics using the preferred skin antiseptic agent of alcohol-based chlorhexidine solution.<sup>5-10</sup> (I)
  - 1. If there is a contraindication to chlorhexidine solution, an iodophor (eg, povidone-iodine) or 70% alcohol may also be used.<sup>5,6,10</sup> (IV)
  - 2. Aqueous chlorhexidine may be considered if there is a contraindication to alcohol-based chlorhexidine.<sup>3</sup> (IV)

- 3. For preterm neonates, low-birth-weight infants, and within the first 14 days of life:
  - a. Use povidone-iodine, alcohol-based or aqueous chlorhexidine solution.<sup>4,11-17</sup> (I)
  - b. Use both aqueous and alcohol-based chlorhexidine with caution due to risks of chemical burns to the skin. Systemic absorption has been reported due to skin immaturity; however, systemic effects are not documented. Studies have not established one antiseptic solution as superior for safety or efficacy in neonates.<sup>11-17</sup> (IV)
  - c. Avoid the use of tincture of iodine due to the potential deleterious effect on the neonatal thyroid gland.<sup>18-20</sup> (II)
  - d. Remove antiseptics after the procedure is complete using sterile water or saline.<sup>11,16</sup> (IV)
- D. Use a single-use sterile applicator containing sterile solution, not a multiple use product (eg, bottle of antiseptic solution).<sup>3,5</sup> (IV)
  - 1. Follow manufacturers' directions for use to determine appropriate product application and dry times; always allow product to naturally dry without wiping, fanning, or blowing on skin.<sup>3</sup> (V)



## SHEA/IDSA/APIC Practice Recommendation

# Strategies to prevent central line-associated bloodstream infections in acute-care hospitals: 2022 Update

Niccolò Buetti MD, MSc, PhD<sup>1,2,a</sup> , Jonas Marschall MD, MSc<sup>3,4,a</sup> , Marci Drees MD, MS<sup>5,6</sup> ,  
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Elizabeth Monsees PhD, MBA, RN, CIC<sup>10,11</sup> , Shannon Novosad MD MPH<sup>12</sup>, Naomi P. O'Grady MD<sup>13</sup>,  
Mark E. Rupp MD<sup>14</sup> , Joshua Wolf MBBS, PhD, FRACP<sup>15,16</sup> , Deborah Yokoe MD, MPH<sup>17</sup> and  
Leonard A. Mermel DO, ScM<sup>18,19</sup> 

<sup>1</sup>Infection Control Programme, University of Geneva Hospitals and Faculty of Medicine, Geneva, Switzerland, <sup>2</sup>University of Paris, Paris, France, <sup>3</sup>Department of Infectious Diseases, Bern University Hospital and University of Bern, Bern, Switzerland, <sup>4</sup>Division of Infectious Diseases, Department of Medicine, Washington University School of Medicine, St. Louis, Missouri, United States, <sup>5</sup>ChristianaCare, Wilmington, Delaware, United States, <sup>6</sup>Sidney Kimmel Medical College at Thomas Jefferson University, Philadelphia, Pennsylvania, United States, <sup>7</sup>Ascension Healthcare and Wayne State University School of Medicine, Detroit, Michigan, United States, <sup>8</sup>Lynn Hadaway Associates, Milner, Georgia, United States, <sup>9</sup>Johns Hopkins University School of Medicine, Baltimore, Maryland, United States, <sup>10</sup>Children's Mercy Hospital, Kansas City, Missouri, United States, <sup>11</sup>University of Missouri–Kansas City School of Medicine, Kansas City, Missouri, United States, <sup>12</sup>Division of Healthcare Quality Promotion, Centers for Disease Control and Prevention, Atlanta, Georgia, United States, <sup>13</sup>National Institutes of Health, Bethesda, Maryland, United States, <sup>14</sup>University of Nebraska Medical Center, Omaha, Nebraska, United States, <sup>15</sup>Department of Infectious Diseases, St. Jude Children's Research Hospital, Memphis, Tennessee, United States, <sup>16</sup>Department of Pediatrics, University of Tennessee Health Science Center, Memphis, Tennessee, United States, <sup>17</sup>University of California–San Francisco, San Francisco, California, United States, <sup>18</sup>Warren Alpert Medical School of Brown University, Providence, Rhode Island, United States and <sup>19</sup>Rhode Island Hospital, Providence, Rhode Island, United States

7. **Use an alcoholic chlorhexidine antiseptic for skin preparation (Quality of Evidence: HIGH)<sup>42,129–134</sup>**
  - a. Before catheter insertion, apply an alcoholic chlorhexidine solution containing at least 2% chlorhexidine gluconate to the insertion site.
    - i. The antiseptic solution must be allowed to dry before making the skin puncture.
    - ii. Alcoholic chlorhexidine for skin antisepsis to prevent CLABSI in NICU patients should be used when the benefits are judged to outweigh potential risk.



**ChloraPrep®**

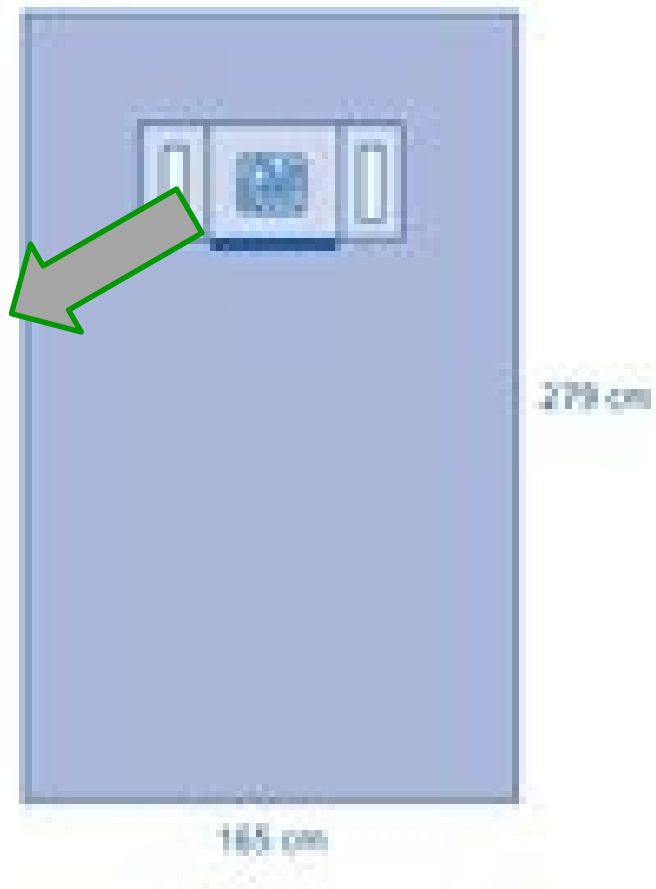


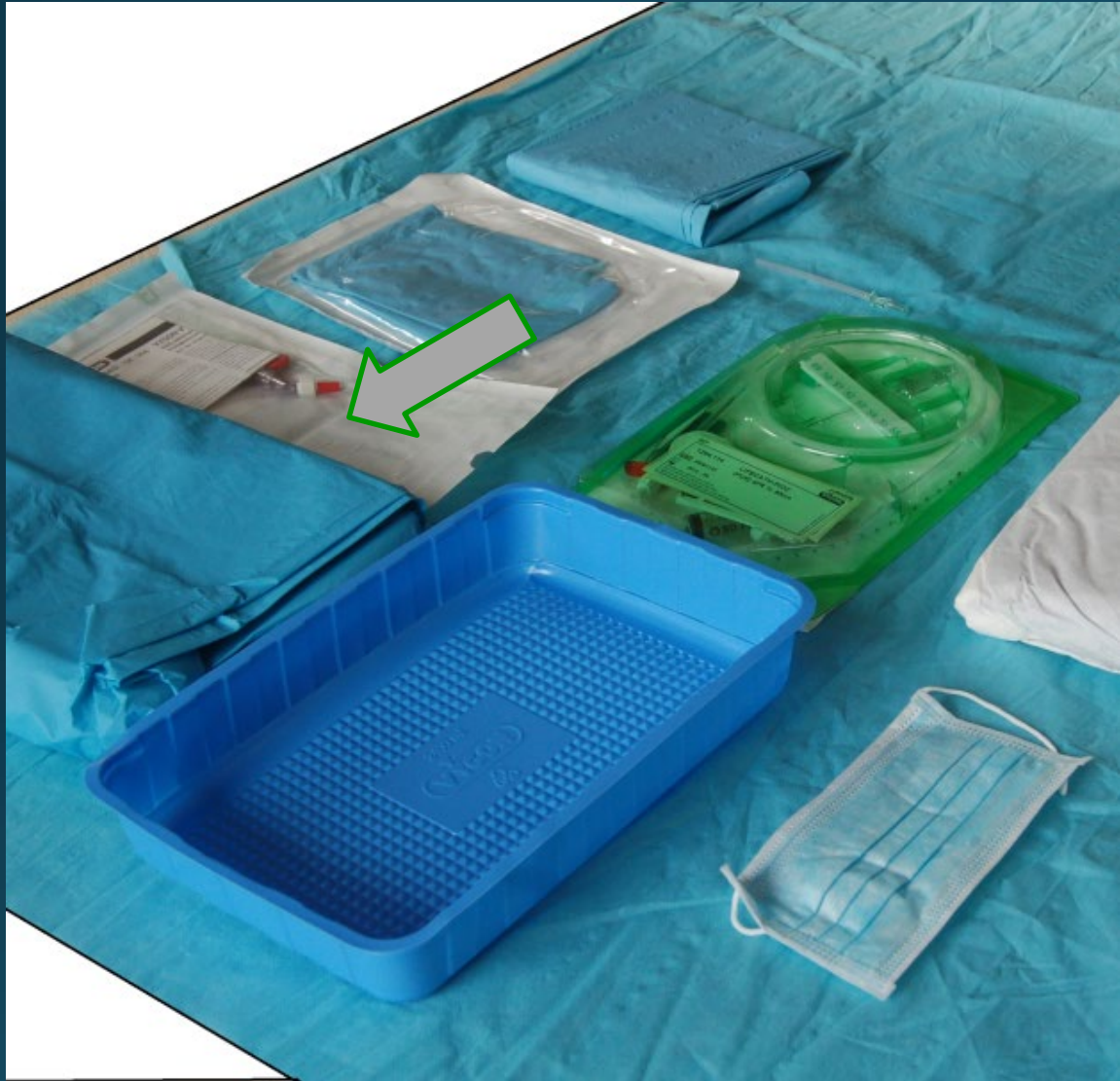
# KEYPOINTS

- IMPIANTO ECOGUIDATO
- ANTISETTICI (CLOREXIDINA GLUCONATO 2% IN IPA 70%)
- **PACK PROCEDURALI**
- SISTEMI DI FISSAGGIO
- PORT PROTECTORS
- SIRINGHE PRERIEMPITE STERILI PER FLUSH E LOCK DEL CATETERE
- LOCK PROPHYLAXIS









# The ErgoPack<sup>®</sup> System:

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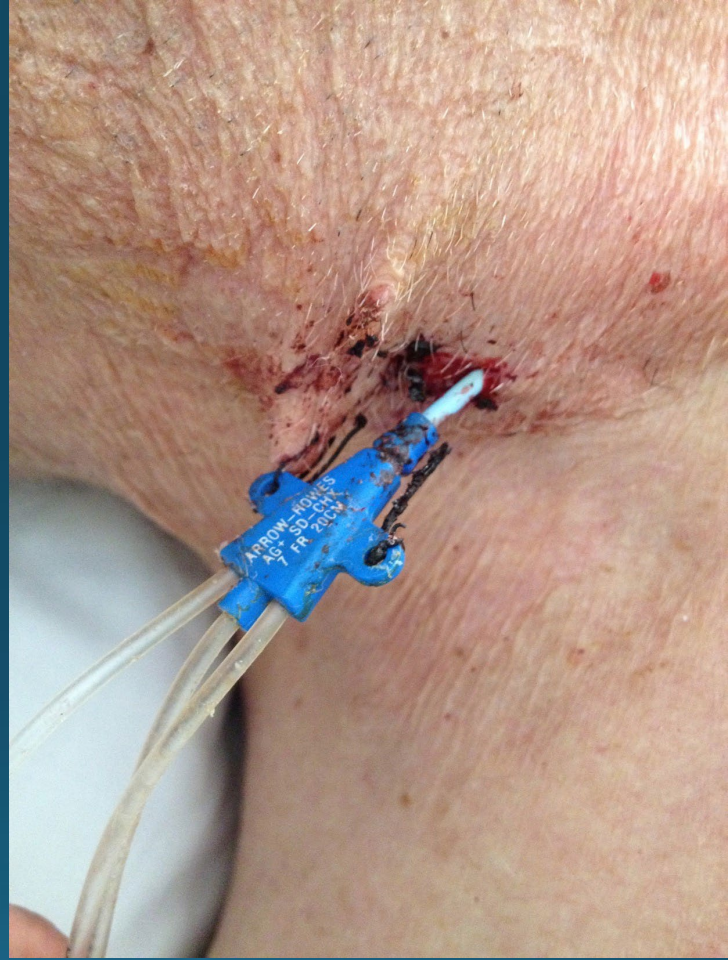


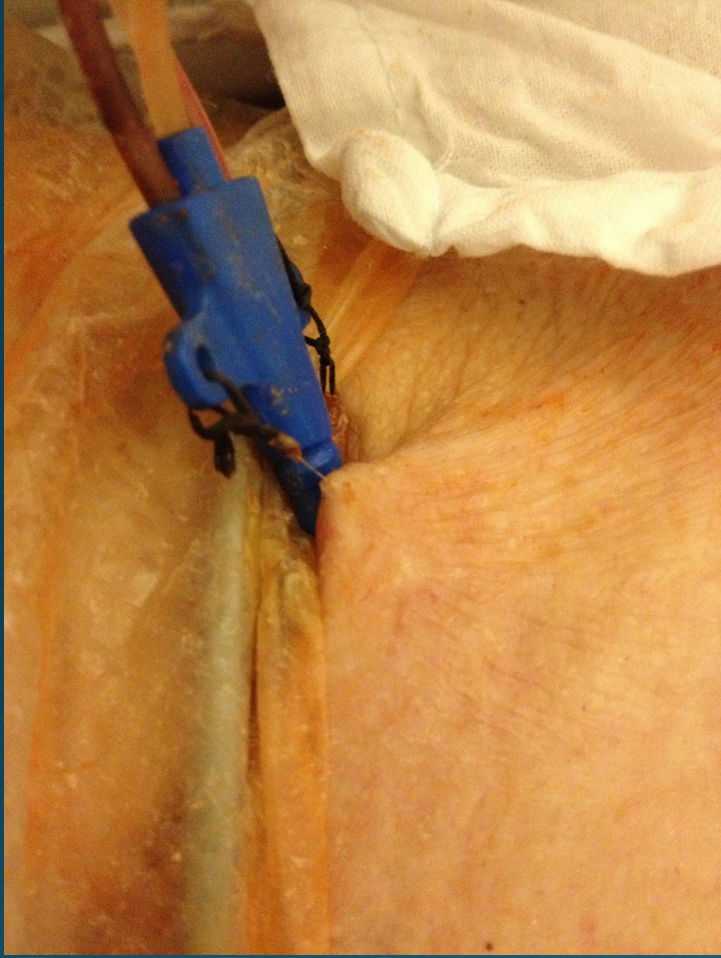
# KEYPOINTS

- IMPIANTO ECOGUIDATO
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- **SISTEMI DI FISSAGGIO**
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- LOCK PROPHYLAXIS

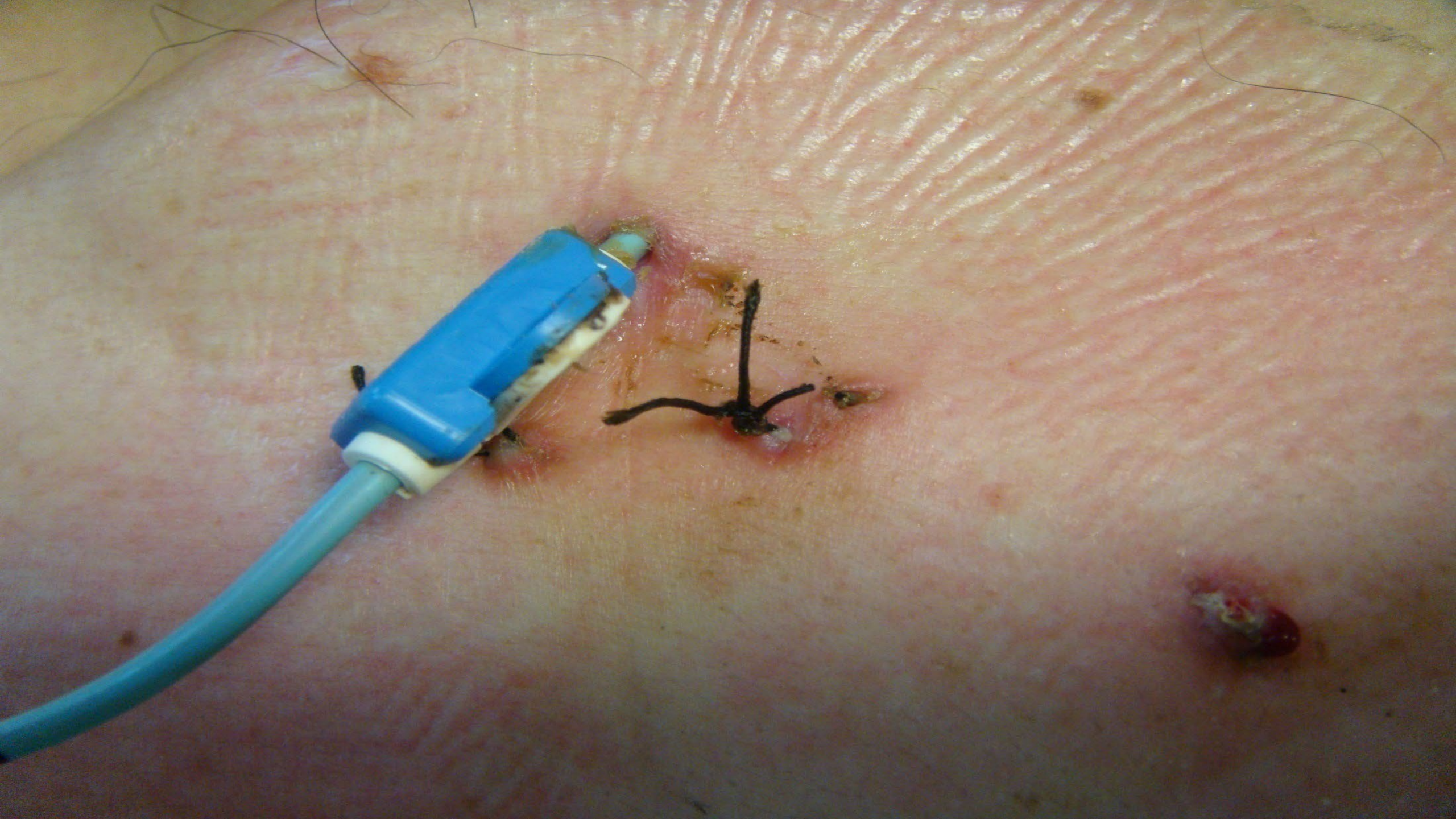
# THE RISK OF SUTURE...

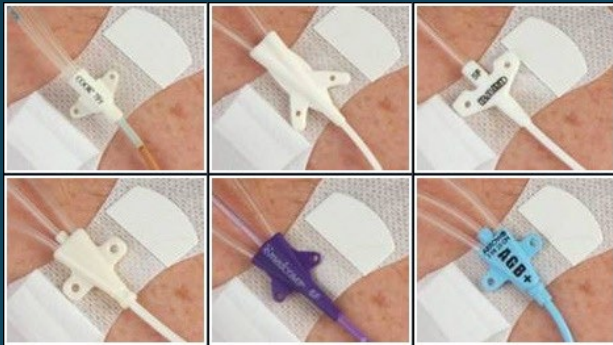










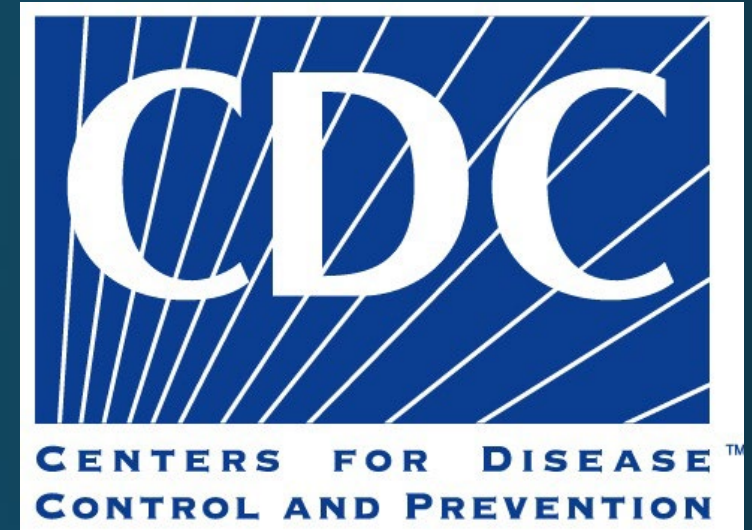




## Dispositivi per il Fissaggio del Catetere

### Raccomandazioni

Usare un dispositivo di fissaggio *sutureless* al fine di ridurre il rischio di infezione per i cateteri intravascolari [105]. Categoria II



## 547 Catheter Securement Devices

### 548 Recommendation

549 Use a sutureless securement device to reduce the risk of infection for PICCs [163].

550 Category II



Avoid use of tape or sutures, as they are not effective alternatives to an ESD. Rolls of nonsterile tape can become contaminated with pathogenic bacteria, although its contribution to VAD infection has not been quantified. Sutures are associated with needle-stick injury, in addition to supporting the growth of biofilm and increasing the risk of catheter-related bloodstream infection.<sup>7-10</sup> (II, Regulatory)

F. Subcutaneous ESDs have been successful in stabilizing PICCs and CVADs inserted through the internal jugular vein of adults. Patient outcomes and patient and inserter satisfaction have been favorable; however, additional studies with other CVADs are needed.<sup>21-23</sup> (V)

INS 2024

# Clinical experience of a subcutaneously anchored sutureless system for securing central venous catheters

Mauro Pittiruti, Giancarlo Scoppettuolo, Laura Dolcetti, Davide Celentano, Alessandro Emoli, Bruno Marche and Andrea Musarò

## ABSTRACT

This article reports the results of three prospective clinical studies conducted in a university hospital regarding the efficacy, safety and cost effectiveness of a subcutaneously anchored sutureless system for securing central venous catheters. The results were favourable to the adoption of such a device, and the analysis of the data allowed the authors to define those categories of patients where the device should have the most benefit: neonates, children, non-compliant older patients with cognitive difficulties, patients with skin abnormalities that may reduce the effectiveness of a skin-adhesive sutureless securement system, patients who are candidates for having a peripherally inserted central catheter (PICC) in place for more than 8 weeks, and any other category of patients with a recognised high risk of catheter dislodgement.

**Key words:** Sutureless securement ■ Central venous catheters  
■ Peripherally inserted central catheter ■ Subcutaneously anchored securement ■ Stabilisation device

the use of such a device for the purpose of reducing or eliminating the clinical occurrence of catheter dislodgement, an often-neglected complication. In the authors' experience, CVC dislodgement is only partially prevented by current securement strategies: adoption of semipermeable transparent dressings (Tegaderm, 3M; IV 3000, Smith & Nephew); application of cyanoacrylate glue (Hystoacryl, BBraun) on the exit site; consistent use of sutureless devices, either adhesive to the skin (StatLock, BD; Grip-Lok, Zefon International; WingGuard, Centurion Medical Products) or integrated in the transparent dressing (SorbaShield, Centurion). In some patient populations within the authors' hospital (such as in children), partial or total dislodgement of the catheter is the most frequent reason for losing venous access (far more frequent than infection or venous thrombosis or lumen occlusion).

Before introducing this new device into clinical practice, the authors have considered several aspects: its applicability, its

# GAVeCeLT-WoCoVA Consensus on subcutaneously anchored securement devices for the securement of venous catheters: Current evidence and recommendations for future research

The Journal of Vascular Access  
2021, Vol. 22(5) 716–725  
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DOI: 10.1177/1129729820924568  
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Fulvio Pinelli<sup>1</sup>, Mauro Pittiruti<sup>2</sup>, Ton Van Boxtel<sup>3</sup>, Giovanni Barone<sup>4</sup>, Roberto Biffi<sup>5</sup>, Giuseppe Capozzoli<sup>6</sup>, Alessandro Crocoli<sup>7</sup>, Stefano Elli<sup>8</sup>, Daniele Elisei<sup>9</sup>, Adam Fabiani<sup>10</sup>, Cristina Garrino<sup>11</sup>, Ugo Graziano<sup>12</sup>, Luca Montagnani<sup>13</sup>, Alessio Pini Prato<sup>14</sup>, Giancarlo Scoppettuolo<sup>15</sup>, Nicola Zadra<sup>16</sup>, Clelia Zanaboni<sup>17</sup>, Pietro Zerla<sup>18</sup>, Evangelos Konstantinou<sup>19</sup>, Matt Jones<sup>20</sup>, Hervé Rosay<sup>21</sup>, Liz Simcock<sup>22</sup>, Marguerite Stas<sup>23</sup> and Gilda Pepe<sup>15</sup>

## Abstract

**Background:** Subcutaneously anchored securement devices (or subcutaneous engineered securement devices) have been introduced recently into the clinical practice, but the number of published studies is still scarce. The Italian Group of Long-Term Central Venous Access Devices (GAVeCeLT)—in collaboration with WoCoVA (World Congress on Vascular Access)—has developed a Consensus about the effectiveness, safety, and cost-effectiveness of such devices.

**Methods:** After the definition of a panel of experts, a systematic collection and review of the literature on subcutaneously anchored securement devices was performed. The panel has been divided in two working groups, one focusing on adult patients and the other on children and neonates.

**Results:** Although the quality of evidence is generally poor, since it is based mainly on non-controlled prospective studies, the panel has concluded that subcutaneously anchored securement devices are overall effective in reducing the risk of dislodgment and they appear to be safe in all categories of patients, being associated only with rare and negligible local adverse effects; cost-effectiveness is demonstrated—or highly likely—in specific populations of patients with long-term venous access and/or at high risk of dislodgment.

**Conclusion:** Subcutaneously anchored securement is a very promising strategy for avoiding dislodgment. Further studies are warranted, in particular for the purpose of defining (a) the best management of the anchoring device so to avoid local

# KEYPOINTS

- IMPIANTO ECOGUIDATO
- ANTISETTICI (CLOREXIDINA GLUCONATO 2% IN IPA 70%)
- PACK PROCEDURALI
- SISTEMI DI FISSAGGIO
- **PORT PROTECTORS**
- SIRINGHE PRERIEMPITE STERILI PER FLUSH E LOCK DEL CATETERE
- LOCK PROPHYLAXIS

DISINFEZIONE DEI PUNTI DI ACCESSO (HUB O NEEDLEFREE CONNECTORS)  
DI UN CVC MEDIANTE SCRUBBING CON SOLUZIONI ALCOLICHE  
(PREFERIBILMENTE CLOREXIDINA 2% IN SOLUZIONE ALCOLICA) OPPURE DISINFEZIONE  
PASSIVA  
DEI NFC MEDIANTE PORT PROTECTORS.



F. Perform a vigorous mechanical scrub for manual disinfection of the needleless connector prior to each VAD access and allow it to dry.

1. Acceptable disinfecting agents include 70% isopropyl alcohol, iodophors (ie, povidone-iodine), or >0.5% chlorhexidine in alcohol solution.<sup>7,16</sup>

(II)

G. Use of passive disinfection caps containing disinfecting agents (eg, isopropyl alcohol) has been shown to reduce intraluminal microbial contamination and reduce the rates of central line-associated bloodstream infection (CLABSI). Use of disinfection caps on peripheral catheters has limited evidence but should be considered.

RESEARCH ARTICLE

Open Access

# Educational interventions alone and combined with port protector reduce the rate of central venous catheter infection and colonization in respiratory semi-intensive care unit



Riccardo Inchingolo<sup>1\*</sup> , Giuliana Pasciuto<sup>1</sup>, Daniele Magnini<sup>1</sup>, Manuela Cavalletti<sup>1</sup>, Giancarlo Scoppettuolo<sup>2</sup>, Giuliano Montemurro<sup>1</sup>, Andrea Smargiassi<sup>1</sup>, Riccardo Torelli<sup>3</sup>, Maurizio Sanguinetti<sup>3,4</sup>, Teresa Spanu<sup>3,4</sup>, Giuseppe Maria Corbo<sup>1,5</sup> and Luca Richeldi<sup>1,5</sup>



## Strategies to Prevent Central Line–Associated Bloodstream Infections in Acute Care Hospitals: 2014 Update

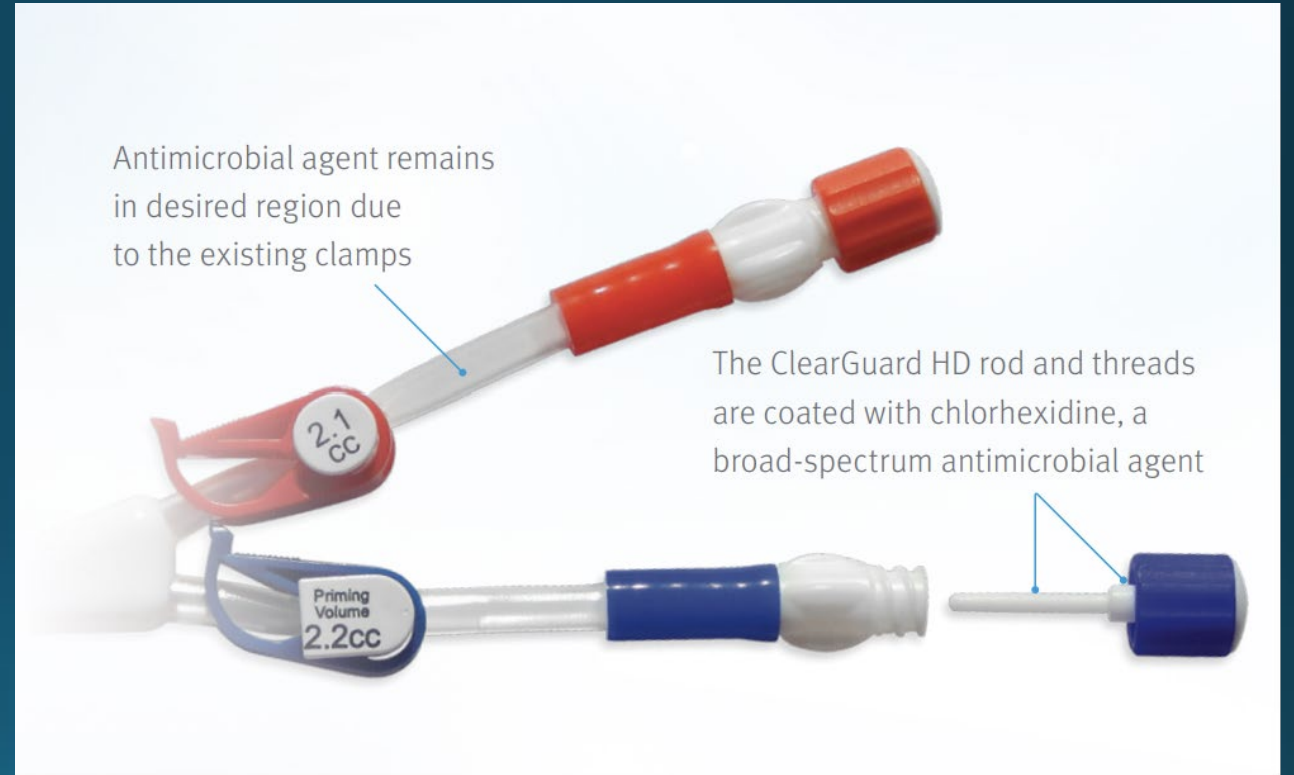
Jonas Marschall, MD;<sup>1,2,a</sup> Leonard A. Mermel, DO, ScM;<sup>3,a</sup> Mohamad Fakih, MD, MPH;<sup>4</sup>  
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Lisa L. Maragakis, MD, MPH;<sup>11</sup> Deborah S. Yokoe, MD, MPH<sup>12</sup>

3. Use an antiseptic-containing hub/connector cap/port protector to cover connectors (quality of evidence: I).<sup>161-165</sup>

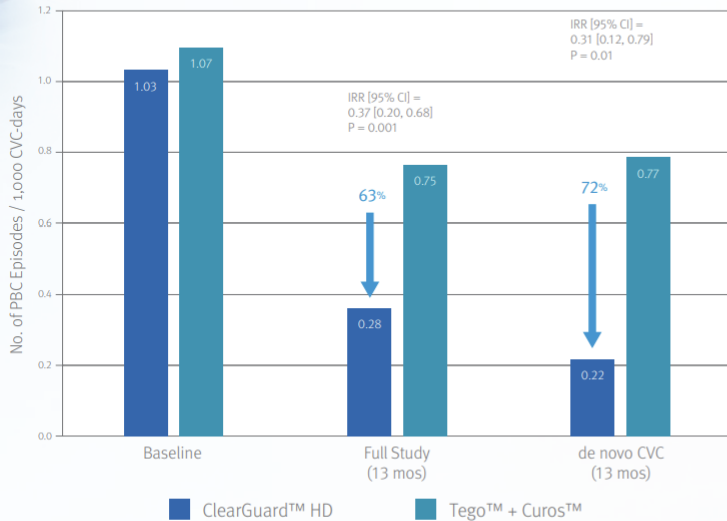
# INS 2021 : port protectors

G. Use of passive disinfection caps containing disinfecting agents (eg, isopropyl alcohol) has been shown to reduce intraluminal microbial contamination and reduce the rates of central line-associated bloodstream infection (CLABSI). Use of disinfection caps on peripheral catheters has limited evidence but should be considered.

# HD ANTIMICROBIAL BARRIER CAPS



### Cluster-Randomized Trial of Devices to Prevent Catheter-Related Bloodstream Infections



### ClearGuard HD Caps vs. Tego™ + Curoso™

Brunelli, SM et al. Cluster-randomized trial of devices to prevent catheter-related bloodstream infection. J Am Soc Nephrol 2018 Apr; 29(4):1336-1343.

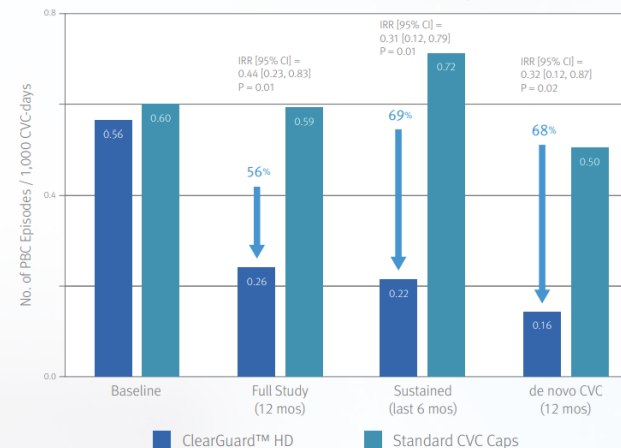
- › 13-month prospective, cluster-randomized multicenter open-label trial
- › 1,671 patients (826 treatment, 845 control) accruing ~183,000 CVC days
- › 40 centers across the US
- › Primary endpoint was PBC rate as an indicator of BSI rate

**Results:** Use of the ClearGuard HD caps for 13 months was associated with a 63% lower BSI rate vs. use of Tego + Curoso.



Recommended in the UK's NICE National Guidance for hemodialysis catheter-related bloodstream infections: 2021\*\*  
<https://www.nice.org.uk/guidance/mtg62/>

### Dialysis Catheter-Related Bloodstream Infections: A Cluster-Randomized Trial of the ClearGuard HD Antimicrobial Barrier Cap



### ClearGuard HD Caps vs. Standard Dialysis Caps

Hymes, JL et al. Dialysis catheter-related bloodstream infections: A cluster-randomized trial of the ClearGuard HD antimicrobial barrier cap. Am J Kidney Dis. 2017; 69(2):220-227.

- › 12-month prospective, cluster-randomized, multicenter, open-label comparative effectiveness trial in hemodialysis patients with central venous catheters
- › 2,470 patients (1,245 treatment, 1,225 control) accruing ~350,000 CVC days
- › 40 centers across the US
- › Primary endpoint was PBC rate as an indicator of BSI rate

**Results:** Use of the ClearGuard HD caps for 12 months was associated with a 56% lower BSI rate vs. use of standard caps. When considering sustained use (defined as 6 months of the study), the intervention vs. control was associated with a 69% lower BSI rate.

**National Kidney Foundation** **KDOQI** 2019 Update

ClearGuard HD caps recommended in NKF's KDOQI Clinical Practice Guideline for Vascular Access: 2019<sup>6</sup>

21.3 KDOQI considers it reasonable to use an antimicrobial barrier cap to help reduce CRBSI in high-risk patients or facilities; the choice of connector should be based on clinician's discretion and best clinical judgment. (Expert Opinion)

# KEYPOINTS

- IMPIANTO ECOGUIDATO
- ANTISETTICI (CLOREXIDINA GLUCONATO 2% IN IPA 70%)
- PACK PROCEDURALI
- SISTEMI DI FISSAGGIO
- PORT PROTECTORS
- SIRINGHE PRERIEMPITE STERILI PER FLUSH E LOCK DEL CATETERE
- LOCK PROPHYLAXIS

- A. Use single-dose systems (eg, single-dose vials or pre-filled labeled syringes) for all VAD flushing and locking.
1. Commercially available prefilled syringes may reduce the risk of CR-BSI and save staff time for syringe preparation.<sup>1-3</sup> (IV)
  2. If multiple-dose vials must be used, dedicate a vial to a single patient (see Standard 49, *Infection*).<sup>4</sup> (V)
  3. Do not use intravenous (IV) solution containers (eg, bags or bottles) as a source for obtaining flush solutions.<sup>3-6</sup> (IV)

The logo features a green curved line above the text "Infusion Therapy Standards of Practice" in a bold, white, sans-serif font. The background is a light blue grid pattern.

## Infusion Therapy Standards of Practice

UTILIZZO DI SIRINGHE PRERIEMPITE STERILI PER IL FLUSH  
E IL LOCK DEI CVC

# SwabFlush™



## SwabFlush

Offers clinicians the convenience of having SwabCap **there when they need it**, after the final flush!

### When to use SwabFlush

- When a saline flush/lock is required to finish a patient's IV therapy
- Use SwabFlush to administer the final saline:
  - After catheter insertion
  - After medication delivery
  - After blood is withdrawn or delivered through the catheter



### SwabCap

NON-VENTING  
DISINFECTION CAP  
CONTAINS 70% IPA

Single Use Only  
Sterile Packaging  
Luer Lock Design

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# IDEAL LOCK SOLUTION

- Spectrum of activity should include common or targeted pathogens
- Ability to penetrate or disrupt a biofilm
- Compatibility with anticoagulants
- Prolonged stability
- Low risk of toxicity and adverse events
- Low potential for resistance
- Cost effectiveness

# SOSTANZE AD ATTIVITA' ANTIMICROBICA (NON ANTIBIOTICI)

- ETANOLO
- **TAUROLIDINA (anche coniugata con eparina o citrato 4% oppure urokinasi + citrato 4%)**
- CITRATO
- EDTA



*Grazie per l'attenzione!*

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