

Catheter securement in 2022

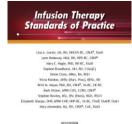
Subcutaneous anchorage of PICCs

Fabrizio Brescia



## OBJECTIVES OF ADEQUATE STABILIZATION

Ensure the integrity of the device



RTM EDITION
REVISED 2021

INS
STATES A SELECT SOCIETY
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Cox Editoriol Does, Mol 000
Cox Editoriol Doe

Minimize the movement of the catheter at the exit site

Prevent dislocation of the catheter



## OBJECTIVES OF ADEQUATE STABILIZATION

No interference with the evaluation and control of the exit site

No impact on blood flow or therapy infusion

Infusion Therapy
Standards of Practice

The A Great, 19, 10, 10016 St., CHP\* SAM!
Lym Habers, 162, 10, 1100 St., CHP\*
Bryt Stepp, 16, 100, 100, 100, 100
Step Comp. 100, 100, 100, 100, 100
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An integral part of the clinicaltherapeutic path of patients



## INSERTION BUNDLE

Global use of US

US-guided venipuncture + micro-introducer kit

MINIMIZE THE RISK OF COMPLICATIONS

Appropriate catheter/vein ratio

Appropriate aseptic techinique

Optimal stabilization of the device

Tip location Tip navigation

#### DISLOCATION - DEFINITION AND INCIDENCE

## Loss of the device or otherwise loss of function and central position of the catheter tip

Support Care Cancer DOI 10.1007/s00520-012-1554-0

ORIGINAL ARTICLE

Peripherally inserted central catheters (PICCs) in the management of oncohematological patients submitted to autologous stem cell transplantation

Silvia Bellesi · Patrizia Chiusolo · Gennaro De Pascale · Mauro Pittiruti · Giancarlo Scoppettuolo · Elisabetta Metafuni - Sabrina Giammarco -Federica Sorà · Luca Laurenti · Giuseppe Leone ·

Received: 29 March 2012 / Accepted: 23 July 2012 © Springer-Verlag 2012





Elsevier Masson France EM consulte www.em-consulte.com



Médecine et maladies infectiesses, 43 (2013) 350, 355

Original article

Prospective follow-up of complications related to peripherally inserted central catheters 年, 章章

Suivi prospectif des complications associées aux cathéters veineux centraux insérés par voie périphérique

C. Leroyer a.\*, A. Lashéras a, V. Marie a, Y. Le Bras b, T. Carteret c, M. Dupon d, A.-M. Rogues a

- \* Service d'hygiène hospitalière, groupe hospitalier Pellegrin, CHU de Bordeaux, bâtiment POR 1et étage, place Amélie-Raba-Léon, 33076 Bordeaux cedex, France \* Service d'imagerie diagnostique et interventionnelle, groupe hospitalier Pellegrin, CHU de Bordeaux, pluce Amélie-Raba-Léon, 33076 Bordeaux cedex, France
- Service d'imagerie diagnostique et interventionnelle, groupe hospitalier Saint-André, CHU de Bordeaux, place Amélie-Raba-Léon, 33076 Bordeaux cedex, France

<sup>4</sup> Service des maladies infecticuses et médecine tropicale, groupe hospitalier Pellegrin, CHU de Bonteaux, place Amélie-Raba-Léon, 33076 Bonteaux cedex, France Received 5 October 2012: received in revised form 30 April 2013: accepted 18 June 2013

5-15%

**CLINICAL STUDY** 

Peripherally Inserted Central Catheters: Use at a **Tertiary Care Pediatric Center** 

Craig Gibson, MBBS, Bairbre L. Connolly, MD, FRCPC, Rahim Moineddin, PhD, Sanjay Mahant, MD, Doina Filipescu, BS, and Joao G. Amaral, MD



I Vasc Access 2017; 18 (5): 408-414 DOI: 10.5301/jva.5000738

ORIGINAL RESEARCH ARTICLE

Impact of arm selection on the incidence of PICC complications: results of a randomized controlled trial

France Paquet1,2, Louis-Martin Boucher1,2, David Valenti1,2, Richard Lindsay3

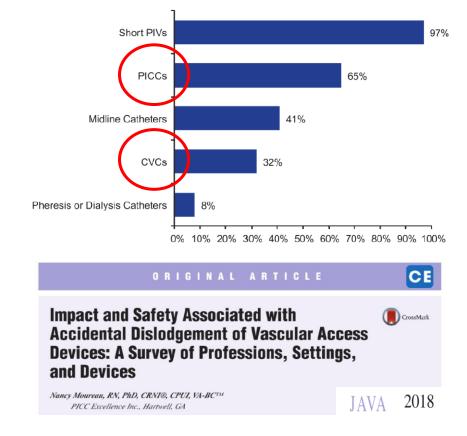
<sup>1</sup>McGill University Health Centre, Montreal, Quebec - Canada

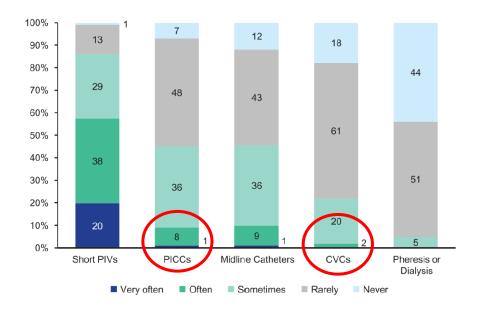
<sup>2</sup>McGill University, Montreal, Quebec - Canada

<sup>5</sup> Belfast Health and Social Care Trust, Knockbracken Healthcare, Belfast - UK



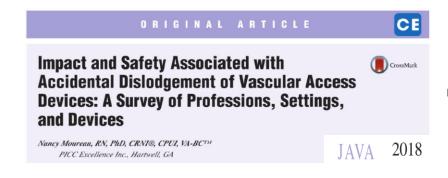
#### **DISLOCATION - INCIDENCE**

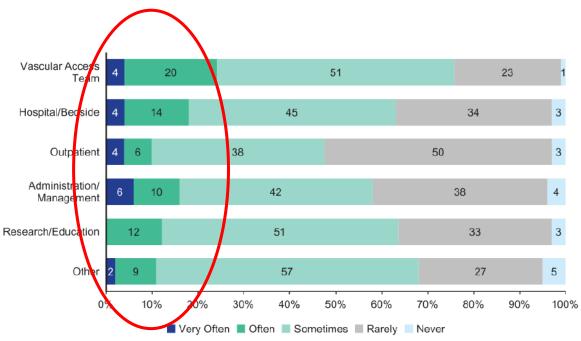






#### **DISLOCATION**

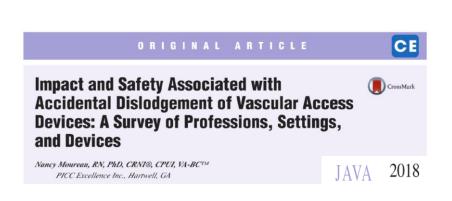


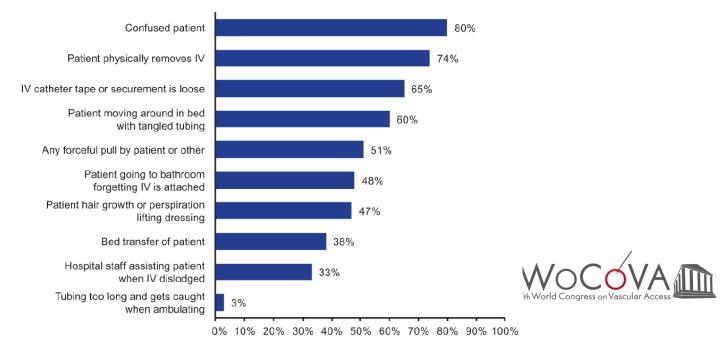




#### **DISLOCATION - CAUSES**

- factors dependent on the patient's cognitive status
- factors related to the actual efficacy/inefficacy of the securement
- factors related to the active or passive mobilization of the patient





#### DISLOCATION - CLINICAL IMPACT

Interruption of therapy

Loss of vascular access

Need for repositioning of the VAD

Discomfort for the patient

Additional costs



### VAD STABILIZATION SYSTEMS

Adhesive Engineered Securement Devices (AESDs) Engineered Securement Devices (ESDs)

Adhesive Sutereless Devices Stabilization systems integrated into the dressing





#### Sutureless Securement Device Reduces Complications of Peripherally Inserted Central Venous Catheters

Alvin J. Yamamoto, MD, Jeffrey A. Solomon, MD, Michael C. Soulen, MD, James Tang, MD, Kim Parkinson, RN, Richard Lin, MD, and Gregory J. Schears, MD

J Vasc Interv Radiol 2002; 13:77-81

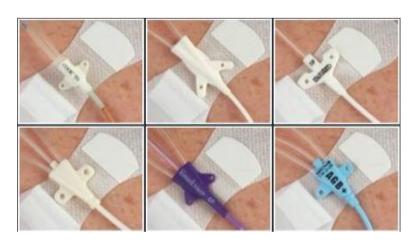
### Never use sutures



- High infectious risk
- High dislocation rates
- Major safety concern for operators needle-stick injury

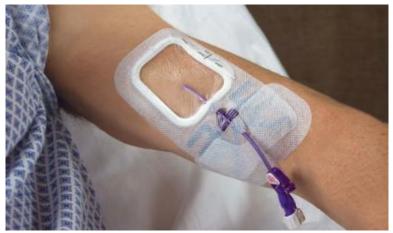


### ADHESIVE SUTURELESS DEVICES





Overcome the poor performance of the sutures...







#### ADHESIVE SUTURELESS DEVICES

- Weekly change of the system
- Mobilization of the device from reduced adhesiveness
- Strong adhesives may require the use of specific solvents to remove residues
- Movement "in and out" at the exit-site
- Risk of partial dislocation during dressing and AESD changes
- Local skin irritation during changes, repositioning in a different area: dislocation



#### Results

In the 2013 period, 1111 PICCs were placed. During this period, 66 (5.94%) PICCs had migration or dislodgment issues that required replacement of the catheter. Data collection did not include small migratory incidences that did not require device replacement. Additional costs associated with PICC migration such as confirmatory radiography and complication interventions for occlusion management were not recorded. PICC migration is often underreported and was difficult to capture through standard documentation methods despite observation from the IST.



#### ADHESIVE SUTURELESS DEVICES

MARSI has been defined as "an occurrence in which erythema and/or other manifestation of cutaneous abnormality (including, but not limited to, vesicle, bulla, erosion, or tear) persists 30 minutes or more after removal of the adhesive"

Medical Adhesive Related Skin Injury

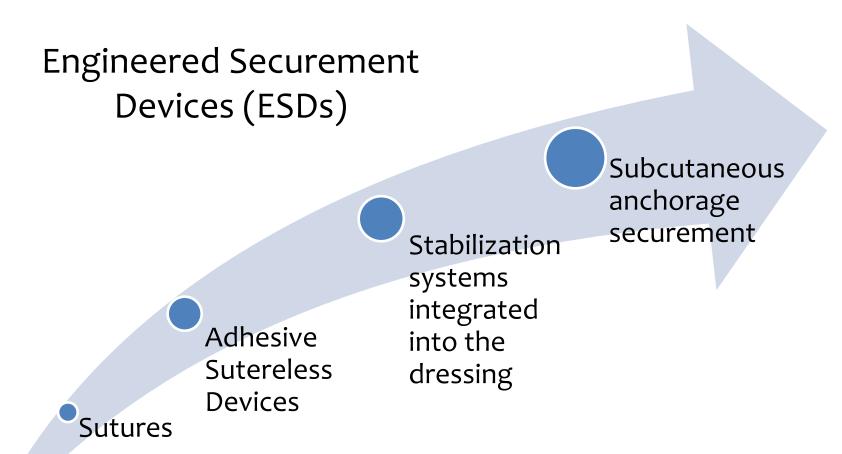
**MARSI** 





#### VAD STABILIZATION SYSTEMS

Subcutaneous Engineered Securement Device (SESD)

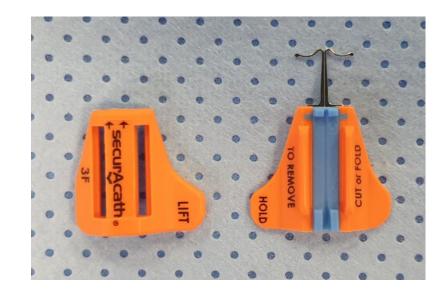




#### SUBCUTANEOUS ANCHORAGE SECUREMENT

SAS devices allows a stabilization of the catheter through the use of nitinol bars anchored in the subcutaneous tissue

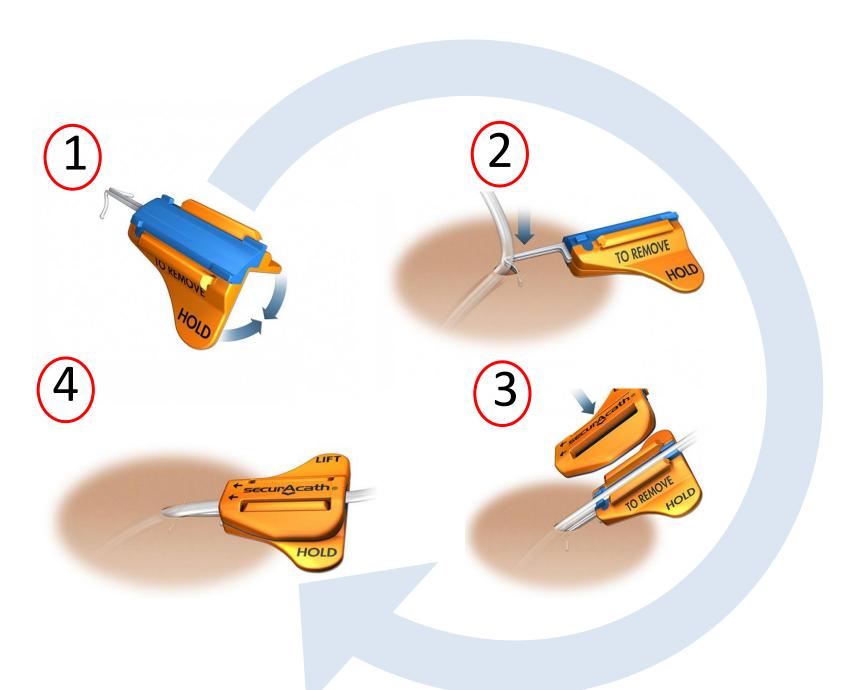




One device

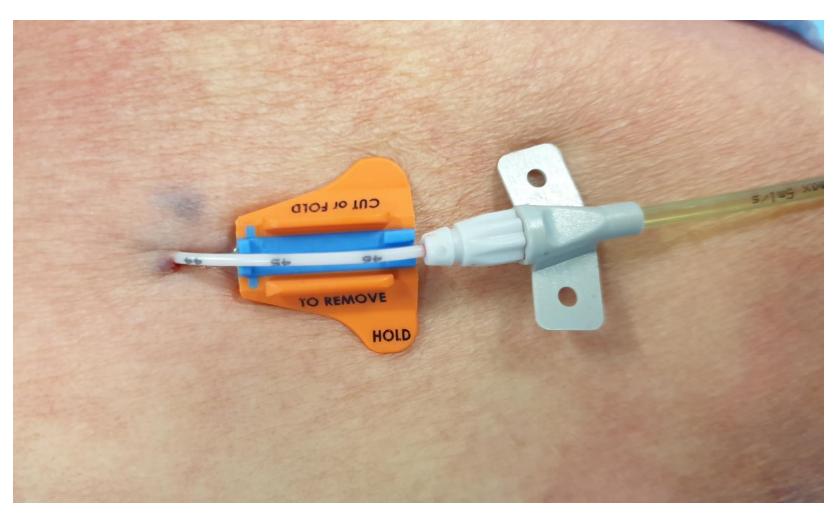
... from placement to removal





## SUBCUTANEOUS ANCHORAGE SECUREMENT

## SUBCUTANEOUS ANCHORAGE SECUREMENT





#### SAS - ADVANTAGES

- not require periodic replacement
- complete disinfection of the exit site
- "in and out" micromovements of the catheter at the exit site are eliminated
- efficacy is not affected by characteristics of the skin





Study	Design	Population/ no. of patients	Catheter type	Primary outcome	Results	Dislodgment N (%)	Success rate	Adverse events	Cost analysis
and	Multicentre Prospective Observational	Adult/74	CICC 7 Fr	Successful securement	72(97%)	2(2.7%)	100%	None	No
Egan et al. <sup>2</sup>	Multicentre Prospective Observational	Adult/68	PICC 5 Fr	Successful securement	62(91.2%)	None	100%	6(8.8%)	No
Hughes <sup>4</sup>	Observational	Adult/3 I	PICC	Successful securement	30(96.7%)	One moved out of I cm (3.3%)	100%	25% difficult removal; 22.5% other	Yes
Dolcino et al. <sup>6</sup>	Prospective Observational retrospectively controlled	Pediatric/ 51	Tunneled- cuffed CICC	Incidence of dislodgment within first 30 days	2(1.1%)	2(1.1%)	NR	NR	No
Zerla et al. <sup>5</sup>	Prospective Observational	Adult/30	PICC 4 Fr	Incidence of dislodgment	None	None	NR	NR	Yes
Goossens et al. <sup>7</sup>	RCT (StatLock™ vs Securacath™)	Adult/105	PICC 4-5 Fr	Nursing time for dressing change	7.3 vs 4.3 min (p < 0.0001)				No
Pittiruti et al. <sup>8</sup>	Prospective Observational	Neonatal; Pediatric; Adult/190	PICC; CICC; FICC Tunneled/ untunneled	Successful securement	187(98.4%)	3(1.6%)	99%	5(2.6%) Local inflammation	Yes

SAS: subcutaneously anchored securement; NR: not reported; CICC: centrally inserted central catheters; PICC: peripherally inserted central catheters; FICC: femorally inserted central catheters.

Original research article

JVA The Journal of Vascular Access

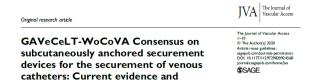
GAVeCeLT-WoCoVA Consensus on subcutaneously anchored securement devices for the securement of venous catheters: Current evidence and recommendations for future research The Journal of Vascular Access I-10

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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 Konstatinou<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>





recommendations for future research

Attention to...

adequate choice of the size of the SAS



Original research article

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

recommendations for future research

Attention to...

- adequate choice of the size of the SAS
- subcutaneous placement





Original research article

GAVECELT-WoCoVA Consensus on subcutaneously anchored securement devices for the securement of venous

catheters: Current evidence and recommendations for future research

Attention to...

- adequate choice of the size of the SAS
- subcutaneous placement

(S)SAGE

adequate training for positioning

Original research article

Intravascular catheter migration: A cross-sectional and health-economic comparison of adhesive and subcutaneous engineered stabilisation devices for intravascular device securement

Dympna McParlan, L Edgar, M Gault, S Gillespie, R Menelly and M Reid

#### Method

JVA The Journal of Vascular Access

A before and after analysis was performed comparing all PICCs placed in 2013 to the post-device implementation period, beginning in 2015, providing 2 years of data. The subcutaneous ESD was introduced to staff during June 2014, allowing the IST to provide 6 months of dedicated education before device implementation and data collection started. During 2014, the currently used AESD was maintained for device securement. All patients were monitored for catheter migration and dislodgement using the PICC History Sheet (See Supplemental Material) and results were kept in an electronic patient database.

#### Results

In the 2013 period, 1111 PICCs were placed. During this period, 66 (5.94%) PICCs had migration or dislodgment issues that required replacement of the catheter. Data collection did not include small migratory incidences that did not require device replacement. Additional costs associated with PICC migration such as confirmatory radiography and complication interventions for occlusion management were not recorded. PICC migration is often underreported and was difficult to capture through standard documentation methods despite observation from the IST.

By the end of the subcutaneous device implementation period (January–December 2015), 1139 PICCs were successfully inserted with zero (0.0%) catheter migrations requiring replacement reported.

1139 patients: no dislocation



#### SAS - TUNNELING

 Table I. RAVESTO—Rapid Assessment of Vascular Exit Site and Tunneling Options.

Central venous access device	Type and path of tunnel	Indications for tunneling		
PICC	Tunnel to Dawson's green area	Puncture site in Dawson's yellow area; non-hospitalized patients with expected long intravenous treatment		
CICC (supraclavicular puncture)	Tunnel to infraclavicular area	Long term intravenous treatment in non-hospitalized patients (antibiotics, parenteral nutrition, chemotherapy); expected difficulties in management of the exit site in hospitalized patients (beard, humidity, tracheostomy, instability, etc.)		
	Tunnel to arm	Compromised skin integrity of the chest area; oral or endotracheal secretions over chest; implanted device on ipsilateral chest; chest surgery; contracted shoulder; etc.		
	Tunnel to back	Cognitive disorder resulting in device removal; contraindication to chest or arm exit site		
CICC (infraclavicular puncture)	Tunnel to lower chest	Long term intravenous treatment in non-hospitalized patients (antibiotics, parenteral nutrition, chemotherapy); expected problems in management of the exit site in hospitalized patients (tracheostomy, etc.)		
	Tunnel to arm	Compromised skin integrity of the chest area; oral or endotracheal secretions over chest; implanted device on ipsilateral chest; chest surgery; contracted shoulder; etc.		
	Tunnel to back	Cognitive disorder resulting in device removal; contraindication to chest or arm exit site		
FICC (puncture at the groin)	Tunnel to the abdomen	Non-emergency line in walking patients with contraindication to PICC/		
	Tunnel to mid-thigh	Non-emergency line in bedridden patients with contraindication to PICC/CICC		
FICC (puncture at mid-thigh)	Tunnel to the abdomen	Non-emergency line in walking patients with contraindication to PICC/		
	Tunnel to distal thigh	Long term intravenous treatment in bedridden patients with contraindication to PICC/CICC		

Techniques in vascular access



Rapid Assessment of Vascular Exit Site and © The Author(s) 2021 Tunneling Options (RAVESTO): A new decision tool in the management of the complex vascular access patients

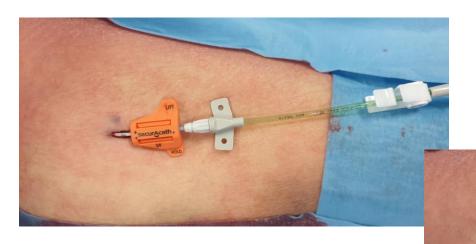
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\$SAGE

Matthew D Ostroff<sup>1</sup>, Nancy Moureau<sup>2</sup> and Mauro Pittiruti<sup>3</sup>



#### SAS - GLUE



Stop bleeding

Synergistic effect

Reduction of infectious risk

Reduction of the risk of dislocation



## SAS – COVID-19

Pittiruti and Pinelli Critical Care (2020) 24:269 https://doi.org/10.1186/s13054-020-02997-1

Critical Care

Editorial

DOI: 10.1177/1129729820923935

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Choice and management of vascular access in the context of COVID-19 outbreak in Italy: Recommendations from clinical practice

Intensiva, SIAARTI)

(Società Italiana di Anestesia, Analgesia, Rianimazione e Terapia

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The Journal of Vascular Access

SSAGE

Davide Vailati<sup>1</sup>, Giorgia Montrucchio<sup>2</sup>, Vittorio Cerotto<sup>3</sup> Giuseppe Capozzoli<sup>4</sup>, Fabio Gori<sup>5</sup>, Flavia Petrini<sup>6,7</sup> and Luca Brazzi<sup>2,8</sup>; on behalf of the Italian Society of Anesthesia and Intensive Care

#### COMMENTARY

**Open Access** 

Check for updates

Recommendations for the use of vascular access in the COVID-19 patients: an Italian perspective

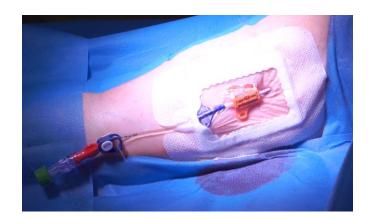
Mauro Pittiruti<sup>1\*</sup>, Fulvio Pinelli<sup>2</sup> on behalf of the GAVeCeLT Working Group for Vascular Access in COVID-19

As the risk of central venous catheter dislodgment is particularly high in the COVID-19 patient, particularly during the maneuvers of pronation-supination, consider the use of subcutaneously anchored securement.

Vascular access in COVID-19 patients: Smart decisions for maximal safety

Giancarlo Scoppettuolo<sup>1</sup>, Daniele Guerino Biasucci<sup>2</sup>

and Mauro Pittiruti<sup>3</sup>®



- complexity of clinical cases
- frequent pronation of patients
- exceptional workload in ICU





Fabrizio Brescia D, Mauro Pittiruti D, Laura Roveredo, Chiara Zanier, Antonietta Morabito, Elisabetta Santarossa, Valentina Da Ros, Marcella Montico and Fabio Fabiani

#### Methods

Study design and setting

This is a retrospective cohort study conducted in the Unit of Anesthesia, Intensive Care Medicine and Vascular Access Team of CRO National Cancer Institute, a Clinical and Research Cancer Institute located in Aviano (PN), Italy. We analyzed all PICCs secured with SAS in cancer patients, during the last 3 years (2018–2020). The SAS device used was the only currently available for clinical use, Securacath (Interrad Medical).



<sup>&</sup>lt;sup>1</sup>Unit of Anesthesia and Intensive Care Medicine, Vascular Access Team, Centro di Riferimento Oncologico di Aviano, IRCCS, Aviano, Italy

<sup>&</sup>lt;sup>2</sup>Department of Surgery, Fondazione Policlinico Universitario "A.Gemelli" IRCCS, Rome, Italy

<sup>&</sup>lt;sup>3</sup>Clinical Oncology Department, Centro di Riferimento Oncologico di Aviano, IRCCS, Aviano, Italy

<sup>&</sup>lt;sup>4</sup>Clinical Trial Office, Centro di Riferimento Oncologico di Aviano, IRCCS, Aviano, Italy

Fabrizio Brescia D, Mauro Pittiruti D, Laura Roveredo, Chiara Zanier, Antonietta Morabito, Elisabetta Santarossa, Valentina Da Ros, Marcella Montico and Fabio Fabiani



#### Technique of PICC insertion

PICC insertion was performed by experienced practitioners of our Vascular Access Team. In all patients, the procedure was performed according to our local insertion bundle for PICC insertion, which includes: pre-procedural ultrasound vascular assessment following the RaPeVA protocol (RaPeVA=Rapid Peripheral Vein Assessment), 11 measurement of vein diameter and respect for a catheter/vein ratio less than or equal to 1:3, skin antisepsis with 2% chlorhexidine, maximal barrier precautions, ultrasound-guided venipuncture, use of intracavitary electrocardiography method to verify the correct position of the tip of the catheter at the cavo-atrial junction, location of the exit site in Dawson's green zone (adopting tunneling from the yellow to the green zone, if necessary), 12 catheter securement with SAS, and sealing of the exit site with cyanoacrylate glue. Subsequent dressings and saline flushing of the PICC were performed weekly. The care and maintenance of the devices was entrusted to specialized nurses of our Access Vascular Team, according to institutional protocols.

#### **Outcomes**

Primary endpoints were (a) the efficacy of SAS, in terms of reducing the risk of dislocation and the need to reposition the vascular access device, (b) as well as its safety, evaluated investigating the incidence of immediate complications during SAS placement (difficulty, pain, etc.), early complications, that is, within 48 h (pain, local bleeding, etc.) and late complications (pain, malfunction, local or systemic infection, reversible or irreversible occlusion, catheter-related venous thrombosis, skin lesions due to the nitinol anchors, pressure ulcer of the device on the skin, etc.).



Fabrizio Brescia D, Mauro Pittiruti D, Laura Roveredo, Chiara Zanier, Antonietta Morabito, Elisabetta Santarossa, Valentina Da Ros, Marcella Montico and Fabio Fabiani



#### Results

A total of 639 patients had a PICC inserted and secured with SAS in the last 3 years (2018–2020) (Table 1).

PICCs of different brands and calibers were inserted: 254 Lifecath PICC Easy 4 Fr (Vygon), 114 Lifecath PICC Easy 5 Fr (Vygon), 97 HealthPICC 4 Fr single-lumen (Plan-1-Health), 153 HealthPICC 5 Fr single-lumen (Plan-1-Health), 21 HealthPICC 5 Fr double-lumen (Plan-1-Health). Indications for PICC insertion was chemotherapy in 120 patients, parenteral nutrition and chemotherapy in 410 patients, and parenteral nutrition in 109 patients.

DISLOCATIONS: 3 patients with psychomotor agitation 4 mismatch between the size of the catheter and the size of the SAS

As regards the effectiveness of securement with SAS, we recorded dislodgment only in seven patients (1.1%): three of these patients were non-collaborative patients with psychomotor agitation. In the remaining four cases, dislodgment occurred due to a mismatch between the size of the catheter and the size of the SAS. This was not related to an error of the operator but to an actual inconsistency of the caliber of the catheter as stated by the manufacturer. In fact, all four cases of dislodgment occurred with 4 Fr LifeCath PICC Easy (Vygon) secured with 4 Fr SAS or with 5 Fr LifeCath PICC Easy (Vygon) secured with 5 Fr SAS. In the early phase of our experience, noting this issue, we understood that the actual size of these catheters is slightly smaller than that the figure declared by the manufacturer. After these unexpected dislodgments, we have been using 3 Fr SAS for 4 Fr Lifecath PICC Easy and 4 Fr SAS for 5 Fr Lifecath PICC Easy, thus eliminating the risk of dislodgment.

No significant immediate complication during SAS placement was reported.

Fabrizio Brescia D. Mauro Pittiruti D. Laura Roveredo, Chiara Zanier, Antonietta Morabito, Elisabetta Santarossa, Valentina Da Ros, Marcella Montico and Fabio Fabiani



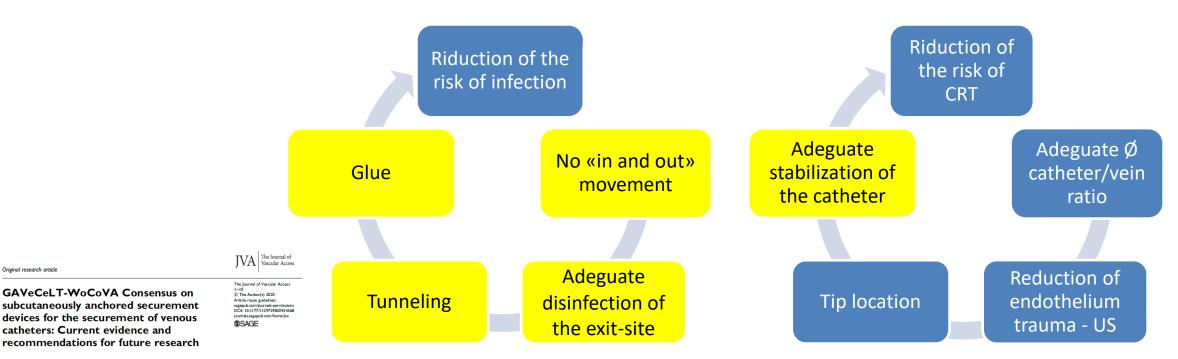
Table 2. Early complications (within 48h).

	N (%)	95% CI
	14 (70)	75% CI
PICC malfunction	0 (0.00)	_
Bleeding or hematoma at the exit-site	0 (0.00)	_
Skin ecchymosis	24 (3.8%)	2.4%-5.5%
Local pain at the exit-site	0 (0.00)	_
Early infection	0 (0.00)	_

Table 3. Late complications (analysis of 93,078 catheter days).

	N (%)	95% CI
Catheter-related bloodstream infection (CRBSI)	16 (2.5)	1.4-4.0
Symptomatic catheter-related venous thrombosis	12 (1.9)	1.0-3.3
Reversible lumen occlusion	15 (2.3)	1.3-3.8
Irreversible lumen occlusion	0 (0.00)	_
Pressure ulcers and painful inflammation	17 (2.66)	1.6-4.2
	Incidence/1000 catheter days	•
Catheter-related bloodstream infection (CRBSI)	0.17	0.10-0.28

#### SAS – INFECTION AND THROMBOSIS



#### Conclusion

There is no evidence that SAS may be effective in reducing other catheter-related complications such as venous thrombosis or exit site infection or blood-stream infection. There is no evidence either that they might increase the incidence of these or other major catheter-related complications.

Theoretical advantage



#### SAS – INFECTION

> Am J Infect Control. 2020 Jun 17;S0196-6553(20)30560-5. doi: 10.1016/j.ajic.2020.06.178. Online ahead of print.

## Catheter Securement Impact on PICC-related CLABSI: A University Hospital Perspective

M S Rowe <sup>1</sup>, K Arnold <sup>2</sup>, T R Spencer <sup>3</sup>



Studio retrospettivo osservazionale: 7779 PICC 2015-2018

Device	CLABSI (n=47)	No CLABSI (n=7732)	Total	Cumulative Incidence
AESD	15	823	838	1.79%
SESD	32	6909	6941	0.46%
			Risk Ratio	3.88
			Percent Relative Effect	288%

Table 3. CLABSI by stabilization device.



#### SAS – COST EFFECTIVENESS



Evaluating safety, efficacy, and cost-effectiveness of PICC securement by subcutaneously anchored stabilization device

Pietro Antonio Zerla<sup>1</sup>, Antonio Canelli<sup>1</sup>, Lidia Cerne<sup>1</sup>, Giuseppe Caravella<sup>2</sup>, Alessandra Gilardini<sup>2</sup>, Giuseppe De Luca<sup>3</sup>, Ana Maria Aricisteanu<sup>4</sup>, Raffaele Venezia<sup>4</sup>

TABLE II - Cost comparison between adhesive stabilization and subcutaneously anchored sutureless device (SAS)

	SAS	Adhesive stabilization device
Maintenance performed	709	709
No. devices used	30	709
Device cost (€)	30	6
Stabilization total cost (€)	900	4.254
SAS savings (€)	3.354	

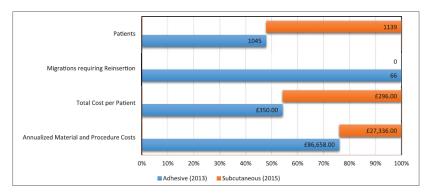
Original research article

Intravascular catheter migration: A cross-sectional and health-economic comparison of adhesive and subcutaneous engineered stabilisation devices for intravascular device securement

JVA The Journal of Vascular Access

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**Graph 1.** Comparison between adhesive and subcutaneous devices groups, highlighting costs per patient, reinsertions and material costs.

when the catheter is meant to remain in place for more than 5-6 weeks or when a high risk of dislodgment is anticipated



#### CONCLUSIONS



Subcutaneously anchored securement for peripherally inserted central catheters: Immediate, early, and late complications

Fabrizio Brescia<sup>1</sup>, Mauro Pittiruti<sup>2</sup>, Laura Roveredo<sup>1</sup>, Chiara Zanier<sup>1</sup>, Antonietta Morabito<sup>1</sup>, Elisabetta Santarossa<sup>1</sup>, Valentina Da Ros<sup>3</sup>, Marcella Montico<sup>4</sup> and Fabio Fabiani<sup>1</sup>

### SAFETY AND PROTECTION

- Subcutaneously anchored securement of PICCs is associated with very low risk of dislodgment and that this risk is limited to
- <u>Non-collaborative patients</u>: SAS must surely be used as securement, but they are not enough, since some other strategies should be addes, such as **tunneling** the catheter so to move the exit site far from the hands away to an area not reachable by the patient and safer during mobilization of the patient RAVESTO protocol

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### SAFETY AND PROTECTION

- Patients at high risk of dislocation regardless of duration (ICU, pronation, history of previous dislocation)
- More than 5-6 weeks of catheter stay duration
- Patients at risk of MARSI or with skin problems (allergies, burns, hyperhidrosis)
- Home management problems



#### CONCLUSIONS



Subcutaneously anchored securement for peripherally inserted central catheters: Immediate, early, and late complications

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## SAFETY AND PROTECTION

- Choice of a SAS of adequate size
- Subcutaneous positioning of the nitilon bars
- Adeguate training before clinicla use
- Use SAS and **glue** to simultaneously achieve adequate catheter safety and optimal exit site protection





## Thank you for your attention