

Selection of the vascular catheter: can it minimize the risk of infection?

E. Bouza, M. Guembe, P. Muñoz

▶ To cite this version:

E. Bouza, M. Guembe, P. Muñoz. Selection of the vascular catheter: can it minimize the risk of infection?. International Journal of Antimicrobial Agents, 2010, 36, 10.1016/j.ijantimicag.2010.11.002 . hal-00650365

HAL Id: hal-00650365 https://hal.science/hal-00650365

Submitted on 10 Dec 2011

HAL is a multi-disciplinary open access archive for the deposit and dissemination of scientific research documents, whether they are published or not. The documents may come from teaching and research institutions in France or abroad, or from public or private research centers. L'archive ouverte pluridisciplinaire **HAL**, est destinée au dépôt et à la diffusion de documents scientifiques de niveau recherche, publiés ou non, émanant des établissements d'enseignement et de recherche français ou étrangers, des laboratoires publics ou privés.

Accepted Manuscript

Title: Selection of the vascular catheter: can it minimize the risk of infection?

Authors: E. Bouza, M. Guembe, P. Muñoz



PII: DOI: Reference:	S0924-8579(10)00463-2 doi:10.1016/j.ijantimicag.2010.11.002 ANTAGE 3456	

To appear in: International Journal of Antimicrobial Agents

Please cite this article as: Bouza E, Guembe M, Muñoz P, Selection of the vascular catheter: can it minimize the risk of infection?, *International Journal of Antimicrobial Agents* (2010), doi:10.1016/j.ijantimicag.2010.11.002

This is a PDF file of an unedited manuscript that has been accepted for publication. As a service to our customers we are providing this early version of the manuscript. The manuscript will undergo copyediting, typesetting, and review of the resulting proof before it is published in its final form. Please note that during the production process errors may be discovered which could affect the content, and all legal disclaimers that apply to the journal pertain.

Selection of the vascular catheter: can it minimize the risk of infection?

E. Bouza^{a,b, *}, M. Guembe^a, P. Muñoz^{a,b}

^a Department of Clinical Microbiology and Infectious Diseases, Hospital General Universitario Gregorio Marañón, Universidad Complutense,

Madrid, Spain

^b Red Española de Investigación en Patología Infecciosa (REIPI),

RD06/0008/1025, Spain

* Corresponding author. Address for correspondence: Servicio de Microbiología Clínica y Enfermedades Infecciosas; Hospital General Universitario Gregorio Marañón, C/. Dr Esquerdo 46, 28007 Madrid, Spain. Tel.: +34 91 586 84 53; fax: +34 91 504 49 06.

E-mail address: ebouza@microb.net (E. Bouza).

Abstract

Data regarding the prevention of catheter-related bloodstream infection (CRBSI) by making the correct decisions about when to place a central line, the appropriate selection of catheter composition and the size and number of lumens, a suitable choice of insertion site and the technique used are not well reported in recent medical literature. There is no clear evidence that the composition of the catheters presently on the market makes a significant difference to the risk of infection. Several prospective studies suggest that femoral vein location represents the highest risk of infection, followed by jugular vein and subclavian vein positioning, however, most articles do not correct for basic confounding variables. Several papers have reported that arterial catheters have a similar risk of infection as central venous catheters (CVCs). The slight increase in infection risk when using multi-lumen catheters is probably offset by their improved convenience. Current evidence does not support routine tunnelling of short-term catheters until its efficacy is evaluated at different placement sites, using specific catheters and situations and in relation to other preventive interventions. Cuffing is usually applied only to long-term tunnelled catheters. The available evidence suggests that chlorhexidine-silver sulfadiazine, minocycline-rifampicin CVCs and antifungal-coated catheters are useful in decreasing the incidence of CRBSI when other measures are not effective.

Keywords:

Catheter-related bloodstream infections

Selection of catheters

Bacteraemia

Colonized lumens

1. Introduction

Catheter-related bloodstream infections (CRBSI) are a major problem in the area of infection control [1–5]. Programmes to minimize the risk of CRBSI are founded in the systematic application of bundles of care and the implementation of well-established procedures in both the placement of the catheter and its subsequent care [6].

Daily assessment of the need for a central line and adequate and early removal of the catheter are important aspects of these bundles. However, data regarding the prevention of infection achieved by correct decisions about when to place a central line, the appropriate selection of catheter composition and the size and number of lumens, a suitable choice of insertion site and the technique used are not so well reported in recent medical literature [7]. Our aim is to review several issues regarding the prevention of CRBSI, mainly those around the catheter and its characteristics.

2. Catheter material

There is no clear evidence that the composition of the catheters presently on the market makes a significant difference to the risk of infection. Selection of the material is mainly performed on the basis of elasticity, resistance to breakage, risk of obstruction and clinical parameters. Moretti et al. stated [8] that infection rates may depend more on non-catheterrelated factors, such as adherence to infection control standards, selection

of the insertion site, duration of catheter placement and frequency of dressing changes.

3. Location of catheter insertion

Central venous catheters (CVCs) are usually indicated when peripheral veins are inaccessible or when administering vasoactive drugs, irritating solutions or parenteral alimentation. They are also required for haemodialysis or haemofiltration and for hemodynamic monitoring. The selection of the location for CVC insertion is not generally taken on the basis of the risk of infection but on the medical need and the risk of complications during the insertion or maintenance of the catheter. It is not clear if clinical experience in catheter implantation and the number of insertion attempts may influence the risk of infection. Whenever possible, catheters should not be inserted through the site of burns or infection. No randomized trials have been performed to assess a clear superiority of one location over another [9]. However, several prospective studies using multivariate analysis suggest that femoral vein location represents the highest risk of infection, being greater than jugular vein location, and the risk with jugular insertion being greater than location in the subclavian vein [10–16]; however, most articles do not correct for basic confounding variables. Deshpande et al. [16] assessed the risk of CVC infection with respect to the site of insertion in an intensive care unit (ICU) population. The subclavian, internal jugular and femoral sites were studied in a prospective epidemiological observation study. The optimal insertion site

for each patient was selected by the intensive care physicians. A total of 831 CVCs and 4735 catheter days in 657 patients were studied. The incidence of catheter infection and colonization was low overall and no statistically significant differences in the incidence of infection and colonization or duration of catheters among the insertion sites were found.

Until recently, the literature suggested that arterial catheters may have a lower infection risk than catheters located in central veins [17–19]. However, since 2005, several papers have reported that arterial catheters have a similar risk of infection as CVCs [20–22]. Lucet et al. [20] reported a study from seven French ICUs collecting 3532 catheters and 27 541 days of observation. The rates of colonization for arterial catheters and CVCs were 11.4 and 11.1/1000 days of exposure, respectively. Regarding the risk of infection, the figures were 1.0 and 1.1/1000 days for arterial catheters and CVCs, respectively.

Koh et al. [21] prospectively measured colonization and rates of CRBSI in arterial catheters in comparison with CVCs cared for in the same way. Overall, 301 arterial catheters in 252 adult and paediatric patients were observed for 1082 catheter days, and 618 central venous catheters in 410 patients were observed for 4040 catheter days. Arterial catheter colonization was not significantly different to that in CVCs. The arterial catheter should be accorded the same degree of importance as the CVC as a potential source of sepsis.

Table 1, modified from a meta-analysis published by Maki et al. in 2006 [7], compares the incidence and incidence density of infections in different catheter locations. Overall, the authors demonstrated the need to express the risk of catheter infection as incidence density, avoiding the bias produced by comparing plain incidence. For instance, comparing incidence densities, steel needles carry a higher risk for 1000 days of exposure (8.6) than peripherally inserted CVCs (0.5).

[Table 1 here]

4. Number of lumens in the catheter

The number of lumens should be chosen to meet clinical needs. Multilumen catheters are commonly selected to avoid multiple CVC sites. The higher the number of lumens and the longer the catheter, the higher the resistance to flow. Dialysis catheters are typically double-lumen largebore catheters because of the high flow rates required for dialysis.

Multi-lumen central venous catheters may be associated with a slightly higher risk of infection when compared with single-lumen catheters. However, this relationship diminishes when only high-quality studies that control for patient differences are considered. The slight increase in infection risk when using multi-lumen catheters is probably offset by their improved convenience, thereby justifying the continued use of multilumen vascular catheters [23].

Nevertheless, the use of catheters with multiple lumens should be restricted, and if a catheter cannot be removed the permanent closure of unneeded lumens may reduce the risk of CRBSI [24,25].

5. Tunnelled versus non-tunnelled catheters

A meta-analysis of randomized controlled trials comparing tunnelling short-term CVCs to prevent catheter-related infection showed that tunnelling may decrease the infection rate. However, current evidence does not support routine tunnelling until its efficacy is evaluated at different placement sites, using specific catheters and situations and in relation to other preventive interventions [26].

Most information regarding tunnelling of catheters has been collected in adult patients. Nahum et al. [27], in a study of bacterial colonization of femoral CVCs in critically ill children, found that the tip colonization rate was higher in tunnelled catheters, but they were unable to demonstrate significant differences in the rates of CRBSI.

6. Cuffed vs non-cuffed catheters

There is no clear evidence that cuffing of short-term catheters significantly decreases the incidence of CRBSI. Cuffing is usually applied to long-term tunnelled catheters.

Flowers et al. in 1989 [28] reported a randomized controlled trial of an attachable subcutaneous cuff for the prevention of CVC-related infection among patients receiving intensive care. Catheters were placed into new sites with or without a cuff and were dressed with antibiotic ointment

containing polymyxin, neomycin and bacitracin. Microbial colonization developed in 34.5% of 29 control catheters and 7.7% of 26 cuffed catheters. CRBSI occurred with 13.8% of control vs. 0% of cuffed catheters. These data suggested that the cuff could reduce the incidence of catheter-related infection among high-risk patients receiving cathetersite care with an antibacterial ointment.

By contrast, Groeger et al. [29] evaluated the effect of a silverimpregnated cuff on the incidence of catheter-related bacteraemia/fungaemia or tunnel tract infection in 200 randomly selected cancer patients with chronic dual-lumen tunnelled venous access catheters. The hazard rate for infection/day was not significantly different and the study indicated that a silver-impregnated cuff did not decrease the incidence of catheter-related bacteraemia/fungaemia.

More recently, Trerotola et al. [30] randomized 84 patients to receive a 5F single- or 6F dual-lumen small-bore central catheter with or without a polyester cuff. The overall infection rate was 0.4 per 1000 catheter days and cuffing did not make any difference to the risk of infection in jugular small-bore catheters. Cuffed catheters were no more difficult to insert but took slightly longer to remove and often required local anaesthesia for removal.

7. Totally implanted vs non-implanted catheters

Totally implanted catheters are generally accepted as carrying a lower risk for CRBSI [7,31,32]. However, no properly performed comparative

studies permit us accurately to assess the difference. In general, totally implanted catheters are accessed less frequently, are usually associated with more precautions and are inserted by more specialized healthcare workers than open catheters. An infection incidence density of 0.1 episodes/1000 days in place has been reported [7].

Prospective studies suggest that there are no major differences in risk of infection between valved and non-valved totally implantable catheters or between low-profile and conventional devices [33,34].

A randomized comparison of complications from three different central venous access systems was not able to demonstrate significant differences between polyurethane ChemoSite, polyurethane Port-a-Cath and silicone Port-a-Cath devices [35].

8. Antimicrobial-coated catheters

The greatest amount of literature regarding catheter types in recent years has been devoted to the potential decrease of the risk of infection in catheters coated with various antimicrobial substances [36,37]. Ramritu et al. [38] performed a systematic review comparing the relative effectiveness of antimicrobial-coated catheters in ICUs. Another excellent review is by Casey et al [39]. In summary, the first-generation externally coated chlorhexidine–silver sulfadiazine CVCs reduced colonization and CRBSI when compared with non-coated catheters. Externally and internally coated catheters were superior to those coated only externally. Antibiotic-coated catheters (minocycline–rifampicin) were superior to

those coated with chlorhexidine-silver sulfadiazine in reducing colonization and CRBSI.

A cohort study by Lorente et al. [40] showed a drastic reduction of CRBSI in femoral and jugular CVCs using rifampicin–miconazole-impregnated catheters compared with standard catheters (0 vs 8.62 cases/1000 catheter days for femoral catheters and 0 vs 4.93 cases/1000 catheter days for jugular catheters).

Despite the shortcomings of the studies of antimicrobial-coated catheters, the available evidence suggests that chlorhexidine–silver sulfadiazine, minocycline–rifampicin CVCs and antifungal-coated catheters are useful if the incidence of CRBSI is above institutional goals despite full implementation of infection prevention interventions.

The role of silver particles in protecting CVCs from colonization and reducing the risk of CRBSI is not yet clear [41,42,43]. In a prospective, randomized, non-blinded, controlled study Hagau et al. [44] found no significant difference in colonization and CRBSI per 1000 catheter days but a potential delay in colonization in silver-impregnated vs standard catheters. Whereas standard catheters were first colonized 3 days after insertion, silver-impregnated catheters were first colonized 5 days after insertion.

Funding: E. Bouza received an honorarium for writing this article. The funds for the honorarium were provided by Novartis AG, Switzerland and were handled by the organizing committee of the 4th European

Conference on Bloodstream Infections for the publication of this

supplement.

Competing interests: None.

Ethical approval: Not required.

References

[1] Mermel LA, Allon M, Bouza E, Craven DE, Flynn P, O'Grady NP, et al. Clinical practice guidelines for the diagnosis and management of intravascular catheter-related infection: 2009 Update by the Infectious Diseases Society of America. Clin Infect Dis 2009; 49: 1–45.

[2] O'Grady NP, Alexander M, Dellinger EP, Gerberding JL, Heard SO, Maki DG, et al. Guidelines for the prevention of intravascular catheterrelated infections. Centers for Disease Control and Prevention. MMWR Recomm Rep 2002; 51 (RR-10): 1–29.

[3] Bouza E, Burillo A, Munoz P. Empiric therapy for intravenous central line infections and nosocomially-acquired acute bacterial endocarditis. Crit Care Clin 2008;24:293–312, viii-ix.

[4] Bouza E, Hortal J, Munoz P, Perez MJ, Riesgo MJ, Hiesmayr M.
Infections following major heart surgery in European intensive care units: there is room for improvement (ESGNI 007 Study). J Hosp Infect 2006; 63: 399–405.

[5] Bouza E, Burillo A, Munoz P. Catheter-related infections: diagnosis and intravascular treatment. Clin Microbiol Infect 2002;8:265–74.

[6] Pronovost P, Needham D, Berenholtz S, Sinopoli D, Chu H, Cosgrove
 S, et al. An intervention to decrease catheter-related bloodstream
 infections in the ICU. N Engl J Med 2006; 355: 2725–32.

[7] Maki DG, Kluger DM, Crnich CJ. The risk of bloodstream infection in adults with different intravascular devices: a systematic review of 200 published prospective studies. Mayo Clin Proc 2006; 81: 1159–71.

[8] Moretti EW, Ofstead CL, Kristy RM, Wetzler HP. Impact of central venous catheter type and methods on catheter-related colonization and bacteraemia. J Hosp Infect 2005;61:139–45.

[9] Ruesch S, Walder B, Tramer MR. Complications of central venous
 catheters: internal jugular versus subclavian access—a systematic review.
 Crit Care Med 2002; 30: 454–60.

[10] Harden JL, Kemp L, Mirtallo J. Femoral catheters increase risk of infection in total parenteral nutrition patients. Nutr Clin Pract
1995; 10:60–6.

[11] Ishizuka M, Nagata H, Takagi K, Kubota K. Femoral venous catheterization is a major risk factor for central venous catheter-related bloodstream infection. J Invest Surg 2009; 22: 16–21.

[12] Ishizuka M, Nagata H, Takagi K, Horie T, Furihata M, Nakagawa A, et al. External jugular Groshong catheter is associated with fewer complications than a subclavian Argyle catheter. Eur Surg Res 2008; 40: 197–202.

[14] Ameh V, Jones S. Best evidence topic report. Central venouscatheterisation: internal jugular or subclavian approach? Emerg Med J2007; 24:662–3.

[15] Nagashima G, Kikuchi T, Tsuyuzaki H, Kawano R, Tanaka H, Nemoto H, et al. To reduce catheter-related bloodstream infections: is the subclavian route better than the jugular route for central venous catheterization? J Infect Chemother 2006; 12:363–5.

[16] Deshpande KS, Hatem C, Ulrich HL, Currie BP, Aldrich TK, Bryan-Brown CW, et al. The incidence of infectious complications of central venous catheters at the subclavian, internal jugular, and femoral sites in an intensive care unit population. Crit Care Med 2005; 33: 13–20; discussion 234–5.

[17] Furfaro S, Gauthier M, Lacroix J, Nadeau D, Lafleur L, Mathews S.
 Arterial catheter-related infections in children. A 1-year cohort analysis.
 Am J Dis Child 1991;145:1037–43.

[18] Thomas F, Burke JP, Parker J, Orme JF Jr, Gardner RM, Clemmer TP, et al. The risk of infection related to radial vs femoral sites for arterial catheterization. Crit Care Med 1983; 11:807–12.

[19] Raad I, Abi-Said D, Carrasco CH, Umphrey J, Hill LA. The risk of infection associated with intra-arterial catheters for cancer chemotherapy. Infect Control Hosp Epidemiol 1998;19:640–2.

[20] Lucet JC, Bouadma L, Zahar JR, Schwebel C, Geffroy A, Pease S, et al. Infectious risk associated with arterial catheters compared with central venous catheters. Crit Care Med 2010; 38: 1030–5.

[21] Koh DB, Gowardman JR, Rickard CM, Robertson IK, Brown A. Prospective study of peripheral arterial catheter infection and comparison

with concurrently sited central venous catheters. Crit Care Med 2008; 36: 397–402.

[22] Traore O, Liotier J, Souweine B. Prospective study of arterial and central venous catheter colonization and of arterial- and central venous catheter-related bacteremia in intensive care units. Crit Care Med 2005; 33: 1276–80.

[23] Dezfulian C, Lavelle J, Nallamothu BK, Kaufman SR, Saint S. Rates of infection for single-lumen versus multilumen central venous catheters: a meta-analysis. Crit Care Med 2003;31:2385–90.

[24] Templeton A, Schlegel M, Fleisch F, Rettenmund G, Schobi B, Henz S, et al. Multilumen central venous catheters increase risk for catheterrelated bloodstream infection: prospective surveillance study. Infection 2008; 36: 322–7.

[25] Zurcher M, Tramer MR, Walder B. Colonization and bloodstream infection with single- versus multi-lumen central venous catheters: a quantitative systematic review. Anesth Analg 2004;99:177–82.

[26] Randolph AG, Cook DJ, Gonzales CA, Brun-Buisson C. Tunneling short-term central venous catheters to prevent catheter-related infection: a meta-analysis of randomized, controlled trials. Crit Care Med 1998; 26: 1452–7.

[27] Nahum E, Levy I, Katz J, Samra Z, Ashkenazi S, Ben-Ari J, et al. Efficacy of subcutaneous tunneling for prevention of bacterial colonization

of femoral central venous catheters in critically ill children. Pediatr Infect Dis J 2002; 21: 1000–4.

[28] Flowers RH 3rd, Schwenzer KJ, Kopel RF, Fisch MJ, Tucker SI, Farr BM. Efficacy of an attachable subcutaneous cuff for the prevention of intravascular catheter-related infection. A randomized, controlled trial. JAMA 1989;261:878–83.

[29] Groeger JS, Lucas AB, Coit D, LaQuaglia M, Brown AE, Turnbull A, et al. A prospective, randomized evaluation of the effect of silver impregnated subcutaneous cuffs for preventing tunneled chronic venous access catheter infections in cancer patients. Ann Surg 1993;218:206–10.

[30] Trerotola SO, Patel AA, Shlansky-Goldberg RD, Solomon JA, Mondschein JI, Stavropoulos SW, et al. Short-term infection in cuffed versus noncuffed small bore central catheters: a randomized trial. J Vasc Interv Radiol 2010; 21: 203–11.

[31] Zhang Q, Jiao L, Zhou H. Comparison of implantable central venous ports with catheter insertion via external jugular cut down and subclavian puncture in children: single center experience. Pediatr Surg Int 2009; 25: 499–501.

[32] Adler A, Yaniv I, Steinberg R, Solter E, Samra Z, Stein J, et al. Infectious complications of implantable ports and Hickman catheters in paediatric haematology–oncology patients. J Hosp Infect 2006;62:358– 65.

[33] Carlo JT, Lamont JP, McCarty TM, Livingston S, Kuhn JA. A prospective randomized trial demonstrating valved implantable ports have fewer complications and lower overall cost than nonvalved implantable ports. Am J Surg 2004; 188: 722–7.

[34] Teichgraber UK, Streitparth F, Cho CH, Benter T, Gebauer B. A comparison of clinical outcomes with regular- and low-profile totally implanted central venous port systems. Cardiovasc Intervent Radiol 2009; 32:975–9.

[35] Vandoni RE, Guerra A, Sanna P, Bogen M, Cavalli F, Gertsch P. Randomised comparison of complications from three different permanent central venous access systems. Swiss Med Wkly 2009; 139: 313–6.

[36] Ostendorf T, Meinhold A, Harter C, Salwender H, Egerer G, Geiss HK, et al. Chlorhexidine and silver-sulfadiazine coated central venous catheters in haematological patients—a double-blind, randomised, prospective, controlled trial. Support Care Cancer 2005;13:993–1000.

[37] Carrasco MN, Bueno A, de las Cuevas C, Jimenez S, Salinas I, Sartorius A, et al. Evaluation of a triple-lumen central venous heparincoated catheter versus a catheter coated with chlorhexidine and silver sulfadiazine in critically ill patients. Intensive Care Med 2004; 30: 633–8.

[38] Ramritu P, Halton K, Collignon P, Cook D, Fraenkel D, Battistutta D, et al. A systematic review comparing the relative effectiveness of antimicrobial-coated catheters in intensive care units. Am J Infect Control 2008; 36: 104–17.

[39] Casey AL, Mermel LA, Nightingale P, Elliott TS. Antimicrobial central venous catheters in adults: a systematic review and meta-analysis. Lancet Infect Dis 2008; 8:763–76.

[40] Lorente L, Lecuona M, Ramos MJ, Jimenez A, Mora ML, Sierra A. The use of rifampicin-miconazole-impregnated catheters reduces the incidence of femoral and jugular catheter-related bacteremia. Clin Infect Dis 2008; 47: 1171–5.

[41] Bong JJ, Kite P, Wilco MH, McMahon MJ. Prevention of catheter related bloodstream infection by silver iontophoretic central venous catheters: a randomised controlled trial. J Clin Pathol 2003; 56: 731–5.

[42] Khare MD, Bukhari SS, Swann A, Spiers P, McLaren I, Myers J. Reduction of catheter-related colonisation by the use of a silver zeoliteimpregnated central vascular catheter in adult critical care. J Infect 2007; 54: 146–50.

[43] Kalfon P, de Vaumas C, Samba D, Boulet E, Lefrant JY, Eyraud D, et al. Comparison of silver-impregnated with standard multi-lumen central venous catheters in critically ill patients. Crit Care Med 2007; 35:1032–9.
[44] Hagau N, Studnicska D, Gavrus RL, Csipak G, Hagau R, Slavcovici AV. Central venous catheter colonization and catheter-related bloodstream infections in critically ill patients: a comparison between standard and silver-integrated catheters. Eur J Anaesthesiol 2009; 26: 752–8.

Table 1

Incidence and incidence density of catheter-line associated bloodstream infection

(CLABSI) with different types and location of catheter (modified from Maki et al. [7])

Devices	No. of studies	CLABSI rate	
		Per 100 devices	Per 1000 days.
		Pooled mean	Pooled mean
Plastic peripheral	110	0.1	0.5
catheters			
Steel needles	1	2	8.6
Venous cutdown	1	3.7	9
Midline catheters	3	0.4	0.2
Arterial catheters	14	0.8	1.7
CVCs short-term, non-	79	4.4	2.7
cuffed, non-tunnelled			
CVCs short-term, non- cuffed, tunnelled	9	4.7	1.7
CVCs chlorhexidine- silver sulfadiazine	18	2.6	1.6
CVCs minocycline-	3	1	1.2

rifomniain

rifampicin

CVCs silver	2	5.2	4.7
impregnated			
CVCs silver iontophoretic	2	4	3.3
Haemodialysis,	16	8	4.8
temporary, non-cuffed			
Haemodialysis, long- term, cuffed, tunnelled	16	21.2	1.6
CVCs cuffed, tunnelled	29	22.5	1.6
Ports, central	14	3.6	0.1
Ports, peripheral	3	4	0.1

CVC, central venous catheter.