

Comment gérer les complications des cathéters permanents

SUMMARY



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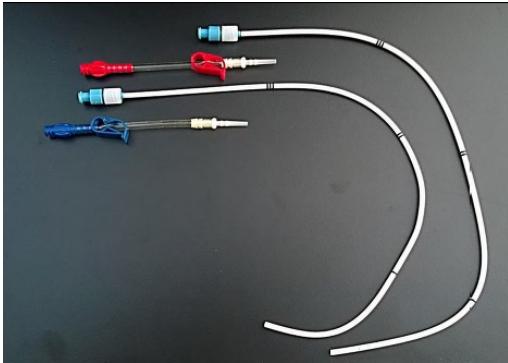


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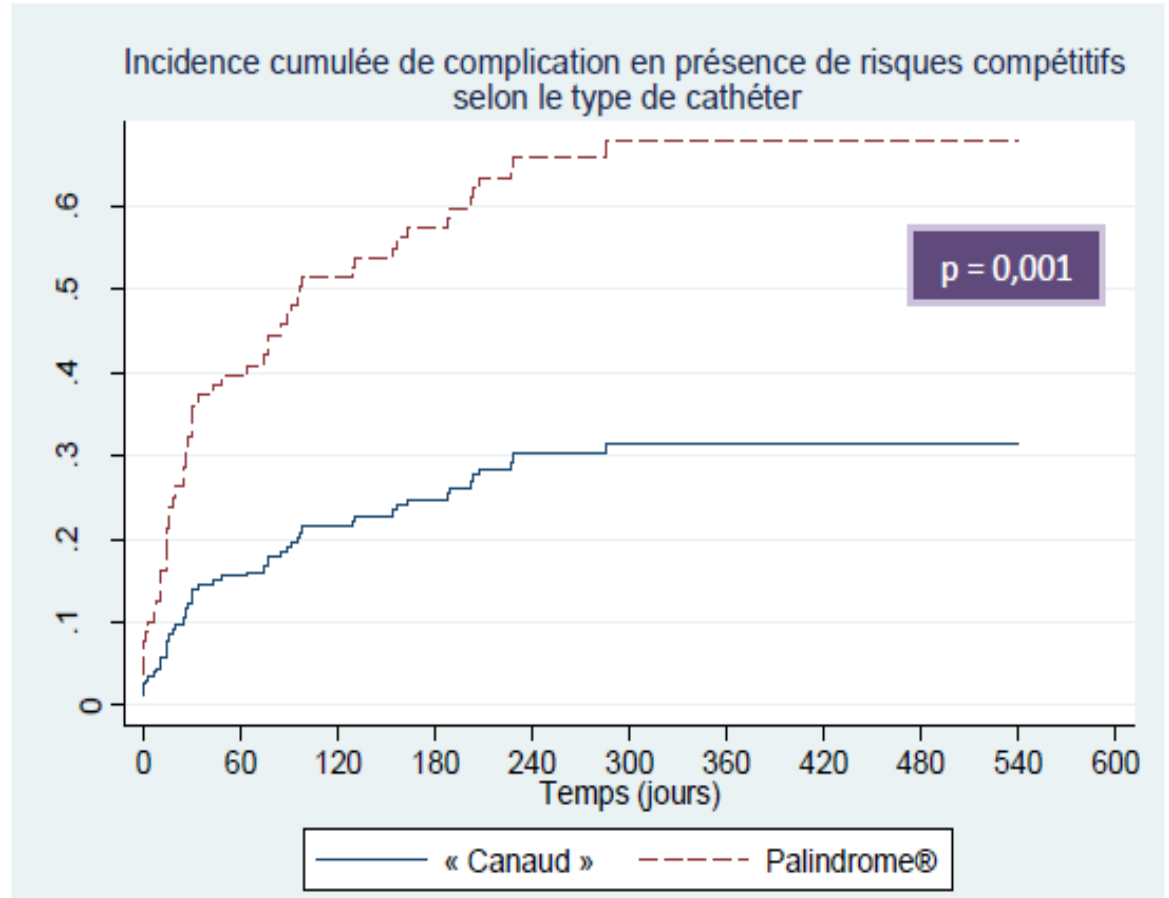
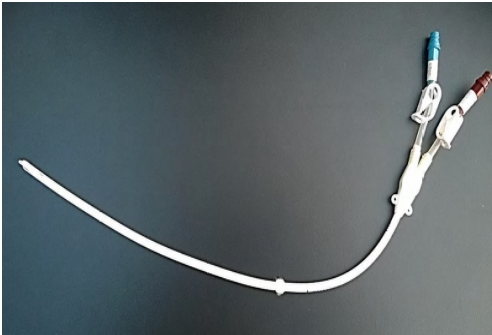
Faculté de Médecine de Sousse



Bi-cathéters Indépendants



Cathéter monobloc Bi- Lumières



Dysfunction
Thromboses
Infection

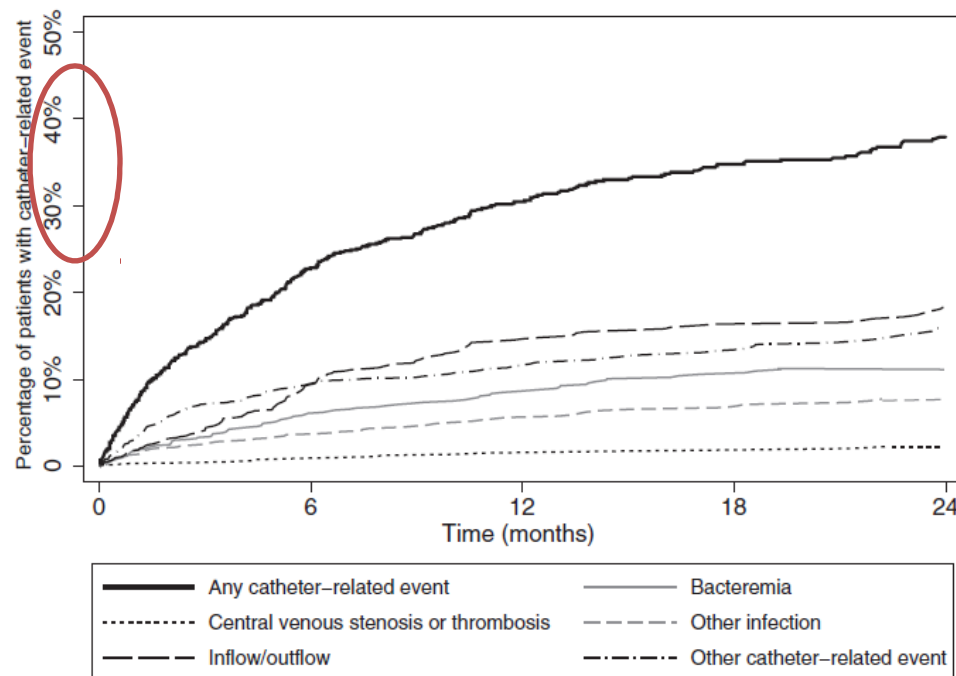


Figure 2. Cumulative incidence of catheter-related events. Percentage of patients who have experienced at least 1 catheter-related event from the time of tunneled catheter insertion (in months). The top black line is the first of any catheter-related event (eg, bacteremia or inflow/outflow) while the other lines are cause specific (eg, first bacteremia irrespective of other complications). Death, transplantation, recovery, and starting peritoneal dialysis therapy were considered competing events. Censoring events were transfers, losses to follow-up, or end of study follow-up.

Dysfonction mécanique

incapacité à fournir un débit suffisant pour une dialyse adéquate

KDOQI: failure to attain and maintain an extracorporeal blood flow, Q_b , of 300 mL/min or greater at a prepump arterial pressure more

Table I. Proposed Definitions for Catheter Dysfunction.^{3,5,8}

| |
|--|
| High arterial pressures (<-250 mm Hg) |
| High venous pressures (>250 mm Hg) |
| High pressure alarms |
| Decreased blood flow rates |
| Inability to withdraw and/or flush catheter lumens |
| Need to reverse lines |
| Reduced urea clearance ($Kt/V < 1.2$, or urea reduction ratio < 65%) |

negative than -250 mm Hg.
Incidence 30% des patients une dysfonction/séance/mois

Étiologies de la dysfonction des cathéters

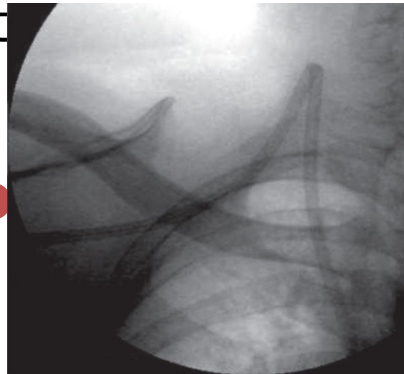
Précoces: première semaine

Mécaniques

- Position incorrecte:
 - bout de KT contre une paroi vasculaire
 - Loin de la jonction VC/Oreillette

- Plicature

=> **Repositionnement**



Tardives

- Agressions répétées/turbulences
- + Inflammation + coagulation

=> **Thrombose**
:Catheter-related thrombosis CRT

Prévention des thrombi liés au cathéter

- Verrou à base d'anticoagulant: Citrate de sodium 4% ou Héparine ou Bicarbonate
- Pré-CLOT trial: tPA *tissue Plasminogène Activator*

Plus efficace mais plus coûteux

- Antiagrégants plaquettaires: effet controversé
- Anticoagulation systémique: risque hémorragique > bénéfique



Canadian Society of Nephrology/
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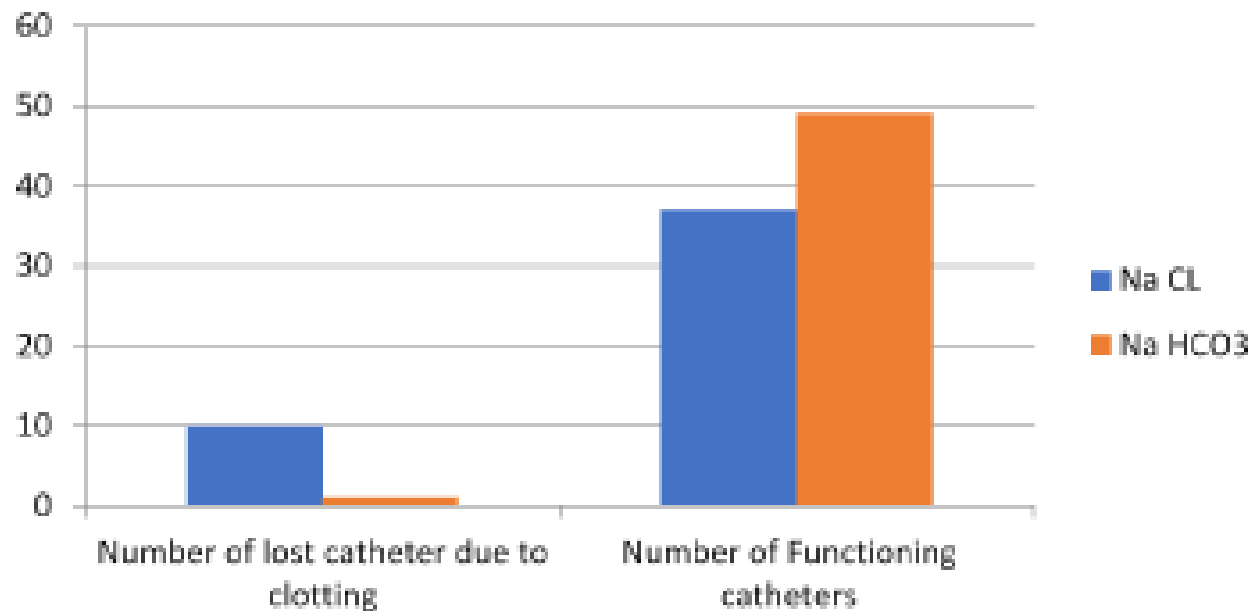


Figure 2. Shows the Kaplan-Meier survival curves for the time course of the lost and functioning catheters in both Na CL Lock solution and Na HCO₃ Lock solution groups due to clot formation during a period of 6 months. P = 0.003.

Ttt des thrombi liés aux cathéters

- Purge par sérum physiologique en flash (rupture du KT et d'embolies)
- Enlever?
 - Oui si alternative vasculaire
- Sinon
- Injection possible mais pas l'aspiration:
Anticoagulation systémique via le KT
- Occlusion totale: thrombolyse
- Échec et pas d'infection: changement sur guide

CRT: place de l'anticoagulation systémique

Cathéter en place

- HNF en IV, bolus 60ui/Kg puis 12 ui/kg/h TCA 1.5 à 2.5
- HBPM: risque hémorragique
- AVK: INR 2,5 à 3,5
- Si efficace: à garder tant que le KT est en place

Cathéter enlevé

- Anticoagulation x 6 sem
- Début / risque d'embolie
 - Faible: Au moment de l'ablation
 - Élevé: 5 jours avant
- Thrombus < 1.5cm: arrêt précoce

Traitement des CRT: Thrombolyse

- Activateur tissulaire du plasminogène t-PA (altéplase et reteplase) => succès ds 83% à 98%

| Type | Dose | Time | Delivery |
|--|--|--|---|
| Nonfunctioning catheter (Qb < 150/200 mL/min or inability to withdraw) (short or long dwell) | Typical dose: 2 mg tPA instillation (1 mg in each lumen) Alternative dose: 2 mg tPA per lumen | 30 min after instillation (short dwell) Alternative: wait additional 2 h (long dwell) if 30 min fails ²⁸ | tPA instilled as dwell for 30-60 min Alternatives: push one-third of dose every 10 min; advance the initial pushed tPA at 10- to 15-min intervals with 0.3 mL saline, or deliver via infusion pump over 30 min |
| Poor functioning catheter with dialysis stopped | 2-4 mg tPA | As above | As above |
| Poor functioning catheter with dialysis ongoing (eg, Qb > 200 mL/min) but frequent alarms (intradialytic infusion) | 2-4 mg tPA | Over 1 h | Via infusion pump—delivered in the dialysis circuit with catheter lumens reversed for 30 min, then with normal positions for next 30 min |
| Poor functioning catheter post dialysis (lock, post hemodialysis dwell, overnight dwell) | 1-2 mg tPA per lumen | Until next dialysis | tPA instilled as push; allowed to dwell in catheter lumen for 48-72 h, until the next hemodialysis session |

Note. tPA = tissue plasminogen activator.

Ann Vasc Surg. 2018 Aug;51:298-305.

t-PA > Urokinase

Table 2. Catheter-locking solutions in the management of catheter dysfunction

| Study (Reference)/Design | Intervention | Results |
|--|---|---|
| Haire <i>et al.</i> , 1994 (55) Randomized (<i>n</i> =50) | Urokinase (10,000 IU/ml) versus tPA (2 mg) | tPA significantly better in restoring catheter function |
| Savader <i>et al.</i> , 2001 (52) Prospective (<i>n</i> =55) | rtPA infusion (2.5 mg over 3 h) | 91% technical success rate; primary patency low but improved secondary patency |
| Little and Walshe, 2002 (54) Prospective (<i>n</i> =336) | tPA (1 mg/ml); dwell time 2–8 h | Patency declined with successive use; median patency only 5–7 treatments |
| Haire <i>et al.</i> , 2004 (56) Randomized (<i>n</i> =180) | Urokinase (5000 IU/ml) versus placebo | Urokinase significantly better in restoring catheter function |
| Falk <i>et al.</i> , 2004 (49) Retrospective (<i>n</i> =19) | Reteplase (0.4 units); dwell time 30 min–1 h | Initial patency 88%; no adverse events |
| Macrae <i>et al.</i> , 2005 (50) Randomized (<i>n</i> =60) | tPA (1 mg/ml); dwell time 1 h versus 48–96 h | Initial patency 78% but short-lived; median patency only 6 treatments |

tPA, tissue plasminogen activator; rtPA, recombinant tPA.

Échec ???

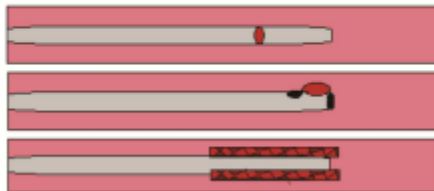
Engainement de fibrine « *fibrin sheath* »



Biofilm: Fibrinogen, Lipoprotéine, Albumine, Facteurs de coagulation Adhésion de leucocytes, microorganismes Puis dépôt de collagène et migration de cellules musculaires lisses

- Ttt endovasculaire/ stripping, rupture
- KDOQI: angioplastie au ballon puis changement du KT sur guide

Intrinsic thrombosis



Intraluminal thrombosis

Catheter tip thrombosis

Fibrin sheath thrombosis

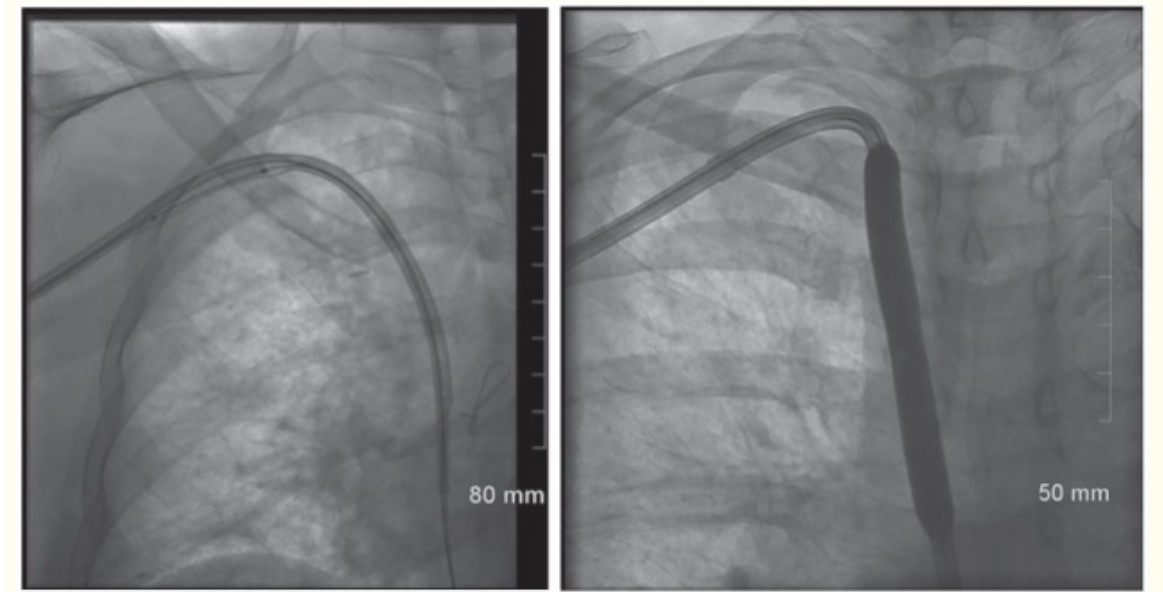
Table 3. Endovascular procedures in the management of catheter dysfunction

| Study (Reference)/ Design | Intervention | Results |
|---|---|--|
| Duszak <i>et al.</i> , 1998 (61) Retrospective (n=68) 56 | <i>De novo</i> catheter placements versus catheter exchange | No significant difference in patency or infections |
| Merport <i>et al.</i> , 2000 (62) Randomized (n=30) 57 | Fibrin sheath stripping versus catheter exchange | Immediate technical success 97%, but primary patency significantly better in the exchange group |
| Janne d'Othée <i>et al.</i> , 2006 (63) Retrospective (n=66) | Fibrin sheath stripping versus catheter exchange versus angioplasty disruption and exchange | All 3 equivalent in terms of technical success, patency, and complications |
| Oliver <i>et al.</i> , 2007 (64) Randomized (n=47) | Catheter exchange versus angioplasty disruption and exchange | Improved patency and statistically significant increased blood flow and urea replacement ratio in the exchange group |

- Nouveau KT =
changement sur guide
- Stripping: bon résultat
initial mais risque de
récidive

- Angioplastie puis
changement KT >
changement KT

Cathéter « incrusté » : *Embadded catheter*



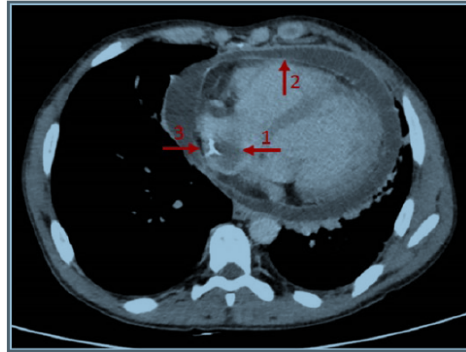
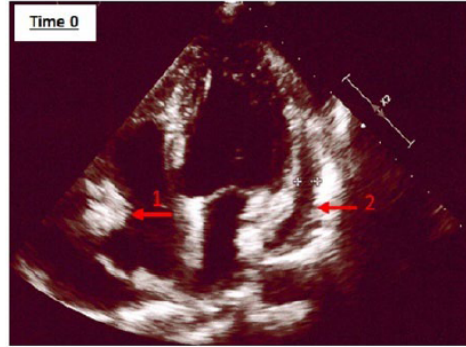
Balloon-assisted endoluminal dilatation for the removal of embedded catheters.

Source. Photo courtesy of Dr Adnan Hadziomerovic, Department of Medical Imaging, The Ottawa Hospital and University of Ottawa.

Thrombus de l'oreillette droite

Catheter-related right atrial thrombosis (CRAT)

- H, 42 ans
- HD/KT tunnelisé
- Fièvre,
- Dysfonction KT
- AVK + thrombolyse loco-régionale: Lock/Urokinase 50 000 ui/ml + Corticoïdes
- Disparition du thrombus à J42
- KT fonctionnel



- 5.4 % à 18% des KT
- Bout de KT /oreillette
- Pronostic vital
- Ttt: Pas de guidelines
 - Ablation du KT
 - Anticoagulation
 - Thrombolyse systémique
 - Thrombectomie chirurgicale

Figure 2. Trans-thoracic echocardiogram 1: right atrial thrombus (2.8 x 2.2 cm)- 2: pericardial effusion - 3: catheter tip placed within the right atrium.

Thrombus de l'oreillette droite
Catheter-related right atrial thrombosis (CRAT)

Thrombus < 6 cm

- Enlever KT
- Anticoagulation

Thrombus > 6 cm

Anticoagulation CI

Anomalies
cardiaques

- Thrombectomie
chirurgicale

Sténoses des veines centrales

Symptômes

Tardifs

Dépendent de la localisation

- Œdème brachial
- Douleurs
- Circulation veineuse collatérale
- Syndrome cave sup
- Rarement: pleurésie
- Dgc: Doppler, Angioscanner ou angioIRM

Traitement

- Sténoses symptomatiques
- Angioplastie +++ svt sans stent
- Chirurgie: si échec ou récurrence < 3 mois

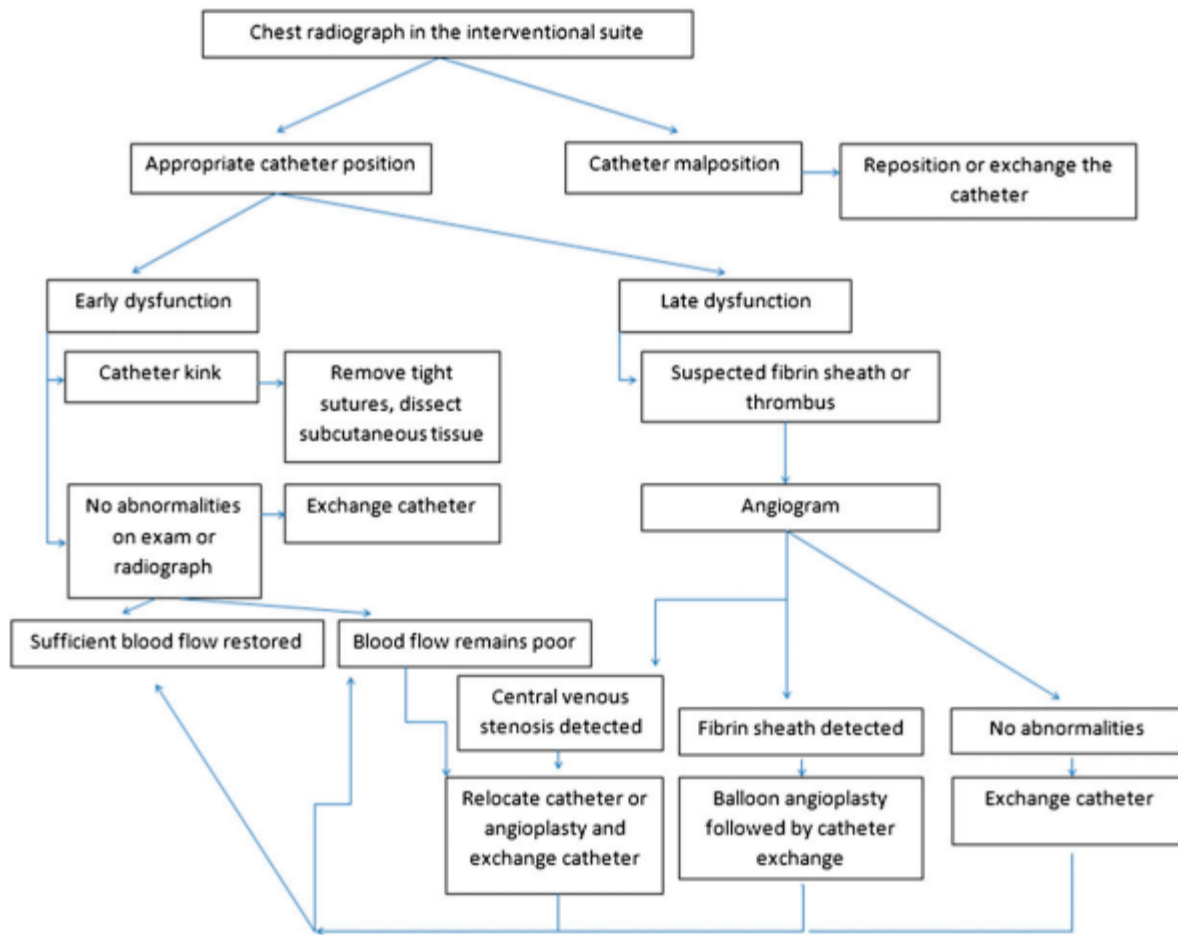


Figure 5. | Algorithm for the endovascular management of dysfunctional catheters in the interventional suite.

Complications Infectieuses

ILC: Infections liées au
Cathéter

*CRI Cathéter-related
infection*



Figure 2. Exit site infection.

infection de
l'orifice
Erythème,
tuméfaction et/ou



Figure 3. Tunnel infection.

Tunnelite
Erythème,
tuméfaction et/ou
douleur > 2cm

bactériémie liée au cathéter
Fièvre > 38°C ou frissons ou hypotension
sans aucune cause évidente

Diagnostic infection liée au cathéter quels prélèvements ?

Cathéter en place

- **Ecouvillonnage site d'insertion**
- Hémocultures standard
- **Hémocultures appariées:**
 - **centrales**
 - **périphériques**



Cathéter enlevé

- **Extrémité distale cathéter**
- Hémocultures (48h)



Hémocultures appariées qualitatives

- Sur KT tunnelisé
 - 1 Hc sur KT > 100 UFC/ml évoque une infection même isolée
- Comparer temps de positivité des hémocult.

Capdevilla, EJCA



Si HC sur KT positive > 2 heures avant HC périph



Bactériémie liée au KT

Se = 91%, Sp = 94% Blot, Lancet 1999



**Treatment Guidelines for Dialysis Catheter–Related Bacteremia:
An Update**

Michael Allon, MD

University of Alabama at Birmingham Birmingham, Alabama

Diagnostic en HD (ambulatoire)

- HC périphérique souvent non réalisable:
 - Épuisement vasculaire
 - Épargne vasculaire
- CEC: pas de différence entre sang périphérique et celui du cathéter
- Centres d'hémodialyse à distance des laboratoires: délai entre prélèvement et incubation variable



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Diagnostic en HD (ambulatoire)

- Limites => exiger moins de preuves
- Définition pratique: HC positive chez un patient symptomatique en absence d'autre source d'infection

Because of these limitations, diagnosis of catheter-related bacteremia in a dialysis outpatient setting may require a lower standard of proof than in hospitalized patients. One practical definition used in some dialysis studies has been the requirement of positive blood cultures (drawn from blood tubing) in a symptomatic patient (fever or chills) in the absence of clinical evidence of an alternate source of infection.⁶⁻⁸ There is no consensus about the

Table 1 Organisms responsible for hemodialysis central venous catheter-related bloodstream infections.

| Organism | Percentage reported* |
|---|----------------------|
| Gram-positive cocci | 52–85% |
| <i>Staphylococcus aureus</i> | 22–60% |
| <i>Staphylococcus epidermidis</i> | 9–13% |
| Meticillin-resistant <i>Staphylococcus aureus</i> | 6–29% |
| <i>Enterococcus faecalis</i> | 2–16% |
| Gram-negative bacilli | 20–28% |
| <i>Pseudomonas aeruginosa</i> | 2–15% |
| <i>Enterobacter cloacae</i> | 9% |
| <i>Escherichia coli</i> | 10% |
| <i>Acinetobacter</i> species | 13% |
| <i>Serratia marcescens</i> | 1–2% |
| <i>Klebsiella pneumoniae</i> | 6% |
| Polymicrobial | 16–20% |
| Acid-fast organisms | Rare |
| Fungi | Rarely reported |

*Percentages do not add up to 100% because data are drawn from different sources.

Traitement empirique

- D.1.8 When haemodialysis catheter infection is suspected, primary antibiotic approach should be inspired by the previously recorded responsible organisms in the unit. If both gram-positive and gram-negative organisms are registered on a regular basis, both types should be covered with eventual refining of the antibiotic regime once the organisms and their sensitivities are known.
- D.1.11 For methicillin-sensitive *S. aureus*, haemodialysis patients should receive cefazolin.
- D.1.10 Vancomycin or teicoplanin is the first choice for empirical therapy of gram positives in settings where MRSA is highly prevalent.

- Inspirée de l'écologie de l'unité d'HD
- Si les CGP et les BGN sont enregistrés avec des prévalences équivalentes, l'antibiothérapie empirique doit couvrir les deux types de spectres par Vancomycine + aminoside ou Vanco + CGIII
- En cas de faible prévalence de staph métiR, la céfazoline est une bonne alternative

Infection de l'orifice sans fièvre

- D.1.7 For exit-site infection without fever, topical antibiotic application can be considered as an alternative. If infection does not resolve, systemic antibiotics should be administered. For tunnel infection without fever, systemic antibiotics are the preferred option, although peroral treatment may be sufficient. If these treatments fail, the catheter should be removed.

- Antibiothérapie locale
- Évolution défavorable: antibiothérapie générale
- Pour la tunnelite:
 - ATB générale,
 - Si l'évolution est défavorable, il faut enlever le cathéter



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Diagnosis, prevention and treatment of haemodialysis catheter-related bloodstream infections (CRBSI): a position statement of European Renal Best Practice (ERBP)

Raymond Vanholder¹, Bernard Canaud², Richard Fluck³, Michel Jadou⁴, Laura Labriola⁴,
A. Marti-Monnes⁵, J. Tordoir⁶ and W. Van Biesen¹

Γ Après culture

- Adaptation de l'antibiothérapie selon l'antibiogramme
- Préférer Céfazoline à la Vancomycine en cas de staph Métis
- Privilégier les antibiotiques dont la pharmacocinétique permet leur administration uniquement après chaque séance d'hémodialyse: vancomycine, céfazoline, aminosides, ceftazidime et daptomycine
- Les BGN sont généralement sensibles aux aminosides et aux CGIII, les CGIII sont préférées pour éviter le risque d'ototoxicité



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Durée du traitement

- Patients non compliqués: 3 semaines
- Localisation secondaire: 6 semaines

Si on garde le cathéter ...

- D.1.6 If a catheter is not removed, blood cultures should be checked 1 week after completion of antibiotic treatment, and if those cultures remain positive, the catheter should be removed.

- HC une semaine après la fin de l'antibiothérapie
- Si HC positive => ablation du cathéter

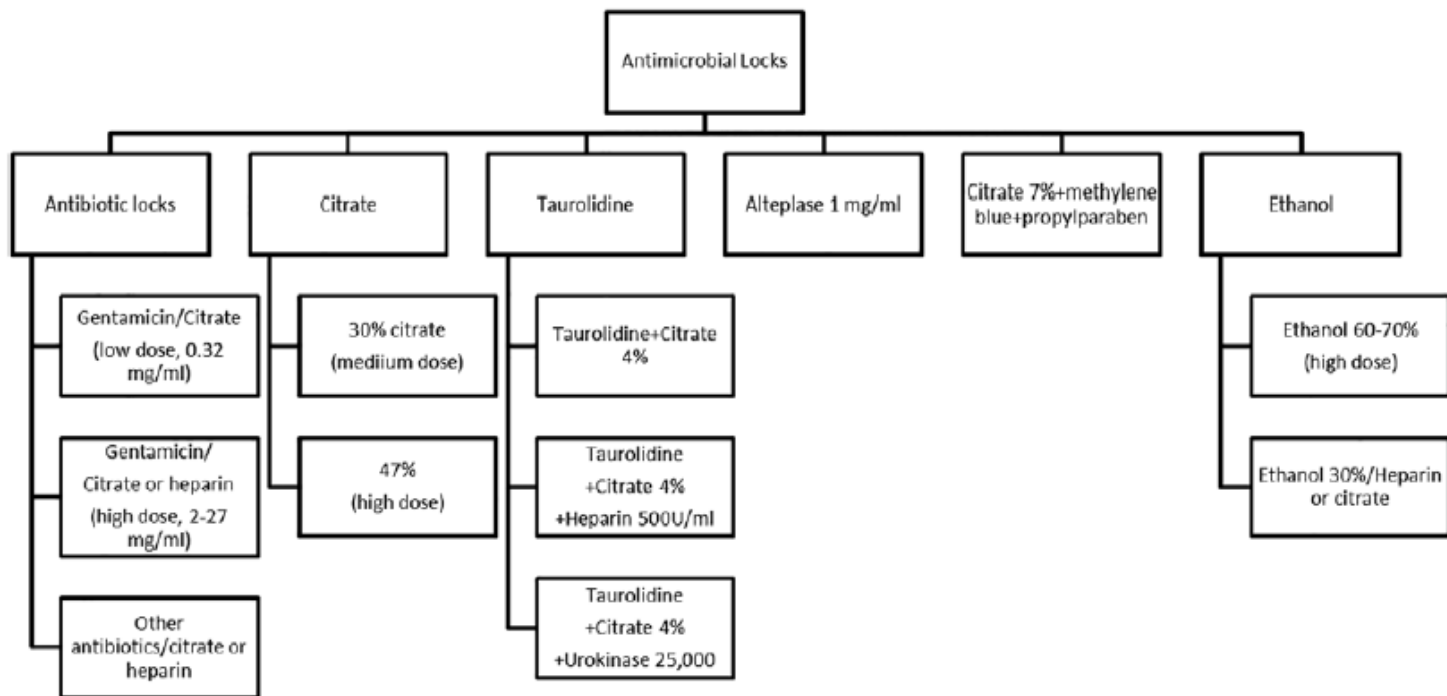


Figure 3 Antimicrobial locks used for the prevention of catheter associated blood stream infection in the hemodialysis catheters.

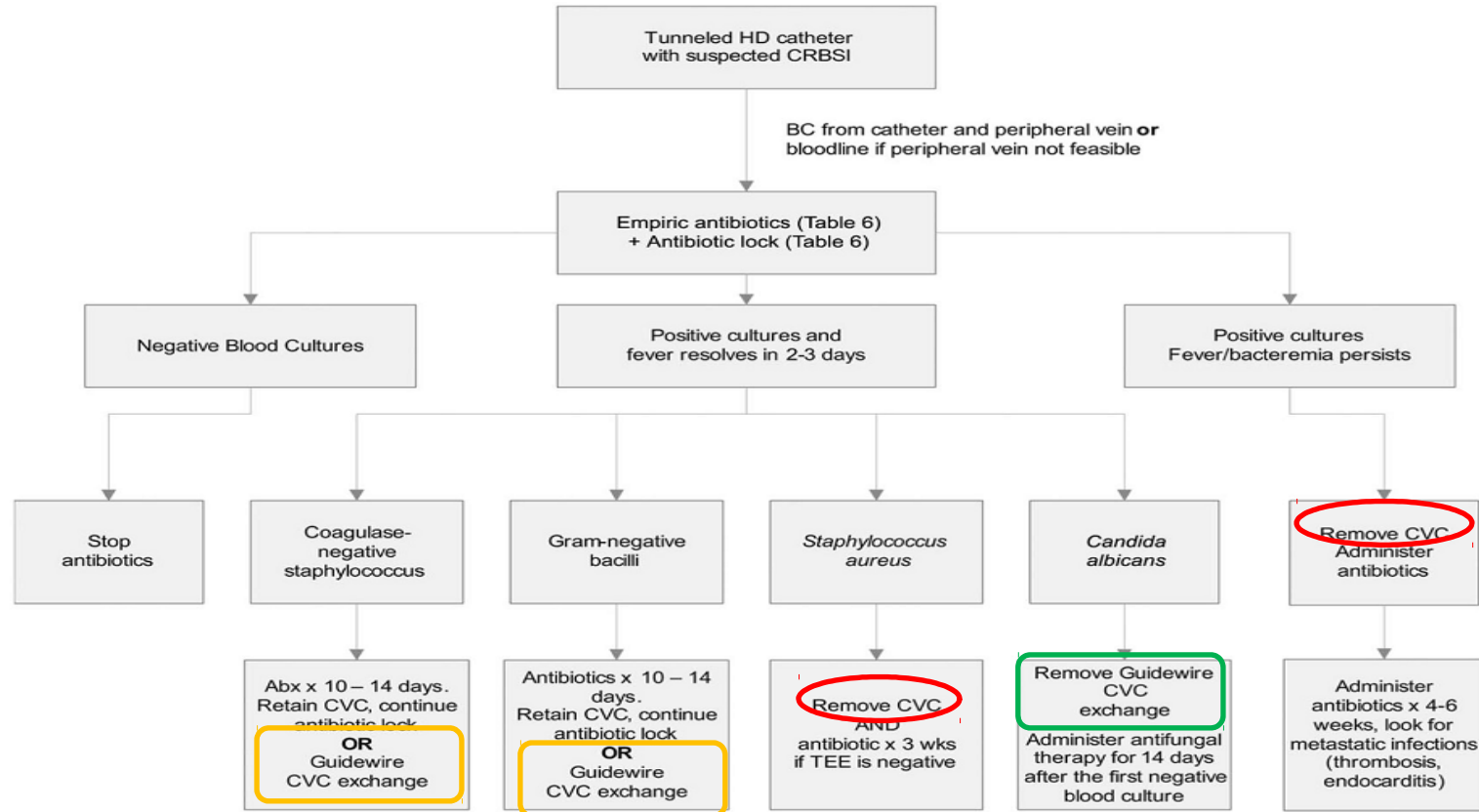
Table 3 | Summary of clinical trials using prophylactic antimicrobial lock (AML) for catheter-related bacteremia (CRB) associated with hemodialysis catheters (CVC)

| First author, year | Subject number | Antimicrobial lock (AML) | Controls (C) (U/ml) | CRB rate AML vs C/1000 CVC days | P-value |
|--------------------------------|----------------|---|---------------------|---------------------------------|---------|
| Pervez, 2002 ⁹⁷ | 36 | Gentamicin 20 mg/ml+citrate 4.67% | Heparin 1000 | 0.62 vs 2.11 | NA |
| Dogra, 2002 ¹⁴² | 83 | Gentamicin 27 mg/ml+citrate 1% | Heparin 5000 | 0.3 vs 4.2 | 0.0003 |
| McIntyre, 2004 ¹⁴³ | 50 | Gentamicin 5 mg/ml+heparin 5000 U/ml | Heparin 5000 | 0.3 vs 4 | 0.02 |
| Nori, 2006 ¹⁴⁴ | 30 | Gentamicin 4 mg/ml+citrate 3.13%; or | Heparin 5000 | 0 vs 4 | 0.008 |
| Venditto, 2010 ¹⁰⁹ | 265 | Gentamicin 2 mg/ml+heparin 5000 U/ml | Heparin 5000 | 0.4 vs 2.9 | 0.06 |
| Onder, 2009 ¹⁴⁵ | 43 | Tobramycin 5 mg/dl+TPA 1 mg/ml | Heparin 5000 | 6.2 vs 16.8 | 0.2 |
| Bleyer, 2005 ¹⁴⁶ | 60 | Minocycline 3 mg/ml+EDTA 30 mg/ml | Heparin (dose NA) | 0 vs 0.47 | 0.35 |
| | 30 | Minocycline 3 mg/ml+EDTA 30 mg/ml | Heparin 5000 | 0.4 vs 4 | 0.02 |
| Saxena, 2005 ¹⁴⁷ | 96 | Cefotaxime 10 mg/ml+heparin 5000 U/ml | Heparin 5000 | 1.65 vs 3.13 | NA |
| Saxena, 2006 ¹⁴⁸ | 113 | Cefotaxime 10 mg/ml+heparin 5000 U/ml | Heparin 5000 | 1.44 vs 3.15 | <0.001 |
| Al-Hwiesh, 2007 ¹⁴⁹ | 63 | Vancomycin 25 mg/ml+gentamicin 40 mg/ml+heparin 5000 U/ml | Heparin 5000 | 4.54 vs 13.11 | 0.05 |
| Kim, 2006 ¹⁵⁰ | 120 | Cefazolin 10 mg/ml+gentamicin 5 mg/ml+heparin 1000 U/ml | Heparin 1000 | 0.44 vs 3.12 | 0.031 |
| Allon, 2003 ³⁵ | 50 | Taurolidine 1.35%+citrate 4% | Heparin 5000 | 0.6 vs 5.9 | <0.001 |
| Betjes, 2004 ¹⁰⁸ | 58 | Taurolidine 1.35%+citrate 4% | Heparin 5000 | 0 vs 2.1 | 0.047 |
| Solomon, 2010 ³³ | 110 | Taurolidine 1.35%+citrate 4% | Heparin 5000 | 1.4 vs 2.4 | 0.1 |
| Weijmer, 2005 ³⁰ | 291 | Trisodium citrate 30% | Heparin 5000 | 1.1 vs 4.1 | <0.001 |
| Winnett, 2008 ¹¹⁰ | 413 | Trisodium citrate 46.7% | Heparin 5000 | 0.81 vs 2.13 | <0.001 |
| Power, 2009 ³¹ | 232 | Trisodium citrate 46.7% | Heparin 5000 | 0.7 vs 0.7 | 0.9 |
| Venditto, 2010 ¹⁰⁹ | 265 | Trisodium citrate 46% | Heparin 5000 | 3.4 vs 2.9 | NS |

Abbreviations: CRB, catheter-related bacteremia; CVC, central venous catheter; EDTA, ethylenediaminetetraacetic acid; NA, not available; NS, non-significant; TPA, tissue plasminogen activator.

Catheter-related blood stream infection (CRBSI) among patients who are undergoing hemodialysis (HD) with tunneled catheters. (IDSA Guidelines 2009)

BC, blood culture; CVC, central venous catheter; TEE, transesophageal echocardiograph.



Lecture



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VAWG Vascular Access Series

Hemodialysis Tunneled Catheter Noninfectious Complications

Lisa M. Miller¹, Jennifer M. MacRae², Mercedesh Kiai³,
Edward Clark⁴, Christine Dipchand⁵, Joanne Kappel⁶,
Charmaine Lok⁷, Rick Luscombe⁸, Louise Moist⁹,
Matthew Oliver¹⁰, Pamela Pike¹¹, and Swapnil Hiremath¹;
on behalf of the Canadian Society of Nephrology Vascular
Access Work Group

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Moving Points in Nephrology

Interventional Nephrology: Catheter Dysfunction— Prevention and Troubleshooting



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Nephrology Dialysis Transplantation

Special Feature

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bloodstream infections (CRBSI): a position statement of European
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