VASCULAR ACCESS

Monitoring and Prevention of CVC Related Infections



Checklists for CVC complications

Monitoring/Surveillance of CVC Complications

At each dialysis session:



Perform a basic medical history focused on signs and symptoms of CVC-related complications (e.g., dysfunction, infection)

Perform a physical examination or check of the dialysis catheter, exit site, tunnel, and surrounding area at each catheter dressing change or dialysis session. CPG 20.1



Checklists for CVC complications – cont.

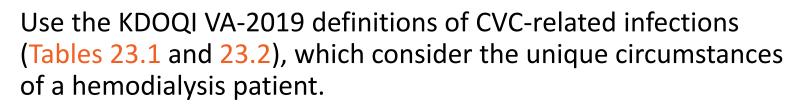
Catheter Dysfunction



Assess for CVC dysfunction at each HD session



Use standardized definitions for CVC-related infections to allow for comparisons across programs/jurisdictions.





Connect the vascular access to circuit properly (see Vascular Access Connection)



Monitor and track vascular access related infections in your unit





General Prevention of CVC Infection and Use of Infection Surveillance Programs and Infection Control Teams

General Checklist: How to monitor and prevent hemodialysis catheter related infections?

CPG 24.1



Educate staff and patients on how to monitor their vascular access (e.g. CVC) for complications



Educate staff and patients on how to prevent vascular access (e.g. CVC) infections



Incorporate an infection control program



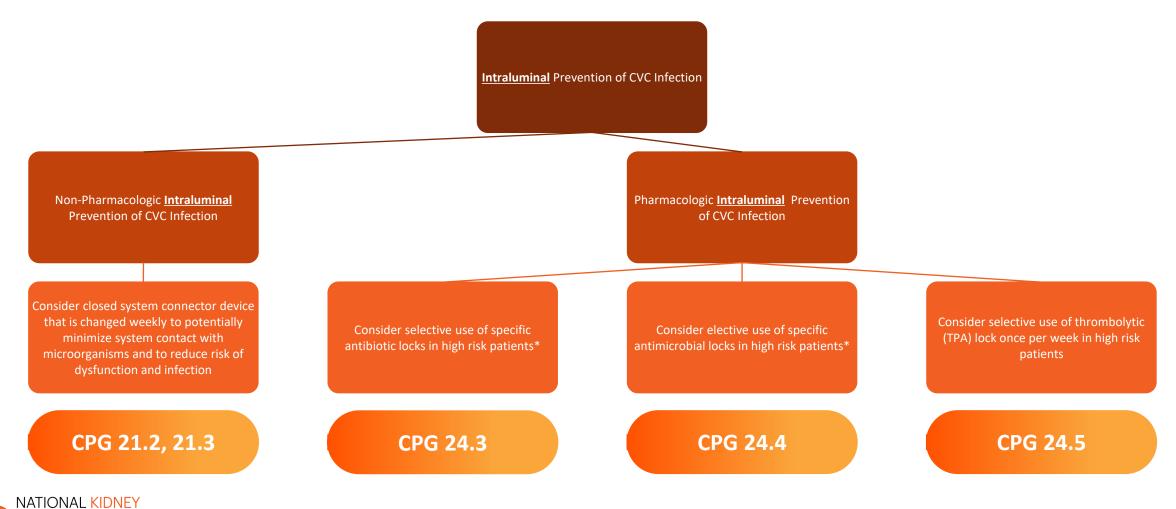
Include an infection surveillance team to monitor, track (in an electronic database), help prevent, and evaluate outcomes of vascular access infections and, in particular, CVC related infections



Flow Diagram 24.a. Prevention of CVC Related Infection

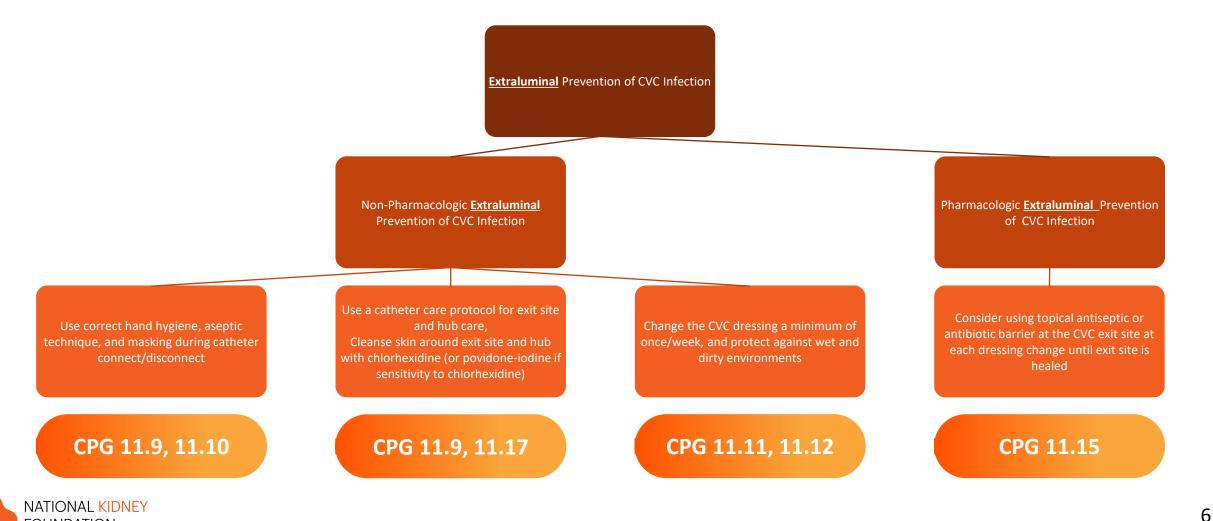
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Note: High risk patients include those with multiple prior catheter related bloodstream infections (CRSBI), especially in facilities with high rates of CRBSI e.g., >3.5/1,000 days.



Flow Diagram 24.b. Prevention of CVC Related Infection – cont.

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(DOQI-2019	KDOQI-2006 ¹³	CDC ²⁹⁷	IDSA ³¹⁹	
Clinical manifestations and at least 1 positive BC from a peripheral source (dialysis circuit or vein) and no other apparent source, with either positive semiquantitative (>15 CFU/ catheter segment, hub or tip) or quantitative (>10 ² CFU/catheter segment, eg, hub or tip) culture, whereby the same organism (species and antibiogram) is isolated from the catheter segment (eg, hub or tip) and a peripheral source (dialysis circuit or vein) blood sample. If available, the following would be supportive: Simultaneous quantitative cultures of blood samples with a ratio of \geq 3:1 (catheter hub/tip vs peripheral [dialysis circuit/vein]); differential period of catheter culture versus peripheral BC positivity of 2 hours.	Definite: Same organism from a semiquantitative culture of the catheter tip (>15 CFU/catheter segment) and from a BC in a symptomatic patient with no other apparent source of infection. Probable: Defervescence of symptoms after antibiotic therapy with or without removal of the catheter, in the setting in which BC confirms infection, but catheter tip does not (or catheter tip does, but blood does not) in a symptomatic patient with no other apparent source of infection. Possible: Defervescence of symptoms after antibiotic treatment or after removal of catheter in the absence of laboratory confirmation of BSI in a symptomatic patient with no other apparent source of infection.	Clinical manifestations and at least 1 positive BC from a peripheral vein and no other apparent source, with either positive semiquantitative (>15 CFU/catheter segment) or quantitative (>10 ² CFU/catheter segment) culture, whereby the same organism (species and antibiogram) is isolated from the catheter segment and a peripheral blood sample. Simultaneous quantitative cultures of blood samples with a ratio of \geq 3:1 (catheter vs peripheral) Differential period of catheter culture versus peripheral BC positivity of 2 hours OR Isolation of the same organism from semiquantitative or quantitative culture segment and from blood (preferably from a peripheral vein) of a patient with accompanying symptoms of BSI and no other apparent source of infection.	Bacteremia/fungemia in a patien with an intravascular catheter wi at least 1 positive BC and with clinical manifestations of infections (ie, fever, chills, and/o hypotension) and no apparent source for the BSI except the catheter AND One of the following shou be present: A positive semiquantitative (>15 CFU/ catheter segment) or quantitative (>10 ² CFU/catheter segment) culture whereby the same organism (species and antibiogram) is isolated from the catheter segment and periphera blood. Simultaneous quantitative BC with a >5:1 ratio catheter versu peripheral. Differential time period of cathete culture versus peripheral BC positivity of >2 hours.	

Table 23.1. Definitions of CVC-Related Blood Stream Infections

Abbreviations: BC, blood culture; BSI, bloodstream infection; CDC, Centers for Disease Control and Prevention; CFU, colony-forming unit; KDOQI, Kidney Disease Outcomes Quality Initiative; IDSA, Infectious Diseases Society of America.



DOQI 2019	KDOQI 2006 ¹³	CDC ²⁹⁷	IDSA ³¹⁹				
	Exit Site	Infection		Tunnel Infection			
Hyperemia, induration, and/ or tenderness ≤2 cm from catheter exit site. May be associated with drainage from the exit site. It may or may not be associated with bacteremia. If there is exit site drainage, it should be collected and sent for Gram staining,	site, not extending superiorly beyond the cuff if the catheter is tunneled, with exudate culture result confirmed to be positive.	Erythema or induration within 2 cm of the catheter exit site, in the absence of concomitant BSI and without concomitant purulence.	cm from catheter exit site.	Tenderness, hyperemia, and/ or induration that extends along the subcutaneous tunnel. It may or may not be associated with bacteremia. If there is drainage, it should be collected and sent for Gram staining, culture, and sensitivities.		Tenderness, erythema, or site induration >2 cm from the catheter site along the subcutaneous tract of a tunneled catheter, in the absence of concomitant BSI.	Tenderness, hyperemia, and or induration that extends > cm from the e site and along the subcutaneous tunnel. It may may not be associated wit bacteremia. If there is purule drainage, it should be collected and sent for Gram staining and culture.

Table 23.2. Definitions of CVC Exit Sites and Tunnel Infections

