## 2023 NHSN Cardiovascular System Infection (CVS) Checklist

Documentation Review Checklist		
CVS - CARDIOVASCULAR SYSTEM INFECTION		
CARD-Myocarditis or pericarditis		
Element	Element Met	Date
Myocarditis or pericarditis must meet at least <u>one</u> of the following criteria:		
<ol> <li>Patient has organism(s) identified from pericardial tissue or fluid by a culture or non- culture based microbiologic testing method, which is performed for purposes of clinica diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST)</li> </ol>		
2. Patient has at least <u>two</u> of the following signs or symptoms:		
• Fever (>38.0°C)		
Chest pain*		
Paradoxical pulse*		
Increased heart size*		
AND at least one of the following:		
a. Abnormal EKG consistent with myocarditis or pericarditis.		
b. Evidence of myocarditis or pericarditis on histologic exam of heart tissue.		
c. 4-fold rise in paired sera from IgG antibody titer.		
d. Pericardial effusion identified by echocardiogram, CT scan, MRI, or angiography.		
3. Patient ≤1 year of age has at least <u>two</u> of the following signs or symptoms:	L.	
• Fever (>38.0°C)		
• Hypothermia (<36.0°C)		
Apnea*		
Bradycardia*		
Paradoxical pulse*		
<ul> <li>Increased heart size*</li> </ul>		
AND at least <u>one</u> of the following:		
a. Abnormal EKG consistent with myocarditis or pericarditis.		
b. Histologic examination of heart tissue shows evidence of myocarditis or pericardi	tis. 🗆	
c. 4-fold rise in paired sera from IgG antibody titer.		
d. Pericardial effusion identified by echocardiogram, CT scan, MRI, or angiography.		
*With no other recognized cause documented by physician		



#### **CVS - CARDIOVASCULAR SYSTEM INFECTION**

#### **ENDO-Endocarditis**

#### Note: When meeting the Endocarditis (ENDO) definition:

- The ENDO Infection Window Period is defined as the 21 days during which all site-specific infection criteria must be met. It includes the date the first positive diagnostic test that is used as an element of the ENDO criterion was obtained, the 10 calendars days before and the 10 calendar days after. The Infection Window Period is lengthened for this event to accommodate the **extended** diagnostic timeframe that is frequently required to reach a clinical determination of endocarditis.
- The RIT for Endocarditis (ENDO) is extended to include the remainder of the patient's current admission.
- When meeting the Endocarditis (ENDO) definition, the secondary BSI attribution period includes the 21-day infection window period and all subsequent days of the patient's current admission.
  - As a result of this lengthy secondary BSI attribution period, secondary BSI pathogen assignment for ENDO is limited to organism(s) identified in blood specimen that match the organism(s) used to meet the ENDO definition.

Element	Element	Date
	Met	
Endocarditis of a natural or prosthetic heart valve must meet at least <u>one</u> of the following criteria:		
<ol> <li>Organism(s) identified from cardiac vegetation*<sup>†</sup>, embolized vegetation (for example, solid-organ abscess) documented as originating from cardiac source, or intracardiac abscess by a culture or non-culture based microbiologic testing method, which is performed for purposes of clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST).</li> </ol>		
<ol> <li>Organism(s) seen on histopathologic examination of cardiac vegetation*, embolized vegetation (for example, solid organ abscess) documented as originating from cardiac source, or intracardiac abscess.</li> </ol>		
<ol> <li>Endocarditis seen on histopathologic examination of cardiac vegetation* or intracardiac abscess.</li> </ol>		
4. At least <u>one</u> of the following echocardiographic evidence of endocarditis* <sup>‡</sup> :		•
i. Vegetation on cardiac valve or supporting structures		
ii. Intracardiac abscess		
iii. New partial dehiscence of prosthetic valve		
AND at least one of the following:		
<ul> <li>a. Typical infectious endocarditis organism(s) (specifically, Viridans group streptococci, Streptococcus bovis, Haemophilus spp., Actinobacillus actinomycetemcomitans, Cardiobacterium hominis, Eikenella corrodens, Kingella spp., Staphylococcus aureus, Enterococcus spp.) identified from ≥2 matching blood collections drawn on separate occasions with no more than 1 calendar day between specimens by a culture or non-culture based microbiologic testing method, which is performed for purposes of clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST).</li> </ul>		
<ul> <li>b. Coxiella burnetii identified by anti-phase I IgG antibody titer &gt;1:800 or identified from blood by a culture or non-culture based microbiologic testing method, which is performed for purposes of clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST).</li> </ul>		



	east <u>three</u> of the following <b>(Note: Meaning one element from i, ii, iii, or iv and only one</b>	condition v	within
-	n element can be used.):	_	
i.	Prior endocarditis, prosthetic valve, uncorrected congenital heart disease, history of		
	rheumatic heart disease, hypertrophic obstructive cardiomyopathy, or known IV		
	drug use <sup>§</sup>		
ii.	Fever (>38.0°C)		
iii.	Vascular phenomena: major arterial emboli (specifically, embolic stroke, renal		
	infarct, splenic infarct or abscess, digital ischemic/gangrene from embolic source),		
	septic pulmonary infarcts, mycotic aneurysm (documented by imaging, seen in		
	surgery, or described in gross pathological specimen), intracranial hemorrhage,		
	conjunctival hemorrhage, or Janeway's lesions documented		
iv.	Immunologic phenomena: glomuleronephritis (documented in chart, or white cell or		
IV.			
	red blood cell casts on urinalysis), Osler's nodes, Roth's spots, or positive		
	rheumatoid factor		
	least <u>one</u> of the following:		
a.	Typical infectious endocarditis organism(s) (specifically, Viridans group streptococci,		
	Streptococcus bovis, Haemophilus spp., Actinobacillus actinomycetemcomitans,		
	Cardiobacterium hominis, Eikenella corrodens, Kingella spp., Staphylococcus aureus,		
	<i>Enterococcus</i> spp.) identified from ≥2 matching blood collections drawn on separate		
	occasions with no more than 1 calendar day between specimens by a culture or		
	non-culture based microbiologic testing method, which is performed for purposes of		
	clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing		
	(ASC/AST).		
h			
b.	, , , , , ,		
	from blood by a culture or non-culture based microbiologic testing method, which is		
	performed for purposes of clinical diagnosis or treatment, for example, not Active		
	Surveillance Culture/Testing (ASC/AST).		
6. At le	east <u>one</u> of the following <sup>*‡</sup>		
i.	Vegetation on cardiac valve or supporting structures seen on echocardiogram		
ii.	Intracardiac abscess seen on echocardiogram		
iii.	New partial dehiscence of prosthetic valve seen on echocardiogram		
AND at	least three of the following (Note: Meaning one element from a, b, c, d, or e and only	one conditi	on within
	ement can be used.):		
	Prior endocarditis, prosthetic valve, uncorrected congenital heart disease, history of		
	rheumatic heart disease, hypertrophic obstructive cardiomyopathy, or known IV		
	drug use <sup>§</sup>		
b.	Fever (>38.0°C)		
	Vascular phenomena: major arterial emboli (specifically, embolic stroke, renal		
C.			
	infarct, splenic infarct or abscess, digital ischemic/gangrene from embolic source),		
	septic pulmonary infarcts, mycotic aneurysm (documented by imaging, seen in		
	surgery, or described in gross pathological specimen), intracranial hemorrhage,		
	conjunctival hemorrhage, or Janeway's lesions documented		
d.	Immunologic phenomena: glomuleronephritis (documented in chart, or white cell or		
	red blood cell casts on urinalysis), Osler's nodes, Roth's spots, or positive		
	rheumatoid factor		
e.	Identification of organism(s) from the blood by at least <u>one</u> of the following		
	methods:		
	<ul> <li>Recognized pathogen(s) identified from blood by a culture or non-culture</li> </ul>		
	based microbiologic testing method, which is performed for purposes of		



clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST).		
<ul> <li>Same common commensal organism(s) identified from ≥2 blood collections drawn on separate occasions on the same or consecutive days by a culture or non-culture based microbiologic testing method, which is performed for purposes of clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST).</li> </ul>		
7. <u>All</u> of the following criteria (Note: Meaning one element from a, b, c, d, or e and only one of	condition w	vithin each
element can be used.):		
<ul> <li>Prior endocarditis, prosthetic valve, uncorrected congenital heart disease, history of rheumatic heart disease, hypertrophic obstructive cardiomyopathy, or known IV drug use<sup>§</sup></li> </ul>		
b. Fever (>38.0°C)		
<ul> <li>c. Vascular phenomena: major arterial emboli (specifically, embolic stroke, renal infarct, splenic infarct or abscess, digital ischemic/gangrene from embolic source), septic pulmonary infarcts, mycotic aneurysm (documented by imaging, seen in surgery, or described in gross pathological specimen), intracranial hemorrhage, conjunctival hemorrhages, or Janeway's lesions documented</li> </ul>		
<ul> <li>Immunologic phenomena: glomuleronephritis (documented on chart, or white cell or red blood cell casts on urinalysis), Osler's nodes, Roth's spots, or positive rheumatoid factor</li> </ul>		
<ul> <li>e. Identification of organism(s) from the blood by at least <u>one</u> of the following methods:</li> <li>Recognized pathogen(s) identified from blood by a culture or non-culture based microbiologic testing method, which is performed for purposes of clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST).</li> <li>Same common commensal organism(s) identified from ≥2 blood collections drawn on separate occasions on the same or consecutive days by a culture or non-culture based microbiologic testing method, which is performed for purposes of clinical diagnosis or treatment, for example, not Active Surveillance Culture based microbiologic testing method, which is performed for purposes of clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST).</li> </ul>		
* Cardiac vegetation can be found on a cardiac valve, pacemaker/defibrillator lead, or ventricular a components within the heart.	ssist device	(VAD)

<sup>†</sup> The following can also meet the definition of a "cardiac vegetation":

• Positive culture from a cardiac valve, pacemaker/defibrillator lead, or ventricular assist device (VAD) components within the heart

<sup>‡</sup> Which if equivocal is supported by clinical correlation (specifically, physician documentation of antimicrobial treatment for endocarditis).

<sup>§</sup> Elements of 5i, 6a, and 7a documented during the current admission:

- May be documented outside of the ENDO infection window period or SSI surveillance period.
- Should not be used to set the ENDO date of event.



CVS - CARDIOVASCULAR SYSTEM INFECTION		
MED-Mediastinitis		
Element	Element Met	Date
Mediastinitis must meet at least one of the following criteria:		
1. Patient has organism(s) identified from mediastinal tissue or fluid by a culture or non-		
culture based microbiologic testing method, which is performed for purposes of clinica		
diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST	).	
2. Patient has evidence of mediastinitis on gross anatomic or histopathologic exam.		
3. Patient has at least <u>one</u> of the following signs or symptoms:		•
• Fever (>38.0°C)		
Chest pain*		
Sternal instability*		
AND at least <u>one</u> of the following:		
a. Purulent drainage from mediastinal area.		
b. Mediastinal widening on imaging test.		
<ol> <li>Patient ≤1 year of age has at least <u>one</u> of the following signs or symptoms:</li> </ol>		
• Fever (>38.0°C)		
• Hypothermia (<36.0°C)		
Apnea*		
Bradycardia*		
Sternal instability*		
AND at least <u>one</u> of the following:		-
a. Purulent drainage from mediastinal area.		
b. Mediastinal widening on imaging test.		
*With no other recognized cause documented by physician	I	1
Comment:		
• The mediastinal space is the area under the sternum and in front of the vertebral colu and its large vessels, trachea, esophagus, thymus, lymph nodes, and other structures a into anterior, middle, posterior, and superior regions.	-	
<ul> <li>Reporting instruction:</li> <li>Report mediastinitis (MED) following cardiac surgery that is accompanied by osteomyothan SSI-BONE.</li> </ul>	elitis as SSI-MED	rather





### **CVS - CARDIOVASCULAR SYSTEM INFECTION**

VASC-Arterial or venous infection, excluding infections involving vascular access devices with organisms identified in the blood

# Note: If a patient meets the criteria for an LCBI in the presence of an arterial or vascular infection (VASC) report as an LCBI not as a VASC.

\*\*Occasionally, a patient with both an eligible central line and another vascular access device will have pus at the other access site. If there is pus at the site of one of the following vascular access devices and a specimen collected from that site has at least one matching organism to an organism identified in the blood during the BSI IWP, report such events marking the "pus at the vascular access site" field as "Yes." Vascular access devices included in this exception are limited to:

- Arterial catheters unless in the pulmonary artery, aorta or umbilical artery
- Arteriovenous fistulae
- Arteriovenous grafts
- Atrial catheters (also known as transthoracic intra-cardiac catheters, those catheters inserted directly into the right or left atrium via the heart wall)
- Hemodialysis reliable outflow (HERO) dialysis catheters
- Intra-aortic balloon pump (IABP) devices
- Non-accessed CL (those neither inserted nor used during current admission)
- Peripheral IV or Midlines

Eleme	nt	Element Met	Date
Arteria	al or venous infection must meet at least <u>one</u> of the following criteria:		
1.	Patient has organism(s) from extracted arteries or veins identified by a culture or non- culture based microbiologic testing method, which is performed for purposes of clinical diagnosis or treatment, for example, not Active Surveillance Culture/Testing (ASC/AST).		
2.	Patient has evidence of arterial or venous infection on gross anatomic or histopathologic exam.		
3.	Patient has at least <u>one</u> of the following signs or symptoms:		
	• Fever (>38.0°C)		
	• Pain*		
	Erythema*		
	Heat at involved vascular site*		
<u>A</u>	<u>ND</u>		
	<ul> <li>More than 15 colonies cultured from intravascular cannula tip using semi- quantitative culture method.</li> </ul>		
4.	Patient has purulent drainage at involved vascular site.		
5.	Patient <b>≤1 year of age</b> has at least <u>one</u> of the following signs or symptoms:		
	• Fever (>38.0°C)		
	Hypothermia (<36.0°C)		
	Apnea*		
	Bradycardia*		
	<ul> <li>Lethargy*</li> </ul>		

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Pain*		
Erythema*		
<ul> <li>Heat at involved vascular site*</li> </ul>		
AND		
<ul> <li>More than 15 colonies cultured from intravascular cannula tip using semi- quantitative culture method.</li> </ul>		
*With no other recognized cause documented by physician		
Reporting instructions:		
<ul> <li>Report infections of an arteriovenous graft, shunt, fistula, or intravascular cannulation site without organism(s) identified from blood as CVS-VASC.</li> </ul>		
<ul> <li>Report Organ Space VASC infections as an SSI and not an LCBI when you have an SSI with secondary BSI.</li> </ul>		

 Report intravascular infections with organism(s) identified from the blood and meeting the LCBI criteria as BSI-LCBI.



