

Ventilator-Associated Event (VAE)

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*required for saving **required for completion

Facility ID:	Event #:
*Patient ID:	Social Security #:
Secondary ID:	Medicare #:
Patient Name, Last:	First: Middle:
*Gender: F M Other	*Date of Birth:
Ethnicity (Specify):	Race (Specify):
*Event Type: VAE	*Date of Event:
Post-procedure VAE: Yes No	Date of Procedure:
NHSN Procedure Code:	ICD-10-PCS or CPT Procedure Code:
*MDRO Infection Surveillance:	
<input type="checkbox"/> Yes, this infection's pathogen & location are in-plan for Infection Surveillance in the MDRO/CDI Module <input type="checkbox"/> No, this infection's pathogen & location are not in-plan for Infection Surveillance in the MDRO/CDI Module	
*Date Admitted to Facility:	*Location:
* Location of Mechanical Ventilation Initiation: _____ *Date Initiated: ___ / ___ / ___ APRV: Yes No	

Event Details

*Specific Event: VAC IVAC PVAP

*Specify Criteria Used:

STEP 1: VAC (≥ 1 REQUIRED)

Daily min FiO_2 increase ≥ 0.20 (20 points) for ≥ 2 days[†] **OR** Daily min PEEP increase $\geq 3 \text{ cm H}_2\text{O}$ for ≥ 2 days[†]
[†]after 2+ days of stable or decreasing daily minimum values.

STEP 2: IVAC

Temperature $> 38^\circ\text{C}$ or $< 36^\circ$ **OR** White blood cell count $\geq 12,000$ or $\leq 4,000 \text{ cells/mm}^3$
AND

A new antimicrobial agent(s) is started, and is continued for ≥ 4 days

STEP 3: PVAP

Criterion #1: Positive culture of one of the following specimens, meeting quantitative or semi-quantitative thresholds as outlined in protocol, [‡] without requirement for purulent respiratory secretions:

<input type="checkbox"/> Endotracheal aspirate	<input type="checkbox"/> Lung tissue
<input type="checkbox"/> Bronchoalveolar lavage	<input type="checkbox"/> Protected specimen brush

OR

Criterion #2: Purulent respiratory secretions[‡] (defined in the protocol) plus organism(s) identified from one of the following specimens:[‡]

<input type="checkbox"/> Sputum	<input type="checkbox"/> Lung tissue
<input type="checkbox"/> Endotracheal aspirate	<input type="checkbox"/> Protected specimen brush
<input type="checkbox"/> Bronchoalveolar lavage	

OR

Criterion #3: One of the following positive tests (as outlined in the protocol):[‡]

<input type="checkbox"/> Organism(s) identified from pleural fluid	<input type="checkbox"/> Diagnostic test for <i>Legionella</i> species
<input type="checkbox"/> Lung histopathology	<input type="checkbox"/> Diagnostic test for selected viral pathogens

[†]collected after 2 days of mechanical ventilation and within +/- 2 days of onset of increase in FiO_2 or PEEP.

*Secondary Bloodstream Infection: Yes No

**Died: Yes No

VAE Contributed to Death: Yes No

Discharge Date:

*Pathogens Identified: Yes No *If Yes, specify on pages 2-3

Assurance of Confidentiality: The voluntarily provided information obtained in this surveillance system that would permit identification of any individual or institution is collected with a guarantee that it will be held in strict confidence, will be used only for the purposes stated, and will not otherwise be disclosed or released without the consent of the individual, or the institution in accordance with Sections 304, 306 and 308(d) of the Public Health Service Act (42 USC 242b, 242k, and 242m(d)).

Public reporting burden of this collection of information is estimated to average 28 minutes per response, including the time for reviewing instructions, searching existing data sources, gathering and maintaining the data needed, and completing and reviewing the collection of information. An agency may not conduct or sponsor, and a person is not required to respond to a collection of information unless it displays a currently valid OMB control number. Send comments regarding this burden estimate or any other aspect of this collection of information, including suggestions for reducing this burden to CDC, Reports Clearance Officer, 1600 Clifton Rd., MS D-74, Atlanta, GA 30333, ATTN: PRA (0920-0666).

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Pathogen #	Gram-positive Organisms								
_____	<i>Staphylococcus coagulase-negative</i> VANC SIRN (specify species if available):								
_____	<i>Enterococcus faecium</i>		DAPTO S NS N	GENTHL [§] SRN	LNZ SIRN	VANC SIRN			
	<i>Enterococcus faecalis</i>								
	<i>Enterococcus</i> spp. (Only those not identified to the species level)								
_____	<i>Staphylococcus aureus</i>	CIPRO/LEVO/MOXI SIRN	CLIND SIRN	DAPTO S NS N	DOXY/MINO SIRN	ERYTH SIRN	GENT SIRN	LNZ SRN	
		OX/CEFOX/METH SIRN	RIF SIRN	TETRA SIRN	TIG S NS N	TMZ SIRN	VANC SIRN		
Pathogen #	Gram-negative Organisms								
_____	<i>Acinetobacter</i> (specify species)	AMK SIRN	AMPSUL SIRN	AZT SIRN	CEFEP SIRN	CEFTAZ SIRN	CIPRO/LEVO SIRN	COL/PB SIRN	
		GENT SIRN	IMI SIRN	MERO/DORI SIRN	PIP/PIPTAZ SIRN		TETRA/DOXY/MINO SIRN		
		TMZ SIRN	TOBRA SIRN						
_____	<i>Escherichia coli</i>	AMK SIRN	AMP SIRN	AMPSUL/AMXCLV SIRN	AZT SIRN	CEFAZ SIRN	CEFEP S I/S-DD R N	CEFOT/CEFTRX SIRN	
		CEFTAZ SIRN	CEFUR SIRN	CEFOX/CETET SIRN	CIPRO/LEVO/MOXI SIRN		COL/PB [†] SRN		
		ERTA SIRN	GENT SIRN	IMI SIRN	MERO/DORI SIRN	PIPTAZ SIRN	TETRA/DOXY/MINO SIRN		
		TIG SIRN	TMZ SIRN	TOBRA SIRN					
_____	<i>Enterobacter</i> (specify species)	AMK SIRN	AMP SIRN	AMPSUL/AMXCLV SIRN	AZT SIRN	CEFAZ SIRN	CEFEP S I/S-DD R N	CEFOT/CEFTRX SIRN	
		CEFTAZ SIRN	CEFUR SIRN	CEFOX/CETET SIRN	CIPRO/LEVO/MOXI SIRN		COL/PB [†] SRN		
		ERTA SIRN	GENT SIRN	IMI SIRN	MERO/DORI SIRN	PIPTAZ SIRN	TETRA/DOXY/MINO SIRN		
		TIG SIRN	TMZ SIRN	TOBRA SIRN					
_____	<i>Klebsiella pneumonia</i>	AMK SIRN	AMP SIRN	AMPSUL/AMXCLV SIRN	AZT SIRN	CEFAZ SIRN	CEFEP S I/S-DD R N	CEFOT/CEFTRX SIRN	
		CEFTAZ SIRN	CEFUR SIRN	CEFOX/CETET SIRN	CIPRO/LEVO/MOXI SIRN		COL/PB [†] SRN		
		ERTA SIRN	GENT SIRN	IMI SIRN	MERO/DORI SIRN	PIPTAZ SIRN	TETRA/DOXY/MINO SIRN		
		TIG SIRN	TMZ SIRN	TOBRA SIRN					

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Pathogen #	Gram-negative Organisms (continued)									
	<i>Pseudomonas aeruginosa</i>	AMK SIR N	AZT SIR N	CEFEP SIR N	CEFTAZ SIR N	CIPRO/LEVO SIR N	COL/PB SIR N	GENT SIR N		
		IMI SIR N	MERO/DORI SIR N		PIP/PIPTAZ SIR N	TOBRA SIR N				
Pathogen #	Fungal Organisms									
	<i>Candida</i> (specify species if available)	ANID SIR N	CASPO S NS N	FLUCO S S-DD R N	FLUCY SIR N	ITRA S S-DD R N	MICA S NS N	VORI S S-DD R N		
Pathogen #	Other Organisms									
	Organism 1 (specify)	Drug 1 SIR N	Drug 2 SIR N	Drug 3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 7 SIR N	Drug 8 SIR N	Drug 9 SIR N
	Organism 1 (specify)	Drug 1 SIR N	Drug 2 SIR N	Drug 3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 7 SIR N	Drug 8 SIR N	Drug 9 SIR N
	Organism 1 (specify)	Drug 1 SIR N	Drug 2 SIR N	Drug 3 SIR N	Drug 4 SIR N	Drug 5 SIR N	Drug 6 SIR N	Drug 7 SIR N	Drug 8 SIR N	Drug 9 SIR N

Result Codes

S = Susceptible I = Intermediate R = Resistant NS = Non-susceptible S-DD = Susceptible-dose dependent N = Not tested

[§] GENTHL results: S = Susceptible/Synergistic and R = Resistant/Not Synergistic

[†] Clinical breakpoints have not been set by FDA or CLSI, Sensitive and Resistant designations should be based upon epidemiological cutoffs of Sensitive MIC ≤ 2 and Resistant MIC ≥ 4

Drug Codes:

AMK = amikacin	CEFTRX = ceftriaxone	FLUCY = flucytosine	OX = oxacillin
AMP = ampicillin	CEFUR= cefuroxime	GENT = gentamicin	PB = polymyxin B
AMPSUL = ampicillin/sulbactam	CETET= cefotetan	GENTHL = gentamicin –high level test	PIP = piperacillin
AMXCLV = amoxicillin/clavulanic acid	CIPRO = ciprofloxacin	IMI = imipenem	PIPTAZ = piperacillin/tazobactam
ANID = anidulafungin	CLIND = clindamycin	ITRA = itraconazole	RIF = rifampin
AZT = aztreonam	COL = colistin	LEVO = levofloxacin	TETRA = tetracycline
CASPO = caspofungin	DAPTO = daptomycin	LNZ = linezolid	TIG = tigecycline
CEFAZ= cefazolin	DORI = doripenem	MERO = meropenem	TMZ = trimethoprim/sulfamethoxazole
CEFEP = cefepime	DOXY = doxycycline	METH = methicillin	TOBRA = tobramycin
CEFOT = cefotaxime	ERTA = ertapenem	MICA = micafungin	VANC = vancomycin
CEFOX= cefoxitin	ERYTH = erythromycin	MINO = minocycline	VORI = voriconazole
CEFTAZ = ceftazidime	FLUCO = fluconazole	MOXI = moxifloxacin	

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Custom Fields

Label	/ /	Label	/ /

Comments

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