

Pediatric Ventilator-associated Event (PedVAE)

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2022 NHSN Training

Training Objectives

- Explain the history and development of PedVAE
- Review PedVAE key terms
- Explain how to Select Daily Minimum Values
- Describe the PedVAE Surveillance Algorithm
- Demonstrate use of the PedVAE Calculator



Ventilated Patients and the Need for Surveillance

- Ventilated patients are at high risk for complications and poor outcomes
 - Ventilator-associated pneumonia (VAP), sepsis, Acute Respiratory Distress
 Syndrome (ARDS), pulmonary embolism, barotrauma, and pulmonary edema
- Such complications can lead to longer duration of mechanical ventilation, longer stays in the ICU and hospital, increased healthcare costs, and increased risk of disability and death
- In preterm neonates, prolonged mechanical ventilation for respiratory distress syndrome can contribute to the development of chronic lung disease
- Prolonged mechanical ventilation in extremely low birthweight infants is also associated with neurodevelopmental delay

PedVAE Surveillance: Development

- Neonatal & Pediatric VAE Surveillance Working Group convened in 2012 to explore use of VAE (adult algorithm) in pediatric and neonatal inpatient locations
 - Insufficient data to use the same approach as used for adults

PedVAE Surveillance: Development

Publication* in 2016 on the use of a pediatric VAE-like definition demonstrated detection of events defined by changes in FiO₂ and Mean Airway Pressure were associated with increases in length of stay and mortality

Ventilator-Associated Events in Neonates and Children—A New Paradigm*

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- This research was standed by the Agency for Healthcare Research and Quality (AHRQ) R18 grant (#1R18HS0216SR) to Dt. Lee. Dr. Gray has received sunding from the Vermont Orders Network. Dr. Logan also receives from the National Institutes of Health (NIH) (BK08A11250S-01; not related to this grant).
- DC Cocored Institution received funding from the Agency for Healthcare Research and Clustilly (AHRCI). Dr. Kleinman's Institution received funding strem AHRCI. Dr. Peteor's Institution received funding from AHRCI. Dr. Glasy received funding from the Verment Clettor Notwork. His Institucen received funding from the Verment Clettor Notwork - His Institucen received funding from AHRCI. Dr. II your received support for article

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Objectives: To identify a pediatric ventilator-associated condition definition for use in neonates and children by exploring whether potential ventilator-associated condition definitions identify a patients with worse outcomes.

Design: Retrospective cohort study and a matched cohort analysis.

Setting: Pediatric, cardiac, and neonatal ICUs in five U.S. hospitals.

Patients: Children 18 years old or younger ventilated for at least

Interventions: None

Measurements and Main Results: We evaluated the evidence of worsering oxygenation via a range of thresholds for increases in daily minimum traction of inspired oxygen (by 0.20, 0.25, and 0.30), and daily minimum mean airway pressure (by 4.5, 6, and 7 cm H₂O). We required worsering oxygenation be sustained for at least 2 days after at least 2 days of stability. We matched patients with a verifiator-associated condition to those without and used Oxygenoportional

^{*}Cocoros NM, Kleinman K, Priebe GP, et al. Ventilator-Associated Events in Neonates and Children--A New Paradigm. Crit Care Med. 2016 Jan;44:14-22.

PedVAE Surveillance: Available as of 2019

- Neonatal & Pediatric VAE Surveillance
 Working Group consensus reached to
 begin development with a plan to
 implement PedVAE as an available event in
 NHSN
- PedVAE field testing conducted in 2017
- PedVAE available as an NHSN event beginning January 2019



Pediatric Ventilator-Associated Event (PedVAE)

For use in neonatal and pediatric locations only

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Introduction

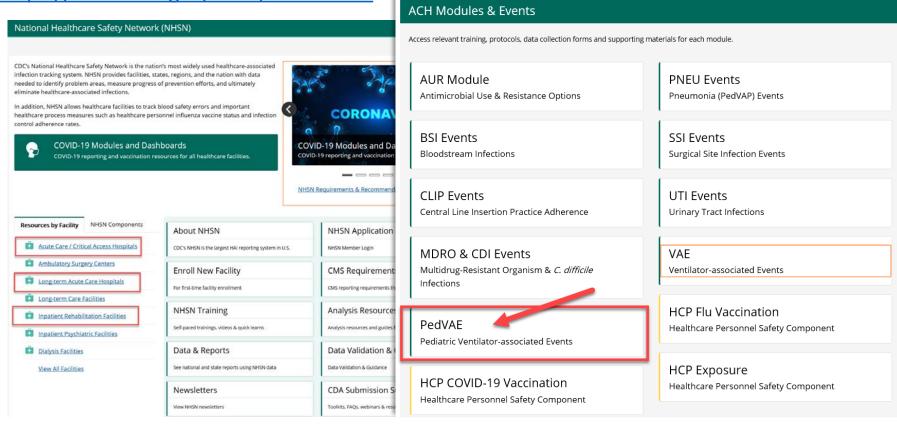
Mechanical ventilation is an essential, life-saving therapy for patients with critical illness and respiratory failure. Hundreds of thousands of patients receive mechanical ventilation in the United States each year [1-3]. These patients are at high risk for complications and poor outcomes, including death [1-5]. Ventilator-associated pneumonia (VAP), sepsis, Acute Respiratory Distress Syndrome (ARDS), pulmonary embolism, barotrauma, and pulmonary edema are among the complications that can occur in patients receiving mechanical ventilation. Such complications can lead to longer duration of mechanical ventilation, longer stays in the ICU and hospital, increased healthcare costs, and increased risk of disability and death. In preterm neonates, prolonged mechanical ventilation for respiratory distress syndrome can contribute to the development of chronic lung disease [6]. Prolonged mechanical ventilation in extremely low birthweight Infants is also associated with neurodevelopmental delay [7].

Surveillance for ventilator-associated events in the National Healthcare Safety Network (NHSN) prior to 2013 was limited to VAP. Traditional VAP definitions, including the NHSN PNEU definitions (revised in 2002), have well-described limitations [8-11]. They typically require radiographic evidence of pneumonia, although data suggest that chest radiograph findings do not accurately identify VAP. The subjectivity and variability inherent in chest radiograph technique, interpretation, and reporting make chest imaging ill-suited for inclusion in a definition algorithm to be used for the potential purposes of public reporting, inter-facility comparisons, and pay-for-reporting and pay-for-performance programs. Another major limitation of the available VAP definitions is their reliance on specific clinical signs or symptoms, which are subjective and may be poorly or inconsistently documented in the medical record.

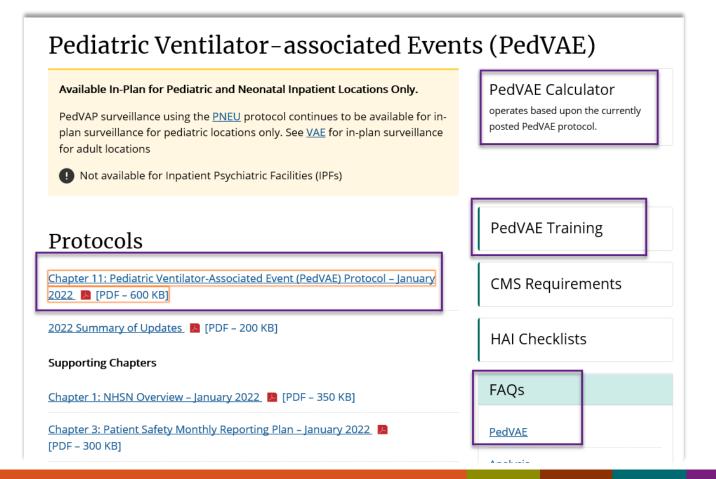
PedVAE Surveillance – Before you begin

Where do I find the PedVAE surveillance guidance?

https://www.cdc.gov/nhsn/index.html



https://www.cdc.gov/nhsn/psc/pedvae/index.html



Who is eligible for PedVAE surveillance?

- Ventilated inpatients of acute care hospitals, long term acute care hospitals, inpatient rehabilitation facilities
- Patients in pediatric and neonatal locations only where denominator data (patient days and ventilator days) can be collected
 - Ventilated adults in pediatric locations are included in PedVAE surveillance – regardless of age



Who is NOT eligible for PedVAE surveillance?

- Patients on extracorporeal life support or paracorporeal membrane oxygenation are <u>not eligible</u> for VAE surveillance
 - Ineligibility only applies to periods of time while receiving this form of support
- Non-acute care locations in acute care facilities are not eligible to participate in PedVAE surveillance.

What about other specific modes of mechanical ventilation?

- INCLUDE patients on:
 - High Frequency Oscillatory or jet ventilation (HFO)
 - Airway Pressure Release Ventilation (APRV)
 - Volumetric Diffusive Respiration (VDR) FiO₂ parameter only
- INCLUDE patients who are receiving a conventional mode of mechanical ventilation while receiving:
 - Surfactant
 - Corticosteroids
 - Prone positioning
 - Nitric oxide therapy
 - Helium-oxygen mixture
 - Epoprostenol therapy

PedVAE Algorithm Overview

Remember: The PedVAE definition algorithm is for use in surveillance; it is not a clinical definition algorithm and is not intended for use in the clinical management of patients.

PedVAE Definition Algorithm Summary

Patient on mechanical ventilation > 2 days



Baseline period of stability or improvement, followed by sustained period of worsening oxygenation



Pediatric Ventilator-Associated Event (PedVAE)

PedVAEs are determined by identification of deterioration in respiratory status after a period of stability or improvement on the ventilator

- Using either of two key parameters that demonstrate effective oxygenation in ventilated patients
 - FiO₂
 - Mean Airway Pressure (MAP)



FiO₂

- Fraction of Oxygen in inspired gas
 - FiO₂ of room air is 0.21
 - Oxygen concentration of room air is 21%
- FiO₂ is a setting on the ventilator and is one of the key parameters that can be adjusted depending on the patient's oxygenation requirement

MAP

- Mean Airway Pressure
 — Mean (average) pressure exerted on the airway and lungs from the beginning of inspiration until the beginning of the next inspiration (inspiratory cycle)
- MAP is a measured/calculated value (not a ventilator setting) that is determined by
 - PEEP Peak End-Expiratory Pressure
 - PIP- Peak Inspiratory Pressure
 - Inspiratory time
 - Other parameters like flow or frequency
- MAP for purposes of PedVAE surveillance is <u>NOT Mean Arterial</u> <u>Pressure</u>

FiO₂ and MAP

- FiO₂ ventilator settings and MAP values documented across the calendar day are used to identify the <u>daily minimum</u> FiO₂ and <u>daily minimum</u> MAP values
- The daily minimum FiO₂ and daily minimum MAP values are used to determine both the period of stability or improvement and the period that indicates worsening oxygenation.
- Stability, improvement or worsening is not determined by comparing FiO₂ settings and MAP values that occur <u>during</u> a calendar day but by comparing the daily minimum values from calendar day to calendar day
- Use a calendar day not any other "24-hour capture period"

Daily Minimum FiO₂ and MAP

Daily Minimum FiO₂ and MAP

- FiO₂ settings and MAP readings are typically recorded in the paper or electronic medical record, on respiratory therapy and/or nursing flow sheets, in the section of the flow sheet that pertains to respiratory status/mechanical ventilation
- When choosing the daily minimum FiO₂ and MAP, use all documented findings that are recorded throughout the calendar day during times when the patient is <u>receiving support from an eligible mode of mechanical ventilation</u>
 - Include FiO₂ and MAP values documented during weaning/mechanical ventilation
 liberation trials as long as the patient is receiving ventilator support during those trials
 - Excludes FiO₂ and MAP values documented during periods of time when the patient is on extracorporeal or paracorporeal membrane oxygenation

Daily Minimum FiO₂

- The daily minimum FiO₂ is defined as the lowest documented FiO₂ setting that was maintained for > 1 hour during a calendar day
- Protocol provides examples of how > 1 hour is to be determined to ensure standardization across all facilities
 - If tracking every 15 minutes, 5 consecutive recordings of a certain level would be needed (e.g., at 09:00, 09:15, 09:30, 09:45 and 10:00)
 - If tracking every 30 minutes, 3 consecutive recordings at a certain level would be needed (e.g., at 09:00, 09:30, and 10:00)
 - If tracking PEEP every hour, 2 consecutive recordings at a certain level (e.g., at 09:00 and 10:00)
- If there is no setting that has been maintained for > 1 hour then select the lowest setting regardless of the period of time in which the setting was maintained

Identify the Daily Minimum FiO₂ for Monday

Select the lowest value recorded for the calendar day that is maintained for >1 hour

	Monday 12am	3am	4am	6am	9am	12pm	3pm	9pm	11pm
FiO ₂	0.80	0.70	0.90	0.80	0.80	0.75	0.75	0.75	0.75

Identify the Daily Minimum FiO₂

Select the lowest value recorded for the calendar day that is maintained for >1 hour

	Monday 12am	3am	4am	6am 	9am	12pm	3pm	9pm	11pm
FiO ₂	0.80	0.70	0.90	0.80	0.80	0.75	0.75	0.75	0.75

For Monday: 0.75 is the Daily Minimum FiO₂. The lowest value of 0.70 was not maintained for > 1 hour

Identify the Daily Minimum FiO₂ for Monday and Tuesday

Ventilation is initiated late in the calendar day

	Monday 2300	2330	Tuesday 0030	0100	0300	0600	0900	1200
FiO ₂	0.70	0.80	0.80	0.80	0.80	0.75	0.75	0.80

Identify the Daily Minimum FiO₂ for Monday and Tuesday

Ventilation is initiated late in the calendar day

 SS	Monday 2300	2330	Tuesday 0030	0100	0300	0600	0900	1200
FiO ₂	0.70	0.80	0.80	0.80	0.80	0.75	0.75	0.80

- Monday: 0.70 is the Daily Minimum FiO₂, there was no value maintained for > 1 hour
- Do not look to the next calendar day to determine if a setting was maintained > 1 hour
- Tuesday: 0.75 is the Daily Minimum FiO₂, the lowest value maintained for > 1 hour

Daily Minimum MAP

 The daily minimum MAP is the lowest value documented during a calendar day regardless of how long the value is maintained

- When determining the daily minimum MAP, if MAP values include a decimal place, then round the MAP value to the nearest whole number. For example:
 - A MAP of 10.00 10.49 is rounded to 10
 - A MAP of 10.50 10.99 is rounded to 11

Daily Minimum MAP

- For <u>patients < 30 days</u> MAP values of 0-8 cmH₂O are considered equal to 8 cmH₂O
 - Any day where daily minimum MAP is 0-8 cmH₂0 will be assigned a daily minimum MAP value of 8 cmH₂0.
- For <u>patients ≥ 30 days</u> MAP values 0-10 cmH₂O are considered equal to 10 cmH₂O
 - Any day where daily minimum MAP is 0-10 cmH₂0 will be assigned a daily minimum MAP value of 10 cmH₂0.

Identify the Daily Minimum MAP for a Patient < 30 Days

	Monday 12am	3am	6am	9am	12pm	3pm	6pm	9pm
MAP	8	6	8	5	5	8	10	10

Identify the Daily Minimum MAP for a Patient < 30 Days

Select the lowest value recorded for each calendar day

	Monday 12am	3am	6am	9am	12pm	3pm	6pm	9pm
MAP	8	6	8	5(8)	5	8	10	10

Explanation: The lowest value is $5 \text{ cmH}_2\text{O}$ on Monday, but remember that for patients < 30 days, values 0-8=8.

Monday: Daily Minimum MAP is 8 cmH₂O

Identify the Daily Minimum MAP for a Patient ≥ 30 Days

	Monday 12am		6am	9am	12pm	3pm	6pm	9pm
MAP	8	6	8	5	5	8	10	10

Identify the Daily Minimum MAP for a Patient ≥ 30 Days

Select the lowest value recorded for each calendar day regardless of how long it was maintained

	Monday 12am	3am	6am	9am	12pm	3pm	6pm	9pm
MAP	8	6	8	5(10)	5	8	10	10

Explanation: The lowest value is $5 \text{ cmH}_2\text{O}$ on Monday, but remember that for patients $\geq 30 \text{ days}$, values 0-10 = 10.

Monday: Daily Minimum MAP is 10 cmH₂O

PedVAE Surveillance

Figure 1: Pediatric Ventilator-Associated Events (PedVAE) Surveillance Algorithm

Patient has a baseline period of stability or improvement on the ventilator, defined by ≥ 2 calendar days of stable or decreasing daily minimum* FiO₂ or MAP values. The baseline period is defined as the 2 calendar days immediately preceding the first day of increased daily minimum MAP or FiO₂.

*Daily minimum FiO₂ is defined as the lowest value of FiO₂ documented during a calendar day that is maintained for > 1 hour. Daily minimum MAP is the lowest value documented during the calendar day.

For patients < 30 days old, daily minimum MAP values 0-8 cm H₂O are considered equal to 8 cmH₂O for the purposes of surveillance.

For patients ≥ 30 days old, daily minimum MAP values 0-10 cmH₂O are considered equal to 10 cmH₂O for the purposes of surveillance.



After a period of stability or improvement on the ventilator, the patient has at least one of the following indicators of worsening oxygenation:

- 1) Increase in daily minimum FiO_2 of ≥ 0.25 (25 points) over the daily minimum FiO_2 of the first day in the baseline period, sustained for ≥ 2 calendar days.
- 2) Increase in daily minimum MAP values of ≥ 4 cmH₂O over the daily minimum MAP of the first day in the baseline period, sustained for ≥ 2 calendar days.



Pediatric Ventilator-Associated Event (PedVAE)

Meeting PedVAE

- Patient must be ventilated > 2 days to be eligible for PedVAE surveillance
 - However, the first two days of mechanical ventilation can establish a baseline period
- Patients must be mechanically ventilated for at least 4 calendar days to fulfill PedVAE criteria (where the day of intubation or initiation of mechanical ventilation is day 1)
 - 2 days of stability or improvement
 - 2 days of evidence of worsening oxygenation

What do we mean when we use "Baseline" and "Worsening" in the PedVAE definition?

- Baseline: ≥ 2 calendar days of stable or decreasing daily minimum FiO2 or MAP values and immediately precedes the first day of increased daily minimum MAP or FiO2.
- Worsening: After a period of stability or improvement on the ventilator, the patient has at least one of the following indicators of worsening oxygenation:
 - Increase in daily minimum FiO2 of ≥ 0.25 (25 points) over the daily minimum FiO2 of the first day in the baseline period, sustained for ≥ 2 calendar days.

OR

 Increase in daily minimum MAP values of ≥ 4 cmH2O over the daily minimum MAP of the first day in the baseline period, sustained for ≥ 2 calendar days.

Determining if a PedVAE is identified using the Daily Minimum FiO2 and MAP Values

FiO2: a baseline period of stability or improvement immediately followed by an increase in daily minimum FiO_2 of \geq 0.25 (25 points) over the daily minimum FiO_2 of the first day in the baseline period, sustained for \geq 2 calendar days

MAP: a baseline period of stability or improvement immediately followed by an increase in daily minimum MAP values of ≥ 4 cmH₂O over the daily minimum MAP of the first day in the baseline period, sustained for ≥ 2 calendar days

Operationalizing PedVAE

(patient is <30 days - MAP values 0-8 = 8)

First, determine if a baseline period of stability or improvement is established in either the Daily Minimum MAP or Daily minimum FiO₂ parameter.

Vent Day	Daily Minimum MAP	Daily Minimum FiO ₂
1	13	60
2	8 (7)	40
3	8 (7)	40
4	12	65
5	12	50
6	10	40
7	8	40
8	8	40

Operationalizing PedVAE

(patient is < 30 days)

Vent Day	Daily Minimum MAP	Daily Minimum FiO ₂
1	13	60
2	8 (7)	40
3	8 (7)	40
4	12	65
5	12	50
6	10	40
7	8	40
8	8	40

Operationalizing PedVAE

(patient is < 30 days)

	Vent Day	Daily Minimum MAP	Daily Minimum FiO ₂	
	1	13	60	
	2	8 (7)	40	l
	3	8 (7)	40	≥ 2-day period of
	4	12	65	stability
	5	12	50	(MAP or FiO ₂)
	d of worsening in ameter with an	10	40	
	cm H ₂ O over the	8	40	
first day of the	e baseline period	8	40	

A PedVAE is identified in the MAP parameter

	Vent Day	Daily Minimum MAP	Daily Minimum FiO ₂	
	1	13	60	
	2	8 (7)	40	1
	3	8 (7)	40	≥ 2-day period of
	4	12	65	stability
	5	12	50	(MAP or FiO ₂)
	d of worsening in ameter with an	10	40	
	cm H ₂ O over the	8	40	
first day of the	e baseline period	8	40	

What if we take the same example in a patient ≥ 30 days?

Vent Day	Daily Minimum MAP	Daily Minimum FiO ₂
1	13	60
2	10 (7)	40
3	10 (7)	40
4	12	65
5	12	50
6	10	40
7	8	40
8	8	40

A PedVAE is NOT identified

Remember – for patients ≥ 30 days, a MAP of 1-10 are equal to 10

Vent Day	Daily Minimum MAP	Daily Minimum FiO ₂	
1	13	60	≥ 2-day period of worsening in the MAP
2	10 (7)	40	parameter with an increase of ≥ 4 cm H ₂ O
3	10 (7)	40	over the baseline period <u>is</u>
4	12	65	not identified
5	12	50	
6	10	40	
7	8	40	
8	8	40	

A PedVAE is identified Patients ≥ 30 days – MAP values of 1-10 are equal to 10

Vent Day	Daily Minimum MAP	Daily Minimum FiO ₂	
1	13	60	≥ 2-day period of worsening in the MAP
2	10 (7)	40	parameter with an increase of ≥ 4 cm H ₂ O
3	10 (7)	40	over the first day in the
4	14	65	baseline period – 10 to 14
5	14	50	
6	10	40	
7	8	40	
8	8	40	

Date of Event

 The date of onset of worsening oxygenation (day 1 of the required ≥ 2-day period of worsening oxygenation following a ≥ 2-day period of stability or improvement on the ventilator)

- Earliest date of event for VAE is mechanical ventilation day 3 (first day of worsening oxygenation)
- First possible day that PedVAE criteria can be <u>fulfilled</u> is mechanical ventilation day 4

Date of Event continued

- Sets the 14-day PedVAE Event Period
 - Each PedVAE is 14 days in duration (arbitrary—to standardize).
 - The Date of Event is day 1 of the PedVAE Event Period. So, for example, if June 1 is the date of onset of worsening oxygenation and a PedVAE is reported, a second PedVAE cannot be detected and reported until June 15.
- Defines the period during which the antimicrobial and pathogen optional questions apply

Reporting Exception related to Date of Event

• If the date of event (date of onset of worsening oxygenation) is on or after the date of documentation of evidence of consent AND the patient is being supported for organ donation purposes, the event should not be reported as a PedVAE.



Operationalizing PedVAE – Date of Event

(patient is < 30 days)

Vent Day	MAP min	FiO ₂ min	
1	13	60	
2	8 (7)	40	
3	8 (7)	40	
4	12	Event Date = Vent Da	y 4 (first day of
5	12	worsening oxy	genation)
6	10	14 Day event period Vent Day	
7	8	40	
8	8	40	

PedVAE Data Collection and Reporting

Where to find the PedVAE Data Collection forms?

https://www.cdc.gov/nhsn/ps c/pedvae/index.html

Data Collection Forms & Instructions

All Data Collection Forms are Print-only

PedVAE

Pediatric Ventilator-associated Event (PedVAE) form – January 2021 (57.113)
[PDF – 190 KB]

- Customizable form [DOCX 60 KB]
- Table of Instructions
 [PDF 150 KB]

Denominator Forms

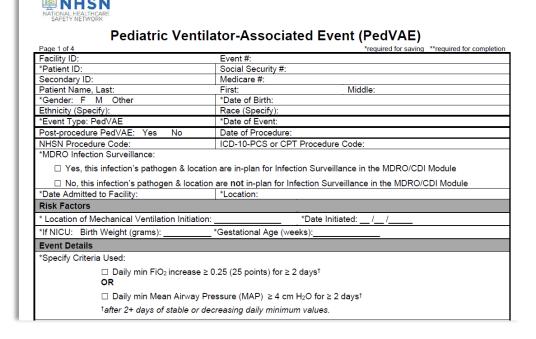
ACH

- Table of Instructions 🔼 [PDF 200 KB]

PedVAE Data Collection Form

All fields with an asterisk (*) are required:
 https://www.cdc.gov/nhsn/ forms/57.113-p.pdf

Refer to the Table of
 Instructions for details for accurate completion:
 https://www.cdc.gov/nhsn/forms/instr/57 113-508.pdf





Pediatric Ventilator-Associated Event (PedVAE)

Page 1 of 4	*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: PedVAE	*Date of Event:			
Post-procedure PedVAE: Yes No	Date of Procedure:			
NHSN Procedure Code:	ICD-10-PCS or CPT Procedure Code:			
*MDRO Infection Surveillance:				
☐ Yes, this infection's pathogen & location	are in-plan for Infection Surveillance in the MDRO/CDI Module			
☐ No, this infection's pathogen & location are not in-plan for Infection Surveillance in the MDRO/CDI Module				
*Date Admitted to Facility:	*Location:			
Risk Factors				
* Location of Mechanical Ventilation Initiation:	*Date Initiated: //			
*If NICU: Birth Weight (grams):	*Gestational Age (weeks):			
Event Details				
*Specify Criteria Used:				
 □ Daily min FiO₂ increase ≥ 0.25 (25 points) for ≥ 2 days[†] OR 				
□ Daily min Mean Airway Pressure (MAP) ≥ 4 cm H ₂ O for ≥ 2 days [†]				
†after 2+ davs of stable or decreasing daily minimum values.				

Data Collection Form (optional data)

Clinical event associated with the PedVAE? $\ \square$ Yes $\ \square$ I	No □ Unknown If Yes, check all that apply:	
□ Ventilator-associated Pneumonia	□ Sepsis or Septic Shock	
☐ Atelectasis	□ Neonatal Respiratory Distress Syndrome (RDS)	
☐ Acute Respiratory Distress Syndrome (ARDS)	☐ Bronchopulmonary Dysplasia/Chronic Lung Disease	
☐ Pulmonary Hypertension	☐ Reopened Patent Ductus Arteriosus (PDA)	
□ Pulmonary Edema	☐ Weaning from mechanical ventilation or other change in mechanical ventilation approach <u>without</u> clinical worsening	
☐ Pulmonary Hemorrhage	□ Other (specify)	
Antimicrobial agent(s) administered?		
☐ Yes ☐ No If Yes, select up to 3 antimicrobia	al agents:	
Drug1:; Dru	ug1 start date: / /	
Drug2:; Drug2 start date: / /		
Drug3:; Drug3 start date: / /		
Pathogen identified from one or more of the listed specir If Yes, which specimen type? (check all that apply)	mens? □ Yes □ No If Yes, specify pathogen on pages 2-3	
\square Lower Respiratory \square Upper Respiratory \square	Lung Tissue □ Pleural Fluid	
☐ Urine for Legionella or Streptococcus pneum	noniae antigen testing	
Pathogen identified from BLOOD? ☐ Yes ☐ No	_	
**Died: Yes No PedVAE contributed t	to death: Yes No Discharge Date:	
COVID-19: Yes No		
If Yes: Confirmed Suspected Assuance 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	
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Assuance 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, 308 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 25 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 sponsors, and a person is not required to respond to a collection of information unersity data OMB control numbers 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 (0620-0868). CDC 57.113 (Great 181-14).

Instructions for Completion of Pediatric Ventilator-Associated Event (PedVAE) Form (TOI)

https://www.cdc.gov/nhsn/forms/instr/57 113-508.pdf (5 pages total)

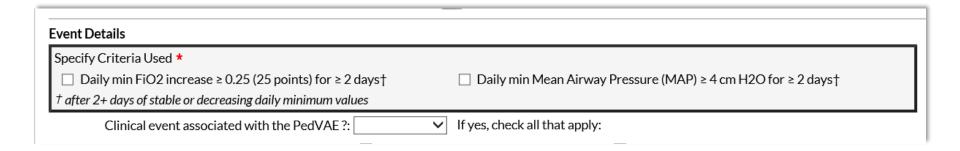


January 2022

Instructions for Completion of Pediatric Ventilator-Associated Event (PedVAE) Form (57.113)

Data Field	Instructions for Data Collection
Facility ID	The NHSN-assigned facility ID will be auto entered by the computer.
Event #	Event ID number will be auto entered by the computer.
Patient ID	Required. Enter the alphanumeric patient ID number. This is the patient identifier assigned by the hospital and may consist of any combination of numbers and/or letters.
Social Security #	Optional. Enter the 9-digit numeric patient Social Security Number.
Secondary ID	Optional. Enter the alphanumeric ID number assigned by the facility.

PedVAE: NHSN Application



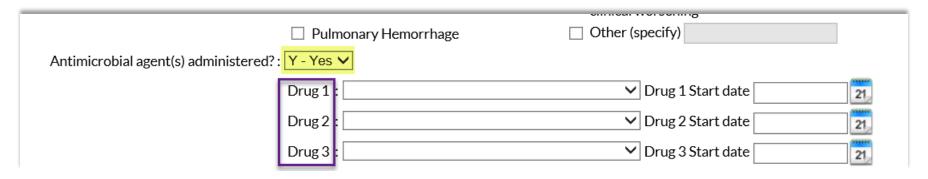
Select the specific criteria (FiO₂, MAP) used to meet PedVAE definition.

PedVAE Event: Clinical Event Associated with the PedVAE

Event Details			
Specify Criteria Used *			
☐ Daily min FiO2 increase ≥ 0.25 (25 points) for ≥ 2	days†	Daily min Mean Airway	Pressure (MAP) ≥ 4 cm H2O for ≥ 2 days†
† after 2+ days of stable or decreasing daily minimum val	lues		
Clinical event associated with the PedVAE?:	Y - Yes 💙	If yes, check all that apply:	
	☐ Ventilator-a	ssociated Pneumonia	☐ Sepsis or Septic Shock
	Atelectasis		☐ Neonatal Respiratory Distress Syndrome (RDS)
	☐ Acute Respi (ARDS)	ratory Distress Syndrome	☐ Bronchopulmonary Dysplasia/Chronic Lung Disease
	☐ Pulmonary Hypertension		☐ Reopened Patent Ductus Arteriosus (PDA)
	☐ Pulmonary I	Edema	Weaning from mechanical ventilation or other change in mechanical ventilation approach <u>without</u> clinical worsening
	☐ Pulmonary Hemorrhage		Other (specify)

Optional. Select Yes if the PedVAE is associated with any clinical diagnoses or events. Otherwise check No or Unknown. If Yes, check all that apply from the list above. Please note that there is a 200 character limit for "Other."

PedVAE Event: Antimicrobial agent(s) administered



- This field is <u>optional</u>. Select **Yes** if an antimicrobial agent(s) listed in the Appendix was administered on the date of event or within the 2 days before or 2 days after the date of event. Otherwise select **No**.
 - If antimicrobial agent(s) administered = Y Record Drug (up to 3) and enter administration start date.
 - Administration start date is limited to 1 year prior to current admission date.

PedVAE Event: Pathogen identified

Pathogen identified from one or more of the listed specimens?	es VIII Yes, which specimen types	pe? (check all that apply)	
□ Lo	ower Respiratory	Upper Respiratory	
☐ Lu	ung Tissue	Pleural Fluid	
☐ Ur	rine for Legionella or Streptococo	cus pneumoniae antigen testing	
Pathogen identified from BLOOD? : Y - Ye	es 🗸		
Died **: Y - Ye	es 🗸		PedVAE contributed to death
Discharge Date :	21		
COVID-19 *:	~		
Pathogens			
Pathogen 1:			
Pathogen 2:			
Pathogen 3:			

PedVAE Event: TOI for Pathogen identified

https://www.cdc. gov/nhsn/forms/i nstr/57 113-508.pdf

Event Details: Pathogen identified	Optional. Check Y if any pathogen was detected by culture or non-culture-based microbiological testing of upper or lower respiratory specimens, or <i>Legionella</i> or <i>Streptococcus pneumoniae</i> detected by urine antigen testing on the date of event or within the 2 days before or 2 days after the event; otherwise, check N. Specify pathogens on reverse form.
Event Details: Source of Pathogen Identified	Optional. If pathogen identified = Y select all specimen sources that apply: Lower Respiratory (for example, sputum, tracheal aspirate, bronchial washing, bronchoalveolar lavage), Upper Respiratory (for example, nasopharyngeal wash or swab), Lung Tissue, Pleural Fluid, Urine for Legionella or Streptococcus pneumoniae antigen testing; otherwise, check N.
Event Details: Pathogen identified in Blood	Optional. Check Y if pathogen was identified from blood with a specimen collection date within 2 days before the date of event to 13 days after the date of event; otherwise, check N. Specify pathogens on reverse form.
Event Details: Died	Required. Check Y if patient died during the hospitalization; otherwise, check N.
Event Details: PedVAE Contributed to Death	Conditionally required. If the patient died, check Y if such evidence is available (for example, death/discharge note, autopsy report, etc.); otherwise, check N.

PedVAE Event: COVID-19

COVID – 19

Required. Check Y if the patient met the definition of confirmed COVID-19 on the date of event; otherwise, check N.

Confirmed: A patient with a positive COVID-19 (SARS CoV-2) laboratory viral test indicating current infection (**NOTE:** this does not include serology testing for antibody).

- Required for all events occurring on or after January 1, 2022
- Lab finding of the most recent COVID-19 viral test prior to or on the date of event should be used for the response
 - Answer COVID-19 as 'YES' if the patient is lab test confirmed COVID-19 on the date of event
 - Answer COVID-19 as 'NO' if the most recent lab test prior to or on the date of event is negative

Denominator Data

- Patient Days (required)
- Ventilator Days (required)
- NICU denominator data (patient days and ventilator days)
 - NICU Birthweight (required)

701 1000 g 1001 1000 g 1001 2000 g	≤750 g	751-1000 g	1001-1500 g	1501-2500 g	>2500 g
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NICU Gestational Age (optional)

Extremely preterm (<28 weeks)	Very preterm (28 to <32 weeks)	Moderate to late preterm (32 to <37 weeks)	Term (≥37 weeks)
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Episodes of Mechanical Ventilation (optional)

Monthly Reporting Plan – Pediatric Location

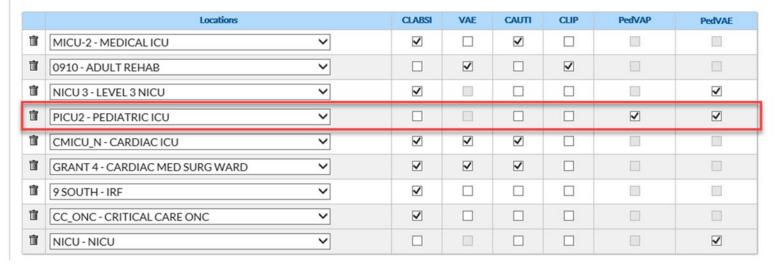


Add Monthly Reporting Plan

Mandatory fie	elds marked with *
Facility ID *	
Month *	
Year *	
	☐ No NHSN Patient Safety Modules Followed this Month

Note that in pediatric locations, both PedVAP and PedVAE are eligible for selection.

Device-Associated Module



Summary Data - Pediatric Location



Denominators for Intensive Care Unit (ICU)/Other locations (not NICU or SCA)

Denominator Data	
	Report No Events
Total Patient Days *:	
Central Line Days:	CLABSI:
Urinary Catheter Days:	CAUTI:
Ventilator Days ★:	VAE: PedVAE: PedVAP: PedVAP:
APRV Days:	
Episodes of Mechanical Ventilation:	

Sample Values For Estimating Denominator Data					
	Check Box(es) if Sampling Used				
Sample Patient Days:					
Sample Central Line Days:					
Sample Urinary Catheter Days:					

When PedVAE is selected in the monthly reporting plan, both patient days and ventilator days are required when entering monthly summary data.

Monthly Reporting Plan – NICU Location



Add Monthly Reporting Plan

Note that only PedVAE is eligible for selection in the monthly reporting plan for a neonatal location.

Device-Associated Module

	Locations	CLABSI	VAE	CAUTI	CLIP	PedVAP	PedVAE
Î	MICU-2 - MEDICAL ICU	V		✓			
T	0910 - ADULT REHAB		✓		✓		
Î	NICU 3 - LEVEL 3 NICU	✓					✓
Ì	PICU2 - PEDIATRIC ICU					✓	✓
ì	CMICU_N - CARDIAC ICU ✓	✓	✓	✓			
ì	GRANT 4 - CARDIAC MED SURG WARD	V	✓	✓			
Û	9 SOUTH - IRF	✓					
Û	CC_ONC - CRITICAL CARE ONC	V					
Î	NICU - NICU						V

Summary Data - NICU Location



Mandatory fields ma	rked with *	
Facility ID *:	DHQP Memorial Hospital (ID 10000) V	•
Location Code *:	NICU - NICU	~

Birth Weights							
Birth Weight	Patient Days *	CL Days CLABSI	Vent Days *	No No PedVAE PedVAP	EMV	UrC Days	
<=750							
751-1000							
1001-1500							
1501-2500							
>2500							

Gestational Ages				
Gestational Age	Patient Days	Vent Days	No PedVAE	EMV
Extremely preterm (<28 weeks)				
Very preterm (28 to <32 weeks)				
Moderate to late preterm (32 to <37 weeks)				
Term (>=37 weeks)				

PedVAE Calculator

PedVAE Calculator

https://www.cdc.gov/nhsn/psc/pedvae/index.html

Pediatric Ventilator-associated Events (PedVAE)

Available In-Plan for Pediatric and Neonatal Inpatient Locations Only.

PedVAP surveillance using the <u>PNEU</u> protocol continues to be available for inplan surveillance for pediatric locations only. See <u>VAE</u> for in-plan surveillance for adult locations

Not available for Inpatient Psychiatric Facilities (IPFs)

PedVAE Calculator

operates based upon the currently posted PedVAE protocol.

Pediatric Ventilator-Associated Event Calculator

Version 1.0

Welcome to Version 1.0 of the PedVAE Calculator. Version 1.0 operates based upon the currently posted PedVAE protocol.

The Calculator is a web-based tool that is designed to help you learn how the PedVAE surveillance definition algorithm works and assist you in making PedVAE determinations.

Please note that the PedVAE Calculator will not ask you to enter any patient identifiers (other than dates of mechanical ventilation, which you can change as you see fit). The PedVAE Calculator does not store any patient data that you enter, and it will not report any data that you enter or any PedVAE determinations to the NHSN. You will not be able to export data entered into the Calculator.

If you have questions or suggestions about the Calculator, please feel free to send them to the NHSN mailbox, nhsn@cdc.gov.



Pediatric Ventilator-Associated Event Calculator Version 1.0 (must have javascript enabled)

https://www.cdc.gov/nhsn/pedvae-calculator/index.html

National Healthcare Safety Network (NHSN)

CDC > NHSN > Materials for Enrolled Facilities

NHSN Pediatric Ventilator-Associated Event (PedVAE) Calculator Ver. 1.0

Welcome to the Pediatric Ventilator-Associated Event Calculator. Version 1.0 operates based upon the currently posted PedVAE protocol. It is strongly encouraged that you thoroughly review the PedVAE protocol.

- The calculator recognizes Mean Airway Pressure (MAP) values 0-8 cm H_2 0 as equal to 8 for patients < 30 days of age and MAP values 0-10 cm H_2 0 as equal to 10 for patients \geq 30 days of age and corrects entries according to the PedVAE protocol prior to making a PedVAE determination.
- Daily minimum MAP readings are to be rounded to the nearest whole number using the following method as an example: A MAP value 10.00 10.49 is rounded to 10 and a MAP value 10.50 10.99 is rounded to 11.
- The calculator finds multiple PedVAEs per patient as long as they conform to the 14 day rule.

To get started, enter a date below that corresponds to the first day the patient was placed on mechanical ventilation during the mechanical ventilation episode of interest. You may type in a date or use the popup calendar when it appears. You may only enter dates within the past year. If the patient has been on mechanical ventilation for more than one year during the current mechanical ventilation episode, choose a start date that is more recent but is at least 7 days before the period of interest, more...

mechanical ventilation episode, choose a start date that is more recent but is at least 7 days before the period of interest. more...

Click "more" for additional instructions on using the PedVAE Calculator.

Is the patient's day of life (where date of birth = day of life 1)

less than 30 days on the Mechanical Ventilation Start Date?

NHSN Pediatric Ventilator-Associated Event (PedVAE) Calculator Ver. 1.0

Welcome to the Pediatric Ventilator-Associated Event Calculator. Version 1.0 operates based upon the currently posted PedVAE protocol. It is strongly encouraged that you thoroughly review the PedVAE protocol.

- The calculator recognizes Mean Airway Pressure (MAP) values 0-8 cmH₂0 as equal to 8 for patients < 30 days of age and MAP values 0-10 cmH₂0 as equal to 10 for patients \geq 30 days of age and corrects entries according to the PedVAE protocol prior to making a PedVAE determination.
- Daily minimum MAP readings are to be rounded to the nearest whole number using the following method as an example: A MAP value 10.00 10.49 is rounded to 10 and a MAP value 10.50 10.99 is rounded to 11.
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The calculator runs locally on your machine. Data that you enter are not stored, nor are they transmitted to NHSN. Feel free to enter or change as much data as you like. If you don't understand something, there are several mechanisms for getting help. Most of the buttons and table headings will give an expanded description if you hover your mouse over the item in question. Also the explain button will pop up an explanation of the reasoning behind the calculation. The explanation box is movable as are all the popup windows. That allows you to open one up and drag it to the side as you work. The explanation will automatically update itself as you work through the protocol.

less...

Welcome to the Pediatric Ventilator-Associated Event Calculator. Version 1.0 operates based upon the currently posted PedVAE protocol. It is strongly encouraged that you thoroughly review the PedVAE protocol.

- The calculator recognizes Mean Airway Pressure (MAP) values 0-8 cmH₂0 as equal to 8 for patients < 30 days of age and MAP values 0-10 cmH₂0 as equal to 10 for patients ≥ 30 days of age and corrects entries according to the PedVAE protocol prior to making a PedVAE determination.
- Daily minimum MAP readings are to be rounded to the nearest whole number using the following method as an example: A MAP value 10.00 10.49 is rounded to 10 and a MAP value 10.50 10.99 is rounded to 11.
- The calculator finds multiple PedVAEs per patient as long as they conform to the 14 day rule.

To get started, enter a date below that corresponds to the first day the patient was go a date or use the popup calendar when it appears. You may only enter dates within the mechanical ventilation episode, choose a start date that is more recent but is at least. The calculator runs locally on your machine. Data that you enter are not stored, nor a understand something, there are several mechanisms for getting help. Most of the bu question. Also the explain button will pop up an explanation of the reasoning behind t up and drag it to the side as you work. The explanation will automatically update itself

less...



mechanical ventilation episode of interest. You may type in nical ventilation for more than one year during the current

23 4 5 er or change as much data as you like. If you don't d description if you hover your mouse over the item in le as are all the popup windows. That allows you to open one

Mechanical Ventilation Start Date:

(mm/dd/yyyy)

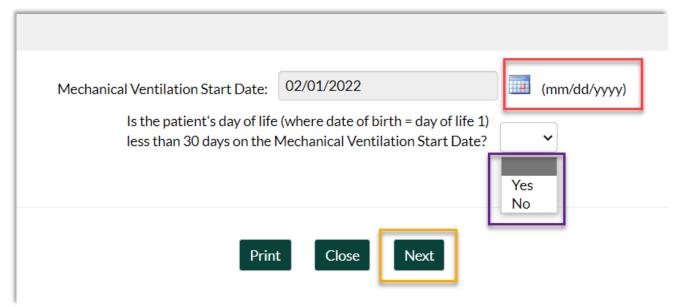
Is the patient's day of life (where date of birth = day of life 1) less than 30 days on the Mechanical Ventilation Start Date?



Enter the date of mechanical ventilation initiation and respond to the day of life question.

This allows the calculator to determine which Daily Minimum MAP value interpretation to use on the next screen.

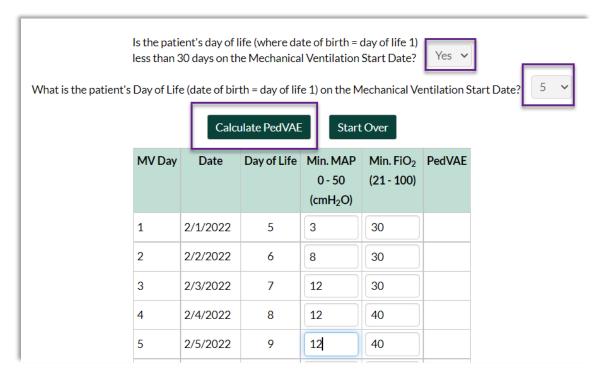
Click NEXT.



PedVAE Calculator

- Note the patient is < 30
 days on Mechanical
 Ventilation Start Date
 - the age category and day of life are not able to be edited on this screen

Enter the Daily Minimum
 Values and click on
 Calculate PedVAE.



NHSN Pediatric Ventilator-Associated Event (PedVAE) Calculator Ver. 1.0

A Pediatric Ventilator-Associated Event (PedVAE) based on MAP values occurred on 2/3/2022.

Click on the "Explain" button to see how this determination was made.

Is the patient's day of life (where date of birth = day of life 1) less than 30 days on the Mechanical Ventilation Start Date?



What is the patient's Day of Life (date of birth = day of life 1) on the Mechanical Ventilation Start Date?



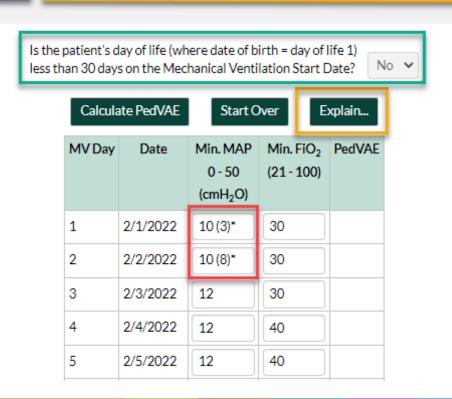
- The Calculator interprets MAP values of 0-8 cmH₂O as equal to 8 cmH₂O since the patient < 30 days</p>
- A PedVAE is identified

	Calculate Pe	edVAE	Start Over	Explair	1
MV Day	Date	Day of Life	0 - 50 (cmH ₂ O)	Min. FiO ₂ (21 - 100)	PedVAE
1	2/1/2022	5	8 (3)*	30	
2	2/2/2022	6	8	30	
3	2/3/2022	7	12	30	‡ PedVAE
4	2/4/2022	8	12	40	

NHSN Pediatric Ventilator-Associated Event (PedVAE) Calculator Ver. 1.0

No Pediatric Ventilator-Associated Event (PedVAE) detected. Click on the "Explain" button to see an explanation of the PedVAE definition.

- The Calculator interprets MAP values of 0-10 cmH₂O as equal to 10 cmH₂O since the patient ≥ 30 days
- No PedVAE is detected because the increase over the baseline (10) is not ≥ 4 cmH₂0
- Select Explain button for further explanation



Questions: NHSN@cdc.gov

NHSN Website: http://www.cdc.gov/nhsn/

For more information, contact CDC 1-800-CDC-INFO (232-4636) TTY: 1-888-232-6348 www.cdc.gov



The findings and conclusions in this report are those of the authors and do not necessarily represent the official position of the Centers for Disease Control and Prevention.

