2022 NHSN Pediatric Ventilator-Associated Event (PedVAE) Checklist

Pediatric Ventilator-Associated Event (PedVAE) Summary			
Criterion	Criterion Met	Date of Event (DOE)	
PedVAE			
Please refer to Chapter 11 Pediatric Ventilator-Associated Event (PedVAE) of the Patient Safety			
Manual for additional information.			

Documentation Review Checklist			
Pediatric Ventilator-Associated Event (PedVAE)			
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Element	Element Met	Date	
Patient has at least <u>one</u> of the following:			
Baseline period of stability* on the ventilator			
Baseline period of improvement* on the ventilator			
<u>AND</u> immediately following a period of stability or improvement (as above), patient has at least <u>one</u> 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 \geq 4 cm H ₂ O over the daily minimum MAP of the first day in the baseline period, sustained for \geq 2 calendar days			
Note: *Stability or improvement on the ventilator is defined by ≥ 2 calendar days of stable or decreasing daily minimum FiO ₂ or MAP values. The baseline period is defined as the 2 calendar days immediately preceding the first day of increased daily minimum FiO ₂ or MAP. **Daily minimum FiO ₂ is the lowest value of FiO ₂ documented during a calendar day that is maintained for > 1 hour. †Daily minimum MAP is the lowest value documented during the calendar day. For the purposes of surveillance, in patients < 30 days old, daily minimum MAP values of 0-8 cm H ₂ O are considered equal to 8 cm H ₂ O; in patients ≥ 30 days old, daily minimum MAP values of 0-10 cm H ₂ O are considered equal to 10 cm H ₂ O.			
Comments/Notes:			



REPORTING INSTRUCTIONS:

- Conducting in-plan PedVAE surveillance means assessing patients for the presence of events meeting the PedVAE definition.
- 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.
- Secondary BSIs are not reported or attributable to a PedVAE.
- Clinical findings associated with a PedVAE may assist in better understanding the etiology and focusing efforts to
 prevent PedVAEs. Should a facility choose to provide the following information, the PedVAE form includes
 optional data fields to report:
 - O Clinical diagnoses or events that were associated with the PedVAE. Note that multiple events may be reported for a single PedVAE.
 - Antimicrobial agents listed in the Appendix (<u>PedVAE Protocol</u>) that are administered on the date of event or within the 2 days before or 2 days after the event. The name of the specific antimicrobial agent and the administration initiation date may also be reported.
 - Pathogens detected by culture or non-culture-based microbiological testing of upper or lower respiratory specimens with a specimen collection date on the date of event or within the 2 days before or 2 days after the date of event or in blood with a specimen collection date within the 2 days before the date of event and up to 13 days after the date of event.
 - <u>Note</u>: Because organisms belonging to the following genera are typically causes of community-associated respiratory infections and are rarely or are not known to be causes of healthcare-associated infections, they are excluded, and cannot be reported: *Blastomyces*, *Histoplasma*, *Coccidioides*, *Paracoccidioides*, *Cryptococcus*, and *Pneumocystis*.
 - Legionella or Streptococcus pneumoniae detected by urine antigen testing with a date of specimen collection on the date of event or within the 2 days before or 2 days after the event.

