

# **Prevention of Ventilator-Associated Pneumonia in Pediatric Intensive Care Unit**

*Thesis*

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of the M.Sc. degree  
in Pediatrics**

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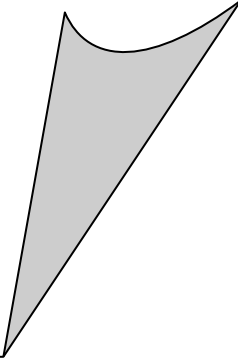
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## *List of Abbreviation*

<b>A/C</b>	Assist –control
<b>BAL</b>	Bronchoalveolar lavage
<b>CA-MRSA</b>	Community acquired-Methicillin resistant staphylococcus auras
<b>CDC</b>	Centers of Disease Control and Prevention
<b>CDC</b>	Centers of Disease Control and Prevention
<b>CMV</b>	Continuo mandatory ventilation
<b>CONS</b>	Coagulase negative staphylococcus aueus
<b>CPAP</b>	Continuous positive airway pressure
<b>CPIS</b>	Clinical pulmonary infection score
<b>CPIS</b>	Clinical Pulmonary Infection Score
<b>CXR</b>	Chest x ray
<b>ESBLs</b>	Extended spectrum B-lactamases
<b>ETT</b>	Endotracheal tube
<b>FDA</b>	Food and drug American Association
<b>FRC</b>	Functional residual capacity
<b>HAI</b>	Hospital acquired infection
<b>HA-MRSA</b>	Hospital acquired-Methicillin resistant staphylococcus auras
<b>HCU</b>	High- care unit
<b>HDUs</b>	High-density units
<b>HIV</b>	Human immunodeficiency virus
<b>HOB</b>	Head of bed
<b>ICU</b>	Intensive care unit
<b>IMV</b>	Intermittent mandatory ventilation
<b>MDR</b>	Multi-drug resistant
<b>MRSA</b>	Methicillin-Resistant Staphylococcus Auras
<b>NI</b>	Nosocomial infection
<b>NIPPV</b>	Non invasive positive pressure ventilation
<b>NNIS</b>	National Nosocomial Infection Surveillance system
<b>PEEP</b>	Positive end expiratory pressure
<b>PICU</b>	Pediatric intensive care unit
<b>PPV</b>	Positive pressure ventilation
<b>PSB</b>	Protected specimen brush
<b>SDD</b>	Selective decontamination of the digestive tract
<b>SICU</b>	Surgical intensive care unit
<b>SIMV</b>	Synchronized intermittent mandatory ventilation
<b>SSD</b>	Subglottic secretion drainage
<b>TICU</b>	Trauma Intensive Care Unit
<b>VAP</b>	Ventilator-associated pneumonia
<b>VISA</b>	Vancomycin-intermediate staphylococcus auras
<b>VRSA</b>	Vancomycin-resistant staphylococcus auras

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## ***Introduction***

Ventilator-associated pneumonia (VAP) is defined as pneumonia in mechanically ventilated patients that develops later than or at  $\geq 48$  hrs after the patient has been placed on mechanical ventilator. It is the second most common hospital-acquired infection among pediatric and neonatal intensive care patients occurring in 6% to 22% of ventilated pediatric intensive care unit (PICU) patients (*Fogelia, et al., 2007*)

VAP is associated with increased morbidity and mortality in PICU patients. It is the leading cause of death among hospital-acquired infections in the intubated patients. Hospital mortality in ventilated patients who develops VAP is 46 %, compared to 32% for ventilated patient who do not develop VAP along the duration of mechanical ventilation. Moreover, VAP adds an estimated cost of \$ 40,000 to a typical hospital admission (*Ibrahim, et al., 2007*)

VAP is diagnosed in patients who are mechanically ventilated for more than or equal to  $\geq 48$  hrs and have developed abnormal chest radiography with either one of the following symptoms: fever ( $\geq 38$  or more) with no other recognized cause, leucopenia ( $< 4,000/mm^3$  or less), leucocytosis ( $> 13,000/mm^3$  or more). In addition, patients should have at least two changes either in the sputum character, respiratory secretions, cough or worsening of gas exchange ( $O_2$  desaturation, increased  $O_2$  requirements or increased ventilation demand) (*Cordero, et al., 2007*)

The risk factors for developing VAP were found to be genetic syndromes, transport out of the PICU, re-intubation, prior antibiotic use, continuous enteral feeding, bronchoscope, immunosuppressants, immunodeficiency, neuromuscular blockade, some medications such as steroids and  $H_2$  blockers. The most commonly isolated organisms in

VAP were staphylococcus aureus and pseudomonas aeruginosa (*Elward, et al., 2002*)

Because of higher incidence and costs of VAP, there are several recommendations to decrease it. The health care infection control practices advisory committee suggests using oro-tracheal tubes instead of naso-tracheal tubes when the patients require mechanical ventilation, changing breathing circuits of ventilator only if malfunction or visibly contaminated and using endo-tracheal tubes with dorsal lumen to allow respiratory secretions to drain. (*Tablan, et al., 2004*) Lately, health care infection control practices advisory committee suggested also implementing ventilator bundle which is a group of evidence-based practices that when implemented together for ventilated patients resulted in dramatic reductions in the incidence of VAP.

The ventilator bundle has four key components:

- Elevation of the head of the bed to between 30 and 45 degrees.
- Daily "sedation vacation" and daily assessment of readiness for extubation.
- Peptic ulcer disease prophylaxis (unless contraindicated).
- Deep venous thrombosis (DVT) prophylaxis (unless contraindicated). (*Curley, et al., 2007*)



## *Aim Of The Work*

The aim of this work is to study the prevalence and risk factors of VAP in ventilated patients admitted in PICU. We also aim to assess the impact of implementing head of bed elevation, sedation vacation, weaning and assessment of spontaneous breathing and peptic ulcer prophylaxis on VAP rates.

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## ***Overview On Mechanical Ventilation***

### **What is meant by mechanical ventilation?**

Assisted ventilation can be defined as the movement of gas into and out of the lung by an external source connected directly to the patient. The external source may be a resuscitation bag, a continuous distending pressure device, or a mechanical ventilator. Attachment to the patient can be via a face mask, endotracheal tube, nasal prongs, or tracheotomy. Although not in general use today in modern intensive care nurseries, negative-pressure ventilation can be applied by an apparatus surrounding the infant's thorax. (*Goldsmith and Karotkim, ۲۰۰۳*)

### **Indication of mechanical ventilation**

The principal indication for mechanical ventilation (MV) is respiratory failure. (*Amitai, et al., ۲۰۰۴*) It can be also indicated when other simple measures of respiratory support (oxygen, aerosol, chest physiotherapy and suctioning) are not effective to improve oxygenation and/ or ventilation. (*Viana, et al., ۲۰۰۴*)

MV is indicated for both hypercapnic respiratory failure and hypoxemic respiratory failure. It is also indicated for intentional hyperventilation in the setting of major head injury with elevated intracranial pressure and for suspicion of clinical brain herniation from any cause. (*Amitai, et al., ۲۰۰۴*)

Respiratory failure is almost always and most appropriately a clinical diagnosis. The decision to intubate and mechanically ventilate is generally made purely on clinical grounds without delay for laboratory evaluation or pulmonary function data. (*Amitai, et al., ۲۰۰۴*) **As shown in Table ۱.**

**Table (1): Laboratory and clinical criteria for mechanical ventilation**

**Persistent arterial hypoxemia**

Arterial oxygen saturation below 80% inspite of 100% oxygen therapy.

PaO<sub>2</sub> below 60 mmHg inspite of 100% oxygen therapy.

Cyanosis inspite of 100% oxygen therapy.

**Alveolar hypoventilation**

Altered consciousness or PaCO<sub>2</sub> above 70 mmHg (due to severe ventilation-perfusion mismatch or respiratory muscle fatigue).

(*Slutsky, 2009*)

Respiratory failure may be caused by disorders at any point in the respiratory system: respiratory center in the brain stem, spinal cord, motor nerve roots, respiratory muscles, the thoracic cage, airways and the lung interstitium and pulmonary vessels. (*Henning and South, 1999*) As shown in **Table 2**

**Table (2): Causes of Respiratory Failure:**

**Pulmonary causes:**

- Respiratory distress syndrome.
- Aspiration syndromes.
- Pulmonary hemorrhage.
- Pneumonia.
- Pulmonary alveolar proteinosis.
- Wilson-Mikity syndrome.
- Broncho-pulmonary dysphasia.
- Pulmonary insufficiency of prematurity.
- Pneumothorax.
- Tumors.
- Diaphragmatic hernia.
- Cylothorax.
- Congenital malformations (Lobar emphysema, cystic adenomatoid malformation, lymphangiectasis).
- Cystic fibrosis.
- Vasculitis, collagen vascular diseases.

**Central causes:**

- Status epilepticus.
- Severe static encephalopathy.

- Brain stem insult.
- Brain abscesses, hematoma and tumors.
- Apnea of prematurity.
- Drugs: morphine, Magnesium sulfate, Mepivcaine, meperidine).
- Arnold-Chiari malformation (Central type).
- Abnormalities of muscles of respiration.
- Phrenic nerve palsy.
- Spinal cord injury.
- Werdnig-Hoffmann syndrome.
- Myasthenia gravis.
- Myopathy.
- Neuropathy.
- Guillan-Barre syndrome.
- Botulism.

**Airway obstruction.**

- Laryngomalacia.
- Chonal atresia.
- Pierre Robin syndrome.
- Micrognathia.
- Nasopharyngeal tumor.
- Adenotonsillar hypertrophy.
- Retropharyngeal abscess.
- Subglottic stenosis.
- Acute epiglottitis
- Foreign body aspiration

**Miscellaneous**

- Congestive heart failure.
- Persistent fetal circulation.
- Post operative anesthesia, sedation.
- Extreme immaturity.
- Shock.
- Sepsis.
- Hypoglycemia.
- Electrolyte abnormalities.
- Acid-base imbalance (Sepsis, renal failure, diabetic ketoacidosis, hepatic disease)

*(Baker and Ruddy, ٢٠٠٠ and Goldsmith and Karotkim, ٢٠٠٢)*

To summarize, absolute indications of MV include apnea, persistent hypoxemia, severe hypoventilation and markedly elevated intracranial pressure while relative indications include shock state, deep coma and refractory status epilepticus. (*Khilnani and Uttam, २००३*) As shown in **Table २**.

**Table (२): Indications of Mechanical ventilation:**

Resuscitation from circulatory arrest; shock
Hypoventilation and apnea;
Respiratory failure due to hypoxemic and intrinsic pulmonary disease,
Respiratory assistance for super normal gas exchange, including both
persistent pulmonary hypertension (newborn or post-cardiac) and
increased intracranial pressure;
Loss of mechanical integrity of the respiratory apparatus (e.g., muscle
weakness, paralysis);
Prophylactic indication as post-surgical recovery and to reduce work of
breathing.

*Khilnani and Uttam, २००३*

## **Means and Modes of Mechanical Ventilation:**

Ventilators can be classified according to the manner in which they control ventilation, often termed the ventilator mode. To start inspiration, the machine can be triggered by the patient (Assistor type), by ventilator only (controller type), or by both the patient and the ventilator (assistant-controller type). In assistor-controller ventilator, a device allows the patient to initiate some respirations; however, it also has a pre-determined frequency of intermittent mandatory ventilation (IMV) that can be used as backup. (*Goldsmith and Kartkim, २००२*)

**Continuous mandatory ventilation (CMV)**

Breaths are delivered at preset intervals, regardless of patient effort. This mode is used most often in the paralyzed or apneic patient because it can increase the work of breathing if respiratory effort is present. Many ventilators do not have a true CMV mode and offer A/C instead. (*Amitai, 2007*)

**Intermittent mandatory ventilation (IMV)**

With IMV, breaths are delivered at preset interval, and spontaneous breathing is allowed between ventilator-administered breaths. Spontaneous breathing occurs against the resistance of the airway tubing and ventilator valves which may be formidable. This mode has given way to synchronous intermittent mandatory ventilation (SIMV). (*Amitia, 2007*)

**Assist-control (A/C)**

The ventilator delivers preset breaths in coordination with the respiratory effort of the patient. With each inspiratory effort, the ventilator delivers a full assisted tidal volume. Spontaneous breathing independent of the ventilator between A/C breaths is not allowed. As might be expected, this mode is better tolerated than CMV in patients with intact respiratory effort. (*Amitai, 2007*)

If the patient does not trigger the ventilator frequently enough, the ventilator initiates breaths, ensuring the desired minimum respiratory rate. (*Porter and Kaplan, 2007*) A high sensitivity (shallow inspiratory effort) may cause unintentional triggering of mandatory breaths, and a low sensitivity may increase work of breathing needed to open a demand valve to increase gas flow to the patient. (*Rotta, 2007*)

**Synchronized intermittent mandatory ventilation (SIMV)**

SIMV delivers breaths at a set rate and volume that is synchronized to the patient efforts. In contrast to A/C, patient efforts beyond the set

respiratory rate are unassisted, although the intake valve opens to allow a breath. This mode remains popular, despite the fact that it neither provides full ventilator support as does A/C nor is an effective means of liberating the patient from mechanical ventilation. (*Porter and Kaplan, २००१*)

Synchronization attempts to limit barotraumas that may occur with IMV when a preset breath is delivered to a patient who is already maximally inhaled (breath staking) or is forcefully exhaling. (*Amitai, २००१*)

### **CPAP and PEEP (Continuous positive airway pressure and positive end expiratory pressure)**

CPAP refers to the maintenance of positive pressure throughout the ventilatory cycle. It implies that ventilation is occurring spontaneously without mechanical pressure breath, it is referred to as PEEP. (*Khilnani and Uttam, २००६*)

It is a mode of operation (debated as to whether truly a mode, because no tidal volume is generated by the ventilator) in which a preset pressure is maintained while the patient is allowed to breathe spontaneously. Patient determines his or her own rate and tidal volume. It increases mean airway pressure and; therefore oxygenation. Possibly decreases work of breathing if optimizes functional residual capacity (FRC) and ventilation/perfusion matching. (*Goldsmith and Karotkim, २००३*)

In the pediatric patient, it is used most commonly in the weaning of the chronically ventilated patient with chronic lung disease, or in patients in whom malacia of the airway is the predominant factor. In a neonate CPAP is commonly used as a mode of assisting the respiratory status for lung disease to avoid mechanical ventilation, as well as a weaning mode. (*Khilnani and Uttam, २००६*)