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MANAGEMENT OF PATHOLOGICAL CONDITIONS INDUCED BY MECHANICAL VENTILATION

An Essay submitted for Partial Fulfillment of Master Degree In Intensive Care

By

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IN THE NAME OF ALLAH THE COMPASSIONATE, THE MERCIFUL

"And say (O' my lord) advance me in knowledge" (Sura
Taha 114)

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List of abbreviations

(A/C)Assist-control ventilation.
(APRV)airway pressure release ventilation.
(AMV) Assisted mechanical ventilation.
(ARDS) acute respiratory distress syndrome.
(ATS)American Thoracic Society.
(BAL)bronchoalveolar lavage.
(BVV)Biologically variable ventilation.
(CIM)critical illness myopathy.
(CINMA)critical illness neuromuscular abnormality.
(CMV) Controlled-mechanical ventilation.
(CO)cardiac output.
(COPD) chronic obstructive pulmonary disease.
(CPAP). Spontaneous continuous positive airway pressure.
(CPIS)clinical pulmonary infection score.
(C _{RS}) respiratory-system compliance.
(CT)

(DO_2) oxygen delivery.
(ERV)The expiratory reserve volume.
(ETT) endotracheal tube.
(FRC) The functional residual capacity.
(GI) Gastrointestinal.
(HAP)Hospital acquired pneumonia .
(HCAP) Heath care associated pneumonia.
(HFOV) High frequency oscillatory ventilation.
(IC)The inspiratory capacity.
(ICP)intracranial pressure.
(ICU) the intensive care unit.
(IMV) Intermittent mandatory ventilation.
(IRV) The inspiratory reserve volume.
(IV)Intravenous.
(LTMV)long-term mechanical ventilation.
(MRSA)Methicillin-resistant Staphylococcus aureus.
(MV)Minute volume.
(NIV) Non-invasive ventilation.

(NMB)Neuromuscular bloker
(NPPV) non-invasive positive pressure ventilation.
(PAV)Proportional assist ventilation.
(P _{aw}) airway pressure.
(PCV) Pressure-controlled ventilation.
(PC-IRV) pressure-controlled inverse-ratio ventilation.
(PEEP)positive end-expiratory pressure.
(PIP) Peak Inspiratory Pressures.
(PNCI) polyneuropathy of critical illness.
(PSV) Pressure support ventilation.
(PTX)Pneumothorax.
(QEA) quantitative-culture of endotracheal aspirate.
(RAP)right atrial pressur.
(RR) respiratory rate.
(RV) The residual volume.
(SENIC) study on the efficacy of the nosocomial infection
control.
(SIMV). Synchronized intermittent mandatory ventilation.

(SIRS)Systemic Inflammatory Response Syndrome.
(TLC) The total lung capacity .
(UTI)Urinary Tract Infection.
(V_A) volume of alveolar ventilation/minute.
(VAIs) ventilator assisted individuals.
(VALI) Ventilator-associated lung injury.
(VAP)Ventilator-associated pneumonia.
(VC) The vital capacity.
(V _D)dead space air.
(· D)
(V_E) the total volume of expired air.
(V_{E})the total volume of expired air.
$(V_E)the\ total\ volume\ of\ expired\ air.$ $(VILI)ventilation-induced\ lung\ injury.$

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Introduction 1

INTRODUCTION

Mechanical ventilation is one of the life support procedures closely associated with the development of modern Intensive Care Medicine. Although mechanical ventilation is a life-saving intervention for patients suffering from acute respiratory failure, it is associated with numerous grave complications and should be discontinued at earliest possible time (**Bilan et al., 2009**).

The indications for mechanical ventilation, rather than acute respiratory failure are, coma, acute exacerbation of chronic obstructive pulmonary disease, neuromuscular disorders, and other disorders including; acute respiratory distress syndrome, heart failure, pneumonia, sepsis, complications of surgery, and trauma. The objectives of mechanical ventilation are primarily to decrease the work of breathing and reverse life-threatening hypoxemia or acute progressive respiratory acidosis (Bilan et al., 2009).

The most serious potential complication of mechanical ventilation is ventilation-induced lung injury (VILI). VILI was, for years, synonymous with clinical barotrauma, the leakage of air due to disruption of the airspace wall. The extra-alveolar accumulation of air causes several manifestations of which the most threatening is tension pneumothorax (Han et al., 2005).

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However, mechanical force-induced signal transduction (mechano-transduction) is responsible for many physiological processes in lung development_in maintaining lung functions and in pathological conditions related to lung disease, such as asthma, chronic obstructive pulmonary disease (COPD), and acute respiratory distress syndrome (ARDS), especially related to VILI. However, many signal transduction pathways are shared by both the injury and repair processes (Han et al., 2005).

On other hand, acute lung injury and its more severe manifestation, ARDS, continue to represent significant clinical challenges with mortality rates of about 60%. Treatment in this patient population remains largely supportive, with mechanical ventilation until the acute insult subsides. Although necessary, positive pressure mechanical ventilation has been implicated as a cause of secondary lung injury acting to exacerbate the primary lung injury. Three main mechanisms of VILI have been postulated; volutrauma, or alveolar over-distension, atelectrauma or repetitive shear stresses of the alveolar epithelium caused by unstable alveoli recruiting, and derecruiting, and biotrauma, or inflammation secondary to the mechanical injury induced by volutrauma and atelectrauma. Ventilator-associated pneumonia (VAP) is a major infectious complication in critically ill patients in terms of its incidence and associated mortality and morbidity (Aarts et al., 2008).