



ARDS

Dr Hussam Bachir Al Bardan
MS.c. internal and pulmonary medicine
Ph.D. critical care medicine

3-10-2024

ARDS

- life-threatening form of respiratory failure
- characterized by
- 1. acute hypoxemia
- 2. bilateral radiographic infiltrates

- ARDS management remains largely supportive focusing on strategies intended to limit further lung injury
- high mortality rates persist, with those who survive often facing longterm impairments



 Should Patients with ARDS Receive Systemic Corticosteroids?

- Recommendation. We suggest using corticosteroids for patients with ARDS
- (conditional recommendation, moderate certainty of evidence).

Background

- Corticosteroids are anti-inflammatory medications that inhibit the synthesis of proinflammatory mediators present in ARDS.
- They are widely administered to patients with ARDS for the management of ARDS specifically and for concurrent conditions such as septic shock or pneumonia

 More recently, corticosteroids have been found to reduce mortality in COVID-19-related acute hypoxemic respiratory failure and severe community-acquired pneumonia

- May be associated with increased risk of harm when initiated after
 - > 14 days of mechanical ventilation
- Monitor more closely for adverse effects in patients with immunosuppressed conditions, metabolic syndrome, or known or increased risk of fungal, parasitic, or mycobacterial infections

- Optimal regimen, including type of corticosteroid, is unknown
- For patients with corticosteroid-responsive etiologies, regimen should be tailored to the specific condition
- · For other patients, regimens used in prior RCTs may be used
- For patients that improve rapidly, consider discontinuation at time of extubation

Dexa Ards trial

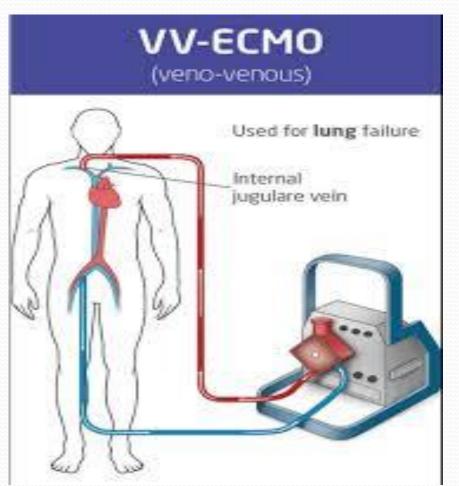
Original Article

Pierre-François Dequin, M.D., Ph.D., Ferhat Meziani, M.D., Ph.D., Jean-Pierre Quenot, M.D., Ph.D., Toufik Kamel, M.D., Jean-Damien Ricard, M.D., Ph.D., Julio Badie, M.D., Jean Reignier, M.D., Ph.D., Nicholas Heming, M.D., Ph.D., Gaëtan Plantefève, M.D., Bertrand Souweine, M.D., Ph.D., Guillaume Voiriot, M.D., Ph.D., Gwenhaël Colin, M.D., Jean-Pierre Frat, M.D., Ph.D., Jean-Paul Mira, M.D., Ph.D., Nicolas Barbarot, M.D., Bruno François, M.D., Guillaume Louis, M.D., Sébastien Gibot, M.D., Ph.D., Christophe Guitton, M.D., Ph.D., Christophe Giacardi, M.D., Sami Hraiech, M.D., Ph.D., Sylvie Vimeux, M.D., Erwan L'Her, M.D., Ph.D., Henri Faure, M.D., Jean-Etienne Herbrecht, M.D., Camille Bouisse, M.D., Aurélie Joret, M.D., Nicolas Terzi, M.D., Ph.D., Arnaud Gacouin, M.D., Charlotte Quentin, M.D., Mercé Jourdain, M.D., Ph.D., Marie Leclerc, M.Sc., Carine Coffre, M.Sc., Hélène Bourgoin, Pharm.D., Céline Lengellé, Pharm.D., Caroline Caille-Fénérol, M.Sc., Bruno Giraudeau, Ph.D., Amélie Le Gouge, M.Sc., for the CRICS-TriGGERSep Network March 2023

 Among patients with severe community-acquired pneumonia being treated in the ICU, those who received hydrocortisone had a lower risk of death by day 28 than those who received placebo.



 Should Patients with ARDS Receive VV-ECMO?





Recommendation

- We suggest the use of VV-ECMO in selected patients with severe ARDS
- (conditional recommendation, low certainty of evidence)

- Less invasive therapies, including lung protective ventilation, prone positioning, and neuromuscular blockade, should be initiated prior to ECMO consideration
- Resource limitations should be considered, with an emphasis on maximizing access for patients most likely to benefit from ECMO
- For patients meeting these criteria at hospitals without ECMO capabilities, consider transfer to ECMO centers when feasible

Conditions associated with increased risk for futility of treatment

- Irreversible etiology of respiratory failure
- Mechanical ventilation > 7 days
- Immunosuppression
- Multi-organ failure
- Older age
- Systemic bleeding or other contraindication to anticoagulation
- Chronic medical condition and life expectancy <1yr
- CNS hemorrhage or irreversible and incapacitating CNS pathology

 Should Patients with ARDS Receive
 Neuromuscular Blockade?

Recommendation

 We suggest using neuromuscular blockade in patients with early severe ARDS -(conditional recommendation, low certainty

- Unknown and potentially increased incidence of neuromuscular weakness with infusions of > 48 hours duration
- Use caution in patients with prior neuromuscular conditions

- Reduced mortality when compared to deep sedation. No mortality benefit when compared to light sedation
- May have greater utility in patients with ventilator dyssynchrony not mitigated by ventilator changes
- · Either bolus dosing or continuous infusion may be appropriate
- Consider cessation after 48 hours or earlier for patients that are improving rapidly
- Cisatracurium most frequently used in clinical trials; optimal agent unknown

Should Patients with ARDS
 Receive Higher Compared with
 Lower PEEP, with or without
 LRMs?

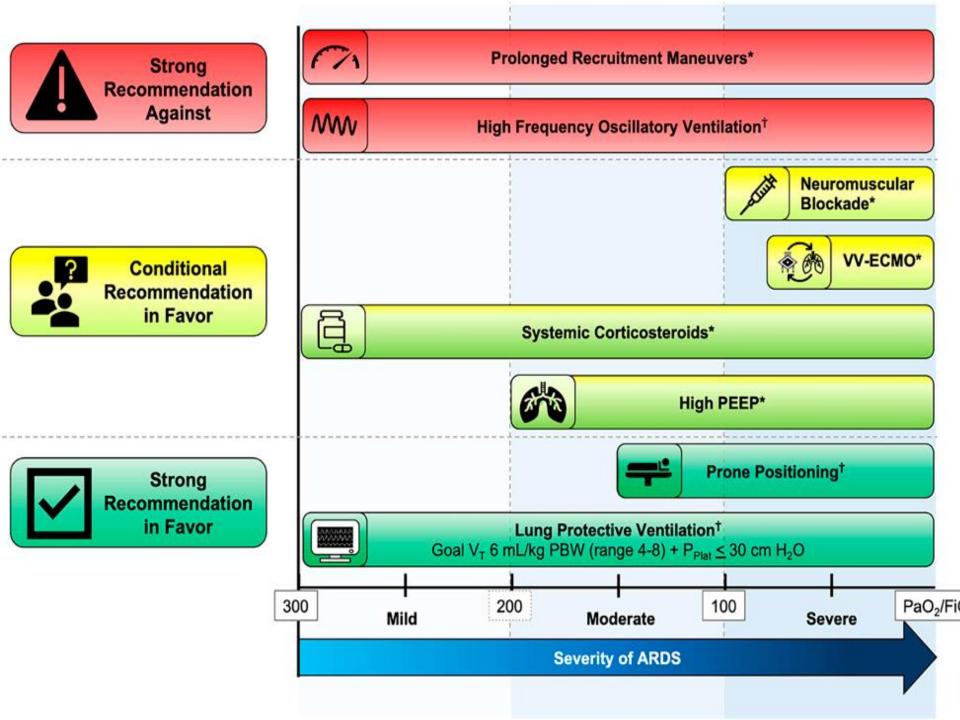


Recommendation

- We suggest using higher PEEP without LRMs rather than lower PEEP in patients with moderate to severe ARDS (conditional recommendation, low-moderate certainty).
- We recommend against using prolonged (PEEP >35 cm H2O for .60 s) LRMs in patients with moderate to severe ARDS (strong recommendation, moderate certainty).

- Respiratory mechanics, hemodynamics, and response to PEEP should be continuously monitored
- Use additional caution in patients with severe hemodynamic instability or increased risk of barotrauma
- · Prolonged recruitment maneuvers should be avoided

- Optimal strategy is unknown; selected strategy should be tailored to clinician expertise
- Potential strategies may include oxygenation-based titration or titration to maximal compliance or maximal safe plateau pressure
- Deleterious clinical response to higher PEEP (i.e. worsened oxygenation, dead space, compliance, or hemodynamics) should prompt re-evaluation of PEEP level



Thank you



