Negative Pressure Ventilation for Acute and Acute-on-Chronic Respiratory Failure

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Introduction

Respiratory failure is the commonest organ failure seen in the ICU¹ and is managed with non-invasive or invasive positive pressure ventilation (PPV). Negative pressure ventilation (NPV) could offer a safe and effective alternative, however existing devices, such as the iron lung, are heavy and access to the patient for ongoing care is a limitation. However, a new, light-weight negative pressure ventilator is now available that is safe and effective in a healthy volunteer trial². This device offers a practical alternative to PPV. They are cheaper to manufacture and do not require a pressurised gas supply. Therefore, they could potentially be used in Low- and Middle-Income Countries where acute and acute-on-chronic respiratory failure continue to cause significant morbidity and mortality³.

Potential Benefits of NPV: less ventilator-induced lung injury⁴; an increased cardiac output⁵; decreased pulmonary vascular resistance⁶

Study Objective: To address whether acute or acute-on-chronic respiratory failure in hospitalised adults can be safely and effectively managed with NPV.



Methodology

The protocol for this systematic review was registered with the international prospective register of systematic reviews (ID CRD420200220881). MEDLINE, EMBASE, CENTRAL, medRxiv, bioRxiv and Trip databases were searched (inception to 22nd April 2021).

Figure 1: Modern negative pressure ventilator

Two authors (RE, JVP) independently screened for potentially eligible studies using Rayyan Intelligent Systematic Review Software and the same two authors then reviewed the full texts of those deemed potentially eligible. Data extraction was then carried out by RE and independently checked by JVP.

Where appropriate a pooled effect estimate was carried out with variable summarised as weighted mean difference and results displayed as forest plots.

Results

575 unique citations were screened with 14 meeting inclusion criteria. 1032 acute episodes (888 patients) of respiratory failure were managed with NPV, with 234 receiving PPV as a comparator. The majority (n=845, 66.7%) were treated for an acute exacerbation of COPD. 417 patients from four studies were included in the meta-analysis.

The effect of NPV on $PaCO_2$ (see figure 2), pH and PaO_2/FiO_2 was similar to PPV with a mean difference -0.39kPa (95% confidence interval (CI): -0.95, 0.18), 0.01 (95% CI: 0.00, 0.02), and -0.16 (95% CI: -1.98, 1.66) respectively.

Of those studies not included in the meta-analysis six showed a statistically significantly increase in PaO_2 with the use of NPV and five showed a statistically significant improvement in $PaCO_2$. Rates of complications were similar with NPV in those studies that compared it to PPV, and NPV appeared to be well tolerated by patients. This systematic review study was limited by a wide range of study designs.

	NPV			PPV			Mean Difference			Mean Difference	
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI	Year	IV, Random, 95% CI	
Corrado 2004	10.3	2.1	22	9.7	3	22	11.7%	0.60 [-0.93, 2.13]	2004	+	
Gorini 2004	9.9	2.1	137	10.1	2.1	79	43.7%	-0.20 [-0.78, 0.38]	2004	•	
Corrado 2009	9.3	1.7	70	10.2	2.2	71	39.3%	-0.90 [-1.55, -0.25]	2009	•	
Chaturvedi 2011	7.3	2.4	8	7.6	2.4	8	5.4%	-0.30 [-2.65, 2.05]	2011	+	
Total (95% CI)			237			180	100.0%	-0.39 [-0.95, 0.18]			
Heterogeneity: $Tau^2 = 0.10$; $Chi^2 = 4.37$, $df = 3$ (P = 0.22); $I^2 = 31\%$ Test for overall effect: Z = 1.34 (P = 0.18)										-100 -50 0 50 100 Favours NPV Favours PPV	

Figure 2: Forest plot comparing effects of NPV and PPV on PaCO2

Conclusions

References

NPV appears to be a safe and effective alternative to PPV in the management of acute exacerbation of COPD Evidence for its use in other forms of respiratory failure is limited but warrants further investigation Soo A, Zuege DJ, Fick GH et al. Describing organ dysfunction in the intensive care unit: a cohort study of 20,000 patients. Critical Care 2019; 23: 186-200.
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