CJEM Journal Club

Apneic oxygenation provides incremental benefit during intubation of patients in the emergency medicine and critical care settings

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Clinical question

Should I use apneic oxygenation when intubating patients in the emergency department to increase periintubation oxygen saturation and first-pass success?

Article chosen

Oliveira JE, Silva L, Cabrera D, Barrionuevo P, et al. Effectiveness of apneic oxygenation during intubation: a systematic review and meta-analysis. *Ann Emerg Med* 2017;70(4):483-94.

Objective

The purpose of this study was to synthesize and appraise the apneic oxygenation literature for its effect on peri-intubation outcomes.

Keywords: critical care, emergency medicine, intubation, meta-analysis

BACKGROUND

In the emergency department (ED), providers use rapid sequence intubation (RSI) as their preferred method of airway control.¹ RSI involves the simultaneous administration of induction and paralytic agents, which induces favorable intubating conditions and minimizes aspiration risk.² On the other hand, patient ventilation is interrupted until mechanical ventilation begins. Successful RSI includes the prevention of hypoxia during this apneic period, because hypoxia increases the risk of cardiac arrest and adverse outcomes.³ One of the primary strategies to avoid hypoxia is preoxygenation prior to intubation.⁴ Preoxygenation of the patient creates an oxygen reservoir for the patient to use during the apneic period. Despite preoxygenation, however, hypoxia can still occur.⁵

Apneic oxygenation has been touted as a method to prolong safe apnea time. The technique is performed by leaving nasal cannula on the patient during the apneic period, while an operator performs laryngoscopy and places the endotracheal tube.⁴ Apneic oxygenation works because alveoli are able to exchange oxygen even without diaphragmatic movements.⁴ When providers give supplemental oxygen to the upper airway, they create a negative pressure gradient between the pharynx and lungs, promoting movement of oxygen into the bloodstream. In the operating room setting, apneic oxygenation prolongs time to desaturation from 3.5 to 5.3 minutes.⁶ Apneic oxygenation has received rapid uptake among the emergency medicine community^{7,8} and has the advantages of being inexpensive, practical, and safe in most patients.⁹

Nevertheless, patients who are intubated in the ED and intensive care unit (ICU) settings are acutely and often critically ill. Benefits of apneic oxygenation in this patient population have been conflicting with some studies showing benefit^{7,10} and others showing no benefit.¹¹⁻¹⁴ Oliveira et al.¹⁵ conducted a systematic review and meta-analysis to provide clarity on the overall state of the evidence.

POPULATION STUDIED

The study authors included studies in both the ED and ICU setting because those intubations occur on an

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urgent or emergent basis in a controlled environment. The authors excluded studies in the operating room or out-of-hospital environments. For studies to be included, patients had to receive oxygen during the apneic period, using any technique. Both randomized controlled trials (RCTs) and observational studies were included because there are few studies available.

STUDY DESIGN

A trial protocol was listed on the PROSPERO website prior to the systematic review. A medical librarian performed a comprehensive search of literature for studies on apneic oxygenation from 2006 until 2016 using Ovid MEDLINE, EMBASE, CENTRAL, and Scopus, without language restriction. The authors searched gray literature, ongoing clinical trials, and reference lists. Two investigators independently performed study selection and data extraction. Included studies were described in detail, and reasons for study exclusion were enumerated. The authors assessed risk of bias using standardized tools. They performed a meta-analysis with randomeffects models using Review Manager (Version 5.3; The Nordic Cochrane Centre, The Cochrane Collaboration, Copenhagen, Denmark). They used odds ratios (OR) and weighted mean differences with 95% confidence intervals (CI) to measure pooled-effect estimates, as well as the I^2 statistic to judge study heterogeneity. The subgroup analysis included risk of bias. The authors adhered to the Preferred Reporting Items for Systematic Reviews and Meta-Analyses (PRISMA) statement.¹⁶

OUTCOME MEASURES

The authors considered the lowest SpO2 peri-intubation, hypoxemia (SpO2 < 93%), severe hypoxemia (SpO2 < 80%), and life-threatening hypoxemia (SpO2 < 70%). First-pass success was also reported. Furthermore, the authors examined some ICU outcomes, including duration of mechanical ventilation, ICU length of stay, and ICU mortality.

RESULTS

The authors identified 1386 potential studies, eventually narrowing it down to 14 for qualitative and 8 for quantitative analysis. The included studies involved 2023 patients, 369 of which were studied in the RCT context. Six of the studies were done in the ED and two in a mixed ED-ICU setting. Five trials were RCTs, with one done in the ED. The ED trial included only 13 patients¹⁷ and was not quantitatively analysed.

The primary sources of heterogeneity were the setting and context for intubation, approaches to preoxygenation, approaches to apneic oxygenation, and proceduralist experience. Outcomes of the metaanalysis are listed in Table 1.

During a subgroup analysis, the authors found that, when they exclusively combined studies at low risk of bias,^{10,11,14} apneic oxygenation had no benefit.

COMMENTARY

We used the AMSTAR 2 checklist,¹⁸ which rated a high overall confidence in the systematic review. The authors did not perform a funnel plot to evaluate publication bias, which, in this case, is appropriate because the number of studies were few, most studies had few patients, and the overall treatment effect is small.¹⁹

The most significant limitation of the meta-analysis is the overall low-to-moderate certainty in the evidence with small sample sizes among the included studies. Only three studies were deemed to be of low risk of bias, none among the ED population. Furthermore,

Favors control	No difference	Favors apneic oxygenation				
None	Severe hypoxemia, SpO2 < 80% (OR 0.86; 95% Cl 0.47 to 1.57, l ² = 44%)	Higher lowest peri-intubation SpO2 (Difference 2.21%, 95% Cl 0.81% to 3.61%, $l^2 = 0\%$				
	Life-threatening hypoxemia, SpO2 < 70% (OR 0.90; 95% Cl 0.52 to 1.55, $l^2 = 2\%$)	First-pass success (OR 1.59, 95% Cl 1.04 to 2.44, I ² = 48%)				
	Duration of mechanical ventilation (Difference 1.42 days; 95% Cl 1.40 to 4.37, $l^2 = 63\%$)	Hypoxemia, SpO2 < 90%-93% (OR 0.66, 95% Cl 0.52 to 0.84, I ² = 0%)				
	ICU mortality (OR 0.82; 95% Cl 0.38 to 1.76, l ² =63%)	Decreased ICU length of stay (2.88 days, 95% Cl 1.40 to 4.37, $l^2 = 0\%$)				

	Apneic Oxygenation Con					Mean Difference			Mean Difference				
Study or Subgroup	Mean	SD	Total	Mean	SD	Total	Weight	IV, Random, 95% CI		IV, Ra	ndom, 9	5% CI	
Simon M 2016	89	18	20	86	11	20	6.3%	3.00 [-6.25, 12.25]				•	
Jaber S 2016	97.4	4.1	23	91.5	12.5	24	15.0%	5.90 [0.63, 11.17]			I-		
ourc'h M 2015	91.5	11.9	62	89.5	10.4	57	20.9%	2.00 [-2.01, 6.01]					
Semler MW 2016	92	11.1	77	90	11.9	73	22.8%	2.00 [-1.69, 5.69]			-		
Caputo N 2017	92	7.2	100	93	7.3	100	35.1%	-1.00 [-3.01, 1.01]		-	-		
Total (95% CI)	282 274					274	100.0%	1.59 [-0.90, 4.08]					
Heterogeneity: Tau ² =	3.54; Chi	$^{2} = 7.59$, df = 4	(P = 0)	11); I ²	= 47%							
Test for overall effect:									-10	-5	0	5	10
									Favours Control Favours ApOx				

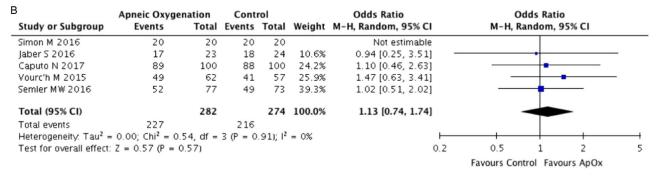


Figure 1. Forest plots of randomized controlled trials from Oliveira et al.'s meta-analysis on periprocedural outcomes, with Caputo et al.'s data added. A) Lowest oxygen saturation (SpO2) peri-intubation. B) First-pass success.

both ICU and ED settings were combined. The ICU studies primarily used high-flow apneic oxygenation devices, which provide additional physiologic benefits compared to the low-flow devices that are available in most EDs. Therefore, the effect size for low-flow devices is likely more modest than reported in the meta-analysis.

Since the publication of the meta-analysis, the ENDAO trial was performed, the first high-quality RCT comparing apneic oxygenation against usual care among ED patients.²⁰ This study enrolled 206 patients and found no difference in the primary outcome of lowest peri-intubation oxygen saturation.

We contacted the lead author of the ENDAO trial, who supplied nonpublished data for lowest periintubation oxygen saturation and first-pass success, which we used in a Forest plot with the other RCTs from the Oliveira et al. meta-analysis (Figure 1). The benefit of apneic oxygenation for improving lowest oxygen saturation loses statistical significance, although moderate study heterogeneity is introduced (difference 1.59%; 95% CI -0.90% to 4.08%, $I^2 = 47\%$) (see Figure 1). The heterogeneity may be because this was the first high-quality RCT in the ED setting, or perhaps because patients had a mean of only 60 seconds of apnea. Similarly, the benefit of first-pass success is not maintained, without a significant change in heterogeneity (OR 1.13; 95% CI 0.74 to 1.74, $I^2 = 0\%$) (see Figure 1). Overall, we caution against over-enthusiasm for apneic oxygenation and agree with the measured conclusion of the meta-analysis, that "apneic oxygenation is a potentially important adjunct for emergency airway management." To prevent hypoxia, ensuring proper preoxygenation is likely more valuable than apneic oxygenation.^{4,9}

CONCLUSION

The use of apneic oxygenation may provide incremental benefit to intubation in the ED. Apneic oxygenation does not replace basics in airway management, such as preoxygenation or technical skill. Further studies may help define which patient groups benefit from apneic oxygenation and be powered to examine potential adverse effects.

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Author contributions: Both authors contributed to the conception, drafting, and revising of the article. Both authors approved the final version to be published and agreed to act as guarantors of the results.

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