

# Comparison of blind intubation via supraglottic airway devices versus standard intubation during different airway emergency scenarios in inexperienced hand

## Randomized, crossover manikin trial

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### Abstract

**Background:** Securing the airway and enabling adequate oxygenation and ventilation is essential during cardiopulmonary resuscitation (CPR). The aim of the study was to evaluate the success rate of blind intubation via the I-Gel and the Air-Q compared with direct laryngoscopy guided endotracheal intubation by inexperienced physician and to measure time to successful intubation.

**Methods:** The study was designed as a randomized, cross-over simulation study. A total of 134 physicians, from specialties other than Anesthesia or Emergency Medicine, who considered themselves skilled in endotracheal intubation but who have never used any kind of supraglottic airway device performed blind intubation via the I-Gel and Air-Q and direct laryngoscopy guided endotracheal intubation in 3 randomized scenarios: normal airway without chest compression during intubation attempt; normal airway with continuous chest compression during intubation attempt; difficult airway with continuous chest compression.

**Results:** Scenario A: Success rate with initial intubation attempt was 72% for endotracheal intubation, 75% in Air-Q, and 81% in I-Gel. Time to endotracheal intubation and ease of intubation was comparable with all 3 airway devices used. Scenario B: Success rate with the initial intubation attempt was 42% for endotracheal intubation, compared with 75% in Air-Q and 80% in I-Gel. Time for endotracheal intubation was significantly prolonged in endotracheal intubation (42 seconds, 35–49), compared with Air-Q (21 seconds, 18–32) and I-Gel (19 seconds, 17–27). Scenario C: The success rate with the initial intubation attempt was 23% in endotracheal intubation, compared with 65% in Air-Q and 74% in I-Gel. Time to intubation was comparable with both supraglottic airway devices (20 vs 22 seconds) but was significantly shorter compared with endotracheal intubation (50 seconds,  $P < .001$ ).

**Conclusions:** Less to moderately experienced providers are able to perform endotracheal intubation in easy airways but fail during ongoing chest compressions and simulated difficult airway. Consequently, less to moderately experienced providers should refrain from endotracheal intubation during ongoing chest compressions during CPR and in expected difficult airways. Supraglottic airway devices are reliable alternatives and blind intubation through these devices is a valuable airway management strategy.

**Abbreviations:** CPR = cardiopulmonary resuscitation, ID = internal diameter, IQR = interquartile range, IRB = Institutional Review Board, SAD = supraglottic airway device, VAS = Visual Analog Scale.

**Keywords:** blind intubation, cardiopulmonary resuscitation, endotracheal intubation, medical simulation, physician

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Ethical approval: The study protocol was approved by the Institutional Review Board of the Polish Society of Disaster Medicine (Approval no. 43.12.2017.IRB).

Informed consent: Informed consent was obtained from all individual participants included in the study.

The authors have no conflicts of interest to disclose.

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## 1. Introduction

More than 350,000 people suffer an out-of-hospital cardiac arrest in the United States annually and early initiation of advanced cardio-pulmonary resuscitation (CPR) is the key to favorable outcomes.<sup>[1–4]</sup> High-quality chest compressions are the most single important component of CPR. Even minor interruptions of chest compressions during CPR, represented by the so-called hands-off time, reduces systemic blood flow and increases morbidity and mortality.<sup>[5]</sup> Consequently, international CPR guidelines advocate to keep the hands-off time as short as possible.<sup>[2,6]</sup>

Securing the airway and enabling adequate oxygenation and ventilation is essential during CPR, and although controversially discussed, endotracheal intubation is considered the airway of choice, at least by experienced providers.<sup>[7,8]</sup> Endotracheal intubation requires regular training and high level of individual skills and experience.<sup>[9–11]</sup> Supraglottic airway devices are less invasive, are easier to place, and have previously been reported to decrease the time needed to intubate, even by unexperienced providers and during ongoing chest compression during CPR.<sup>[12–14]</sup> Supraglottic airway devices do not prevent pulmonary aspiration of gastric content, a common and serious clinical concern.<sup>[15]</sup> However, once the airway is secured by a supraglottic airway and ventilation is restored, the trachea can be intubated through some types of supraglottic airway devices. Intubation through these devices can be performed using a fiberoptic bronchoscope or blindly, meaning that the tube is inserted through the supraglottic airway device without direct visualization of the airway.<sup>[16,17]</sup> Success rate of blind intubation ranges between 15% and 97%, mostly depending on the type of the supraglottic airway device used, patient characteristics and operator skills.<sup>[18,19]</sup>

The I-Gel and the Air-Q are recently introduced supraglottic airway devices designed to permit fiberoptic-guided and blind endotracheal intubation.<sup>[19,20]</sup> Whether these airway devices can be used successfully by non-experienced providers like physicians within acceptable time frame remains currently unclear. We, therefore, performed this manikin study to evaluate the success rate of blind intubation via the I-Gel and the Air-Q compared with direct laryngoscopy guided endotracheal intubation by unexperienced physician. Additionally, we investigated the number of intubation attempts and the time to successful intubation. In particular, we tested the hypothesis, that success rates of blind-intubation via I-Gel and Air-Q are non-inferior to direct laryngoscopy guided endotracheal intubation.

## 2. Material and methods

The study was designed as a randomized, cross-over simulation study. The study protocol was approved by the Institutional Review Board of the Polish Society of Disaster Medicine (Approval no. 43.12.2017.IRB) and was performed at the International Institute of Rescue Research and Education (Warsaw, Poland) between December 2017 and April 2018.

### 2.1. Participants

After informed written consent was obtained, 134 physicians voluntarily participated in the study (Table 1). All participating physicians considered themselves as skilled in endotracheal intubation, but never used any kind of supraglottic airway device. Physicians were excluded in case of pregnancy or any physical injury, affecting probability to perform endotracheal intubation.

**Table 1**

#### Baseline characteristics.

Characteristics	Subjects (n, 134)
Sex, n (%)	
Male,	55 (41%)
Female	79 (59%)
Age, y	35 [IQR; 29–41]
Working experience, y	8.5 [IQR; 3.0–12.5]

Data expressed in percentage or median and interquartile range (IQR).

Anesthesiologists or emergency physicians were excluded from the study.

### 2.2. Protocol

Prior to the study, participating physicians received a standardized audio-visual lecture lasting 30 minutes covering fundamental airway anatomy, physiology, and pathophysiology, and common methods of airway management including several supraglottic airway devices used for blind intubation. This lecture was followed by a 30 minutes lasting hands-on training. Blind intubation via the I-Gel and Air-Q and direct laryngoscopy guided endotracheal intubation were demonstrated and physicians were able to practice the devices on an airway manikin. Endotracheal intubation was performed by using a laryngoscope with a Macintosh's blade size 3 (HEINE Optotechnik GmbH & Co. KG, Herrsching, Germany). Blind intubation was performed using an I-Gel supraglottic airway size 4 (Intersurgical Ltd., Wokingham, UK), and Air-Q supraglottic airway device size 4.5 (Mercury Medical, Florida) (Fig. 1). A standard single lumen endotracheal tube with 7.0 mm internal diameter (ID) was used for all intubation attempts. Both, the endotracheal tube and respective supraglottic airway devices were thoroughly lubricated with a lubricant recommended by the producer of the manikin.

A Resusci Anne QCPR (Laerdal, Stavanger, Norway) simulator was placed on the floor in a well-lit room. Subjects were asked to perform airway management with all 3 airway devices in 3 separate scenarios:

- (1) Scenario A: normal airway without chest compression during intubation attempt.
- (2) Scenario B: normal airway with continuous chest compression during intubation attempt.
- (3) Scenario C: difficult airway with continuous chest compression. The simulator control software generated a Mallampati grade 3 through tongue edema.

The physicians were asked to perform airway management as fast as possible. Chest compressions were performed at a rate of 100 compressions per minute, and compression depth of 5 cm by using the CORPULS CPR (GS Elektromedizinische Geräte G. Stemple GmbH, Kaufering, Germany) chest compression system.

Both, the intubation methods and scenarios sequence were randomized by using a Research Randomizer program (randomizer.org) (Fig. 2). Physicians performed airway management with all 3 devices in each scenario. Afterwards, physicians rested for about 10 minutes, before they were asked to handle the second, respectively the third airway scenario.

### 2.3. Data collection

Primary endpoint was overall rate of successful intubation, defined as successful placement of an endotracheal tube in the

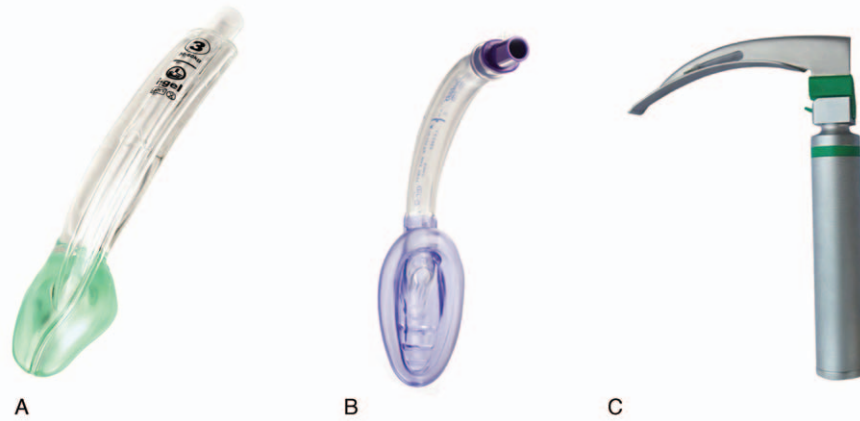


Figure 1. Devices used in the study: (A) iGEL mas; (B) Air-Q maks; (C) Macintosh laryngoscope.

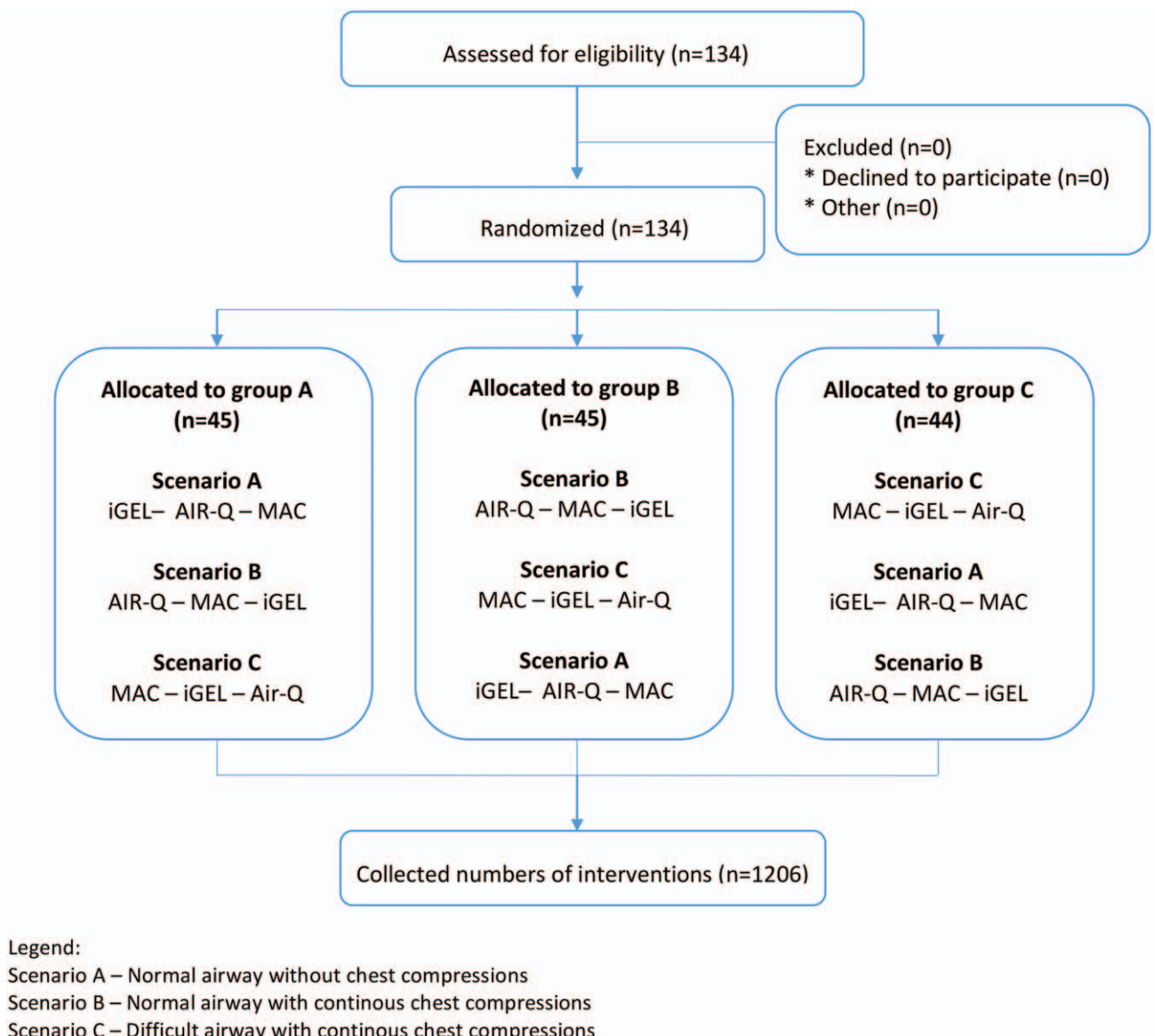


Figure 2. Randomization flow chart.

simulators trachea. Correct placement was confirmed by successful ventilation of the manikin's lungs with an anesthesia bag. Intubation attempts lasting longer than 120 seconds, or unrecognized esophageal intubation were defined as failure.

Secondary endpoints included: rate of successful intubation with 1 single intubation attempt. Time to successful supraglottic airway device (SAD) placement, defined as the time from the initial insertion of the SADs in the manikin's mouth until stabilization of the supraglottic airway device. Time to successful endotracheal intubation was defined as time from the initial insertion of the laryngoscope or SAD in the manikin's mouth to confirmation of lung inflation with positive pressure breaths through the endotracheal tube. Time to successful blind intubation was defined as time from the initial insertion of the Supraglottic airways (SGAs) in the manikin's mouth to confirmation of lung inflation with positive pressure breaths through the tracheal tube.

Once physicians performed all 3 intubations during a scenario, physicians were asked to rate difficulty of intubation of each device using a visual analog scale (VAS) ranging from 1 (extremely easy) to 100 (extremely difficult).

#### 2.4. Statistical analysis

The statistical package Statistica 13.0 EN (Tibco, Tulsa, OK) was used for all statistical analysis. The type I error was set to  $\alpha = 0.05$ . To account for multiple testing, critical was adjusted using the Bonferroni method ( $\alpha = 0.0125$ ). To detect possible differences in procedure time and success rate of endotracheal intubation, the analysis of variance (ANOVA) Kruskal Wallis was used. The participants' subjective opinions were compared with the use of the Stuart–Maxwell test. Data were presented as medians and interquartile range (IQR), or percentages (%). The  $\alpha$ -error level for all analyses was set at  $P < .05$ .

### 3. Results

One hundred and thirty-four physicians (79 women) with a median age 35 [IQR; 29–41] years and work experience of 8.5 [3.0–12.5] years participated in this study. All physicians declared the ability to perform endotracheal intubation using direct laryngoscopy but had not used any form of any supraglottic airway device beforehand.

#### 3.1. Scenario A: normal airway without chest compression

Success rate with initial intubation attempt was 72% for endotracheal intubation, 75% in Air-Q, and 81% in I-Gel (Table 2). The final success rate within 3 intubation attempts was 95% in endotracheal intubation, 99% in Air-Q, and 100% in I-Gel (Table 2). Time to endotracheal intubation and ease of intubation was comparable with all 3 airway devices used (Figs. 3A and 4A, respectively).

#### 3.2. Scenario B: normal airway with continuous chest compression

Success rate with the initial intubation attempt was 42% for endotracheal intubation, compared with 75% in Air-Q and 80% in I-Gel (Table 2). Overall success rate was 42% for endotracheal intubation, 95% in Air-Q, and 100% in I-Gel (Table 2). Time for endotracheal intubation was significantly prolonged in endotracheal intubation (42 seconds, 35–49), compared with Air-Q (21 seconds, 18–32) and I-Gel (19 seconds, 17–27) (Fig. 3B).

Endotracheal intubation was also rated to be significantly more difficult compared with both supraglottic airway devices (69 vs 38 for Air-Q and 36 for I-Gel; Fig. 4B).

#### 3.3. Scenario C: difficult airway with continuous chest compression

During this scenario, the success rate with the initial intubation attempt was 23% in endotracheal intubation, compared with 65% in Air-Q and 74% in I-Gel (Table 2). Overall success rate was significantly higher in I-Gel compared with Air-Q (95% vs 86%,  $P = .003$ ). Both supraglottic airway devices offered significant better overall success rate, compared with endotracheal intubation (95% vs 86% vs 31%,  $P < .001$ ). Time to intubation was comparable with both supraglottic airway devices (20 vs 22 seconds) but was significantly shorter compared with endotracheal intubation (50 seconds,  $P < .001$ ; Fig. 3C). Again, both supraglottic airway devices were rated comparable (36 vs 40), but significantly easier compared with endotracheal intubation (80,  $P < .001$ ; Fig. 4C).

### 4. Discussion

Results of this manikin study indicate, that relatively skilled and experienced providers were able to intubate a manikin with an easy regular airway. However, these providers failed during ongoing chest compressions or with a simulated difficult airway (success rate of 42% and 31%, respectively). Contrary, placement of supraglottic airway devices was easy, and success rate of blind intubation via these supraglottic airway devices were between 86% and 100% in all intubation attempts and all 3 airway scenarios.

The I-Gel as well as the Air-Q devices have been previously reported to be easy to insert and providing adequate ventilation in several clinical scenarios.<sup>[20–23]</sup> Placement of both supraglottic airway devices were possible in all intubation attempts in all 3 intubation scenarios. The corresponding success rate of 100% is in line with several previous reports.<sup>[19,20,24]</sup> Placement of supraglottic airway devices are easy to place, quickly inserted, and nearly always provide adequate ventilation.<sup>[14,20,25,26]</sup> However, supraglottic airway devices are generally limited due their ability to prevent pulmonary aspiration of gastric content, a significant clinical concern in patients undergoing both, regular anesthesia for a surgical procedure or CPR.<sup>[8,27,28]</sup>

Once the airway is secured with a supraglottic airway device, several devices including the I-Gel and the Air-Q offer the possibility to intubate the trachea through the supraglottic airway device. This can be done without direct visualization of the airway, the so called blind intubation, or guided by using a fiberoptic bronchoscope. Especially in the emergency setting, blind intubation might be more reasonable, as it is much less time consuming and a fiberoptic might not be regularly available.

Success rate of blind intubation via the supraglottic airway devices with the initial intubation attempt differed between the 2 different devices used in this study and within the 3 airway scenarios. As expected, success rate was highest in scenario 1 with an easy airway anatomy (81% in I-Gel and 75% in Air-Q), and somewhat decreased during ongoing chest compression (scenario 2, 80% in I-Gel and 75% in Air-Q) and difficult airway (scenario 3, 74% in I-Gel and 64% in Air-Q). Overall, the success rate with initial intubation attempt is well within the previously reported range of success rates ranging between 57% and 70%.<sup>[18,20,29]</sup> The obvious reason is the use of airway manikins in this study instead of patients undergoing real-world intubations.

**Table 2****Study outcomes by supraglottic airway devices with unexperienced physician (N=134).**

Outcome	iGEL	Air-Q	MAC	P-value
Scenario A: normal airway without chest compression				
Success of intubation attempts (%)				iGEL vs Air-Q = .043
1st	109 (81.3%)	100 (74.6%)	96 (71.7%)	iGEL vs MAC < .001
2nd	24 (18.0%)	21 (15.7%)	27 (20.1%)	Others: NS
3rd	1 (0.7%)	11 (9.7%)	4 (3.0%)	
Overall intubation success rate (%)	134 (100%)	132 (98.5%)	127 (94.8%)	iGEL vs MAC = .038
				Others: NS
Time to supraglottic airway device placement, s	8.5 [7.5–12]	14.5 [11–17]		iGEL vs Air-Q = .011
Time to endotracheal intubation, s	19 [16–25]	23 [16.5–31]	24.5 [21–32]	iGEL vs Air-Q = .034
				iGEL vs MAC = .007
				Air-Q vs MAC = .048
Ease of use, 1–100 scale	35 [27–37]	36 [27.5–40]	42 [36–47]	iGEL vs MAC = .046
				Others: NS
Scenario B: normal airway with continuous chest compression				
Success of intubation attempts (%)				iGEL vs Air-Q = .032
1st	107 (79.9%)	100 (74.6%)	56 (41.8%)	iGEL vs MAC < .001
2nd	25 (18.6%)	21 (15.7%)	0 (0%)	Air-Q vs MAC < .001
3rd	2 (1.5%)	6 (4.5%)	0 (0%)	
Overall intubation success rate (%)	134 (100%)	127 (94.8%)	56 (41.8%)	iGEL vs Air-Q = .014
				iGEL vs MAC < .001
				Air-Q vs MAC < .001
Time to supraglottic airway device placement, s	9 [8–12]	15.5 [12–18]		iGEL vs MAC = .001
Time to endotracheal intubation, s	18.5 [17–27]	20.5 [18–32]	41.5 [35–49]	iGEL vs Air-Q = .042
				iGEL vs MAC < .001
				Air-Q vs MAC < .001
Ease of use, 1–100 scale	36 [26–38]	38 [27–44]	69 [51–77]	iGEL vs MAC < .001
				Air-Q vs MAC < .001
				Others: NS
Scenario C: difficult airway with continuous chest compression				
Success of intubation attempts [%]				iGEL vs Air-Q = .007
1st	99 (73.9%)	87 (64.9%)	31 (23.1%)	iGEL vs MAC < .001
2nd	27 (20.2%)	20 (15.0%)	7 (6.0%)	Air-Q vs MAC < .001
3rd	1 (0.7%)	8 (5.9%)	3 (2.2%)	
Overall intubation success rate [%]	127 (94.8%)	115 (85.8%)	42 (31.3%)	iGEL vs Air-Q = .003
				iGEL vs MAC < .001
				Air-Q vs MAC < .001
Time to supraglottic airway device placement, s	9.5 [7.5–13.5]	15.5 [13–18.5]		iGEL vs Air-Q = .005
Time to endotracheal intubation, s	19.5 [17.5–27.5]	22 [19.5–35]	49.5 [44–67]	iGEL vs Air-Q = .019
				iGEL vs MAC < .001
				Air-Q vs MAC < .001
Ease of use, 1–100 scale	36 [27–39]	40 [29–46]	80 [63–84]	iGEL vs Air-Q = .049
				iGEL vs MAC < .001
				Air-Q vs MAC < .001

Data expressed in percentage (%) or median and interquartile range [25<sup>th</sup> percentile–75<sup>th</sup> percentile].

Previous studies reported an overall success rate of blind intubation via a supraglottic airway devices, ranging from 24% to 97%.<sup>[19,20]</sup> Results of our study indicate, that blind intubation via a supraglottic airway devices with a maximum of 3 intubation attempt is highly success full in easy airways (scenario 1, 100% in I-Gel and 99% in Air-Q) and is relatively not altered by ongoing chest compression (scenario 2, 100% in I-Gel and 95% in Air-Q). Blind intubation in difficult airways is less successful (scenario 3, 95% in I-Gel and 86% in Air-Q). In general, blind intubation is often considered an airway technique, potentially associated with serious complications including airway trauma and trachea rupture. Although, complications were not investigated in this study, a most recent large clinical trial did not report any major complication.<sup>[20]</sup> Given the fact, that success rate of insertion of the supraglottic airway devices was promising and even success rate of blind intubation was acceptable high, less experienced providers

should refrain from endotracheal intubation and should place a supraglottic airway device instead.

Ease of intubation, as determined by subjective evaluation of the physicians participating in this study, was comparable with both supraglottic airway devices throughout all 3 airway scenarios. In contrast, direct laryngoscopy guided endotracheal intubation was rated much more difficult.

Participating physicians classified themselves as experienced with direct laryngoscopy guided endotracheal intubation however anesthesiologists or emergency physicians were excluded from the study. In Poland endotracheal intubation is performed mainly by anesthesiologists and emergency physicians. Physicians of other specialties or specialists in other fields of medicine are not experienced in endotracheal intubation but in several clinical situation may decide to perform endotracheal intubation especially in sudden cardiac arrest. This is especially important, as success rate in the easy airway scenario was 95%, but

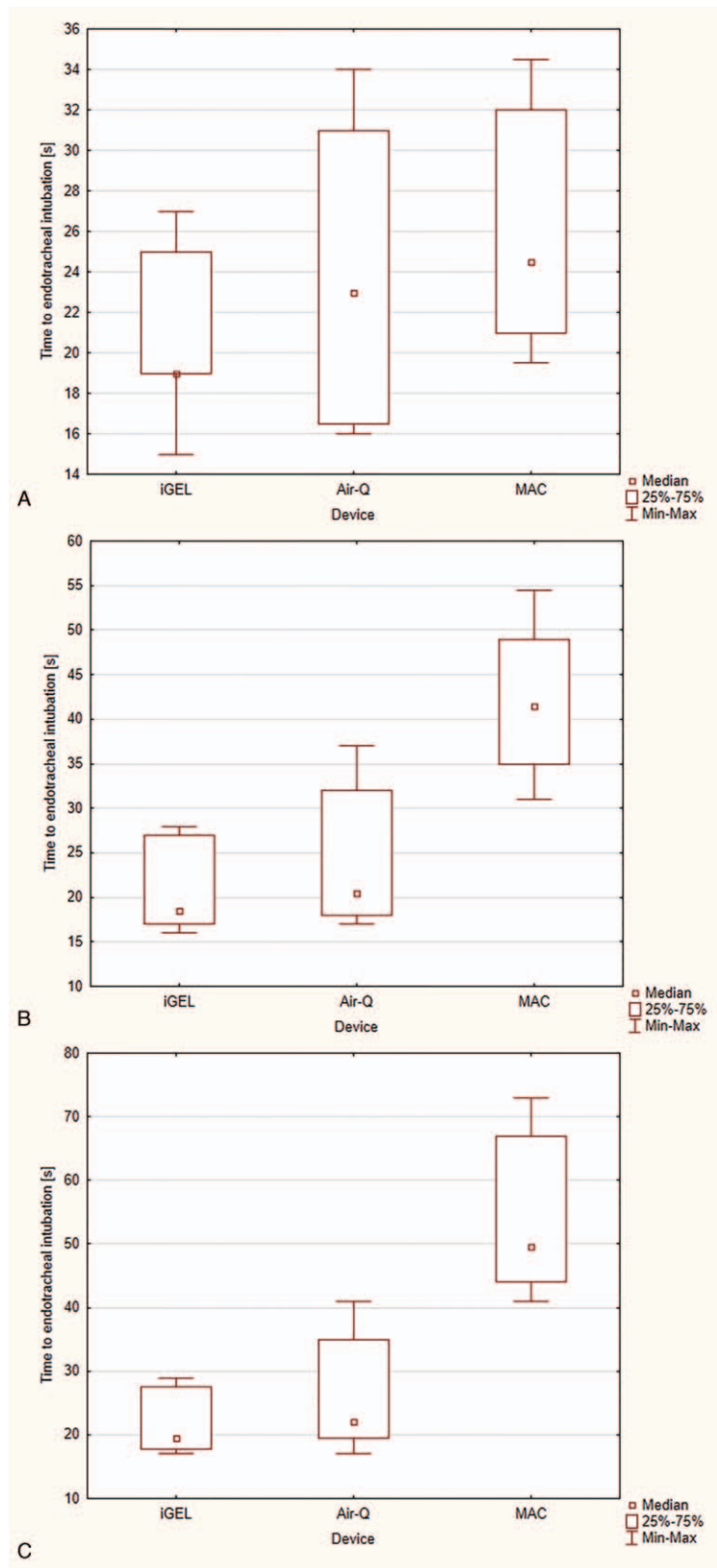


Figure 3. Time to endotracheal intubation during: (A) Scenario A; (B) Scenario B; (C) Scenario C).

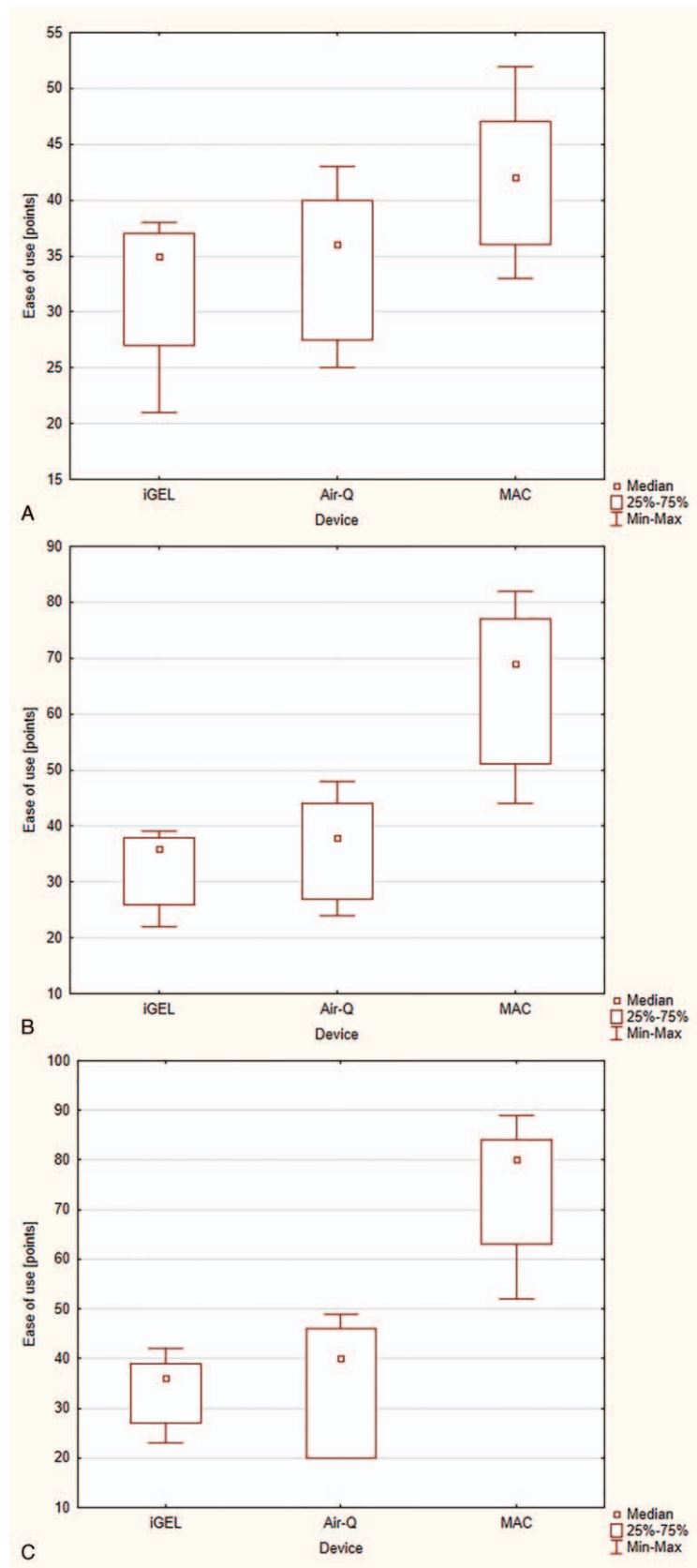


Figure 4. Ease of use during: (A) Scenario A; (B) Scenario B; (C) Scenario C.

significantly dropped during scenario 2 and 3 (42% and 24%, respectively). This finding is significant, as success rate during easy airway indicates, that physicians were truly at least moderately experienced with this technic. Success rates during chest compressions or difficult airway scenario confirm previous findings, that moderately physicians should refrain from endotracheal intubation during CPR and should consider a supraglottic airway devices instead.<sup>[12,13,24,30]</sup>

Our study has several limitations. First, this study was performed in airway manikins, which might not mimic real conditions during real CPR. However, airway manikins are considered reliable airway teaching tools, and based on ethical considerations, cross-over studies like that might not be feasible in real patients undergoing real CPR. Nevertheless, results of airway manikin's studies are generally considered reliable.<sup>[31,32]</sup> Second, our participating physicians considered themselves are skilled in endotracheal intubation. We were not able to assess the number of endotracheal intubations in real patients. Consequently, this opens the possibility for self-overestimating of personal skills and experience.

## 5. Conclusions

In summary, our study indicates, that less to moderately experienced providers are able to perform endotracheal intubation in easy airways but fail during ongoing chest compressions and simulated difficult airway. Consequently, less to moderately experienced providers should refrain from endotracheal intubation during ongoing chest compressions during CPR and in expected difficult airways. Supraglottic airway devices are reliable alternatives and blind intubation through these devices is a valuable airway management strategy.

## Author contributions

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**Formal analysis:** Andrzej Bielski, Eva Rivas, Kurt Ruetzler, Oliver Robak, Lukasz Szarpak.

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