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I-Gel and LMA classic in the hands of novice: A comparative study



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ABSTRACT

Background: Supraglottic airway devices (SADs) such as LMA classic (cLMA), I-Gel, etc. are indispensable tool for the anaesthesiologists experienced in airway management. But studies evaluating the performance of these devices in the hands of unskilled personnel are scarce. Aims and Objective: To determine the procedure time and the proportion of patients having successful placement of I-gel and LMA classic by first-year Post Graduate Trainees (PGTs) of Anaesthesiology who tried insertion of those devices after a short training in mannequins without any hands-on training regarding placement of the devices in human. Materials and Methods: After getting Institute's Ethics Committee's approval for this interventional study, forty adult patients, posted for short surgical or gynaecological surgery, were randomly allocated in to two groups to have placement of either I-Gel (group 'I', n=20) or cLMA (group 'C', n = 20) by first-year PGTs. The procedure time (Primary outcome) i.e. the time taken for successful placement of either device was determined and compared. A standard technique of anaesthesia was followed in every patient. Any adverse event such as sore throat, odynophagia, blood stain on the device, etc. was also recorded. Results: All patients were comparable with respect to demographic data and Mallampati scores. The mean procedure time (seconds) was considerably lower in I-Gel group compared with cLMA (63.3 \pm 57.2 versus 163.0 \pm 158.3, respectively, *P* value < 0.001). The incidence of successful placement at first attempt was significantly higher for I-Gel group. The incidence of adverse events was comparable. Conclusion: Procedure time for I-Gel insertion is significantly shorter than with LMA Classic along with a higher success rate with first attempt for the former. I-Gel may be a better alternative as airway device for the unskilled anaesthesiologist.

Key words: Adult; Anesthesiology; Airway Management; Critical Care; Humans; Intubation, Intratracheal; Outcome Assessment, Health Care; Manikins; Students

INTRODUCTION

Nowadays, the supraglottic airway devices (SADs) are widely used as an intermediate device between facemask and endotracheal tube for administration of general anaesthesia in short surgical procedures. The second generation SADs has achieved a solid and proven place as rescue device for managing difficult airways^{1,2} and as conduit ^{3,4} to aid endotracheal intubation. These are also frequently used as resuscitation device to secure the airway especially in the out-of-hospital emergencies. Availability of experienced anaesthesiologist or and skilled airway manager cannot be ensured every time at all such fields for securing and maintaining airway by intubation. The doctors and paramedics working outside the discipline of anaesthesia, critical care, trauma care and emergency medicine often face difficulties to acquire the intubation skill and subsequently to maintain that. About 25% of endotracheal tubes placed by paramedical staffs in emergency department were found to be improper.⁵ SADs

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can be used in a safer way by novice occasional airway managers, such as medical students, unskilled nurses, allied health care professionals and physicians working outside the field of anaesthesiology. In the initial phase, the Post Graduate Trainees in anaesthesiology can also be considered as novice regarding placement of SADs.

The performance of I-Gel has been compared with the LMA Proseal⁶⁻⁸ or with LMA Supreme.⁹ Several studies have been reported comparing I-Gel and LMA classic (cLMA) in adult,^{1,10-16} in child¹⁷⁻¹⁹ and in mixed age group²⁰ population. Some of these studies report the comparative performance of I-Gel and cLMA in the hands of experienced or skilled anaesthesiologists.^{1,10-12,17-19} The performance of only I-Gel in both mannequins and anaesthetized patients was reported in a single study²¹ when the device was used by novice users who all were unfamiliar with the use of I-Gel, cLMA or other SADs.

In a few studies the procedure time and success rate of both I-Gel and LMA have been determined and compared during the use of these devices by novice.²²⁻²⁷ Some of these studies have evaluated the performance of novice users during their insertion of devices in the mannequin²²⁻²⁴ or in the clinical scenarios.²⁵⁻²⁷ This paucity in clinical studies evaluating the performance of I-Gel and cLMA in the hands of novice²⁵⁻²⁷ was the stimulus for designing the present study. Hence, the present study was designed to compare the procedure time (Primary outcome) during placement of I-Gel or LMA Classic in the hands of firstyear Post Graduate Trainees (PGTs) of Anaesthesiology who had no previous experience in placing these devices in humans. Other outcome measures were to determine the success rate of placement of either device within single attempt. Additionally, the incidence of adverse events with the use of either device was noted and compared.

MATERIALS AND METHODS

The present study was designed to compare the procedure time regarding the placement of I-Gel and LMA classic (cLMA) by a first-year PGT of Anaesthesiology. This interventional, single-blind study was performed in a tertiary care center, a Government Medical College. After receiving permission from the Institute's Ethics Committee, 40 patients aged 18-50 years of ASA I and II, planned for short surgical and Gynaecological procedures (surgical procedures not exceeding one hour) were recruited for this study. The patients and their family members were explained about the procedure and the associated risk and benefit, in their own language. Patients with airway status of Mallampati Grade 1 and 2 were included for the present study. The patients who had anticipated difficult intubation status during pre-anaesthetic evaluation were excluded from the present study. Similarly, the patients having an increased chance for gastric regurgitation and aspiration were also excluded. Thus, the patients with simpler airway were selected owing to our nature of study evaluating the performance of first-year residents.

For this interventional study, 40 patients were selected based on inclusion and exclusion criteria and informed consent was taken from each of them. The group allocation was performed after the induction of anaesthesia. There were 40 sealed envelopes each containing a piece of paper marked as 'T' or 'C' (20 papers marked as 'T' and 20 as 'C'). After induction of anaesthesia an envelope was randomly picked up and opened. The alphabet displayed in the paper slip corresponded to the group allocation of the patient. The patients were thus randomly divided into two groups, Group 'T' where the patients received I-Gel and Group 'C' where the patients received cLMA. The used envelope and the paper were discarded then.

In this interventional study the procedure time for successful placement of I-Gel or cLMA was determined and compared (Primary outcome). The success rate of insertion of either device in a single attempt and the adverse events occurring in either case was also compared. Correct positioning was confirmed by bilateral chest rise, bilateral equal breath sounds, 'square-head' capnograph and peripheral arterial oxygen saturation (SpO₂) more than 95%. The case was considered 'successful' if the device was inserted in a single or two attempts. If it could not be done within two attempts it was taken as a 'failure' and tracheal intubation was done. The procedure time was calculated as the time taken from 'picking up the device to the appearance of square-head capnograph'. The adverse events included sore throat, difficulty in swallowing (dysphagia), pain on swallowing (odynophagia) and blood on devices.

In the present study, I-Gel and cLMA were inserted in the respective groups by a first-year resident without any previous exposure to use of these airway devices. The resident placed the device under the close supervision of a skilled anaesthesiologist. All the first-year residents received formal teaching and demonstration about the procedure previously with the help of a mannequin. They individually practiced 30 insertions of the devices on mannequins. The residents were allowed to perform in the present study only after the facilitator had been satisfied with their performance on mannequins. Subsequently the residents observed seniors performing the procedure on patients. However, the residents did not have any hands-on experience on patients before the start of the study. After receiving any patient in operating room (OR), monitor was attached and the baseline vitals such as blood pressure, SpO₂ and ECG were recorded. Subsequently, inj. fentanyl (2 microgram/kg) and midazolam (0.03 mg/kg) were administered via intravenous (i.v.) route as premedication. The patient was induced with inj. propofol (2 mg/kg) i.v. and adequate anaesthesia was determined by no response to verbal commands. Once deep plane of anaesthesia was achieved, the I-Gel or cLMA was placed according to group assignment. Anaesthesia was maintained using nitrous oxide, oxygen and sevoflurane. Throughout the procedure the hemodynamic parameters such as heart rate, systolic and diastolic blood pressure, and SpO2 were monitored. Once the procedure was over, the device was taken out after oxygenation and proper recovery. The procedure time i.e. the time required for successful insertion of each device, the proportion of patients with successful placement within single attempt and the adverse events -- all were recorded and analyzed subsequently.

Based on previous study²⁶ and also complying with the expert opinion, a five second difference in procedure time was considered to be clinically significant. This was the effect size. For calculation of sample size by 'comparing two means', the present researcher consulted the methods and formulas as mentioned in articles of Das S et al.,²⁸ and Hazra A et al.²⁹ Setting the power of the study at 80% and permitting a type I error at 5% (alpha value at 0.05) the sample size was calculated to be 16. Assuming a 20% dropout possibility, 20 patients were recruited in each group. Hence, a total of 40 patients were recruited for the study.

The data was decoded, tabulated and analyzed with the help of a biostatistician. The data was entered in Microsoft Excel spreadsheet and analysis was done using Statistical Package for Social Studies (SPSS) version 21.0. Categorical variables were presented in number and percentage (%) while continuous variables were expressed as mean \pm SD and median. Normality of data was tested by Kolmogorov-Smirnov test. If the normality was rejected then non parametric test was used. Quantitative variables were compared using independent t test/Mann Whitney test (when the data sets were not distributed normally between the two groups). Qualitative variables were compared using Chi-square test/Fischer's Exact test. A *P* value of <0.05 was considered statistically significant.

RESULTS

The study spanned over one year approximately, from May 2019 to March 2020. Data from all 40 patients was available for analysis (Figure 1). All data were found to have normal distribution except two quantitative variables, the

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body mass index (kg/m^2) and procedure time (in seconds) which were found to have a non-normal distribution. Hence, Mann Whitney test (non-parametric) was used for comparison of body mass index (Table-1) and procedure time. (See later, table-3). In group I the patients have received I-Gel and in group C the patients have received LMA classic.

There was no significant difference between the patients of the two groups regarding the demographic parameters (age, gender, BMI and ASA status). However, the majority of the patients were females. This was because almost half of the patients were recruited from Gynecology department. The Mallampati scores were comparable between the groups and thus it can be said that the level of airway difficulty was comparable between the groups.

The variable body mass index (kg/m^2) was not normally distributed. Thus, non-parametric test was used for the comparison. No significant difference was seen in body mass index (kg/m^2) between I-Gel and cLMA (*P* value >0.05).



Figure 1: Flow diagram showing patient selection, randomization and lost to follow up

Table 1: Domographic data					
Table 1. Demographic data					
Parameters	Group I (n=20)	Group C (n=20)	P value		
Age	27.1 ± 8.6	32.8 ± 9.8	0.058		
Gender* (M/F)	4/16	7/13	0.48		
ASA-PS* (I/II)	18/2	17/3	1		
BMI (kg/m ²)	22 (20-24.2)	22 (21-23.3)	0.753		
[Median (IQR)]					

The gender and ASA-PS were analysed according to *Fisher's Exact test. Age was tested with Student's *t*-test. All were found comparable. BMI is expressed as median (interquartile range). Mallampati score were comparable between the groups (Table-2).

As the data of procedure times was having a non-normal distribution, the Median (IQR) value of procedure time (seconds) in cLMA was found to be significantly higher as compared with that of I-Gel (Table-3).

The variable duration of insertion (seconds) was not normally distributed. Thus, non-parametric test was used for the comparison. (Table 3) Significant difference was seen in procedure time (seconds) between I-Gel and cLMA (P value <0.05). Median (IQR) value of procedure time (seconds) in cLMA was 100.5 (89-117.5) which was found to be considerably higher as compared with that of I-Gel [47(43.0-57.8)]. The 'box and whisker' plot (Figure 2) depicts the distribution of duration of insertion (seconds) in the 2 groups. The middle horizontal line represents the median duration of insertion (seconds), the upper and lower bounds of the box represent the 75thand the 25thcentile of duration of insertion (seconds) respectively, and the upper and lower extent of the whiskers represent the maximum



Figure 2: Box-whisker plot showing procedure times (in seconds) for placement of devices

Table 2: Distribution of Mallampati scoresbetween the two groups					
Mallampati score	Group I (n=20)	Group C (n=20)	Total	P value	
1	9 (45%)	12 (60%)	21 (52.5%)		
2	11 (55%)	8 (40%)	19 (47.5%)	0.342	
Iotal	20 (100%)	20 (100%)	40 (100%)		

Chi square test

Table 3: Procedure times (in seconds)					
ProcedureGroup IGroup C (n=20)P valutime (seconds)(n=20)					
Mean ± SD	63.3 ± 57.2	163.0 ± 158.3			
Median (IQR)	47 (42.9-57.8)	100.5 (89-117.5)	<0.0001		
Range	36-300	59-660			
Test performed. Mann Whitney test IOR interquartile range					

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and the minimum duration of insertion (seconds) in each of the groups.

Significant difference was seen in the distribution of number of attempts between I-Gel and LMA Classic (*P* value < 0.05). Placement of the device was possible within single attempt in considerably higher number of patients in I-Gel group compared with LMA group (95% versus 55%, respectively). On the other hand, 2nd attempt was required in lesser number of patients in I-Gel group compared with LMA group (5% versus 45%, respectively). It is shown in table 4.

Apparently, lesser number of patients had adverse events in I-Gel group compared with cLMA group. Blood staining on the device was found in majority of patients (15% in I-Gel and 25% in LMA classic). Other adverse events were dysphagia (5% in I-Gel versus 15% in cLMA), odynophagia (5% in I-Gel versus 15% in cLMA) and sore throat in very few patients (0% in I-Gel versus 10% of patients in cLMA group). All such adverse events were found comparable (*P* value>0.05) between the groups, when analyzed (Table 5).

DISCUSSION

This interventional study has compared the performance of one second generation SAD (I-Gel) with one first generation SAD (LMA classic) while those have been introduced in patients by novice performers. The present study finds that the procedure time for I-Gel insertion is considerably shorter than for the time for placement of cLMA in the hands of beginners. The procedure time for I-Gel was almost 2.5 times less than that of cLMA (63 seconds versus 163 seconds). This observation of the present study is in line with that of Kwak DI et al.,²⁴ where shorter procedure times were required using I-Gel compare with cLMA in mannequins with normal airway as well as difficult airway settings while the insertions were performed by novice i.e. with no prior experience in using SADs (Table 6). In the present study, insertion of cLMA required second attempts in many patients and thus further inflated the procedure time in the LMA group. Procedure time for I-Gel was also found to be almost a half to that of cLMA in several other studies (Table 6). For instance,

Table 4: Number of attempts required forplacement of devices						
Number of attempts	Group I (n=20)	Group C (n=20)	Total	P value		
1	19 (95%)	11 (55%)	30 (75%)			
2	1 (5%)	9 (45%)	10 (25%)	0.008		
Total 20 (100%) 20 (100%) 40 (100%)						
Analyzed with Fischer's Exact test						

the procedure time was found to be approximately 16 sec for I-Gel and 26 sec for cLMA.^{10,26} Shorter procedure times for I-Gel insertion were also found in comparison with cLMA in many other studies.^{11-14,16,27}

During evaluation of the performance of I-Gel in mannequins and anaesthetized patients it was observed that

Table 5: Adverse events					
Adverse events	Group I (n=20)	Group C (n=20)	<i>P</i> value		
Blood on device	3 (15%)	5 (25%)	0.695		
Dysphagia	1 (5%)	3 (15%)	0.605		
Odynophagia	1 (5%)	3 (15%)	0.605		
Sore throat	0 (0%)	2 (10%)	0.487		
Analyzed with Fischer's Exact test					

I-Gel can be easily inserted in both mannequins and patients by an inexperienced or novice person, thereby leading to shorter procedure time and higher success rate.²¹ In a recent study 30, the efficacy of two commonly available SADs (cLMA and I-gel) has been determined when inexperienced persons (58 paramedics and 46 medical students), after a brief training, inserted the devices in adult mannequin. The authors concluded that inexperienced persons could learn insertion of I-Gel and cLMA successfully in the mannequin after a brief training on mannequin. The firstattempt success rate and insertion of I-gel was found to be easier and faster than that of cLMA by both groups of performers and majority of participants preferred I-gel due to ease of handling.³⁰The performance of both the experienced and the novice physicians was assessed regarding I-Gel and cLMA insertion in mannequins in

Table 6: Procedure time and success	within 1 st attempts fo	or I-Gel and cLMA: Prese	ent study is compared
with other studies			

Studies	udies Procedure time (Seconds) Success rate (%)		ess (%)	Performer	Nature of study	
	I-Gel	cLMA	I-Gel	cLMA	-	
Present Study (2020)	63.3 ± 57.2	163.0 ± 158.3	95	55		Human, Adult
Studies with Novice perf	ormer					
Castle N, et al.	12.3 (11.5-	33.8 (30.9-	-	-	Paramedic student	Mannequin
(2010) 22	13.1)	36.7)				study
Stroumpoulis K, et al. (2012) ²³	15.2 ± 3.4	22.0 ± 4.4	90.1	47	Novice	Mannequin study
Kwak DI, et al. (2013) 24	10.0 ± 3.7	28.3 ±8.3	-	-	Novice, Normal airway	Mannequin
	10.3 ± 3.0	29.1 ± 8.2	-	-	Novice, Difficult airway	study
Pratheeba N, et al. (2016) ²⁶	15.9 ± 1.6	26.1 ± 5.1	100	84	Final year PGTs (Trained on mannequin)	Human, Adult
Alex S. et al. (2017) 27	13.6 ± 3.9	23.2 ± 7.9	96	80	First time user	Human, Adult
Kannaujia AK, et al.	16.8 ± 9.1	25.5 ± 19.4	69	53	Paramedics	Mannequin
(2020) ³⁰	15 (10-18)	15 (15-26)				study
	13.1 ± 6.8	19.2 ± 10.1	74	70	Medical students	,
	10 (9-15)	15 (14-20)				
Studies with Experience	d performer					
Singh J. et al. (2012) ¹	19.3	23.5	91.7	79.2	Experienced (cLMA>1000 and I-Gel >20)	Human, Adult
Stroumpoulis K, et al. (2012) 23	11.3 ± 5.6	13.7 ± 6.6	90	84	Experienced (>20 insertions)	Mannequin study
Helmy AM, et al. (2012) ¹⁰	15.6 ± 4.9	26.2 ± 17.7	90	80	'Senior Anaesthesiologist'	Human, Adult
Lee JR, et al. (2012) ¹⁷	17 (13.8-20) [10-40]	21 (17.5-25) [15-70]	96	92	Experienced (I-Gel>20; cLMA>200 insertions)	Human, Child
Kim MS, et al. (2014) ¹⁸	15 (13-16) [9-22]	17 (12-18) [9-69]	100	84	Experienced (> 100 uses, each device)	Human, Child
Gupta P, et al. (2015) ¹³	29.3 ± 6.9	36.7 ± 7.3	87.5	77.5	NA	Human, Adult
Ari DE, et al. (2015) ¹⁴	21 ± 4.2	30.4 ± 12.2	88	88	'Anaesthesiologist'	Human, Adult
Polat R, et al. (2015) 11	11.6 ± 2.4	13.1 ± 1.8	89.8	89.8	Experienced. Both devices (>200 uses)	Human, Adult
Rao GS.et al. (2016) 15	17.3 ± 2.9	24.9 ± 4.8	98	90	ŇA	Human, Adult
Engineer SR, et al. (2016) ²⁰	53.1 ± 6.0	57.8 ± 9.8	88	64	NA	Human, 5-60 vears
Sivasamy G (2018) ¹⁶	9.7 ± 1.0	17.2 ± 2.0	95	92	NA	Human, Adult
Arora V. et al. (2018) ¹²	5 (5-6)	23 (11-27)	92.5	80	Skilled	Human, Adult
ElGohary MM, et al. (2018) ¹⁹	78 ± 39.6	153 ± 63	80	40	NA	Human, Child

N.B. Success rates within 1st attempt are presented as proportion of patients. Procedure times are tabulated as median (IQR) or [range], values are in seconds). Considering the time in seconds, the values are rounded off to values up to one place of decimal only. NA, data not available. "I-Gel>20, cLMA>200" to be read as follows: I-Gel insertions of more than 20 occasions or LMA Classic insertions of more than 200 occasions or LMA Classic insertions of more than 200 occasions is the criteria for 'experienced' one, and so on.

a single study where first-pass success rate for I-Gel was found to be high among the novice doctors, equal to those achieved by the experienced group.²³ In another study²⁴ the researchers found that easier and quicker placement by novice performers (nurses and interns) was possible with I-Gel in comparison with cLMA in both normal and difficult airway conditioned mannequins. It was also observed in that study²⁴ that the I-gel had a higher positive response for attitude and preference.

In a clinical study²⁵ fifty doctors and paramedical staffs were divided into two groups- skilled (more than 30 insertions of SADs) or novice on the basis of their experience, and then were timed to insert the two SADs, I-Gel and LMA. The insertion of I-Gel was found to be 67% faster than the LMA in the hands ofnovice.²⁵ However, the procedure times of I-Gel insertion by skilled and novice groups were comparable in that study.²⁵ In other words, novice can perform as good as a skilled person if the SAD is the I-Gel, and not the LMA.

In the present study the post graduate trainees who had no hand-on experience in placement of SADs in human have been designated as novice. What constitutes true 'experience' in handling the SADs has not been accurately standardized; different researchers have used their own criteria.^{1,10-12,17,18,23,25} It was mentioned in the literature with quite variation, to mention a few, device insertions on more than 20 occasions,²³ insertion of each device on more than 30 occasions,²⁵ minimum 100 insertions of each device,¹⁸ insertions of both devices on more than 200 occasions,¹¹ insertion of I-Gel on more than 20 and cLMA on more than 200 occasions,¹⁷ placement of cLMA on more than 1000 and I-Gel on more than 20 occasions,¹ and nonspecific mention of 'senior anaesthesiologists'¹⁰ or 'skilled'¹² only.

While comparing with other studies in the related field, the present study also shows an inflation of mean values of procedure times in both the groups (Table 3). The higher values of procedure times in both the groups in the present study may be attributed to the following facts. According to the study design, the placement of the airway devices was done by the first-year PGTs with no prior hands-on experience of placement of the devices in human subjects. This probably has inflated the procedure time unnecessarily. In the present study, the high values of standard deviation in both the groups indicate that there was a high inter-personal variation of procedure time which should get shortened a bit with improvement of performance by training.

The definition of 'procedure time' (time needed for successful insertion) varies from studies to studies and

that may also explain difference in the results of various studies.^{17,31} For example in one study the procedure time was defined in a different way such as from the 'moment of face mask removal up to the first capnograph upstroke'¹⁷ while in the present study the procedure time was defined as the time from 'picking the device to the appearance of square waves of $EtCO_2$ '. Here, although the end point is same, the start points differ for the calculation of procedure time.

The present study finds a 95% success rate with first attempt during use of I-Gel while it was 55% with first attempt during use of cLMA, the difference being statistically significant (Table 3). In the second attempt another 5% success was achieved with I-Gel and another 45% success was achieved with the use of cLMA. Thus, considering the 2nd attempt, there was no failure in proper placement of both the devices. Pratheeba N et al.,26 found 100% success rate for proper placement of I-Gel in the first attempt while it was 84% for cLMA in the first attempt and further 16% in the second attempt. A considerably higher success rate in the first attempt was observed with the use of I-Gel in comparison with cLMA in many other studies^{10,12,27} (Table 3). However, there are other studies^{11,14} reporting high success rate for each of the devices without considerable differences between them (Table 3).

The shorter procedure time and higher success rate of I-Gel in the first attempt in comparison with cLMA in the present study may be due to technical and ergonomic reason such as the absence of cuff, less flexible stem (robust conducting channel),¹¹ etc., which makes its insertion easier and there is no need for cuff inflation. The mask shape of the cLMA resembles a wedge-shaped doughnut in overall design. Their inflatable cuffs provide airway seal but can affect their insertion and position. The leading edge of deflated mask of LMA can catch the epiglottis edge and cause it to bend downwards or impede proper placement under the tongue.³²⁻³⁴ The mask of I-Gel is uncuffed, thereby leads to easier insertion, minimal risk of tissue compression, and offers a stability of position as there is no issue of change in position owing to cuff inflation.^{11,33}

In the present study, the post-procedure hemodynamic parameters were found comparable in the intra- and postoperative period in both the groups.

In the present study, the adverse events such as sore throat, dysphagia, odynophagia and blood staining on the devices were higher in cLMA than I-Gel. However, on analysis, the difference was not statistically significant. Comparable adverse events between the uses of these two devices were also reported in some recent study.¹⁴ The adverse events such as cough, sore throat, dysphagia/dysphonia and blood

stain on device were reported to be lower with the use of I-Gel compared with cLMA in many other studies.^{12,13,19,20,27}

The study has a few limitations. Here, a second-generation SAD (I-gel) is compared with a first-generation SAD (cLMA). It was carried out in elective cases with people having Mallampatti score I and II, without any airway difficulty. Hence, the trend of procedure time and success rate of I-Gel and LMA Classic as determined in the present study may not be totally applicable in emergency settings or patients with difficult airway. The leak pressure was not measured in the present study. The fiberoptic confirmation of proper placement of airway devices was not done. The study setting was also set in comparatively shorter duration surgeries and avoided surgical areas like airway, neurosurgery, pediatric surgery etc. Hence, the inference drawn about the performance of such airway devices may not be generalized.

To summarize, the present study found that procedure time was considerably shorter (2.5 times) with the use of I-Gel. The present study also found a 95% success rate for I-Gel which was considerably higher than LMA Classic. The adverse events were found apparently higher in LMA Classic, though it was not statistically significant.

CONCLUSION

To conclude, I-Gel is a better alternative to LMA Classic in the hands of novice residents for securing the airway in terms of shorter procedure time as well as higher success rate in the first attempt. The changes in hemodynamic parameters in the post-procedure period are comparable between the devices. The adverse events related to placement of both the devices are also comparable. Thus, I-Gel appears to be a better alternative to LMA Classic as an airway management tool for the beginners.

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