# I-Gel versus Proseal Laryngeal Mask Airway in Pediatric Airway Management: A Comparative Study

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### ABSTRACT

**Aim:** To compare the insertion characteristics of supraglottic airway devices I-Gel and Proseal laryngeal mask airway (PLMA) in pediatric airway management during elective surgeries under general anesthesia. **Methodology:** This prospective randomized comparative study was conducted in 60 pediatric patients divided into two groups of 30 each (Group I and Group P), aged 1 to 5 years and belonging to American Society of Anesthesiologists (ASA) Class 1 and 2 posted for elective surgeries under general anesthesia. In Group I, I-Gel was used and in Group P, PLMA was used. The primary outcome of the study was to assess proper placement of airway devices with adequate oropharyngeal sealing and the secondary outcomes were time taken for insertion, ease of insertion, number of attempts, hemodynamic changes associated with insertion of the device, ease of gastric tube passage and complications. Statistical analysis was done by SPSS version 25. Quantitative variables were analyzed through independent sample *t*-test and categorical variables were analyzed by Chi-square test. P value <0.05 was taken as statistically significant. **Results:** The demographic data, insertion time and number of attempts were comparable in both the groups. Placement of I-Gel was better in comparison with that of PLMA and was statistically significant (p - 0.010). **Conclusion:** I-Gel is a better supraglottic airway device when compared to PLMA in terms of ease of insertion and proper placement and there are no significant hemodynamic changes with insertion of both devices.

Keywords: Supraglottic airways, I-Gel, PLMA, pediatric patient

aintenance of a patent airway remains as one of the important duties of anesthesiologists. At times, airway management becomes challenging for the anesthesiologist, specifically in pediatric age groups. Though endotracheal intubation is the gold standard technique, it has its disadvantages like reflex sympathetic stimulation and is accompanied with elevated levels of plasma catecholamines, hypertension, tachycardia, myocardial ischemia and depression, ventricular arrhythmias and intracranial hypertension. So, these days a wide variety of supraglottic airway devices (SADs) are being used to protect the airway in both elective and emergency situations, so that endotracheal intubation could be avoided in pediatric patients. Advanced airway devices like Proseal laryngeal mask airway (PLMA) and I-Gel are now considered as alternatives to endotracheal intubation for securing the airway and providing adequate ventilation even in difficult intubation and in emergency situations.<sup>1,2</sup>

Many individual studies have been done to compare the advantages and disadvantages of both these airway devices in adults. But a search through the literature reveals few studies comparing PLMA and I-Gel in routine anesthetic practice for airway management in pediatric patients. In this study, we have made an attempt to compare both these airway devices with respect to the insertion conditions and hemodynamic responses in pediatric patients posted for elective surgery under general anesthesia.

The aim of the present study was to compare the clinical performance of the PLMA (Teleflex Medical Europe Ltd, County Westmeath, Ireland) with I-Gel

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(Intersurgical, UK) in pediatric patients posted for elective lower abdominal and lower limb surgeries under general anesthesia. The primary outcome of the study was to assess proper placement of airway devices with adequate oropharyngeal sealing and the secondary outcomes were time taken for insertion, ease of insertion, number of attempts, hemodynamic changes associated with insertion of the device, ease of gastric tube passage and complications.

#### MATERIAL AND METHODS

This prospective randomized double-blind comparative study was conducted at the Institute of Medical Sciences and SUM Hospital, Bhubaneswar during the period of July 2019 to June 2020. After obtaining approval of Institutional Ethical Committee, 60 pediatric patients were selected and enrolled for the study. The parents of the patients were explained about the purpose of the study, the procedure, the intended study methods and any adverse outcome associated with it and informed written consent was obtained from them. Patients aged 1 to 5 years of either sex belonging to the American Society of Anesthesiologists (ASA) Class 1 and 2 posted for elective lower abdominal and lower limb surgeries under general anesthesia (GA) were included in the study. Patients who were not willing to participate in the study, belonging to ASA Class  $\geq$ 3, patients with anticipated difficult airway, those who required surgery in prone position and patients having risk of aspiration were excluded from the study.

Thorough preanesthetic evaluation was done including proper history, general and systemic examinations for categorizing into ASA class and inclusion into the study. Patients were randomly assigned into two groups of 30 each with help of computer-generated randomization table - Group P, for whom PLMA was used, and Group I, for whom I-Gel was used. All children fasted 6 hours preoperatively for solids and 2 hours for clear fluids. The patients were brought into the operation theater and intravenous access was obtained with appropriate size intravenous cannula. Intravenous Ringer's lactate was started. Standard monitors like pulse oximeter, automated noninvasive blood pressure, ECG, precordial stethoscope were connected and baseline values were recorded. All patients were premedicated with injection glycopyrrolate 10 µg/kg IV, injection midazolam 0.02 mg/kg IV, injection fentanyl 2 µg/kg IV, 5 minutes before induction of anesthesia. Preoxygenation was done with 100% oxygen for 3 minutes.

Induction was achieved with injection propofol 2 mg/kg IV. Facemask ventilation was done with 2% sevoflurane and oxygen. After checking for adequacy of mask ventilation, neuromuscular blockade was achieved with IV atracurium 0.5 mg/kg. Patients were allocated just before device insertion to either Group P or Group I based on sequential computer-generated numbers in opaque sealed envelopes.

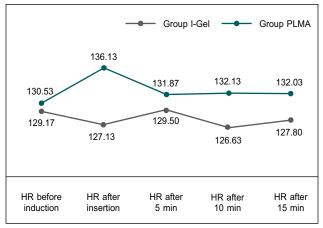
Anesthesiologist not involved in the study generated the random number table. The Anesthesiologist was blinded to the group allocation. The Anesthesiologist who inserted the airway devices had performed at least 50 PLMA and 50 I-Gel device insertions. An opaque screen was used to separate the head end from the monitor so that the observer will not be able to see, which supraglottic device is being used, to eliminate the bias. After 3 minutes of atracurium injection maintaining the patients head in sniffing position, jaw was opened and appropriate sized (based on the weight of the patient according to manufacturer's recommendation) supraglottic airway device was inserted. The I-Gel was inserted by firmly holding the device such that the cuff outlet was facing the chin of the patient and it was then guided along the hard palate until definitive resistance was felt. The insertion of PLMA was performed using the digital method. The PLMA cuff was inflated with appropriate amount of air as per manufacturer's instructions. Effective ventilation was judged using a square wave capnograph tracing and bilateral chest movements on gentle manual ventilation. In the event of partial or complete airway obstruction or a significant air leak, the device was removed, and reinsertion was attempted till a maximum of three attempts before the device was considered a failure. Endotracheal tube was used in such a situation.

The time interval between picking up the device and advancing it beyond the central incisors till it is fully inserted and total resistance has been encountered was recorded as insertion time. The number of insertion attempts to proper placement was recorded. The ease of insertion was graded as: easy – as no resistance to insertion of airway into the pharynx in a single movement, and difficult – as the resistance to insertion of airway requiring adjustment for the correct placement of the device. A lubricated orogastric tube (OGT) was inserted through the drain tube after insertion of SAD. Correct OGT placement was determined by suction of fluid or detection of injected air by listening with a stethoscope over the epigastrium. Proper placement of the device was assessed during manual ventilation,

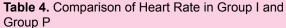
Both the Study Groups					
Parameters	Group I	Group P	P value		
Insertion time in secs	11.67 ± 3.80	10.77 ± 6.10	0.496		
Number of attempts					
First	29 (96.7%)	27 (90%)	0.612		
Second	1 (3.3%)	3 (10%)			
Ease of insertion					
Easy	28 (93.3%)	16 (53.3%)	0.001		
Difficult	2 (6.7%)	14 (46.7%)			
Proper placement					
Good	4 (13.3%)	14 (46.7%)	0.01		
Excellent	26 (86.7%)	16 (53.3%)			
Complications					
Present	1 (3.3%)	0 (0%)	1.00		
Absent	29 (96.7%)	30 (100%)			

Table 3. Insertion Time, Number of Attempts, Ease

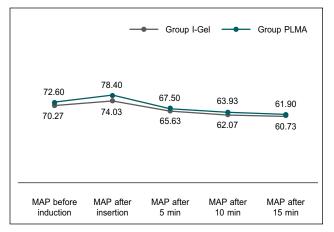
of Insertion, Proper Placement and Complications in



**Figure 1.** Changes in heart rate in Group I and Group P. HR = Heart rate.



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	I-Gel (Mean ± SD)	PLMA (Mean ± SD)	P value
HR before induction	129.17 ± 20.35	130.53 ± 19,87	0.793
HR immediately after insertion	127.13 ± 29.92	136.13 ± 21.73	0.188
HR after 5 min	129.50 ± 18.90	131.87 ± 21.25	0.650
HR after 10 min	126.63 ± 18.43	132.13 ± 20.03	0.273
HR after 15 min	127.80 ± 19.61	132.03 ± 21.27	0.426



**Figure 2.** Changes in MAP in Group I and Group P. MAP = Mean arterial pressure.

Table 5. Comparison	of MAP in	Group I	and Group P

	l-Gel (Mean ± SD)	PLMA (Mean ± SD)	P value
MAP before induction	70.27 ± 9.88	72.60 ± 8.42	0.329
MAP immediately after insertion	74.03 ± 10.22	78.40 ± 9.58	0.093
MAP after 5 min	65.63 ± 7.19	67.50 ± 5.81	0.273
MAP after 10 min	62.07 ± 5.98	63.93 ± 4.81	0.188
MAP after 15 min	60.73 ± 5.72	61.90 ± 4.71	0.392

difference was statistically significant (p = 0.001). The proper placement of the airway device in Group I was excellent in 26 (86.7%) patients, and good in 4 (13.3%) patients. The proper placement of airway device in Group P was excellent in 16 (53.3%) patients, and good in 14 (46.7%) patients. So, proper placement of the airway device was better in Group I in comparison with that of Group P and was statistically significant (p = 0.01). The placement of gastric tube was easy in all the patients in whom the study was conducted and both the groups were comparable (Table 3).

Though the heart rate was slightly on higher side in Group P (Fig. 1) compared to Group I throughout the monitoring period, the difference was not statistically significant (p > 0.05) as mentioned in Table 4. Similarly, the changes in MAP in both groups were similar (Fig. 2) and statistically not significant as shown in Table 5, where p value is >0.05 throughout the study period.

## DISCUSSION

Although it has been studied that use of SADs avoids the need for laryngoscopy resulting in less painful stimulation of the airway and hence lesser degree of pressor response, there are very few studies comparing insertion characteristics of I-Gel and PLMA in pediatric patients. In our study, we have compared I-Gel with PLMA with respect to the ease of insertion, proper placement of the airway device and hemodynamic changes during insertion of the device. In this study, the demographic data of patients, like age, sex and body weight, were similar and were comparable in both groups.

Though the insertion time for supraglottic devices in this study was more in Group I (11.67 seconds) compared to Group P (10.77 seconds), the difference was statistically not significant (p = 0.496). Insertion time for I-Gel in our study was longer than a study which achieved I-Gel insertion within 5 seconds.<sup>3</sup> The success rate of insertion of I-Gel was 96.7% in the first attempt which was better than PLMA, for which success rate in first attempt was 90%. But the difference was statistically insignificant (p = 0.612). In a study by Kannaujia et al,4 the success rate at first attempt was 90%. Similar to our study, the study by Goyal et al<sup>5</sup> showed success rate at first attempt was 95% and success rate at second attempt was 100%. A study by Francksen et al<sup>6</sup> reported that success rate of insertion of I-Gel was 90% in first attempt and overall success rate was 100%. Arslan et al<sup>7</sup> reported a success rate of 100% with PLMA.

Ease of insertion in Group I was easy in 93.3% patients and difficult in 6.7% patients, and in Group P it was easy in 53.3% patients and difficult in 46.7% patients. Insertion of I-Gel was easy compared to PLMA and was statistically significant (p = 0.001). Similar to our study, Singh et al<sup>8</sup> and Goyal et al<sup>5</sup> concluded in their studies that insertion of I-Gel was easier than any other currently available supraglottic devices. But the studies by Theiler et al<sup>9</sup> and Michalek et al<sup>10</sup> concluded that insertion of I-Gel was difficult, likely due to bulky design of I-Gel.

The proper placement of the airway device was better with I-Gel than PLMA. In Group I, the placement of airway device was excellent in 86.7% patients and was good in 13.3% patients. In Group P, the placement of airway device was excellent in 53.3% patients and was good in 46.7% patients. The I-Gel showed higher leak pressures when compared to PLMA by adequate sealing with perilaryngeal structures. This could be attributed to unique noninflatable cuff of I-Gel, which mirrors the perilaryngeal anatomy. The leak pressure of I-Gel improves with time due to thermoelastic material, which forms more efficient airway seal after warming to the body temperature. To obviate this effect, we checked for airway seal after 5 minutes of insertion of I-Gel. The placement of gastric tube was easy in both Group I and Group P and was 100% successful overall. A study conducted by Helmy et al<sup>11</sup> showed success rate of gastric tube insertion was high in I-Gel group.

In our study, the mean heart rate at preinsertion, immediately after insertion, at 5, 10 and 15 minutes were compared. Though the heart rate in PLMA group was on slightly higher side throughout the study period as compared to I-Gel group, the difference in both groups was negligible and when compared with preinsertion value and it was statistically insignificant (p > 0.05).

Similarly, in both the groups, the changes in MAP were not statistically significant (p > 0.05). Similar to our study, Mitra et al<sup>12</sup> and Chauhan et al<sup>13</sup> in their studies concluded that hemodynamic changes with insertion of I-Gel were comparable to that of PLMA. Contrary to our study, Jindal et al<sup>14</sup> concluded that I-Gel insertion causes less hemodynamic changes as compared to other supraglottic devices. In this study, we observed complications of insertion of both airway devices. In Group I, one case of laryngospasm had been observed which was managed by deepening plane of anesthesia and positive pressure mask ventilation with 100% oxygen. There were no complications in Group P. Goyal et al<sup>5</sup> found that the incidence of complications, both in PLMA and I-Gel groups, was low. Helmy et al<sup>11</sup> reported that airway trauma was minimal with I-Gel. Our study findings are in consistence with these studies.

So, in our study, we observed that insertion of I-Gel was easier with proper placement compared to PLMA. But when we compared the insertion time, number of attempts needed for insertion and hemodynamic changes with insertion in both our study groups, the results were comparable as seen in many published studies like Mitra et al<sup>12</sup> and Chauhan et al.<sup>13</sup>

## CONCLUSION

In this study, based on the results, we concluded that I-Gel is a better supraglottic airway device when compared to PLMA in terms of ease of insertion and proper placement and there are no significant hemodynamic changes with insertion for both devices. But both the airway devices can be safely used to provide anesthesia in elective surgical procedures in pediatric patients.

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