International Journal of Biomedical and Advance Research

ISSN: 2229-3809 (Online); 2455-0558 (Print) Journal DOI: <u>https://doi.org/10.7439/ijbar</u> CODEN: IJBABN e5428

Comparison of Laryngeal Mask Airway Supreme, i-gelTM and Ambu Auragain in Children for Airway Management

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Abstract

Background and Aims: This was a prospective, randomised study performed on 180 children of ASA I/II, aged 5-14 years, planned for elective surgery and requiring general Anaesthesia in the Department of Anaesthesiology, critical care, Pain and Palliative Medicine, Dr Sushila Tiwari Government Medical College, Haldwani. To compare the performance of three airway devices, the laryngeal mask airway supreme, i-gelTM and Ambu Auragain in children for airway management. **Methods:** children were randomized into three groups (60 each): Group S (LMA Supreme), Group I (i-gel) and Group A (Ambu Auragain). The primary outcome was the insertion time. We also assessed the number of insertion attempts, ease of insertion, haemodynamic parameters and complications. Intergroup differences were compared using one-way analysis of variance (ANOVA) post - hoc correction for continuous data and Kruskall Wallis test for categorical variables.

Results: Demographic data did not differ between the three groups. Insertion time for i-gel (18.5 (18-20) sec) was shorter than for the LMA Supreme (22 (20-22) sec) and Ambu Auragain (20.5 (19-23) sec) (P = 0.02). There were no differences in the number of attempts, ease of insertion, haemodynamic parameters and complications between all three groups.

Conclusion: LMA Supreme, Ambu Auragain and i-gel provided a similar performance of airway management in children. The success rate of insertion and ease of intubation of LMA Supreme, Ambu Auragain and i-gel were comparable. i-gel has a lesser time of insertion than LMA Supreme and Ambu Auragain.

Keywords: Laryngeal mask airway, Supreme, Ambu auragain, i-gelTM, children.

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How to cite: Madeshia A, Bhandari G, Shahi K. S and Chand G. Comparison of Laryngeal Mask Airway Supreme, i-gelTM and Ambu Auragain in Children for Airway Management. *International Journal of Biomedical and Advance Research* 2020; 11(06): e5428. Doi: 10.7439/ijbar.v11i6.5428 Available from: https://ssjournals.com/index.php/ijbar/article/view/5428

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1. Introduction

For the Anaesthesiologists and critical care providers, airway management remains one of the most challenging tasks, whether it is adult or paediatric patients. One should also consider the uniqueness of paediatric airway management. Dr Archie Brain developed an airway device that was less stressful to the patient as compared to tracheal intubation, and as safe as facemask and airway. He hoped that his device would benefit for cases where mask ventilation and intubation was particularly difficult. Thus it gives anaesthesiologists a safer alternative to complex intubation, especially in emergency scenarios. [1] Supraglottic airway devices (SADs) are the part of routine and emergency paediatric airway management, including use in the difficult airway management and neonatal resuscitation [2]. First-generation devices are simple airway tubes attached to a mask that rests over the glottic opening (e.g. LMA classic, flexible, cobra PLA). A second-generation device has a gastric access channel that allows for gastric venting and gastric tube placement (LMA Proseal, supreme, i-gelTM, Ambu Auragain).

Introduction of LMA (laryngeal mask airway) was one such advancement. Laryngeal mask airway (LMA) offers several advantages, including ease of placement, lower drug requirement, reduced haemodynamic response, reduced intracranial and intraocular tension, smoother emergence and a lower incidence of airway trauma and complication.

LMA Supreme is a single-use supraglottic airway device, easy insertion without the need for digital or introducer tool guidance. It forms two seals: one at the upper oesophagal sphincter and the other over the glottic opening. It also has a fixation tab; a rectangular structure which projects over the patient's upper lip. LMA supreme designed to facilitate easy insertion and fixation [3].

i-gelTM is a soft, gel-like, non-inflatable cuff, made of thermoplastic elastomer. It has widened, flattened stem with a rigid bite-block that acts as a buccal stabiliser to reduce axial rotation and malpositioning. It has oesophageal vent through which a gastric tube can be passed [4-6].

Ambu Aura Gain is a newly introduced supraglottic airway device. It is anatomically curved with integrated gastric access and can use as a conduit for direct endotracheal intubation assisted by a flexible scope. [7]

2. METHODS

After ethical committee Clearance 180 ASA grade I/II patients aged between 5-14 years children undergoing surgery of duration 1-2 hours under general anaesthesia with controlled ventilation included in the study. Patients randomly divided into three equal groups: Group S – (LMA supreme, n=60), Group I- (i-gelTM, n=60) and Group A- (Ambu Auragain, n=60). Randomization procedure done by using closed envelops method. All patients were pre-medicated with injection Glycopyrrolate (0.01mg/kg) i.v., Tramadol (1.5mg/kg)i.v., injection ranitidine (1mg/kg) i.v. All the patients were pre oxygenated with O2 via facemask for 3 min. General Anaesthesia (GA) was induced with injection propofol 2mg/kg i.v., injection succinylcholine 1.5mg/kg i.v. and were manual ventilation done.

Once the adequate depth of anaesthesia achieved either of LMA supreme or $i-gel^{TM}$ or Ambu AuraGain

appropriate for weight inserted. The selection of the SAD size based on the children's actual body weight (size 2 for 10–20 kg, size 2.5 for 20–30 kg). The device insertion technique was based on manufacturer recommendations. Once in place, the cuff was inflated according to the size of the SAD, as per the manufacturer's instruction manual (2.0 size: 10 ml, and 2.5 sizes: 14 ml). Correct placement of LMA confirmed by bilateral symmetrical chest expansion on manual ventilation and square waveform on capnography. After fixing the device, patient's lungs were ventilated with a tidal volume of 8-10ml/kg and anaesthesia maintained with 50% Oxygen, 50% Nitrous oxide and sevoflurane (1-3%) and injection rocuronium (0.8mg/kg).

Haemodynamic parameters- Heart rate (HR), systolic blood pressure (SBP), diastolic blood pressure (DBP), Mean arterial pressure (MAP), arterial oxygen saturation (Spo2), end-tidal CO2 (ETCO2) monitored.

At the end of the surgical procedure, anaesthesia reversed with a standard dose of Neostigmine (0.05mg/kg) and Glycopyrrolate (0.2mg/1mg Neostigmine), and the device removed. The tip of the LMA examined for the presence of blood. In the evening of the same postoperative day, we recorded any discomfort while swallowing water or food, sore throat and hoarseness if present.

More than three attempts taken as a failure and the child were intubated by conventional laryngoscope with ET tube, and surgery performed.

We analysed data using IBM SPSS Statistics 24 software. Continuous numerical variables presented as mean (SD) or Median (IQR) and intergroup differences were compared using one-way analysis of variance (ANOVA) with posthoc correction. Categorical variables were presented as a ratio or as n (%), and between-group differences compared using the Kruskall-Wallis test. P<0.05 were considered statistically significant.

3. Results

Demographic data did not differ between the three groups. Males were predominant in our study (Table 1). Insertion success rate was similar in all three groups. Insertion time for the i-gel (18.5 (18-20) sec) was shorter than for the LMA Supreme (22 (20-22) sec) and Ambu Auragain (20.5 (19-23) sec) (P = 0.02) and there were no differences in number of attempts, ease of insertion (Table 2), haemodynamic parameters (Table 3, 4), other parameters (Table 5, 6) and complications between all three groups. This study demonstrated that all three supraglottic airway devices provided almost similar performance for airway management in children, best being i-gel.

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|---|--------------------|--------------|--------------|----------|--|
| Patient characteristics | Group S | Group A | Group I | P Value | |
| Age (Yrs) | 8.98±3.40 | 9.52±3.44 | 9.40±3.13 | P > 0.05 | |
| Weight (kg) | 26.02±12.48 | 27.18±13.09 | 27.03±12.25 | P > 0.05 | |
| Height (cm) | 124.08 ± 20.27 | 126.55±21.37 | 124.88±21.04 | P > 0.05 | |
| Sex (M/F) | 45/15 | 48/12 | 45/15 | P>0.05 | |
| | 0 . | | | | |

Data presented as mean \pm SD or (range) or number of patients Group S- Supreme, Group A- Ambu Auragain, Group I- i gelTM

Table 2: Clinical performance of all three Supraglottic airway devices

| Parameters | Group S | Group A | Group I | P Value | |
|--------------------------|------------|--------------|--------------|---------|--|
| Number of attempts (1/2) | 54/6 | 56/4 | 56/4 | 0.73 | |
| Time of insertion of SAD | 20 (20-22) | 20.5 (19-23) | 18.5 (18-20) | 0.02 | |
| Ease of insertion (1/2) | 54/6 | 54/6 | 56/4 | 0.76 | |
| Complication | | | | | |
| Sore throat | 1 | 4 | 2 | 0.35 | |
| Dysphagia | 1 | 3 | 2 | 0.61 | |
| Hoarseness | 2 | 1 | 3 | 0.61 | |
| Blood stained | 3 | 2 | 1 | 0.61 | |

Values expressed as mean \pm SD, median (IQR) or a number of patients. Ease of insertion of the device as grade 1=no resistance, 2=mild resistance

Table 3: Comparison of Heart rate at different period of time between the three groups

| Heart rate | Group S (n=60) | Group A (n=60) | Group I (n=60) | P value |
|------------------|----------------|----------------|--------------------|---------|
| Baseline | 107.03±12.15 | 105.45±12.25 | 106.38 ± 10.21 | 0.75 |
| After induction | 111.08±11.76 | 108.97±11.99 | 109.68 ± 10.07 | 0.58 |
| After intubation | 115.50±11.43 | 113.57±11.85 | $113.77{\pm}10.07$ | 0.58 |
| At 1 minutes | 113.45±11.47 | 110.85±11.73 | 111.70±10.26 | 0.43 |
| At 3 minutes | 111.92±11.26 | 109.75±11.62 | 110.25 ± 10.02 | 0.53 |
| At 5 minutes | 109.57±11.48 | 107.63±12.16 | 108.23±9.75 | 0.62 |
| At 10 minutes | 107.12±12.03 | 105.65±12.20 | 106.55 ± 9.90 | 0.78 |
| At 15 minutes | 106.40±12.09 | 104.83±12.29 | $106.02{\pm}10.01$ | 0.74 |

Table 4: Comparison of BP at different period of time between the three groups

| | | at different period of time between the three gi | | | |
|------------------|-----|--|-------------------|-------------------|---------|
| Blood Pressure | | Group S | Group A | Group I | P Value |
| | SBP | 98.28 ± 7.08 | $98.60{\pm}7.01$ | 99.10±7.47 | 0.82 |
| Baseline | DBP | 54.45 ± 4.06 | 54.98±3.52 | 54.67±3.81 | 0.74 |
| | MAP | 68.90 ± 4.28 | 69.48±4.34 | 69.25±4.24 | 0.75 |
| | SBP | 99.35±6.95 | 99.50±6.99 | 100.37±7.53 | 0.70 |
| After induction | DBP | $54.10{\pm}4.08$ | 54.77±3.69 | 54.37±3.92 | 0.64 |
| | MAP | 69.12±4.29 | 69.55±4.43 | 69.53±4.47 | 0.83 |
| | SBP | 101.83±6.79 | 101.88 ± 6.88 | 102.57±7.59 | 0.82 |
| After intubation | DBP | $55.80{\pm}4.08$ | 56.42±3.82 | 55.98±3.74 | 0.67 |
| | MAP | 70.90±4.19 | 71.40±4.43 | 71.40±4.42 | 0.76 |
| | SBP | 100.33 ± 6.77 | 100.27±6.76 | 101.10±7.37 | 0.77 |
| At 1 min | DBP | 54.52 ± 3.98 | 55.25±3.74 | 54.70±3.70 | 0.55 |
| | MAP | 69.68±4.25 | 70.23±4.34 | 70.10±4.33 | 0.76 |
| At 3 min | SBP | 99.13±6.68 | 99.33±6.96 | 99.93±7.39 | 0.81 |
| | DBP | 53.38±3.79 | 54.10±3.93 | 53.40±3.61 | 0.49 |
| | MAP | 68.67±4.25 | 69.45±4.97 | 68.97±4.29 | 0.63 |
| | SBP | 99.48 ± 6.90 | 99.23±6.76 | 100.25±7.24 | 0.71 |
| At 5min | DBP | 52.48±3.49 | 52.98±3.57 | 52.48±3.13 | 0.65 |
| | MAP | 68.18 ± 4.04 | 68.55±4.15 | 68.43±3.96 | 0.88 |
| At 10min | SBP | 100.08 ± 6.01 | 100.37 ± 5.75 | 101.67±6.36 | 0.31 |
| | DBP | 51.78±2.85 | 52.28±3.01 | 51.67±2.68 | 0.45 |
| | MAP | 67.87±3.29 | 68.28±3.57 | 68.43±3.53 | 0.65 |
| At 15min | SBP | 100.20 ± 5.90 | 100.40 ± 5.69 | 101.62 ± 6.88 | 0.39 |
| | DBP | 51.30±2.36 | 51.43±2.35 | 51.23±2.06 | 0.88 |
| | MAP | 67.88±2.99 | 67.75±3.08 | 68.03±3.19 | 0.88 |

| SPO ₂ | Group S (n=60) | Group A (n=60) | Group I (n=60) | P value |
|------------------|----------------|------------------|----------------|---------|
| Baseline | 98.68±1.04 | 98.88 ± 0.84 | 98.92±0.94 | 0.35 |
| After induction | 98.98±0.77 | 99.07±0.78 | 99.10±0.73 | 0.68 |
| After intubation | 98.40±0.64 | 98.60±0.58 | 98.60±0.74 | 0.16 |
| At 1 minute | 98.88±0.69 | 98.90±0.66 | 98.98±0.70 | 0.69 |
| At 3 minutes | 98.87±0.65 | 98.98±0.59 | 98.93±0.68 | 0.61 |
| At 5 minutes | 98.93±0.68 | 99.05±0.65 | 98.98±0.74 | 0.65 |
| At 10 minutes | 99.13±0.81 | 99.07±0.82 | 98.97±0.80 | 0.53 |
| At 15 minutes | 99.15±0.75 | 99.25±0.71 | 99.20±0.73 | 0.75 |

Table 5: Comparison of SPO₂ at different period of time between the three groups

| Table 6: Comparison of ETCO ₂ at different period of time between the three groups |
|---|
|---|

| ETCO ₂ | Group S (n=60) | Group A (n=60) | Group I (n=60) | P value |
|-------------------|----------------|----------------|----------------|---------|
| Baseline | 36.62±2.08 | 37.20±1.78 | 37.23±1.64 | 0.12 |
| After induction | 36.92±1.85 | 37.37±1.65 | 37.43±1.38 | 0.17 |
| After intubation | 37.15±1.76 | 37.58±1.73 | 37.78±1.40 | 0.10 |
| At 1 minute | 37.32±1.57 | 37.80±1.64 | 37.68±1.44 | 0.21 |
| At 3 minutes | 37.62±1.63 | 37.95±1.69 | 37.88±1.48 | 0.07 |
| At 5 minutes | 38.04±2.24 | 38.17±2.18 | 38.53±1.89 | 0.06 |
| At 10 minutes | 37.47±2.36 | 37.92±2.25 | 38.25±2.18 | 0.16 |
| At 15 minutes | 37.35±1.89 | 37.70±1.83 | 37.82±1.69 | 0.34 |

3. Discussion

There were various studies on the safety and efficacy of airway maintenance when using supraglottic airway devices on children as anatomy and physiology of children differ from adults. So, we conducted a clinical study comparing safety and efficacy of the three supraglottic devices LMA Supreme, i-gelTM and Ambu AuraGain in anaesthetised patients on mechanical ventilation undergoing elective surgical procedures.

LMA Supreme was successfully inserted on the first attempt in 90% of patients, while in i-gelTM and Ambu Auragain, it was 93.3% of patients (p=0.73). All three groups are comparable and statistically not significant in our study, so all three SGA are equally efficacious. Lee *et al*[8] compared i-gelTM and Laryngeal Mask Airway Supreme during general anaesthesia in 60 infants in which 100% insertion success rate during 1st attempt with i-gelTM and 96% with supreme (p=1.00), which is not comparable with our study but the difference is statistically not significant.

A study by Jagannathan *et al*[9] compared Ambu Auragain and LMA Supreme in children with a success rate of insertion on 1st attempt at 96% and 100% respectively (p=0.5). However, it is not comparable with our study, and the difference is statistically not significant. In our study 2nd attempt was successful in 10% patients in LMAsupreme, while in i-gelTM and Ambu Auragain it was successful in 6.7% patients (p=0.73). A study conducted by Jagannathan *et al*[9] 2nd attempt was successful in 2% patient in LMA supreme and 2% patient in Ambu Auragain (p=0.5). Although it is not comparable with our study and difference is statistically not significant.

90% of children had no resistance during insertion of LMA Supreme and Ambu Auragain while in i-gelTM, 93% of children have no resistance during insertion. There was moderate resistance during insertion in 10% of children in LMA supreme and LMA Auragain while in 6.67% children in i-gelTM respectively (p=0.7). A study conducted by Arslan et al[10] compared LMA Proseal and LMA Supreme in children. In LMA Proseal they did not have resistance while insertion and in LMA Supreme, 93% of children had no resistance but in 6.67% of children had moderate resistance (p=0.2). Although it is not comparable with our study and the difference is statistically not significant. Jagannathan et al[9] compared LMA Ambu Auragain and supreme in children, no resistance was seen in 76% of children in LMA Ambu Auragain and 90% children in LMA supreme (p=0.09). It is comparable with LMA supreme with our study but not comparable with LMA Ambu Auragain. 22% of children had moderate resistance in LMA Ambu Auragain and 8% children in supreme. 2% of children in both LMA Ambu Auragain and supreme had severe resistance, respectively. The difference is statistically, not significant.

The median time for effective placement of device was lowest for i-gelTM, i.e. 18.5 seconds (range 18- 20 secs), followed by 20.5 secs for Ambu Auragain (range 19- 23 secs). LMA Supreme had the maximum median insertion time of 22 secs (range 20-22 secs) of all the three devices. Statistical significance was seen between effective insertion time of all the three devices (p=0.02).

Mihara *et al*[11] compared the clinical performance of i-gelTM and Ambu Auragain in children with insertion time 17.1 ± 4.5 sec and 21.3 ± 6.5 sec

respectively for i-gelTM and Auragain(p=<0.001) which is comparable with our study and statically significant. In Jagannathan *et al*[9] the insertion time for LMA Ambu Auragain and LMA Supreme were 13 (12 to15) and 13(12 to 14) sec respectively, which is not comparable with our study.

We also compared systolic blood pressure, diastolic blood pressure, mean arterial pressure, heart rate, SPO_2 and $ETCO_2$ between the three groups at different time intervals, i.e. Baseline, Induction, intubation, 1 min, 3 mins, 5 mins, 10 mins and 15mins after induction. We found all the three groups were comparable to the parameters as mentioned above, and the results were statistically insignificant.

The fact that all three are Supraglottic devices sit in the hypopharynx and stimulate the stretch receptors, which are present above the vocal cords. Hence, changes in the blood pressure and the heart rate at different time intervals were comparable between the three groups. SBP mean value was 100.33±6.77 for LMA Supreme, 100.27±6.76 for Ambu Auragain, 101.10±7.37 for i-gelTM (p=0.77) and DBP mean value was 53.38±3.79 for LMA Supreme, 54.10±3.93 for Ambu Auragain, 53.40±3.61 for igelTM (p=0.49), respectively which were comparable and statistically not significant. In our study, the MAP mean value was 68.67±4.25 for LMA supreme, 68.28±3.57 for LMA Auragain and 68.43 ± 3.53 for i-gelTM (p=0.65) which was statistically not significant. It was comparable to the study conducted by Gu et al.[12] had 69.14±7.21 for LMA supreme, 70.98±5.67 for

LMA AuraOnce and 69.98±7.51 for i-gelTM (p=0.269). Heart rate mean value in our study was 109.57±11.48 for LMA Supreme, 107.63±12.16 for LMA Auragain and 110.25 ± 10.02 for i-gelTM (0.78) respectively, which are statistically not significant. A study by Gu et al.[12] had 114.97±5.35 for LMA supreme, 114.64±6.62 for LMA AuraOnce and 112.96±7.31 for i-gelTM (P=0.148) which were comparable with our result but statistically not significant. In our study, the SPO₂ mean value was 99.15±0.75 for LMA supreme, 99.25±0.71 for LMA Auragain and 99.20±0.73 for i-gelTM (p=0.75). Similar result found in a study conducted by Arslan et al.[10] SpO₂ 99.9±4.3 for LMA Proseal and 99.9±0.3 for supreme (p=0.7). The mean value of ETCO₂ for LMA Supreme was 37.35±1.89, 37.70±1.83 for LMA Auragain and 37.82±1.69 for i-gelTM (p=0.34) respectively, which were comparable and statistically not significant. In a study by Arslan et al.[10] had ETCO2 40±5.2 for Proseal and 40.5±4.5 for Supreme (p=0.7), which is comparable with our study but statistically not significant.

Incidence of sore throat was seen in 1.6%, 6.6% and 3.3% patients with LMA Supreme, Auragain and i-

gelTM (p=0.35) respectively. In Gu et al.[12] study the incidence of sore throat was 16.1%, 3.1% and 18.7% for LMA supreme, AuraOnce and i-gelTM (p=0.12). Incidence of sore throat with i-gelTM was comparable but not with LMA Supreme, and the difference is statistically not significant. In our study incidence of dysphagia was seen in 1.6%, 5%, 3.33% patients with LMA Supreme, Auragain and i-gelTM respectively (p=0.61). Hoarseness of voice was seen in 3.3%, 1.67% and 5% (p=0.61) patients with LMA Supreme, Ambu AuraGain and i-gelTM insertion respectively in our study. A study conducted by Gu et al.[12] 3.1%, 9.3% and 3.2% (p=0.614) Hoarseness were noted in patient with LMA Supreme, AuraOnce and igelTM. The study was comparable with LMA Supreme but not with i-gelTM.

Blood-stained on LMA was seen in 5 %, 3.3% and 1.67% patient in LMA Supreme, Auragain and i-gelTM respectively (p=0.61). Gu *et al*[12] did a study had 3.2%, 3.1% and 0% in LMA supreme, AuraOnce and i-gelTM (p=0.7). Incidence of blood-stained in LMA Supreme and i-gelTM noted in our study was not comparable with the study done by Gu *et al.* [12] There was no incidence of aspiration, bronchospasm and Laryngospasm in our study. Majority of the patients from our study did not have postoperative complications which could be due to the high success rate in first insertion attempts and results were comparable in all the three groups.

4. Conclusion

This study demonstrated that LMA Supreme, Ambu Auragain and i-gel provided a similar performance of airway management in children. Success rate of insertion of LMA Supreme, Ambu Auragain and i-gel were comparable and ease of intubation was also comparable. In terms of total time taken for insertion, i-gel has lesser time of insertion than LMA Supreme and Ambu Auragain. All three LMA Supreme, Ambu Auragain and i-gel are equally efficacious in children for securing airway in controlled ventilation. i-gel requires less manipulation , no cuff inflation and hence securing an airway is rapid with i-gel in most of the children.

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