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SHORT REPORT

Characteristics of a New Device for Intubation Through a Laryngeal Mask Airway Over a Flexible Endoscope

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Keywords: laryngeal mask airway, intubation, fiberoptic, emergency, airway, scope, vortex

Introduction

Archie Brain introduced the laryngeal mask airway, ie, LMA-Classic in 1987.¹ Soon, the need for an LMA to be used as an intubation conduit for endotracheal intubation was recognized. The LMA-Classic was unsuitable, the airway tube was too narrow and too long to allow the endotracheal tube (ETT) to be reliably placed distal to the vocal cords, and it was subsequently difficult to remove the LMA over the ETT after tracheal intubation. In recognition of the importance of securing the airway with an ETT after placement of an LMA, a decade later in 1997, Brain produced the intubating LMA and a specific endotracheal tube,² the LMA-Fastrach [™] and Fastrach[™] ETT (both Teleflex[®], Westmeath, Ireland). Since then, multiple laryngeal mask airways dedicated for this purpose have been produced (Ambu AuraGain[™] Air-Q[™], C Trach[™], and Unique Evo[™]), the evolution of the LMA has been eloquently described by Sorbello in "Evolution of the supraglottic airway devices: the Darwinian perspective."³ The Fastrach[™] ETT designed by Brain remains the only widely commercially available ETT designed for intubation through an LMA. The development of the ETT used to intubate through an LMA may have been neglected and needs to be subjected to the same evolutionary pressures.

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Received: 21 April 2021 Accepted: 16 June 2021 Published: 6 July 2021 Intubation through an LMA over a flexible endoscope in a modified Baileys maneuver⁴ is gaining acceptance as a primary technique when poor tolerance to apnea is expected. Clinicians will be increasingly faced with the dilemma (Table 1) of managing the patient who has an LMA in place, but still requires the airway secured by an ETT.

The Ideal Characteristics of an Endotracheal Tube to Intubate Through an LMA Over a Flexible Endoscope

When Brain was designing the ETT to be compatible with the intubating LMA, a major concern was the angle at which the ETT exited from the LMA mask aperture so that it lined up with the vocal cords and trachea to assist in blind intubation. The ideal tube was found to be a straight, soft wire reinforced, silicone ETT with a soft conical Tuohy-like tip – the FastrachTM ETT (Teleflex®, Westmeath, Ireland).

Although blind intubation through a laryngeal mask airway is feasible to reduce trauma to the larynx and increase first pass intubation, using a flexible endoscope to achieve endotracheal intubation is currently recommended.^{5,6}

In the design of an ETT suitable for intubation over a flexible endoscope through an LMA, the following characteristics were considered important:

Table I A Comparison of Two Current Devices i.e., the Fastrach ETT[™] and Aintree Intubating Catheter[™] (AIC) and the Rescue ETT[™] a New Device Proposed for Intubation Through a Laryngeal Mask Airway Over a Flexible Endoscope

	Fastrach™ ETT	Aintree Intubation Catheter™	Rescue™ ETT
Design of tip/bevel to assist successful first pass intubation	Conical Tuohy-like tip	n/a	Posterior facing bevel + curved hemispheric tip
Length of device	25 cm ETT sizes 8, 7.5, and 7 mm measure 25 cm to upper border of cuff	56cm	ETT sizes 5.5 to 9 mml 7 to 8.5 mm measure 25 cm to upper border of cuff
Material used for manufacture	Wire reinforced silicone and PVC. Soft and malleable	Polyethylene Semi-rigid *Haemorrhage, pneumothorax, mediastinitis and deaths, have all been reported with the use of airway exchange catheters. ^{16,17}	PVC and wire reinforced PVC. Soft and malleable
Ease of removal of the LMA over the ETT	Withdrawn over solid stabilizer rod – danger of tube migrating/ extubation. FOB needs to be reintroduced to verify position of ETT.	Easily withdrawn over narrow diameter AIC	Withdrawn over airway tube connected to ETT. Continued ventilation. FOB only removed after LMA is withdrawn, allowing ETT position to be verified.
Continuous ventilation possible	No Interruption occurs when LMA is withdrawn over ETT + solid stabilizer rod	Ventilation through AIC only recommended when oxygen requirements are high due to the risk of barotrauma	Yes
Suitable long-term intubation	No • Low volume high-pressure cuff • Wire reinforced ETT	n/a	Yes • ETT available in PVC • High-volume low-pressure cuff
Interruption to airway security	Nil Achieved with an LMA and then an ETT	Yes After the LMA is withdrawn over the AIC the airway is not secure until the ETT is railroaded over the AIC	Nil Achieved with an LMA and then an ETT

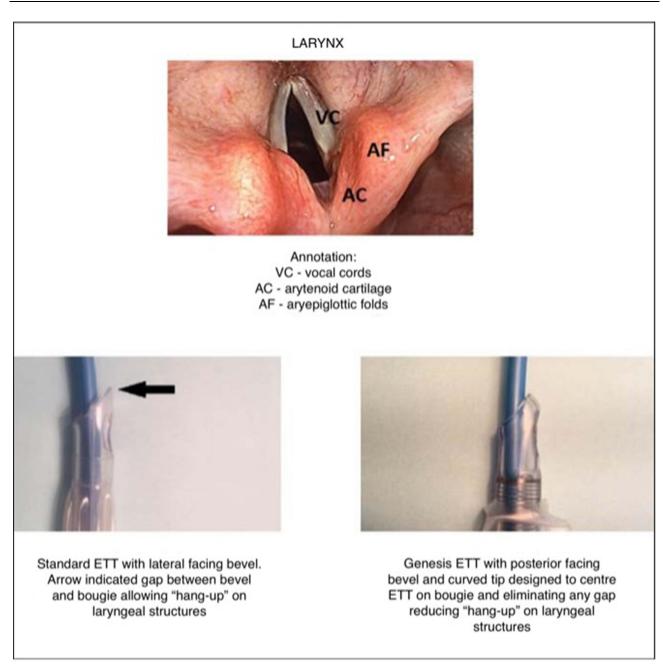


Figure I Comparison of the bevels of a standard ETT with lateral facing bevel (left) and Rescue™ ETT (right) to illustrate the mechanisms to reduce laryngeal "hang-up". Notes: A bougie is photographed inside the lumen of the ETTs to simulate the similar mechanism that occurs with a flexible endoscope.

• The tip and bevel must be suitable for first pass successful intubation. Hang-up (Figure 1) may occur at the level of the arytenoids/aryepiglottic folds, or on the anterior tracheal wall.⁷ Studies suggest that a posterior facing bevel,^{8–11} conical or hemispheric shapes¹² as well as a softer tip are superior.¹² When using an ETT with a lateral facing bevel rotating the tube 90 degrees in an anticlockwise direction (the bevel becomes posterior facing) relieves the hang-up

but may cause significant trauma and even complete airway obstruction in awake patients.¹³ For this reason many clinicians recommend the 90-degree anticlockwise rotation of the endotracheal tube before intubation is attempted.

• The length of the ETT needs to be sufficient to be placed below the cords proximal to the carina. Standard ETTs are unsuitable. The shaft length of most laryngeal mask airways sizes numbers 3 to 5

is 19–20 cm. The distance from the airway tube aperture in the mask to the vocal cords varies from 2 to 4.7 cm in adults.¹⁴ For the lower border of the ETT cuff to be 2 cm below the vocal cords, the length of the ETT to the proximal border of the cuff should be 20+4.7+2=26.4 cm. A preformed nasal RAE and a microlaryngeal ETT are longer than standard oral ETTs and have been advocated for intubation through a laryngeal mask airway.¹⁵

- The ETT is preferably manufactured from a soft material (eg, silicone). A softer ETT reduces trauma to the larynx and trachea and allows the tube to conform to the flexible endoscope reducing hang-up particularly at the anterior tracheal wall.^{11,12}
- The LMA must be easily withdrawn over the ETT.
- The 15 mm connector for the ETT must be detachable.
- Continued airway security with minimal interruption to ventilation and oxygenation for the duration of the procedure is vital.
- The technique should be simple with minimal steps to the procedure, easy to learn and perform without regular practice.
- The ability to verify ETT position with the flexible endoscope until after the laryngeal mask airway is withdrawn would be advantageous.
- The ETT should be suitable for long-term intubation with a high-volume low-pressure cuff manufactured from polyvinyl chloride (PVC) without wire reinforcing.
- The device should be affordable to allow regular practice and universal availability.

Discussion Internal Diameter (ID) of ETT that May

be Used

The ID of the Rescue[™] and Fastrach[™] ETT (see Table 1) is limited to a maximum of 8 mm with passage through an airway tube of an LMA. Evidence suggests that with current ventilators smaller ID ETTs are adequate,¹⁸ except for critically ill patients requiring long-term intubation when an 8 mm ID ETT for women and a 9 mm ID ETT for men is required. The use of large diameter ETTs can cause trauma to the larynx and trachea.¹⁸ When using an Aintree Intubation Catheter (AIC) an ETT larger than 8 mm ID may be used, however, 7 mm is the minimum ID that fits over an AIC, limiting its use in pediatric patients. The Rescue[™] ETT is available from ID size 5.5 to size 8 mm and the Fastrach ETT[™] in size 6 to size 8 mm allowing use both in pediatric patients and adults.

Limitations Related to the Length of the Devices

A size 8 mm ID Fastrach[™] ETT measures 25 cm from the proximal end of the tube to the upper border of the cuff. An i-gel[™] laryngeal mask airway (size 5) as an airway tube of 20 cm, in a tall patient add 4.7 cm¹⁴ for the distance between the aperture in the mask to the vocal cords makes 24.7 cm. This makes the Fastrach[™] ETT too short to be reliably placed below the vocal cords in the trachea. The Rescue[™] ETT (Figure 2A) when combined with the airway tubing has a minimal length of 34 cm (Figure 3) to the upper border of the cuff allowing it to be placed at an adequate depth even with an extremely tall patient, The AIC length (56 cm) may be too short to railroad an ETT over in very tall patients.¹⁹

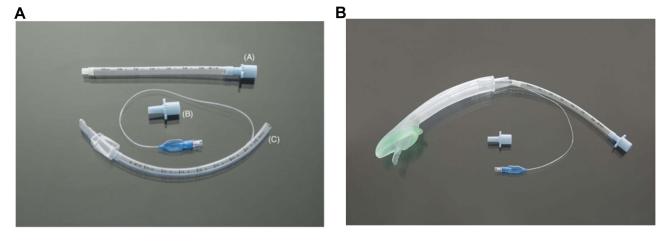


Figure 2 (A) Components of the Rescue™[™] ETT (PVC). (A) Airway tubing. (B) 15 mm ISO connector. (C) ETT with posterior-facing bevel. (B) THE RESCUE™ ETT inserted through an I-GEL™ laryngeal mask airway.

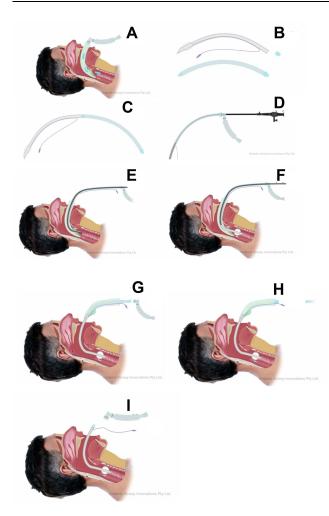


Figure 3 A schematic diagram to demonstrate the method of inserting the Rescue™ ETT through an i-gel™ laryngeal mask airway.

Notes: Rescue[™] ETT; A method of use with a flexible endoscope. (A) An LMA is used to secure the airway. (B) An appropriately sized Rescue™ ETT is selected (the ETT selected must not be forced through the shaft of the LMA as damage to the cuff of the ETT may occur). (C) The 15 mm ISO connector is removed from the ETT and is connected to the airway tubing. The proximal end of the ETT and cuff is well lubricated. (D) A self-sealing connector with a sidearm is placed over the 15 mm connector of the airway tubing. An intubating flexible endoscope with a working length of at least 60 cm and sufficient diameter to abut against the curved atraumatic tip of the ETT is chosen. This complex is then loaded over the endoscope with the endoscope tip just proximal to the tip of the ETT. (E) The LMA is disconnected. The ETT/airway tube is then connected to the anesthetic machine and the distal ETT inserted into the shaft of the LMA. The cuff of the ETT is inflated in the shaft of the LMA once the ETT tube tip is 2 cm out of the distal LMA aperture allowing continued ventilation. (F) The flexible endoscope is then advanced through the vocal cords and into the trachea. The cuff of the ETT is deflated and the ETT/airway tubing is railroaded over the endoscope until the cuff of the ETT is a safe distance below the vocal cords and the cuff is reinflated. (G) The ETT/airway tubing is stabilized. The LMA cuff is deflated, and it is withdrawn carefully over the ETT/airway tube. (H) The ETT and airway tube are disconnected, and the airway tube is removed from the shaft of the LMA and then the inflation line, pilot balloon and inflation valve are delivered through the shaft of the LMA. (I) The 15 mm connector is inserted into the proximal ETT tubing, and the patient is connected to the ventilator.

Safety Concerns with the AIC

The AIC is an airway exchange catheter (AEC), which is often recommended for intubation over an FOB. Of concern is the safety profile. The literature shows that these products can have a high rate of complications, as high as 60%.^{20–22} Pneumothorax, which can occur in up to 11%,²³ is a known complication of high-pressure jet ventilation as a result of barotrauma, auto-PEEP, and dynamic hyperinflation. However, pneumothorax has been reported with the use of an AEC without high-pressure jet ventilation.^{24,25}

We suggest that AEC should only be used by teams that are versatile and trained in their use.

Potential of Uncoupling While Using the Rescue[™] ETT

A potential concern with the RescueTM ETT is the airway tube and ETT uncoupling in the procedure (Figures 2A and B), however, the forces applied while intubating and then withdrawing the LMA over the ETT/airway complex, act to keep this coupling together.

Ability to Maintain Ventilation

When using the Rescue[™] ETT (Figure 3) a self-sealing connector with a sidearm placed over the 15mm connector of the airway tubing that is joined to the ETT. The flexible endoscope is then inserted through the complex with the tip of the endoscope just proximal to the end of the ETT. The sidearm of the self-sealing connector is then attached to a ventilator. The lubricated cuff of the ETT usually forms an adequate seal with the airway tube of the LMA to allow continuous ventilation while the trachea is intubated, alternatively if the seal is not adequate once the tip of the endoscope is just out of the laryngeal mask airway tube the cuff of the ETT may be inflated in the airway tube of the LMA and ventilation commenced with the cuff being deflated before the ETT is railroaded over the endoscope into the trachea. While withdrawing the LMA ventilation is continued, and the flexible endoscope is left in place until the LMA is fully withdrawn to confirm the ETT remains placed at the correct depth.

The Fastrach[™] ETT requires an interruption to ventilation while the laryngeal mask airway is withdrawn over a solid stabilizer rod and then a repeat scope to confirm that the ETT has not migrated.

Conclusion

On review of the literature, it is abundantly obvious that an ETT with a posterior-facing bevel and curved or tapered tip to center the tube on the scope and reduce hang-up is ideal for intubation over a flexible endoscope and that the use of the current standard ETT with a lateral-facing bevel should be discouraged.

The newly described Rescue[™] ETT is a simple and safe device deployed with a simple technique. The Rescue[™] ETT retains all the advantages of the Fastrach[™] ETT and as demonstrated in this article, offers several advantages over both the Fastrach[™] ETT and the Aintree Intubation Catheter[™]. A clinical study is required to verify these claims before being introduced into clinical practice.

Author Contributions

All authors made a significant contribution to the work reported, whether that is in the conception, study design, execution, acquisition of data, analysis and interpretation, or in all these areas; took part in drafting, revising or critically reviewing the article; gave final approval of the version to be published; have agreed on the journal to which the article has been submitted; and agree to be accountable for all aspects of the work.

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Disclosure

Vernon H Vivian is a director of Genesis Airway Innovations Pty Ltd that holds the patents on these devices. André Van Zundert, Tyson L Pardon and Samuel Vivian report no conflicts of interest in this work.

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