Radiologic Imaging of the In Vivo Position of the New Supraglottic Airway Device Spritztube® in an Adult Patient—A Case Report

Silvia De Rosa 1,*, Massimiliano Sorbello 2, Alessandro Rigobello 1, Lucia Cattin 1, Giuseppe Iannucci 3, Paolo Gennaro 1, Vinicio Danzi 1 and Stefano Checcacci Carboni 1

1 Department of Anesthesiology and Intensive Care Medicine, AULSS 8 Berica, San Bortolo Hospital, Viale Rodolfi 37, 36100 Vicenza, Italy
2 Anesthesia and Intensive Care, AOU Policlinico Vittorio Emanuele, 95123 Catania, Italy
3 Department of Neuroradiology, San Bortolo Hospital, 36100 Vicenza, Italy
* Correspondence: derosa.silvia@gmail.com; Tel.: +39-0444-757710 or +39-3933098583; Fax: +39-0444-753973

Abstract: Spritztube® is a new supraglottic airway device that allows either extraglottic ventilation or orotracheal intubation with the same device. The aim of the present report is to provide the first radiologic images of the Spritztube in situ in a living human and to assess the depth of insertion and its anatomical relationships in vivo. We present the case of a 55-year-old man who was admitted to our centre to perform an interventional neuroradiological procedure. We obtained and analysed radiologic images of the head and neck of an adult patient to ascertain the position of the cuffs of the Spritztube relative to different anatomic structures. The insertion and depth of the device, correct tip positioning, effects of the distal and proximal cuffs on adjacent soft tissues, and the position of the pharyngeal cuff from the cranial to the hyoid bone were evaluated. Our report shows that Spritztube could be helpful in maintaining an adequate airway during radiologic procedures.

Keywords: airway management; difficult airway; supraglottic airway device; Spritztube

1. Background

The number of supraglottic airways (extraglottic devices) has progressively increased in recent years [1]. As a general principle, these devices allow access to the distal airway with a pharyngeal, laryngeal, or pharyngolaryngeal seal, enabling oxygenation and ventilation by creating an adequate seal.

Moreover, some extraglottic devices allow the passage of a tracheal tube, which increases the success and safety of fibreoptic bronchoscopy. A re-usable extraglottic, the Spritztube [®Med Europe s.r.l., San Pietro in Casale, Italy], has been designed, developed, and tested in our centre. It consists of a silicone-based tracheal tube with two low-pressure cuffs. The proximal cuff is designed to seal the oropharynx above the epiglottis and the distal cuff is designed to seal the upper oesophagus (Figures 1 and 2). Cuffs are inflated through independent tubing. The Spritztube comes with a long stylet designed to stiffen the device and allow passage through the oropharyngeal space. The positioning of this device requires the aid of this dedicated stylet, designed to align the two cuffs while stiffening the Spritztube so as to allow passage through the oropharynx down to the hypopharynx. The Spritztube can be used for extraglottic ventilation, for tracheal intubation with the direct and indirect laryngoscopes, or for fibreoptic-assisted tracheal intubation [2].

The Spritztube is designed for blind insertion. One holds it as a pen and the device is inserted into the oral cavity following the soft palate, stopping introduction once the black marker on the Spritztube (teeth mark) reaches the upper incisors (Figure 1).
Precisely, a marker on the device, positioned at the incisors, indicates the adequate depth of insertion so that the distal cuff is above the oesophagus and the proximal cuff is above the laryngeal opening (Figures 2 and 3).

**Figure 1.** Spritztube details. The device, after insertion into the oral cavity, is stopped once the black marker on the Spritztube (teeth mark) reaches the upper incisors.

**Figure 2.** Spritztube characteristics (A): device assembled for positioning; note the stylet inside. (B): details of distal (oesophageal) cuff (solid arrow) and oropharyngeal cuff (empty arrow). (C): colour-coded pilot balloons for oesophageal and oropharyngeal cuffs with respective (10 and 60 cc) inflation volumes.
Figure 3. Spritztube details. The device assembled as if in definitive position; note the removed stylet. Once placed as extraglottic device, the distal (oesophageal) cuff (solid arrow) closes the upper oesophageal sphincter while the oropharyngeal cuff (empty arrow) seals the oropharynx. If intubation is performed, the proximal cuff is left inflated and in place, the oropharyngeal cuff deflated and the tube is addressed in trachea with direct view (flexible scope or video-stylet).

Once in position, the proximal cuff is inflated up to 60 cc, until no air leak is perceived. The distal cuff, inside the upper oesophageal sphincter, is inflated using approximately 20 cc of air, and then the stylet is removed.

This position allows the Spritztube to function as a **extraglottic device**. If endotracheal intubation is needed, the proximal cuff is deflated, and the tube is addressed into the trachea without repositioning the stylet, under direct or indirect laryngoscope, flexible endoscope, or video-stylet view.

Once intubation has been performed, the distal cuff can either be deflated or left in position to act as an oesophageal obturator [2] (Figure 4). As a normal endotracheal tube, average distances such as the 20/22 or 21/23 (cm, female/male) rule have been simply applied.

The Spritztube is available in unique and adult sizes, and it is recently available on the market (given the recent certification mark for conformity with health, safety, and environmental protection standards for products sold within the European market). It was provided for our case report in order to elucidate how it might allow for airway control and whether the two cuffs distort the pharyngeal anatomy.

This is the first report that provides the first images of the Spritztube device in situ in a living human and assesses the depth of insertion and its positioning in vivo. The patient was part of a clinical trial published previously. We obtained and analysed radiologic images of the head and neck of an adult patient under general anaesthesia to ascertain the position of the cuffs of the Spritztube relative to anatomic structures. Written informed consent was obtained from the patient to publish this case report.
2. Case Presentation

A patient enrolled in a first-in-human trial of the Spritztube. This trial compared Spritztube tracheal cannula with the Laryngeal Mask Airway Supreme in anaesthetised adult patients, showing the effectiveness of the Spritztube in maintaining the airway with all patients being successfully ventilated without difficulty. The success rate of achieving a patent airway was comparable between the groups, with a similar occurrence of complications [3].

The patient gave written informed consent for the anaesthetic procedure and for publication of this case report, including accompanying images. Effect on adjacent soft tissue are shown in Table 1. The patient was a 55-year-old male, 109 kg in weight and 183 cm in height [Body Mass Index 32.5 kg/m²], admitted to our centre to perform a diagnostic neuroradiological procedure. He was classified as American Society of Anesthesiologists status of II with no indications of a difficult airway (El-Ganzouri Risk Index 2, Mallampati 1, thyromental distance 6.5 cm, inter-incisor distance 4 cm full neck movement, no history of previous difficult airway management). Radiological images were obtained to assess the supraglottic use of the Spritztube (distal cuff inflated in oesophagus, proximal cuff inflated in oropharynx). Angiography was performed for diagnostic purposes for suspected aneurysm. The general anaesthesia protocol, intra-operative monitoring, and postoperative care performed for the patient included HR, non-invasive blood pressure, peripheral oxygen saturation, and continuous electrocardiography. The premedication used was midazolam 0.03 mg kg⁻¹, followed by fentanyl 2 μg kg⁻¹, and propofol 2 mg kg⁻¹ for induction of anaesthesia. Anaesthesia was maintained with propofol (continuous infusion 6 to 8 mg kg⁻¹ h⁻¹) and remifentanil (continuous infusion 0.25 to 0.75 μg kg⁻¹ min⁻¹). Neither nitrous oxide nor a neuromuscular blocking agent were given, and depth of anaesthesia was monitored with the bispectral index (BIS monitor; Covidien Medical, Louisville, CO, USA), targeting the range 40 to 60. The Spritztube was inserted by trained anaesthesiologist with 10 years of experience.
Table 1. Effect on adjacent soft tissue after device insertion and with cuffs inflated.

| Depth of Insertion Parameters | Measurement (cm) |
|------------------------------|------------------|
| Cervical vertebra to tip      |                  |
| Oesophageal                   | 0.87             |
| Oropharyngeal                 | 1                |
| Hyoid bone to tip            |                  |
| Oesophageal                   | 8.74             |
| Oropharyngeal                 | 8.49             |
| Mandible to tip              |                  |
| Oesophageal                   | 11.8             |
| Oropharyngeal                 | 8.66             |
| Effect on trachea diameter   |                  |
| Oesophageal                   | 1.35             |
| Oropharyngeal                 | 2.85             |
| Position of hyoid bone relative to mental spine of mandible | |
| Oesophageal                   | 8.15             |
| Oropharyngeal                 | 7.02             |
| Position of hyoid bone relative to cervical spine | |
| Oesophageal                   | 7.40             |
| Oropharyngeal                 | 8.95             |

The Spritztube was then inserted based on the blind insertion technique after anaesthetic depth. Once the Spritztube was positioned (marker at the level of the incisors) and effective ventilation obtained, we visually assessed the depth of insertion and the impact of the Spritztube on adjacent soft tissues through fluoroscopic images (Figure 5).

Figure 5. Radiological Localisation of Spritztube (ST) as extraglottic device. Dash lines are used to show the depth of insertion and the effect on trachea diameter. Continuous lines are used to show the effect of the ST on adjacent soft tissues. Held as a pen, the device was inserted into the oral cavity (A) following the soft palate (B,C), stopping introduction once the black marker on the Spritztube (teeth mark) reached the upper incisors (D). Then, distal and proximal cuffs were inflated, the stylet removed and the tube connected to the breathing circuit (E,F). (A–D): notice the presence of radio-opaque ST stylet, subsequently removed in (E,F). (F) shows oropharyngeal cuff inflated. Abbreviations: D1, Distance 1; D2, Distance 2; E, Epiglottis; Es, Oesophagus; H, Hyoid Bone; T, Trachea.
The depth of insertion was assessed:

1. Relative to hyoid bone. As the epiglottis is located behind the hyoid bone, the depth of insertion relative to the glottic opening was defined as the perpendicular distance between the hyoid bone and the tip of the Spritztube device in the upper oesophageal sphincter on sagittal images (Figure 5D,F).

2. Relative to the cervical spine. This distance was defined as the distance between the superior surface of the body of the third cervical vertebra and the tip of the Spritztube on sagittal images (Figure 5D,F).

3. The position of the hyoid bone relative to the cervical spine. This measurement was determined as the distance between the ventral margin of the hyoid body and the anterior margin of the body of third cervical vertebra (Figure 5).

4. The position of the ventral margin of the hyoid body relative to the mental spine of the mandible. This measurement was determined by calculating the distance between the medial portion of the hyoid bone and the lower margin of the mandible.

The effect of cuff inflations on tracheal diameter was also assessed.

All detected measurements are shown in Table 1.

3. Discussion and Conclusions

The Spritztube is a new extraglottic device designed to be used in general anaesthesia during spontaneous or positive pressure ventilation, or as alternative to endotracheal intubation, particularly in emergency situations. In this case report, the radiological position of the Spritztube in a patient was evaluated for the first time using fluoroscopic images in addition to clinical assessments. The insertion and depth of the device, placed as endotracheal tube, correct tip positioning, effects of the distal and proximal cuffs on adjacent soft tissues, and the position of the pharyngeal cuff from the cranial to the hyoid bone were evaluated.

The correct tip positioning could be affected by body weight, oropharyngeal and neck anatomy, and relative positions of the hyoid, mandibular, and other structures which are simultaneously relevant aspects resulting in a difficult or failed intubation [4]. In this case, Body Mass Index suggested moderate obesity but any other parameter influencing airway management was deemed to be normal according to a recently published consensus [5], and no difficulty was recorded or experienced for the Spritztube insertion. The first aim of our report was to assess the relations between the ST and neck anatomy using the fluoroscopy, allowing also possible comparisons with extraglottic devices already in use. Some considerations should be performed. Firstly, the insertion of the Spritztube is mediated by a rigid stylet that is effective when used by experienced operators. In this case, the clinician must be alert to the advancement of the rigid stylet device in the airways as it has a high potential for lacerations and perforations of the oropharyngeal mucosa and subglottic structures. Secondly, although adverse effects due to high intracuff pressures may not commonly cause serious adverse effects, it would be prudent to minimise this risk by ensuring the availability of cuff pressure manometers.

Thirdly, Spritztube is a re-useable device: not all protein material can be removed by routine cleaning and this raises theoretical concerns over cross-infection risk, hence steam autoclaving is the recommended method of sterilising this device. New Spritztube devices will be soon released on the market for a single use.

The role of extraglottic devices is well recognised for non-operating room anaesthesia, and the novelty of our report is in adding our new device to the portfolio of available alternatives. Particularly, some specific features of our device might offer interesting advantages, such as the possibility of early switching from extraglottic position to endotracheal intubation. In addition, since fluoroscopy is available for the neuroradiologic procedure, our case report also describes the depth and position of the Spritztube relative to soft tissues and the cervical spine. The resulting anterior displacement of the hyoid bone is marginal and different when compared to other extraglottic airway devices, such as i-gel™ and LMA-Supreme™ [6].
This specific finding might have important implications, given the evidence that extraglottic device cuff hyper-inflation-related damage to the lingual, hypoglossal, recurrent, and glosso-pharyngeal nerves seems to be a consequence of compression against the hyoid bone [1,7,8]. These findings need to be prudently interpreted taking into account inter-individual anatomical variability of neck anatomy as well as the single case within our report. Although those measurements indicate the effects of Spritztube on the laryngeal structures, magnetic resonance imaging, which would not really be feasible, may better confirm, define the position of the device, and clarify the correlation between these parameters and the anthropological airway anatomy parameters.

Further studies are necessary to evaluate the safety, efficacy, and potential for the Spritztube in airway management.

Author Contributions: G.I., S.C.C., P.G. and A.R. managed the patient; S.D.R., M.S., L.C. and A.R. wrote the first draft of the manuscript and provided revisions; S.C.C., G.I., V.D. and P.G. critically revised the manuscript. All authors have read and agreed to the published version of the manuscript.

Funding: This research received no external funding.

Institutional Review Board Statement: Ethics approval and Consent to participate was obtained from St. Bortolo Hospital Ethics Committee (Study ID Number 29/16).

Informed Consent Statement: Written informed consent was obtained from the patient for publication of this case report and accompanying images. It is available for the journal.

Data Availability Statement: Data sharing is not applicable to this article as no datasets were generated or analysed during the current study.

Acknowledgments: We acknowledge the health personnel of the San Bortolo Hospital who took part in the management of this patient.

Conflicts of Interest: S.D.R. is Associate Editor of BMC Anesthesiology; A.R., E.F., P.G. and G.I. have no conflict of interest; S.C.C. is the inventor of the Spritztube, that is marked CE. There is no financial or personal relationship with people or organisation that could inappropriately influence the work; M.S. research and development and pre-clinical testing of the LMA Protector™. Paid Consultancy with Teleflex Medical, Dublin, Ireland. Patent owner (no royalties), paid consultancy with DEAS, Castelbolognese, Italy. M.S. is Associate Editor of BMC Anesthesiology.

Abbreviations

ST Spritztube

References

1. Sorbello, M.; Petrini, F. Supraglottic Airway Devices: The Search for the Best Insertion Technique or the Time to Change Our Point of View? Turk. J. Anaesthesiol. Reanim. 2017, 45, 76–82. [CrossRef] [PubMed]
2. DE Rosa, S.; Ferrari, F.; Carboni Checcacci, S.; Rigobello, A.; Gennaro, P.; DE Luca, D.; Primadei, M.; Politi, F.; Pellizzari, A.; Bonato, R. Airway management with Fastrach laryngeal mask versus Spritztube: A prospective randomized manikin-based study. Minerva Anestesiol. 2018, 84, 455–462. [CrossRef] [PubMed]
3. De Rosa, S.; Messina, A.; Sorbello, M.; Rigobello, A.; Colombo, D.; Piccolo, A.; Bonaldi, E.; Gennaro, P.; Urukalo, V.; Pellizzari, A.; et al. Laryngeal Mask Airway Supreme vs. the Spritztube tracheal cannula in anaesthetised adult patients: A randomised controlled trial. Eur. J. Anaesthesiol. 2019, 36, 955–962. [CrossRef] [PubMed]
4. Petrini, F.; Accorsi, A.; Adrario, E.; Agrò, F.; Amicucci, G.; Antonelli, M.; Azzeri, F.; Baronzini, S.; Bettelli, G.; Caffaggi, C.; et al. Recommendations for airway control and difficult airway management. Minerva Anestesiol. 2005, 71, 617–657. [PubMed]
5. Petrini, F.; Di Giacinto, I.; Cataldo, R.; Esposito, C.; Pavoni, V.; Donato, P.; Troilo, A.; Merli, G.; Sorbello, M.; Obesity Task Force for the SIAARTI Airway Management Study Group; et al. Perioperative and periprocedural airway management and respiratory safety for the obese patient: 2016 SIAARTI Consensus. Minerva Anestesiol. 2016, 82, 1314–1335. [PubMed]
6. Russo, S.; Cremer, S.; Eich, C.; Jipp, M.; Cohnen, J.; Strack, M.; Quintel, M.; Mohr, A. Magnetic resonance imaging study of the in vivo position of the extraglottic airway devices i-gel™ and LMA-Supreme™ in anaesthetized human volunteers. Br. J. Anaesth. 2012, 109, 996–1004. [CrossRef] [PubMed]
7. Brimacombe, J. Problems. In Laryngeal Mask Anesthesia; Brimacombe, J., Ed.; Saunders: Philadelphia, PA, USA, 2005; pp. 551–575.
8. Renes, S.H.; Zwart, R.; Scheffer, G.J.; Renes, S. Lingual nerve injury following the use of an i-gel laryngeal mask. Anaesthesia 2011, 66, 226–227. [CrossRef] [PubMed]