Post-tracheostomy tracheo-oesophageal fistula – an unusual presentation

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Tracheostomy, one of the oldest surgical procedures known to mankind. It was mentioned in the ancient Sanskrit script Rigveda, dating as far back as 2000 BC. The first successful elective human tracheostomy was performed in 1869 for the administration of general anaesthesia.[1] It is a commonly performed bedside procedure in the modern intensive care unit (ICU); however, it is not devoid of complications. Acquired ulcerative tracheo-oesophageal fistula (TOF) is a rare, but potentially fatal, complication of tracheostomy. We report a case of ulcerative TOF with an unusual yet characteristic presentation, in a ventilator-dependent tracheostomised patient with Guillain-Barré syndrome.

Case report

A 28-year-old man with marfanoid habitus was admitted to our ICU with GBS. He was intubated and mechanically ventilated due to respiratory failure. His muscle power did not improve despite intravenous immunoglobulin and he continued to be ventilator-dependent. Surgical tracheostomy using a size 8.0 Portex (Smiths Medical, USA) polyvinylchloride (high volume, low pressure cuff) tracheostomy tube (TT) was performed after 2 weeks. He had anaemia and weight loss despite enteral nutrition through a 12F nasogastric (NG) tube. He developed a bronchopleural fistula on the right side following intercostal drain insertion for spontaneous pneumothorax. He acquired ventilator-associated tracheobronchitis and pneumonia, and worsened into septic shock. His airway pressures were continually maintained below 30 cm H2O. The tracheostomy tube was changed every 3 weeks as per the ICU protocol. As there was no cuff pressure manometer available in our ICU, we were unable to monitor the TT cuff pressure accurately. His mode of communication with the unit staff and relatives was through lip, head and neck movements.

From the 76th post-tracheostomy day onwards, there was an audible air leak around the TT cuff and increasing volumes of air seal (10 mL, increasing to 14 mL) were required daily. This raised suspicion of tracheal dilatation; however, bedside radiograph of the neck showed a normal cuff air shadow. On the 83rd post-tracheostomy day, we noticed rapidly progressive abdominal distension with each mechanical breath from the ventilator. The epigastric distension was relieved immediately on opening the proximal end of the NG tube. A clinical diagnosis of TOF was made. A bedside radiograph showed a TT cuff air shadow extending beyond the confines of the trachea and a grossly distended stomach (Fig. 1).

Fibreoptic bronchoscopy (FOB) revealed a TOF (3 cm long and 1 cm wide) in the right posterolateral tracheal wall, situated distal to the equator of the cuff and extending almost to the tip of the TT. After removal of the TT, orotracheal intubation with endotracheal tube (ET) size 8.0 was performed under FOB guidance and the cuff was positioned distal to the TOF. The tracheostomy stoma was then closed by suturing. Definitive surgical repair was planned once sepsis was treated. Unfortunately, the patient succumbed to septic shock 10 days after diagnosis of TOF.
inflation pressures, increasing minimal occluding cuff volumes (more than 10 mL of air) and unexplained persistent air leak around the TT cuff should raise concerns about tracheal dilatation and TOF. FOB is the gold standard for the diagnosis of TOF. Postero-anterior radiographs of the neck and superior mediastinum may show marked bulging of the cuff beyond the normal confines of the trachea. A special TT with the cuff positioned distal to the fistula may help to re-establish adequate ventilation and prevent soiling of the tracheobronchial tree.[10] Feeding can be managed through gastrostomy or jejunostomy tube.

Regular monitoring of tracheal tube cuff volume and pressure is important. It is advisable to maintain cuff pressures below 20 cm H₂O[11] and avoid using more than 8 mL of air for cuff inflation.[10] Routine cuff deflation and re-inflation are no longer recommended.[12,13] Proper placement of the tracheostomy and tracheostomy tube, use of thin-walled cuffs to minimise movement of the tracheostomy tube relative to the trachea and initiation of weaning as early as possible may help in curtailing trauma to the trachea. Conscious and well-orientated tracheostomised patients should be educated and advised about avoiding excessive neck movements. Measures to minimise gastro-oesophageal reflux, such as head-end elevation to 30°, post-pyloric feeding, administration of enteral feeds as continuous infusion and use of prokinetics, should be implemented. NG tubes, if used, should be small and soft and removed as soon as they are no longer indicated. Predisposing factors such as hypotension, hypoxaemia, anaemia, infection, malnutrition and metabolic acidosis should be prevented and, if present, addressed early.

Conclusion

TOF is a rare but potentially fatal complication of tracheostomy. Rhythmic and rapid distension of the epigastrium, though unusual, is a dangerous manifestation of TOF. Regular monitoring of tracheal tube cuff pressures and cuff volumes is advisable for the prevention and early recognition of this adverse event.

References

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