Upper airway obstruction due to inoperable intrathoracic goitre treated by tracheal endoprosthesis

M Noppen, M Meysman, E Dhondt, L Gepts, B Velkeniers, L Vanhaelst, W Vincken

Abstract
Thyroidectomy is the treatment of choice in patients with thyroid enlargement complicated by compression or displacement of the trachea because of the risk of complete airway obstruction due to sudden enlargement of the goitre by, for example, haemorrhage. In patients who are medically inoperable an endoscopically inserted tracheal endoprosthesis may provide longstanding airway patency, as reported here.

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Insertion of tracheal, bronchial, or tracheobronchial stents can offer excellent palliation in patients with central airway obstruction who are not amenable to surgery.12 Tracheal or bronchial malignant processes with extrinsic airway compression constitute the major indication for airway stenting, although longstanding success has been achieved in patients with benign tracheostenosis following intubation.3 We therefore attempted tracheal stenting in two patients with acute upper airway obstruction due to non-malignant intrathoracic goitre in whom thyroidectomy was contraindicated.

Case 1
A 90 year old woman was surgically treated for a left femoral fracture in a general hospital under locoregional anaesthesia. Postoperatively she developed acute severe inspiratory and expiratory dyspnoea. She was known to have a large intrathoracic goitre, causing tracheal displacement and narrowing, which was responsible for her acute respiratory distress.

The patient, in otherwise good general condition, was transferred to our department for endoscopic treatment. On examination she was conscious, tachypnoeic (25 breaths/min) and anxious. Both inspiration and expiration were shallow and accompanied by a permanent wheeze. Physical examination was otherwise normal except for the scar of the recent hip intervention. There were no signs of venous thrombosis.

A radiograph of the trachea showed severe displacement and narrowing by the intrathoracic goitre. After topical anaesthesia of the oropharynx (10% lignocaine spray) and vocal cords (2% lignocaine, 5 ml) a fibreoptic bronchoscopy was performed which showed severe compression of the trachea from both sides (residual lumen 6 mm diameter) over a length of 4.5 cm (fig 1a). The distal airways (which were quickly inspected) appeared normal.

Since thyroidectomy was considered to be contraindicated in this elderly patient, endoscopic placement of a tracheal
endoprosthesis was attempted. Under light general anaesthesia with intravenous propofol rigid bronchoscopy (Storz, Germany) was performed. Ventilation and oxygenation were maintained with high frequency jet ventilation delivered via a side port of the bronchoscope. A silicone Dumon tracheal endoprosthesis (Endoxane TD, diameter 12 mm, length 4 cm; Cometh Laboratoire, Marseille, France) was longitudinally folded and pushed through the siliconised lumen of the properly positioned bronchoscope through the stenosis. Under direct vision the prosthesis was unfolded with forceps. After ensuring proper placement and fixation of the prosthesis, the bronchoscope was removed. The entire procedure lasted 20 minutes and the postoperative course was uneventful. Follow up fibreoptic bronchoscopy the next day showed good airway patency (fig 1b); the patient was discharged the same day.

There was no recurrence of upper airway obstruction four months after insertion of the stent.

Case 2
An 89 year old woman with a large multinodular goitre presented with rapidly progressive dyspnoea, stridor, and dysphagia due to a sudden increase in size of her goitre following an intrathyroid haemorrhage. On examination she was tachypnoeic with stridors at rest; vital parameters were normal. Radiographic examination and tomographic scanning showed narrowing and displacement of the trachea. Spirometry and a flow-volume loop was suggestive of a fixed mixed intrathoracic and extrathoracic upper airway obstruction. Fibreoptic bronchoscopy showed bilateral wall narrowing and tortuous displacement of the trachea over 4 cm; the diameter of the tracheal lumen was about 5 mm (fig 2a).

Surgery was considered to be contraindicated. A silicone Dumon tracheal endoprosthesis (Endoxane TD, diameter 14 mm, length 4 cm) was inserted into the stenosis via a rigid bronchoscope under general anaesthesia using high frequency jet ventilation (fig 2b). The immediate postoperative course was uneventful with complete disappearance of dyspnoea and stridor. However, 48 hours after the procedure acute stridor recurred; urgent bronchoscopy showed a stent obstruction due to fibropurulent debris. After bronchial toilet, oedematous tracheal mucosa was visualised just distal to the stent. The stent was removed and the mucosal obstruction was vapourised with Nd:YAG laser photodestruction. The stent was reinserted and stable airway patency and respiration were restored. Daily inhalations of a mucolytic agent (mesna) were prescribed.

Over the following days there were occasional short episodes of dyspnoea, probably due to partial stent obstruction by secretions in the very tortuous trachea. A reintervention one month later was therefore performed with removal of the first stent which was replaced by a longer (5 cm) model.

Thereafter the patient was completely asymptomatic. She died two months later due to an acute myocardial infarction. At necropsy the stent was completely patent.

Discussion
Thyroidectomy is the treatment of choice for most intrathoracic goitres because of the risk of intrathyroidal haemorrhage, especially when symptoms or signs of tracheal obstruction are present. Surgery is recommended when peak expiratory flows decrease to <100 l/min.7 In high risk patients, or those in whom surgery is contraindicated, alternative means of airway protection must be sought.

In patients with intrinsic or extrinsic tracheal obstruction who are not amenable to surgery, placement of tracheal endoprostheses (stents) can offer palliation.1 Of the various stents commercially available, the silicone Dumon2 and the expandable metal Gianturco Z stent3 are the most widely used. Both are easily inserted, are efficacious in stenoses up to 5 cm in length, and are well tolerated. Advantages of the silicone Dumon stents are the possibility of adjustment and removal (which proved to
Massive haemothorax secondary to angiosarcoma

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Abstract
A patient who presented with recurrent haemoptyses was found to have an angiosarcoma of the adrenal gland which disseminated throughout the pleural space.

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Case report
A 44 year old man presented with daily haemoptysis for one week, but was otherwise well. He had smoked 20 cigarettes per day for 20 years and had not been exposed to asbestos, arsenic, thorotrust, vinyl chloride, or radiotherapy. Examination was unremarkable, with no finger clubbing, a clear chest, and no carotid or vertebral bruits. Chest radiography and fiberoptic bronchoscopy were normal. He continued to have haemoptysis and four weeks later he was admitted as an emergency, acutely short of breath. A chest radiograph now showed large bilateral pleural effusions. Aspiration proved these to be haemorrhages. His haemoglobin concentration was 6.0 g/dl. He was transferred to the regional cardiothoracic centre and four and one litres of blood were drained via left and right chest drains respectively. A computed tomographic (CT) scan of the thorax demonstrated the haemorrhages but was otherwise normal. However, a CT scan of the abdomen revealed a left adrenal mass measuring 5 × 4 × 4 cm (fig 1).

A further 20 litres of blood were drained over the following five days. Clotting screens were normal and the adrenal tumour appeared to be non-functioning (urinary levels of 17-ketosteroids and vanillylmandelic acid were normal). Pulmonary angiography and selective angiography of the intercostal arteries failed to demonstrate a definite bleeding point.

On the sixth day adrenalectomy and exploratory thoracotomy were performed. The adrenal mass was a well circumscribed cystic lesion filled with organising blood clot. Two litres of blood were removed from the left hemithorax to reveal visceral and parietal pleura covered in organising blood clot and studded with multiple haemorrhagic cysts 5–30 mm in diameter. The lung parenchyma appeared to be normal. Frozen sections of the lesions were reported as angiosarcoma. Bleeding was controlled and the chest closed.

Postoperatively the patient had a dense right sided hemiplegia. However, he remained stable.
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