The role of magnetic resonance imaging in the assessment of suspected extrinsic tracheobronchial compression due to vascular anomalies

T H Malik, I A Bruce, V Kaushik, D J Willatt, N B Wright, M P Rothera


METHODS

A database review at our paediatric tertiary referral centre was undertaken. Cases were identified where there had been a clinical suspicion of extrinsic tracheobronchial vascular compression recorded at DLTB. Typically this suspicion would involve a pulsatile indentation or constriction of the tracheobronchial wall with reduction of the cross-sectional area at characteristic sites and configurations described later. The DLTB was performed using a rigid laryngoscope through which a 2.7 mm 0° rigid endoscope (Hopkin’s rod) was introduced. By performing a dynamic procedure any malacic segments would also become apparent. The level of anaesthesia was such that it prevented coughing but allowed for spontaneous ventilation using nasopharyngeal intubation, with the end of the tube situated in the hypopharynx. The vocal folds were sprayed with local anaesthetic spray.

These children then routinely underwent thoracic MRI (T1 weighted, echo planar imaging in three orthogonal planes (spin-echo with EPI) and additional turbo field echo transverse images to provide angiographic information) within 10 days of the DLTB. Sedation was given during MRI by means of oral chloral hydrate 100 mg/kg (maximum 2.0 g) with or without rectal paraldehyde 0.3 ml/kg for children weighing <20 kg, and oral quinalbarbitone (secobarbital) 10 mg/kg (maximum 200 mg) for those weighing >20 kg.

Abbreviations: DLTB, dynamic laryngotraceobronchoscopy; MRI, magnetic resonance imaging
A case note review was conducted and the findings on endoscopy were then compared to those of MRI, and where performed, echocardiography, aortography, and surgery.

RESULTS
We identified 29 cases where children had a clinical suspicion of external vascular compression on DLTB. All subsequently underwent thoracic MRI as is the policy at our unit. There were 17 males and 12 females, with a mean age of 5 months (range 28 weeks gestation to 60 months). The presenting features are summarised in table 1.

Of the 29 patients, MRI was normal in eight children (27.5%). None of the patients with normal magnetic resonance scans were later diagnosed as having vascular compression, either clinically or using other radiological modalities, during the follow up period of at least 18 months. Anomalies found on MRI are shown in table 2, and depicted in figs 1–8 (see the ADC website for figs 1, 2, and 5–8: www.archdischild.com/supplemental).

Echocardiography was performed in 19 of the 29 cases. Apart from hinting at one case of a right aortic arch, and another of left pulmonary artery aneurysm, this investigation was found to be unhelpful in the diagnosis of extra-cardiac vascular anomalies. Barium swallow was performed in two of the five cases later diagnosed as vascular rings: one was correctly diagnosed as a right aortic arch, but the other failed to visualise a double aortic arch.

Eight of the twenty one patients with abnormalities identified on MRI went on to have aortography. The findings at aortography were in agreement with the MRI findings in all five vascular rings, two cases of innominate artery compression (causing >50% tracheal compression), and the case of aneurysmal left pulmonary artery.

Surgery was performed in seven cases, comprising all five vascular rings, one innominate artery compression (causing >50% tracheal compression), and the aneurysmal left pulmonary artery. The findings at operation were in concordance with the preoperative MRI in all cases.

DISCUSSION
Developmental anomalies of the aortic arch and its branches have two effects on the upper airway. Firstly they cause direct extrinsic compression with reduction of the luminal area, and secondly they cause malacia because the airway wall weakens and becomes more compliant and collapsible. Mucociliary clearance of secretions can also be impaired, compounding the functional airway obstruction and leading to infections and chronic cough. 7

The most common presenting complaint in our series was stridor, which occurred in 69% of patients, in agreement with the literature. 2 9 10 Fifteen patients presented with either episodic apnoea or cyanotic episodes. Such episodes are also referred to as reflex apnoea or “death spells”. 9 It has been hypothesised that such episodes of respiratory arrest result from tracheal irritation at the site of compression. 10

Airway obstruction resulting from innominate artery compression (IAC) (see figs 1 and 2 on the ADC website: www.archdischild.com/supplemental) may result in an expiratory or biphasic wheeze, or stridor. Patients may also present with a “croupy” barking cough, reflex apnoea, and recurrent lower respiratory tract infections. 11 Any compromise associated with IAC is often self-limiting and improves with increasing age. 11 At DLTB a right anterior compression is seen above the carina, synchronous with the pulse. IAC has been shown to be associated with oesophageal atresia and severe gastro-oesophageal reflux disease. 1 IAC with up to 90% tracheal compression can usually be treated expectantly, but severe or complicated cases require aortopexy whereby the innominate artery is secured to the posterior surface of the sternum, thus relieving the anterior tracheal compression. 3 While MRI elegantly shows IAC, it does not predict the need for surgical intervention: 4 in our series only one child required aortopexy.

A vascular ring refers to an anomaly of the aortic arch system and its major branches that encircles and compresses...
What is already known on this topic

- Approximately 3% of the general population have a congenital anomaly involving the aortic arch system. However, only a small proportion of these result in symptomatic vascular compression of the airway.
- The evaluation of children suspected of having such a problem may include the use of chest radiography, barium oesophagography, echocardiography, CT, MRI, endoscopy, or angiography.

What this study adds

- Thoracic MRI can be used successfully to investigate children with endoscopically suspected vascular compression of the upper airway. It provides excellent detail without radiation or contrast exposure.

The trachea and oesophagus. The most common symptomatic complete vascular ring is the double aortic arch (figs 3 and 4), followed by a right aortic arch in which the ring is completed by the ligamentum arteriosum (see figs 5 and 6 on the ADC website: www.archdischild.com/supplemental). The most common incomplete ring is an aberrant right subclavian artery. The complete rings tend to present earlier.

A double aortic arch (figs 3 and 4) is formed when the embryonic fourth aortic arches and dorsal aortic roots persist and remain patent on both sides. The resultant ring is tight and symptoms usually begin at birth. The stridor is biphasic in nature and exacerbated by feeding. Recurrent respiratory tract infections and cyanotic episodes are common. Significant dysphagia is not usually seen until solid food is introduced. One of the arches is usually of larger calibre, the right arch usually predominating as shown in our three cases. The findings on endoscopy are a concentric or triangular compression surrounding the trachea, main bronchi, and oesophagus. Double aortic arches are treated by ligation and division of the smaller, less dominant arch, which in 80% of cases corresponds to the left aortic arch. The development of the right aortic arch (see figs 5 and 6 on the ADC website: www.archdischild.com/supplemental) is similar to the double aortic arch, but the left arch regresses. Several variations exist, including right aortic arch with aberrant left subclavian artery and mirror image branching. Unlike a double aortic arch anomaly, a right aortic arch with mirror image branching is often associated with congenital heart disease such as tetralogy of Fallot or pulmonary atresia. Persistence of the dorsal segment of the right aortic arch results in an aberrant origin of the right subclavian artery (see figs 7 and 8 on the ADC website: www.archdischild.com/supplemental). The latter now passes behind the trachea and oesophagus, indenting the oesophagus, to reach the right upper limb. This anomaly is not a true complete vascular ring and is often asymptomatic. It is the most common congenital arch anomaly occurring in 0.4–2.3% of the population. Significant airway compromise is unlikely, but this anomaly can be associated with dysphagia.

A pulmonary artery sling is produced as a result of an anomalous origin of the left pulmonary artery. In this anomaly the normal left pulmonary artery is absent and the resultant aberrant collateral vessel arises from the right pulmonary artery, tightly encircling the right bronchus and passing between the trachea and oesophagus. Significant airway compromise is common and therefore presentation occurs shortly after birth. Tracheomalacia and tracheal stenosis may also result.

Non-surgical management has been shown to be effective in selected mildly symptomatic cases of vascular tracheobronchial compression, especially innominate artery compression. Outcomes following surgical treatment are good, with complete resolution of symptoms in most cases.

Traditionally the gold standard investigation has been aortography with its inherent risks associated with exposure to ionising radiation and the use of contrast media, as well as the potential complications related to the puncture site. Other imaging modalities have been advocated in the investigation of suspected vascular tracheobronchial compression, such as contrast imaging of the oesophagus, echocardiography, and computed tomography (CT).

A high degree of clinical suspicion remains the most important factor in the diagnosis of paediatric airway obstruction resulting from vascular compression. Our preferred diagnostic strategy for cases of airway obstruction involves the primary use of DLTB. It provides an unrivalled view of the entire airway, has the capability to diagnose a variety of causes of airway obstruction (for example, laryngomalacia, vocal cord palsy, and subglottic haemangioma), and permits a functional assessment of airway compromise. At present, technology is not advanced enough for MRI to be able to perform useful real-time “virtual endoscopy” and thus be considered a viable first line alternative.

The combination of DLTB and MRI proved to be a robust method of diagnosis in all the patients in this study. This study supports the first line use of MRI to investigate all endoscopically suspected extrinsic vascular compression of the upper airway.
Vascular anomalies

We found the T1 EPI “black blood” sequence to be the most useful sequence at our institution. If MRI confirms an anomaly which would require surgical intervention or is indeterminate, then one should proceed to angiography.

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REFERENCES


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