A 78 year old man with a history of hypertension, arteriosclerosis, and a myocardial infarction woke up one morning to realise that he had lost his voice. He did not feel ill nor had he experienced symptoms of cough or fever. He had smoked cigarettes for over 60 years and had a brother with lung cancer. Left vocal cord paralysis was assessed at laryngoscopy.

A contrast enhanced computed tomographic (CT) scan of the chest showed a mass with a central opacity in the aortopulmonary window (fig 1, arrow). In the diagnostic workup densities of this mass were measured which did not correspond with contrast as seen in the aorta. A mediastinal metastasis or mediastinal tumour was the most likely diagnosis given by the radiologist. As he was not aware of the clinical entity of Ortner’s syndrome, an aneurysm was considered less likely. Transoesophageal endoscopic ultrasonography revealed a saccular aneurysm of the aortic arch surrounded by a haematoma (fig 2). Using colour Doppler, flow was detected through a 5 mm connection between the aorta and the aneurysm (fig 2, arrow).

Only a few such cases have been described, possibly not as well documented as this one. An aortic arch aneurysm is a rare cause of cardiovocal hoarseness, also known as Ortner’s syndrome, first described in 1897 in patients with left atrial enlargement due to mitral valve stenosis.1 In this syndrome the left recurrent laryngeal nerve is thought to be injured due to compression caused by changes in the anatomy of the heart or great vessels, most often by left atrial enlargement due to mitral valve stenosis. Endoscopic ultrasonography may be the diagnostic procedure of choice in establishing the diagnosis.

REFERENCE

A rare cause of Ortner's syndrome (cardiovocal hoarseness)

J T Annema, J J Brahim and K F Rabe

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