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A 55-year-old man with mild shortness of breath

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Answer

The correct answer is C.

The chest radiograph shown in Fig. 1a of the Question shows the silhouette of a right-sided aortic arch coursing to the right of the trachea. This is a rare, often asymptomatic and incidental finding in the setting of an otherwise structurally normal heart, which was first described in 1763 [1]. It is estimated to occur in 0.04–0.1% of the population. About half of the cases are associated with an aberrant left subclavian artery, as is illustrated in our patient (type II right-sided aortic arch variant). Other types of right-sided aortic arch variants include mirror-image arch branches (type I) and an isolated left subclavian artery communicating with the pulmonary artery (type III). Type I right aortic arch in particular is strongly associated with congenital heart disease, most commonly tetralogy of Fallot.

There is no evidence of acute aortic pathology (associated with a second 'false' lumen and dissection flap in the aortic wall), left persistent vena cava superior (associated with a dilated coronary sinus) or a double aortic arch on the images presented. Therefore, we can exclude these options.

A three-dimensional reconstruction of the CT scan (Fig. 1a, b) facilitates our comprehension of the anatomy of the right-sided aortic arch giving rise to a left and right common carotid artery, a right subclavian artery and an aberrant left subclavian artery. The left subclavian artery originates from a vascular pouch, referred to as the Kommerell diverticulum (alternatively called remnant diverticulum or lusoria root) posterior to the trachea and oesophagus, poten-

A. D. Egorova $(\boxtimes) \cdot J. M. Smit \cdot P. Kiès$ Department of Cardiology, Heart Lung Centre, Leiden University Medical Centre, Leiden, The Netherlands a.egorova@lumc.nl tially compressing these structures. Its aneurysmatic enlargement can lead to dissection and rupture, making it an important clinical phenomenon to recognise [2]. CT and MRI are essential tools for evaluation of this pathology. The timing of the (often complex surgical) management of this congenital pathology remains a matter of debate.

Conflict of interest A.D. Egorova, J.M. Smit and P. Kiès declare that they have no competing interests.

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Fig. 1 A three-dimensional reconstruction of the CT angiography showing the frontal view (**a**) and the left lateral view (**b**) of the right-sided aortic arch giving rise to the aberrant left subclavian artery from the Kommerell diverticulum. (*AoAsc* ascending aorta, *AoDes* descending aorta, *Tra* trachea, *LCC* left common carotid artery, *RCC* right common carotid artery, *RSA* right subclavian artery, *LSA* aberrant left subclavian artery, *KD* Kommerell diverticulum)

