Case

In May 2015, a 59-year-old woman presented to Princess Margaret Hospital, Hong Kong, with chronic intermittent vertigo and syncope, aggravated by head rotation to the right. Physical, otoscopy, and nasal endoscopy examination results were unremarkable. Cervical spine plain radiographs demonstrated cervical spondylosis with marginal osteophytosis (Fig 1). In May 2015, computed tomography angiography of the head and neck revealed focal moderate (50%) stenosis at bilateral vertebral arteries at C5/6 levels due to extrinsic compression from hypertrophied uncovertebral joints (Fig 2). In January 2016, magnetic resonance imaging demonstrated disco-osteophytic protrusion at C5/6 level without evidence of cord compression. In March 2016, digital subtraction angiography, performed with the patient's head in neutral and bilateral rotated positions, demonstrated dynamic deterioration of focal stenosis of right vertebral artery at C5/6 level to up to 80% stenosis during head rotation to the right (Figs 3 and 4).

A static focal moderate (50%) stenosis of the left vertebral artery at C5/6 was also present. Overall findings were compatible with bow hunter's syndrome (BHS) with dynamic deterioration of right vertebral artery stenosis on head rotation, related to extrinsic compression by hypertrophied facet joint and disc protrusion. In April 2016, the patient underwent anterior C5/6 cervical discectomy and anterior spinal fusion with smooth recovery and symptomatic resolution.

Discussion

Bow hunter's syndrome was first reported in 1978 when a patient developed lateral medullary syndrome during archery practice with lateral head rotation. It refers to symptomatic vertebrobasilar insufficiency secondary to mechanical occlusion or stenosis of vertebral arteries upon head and neck rotation.

The pathogenesis of BHS is related to the tortuous anatomical course of the vertebral artery along the cervical spine, which renders the artery susceptible to extrinsic compression, repetitive shear stress resulting in haemodynamic events in
Osteophytes, disc herniation, ligamentous or neck muscle hypertrophy are risk factors for BHS. Though rare, BHS is a not to be missed cause of vertigo, owing to its specific relationship with head and neck rotation and its potential risk of posterior circulation ischaemic stroke. Bow hunter’s syndrome is more common among males and those aged between 50 and 70 years old. Common clinical manifestations include vertigo and syncope. Other symptoms include nystagmus, emesis, Horner’s syndrome, and rarely motor and sensory deficits.

Imaging is crucial in establishing the diagnosis of BHS, delineating the cause and site of extrinsic compression and evaluating complications such as infarction. Dynamic digital subtraction angiography remains the preferred modality for prompt and accurate localisation of stenotic segment and establishing causal relationship with head rotation. Non-invasive computed tomography or magnetic resonance angiography in both neutral and rotated head positions are also used. Computed tomography can delineate the relationship with surrounding compressive skeletal structures and magnetic resonance would be sensitive in documenting early ischaemic event.

In this case, the patient’s complaint of vertigo exacerbation with specific direction of head rotation should raise the suspicion of BHS. Surgical treatment was offered in view of failed conservative approach, repeated fall related to syncope, and underlying uncovertebral joint hypertrophy and disc protrusion as the aetiological factors of vertebral artery compression.

Author contributions
All authors had full access to the data, contributed to the study, approved the final version for publication, and take responsibility for its accuracy and integrity.

Concept or design: All authors.
Acquisition of data: TS Chan, JKF Ma.
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Drafting of the manuscript: SC Wong. TS Chan.
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All patients were treated in accordance with the Declaration of Helsinki and provided consent for all investigations and procedures.

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