


Eagle Syndrome as a Cause of Cerebral Venous Sinus Thrombosis

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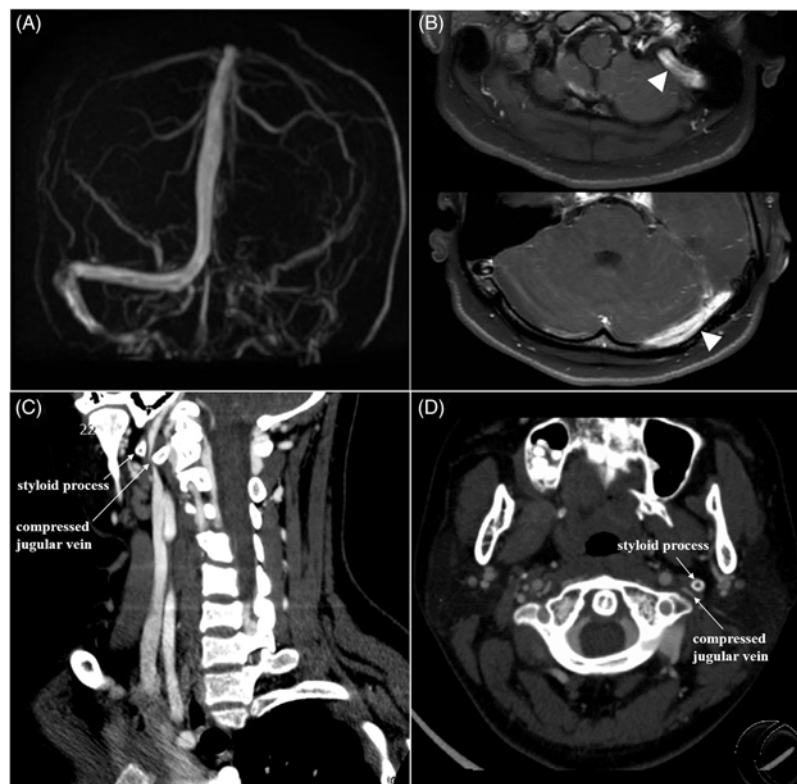


Figure 1: Neurological imaging examinations of the patient. PC-MRV (A) showed absent signal in the left transverse-sigmoid sinuses. Contrast-enhanced 3D fat-saturated T1 VISTA MRI (B; white arrowheads) demonstrated gadolinium enhancement of the left transverse-sigmoid sinuses, indicating chronic venous sinus thrombosis. The curved planar reformation (CPR) and axial source images from CT venography images (C and D; white arrow) showed styloid process compressing the left jugular vein.

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A 15-year-old teenager presented with a 2-month history of headache. Neurological examination was normal except for papilledema. Further lumbar puncture indicated intracranial hypertension (330 mm H₂O). Brain magnetic resonance imaging (MRI) was normal but phase contrast-magnetic resonance venography (PC-MRV) (Figure 1(A)) suggested possible left transverse-sigmoid sinus thrombosis; subsequent contrast-enhanced 3D fat-saturated T1 volumetric isotropic turbo spin echo acquisition (VISTA) MRI (Figure 1(B)) confirmed the pathology. Hyper-coagulable panel results (including six steroid sex hormones, antithrombin III, protein C, protein S, lupus anticoagulant, and anticardiolipin antibodies) were all within normal range. In further examination, computed tomography (CT) venography images (Figure 1(C) and (D)) showed that the left jugular vein was compressed by the styloid process, consistent with Eagle syndrome.¹ The patient who refused the recommended surgical treatment, however, chose anticoagulant therapy consisting of low-molecular weight heparin subcutaneous injection in addition to new oral anticoagulant. At 18-month follow-up, the patient reported no symptoms remained.

Eagle syndrome, also known as styloid syndrome, was initially described by Watt Weems Eagle in 1937; it often presented with cervical pain on the side of the elongated styloid process, foreign-body sensation in the throat, and dysphagia.² In addition, it can cause transient ischemic attacks or stroke due to cervical internal carotid artery compression.¹ However, cases of Eagle syndrome causing cerebral venous sinus thrombosis were seldom reported.³ In our report, we want to remind the neurologists that

Eagle syndrome should be taken into account when dealing with cases of cerebral venous sinus thrombosis of unknown origin, and CT venography may be a useful diagnostic tool.

STATEMENT OF AUTHORSHIP

Study concept and design: Yang; acquisition of data: Zhang, Zhou, and Guo; image analysis: Zhou; drafting of the manuscript: Zhang, Zhou; critical revision of the manuscript for important intellectual content: Yang; obtained funding: Yang.

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DISCLOSURES

The authors declare no conflicts of interest.

REFERENCES

1. Badhey A, Jategaonkar A, Anglin Kovacs AJ, et al. Eagle syndrome: a comprehensive review. *Clin Neurol Neurosurg.* 2017;159:34–8.
2. Eagle WW. Elongated styloid processes: report of two cases. *Arch Otolaryngol.* 1937;25(5):584–7.
3. Aydin E, Quliyev H, Cinar C, Bozkaya H, Oran I. Eagle syndrome presenting with neurological symptoms. *Turk Neurosurg.* 2018;28(2):219–25.