Dural Arteriovenous Fistula with Hypoglossal **Nerve Paralysis**

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A male in his early 30s presented with a swelling just below the angle of his left mandible for the last 2 months. The swelling had insidious onset, progressive and painless. Examination found 4×4 cm, diffuse, soft to firm, pulsatile swelling in the upper part of the neck on left side (**Fig. 1**). An oral examination revealed left hypoglossal nerve paralysis. Further examination was noncontributory, including vagus and accessory spinal nerve examination. A probable diagnosis of carotid body tumor was kept, and the patient was subjected to computerized angiography.

Angiography revealed dural arteriovenous fistula (DAVF), a rare clinical entity leading to hypoglossal nerve paralysis (>Figs. 2 (A, B)). Patient was advised to undergo digital subtraction angiography for further management, but he refused and chose conservative care with regular follow-up.

Hypoglossal nerve paralysis due to DAVF is rare. Digital subtraction angiography is the gold standard investigation, whereas endovascular embolization is the treatment of choice.² Regular follow-up with imaging is the option in a few selective cases.²⁻⁴





Fig. 1 A diffuse swelling in the neck (white arrow) with left hypoglossal nerve paralysis (black arrow).

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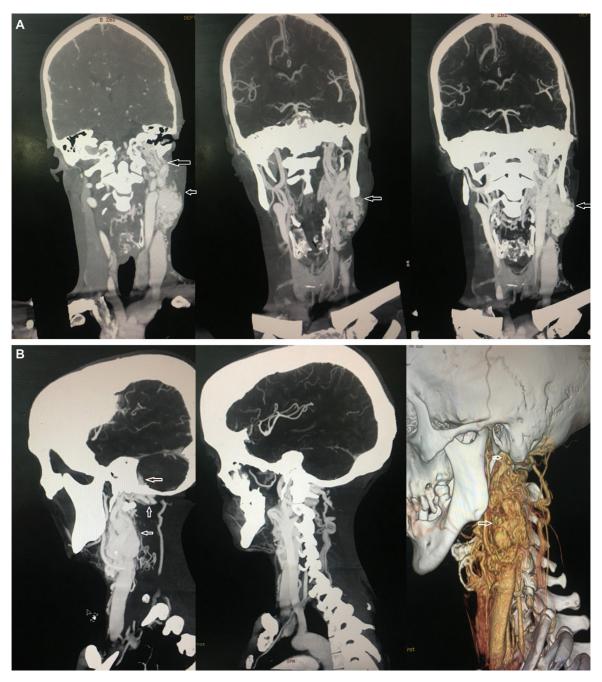


Fig. 2 (A, B) Computed tomography angiography showing arteriovenous malformation in the neck and its extension into the cranial cavity with dural arteriovenous fistula (white arrows).

Conflict of Interest None declared.

References

1 Chan NHHL. Hypoglossal dural arteriovenous fistula: a rare cause of unilateral hypoglossal nerve palsy. BJR Case Rep 2017;3(03): 20160144

- 2 Mayercik VA, Sussman ES, Pulli B, et al. Efficacy and safety of embolization of dural arteriovenous fistulas via the ophthalmic artery. Interv Neuroradiol 2021;27(03):444-450
- 3 Baharvahdat H, Ooi YC, Kim WJ, Mowla A, Coon AL, Colby GP. Updates in the management of cranial dural arteriovenous fistula. Stroke Vasc Neurol 2019;5(01):50-58
- 4 Hiramatsu M, Sugiu K, Hishikawa T, et al. Results of 1940 embolizations for dural arteriovenous fistulas: Japanese Registry of Neuroendovascular Therapy (JR-NET3). J Neurosurg 2019;133(01):166-173