



A Rare Case of Multiple Intracranial, Intraspinal, and Peripheral Schwannoma with Intracranial Meningioma

Sumit Bansal¹ Rabi Narayan Sahu¹ Ashis Patnaik¹

¹Department of Neurosurgery, All India Institute of Medical Sciences, Bhubaneswar, Orissa, India

Address for correspondence Sumit Bansal, MCh, Associate Professor, Room No. 419, Department of Neurosurgery, All India Institute of Medical Sciences, Bhubaneswar, Orissa 751019, India (e-mail: drsumitbansal@gmail.com).

Indian | Neurosurg 2019;8:89-90

Bilateral vestibular schwannoma (VS) are found in 90 to 95% of patients with neurofibromatosis 2 (NF2). It is reported that more than 99% of VS cases in NF2 are benign, but they remain an important cause of mortality due to their location. Schwannomas can develop along the course of the cranial, spinal, and peripheral nerves, differently than

vestibular. Often, it arises from the oculomotor, trigeminal, and facial nerves.² Bilateral trigeminal schwannoma with NF2 is rare.^{3,4} Intracranial meningiomas appear in 45 to 58% of patients with this disorder.⁵ We here present a case of NF2 with intracranial, intraspinal, and peripheral involvement.

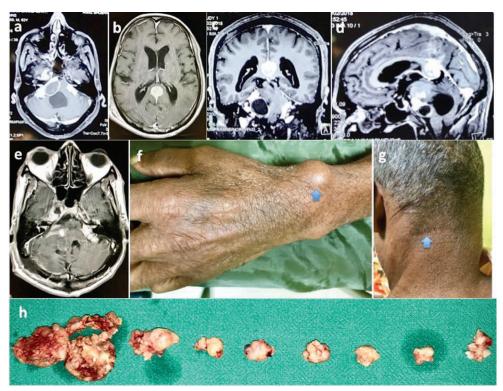


Fig. 1 Contrast T1-weighted MRI images (axial, coronal, and sagittal) showing bilateral trigeminal schwannoma (right is larger than left) with bilateral multiple small schwannoma of seventh, eighth, and lower cranial nerves with velum interpositum meningioma (a-d). Postoperative axial T1-weighted contrast MRI showing excision of right-sided tumors with residual middle fossa component of right trigeminal schwannoma (e). Large subcutaneous nodule on radial aspect of left wrist (f) and old scar of previously operated schwannoma on left side of nape of neck (g). Excised tumors (eight in number) from right cerebellopontine angle region, largest was of trigeminal schwannoma (h).

received May 15, 2018 accepted May 24, 2018 published online April 22, 2019 **DOI** https://doi.org/ 10.1055/s-0039-1678466 **ISSN** 2277-954X. ©2019 Neurological Surgeons' Society of India

License terms









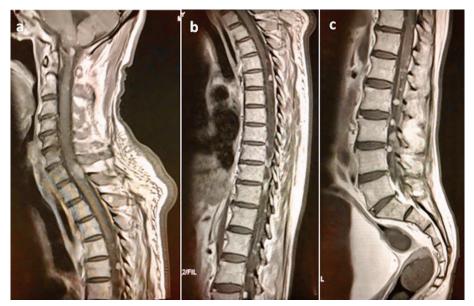


Fig. 2 Contrast MRI of spine showing scattered multiple schwannoma along the whole length of spinal cord (a-c).

A 63-year-old man presented with 32-year history of hearing impairment in both the ears, with a 20-year history of multiple subcutaneous nodules all over the body with imbalance while walking for 18 months. He also had nasal regurgitation of fluid for 6 months. On examination he had multiple subcutaneous skin swellings all over the body along with diminution of vision in both the eyes (6/60), with bilateral fifth, seventh, eighth, and lower cranial nerves involvement. Bilateral cerebellar signs were present.

On magnetic resonance imaging (MRI) of the brain and spine, there were bilateral trigeminal schwannoma (right > left) along with bilateral multiple small schwannoma of the seventh, eighth, and lower cranial nerves with velum interpositum meningioma (**Fig. 1**). Whole-spine screening MRI showed multiple schwannoma along the length of the spinal cord (**Fig. 2**).

Right-sided trigeminal schwannoma along with other right-sided small schwannoma were excised. Pathologic examination of tumors revealed schwannoma. Postoperative course was uneventful. Other lesions were managed conservatively and referred for gamma knife radiosurgery.

Financial Support

None.

Conflict of Interest

None.

References

- 1 Asthagiri AR, Parry DM, Butman JA, et al. Neurofibromatosis type 2. Lancet 2009;373(9679):1974–1986
- 2 Fisher LM, Doherty JK, Lev MH, Slattery WH III. Distribution of nonvestibular cranial nerve schwannomas in neurofibromatosis 2. Otol Neurotol 2007;28(8):1083–1090
- 3 Yamada K, Ohta T, Miyamoto T. Bilateral trigeminal schwannomas associated with von Recklinghausen disease. AJNR Am J Neuroradiol 1992;13(1):299–300
- 4 Goel A, Muzumdar D, Raman C. Trigeminal neuroma: analysis of surgical experience with 73 cases. Neurosurgery 2003;52(4):783–790, discussion 790
- 5 Patronas NJ, Courcoutsakis N, Bromley CM, Katzman GL, MacCollin M, Parry DM. Intramedullary and spinal canal tumors in patients with neurofibromatosis 2: MR imaging findings and correlation with genotype. Radiology 2001;218(2):434–442