

Images in Hospital Medicine

Vagus Nerve Schwannoma

Andrew J Aultman II¹, Matthew Genelin¹, Jessica Gonzalez¹

¹ Internal Medicine, Brown University

Journal of Brown Hospital Medicine

Vol. 3, Issue 1, 2024

Article Information

Keywords: vagal nerve schwannoma, schwannoma, carotid sheath, vagus nerve

https://doi.org/10.56305/001c.89988

Submitted: September 20, 2023 EST

Accepted: November 07, 2023

EST

Abstract

Schwannomas are peripheral nerve tumors that may impact cranial nerves. We present the case of a 21-year-old male with imaging findings consistent with vagal nerve schwannoma with an atypical symptomatic presentation. A review of the presentation, characteristic imaging findings, differential diagnosis, and treatment of vagal nerve schwannoma is provided.

VAGUS NERVE SCHWANNOMA

A 21-year-old previously healthy male presented with sudden-onset headache, photophobia, and neck pain accompanied by nausea and anxiety. Examination revealed right-sided neck tenderness to palpation and right-sided pain with active and passive movement. He had a normal cranial nerve and neurological exam without nuchal rigidity. Computerized tomography (CT) of the brain and neck showed a large cystic lesion in the carotid space (<u>Figure 1</u>). The lesion was further characterized with magnetic resonance imaging (MRI) that showed a 7.7 cm x 1.6 cm x 4.1 cm lobulated soft tissue mass involving the right carotid space with anterior displacement of the carotid artery. The lesion was isointense on T1-weighted imaging (T1WI), hyperintense on T2-weighted imaging (T2WI), and showed patchy internal enhancement concerning for a vagal nerve schwannoma (Figure 2).

Schwannomas are rare peripheral nerve tumors arising from the myelinating Schwann cells of the peripheral nervous system.1 Less frequently, they may impact cranial nerves. When affecting the vagus nerve, the patient may experience hoarseness or cough or may remain asymptomatic.² Growth is usually limited to 2-3mm per year, and malignant transformation is possible but rare. Differential diagnosis of carotid space masses is broad but includes schwannoma of the vagus, glossopharyngeal, or spinal accessory nerves; congenital branchial cyst; paraganglioma; and infectious process.^{3,4} On MRI, paragangliomas usually show "salt and pepper" appearance due to flow voids, while branchial cleft cysts may show wall thickening, hypointensity on T1WI tending towards hyperintensity with infection or proteinaceous content, and variable intensity on T2WI.^{5,6} Schwannomas may appear cystic with isointensity on T1WI and hyperintensity on T2WI.¹ Once schwannoma is suspected, the location of the internal carotid artery can provide diagnostic clues.⁷ Displacement of the internal carotid artery anteriorly away from the internal jugular vein is consistent with vagal nerve origin, and extension towards the jugular foramen of the skull further supports this diagnosis.^{2,7,8} Given encasement of upper cervical right internal carotid artery with anterior displacement, extension to the skull base, lack of internal flow voids disfavoring paraganglioma, and internal enhancement pattern disfavoring branchial cleft cyst, vagal nerve schwannoma was suspected and the patient was referred for surgical intervention.

Treatment is with surgical excision or debulking with nerve-sparing techniques when complete excision is not possible.² Potential complications of excision include vocal cord palsy due to vagus nerve damage which may require subsequent nerve reconstruction.² Progression may also lead to nerve deficit, pain, and compression of the carotid sheath structures. Prognosis is typically favorable with treatment.

Author contributions

All authors have reviewed the final manuscript prior to submission. All the authors have contributed significantly to the manuscript, per the International Committee of Medical Journal Editors criteria of authorship.

- Substantial contributions to the conception or design of the work; or the acquisition, analysis, or interpretation of data for the work; AND
- Drafting the work or revising it critically for important intellectual content; AND
- Final approval of the version to be published; AND
- Agreement to be accountable for all aspects of the work in ensuring that questions related to the accu-

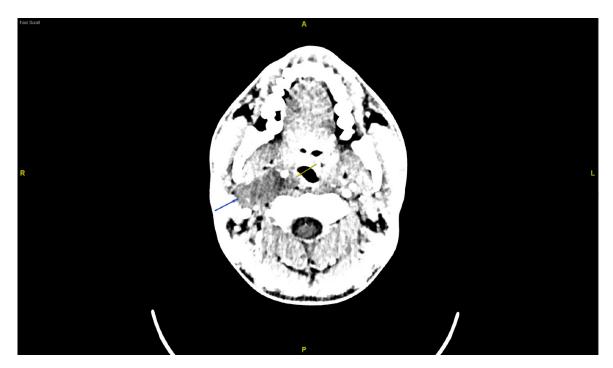


Figure 1. CT scan showing large cystic lesion in the carotid space

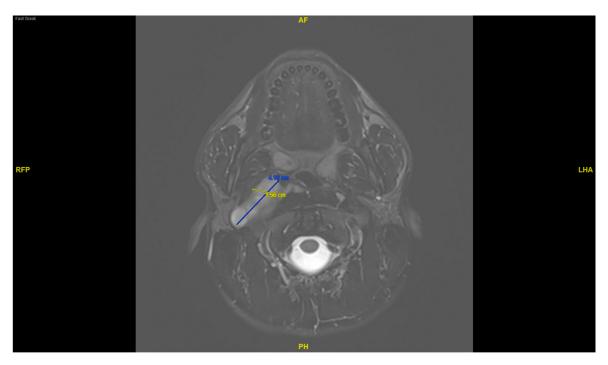


Figure 2. Axial T2-weighted MRI showing hyperintensity

racy or integrity of any part of the work are appropriately investigated and resolved.

Conflicts of interest

The authors have no conflicts of interest to disclose.

Corresponding Author

Andrew Aultman Warren Alpert Medical School at Brown University Providence, RI, 02903

Email: Andrew aultman@brown.edu



This is an open-access article distributed under the terms of the Creative Commons Attribution 4.0 International License (CCBY-NC-4.0). View this license's legal deed at https://creativecommons.org/licenses/by-nc/4.0/legalcode for more information.

REFERENCES

- 1. Hilton DA, Hanemann CO. Schwannomas and their pathogenesis. *Brain Pathol*. 2014;24(3):205-220. doi:10.1111/bpa.12125
- 2. Cavallaro G, Pattaro G, Iorio O, Avallone M, Silecchia G. A literature review on surgery for cervical vagal schwannomas. *World J Surg Onc.* 2015;13(1):130. doi:10.1186/s12957-015-0541-6
- 3. Geethapriya S, Govindaraj J, Raghavan B, et al. Cranial nerve schwannoma A pictorial essay. *Indian J Radiol Imaging*. 2020;30(2):116-125. doi:10.4103/ijri.ijri_17_20
- 4. Hosalkar RM, Khivasara JS, Swain N. Carotid body paraganglioma. *Ann Maxillofac Surg.* 2019;9(2):423-428. do i:10.4103/ams.ams_183_18

- 5. Itani M, Mhlanga J. Imaging of Pheochromocytoma and Paraganglioma. In: Mariani-Costantini R, ed. *Paraganglioma: A Multidisciplinary Approach*. Codon Publications; 2019:41-61. doi:10.15586/paraganglioma.2019.ch3
- 6. Kawaguchi M, Kato H, Aoki M, Kuze B, Hara A, Matsuo M. CT and MR imaging findings of infection-free and benign second branchial cleft cysts. *Radiol Med*. 2018;124(3):199-205. doi:10.1007/s11547-018-0959-3
- 7. Sreevatsa MR, Srinivasarao RV. Three cases of vagal nerve schwannoma and review of literature. *Indian J Otolaryngol Head Neck Surg.* 2011;63(4):310-312. doi:10.1007/s12070-011-0220-z
- 8. Chiofalo MG, Longo F, Marone U, Franco R, Petrillo A, Pezzullo L. Cervical vagal schwannoma. A case report. *Acta Otorhinolaryngol Ital*. 2009;29(1):33-35.